

Swiss National Hip & Knee Joint Registry

Report 2025

Annual Report of the SIRIS Registry Hip & Knee, 2012 – 2024

Hip and knee replacement results 2012 – 2024

SIRIS Report 2025

Annual Report of the Swiss National Joint Registry, Hip and Knee

SIRIS – Foundation for Quality Assurance in Implant Surgery

swiss orthopaedics – Swiss Society of Orthopaedics and Traumatology

ANQ – National Association for the Development of Quality in Swiss Hospitals and Clinics

SwissRDL – Medical Registries and Data Linkage, Institute of Social and Preventive Medicine,
University of Bern



Preface

Among those who worked assiduously on establishing the partnership between SIRIS (the Foundation for Quality Assurance in Implant Surgery), swiss orthopaedics and ANQ was Regula Heller. On the occasion of her retirement as Head of the Department of Acute Care and Deputy Managing Director of ANQ in September 2025, she now looks back at what happened in the beginning and talks about the relevant factors that determined the success of the undertaking.

The 2025 SIRIS Report is already the 12th annual report concerning the Swiss National Hip & Knee Joint Registry. The continually expanding database and the excellent quality of the data make possible increasingly informative evaluations. This year's report also contains some fascinating innovations. For the first time, it introduces analyses of regional differences, thus providing additional helpful insights for funding agencies, service providers, policymakers and the authorities.

A SPECIAL PARTNERSHIP

When the SIRIS Foundation and ANQ initiated their collaboration in 2011, they had quite specific expectations. SIRIS was anticipating the nationwide collection of implant data while ANQ was looking forward to the extension of its quality reviews to the realm of orthopaedics. The implant registry was made part of ANQ's review plan in 2012. Since then, all acute care hospitals and clinics have been required to enter the knee and hip implant procedures they carry out in the SIRIS registry. With its data collection rate of greater than 98%, the registry today also stands out in international comparison. This success is attributable to the close cooperation between the SIRIS Foundation, swiss orthopaedics and ANQ, sharing as they do a common ambition to continuously improve implantation medicine for the benefit of patients. Although the missions of the three organisations differ, they have managed to find ways of reconciling their individual requirements with their shared goals. The SIRIS implant registry thus represents a shining example of how an interprofessional partnership in the healthcare sector can function superlatively - not only that, but the registry itself is an outstanding instance of what can be achieved by means of joining forces.

EXTENSIVE REPORTING FRAMEWORK – TARGETED IMPROVEMENTS

Considerable planning and persuasion efforts were necessary to both ensure that the data pool was such that risk-adjusted hospital/clinic comparisons were possible and that the initial resistance to disclosing outcomes was overcome. By 2021, all was ready and, for the first time, ANQ was able to publish the two-year revision rates for primary total hip and knee replacements in transparent form.

The registry may have originally served to essentially identify faulty joint implants, but today the focus is also on the surgical services provided and the procedures themselves. Increasing numbers of hospitals and clinics are using the registry evaluations as the basis for the introduction of targeted improvements. This is made possible by the extensive analyses provided, including the quarterly reports on in-house data and the reports of problems with joint replacements. These changes are to be welcomed as the two-year revision rates for primary total hip and knee joint replacements are now exhibiting declining trends.

FACTORS FOR FUTURE SUCCESS

Thanks to the ANQ National Quality Agreement in 2011 and the commitments entered into by SIRIS and ANQ, the targets, processes, divisions of responsibilities and structures are clearly defined while the financing is guaranteed. SIRIS is well-positioned to continue in future with what has been achieved. There are three factors that are of particular relevance:

1. Easing of the burden on hospitals and clinics: the project designed to exploit synergies in data collection processes drawn up in cross-disciplinary partnership with Swissnoso indicates the way ahead.

2. Patient-reported outcome measures (PROMs): the point of view of patients must be taken into account if care is to be more patient-oriented and there is increasing focus on this aspect in considerations of cost-benefit factors. With the optional collection of PROMs, the required basics are in place.

3. Secure positioning in the discipline of orthopaedics in close cooperation with swiss orthopaedics: so that the registry can develop its full potential, it also needs in future to be closely linked with practice and with the world of the professionals. In this context, swiss orthopaedics can play a key role as both a mediator and as an ambassador.

PERSONAL THOUGHTS AS I RETIRE

As Head of the Department of Acute Care at ANQ, I was able to work with SIRIS for 14 years. I considered this to be a privilege. I was constantly impressed by the specialist expertise and constructive solidarity that was evident in the committees and the dedication shown by the hospitals and clinics. I should like to express my profound gratitude to all involved for the way we were able to work together. I am convinced that the registry will continue to play an important role in implantation medicine in future and would like to take this opportunity to wish the 'SIRIS collective' every success in the coming years!

Regula Heller

MNSc, MPH

Acknowledgements

SIRIS Scientific Advisory Board (SSAB)

Prof. Martin Beck, MD

Orthopädische Klinik Luzern AG, Lucerne
swiss orthopaedics Hip Expert Group
Author of the scientific SIRIS-Report

Stefan Bauer, MD

Ensemble Hospitalier de la Côte, Morges
swiss orthopaedics Shoulder Expert Group

Lilianna Bolliger, MSc ETH

Data Quality and Monitoring
SwissRDL, ISPM, University of Bern

Christian Brand, PhD, MSc

Statistical Analysis and Research
SwissRDL, ISPM, University of Bern
Author of the scientific SIRIS-Report

Bernhard Christen, MD, MHA

articon AG, Salem-Spital Berne,
swiss orthopaedics Knee Expert Group
Author of the scientific SIRIS-Report

PD Karlmeinrad Giesinger, MD

HOCH Health Ostschweiz,
Department of Orthopaedics and Traumatology,
Kantonsspital St.Gallen

Vilijam Zdravkovic, MD, MSc

HOCH Health Ostschweiz,
Department of Orthopaedics and Traumatology,
Kantonsspital St.Gallen,
Autor of the scientific SIRIS-Report

SIRIS Expert Group ANQ

all Members of the SSAB and:

Prof. Claudio Dora, MD

Schulthess Clinic, Zurich, Senior Consultant Hip
Surgery
SIRIS Board Member

Regula Heller, MNSc, MPH

Chair Akutsomatik ANQ
Lead and moderation SIRIS Expert Group ANQ

PD Hermes Miozzari, MD

Médecin adjoint, HUG
swiss orthopaedics Board Member

Jasmin Vonlanthen

SIRIS Administration Lead

Florian Rütter, MD

Head Quality Management & Value Based
Healthcare, University Hospital Basel

Thomas Schärer

Enovis Surgical Switzerland GmbH,
General Manager
Representative of Swiss Medtech

Prof. Moritz Tannast, MD

Head Clinic of Orthopaedics and
Traumatology, Inselspital
swiss orthopaedics Board Member

PD Peter Wahl, MD

Staff Surgeon, ENDO-Team, Birshof Hospital
Lector of the scientific SIRIS-Report

Partner Associations

ANQ

National Association for the Development
of Quality in Swiss Hospitals and Clinics
info@anq.ch

EUROSPINE

the Spine Society of Europe
info@eurospine.org

H+

Association of the Hospitals of Switzerland
geschaefsstelle@hplus.ch

santésuisse

Association of Swiss Medical Insurers
mail@santesuisse.ch

SGNC

Swiss Society of Neurosurgery
sgnc@imk.ch

SGS

Swiss Society of Spinal Surgery
office@spinecociety.ch

Swiss Medtech

Association of the Swiss Medtech Industry
office@swiss-medtech.ch

swiss orthopaedics

Swiss Society of Orthopaedics and Traumatology
welcome@swissorthopaedics.ch

SwissRDL

ISPM, University of Bern
swissrdl.ispm@unibe.ch

SIRIS Foundation Board Members

Prof. Claudio Dora, MD, President
Representative of swiss orthopaedics

Verena Nold, Vicepresident
Representative of santésuisse

Dr. med. Stefan Grunder, MD
Representative of santésuisse

Prof. Bernhard Jost, MD
Representative of swiss orthopaedics

Prof. Christoph Meier, MD
Representative of Hplus

Prof. Andreas Raabe, MD
Representative of swiss neuro surgery

Adriano Salvisberg
Representative of Swiss Medtech

Eduardo Stadelmann
Representative of Swiss Medtech

Thomas Straumann
Representative of Hplus

Foundation for quality assurance in implant surgery – SIRIS

c/o Core Treuhand AG, 3007 Bern
info@siris-implant.ch, www.siris-implant.ch

All information in this report was composed with the utmost care. If any changes or modifications are made after publication, these will be published on our website www.siris-implant.ch, where you can also download the SIRIS Report 2025 and all previous reports.

Published December 2025

Content

1. Introduction	
1.1 Purpose of the registry	9
1.2 Strong commitment	12
2. Methods	
2.1 Maintenance and hosting of the registry	13
2.2 Data quality and completeness	14
2.3 Coverage	15
2.4 Statistical precision, evaluation and outlier detection	18
2.5 Patient-reported outcome measures (PROMs)	19
3. Demography and Epidemiology	
3.1 Introduction	21
3.2 Hip arthroplasty	22
3.3 Knee arthroplasty	40
4. Hip arthroplasty	
Overview of data structure	61
4.1 Primary total hip arthroplasty	63
4.2 First revision within 2 years of primary total hip arthroplasty	71
4.3 Re-revision of total hip arthroplasty	88
4.4 Results of implants in total hip arthroplasty	94
4.5 Treatment of hip fractures	117
4.6 Early revision after arthroplasty for fracture of the hip	120
4.7 Results of implants after hip fracture	125
5. Knee arthroplasty	
Overview of data structure	133
5.1 Primary total knee arthroplasty	135
5.2 Revision of primary total knee arthroplasty implanted before 2012	140
5.3 First revision within two years after primary TKA	140
5.4 Re-revision of total knee arthroplasty	152
5.5 Results of implants in total knee arthroplasty	157
5.6 Primary partial knee arthroplasty	167
5.7 First revision within two years after primary partial knee arthroplasty	170
5.8 Results of implants in unicondylar partial knee arthroplasty	176
5.9 Results of implants in patellofemoral joint partial knee arthroplasty	181
6. Patient reported outcome measures (PROMs)	
6.1 Joint-specific pain	184
6.2 Quality of life	186
6.3 Joint-specific satisfaction	188
6.4 Treatment effects	188
6.5 Conclusion	191
SIRIS outlier watch list – hip implants	192
SIRIS outlier watch list – knee implants	195
List of manufacturers and distributors	196
Definitions	197
Participating hospitals 2024	199
List of tables and figures	201



**Introduction, Methods,
Demography and Epidemiology**

1. Introduction

1.1 Purpose of the registry

The Swiss National Implant Registry (SIRIS) was implemented in September 2012 to evaluate hip and knee joint replacements, introducing also shoulder joint replacements since January 2025. Registration in SIRIS is mandatory for all Swiss hospitals performing knee and hip arthroplasties, as they are bound to the ANQ's national quality agreement.

To ensure that all contributors and participants pursue a common goal, it is essential to clearly define the aim of the SIRIS registry. This also influences the data architecture within the registry, since there will be different requirements for each stakeholder. The multi-partner association required to set up the registry meant that various points of view had to be considered to ensure success and support. Although each partner naturally tends to focus preferentially on their particular interest, one fundamental interest is common to all, namely the long-term well-being of the patient after a prosthetic joint replacement. The following paragraphs will explain the various perspectives of key stakeholders that were considered during development of SIRIS.

The patient's perspective. Since patients expect joint replacement surgery to provide them with long-lasting, pain-free mobility, the surgical procedure must be adapted to their level of activity and should be followed by rapid recovery without complications. Ideally, after a prosthetic joint replacement and successful rehabilitation process, patients should feel well and report about excellent outcome (PROMs – Patient Reported Outcome Measures). From the patient's perspective, the registry data should be presented as to be readily comprehensible, allowing patients to find information of interest despite the complexity of the data and of the methodology behind the analysis. The growing importance of functional results beyond revision rates is reflected the inclusion of PROMs in the annual report since 2022. Nationwide registration of

PROMs within SIRIS is planned to be implemented in 2025, but data from the pilot project of the Canton of Zurich already provided a basis for the development of reporting standards of PROMs and delivered first insights into patient-related outcomes. Patients who will read the report may find the information helpful to better understand their past or future surgery and thus be able to better discuss it with their surgeons.

The surgeon's perspective. For surgeons, a priority is to avoid surgical complications and shortcomings for their patients. In fact, the needs of patients and the goals of surgeons are fully aligned: obtaining functional, long-lasting and pain-free joint replacement. Introducing shoulder joint replacement added a new group of SIRIS users – shoulder surgeons. By choosing a particular prosthesis, surgeons integrate the performance of the implant with their expertise. The implants must be well designed and impeccably manufactured to avoid problems such as wear particle disease or breakage. Hence, to add value from the surgeon's perspective, the registry should be able to identify problematic implants within a relatively short time frame and provide valuable early warning to surgeons when required. However, entering individual clinical results into a registry is not a welcome addition to an already intense daily schedule of activities. Furthermore, although surgeons certainly appreciate benchmarking their results, the public availability of the information at the individual surgeon's level remains a controversial aspect. PROMs are also adding a new perspective for surgeons – their patients' reports benchmarked relative to PROMs of patients treated by other surgeons in Switzerland.

The industry's perspective. The industry's main activity is manufacturing and sales, driven by a legitimate profit-orientation. Designing and providing first-rate, problem-free implants is the most enduring strategy in this domain, as a series of failures may lead to allegations of negligence that could ultimately destabilise the company. Hence, the aspect

of economic viability coincides with the primary interest of the patients, namely successful prosthetic joint replacement on the long term. Since progress and technical innovation are extremely important for an industry dedicated to providing safe high-performance implants, the registry is regarded as an essential tool for post-market surveillance that validates improvements in materials, designs, and concepts in the real-life clinical setting. If the industry accepts quality as being the principal market-regulating factor, then the registry is a welcome tool and motivates participation. To date, the publication of two-year revision rates for registered implants in the SIRIS reports was met with great interest from involved industry and orthopaedic surgeons. Adding the medium-term follow-up at 5 and 10-years to the report further improves the function of the registry as quality assessment tool. Introduction of shoulder implants and PROMs enlarges the portfolio of SIRIS registry and offers additional information to industry.

The hospital's perspective. It is in hospitals where the surgeons and patients interact, with all parties sharing a common interest. Hospitals aim to provide high-quality healthcare at reasonable costs, integrating volume of patients into considerations. However, hospitals or departments also have an interest in ensuring that patients do not forget the institution where they were treated successfully, making them return to the same hospital if necessary, including for reasons other than a prosthetic joint replacement. The registry is perceived as an instrument for quality control, not only for the implants used but for the entire process, ranging from the preoperative consultation to the procedures in the operating room, as well as the postoperative follow-up. Personal recommendations from satisfied patients are the very best advertising for hospitals and related medical institutions. As institutions providing healthcare in today's competitive environment, hospitals are also very keen to uphold their reputation, and the registry is an invaluable tool for this purpose. Additionally,

since certain Swiss cantons require reaching a sufficient volume of procedures in order to keep hospitals on contract lists, it appears that participating in the registry might be crucial for the survival of some institutions, a strong motivation for participation in an environment where hospital mergers and closures are frequently discussed. Performance benchmarks containing the two-year revision rates of institutions registered in SIRIS have been published online since 2020 ([LINK to ANQ](#)) and are updated with every new report.

The healthcare insurances's perspective. Insurers and third-party payers are concerned about healthcare costs, and thus aim to reduce hospitalization costs, to avoid re-admissions for complications, and obtain return to work of the patient as early as possible, if applicable. Insurers are very conscious of costs regarding implant pricing, medical fees, and hospital bills. Because the insurer's objective is to provide equal benefits to all its clients within the available budget, the registry is perceived as an instrument that can provide information regarding the performance of surgeons and institutions, functioning as a cost-quality assessment tool. Since costs for primary prosthetic replacement are constricted within DRG – reimbursement system, but revision surgeries may cause significant additional and potentially avoidable costs, the focus of insurers could be directed to revision rates and re-revisions.

The government's perspective. The government organises the healthcare system on behalf of all citizens. Therefore, the main challenge it faces is having to consider and bring together the needs and preferences of all players in the healthcare economy. One specific characteristic of the Swiss healthcare system is that cantons are independent and are the principal political and financial authorities for their healthcare systems. Furthermore, the healthcare system of the Principality of Liechtenstein (FL) interacts closely with the Swiss healthcare system and also participates in

SIRIS. Although the federal government may not have any inherent financial interest in the healthcare system, the cantonal governments directly bear a major share of hospital costs and are very active participants in all debates concerning hospital treatments, including their outcomes and costs. Governments also have an interest in assessing the overall situation concerning the quality of healthcare provision. While patients understandably tend to place their primary focus on receiving treatment that provides optimal and long-lasting results, the government certainly shares this aim but also has to ensure cost-effectiveness, as resources are limited. Nevertheless, political decision makers in Switzerland at all levels have arguably so far largely avoided that aspect, currently leaving the Swiss healthcare system somewhat adrift when it comes to actually allocating the limited resources in a rational and transparent manner. Despite this, governments require data on the overall surgical performance to assess requirements and subsequently plan the macroeconomic policies related to healthcare. Hence, government health agencies are commissioned to ensure that the institutions under their supervision provide high-quality healthcare to the population, whereby the agencies also have an interest in benchmarking hospitals and keeping insurance and third-party payer costs down to a reasonable minimum. Although the fragmentation of the dataset down to the cantonal level may sometimes preclude meaningful statistical analysis, the information can still be of interest to the Swiss cantonal and FL governments, as well as to the general public.

1.2 Strong commitment

The annual report 2025 represents a collaborative data collection involving all the institutional partners of SIRIS and includes the surgeons and operating teams of orthopaedic or surgical units performing hip (146 units) or knee (142 units) replacement surgery. Streamlining, improving, and optimising data collection is a continuous effort involving expert groups and all partners. Crucially, this report also contains evidence that hospitals and surgeons are making progress beyond improving data collection. Decreasing early revision rates, fewer outlier implants in the market, and more homogenous early revision rates in general testify real improvements in treatment quality that coincide with SIRIS reporting.

Coverage is one important indicator of the commitment of all parties involved in SIRIS, and it correlates with reporting accuracy. However, it is difficult to assess coverage, because any benchmark registration system will have weaknesses. For SIRIS, only arthroplasties performed and submitted to the registry as closed records can be used in the analysis, as open cases may experience later modification. In 2024, the coverage of SIRIS was over 98% for primary hip and knee replacements. This not only demonstrates the strong commitment to the project by the surgeons and their teams, both in public and private hospitals, but also the high quality of the data collection, coaching, and organisation by the SIRIS team. Further details regarding the coverage are provided in Chapter 2 Methods, Part 2.3 Coverage.

SIRIS has thus achieved excellent coverage within a relatively short period of time since implementation in 2012 and continues to improve the content of the reports. The SIRIS annual report 2025 provides information on the state of hip and knee replacements in Switzerland and the Principality of Liechtenstein and presents a wealth of new information. The report also provides important and verifiable information that we hope patients, healthcare providers, third-party payers, and healthcare regulators will find useful.

Factors	Variables
Patient related	Name
	Surname
	Date of birth
	Gender
	Height
	Weight
Surgery related	Main diagnosis
	Previous surgery
	Date and place of surgery
	Morbidity state
	Charnley class
	Intervention
	Approach
	Positioning
	Component fixation
Cementing technique	
Implant related	Type of implant
	Article number
	LOT number
	Company name
	Brand name

Table 1.1
Variables collected by the SIRIS registry

2. Methods

2.1 Maintenance and hosting of the registry

The SIRIS registry is hosted and maintained by SwissRDL at the Institute for Social and Preventive Medicine (ISPM), University of Bern. A dedicated team comprising a statistician/methodologist, a data monitoring team, data management/IT specialists, as well as support staff, is responsible for management and maintenance, technical support, analysis, and reporting of the data. The data monitoring team supervises data quality, provides user support and trains collaborators at the participating hospitals to ensure correct and efficient data entry into the registry. The overall project management support at SwissRDL is provided jointly by the data monitoring team and the statistician/methodologist, and both are present in the SIRIS Scientific Advisory Board that directs and oversees the registry and produces this annual report, among other things.

SIRIS data are collected via an online documentation platform (accessible at siris.memdoc.org) to record data on primary arthroplasties, reoperations and component revisions. Furthermore, clinics may also register post-operative follow-up data at their discretion. All individual implants used (including minor components) are registered alongside clinical data from the surgical procedures. The current versions of the SIRIS CRF (v2021) can be downloaded from www.siris-implant.ch. While most participating surgical units use the standard online interface or dedicated apps (mostly on tablet devices) for data registration, some (mostly large) centres send data exports directly from their hospital information systems to SwissRDL via a web service client. Implant data are mostly entered into SIRIS by scanning the barcodes on the implant tags, which also is the recommended procedure, being most reliable. Manual data entry of implants is also possible with multiple-choice drop-down menus containing known implants. New implants first need to be registered by SwissRDL at the request of SIRIS users or upon notification by a producer.

The data of the SIRIS registry are stored on dedicated and protected servers at the University of Bern. Patient identification data (e.g. medical record number, name, and date of birth) are stored on a specific, physically separate, module server, for reasons of data protection and anonymisation. The patient's identification information is encrypted into a salted hash code to facilitate the linking of revisions performed at different healthcare facilities. This is needed to ensure continuity of implant follow-up and to calculate revision rates. Data protection complies with current standards and the methodology of separating the clinical data from the patient-identifying information was reviewed and approved by data protection delegates from the Swiss federal authority. Patients must provide written informed consent for data registration into SIRIS, which is ensured by participating surgeons and hospitals. Furthermore, patients have the right at any time to withdraw participation, check their data, or have their data deleted. Surgeons and hospitals may also use their specific SIRIS data, with certain restrictions applied, for internal quality assessment. The SIRIS Foundation, as the holder of the entire dataset, makes selected parts available in anonymised form upon request for independent academic research.

It is essential that only patients confirmed alive and residing in Switzerland are considered at risk for the analysis of revision rates. Patients who have died or left the country during the observation period are accounted for until death or until leaving Switzerland. For this purpose, patient data from SIRIS were cross-checked with both the database of the Swiss Central Compensation Office (ZAS Geneva) and the Federal Statistical Office (FSO Neuchâtel). Until 2023, SIRIS could verify annually whether a patient had died or left Switzerland. Fewer than 5% of patients had an unknown status or were foreigners operated on in Switzerland without being registered residents. These patients were considered lost to follow-up, unless later revised again in Switzerland, and were subsequently excluded from the analysis of (long-term) re-

vision rates. However, the agreement with the Federal Statistical Office regarding data exchange ended in 2024 and the last update of mortality and migration data was received in April 2023. For the continued application of censoring events beyond April 2023 a simulation model based on 500+ patient typologies per joint was developed in order to project censoring events for a limited time horizon. The purpose of this simulation is not to predict a death or an emigration event (which would be entirely unfeasible), but merely to apply known attrition rates from observed life table information to specific homogeneous cohorts of patients. Patients with unknown status after April 2023 are selected at random for simulated censoring events, replicating the original distributional properties of these events and taking into account known revision or other primary operations. In other words, these simulated events are just “standing in” for the real censoring events until they can be updated again, but the impact on estimates revision rates is entirely comparable to that of the real events if the assumed event proportions per group and year are reasonable estimates. It should be noted that not taking into account censoring events beyond a certain date is certainly introducing bias, especially in groups with high mortality, and therefore even an imperfect adjustment is preferable to no adjustment. Work is ongoing regarding a new procedure for linking again these crucial data to SIRIS.

2.2 Data quality and completeness

The data for this report were exported from the database in June 2025. The consistency and completeness of SIRIS data are checked in part through systematic software-generated validation tests, and additionally, quarterly by the registry’s statistician/methodologist, running automated analysis scripts for identification of likely data errors. These are then fed back to the data monitoring team, which analyses root causes of confirmed problems and provides feedback to affected hospitals.

In addition to the ongoing data quality checking routines, several specific methodological decisions have been made to increase accuracy of reporting. For example, when the information provided on a form and the registered implants contradict each other (e.g. hemiarthroplasty is selected on the form, while THA components were registered), and it has not yet been possible to verify the case, then the implant registration information is given priority (in this example, the case is provisionally accounted for as a THA). This information is indicated in the relevant tables or figures if such decisions are likely to impact the overall results.

Two updates of the CRF have been implemented since the launch of SIRIS. The first version was used from 2012 to 2014, and an updated version was in use between 2015 and 2020. Some changes to the definition of existing variables (particularly for the type of arthroplasty of the knee) were introduced with this update, as well as some new variables, notably the body mass index (BMI) and the morbidity state according to the ASA classification. Inconsistent use among providers of the answer option “unknown” (one institution even reporting unknown ASA status in almost all cases) indicates issues with data collection. Other common problems included impossible or inconsistent responses, which are more frequently observed in some sections of the forms than in others, like revisions relating to acetabular components in hemiarthroplasties. This could be due to a system-

atic misunderstanding of the meaning of certain response categories (i.e. confusion between revision of the acetabular component and conversion to THA after hemiarthroplasty) or because of random data entry errors likely aggravated by design issues such as long drop-down lists. The latest CRF version 2021 has successfully addressed some of these problems and has led to clear improvements in data quality. SIRIS conducted organisational surveys in 2023 and 2024. Hospitals were surveyed on several issues including data entry mode, local responsibilities, coverage and availability of PROMs. A noteworthy finding of the surveys was that paper forms still play a major role in data collection. Two thirds of hospitals use them, typically to collect registry data on paper first before registration in the online system. Only a minority of hospitals register cases directly from the operation theatre or shortly after the operation. This is reflected in the relatively long delay in collection of SIRIS data. Only 26% of procedures were registered and completed within 24 hours and 65% within four weeks of operation. Nine percent of cases were registered with a delay of six months or longer. Other noteworthy findings were that most hospitals do not yet use specific PROMs, and most of those which do, already use the ones that SIRIS has decided to use in the near future.

2.3 Coverage

Reliable reference data from other sources are needed to estimate the coverage of SIRIS. Nevertheless, any benchmarking system has its specific weaknesses and disadvantages, resulting in a certain degree of incompleteness.

One option is to compare the annual number of cases reported in the registry with the numbers from quality indicators for Swiss acute care hospitals as published by the Federal Office of Public Health (FOPH/BAG). This encompasses a complete annual survey of all hospital discharges in Switzerland, whereby each entry represents the hospital discharge and includes information about the patient's socio-demographic characteristics, diagnosis, and treatment. Codes I.1.8.F, I.1.9.F, and I.1.10.F can be used to identify primary hip prostheses of any kind and for any diagnosis, while codes I.1.15.F and I.1.16.F are used for total and partial knee prostheses. At the time of writing the 2025 annual report, only figures up to 31.12.2023 were available. These figures are published online, but only with a considerable time lag (detailed information provided at www.bag.admin.ch). As shown in **Table 2.1**, primary coverage peaked at 98.7% in 2022, which includes the best value ever for hip arthroplasties and the third best for knee arthroplasties. Having to rely on figures publicly available with a considerable time lag is suboptimal for a registry, but efforts to secure timelier access to the actual raw figures reported by individual hospitals to FOPH/BAG have been rejected on legal grounds.

SIRIS also accesses annual implant sales figures from the participating industry partners in Switzerland, specifically the number of femoral stems (indicator for hip arthroplasties) and tibia plateaus (indicator for knee arthroplasties) sold per year. In previous years, the two different ways of calculating coverage rates were mostly in agreement. However, starting in 2021 and observed increasingly in 2022/2023, we find that these figures are no longer in agreement with registry data or other sources, at least not on a

calendar year basis and for specific hospitals. Analysis strongly suggests that yearly sales and implant use figures in hospitals do not always agree. In other words, hospitals can report more procedures per year than implant sales in Switzerland suggest, resulting in a coverage rates above 100%. We also became aware of the possibility that implants may be imported directly from foreign suppliers and are therefore not counted among official sales in Switzerland. However, it is reasonable to assume that such discrepancies tend to even out over time and across hospitals, or appear to be relatively small. For this type of analysis, we thus consider coverage rates between 90% and 110% as the target zone. Feedback from individual manufacturers to implant reports provided by SIRIS indicates that these high coverage rates are realistic. For instance, in specific implant reports, coverage rates tended to be as high as 100% for typical standard implants such as primary hip stems and as low as 80% for less common implants such as acetabular reinforcement cages.

Nevertheless, feedback on implant figures confirms the observed general upward trend in coverage. We also observed a clear progression of coverage at the hospital level since 2017, as all eligible units are currently submitting cases to SIRIS. In previous years, we had reasons to believe that the registry already had a higher, albeit not officially counted, coverage rate, as cases created in the online registration system need to be completed before they can be included in the analysis. For most procedures, at least one implant must be registered before a case may be recognised as completed by the system. A certain number of incomplete and unsubmitted cases are left in the system every year, mostly caused by missing implant entries. The improvements in coverage since 2017 are, to a certain extent, due to monitoring of and support to hospitals by SIRIS to solve submission problems. As a direct result of these efforts, the number of registered cases keeps increasing even after closure of each reporting period.

	2017	2018	2019	2020	2021	2022	2023
Primary hip prostheses							
BAG	22,970	23,160	23,619	23,310	24,834	26,435	27,691
SIRIS*	20,992	21,739	22,462	22,747	24,344	26,066	27,266
Primary coverage (%)	91.4	93.9	95.1	97.6	98.0	98.6	98.5
Primary knee prostheses							
BAG	18,558	18,325	19,181	18,837	20,280	23,070	24,569
SIRIS**	17,108	17,440	18,546	18,588	19,850	22,686	24,140
Primary coverage (%)	92.2	95.2	96.7	98.7	98.0	98.3	98.3
All primary hip and knee prostheses							
BAG	41,528	41,485	42,800	42,147	45,114	49,505	52,260
SIRIS	38,100	39,179	41,008	41,335	44,194	48,752	51,406
Primary coverage (%)	91.7	94.4	95.8	98.1	98.0	98.5	98.4

Table 2.1

Retrospective coverage analysis 2017–2023 based on National Office of Public Health figures (BAG)

All SIRIS figures excluding Liechtenstein

* I.1.8.F/I.1.9.F/I.1.10.F (all first hip prostheses, all diagnoses)

** I.1.15.F/I.1.16.F (all first knee prostheses, all diagnoses)

In recent years, however, at least part of the gap in data entries could be explained by increasing difficulties in obtaining informed consent from patients. This is a topic to observe in the future, as participation refusal poses a direct threat to the quality of any implant registry, as very high – ideally complete – coverage of all primary and revision procedures is needed for reliable analysis. The very principles of consent-based participation and general data protection rules denying the registry access to certain data without explicit consent are in conflict with the registry’s mission of nationwide quality control.

In this report we added a new type of geographic analysis based on the BAG figures reported in **Table 2.1**. **Figures 2.1 and 2.2** show the cantonal coverage rates for primary hip and knee procedures. In several Cantons SIRIS achieved 2023 coverage rates above 99%, whilst in others they were below 90%. This picture is undoubtedly subject to annual variation as specific data submission problems in important hospitals have the potential to easily distort submission rates by several percentage points. It is also noteworthy that submission rates for hip and knee are not necessarily in agreement at the cantonal and therefore at the hospital level. This highlights the importance of ongoing monitoring in each participating surgical unit.

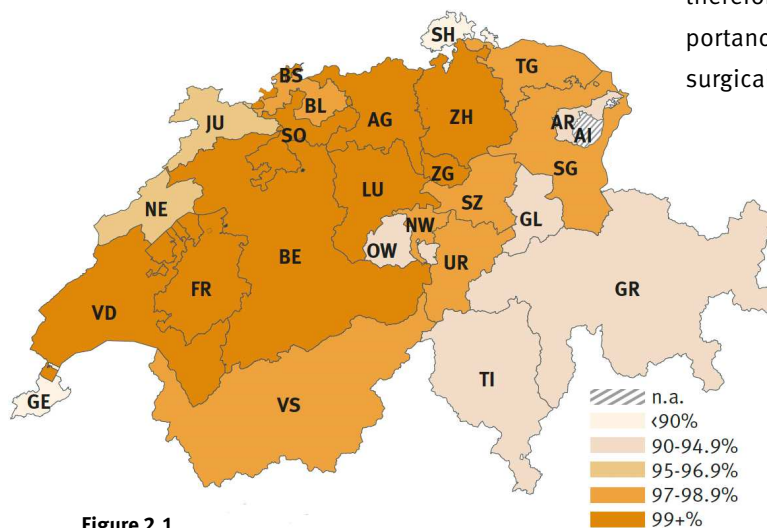


Figure 2.1
Coverage rates for primary hip prostheses in the SIRIS registry 2023

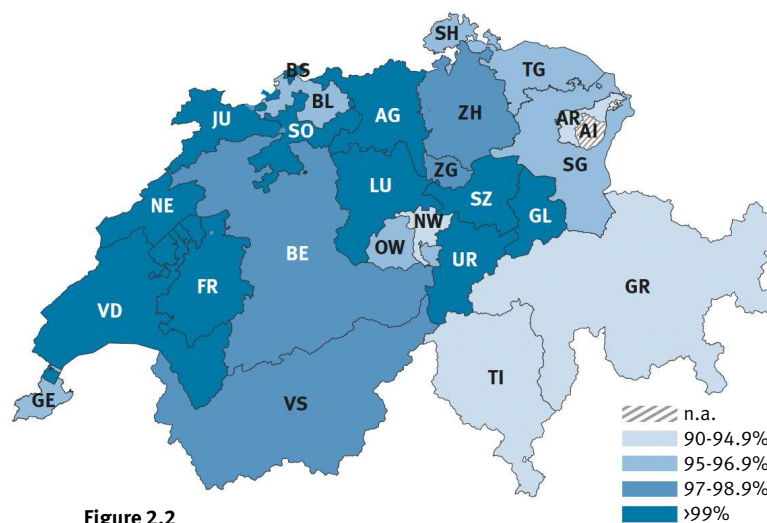


Figure 2.2
Coverage rates for primary knee prostheses in the SIRIS registry 2023

2.4 Statistical precision, evaluation and outlier detection

The figures in this report are, whenever appropriate, accompanied by 95% confidence intervals. A confidence interval indicates the plausible range within which the true value should lie with the indicated probability, considering random variation of samples of limited size. For an implant combination with large numbers, the confidence interval is usually narrow and, as numbers get smaller, the statistical precision decreases, which results in wider confidence intervals. For practical purposes, any position within the confidence interval should be seen as a plausible value and if confidence intervals overlap, they should be regarded as not statistically different. All confidence intervals are unadjusted for the various forms of clustering that may also affect precision, especially when results depend on small numbers of surgeons, hospitals or implants. The latter aspect is a particular challenge for a medical registry in a small yet diverse country such as Switzerland and must be evaluated on a case-by-case basis during any outlier detection. In 2022, we started reporting a simple metric that we called the “case concentration score” (CCS) to address this issue. The CCS represents the share of a particular implant combination or system accounted for by the hospital using it most often. We provide this information in the interest of transparency as the performance of implants used in few places may not be an unbiased estimate of their true performance. Since the diversity of knee systems has also been increased to include more complicated designs, we also provide mean age figures for each system, as long as these designs were used for primary osteoarthritis. This is done in the interest of transparency, as knee systems used in younger patients tend to have higher revision risks not necessarily related to implant quality. Several means are used for detection of outliers. For clinics and individual surgeons (the latter not being part of the scope of this report), we rely on risk-adjusted funnel plots and use the 99.8%

limit as the relevant threshold. In other words, a clinic is deemed an outlier if the 2-year revision rate is higher than the range of plausible observations in which 99.8% of observations would fall in case of random variation. The likelihood to observe a value outside of this range (above or below the limits) just due to chance would be 1 in 500. Indeed, the specific likelihood of exceeding such an upper boundary solely by chance would be only 1 in 1,000.

For implants, we use a much simpler method but report the results with several caveats and additional context. In this report, we continue with a distinction between the 2-year revision rate and the mid-term evaluation, whereby the latter starts at 5 years follow-up and currently ends at 11 years follow-up. All implant combinations or systems with at least 500 cases initially are evaluated. In the short-term (2-year) evaluation, we rate an implant as potential outlier if the observed 2-year revision rate is more than twice as high as for the average of the reference group (“outlier alert boundary”). The threshold for inclusion in this analysis was at least 50 cases reported in the current evaluation time frame (i.e. all primary operations between 1.1.2019 and 31.12.2022). We benchmark implants within a moving time window of 4 years to ensure that results represent current performance, not being affected by previous period effects. In this report, we refrained from ranking the implants by their 2-year revision rates and we excluded any potential outliers with confidence intervals overlapping with the 95% confidence interval of the reference group average. In other words: the outlier status comes with varying degrees of statistical probability and is considered highly likely when both the estimated revision rate and the complete confidence interval exceed the outlier alert boundary. For this reason, an implant whose revision rate exceeds double the mean revision rate, while the confidence interval overlaps with the outlier boundary, is defined as a potential outlier. If the lower bound of the confidence interval exceeds twice the mean revision rate, it is considered a definitive outlier. Furthermore, we

identify three possible deviations from normal mid-term performance:

- 1.** Implants with elevated revision risk, i.e. those whose revision rate is increased by a factor ranging from 1.5 to 1.99 relative to the group average at any time point of at least 5 years follow-up;
- 2.** Implants with long-term outlier status, i.e. those with a revision rate that is increased by at least a factor 2 relative to the group average at any time point of at least 5 years follow-up; and
- 3.** Implants with below average revision risk, i.e. those with a revision rate that is decreased by a factor of at least 0.66 relative to the group average at longest available follow-up. In other words, the revision rate of the implant cannot be more than 66% of the relevant group average to be rated as having revision risk below average. All these analyses are subject to further limitations linked to remaining numbers at risk over time and consecutive spread of confidence intervals as specified in the relevant chapters.

We thus benchmark implants directly against the relatively narrow field of comparable products in their normal variety of uses. There is no further risk adjustment, as similar products are already meant to be used for a particular range of comparable patients and diagnoses. However, detailed outlier reports are produced for manufacturers and affected hospitals, and we also provide additional analytical information such as risk-adjusted hazard ratios in this context. As implant group sizes vary markedly, readers must pay attention to the reported confidence intervals and any other contextual information – especially relating to small numbers of clinics involved – indicated on the outlier watch board in this report.

To help readers understand the grouping decisions of implant combinations or systems, an additional online appendix to this report is provided. The appendix lists all implants and provides additional information on the group composition (e.g. included stem or cup variants), and whether or not there is a likelihood of so-called camouflage effects, where unreported subgroups may influence the overall result.

2.5 Patient-reported outcome measures (PROMs)

SIRIS benefitted from two local PROMs initiatives that were conducted using the registry's platform. National evaluation of PROMs, with the Swiss Orthopaedics minimal dataset (SO-MDS) at its core, is planned for autumn 2025 on a voluntary basis.

Between 2017 and 2020, nine hospitals in the Cantons of Basel Stadt, Basel Landschaft, and Solothurn collected PROMs for elective THA and TKA using the Core Outcome Measures Index (COMI). Patients were followed up after 6 months and 2 years postoperatively. Another PROMs project in collaboration with SIRIS was initiated in 2019 by the Cantonal Health Authority of Zurich in conjunction with Swiss Orthopaedics. All hospitals receiving public funding for elective hip and knee arthroplasties in the Canton of Zurich had to collect the Swiss Orthopaedics minimal dataset (SO-MDS), consisting of the 5-item version of the EQ-5D quality of life questionnaire and a small selection of additional questions on joint-specific pain and satisfaction. Patients were followed up one year postoperatively. Several other hospitals in other cantons also provided the same data. In the previous years, we reported both PROMs pilot studies for hip and knee results separately. In this report, we continue to report only on the SO-MDS results from all available hospitals and contrast hip and knee results.

The primary patient inclusion rate of all participating hospitals from the Canton of Zurich reached 73% in 2023 and the 1-year follow-up rate for primary cases from 2022 with valid SO-MDS was 79% for hip prostheses and 73% for knee prostheses. These rates remained stable in 2024. The SO-MDS forms, currently provide data with follow-up at 1 year from 9,065 THA patients with primary osteoarthritis, 974 THA patients with secondary osteoarthritis, 7,087 TKA patients with primary osteoarthritis and a further 991 with secondary osteoarthritis. They also provide data with complete 1-year follow-up for 1,351 PKA patients with primary osteoarthritis.

The analysis focused on the so-called treatment effect (TE). This was expressed on one hand as pain reduction and on the other hand as general quality of life improvement. The calculations are performed on converted scales, with 0 defined as no symptoms and positive values designating levels of symptoms. TE is based on the following formula:

$$TE = \frac{\text{preoperative score} - \text{postoperative score}}{\text{preoperative score}}$$

In other words, this metric is the relative symptom reduction expressed on a numerical scale, whereby 1 equals complete symptom regression, 0 representing the complete absence of effect (e.g. same pain reported as before treatment), and a negative value represents an outcome worse than the pre-operative state (e.g. more pain than before treatment). The analysis excludes patients who did not report pre-operative symptoms (= 0). The scale can be presented as approximate percentage categories for comparison between settings or types of procedures. However, readers are advised that this purely numerical analysis may differ from other clinical evaluations of treatment success. We used TE for both the pain score and the EQ-5D summary scores.

¹ Huber J, Irlenbusch U, Käähb MJ, Reuther F, Kohut G, Judge A.

Treatment effects of reverse total shoulder arthroplasty – a simple method to measure outcomes at 6, 12, 24 and 60 months for each patient. *BMC Musculoskelet Disord.* 2020 Jun 22;21(1):397. doi: 10.1186/s12891-020-03427-7. PMID: 32571282; PMCID: PMC7310507.

² Huber J, Dieppe P, Dreinhoefer K, Günther KP, Judge A.

The Influence of Arthritis in Other Major Joints and the Spine on the One-Year Outcome of Total Hip Replacement: A Prospective, Multicenter Cohort Study (EUROHIP) Measuring the Influence of Musculoskeletal Morbidity. *J Bone Joint Surg Am.* 2017 Sep 6;99(17):1428-1437. doi: 10.2106/JBJS.16.01040. PMID: 28872524; PMCID: PMC5685421.

3. Demography and Epidemiology

3.1 Introduction

Switzerland has an intricate and arguably unique healthcare system. In addition to a national health department, which plays a coordinating role, each canton holds full responsibility for its own healthcare services. In fact, the federal government of Switzerland acts on the principle of subsidiarity, intervening only for questions of nationwide importance – as it was the case during the COVID-19 pandemic. Even university hospitals are not part of a centralised, national healthcare system. Instead, they are owned and managed by the individual cantons in which they are located. Furthermore, alongside the public healthcare service overseen by the cantons, a significant portion of the healthcare provision is organised privately. However, most private hospitals hold a public service mandate of the cantons and take part in the public healthcare system of the cantons.

Healthcare insurance is mandatory in Switzerland, but it is likely as complex as the organisation of the healthcare services. Currently, there are three levels of inpatient healthcare insurance coverage: general/public, semi-private and private. In the event of a hospital stay, the patient pays only a small proportion of the costs, while the large majority is covered jointly by the insurance company and the canton, the exact amount and distribution of costs depending on the insurance model. A nationwide insurance system exists for accidents, covering all related expenses, including compensation for work incapacity or a pension, if necessary.

Traditionally, the French speaking, western part of Switzerland has maintained close ties with France, just as the canton Ticino has with northern Italy, or the German speaking part has with Germany and Austria. This has made Switzerland something of a melting pot of different concepts and approaches in orthopaedic surgery, particularly in hip and knee arthroplasty. It may also explain why we are able to report on nearly every existing brand and technology, even only in relatively small numbers due to the country's small size. In this context, it is a fortunate circumstance that the Swiss joint registry is organised at the national level and, thanks to mandatory reporting, achieves a participation rate of 98.5%.

Because of the abovementioned fragmentation of the healthcare system, the chapter Epidemiology will highlight results at the national level, as well as by canton or region, and finally at the level of individual hospitals. The Principality of Lichtenstein is included in the analysis.

3.2 Hip arthroplasty

3.2.1 Nationwide data, Switzerland and Principality of Liechtenstein

Incidence of THA and HA

Since introduction of SIRIS in 2012, a total of 251,930 primary THA and 28,039 primary HA were registered, along with 33,201 revision arthroplasties of the hip. Overall, 41.2% of the revisions could be linked to a previously registered primary or revision procedure. The proportion of linked revisions increased from 13.1% in 2012 to 57.2% in 2024. However, 43.8% of the revisions remain unlinked, mainly because the index procedure was performed prior to implementation of SIRIS in 2012, or because the index arthroplasty was not reported (Table 3.1).

The incidence of THA has steadily increased over the years, from 228 per 100,000 inhabitants in 2013 to 286 per 100,000 inhabitants in 2024. This corresponds to an average annual increase of approximately 1%. Among the population most at risk – the age group between 50 to 89 years, who account for 98% of the THA recipients – the incidence reached 672 per 100,000 inhabitants at risk in 2024 (Figure 3.1). This corresponds to a yearly growth rate of approximately 2%. As this exceeds the overall growth rate of the incidence, it indicates an age-biased distribution of the population.

Year	Primary THA	Primary HA	Primary others or type unclear	Primary total ¹	Annual growth rate THA	Annual growth rate HA	Linked Rev./ Reop. of THA ²	Linked Rev./ Reop. of HA ²	Unlinked Rev./ Reop. ³	Rev./ Reop. total ⁴	% Linked Rev./ Reop.
2012*	6,704	637	3	7,344			113	6	792	911	13.1
2013	16,900	1,935	3	18,838			409	39	1,871	2,320	19.3
2014	17,184	2,030	1	19,215	1.7%	4.9%	572	60	1,900	2,533	25.0
2015	17,696	1,991	5	19,692	3.0%	-1.9%	726	65	1,812	2,604	30.4
2016	18,712	2,007	4	20,723	5.7%	0.8%	843	85	1,714	2,644	35.1
2017	18,888	2,098	9	20,995	0.9%	4.5%	865	78	1,681	2,629	35.9
2018	19,530	2,263	5	21,798	3.4%	7.9%	970	101	1,566	2,638	40.6
2019	20,201	2,359	8	22,568	3.4%	4.2%	1,109	106	1,529	2,747	44.2
2020	20,362	2,430	5	22,797	0.8%	3.0%	1,250	108	1,461	2,825	48.1
2021	22,117	2,409	8	24,534	8.6%	-0.9%	1,356	118	1,340	2,815	52.4
2022	23,492	2,629	1	26,122	6.2%	9.1%	1,342	138	1,304	2,789	53.1
2023	24,669	2,605	5	27,279	5.0%	-0.9%	1,472	125	1,339	2,936	54.4
2024	25,475	2,646	1	28,122	3.3%	1.6%	1,468	140	1,200	2,810	57.2
All	251,930	28,039	58	280,027			12,495	1,169	19,509	33,201	41.2

Table 3.1

Total and partial hip arthroplasty (THA and HA), primary and revisions/reoperations

All documented operations

¹ Does not represent a full year of data, as data collection in most hospitals started only in October 2012

² i.e. primaries already registered in SIRIS

³ can be of THA and HA

⁴ including linked revisions/reoperations of procedures that were classified as „primary others“ or of unclear type

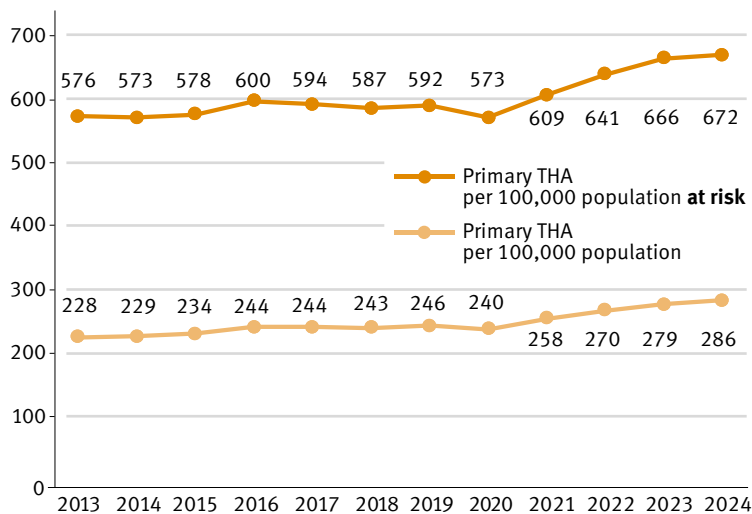


Figure 3.1
Incidence of primary total hip arthroplasties registered in SIRIS

Per 100,000 residents and per 100,000 residents at-risk*.
 Adjusted for estimated coverage. SIRIS figures excluding Liechtenstein.
 Coverage rates 2013–2016 estimated at 91% ; 2017–2023 based on federal health office data; 2024 estimated at 98.5%.
 *Age group 50–89 years accounts for 93% of all recipients of THA

Looking at the incidence per 100,000 population, a significant drop occurred in 2020 due to restrictions linked to the COVID-19 pandemic. Before 2020, the average annual increase was approximately 0.5%. The growth rate peaked at 6.3% in 2021 and 5.3% in 2022, then declined to 3.9% in 2023 and further to 0.9% in 2024. Much of the surge in 2021 and 2022 appears to be a rebound effect following the COVID-19 pandemic.

Growth rates of primary hip arthroplasty by cantons

Between 2019 and 2024, the absolute growth rate was 24.6% nationwide. The growth rate per canton shows a heterogeneous pattern (Figure 3.2). However, the figure illustrates the increase of THA performed by each canton, but the origin of the patient remains unknown. Patient flows may not be analysed from the data available to SIRIS.

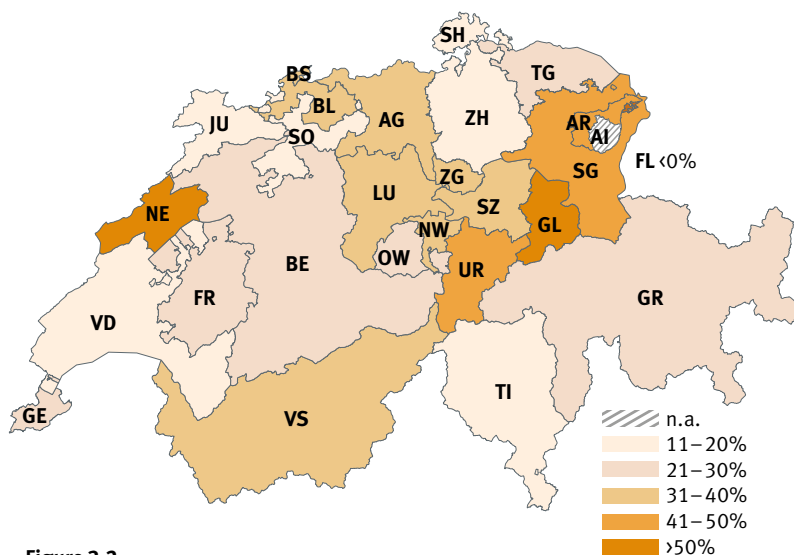
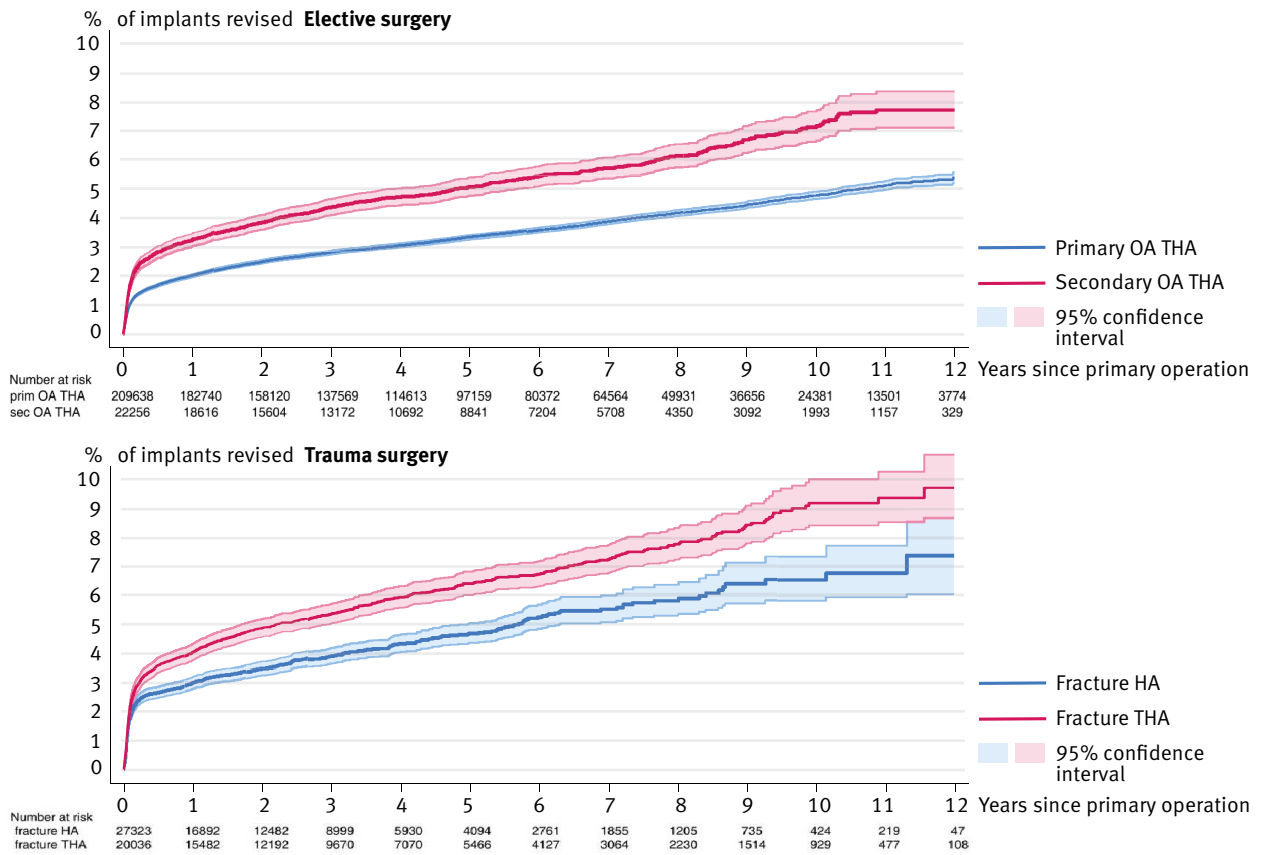


Figure 3.2
Growth rates of primary hip arthroplasties by canton: 2019 – 2024

Revision rates

Revision rates increased over time since the index arthroplasty. The revision rate differs between reasons leading to surgery. Analysing data from 2012 onwards, at 2 years, the average revision rate for primary THA was 2.5% (CI 2.4 – 2.6%) for primary OA, whereas it was 3.9% (CI 3.6 – 4.1%) for secondary

OA and as high as 4.9% (CI 4.6 – 5.2%) for primary THA performed for fracture. The 2-year revision rate for HA was 3.5% (CI 3.2 -3.7%). At 12 years, the revision rates were 5.4% (CI 5.2 – 5.6%), 7.7% (CI 7.1 – 8.4%), 9.7% (8.7 – 10.9%) and 7.4% (6.0 – 9.0%), respectively. Further details are provided in **Figure 3.3**.



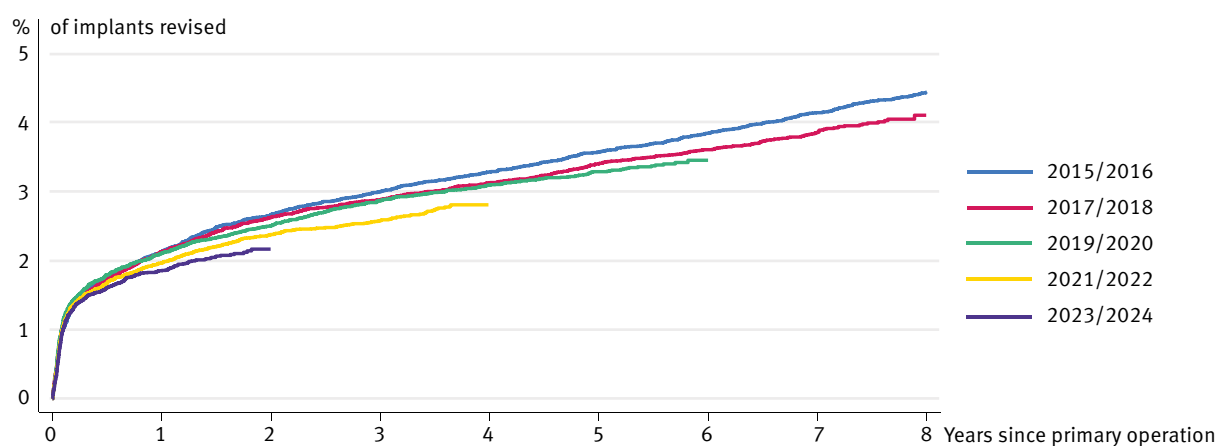
Failure rate	1 year	2 years	3 years	5 years	6 years	8 years	9 years	10 years	12 years
Prim OATHA	2.0 (2.0-2.1)	2.5 (2.4-2.6)	2.8 (2.7-2.9)	3.3 (3.3-3.4)	3.6 (3.5-3.7)	4.2 (4.1-4.3)	4.5 (4.3-4.6)	4.8 (4.7-4.9)	5.4 (5.2-5.6)
Sec OA THA	3.2 (3.0-3.5)	3.9 (3.6-4.1)	4.4 (4.1-4.7)	5.1 (4.8-5.4)	5.4 (5.1-5.8)	6.1 (5.7-6.5)	6.7 (6.2-7.2)	7.1 (6.6-7.7)	7.7 (7.1-8.4)
Fracture HA	3.0 (2.8-3.2)	3.5 (3.2-3.7)	3.9 (3.7-4.2)	4.7 (4.4-5.1)	5.3 (4.8-5.7)	5.8 (5.3-6.4)	6.4 (5.7-7.1)	6.5 (5.8-7.3)	7.4 (6.0-9.0)
Fracture THA	4.0 (3.8-4.3)	4.9 (4.6-5.2)	5.4 (5.0-5.7)	6.4 (6.0-6.8)	6.7 (6.3-7.2)	7.8 (7.2-8.3)	8.4 (7.8-9.1)	9.2 (8.4-10.0)	9.7 (8.7-10.9)

Figure 3.3

Kaplan Meier estimate of cumulative postoperative revision risk after primary hip arthroplasty

Time since operation, 2012 – 2024, all services, all diagnoses

The comparison of different periods since 2015 shows a trend of decreasing revision rates over time, particularly since 2021/2022 (Figure 3.4). A statistically significant reduction is present for the one-year revision rate between the periods 2015/2016 and 2023/2024, and also for the three-year revision rates between 2015/2016 and 2021/2022. Such a decrease over time is one of the desired effects of a registry.



Failure rate	1 year	2 years	3 years	4 years	5 years	6 years	7 years	8 years
2015/2016	2.1 (2.0-2.3)	2.7 (2.5-2.9)	3.0 (2.8-3.2)	3.3 (3.1-3.5)	3.6 (3.4-3.8)	3.8 (3.6-4.1)	4.1 (3.9-4.4)	4.4 (4.2-4.7)
2017/2018	2.1 (2.0-2.3)	2.6 (2.5-2.8)	2.9 (2.7-3.1)	3.1 (2.9-3.3)	3.4 (3.2-3.6)	3.6 (3.4-3.8)	3.9 (3.6-4.1)	
2019/2020	2.1 (2.0-2.3)	2.5 (2.3-2.7)	2.9 (2.7-3.0)	3.1 (2.9-3.3)	3.3 (3.1-3.5)			
2021/2022	2.0 (1.8-2.1)	2.4 (2.2-2.5)	2.6 (2.4-2.7)					
2023/2024	1.9 (1.7-2.0)							

Figure 3.4

Kaplan Meier estimate of cumulative postoperative revision risk after THA by time period

Time since operation, 2015 – 2024, primary OA

3.2.2 Primary total hip arthroplasty

Since the introduction of SIRIS in 2012, a total of 251,930 primary THA have been registered. Over the last six years, 136,316 primary THA were registered, forming the base of the current analysis (Table 3.2). The registry discriminates between THA performed for primary OA (81.4%, the largest group), for secondary

OA (9.3%, which includes post-traumatic OA, OA due to inflammatory diseases, avascular necrosis, and sequels of childhood diseases such as developmental dysplasia of the hip or Perthes' disease), and for fractures of the hip (9.3%). The proportion of primary OA continuously declined from 84.4% in 2018 and to 80.5% in 2024, while the proportions of secondary OA and fracture cases increased proportionally.

		2019	2020	2021	2022	2023	2024	2019 – 2024
N		20,201	20,362	22,117	23,492	24,669	25,475	136,316
Diagnosis [%]	Primary OA	83.6	82.1	81.0	80.6	81.0	80.5	81.4
	Secondary OA	8.5	9.1	9.4	9.6	9.4	9.8	9.3
	Fracture	7.9	8.8	9.6	9.7	9.6	9.7	9.3
Women [%]		53.1	52.3	53.8	53.3	53.5	53.5	53.3
Mean age (SD)	All	69.1 (11.5)	69.0 (11.6)	69.2 (11.7)	69.5 (11.5)	69.4 (11.4)	69.4 (11.5)	69.3 (11.5)
	Women	70.8 (11.2)	70.6 (11.4)	70.8 (11.5)	71.0 (11.2)	70.8 (11.0)	70.9 (11.2)	70.8 (11.3)
	Men	67.1 (11.6)	67.1 (11.6)	67.4 (11.7)	67.7 (11.6)	67.6 (11.5)	67.7 (11.7)	67.5 (11.6)
Age group [%]	<45	2.5	2.5	2.7	2.4	2.5	2.7	2.5
	45–54	8.6	8.9	8.6	7.9	7.5	7.5	8.1
	55–64	21.6	21.9	21.3	22.0	22.2	22.3	21.9
	65–74	32.3	31.6	30.9	30.6	31.4	30.4	31.2
	75–84	27.7	27.8	28.7	29.5	29.0	29.6	28.8
	85+	7.3	7.3	7.8	7.7	7.4	7.6	7.5
N unknown BMI (%)		2,925 (14)	2,515 (12)	1,964 (9)	1,333 (6)	1,275 (5)	927 (4)	10,939 (8)
N known BMI		17,276	17,847	20,153	22,159	23,394	24,548	125,377
Mean BMI (SD)		27.0 (5.0)	26.9 (5.1)	26.9 (5.2)	26.9 (5.2)	26.9 (5.2)	26.9 (5.2)	26.9 (5.1)
BMI [%]	<18.5	2.1	2.3	2.2	2.2	2.2	2.1	2.2
	18.5–24.9	35.6	36.5	36.2	36.6	36.5	36.9	36.4
	25–29.9	39.1	38.1	37.4	36.8	37.4	37.5	37.6
	30–34.9	16.6	16.6	17.3	17.6	16.9	16.5	16.9
	35–39.9	5.2	4.8	5.1	5.2	5.2	5.0	5.1
	40+	1.5	1.7	1.9	1.7	1.8	1.9	1.8
N unknown ASA (%)		1,497 (7)	1,238 (6)	738 (3)	385 (2)	378 (2)	334 (1)	4,570 (3)
N known ASA		18,704	19,124	21,379	23,107	24,291	25,141	131,746
Morbidity state [%]	ASA 1	12.1	11.6	11.2	9.9	9.6	8.9	10.4
	ASA 2	59.1	59.0	57.8	58.9	59.3	58.8	58.8
	ASA 3	27.9	28.3	29.8	29.9	29.9	31.2	29.6
	ASA 4/5	0.8	1.0	1.2	1.3	1.2	1.1	1.1

Table 3.2

Primary total hip arthroplasty: Baseline patient characteristics by year

Sex and age

Both the age and the male/female ratio remained stable over time for primary THA. Women represented 53.3% of the primary THA registered, and their mean age of 70.8 years was higher than the 67.5 years of men.

The distribution among the age groups showed minimal changes over the last 6 years. Overall, 67.5% of THA were performed in patients older than 65 years of age. Patients aged over 85 years represented 7.5%, while patients aged <55 years constituted 10.6% of the recipients. The proportion of THA performed on patients <45 years of age remained constant over the years, oscillating between 2.5 and 2.7%. This corresponds to 3,470 patients younger than 45 years. In contrast, the largest group were patients between 65

and 74 years of age, representing 31.2% or 42,473 patients (Table 3.2). For fractures, most patients were 85 years and older (Figure 3.5).

BMI and ASA score

BMI and ASA scores are recorded in SIRIS only since 2015. Data collection is still improving. Pleasingly, the share of unknown BMI decreased continuously from 16% in 2018 to 5% in 2023 and finally 4% in 2024. This is due to the ongoing efforts of the support team supervising the clinics.

The mean BMI was 26.9 kg/m² for all primary THA, whereby 37.6% were performed in overweight patients (BMI 25 – 29.9 kg/m²) and 23.8% in obese patients (BMI >30 kg/m²) (Table 3.2). Younger patients were observed to have higher BMI, and this observation could be made in both male and female patients (Figure 3.6). The distribution of BMI remained constant during the observation period between 01.01.2019 and 31.12.2024.

Most primary THA were performed on healthy individuals (ASA class 1 and 2), but 32.3% of the THA were performed on patients ASA class ≥3. The proportion of ASA class 3 patients increased from 27.9% in 2019 to 31.2% in 2024. During the same period, the proportion of ASA class 1 patients continued to decrease from 12.1% in 2019 to 8.9% in 2024.

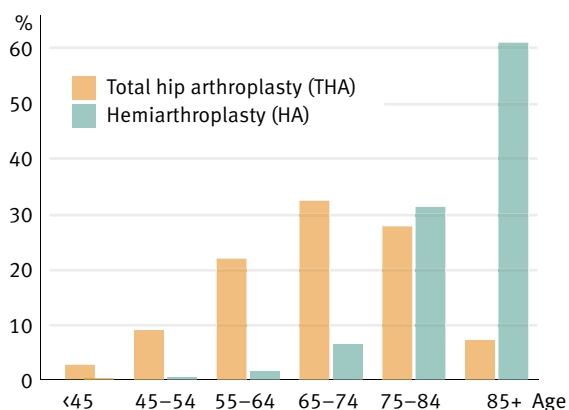


Figure 3.5
Age distribution at surgery of primary total hip arthroplasty and hemiarthroplasty

All documented operations

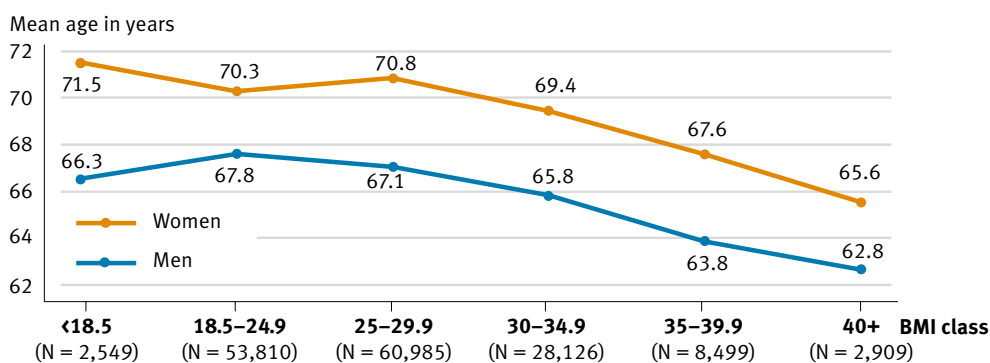


Figure 3.6
Primary total hip arthroplasty: Mean age at primary arthroplasty depending on BMI class

Primary and secondary osteoarthritis patients only. Please note that group sizes vary considerably. All documented operations

		Primary OA	Secondary OA	Fracture
N (2019 – 2024)		110,954	12,706	12,656
Women [%]		51.7	57.4	63.1
Mean age (SD)	All	69.2 (10.8)	63.9 (15.3)	74.9 (10.9)
	Women	70.8 (10.4)	65.5 (15.2)	76.1 (10.3)
	Men	67.6 (10.9)	61.7 (15.3)	73.0 (11.7)
Age group [%]	<45	1.8	11.2	0.8
	45–54	7.7	16.0	3.5
	55–64	22.8	22.3	13.3
	65–74	32.8	22.1	25.6
	75–84	28.7	20.9	37.3
	85+	6.1	7.6	19.5
Diagnosis [%]	Osteoarthritis	99.9	0.0	0.0
	Inflammatory arthritis	0.0	4.0	0.0
	Developmental dysplasia	0.0	27.5	0.0
	Osteonecrosis	0.0	55.4	0.0
	Miscellaneous*	0.1	13.1	2.3
	Fracture	0.0	0.0	97.7
N unknown BMI (%)		8,487 (8)	826 (7)	1,626 (13)
N known BMI		102,467	11,880	11,030
Mean BMI (SD)		27.2 (5.1)	26.7 (5.5)	24.2 (4.5)
BMI [%]	<18.5	1.5	2.7	7.4
	18.5–24.9	34.0	39	55.2
	25–29.9	39.0	34.4	28.3
	30–34.9	18.1	16.2	7.0
	35–39.9	5.5	5.1	1.6
	40+	1.8	2.3	0.5
N unknown ASA		3,752 (3)	338 (3)	480 (4)
N known ASA		107,202	12,368	12,176
Morbidity state [%]	ASA 1	10.7	12.1	6.0
	ASA 2	61.5	53.5	41.3
	ASA 3	27.2	32.8	48.0
	ASA 4/5	0.6	1.6	4.8

Table 3.3

Primary total hip arthroplasty: Baseline patient characteristics by main diagnostic group

* Miscellaneous diagnoses are free text entries that typically describe more complex situations. Most of those must be classified as secondary arthroses, but there are also entries that describe fractures or directly fracture-related conditions such as pathological fractures or complications after osteosynthesis.

Underlying diagnosis

Patients treated for secondary OA were on average 5.3 years younger than those treated for primary OA. The proportion of developmental dysplasia of the hip among all primary THA performed for secondary OA increased from 20.5% in 2015 to 27.5% in 2024, while the majority of this subgroup (55.4%) were treated for avascular necrosis. Compared to the other main diagnostic groups, an increasing number of young patients (<45 years) were treated for secondary OA, reaching 11.2% in 2024 (**Table 3.3**).

Among the patients treated for fractures, there were considerably more women than men, representing close to two-thirds (63.1%). The average age of these women was 76.1 years, compared to 73.0 years for men. More than 80% of fractures occurred in patients over 65 years of age, and more than 55% in patients over 75 years. There was also a much higher proportion of patients in the fracture group belonging to ASA class ≥ 3 (48%).

Regional differences in bearing selection

The choice of bearing surfaces has an important impact on durability of an implant combination. The choice of the material of the femoral head (metal, ceramic, ceramicised metal) and acetabular side (ceramic, standard polyethylene, highly crosslinked polyethylene) is based on patient, surgeon and economic factors. In addition to the material, different head sizes can be used. There are distinct regional differences.

The two most common cup materials are highly cross-linked polyethylene (XLPE) and ceramic inlays. In the cantons of Bern, Obwalden and St. Gallen XLPE is used in 81–90% of the cases. In Zug almost always XLPE cups are used. In contrast, in the Wallis, XLPE is rarely used, and ceramic is the predominant bearing (Figure 3.7).

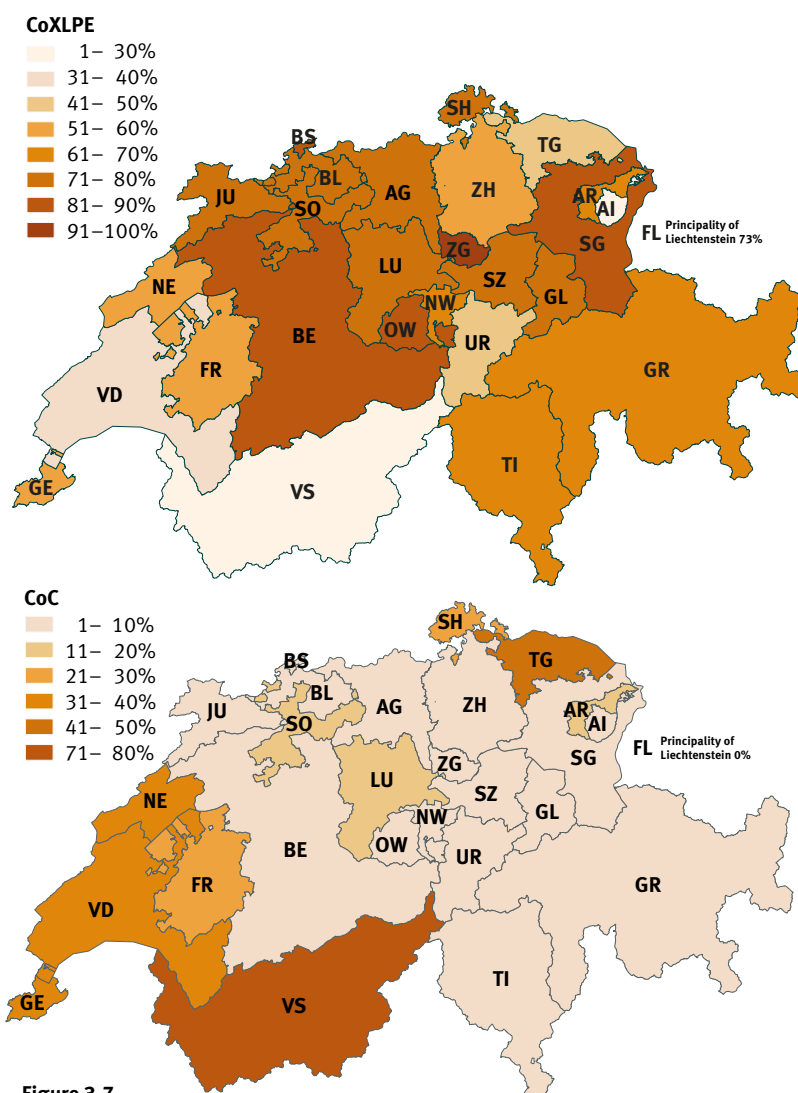


Figure 3.7
Relative share of total hip arthroplasty procedures using different bearing surfaces by canton (2019 – 2024)
Primary OA

In general head sizes of 36 and 32mm are used. Smaller head (22 and 28mm) are used in special situations with very small hips or implants. Large 36mm heads are used in more than 50% of the cases in the cantons of Vaud, Fribourg, Aargau, Thurgau and Appenzell Aussen- and Innerrhoden. In Nidwalden and Uri there is almost no use of 36mm heads – there 32mm heads are used predominantly. Central Switzerland, Bern and Zurich used 36mm head in 30–40% (Figure 3.8).

The use of DMC is also very regional (Figure 3.9). It is used more frequently in the French speaking part of Switzerland, in North-Eastern Switzerland and in the canton Schwyz. The rationale is to reduce the risk of dislocations with the use of large femoral heads and is in part triggered by the approach with posterior approaches having an increased rate of dislocations. On the other hand, there is also an influence through neighboring countries, as in France DMC are more commonly used.

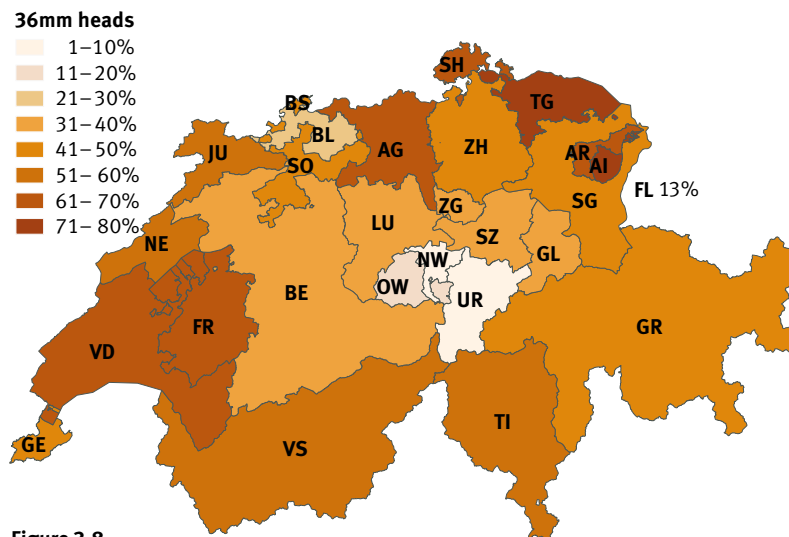


Figure 3.8
Relative share of total hip arthroplasty procedures using 36 mm heads by canton (2019 – 2024)
 Primary OA, only uncemented fixation and regular cups

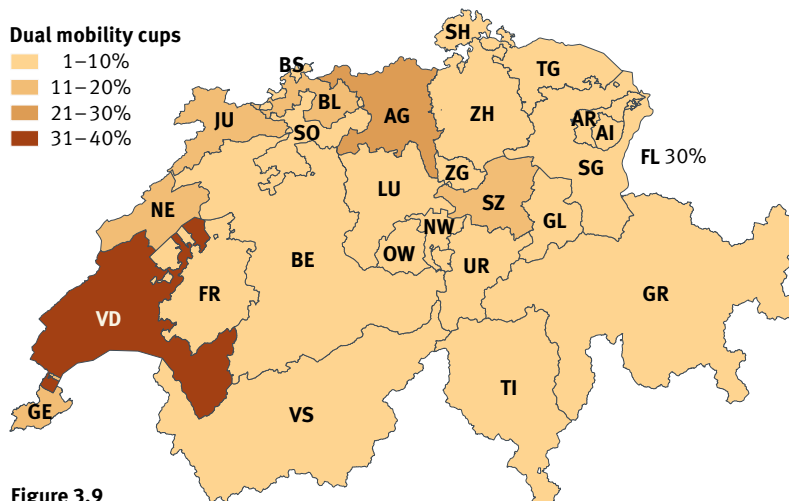


Figure 3.9
Relative share of total hip arthroplasty procedures using dual mobility cups by canton (2019 – 2024)
 Primary OA, only uncemented fixation and regular cups

3.2.3 Revision of primary THA

The current 4-year moving window from 01.01.2019 to 31.12.2022 (with 2 years follow-up) included 86,177 documented primary THA, of which 70,465 had been made for primary OA, 7,900 for secondary OA, and 7,812 for the treatment of fractures. A total of 2,395 of these had been revised within 2 years, resulting in an overall revision rate of 2.8% (CI 2.7 – 2.9%) (Table 4.9). The revision rate for primary OA was 2.4% (CI 2.3 – 2.6%), for secondary OA 4.0% (CI 3.6 – 4.5%) and for fractures 5.0% (CI 4.5 – 5.5%).

Sex and age

There was no significant difference in the 2-year revision rate among men (2.4%, CI 2.2 – 2.6%), compared to women (2.5%, CI 2.3 to 2.7%). The overall revision rate for primary OA improved by 0.1 percentage points compared to the previous 4-year period. The lowest 2-year revision rate (2.0%, CI 1.8 – 2.2%) was observed in the age group 55-64 years old. The age groups <55 and 75– 84 years of age had the highest revision rate of 2.7% (CI 2.4 – 3.1% and 2.5 – 3.0%, respectively).

BMI and ASA-score

BMI had a relevant impact on the 2-year revision rate. The highest revision rates were found in patients with a BMI >35kg/m². In these groups it was almost three times higher than in normal weight patients. The ASA score had a minor influence on the revision rate. The highest revision rate of 3.3% (CI 3.0 – 3.5%) was found in ASA class 3 patients. On the other hand, healthy patients (ASA class 1 and 2) had the lowest revision rate (1.9% CI 1.7 – 2.1% and 2.2% CI 2.0 – 2.4%).

Underling diagnosis

The revision rate differs between the various underlying diagnosis. For primary OA, the 2-year revision rate was 2.4% (CI 2.3 – 2.6%), whereas for secondary OA it was 4.0% (CI 3.6 – 4.5%) and for fractures it increased to 5.0% (CI 4.0 – 5.5%), each significantly different from the other subgroups.

3.2.4 Data by hospitals

In 2024, 146 hospital units performed primary THA, with a yearly average load of 143 THA per unit. The number of units continues to decline from 153 in 2020 to 146 in 2024. However, these numbers also may reflect purely administrative measures, including consolidation of units within hospitals or hospital groups. There is a shift towards larger units performing an increasing number of THA. In 2019, 12.1% of all THA were implanted in units performing <100 cases/year. In 2024, this share decreased to 7.8%. The same trend is observed in units performing 100–199 cases/year. On the other hand, the share of units performing >200 cases/year increased from 54.6% in 2019 to 73.6% in 2024 (Tables 3.4 and 3.5).

Since 2013, the contribution of smaller services with <200 cases/year declined from 51% to 26%. At the same time, the proportion of implantations in larger services increased from 49% to 74%. There is a clear trend for concentration in larger centres throughout the country (Figure 3.10).

Figure 3.11 illustrates the distribution of the workload among the 149 units participating in 2024, including three units performing only HA. The largest unit performed 1,362 primary THA, while nine units did <10 cases/year, two of those performing only one primary THA. Hip revision arthroplasty was performed in 131 units. However, 52 of them did <10 revisions per year (Figure 3.11).

		2019	2020	2021	2022	2023	2024
Primary total hip arthroplasty	N services	150	153	149	150	147	146
	M per service	87	94	117	122	132	143
Primary hemiarthroplasty of the hip	N services	126	125	105	110	105	101
	M per service	10	10	16	17	16	18
Revision arthroplasty (THA or HA)	N services	137	134	140	142	131	131
	M per service	10	12	12	11	12	13

Table 3.4

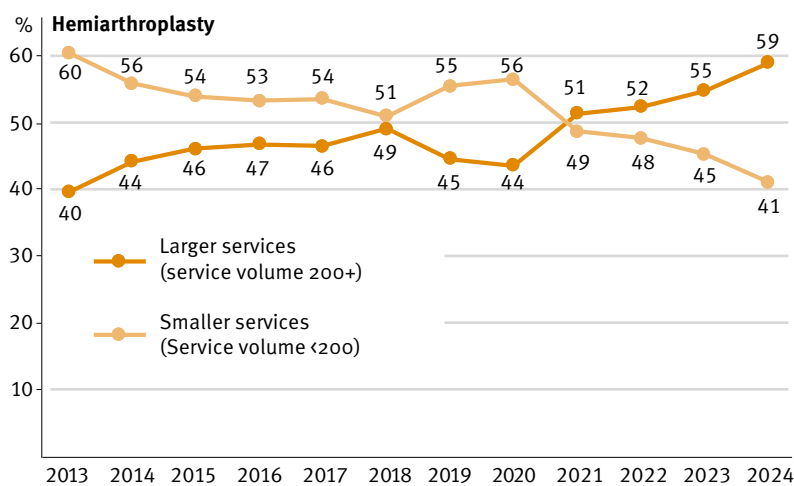
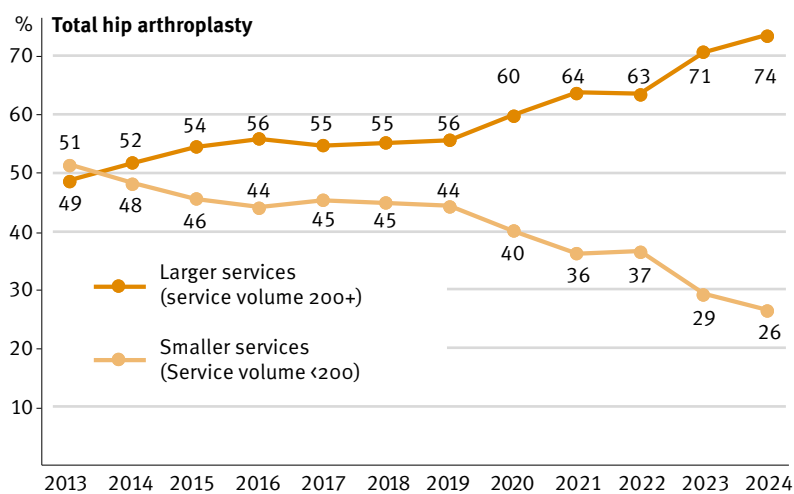
Number of participating hospital services (N) and median procedures (M) per unit per year

		2019	2020	2021	2022	2023	2024
<100	N procedures (%)	2,236 (12.1)	2,829 (14.0)	2355 (10.9)	2,431 (10.4)	2,439 (9.9)	1,996 (7.8)
	N services (%)	64 (42.7)	73 (47.7)	61 (40.9)	56 (37.3)	56 (38.1)	49 (33.6)
100–199	N procedures (%)	6,669 (33.3)	5,551 (27.5)	6,097 (27.9)	6,675 (28.6)	4,887 (19.9)	4,750 (18.7)
	N services (%)	51 (34.0)	43 (28.1)	46 (30.9)	50 (33.3)	37 (25.2)	38 (26.0)
200–299	N procedures (%)	4,424 (22.1)	4,995 (24.8)	5,185 (23.8)	5751 (24.6)	7,473 (30.5)	8,170 (32.1)
	N services (%)	20 (13.3)	22 (14.4)	24 (16.1)	26 (17.3)	34 (23.1)	34 (23.3)
>300	N procedures (%)	6,522 (32.5)	6,800 (33.7)	8,178 (37.4)	8,509 (36.4)	9,717 (39.6)	10,559 (41.5)
	N services (%)	15 (10.0)	15 (9.8)	18 (12.1)	18 (12.0)	20 (13.6)	20 (13.7)

Table 3.5

Number of hospital services and number of primary total hip arthroplasties according to hospital volume*

*Note that service volume is defined as the sum of primary procedures per year



The distinction between larger and smaller services is based on the annual volume of THA. However, it does not necessarily indicate that surgery is performed by high volume surgeons in larger services. There are larger services with numerous attending physicians, each of them contributing various number of surgeries. Some of them may perform hundreds of primary THA, whereas others contribute only few procedures. On the other hand, it may well be that in smaller units, only one or a few surgeons are performing THA, at a comparable frequency as others in large services. Therefore, caution is advised before drawing conclusions about the effect of numbers on the quality of THA.

Figure 3.10

What share of selected procedures is performed in hospital services with different service volumes*?

* Note that service volume is defined as the sum of primary procedures per year

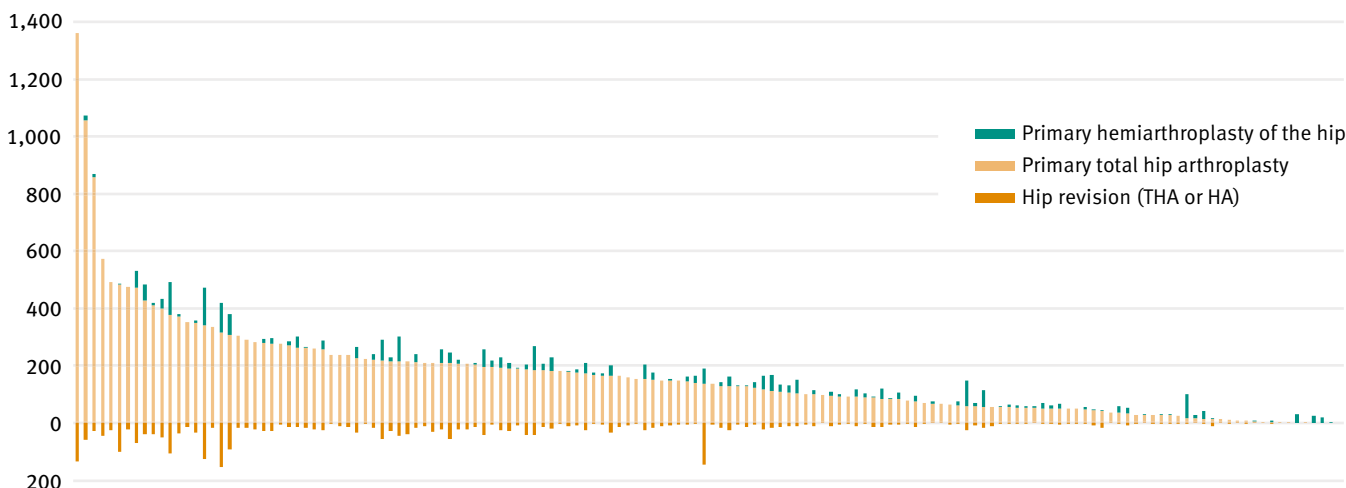


Figure 3.11

Cases per hospital service 2024: Total hip arthroplasty and hemiarthroplasty

In 2024, 14,193 THA were implanted in the 49 units performing <100 THA/year (10%), corresponding to 999 fewer cases than in 2023. Compared to 2023, an increase of 4,079 cases was observed in the 20

services performing more than 300 THA/year, corresponding to 51,378 primary THA. In these larger units, primary THA for secondary OA were performed more frequently, and the share of young patients was higher (Table 3.6).

Hospital service volume		<100	100–199	200–299	300+
N (2019–2024)		14,193	33,492	37,253	51,378
Women [%]		52.7	53.8	52.7	53.5
Mean age (SD)	All	70.4 (11.1)	69.9 (11.3)	69.5 (11.3)	68.4 (12.0)
	Women	71.9 (10.9)	71.5 (10.9)	71.0 (11.0)	70.0 (11.7)
	Men	68.8 (11.1)	68.0 (11.4)	67.8 (11.3)	66.6 (12.0)
Age group [%]	<45	1.6	2.0	2.2	3.4
	45–54	7.2	7.5	7.8	8.9
	55–64	20.3	21.1	21.8	22.9
	65–74	31.1	31.5	31.6	30.6
	75–84	30.7	29.7	29.1	27.5
	85+	9.1	8.2	7.5	6.7
Diagnosis [%]	Primary OA	79.2	80.9	83.5	80.8
	Secondary OA	7.9	7.7	7.4	12.2
	Fracture	12.9	11.5	9.1	7.0
N unknown BMI (%)		2,216 (16)	3,166 (9)	2,362 (6)	3,195 (6)
N known BMI		11,977	30,326	34,891	48,183
Mean BMI (SD)		26.8 (5.0)	27.0 (5.1)	27.1 (5.2)	26.8 (5.1)
BMI [%]	<18.5	2.4	2.3	2.0	2.2
	18.5–24.9	37.2	35.8	35.5	37.2
	25–29.9	37.6	37.4	37.9	37.6
	30–34.9	16.9	17.3	17.2	16.5
	35–39.9	4.5	5.3	5.5	4.8
	40+	1.4	1.8	1.9	1.7
	N unknown ASA (%)		335 (2)	1,382 (4)	1,049 (3)
N known ASA		13,858	32,110	36,204	49,574
Morbidity state [%]	ASA 1	10.8	9.4	10.0	11.3
	ASA 2	59.3	58.9	59.6	58.1
	ASA 3	28.6	30.5	29.4	29.5
	ASA 4/5	1.2	1.1	1.0	1.1

Table 3.6

Primary total hip arthroplasty: Baseline patient characteristics of primary total hip arthroplasty by hospital service volume*

Calculations of hospital service volume based on primary hip surgeries in each included year (2019–2024).

*Note that service volume is defined as the sum of primary procedures per year

Reporting of prostheses-related revision rates by hospitals

The national average revision rate at 2 years for THA for primary OA of 2.5% (CI 2.3 – 2.6%) served as reference for calculating the outlier status of individual hospitals, implants and surgeons. The results for all participating units are shown in funnel plots (Figure 3.12). Results were risk-adjusted for age, sex, BMI, ASA, and Charnley scores, as available, and restricted to THA performed solely for primary OA. Each dot represents a hospital/service. Confidence intervals were centred on the national average. Following convention, the 99.8% control limits were used to define the outlier status, whereas units between the 95% and 99.8% limits are defined as units with an elevated revision rate. In 2024, there were three units, one less than in 2023, that were detected as outliers and 14 institutions with an elevated revision risk, two less than 2023. Such improvements would be part of the desired effect of an arthroplasty registry.

3.2.5 Arthroplasty for fractures of the hip

Fractures of the hip may be treated by internal fixation, HA or THA. The choice of treatment option depends on the underlying pathology, functional requirements of the patient, availability within the department, as well as the experience of the treating surgeon. The patients' age, the activity level and comorbidities influence functional requirements and thus the choice of treatment option. Only arthroplasties are recorded in SIRIS. Therefore, internal fixation is not recorded. While HA is the preferred option in frail, low-demand patients, THA is more commonly performed in healthier and more active patients. SIRIS collects only data on fractures treated with THA or HA. The number of treatments with osteosynthesis is unknown. Therefore, growth rates may also reflect a change in treatment strategies between prostheses and osteosynthesis.

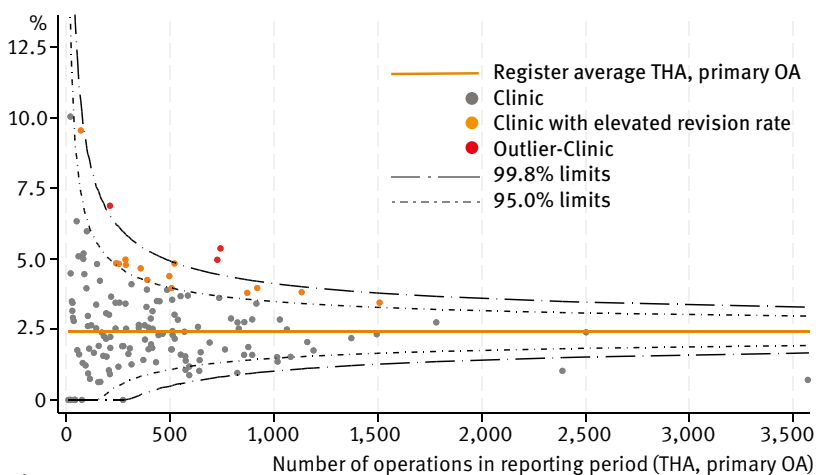


Figure 3.12
2-year revision rate of primary total hip arthroplasty by service*

THA results restricted to patients with primary osteoarthritis (prim OA).
Results are risk-adjusted for age, sex and BMI, ASA, Charnley Score if available
*Number of operations in the reporting period 01/2019–12/2022 (4-year moving average, follow-up to 12/2024)

Interpretation of funnel plots

- The coloured line denotes the Swiss average 2-year revision rate
- Clinics that lie between the 95% limits (grey dots) have revision rates that are within the statistically expected range of observations given their operation volume
- Clinics below the 95/99.8% limits are performing better than the average
- Clinics above the 95% limit and below the 99.8% limit (orange dots) have elevated 2-year revision rates. This could be due to random variation, but we recommend that possible reasons are investigated, in particular if the position should be stable over time or worsen.
- Clinics above the 99.8% limit (red dots) have 2-year revision rates that deviate markedly from the national average (unlikely to be due to random variation alone).

Between 2019 and 2024, the registry recorded a total of 27,384 primary hip arthroplasties (both THA and HA) performed for fracture, with an annual increase of 7–8% per year. During this period, there was a 31% increase in the annual number of registered arthroplasties (both THA and HA) performed for fracture of the hip, from 3,873 to 5,075. The previously observed increase of hip fracture treatment with THA continued in 2024 (Table 3.7). However, growth declined since 2023 and was only 3% for 2024 (Table 3.7). In 2024,

46.2% of the patients treated for fractures of the hip got THA and 53.8% HA. There was a clear trend over the years towards treatment with THA, proportions increasing from 41.1% in 2019 to 48.7% in 2024.

Sex and age

Women represented the majority of the patients treated for fractures, comprising 66.8% of the cases. The vast majority of patients (91.1%) was aged >65 years, while the age group >85 years accounted for 42.7% of

		2019	2020	2021	2022	2023	2024	2019–2024
N		3,873	4,149	4,495	4,865	4,27	5,075	27,384
Treatment with THA* [%]		41.1	43.1	47.5	47.1	48.3	48.7	46.2
Treatment with HA** [%]		58.9	56.9	52.5	52.9	51.7	51.3	53.8
Women [%]		69.1	67.1	67.0	66.9	65.8	65.4	66.8
Mean age (SD)	All	81.0 (10.7)	81.1 (10.7)	80.9 (10.7)	80.8 (10.7)	80.6 (10.7)	80.5 (10.7)	80.8 (10.7)
	Women	81.7 (10.1)	82.3 (10.0)	81.8 (10.3)	82.0 (10.0)	81.7 (10.1)	81.8 (9.7)	81.9 (10.0)
	Men	79.4 (11.7)	78.7 (11.7)	79.2 (11.4)	78.6 (11.7)	78.6 (11.6)	78.1 (12.1)	78.7 (11.7)
Age group [%]	<45	0.4	0.2	0.3	0.3	0.4	0.6	0.4
	45–54	1.8	1.9	1.6	1.8	1.9	1.5	1.8
	55–64	6.1	6.9	6.9	7.0	6.8	6.8	6.8
	65–74	15.3	14.7	14.7	13.9	14.9	14.5	14.7
	75–84	32.2	32.1	32.8	34.2	34.6	35.9	33.8
	85+	44.1	44.2	43.6	42.7	41.5	40.6	42.7
N unknown BMI (%)		895 (23)	781 (19)	715 (16)	609 (13)	549 (11)	395 (8)	3,944 (14)
N known BMI		2,978	3,368	3,780	4,256	4,378	4,680	23,440
Mean BMI (SD)		23.7 (4.3)	23.6 (4.4)	23.8 (4.3)	23.6 (4.2)	23.8 (4.6)	23.7 (4.3)	23.7 (4.4)
BMI [%]	<18.5	9.0	10.1	8.7	9.5	9.1	9.3	9.3
	18.5–24.9	57.4	56.7	56.7	56.5	57.3	57.4	57.0
	25–29.9	26.4	25.9	26.6	26.7	25.7	26.0	26.2
	30–34.9	5.5	5.6	6.4	6.2	5.9	5.7	5.9
	35–39.9	1.4	1.4	1.2	0.9	1.4	1.3	1.2
	40+	0.3	0.3	0.3	0.2	0.7	0.3	0.4
N unknown ASA (%)		278 (7)	247 (6)	202 (4)	135 (3)	135 (3)	84 (2)	1,081 (4)
N known ASA		3,595	3,902	4,293	4,730	4,792	4,991	26,303
Morbidity state	ASA 1	3.3	3.7	3.1	3.4	2.6	2.7	3.1
[%]	ASA 2	30.7	28.9	28.0	27.4	28.6	28.6	28.6
	ASA 3	58.4	59.9	60.2	60.4	60.1	61.0	60.1
	ASA 4/5	7.6	7.4	8.6	8.8	8.7	7.7	8.2

Table 3.7

Fracture of the hip: Baseline patient characteristics by year

*THA= Total Hip Arthroplasty. **HA= Hemiarthroplasty

the hip arthroplasties (Table 3.7). On the other hand, only 2.2% of the patients were younger than 55 years, whereas 6.8% were aged between 55 and 64 years. Patients treated with HA were on average 11 years older than those treated with THA (Table 3.8). As discussed above, younger patients were more likely to receive THA. Overall, there were more HA than THA. A total of 583 patients <55 years of age had hip arthroplasty for hip fractures and 90% of these (N = 536) were treated with THA. Interestingly, 47 of these patients <55 years of age were treated with HA. On the other hand, 62.6% of HA were performed in patients

aged 85 years and older. However, 26.8% of the patients >85 years of age received THA, the other 73.2% being treated with HA (Table 3.8).

BMI and ASA score

Most patients treated for fractures of the hip had a normal BMI. This was the case for 83% for both THA and HA. THA patients were slightly more often obese (9.2% versus 6.2%), whereas more HA patients had a BMI <18.5 kg/m², reflecting more prevalent malnutrition and sarcopenia in this subgroup (Table 3.8). The ASA score had an influence of the treatment modality. An ASA score >3 was prevalent in 81.6% of patients treated with HA but only in 52.8% of those treated with THA. That indicates that frailer patients were treated preferably with HA.

		THA	HA
N (2019–2024)		12,656	14,728
Women [%]		63.1	69.9
Mean age (SD)	All	74.9 (10.9)	85.9 (7.5)
	Women	76.1 (10.3)	86.3 (7.1)
	Men	73.0 (11.7)	84.8 (8.1)
Age group [%]	<45	0.8	0.0
	45–54	3.5	0.3
	55–64	13.3	1.1
	65–74	25.6	5.3
	75–84	37.3	30.7
	85+	19.5	62.6
N unknown BMI (%)		1,626 (13)	2,318 (16)
N known BMI		11,030	12,410
Mean BMI (SD)		24.2 (4.5)	23.3 (4.2)
BMI [%]	<18.5	7.4	11.0
	18.5–24.9	55.2	59
	25–29.9	28.3	24.4
	30–34.9	7.0	4.9
	35–39.9	1.6	1.0
	40+	0.5	0.2
	N unknown ASA		480 (4)
N known ASA		12,176	14,127
Morbidity state [%]	ASA 1	6.0	0.7
	ASA 2	41.3	17.7
	ASA 3	48.0	70.6
	ASA 4/5	4.8	11.0

Table 3.8
Fracture of the hip: Baseline patient characteristics by type of treatment

3.2.6 Data analysis by hospitals for fractures of the hip

Only 18.4% of all patients with a fracture were treated in a hospital with a volume of <100 cases/year (Table 3.9), while 30% were treated in institutions that performed between 100–199 primary arthroplasties per year. Larger units (200–299 and 300+) each treated 26% of all patients. The age distribution in the four

categories (<100 cases/year, 100–199, 200–299, >300) was comparable, with a mean age varying between 80.5 and 81.5 years. Hospitals with smaller caseloads (<100 cases/year) treated more octogenarians and it is interesting to note that the percentage of patients treated by HA in the low-volume institutions was significantly higher (63.7% vs 48.8%) than in the other units (Table 3.9). This may indicate undertreatment, whereby no more detailed analysis is possible

Hospital service volume*		<100	100–199	200–299	300+
N (2019 – 2024)		5,027	8,207	7,099	7,051
Treatment with THA [%]**		36.3	46.8	47.6	51.2
Treatment with HA [%]***		63.7	53.2	52.4	48.8
Women [%]		68.3	68.0	66.6	64.5
Mean age (SD)	All	81.5 (10.0)	80.8 (10.7)	80.7 (10.7)	80.5 (11.3)
	Women	82.2 (9.6)	81.8 (10.0)	81.8 (10.0)	81.7 (10.4)
	Men	80.0 (10.7)	78.6 (11.7)	78.6 (11.6)	78.2 (12.4)
Age group [%]	<45	0.1	0.3	0.3	0.7
	45–54	1.3	1.6	1.7	2.3
	55–64	5.7	7.0	6.8	7.3
	65–74	14.2	14.9	15.3	14.0
	75–84	33.8	34.0	33.8	33.4
	85+	44.9	42.1	42.0	42.4
N unknown BMI (%)		1,204 (24)	1,293 (16)	794 (11)	653 (9)
N known BMI		3,823	6,914	6,305	6,398
Mean BMI (SD)		23.7 (4.3)	23.7 (4.4)	23.7 (4.4)	23.7 (4.4)
BMI [%]	<18.5	8.7	9.6	8.9	9.6
	18.5–24.9	57.8	56.8	57.0	56.8
	25–29.9	26.2	26.2	26.7	25.8
	30–34.9	6.0	5.6	5.8	6.2
	35–39.9	1.0	1.4	1.2	1.3
	40+	0.3	0.4	0.4	0.3
N unknown ASA (%)		179 (4)	461 (6)	249 (4)	192 (3)
N known ASA		4,848	7,746	6,850	6,859
Morbidity state [%]	ASA 1	3.6	2.9	3.2	3.0
	ASA 2	28.3	29.7	29.1	27.3
	ASA 3	58.9	59.4	60.0	61.8
	ASA 4/5	9.2	8.0	7.8	8.0

Table 3.9

Fracture of the hip: Baseline patient characteristics by hospital service volume

Calculations of hospital service volume based on primary hip surgeries in each included year (2019–2024).

* Note that service volume is defined as the sum of primary procedures per year

** THA = Total Hip Arthroplasty

*** HA = Hemiarthroplasty

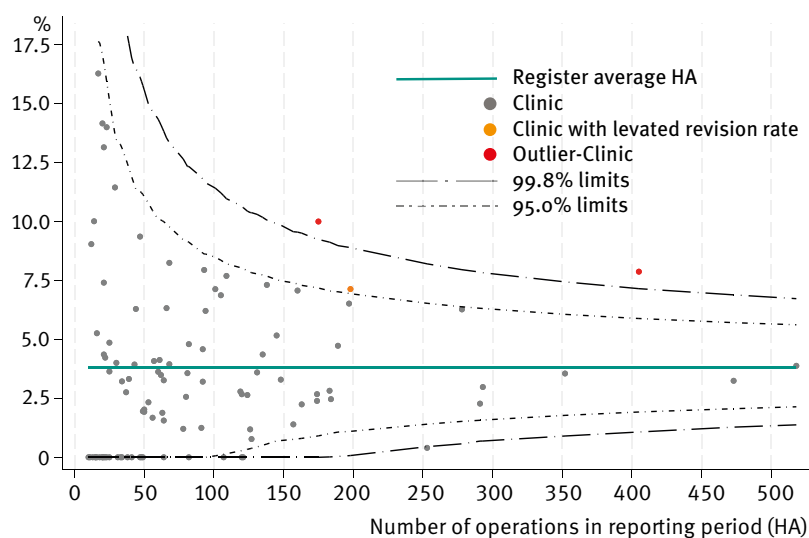
from the data available in SIRIS. One explanation may be that surgeons not trained in THA participated in the treatment of hip fractures in these smaller institutions, therefore performing HA only.

The proportion of hip arthroplasties performed in large and small units has changed significantly since 2013. Since then, the proportion of patients treated in smaller facilities has decreased from 60% to 41%, whereas the proportion of patients treated in larger facilities has risen from 40% to 59%. However, this trend was interrupted during 2019 and 2020. The reasons are not clear for 2019. In 2020 it may be a side effect of the COVID-19 pandemic, smaller hospitals having been less affected by governmental restrictions than larger providers of elective surgery.

Overall, there is a clear trend to treat fractures of the proximal femur in high volume institutions (**Figure 3.10**). The contribution of low volume centres is declining.

Reporting of revision rates of hemiarthroplasties by hospitals

The national average 2-year revision rate for HA of 3.5% (CI 3.2 – 3.7%) served as reference for calculating the outlier status of individual hospitals, implants and surgeons. The results of the participating units are shown in the funnel plot (**Figure 3.13**). For HA, there were two outlier units, one more than 2023, but only one service with an elevated revision risk, three less than 2023.



Interpretation of funnel plots

- The coloured line denotes the Swiss average 2-year revision rate
- Clinics that lie between the 95% limits (grey dots) have revision rates that are within the statistically expected range of observations given their operation volume
- Clinics below the 95/99.8% limits are performing better than the average
- Clinics above the 95% limit and below the 99.8% limit (orange dots) have elevated 2-year revision rates. This could be due to random variation, but we recommend that possible reasons are investigated, in particular if the position should be stable over time or worsen.
- Clinics above the 99.8% limit (red dots) have 2-year revision rates that deviate markedly from the national average (unlikely to be due to random variation alone).

Figure 3.13

2-year revision rate of primary hemiarthroplasty by service*

Results are risk-adjusted for age, sex and BMI, ASA, Charnley Score if available

*Number of operations in the reporting period 01/2019–12/2022 (4-year moving average, follow-up to 12/2024)

3.3 Knee arthroplasty

3.3.1 Nationwide data, including the Principality of Liechtenstein

Incidence of TKA and PKA

Since 2012, 233,166 primary knee arthroplasties were registered in SIRIS, of which 197,143 were TKA and 35,908 were PKA (Table 3.10 and 3.11). The incidence of knee arthroplasties (including both TKA and

PKA) has steadily increased over the past years, growing from 201 per 100,000 inhabitants in 2013 to 284 per 100,000 inhabitants in 2024. Considering only the population most at risk (the age group between 50 to 89 years, accounting for 98% of the recipients of TKA and PKA), the incidence of TKA and PKA together was 699 per 100,000 inhabitants at risk in 2024 (Figure 3.14). This represents one of the highest incidences for TKA in Europe as well as worldwide.

Year	Primary TKA	Primary PKA	Primary others or type unclear	Primary Total	Share of PKA	Annual growth rate primary	Linked Rev./ Reop. of TKA	Linked Rev./ Reop. of PKA	Unlinked Rev./ Reop. can be of TKA or PKA	Rev./ Reop. Total ²	Share of linked Rev./ Reop.
2012 ¹	4,653	939	8	5,600			19	2	510	531	4.0%
2013	12,656	2,402	17	15,075	15.9%		182	51	1,251	1,489	15.6%
2014	13,051	2,339	14	15,404	15.2%	2.2%	397	113	1,119	1,629	31.3%
2015	13,485	2,389	6	15,880	15.0%	3.1%	597	126	1,070	1,794	40.3%
2016	14,606	2,457	7	17,070	14.4%	7.5%	840	195	1,133	2,171	47.7%
2017	14,477	2,618	15	17,110	15.3%	0.2%	948	265	1,087	2,303	52.7%
2018	14,727	2,726	12	17,465	15.6%	2.1%	1,045	291	1,086	2,424	55.1%
2019	15,614	3,086	4	18,704	16.5%	7.1%	1,213	302	1,063	2,579	58.7%
2020	15,468	3,150	5	18,623	16.9%	-0.4%	1,331	402	1,064	2,797	62.0%
2021	16,816	3,208	4	20,028	16.0%	7.5%	1,389	406	1,028	2,826	63.5%
2022	19,316	3,431	5	22,752	15.1%	13.6%	1,567	458	926	2,955	68.5%
2023	20,706	3,438	8	24,152	14.2%	6.2%	1,672	481	1,017	3,173	67.9%
2024	21,568	3,725	10	25,303	14.7%	4.8%	1,732	508	983	3,225	69.5%
All	197,143	35,908	115	233,166	15.4%		12,932	3,600	13,337	29,896	55.3%

Table 3.10

Total and partial knee arthroplasty (TKA, PKA), primary and revisions/reoperations

All documented operations

¹ Does not represent a full year of data, as data collection in most hospitals started only in October 2012

² including linked revisions/reoperations of procedures that were classified as “primary others” or of unclear type

During the first decade of the registry, which had been introduced in 2012, the increase in the incidence could mainly be explained by improvements of coverage, as the number of participating services and the data completeness both increased over time. The high growth rate of 13.6% observed in 2022 was mostly explainable by a considerable rebound effect after the restrictions on elective surgery during the COVID-19 pandemic (Table 3.10, Figure 3.15). While the growth rate decreased from 6.2% in 2023 to 4.8%

in 2024 (Table 3.10), an unbroken and steeper increase since 2020 resulted in an increase in the incidence of both TKA and PKA by 35.3% over the last six years (Figure 3.15 and 3.10). Although indirect signs of widening of the indication for knee arthroplasty, such as increasing numbers in particularly young or old patients, could still not be detected in the SIRIS dataset (Table 3.11), the mean growth rate is clearly above the growth rate of the population at risk for TKA and PKA of 7.4% over the same period. While the

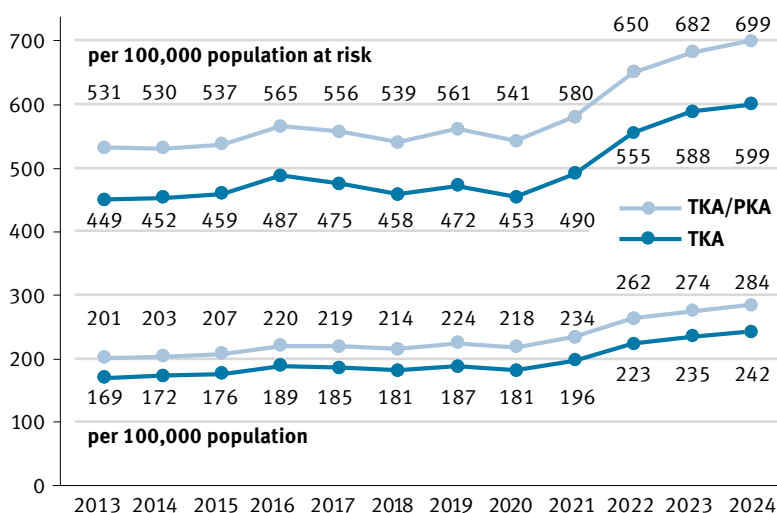


Figure 3.14

Incidence of all primary knee arthroplasties registered in SIRIS

Per 100,000 residents and per 100,000 residents at-risk (Age group 50–89 years accounts for 97% of all recipients of TKA/PKA) Adjusted for estimated coverage. SIRIS figures excluding Liechtenstein. Coverage rates 2013–2016 estimated at 92%; 2017–2023 based on federal health office data; 2024 estimated at 98.3%.

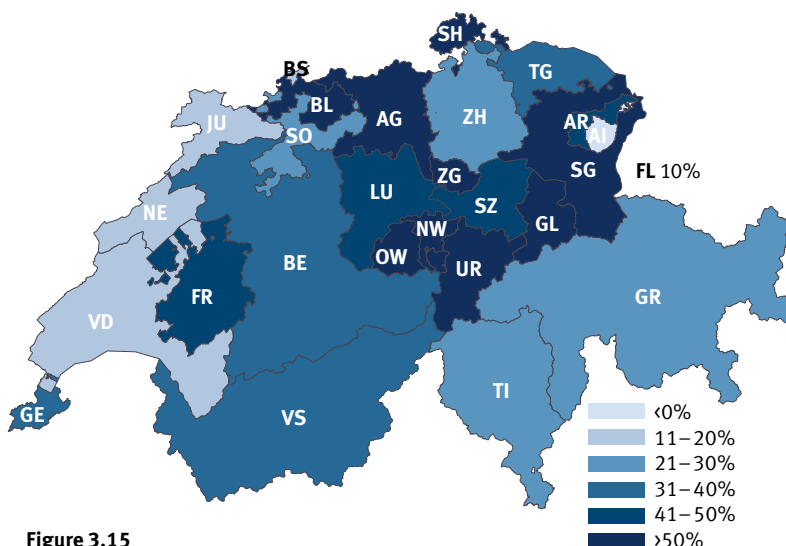


Figure 3.15

Growth rates of primary knee arthroplasties by Canton: 2019 – 2024

effect of the extensive ban of arthroscopy in elderly patients for meniscal pathology or the economic impact of lower reimbursement for arthroscopy in the ambulatory setting remain unclear, there is no doubt

that TKA and PKA are unbowed successful with high request by patients and enthusiastic proposal by surgeons and hospitals.

		2019	2020	2021	2022	2023	2024	2019–2024
N		15,614	15,469	16,816	19,316	20,706	21,568	109,489
Diagnosis [%]	Primary OA*	88.9	88.6	87.0	87.0	87.3	87.4	87.6
	Secondary OA	11.1	11.4	13.0	13.0	12.7	12.6	12.4
	Inflammatory origin	0.9	0.9	1.0	1.0	1.1	1.0	1.0
	Fracture	2.2	2.2	2.4	2.3	2.3	2.1	2.2
	Lesion of ligament	5.2	5.7	5.9	5.9	5.7	6.1	5.8
	Infection	0.2	0.2	0.2	0.2	0.2	0.2	0.2
	Osteonecrosis	1.5	1.4	1.9	1.9	1.8	1.5	1.7
	Other**	1.4	1.2	1.7	1.8	1.8	1.8	1.6
Women [%]		59.7	58.5	59.9	59.1	58.7	58.6	59.0
Mean age (SD)	All	69.8 (9.5)	69.5 (9.5)	69.6 (9.5)	69.8 (9.5)	69.9 (9.3)	69.9 (9.4)	69.8 (9.5)
	Women	70.5 (9.6)	70.1 (9.6)	70.1 (9.7)	70.4 (9.6)	70.3 (9.4)	70.3 (9.5)	70.3 (9.6)
	Men	68.9 (9.3)	68.8 (9.2)	68.8 (9.3)	69.0 (9.2)	69.4 (9.2)	69.3 (9.3)	69.0 (9.2)
Age group [%]	<45	0.4	0.5	0.5	0.5	0.4	0.4	0.5
	45–54	6.0	5.7	5.7	5.4	4.8	5.0	5.4
	55–64	23.0	24.6	24.6	24.5	24.6	25.0	24.4
	65–74	36.3	36.0	35.3	35.3	35.3	34.6	35.4
	75–84	29.3	28.9	29.6	29.8	30.6	30.6	29.9
	85+	5.0	4.2	4.4	4.6	4.3	4.4	4.5
N unknown BMI (%)		2,313 (15)	1,924 (12)	1,535 (9)	1,341 (7)	1,165 (6)	854 (4)	9,132 (8)
N known BMI		13,301	13,545	15,281	17,975	19,541	20,714	100,357
Mean BMI (SD)		29.5 (5.6)	29.2 (5.5)	29.2 (5.6)	29.2 (5.5)	29.2 (5.6)	29.3 (5.6)	29.3 (5.6)
BMI [%]	<18.5	0.6	0.6	0.6	0.5	0.5	0.4	0.5
	18.5–24.9	20.8	22.2	22.1	22.2	22.1	22.2	22.0
	25–29.9	38.8	38.2	38.0	37.8	38.2	37.5	38.0
	30–34.9	24.9	24.6	24.8	24.8	24.6	25.2	24.8
	35–39.9	10.2	10.1	9.9	10.5	10.4	10.1	10.2
	40+	4.8	4.3	4.6	4.2	4.3	4.5	4.4
N unknown ASA (%)		1,166 (7)	1,017 (7)	577 (3)	445 (2)	428 (2)	718 (3)	4,351 (4)
N known ASA		14,448	14,452	16,239	18,871	20,278	20,850	105,138
Morbidity state [%]	ASA 1	8.1	7.9	6.9	6.9	6.3	6.1	6.9
	ASA 2	61.5	62.1	61.8	63.0	63.5	62.1	62.4
	ASA 3	29.9	29.6	30.9	29.5	29.8	31.4	30.2
	ASA 4/5	0.5	0.4	0.4	0.6	0.4	0.5	0.5

Table 3.11

Primary total knee arthroplasty: Baseline patient characteristics by year

* As of SIRIS version 2021, this category includes the category “secondary arthritis after meniscus surgery”.

This category accounts for more than 6% of current entries, but shows large variability between hospitals.

** A small number of cases with “secondary OA caused by patellar instability” were added to this category.

3.3.2 Primary total knee arthroplasty

In 2024, the total number of primary TKA registered in SIRIS reached 197,143 cases (Table 3.10). Between 2019 and 2024 109,489 primary TKA were performed (Table 3.11).

Sex and age

The share of women (59.0%) and the mean age of the patients (69.8 years) remained constant between 2019 and 2024. The proportion of younger patients (0.5% younger than 45 years of age and 5.4% aged 45 to 54 years) and of patients older than 85 years (4.5%) did not change significantly in recent years (Table 3.11 and Figure 3.16). On average, women were older than men when TKA was performed in all BMI classes (Figure 3.17).

Patients with secondary OA were significantly younger (mean age 65.3 years) than those with primary OA

(mean age 70.4 years) when receiving TKA. In addition, the share of men was higher in secondary OA (52.4%) than for women (47.6%) (Table 3.12). Patients older than 65 years of age were less frequently classified as secondary OA.

Younger age was the main reason for differences in revision rates after TKA between primary and secondary OA. When comparing the corresponding age groups, revision rates did not differ between primary and secondary OA (see Chapters 5.3 and 5.4).

BMI and ASA-score

The share of unknown BMI constantly decreased from 15% in 2019 to 4% in 2024. It seems that almost all participating institutions have realised the importance of BMI for risk adjustment, as missing values lead to overestimation of revision rates. These data allow for a risk adjustment most often favourable to the provider. Between 2019 and 2024, the BMI re-

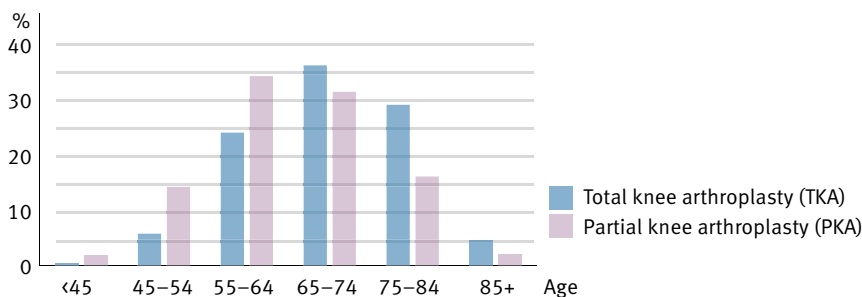


Figure 3.16

Age distribution at surgery of primary total and partial knee arthroplasty

All documented operations

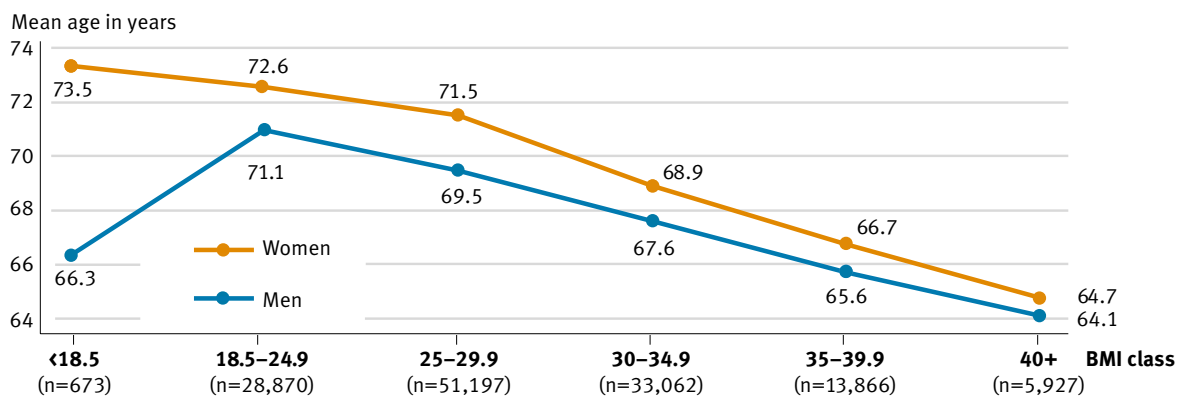


Figure 3.17

Primary total knee arthroplasty: Mean age at primary arthroplasty depending on BMI class

All documented operations. Please note that group sizes vary considerably.

mained constant at mean 29.3 kg/m². The distribution among the BMI subgroups did not change relevantly during this period (Table 3.11).

In all BMI subgroups, women were older than men at primary TKA, although the difference decreased with age and when the BMI exceeded 30 kg/m² (Figure 3.17). The difference in younger patients was mainly due to men’s higher share of post-traumatic OA. Younger patients tended to be obese more frequently. The mean age at surgery was approximate-

ly 70 years for patients with a BMI under 30 kg/m², whereas patients with a BMI of more than 40 kg/m² had surgery performed between 5 and 6 years earlier (Figure 3.11). In posttraumatic (secondary) OA, the mean BMI was significantly lower overall and in all BMI subgroups (Table 3.12).

As with the BMI, the share of unknown ASA-score decreased continuously from 7% in 2019 to 3% in 2024. This score is another important factor for risk adjustments. The distribution of the ASA-score also remained constant over the past 6 years.

BMI and ASA-score did not differ between the groups of primary and secondary OA.

Underlying diagnosis

The most frequent indication for TKA was classified as primary OA, representing 87.6% of the cases during the period from 2019 to 2024 (Table 3.11). This proportion indicates a probable issue of underreporting of secondary causes of OA. As additional causes for secondary OA (such as ligament lesions or infection) were introduced on the CRF in 2015, and as the awareness of risk factors for knee OA has steadily increased during the recent decades, a higher proportion of secondary OA would be expected.

Secondary OA had a share of 12.4% between 2019 and 2024, not varying significantly over time. Ligament lesions formed the most frequent subgroup, followed by sequels of fractures and osteonecrosis, representing 5.8%, 2.2% and 1.7% of all TKA, respectively. All other conditions were rare, the proportion of inflammatory disease as reason for TKA was 1.0%, while the category “other” represented 1.6%.

For more details regarding sex, age, BMI and ASA see above. Of note, there were considerable differences in distribution of diagnosis among high-volume hospitals (please refer to 3.3.5).

	Primary OA*	Secondary OA
N (2019–2024)	95,940	13,549
Women [%]	60.7	47.6
Mean age (SD)		
All	70.4 (9.1)	65.3 (10.5)
Women	70.7 (9.3)	66.5 (11.2)
Men	70.0 (8.9)	64.1 (9.7)
Age group [%]		
<45	0.2	2.1
45–54	4.4	12.2
55–64	22.8	35.7
65–74	36.3	28.9
75–84	31.5	18.1
85+	4.7	3.0
N unknown BMI (%)	8,124 (8)	1,008 (7)
N known BMI	87,816	12,541
Mean BMI (SD)	29.4 (5.6)	28.2 (5.1)
BMI [%]		
<18.5	0.5	0.8
18.5–24.9	21.3	27
25–29.9	37.7	40.2
30–34.9	25.1	22.7
35–39.9	10.7	6.9
40+	4.7	2.5
N unknown ASA (%)	3,884 (4)	467 (3)
N known ASA	92,056	13,082
ASA state [%]		
ASA 1	6.3	11.1
ASA 2	62.4	62.7
ASA 3	30.9	25.7
ASA 4/5	0.4	0.6

Table 3.12
Primary total knee arthroplasty: Baseline patient characteristics by main diagnostic group

* Including “arthritis after meniscus surgery”

3.3.3 Primary partial knee arthroplasty

A total of 35,908 primary PKA were registered since inception of SIRIS in 2012 (Table 3.10). The proportion of PKA among all knee arthroplasties was 15.4% over the past 13 years. It showed a peak at 16.5% in 2019, 16.9% in 2020 and 16.0% in 2021 and tended to decrease from 2022 onwards. In 2024, 21,568 TKA and 3,725 PKA were registered, resulting in a share of

PKA of 14.7%. The mean proportion over the past 6 years was 15.5%. This is clearly less than in Denmark, where PKA rates were 30.6% in 2023. It is important to note that the proportion of PKA varies greatly across Switzerland, changing considerably depending on cantons, regions, hospitals or individual surgeons. Whereas in some units PKA are rarely performed, other centres performed PKA in more than 40% of their primary knee arthroplasty cases.

Hospital service volume*		<100	100–199	200–299	300+
N (2019–2024)		2,304	4,364	4,495	8,874
Women [%]		47.5	45.4	47.1	46.4
Mean age (SD)	All	64.3 (10.3)	64.6 (10.0)	65.3 (10.1)	64.8 (10.1)
	Women	64.1 (11.1)	64.3 (10.5)	64.7 (10.4)	64.6 (10.5)
	Men	64.4 (9.6)	64.8 (9.6)	65.8 (9.8)	65.0 (9.7)
Age group [%]	<45	2.0	1.7	2.0	1.8
	45–54	15.7	14.5	12.5	14.1
	55–64	35.2	36.1	33.6	34.3
	65–74	29.9	29.4	31.9	31.5
	75–84	14.5	16.4	17.8	16.2
	85+	2.7	2.0	2.2	2.1
Diagnosis [%]	Primary OA	90.7	90.1	90.2	88.7
	Secondary OA	9.3	9.9	9.8	11.3
N unknown BMI (%)		359 (16)	340 (8)	155 (3)	870 (10)
N known BMI		1,945	4,024	4,340	8,004
Mean BMI (SD)		28.5 (4.8)	28.8 (4.9)	28.3 (4.9)	28.1 (4.7)
BMI [%]	<18.5	0.4	0.2	0.3	0.5
	18.5–24.9	22.6	22.3	25.2	27.0
	25–29.9	43.7	40.4	42.3	41.7
	30–34.9	24.1	26.3	23.5	22.8
	35–39.9	7.5	8.6	6.9	6.4
	40+	1.8	2.2	1.8	1.6
N unknown ASA (%)		116 (5)	123 (3)	179 (4)	273 (3)
N known ASA		2,188	4,241	4,316	8,601
ASA state [%]	ASA 1	15.2	14.5	10.9	13.3
	ASA 2	66.7	65.7	69.0	66.7
	ASA 3	17.7	19.7	19.6	19.8
	ASA 4/5	0.4	0.2	0.4	0.2

Table 3.13

Baseline patient characteristics of primary partial knee arthroplasty by hospital service volume

Calculations of hospital service volume based on primary knee surgeries in each included year (2019 – 2024).

* Note that service volume is defined as the sum of primary procedures per year

Hospitals with >100 knee arthroplasties per year performed 88.5% of the PKA between 2019 and 2024 (**Table 3.13**).

Sex and age

Age at surgery was lower for PKA than for TKA, with the age peak at 55 – 64 years for the former, compared to 65 – 74 years for the latter (**Table 3.14**). In 15.9% of the PKA, the patients were younger than 54 years of age, whereas the age group 75 – 84 years for accounted 16.4% and patients older than 85 years for 2.2%. Elderly patients are of special interest as surgical and general health risks can be reduced remarkably by PKA compared to TKA.

BMI and ASA-score

The BMI was not recorded in 9% of the cases between 2019 and 2024. The mean BMI of PKA patients was 28.3 kg/m², slightly lower than in TKA (29.3 kg/m²). The ASA-scores not reported in 3% of the cases registered between 2019 and 2024. Over the whole period, patients with an ASA-score of 3 and 4 represented 19.8%. This was significantly less than in TKA, where ASA-score 3 and 4 represented 30.7% of the cases (**Table 3.14**). The distribution of BMI or ASA-score did not vary relevantly over the past six years (**Table 3.14**).

Recipients of PKA were not only healthier but also significantly younger than the TKA cohort (64.8 versus 69.8 years of age) (**Table 3.11 and 3.14**).

Participants are encouraged to provide information about risk factors such as BMI and ASA-score, as it allows risk adjustments for revision rates, which are mostly favourable to the providers.

Underlying diagnosis

Between 2019 and 2024, 89.6% of the OA were classified as primary, the remaining 10.4% being secondary. In 5.3% of the latter, an osteonecrosis was registered, in 2.1% a ligament lesion and 2.0% were classified as “other reason”. Fractures were rare indications (0.8%). In 0.2%, only a PKA was performed when an inflammatory arthritis was present (**Table 3.14**). The distribution of the underlying diagnosis did not vary relevantly over the past six years (**Table 3.13**).

		2019	2020	2021	2022	2023	2024	2019-2024
N		3,086	3,149	3,208	3,431	3,438	3,725	20,037
Diagnosis [%]	Primary OA*	90.6	91.2	89.0	88.8	89.4	88.7	89.6
	Secondary OA	9.4	8.8	11.0	11.2	10.6	11.3	10.4
	Inflammatory origin	0.1	0.2	0.2	0.3	0.3	0.3	0.2
	Fracture	0.6	0.8	0.7	0.9	0.8	0.9	0.8
	Lesion of ligament	2.1	2.1	2.4	2.1	2.0	2.1	2.1
	Infection	0.0	0.0	0.0	0.1	0.0	0.1	0.0
	Osteonecrosis	5.5	4.5	5.5	5.2	5.2	5.6	5.3
	Other**	1.1	1.2	2.1	2.7	2.3	2.4	2.0
Women [%]		48.5	47.9	46.6	44.9	46.5	44.9	46.5
Mean age (SD)	All	64.7 (10.3)	64.6 (10.2)	64.5 (10.0)	65.0 (10.0)	65.0 (10.0)	65.0 (10.1)	64.8 (10.1)
	Women	64.5 (10.9)	64.2 (11.0)	64.2 (10.1)	64.7 (10.4)	64.8 (10.5)	64.4 (10.6)	64.5 (10.6)
	Men	64.8 (9.8)	64.9 (9.5)	64.7 (10.0)	65.3 (9.6)	65.2 (9.6)	65.5 (9.7)	65.1 (9.7)
Age group [%]	<45	2.1	2.3	1.9	1.5	1.7	1.6	1.9
	45–54	14.5	14.4	14.5	13.4	13.5	13.7	14.0
	55–64	34.1	34.1	34.7	35.3	34.8	34.8	34.6
	65–74	30.5	31.1	31.7	30.8	31.0	30.7	31.0
	75–84	16.4	16.0	15.2	16.8	16.8	17.0	16.4
	85+	2.3	2.2	2.0	2.2	2.2	2.2	2.2
N unknown BMI (%)		447 (14)	347 (11)	300 (9)	258 (8)	205 (6)	167 (4)	1724 (9)
N known BMI		2,639	2,802	2,908	3,173	3,233	3,558	18,313
Mean BMI (SD)		28.4 (5.0)	28.5 (4.9)	28.4 (5.0)	28.2 (4.7)	28.3 (4.9)	28.3 (4.6)	28.3 (4.8)
BMI [%]	<18.5	0.5	0.5	0.3	0.2	0.3	0.5	0.4
	18.5–24.9	25.1	24.7	25.7	26.1	24.9	24.0	25.1
	25–29.9	41.5	40.8	39.9	42.5	42.7	42.6	41.7
	30–34.9	23.0	24.8	24.1	23.0	23.4	24.9	23.9
	35–39.9	8.1	7.4	8.0	6.6	6.4	6.4	7.1
	40+	1.8	1.7	1.9	1.5	2.3	1.6	1.8
N unknown ASA (%)		167 (5)	152 (5)	59 (2)	83 (2)	82 (2)	148 (4)	691 (3)
N known ASA		2,919	2,997	3,149	3,348	3,356	3,577	19,346
Morbidity state [%]	ASA 1	16.9	14.5	14.8	12.3	11.1	10.8	13.3
	ASA 2	65.1	68.4	65.9	66.8	67.7	67.8	67.0
	ASA 3	17.8	16.8	19.0	20.7	21.0	21.0	19.5
	ASA 4/5	0.2	0.2	0.3	0.3	0.2	0.4	0.3

Table 3.14

Primary partial knee arthroplasty: Baseline patient characteristics by year

* As of SIRIS version 2021, this category includes “secondary arthritis after meniscus surgery”. This category accounts for more than 6% of current entries, but shows large variability between hospitals.

** A small number of newly added cases with “secondary OA caused by patellar instability” were added to this category.

3.3.4 Revision of primary TKA

During the current 4-year moving window from 1.1.2019 to 31.12.22 with a complete 2-year follow-up until 31.12.2024, a total of 67,217 TKA had been performed. Of those, 2,280 had been revised, resulting in a 2-year revision rate of 3.4% (CI 3.3 – 3.6%) (Table 5.3). The risk of revision within 5 years after primary TKA was 5.7%. Ten years after primary TKA, the global revision rate increased to 7.8%.

When considering revisions after five years 85.6% were first revisions, only 3.8% were revised twice. Third and fourth revisions were rare (Figure 3.18).

Taking revisions after 10 years 81.1% of these were revised once, 14.9% twice and 3.1% three times. Fourth and fifth revisions remained rare (Figure 3.18).

Sex and age

Younger patients were more at risk of early revision, with a two-year revision rate of 5.4% (CI 4.6-6.2%) for the age group <55 years old and of 4.0% (CI 3.7-4.4%) for the age group 55 – 64 years old. On the other hand, older patients (>75 years) were revised significantly less often than all the other age groups. One reason for this difference could be explained by the higher functional demands of younger patients. Ad-

ditionally, one could assume that unsatisfactory results after primary TKA were better accepted by older patients due to other compromising medical reasons and possibly higher tolerance for inferior results.

There was no statistically significant difference in the early revision rates among women (3.2%, CI 3.0 – 3.4%) and men (3.6%, CI 3.3 – 3.8%) (Table 5.3).

BMI and ASA-score

The BMI was moderately associated with short-term revision rates. Normal weight patients had the lowest early revision rate. With BMI >30 kg/m², the revision rate increased to 3.5% (CI 3.2-3.8%), while it reached 4.0% (CI 3.5-4.6%) at BMI >35 kg/m². The latter value represents a statistically significant difference from the normal weight reference group. Small numbers may have influenced the apparently increased revision rate in low BMI patients, as only 247 TKA patients had a BMI <18 kg/m² and accounted for only 11 early revisions. One revision more or less could therefore dramatically influence the outcome. Small group size is also reflected by a wide confidence interval (CI 2.6 – 8.1%).

Patients with ASA-score 4/5 tended to be revised more often, the difference however remaining statistically not significant (Table 5.3).

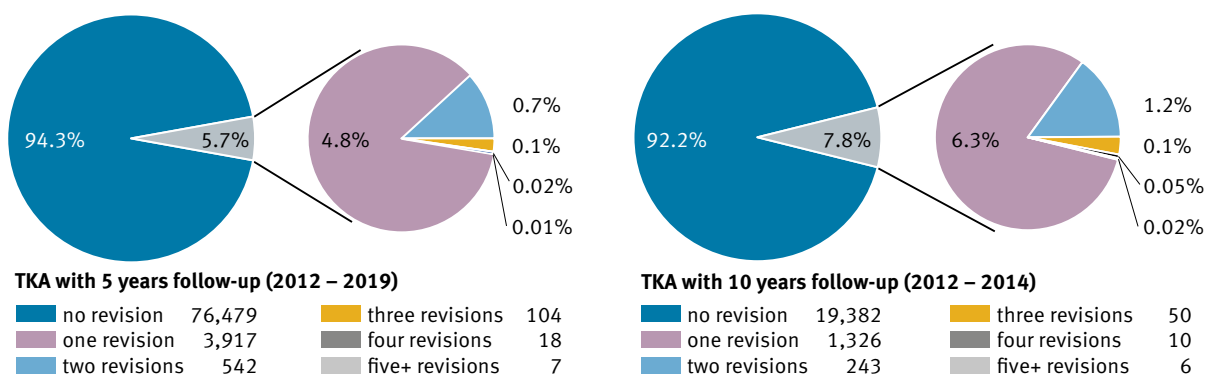


Figure 3.18
Risk of revision within 5 years and 10 years after primary TKA in percent

Multi-stage operations are counted as one revision

Underlying diagnosis

Whereas the 2-year revision rate was 3.3% (CI 3.2 – 3.5%) in primary OA, this increased significantly to 4.2% (CI 3.8 – 4.6%) in secondary OA. This is however mainly an effect of the age difference at the index arthroplasty, which was 70.4 years on average between 2019 and 2024 for primary OA, compared to 65.3 years for secondary OA (Table 3.11). Please refer also to chapter 3.3.2.

TKA revision rates over time

Comparing the revision rates of TKA over different time periods using Kaplan-Meier estimates, a trend towards continuous improvement may be observed since 2021/2022 (Figure 3.19). TKA revision rates for the period 2023/2024 had logically a follow-up limited to two years only but were significantly reduced compared to all previous periods. Such improvements over time would be one of the main goals of an implant registry.

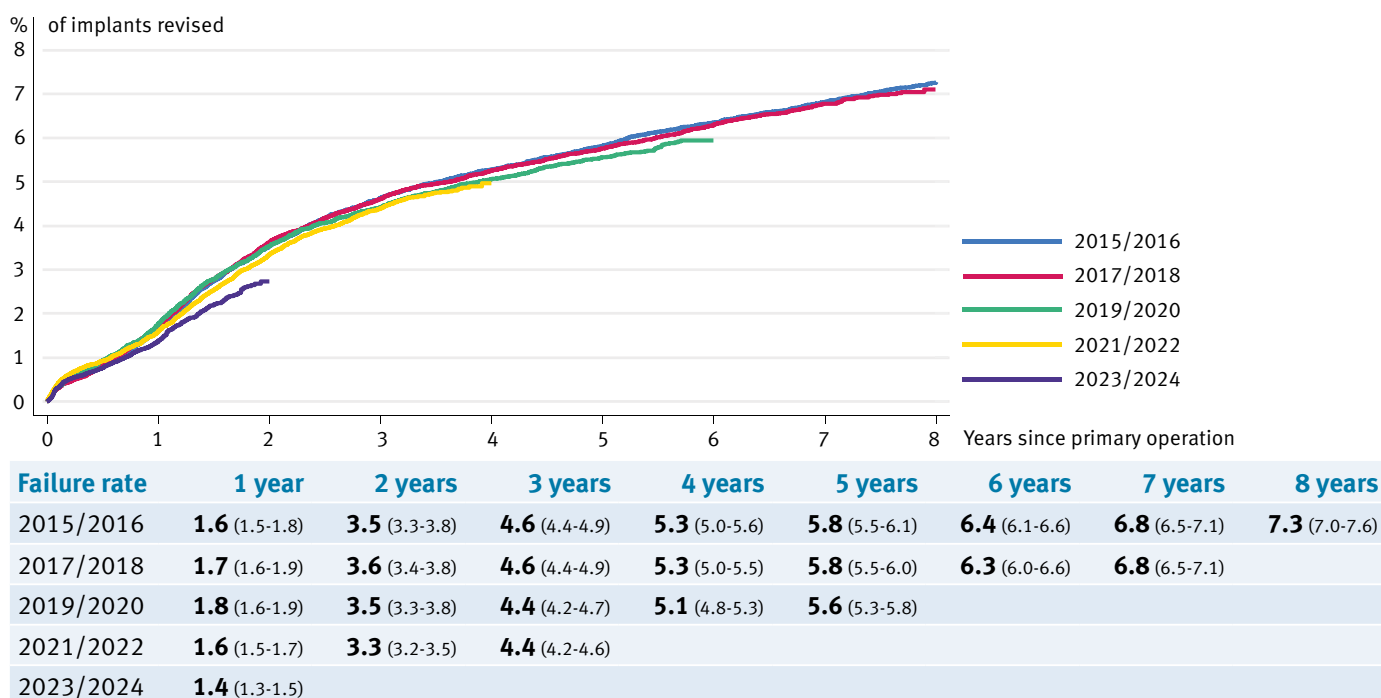


Figure 3.19
Kaplan Meier estimate of cumulative postoperative revision risk after TKA by time period

Time since operation, 2015–2024, all services, all diagnoses

3.3.5 Revision of primary PKA

Compared to previous years up to 2021, the revision rate of PKA has increased, as it did for TKA. The reason for this is likely the improved linkage rate, leading to the detection of formerly unrecognised revisions.

First revisions are revisions linked to primary PKA registered in SIRIS and occurring for the first time. Of the 35,908 PKA documented since 2012, 12,874 had been performed between 01.01.2019 and 31.12.2022 and were at risk for revision in the 4-year moving window used to calculate the current 2-year revision rate. Of the implants in this cohort, 633 were

revised, accounting for a 2-year revision rate of 5.0% (CI 4.6 – 5.3%) (Table 5.11). The risk of revision 5 years after primary PKA was 8.4%.

At revision 5 years after primary PKA 84.9% of those were revised once, 12.2% twice compared to 3.8% 5 years after primary TKA (see 3.3.4), 2% were third revisions (Figure 3.20).

There was a significant difference in survival of TKA compared to PKA, being significant from the first postoperative year onwards, as depicted in a cumulative Kaplan-Meier estimate (Figure 3.21). At 12 years, revision rate for TKA was 8.5% (8.2 – 8.7%), whereas it was 15.2% (14.4 – 16.0%) for PKA.

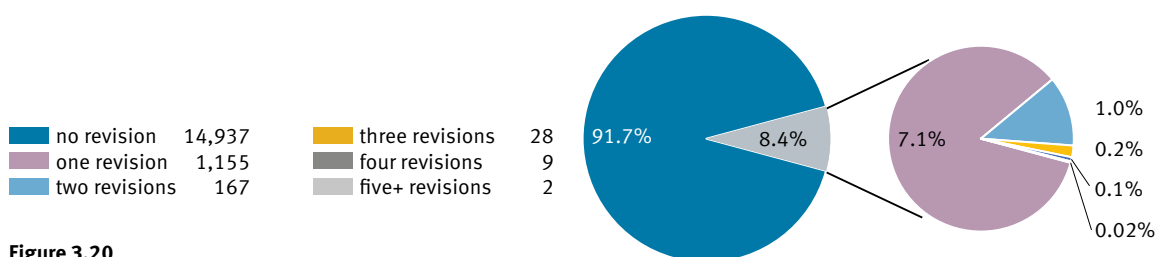
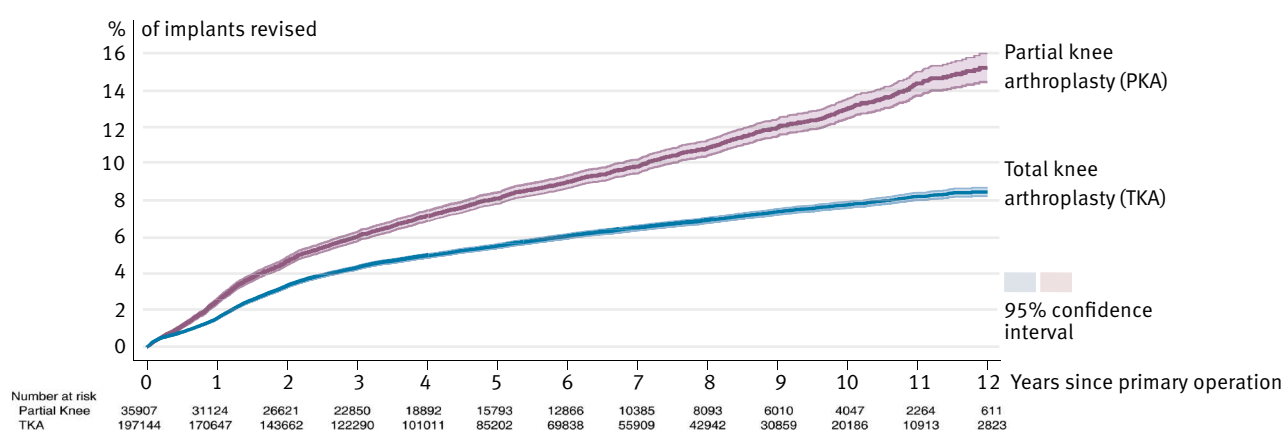


Figure 3.20
Risk of revision within 5 years after primary PKA in percent
2012 – 2019. Multi-stage operations are counted as one revision



Failure rate	1 year	2 years	3 years	5 years	6 years	8 years	9 years	10 years	12 years
TKA	1.6 (1.5-1.6)	3.3 (3.3-3.4)	4.3 (4.2-4.4)	5.5 (5.4-5.6)	6.0 (5.9-6.2)	6.9 (6.8-7.1)	7.4 (7.2-7.5)	7.8 (7.6-7.9)	8.5 (8.2-8.7)
PKA	2.5 (2.3-2.7)	4.7 (4.5-4.9)	6.0 (5.8-6.3)	8.1 (7.8-8.5)	9.0 (8.7-9.4)	10.9 (10.4-11.3)	12.0 (11.5-12.5)	13.0 (12.5-13.5)	15.2 (14.4-16.0)

Figure 3.21
Kaplan Meier estimate of cumulative postoperative revision risk after primary knee arthroplasty
Time since operation, 2012–2024, all services, all diagnoses

Sex and age

Younger patients had a significantly higher revision risk, with two-year revision rates of 7.0% (CI 5.9–8.3%) in the age group under 55 years, compared to 3.1% (CI 2.4–3.9%) in the age group 75–84 years. As in TKA, younger patients were more prone to early revision, reflecting the higher functional demands in this group and perhaps more acceptance of inferior results by older patients on the other hand due to lower activity levels or because of reluctance of any additional surgery.

The 2-year revision rates for women were 5.2% (CI 4.6–5.8%), whereas for men it was 4.4% (CI 3.9–4.9%), a difference not statistically significant (Table 5.11).

3.3.6 Two-year revision rates for TKA and PKA

Funnel plots (Figures 3.22–3.24) illustrate risk-adjusted early revision rates for primary OA, considering age, sex, BMI, ASA, and Charnley scores, as available, for TKA and PKA, as well as for revision of TKA excluding isolated patella resurfacing. Each dot represents a hospital service.

The spread of outcomes in Switzerland was relatively homogeneous, although there were exceptions, and there appears to be more variation with knee than with hip procedures. More clinics with elevated revision rates (orange dots) and outliers (red dots) could be identified for both TKA and PKA than for THA.

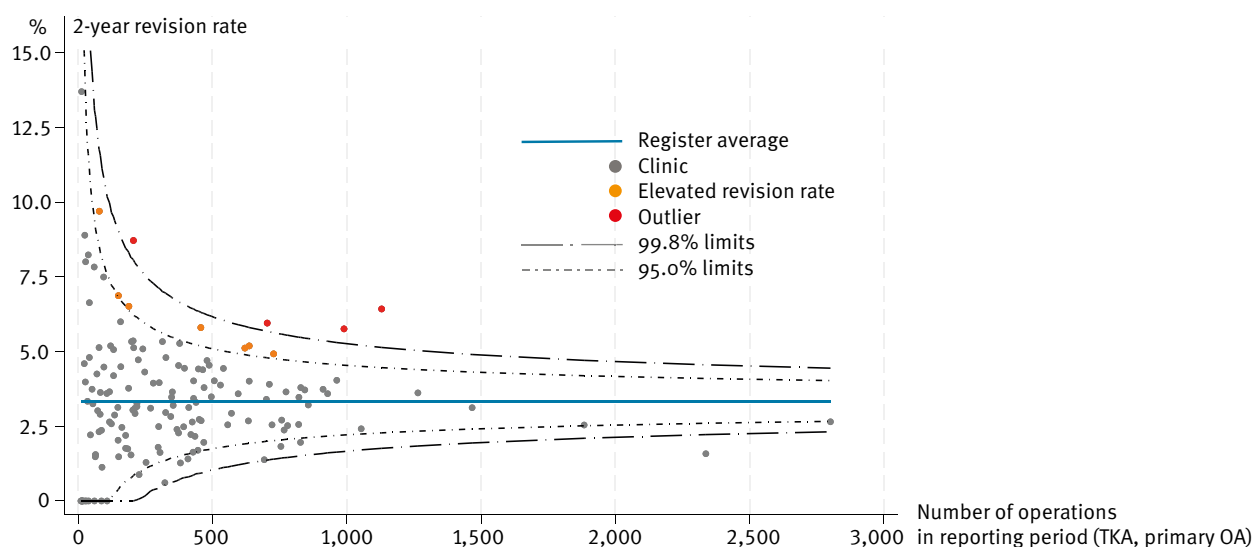


Figure 3.22

2-year revision rate of primary total knee arthroplasty by service*

TKA results restricted to patients with primary osteoarthritis (prim OA). Results are risk-adjusted for age, sex and BMI, ASA, Charnley Score if available

*Number of operations in the reporting period 01/2019–12/2022 (4-year moving average, follow-up to 12/2024)

Interpretation of funnel plots

- The coloured line denotes the Swiss average 2-year revision rate
- Clinics that lie between the 95% limits (grey dots) have revision rates that are within the statistically expected range of observations given their operation volume
- Clinics below the 95/99.8% limits are performing better than the average
- Clinics above the 95% limit and below the 99.8% limit (orange dots) have elevated 2-year revision rates. This could be due to random variation, but we recommend that possible reasons are investigated, in particular if the position should be stable over time or worsen.
- Clinics above the 99.8% limit (red dots) have 2-year revision rates that deviate markedly from the national average (unlikely to be due to random variation alone).

When isolated secondary patella resurfacing in primary TKA is excluded, the spread of results becomes less pronounced, with numbers of outliers decreasing from 4 to 3, unfortunately leaving three mid-volume hospitals (Figure 3.23). However, any addition of an implant, such as a secondary patella resurfacing or an additional PKA, counts by definition as

a revision. If a secondary patella resurfacing has to be performed within two years of the primary TKA, it cannot be considered as a good result. Considering Kaplan Meier estimation of cumulative revision risk a lower revision risk after primary TKA can be assumed in the cohort 2023/2024 in short term, though not being statistically significant (Figure 3.19).

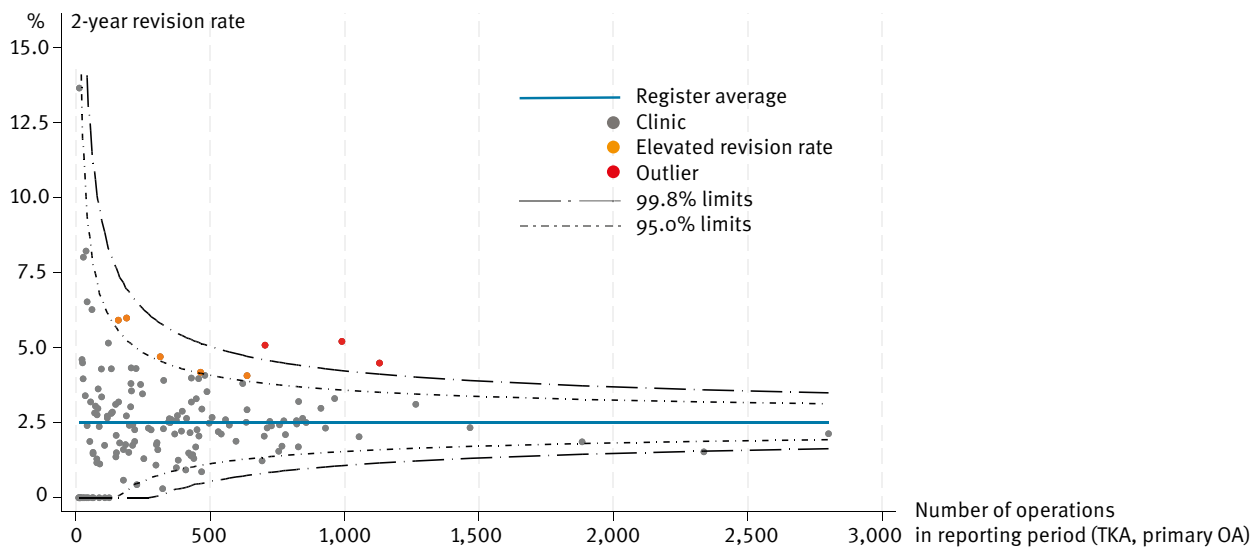


Figure 3.23

2-year revision rate of primary total knee arthroplasty by service*, without isolated secondary patella resurfacing

TKA results restricted to patients with primary osteoarthritis (prim OA). Results are risk-adjusted for age, sex and BMI, ASA, Charnley Score if available

*Number of operations in the reporting period 01/2019–12/2022 (4-year moving average, follow-up to 12/2024)

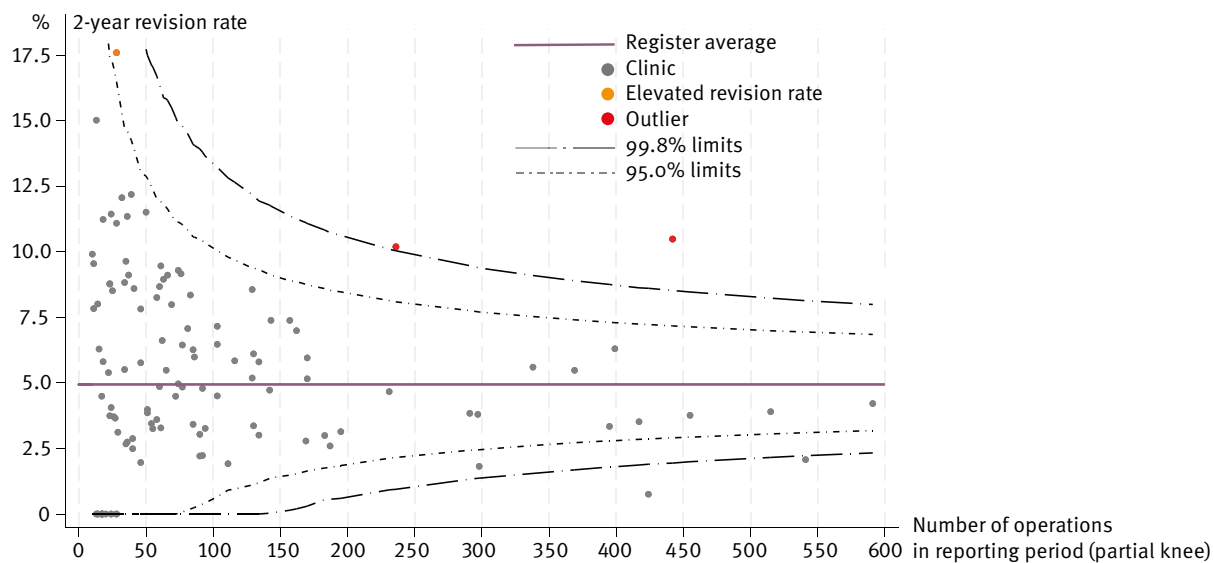


Figure 3.24

2-year revision rate of partial knee arthroplasty by service*

PKA results restricted to patients with primary osteoarthritis (prim OA). Results are risk-adjusted for age, sex and BMI, ASA, Charnley Score if available

*Number of operations in the reporting period 01/2019–12/2022 (4-year moving average, follow-up to 12/2024)

3.3.7 Regional data

As mentioned above (Section 3.3.1), the numbers of knee arthroplasties increased nationwide by 35.3% between 2019 and 2024. Growth rates differed significantly between individual cantons (Figure 3.15). Only in Appenzell Innerrhoden TKA and PKA did not increase in numbers. This is most probably due to closure of a hospital 2021, the patients having to be taken care of in another canton. It can be assumed that patients were treated in Appenzell Ausserrhoden or St. Gallen, both cantons with high growth rates. It is interesting to note that only the cantons Vaud, Neuchâtel and Jura had a growth rate less than the increase of the population at risk for TKA and PKA, followed by Solothurn, Zürich, Ticino and Graubünden, with rates ranging 21 to 30% over the past six years.

Type of knee systems

Of note is the fact that the knee replacement systems used varied significantly between individual hospitals and cantons, respectively regions. Traditionally, posterior stabilised (PS) TKA were used more commonly in the western part of Switzerland, whereas in the German-speaking cantons, cruciate retaining (CR) and cruciate sacrificing (CS), including ultra-congruent (UC) TKA, were favoured. In contrast, the implantation of medial-pivot (MP) knees did not appear to follow a particular regional pattern but seemed to be preferred in specific hospitals. Figure 3.25 illustrates the variability of the different types of TKA used in Switzerland, respectively changes between the periods spanning 2019 – 2024.

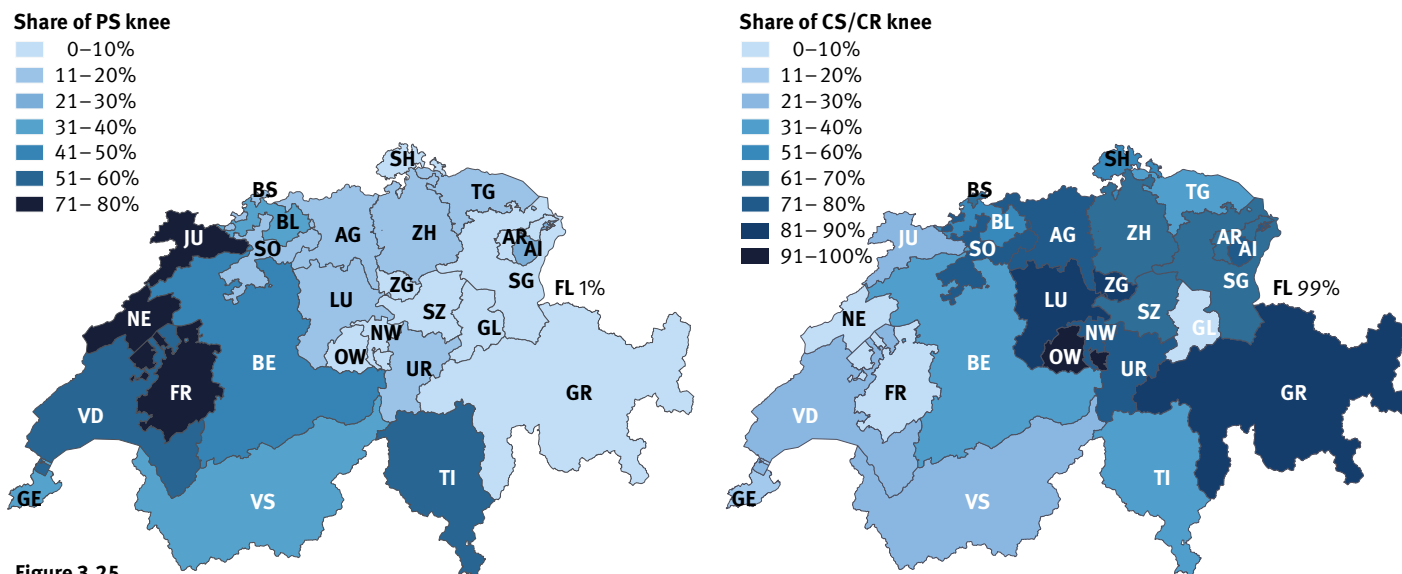
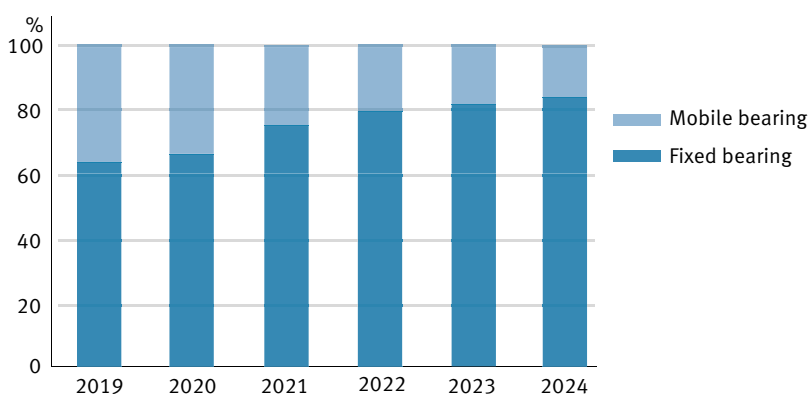


Figure 3.25
Relative share of TKA procedures using PS or CS/CR systems by Canton: 2019 – 2024

Bearing type

The proportion of mobile bearings did rapidly decrease nationwide over the past six years, from 36.5% in 2019 to 16.4% in 2024 (Figure 3.26). However, the bearing type showed again a high regional variability

(Figure 3.27). While cantons like Basel Landschaft, Solothurn, Bern, Ob- and Nidwalden, Schaffhausen, Glarus and the Principality of Liechtenstein had a low share of mobile bearings (ranging from only 0.7% in Obwalden to 8.7% in Nidwalden), this bearing type was used in 79.6% in Ticino.



	2019	2020	2021	2022	2023	2024	2019 – 2024
N	13,762	13,525	16,331	18,931	20,284	21,140	103,973
Mobile bearing	36.5	34.0	25.1	20.9	18.7	16.4	24.0
Fixed bearing	63.5	66.0	74.9	79.1	81.3	83.6	76.0

Figure 3.26

Primary total knee arthroplasty: Type of bearing

Percentage by year, all diagnoses

Share of mobile bearing knee

- 0–10%
- 11–20%
- 21–30%
- 31–40%
- 41–50%
- 51–60%
- 71–80%

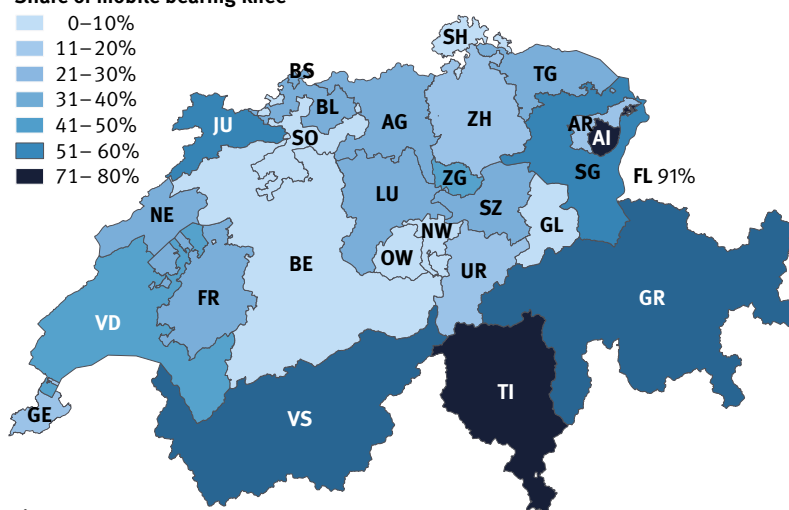


Figure 3.27

Relative share of TKA procedures using mobile bearing by Canton: 2019– 2024

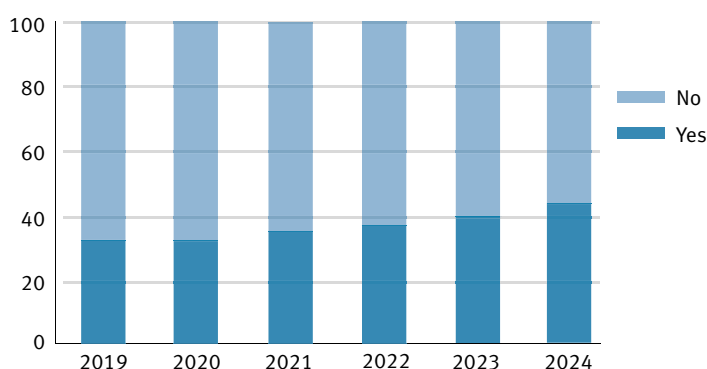
Patella resurfacing

The patella was not resurfaced in 63.1% of the primary TKA performed between 2019 and 2024 (Figure 3.28). However, the nationwide resurfacing rate increased continuously over the last years, from 32.0% in 2019 to 43.2% in 2024. As for the type of knee system or regarding the selection between fixed and mobile bearings, there were considerable differences between the cantons (Figure 3.29).

As an example, patella resurfacing was performed in 0.5% only in Glarus. On the other hand, cantons like Geneva, Vaud, Valais, Fribourg, Jura, Luzern and Thurgau resurfaced the patella between 60.9% (VS) and 70.5% (GE) of the cases. Some of these differences may be explained by the use of PS TKA, traditionally more popular in the western, French-speaking part of Switzerland, as well as in some specific centres in the German-speaking part of Switzerland. In PS TKA resurfacing of the patella is recommended more vigorously than in other designs.

Please refer to the annual report 2021 regarding more details about patella resurfacing. Particularly, it could be demonstrated that whether the patella is resurfaced primarily or not is more dependent on the surgeon's personal preference than on the knee system type or the geographic region.

The observed general trend towards more frequent primary patella resurfacing in the past years may well be explained by a surgeon's intent to prevent early revision and improve the two-year revision rates reported for himself and the hospital, independent of functional outcomes. The same effect could be observed in Australia over the past 15 years.



	2019	2020	2021	2022	2023	2024	2019–2024
N	15,607	15,467	16,814	19,315	20,706	21,568	109,477
No	67.9	68.2	65.1	63.2	60.7	56.8	63.1
Yes	32.0	31.8	34.9	36.7	39.3	43.2	36.8

Figure 3.28

Primary total knee arthroplasty: Patellar component

Percentage by year, all diagnoses

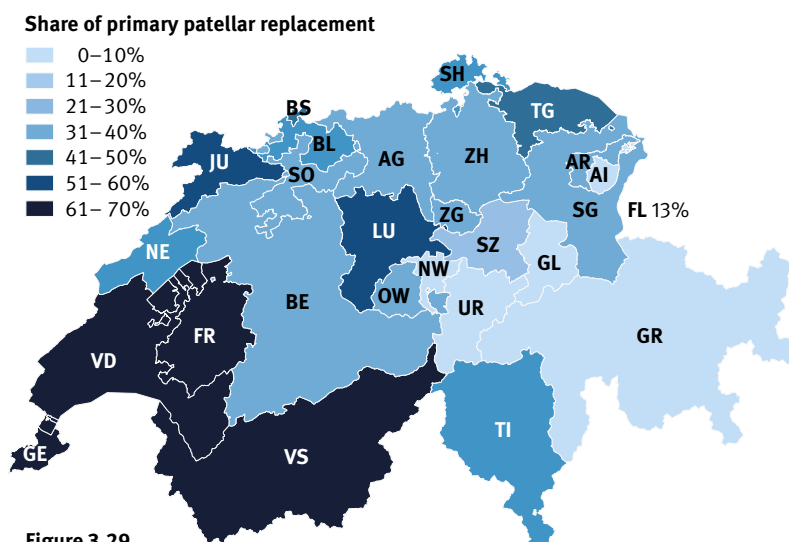


Figure 3.29

Relative share of TKA procedures using primary patellar replacement by Canton: 2019 – 2024

3.3.8 Data by hospitals

In 2024, 142 hospitals registered TKA, 122 PKA, and 134 revision arthroplasties of the knee (Tables 3.15 and 3.16). SIRIS has achieved a 100% participation rate of the hospital services since 2018 for primary TKA and PKA, as well as revision arthroplasty for relevant institutions. As SIRIS is highly dependent on data quality, such a deed merits congratulations of the providers.

Number of procedures per year

The median procedure figures per hospital (Table 3.15) remained relatively stable between 2017 and 2021 but jumped upwards considerably in 2022 and maintained that growth rate until 2024. The median annual number of primary TKA per hospital increased from 67 in 2015 to 119 in 2024, a clear increase in volume per unit. Tables 3.15 and 3.16 highlight the distribution of case numbers within service size categories (yearly numbers of procedures <100, 100 – 199, 200 – 300, >300).

		2019	2020	2021	2022	2023	2024
Primary arthroplasty of the knee (TKA)	N services	147	146	145	145	144	142
	M per service	79	77	86	106	111	119
Primary partial arthroplasty of the knee	N services	127	128	127	127	126	122
	M per service	12	12	13	14	14.5	17.5
Revision arthroplasty of the knee (TKA or partial)	N services	133	130	134	135	135	134
	M per service	9	13	12	13	16	13.5

Table 3.15

Number of participating hospital services (N) and median procedures (M) per unit per year

		2019	2020	2021	2022	2023	2024
<100	N procedures (%)	3,184 (20.5)	2,721 (17.7)	2,551 (15.4)	2,699 (14.1)	2,904 (14.2)	2,299 (10.7)
	N services (%)	81 (54.7)	78 (53.4)	72 (49.3)	65 (44.6)	66 (45.8)	57 (40.1)
100–199	N procedures (%)	4,523 (29.1)	4,698 (30.5)	4,778 (28.9)	5,551 (28.9)	5,235 (25.5)	4,285 (19.9)
	N services (%)	37 (25.0)	39 (26.7)	40 (27.4)	42 (28.9)	38 (26.3)	34 (40.0)
200–299	N procedures (%)	3,461 (22.3)	3,240 (21.0)	4,041 (24.4)	3,452 (18.0)	3,590 (17.5)	5,475 (25.4)
	N services (%)	17 (11.5)	16 (10.9)	19 (13.0)	18 (12.4)	17 (11.8)	27 (19.0)
>300	N procedures (%)	4,352 (28.0)	4,754 (30.8)	5,185 (31.3)	7,493 (39.0)	8,766 (42.8)	9,509 (44.1)
	N services (%)	12 (8.1)	13 (8.9)	14 (9.6)	20 (13.8)	23 (16.0)	24 (16.9)

Table 3.16

Number of hospital services and number of primary total knee arthroplasties according to hospital volume

Simplifying the classification into centres with less than 200 and more than 200 interventions per year shows a continuous concentration over time towards larger centres for TKA. This trend is even more pronounced and occurred earlier for PKA (Figure 3.30). High-volume services tended to perform more PKA

and revision arthroplasties than smaller units, while some centres seemed to focus on PKA and/or revision, perhaps reflecting a sort of sub-specialisation (Figure 3.31). Hospitals with more than 100 knee arthroplasties of all types per year performed 89.3% of the PKA between 2019 and 2024 (Table 3.17).

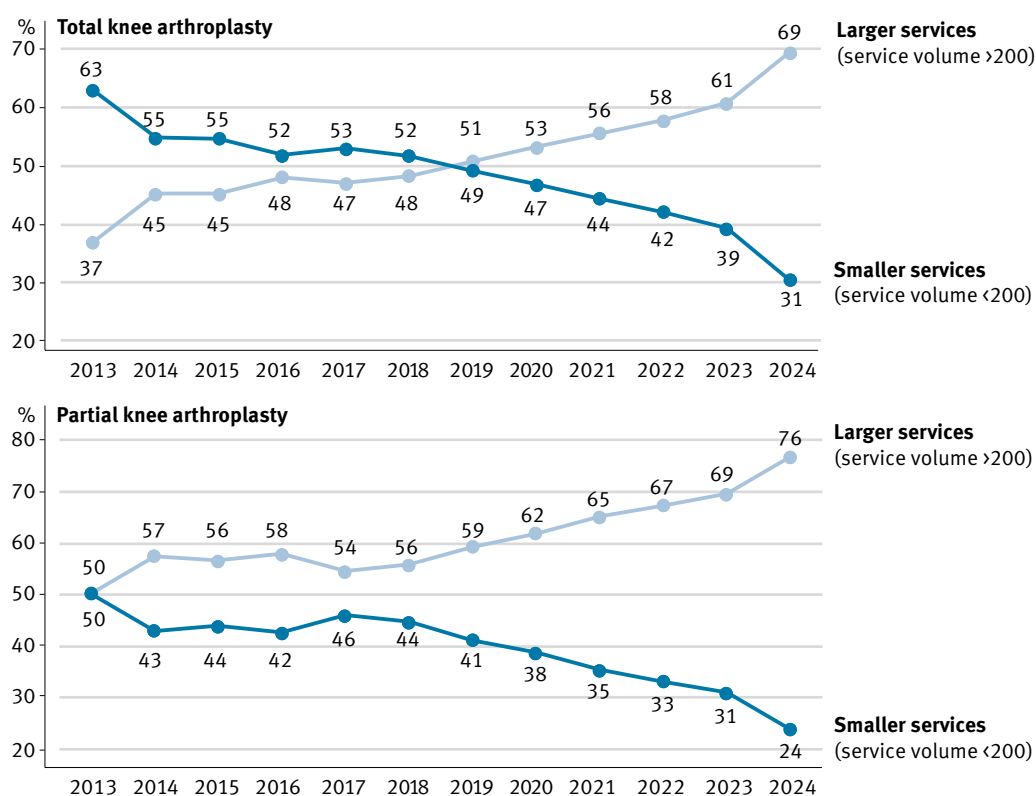


Figure 3.30
What share of selected procedures is performed in hospital services with different service volumes*?

* Note that service volume is defined as the sum of primary procedures per year

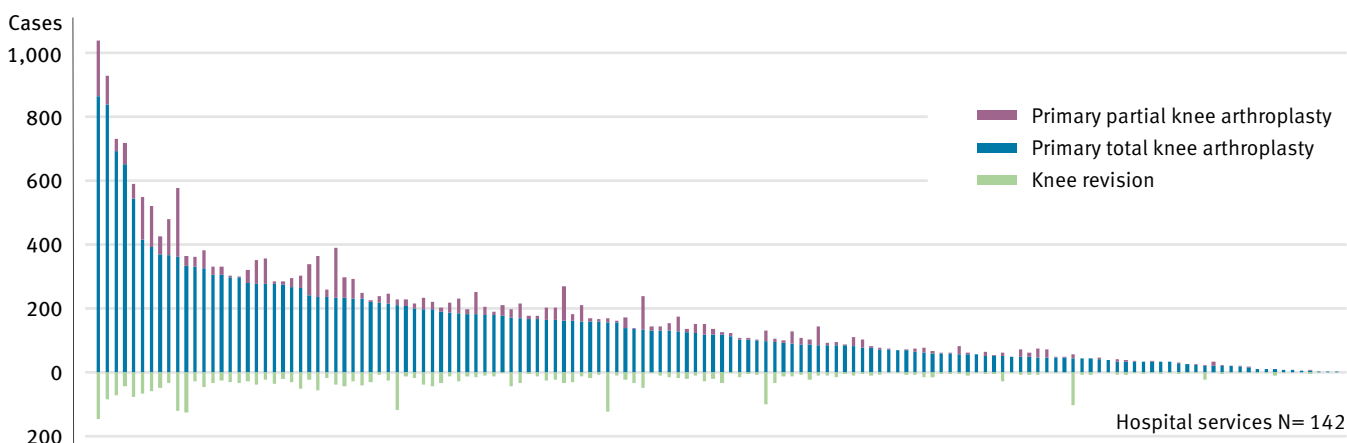


Figure 3.31
Cases per hospital service 2024: Total and partial knee arthroplasty

The distinction between large and small service units is based on the volume of knee arthroplasties performed. However, it does not necessarily indicate that surgery is performed in large services by high volume surgeons. There are services with multiple attending physicians, each of them contributing various number of cases. Some of them may perform hundreds

of implantations, others contribute with only few interventions. On the other hand, it may well be that in small units only one or few surgeons are performing arthroplasties – at a comparable frequency as others in large services. Therefore, conclusions about the effect of numbers on the quality of surgery must be considered carefully.

Hospital service volume		<100	100–199	200–299	300+
N (2019–2024)		15,765	29,472	23,355	40,897
Women [%]		59.0	58.8	60.0	58.7
Mean age (SD)	All	70.1 (9.6)	70.1 (9.4)	69.9 (9.5)	69.3 (9.4)
	Women	70.7 (9.6)	70.6 (9.5)	70.3 (9.6)	69.8 (9.5)
	Men	69.3 (9.6)	69.3 (9.1)	69.4 (9.3)	68.6 (9.1)
Age group [%]	<45	0.5	0.4	0.5	0.5
	45–54	5.5	4.9	5.2	5.7
	55–64	23.1	24.0	23.8	25.6
	65–74	34.7	35.4	35.6	35.5
	75–84	31.1	30.5	30.2	28.7
	85+	5.2	4.8	4.7	3.9
Diagnosis [%]	Primary OA	88.3	88.7	88.6	86.0
	Secondary OA	11.7	11.3	11.4	14.0
N unknown BMI (%)		2,178 (14)	1,882 (6)	1,208 (5)	3,864 (9)
N known BMI		13,587	27,590	22,147	37,033
Mean BMI (SD)		29.3 (5.6)	29.6 (5.7)	29.4 (5.6)	29.0 (5.4)
BMI [%]	<18.5	0.6	0.5	0.5	0.5
	18.5–24.9	21.9	20.6	21.4	23.4
	25–29.9	38.2	37.3	37.6	38.7
	30–34.9	24.3	25.9	25.3	23.9
	35–39.9	10.6	10.9	10.3	9.5
	40+	4.3	4.9	4.8	3.9
N unknown ASA (%)		583 (4)	820 (3)	806 (3)	2,142 (5)
N known ASA		15,182	28,652	22,549	38,755
ASA state [%]	ASA 1	6.9	6.5	5.6	8.0
	ASA 2	61.9	62.6	62.0	62.6
	ASA 3	30.5	30.4	31.8	29.1
	ASA 4/5	0.7	0.5	0.5	0.3

Table 3.17

Primary total knee arthroplasty: Baseline patient characteristics by hospital service volume

Calculations of hospital service volume based on primary hip surgeries in each included year (2019 – 2024).

Demography and type of OA

Sex, mean age, age groups, BMI, and ASA classifications did not differ among low or high-volume (<100, 100 – 199, 200 – 299, 300+ primary TKA per year) hospitals (Table 3.17).

The most frequent indication for TKA had been classified as primary OA, representing 87.6% of the cases over the period 2019 to 2024 (Table 3.11). Classification of primary and secondary OA in the 36 hospitals with more than 200 procedures per year varied remarkably. Primary OA was registered as the diagnosis leading to the operation between 49% and up to more than 91% of the cases, indicating varying practice in coding of diagnosis (Figure 3.32). In units with a high share of secondary OA, a meniscectomy was indicated most frequently as a previous operation, although this diagnosis does not significantly influence demography nor outcome after knee arthroplasty and is thus included among primary OA cases for calculations of revision rates (Figure 3.32). The

increasing share of secondary OA in some hospitals can partially be explained by the introduction of more reporting options with the 2015 version of the CFR. Rates of secondary OA due to other causes also greatly varied among the high-volume centres, similarly to proportions of post meniscectomy OA.

A technical bias towards primary OA is possible, as this diagnosis is on the top in the selection menu and thus possibly decreases the probability of selecting other alternatives, even if more appropriate. Real discrepancies in patient demographics among the 36 high-volume hospitals do not explain these considerable differences, as all other parameters were comparable. Increasing rates of secondary OA risk can influence revision rates, a benchmark established solely on arthroplasty performed for primary OA. Providers are encouraged to improve coding as proper identification of secondary OA is necessary for correct benchmarking, just as comorbidities like BMI or ASA-scores.

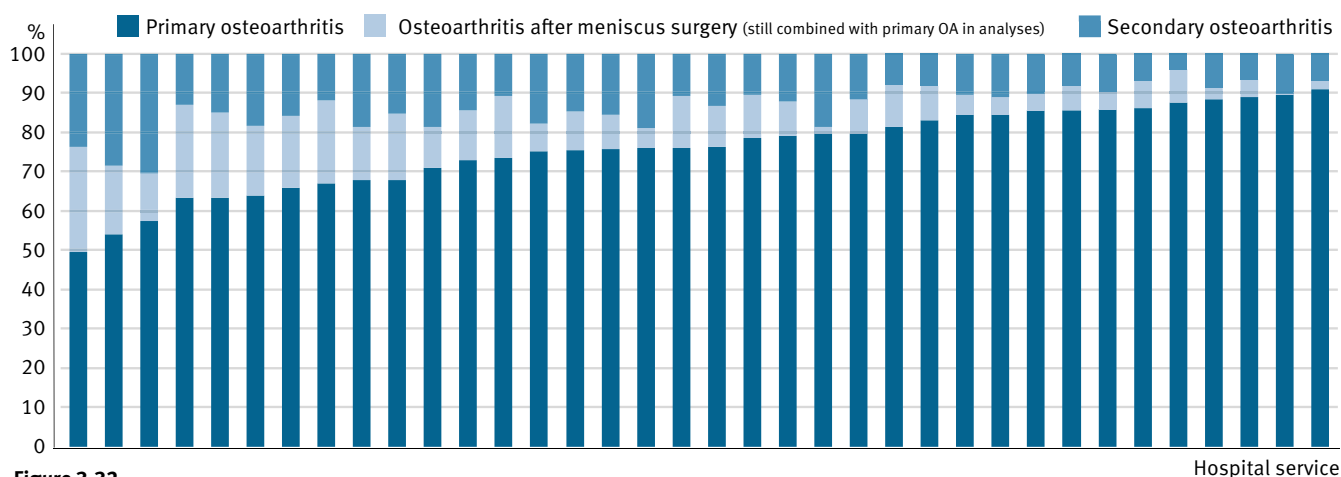


Figure 3.32
Distributions of different diagnoses in 36 hospitals >200 cases (2024)

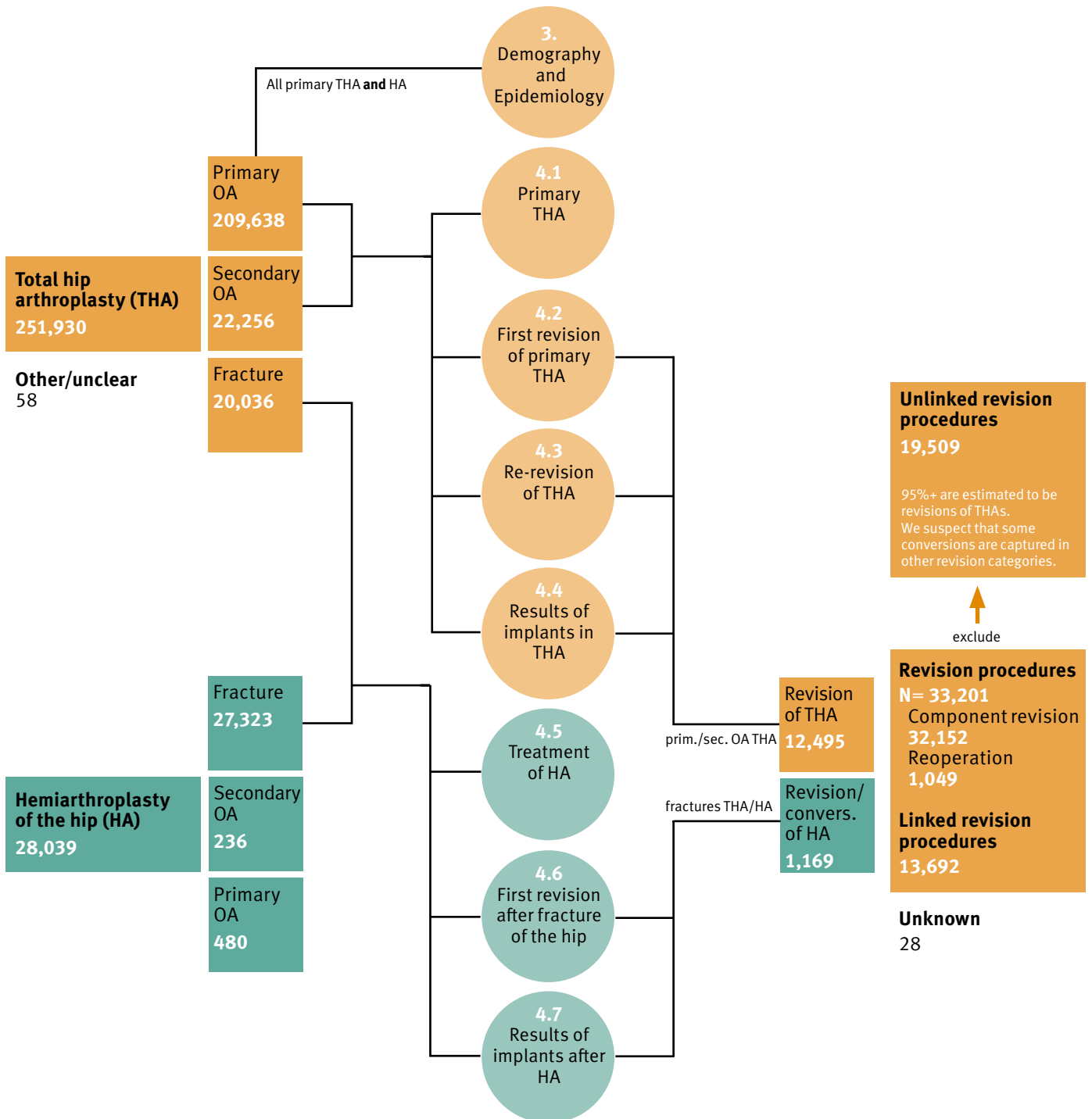
4. Hip arthroplasty

Overview of data structure

Primary procedures

Chapter in the SIRIS report

Revision/reoperation procedures



Overview of types of analyses for determining revision rates

Types of analysis	Kaplan-Meier estimates 2012 – 2024	2-year revision rates (implants 2019 – 2022 with completed 2-year follow-up)	Funnel plots of 2-year hospital revision rates (implants 2019 – 2022 with completed 2-year follow-up)
Report section	Adjusted for censoring events	Adjusted for censoring events	Risk-adjusted and adjusted for censoring events
Hip overview	All total hip arthroplasties (THA) All hemi arthroplasties (HA)		THA after primary osteo- arthritis (primary OA). ANQ online reporting, above 99.8%= outlier status All hemi arthroplasties (HA)
First revision of primary THA	THA for various subgroups	THA for various subgroups	
First revision of THA/HA after fracture of the hip	HA with bipolar versus unipolar heads THA after fracture of the hip	HA for various subgroups THA for various subgroups	
Hip implants (minimal number in group)	Uncemented stem-cup combinations, THA after primary OA (500+) Hybrid fixation stem-cup combinations, THA after primary OA (500+) Uncemented stem-cup combinations, THA after secondary OA (500+)	Uncemented stem-cup combinations, THA after primary OA (50+) Hybrid fixation stem-cup combinations, THA after primary OA (50+) Uncemented stem-cup combinations, THA after secondary OA (50+) Uncemented stem-cup combinations, THA after fracture OA (50+) Hybrid fixation stem-cup combinations, THA after fracture (50+) Cemented stem-head combinations, HA after fracture (50+)	
	Long-term evaluation 5–10 years: elevated revision rate or outlier	2-year evaluation (two times group average= outlier status)	

4.1 Primary total hip arthroplasty

Until 31.12.2024, the total number of primary THA registered in SIRIS reached 251,930 cases (Table 3.1). The current 4-year moving window includes 136,316 THA performed between 01.01.2019 and 31.12.2022, providing a complete 2-year follow-up. The share of women (53.3%) and the mean age of the patients (69.3 years) remained constant throughout the current period of observation from 2019 to 2024. Please consult Chapter 3 Demography and Epidemiology for further details regarding incidence and demographic characteristics.

Main diagnostic groups

The register categories primary THA into three groups, depending on the diagnosis leading to the operation: THA for primary OA, for secondary OA and for the treatment of fractures, as the revisions rates differ significantly between these groups (Table 4.1). The latter group, behaving relevantly differently from the other two groups, is considered more in detail in chapter 4.5 Fractures of the hip.

Primary THA for primary OA accounts for 83% of all primary THA. The share of secondary OA was 8.3% and for fractures it was 8.7%.

Main diagnostic group		Primary OA		Secondary OA		Fracture	
N (2019–2024)		N	%	N	%	N	%
Previous surgery	None	107,608	97.0	10,772	84.8	11,328	89.5
	Internal fixation femur			747	5.9	1,022	8.1
	Osteotomy femur			487	3.8	60	0.5
	Internal fixation acetabulum			95	0.7	120	0.9
	Osteotomy pelvis			310	2.4	8	0.1
	Arthrodesis			6	0.1	3	0.0
	Open impingement surgery (v2021)			69	0.5	5	0.0
	Arthroscopic impingement surgery (v2021)			81	0.6	2	0.0
	Other previous surgery	3,346	3.0	383	3.0	158	1.2
	Intervention	Total hip replacement (as entered on SIRIS form)	110,806	99.9	12,659	99.6	12,570
Hip resurfacing		21	0.0	1	0.0	0	0.0
Other (other cat. and free text entr. recog. as THA)*		127	0.1	46	0.4	86	0.7
Approach	Anterior	61,409	55.3	6,280	49.4	7,311	57.8
	Anterolateral	31,931	28.8	3,846	30.3	2,953	23.3
	Posterior	14,135	12.7	1,719	13.5	1,514	12.0
	Lateral	3,086	2.8	609	4.8	665	5.3
	Other approach	393	0.4	252	2.0	213	1.7
Fixation	All uncemented	97,004	87.4	10,313	81.2	6,317	49.9
	Hybrid (acetabulum uncemented, femur cemented)	12,469	11.2	1,707	13.4	5,289	41.8
	All cemented	877	0.8	329	2.6	645	5.1
	Reverse hybrid (acetabulum cemented, femur uncemented)	400	0.4	173	1.4	200	1.6
	Reinforcement ring, femur uncemented	75	0.1	69	0.5	54	0.4
	Reinforcement ring, femur cemented	129	0.1	115	0.9	151	1.2

Table 4.1 Part one

Primary total hip arthroplasty: Surgery characteristics by main diagnostic group

* in case of inconsistencies between form entry and implant registration, we use the implant in determining the relevant category (e.g. entered „bipolar prosthesis“ but registered stem and dual mobility cup) . Such cases are routinely counted as THAs, but still retained in the „other“ category chosen by the user.

Type of hip prosthesis

In over 99% of the cases, conventional THA had been performed in all three main diagnostic groups. Resurfacing of the hip has largely been abandoned in Switzerland, only 22 cases were registered during the past 5 years (Table 4.1).

The use of dual mobility cups (DMC) has increased over the last 10 years, particularly in the fracture group, where more than one-third of the THA performed in 2024 were made using a DMC. The use of DMC also increased in primary and secondary OA, albeit at a lower level (Figure 4.1).

Main diagnostic group		Primary OA		Secondary OA		Fracture	
N (2021–2024)		N	%	N	%	N	%
Technology	Conventional	49,616	64.7	5,569	61.7	5,294	57.5
	Computer assisted cup	1,450	1.9	88	1.0	36	0.4
	Computer assisted stem	1,651	2.2	96	1.1	20	0.2
	Robotic assisted (image guided, CT based)	654	0.9	83	0.9	12	0.1
	Patient specific cutting blocks	851	1.1	39	0.4	5	0.1
	Intraoperative fluoroscopy/radiography	25,075	32.7	3,301	36.5	3,877	42.1
	Total THA (multiple responses)	76,702		9,033		9,210	
Add. intervention	None	74,579	96.4	8,031	87.9	8,021	86.5
	Acetabular roof reconstruction	890	1.2	257	2.8	105	1.1
	Central osseous reconstruction	598	0.8	202	2.2	178	1.9
	Proximal femur osteotomy	12	0.0	28	0.3	26	0.3
	ORIF/CRIF acetabulum	35	0.0	25	0.3	157	1.7
	Cerclage femur	548	0.7	236	2.6	491	5.3
	ORIF/CRIF femur	47	0.1	31	0.3	119	1.3
	Augments	15	0.0	15	0.2	9	0.1
	Other	796	1.0	481	5.3	453	4.9
	Total THA (multiple responses)	77,341		9,138		9,274	

Table 4.1 Part two

Primary total hip arthroplasty: Surgery characteristics by main diagnostic group

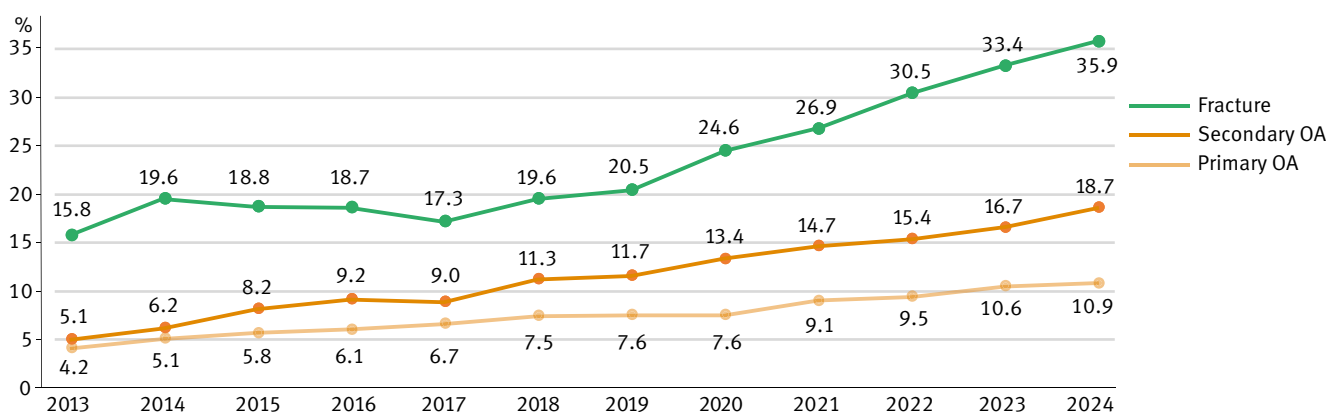


Figure 4.1

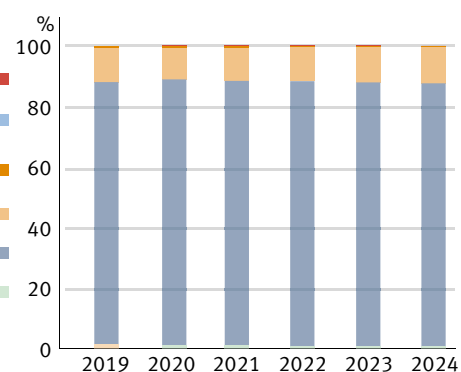
Share of dual-mobility cups over time by main pathology (%)

Fixation of the stem differed significantly among the main diagnostic groups but did not change significantly since 2019 (Table 4.2). Uncemented stems were used in more than 80% of primary and secondary OA, whereas approximately 50% of the stems had cemented fixation when treating fractures. The use of

all uncemented fixation declined slightly since 2022. Cages and reinforcement rings were used noticeably more frequently in the secondary OA and the fracture groups. This is most probably due to more complex deformities or fractures of the acetabulum needing advanced reconstruction techniques.

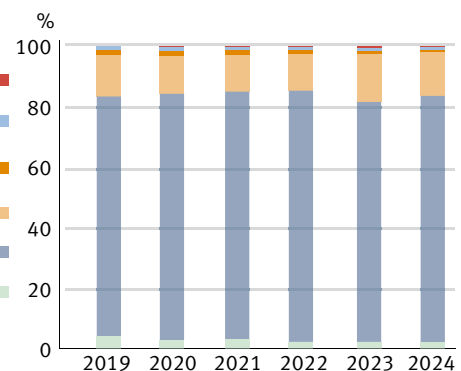
Primary osteoarthritis

2019	2020	2021	2022	2023	2024	
0.1	0.1	0.1	0.1	0.1	0.0	Reinforcement ring femur uncem.
0.1	0.2	0.2	0.1	0.1	0.0	Reinforcement ring femur cemented
0.4	0.4	0.5	0.3	0.3	0.3	Reverse hybrid
11.3	10.4	10.8	11.2	11.6	11.9	Hybrid
86.7	88.0	87.6	87.7	87.4	87.2	All uncemented
1.3	0.9	0.9	0.7	0.6	0.5	All cemented
16,892	16,721	17,907	18,946	19,973	20,515	N



Secondary osteoarthritis

2019	2020	2021	2022	2023	2024	
0.3	0.4	0.5	0.6	0.8	0.5	Reinforcement ring femur uncem.
1.0	1.2	0.8	0.8	0.9	0.8	Reinforcement ring femur cemented
1.6	1.7	1.6	1.3	1.2	0.9	Reverse hybrid
13.8	12.4	12.1	12.1	15.6	14.3	Hybrid
79.4	81.4	81.9	83.1	79.4	81.6	All uncemented
3.9	2.8	3.0	2.2	2.2	1.9	All cemented
1,716	1,852	2,076	2,256	2,316	2,490	N



Fracture

2019	2020	2021	2022	2023	2024	
0.4	0.4	0.4	0.5	0.5	0.3	Reinforcement ring femur uncem.
1.3	1.2	1.2	1.0	1.6	0.9	Reinforcement ring femur cemented
2.1	1.9	1.9	1.5	1.1	1.2	Reverse hybrid
41.2	39.4	39.7	39.8	43.9	45.6	Hybrid
47.6	52.2	51.2	52.3	48.6	47.7	All uncemented
7.3	4.9	5.6	4.9	4.4	4.2	All cemented
1,593	1,789	2,134	2,290	2,380	2,470	N

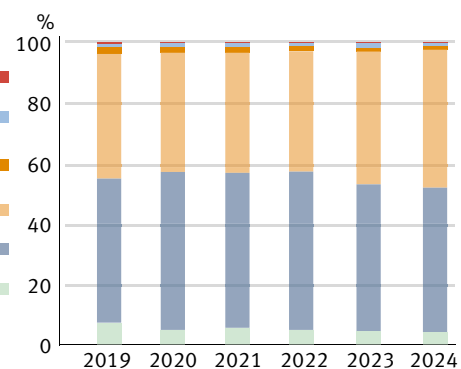


Table 4.2

Primary total hip arthroplasty: Component fixation methods by diagnostic group by year

Approach

For primary OA, the anterior approach was used most often, followed by the anterolateral approach. Approaches first started to be recorded in 2015. Since then, the use of the anterior approach has gradually increased, reaching 58.6% in 2024. The anterolateral approach was quite popular in 2019, being used in 31.2% of the primary THA, but its use declined to 27.6% in 2024. The lateral approach is getting less and less popular. The share of the posterior approach is also declining; it was used in 11.9% of the cases in 2024 (Table 4.3).

The anterior approach is most frequently used in northeastern Switzerland, as well as in the Cantons of Lucerne, Wallis and Geneva. On the other hand, the anterolateral approach is used most often in the Cantons of Bern, Jura, Solothurn, Uri, Nidwalden and Graubünden (Figure 4.2).

Surgical approach	2019	2020	2021	2022	2023	2024	2019–2024
Anterior	50.5	52.8	55.8	56.9	56.4	58.6	55.3
Anterolateral	31.2	30.6	27.7	27.8	28.4	27.6	28.8
Lateral	4.6	3.6	2.9	2.5	1.9	1.6	2.8
Posterior	13.0	12.5	13.3	12.6	13.1	11.9	12.7
Other approach	0.7	0.5	0.3	0.2	0.2	0.2	0.4
Total [N]	16,892	16,721	17,907	18,946	19,973	20,515	110,954

Table 4.3

Surgical approach in total hip arthroplasty for primary osteoarthritis by year (in %)

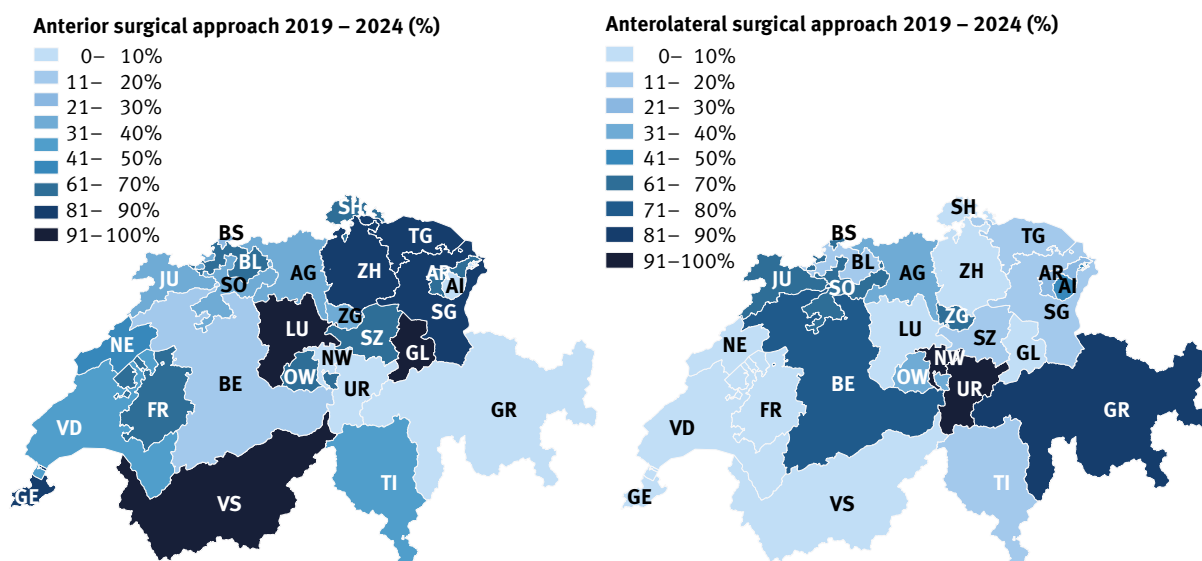


Figure 4.2

Relative share of total hip arthroplasty procedures using different surgical approaches by Canton (2019 – 2024)

Bearing

The bearing is one of the most important key factors for wear resistance and consequently long-term implant survival. Improvement of bearing materials has led to a decrease in osteolysis and loosening, with decreasing necessity for consecutive revisions. The selection of the bearing depends, amongst other criteria, on the activity level and the age of the patient.

Bearings with favourable wear characteristics, e.g., ceramic on highly crosslinked polyethylene (CoXLPE) and ceramic on ceramic (CoC), were most frequently used in younger patients. Currently, the most frequently used bearing in Switzerland was CoXLPE and its use continued to increase. In 2024, this combination was chosen in 69.4% of all primary THA for primary OA (Table 4.4 and Figure 4.3). Metal on conven-

Bearing surface	2019	2020	2021	2022	2023	2024	2019–2024
Metal on conventional polyethylene (MoCPE)	1.8	1.5	2.1	1.3	1.6	1.4	1.6
Ceramic on conventional polyethylene (CoCPE)	4.2	3.9	4.5	4.7	5.6	5.4	4.8
Metal on cross-linked polyethylene (MoXLPE)	11.3	9.6	9.0	7.8	6.7	7.0	8.5
Ceramic on cross-linked polyethylene (CoXLPE)	60.3	62.6	63.9	66.7	67.7	69.4	65.3
Ceramic on ceramic (CoC)	15.2	15.0	14.0	12.9	12.1	10.4	13.1
Ceramicised metal on conventional polyethylene (CMoCPE)	0.4	0.3	0.3	0.3	0.3	0.3	0.3
Ceramicised metal on cross-linked polyethylene (CMoXLPE)	6.8	7.1	6.2	6.4	6.2	6.2	6.5
N (bearing surface known)	16,515	16,455	17,655	18,664	19,524	20,032	108,845
N (bearing surface unknown)	377	266	252	282	449	483	2,109

Table 4.4

Primary total hip arthroplasty: bearing surface* in primary osteoarthritis by year (in %)

* Femoral heads and acetabular inserts/monobloc cups

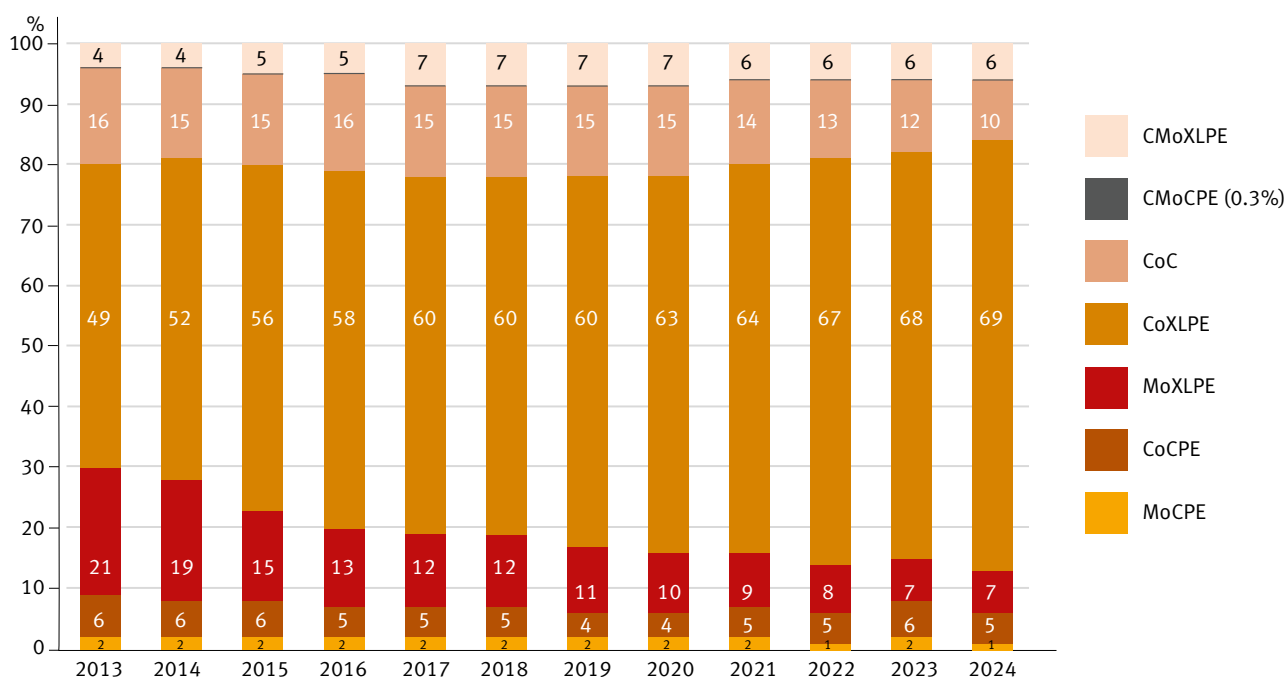


Figure 4.3

Relative shares of bearing surfaces in total hip arthroplasty over time

Primary OA

tional polyethylene (MoCPE) continued to have a very low share, essentially unchanged around 1.6% over the years. Ceramised metal (CM) was used in 6.8% of the cases, mostly in combination with XLPE. The second most frequently used bearing was CoC, accounting overall for 13.1% of all THA for primary OA. However, its use slowly declined over the years and

was used in only 10.4% in 2024. It was used most often in patients <45 years of age, where it was chosen in 24.1% (Table 4.5). XLPE is the most commonly used acetabular bearing in the age group above 75 years of age. However, it is more often combined with metal heads, while in the younger patients CoXLPE or CoC are prevalent. The age-related use of the various bearings is shown in Figure 4.4.

Bearing surface	<45	45-54	55-64	65-74	75-84	85+	All
Metal on conventional polyethylene (MoCPE)	0.1	0.2	0.4	0.8	2.8	7.2	1.6
Ceramic on conventional polyethylene (CoCPE)	1.2	1.4	2.2	4.2	7.4	10.4	4.8
Metal on cross-linked polyethylene (MoXLPE)	3.4	4.9	5.5	7.7	11.1	17.0	8.5
Ceramic on cross-linked polyethylene (CoXLPE)	64.9	65.4	68.2	67.3	63.3	53.4	65.3
Ceramic on ceramic (CoC)	24.1	20.8	16.4	13.0	9.3	7.0	13.1
Ceramicised metal on conventional polyethylene (CMoCPE)	0.00	0.04	0.09	0.16	0.53	0.89	0.29
Ceramicised metal on cross-linked polyethylene (CMoXLPE)	6.37	7.21	7.26	6.91	5.62	4.05	6.46
N (bearing surface known)	1,901	8,435	24,856	35,795	31,220	6,637	108,844
N (bearing surface unknown)**	46	128	466	637	656	176	2,109

Table 4.5

Primary total hip arthroplasty: bearing surface* in primary osteoarthritis by age (in %)**

* Femoral heads and acetabular inserts/monobloc cups

** please note that age is missing in one case

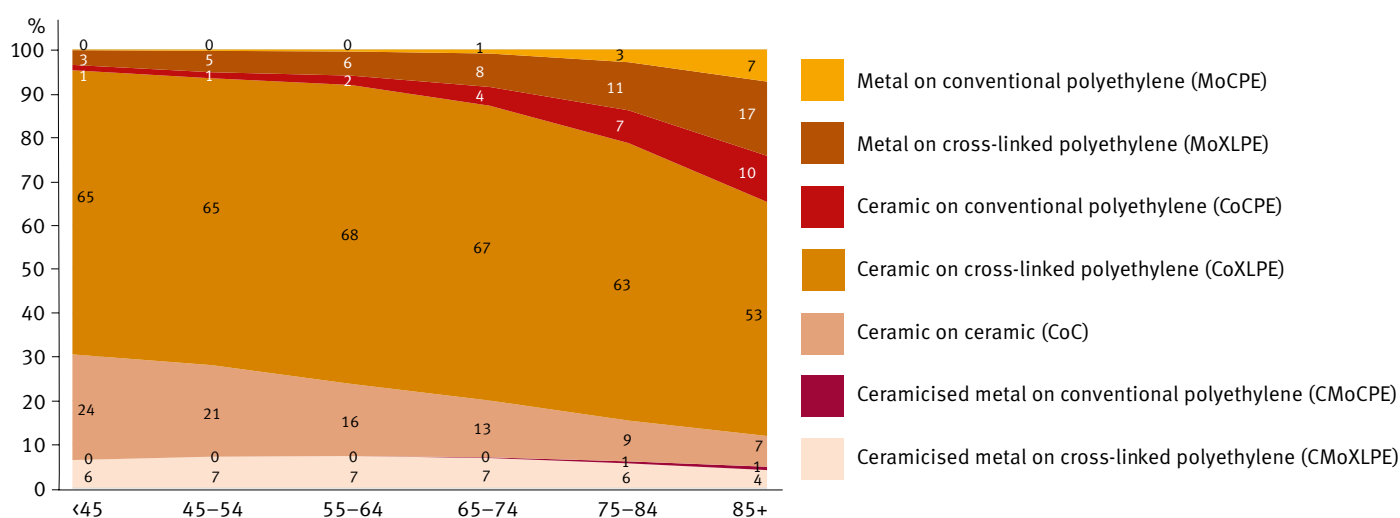


Figure 4.4

Primary total hip arthroplasty: bearing surface in primary osteoarthritis by age (in %)

Primary OA, 2019 – 2024

Fixation

Uncemented fixation of both components was standard for primary THA performed for primary OA, accounting for 87.4% of all the cases and more than 95% of patients under the age of 65. Patients older than 85 years still received cementless implants in

59% of the cases (Table 4.6). Hybrid fixation with cemented stems and uncemented cups was chosen more frequently in females (15.6%) than in males (6.6%) (Table 4.7).

Fixation method	<45	45–54	55–64	65–74	75–84	85+	All
All cemented	0.3	0.2	0.2	0.5	1.2	3.5	0.8
All uncemented	97.6	97.5	96.4	91.3	78.6	59.0	87.4
Hybrid acetabulum uncemented, femur cemented	1.3	1.8	3.0	7.8	19.5	36.1	11.2
Reverse hybrid acetabulum cemented, femur uncemented	0.7	0.3	0.2	0.3	0.5	0.9	0.4
Reinforcement ring, femur cemented	0.00	0.07	0.08	0.07	0.16	0.40	0.12
Reinforcement ring, femur uncemented	0.2	0.2	0.1	0.1	0.0	0.1	0.1
N	1,947	8,563	25,322	36,432	31,876	6,813	110,953

Table 4.6

Primary total hip arthroplasty: fixation methods in primary osteoarthritis by age* (in %)

* please note that age is missing in one case

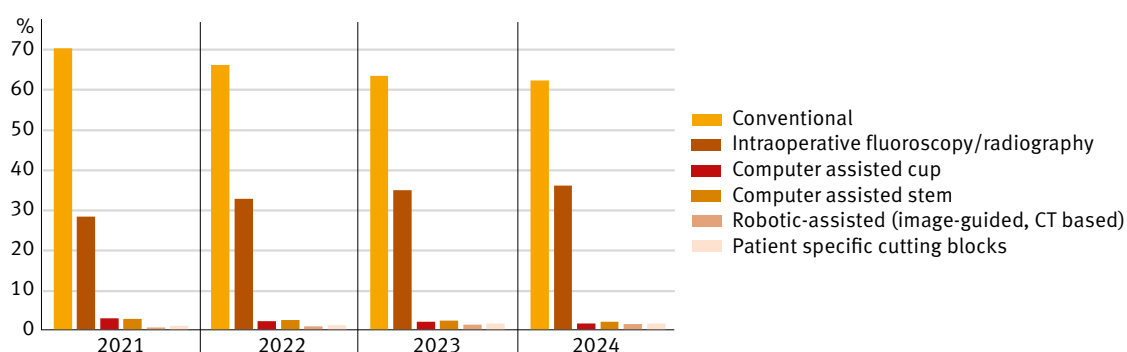
Fixation method	Women	Men	All
All cemented	1.1	0.5	0.8
All uncemented	82.6	92.6	87.4
Hybrid acetabulum uncemented, femur cemented	15.6	6.6	11.2
Reverse hybrid acetabulum cemented, femur uncemented	0.5	0.2	0.4
Reinforcement ring, femur cemented	0.16	0.07	0.12
Reinforcement ring, femur uncemented	0.1	0.1	0.1
N	57,338	53,616	110,954

Table 4.7

Primary total hip arthroplasty: fixation methods in primary osteoarthritis by sex (in %)

Technology

Navigation, robotics or patient specific instruments (PSI) are not widely used in THA in Switzerland, contrary to knee arthroplasty. Such aids were used in 6.1% of all cases only. On the other hand, intraoperative fluoroscopy was reported to be used in about one third of all cases (32.7%). The use of fluoroscopy increased over the last three years (Figure 4.5).



Technology [%]	2021	2022	2023	2024	2021–2024
Conventional	69.6	65.5	62.7	61.7	64.5
Intraoperative fluoroscopy/radiography	27.9	32.2	34.4	35.5	32.7
Computer assisted cup	2.6	2.0	1.7	1.4	1.9
Computer assisted stem	2.6	2.2	2.1	1.8	2.2
Robotic-assisted (image guided, CT based)	0.4	0.6	1.1	1.2	0.9
Patient specific cutting blocks	0.8	1.0	1.4	1.3	1.1
N	17,268	18,946	19,973	20,515	76,702

Figure 4.5

Primary total hip arthroplasty: technologies used in primary OA

Multiple responses possible (percentages do not sum to 100)

4.2 First revision within 2 years of primary total hip arthroplasty

First revisions are revisions that are linked to a primary THA registered in SIRIS, occurring for the first time, as opposed to re-revisions, which are repeated revisions after previous revisions. SIRIS differentiates between early revisions, performed within the first 2 years after implantation and long-term revisions, now up to 12 years after the index operation.

The 2-year revision rates were calculated for a moving 4-year window, which includes the last 4 years with a full 2-year follow-up. For this report, this corresponds to primary THA implanted between 01.01.2019 and 31.12.2022. For long-term outcomes, cumulative revision rates were calculated using KM survival estimations.

For primary OA, the 2-year revision rate was 2.4% (CI 2.3 – 2.6%), whereas for secondary OA it was 4.0% (CI 3.6 – 4.5%) and for fractures 5.0% (CI 4.0 – 5.5%), each significantly different from the others (Table 4.9).

The lowest 2-year revision rate (1.8%) was observed in the age group 55 – 64 years old. The revision rate for primary OA improved by 0.01% compared to previous 4-year period.

		Primary	Revised within 24 months			
			Revised	95% CI		
		N at risk*	N	%**	lower	upper
Overall (moving average)		86,177	2,395	2.8	2.7	2.9
Diagnosis	Primary OA	70,465	1,707	2.4	2.3	2.6
	Secondary OA	7,900	316	4.0	3.6	4.5
	Fracture	7,812	372	5.0	4.5	5.5
Overall Primary OA		70,465	1,707	2.4	2.3	2.6
Sex	Women	36,261	897	2.5	2.3	2.7
	Men	34,204	810	2.4	2.2	2.6
Age group	<55	6,955	189	2.7	2.4	3.1
	55–64	15,894	319	2.0	1.8	2.2
	65–74	23,241	543	2.3	2.2	2.6
	75–84	20,030	543	2.7	2.5	3.0
	85+	4,345	113	2.6	2.2	3.2
BMI group	<18.5	985	20	2.1	1.3	3.2
	18.5–24.9	21,561	401	1.9	1.7	2.1
	25–29.9	24,851	545	2.2	2.0	2.4
	30–34.9	11,603	331	2.9	2.6	3.2
	35–39.9	3,477	153	4.4	3.8	5.2
	40+	1,132	65	5.8	4.6	7.3
Unknown		6,856	192	2.8	2.5	3.2
Morbidity state	ASA 1	7,704	129	1.7	1.4	2.0
	ASA 2	41,218	894	2.2	2.0	2.3
	ASA 3	17,958	581	3.3	3.0	3.5
	ASA 4/5	433	13	3.1	1.8	5.3
	Unknown	3,152	90	2.9	2.3	3.5

Table 4.9

First revision of primary total hip arthroplasty within 24 months overall and according to baseline characteristics

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

* Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

** Rates adjusted for effects of mortality and emigration.

Reason for early first revision of THA for primary OA

A total of 1,707 revisions of primary THA performed for primary OA were registered during the current 4-year moving window. This is almost the same number (1,708 revisions) as during the window 2018 – 2021 used in the last annual report. However, the number of primary THA has increased from 67,616 to 70,465. The most frequent cause of early first revision of primary THA for primary OA was infection (30.5%), followed by periprosthetic fracture (19.6%), dislocation (17.0%), and femoral loosening (16.0%) (Table 4.10). There was a substantial increase in the share of revisions attributed to infections from 27.9% in the previous report to currently 30.5%. Approximately one-sixth of all revisions (14.9%) were undertaken for malposition of either acetabular or femoral components.

Reason	revised	%
Infection	520	30.5
Periprosthetic fracture	334	19.6
Dislocation	291	17.0
Loosening femoral	273	16.0
Position/orientation of cup	143	8.4
Loosening acetabular	134	7.9
Position/orientation of stem	112	6.6
Wear	17	1.0
Spacer	14	0.8
Implant failure	13	0.8
Impingement	8	0.5
Osteolysis FE	7	0.4
Trochanter pathology	6	0.4
Osteolysis AC	5	0.3
Acetabular protrusion	4	0.2
Squeaking	3	0.2
Other	201	11.8
Total Revisions	1,707	

Table 4.10

Reason for early first revision of primary total hip arthroplasty (primary OA)

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

Early first revisions are those occurring within 2 years of the primary arthroplasty. Multiple responses possible (percentages do not sum to 100).

Type of revision surgery

During the current 4-year moving period, there were in total 1,707 first revisions. The spectrum reached from sole exchange of the head or of the inlay to complete exchange of all components (the latter making 13.9% of all revisions) (Table 4.11). The most frequently performed operation was the exchange of the femoral component only (24.7%). An exchange of the head and of the inlay was performed in 16.3%. There were 47 (2.8%) component reimplantations after Girdlestone or spacer implantation registered as first revision, despite no previous component removal being registered. Obviously, these were not first revisions but appeared as first revisions because no other revision such as spacer implantation or Girdlestone resection had been documented. The registry does not correct this faulty entry, as it documents that there was a revision performed, which is sufficient for the analysis. Surgeons are invited to document all the revisions performed, and to carefully choose among the options available during data entry to avoid such mistakes.

Type	N revised	%
Exchange femoral component	422	24.7
Exchange head and inlay	279	16.3
Exchange acetabular and femoral components	237	13.9
Exchange acetabular component and head	227	13.3
Exchange head	198	11.6
Exchange femoral component and inlay	92	5.4
Exchange acetabular component	79	4.6
Component reimplantation (after spacer or Girdlestone)	47	2.8
Other intervention	44	2.6
Component removal, spacer implantation	30	1.8
Exchange inlay	22	1.3
Exchange femoral component, inlay and osteosynthesis	19	1.1
Girdlestone	11	0.6
Total Revisions	1,707	100

Table 4.11

Type of early first revision of total hip arthroplasty (primary OA)

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

Early first revisions are those occurring within 2 years of the primary arthroplasty.

Implants and fixation

Three different situations of early first revision were analysed more in detail, revision of both the acetabular and the femoral implants, femoral revision only, and component reimplantation after spacer/Girdlestone procedures. The type of fixation, whether primary or revision implants were used, and the patient's age at revision were considered in the analysis. The distribution of implants used differed among these three revision situations (Table 4.12, see next page). Overall, primary stems were used in 65.3% of all these revisions. Among the cases with primary stems, uncemented stems (including uncemented

short stems) were used in 40% of all revisions and cemented primary stems in 26%. Uncemented revision stems were implanted in 33.8%. Uncemented stems were used more frequently in younger patients, the share of cemented stems increasing with age. The use of cemented revision stems was exceptional. For acetabular revisions, standard uncemented acetabular cups were used in 79.5% of the revisions, while cemented cups without cages were used in 9.7%. Reconstruction cages were used in 9.1%. Specific revision cups were used only rarely. An overview of the implants used is provided in Table 4.13.

Femoral components		N	N		Acetabular comp.		N	N
Cem. primary stems 10+			Short stems* 10+		Cem. primary cups 10+		Delta TT DM	13
Twinsys	61		Optimys	40	Polarcup	27	Liberty	12
SPII Lubinus	43		Amistem-C	25	Versacem	16	Mpact DM	10
Corail	37		Fitmore	21	DS evolution (cem)	15	Other cups	82
Quadra-C	27		Other stems	25	Avantage	10	Revision cups (7+)	
Quadra-P (cem)	27		Cem. revision stems 10+		Other cups	20	Pinnacle	7
Avenir	19		Arcad L XL	10	Uncem. primary cups 10+		Other cups	8
Weber	18		Other stems	3	RM pressfit vitamys	80	AC roof ring or cage 10+	
Centris	12		Uncem. revision stems 10+		Pinnacle	72	ZB reinforcement	51
MS-30	11		Corail collared	122	Allofit	58	rings	
Other stems	57		Revitan	55	Versafitcup DM	56	Burch-Schneider cage	16
Uncem. primary stems 10+			Wagner SL	39	Bi-Mentum	54	Other cages	12
Corail collared	83		MRP-titan	29	Polarcup	48		
Avenir	50		Quadra-R	29	Versafitcup trio/ccl.	40		
Polarstem	40		Lima revision	25	Symbol DMHA	34		
Twinsys	40		M-Vizion	21	Avantage	25		
Quadra-P	34		Mathys modular	21	Mpact	24		
Quadra-H	30		revision		DS evolution	23		
Corail	24		Restoration modular	18	G7 DM hemispherical	22		
CLS Spotorno	20		Alloclassic SLL	17	TM	20		
Stellaris	10		Redapt	17	Fitmore	19		
Other stems	47		Other stems	21	R3	15		

Table 4.13

Hip early first revision of primary OA THA: main brands used (2019 – 2024)

e-class categories used: 34-32-10-01, 34-32-10-02, 34-32-10-03, 34-32-10-05, 34-32-10-06, 34-32-10-08, 34-32-10-09, 34-32-10-10, 34-32-10-11. A small proportion of tumor systems such as MUTARS is excluded.

All registered component revisions of four main types 2019-2024 with at least one FE/AC revision component with a known e-class

* Please note that both Fitmore and Amistem are originally classified as a regular primary stems. We reclassified them as short stems.

Category of implant		Age at revision						N
		<45	45-54	55-64	65-74	75-84	85+	
Type of revision of femoral components		%	%	%	%	%	%	
AC + FE revision	cem. primary stems	0.0	14.3	14.8	18.1	27.1	41.2	72
	uncem. primary stems	55.6	54.8	45.7	37.1	33.3	29.4	145
	short stems	11.1	7.1	7.4	10.3	10.4	5.9	33
	cem. revision stems	0.0	0.0	1.2	0.0	0.0	0.0	1
	uncem. revision stems	33.3	23.8	30.9	34.5	29.2	23.5	110
		100%	100%	100%	100%	100%	100%	361
FE revision (with or without inlay)	cem. primary stems	10.0	26.3	22.6	31.3	26.1	46.2	203
	uncem. primary stems	90.0	47.4	33.1	23.2	22.3	13.5	189
	short stems	0.0	13.2	13.5	6.6	10.7	1.9	68
	cem. revision stems	0.0	0.0	0.0	1.0	2.4	0.0	9
	uncem. revision stems	0.0	13.2	30.8	37.9	38.5	38.5	253
		100%	100%	100%	100%	100%	100%	722
Component reimplantation (after spacer)	cem. primary stems	0.0	0.0	11.1	24.0	38.5	0.0	12
	uncem. primary stems	100.0	100.0	22.2	36.0	23.1	66.7	18
	short stems	0.0	0.0	0.0	4.0	0.0	0.0	1
	cem. revision stems							0
	uncem. revision stems	0.0	0.0	66.7	36.0	38.5	33.3	21
		100%	100%	100%	100%	100%	100%	52
Type of revision of acetabular components								
AC + FE revision	cem. primary cups	0.0	0.0	6.3	8.7	11.6	18.8	29
	uncem. primary cups	100.0	100.0	89.9	83.5	78.9	68.8	301
	revision cups	0.0	0.0	1.3	2.6	1.1	0.0	5
	AC roof ring or cage	0.0	0.0	2.5	5.2	8.4	12.5	18
		100%	100%	100%	100%	100%	100%	353
AC revision (with or without head)	cem. primary cups	0.0	6.1	5.6	6.9	19.1	10.3	45
	uncem. primary cups	100.0	87.9	82.0	79.4	63.2	69.0	322
	revision cups	0.0	0.0	4.5	1.5	0.7	0.0	7
	AC roof ring or cage	0.0	6.1	7.9	12.2	16.9	20.7	54
		100%	100%	100%	100%	100%	100%	428
Component reimplantation (after spacer)	cem. primary cups	0.0	0.0	0.0	8.7	30.0	33.3	6
	uncem. primary cups	100.0	100.0	66.7	78.3	70.0	66.7	35
	revision cups	0.0	0.0	11.1	8.7	0.0	0.0	3
	AC roof ring or cage	0.0	0.0	22.2	4.3	0.0	0.0	3
		100%	100%	100%	100%	100%	100%	47

Table 4.12

Hip early first revision of primary OA THA: main components used by age at revision (2019 – 2024)

e-class categories used: 34-32-10-01, 34-32-10-02, 34-32-10-03, 34-32-10-05, 34-32-10-06, 34-32-10-08, 34-32-10-09, 34-32-10-10, 34-32-10-11. A small proportion of tumor systems such as MUTARS is excluded.

All registered component revisions of four main types 2019-2024 with at least one FE/AC revision component with a known e-class

Early revision rate according to stem fixation, bearing and approach

Table 4.14 gives an overview of the revision rates depending on implant fixation, bearing and surgical approach, whereby the overall 2-year revision rate was 2.4% (1,7087 of 70,465 primary THA performed for primary OA). The parameters that were associated with above average revision rates included all cemented fixation (2.9%) and bearings using CPE (3.4% – 4.9%). The use of a posterior approach and other unspecified approaches accounted for 3.3% and 3.8%, respectively.

Timing of revision

Figures 4.6 shows the cause and frequency distribution (Kernel density estimation) of early revision overall and separately for cemented and uncemented femoral implants. With the exception of aseptic loosening, most revisions occurred during the first 3 months after primary THA, including high and early peaks of periprosthetic fractures and dislocation as reasons for revision. Although periprosthetic joint infection and aseptic loosening were more frequent complications, their curves were flatter and remained elevated over a longer period. In case of cemented

	N at risk*	Revised		95% CI	
		N	%**	lower	upper
Overall (moving average)	70,465	1,707	2.4	2.3	2.6
Fixation					
All cemented	639	18	2.9	1.8	4.5
All uncemented	61,666	1,474	2.4	2.3	2.5
Hybrid	7,999	206	2.6	2.3	3.0
Articulation					
Metal on conventional polyethylene (MoCPE)	1,152	39	3.4	2.5	4.7
Ceramic on conventional polyethylene (CoCPE)	3,009	103	3.5	2.9	4.2
Metal on cross-linked polyethylene (MoXLPE)	6,499	192	3.0	2.6	3.4
Ceramic on cross-linked polyethylene (CoXLPE)	43,989	1,007	2.3	2.2	2.4
Ceramic on ceramic (CoC)	9,858	248	2.5	2.2	2.9
Ceramicised metal on conventional polyethylene (CMoCPE)	209	10	4.9	2.7	8.9
Ceramicised metal on cross-linked polyethylene (CMoXLPE)	4,572	74	1.6	1.3	2.0
Approach					
Anterior	38,120	869	2.3	2.1	2.4
Anterolateral	20,612	476	2.3	2.1	2.5
Lateral	2,371	57	2.4	1.9	3.1
Posterior	9,066	294	3.3	2.9	3.7
Other approach	296	11	3.8	2.1	6.8

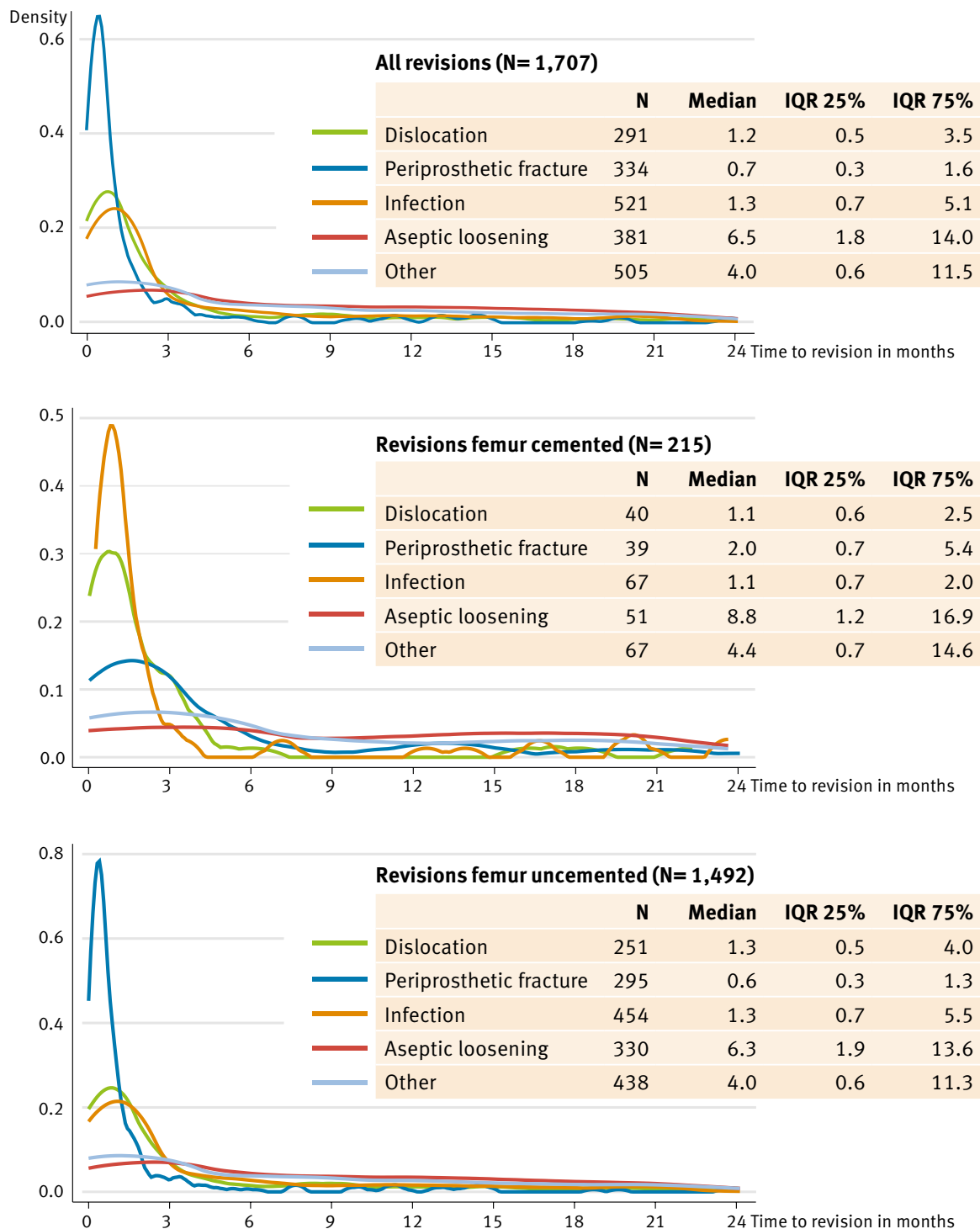
Table 4.14

First revision of primary total hip arthroplasty within 24 months according to stem fixation, articulation and approach

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

* Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

** Rates adjusted for effects of mortality and emigration.



Figures 4.6

Reason for early first revision by time interval since primary total hip arthroplasty

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

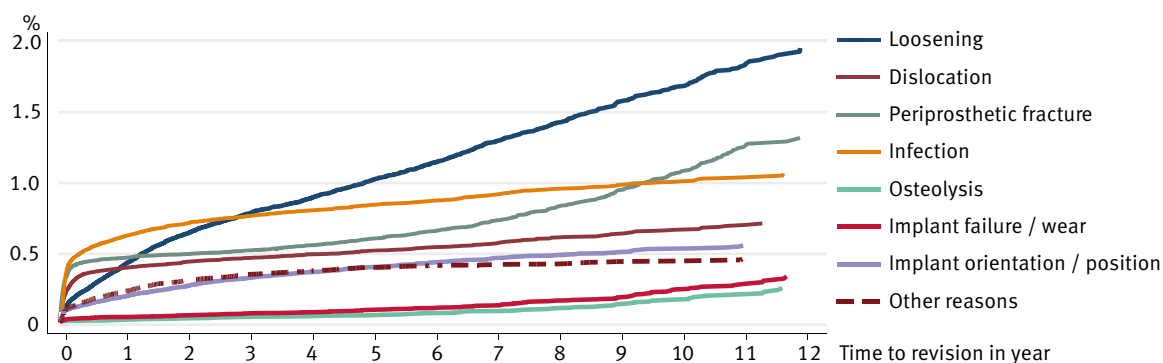
Early first revisions are those occurring within 2 years of the primary arthroplasty

stems, dislocation and infection were the predominant reasons for early revision, whereas other complications occurred later and were distributed over a longer period. In uncemented stems, periprosthetic fractures predominated the early postoperative period and occurred at a higher frequency.

The cumulative incidence rates show the long-term behaviour of implants, illustrating the proportion of implants having experienced at least one revision. **Figures 4.7** presents the cumulative incidence rates up to 12 years postoperatively for all THA performed for primary OA, overall and separately for cemented and uncemented femoral components. Cumulative revision rates for infection, dislocation and osteoly-

sis did not differ between cemented and uncemented stems, whereas periprosthetic fractures occurred more frequently with uncemented stems, both on the short term and up to eleven years postoperatively. Loosening and periprosthetic fractures were the dominant reasons for revisions over the whole observation period. The curve for loosening shows linear increase. The curve for periprosthetic fractures rises faster in the three first months for uncemented stems but flattens out until the fifth year for uncemented and the seventh year for cemented stems. Thereafter, the curves start to rise until it becomes the second most important cause for revision after nine years for uncemented stems.

Primary osteoarthritis (OA) total hip arthroplasty – all fixation methods



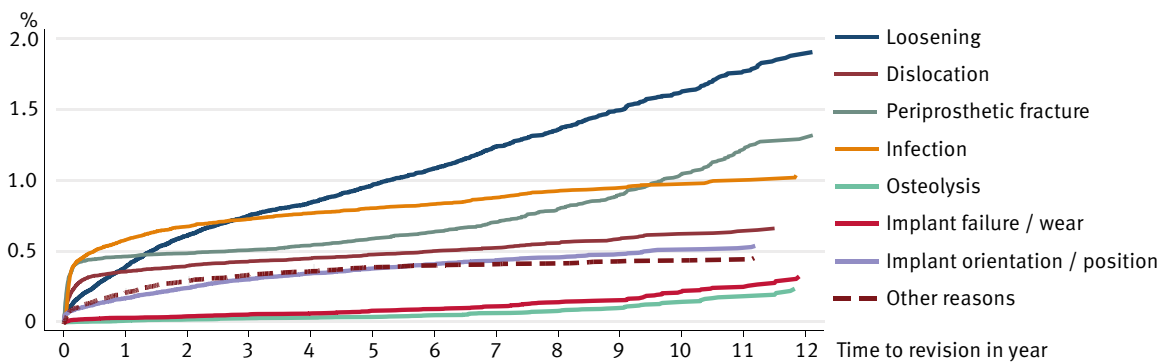
Diagnoses	1 year	2 years	3 years	5 years	9 years	10 years	12 years
Loosening	0.3 (0.3-0.4)	0.6 (0.5-0.7)	0.7 (0.6-0.9)	1.1 (1.0-1.3)	1.7 (1.5-1.9)	1.8 (1.6-2.1)	2.2 (1.6-2.8)
Dislocation	0.5 (0.4-0.6)	0.5 (0.5-0.6)	0.6 (0.5-0.7)	0.6 (0.5-0.8)	0.8 (0.7-0.9)	0.8 (0.7-1.0)	0.9 (0.7-1.2)
Periprosthetic fracture	0.3 (0.3-0.4)	0.4 (0.3-0.5)	0.4 (0.4-0.5)	0.5 (0.4-0.6)	0.9 (0.7-1.1)	1.0 (0.8-1.3)	1.1 (0.9-1.4)
Infection	0.7 (0.6-0.8)	0.8 (0.7-0.9)	0.8 (0.7-0.9)	0.9 (0.8-1.1)	1.0 (0.9-1.2)	1.1 (0.9-1.2)	1.1 (0.9-1.3)
Osteolysis	0.0 (0.0-0.0)	0.0 (0.0-0.1)	0.1 (0.0-0.1)	0.1 (0.0-0.1)	0.2 (0.1-0.3)	0.2 (0.1-0.4)	0.2 (0.1-0.4)
Implant failure / wear	0.0 (0.0-0.1)	0.0 (0.0-0.1)	0.0 (0.0-0.1)	0.1 (0.0-0.1)	0.2 (0.1-0.3)	0.3 (0.2-0.4)	0.3 (0.2-0.4)
Implant orientation / position	0.2 (0.1-0.3)	0.3 (0.2-0.3)	0.3 (0.2-0.4)	0.4 (0.3-0.5)	0.5 (0.4-0.6)	0.5 (0.4-0.6)	0.5 (0.4-0.6)
Other reasons	0.2 (0.1-0.2)	0.2 (0.2-0.3)	0.3 (0.2-0.3)	0.3 (0.2-0.4)	0.3 (0.3-0.4)	0.3 (0.3-0.4)	0.3 (0.3-0.4)

Figure 4.7 Part one

Cumulative incidence rates for different revision diagnoses (primary OA THA)

Time since operation, 2012–2024, % of implants revised

Primary osteoarthritis (OA) total hip arthroplasty – uncemented femur



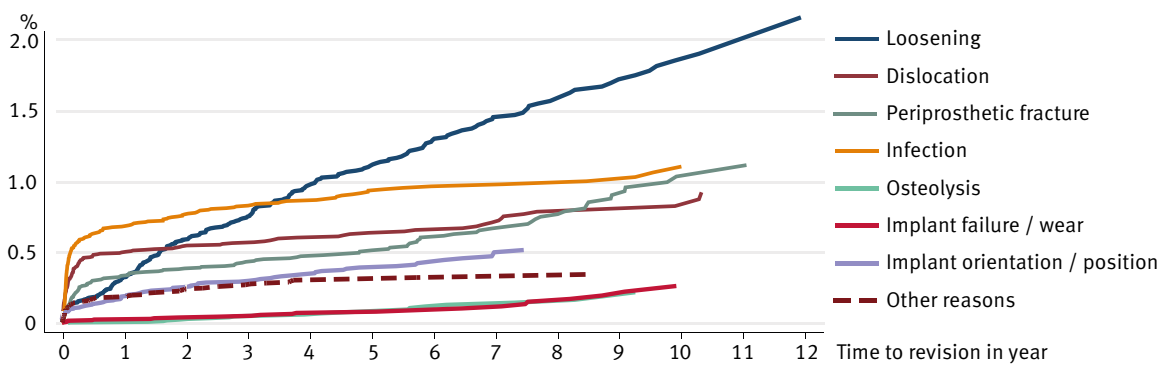
	1 year	2 years	3 years	5 years	9 years	10 years	12 years
Loosening	0.4 (0.4-0.4)	0.6 (0.6-0.7)	0.8 (0.7-0.8)	1.0 (0.9-1.0)	1.5 (1.4-1.6)	1.6 (1.5-1.7)	1.9 (1.8-2.0)
Dislocation	0.4 (0.3-0.4)	0.4 (0.4-0.4)	0.4 (0.4-0.5)	0.5 (0.4-0.5)	0.6 (0.5-0.6)	0.6 (0.6-0.7)	0.7 (0.6-0.7)
Periprosthetic fracture	0.5 (0.4-0.5)	0.5 (0.5-0.5)	0.5 (0.5-0.5)	0.6 (0.6-0.6)	0.9 (0.9-1.0)	1.1 (1.0-1.1)	1.3 (1.2-1.4)
Infection	0.6 (0.5-0.6)	0.7 (0.6-0.7)	0.7 (0.7-0.8)	0.8 (0.8-0.8)	1.0 (0.9-1.0)	1.0 (0.9-1.0)	1.0 (1.0-1.1)
Osteolysis	0.0 (0.0-0.0)	0.0 (0.0-0.0)	0.0 (0.0-0.0)	0.0 (0.0-0.1)	0.1 (0.1-0.1)	0.1 (0.1-0.2)	0.2 (0.2-0.3)
Implant failure / wear	0.0 (0.0-0.0)	0.0 (0.0-0.1)	0.1 (0.0-0.1)	0.1 (0.1-0.1)	0.2 (0.1-0.2)	0.2 (0.2-0.3)	0.3 (0.3-0.4)
Implant orientation / position	0.2 (0.2-0.2)	0.2 (0.2-0.3)	0.3 (0.3-0.3)	0.4 (0.4-0.4)	0.5 (0.4-0.5)	0.5 (0.5-0.6)	0.5 (0.5-0.6)
Other reasons	0.2 (0.2-0.3)	0.3 (0.3-0.3)	0.4 (0.3-0.4)	0.4 (0.4-0.4)	0.5 (0.4-0.5)	0.5 (0.4-0.5)	0.5 (0.4-0.5)

Figure 4.7 Part two

Cumulative incidence rates for different revision diagnoses (primary OA THA)

Time since operation, 2012–2024, % of implants revised

Primary osteoarthritis (OA) total hip arthroplasty – cemented femur



	1 year	2 years	3 years	5 years	9 years	10 years	12 years
Loosening	0.3 (0.3-0.4)	0.6 (0.5-0.7)	0.7 (0.6-0.9)	1.1 (1.0-1.3)	1.7 (1.5-1.9)	1.8 (1.6-2.1)	2.2 (1.6-2.8)
Dislocation	0.5 (0.4-0.6)	0.5 (0.5-0.6)	0.6 (0.5-0.7)	0.6 (0.5-0.8)	0.8 (0.7-0.9)	0.8 (0.7-1.0)	0.9 (0.7-1.2)
Periprosthetic fracture	0.3 (0.3-0.4)	0.4 (0.3-0.5)	0.4 (0.4-0.5)	0.5 (0.4-0.6)	0.9 (0.7-1.1)	1.0 (0.8-1.3)	1.1 (0.9-1.4)
Infection	0.7 (0.6-0.8)	0.8 (0.7-0.9)	0.8 (0.7-0.9)	0.9 (0.8-1.1)	1.0 (0.9-1.2)	1.1 (0.9-1.2)	1.1 (0.9-1.3)
Osteolysis	0.0 (0.0-0.0)	0.0 (0.0-0.1)	0.1 (0.0-0.1)	0.1 (0.0-0.1)	0.2 (0.1-0.3)	0.2 (0.1-0.4)	0.2 (0.1-0.4)
Implant failure / wear	0.0 (0.0-0.1)	0.0 (0.0-0.1)	0.0 (0.0-0.1)	0.1 (0.0-0.1)	0.2 (0.1-0.3)	0.3 (0.2-0.4)	0.3 (0.2-0.4)
Implant orientation / position	0.2 (0.1-0.3)	0.3 (0.2-0.3)	0.3 (0.2-0.4)	0.4 (0.3-0.5)	0.5 (0.4-0.6)	0.5 (0.4-0.6)	0.5 (0.4-0.6)
Other reasons	0.2 (0.1-0.2)	0.2 (0.2-0.3)	0.3 (0.2-0.3)	0.3 (0.2-0.4)	0.3 (0.3-0.4)	0.3 (0.3-0.4)	0.3 (0.3-0.4)

Figure 4.7 Part three

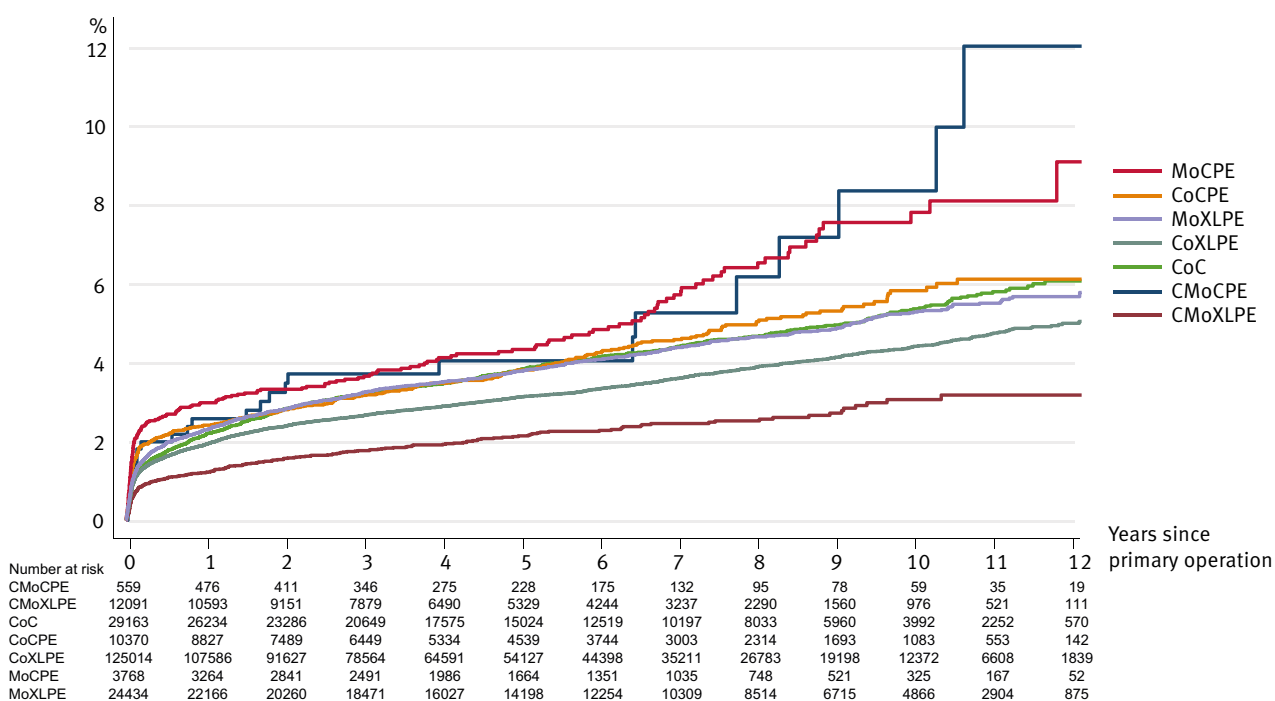
Cumulative incidence rates for different revision diagnoses (primary OA THA)

Time since operation, 2012–2024, % of implants revised

Bearings and head sizes

The 2-year revision rate for the current 4-year moving window was lowest for MCoXLPE (1.6%) and CoXLPE (2.4%), followed by CoC and MoXLPE (2.8%) (Figure 4.8). At 12 years, the estimated cumulative revision rate for CMoXLPE had the lowest revision rate of 3.2% (CI 2.7 – 3.8%) and CoXLPE had a revision

rate of 5.0% (CI 4.8 – 5.3%). In contrast, the highest cumulative revision rate was observed for CMoCPE with 12.0% (CI 7.0 – 20.2%), followed by MoCPE with 9.1% (CI 6.9 – 11.8) (Figure 4.8). However, this observation has to be taken with care for both bearings, considering small numbers with consecutively wide confidence intervals.



Bearing surf.	1 year	2 years	3 years	5 years	6 years	8 years	9 years	10 years	12 years
MoCPE	3.0 (2.5-3.6)	3.3 (2.8-3.9)	3.6 (3.1-4.3)	4.3 (3.7-5.1)	4.8 (4.1-5.7)	6.5 (5.5-7.7)	7.5 (6.3-8.9)	7.8 (6.5-9.3)	9.1 (6.9-11.8)
CoCPE	2.4 (2.1-2.7)	2.8 (2.5-3.1)	3.2 (2.8-3.5)	3.8 (3.4-4.2)	4.3 (3.9-4.8)	5.1 (4.5-5.6)	5.3 (4.8-6.0)	5.8 (5.2-6.5)	6.1 (5.4-6.9)
MoXLPE	2.3 (2.1-2.5)	2.8 (2.6-3.0)	3.2 (3.0-3.5)	3.8 (3.5-4.1)	4.1 (3.8-4.4)	4.6 (4.3-4.9)	4.9 (4.6-5.2)	5.3 (4.9-5.6)	5.8 (5.3-6.2)
CoXLPE	1.9 (1.8-2.0)	2.4 (2.3-2.5)	2.7 (2.6-2.7)	3.1 (3.0-3.2)	3.3 (3.2-3.5)	3.9 (3.8-4.0)	4.2 (4.0-4.3)	4.4 (4.2-4.6)	5.0 (4.8-5.3)
CoC	2.2 (2.0-2.3)	2.8 (2.6-3.0)	3.2 (3.0-3.4)	3.8 (3.6-4.1)	4.1 (3.9-4.4)	4.7 (4.4-5.0)	5.0 (4.6-5.3)	5.4 (5.0-5.7)	6.0 (5.6-6.5)
CMoCPE	2.6 (1.5-4.3)	3.5 (2.2-5.5)	3.7 (2.4-5.8)	4.0 (2.6-6.2)	4.0 (2.6-6.2)	6.2 (3.8-9.9)	8.3 (5.0-13.6)	8.3 (5.0-13.6)	12.0 (7.0-20.2)
CMoXLPE	1.2 (1.0-1.4)	1.6 (1.4-1.8)	1.8 (1.5-2.0)	2.1 (1.9-2.4)	2.3 (2.0-2.6)	2.6 (2.2-2.9)	2.8 (2.4-3.2)	3.0 (2.6-3.6)	3.2 (2.7-3.8)

Figure 4.8

Estimated failure rates of primary total hip arthroplasty for different bearing surfaces

Time since operation, 2012–2024, % of implants revised

Head size and bearing type influence wear, which can lead in the long term to osteolysis and loosening. Head size has an impact on stability, with larger heads being more stable, reducing the risk of dislocation, compared to smaller diameters. However, large heads bear the risk of increased wear, an issue particularly relevant with CPE. Heads with a diameter ≥ 40 mm had an excessively high revision rate of 12.3% at 12 years (Figure 4.9). However, these were used mostly in CoC bearings. Bearings with diameter 32 or 36 mm had an almost identical 12-year revision rate

of 5.1% (CI 4.8 – 5.4%) and 5.2% (CI 4.9 – 5.6%), respectively. The 28 mm heads were associated with a significantly higher revision rate at 12 years of 6.1% (CI 5.5 – 6.8%), compared to 32 mm heads 5.1% (CI 4.8 – 5.4%).

The relationship between head size and bearing type was further analysed, whereby results were broken down between head sizes of 28, 32, and 36 mm and the bearings MoCPE, CoCPE, MoXLPE, CoXLPE, CMoXLPE, and CoC. The results are presented in Table 4.15. MoCPE was used only in combination with 32 mm

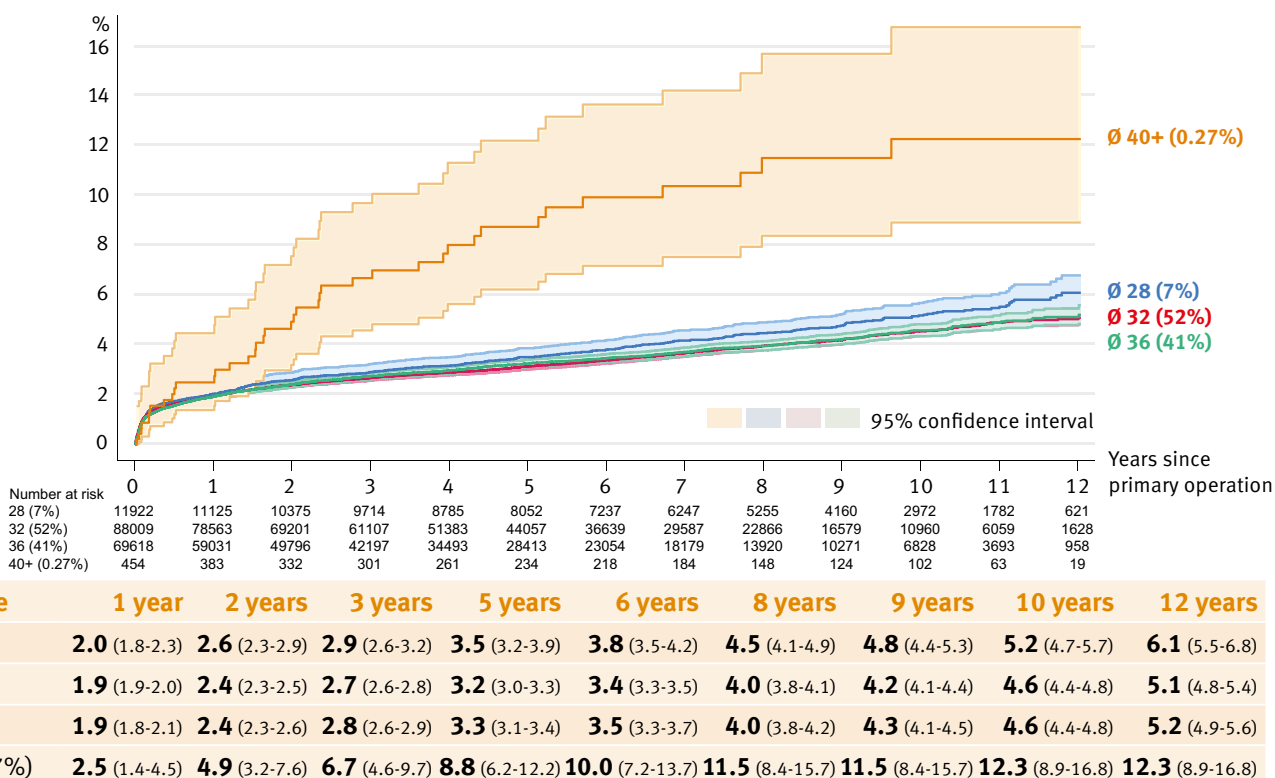


Figure 4.9
Estimated failure rates of primary total hip arthroplasty for different types of head sizes (standard cups: all uncemented fixation)

Time since operation, 2012–2024, diagnosis primary OA

heads (32/MoCPE) and appeared to have the highest revision rate at 11 years of 11.5% (CI 8.3 – 15.9%). MoCPE with 28 mm heads (28/MoCPE) were mainly used in fractures, cemented stems and with dual mobility cups. These subgroups were not part of this analysis.

For the head diameters 32 and 36 mm, ceramicised metal on cross-linked PE (CMoXLPE) had the lowest revision rates with 2.6% (CI 2.1 – 3.3%) and 3.2% (CI 2.3 – 4.4%) respectively at 12 years. Although 32/CMoXLPE had the lowest revision rates, these results

should be interpreted with caution. It is used with a limited number of stems and cups from one company only, all of which perform very well in the registry. In addition, CM is not widely used but concentrates on few centres and few surgeons, further introducing a potential bias.

At 12 years 32/CoXLPE had a revision rate of 4.8% (CI 4.4 -5.2%). The revision rate for 36/CoXLPE was not significantly different with 4.7% (CI 4.3 - 5.1%). In contrast, the 36 diameter heads for CoC (6.2%, CI 5.6-7.0%) and MoXLPE (5.4%, CI 4.0-7.2%) showed higher revision rates.

Head size	1 year	2 years	3 years	5 years	6 years	8 years	9 years	10 years	12 years
28mm									
CoCPE	2.7 (1.8-4.0)	2.8 (1.9-4.2)	3.3 (2.3-4.8)	4.0 (2.9-5.6)	4.2 (3.0-5.7)	4.8 (3.5-6.6)	4.8 (3.5-6.6)	4.8 (3.5-6.6)	5.5 (4.0-7.6)
MoXLPE	2.5 (1.9-3.1)	2.9 (2.3-3.6)	3.4 (2.8-4.2)	3.9 (3.2-4.7)	4.4 (3.6-5.3)	5.1 (4.3-6.1)	5.4 (4.5-6.4)	5.9 (4.9-7.0)	6.2 (5.2-7.5)
CoXLPE	1.8 (1.5-2.1)	2.4 (2.1-2.8)	2.6 (2.2-3.0)	3.2 (2.8-3.7)	3.5 (3.0-4.0)	4.1 (3.6-4.7)	4.4 (3.8-5.0)	4.7 (4.1-5.4)	6.0 (5.0-7.0)
CoC	2.1 (1.3-3.3)	3.0 (2.0-4.4)	3.4 (2.3-4.9)	4.1 (2.9-5.8)	4.5 (3.2-6.3)	4.9 (3.6-6.8)	5.3 (3.8-7.3)	5.3 (3.8-7.3)	5.3 (3.8-7.3)
32mm									
MoCPE	5.0 (3.4-7.3)	5.6 (3.9-8.1)	5.9 (4.1-8.3)	6.7 (4.8-9.4)	7.3 (5.3-10.2)	10.0 (7.3-13.7)	11.5 (8.3-15.9)	11.5 (8.3-15.9)	
CoCPE	2.7 (2.1-3.5)	3.1 (2.4-3.9)	3.5 (2.8-4.4)	4.1 (3.3-5.0)	4.6 (3.8-5.7)	5.4 (4.4-6.6)	6.0 (4.9-7.4)	7.4 (5.9-9.2)	7.7 (6.1-9.7)
MoXLPE	2.6 (2.3-3.0)	3.2 (2.9-3.6)	3.7 (3.3-4.1)	4.3 (3.9-4.7)	4.5 (4.1-5.0)	5.1 (4.6-5.6)	5.3 (4.9-5.9)	5.7 (5.2-6.3)	6.2 (5.6-6.8)
CoXLPE	1.8 (1.7-2.0)	2.3 (2.2-2.4)	2.5 (2.4-2.7)	2.9 (2.8-3.1)	3.1 (3.0-3.3)	3.7 (3.5-3.9)	4.0 (3.8-4.2)	4.2 (4.0-4.4)	4.8 (4.4-5.2)
CoC	2.0 (1.7-2.3)	2.6 (2.3-3.0)	2.9 (2.6-3.3)	3.5 (3.1-3.9)	3.9 (3.4-4.3)	4.3 (3.8-4.8)	4.5 (4.0-5.1)	4.9 (4.3-5.5)	5.3 (4.6-6.1)
CMoXLPE	0.9 (0.7-1.2)	1.3 (1.0-1.6)	1.5 (1.2-1.8)	1.8 (1.5-2.3)	2.0 (1.6-2.4)	2.3 (1.9-2.9)	2.5 (2.0-3.1)	2.6 (2.1-3.3)	2.6 (2.1-3.3)
36mm									
MoXLPE	1.8 (1.5-2.2)	2.4 (2.0-2.9)	2.8 (2.3-3.2)	3.2 (2.7-3.7)	3.4 (3.0-4.0)	3.8 (3.3-4.5)	4.1 (3.6-4.8)	4.4 (3.7-5.1)	5.4 (4.0-7.2)
CoXLPE	2.0 (1.9-2.2)	2.4 (2.3-2.6)	2.7 (2.6-2.9)	3.2 (3.0-3.4)	3.3 (3.1-3.5)	3.8 (3.6-4.1)	4.1 (3.8-4.4)	4.3 (4.0-4.7)	4.7 (4.3-5.1)
CoC	2.1 (1.9-2.3)	2.7 (2.5-3.0)	3.1 (2.9-3.4)	3.8 (3.5-4.1)	4.1 (3.8-4.4)	4.6 (4.2-5.0)	4.9 (4.5-5.3)	5.4 (4.9-5.9)	6.2 (5.6-7.0)
CMoXLPE	0.9 (0.7-1.3)	1.3 (1.0-1.7)	1.6 (1.3-2.1)	2.0 (1.6-2.5)	2.0 (1.6-2.5)	2.2 (1.7-2.8)	2.4 (1.9-3.2)	2.9 (2.2-3.9)	3.2 (2.3-4.4)

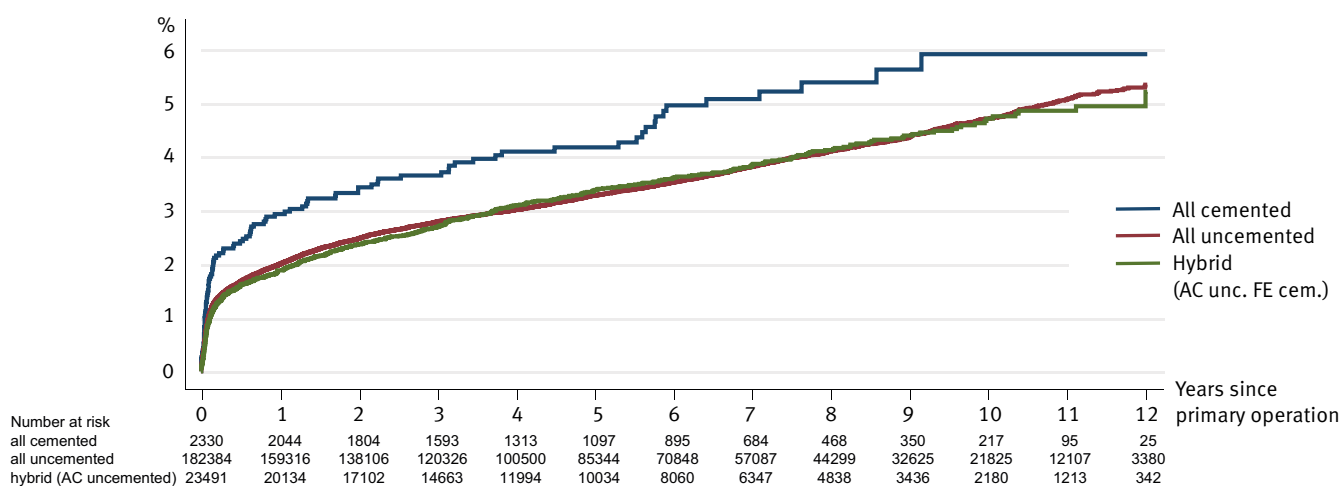
Table 4.15
Estimated failure rates of primary total hip arthroplasty for different types of head sizes by bearing surface (standard cups: all uncemented fixation)

Time since operation, 2012–2024, diagnosis primary OA. Only showing combinations with 500+ cases.

Fixation

Component fixation also affected revision rates (Figure 4.10). Hybrid fixation showed slightly better revision rates at 12 years with 5.2% (CI 4.6 – 6.0%) than uncemented fixation with 5.4% (CI 5.2 – 5.6%) or all

cemented with 5.9% (CI 4.7 – 7.4%). Although the revision rates for hybrid fixation tended to run below the revision rates for uncemented fixation for most of the observation time, confidence intervals remained largely overlapping, indicating no statistical significance (Figure 4.11).



Fixation	1 year	2 years	3 years	5 years	6 years	8 years	9 years	10 years	12 years
all cemented	2.9 (2.3-3.7)	3.4 (2.8-4.3)	3.7 (3.0-4.5)	4.2 (3.4-5.1)	5.0 (4.1-6.1)	4.2 (3.4-5.1)	5.0 (4.1-6.1)	5.9 (4.7-7.4)	5.9 (4.7-7.4)
all uncemented	2.0 (1.9-2.1)	2.5 (2.4-2.6)	2.8 (2.7-2.9)	3.3 (3.2-3.4)	3.5 (3.4-3.6)	3.3 (3.2-3.4)	3.5 (3.4-3.6)	4.7 (4.6-4.9)	5.4 (5.2-5.6)
hybrid*	1.9 (1.7-2.1)	2.4 (2.2-2.6)	2.7 (2.5-2.9)	3.4 (3.1-3.6)	3.6 (3.3-3.9)	3.4 (3.1-3.6)	3.6 (3.3-3.9)	4.7 (4.3-5.2)	5.2 (4.6-6.0)

Figure 4.10

Estimated failure rates of primary total hip arthroplasty for different fixation methods

Time since operation, 2012–2024, diagnosis primary OA. * acetabulum uncemented, femur cemented.

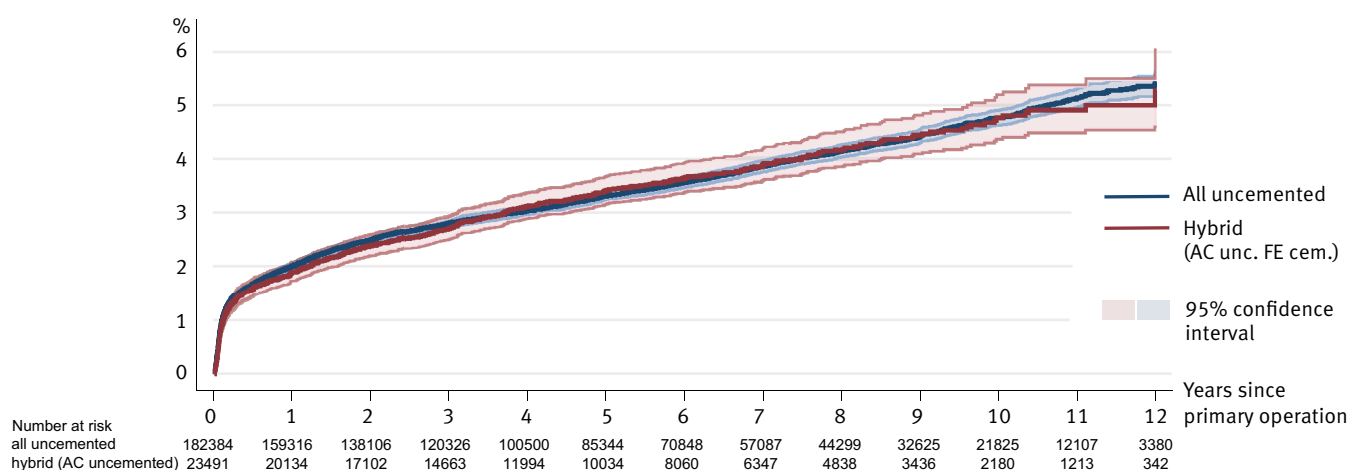


Figure 4.11

Estimated failure rates of primary total hip arthroplasty for different fixation methods

Time since operation, 2012–2024, diagnosis primary OA. * acetabulum uncemented, femur cemented.

Fixation	1 year	2 years	3 years	5 years	6 years	8 years	9 years	10 years	12 years
all uncemented	2.0 (1.9-2.1)	2.5 (2.4-2.6)	2.8 (2.7-2.9)	3.3 (3.2-3.4)	3.5 (3.4-3.6)	3.3 (3.2-3.4)	3.5 (3.4-3.6)	4.7 (4.6-4.9)	5.4 (5.2-5.6)
hybrid*	1.9 (1.7-2.1)	2.4 (2.2-2.6)	2.7 (2.5-2.9)	3.4 (3.1-3.6)	3.6 (3.3-3.9)	3.4 (3.1-3.6)	3.6 (3.3-3.9)	4.7 (4.3-5.2)	5.2 (4.6-6.0)

BMI

Data on BMI is collected by SIRIS since 2015. The observation time for this parameter therefore is limited to nine years. BMI had a high impact on the risk of revision (**Figure 4.12**), with a positive correlation (meaning increasing risk with increasing BMI).

The 2-year revision rate for patients with BMI >40 kg/m² was 6.4% (CI 5.6 – 7.4%), more than three times higher than in patients with normal weight. Most re-

visions occurred within the first 2 to 3 months and the most frequent reason for revision was infection, accounting for up to one-third of all revisions in this population. This was followed by periprosthetic fracture, femoral loosening, and dislocation. However, only infections were more frequent in this subgroup, whereas revisions for periprosthetic fractures and dislocations occurred approximately at the same rates as in patient with normal weight, while femoral and acetabular loosening were even less frequent.

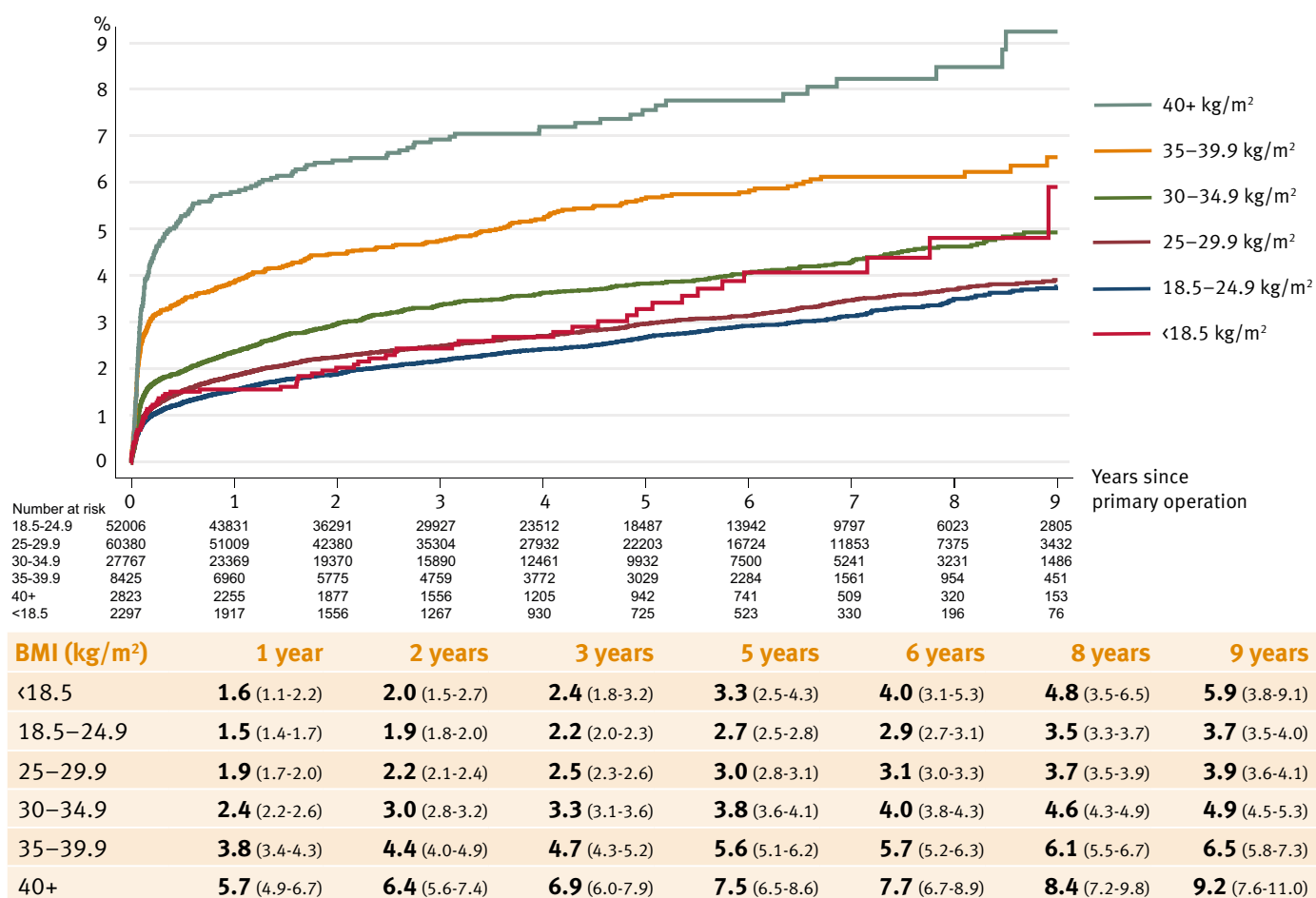


Figure 4.12
Estimated failure rates of primary total hip arthroplasty for different BMI

Time since operation, 2015–2024, diagnosis primary OA

While underweight patients initially had a lower revision risk, the revision rate started to rise at 5 years. At 9 years, the revision rates were comparable to those patients with a BMI between 35 and 39.9 kg/m². **Figure 4.13** shows the estimated failure rates with confidence intervals for the different groups. The differences are highly significant.

Dual mobility cups

Dual mobility cups (DMC) were increasingly used both for primary THA (**Figure 4.1**) as well as at revision. The main indication is to reduce the risk of dislocation, respectively the risk of revision for instability. The exact role of DMC is still debated, and several questions concerning their use are not yet fully answered.

Compared to regular cups, the cumulative revision rate for all DMC in the presence of an uncemented stem was elevated for the whole observation period (**Figure 4.14**).

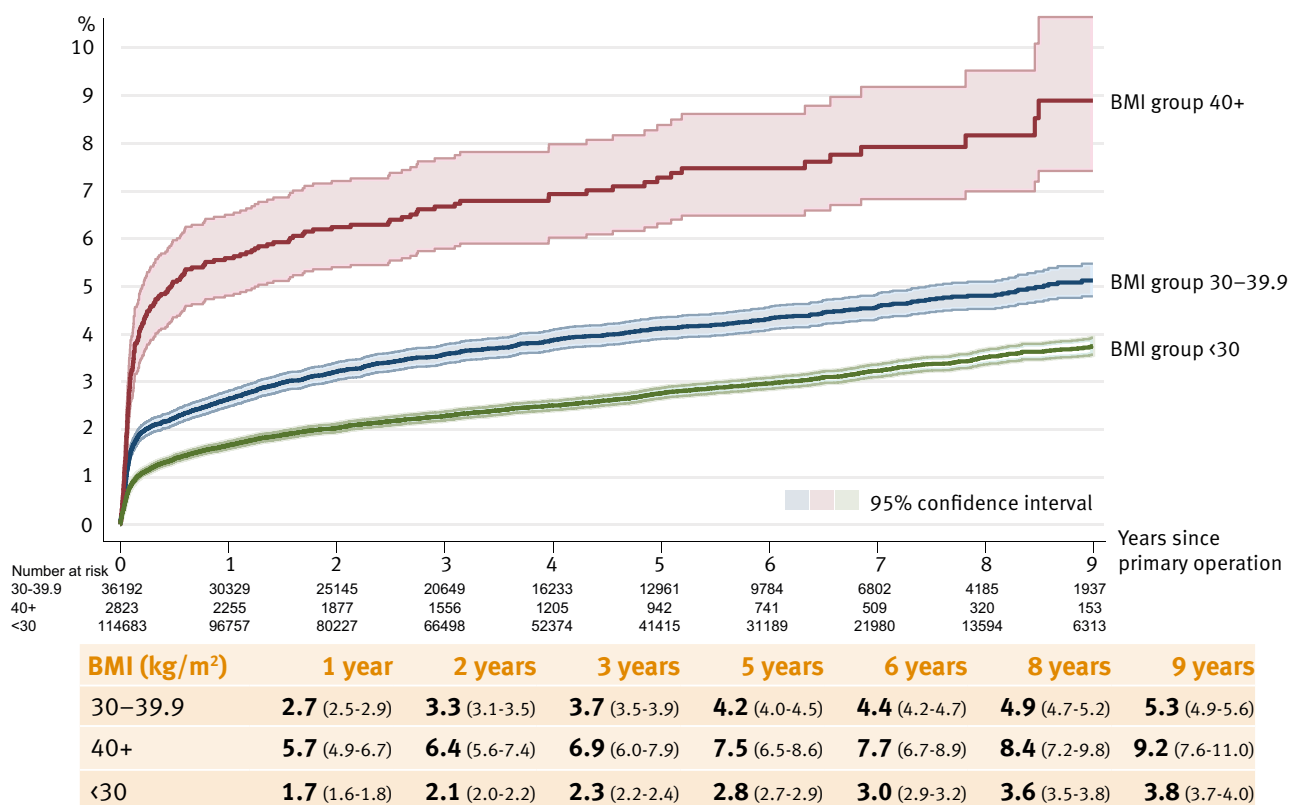


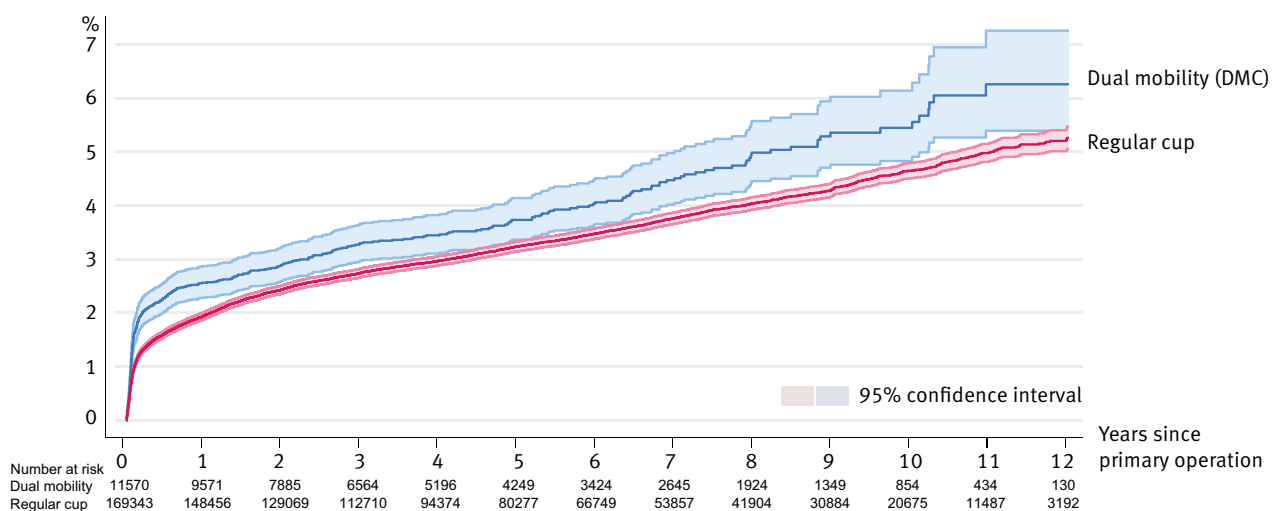
Figure 4.13

Estimated failure rates of primary total hip arthroplasty for different BMI

Time since operation, 2015–2024, diagnosis primary OA

The difference is statistically significant for the entire time period. The revision rate for DMC with cemented stem fixation (hybrid fixation) was lower, but still increased compared to THA with regular cups, with both

cemented and uncemented stems (Figure 4.15). This is most probably to be explained by a negative selection bias of patients.

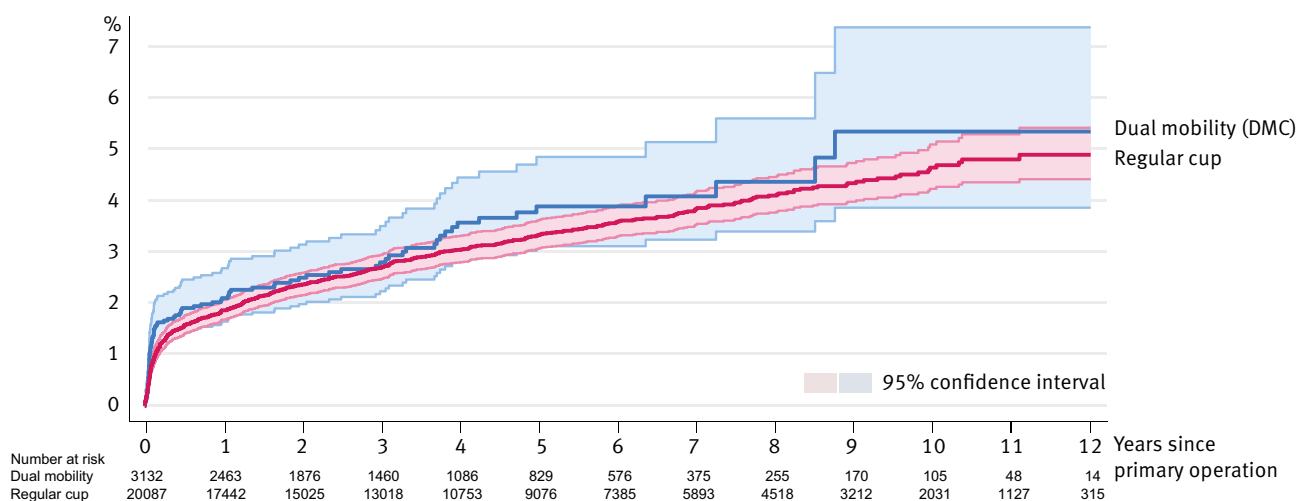


Typ of cup	1 year	2 years	3 years	5 years	6 years	8 years	9 years	10 years	12 years
Regular	2.0 (1.9-2.0)	2.4 (2.4-2.5)	2.8 (2.7-2.8)	3.2 (3.2-3.3)	3.5 (3.4-3.6)	4.0 (3.9-4.2)	4.3 (4.2-4.4)	4.6 (4.5-4.8)	5.3 (5.1-5.5)
Dual mobility	2.6 (2.3-2.9)	2.9 (2.6-3.2)	3.3 (3.0-3.7)	3.7 (3.4-4.1)	4.1 (3.7-4.5)	5.0 (4.5-5.6)	5.4 (4.8-6.0)	5.4 (4.8-6.1)	6.3 (5.4-7.3)

Figure 4.14

Estimated failure rates of primary total hip arthroplasty for different types of cups (all uncemented fixation)

Time since operation, 2012–2024, diagnosis primary OA



Typ of cup	1 year	2 years	3 years	5 years	6 years	8 years	9 years	10 years	12 years
Regular	1.9 (1.7-2.1)	2.4 (2.1-2.6)	2.7 (2.5-2.9)	3.3 (3.0-3.6)	3.6 (3.3-3.9)	4.1 (3.7-4.4)	4.3 (4.0-4.7)	4.6 (4.2-5.1)	4.9 (4.4-5.4)
Dual mobility	2.1 (1.6-2.7)	2.5 (2.0-3.1)	2.8 (2.2-3.5)	3.9 (3.1-4.8)	3.9 (3.1-4.8)	4.3 (3.4-5.6)	5.3 (3.8-7.3)	5.3 (3.8-7.3)	5.3 (3.8-7.3)

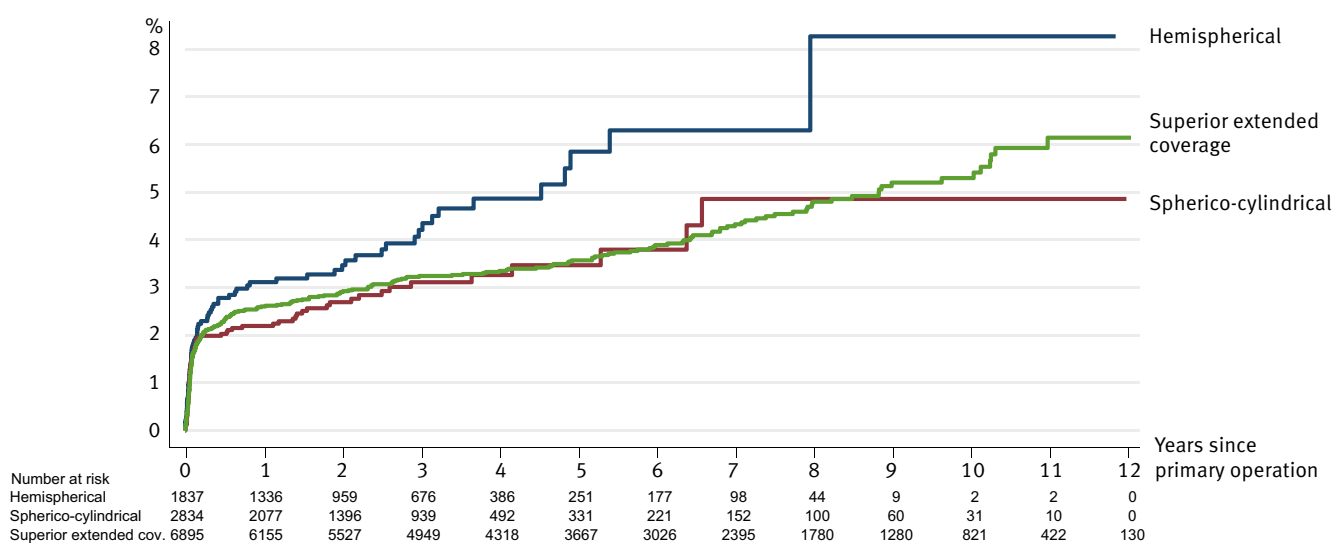
Figure 4.15

Estimated failure rates of primary total hip arthroplasty for different types of cups (hybrid fixation)

Time since operation, 2012–2024, diagnosis primary OA

DMC are provided in three different design philosophies: hemispherical, spherico-cylindrical, and with superior extended coverage. Comparison of these three types is possible until 8 years of follow-up, limited by the shorter observation time of the hemispher-

ical cups. Uncemented dual mobility cups with superior extended coverage had the lowest revision rate with 4.7% (CI 4.2– 5.7%) at eight years, increasing to 6.2% (CI 5.2 – 7.2%) at 12 years. Details regarding the DMC models are displayed in **Figure 4.16**.



Typ of DM-cup	1 year	2 years	3 years	5 years	6 years	8 years	9 years	10 years	12 years
Hemispherical	3.1 (2.4-4.0)	3.4 (2.7-4.4)	4.2 (3.2-5.4)	5.8 (4.4-7.7)	6.2 (4.6-8.4)	8.2 (4.9-13.6)			
Spherico-cylindrical	2.2 (1.7-2.8)	2.7 (2.1-3.4)	3.1 (2.4-3.9)	3.4 (2.7-4.4)	3.8 (2.8-5.0)	4.8 (3.3-7.0)	4.8 (3.3-7.0)	4.8 (3.3-7.0)	
Sup. extended cov.	2.6 (2.2-3.0)	2.9 (2.5-3.3)	3.2 (2.8-3.7)	3.5 (3.1-4.0)	3.9 (3.4-4.4)	4.7 (4.2-5.4)	5.1 (4.5-5.9)	5.2 (4.6-6.0)	6.1 (5.2-7.2)

Figure 4.16

Estimated failure rates of primary total hip arthroplasty for different types of dual mobility cups (all uncemented fixation)

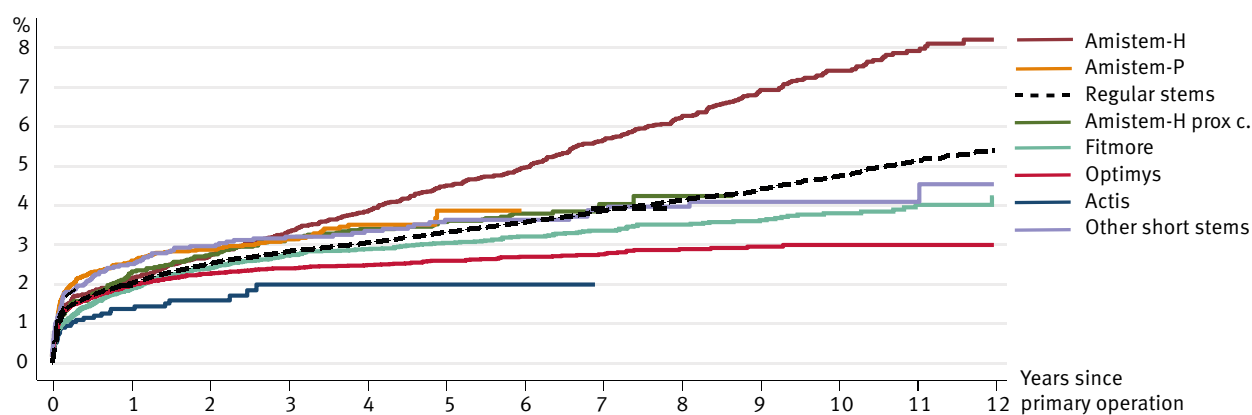
Time since operation, 2012–2024, diagnosis primary OA

* The hemispherical group is not well represented in SIRIS data. It comprises Symbol/DS evolution cups as well as the modular G7 cups

Short stems

The definition of a short stem remains a matter of debate, as no internationally accepted classification is available yet. Particularly, there is no consensus whether shortened stems with diaphyseal fixation should also be considered as short stems. For this analysis, the classification of the French Hip & Knee Society was used, which includes shortened stems with diaphyseal fixation¹. This classification separates short stems into five types, depending on the theoretical zone of fixation: cephalic (type 1), isolated cervical (type 2), calcar femoral (type 3), metaphyseal (type 4), and conventional metaphyseal-diaphyseal shortened stems (type 5). The classification of a wide variety of implants is provided in the publication. There are currently twenty-four different short stems or shortened stems in use in Switzerland. For statistical analysis, only stems with more than 500

implantations were considered individually in this report. The remaining short stems were grouped as “other short stems”. Compared to the standard uncemented stems, the short stems showed a wider range of revision rates (Figure 4.17), whereby the so-called calcar-guided short stems (type 3) performed well. Particularly the type 5 short stems showed a wide range of revision rates with some associated with excellent results, while others performed far less well. It is important to note that short stems did not universally perform well or poorly as a group. The reason for the heterogenous revision rates most likely is multifactorial, including the design of the stem, of any coating, the bearings used, etc. Hence, as in primary uncemented stems, each implant has to be assessed separately for its performance and longevity. However, the well-performing implants had a flatter revision curve than regular stems.



Typ of short stem	1 year	2 years	3 years	5 years	6 years	8 years	9 years	10 years	12 years
Actis	1.2 (0.8-1.8)	1.5 (1.0-2.1)	1.9 (1.3-2.7)	1.9 (1.3-2.7)	1.9 (1.3-2.7)				
Amistem-H	2.0 (1.7-2.4)	2.6 (2.3-3.0)	3.2 (2.9-3.6)	4.4 (4.0-4.9)	4.9 (4.4-5.4)	6.2 (5.7-6.8)	6.9 (6.3-7.5)	7.4 (6.8-8.1)	8.2 (7.5-9.0)
Amistem-H prox c.	2.2 (1.7-2.7)	2.6 (2.2-3.2)	3.1 (2.5-3.7)	3.5 (2.9-4.1)	3.7 (3.1-4.4)	4.2 (3.4-5.0)			
Amistem-P	2.4 (2.1-2.8)	2.8 (2.4-3.2)	3.0 (2.6-3.5)	3.8 (3.1-4.6)					
Fitmore	1.8 (1.6-2.0)	2.3 (2.1-2.5)	2.6 (2.4-2.9)	3.0 (2.7-3.2)	3.1 (2.8-3.4)	3.4 (3.1-3.8)	3.5 (3.2-3.9)	3.7 (3.4-4.1)	4.1 (3.6-4.7)
Optimys	1.8 (1.7-2.0)	2.2 (2.0-2.3)	2.3 (2.1-2.5)	2.5 (2.3-2.7)	2.6 (2.4-2.8)	2.8 (2.6-3.0)	2.9 (2.6-3.1)	2.9 (2.6-3.2)	2.9 (2.6-3.2)
Other short stems	2.4 (1.9-3.0)	2.9 (2.3-3.5)	3.1 (2.5-3.8)	3.5 (2.9-4.2)	3.6 (2.9-4.3)	3.9 (3.2-4.7)	4.0 (3.3-4.9)	4.0 (3.3-4.9)	4.5 (3.4-5.8)
Regular stems	2.0 (2.0-2.1)	2.6 (2.5-2.7)	2.9 (2.8-3.0)	3.4 (3.3-3.5)	3.6 (3.5-3.8)	4.2 (4.1-4.4)	4.5 (4.3-4.6)	4.8 (4.6-5.0)	5.5 (5.3-5.8)

Figure 4.17

Estimated failure rates of primary total hip arthroplasty for different types of short stems (all uncemented fixation)

Time since operation, 2012–2024, diagnosis primary OA

¹ Erivan R et al. French Hip & Knee Society classification of short-stem hip prostheses: Inter- and intra-observer reproducibility. Orthop Traumatol Surg Res. 2022, 108(1):103126.

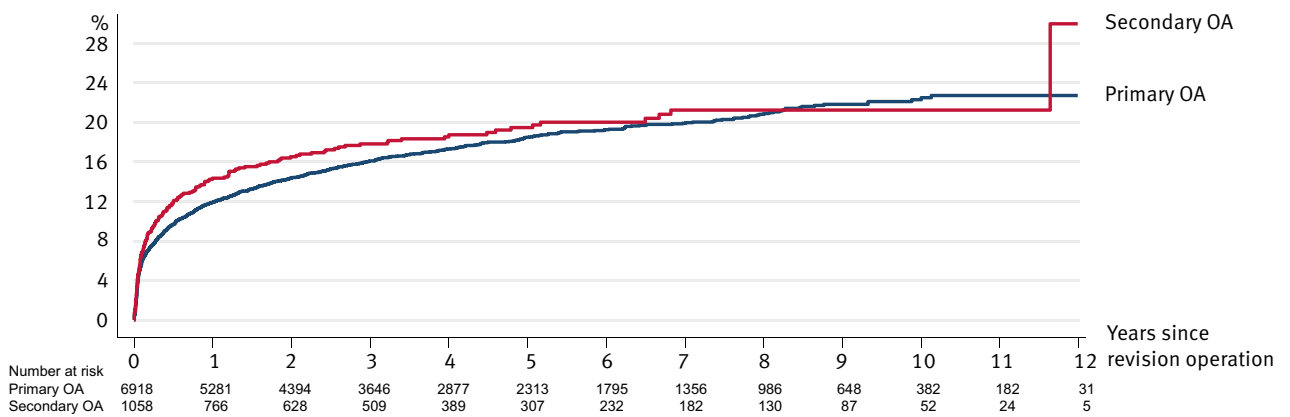
4.3 Re-revision of total hip arthroplasty

Re-revisions are revisions occurring after the first revision of a THA. The first or index revision includes partial or complete revisions and sometimes re-implantations after Girdlestone, when the removal of the implant was not documented in the register.

The re-revision rate is excessively high and reaches 11.9% (CI 11.1 – 12.7%) for primary OA after the first year and 14.4% (CI 12.3 – 16.7%) for secondary OA. That means that every 7th to 8th revision is revised within the first year. After the first year the curves flatten out and after the 7th year the revision rate barely

increases. At eleven years the revision rate is approximately 22% for both primary and secondary OA (Figure 4.18).

However, there is a number of patients who undergo additional revisions that may sum up to five or more revisions in selected cases. After five years 3.5% have been revised at least once, of which 0.46% had two revisions, 0.1% had three revisions, 0.02% had four revisions and 0.01% had five revisions or more (Figure 4.19). After 10 years overall 5.2% were revised. 4.2% had one re-revision, 0.72% two, 0.12% three, 0.07% four and 0.02% more than five revisions (Figure 4.19).



OA	1 year	2 years	3 years	5 years	6 years	8 years	10 years	12 years
Primary OA	11.9 (11.1-12.7)	14.4 (13.6-15.3)	16.1 (15.2-17.1)	18.5 (17.5-19.6)	19.3 (18.2-20.4)	20.8 (19.7-22.1)	22.3 (20.9-23.8)	22.8 (21.3-24.3)
Sec. OA	14.4 (12.3-16.7)	16.5 (14.4-19.0)	17.8 (15.5-20.4)	19.5 (17.0-22.3)	20.0 (17.5-23.0)	21.3 (18.4-24.5)	21.3 (18.4-24.5)	

Figure 4.18

Estimated failure rates after revision of total hip arthroplasty: primary and secondary OA

Time since revision, 2012–2024. Start point of analysis: first registered component revision in SIRIS that meets the inclusion criteria. End point of analysis: next registered component revision.

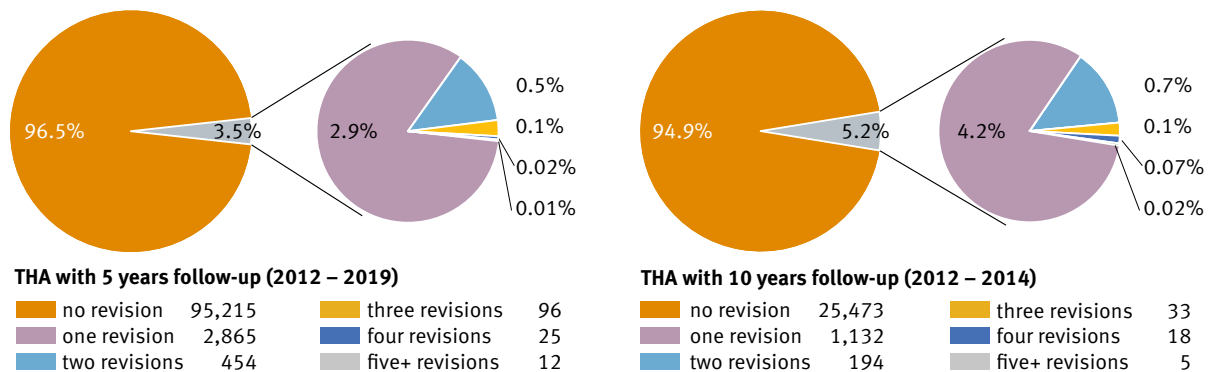


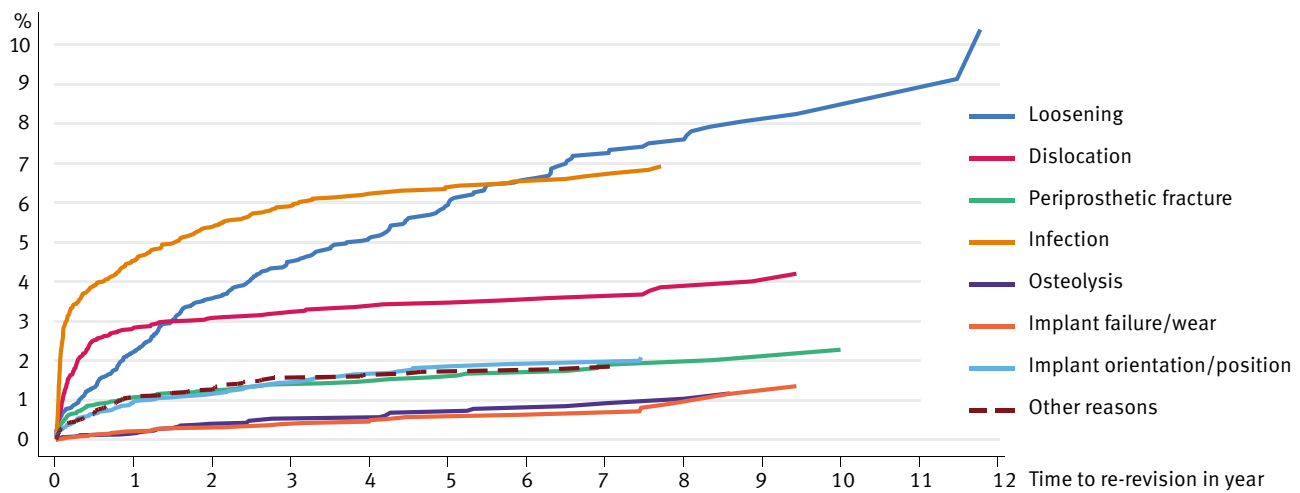
Figure 4.19

Risk of revision within 5 years and 10 years after primary THA (primary OA or secondary OA) in percent

Multi-stage operations are counted as one revision

4.3.1 Reasons for re-revision THA

In the first four years infection is the predominant diagnosis leading to a re-revision, followed by loosening and dislocations. From the sixth year onwards, loosening is the main reason for re-revisions. The cumulative incidence rate for loosening reaches 10.4% after 12 years (CI 7.6-14.0%) (Figure 4.20).



Diagnoses	1 year	2 years	3 years	5 years	9 years	10 years	12 years
Loosening	2.2 (1.9-2.6)	3.6 (3.1-4.1)	4.5 (4.0-5.1)	6.1 (5.4-6.8)	8.0 (7.1-9.1)	8.2 (7.2-9.4)	10.4 (7.6-14.0)
Dislocation	2.8 (2.5-3.3)	3.1 (2.7-3.5)	3.2 (2.8-3.7)	3.5 (3.0-4.0)	4.0 (3.4-4.7)	4.2 (3.5-5.0)	4.2 (3.5-5.0)
Periprosthetic fracture	1.1 (0.8-1.4)	1.3 (1.0-1.6)	1.4 (1.1-1.7)	1.6 (1.3-2.0)	2.0 (1.6-2.6)	2.3 (1.7-3.1)	2.3 (1.7-3.1)
Infection	4.5 (4.0-5.1)	5.4 (4.9-6.0)	5.9 (5.4-6.6)	6.4 (5.8-7.1)	6.9 (6.2-7.7)	6.9 (6.2-7.7)	6.9 (6.2-7.7)
Osteolysis	0.2 (0.1-0.3)	0.4 (0.3-0.6)	0.5 (0.4-0.8)	0.7 (0.5-1.0)	1.2 (0.8-1.8)	1.2 (0.8-1.8)	1.2 (0.8-1.8)
Implant failure / wear	0.2 (0.1-0.4)	0.3 (0.2-0.5)	0.4 (0.3-0.6)	0.6 (0.4-0.9)	1.2 (0.7-1.8)	1.4 (0.8-2.2)	1.4 (0.8-2.2)
Implant orientation / position	1.0 (0.8-1.3)	1.2 (0.9-1.5)	1.4 (1.2-1.8)	1.9 (1.5-2.3)	2.1 (1.7-2.6)	2.1 (1.7-2.6)	2.1 (1.7-2.6)
Other reasons	1.1 (0.9-1.4)	1.3 (1.1-1.7)	1.6 (1.3-2.0)	1.7 (1.4-2.2)	1.9 (1.5-2.4)	1.9 (1.5-2.4)	1.9 (1.5-2.4)

Figure 4.20

Cumulative incidence rates for different re-revision diagnoses after main component changes

Time since operation, 2012 – 2024, % of implants re-revised

4.3.2 Influence of type of previous THA revision

The re-revision rate after two years accounts for 11.8% (CI 10.2–13.7%). After eleven years the revision rate rises to 18.3% (CI 15.5–21.6%). The lowest revision rate is seen after complete revisions, including revision of both stem and cup. Revision

of the cup or stem alone and re-implantations have similar revision rates around 13% at two years and 22% at eleven years (for precise numbers consult **Figure 4.21**). However, head and inlay exchange have a significantly higher revision rate at two years (19.4%, CI 17.8–21.2%) that continues until eight years. The main reason for re-revision after isolated head and inlay exchange at one year is infection with a re-revision

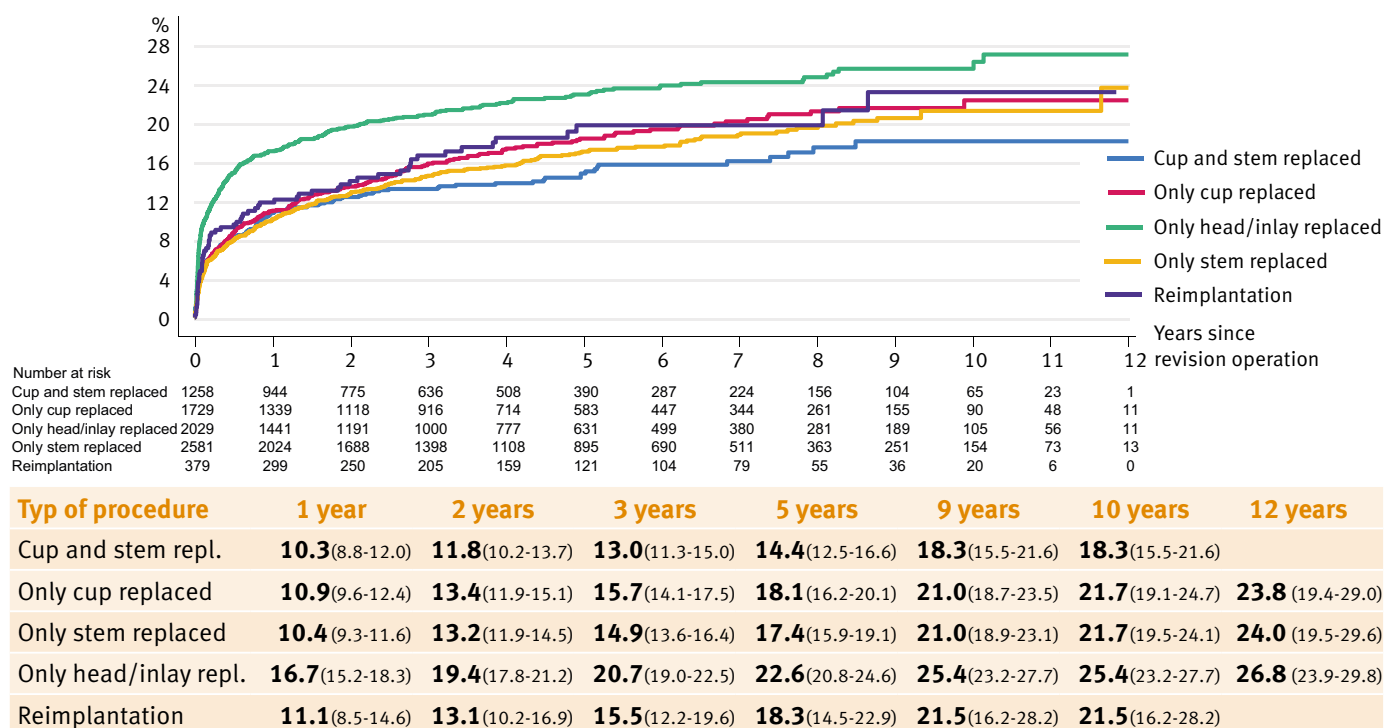


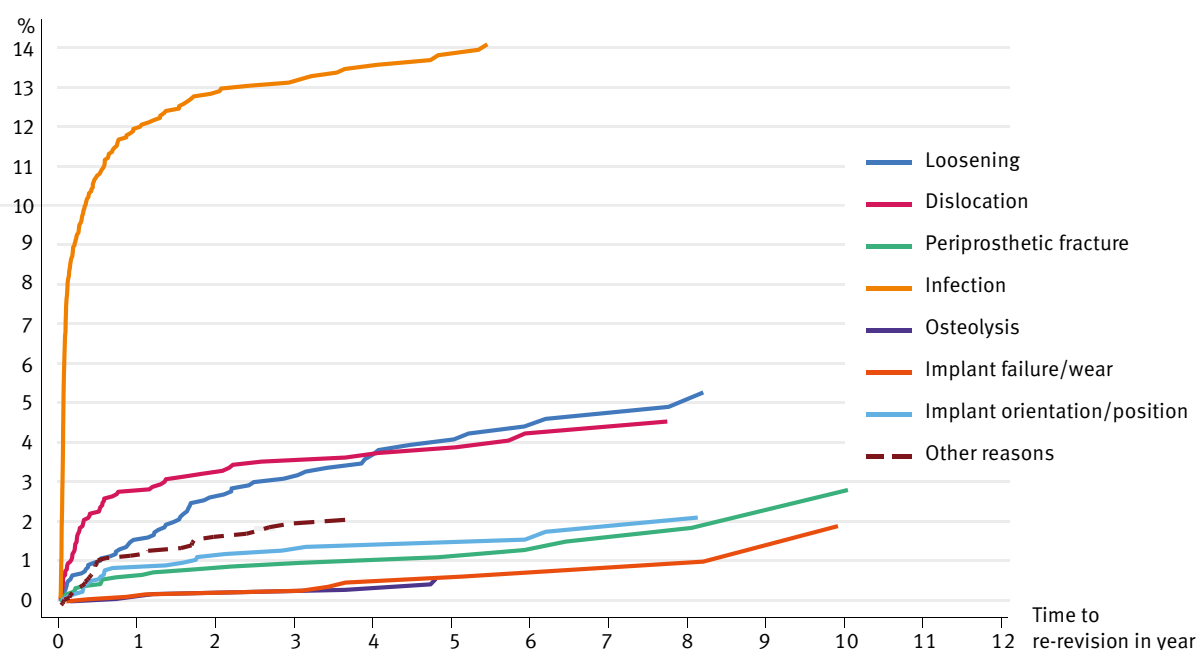
Figure 4.21

Estimated failure rates after revision of total hip arthroplasty (primary and secondary OA): types of procedures

Time since revision, 2012–2024. Start point of analysis: first registered component revision in SIRIS that meets the inclusion criteria.

End point of analysis: next registered component revision

rate of 11.9% (CI 10.6-13.3%) (Figure 4.22). Head and inlay exchange mainly is performed in prosthetic joint infections as DAIR procedure (Debridement, Antibiotics and Implant Retention). These findings indicate that this concept is not successful in every 10th DAIR procedure for acute infections.



Diagnoses	1 year	2 years	3 years	5 years	9 years	10 years	12 years
Loosening	1.6 (1.1-2.3)	2.7 (2.0-3.5)	3.1 (2.4-4.1)	4.0 (3.1-5.1)	5.3 (4.0-7.0)	5.3 (4.0-7.0)	5.3 (4.0-7.0)
Dislocation	2.8 (2.2-3.6)	3.3 (2.6-4.1)	3.6 (2.8-4.5)	3.8 (3.0-4.8)	4.6 (3.5-5.9)	4.6 (3.5-5.9)	4.6 (3.5-5.9)
Periprosthetic fracture	0.7 (0.4-1.1)	0.8 (0.5-1.4)	0.9 (0.6-1.5)	1.2 (0.7-1.9)	1.9 (1.1-3.2)	1.9 (1.1-3.2)	2.8 (1.3-6.0)
Infection	11.9(10.6-13.3)	12.8(11.5-14.3)	13.1(11.7-14.6)	13.8(12.3-15.4)	14.0(12.6-15.7)	14.0(12.6-15.7)	14.0(12.6-15.7)
Osteolysis	0.2 (0.1-0.5)	0.2 (0.1-0.6)	0.2 (0.1-0.6)	0.6 (0.3-1.4)	0.6 (0.3-1.4)	0.6 (0.3-1.4)	0.6 (0.3-1.4)
Implant failure /wear	0.2 (0.1-0.5)	0.2 (0.1-0.6)	0.2 (0.1-0.6)	0.5 (0.2-1.1)	1.0 (0.4-2.4)	1.0 (0.4-2.4)	1.9 (0.7-5.3)
Implant orientation / position	0.9 (0.6-1.4)	1.2 (0.8-1.8)	1.3 (0.9-2.0)	1.4 (0.9-2.1)	2.2 (1.3-3.5)	2.2 (1.3-3.5)	2.2 (1.3-3.5)
Other reasons	1.3 (0.9-1.9)	1.8 (1.3-2.5)	2.1 (1.5-2.9)	2.2 (1.6-3.1)	2.2 (1.6-3.1)	2.2 (1.6-3.1)	2.2 (1.6-3.1)

Figure 4.22

Cumulative incidence rates for different re-revision diagnoses after head/inlay changes

Time since operation, 2012–2024, % of implants re-revised

4.3.3 Implant selection and impact on re-revision

Not only the type of revision but also the choice of the implant influences the re-revision rate. Often the choice of the implant is limited by the underlying pathology. Therefore, the revision rates may be biased, and interpretations must be carefully made. As a general indication what implant should be used, the presented data may be helpful.

The use of primary uncemented stems for a first-time revision comes with the lowest revision rate of 8.9% (CI 7.5-10.4%) at one year. Uncemented revision stems have a slightly higher revision rate over the entire period of 11 years. Primary cemented stems have initially a lower revision rate, which then increases to 25.8% (CI 22.6-29.3%) at eleven years. The difference is statistically significant (Figure 4.23).

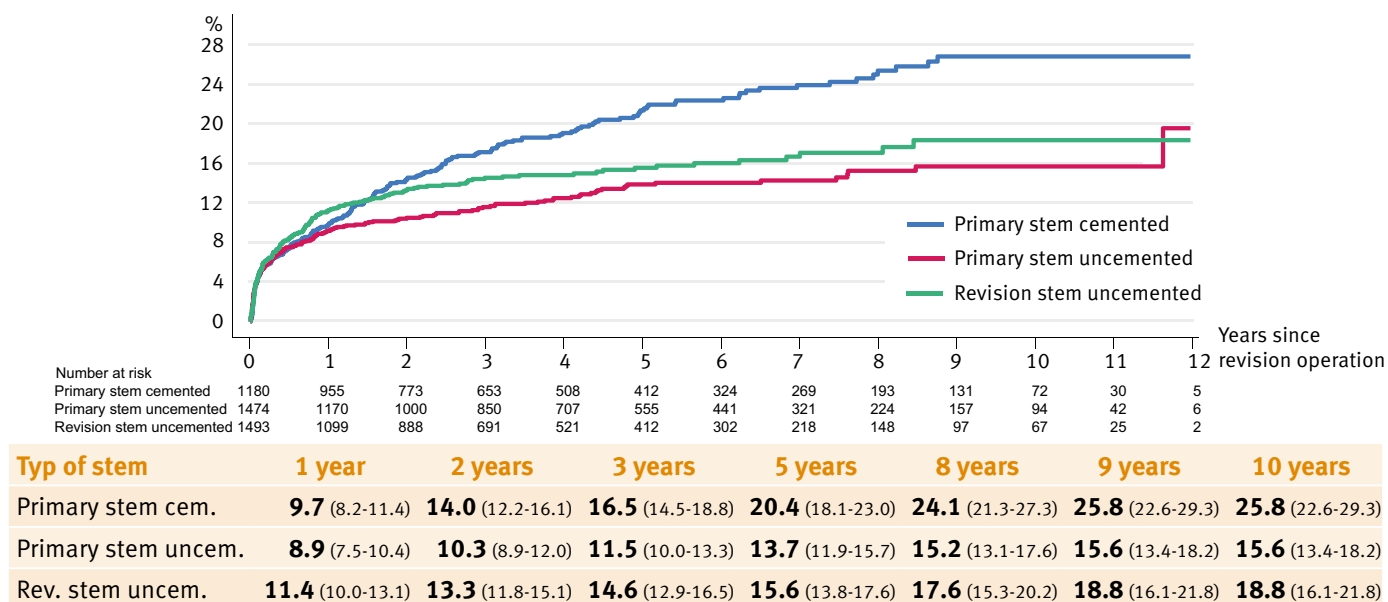


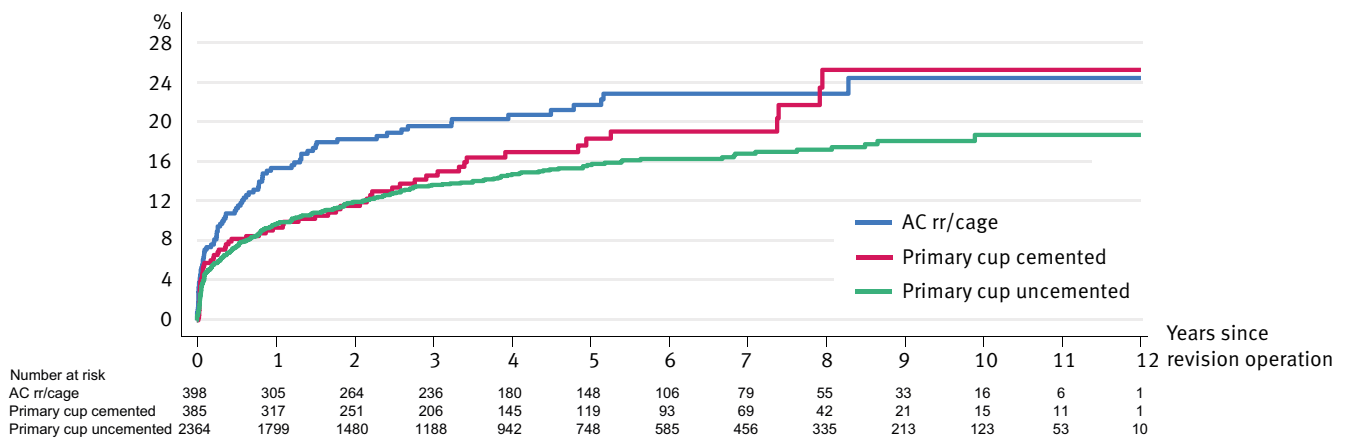
Figure 4.23

Estimated failure rates after revision of total hip arthroplasty (primary and secondary OA): types of stems used in revision

Time since revision, 2012–2024. Start point of analysis: first registered component revision in SIRIS that meets the inclusion criteria.

End point of analysis: next registered component revision.

On the acetabular side, revisions with primary uncemented cups yields the lowest revision rates (9.4%, CI 8.3-10.6%) at one year, whereas reconstruction rings come with the highest revision rates after one year (14.3%, CI 11.5-17.9%) (Figure 4.24).



Typ of cup	1 year	2 years	3 years	5 years	8 years	10 years	12 years
Primary cup cemented	9.6 (7.2-12.7)	11.7 (9.0-15.1)	14.7 (11.6-18.6)	18.0 (14.3-22.6)	24.1 (18.2-31.4)	24.1 (18.2-31.4)	
Primary cup uncem.	9.4 (8.3-10.6)	11.6 (10.3-12.9)	13.4 (12.1-14.9)	15.2 (13.7-16.9)	16.8 (15.1-18.7)	18.2 (16.0-20.7)	18.2 (16.0-20.7)
AC rr/cage	14.3 (11.5-17.9)	17.3 (14.1-21.1)	18.4 (15.1-22.3)	20.3 (16.7-24.5)	23.0 (18.7-28.0)	24.3 (19.5-30.0)	

Figure 4.24

Estimated failure rates after revision of total hip arthroplasty (primary and secondary OA): types of cups used in revision

Time since revision, 2012–2024. Start point of analysis: first registered component revision in SIRIS that meets the inclusion criteria.

End point of analysis: next registered component revision.

4.4 Results of implants in total hip arthroplasty

One of the key elements of an arthroplasty registry is to analyse the performance of implants regarding revision rates over time. While short-term results largely reflect a surgeon's or a hospital's performance, long-term results depend more on the design and quality of the implants. A total hip replacement comprises at least three components: the stem, the cup, and the head. Considering the modularity of the cup, including dual mobility systems, it is sensible to focus investigations on combinations in current use and to compare those with each other as it could be that a cup works well with one stem but poorly with another – and vice versa. THA performed for primary and for secondary OA are considered separately. The internationally recognised benchmark considers only primary OA. The reader is reminded to consider statistical precision of the results. A single revision weighs much more in a small group than in a large group. Hence, there is always a trade-off between statistical stability and the necessity to identify possible low-volume outliers. Methodological details are provided in Chapter 2 Methods.

Currently, SIRIS documents 169 different brands of stems (including all currently identified sub-variants), of which 36 were implanted less than 10 times, while another 32 were used in 10 to 49 cases only. There were 136 different brands of cups, of which 26 were implanted less than 10 times and another 25 were used in 10 to 49 cases. This resulted in 1,377 different stem/cup combinations, of which only 275 were used in more than 50 cases. It is noteworthy that almost half of all recognised combinations (47.9%) were registered less than 5 times. Yet, this remarkable diversity accounted for 0.46% of all THA. In contrast, the ten most used implant combinations covered 51.5% of all THA. The current 4-year moving window covers 80 combinations with more than 50 cases. This is a significant reduction from the 108 combinations in 2024.

A so-called case concentration score (CCS) was introduced since the annual report 2022. It indicates the percentage of implantations performed by the main user hospital service. A higher value signifies an increased likelihood of bias due to local effects induced by a single provider unit. Hence, a share of >50% would suggest that reported results are likely dominated by data from one hospital service while a score of 100% indicates that the implant is used in one hospital only.

Currently documented in SIRIS:
169 brands of hip stems
and
136 brands of hip cups

This resulted in 1,377
stem/cup combinations

only 275
combinations are
used in more
than 50 cases

4.4.1 Twelve-year revision rates

Uncemented combinations for primary OA

The register now can provide an overview of the 12-year performance of implant combinations. The revision rates are shown for 1, 2-, 3-, 5-, 6-, 8-, 9-, 10- and 12-years follow-up.

Table 4.16 shows the stem/cup combinations used in 75% of the uncemented THA performed for primary OA and their evolution between 2019 and 2024. This list is shorter by two combinations than in the 2024 report, which indicates a concentration to fewer implant combinations. **Table 4.17** shows the revision

rates since 2012 for THA conducted for primary OA, whereby only uncemented stem/cup combinations with N>500 are presented. At 12 years, the average revision rate for all uncemented stem/cup combinations was 5.3% (CI 5.1 – 5.5%). The lowest revision rate at 12 years was documented for the combination SL-plus/R3 with 2.4% (CI 1.6 – 3.4%). The highest revision rate was found in the combination Exception/Avantage with 9.2% (CI 7.1 – 12.0%). Interestingly, both had a rather high case concentration score (CCS) of 64 and 78 respectively, indicating the dominant use in one hospital only.

Stem component	Cup component	2019	2020	2021	2022	2023	2024	2019 – 2024
Actis	Pinnacle	119	185	222	406	538	606	2,076
Amistem-P	Versafitcup trio/ccl.	383	1,183	1,229	1,213	1,149	1,032	6,189
Avenir	Allofit	1,143	1,040	714	619	518	542	4,576
Corail	Pinnacle	1,158	1,222	1,135	936	848	638	5,937
Corail collared	Novae TH/Bi-Mentum	45	97	343	357	519	521	1,882
Corail collared	Pinnacle	1,406	1,610	2,004	2,291	2,361	2,421	12,093
Fitmore	Allofit	527	561	617	681	850	1,097	4,333
Fitmore	Fitmore	622	625	596	607	541	230	3,221
Optimys	RM pressfit vitamys	1,845	2,107	2,488	3,045	3,444	3,780	16,709
Polarstem	Polarcup	189	209	173	217	205	196	1,189
Polarstem	R3	687	766	809	935	1,006	1,237	5,440
Quadra-H	Versafitcup trio/ccl.	983	752	482	182	16	0	2,415
Quadra-P	Versafitcup trio/ccl.	33	244	571	866	1,078	1,179	3,971
Twinsys	RM pressfit vitamys	409	397	416	424	458	492	2,596
other combinations	-	4,870	3,573	3,728	3,637	3,668	3,732	23,208
Total		14,419	14,571	15,527	16,416	17,199	17,703	95,835

Table 4.16

Top 75% of primary total hip arthroplasty uncemented combinations (primary OA) 2019 – 2024

Stem component	Cup component	Total CCS*		1 year	3 years	5 years	10 years	12 years
		N		(95% CI)	(95% CI)	(95% CI)	(95% CI)	(95% CI)
Actis	Pinnacle	2,104	29	1.3 (0.9-1.9)	1.8 (1.2-2.6)	1.8 (1.2-2.6)		
Alloclassic	Fitmore	729	66	2.2 (1.4-3.6)	4.2 (2.9-5.9)	4.9 (3.5-6.8)	6.0 (4.5-8.2)	6.0 (4.5-8.2)
Amistem-H	Versafitcup trio/ccl.	7,350	15	2.0 (1.7-2.3)	3.1 (2.8-3.6)	4.3 (3.9-4.8)	7.2 (6.6-7.9)	8.1 (7.3-8.9)
Amistem-H collared	Versafitcup trio/ccl.	556	100	0.9 (0.4-2.2)	1.8 (1.0-3.4)	1.8 (1.0-3.4)	2.7 (1.6-4.7)	
Amistem-H prox coating	Versafitcup trio/ccl.	3,250	12	2.2 (1.7-2.7)	3.0 (2.4-3.6)	3.4 (2.8-4.1)		
Amistem-P	Mpact	690	45	2.9 (1.8-4.5)	3.6 (2.3-5.7)	5.9 (3.0-11.6)		
Amistem-P	Versafitcup trio/ccl.	6,191	14	2.4 (2.1-2.9)	3.1 (2.6-3.5)	3.7 (3.0-4.4)		
Avenir	Alloclassic	591	68	1.9 (1.0-3.3)	2.4 (1.4-4.0)	2.8 (1.7-4.5)	3.7 (2.4-5.8)	4.6 (2.7-7.5)
Avenir	Allofit	11,364	11	2.0 (1.8-2.3)	2.7 (2.4-3.0)	3.0 (2.7-3.4)	4.0 (3.5-4.4)	4.2 (3.7-4.7)
Avenir	Fitmore	2,765	16	3.2 (2.6-3.9)	4.0 (3.3-4.8)	4.2 (3.5-5.0)	4.9 (4.0-5.8)	5.4 (4.2-6.9)
CLS Spotorno	Allofit	1,583	31	2.7 (2.0-3.7)	4.0 (3.2-5.2)	4.5 (3.6-5.7)	6.0 (4.7-7.5)	6.2 (4.9-7.9)
CLS Spotorno	Fitmore	1,910	26	1.6 (1.1-2.3)	2.2 (1.6-3.0)	2.9 (2.2-3.7)	3.7 (2.8-4.9)	4.3 (3.2-5.8)
Corail	Pinnacle	13,501	12	2.2 (1.9-2.4)	3.1 (2.8-3.4)	3.7 (3.3-4.0)	5.3 (4.8-5.9)	6.0 (5.4-6.7)
Corail collared	Gyros	967	65	2.1 (1.3-3.2)	2.7 (1.9-4.0)	2.8 (2.0-4.1)	4.9 (3.1-7.7)	
Corail collared	Novae TH/Bi-Mentum	1,927	26	1.6 (1.1-2.3)	2.4 (1.7-3.5)	2.4 (1.7-3.5)		
Corail collared	Pinnacle	16,879	21	1.4 (1.3-1.6)	2.0 (1.8-2.3)	2.3 (2.1-2.6)	3.1 (2.6-3.7)	3.1 (2.6-3.7)
Exception	Avantage	1,136	78	3.4 (2.4-4.6)	4.4 (3.4-5.8)	5.3 (4.1-6.8)	7.8 (6.2-9.9)	9.2 (7.1-12.0)
Fitmore	Allofit	8,750	66	1.8 (1.5-2.1)	2.7 (2.3-3.0)	3.0 (2.6-3.4)	3.7 (3.3-4.3)	4.0 (3.5-4.6)
Fitmore	Fitmore	6,211	26	1.8 (1.5-2.2)	2.6 (2.2-3.1)	3.1 (2.6-3.5)	4.0 (3.4-4.7)	5.0 (3.5-7.3)
Fitmore	RM pressfit vitamys	1,684	84	1.3 (0.8-1.9)	2.0 (1.4-2.8)	2.2 (1.5-3.1)	2.5 (1.7-3.8)	
H-Max S	Delta TT	531	23	1.5 (0.8-3.0)	1.9 (1.1-3.6)	2.4 (1.4-4.2)	6.0 (3.6-9.9)	
Individual/custom hip	April ceramic	1,439	24	1.9 (1.3-2.7)	2.9 (2.1-4.0)	3.5 (2.6-4.7)	3.7 (2.7-5.1)	4.6 (2.9-7.0)
Optimys	Anexys	577	25	1.4 (0.7-2.9)	1.9 (1.0-3.4)	1.9 (1.0-3.4)		
Optimys	RM pressfit	819	19	2.6 (1.7-4.0)	2.9 (1.9-4.4)	3.4 (2.3-5.0)	4.2 (2.7-6.3)	
Optimys	RM pressfit vitamys	24,591	9	1.8 (1.7-2.0)	2.3 (2.1-2.5)	2.5 (2.3-2.7)	2.8 (2.5-3.1)	2.8 (2.5-3.1)
Polarstem	EP-fit	823	53	3.8 (2.7-5.3)	4.5 (3.3-6.2)	5.0 (3.7-6.7)	6.4 (4.6-8.7)	6.4 (4.6-8.7)
Polarstem	Polarcup	2,573	74	2.2 (1.7-2.9)	2.4 (1.9-3.1)	2.5 (1.9-3.2)	3.0 (2.3-3.9)	3.8 (2.6-5.3)
Polarstem	R3	9,240	62	1.2 (1.0-1.4)	1.7 (1.4-2.0)	1.9 (1.6-2.3)	2.6 (2.1-3.1)	2.7 (2.2-3.4)
Quadra-H	Mpact	540	43	2.2 (1.3-3.9)	2.8 (1.7-4.6)	3.0 (1.9-4.9)		
Quadra-H	Versafitcup trio/ccl.	7,219	19	2.0 (1.7-2.4)	3.0 (2.6-3.4)	3.8 (3.3-4.2)	6.7 (5.9-7.6)	7.8 (6.6-9.1)
Quadra-P	Mpact	789	59	2.0 (1.2-3.3)	2.8 (1.7-4.6)			
Quadra-P	Versafitcup trio/ccl.	3,971	21	1.5 (1.2-2.0)	1.9 (1.5-2.5)	2.3 (1.7-3.1)		
SBG	R3	1,668	42	1.3 (0.9-2.0)	1.9 (1.3-2.7)	2.4 (1.7-3.3)	3.9 (2.6-5.7)	
SL-plus MIA	EP-fit	1,254	30	2.0 (1.4-2.9)	2.2 (1.6-3.2)	2.6 (1.8-3.6)	2.9 (2.1-4.2)	2.9 (2.1-4.2)
SL-plus MIA	HI	882	43	2.1 (1.3-3.2)	3.9 (2.8-5.5)	5.2 (3.9-7.0)	9.3 (7.0-12.3)	10.2 (7.5-14.0)
SL-plus MIA	R3	2,024	64	1.0 (0.7-1.6)	1.4 (0.9-2.0)	1.5 (1.0-2.1)	2.2 (1.5-3.1)	2.4 (1.6-3.4)
SPS evolution	April ceramic	1,712	40	5.0 (4.1-6.2)	6.5 (5.4-7.8)	6.8 (5.7-8.2)	7.3 (6.1-8.8)	7.8 (6.3-9.5)
Tri-Lock	Pinnacle	768	65	1.3 (0.7-2.4)	2.8 (1.8-4.3)	3.3 (2.2-4.8)	3.6 (2.5-5.3)	3.6 (2.5-5.3)
Twinsys	RM pressfit vitamys	5,040	13	2.3 (2.0-2.8)	3.0 (2.6-3.5)	3.4 (2.9-4.0)	5.1 (4.3-6.0)	5.9 (4.8-7.2)
other combinations	-	23,126		2.6 (2.4-2.8)	3.8 (3.5-4.1)	4.6 (4.3-4.9)	6.5 (6.1-7.0)	7.6 (7.0-8.3)
CH average for group				2.0 (1.9-2.1)	2.8 (2.7-2.9)	3.3 (3.2-3.4)	4.7 (4.6-4.9)	5.3 (5.1-5.5)

Table 4.17

Long term evaluation: Failure rates of primary total hip arthroplasty uncemented combinations (primary OA)

Four implant combinations with an elevated revision rate were detected. The corresponding KM estimates are illustrated in **Figure 4.25**. Exception/Avantage and Polarstem/EP fit already appeared in 2023 as implant combinations with elevated revision rates. Quadra-H/Versafitcup trio/ccl, which appeared as outlier in 2023, improved and no longer had an elevated revision rate. The combination Alloclassic/Fitmore and Fitmore/Allofit reached the status of elevated revision rate in 2024. However, Amistem-H was only implanted until 2019 and no longer is in ac-

tive use and has been replaced by new evolutions of the stem. There are two implant combinations with outlier level revision rates during the 12 years observation period. SL plus MIA/Hi and April ceramic, which already had an outlier status at 2 years. After a steep early rise of the cumulative revision rate, the curve flattened over the subsequent years, although the outlier boundary was still exceeded at 5 years follow-up. The revision rate of this combination amounts to 7.8% (CI 6.3 – 9.6%) at 12 years (**Figure 4.26**).

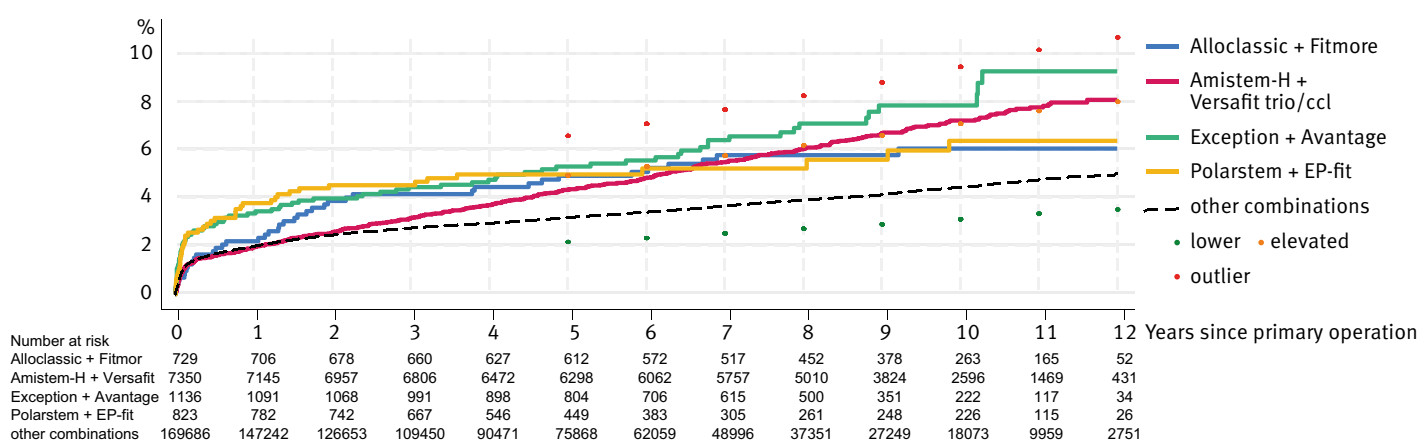


Figure 4.25

Implant combinations with elevated long-term revision rates (primary OA, uncemented THA)

An elevated revision rate was defined as a deviation of at least 50% above the group average at any time between year 5 and year 12 (and lower bounds of the 95% confidence interval exceeding the upper bound of the group average; and at least 50 cases at risk at 5 years). The dots indicate upper and lower limits.

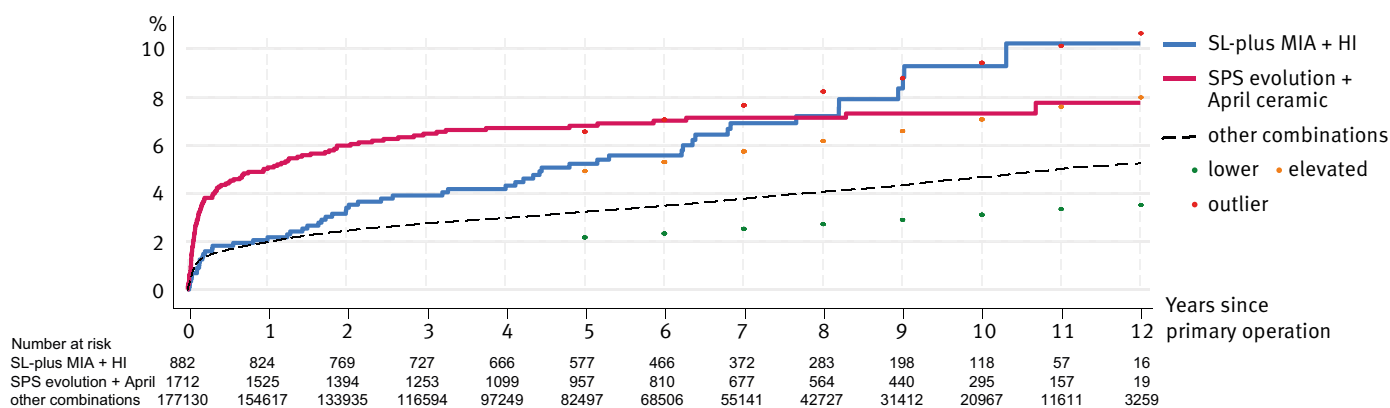


Figure 4.26

Implant combinations with long-term evaluation outlier status (primary OA, uncemented THA)

Outlier status was defined as a revision rate of twice the group average at any time between year 5 and year 12 (and lower bounds of the 95% confidence interval exceeding the upper bound of the group average; and at least 50 cases at risk at 5 years). The dots indicate upper and lower limits.

There are six uncemented implant combinations with a below-average revision rate to be reported in 2024, two more than in 2023 (Figure 4.27). The curves of these well-performing implant combinations have an almost horizontal continuation after the initial rise in the first 3 months. The combination Optimys/RM pressfit Vitamys, Polarstem/R3 and SL-plus MIA/R3 belonged already 2023 to the best performing implant combinations. In addition, Corail collared/Pin-

nacle, Fitmore/RM pressfit Vitamys and SL-plus MIA/EP fit are newly identified as implants with a below average revision rate.

The KM estimate of the cumulative revision risks for all other uncemented implant combinations are shown in Figures 4.28. These curves run between the upper and lower limits corresponding to the elevated revision risk at 150%, respectively the below-average revision risk at 66% from the groups' average.

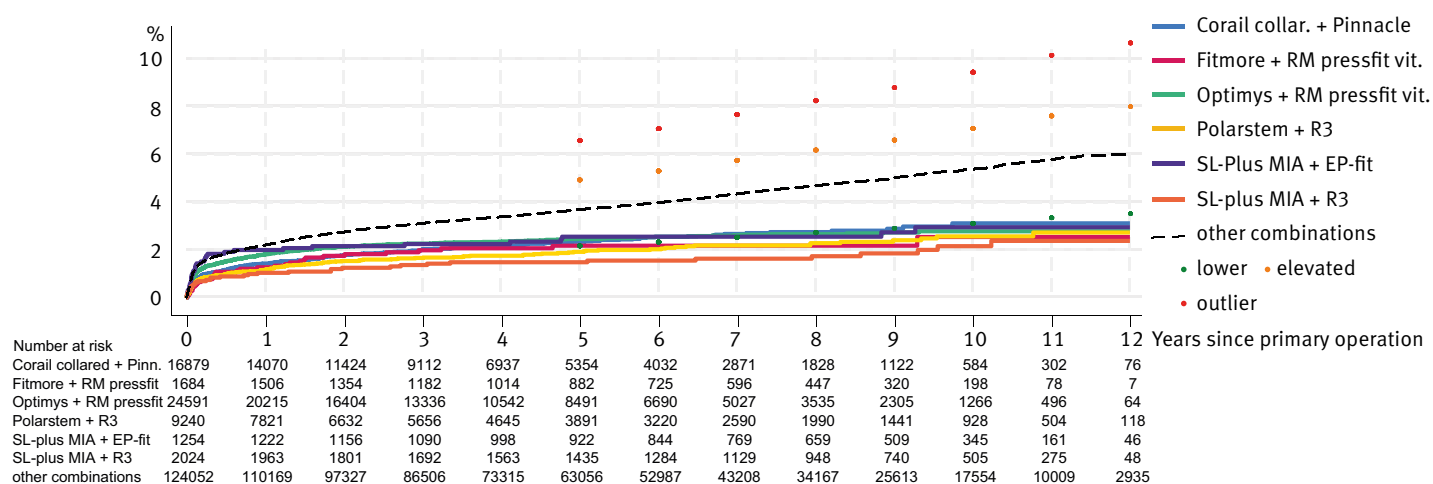


Figure 4.27

Implant combinations with below-average long-term revision rates (primary OA, uncemented THA)

Below-average was defined as an 9-year/10-year revision rate of up to 66% of the group average (and upper bounds of the 95% confidence interval staying below the lower bound of the group average; and at least 25 cases at risk at 9 years/10 years). The dots indicate upper and lower limits.

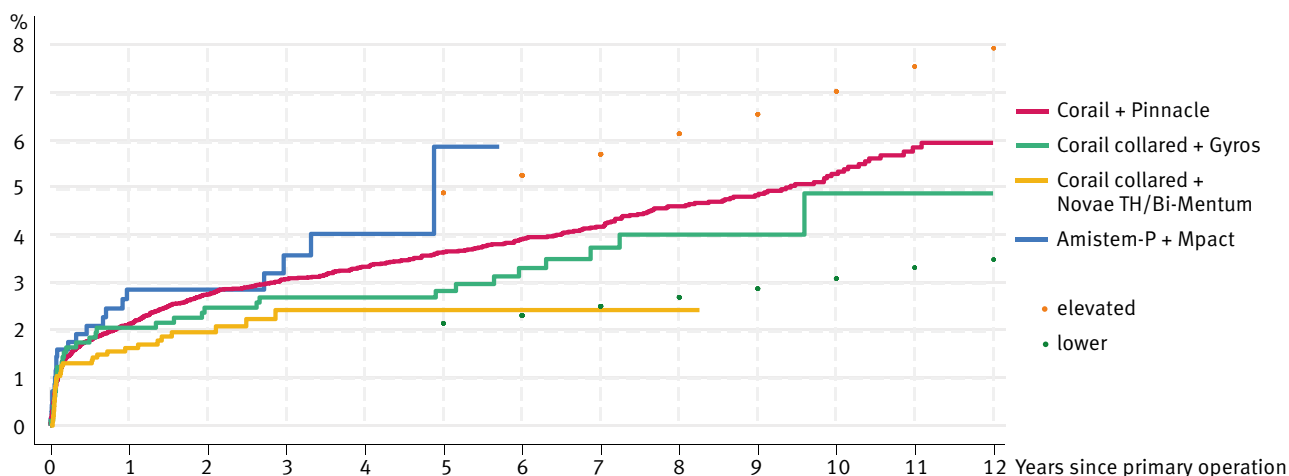
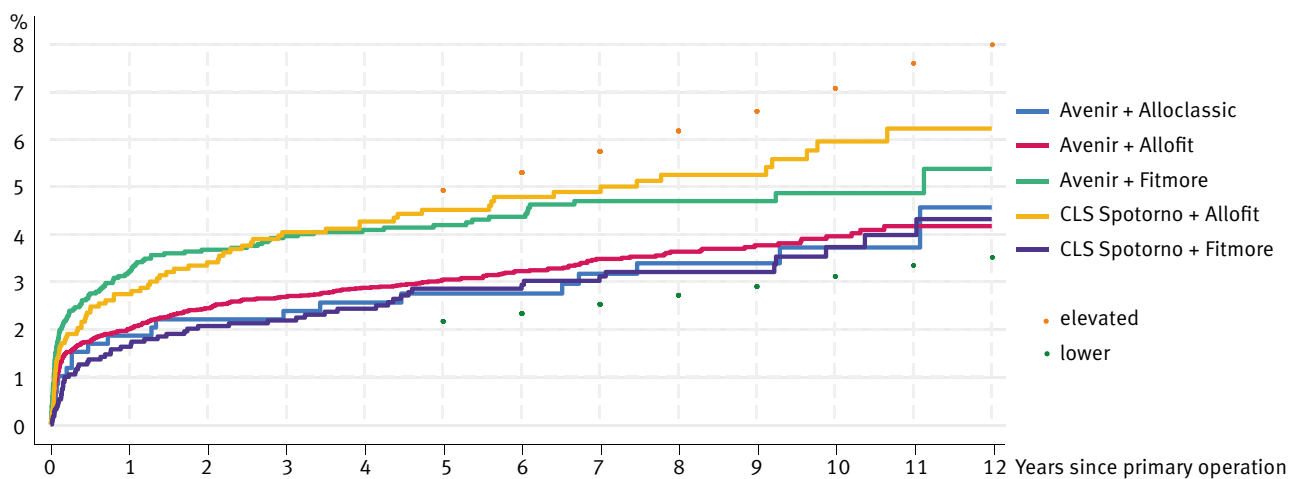
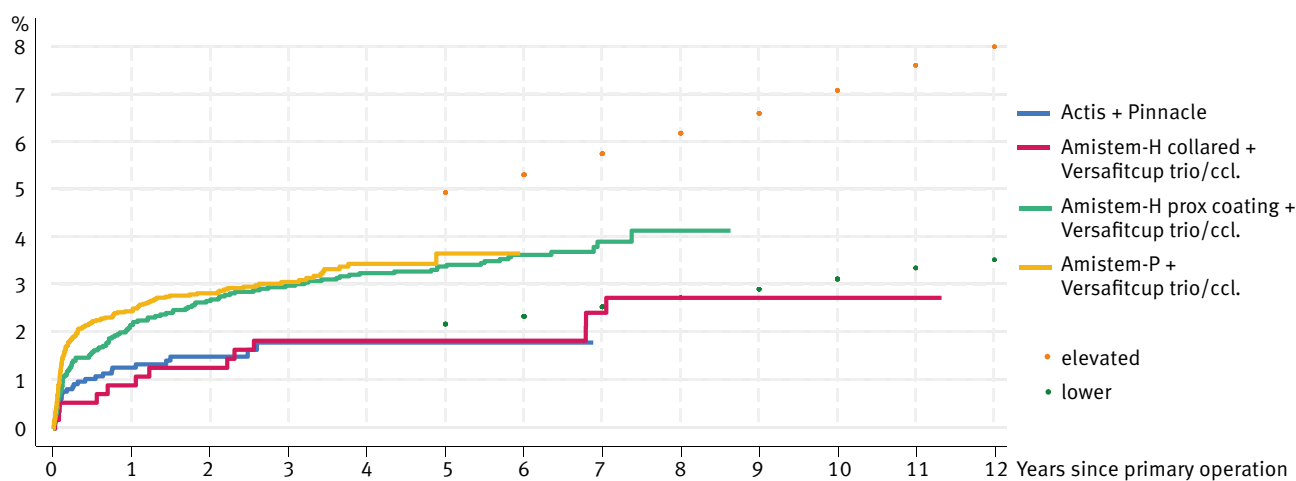


Figure 4.28 – Part one

All remaining implant combinations with average revision risks (primary OA, uncemented THA)

Also showing upper and lower limits (corresponding to elevated and below-average version risk at 150% and 66% of the group average respectively).

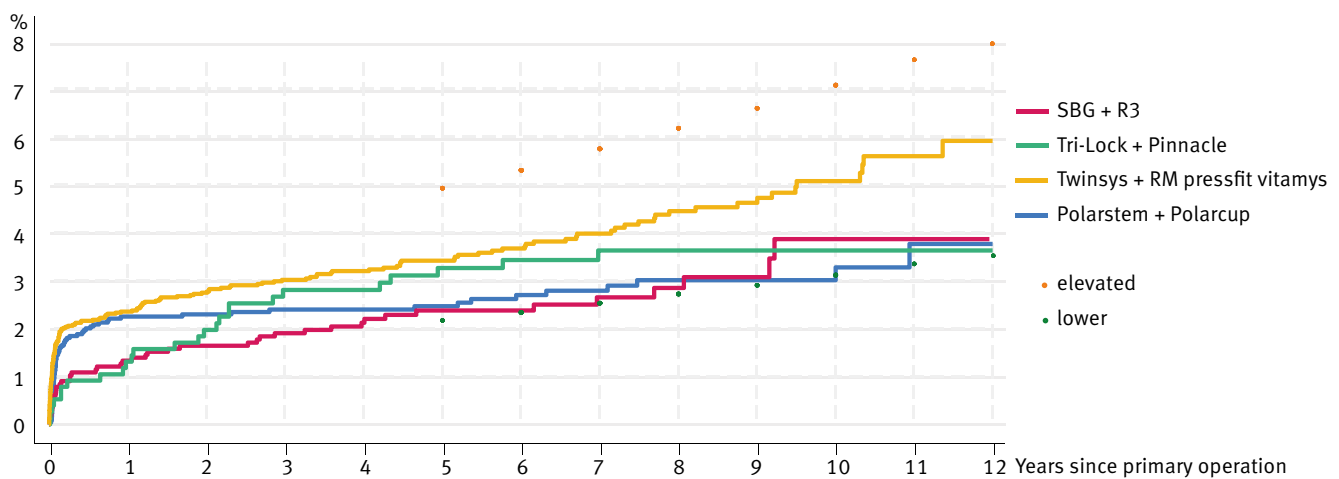
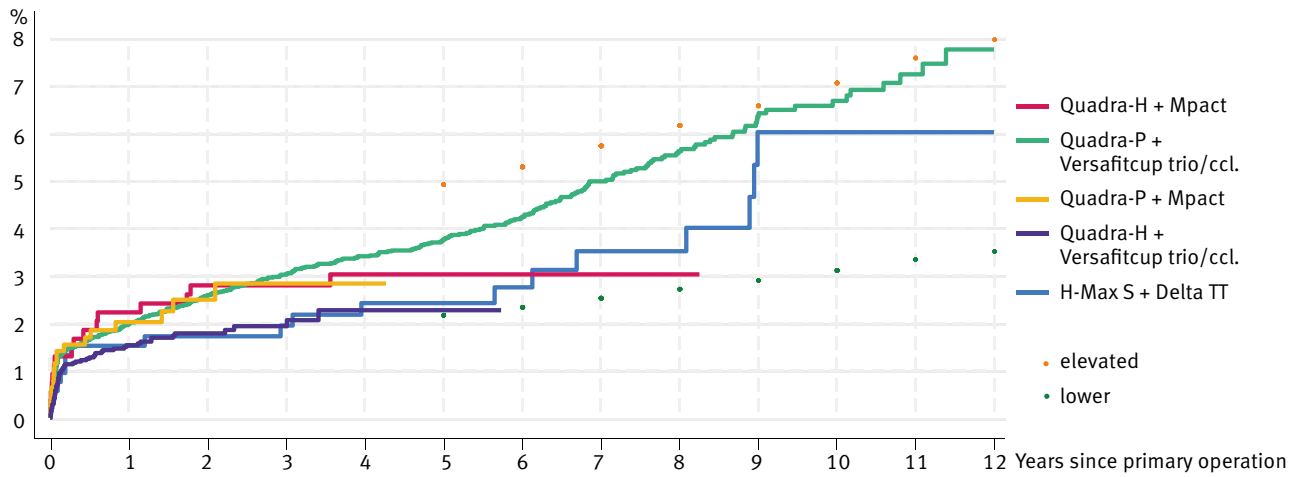
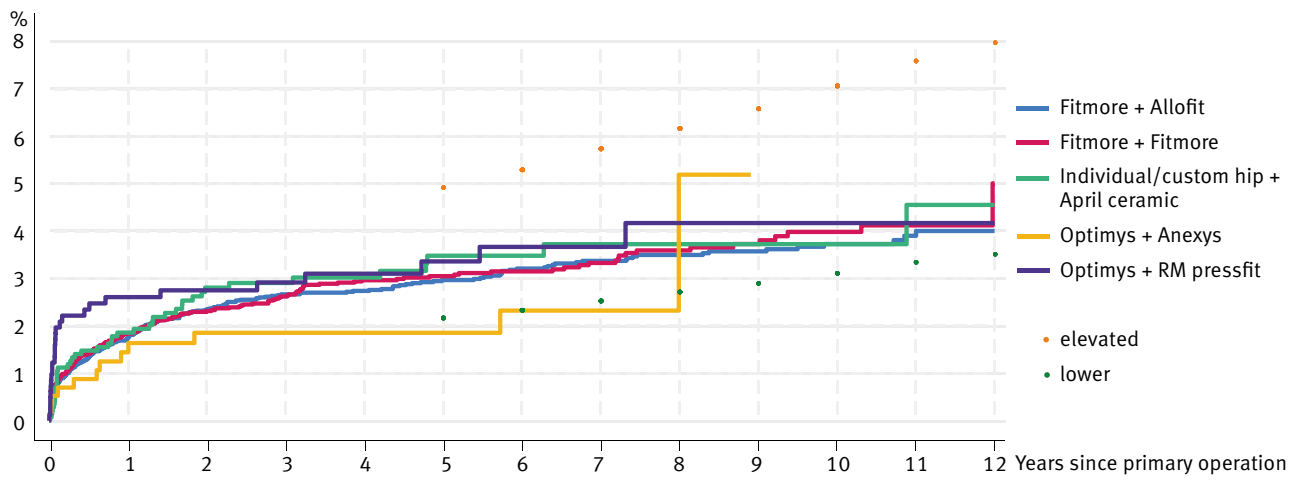


Figure 4.28 – Part two

All remaining implant combinations with average revision risks (primary OA, uncemented THA)

Also showing upper and lower limits (corresponding to elevated and below-average version risk at 150% and 66% of the group average respectively).

Most cup systems are modular allowing the use of different bearings. For the most commonly used 75% of implant combinations, the revision rates were calculated depending on the bearing surface. The results, sorted by implant combination, are shown in **Table 4.18**. Although there were differences between

the various bearings, these did not reach statistical significance. Overall, the pattern is not uniform as shown by examples in which CoXLPE had more revisions than MoXLPE.

Stem component	Cup component	Total N	Bearing surface	1 year (95% CI)	3 years (95% CI)	5 years (95% CI)	10 years (95% CI)	12 years (95% CI)
Actis	Pinnacle	621	CoC	1.2 (0.6-2.5)	1.5 (0.7-3.0)	1.5 (0.7-3.0)		
Actis	Pinnacle	1,457	CoXLPE	1.3 (0.8-2.1)	2.0 (1.2-3.1)	2.0 (1.2-3.1)		
Alloclassic	Fitmore	681	CoXLPE	1.9 (1.1-3.3)	3.8 (2.6-5.6)	4.6 (3.3-6.5)	5.9 (4.3-8.0)	5.9 (4.3-8.0)
Amistem-H	Versafitcup trio/ccl.	2,421	CoC	1.6 (1.1-2.2)	2.8 (2.2-3.5)	3.7 (3.0-4.5)	5.7 (4.8-6.8)	6.0 (5.0-7.2)
Amistem-H	Versafitcup trio/ccl.	3,159	CoXLPE	1.5 (1.1-2.0)	2.8 (2.3-3.4)	3.9 (3.3-4.6)	6.6 (5.7-7.6)	7.7 (6.7-9.0)
Amistem-H	Versafitcup trio/ccl.	1,292	MoXLPE	2.9 (2.1-3.9)	4.1 (3.1-5.3)	5.8 (4.7-7.3)	9.5 (7.9-11.5)	10.1 (8.3-12.3)
Amistem-H collared	Versafitcup trio/ccl.	546	CoC	0.9 (0.4-2.2)	1.9 (1.0-3.4)	1.9 (1.0-3.4)	2.8 (1.6-4.8)	
Amistem-H prox coating	Versafitcup trio/ccl.	1,390	CoC	1.5 (1.0-2.3)	2.2 (1.6-3.2)	2.6 (1.9-3.6)		
Amistem-H prox coating	Versafitcup trio/ccl.	1,339	CoXLPE	2.7 (2.0-3.7)	3.5 (2.6-4.6)	3.8 (2.9-5.0)		
Amistem-P	Versafitcup trio/ccl.	3,087	CoC	1.8 (1.4-2.4)	2.4 (1.9-3.0)	2.7 (2.1-3.4)		
Amistem-P	Versafitcup trio/ccl.	2,420	CoXLPE	2.5 (1.9-3.2)	3.0 (2.3-3.8)	3.9 (2.9-5.4)		
Amistem-P	Versafitcup trio/ccl.	568	MoXLPE	5.3 (3.7-7.5)	6.8 (4.9-9.5)	7.3 (5.3-10.1)		
Avenir	Allofit	8,620	CoXLPE	1.8 (1.6-2.1)	2.5 (2.1-2.8)	2.8 (2.5-3.2)	3.7 (3.2-4.2)	3.9 (3.4-4.5)
Avenir	Allofit	2,283	MoXLPE	2.3 (1.8-3.0)	3.1 (2.5-3.9)	3.3 (2.7-4.2)	4.0 (3.2-5.1)	4.3 (3.3-5.5)
Avenir	Fitmore	2,080	CoXLPE	3.3 (2.6-4.1)	3.9 (3.2-4.9)	4.3 (3.5-5.3)	5.0 (4.1-6.3)	5.0 (4.1-6.3)
Avenir	Fitmore	566	MoXLPE	2.7 (1.6-4.4)	3.7 (2.5-5.7)	3.7 (2.5-5.7)	4.3 (2.9-6.4)	5.8 (3.2-10.3)
CLS Spotorno	Allofit	1,203	CoXLPE	3.2 (2.4-4.4)	4.6 (3.6-6.0)	5.0 (3.9-6.5)	6.8 (5.3-8.8)	7.4 (5.6-9.7)
CLS Spotorno	Fitmore	932	CoXLPE	1.9 (1.2-3.1)	2.5 (1.7-3.8)	3.5 (2.5-5.0)	4.1 (2.9-5.7)	4.8 (3.2-7.2)
CLS Spotorno	Fitmore	952	MoXLPE	1.4 (0.8-2.3)	1.9 (1.2-3.0)	2.3 (1.5-3.5)	3.5 (2.2-5.4)	4.0 (2.5-6.2)
Corail	Pinnacle	1,859	CoC	2.1 (1.5-2.9)	3.5 (2.7-4.4)	4.4 (3.6-5.5)	5.9 (4.8-7.3)	6.8 (5.5-8.5)
Corail	Pinnacle	10,557	CoXLPE	2.2 (1.9-2.5)	3.1 (2.7-3.4)	3.5 (3.1-3.9)	5.1 (4.5-5.7)	5.6 (4.9-6.4)
Corail	Pinnacle	917	MoXLPE	1.9 (1.2-3.0)	2.6 (1.7-3.9)	3.0 (2.0-4.4)	6.6 (4.5-9.4)	7.9 (5.0-12.2)
Corail collared	Gyros	816	CoCPE	2.1 (1.3-3.3)	2.8 (1.9-4.2)	3.0 (2.0-4.5)	5.1 (3.1-8.2)	
Corail collared	Novae TH/Bi-Mentum	1,554	CoCPE	1.8 (1.2-2.6)	2.6 (1.8-3.8)	2.6 (1.8-3.8)		
Corail collared	Pinnacle	2,283	CoC	1.9 (1.4-2.5)	2.5 (1.9-3.3)	3.0 (2.4-3.9)	3.9 (2.9-5.3)	3.9 (2.9-5.3)
Corail collared	Pinnacle	14,002	CoXLPE	1.3 (1.1-1.5)	1.8 (1.6-2.1)	2.1 (1.9-2.4)	2.8 (2.3-3.4)	2.8 (2.3-3.4)
Exception	Avantage	904	CoXLPE	3.5 (2.5-5.0)	4.5 (3.3-6.0)	5.3 (4.0-7.0)	7.4 (5.7-9.6)	9.0 (6.7-12.0)

Table 4.18 – Part one

Long term evaluation: Failure rates of primary total hip arthroplasty uncemented combinations and different bearing surfaces (primary OA)

Time since operation, 2012–2024. Please note that if reported stem-cup combinations involve multiple sub-variants, it is possible that the long-term performance of these sub-variants may be significantly different from their combined performance.

Stem component	Cup component	Total N	Bearing surface	1 year (95% CI)	3 years (95% CI)	5 years (95% CI)	10 years (95% CI)	12 years (95% CI)
Fitmore	Allofit	6,270	CoXLPE	1.5 (1.2-1.9)	2.3 (2.0-2.8)	2.6 (2.2-3.0)	3.1 (2.7-3.7)	3.5 (2.7-4.4)
Fitmore	Allofit	2,340	MoXLPE	2.2 (1.7-2.9)	3.2 (2.5-4.0)	3.6 (2.9-4.5)	4.3 (3.5-5.3)	4.6 (3.7-5.7)
Fitmore	Fitmore	3,108	CoXLPE	1.6 (1.2-2.2)	2.2 (1.7-2.8)	2.9 (2.3-3.6)	3.3 (2.5-4.2)	3.3 (2.5-4.2)
Fitmore	Fitmore	3,044	MoXLPE	2.0 (1.6-2.6)	3.0 (2.5-3.7)	3.3 (2.7-4.0)	4.4 (3.6-5.4)	5.7 (3.8-8.5)
Fitmore	RM pressfit vitamys	1,545	CoXLPE	1.1 (0.7-1.7)	1.5 (1.0-2.3)	1.6 (1.1-2.5)	2.2 (1.2-3.8)	
Individual/custom hip	April ceramic	1,410	CoC	1.8 (1.2-2.7)	2.9 (2.1-4.0)	3.5 (2.5-4.7)	3.7 (2.7-5.1)	4.6 (2.9-7.1)
Optimys	RM pressfit vitamys	24,054	CoXLPE	1.8 (1.7-2.0)	2.3 (2.1-2.5)	2.5 (2.2-2.7)	2.8 (2.5-3.1)	2.8 (2.5-3.1)
Optimys	RM pressfit	662	CoCPE	1.7 (0.9-3.0)	1.9 (1.1-3.3)	2.4 (1.4-4.1)	3.3 (1.9-5.7)	
Polarstem	Polarcup	1,351	CMoXLPE	2.4 (1.7-3.4)	2.5 (1.8-3.5)	2.5 (1.8-3.5)		
Polarstem	Polarcup	862	CoXLPE	2.2 (1.4-3.4)	2.5 (1.6-3.7)	2.6 (1.7-3.9)	3.2 (2.2-4.7)	4.1 (2.7-6.3)
Polarstem	R3	6,165	CMoXLPE	0.9 (0.7-1.1)	1.4 (1.1-1.8)	1.7 (1.4-2.2)	2.4 (1.7-3.3)	
Polarstem	R3	2,837	CoXLPE	1.8 (1.3-2.3)	2.2 (1.7-2.8)	2.4 (1.9-3.1)	2.9 (2.3-3.7)	3.1 (2.4-4.0)
Quadra-H	Versafitcup trio/ccl.	1,130	CoC	1.5 (0.9-2.4)	2.3 (1.6-3.4)	2.8 (2.0-4.0)	4.4 (3.1-6.3)	
Quadra-H	Versafitcup trio/ccl.	4,509	CoXLPE	2.1 (1.7-2.6)	3.2 (2.7-3.8)	4.0 (3.5-4.6)	7.2 (6.2-8.5)	8.6 (7.0-10.5)
Quadra-H	Versafitcup trio/ccl.	1,429	MoXLPE	2.2 (1.6-3.2)	3.0 (2.3-4.1)	3.8 (2.9-5.0)	6.5 (5.1-8.2)	6.9 (5.3-8.9)
Quadra-P	Versafitcup trio/ccl.	994	CoC	1.4 (0.8-2.4)	2.2 (1.3-3.7)	2.2 (1.3-3.7)		
Quadra-P	Versafitcup trio/ccl.	2,711	CoXLPE	1.6 (1.2-2.1)	1.8 (1.3-2.4)	2.1 (1.4-3.1)		
SBG	R3	778	CMoXLPE	1.3 (0.7-2.4)	2.2 (1.4-3.6)	3.0 (2.0-4.6)	3.7 (2.3-5.9)	
SBG	R3	857	CoC	1.4 (0.8-2.5)	1.7 (1.0-2.8)	1.8 (1.1-3.1)	4.5 (2.4-8.4)	
SL-plus MIA	EP-fit	572	CoC	2.8 (1.7-4.5)	2.8 (1.7-4.5)	3.0 (1.9-4.8)	3.4 (2.1-5.5)	3.4 (2.1-5.5)
SL-plus MIA	R3	1,981	CMoXLPE	1.0 (0.7-1.6)	1.3 (0.9-2.0)	1.5 (1.0-2.1)	2.2 (1.5-3.1)	2.4 (1.6-3.5)
SPS evolution	April ceramic	1,694	CoC	5.0 (4.1-6.2)	6.5 (5.4-7.8)	6.8 (5.7-8.2)	7.3 (6.1-8.8)	7.8 (6.3-9.6)
Tri-Lock	Pinnacle	737	CoXLPE	1.1 (0.5-2.2)	1.9 (1.1-3.2)	2.4 (1.5-3.8)	2.8 (1.8-4.3)	2.8 (1.8-4.3)
Twinsys	RM pressfit vitamys	4,920	CoXLPE	2.3 (2.0-2.8)	3.0 (2.5-3.5)	3.4 (2.9-4.0)	5.1 (4.3-6.0)	5.9 (4.8-7.3)

Table 4.18 – Part two

Long term evaluation: Failure rates of primary total hip arthroplasty uncemented combinations and different bearing surfaces (primary OA)

Time since operation, 2012–2024. Please note that if reported stem-cup combinations involve multiple sub-variants, it is possible that the long-term performance of these sub-variants may be significantly different from their combined performance.

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

Hybrid combinations for primary OA

There were 23 hybrid implant combinations, indicating an uncemented cup combined with a cemented stem, covering 75% of all primary THA for primary OA (Table 4.19).

The combinations are based on eleven stems which were combined with twelve different cups. At 12 years, the average revision rate for all hybrid stem/cup combinations was 4.9% (CI 4.5– 5.4%) (Table 4.20).

Stem component	Cup component	2019	2020	2021	2022	2023	2024	2019-2024
Amistem-C	Mpact	26	31	15	20	22	37	151
Amistem-C	Versafitcup DM	29	28	27	24	44	49	201
Amistem-C	Versafitcup trio/ccl.	206	160	181	191	163	183	1,084
Avenir (cem)	Allofit	100	95	94	96	103	197	685
Avenir (cem)	Fitmore	53	53	77	132	111	13	439
Centris	RM pressfit vitamys	32	56	65	0	0	0	153
Corail (cem)	Novae TH/Bi-Mentum	1	14	41	74	96	104	330
Corail (cem)	Pinnacle	130	151	170	189	232	232	1,104
Exeter V40	Symbol DMHA/DS evol.	0	0	25	38	40	22	125
MS-30	Allofit	48	43	66	235	224	215	831
MS-30	Fitmore	71	56	16	32	35	11	221
Original Mueller	Fitmore	30	20	19	5	16	0	90
Polarstem (cem)	Polarcup	10	15	18	18	23	15	99
Quadra-C	Versafitcup trio/ccl.	211	155	80	13	1	0	460
Quadra-P (cem)	Versafitcup DM	0	1	11	20	28	38	98
Quadra-P (cem)	Versafitcup trio/ccl.	0	10	49	71	87	65	282
Twinsys (cem)	RM pressfit	18	19	34	13	10	8	102
Twinsys (cem)	RM pressfit vitamys	198	199	283	317	417	526	1,940
Twinsys (cem)	Symbol DMHA/DS evol.	12	20	14	31	19	27	123
Weber	Allofit	49	38	31	30	29	23	200
Weber	Fitmore	180	162	148	104	39	9	642
other combinations	-	539	429	497	483	580	692	3,220
Total		1,943	1,755	1,961	2,136	2,319	2,466	12,580

Table 4.19

Top 75% of primary total hip arthroplasty hybrid combinations (primary OA) 2019 – 2024

There were no combinations with elevated mid-term revision rates, nor outliers, to be observed. Two implant combinations (Corail [cem]/Pinnacle and MS-30/Allofit) had a below-average long-term revision rate (Figure 4.29). All remaining implants were within the upper and lower limits (Figures 4.30). Some curves did run below the lower limit but were not implant combinations with below-average long-term revision rates because their confidence intervals were wide due to small numbers and overlapping with the reference group and therefore were not statistically different.

Stem component	Cup component	Total N	CCS*	1 year (95% CI)	3 years (95% CI)	5 years (95% CI)	10 years (95% CI)	12 years (95% CI)
Amistem-C	Versafitcup trio/ccl.	2,530	21	2.5 (2.0-3.2)	3.1 (2.5-3.9)	3.7 (3.0-4.5)	5.4 (4.3-6.8)	5.4 (4.3-6.8)
Avenir (cem)	Allofit	985	18	1.7 (1.1-2.8)	2.5 (1.6-3.8)	3.2 (2.1-4.9)		
Avenir (cem)	Fitmore	501	58	2.0 (1.1-3.7)	3.0 (1.8-5.0)	3.0 (1.8-5.0)		
Corail (cem)	Pinnacle	1,977	16	1.3 (0.9-1.9)	1.9 (1.3-2.6)	2.2 (1.6-3.0)	2.5 (1.8-3.5)	2.5 (1.8-3.5)
MS-30	Allofit	1,102	50	1.2 (0.7-2.1)	1.8 (1.1-2.8)	1.8 (1.1-2.8)	1.8 (1.1-2.8)	1.8 (1.1-2.8)
MS-30	Fitmore	863	52	1.3 (0.7-2.3)	1.7 (1.0-2.8)	1.8 (1.1-3.0)	3.0 (1.8-5.0)	3.0 (1.8-5.0)
Quadra-C	Versafitcup trio/ccl.	1,051	32	2.5 (1.7-3.6)	3.3 (2.4-4.6)	4.0 (2.9-5.4)	7.5 (3.7-14.9)	
Twinsys (cem)	RM pressfit vitamys	2,366	19	1.1 (0.7-1.6)	1.6 (1.1-2.2)	2.6 (1.8-3.7)	2.8 (1.9-4.1)	
Weber	Allofit	804	27	1.9 (1.1-3.1)	3.0 (2.0-4.5)	3.5 (2.4-5.1)	5.6 (3.8-8.3)	6.4 (4.2-9.6)
Weber	Fitmore	2,416	28	1.6 (1.2-2.2)	2.6 (2.0-3.3)	3.5 (2.8-4.4)	5.7 (4.6-7.2)	6.3 (5.0-8.0)
other combinations	-	9,108		2.2 (1.9-2.5)	3.1 (2.8-3.6)	3.9 (3.4-4.3)	4.9 (4.3-5.6)	5.1 (4.5-5.9)
CH average for group				1.9 (1.7-2.1)	2.7 (2.5-2.9)	3.3 (3.1-3.6)	4.7 (4.3-5.1)	4.9 (4.5-5.4)

Table 4.20

Long term evaluation: Failure rates of primary total hip arthroplasty hybrid combinations (primary OA)

Time since operation, 2012–2024. Please note that if reported stem-cup combinations involve multiple sub-variants, it is possible that the long-term performance of these sub-variants may be significantly different from their combined performance.

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

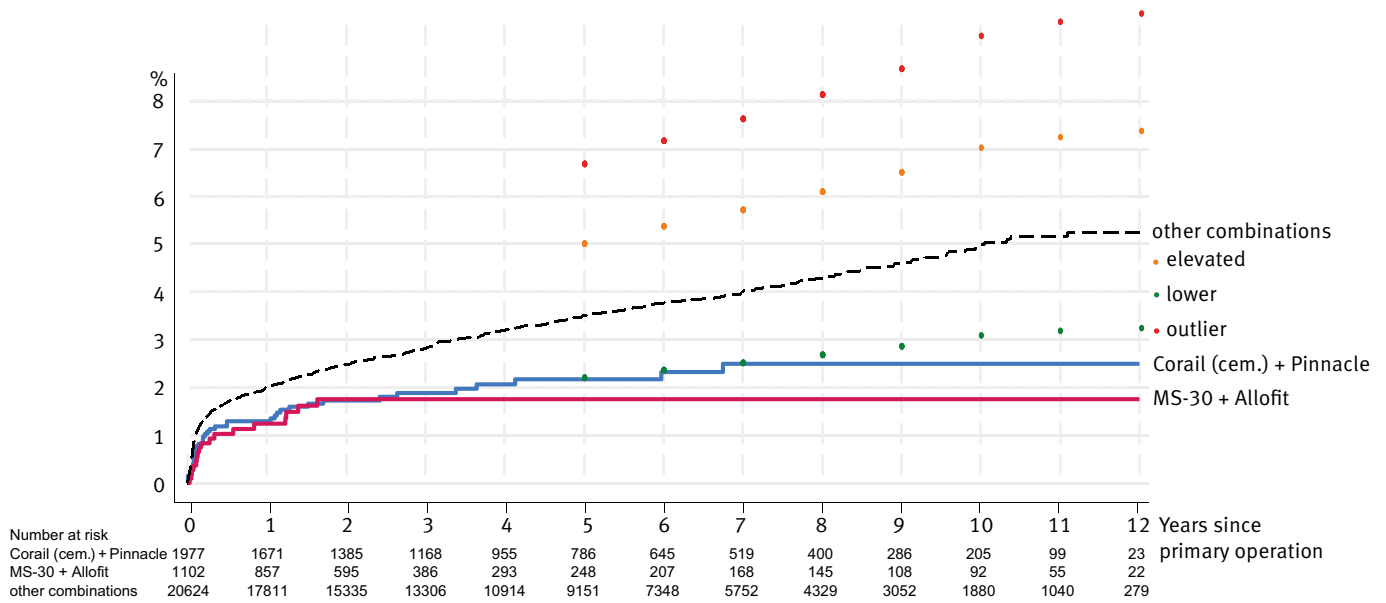
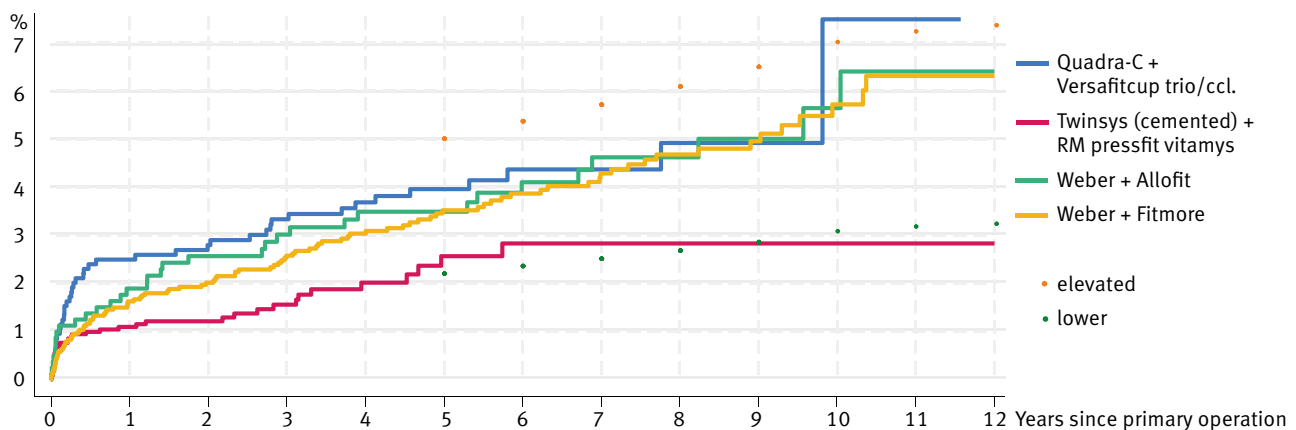
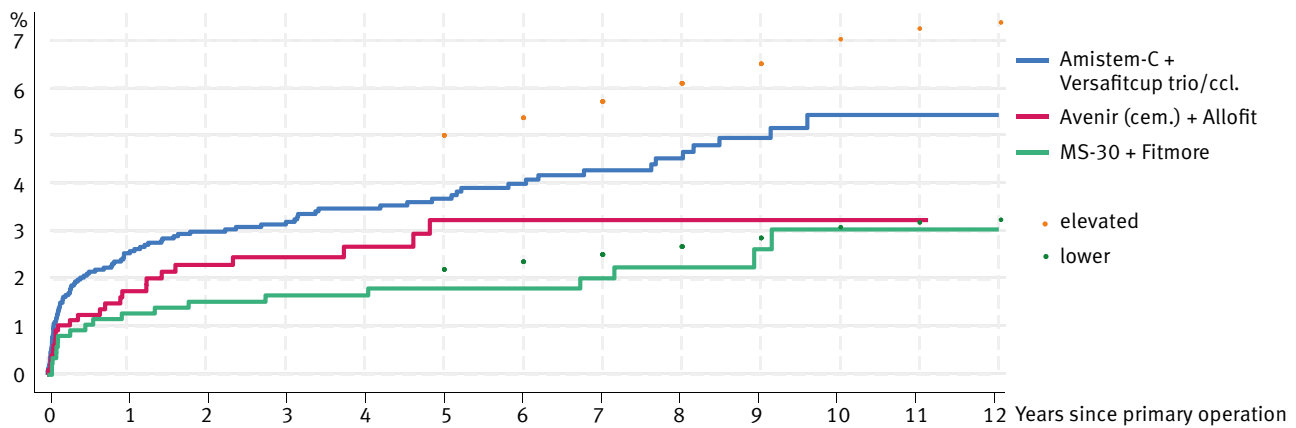


Figure 4.29

Implant combinations with below-average long-term revision rates (primary OA, hybrid THA)

Below-average was defined as an 9-year/10-year revision rate of up to 66% of the group average (and upper bounds of the 95% confidence interval staying below the lower bound of the group average; and at least 25 cases at risk at 9 years/10 years). The dots indicate upper and lower limits.



Figures 4.30

All remaining implant combinations with average revision risks (primary OA, hybrid fixation THA)

Also showing upper and lower limits (corresponding to elevated and below-average version risk at 150% and 66% of the group average respectively).

Uncemented combinations for secondary OA

Table 4.21 shows the 17 implant combinations that cover 75% of all uncemented THA performed for secondary OA and their use over the last 5 years. The 12-year revision rates for the period since 2012 was

7.0% (CI 6.4 – 7.7%) (Table 4.22), being significantly higher than the revision rates observed for uncemented and hybrid THA performed for primary OA, this illustrates the importance of coding the underlying diagnosis correctly for proper analysis of the results.

Stem component	Cup component	2019	2020	2021	2022	2023	2024	2019–2024
Actis	Pinnacle	9	15	36	69	71	84	284
Amistem-P	Versafitcup trio/ccl.	41	111	117	98	78	78	523
Avenir	Allofit	90	101	54	67	62	58	432
Avenir complete	G7 hemispherical	0	0	4	31	27	31	93
Corail	Pinnacle	77	73	99	62	55	43	409
Corail collared	Novae TH/Bi-Mentum	6	17	40	40	57	69	229
Corail collared	Pinnacle	107	128	211	266	215	234	1,161
Fitmore	Allofit	123	131	173	179	163	176	945
Fitmore	Fitmore	58	53	38	46	52	12	259
Fitmore	RM pressfit vitamys	11	33	23	23	16	12	118
Individual/custom hip	April ceramic	21	18	35	23	29	30	156
Optimys	RM pressfit vitamys	145	179	218	266	295	339	1,442
Polarstem	Polarcup	19	30	29	46	37	33	194
Polarstem	R3	74	89	89	91	99	143	585
Quadra-H	Versafitcup trio/ccl.	70	53	43	12	1	0	179
Quadra-P	Versafitcup trio/ccl.	2	21	43	100	130	148	444
Twinsys	RM pressfit vitamys	26	33	40	37	39	43	218
other combinations	-	467	401	385	393	375	466	2,487
Total		1,346	1,486	1,677	1,849	1,801	1,999	10,158

Table 4.21

Top 75% of primary total hip arthroplasty uncemented combinations (secondary OA) 2019– 2024

Although there were no outliers at 12 years, one combination (Quadra-H/Versafitcup Trio/ccl.) continued to have an elevated long-term revision rate (Figure 4.31). There was one combination (Fitmore/Allofit) showing a below-average long-term revision rate until 12 years (Figure 4.32). Figures 4.33 show the results of the implant combination with average revision risk.

Stem component	Cup component	Total N	CCS*	1 year (95% CI)	3 years (95% CI)	5 years (95% CI)	10 years (95% CI)	12 years (95% CI)
Amistem-H	Versafitcup trio/ccl.	557	14	1.8 (1.0-3.3)	2.6 (1.5-4.3)	3.8 (2.5-5.8)	8.0 (5.8-11.1)	10.4 (7.5-14.4)
Amistem-P	Versafitcup trio/ccl.	524	17	3.5 (2.2-5.5)	4.0 (2.6-6.1)	4.0 (2.6-6.1)		
Avenir	Allofit	868	13	3.5 (2.5-5.0)	4.5 (3.3-6.2)	4.9 (3.6-6.6)	5.4 (3.9-7.3)	5.4 (3.9-7.3)
Corail	Pinnacle	966	11	2.8 (1.9-4.1)	3.9 (2.8-5.3)	4.5 (3.3-6.1)	7.4 (5.3-10.2)	8.3 (5.8-11.8)
Corail collared	Pinnacle	1,647	29	2.1 (1.5-3.0)	3.2 (2.4-4.2)	3.7 (2.8-5.0)	4.3 (3.0-6.0)	4.3 (3.0-6.0)
Fitmore	Allofit	1,681	88	1.5 (1.0-2.2)	2.5 (1.8-3.4)	2.6 (1.9-3.5)	4.0 (2.7-5.8)	4.0 (2.7-5.8)
Optimys	RM pressfit vitamys	2,028	18	2.5 (1.9-3.3)	3.2 (2.5-4.1)	3.7 (2.9-4.8)	4.6 (3.5-6.1)	
Polarstem	R3	945	79	2.2 (1.4-3.3)	3.0 (2.1-4.5)	3.3 (2.2-4.8)	4.5 (3.0-6.9)	4.5 (3.0-6.9)
Quadra-H	Versafitcup trio/ccl.	604	26	3.5 (2.3-5.3)	5.2 (3.7-7.4)	7.8 (5.8-10.4)	10.3 (7.5-13.9)	12.9 (8.9-18.5)
other combinations	-	7747		3.5 (3.1-3.9)	4.6 (4.2-5.2)	5.4 (4.9-6.0)	7.2 (6.4-8.0)	7.2 (6.4-8.0)
CH average for group				2.9 (2.7-3.2)	3.9 (3.7-4.3)	4.6 (4.3-5.0)	6.5 (6.0-7.1)	7.0 (6.4-7.7)

Table 4.22

Long term evaluation: Failure rates of primary total hip arthroplasty uncemented combinations (secondary OA)

Time since operation, 2012–2024. Please note that if reported stem-cup combinations involve multiple sub-variants, it is possible that the long-term performance of these sub-variants may be significantly different from their combined performance.

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

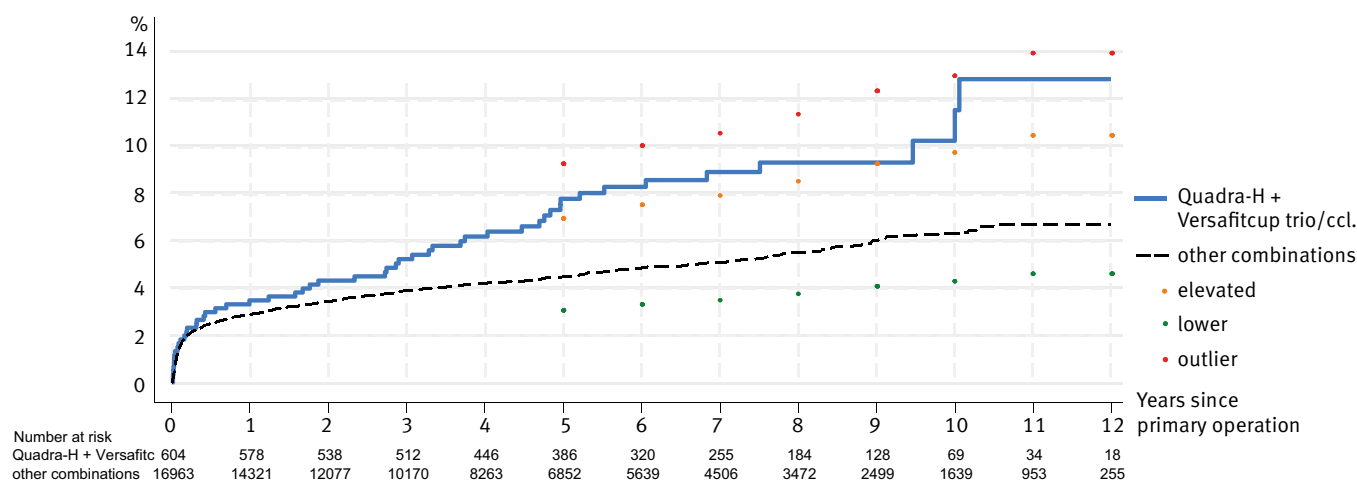


Figure 4.31

Implant combinations with elevated long-term revision rates (secondary OA, uncemented THA)

An elevated revision rate was defined as a deviation of at least 50% above the group average at any time between year 5 and year 12 (and lower bounds of the 95% confidence interval exceeding the upper bound of the group average; and at least 50 cases at risk at 5 years). The dots indicate upper and lower limits.

Hybrid and cemented combinations for secondary OA

Because of the relatively small numbers entered in the database, the data for all cemented and hybrid fixations for secondary OA are not presented, while the results for THA used to treat fractures are presented in Chapter 4.7.

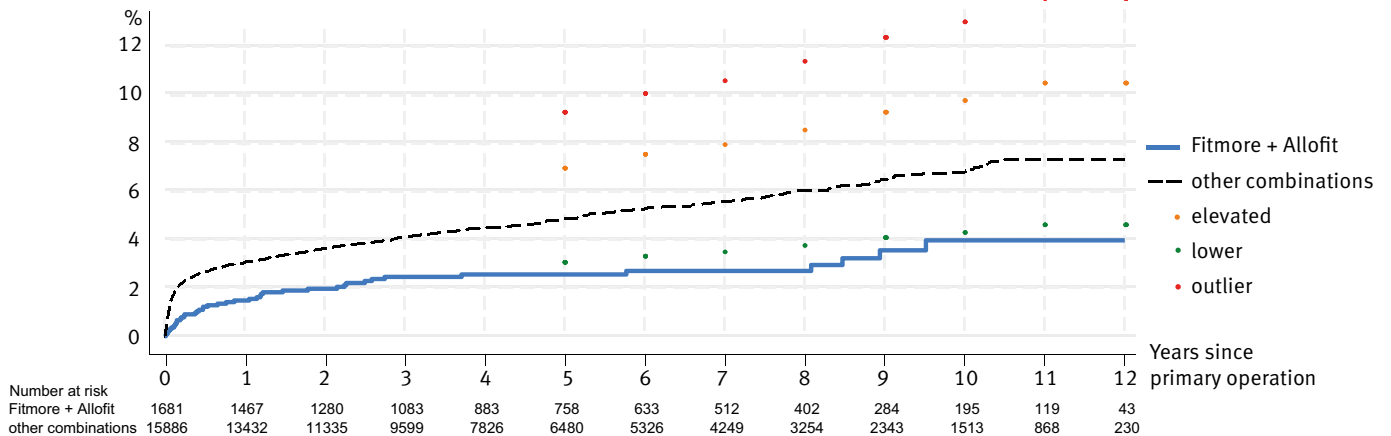
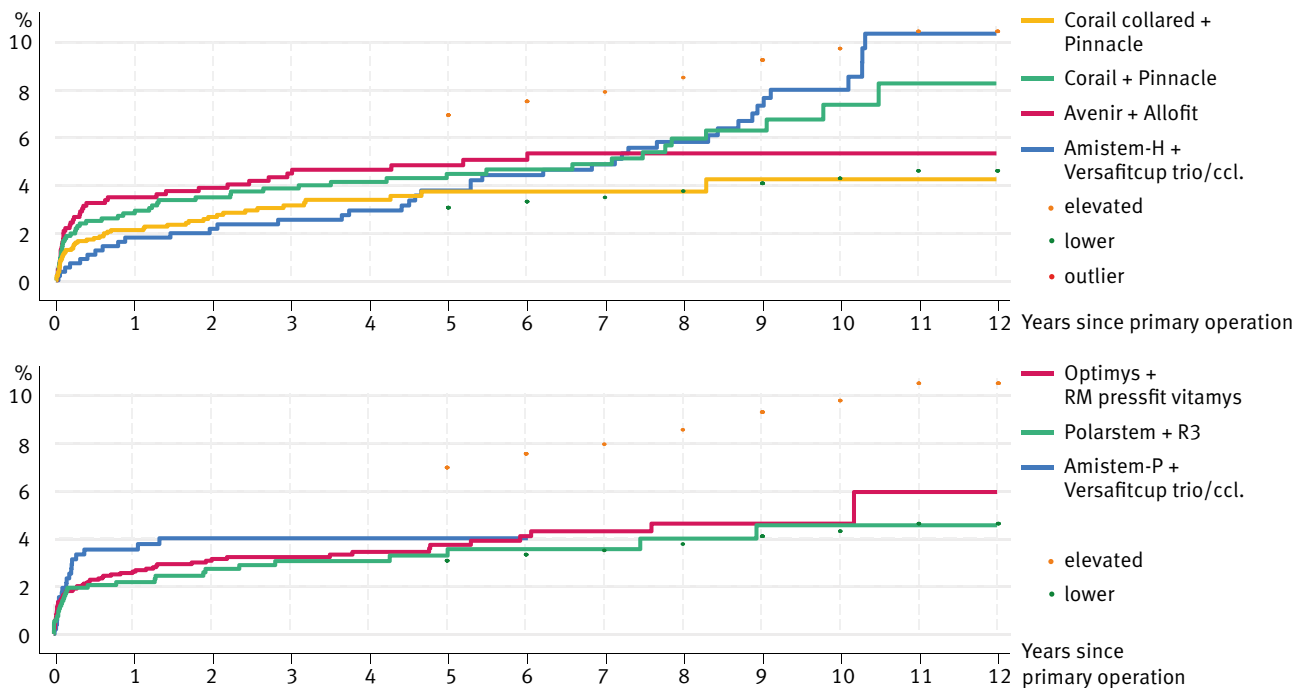


Figure 4.32

Implant combinations with below-average long-term revision rates (secondary OA, uncemented THA)

Below-average was defined as an 9-year/10-year revision rate of up to 66% of the group average (and upper bounds of the 95% confidence interval staying below the lower bound of the group average; and at least 25 cases at risk at 9 years/10 years). The dots indicate upper and lower limits.



Figures 4.33

All remaining implant combinations with average revision risks (secondary OA, uncemented THA)

Also showing upper and lower limits (corresponding to elevated and below-average version risk at 150% and 66% of the group average respectively).

4.4.2 Two-year revision rates

The 2-year revision rate is an important time point for gathering initial results about the early performance of an implant, especially since most early complications occur within the first 3 months after surgery. The average revision rate is calculated for the moving 4-year window period from 01.01.2019 to 31.12.2022. Due to the moving 4-year window for the analysis of the 2-year revision rates, some results may differ from those reported in 2023.

Uncemented combinations for primary OA

A total of 60,853 uncemented THA had been performed during the period of interest for primary OA. Compared to the previous observation period, the average revision rate has improved from 2.5% (CI 2.4 – 2.6%) to 2.4% (CI 2.3 – 2.5%). **Table 4.23** shows the 2-year revision rates of all uncemented implant combinations used more than 50 times for primary OA, representing 96% of all combinations. Overall, 346 implant combinations were used in less than 50 cases during the current 4-year observation period. Eight stem/cup combinations were identified as potential outliers and were further analysed following the protocol in Chapter 2 Methods and presented in the outlier watchlist at the end of this report. Compared to the previous report, two combinations disappeared from the list of outliers, four remained and four implant combinations were newly identified as outliers.

GTS/G7 bispherical, a previous outlier, is not in use anymore. Polarstem/EP-fit was a potential outlier in 2023 but was already improving. It continued to improve and is no outlier anymore. SPS Evolution/April ceramic turned out to be outliers four years in a row, but the results are slightly improving. Symbol/Symbol DMHA/DS evol. identifies for the third year in a row as outlier but has improved. The combination Nanos/R3 is a second-time outlier, however with a worsening revision rate. First time outliers are CLS Spotorno/Allofit, Exception/Avantage, SPS Evolution/April poly and Twinsys/RM pressfit. A detailed assessment of the outliers is presented in the outlier watch list at page 192.

Figure 4.34 shows the alphabetical list of stem/cup combinations with respect to the group average and outlier boundary, being twice the value of the group average of 2.4%.

Hybrid combinations for primary OA

A total of 7,742 hybrid THA had been performed for primary OA within the moving 4-year window period from 01.01.2019 to 31.12.2022. The average 2-year revision rate improved by 0.1% to 2.5% (CI 2.2 – 2.9%) (**Figure 4.35**). There were no outliers regarding the short-term revision rates detected in the current period of observation.

Uncemented combinations for secondary OA

A total of 6,349 uncemented THA had been performed for secondary OA within the current moving 4-year window period. The average 2-year revision rate was 3.6% (CI 3.2–4.1%) (**Figure 4.36**). None of the implant combinations were considered to be outliers. There was one implant combination (Fitmore/Allofit) with a below average revision rate (**Figure 4.32**).

Stem component	Cup component	CCS*	at risk N**	N	Revised % (95% CI)***
Accolade II	Trident II	32	130	8	6.2 (3.1-12.0)
Actis	Pinnacle	33	932	11	1.2 (0.7-2.1)
Alloclassic	Allofit	80	85	1	1.2 (0.2-8.2)
Alloclassic	Fitmore	66	50	0	0.0 (-.)
Amistem-H collared	Versafitcup trio/ccl.	100	68	0	0.0 (-.)
Amistem-H prox coating	Mpact	31	121	3	2.5 (0.8-7.6)
Amistem-H prox coating	Versafitcup trio/ccl.	17	935	30	3.2 (2.3-4.6)
Amistem-P	Mpact	39	385	11	2.9 (1.6-5.1)
Amistem-P	Versafitcup DM	27	106	3	2.8 (0.9-8.6)
Amistem-P	Versafitcup trio/ccl.	15	4,008	107	2.7 (2.2-3.2)
Ana.Nova alpha proxy	Ana.Nova alpha	96	153	2	1.3 (0.3-5.1)
Avenir	Allofit	15	3,516	83	2.4 (1.9-2.9)
Avenir	Avantage	17	77	1	1.3 (0.2-8.9)
Avenir	Fitmore	27	841	37	4.4 (3.2-6.0)
Avenir	G7 DM hemispherical	72	60	0	0.0 (-.)
Avenir	G7 hemispherical	32	60	3	5.2 (1.7-15.3)
Avenir complete	Allofit	50	126	4	3.2 (1.2-8.3)
Avenir complete	G7 DM hemispherical	83	76	2	2.7 (0.7-10.3)
Avenir complete	G7 hemispherical	37	75	3	4.0 (1.3-11.9)
CLS Spotorno	Allofit	44	273	14	5.1 (3.1-8.5)
CLS Spotorno	Fitmore	35	470	7	1.5 (0.7-3.1)
Corail	Fitmore	97	208	3	1.4 (0.5-4.4)
Corail	Pinnacle	17	4,451	129	2.9 (2.5-3.5)
Corail collared	Fitmore	87	111	2	1.8 (0.5-7.0)
Corail collared	Gyros	54	276	8	2.9 (1.5-5.7)
Corail collared	Novae TH/Bi-Mentum	31	842	19	2.3 (1.5-3.6)
Corail collared	Pinnacle	22	7,311	121	1.7 (1.4-2.0)
Corail collared	RM pressfit vitamys	64	70	0	0.0 (-.)
Corehip	Plasmafit	78	198	1	0.5 (0.1-3.6)
Exacta	Jump system/JS traser	87	163	4	2.5 (0.9-6.4)
Exacta S	Jump system/JS traser	52	202	6	3.0 (1.4-6.5)
Exception	Allofit	46	92	1	1.1 (0.2-7.5)
Exception	Avantage	70	304	16	5.3 (3.3-8.5)

Table 4.23 – Part one

2-year evaluation: Revision rates of uncemented primary total hip arthroplasty combinations within 24 months (primary OA)

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

*** Rates adjusted for effects of mortality and emigration.

Stem component	Cup component	CCS*	at risk N**	N	Revised % (95% CI)***
Exception	Exceed	100	63	2	3.2 (0.8-12.1)
Expersus	Primaro	99	73	2	2.7 (0.7-10.5)
Fitmore	Allofit	77	2,386	33	1.4 (1.0-1.9)
Fitmore	Fitmore	37	2,450	52	2.1 (1.6-2.8)
Fitmore	G7 hemispherical	55	64	2	3.2 (0.8-12.0)
Fitmore	RM pressfit vitamys	91	565	6	1.1 (0.5-2.4)
H-Max S	Delta TT	35	160	1	0.6 (0.1-4.4)
H-Max S	Symbol DMHA/DS evol.	85	106	4	3.8 (1.5-9.9)
Harmony	April ceramic	40	52	3	5.8 (1.9-16.8)
Individual/custom hip	April ceramic	28	585	12	2.1 (1.2-3.6)
Individual/custom hip	Pinnacle	53	120	2	1.7 (0.4-6.5)
LCU Hip system	Mobilelink acetabular Cup system	80	169	8	4.8 (2.4-9.3)
Metafix	Trinity	73	215	7	3.3 (1.6-6.7)
Minimax	Versafitcup trio/ccl.	47	144	5	3.5 (1.5-8.1)
Nanos	R3	46	89	7	8.0 (3.9-16.0)
Optimys	Anexys	24	225	7	3.1 (1.5-6.5)
Optimys	RM pressfit	18	297	13	4.4 (2.6-7.5)
Optimys	RM pressfit vitamys	8	9,485	202	2.1 (1.9-2.5)
Optimys	Symbol DMHA/DS evol.	61	157	2	1.3 (0.3-5.0)
Optimys	Trident II	92	152	2	1.3 (0.3-5.2)
Polarstem	EP-fit	94	329	13	4.0 (2.3-6.7)
Polarstem	HI	96	57	0	0.0 (-.)
Polarstem	Novae TH/Bi-Mentum	100	66	4	6.1 (2.3-15.4)
Polarstem	Polarcup	58	788	15	1.9 (1.2-3.2)
Polarstem	R3	52	3,197	49	1.5 (1.2-2.0)
Quadra-H	Allofit	100	50	2	4.0 (1.0-15.1)
Quadra-H	Mpact	36	410	9	2.2 (1.2-4.2)
Quadra-H	Versafitcup DM	48	132	3	2.3 (0.7-6.9)
Quadra-H	Versafitcup trio/ccl.	18	2,399	74	3.1 (2.5-3.9)
Quadra-P	Mpact	72	321	5	1.6 (0.7-3.7)
Quadra-P	Versafitcup DM	61	97	3	3.1 (1.0-9.3)
Quadra-P	Versafitcup trio/ccl.	27	1,714	25	1.5 (1.0-2.2)

Table 4.23 – Part two

2-year evaluation: Revision rates of uncemented primary total hip arthroplasty combinations within 24 months (primary OA)

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

*** Rates adjusted for effects of mortality and emigration.

Stem component	Cup component	CCS*	at risk N**	N	Revised % (95% CI)***
SBG	R3	42	633	10	1.6 (0.9-2.9)
SBG	Xentrax	100	71	1	1.4 (0.2-9.6)
SL-plus	HI	100	93	2	2.2 (0.5-8.3)
SL-plus MIA	EP-fit	47	222	4	1.8 (0.7-4.7)
SL-plus MIA	HI	39	245	7	2.9 (1.4-6.0)
SL-plus MIA	R3	73	383	6	1.6 (0.7-3.5)
SMS	Versafitcup trio/ccl.	56	192	6	3.2 (1.4-6.9)
SPS evolution	April ceramic	39	541	37	6.9 (5.0-9.4)
SPS evolution	April poly	38	61	5	8.3 (3.6-18.9)
SPS evolution	Liberty	40	82	3	3.7 (1.2-10.9)
Stelia-Stem	BSC pressfit	100	93	3	3.2 (1.1-9.7)
Symbol	Symbol DMHA/DS evol.	70	152	9	5.9 (3.1-11.1)
Tri-Lock	Pinnacle	68	94	3	3.2 (1.0-9.6)
Twinsys	Anexys	47	124	3	2.5 (0.8-7.4)
Twinsys	RM pressfit	41	74	7	9.5 (4.6-18.8)
Twinsys	RM pressfit vitamys	16	1,646	39	2.4 (1.7-3.2)
Twinsys	Symbol DMHA/DS evol.	52	67	1	1.5 (0.2-10.1)
other combinations	-		2,188	86	4.0 (3.2-4.9)
CH average for group					2.4 (2.3-2.5)

Table 4.23 – Part three

2-year evaluation: Revision rates of uncemented primary total hip arthroplasty combinations within 24 months (primary OA)

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

*** Rates adjusted for effects of mortality and emigration.

Important information on the use of the implant performance tables below

- Estimated revision rate exceeds the alert boundary, but we do not identify this implant combination as an outlier because the 95% confidence interval overlaps the confidence zone of the reference group.
- Identified as potential outliers. Please note the statistical confidence intervals. The outlier status comes with varying degrees of statistical probability. We consider the potential outlier status „highly likely“ when both the estimated revision rate and the complete confidence interval exceed the outlier alert boundary.

Please be aware that relatively rare implant combinations are frequently used in only a small number or indeed only in one hospital in Switzerland. Observed revision rates may be determined by local factors and performance may differ significantly between locations. Manufacturers of detected outlier implants and the hospitals where they were used (and revisions occurred) have been informed by SIRIS.

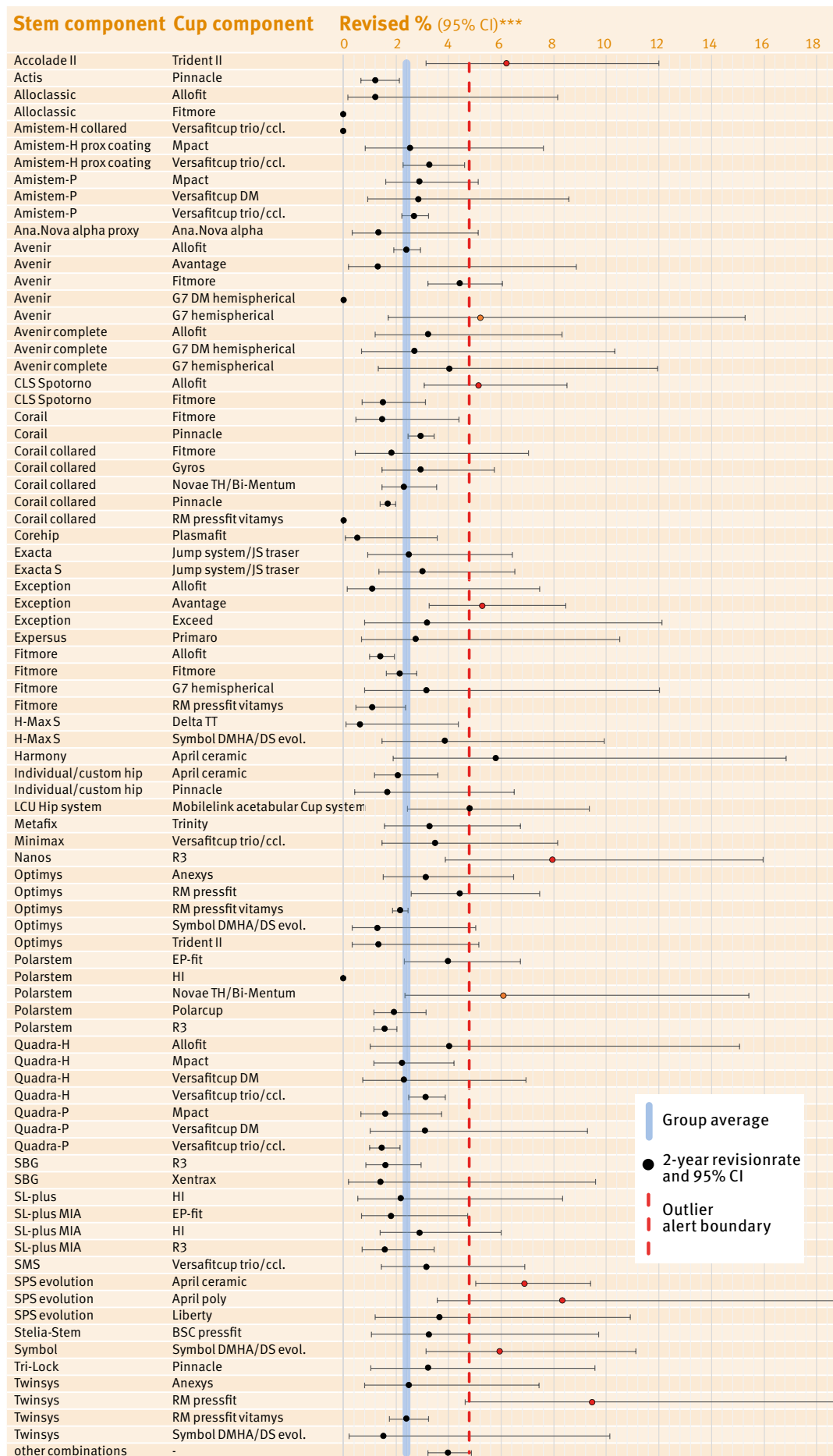


Figure 4.34

2-year evaluation: Revision rates of uncemented primary total hip arthroplasty combin. within 24 months (primary OA)

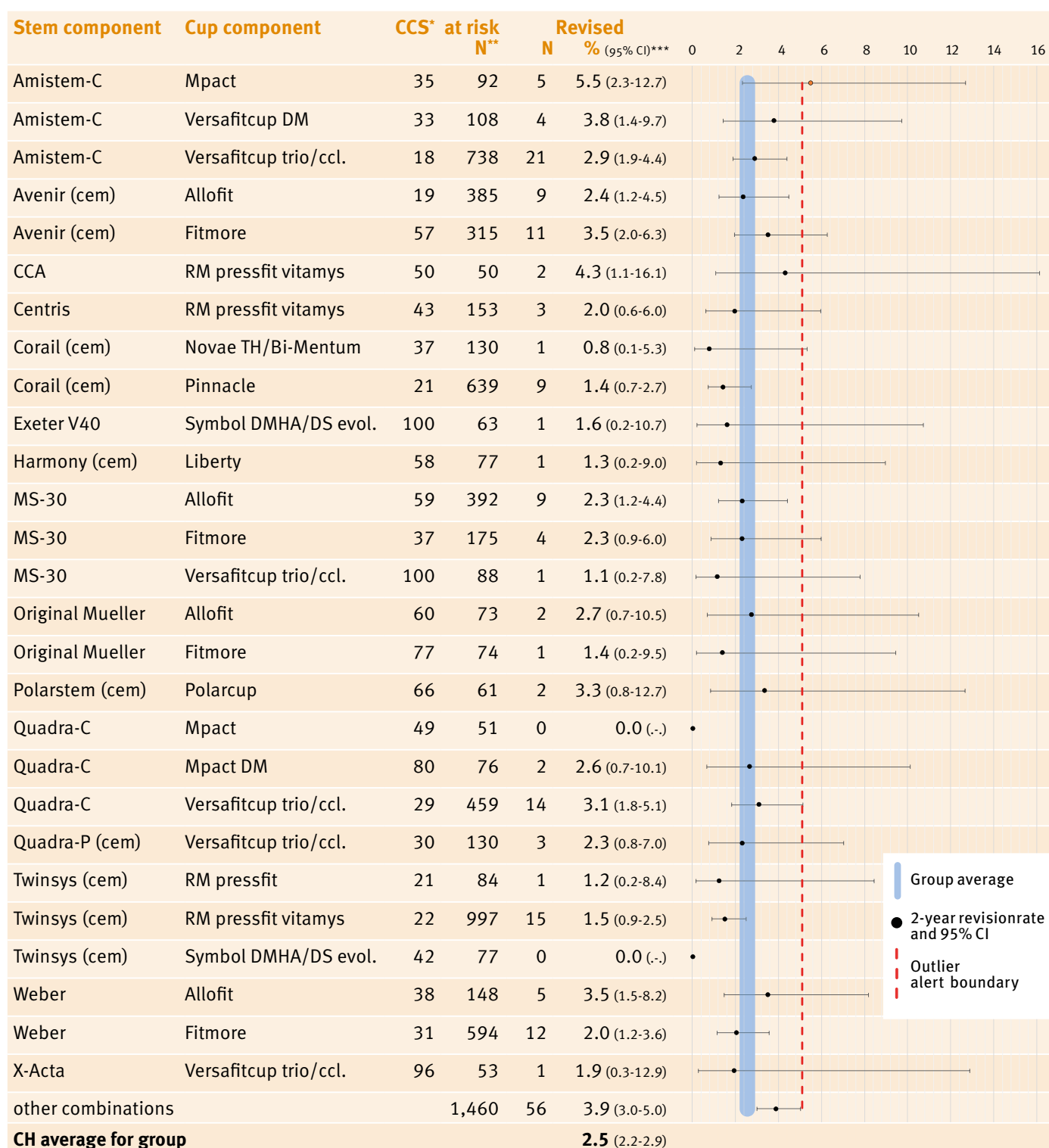


Figure 4.35

2-year evaluation: Revision rates of hybrid primary total hip arthroplasty combinations within 24 months (primary OA)

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

*** Rates adjusted for effects of mortality and emigration.

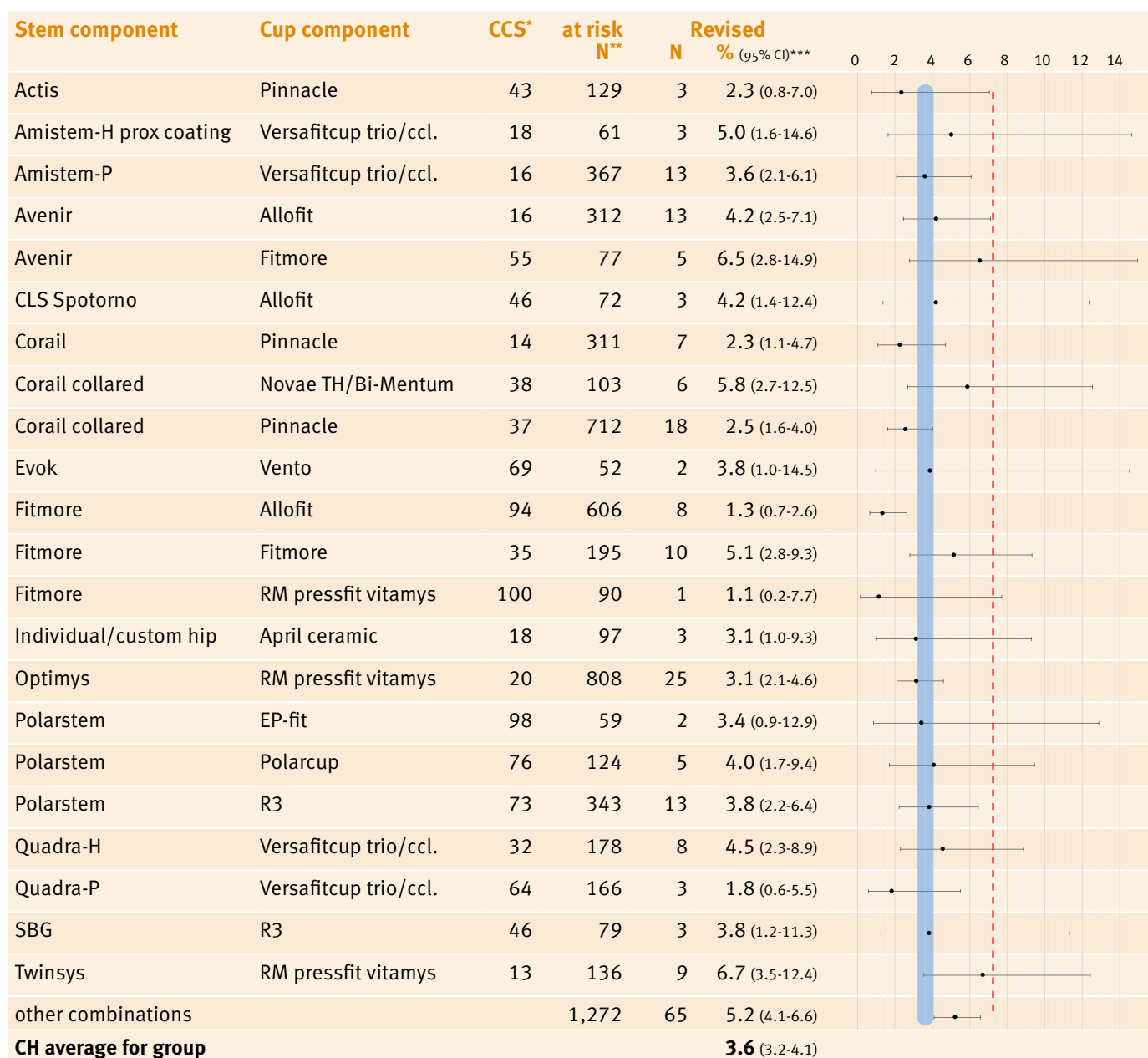


Figure 4.36

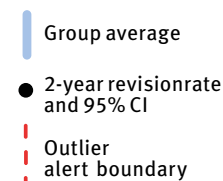
2-year evaluation: Revision rates of uncemented primary total hip arthroplasty combinations within 24 months (secondary OA)

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

*** Rates adjusted for effects of mortality and emigration.





Fracture of the hip

4.5 Treatment of hip fractures

Between 2019 and 2024, the registry recorded a total of 27,384 primary hip arthroplasties performed for fractures, of which 46.2% were THA and 53.8% HA (Table 3.7). Previous internal fixation of the femur

was registered in 8.1% of the THA and 1.0% of the HA. Previous internal fixation of acetabular fractures was recorded in 0.9% of the patients treated with THA. However, the lapse of time between internal fixation and arthroplasty is unknown, as internal fixation is not registered.

Main treatment group		THA		HA	
		N revised	%	N revised	%
Previous surgery ¹	None	11,328	89.5	14,452	98.1
	Internal fixation femur	1,022	8.1	149	1.0
	Osteotomy femur	60	0.5	15	0.1
	Internal fixation acetabulum	120	0.9	5	0.0
	Osteotomy pelvis	8	0.1	1	0.0
	Arthrodesis	2	0.0	0	0.0
	Other previous surgery	166	1.3	108	0.7
Approach	Anterior	7,311	57.8	7,149	48.5
	Anterolateral	2,953	23.3	4,044	27.5
	Posterior	1,514	12.0	2,120	14.4
	Lateral	665	5.3	1,265	8.6
	Other approach	213	1.7	150	1.0
Fixation	All uncemented / uncemented stem	6,317	49.9	1,837	12.5
	Hybrid acetabulum uncemented, femur cemented	5,289	41.8		
	All cemented / cemented stem	645	5.1	12,798	86.9
	Reverse hybrid acetabulum cemented, femur uncemented	200	1.6		
	Reinforcement ring, femur uncemented	54	0.4		
	Reinforcement ring, femur cemented	151	1.2		

Table 4.24

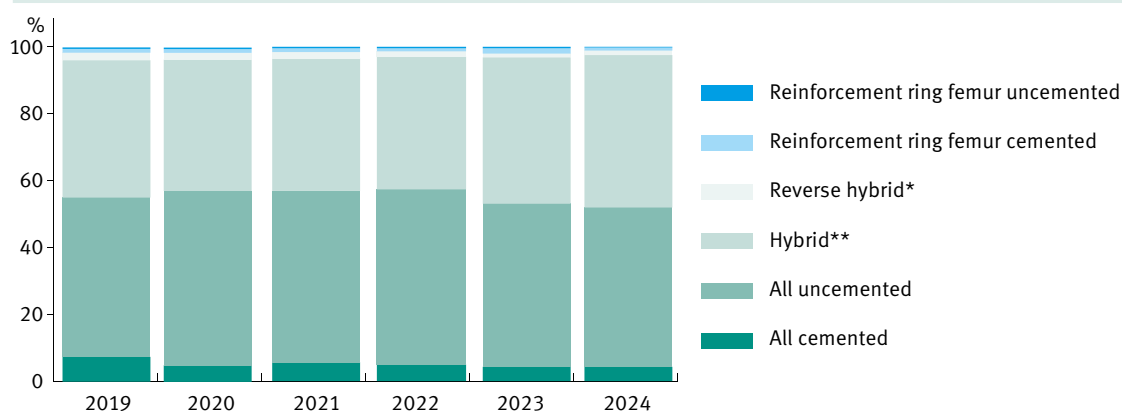
Fractures of the hip: Surgery characteristics by treatment group

¹ multiple responses possible; does not sum to 100%

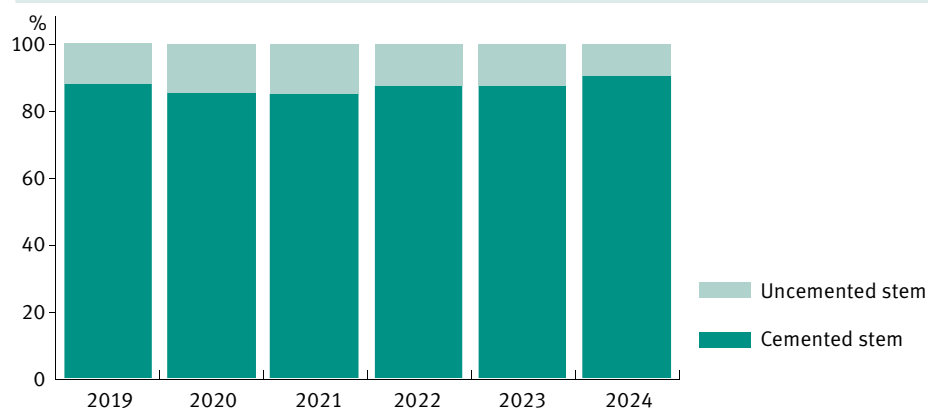
Most HA stems were cemented (86.9%) compared to 49.9% uncemented stems in the THA group (Table 4.24). The share of cemented stems for HA increased from 88.1% in 2019 to 90.5% in 2024. The use of

uncemented stems in the THA group reached a maximum in 2022 with 52.3%, thereafter declining to 47.7% in 2024. Simultaneously, the hybrid fixation showed a compensatory rise (Tables 4.25 and Figures 4.37).

Total hip arthroplasty (THA)	2019	2020	2021	2022	2023	2024
Reinforcement ring, femur uncemented	0.4	0.4	0.4	0.5	0.5	0.3
Reinforcement ring, femur cemented	1.3	1.2	1.2	1.0	1.6	0.9
Reverse hybrid*	2.1	1.9	1.9	1.5	1.1	1.2
Hybrid**	41.2	39.4	39.7	39.8	43.9	45.6
All uncemented	47.6	52.2	51.2	52.3	48.6	47.7
All cemented	7.3	4.9	5.6	4.9	4.4	4.2
Total [N]	1,593	1,789	2,134	2,290	2,380	2,470



Hemiarthroplasty (HA)	2019	2020	2021	2022	2023	2024
Uncemented stem	11.9	14.3	14.7	12.6	12.6	9.5
Cemented stem	88.1	85.7	85.3	87.4	87.4	90.5
Total [N]	2,274	2,351	2,326	2,558	2,536	2,590



Tables 4.25 and Figures 4.37

Fracture of the hip: Component fixation methods by type of treatment by year

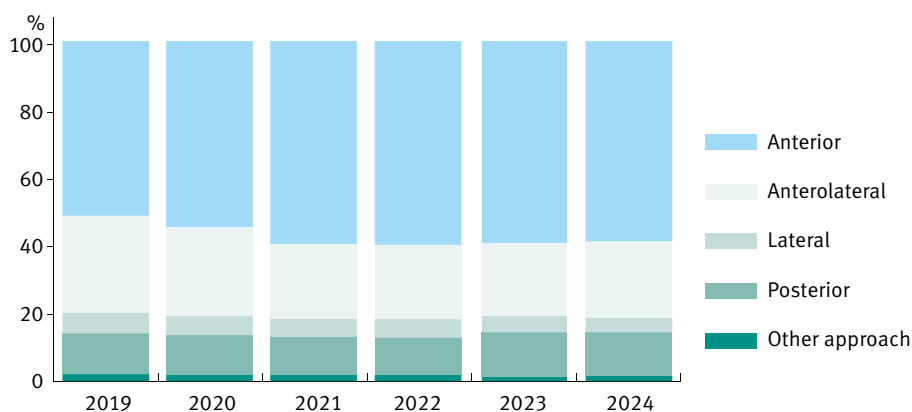
* acetabulum cemented, femur uncemented = Reverse hybrid

** acetabulum uncemented, femur cemented = Hybrid

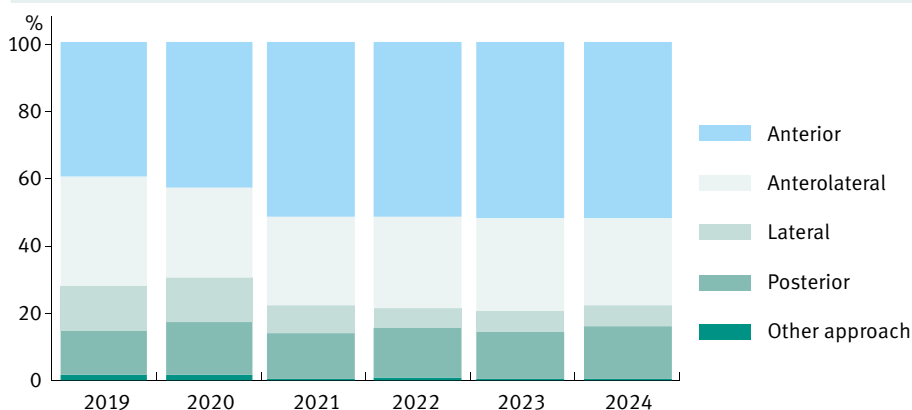
The most common approaches for both procedures were a direct anterior approach, followed by an anterolateral approach. The choice of the approach changed significantly between 2019 and 2024, particularly for HA. The use of the anterior approach for

HA rose from 39.6% in 2019 to 52.1% in 2024. The use of the anterolateral approach declined, as did the other approaches. The development was similar for THA, however less pronounced (Tables 4.26 and Figures 4.38).

Total hip arthroplasty (THA)	2019	2020	2021	2022	2023	2024
Anterior	51.3	54.7	59.7	59.9	59.3	58.9
Anterolateral	28.5	26.0	21.9	21.9	21.5	22.4
Lateral	6.2	5.9	5.3	5.3	5.0	4.3
Posterior	12.1	11.5	11.2	11.1	12.9	12.8
Other approach	1.9	1.9	1.8	1.7	1.3	1.5
Total [N]	1,593	1,789	2,134	2,290	2,380	2,470



Hemiarthroplasty (HA)	2019	2020	2021	2022	2023	2024
Anterior	39.6	42.9	51.4	51.8	52.2	52.1
Anterolateral	32.3	26.8	26.3	26.9	27.1	25.8
Lateral	13.0	12.9	8.3	5.7	6.2	6.2
Posterior	13.1	15.5	13.3	14.7	14.1	15.5
Other approach	1.9	1.9	0.6	0.9	0.5	0.4
Total [N]	2,280	2,360	2,361	2,575	2,547	2,605



Tables 4.26 and Figures 4.38

Fracture of the hip: Surgical approach by type of treatment by year

4.6 Early revision after arthroplasty for fracture of the hip

The 2-year revision rate after THA for fracture of the hip was 5.0% (CI 4.5 – 5.5%) and higher than in HA patients, where 3.7% (CI 3.3 – 4.1%) had been revised. The 2-year revision rate of THA for the treatment of fractures was more than twice that of THA for primary OA (5.0% vs 2.4%).

An elevated BMI was a risk factor for revision, whereby the risk increase for THA already was observed at a BMI >30 kg/m². Higher ASA scores tended to be associated with more frequent revisions in the THA group (Table 4.27).

		Total hip arthroplasty					Hemiarthroplasty				
		At risk*	Revised	95% CI		At risk*	Revised	95% CI			
		N	N	%**	lower	upper	N	N	%**	lower	upper
Overall (moving average)		7,812	372	5.0	4.5	5.5	9,574	310	3.7	3.3	4.1
Gender	Women	4,967	218	4.5	4.0	5.2	6,763	211	3.5	3.1	4.0
	Men	2,845	154	5.7	4.9	6.7	2,811	99	4.3	3.5	5.2
Age group	<55	332	20	6.1	4.0	9.3	31	4	14.4	5.7	34.0
	55–64	1,058	62	6.0	4.7	7.6	119	7	6.2	3.0	12.6
	65–74	2,040	92	4.6	3.8	5.6	508	34	7.6	5.5	10.5
	75–84	2,841	136	5.0	4.2	5.9	2,878	105	4.1	3.4	4.9
	85+	1,541	62	4.4	3.5	5.7	6,038	160	3.1	2.6	3.6
BMI group	<18.5	501	21	4.5	3.0	6.8	842	21	3.0	2.0	4.6
	18.5–24.9	3,633	146	4.2	3.6	4.9	4,533	114	2.9	2.4	3.4
	25–29.9	1,886	109	6.0	5.0	7.2	1,921	86	5.1	4.1	6.2
	30–34.9	463	35	7.9	5.7	10.8	393	24	6.6	4.4	9.6
	35–39.9	92	9	10.0	5.3	18.3	79	10	14.0	7.8	24.5
	40+	27	1	3.7	0.5	23.5	13	1	8.3	1.2	46.1
Morbidity state	Unknown	1,210	51	4.6	3.5	6.0	1,793	54	3.6	2.7	4.7
	ASA 1	486	14	2.9	1.7	4.8	72	5	7.7	3.3	17.6
	ASA 2	3,087	126	4.1	3.5	4.9	1,647	47	3.0	2.3	4.0
	ASA 3	3,498	198	6.0	5.3	6.9	6,386	214	3.9	3.4	4.4
	ASA 4/5	342	15	5.0	3.0	8.2	1,005	28	3.5	2.4	5.1
	Unknown	399	19	5.0	3.2	7.8	464	16	4.0	2.4	6.4

Table 4.27

Fracture of the hip: First revisions within 24 months overall and according to baseline characteristics

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

* Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

** Rates adjusted for effects of mortality and emigration.

In the HA group, uncemented stems had an increased risk for revision caused by a periprosthetic fracture. A posterior approach bore a higher risk of revision for both THA and HA, whereby for THA the effect was higher (7.4% vs 6.3%) as shown in **Table 4.28**. The difference was not significant.

Reasons for early first revision

Dislocation, periprosthetic fractures and infections were the three leading reasons for revision in both the THA and the HA groups (**Table 4.29**). However, infections remained by far the most important cause of revision in the HA group, representing 34.8% of the registered cases, while the most frequent cause for revision in fracture THA was dislocation (26.3%), followed closely by revision for periprosthetic fracture (25.0%).

	Total hip arthroplasty (THA)					Hemiarthroplasty (HA)				
	At risk*		Revised	95% CI		At risk*		Revised	95% CI	
	N	N	%**	lower	upper	N	N	%**	lower	upper
Overall (moving average)	7,812	372	5.0	4.5	5.5	9,574	310	3.7	3.3	4.1
All cemented / cemented stem	438	20	5.1	3.3	7.7	8,235	239	3.4	3.0	3.8
All uncemented / uncem. stem	3,983	199	5.1	4.5	5.9	1,272	68	5.9	4.7	7.4
Hybrid	3,123	137	4.6	3.9	5.4					
Anterior	4,449	195	4.6	4.0	5.3	4,462	142	3.7	3.1	4.3
Anterolateral	1,889	88	4.9	4.0	5.9	2,682	62	2.7	2.1	3.4
Lateral	439	14	3.3	2.0	5.5	946	25	3.2	2.2	4.7
Posterior	892	64	7.4	5.9	9.4	1,358	77	6.3	5.1	7.8
Other approach	143	11	8.1	4.6	14.2	126	4	4.1	1.5	10.6

Table 4.28

Fracture of the hip: First revisions according to stem fixation and approach

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

* Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

** Rates adjusted for effects of mortality and emigration.

	THA		HA			THA		HA	
	N	%	N	%		N	%	N	%
Dislocation	98	26.3	81	26.1	Wear	1	0.3	5	1.6
Infection	96	25.8	108	34.8	Metallosis	1	0.3	0	0.0
Periprosthetic fracture	93	25.0	67	21.6	Acetabular osteolysis	1	0.3	1	0.3
Loosening femoral	49	13.2	32	10.3	Femoral osteolysis	1	0.3	0	0.0
Loosening acetabular	31	8.3			Trochanter pathology	1	0.3	0	0.0
Position/Orientation of cup	28	7.5			Status after spacer	1	0.3	0	0.0
Position/Orientation of stem	20	5.4	11	3.5	Impingement	1	0.3	0	0.0
Implant breakage	6	1.6	1	0.3	Other	38	10.2	34	11.0
Acetabular protrusion	2	0.5	9	2.9	Total revisions	372		310	

Table 4.29

Fracture of the hip: Reasons for early first revisions

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

Early first revisions are those occurring within 2 years of the primary arthroplasty. Multiple responses possible (percentages do not sum to 100)

A more detailed analysis of the reasons for revision of HA, analysing separately unipolar and bipolar heads, showed notable differences. However, small numbers limit any further analysis (Table 4.30).

The revision rates of unipolar and bipolar heads for cemented HA showed that bipolar heads had a higher revision rate in the first year, whereas afterwards the difference was not significant. After 4 years and up to 12 years of follow-up, the revision rates of unipolar heads and bipolar heads remained similar (Figure 4.39).

The analysis again is limited by small numbers remaining at risk at the end of the observation period. The higher early revision rate of bipolar heads was due to revision for dislocation, which occurred significantly more frequently as in unipolar HA. Dislocations also occurred earlier in this subgroup. Periprosthetic fractures were more frequent in HA with unipolar heads, as were infections. The cumulative incidence figures of revision rates over time provide an additional perspective on the reasons for revision (Figures 4.40). It highlights revisions for infection and for dislocation tended to

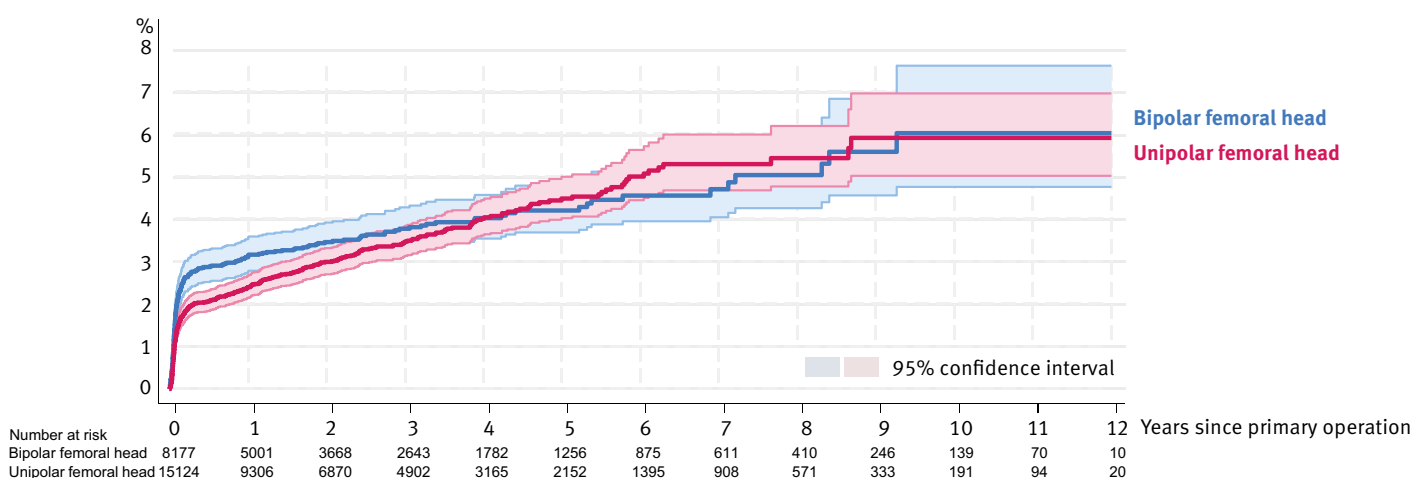
Reason	Unipolar heads		Bipolar heads	
	N	%	N	%
Infection	65	43.6	29	32.6
Dislocation	26	17.4	36	40.4
Periprosthetic fracture	24	16.1	11	12.4
Loosening femoral	14	9.4	13	14.6
Impingement	7	4.7	1	1.1
Acetabular osteolysis	0	0.0	1	1.1
Other	18	12.1	7	7.9
Total revisions	149		89	

Table 4.30

Fracture of the hip: Reasons for early first revisions (unipolar vs. bipolar hemi heads; cemented stems only)

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

Early first revisions are those occurring within 2 years of the primary arthroplasty. Multiple responses possible (percentages do not sum to 100).



Head type	1 year	2 years	3 years	5 years	6 years	8 years	10 years	12 years
Unipolar femoral head	2.4 (2.1-2.7)	3.0 (2.7-3.3)	3.4 (3.1-3.8)	4.5 (4.0-5.0)	5.0 (4.4-5.6)	5.4 (4.8-6.2)	5.9 (5.0-7.0)	5.9 (5.0-7.0)
Bipolar femoral head	3.1 (2.7-3.5)	3.5 (3.1-3.9)	3.8 (3.3-4.3)	4.2 (3.7-4.8)	4.6 (3.9-5.2)	5.0 (4.3-6.0)	6.0 (4.8-7.6)	6.0 (4.8-7.6)

Figure 4.39

Fracture of the hip: Estimated failure rates of hemiarthroplasty of the hip: unipolar heads versus bipolar heads

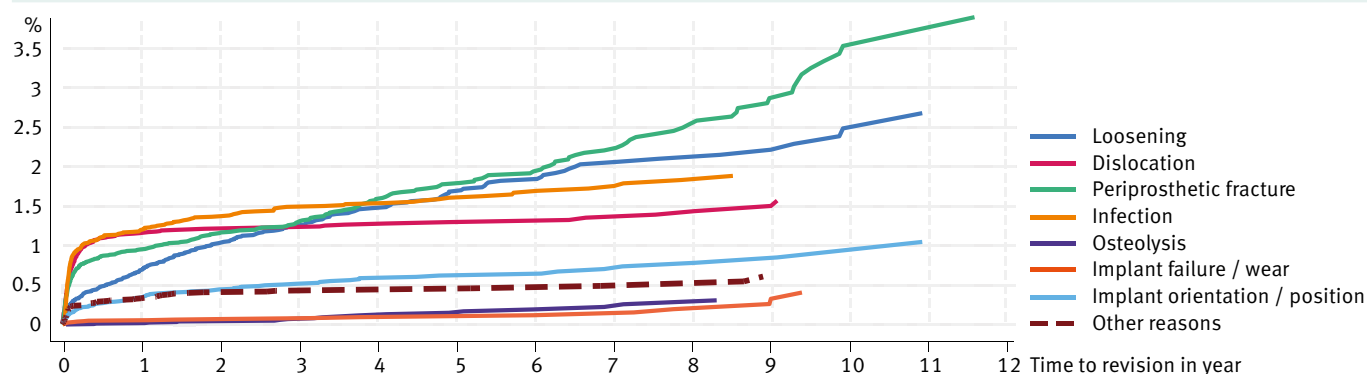
Time since operation, 2012–2024, only cemented stems % of implants revised.

occur rather early on, as indicated by a steep initial increase of the curve, followed by very gradual long-term rise. These observations were more frequent in THA. Incidents of loosening and periprosthetic fractures were the drivers of later revisions.

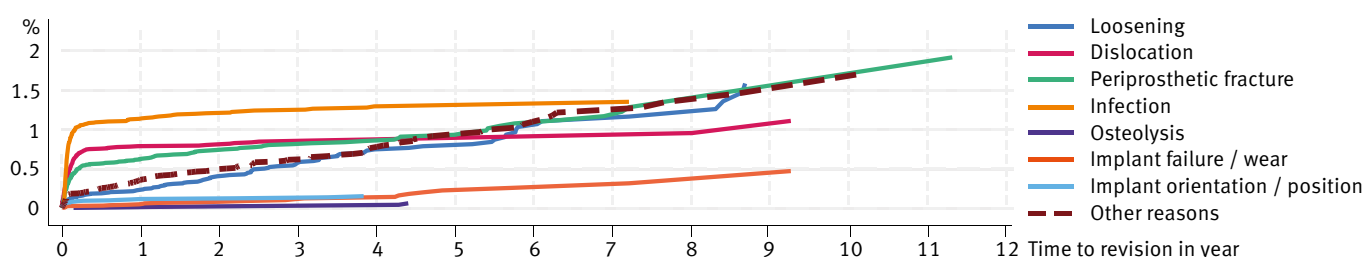
Type of revisions

Among the hip arthroplasties performed for fracture of the hip, 372 revisions of THA and 310 revisions of HA were carried out during the current moving observation period. The most frequent type of revision was

Total hip arthroplasty	1 year	2 years	3 years	5 years	9 years	10 years	12 years
Loosening	0.7 (0.6-0.8)	1.0 (0.9-1.2)	1.3 (1.1-1.4)	1.7 (1.5-1.9)	2.2 (1.9-2.6)	2.5 (2.1-3.0)	2.7 (2.1-3.3)
Dislocation	1.2 (1.0-1.3)	1.2 (1.1-1.4)	1.2 (1.1-1.4)	1.3 (1.1-1.5)	1.5 (1.3-1.8)	1.6 (1.3-1.9)	1.6 (1.3-1.9)
Periprosthetic fracture	1.0 (0.8-1.1)	1.2 (1.0-1.3)	1.3 (1.1-1.5)	1.8 (1.6-2.0)	2.9 (2.4-3.4)	3.5 (2.9-4.2)	3.9 (3.0-5.0)
Infection	1.2 (1.1-1.4)	1.4 (1.2-1.5)	1.5 (1.3-1.7)	1.6 (1.4-1.8)	1.9 (1.6-2.2)	1.9 (1.6-2.2)	1.9 (1.6-2.2)
Osteolysis	0.0 (0.0-0.1)	0.0 (0.0-0.1)	0.1 (0.0-0.1)	0.2 (0.1-0.3)	0.3 (0.2-0.5)	0.3 (0.2-0.5)	0.3 (0.2-0.5)
Implant failure / wear	0.1 (0.0-0.1)	0.1 (0.0-0.1)	0.1 (0.0-0.1)	0.1 (0.1-0.2)	0.3 (0.2-0.6)	0.4 (0.2-0.8)	0.4 (0.2-0.8)
Implant orientation / position	0.3 (0.3-0.4)	0.4 (0.4-0.6)	0.5 (0.4-0.6)	0.6 (0.5-0.8)	0.8 (0.6-1.0)	0.9 (0.6-1.1)	1.0 (0.7-1.6)
Other reasons	0.3 (0.3-0.4)	0.4 (0.3-0.5)	0.4 (0.3-0.5)	0.4 (0.4-0.6)	0.6 (0.4-0.9)	0.6 (0.4-0.9)	0.6 (0.4-0.9)



Hemiarthroplasty	1 year	2 years	3 years	5 years	9 years	10 years	12 years
Loosening	0.2 (0.2-0.3)	0.4 (0.3-0.5)	0.6 (0.5-0.7)	0.8 (0.6-1.0)	1.6 (1.2-2.2)	1.6 (1.2-2.2)	1.6 (1.2-2.2)
Dislocation	0.8 (0.7-0.9)	0.8 (0.7-0.9)	0.8 (0.7-1.0)	0.9 (0.8-1.0)	1.0 (0.8-1.2)	1.1 (0.8-1.5)	1.1 (0.8-1.5)
Periprosthetic fracture	0.6 (0.5-0.7)	0.7 (0.6-0.9)	0.8 (0.7-0.9)	0.9 (0.8-1.1)	1.3 (1.0-1.6)	1.3 (1.0-1.6)	1.9 (1.0-3.7)
Infection	1.1 (1.0-1.3)	1.2 (1.1-1.4)	1.2 (1.1-1.4)	1.3 (1.1-1.5)	1.4 (1.2-1.6)	1.4 (1.2-1.6)	1.4 (1.2-1.6)
Osteolysis	0.0 (0.0-0.0)	0.0 (0.0-0.0)	0.0 (0.0-0.1)	0.1 (0.0-0.2)	0.1 (0.0-0.2)	0.1 (0.0-0.2)	0.1 (0.0-0.2)
Implant failure / wear	0.0 (0.0-0.1)	0.1 (0.0-0.1)	0.1 (0.1-0.2)	0.2 (0.1-0.4)	0.3 (0.2-0.5)	0.5 (0.2-1.0)	0.5 (0.2-1.0)
Implant orientation / position	0.1 (0.1-0.2)	0.1 (0.1-0.2)	0.1 (0.1-0.2)	0.2 (0.1-0.2)	0.2 (0.1-0.2)	0.2 (0.1-0.2)	0.2 (0.1-0.2)
Other reasons	0.4 (0.3-0.4)	0.5 (0.4-0.6)	0.6 (0.5-0.7)	0.9 (0.8-1.1)	1.5 (1.1-1.9)	1.5 (1.1-1.9)	1.7 (1.2-2.5)



Figures 4.40

Fracture of the hip: cumulative incidence rates for different revision diagnoses, THA and HA

Time since operation, 2012–2024, % of implants revised

conversion of HA to THA (almost 35.5% of the revisions). The second most frequent revision in the HA group was the exchange of the head followed by revision of the stem (Table 4.31).

Type of revision	Total hip arthroplasty		Hemiarthroplasty	
	N revised	%	N revised	%
Exchange acetabular and femoral components	55	14.8		
Exchange acetabular component	27	7.3		
Exchange acetabular component and head	52	14.0		
Exchange femoral component	82	22.0	70	22.6
Exchange femoral component and inlay	20	5.4	2	0.6
Exchange head	39	10.5	100	32.3
Exchange inlay	7	1.9	0	0.0
Exchange head and inlay	61	16.4	14	4.5
Conversion of hemi-prosthesis to THA without stem exchange			61	19.7
Conversion of hemi-prosthesis to THA with stem exchange			49	15.8
Component removal, spacer implantation	8	2.2	5	1.6
Component reimplantation (after spacer or Girdlestone)	1	0.3	2	0.6
Girdlestone	4	1.1	2	0.6
Exchange femoral component, inlay and osteosynthesis	5	1.3	2	0.6
Other intervention	11	3.0	3	1.0
Total	372	100.0	310	100.0

Table 4.31

Fracture of the hip: Type of revisions by primary treatment modality, THA versus HA

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

HA: in approx. 11% of cases response categories involving acetabular components were chosen.

These were recoded to conversions.

4.7 Results of implants after hip fracture

Uncemented THA combinations for fractures

There were 19 uncemented stem/cup combinations, accounting for 75% of all uncemented THA performed for fracture of the hip (Table 4.32). The table also shows the distribution of the implants used between 2019 and 2024.

The revision rates for combinations with N>50 are shown in Figure 4.41. None of the implant combinations was conspicuous.

Stem component	Cup component	2019	2020	2021	2022	2023	2024	2019–2024
Amistem-H prox coating	Versafitcup trio/ccl.	63	6	9	0	0	0	78
Amistem-P	Versafitcup trio/ccl.	15	83	87	119	96	76	476
Avenir	Allofit	70	78	69	65	77	86	445
CLS Spotorno	Allofit	15	11	10	2	5	12	55
Corail	Pinnacle	61	66	72	57	51	23	330
Corail collared	Bimobile	0	0	20	122	94	99	335
Corail collared	Liberty	0	13	53	4	1	0	71
Corail collared	Novae TH/Bi-Mentum	4	28	65	74	101	118	390
Corail collared	Pinnacle	49	65	110	155	146	136	661
Fitmore	Allofit	18	15	26	20	39	38	156
Fitmore	Fitmore	15	21	14	26	12	6	94
Fitmore	RM pressfit vitamys	4	10	12	13	6	10	55
Optimys	RM pressfit	12	9	9	12	4	8	54
Optimys	RM pressfit vitamys	91	115	157	158	157	182	860
Optimys	Symbol DMHA/DS evol.	5	11	19	16	25	21	97
Polarstem	R3	13	16	11	23	19	12	94
Quadra-H	Versafitcup trio/ccl.	28	31	19	10	0	0	88
Quadra-P	Versafitcup trio/ccl.	0	10	16	22	31	34	113
Twinsys	RM pressfit vitamys	24	26	34	31	36	31	182
other combinations	-	238	277	259	250	241	265	1,530
Total		725	891	1,071	1,179	1,141	1,157	6,164

Table 4.32

Top 75% of primary total hip arthroplasty uncemented combinations to treat fractures 2019 – 2024

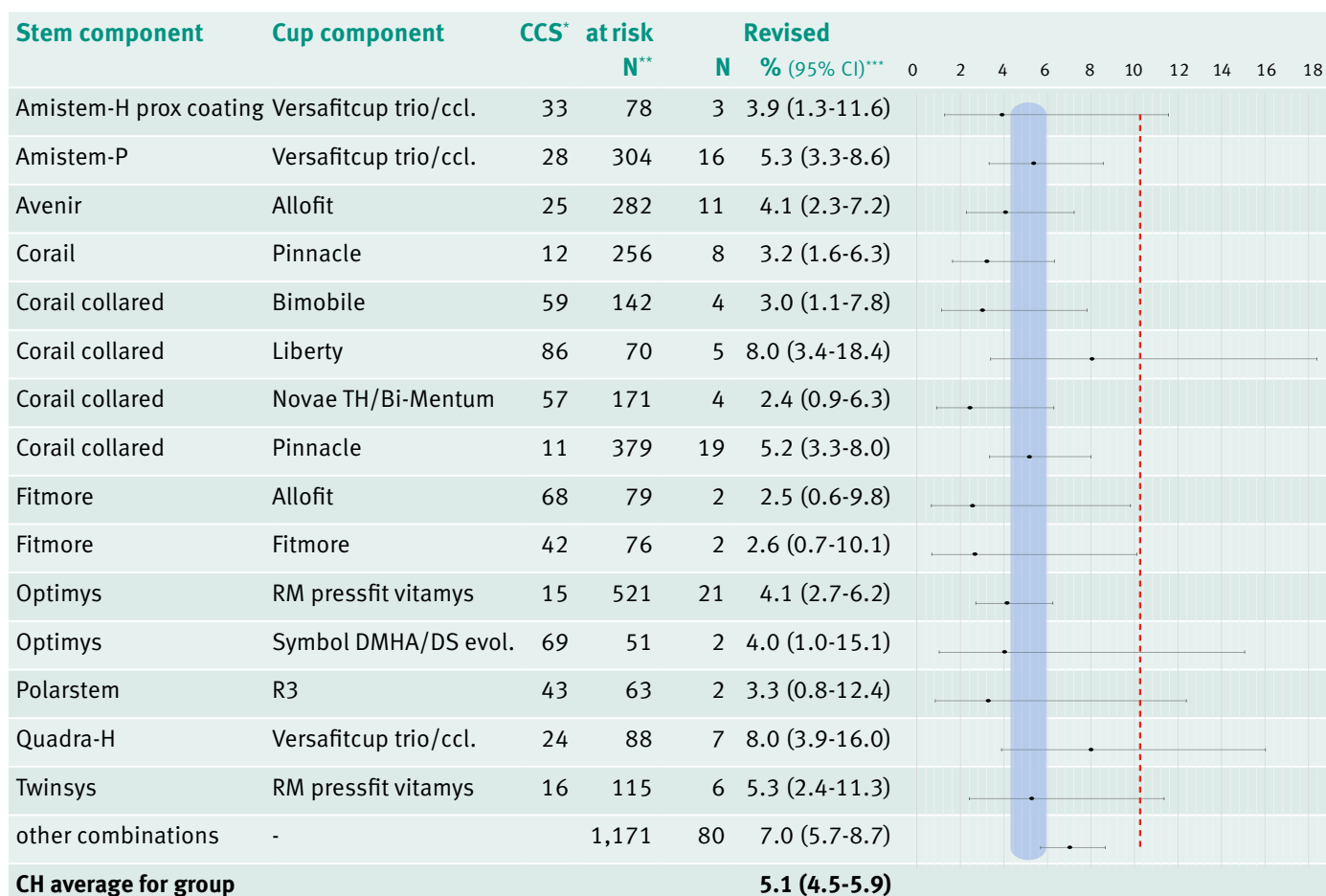


Figure 4.41

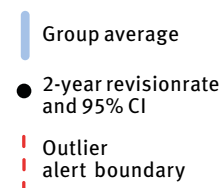
2-year evaluation: Revision rates within 24 months of uncemented primary total hip arthroplasty comb. to treat fractures

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

*** Rates adjusted for effects of mortality and emigration.



Important information on the use of the implant performance tables below

- Estimated revision rate exceeds the alert boundary, but we do not identify this implant combination as an outlier because the 95% confidence interval overlaps the confidence zone of the reference group.
- Identified as potential outliers. Please note the statistical confidence intervals. The outlier status comes with varying degrees of statistical probability. We consider the potential outlier status „highly likely“ when both the estimated revision rate and the complete confidence interval exceed the outlier alert boundary.

Please be aware that relatively rare implant combinations are frequently used in only a small number or indeed only in one hospital in Switzerland. Observed revision rates may be determined by local factors and performance may differ significantly between locations. Manufacturers of detected outlier implants and the hospitals where they were used (and revisions occurred) have been informed by SIRIS.

Hybrid THA combinations for fractures

There were 20 stem/cup combinations covering 75% of the hybrid THA performed for fractures of the hip, but nine of these combinations were used fewer than 100 times during the observed period between 2018 and 2023 (Table 4.33). The revision rates for the 10 combinations with N>50 are presented in Figure 4.42 and show that none of the implants reached potential outlier status.

Stem component	Cup component	2019	2020	2021	2022	2023	2024	2019–2024
Amistem-C	Mpact DM	0	5	4	0	18	15	42
Amistem-C	Versafitcup DM	16	12	9	6	17	11	71
Amistem-C	Versafitcup trio/ccl.	86	94	108	106	81	124	599
Avenir (cem)	Allofit	30	33	39	49	77	137	365
Avenir (cem)	Fitmore	26	37	65	58	45	14	245
CCA	RM pressfit vitamys	19	9	10	19	6	5	68
Centris	RM pressfit vitamys	30	32	53	0	0	0	115
Corail (cem)	Novae TH/Bi-Mentum	2	8	29	53	70	98	260
Corail (cem)	Pinnacle	38	38	79	120	146	99	520
Corail (cem)	RM pressfit vitamys	0	0	0	11	14	33	58
Exacta S	Symbol DMHA/DS evol.	0	0	5	24	39	49	117
Exeter V40	Symbol DMHA/DS evol.	0	0	7	16	29	14	66
MS-30	Allofit	0	0	10	29	43	24	106
Quadra-C	Mpact	26	19	16	8	15	4	88
Quadra-C	Mpact DM	9	32	35	33	14	17	140
Quadra-C	Versafitcup trio/ccl.	72	64	35	21	4	4	200
Quadra-P (cem)	Versafitcup trio/ccl.	0	4	5	3	17	18	47
Twinsys (cem)	RM pressfit	5	6	6	12	5	8	42
Twinsys (cem)	RM pressfit vitamys	69	74	74	119	148	145	629
Twinsys (cem)	Symbol DMHA/DS evol.	4	3	5	13	13	19	57
Weber	Fitmore	51	46	37	33	26	8	201
other combinations	-	185	204	226	186	226	274	1,301
Total		668	720	857	919	1,053	1,120	5,337

Table 4.33

Top 75% of primary total hip arthroplasty hybrid combinations to treat fractures 2019 – 2024

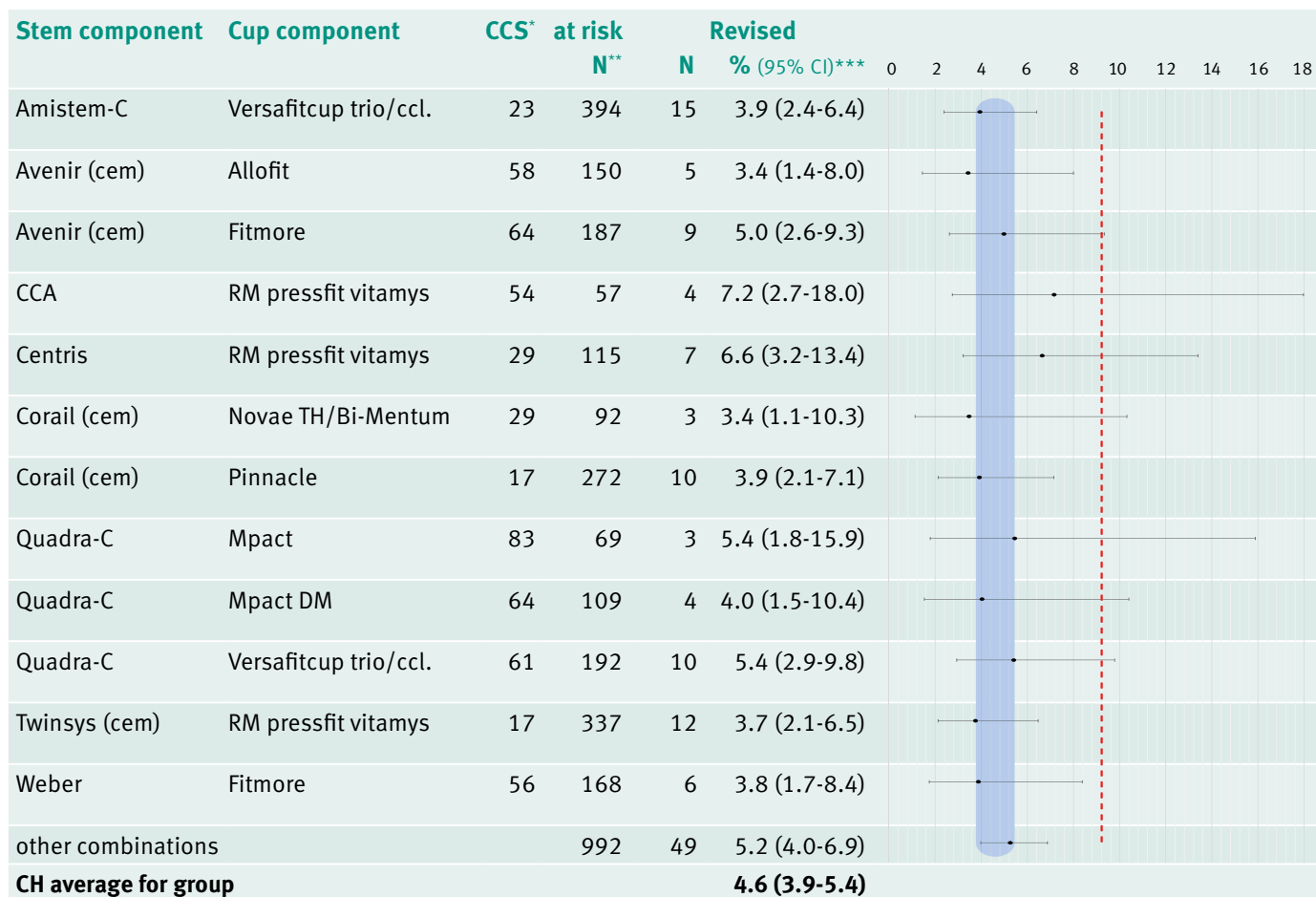


Figure 4.42

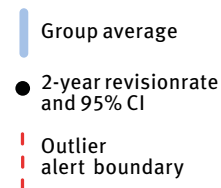
2-year evaluation: Revision rates within 24 months of hybrid primary total hip arthroplasty combinations to treat fractures

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

*** Rates adjusted for effects of mortality and emigration.



Hemiarthroplasty

The choice of implant combination for the treatment of hip fractures with HA was less variable than for THA and there were only nine stem/head combinations accounting for 75% of all implantations (Table 4.34). These combinations were used rather frequently over the last 12 years, and it is worth noting that neither combination was used fewer than 300 times during the last 5 years. Centris was withdrawn from the market. Therefore, it is a matter of time until it disappears

from this analysis. The revision rates for combinations with $n > 50$, which are more than the above-mentioned nine combinations, are shown in Figure 4.43. The average revision rate of this subgroup was 3.3% (CI 2.9–3.8%). There were two implant combinations that were detected as outliers. Among these, the combination Harmony (cem)/Symbios bibop was used in one centre only, illustrating the importance of single centres on the revision rates of specific implants, as a single combination (Table 4.33).

Stem component	Head component	2019	2020	2021	2022	2023	2024	2019–2024
Amistem-C	Medacta bipolar head	92	114	150	171	196	221	944
Amistem-C	Medacta endohead	282	329	377	415	380	345	2,128
Avenir (cem.)	ZB bipolar head	79	99	69	100	118	151	616
Avenir (cem.)	ZB unipolar head	28	44	72	69	69	77	359
CCA	Hemihead SS	437	395	350	289	190	166	1,827
Centris	Hemihead SS	109	103	113	0	0	0	325
Corail (cem)	J&J modular head carthcart	87	106	180	260	277	300	1,210
Twinsys (cem.)	Hemihead SS	97	127	121	238	330	293	1,206
Twinsys (cem.)	Mathys bipolar steel head	38	46	30	65	57	91	327
Weber	ZB unipolar head	225	168	140	151	104	69	857
other combinations	-	524	450	368	459	459	586	2,846
Total		1,998	1,981	1,970	2,217	2,180	2,299	12,645

Table 4.34

Fracture of the hip: top 75% stem/head combinations used in hemiarthroplasty (HA) 2019 – 2024



Figure 4.43 – Part one

2-year evaluation: Revision rates within 24 months of hybrid primary total hip arthroplasty combinations to treat fractures

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

*** Rates adjusted for effects of mortality and emigration.

Stem component	Head component	CCS*	at risk N**	Revised	
				N	% (95% CI)***
Amistem-C	Hemihead SS	54	54	1	2.1 (0.3-14.2)
Amistem-C	Medacta bipolar head	24	526	13	2.7 (1.6-4.7)
Amistem-C	Medacta endohead	29	1,403	45	3.8 (2.8-5.1)
Arcad	OHST bipolar head	73	56	0	0.0 (-.)
Avenir (cem)	ZB bipolar head	30	347	9	2.9 (1.5-5.5)
Avenir (cem)	ZB unipolar head	21	213	4	2.2 (0.8-5.8)
CCA	Hemihead SS	32	1,469	33	2.8 (2.0-3.9)
CCA	Mathys bipolar steel head	27	124	5	5.1 (2.2-12.0)
Centris	Hemihead SS	40	324	6	2.0 (0.9-4.5)
Corail (cem)	J&J modular head carthcart	19	633	18	3.2 (2.0-5.1)
Corail (cem)	S&N bipolar ballhead	89	61	0	0.0 (-.)
Exacta S	Endoheads	100	69	4	7.3 (2.8-18.7)
Harmony (cem)	Acropole bipolar head	97	154	7	5.3 (2.5-10.8)
Harmony (cem)	OHST bipolar head	84	111	8	8.0 (4.1-15.3)
Harmony (cem)	Symbios bibop	100	54	5	10.3 (4.4-23.1)
MS-30	ZB bipolar head	69	59	1	1.9 (0.3-12.9)
Original Mueller	ZB unipolar head	26	152	2	1.6 (0.4-6.4)
Quadra-C	Medacta bipolar head	46	125	8	6.9 (3.5-13.3)
Quadra-C	Medacta endohead	46	68	5	8.5 (3.6-19.3)
Twinsys (cem)	Hemihead SS	21	583	18	3.5 (2.2-5.5)
Twinsys (cem)	Mathys bipolar steel head	22	179	4	2.4 (0.9-6.3)
Weber	ZB bipolar head	50	193	4	2.3 (0.8-5.9)
Weber	ZB unipolar head	34	683	19	3.1 (2.0-4.8)
other combinations	-		500	15	3.4 (2.1-5.6)
CH average for group					3.3 (2.9-3.8)

Figure 4.43 – Part two

2-year evaluation: Revision rates within 24 months of hybrid primary total hip arthroplasty combinations to treat fractures

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

- * Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.
- ** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).
- *** Rates adjusted for effects of mortality and emigration.



Knee arthroplasty

5. Knee arthroplasty

Overview of data structure

Primary procedures

Total knee arthroplasty (TKA)
197,143

Other/unclear
115

Primary OA
176,266
Secondary OA
20,877

Partial knee arthroplasty (PK)
35,908

Primary OA
32,580
Secondary OA
3,328

Revision/reoperation procedures

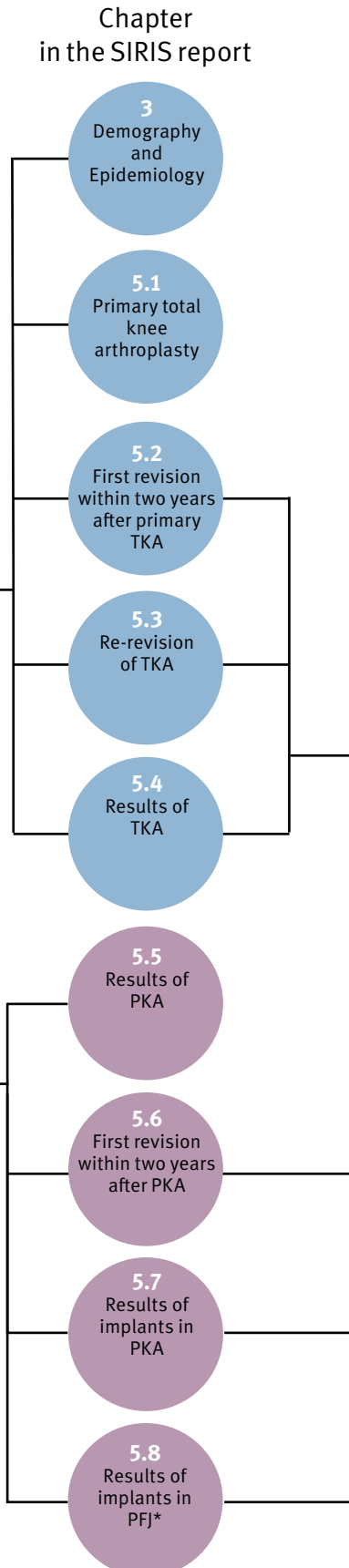
Unlinked revision procedures
13,337

90%+ are estimated to be revisions of TKAs. We suspect that some conversions are captured in other revision categories.

exclude

Revision procedures
N= 29,896
Component revision
28,927
Reoperation
969
Linked revision procedures
16,559

Unknown
27



* Patellafemoral joint partial knee arthroplasty

Overview of types of analyses for determining revision rates

Types of analysis	Kaplan-Meier estimates 2012 – 2024	2-year revision rates (implants 2019 – 2022 with completed 2-year follow-up)	Funnel plots of 2-year hospital revision rates (implants 2019 – 2022 with completed 2-year follow-up)
Report section	Adjusted for censoring events	Adjusted for censoring events	Risk-adjusted and adjusted for censoring events
Knee overview	All total knee arthroplasties (TKA) All partial knee arthroplasties (PKA)		TKA after primary osteoarthritis (primary OA). ANQ online reporting, above 99.8%= outlier status All partial knee arthroplasties (PKA) TKA after primary OA without isolated patella resurfacing
First revision of primary TKA	TKA for various subgroups	TKA for various subgroups	
First revision of primary PKA	PKA for various subgroups	PKA for various subgroups	
Re-revision after revision of TKA/PKA	Re-revision after revised TKA for various subgroups Re-revision after conversion from PKA to TKA		
Knee implants (minimal number in group)	Bicondylar total knee systems, all diagnoses (500+) Unicondylar partial knee systems, all diagnoses (500+) Patellofemoral joint systems, all diagnoses (500+) Long-term evaluation 5–10 years: elevated revision rate or outlier	Bicondylar total knee systems, all diagnoses (50+) Unicondylar partial knee systems, all diagnoses (50+) Patellofemoral joint systems, all diagnoses (50+) 2-year evaluation (two times group average= outlier status)	

5.1 Primary total knee arthroplasty

The total number of primary TKA registered in SIRIS at the end of 2024 reached 197,143 cases (Table 3.10). The share of women (59.0%) and the mean age of the

patients (69.8 years) remained constant throughout the entire registration period (Table 3.11). Please consult also Chapter 3.3.2 for further details regarding incidence and demographic characteristics.

N (2019 – 2024)	N	%	N (2021 – 2024)	N	%
Previous surgery			Additional intervention		
None	74,793	68.3	None	74,327	95.6
Knee arthroscopy	16,658	15.2	Osteosynthesis FE	90	0.1
Meniscectomy	19,786	18.1	Osteosynthesis TI	73	0.1
ACL reconstruction	6,089	5.6	Osteosynthesis PAT	6	0.0
Osteotomy tibia close to knee	2,618	2.4	Removal of metalwork	993	1.3
Osteosynthesis tibia close to knee	1,578	1.4	Operation extensors	462	0.6
Surgery for patella stabilization	1,369	1.3	Reconstruction plasty	82	0.1
Synovectomy	834	0.8	Tibial tubercle osteotomy	1,057	1.4
Osteotomy femur close to knee	449	0.4	Other additional intervention(s)	858	1.1
Osteosynthesis femur close to knee	549	0.5	Total TKA (multiple responses)	77,746	
Surgery for treating infection	198	0.2	Additional components		
Surgery for tumor	44	0.0	Stem FE (cemented)*	1,321	1.7
Other	2,034	1.9	Stem FE (uncemented)**	431	0.6
Total TKA (multiple responses)	109,489		Stem TI (cemented)***	4,652	6.0
Intervention			Stem TI (uncemented)***	711	0.9
CS (cruciate sacrificing) / UCOR	26,065	23.8	Sleeve FE	47	0.1
PS (posterior stabilised)	29,105	26.6	Sleeve TI	140	0.2
PCR (posterior cruciate retaining)	27,851	25.4	Augments FE	98	0.1
BCR (bicruciate retaining)	946	0.9	Augments TI	164	0.2
Hinge type	2,186	2.0	Augments PAT	2	0.0
SC/ CCK (semi-constrained/ constrained)	1,414	1.3	Bone homologous	47	0.1
Other (Medial-Pivot)*	21,291	19.4	Bone autologous	135	0.2
Other	565	0.5	Cone FE	6	0.0
Technology			Cone TI	43	0.1
Conventional (including minimally invasive)	73,540	67.2	Total TKA (multiple responses)	77,935	
Computer assisted / navigation	10,165	9.3			
Patient specific instrumentation	19,861	18.1			
Robotic-assisted (v2021)	6,568	6.0			
Other	1,394	1.3			
Total TKA (multiple responses)	109,489				

* 60% with cement restrictor.

** 32% with coating.

*** 25% with cement restrictor.

**** 35% with coating

Table 5.1

Primary total knee arthroplasty: Surgery characteristics

All diagnoses. *Medial pivot was not available as a response category before SIRIS v2021.

All GMK Sphere knee systems are counted as medial pivot, regardless of the type chosen locally at data entry.

Previous surgery

Between 2019 and 2024, no previous surgery was recorded in 68.3% of the TKA registered, while 18.1% had former meniscectomy and 15.2% arthroscopy. Preceding ACL reconstruction was registered in 5.6%, and osteotomy at tibial level in 2.4%, but at the femur only in 0.4%. All other conditions were rare, in 1.9% “other” former surgeries were reported (Table 5.1). The rate of previous arthroscopies preceding TKA in primary OA was constantly decreasing since 2012 and reached only 13% in 2024. The share of previous arthroscopy was much higher in secondary OA, reaching 25% in 2024, a rate remaining constant over the past 4 years (Figure 5.1).

Type of knee prosthesis

The classification of the TKA systems registered in SIRIS was adapted in 2021 with the last revision of the case report form (CRF), because of previously not unambiguous terminology. Between 2019 and 2024, the share of cruciate-sacrificing/ultra congruent systems (CS/UCOR) was 23.8%, for posterior stabilised (PS) it was 26.6% and posterior cruciate-retaining (PCR/CR) it was 25.4%. A medial pivot (MP) was used in 19.4%, whereas semi-constrained/constrained condylar knees (SC/CCK) or hinged implants were used only in 1.3% and 2.0% of the cases, respectively. Bi cruciate-retaining knees (BCR) were rarely used (0.9%) as well (Table 5.1). The share of MP implants seemed to further increase nationwide and replace more traditional designs such as PS, PCR/CR and CS/UCOR. The proportion of SC/CCK TKA was approximately 1.5% in primary and approximately 3.1% in secondary OA. For hinged systems, the share almost tripled in secondary OA (approx. 4.6% versus approx. 1.6% in primary OA). Of note, the knee replacement systems used varied significantly between cantons, regions, and hospitals (see chapter 3.3.5).

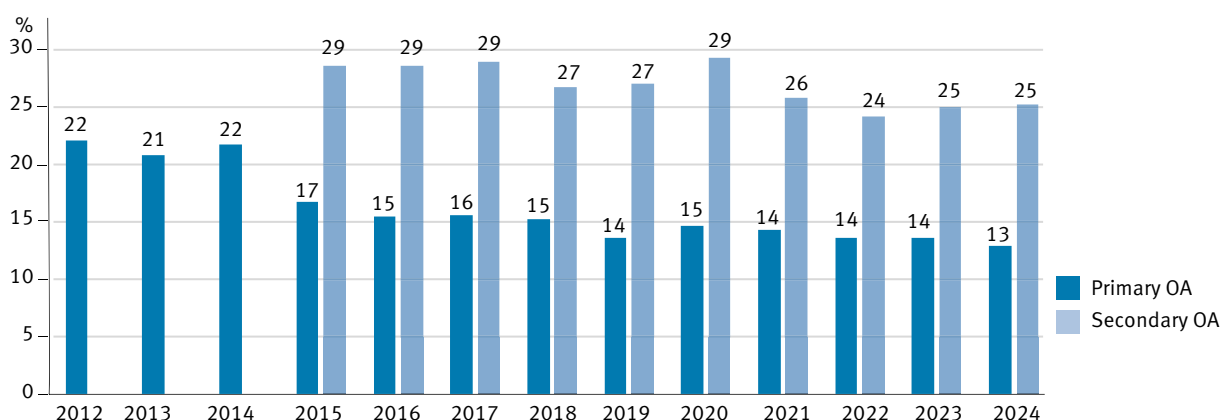


Figure 5.1

Share of TKA patients who had knee arthroscopy prior to arthroplasty (%)

Sec. OA categories were different before 2015 and therefore cannot be compared

* Including “arthritis after meniscus surgery”

Type of bearing

The rate of mobile polyethylene (PE) liners rapidly decreased over the past six years, from 36.5% in 2019 to 16.4% in 2024 (Figure 3.26). However, one must note that the choice of bearing type showed again a high variability among the different cantons, including the Principality of Liechtenstein (Figure 3.27) (see also chapter 3.3.5). As the Swiss joint registry started in 2012, most of the implanted polyethylene (PE) liners were made of highly cross-linked PE (HX-LPE) and comparison to conventional, ultra-high molecular weight PE does not make sense.

Patella resurfacing

The patella was not resurfaced in 63.1% of primary TKA during the period 2019 to 2024 (Figure 3.28). However, the resurfacing rate increased continuously from 32.0% in 2019 to 43.2% in 2024. However, there were again considerable differences between the cantons (see chapter 3.3.5).

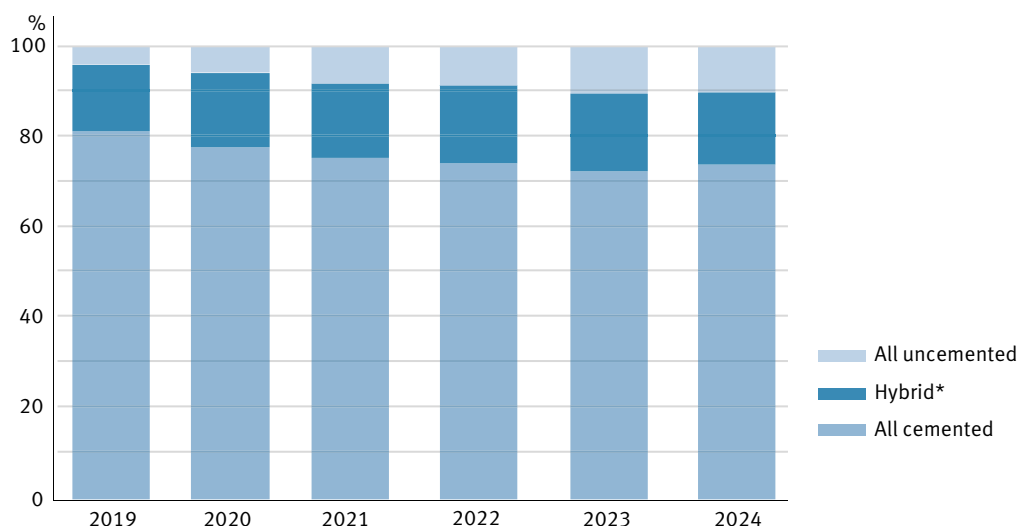
Fixation

Fixation of TKA was predominantly fully cemented, with a proportion of 75.7% over the past six years. Use of hybrid fixation of the components remained constant with 16.1%. Interestingly, cementless fixation represented only 4.0% of the TKA in 2019, but the share more than doubled within 6 years to 10.0% in 2024 (Figure 5.2).

Stems and additional components

More information about components in primary TKA was introduced with the 2021 version of the CRF, improving registering precision.

Stems (femoral and/or tibial) were used in 9.2% of primary TKA. In three quarters (75.4%) of the cases, this was on the tibial side (6.9% of all TKA), and 83.9% of the stems were cemented. Tibial stems were cemented in 86.7%, femoral ones in 75.4% (Table 5.1).



Component fixation [%]	2019	2020	2021	2022	2023	2024	2019 – 2024
N	15,607	15,467	16,814	19,315	20,706	21,568	109,477
All uncemented	4.0	5.5	8.0	8.5	10.2	10.0	8.0
Hybrid*	14.1	16.3	16.2	17.0	17.0	15.7	16.1
All cemented	81.5	77.7	75.6	74.4	72.6	74.1	75.7

Figure 5.2

Primary total knee arthroplasty: Component fixation

All diagnoses. * femur uncemented, tibia cemented

Use of a tibial stem was not associated with obesity (BMI ≥ 30 kg/m²), despite such recommendations from several studies. The use of stems, both on the tibial and femoral side, was mainly associated with higher intrinsic stability of the knee system (SC/CCK or hinge type) causing more stress at the interface between bone and implant. Stems were also used more frequently in PS than in the PCR/CR, CS, or MP designs (Figure 5.3). Over time there was nevertheless a clear increase in using stems in cemented tibial components since 2021 compared to former years (Figure 5.4).

Sleeves, cones, augments, or additional homologous or autologous bone grafting were rarely necessary in primary TKA (Table 5.1).

Additional interventions

Additional interventions were rarely performed during primary TKA (4.4% of the cases on average between 2019 to 2024). The removal of internal fixation devices (1.3%) and osteotomies of the tibial tubercle (1.4%) were the most common additional surgical steps, while 1.1% were reported as “other” (Table 5.1).

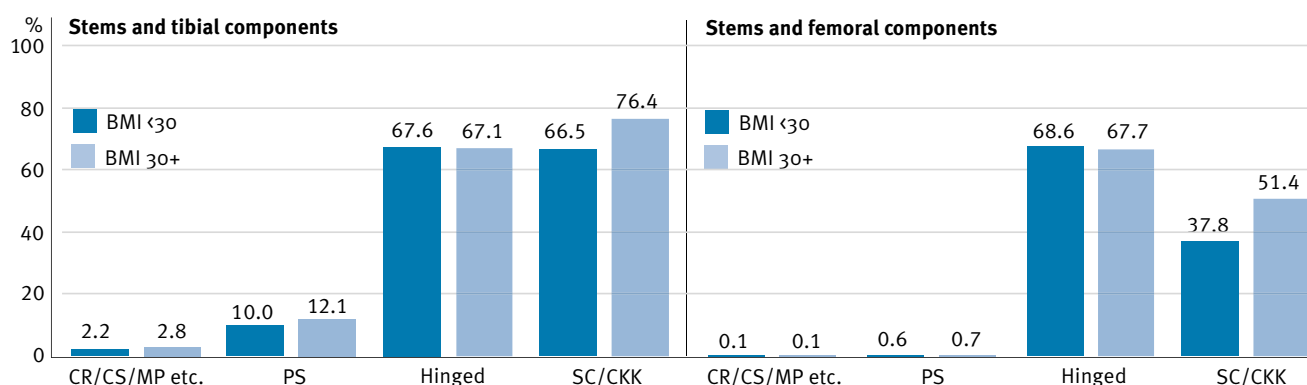


Figure 5.3
Use of stems by BMI and type of knee system (%)

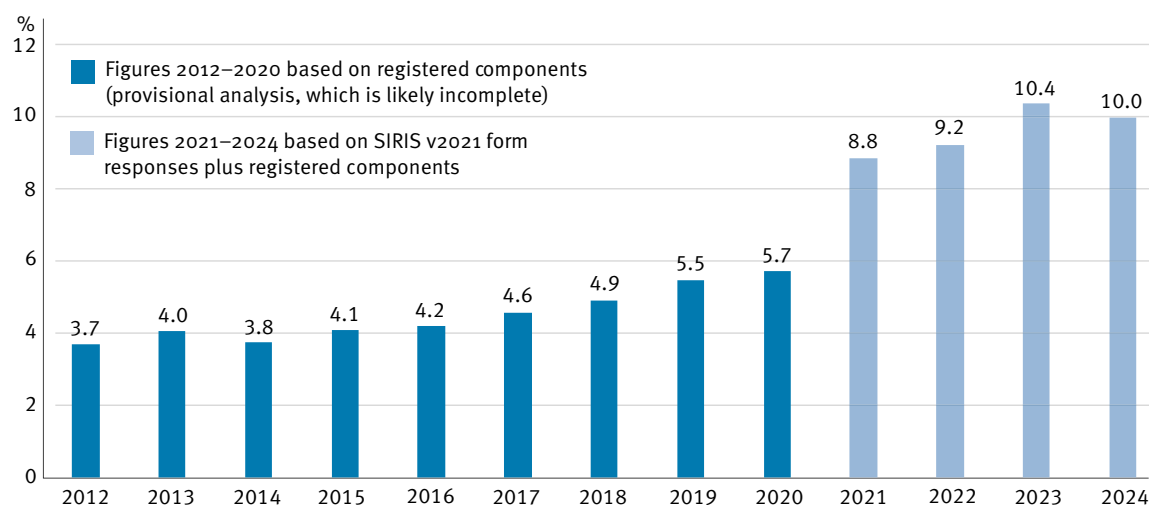


Figure 5.4
Use of stems as a percentage of primary TKAs with cemented tibial components

Can be femoral or tibial stems; form responses suggest 3:1 TI stems to FE stems

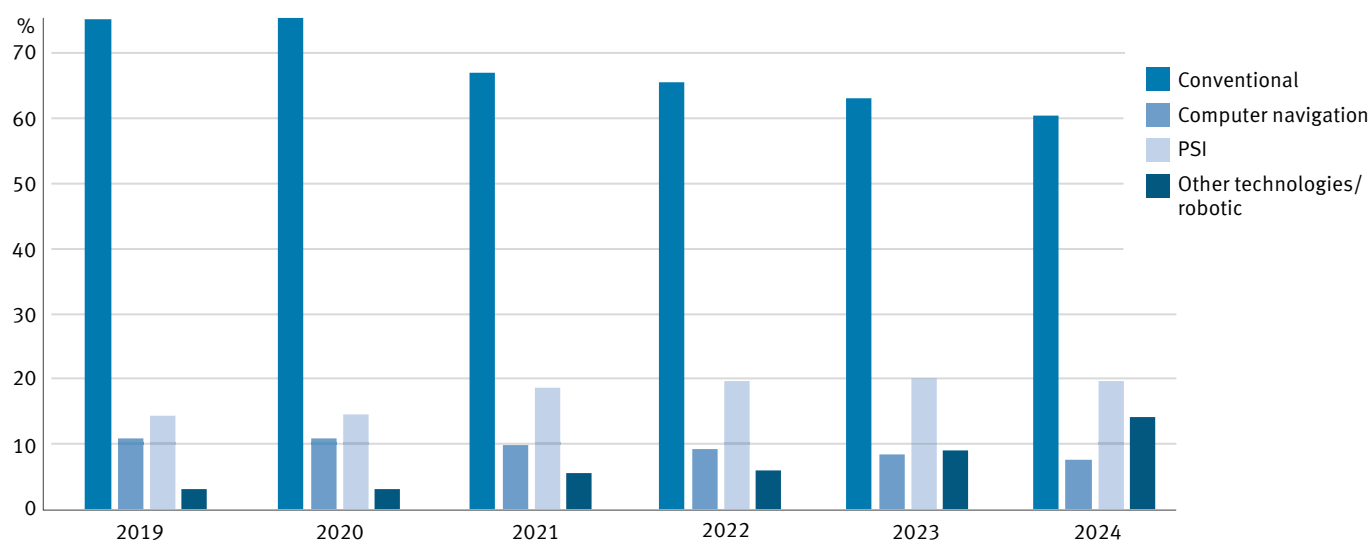
Technology

Between 2019 and 2024, 67.2% of the primary TKA in Switzerland were performed conventionally, without additional technological assistance. The share of conventional TKA decreased continuously from 70.9% in 2019 to 60.5% in 2024 (Table 5.1, Figure 5.5).

The share of computer navigation continuously decreased from 10.8% in 2019 to 7.5% in 2024. On the other hand, the use of patient-specific instrumentation (PSI) increased from 14.4% in 2019 and 14.6% in 2020 to approximately 20% in the following years

reaching 19.7% in 2024. Robotic-assisted TKA (imageless and image-based), formerly classified as “other”, increased from 3.1% in 2019 to 14.0% in 2024 (Figure 5.5). Of note, the different robotic systems are not yet differentiated due to limited numbers.

In summary, surgeons used technical support in 44.2% of TKA over the past 6 years (Table 5.1). Minimal invasive surgery is no longer a topic in Switzerland. It was removed from the CRF in 2021, as there was no uniformly accepted definition, limiting the reliability of the data.



Technology [%]	2019	2020	2021	2022	2023	2024	2019–2024
N	15,614	15,469	16,816	19,316	20,706	21,568	109,489
Conventional	75.8	75.5	66.8	65.5	63.1	60.5	65.8
Computer navigation	10.8	10.8	9.8	9.2	8.5	7.5	9.3
PSI	14.4	14.6	18.7	19.7	20.1	19.7	18.1
Other technologies/robotic	3.1	3.1	5.4	6.0	8.9	14.0	7.2

Figure 5.5

Primary total knee arthroplasty: technologies used

All diagnoses. Multiple responses possible (percentages do not sum to 100).

5.2 Revision of primary total knee arthroplasty implanted before 2012

Revisions that cannot be linked to an index arthroplasty registered in SIRIS are named “unlinked” revisions, whereas index arthroplasties were registered in SIRIS for “linked” revisions (see below).

The share of unlinked revisions steadily decreased from 96% in 2013 to 32.7% in 2023 (848 cases). The

proportion of unlinked TKA revisions over the whole observation period therefore was 46.7%. A total of 5,325 unlinked TKA revisions were performed between 2018 and 2023. The rate of unlinked revision TKA decreases more rapidly than in THA, most probably attributable to the fact that TKA are revised earlier and more frequently than THA.

Because of rapidly decreasing share of unlinked revisions, these older cases are of limited and continuously decreasing value and were not analysed further in this report.

		Primary	Revised within 24 months			
			Revised	95% CI		
		N at risk*	N	%**	lower	upper
Overall (moving average)		67,217	2,280	3.4	3.3	3.6
Diagnosis	Primary OA	59,018	1,942	3.3	3.2	3.5
	Secondary OA	8,199	338	4.2	3.8	4.6
Overall Primary OA		59,018	1,942	3.3	3.2	3.5
Gender	Women	35,999	1,131	3.2	3.0	3.4
	Men	23,018	811	3.6	3.3	3.8
Age group [%]	<55	2,894	154	5.4	4.6	6.2
	55–64	13,406	536	4.0	3.7	4.4
	65–74	21,634	722	3.4	3.1	3.6
	75–84	18,258	468	2.6	2.4	2.8
	85+	2,824	62	2.3	1.8	2.9
BMI group	<18.5	275	11	4.1	2.3	7.3
	18.5–24.9	11,147	310	2.8	2.5	3.1
	25–29.9	19,977	632	3.2	3.0	3.5
	30–34.9	13,185	450	3.5	3.2	3.8
	35–39.9	5,603	223	4.0	3.5	4.6
	40+	2,493	95	3.9	3.2	4.7
	BMI unknown	6,338	221	3.5	3.1	4.0
Morbidity state	ASA 1	3,797	113	3.0	2.5	3.6
	ASA 2	34,889	1,129	3.3	3.1	3.5
	ASA 3	17,181	609	3.6	3.3	3.9
	ASA 4/5	250	11	4.6	2.6	8.2
	ASA unknown	2,901	80	2.8	2.2	3.5

Table 5.3
First revision of primary total knee arthroplasty within 24 months overall and according to baseline characteristics

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

* Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

** Rates adjusted for effects of mortality and emigration.

5.3 First revision within two years after primary TKA

Following international guidelines, a revision is defined as the addition or the exchange of any component. Therefore, a secondary patella resurfacing is a revision by definition. SIRIS has been recording all revision procedures since 2012, irrespective of whether it was the first or any subsequent revision. Linked revisions form the basis for calculations of survival and revision rates and may be separated into first revisions and repeat revisions. The first revision serves to define primary implant survival, but only primary OA is used for this benchmark. Cases with isolated previous meniscus surgery were also included in the group of primary OA.

Other causes of secondary OA, such as previous ligament surgery, fracture fixation, osteotomy or inflammatory arthritis, etc., were excluded, as associated revision rates may be influenced by and increased due to the underlying diagnosis.

In the current 4-year moving window from 1.1.2019 to 31.12.2022 with a complete 2-year follow-up until 31.12.2024, a total of 67,217 TKA have been registered. Of those, 2,280 were revised within two years, resulting in a 2-year revision rate of 3.4% (CI 3.3 – 3.6%) (Table 5.3).

For incidence and demography please consult **chapter 3.3.4**.

5.3.1 Reasons for first revision

The most frequent reason for first revision were patella problems, registered in 36.0%. Additionally, an instability of the patella was reported in 4.4%. Infection (21.5%) and femorotibial instability (18.5%) were the second and third most frequent reasons. Loosening of the tibial component was indicated in 7.7% of the cases, joint stiffness in 8.6% and pain of unclear origin in 3.6%. Wear of the liner was reported rarely (0.7%) but represents a surprising reason for early revision (Table 5.4). Periprosthetic fractures of

the femur, tibia, and/or patella were rarely responsible for early revision. However, most cases treated only with internal fixation were probably not registered, respectively do not represent revisions following the usual definition. Still, 10.5% of the reasons for revision were classified as “other”. This diverse group mostly includes the same reasons as listed above, but with added details and includes wound healing problems and further reasons, such as liner dislocation. Excluding periprosthetic infections and fractures, most of the other reasons appeared to be related to surgeon’s errors in technique and/or indication.

Reason for revision	N	%
Patella problems	821	36.0
Infection	490	21.5
Femorotibial instability	422	18.5
Loosening tibia	176	7.7
Pain (of unclear origin)*	81	3.6
Joint stiffness/arthrofibrosis	195	8.6
Component malposition femur	89	3.9
Component malposition tibia	112	4.9
Loosening femur	86	3.8
Patellar instability	100	4.4
Wear of inlay	15	0.7
Loosening patella	19	0.8
Periprosthetic fracture femur	29	1.3
Sizing femoral component	41	1.8
Periprosthetic fracture tibia	19	0.8
Sizing tibial component	18	0.8
Periprosthetic fracture patella	14	0.6
Other	240	10.5
Total Revisions	2,280	

Table 5.4

Reason for revision* of primary total knee arthroplasty

All diagnoses. 4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024). Early first revisions are those occurring within 2 years of the primary arthroplasty. Multiple responses possible (percentages do not sum to 100)

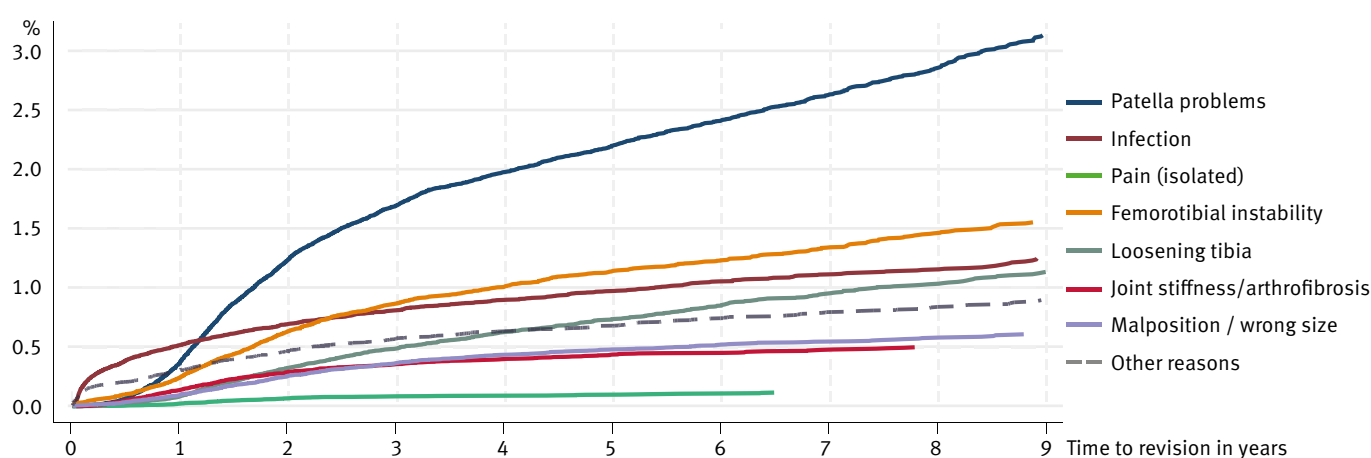
* Pain was frequently reported alongside other reasons. The proportion of “isolated pain” was 3.2%.

The wording was adapted in v2021 and the share dropped accordingly in 2021

Cumulative incidence rates of revision from 2015 to 2024 show patella problems to be the most frequent reason for revision within one year after TKA, and corresponding revision rates increase thereafter more rapidly than for all other reasons (Figure 5.6).

Kernel density estimation – an evaluation of the frequency of an event at a given time – depicts that infections were the most prominent reason for revision early after primary TKA, later declining, whereas

peaks for other reasons such as patella problems, femerotibial instability, isolated pain of unknown origin and loosening of the tibial component, appeared after 9 months follow-up to culminate at 15 months, reflecting the usual pattern in patients with unsatisfactory results after TKA, “wait and see” often being recommended initially. The bell-like pattern shows the increasing revision rates over time, in what might resemble logistic growth curves (slow increase fol-



Revision diagnosis	1 year	2 years	3 years	5 years	6 years	8 years	9 years
Patella problems	0.4 (0.3-0.4)	1.2 (1.2-1.3)	1.7 (1.6-1.8)	2.2 (2.1-2.3)	2.4 (2.3-2.5)	2.9 (2.7-3.0)	3.1 (3.0-3.3)
Infection	0.5 (0.5-0.6)	0.7 (0.7-0.7)	0.8 (0.8-0.9)	1.0 (0.9-1.0)	1.1 (1.0-1.1)	1.2 (1.1-1.2)	1.2 (1.2-1.3)
Pain (isolated)	0.0 (0.0-0.0)	0.1 (0.1-0.1)	0.1 (0.1-0.1)	0.1 (0.1-0.1)	0.1 (0.1-0.1)	0.1 (0.1-0.1)	0.1 (0.1-0.1)
Femerotibial instability	0.2 (0.2-0.3)	0.6 (0.6-0.7)	0.9 (0.8-0.9)	1.1 (1.1-1.2)	1.2 (1.2-1.3)	1.5 (1.4-1.5)	1.6 (1.5-1.7)
Loosening TI	0.1 (0.1-0.1)	0.3 (0.3-0.4)	0.5 (0.5-0.5)	0.7 (0.7-0.8)	0.9 (0.8-0.9)	1.0 (1.0-1.1)	1.1 (1.0-1.2)
Joint stiffness / arthrofibrosis	0.1 (0.1-0.2)	0.3 (0.3-0.3)	0.4 (0.3-0.4)	0.4 (0.4-0.5)	0.5 (0.4-0.5)	0.5 (0.5-0.5)	0.5 (0.5-0.5)
Malposition / wrong size	0.1 (0.1-0.1)	0.3 (0.2-0.3)	0.4 (0.3-0.4)	0.5 (0.4-0.5)	0.5 (0.5-0.6)	0.6 (0.5-0.6)	0.6 (0.6-0.7)
Other reasons	0.3 (0.3-0.3)	0.5 (0.4-0.5)	0.6 (0.5-0.6)	0.7 (0.6-0.7)	0.7 (0.7-0.8)	0.8 (0.8-0.9)	0.9 (0.8-1.0)

Figure 5.6

Cumulative incidence rates for different revision diagnosis of primary total knee arthroplasty

Time since operation, 2015–2024, all services, % of implants revised. Detailed reasons for revisions available since 2015

lowed by steeper growth and finally flattening-out). Patella problems contributed to the observed pattern of revisions, causing a disproportionate number of revisions between 11.3 and 18.8 months after implantation, with the peak at 14.9 months. Revision for joint stiffness showed a flatter pattern, with a maximum at 12.4 months (Figure 5.7 and Table 5.5).

5.3.2 Fixation and first revision

Uncemented TKA were revised significantly more often (4.2%, CI 3.6 – 4.8%) than fully cemented TKA (3.4%, CI 3.2 – 3.5%) within the first two years after the index surgery (Table 5.6). The wider confidence interval of uncemented TKA reflects the smaller number of cases (the share of uncemented TKA was 10.0% only in 2024). The difference among the revision rates of the various fixation types was statistically significant only the first three years after TKA (Figure 5.8). It seems that the higher revision rate of uncemented TKA is mainly due to an increased risk for up to three years after operation, perhaps reflecting failed osteointegration or failures due to malalign-

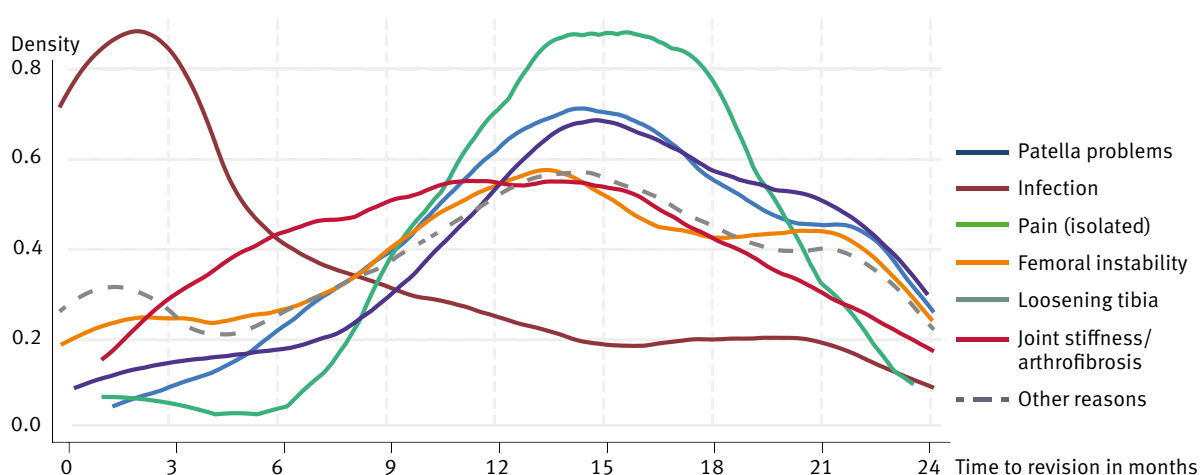


Figure 5.7

Time interval between primary total knee arthroplasty and first revision by reason

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

Reason for revision	N	Median	IQR 25%	IQR 75%
Patella problems	821	14.9	11.3	18.8
Infection	490	4.5	1.2	11.7
Pain (isolated)	24	14.9	12.0	17.4
Femoral instability	422	13.5	8.4	18.9
Loosening tibia	176	15.5	11.8	19.7
Joint stiffness/arthrofibrosis	195	12.4	8.1	17.1
Other	1,290	13.4	7.9	18.0

Table 5.5

Median time interval between primary total knee arthroplasty and early first revision (in months) according to reason

All diagnoses. Early first revisions are those occurring within 2 years of the primary arthroplasty.

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

ment and/or insufficient bone quality. From the 4th year after index surgery onwards, the curve is moving on a higher level than for the cemented and the hybrid fixation TKA but evolving in parallel. All curves seem to converge 8 years after surgery (Figure 5.8). Hybrid fixation performed as well as fully cemented TKA, with a 2-year revision rate of 3.4% (CI 3.1 –

3.8%) and lost therefore advantages regarding revision rates observed in former SIRIS reports. Younger age (<60 years) was correlated with an elevated early revision rate, irrespective of the fixation method (see above). It is important to consider that all TKA brands were included in this KM estimation of cumulative revision risk, mixing in various effects.

		Primary TKA		Revised within 24 months		
			Revised	95% CI		
		N at risk ¹	N	% ²	lower	upper
Component fixation	Overall	67,217	2,280	3.4	3.3	3.6
	All cemented	51,824	1,723	3.4	3.2	3.5
	All uncemented	4,452	185	4.2	3.6	4.8
	Hybrid ³	10,732	363	3.4	3.1	3.8
Patellar replacement	Reverse hybrid ⁴	197	8	4.1	2.1	8.0
	With patellar replacement	22,877	637	2.8	2.6	3.0
	Without patellar replacem.	44,303	1,642	3.8	3.6	3.9
	Status after patellectomy	25	0	0.0		

- ¹ Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).
- ² Rates adjusted for effects of mortality and emigration.
- ³ femur uncemented, tibia cemented
- ⁴ femur cemented, tibia uncemented

Table 5.6
First revision of primary total knee arthroplasty within 24 months overall and according to component fixation

All diagnoses. 4-year moving average covering impl. between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

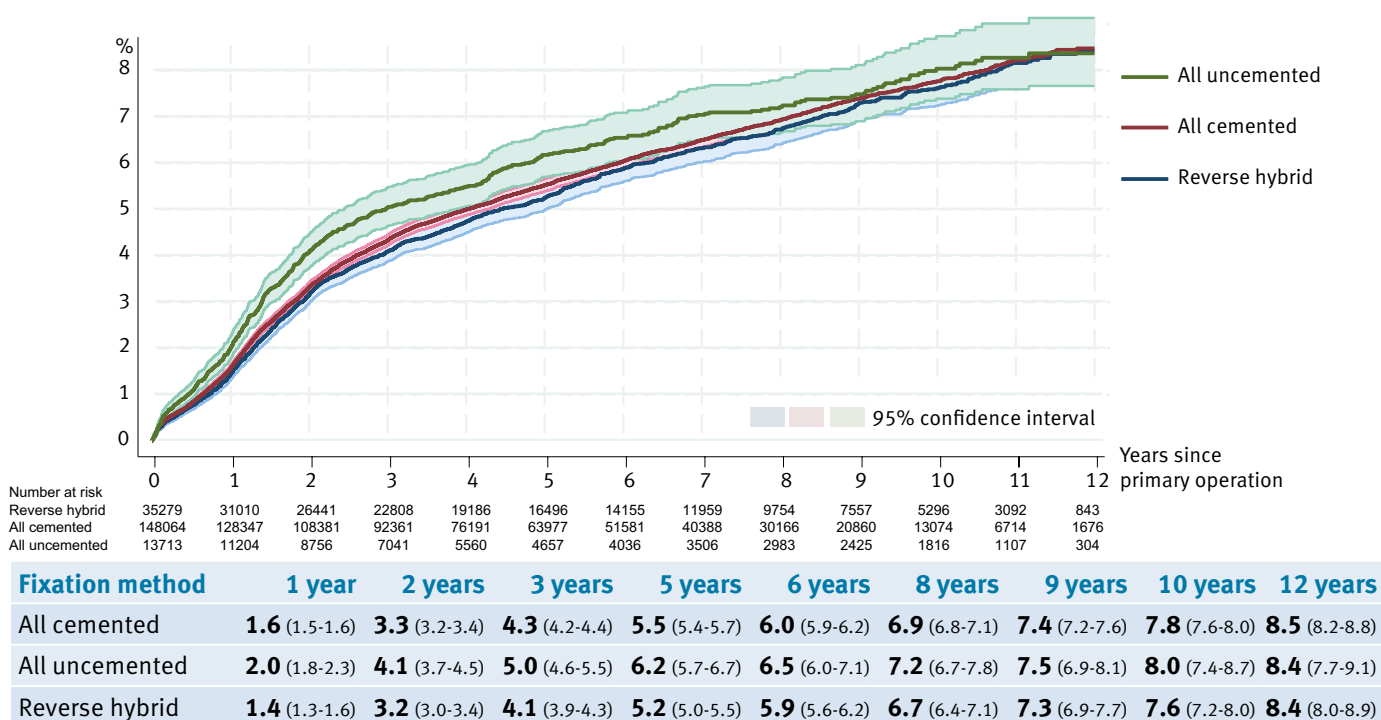


Figure 5.8
Estimated failure rates of primary total knee arthroplasty for different fixation methods

Time since operation, 2012–2024, all diagnoses.

5.3.3 The influence of the type of knee arthroplasty, the use of stems and the BMI on first revision

At two years of follow-up, differences in revision rates among types of TKA were small. Only PS showed a significantly higher revision rate (3.7%, CI 3.5 – 3.9%) than PCR/CR, UCOR or MP (Figure 5.9). Different systems grouped as “other arthroplasty” had a higher

early revision rate than all other TKA systems but this difference remained statistically not significant due to small numbers and therefore wider confidence intervals (4.1%, CI 3.7 – 4.6%). Up to mid-term, CS/UCOR seemed to perform best. This could be partially explained by a certain selection bias, as PS knees are usually selected for more complex cases with more pronounced deformities and ligament insufficiencies.

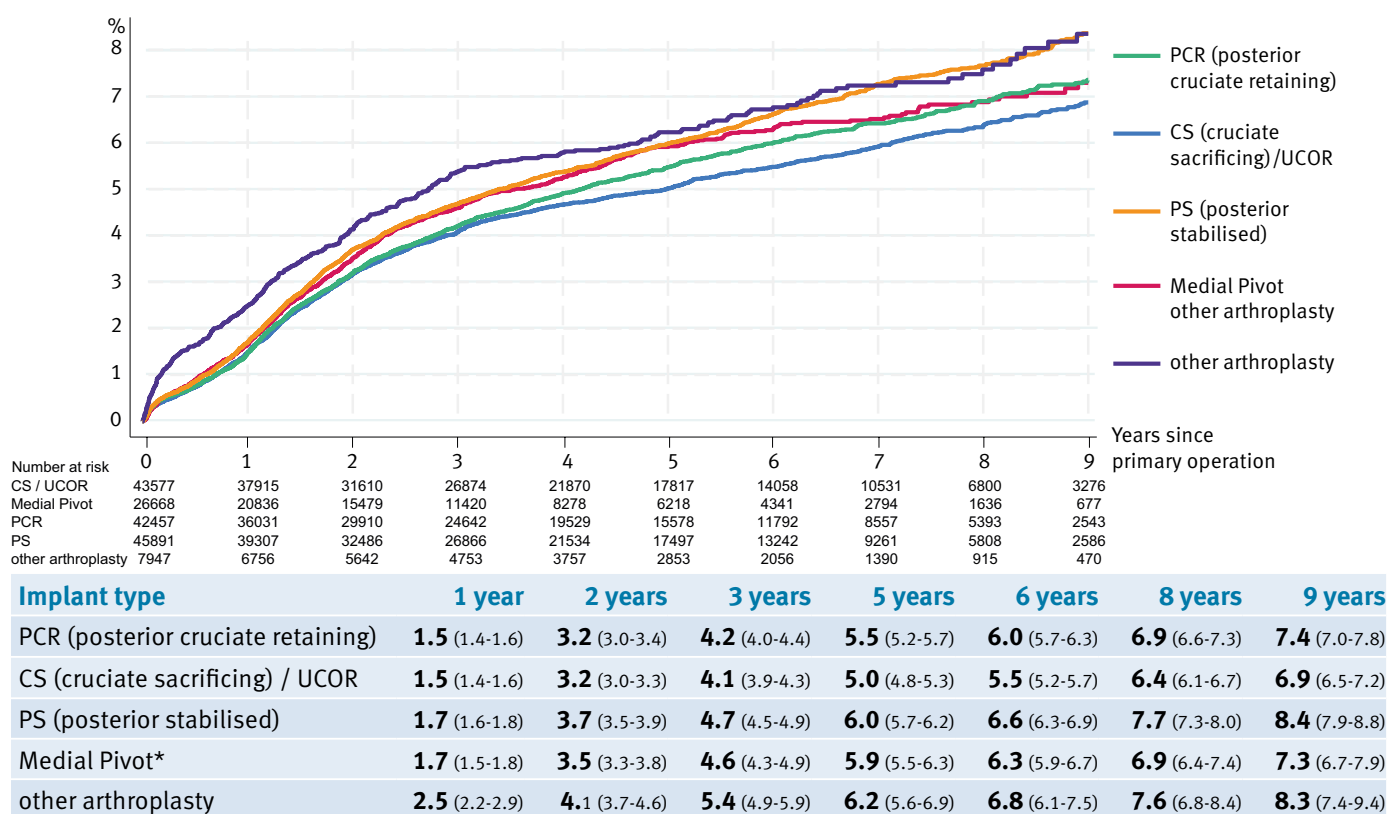


Figure 5.9
Estimated failure rates of primary total knee arthroplasty for different implant types

Time since operation, 2015–2024, all diagnoses. Confidence intervals not shown in graphic for reasons of clarity.

*Medial pivot was not available as a response category before SIRIS v2021. In the annual report 2020, only free text “other” responses were identified as and recoded to medial pivot. However, this missed a number of GMK Sphere total knee systems that were incorrectly registered as other types, mainly CS/UCOR. In this report, all GMK Sphere knee systems are counted as medial pivot, regardless of the type chosen locally at data entry.

As the vast majority of stems were used in primary TKA with cemented tibial components, the analysis was limited to this subgroup for the TKA types PCR/CR, CS, PS or MP. The revision rates did not differ with or without stem when the BMI was less than 30 kg/m². According to the results in the registry, the use of stems in patients with BMI ≥30 kg/m² was associated with an even higher revision rate compared to no stem use (Figure 5.10), though the difference never

was statistically significant. This was not expected, as the literature is reporting significantly lower revision rates when using tibial stems in obese patients. The opposite observation in SIRIS could speak for a certain selection bias with other influencing factors than BMI alone (e.g. osteoporosis). Definitive conclusions however may not be drawn, as further subgrouping would be necessary, which is currently impossible due to small numbers.

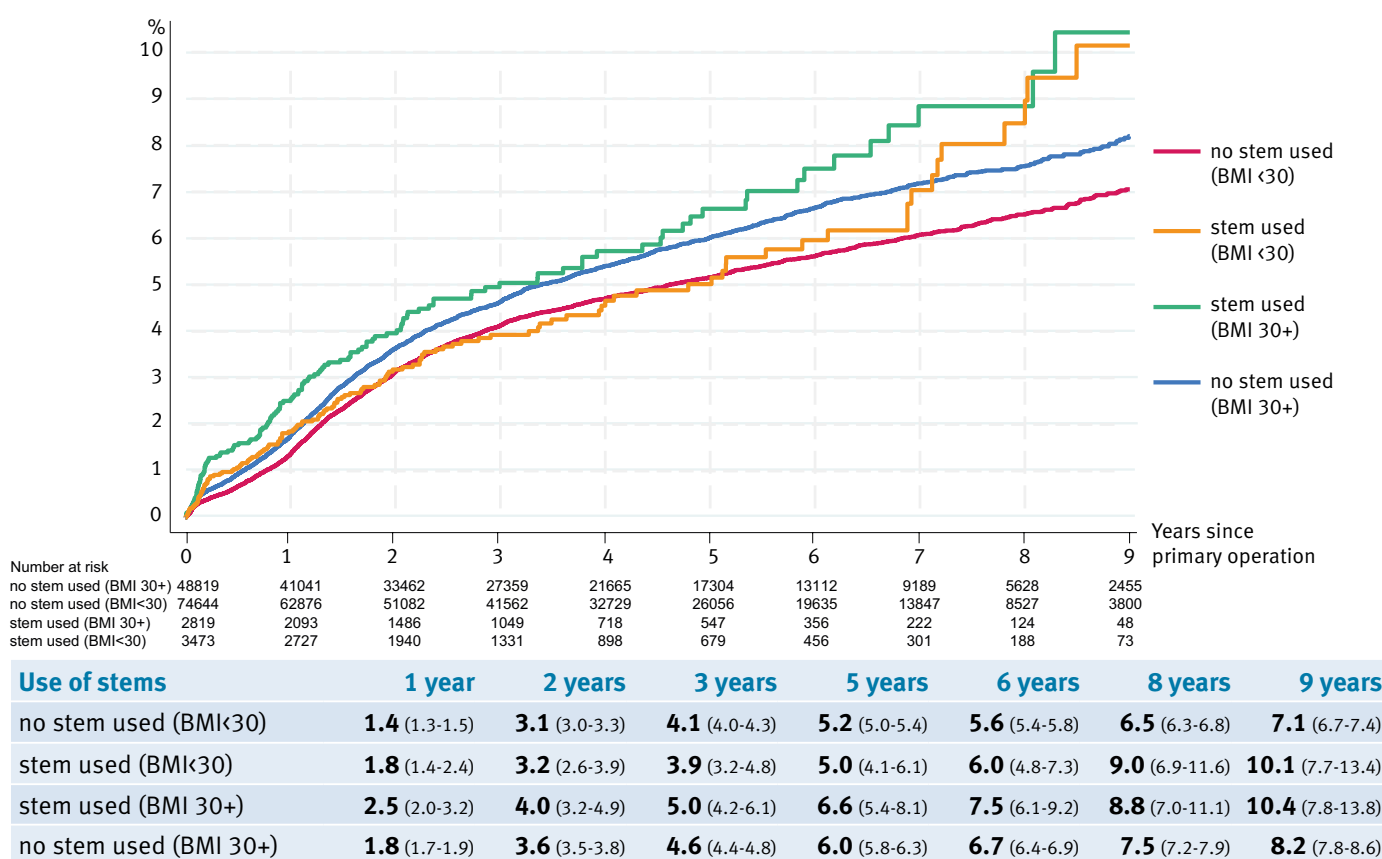


Figure 5.10
Estimated failure rates of primary total knee arthroplasty: use of stems (CR/CS/PS/MP)

Time since operation, 2015–2024, all diagnoses, only cemented tibias.

For primary TKA with higher intrinsic stability (SC/CCK or hinge type) survival up to 9 years after TKA was comparable with or without stems for all types of subsystems when the BMI was lower than 30 kg/m². Again, stemmed constrained TKA with BMI >30 kg/m² was associated with higher revision rates from the 4th year after index surgery but not reaching statistical significance due to small numbers and therefore wide confidence intervals (Figure 5.11).

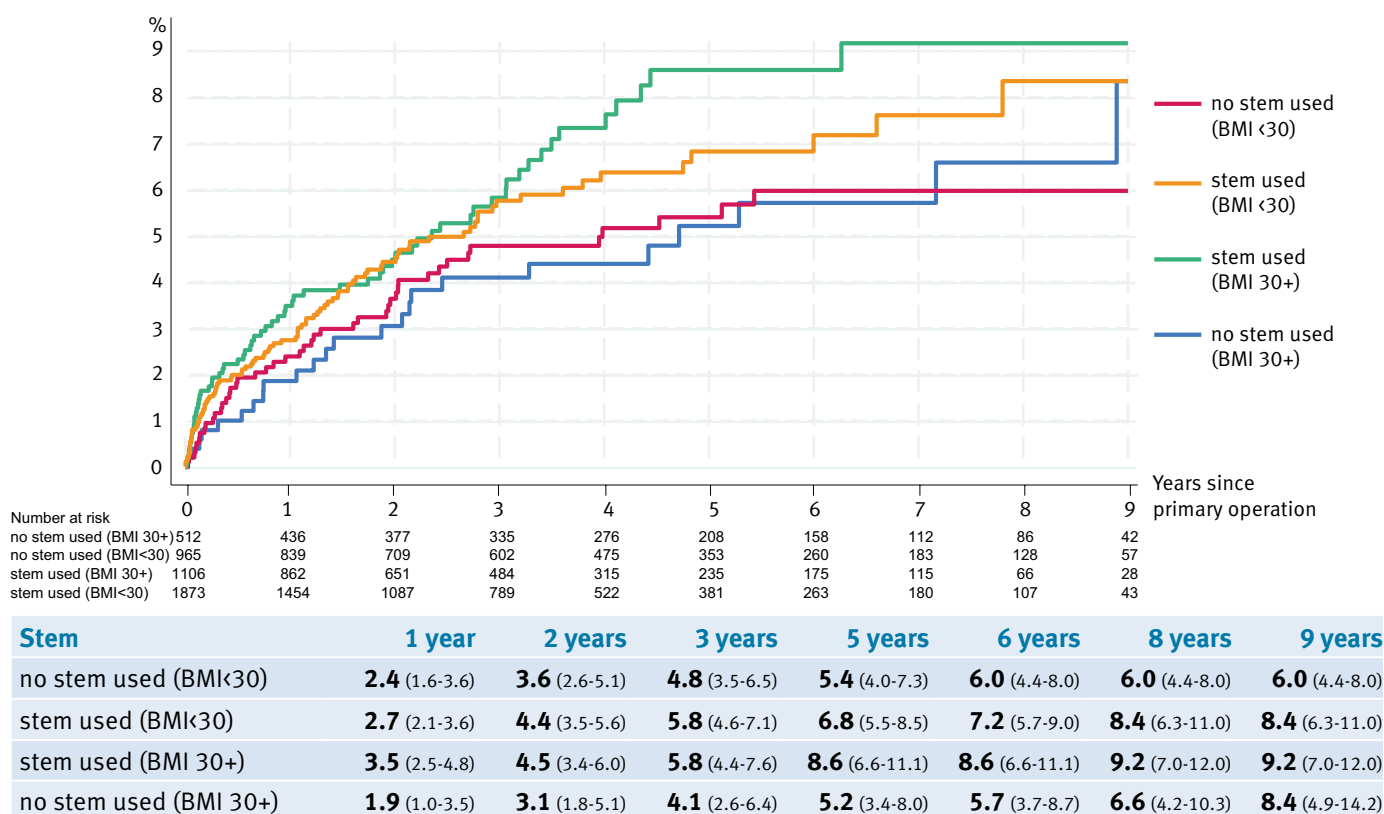


Figure 5.11

Estimated failure rates of primary total knee arthroplasty: use of stems (Hinged, SC/CCK)

Time since operation, 2015–2024, all diagnoses, only cemented tibias of primary TKA with increased intrinsic stability (Hinge, SC/CCK).

5.3.4 Patella resurfacing and first revision

There was clearly a higher probability of revision when the patella was not resurfaced initially, becoming significant from the first year on and remaining significant at up to twelve years of follow-up (Figure 5.12). Early revision rates at two years after primary TKA differed significantly from 2.8% (CI 2.6 – 2.9%) to 3.6% (CI 3.5 – 3.7%), respectively. Between 3 and 12 years after TKA, the revision rates developed in a more linear manner for TKA without and with patella resurfacing, still worsening if patella was not resurfaced (Figure 5.12).

5.3.5 Type of surgery at first early revision

Complete revision of both femoral and tibial components was performed in 22.7% of the cases when revision had to be performed within two years of follow-up. In 26.5% of the cases, only the liner was exchanged. Isolated tibial revision was registered in 5.7%, and isolated femoral revision in 3.7%. Patella surgery is discussed below. Component removal with spacer implantation was recorded in 3.3%, without spacer in 0.2%, reimplantation of prosthesis in 1.4%, all three representing surgical procedures for periprosthetic infections.

Type of implants

With 30.4%, SC/CCK systems formed the largest group of implants used at first TKA revision, followed by PS in 23.5% and hinge-type prosthesis in 21.5%. PCR/CR TKA were used in 6.5% of the revisions, 8.7% were classified as CS or UCOR implants, whereas MP was used in 5.9% of the revisions (Table 5.7).

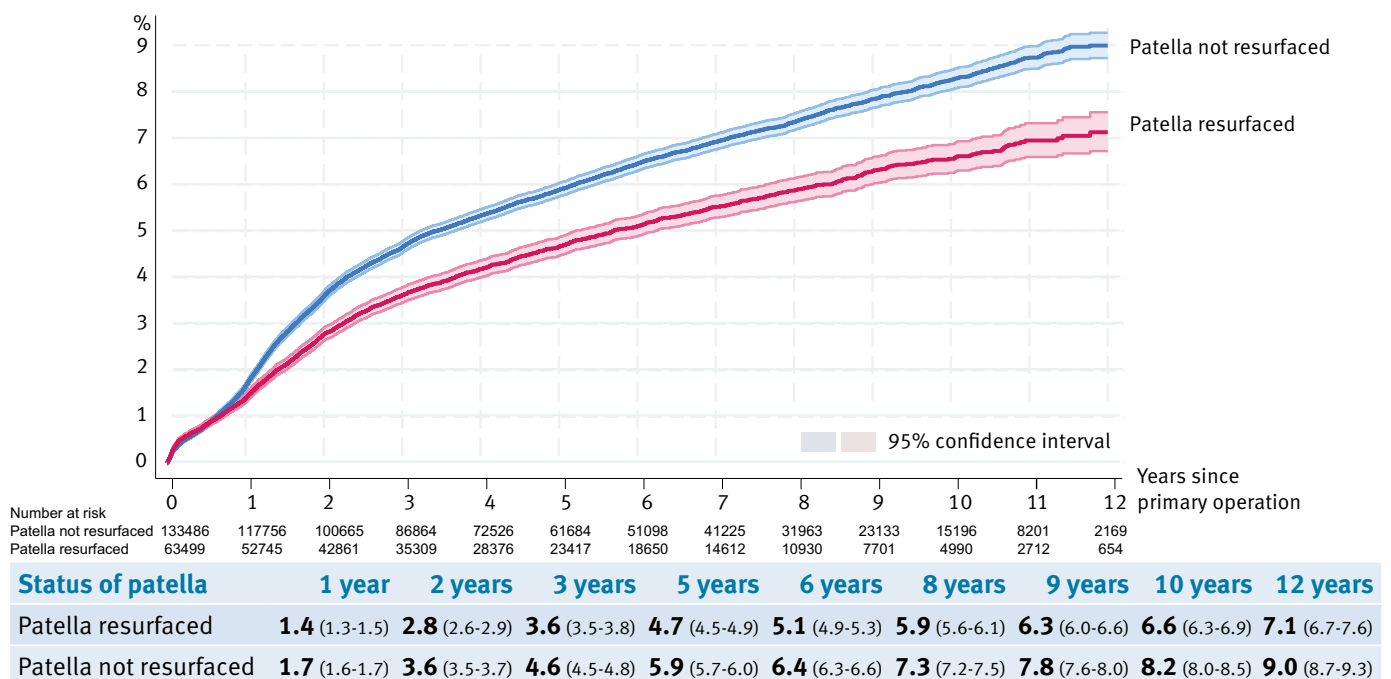


Figure 5.12

Estimated failure rates of primary total knee arthroplasty: status of patella after primary operation

Time since operation, 2012–2024, all diagnoses.

Patella

Early revision TKA was associated with patella resurfacing in 76.7% of cases, but this includes cases where the primary patella component was left in

place. Considering the different types of revisions, the combined percentage of procedures where a patella component could have been replaced or added barely exceeds 60% of first revisions. (i.e. complete

Intervention type ¹	N	%
complete revision	518	22.7
exchange of PE	604	26.5
subsequent patella prosthesis	530	23.2
tibial revision	131	5.7
reimplantation of prosthesis	33	1.4
subsequent patella prosthesis with exchange of PE	148	6.5
patella revision	123	5.4
component removal with spacer implantation	76	3.3
femoral revision	84	3.7
prosthesis preserving revision	2	0.1
osteosynthesis	2	0.1
arthrodesis	29	1.3
component removal without spacer implantation	11	0.2
reconstruction after injury of extensor mechanism	4	0.1
plastic reconstruction	4	0.1
other	104	2.0
Additional intervention	N	%
None	1,323	78.6
Osteosynthesis FE	11	0.7
Osteosynthesis TI	4	0.2
Osteosynthesis PAT	1	0.1
Removal of metalware	18	1.1
Operation extensors	76	4.5
Reconstruction plasty	20	1.2
Tibial tubercle osteotomy	133	7.9
Other additional intervention(s)	137	8.1
Total revisions (multiple responses)	1,683	

Type of arthroplasty	N	%
SC / CCK (semi-constrained / constrained)	238	30.4
Hinge type	168	21.5
PS (posterior stabilized)	184	23.5
CS (cruciate sacrificing) / UCOR	68	8.7
PCR (posterior cruciate retaining)	51	6.5
Medial-Pivot ²	46	5.9
BCR (bicruciate retaining)	8	1.0
Other	20	2.6

Technology	N	%
Conventional	2,123	97.5
Computer assisted / navigation	23	1.1
Patient specific instrumentation	23	1.1
Robotic-assisted (v2021)	2	0.1
Other	11	0.5

Additional components ³	N	%
Stem FE (cemented)*	303	38.8
Stem FE (uncemented)**	150	19.2
Stem TI (cemented)***	349	44.7
Stem TI (uncemented)***	165	21.2
Sleeve FE	32	4.1
Sleeve TI	103	13.2
Augments FE	239	30.6
Augments TI	86	11.0
Augments PAT	0	0.0
Bone homologous	11	1.4
Bone autologous	21	2.7
Cone FE	6	0.8
Cone TI	26	3.3
Total revisions (multiple responses)	780	

Table 5.7

Early first revision of primary total knee arthroplasty: Surgery characteristics

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024)

¹ includes a small proportion of reoperations that are not counted as component revisions in the evaluative parts of this report

² Entered as "other" intervention and then recoded before 2021. As of form version 2021, SIRIS lists Medial Pivot as a separate main category

³ After complete, FE, TI revisions or component reimplantations. Detailed data available since 2021, but main categories available since 2015.

* 61% with cement restrictor

** 27% with coating

*** 52% with cement restrictor

**** 30% with coating

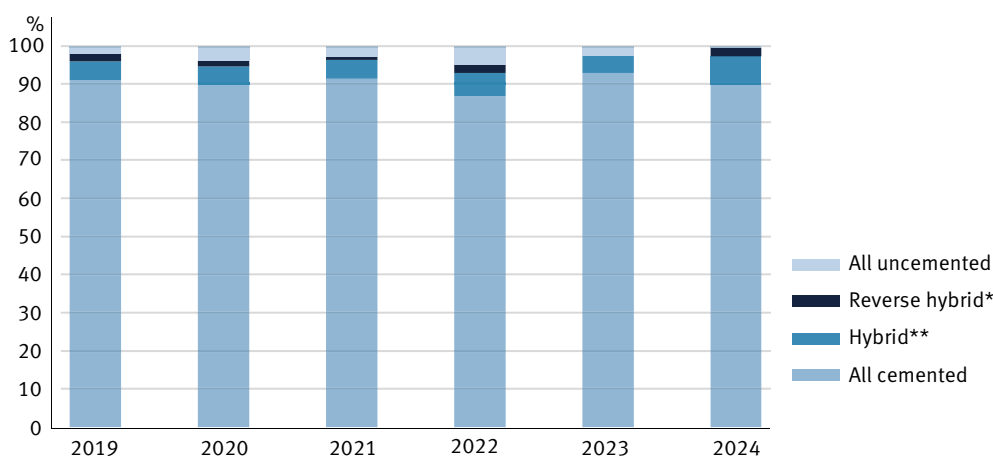
revisions, secondary resurfacing and patella revision). An exact count of how many patella components were added at complete revision is limited by incomplete registration of the primary components, but is likely in the area of 55% or higher.

Isolated patella resurfacing was performed in 23.2% of the first revisions. An additional 6.5% had secondary patella resurfacing combined with liner exchange (Table 5.7). The share of primary patella resurfacing not revised at revision TKA remains unknown, possibly explaining the rather low rate of patella resurfacing at first revision, although patella problems were

the most frequent reason for reoperation. A revision of a patella component was registered in 5.4%, making any patella revision the most frequent component revision. However, reasons for revision cannot be simply added up, as several options can be combined at registration.

Fixation

Fixation of the revision implants was fully cemented in the vast majority, with a mean value of 90.3% over the last 6 years, fluctuating between 86.8% and 92.8% since 2019 (Figure 5.14).



Component fixation [%]	2019	2020	2021	2022	2023	2024	2019–2024
N	200	184	186	182	195	186	1,133
All uncemented	2.0	3.8	2.7	4.9	2.6	0.5	2.7
Reverse hybrid*	2.0	1.6	1.1	2.2	0.0	2.2	1.5
Hybrid**	5.0	4.9	4.8	6.0	4.6	7.5	5.5
All cemented	91.0	89.7	91.4	86.8	92.8	89.8	90.3

* femur cemented, tibia uncemented

** femur uncemented, tibia cemented

Figure 5.14

Early first revision of primary total knee arthroplasty: Component fixation

Component fixation only applicable when new components were implanted. Early first revisions are those occurring within 2 years of the primary arthroplasty.

Stems and additional components

Since 2021, increasing numbers of tibial and femoral stems were used (Figure 5.4 and 5.15). Augments were also used more frequently on the femoral side, stayed on the same level on the tibial side. Tibial and femoral sleeves were constantly used since 2015 (Figure 5.15).

At first TKA revision, 65.9% received a tibial and 58.0% a femoral stem, and 67.9% of the tibial and 66.9% femoral stems were cemented. Femoral augments were used in 30.6%, tibial ones in 11.0%. Reinforcement of the metaphysis by tibial sleeves was registered in 13.2%, femoral ones in 4.1%. Cones were rarely used, in 3.3% at the tibial and in 0.8% at the femoral level. Cones were used less frequently as sleeves. Whereas cones are available in most of the brands providing revisions systems, sleeves are mainly associated with one brand. Therefore, this is influenced mainly by the selection of revision TKA used in Switzerland. Bone grafting was used in 4.1% only of the first revisions, and then autologous bone in 65.6% and can be considered as replaced mostly by stems, augments, cones and sleeves (Table 5.7).

Additional interventions

In 78.6% of first revisions, no additional intervention was necessary. Tibial tubercle osteotomy was performed in 7.3% of the cases, compared to 1.4% in primary TKA. In 4.5% of the cases, an additional intervention at the extensor apparatus was performed, while a formal reconstruction of the extensor apparatus was registered in 1.2%. Removal of orthopaedic or fracture fixation devices was done in combination with revision in only 1.1%. In 8.1%, “other” additional interventions were recorded (Table 5.7).

Technology

Technology was mostly conventional (97.5%), technical assistance was registered rarely (2.3% for all technological together). Computer navigation (1.1%) or PSI (1.1%) did play minor rolls. Robotic assistance (0.1%) would technically be possible in revision TKA but may be hindered by medicolegal constraints in some of the systems available (Table 5.7).

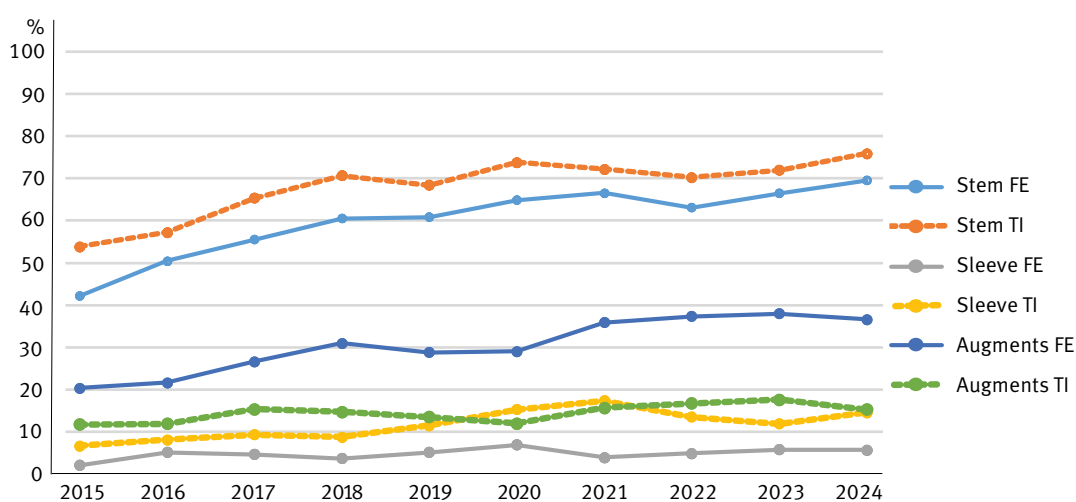


Figure 5.15
Selected additional components used in early first revisions (%)

5.4 Re-revision of total knee arthroplasty

Re-revisions after revision TKA were considered when they occurred after partial or complete revisions, re-implantations or isolated exchange of the liner. We call these first revisions “index revisions” for this analysis. Please note that this does not include revisions after conversions of PKA to TKA, which are presented in a separate analysis (see 5.4.2).

5.4.1 Reasons for TKA re-revision

As multiple reasons for revision may be registered, individual rates do not add up to 100%. This may also lead to under- or overestimation of some topics. Cumulative repeated revision rates after TKA revision from 2015 onwards are depicted in **Figure 5.16**. Infection took the lead early on as reason for re-revision, followed by a clear gap by the equally prevalent problems of femorotibial instability, patella problems, loosening of the tibial component and “other reasons”. Interestingly, patella problems still played an important role in TKA re-revisions, even while one would assume the patella was resurfaced either at primary TKA or at the first revision. The share of unresurfaced patellae at time of re-revision is unknown, as is the rate of revision for malposition or wear/

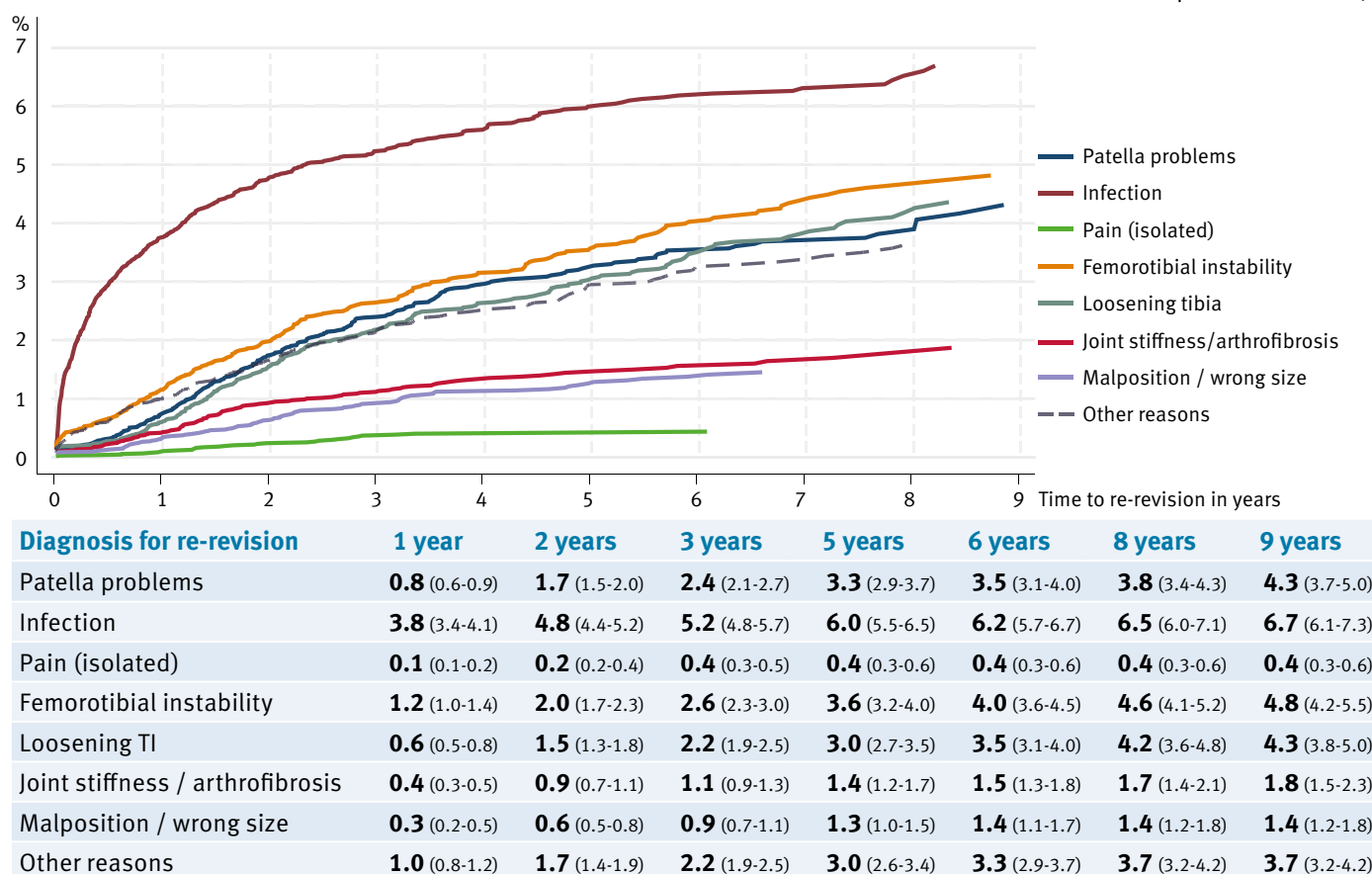


Figure 5.16

Cumulative incidence rates for different re-revision diagnosis of primary total knee arthroplasty

Time since revision, 2015–2024, % of implants re-revised. Detailed reasons for revisions available since 2015.

Comprises of all complete revisions, partial revisions, reimplantations and PE replacements.

loosening of patella buttons. Joint stiffness seemed to play a minor role, as did malposition or wrong sizing of the components. Presumably the frustrating results after revision for a stiff knee after TKA are better known meanwhile, preventing further surgery for this indication. Isolated pain of unknown origin was rarely the reason for a re-revision.

5.4.2 Type of previous TKA revision and TKA re-revision

Early, 2-year re-revision rates reached 10.7% (CI 9.5 – 12.0%) for partial and 8.3% (CI 7.7 – 9.0%) for complete revision respectively, whereas it was 3.4% (CI 3.3 – 3.6%) after revision of primary TKA (Table 5.3). If only the liner was exchanged at the first revision, the early re-revision rate was 16.9% (CI 15.7 – 18.3%) significantly worse compared to partial or complete revision. The re-revision rate at 12 years was 23.3% (CI 21.0 – 25.8%) for partial and 20.8% (CI 19.4 – 22.3%) for complete revision (Figure 5.17). This rate is almost three times higher than after primary TKA (Figure 3.21). Despite cumulative re-revision risk curves appearing to run separately from each other,

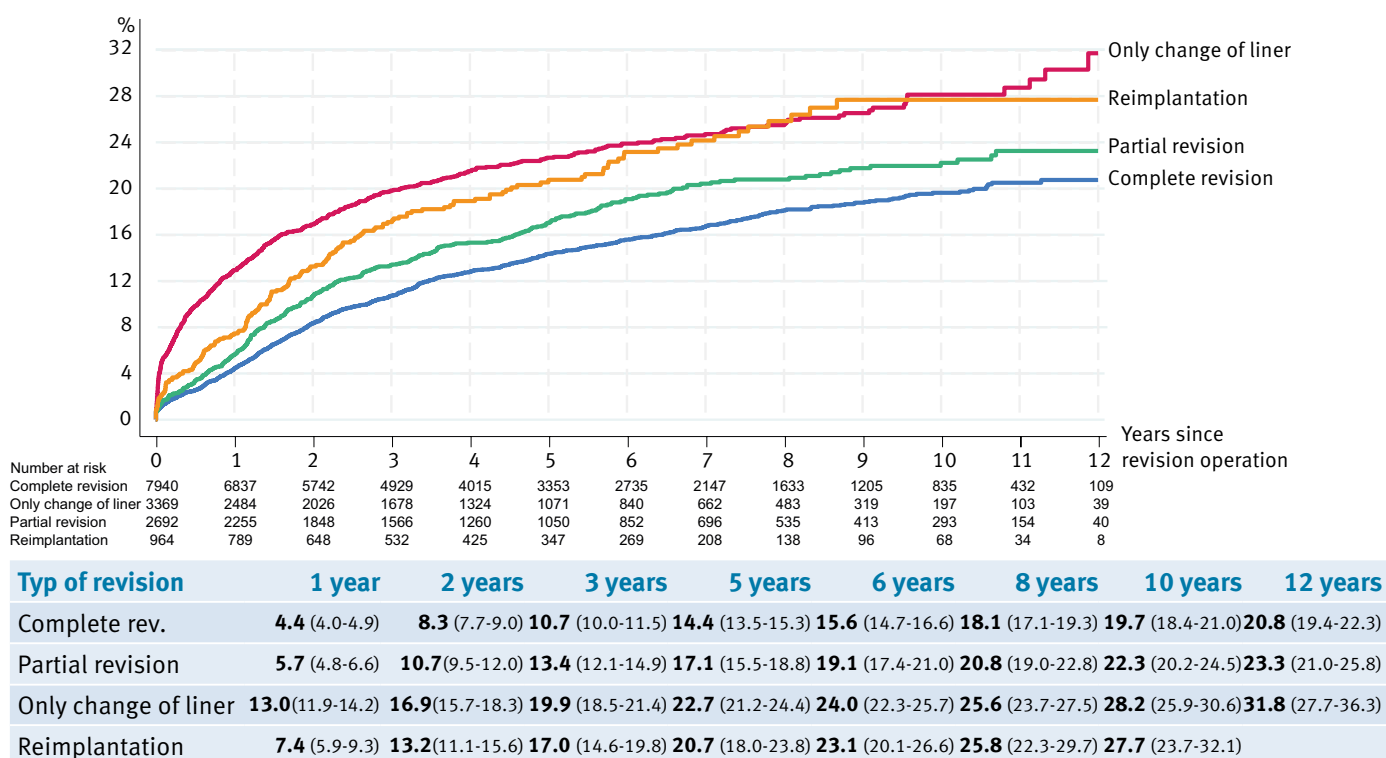


Figure 5.17

Estimated failure rates after revision of total knee arthroplasty: types of revisions

Time since revision, 2012–2024. Start point of analysis: first registered component revision in SIRIS that meets the inclusion criteria.

End point of analysis: next registered component revision. Partial revision comprises femoral revision, tibial revision and patella revision.

Reimplantation refers to implantation of total knee system after spacer (revisions due to infection).

Comprises linked and unlinked revisions. A small proportion of revisions of partial knees may be included because they cannot be reliably excluded when the revision is not linked to a primary SIRIS case.

5.17).

5.4.3 Patella and TKA re-revision

there was no statistically significant difference, as illustrated by overlapping CI. At twelve years exchange of liner alone had a re-revision rate of 31.8% (CI 27.7-36.3%) significantly worse results than partial or complete revision (**Figure 5.17**).

Component reimplantation – standing mostly for staged revision due to periprosthetic joint infection and previous temporary spacer implantation – had an early (two-year) re-revision rate of 13.2% (CI 11.1 – 15.6%). The re-revision rate increased over time up to 27.7% (CI 23.7 – 32.1%) 11 years after reimplantation. Reimplantation performed even better regarding re-revision than partial revision until seven and a half years and then picked up to the same level from the eighth year up to the last follow-up at 12 years (**Figure**

Isolated secondary patella resurfacing was associated with an early re-revision rate of 7.8% (CI 6.9 – 8.7%), which is in the range of the results after complete revision. Secondary patella resurfacing combined with liner exchange led to 9.5% (CI 7.9 – 11.4%) of re-revisions, not significantly worse than isolated secondary patella resurfacing (**Figure 5.18**). At 9 years of follow-up, secondary patella resurfacing in combination with liner exchange performed better than secondary resurfacing alone, again not statistically significantly. This was comparable to the re-revision rates after complete TKA revision (**Figure 5.17**). It is unclear if PE wear played a role in cases of isolated patella resurfacing from 7 years after revision onwards, where the curves are crossing each other, indicating advantages for combined patella resurfacing with liner exchange (**Figure 5.18**). As stated before, the type of PE (CPE vs HXLPE) was not analysed

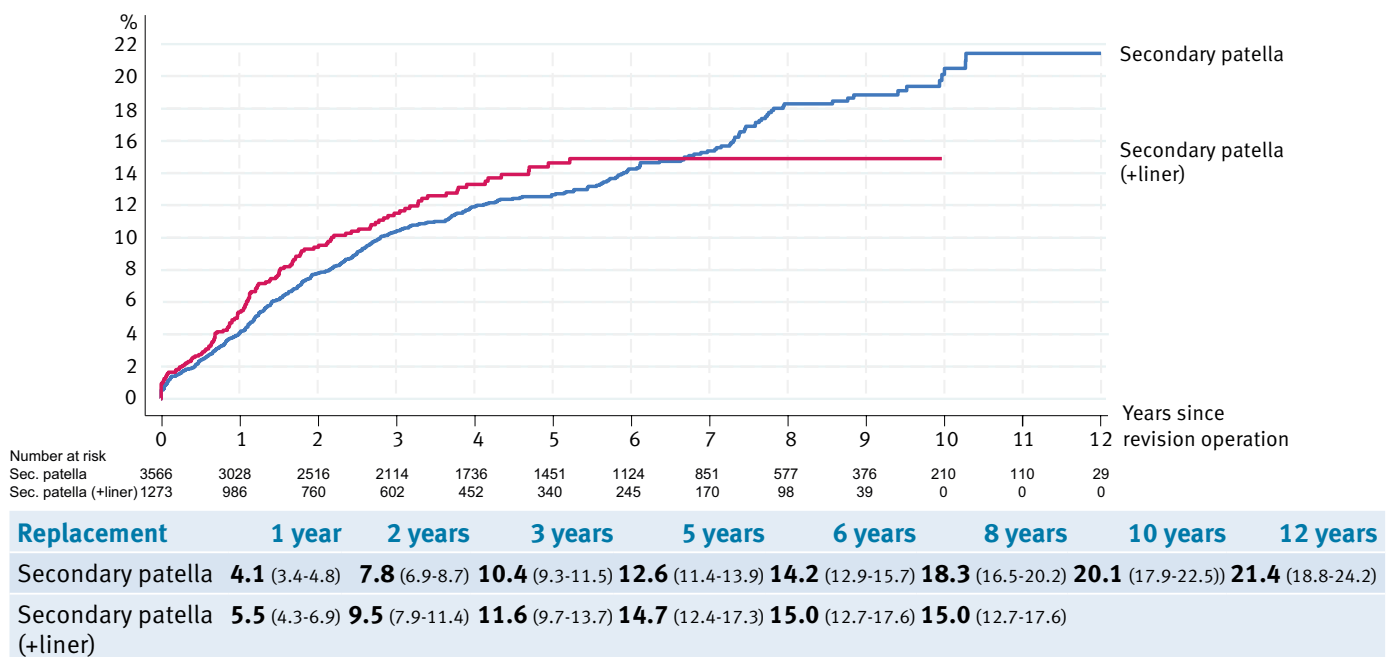


Figure 5.18

Estimated failure rates after revision of total knee arthroplasty: secondary patella replacement

Time since revision, 2012–2024. Start point of analysis: first registered component revision in SIRIS that meets the inclusion criteria.

End point of analysis: next registered component revision. Comprises of linked and unlinked revisions. A small proportion of revisions of partial knees may be included because they cannot be reliably excluded when the revision is not linked to a primary SIRIS case.

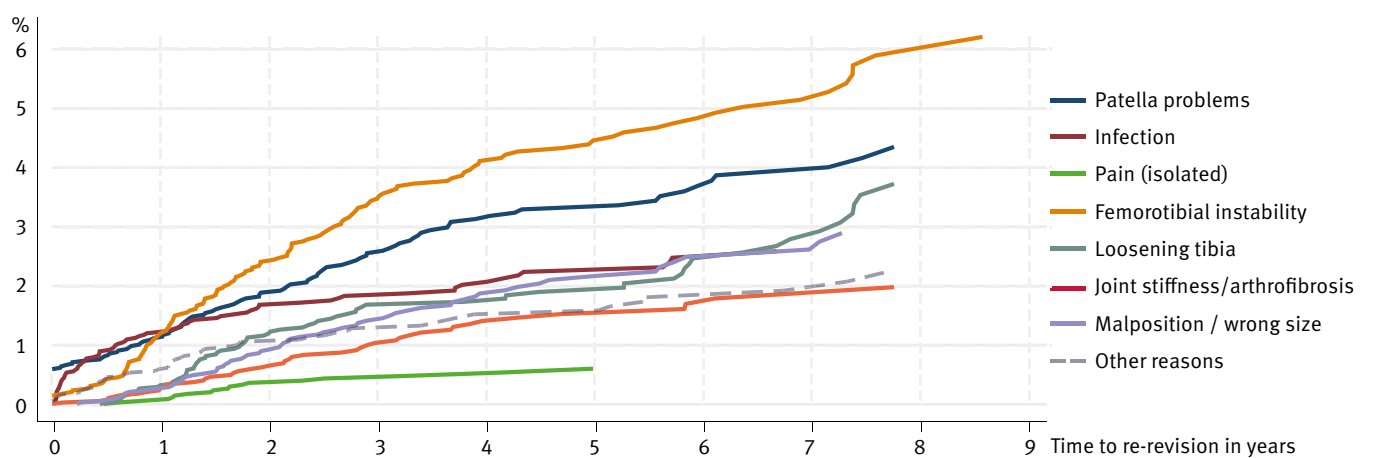
in knee arthroplasty.

The main reason for re-revision after secondary patella resurfacing, with or without liner exchange, was femerotibial instability (Figure 5.19). Astonishingly, (persistent) patella problems were the second most frequent reason, confirming that anterior knee pain after TKA often has other causes, which cannot be solved by secondary patella resurfacing alone. Loosening of the tibial component was an important cause of re-revision, whereas joint stiffness, isolated pain or “other reasons”, fortunately, did not play an important role in this context (Figure 5.19).

5.4.4 Re-revision after Conversion of PKA to TKA

Re-revision rates after conversion of a PKA to TKA reached 9.9% (CI 8.6 – 11.3%) after two years and 21.0% (CI 18.5 – 23.7%) after 11 years, respectively (Figure 5.20). This is comparable to the re-revision rates after revision TKA (Figure 5.17) but far worse than the revision rates after primary TKA (3.4% at two years (CI 3.3 – 3.6%), and 8.5% at 12 years (CI 8.2 – 8.7%) (Table 5.3, Figure 3.21).

Therefore, the widespread opinion that a PKA would be the preferred first surgery as a later conversion to TKA would lead to comparable results as after primary TKA cannot be supported by this report. Obviously, a conversion from PKA to TKA is not equivalent to a



Re-revision diagnosis	1 year	2 years	3 years	5 years	6 years	8 years	9 years
Patella problems	1.1 (0.9-1.5)	1.9 (1.5-2.4)	2.6 (2.1-3.1)	3.3 (2.7-4.0)	3.7 (3.0-4.5)	4.3 (3.5-5.4)	4.3 (3.5-5.4)
Infection	1.2 (0.9-1.6)	1.7 (1.3-2.1)	1.8 (1.5-2.3)	2.2 (1.8-2.8)	2.5 (2.0-3.1)	2.6 (2.0-3.3)	2.6 (2.0-3.3)
Pain (isolated)	0.1 (0.0-0.2)	0.4 (0.2-0.6)	0.5 (0.3-0.7)	0.6 (0.4-1.0)	0.6 (0.4-1.0)	0.6 (0.4-1.0)	0.6 (0.4-1.0)
Femerotibial instability	1.2 (0.9-1.6)	2.4 (2.0-3.0)	3.5 (2.9-4.2)	4.5 (3.8-5.3)	4.8 (4.1-5.7)	5.9 (4.9-7.1)	6.2 (5.1-7.6)
Loosening TI	0.3 (0.2-0.6)	1.2 (0.9-1.6)	1.7 (1.3-2.2)	1.9 (1.5-2.5)	2.5 (1.9-3.2)	3.7 (2.8-4.9)	3.7 (2.8-4.9)
Joint stiffness / arthrofibrosis	0.3 (0.1-0.5)	0.6 (0.4-0.9)	1.0 (0.7-1.4)	1.5 (1.1-2.1)	1.7 (1.2-2.3)	2.0 (1.4-2.8)	2.0 (1.4-2.8)
Malposition / wrong size	0.3 (0.2-0.5)	0.9 (0.6-1.3)	1.4 (1.1-1.9)	2.2 (1.7-2.8)	2.5 (1.9-3.2)	2.9 (2.2-3.8)	2.9 (2.2-3.8)
Other reasons	0.6 (0.4-0.9)	1.1 (0.8-1.5)	1.3 (1.0-1.7)	1.5 (1.2-2.0)	1.8 (1.4-2.4)	2.3 (1.6-3.1)	2.3 (1.6-3.1)

Figure 5.19
Cumulative incidence rates for different re-revision diagnosis after secondary patella replacements (TKA)

Time since revision, 2015–2024, % of implants re-revised. Detailed reasons for revisions available since 2015. Comprises of all secondary patella replacements (with or without PE replacement)

primary TKA but remains a revision with higher re-revision rates than observed after primary TKA.

The main reasons for re-revision after conversion of PKA to TKA were femerotibial instability and tibial

loosening, followed by patella problems. All other reasons were noted less frequently, isolated pain playing only a limited role this time (compare to first revision of PKA in chapter 5.7) (Figure 5.21).

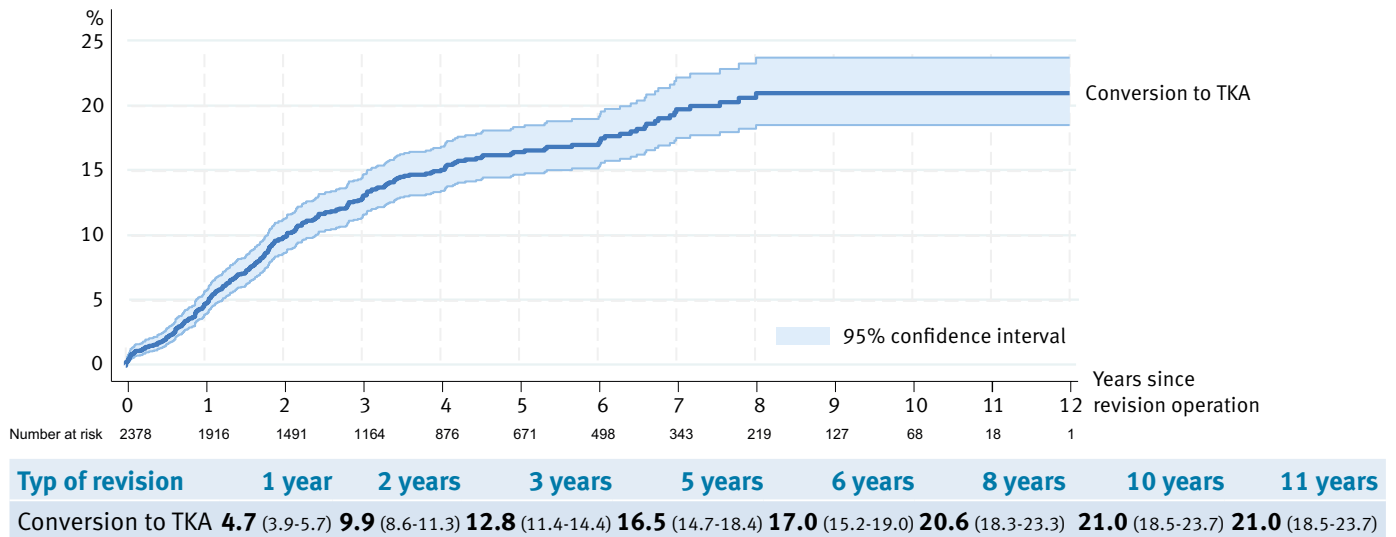


Figure 5.20
Estimated failure rates after conversion from partial knee to total knee arthroplasty

Time since revision, 2012–2024, % of implants re-revised

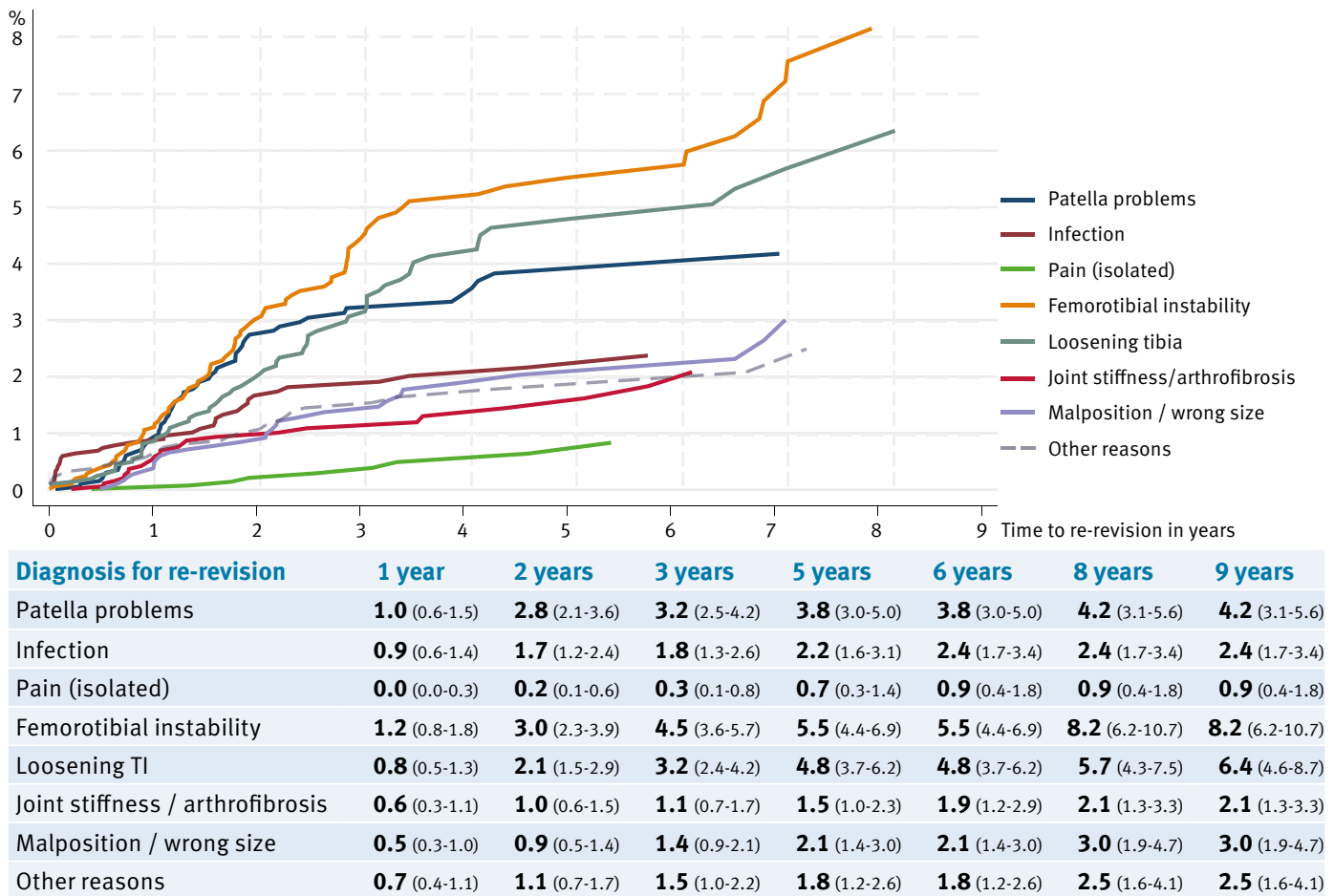


Figure 5.21
Cumulative incidence rates for different re-revision diagnoses of conversions to primary total knee arthroplasty

Time since revision, 2015–2024, % of implants re-revised. Detailed reasons for revisions available since 2015.

5.5 Results of implants in total knee arthroplasty

5.5.1 Two-year revision rates of TKA systems and brands

The two-year revision rate of the individual implants is shown in **Figure 5.27**, reflecting results from TKA performed between 01.01.2019 and 31.12.2022, providing a completed two-year follow-up by 31.12.2024. Of the 58 implant combinations used (the rest being summarised under “other systems”), one system must be considered as a potential outlier. Please refer to **Chapter 2 Methods** regarding methodological details of this definition. As usual, the potential outlier identification results in a more detailed outlier report investigating the reasons for the observed deviations from the national average. The potential outlier system at two years concerns a subtype of an older implant with otherwise solid results overall. A CCS of 52 indicates these short term problems could be related to certain hospitals or surgeons. “Other systems”, summarising TKA with insufficient numbers to

be represented individually, had an elevated 2-year revision rate but within the boundaries of Swiss average. Of interest is that the two mentioned potential long-term outliers (**compare 5.5.1**) had results with only slightly elevated revision rates at two years.

5.5.2 Long-term survival

Table 5.8 shows the TKA systems used most commonly in Switzerland, representing 74.6% (80,916) of the primary TKA from 2019 until 2024. The 25.4% less commonly used implants accounted for 27,551 cases only during this period. Only 355 implant combinations (0.3%) could not be classified, similarly to the rate of missing implants observed in the previous annual reports.

The long-term evaluation for all systems, all diagnoses, and all fixation types is depicted in **Table 5.9**, showing results up to 12 years after surgery. Wider confidence intervals reflect higher variability due to small numbers. Please take note of the case concentration score (CCS), indicating the share of the largest providing hospital, as individual providers may

System*	2019	2020	2021	2022	2023	2024	2019–2024
Attune CR-FB	677	841	1,248	1,693	2,091	2,690	9,240
Attune CR-RP	1,170	1,336	1,424	1,604	1,494	1,411	8,439
Attune PS-FB	543	463	503	675	660	596	3,440
Attune PS-RP	842	750	753	1,036	1,082	1,092	5,555
Balansys CR	294	356	526	923	752	928	3,779
Balansys PS	663	600	623	571	611	672	3,740
Balansys UC	357	387	442	598	759	655	3,198
GMK sphere	2,029	2,084	2,504	3,067	3,273	3,432	16,389
Origin PS	219	354	422	537	553	681	2,766
Persona CR-MC	716	975	1,284	1,805	2,172	2,631	9,583
Persona CR-UC	1,145	1,162	1,108	1,258	1,400	1,080	7,153
Persona PS	600	477	784	937	914	835	4,547
Triathlon PS	184	356	553	516	640	838	3,087
Other systems	5,960	5,193	4,490	3,937	4,141	3,830	27,551
Total	15,399	15,334	16,664	19,157	20,542	21,371	108,467

Table 5.8

Top 75% of primary total knee arthroplasty systems (all diagnoses, all component fixations) 2019 – 2024

*Constrained/hinged systems were included if used for cases of primary OA including OA after meniscectomy

influence results of systems not widely used. A higher share indicates an increased likelihood of bias due to local effects. A CCS of 50% and more means that the results are likely determined by one hospital/ surgeon. Primary TKA subsystems (such as PCR/ CR or PS) were analysed separately if numbers were both sufficient and showed relevantly different revision rates. The 12-year revision rate for all TKA systems was 8.4% (CI 8.2 – 8.6%). However, the various implant systems performed rather differently in the short, medium, and long term (Table 5.9).

5.5.3 Survival of different knee systems

The revision rate after 12 years varied from 4.5% for the best to 11.0% for the worst system of the implant list. “Other systems”, accounting for 25.4% of the TKA and grouped together because of small numbers, had the highest failure rate with a revision rate of 11.4% (CI 10.2–12.7%) after twelve years, significantly worse than average of 8.4% (CI 8.2–8.6%) (Table 5.9). Therefore, it should be recommended to surgeons using more exotic implants should review

System	Total N	CCS*	Mean age**	1 year (95% CI)	3 years (95% CI)	5 years (95% CI)	10 years (95% CI)	12 years (95% CI)
Advance	2,106	19	68	2.2 (1.7-2.9)	5.4 (4.5-6.5)	6.5 (5.5-7.7)	8.5 (7.1-10.0)	9.1 (7.6-11.0)
Attune CR-FB	11,803	12	69	1.3 (1.1-1.5)	3.6 (3.2-4.1)	4.7 (4.2-5.2)	7.5 (6.4-8.8)	
Attune CR-RP	13,883	11	69	1.9 (1.6-2.1)	5.2 (4.8-5.6)	6.4 (6.0-6.9)	8.9 (8.1-9.8)	
Attune PS-FB	6,084	14	70	1.4 (1.1-1.7)	4.0 (3.5-4.6)	5.1 (4.5-5.8)	8.5 (7.4-9.9)	
Attune PS-RP	8,581	15	70	1.6 (1.4-1.9)	4.8 (4.4-5.4)	6.3 (5.7-6.9)	8.9 (7.8-10.1)	
Balansys CR	4,847	15	70	1.1 (0.8-1.4)	3.3 (2.8-4.0)	4.5 (3.8-5.3)	7.3 (5.8-9.1)	7.3 (5.8-9.1)
Balansys PS	5,944	56	70	1.2 (0.9-1.5)	3.6 (3.2-4.2)	4.9 (4.3-5.6)	7.5 (6.3-8.8)	9.4 (7.5-11.8)
Balansys RP	7,171	14	70	1.4 (1.2-1.7)	4.3 (3.8-4.8)	5.5 (5.0-6.1)	8.4 (7.6-9.2)	9.4 (8.4-10.5)
Balansys UC	6,318	22	71	1.4 (1.2-1.8)	4.5 (3.9-5.1)	5.7 (5.1-6.4)	8.7 (7.7-9.8)	9.1 (8.0-10.4)
E.Motion FP/UC	2,042	84	70	1.5 (1.0-2.1)	3.5 (2.8-4.5)	4.8 (3.9-6.0)	7.7 (6.3-9.5)	8.2 (6.7-10.2)
Evolution	904	26	68	1.5 (0.8-2.8)	7.0 (4.6-10.5)	7.8 (5.1-11.9)		
First/First REV	2,812	36	70	1.6 (1.2-2.1)	4.8 (4.0-5.6)	5.8 (4.9-6.7)	7.9 (6.8-9.2)	9.1 (7.1-11.6)
GKS prime flex	516	22	69	1.3 (0.6-2.9)	7.5 (5.0-11.1)	8.0 (5.4-11.9)		
GMK primary CR/UC-RP	2,652	19	70	1.6 (1.2-2.2)	4.1 (3.4-4.9)	5.0 (4.2-5.9)	7.2 (6.1-8.4)	7.6 (6.5-8.9)
GMK primary PS	2,179	23	71	1.2 (0.8-1.8)	3.8 (3.0-4.7)	5.2 (4.3-6.3)	7.8 (6.6-9.2)	8.5 (7.1-10.3)
GMK sphere	22,080	11	69	1.6 (1.5-1.8)	4.5 (4.2-4.9)	5.9 (5.5-6.3)	7.4 (6.8-8.1)	7.8 (6.9-8.8)
ITotal	2,050	24	68	0.8 (0.5-1.3)	3.0 (2.3-3.9)	4.0 (3.2-5.1)	5.6 (4.0-7.9)	5.6 (4.0-7.9)
Innex FB	1,730	42	71	1.4 (0.9-2.1)	4.3 (3.4-5.3)	5.3 (4.3-6.5)	7.7 (6.4-9.2)	8.6 (7.1-10.4)
Innex RP	4,809	17	69	1.7 (1.4-2.1)	4.6 (4.0-5.2)	5.6 (5.0-6.3)	7.7 (6.9-8.6)	9.1 (8.1-10.3)
Journey II	2,630	27	67	3.1 (2.5-3.8)	7.5 (6.5-8.6)	9.2 (8.1-10.4)	12.6 (10.7-14.9)	
LCS compl. cem./hybrid	6,739	22	70	1.5 (1.2-1.8)	4.8 (4.3-5.3)	5.9 (5.3-6.5)	7.2 (6.6-8.0)	7.5 (6.8-8.3)
LCS compl. cementless	3,098	29	69	1.9 (1.5-2.5)	5.3 (4.6-6.2)	6.1 (5.3-7.0)	7.1 (6.2-8.2)	7.4 (6.3-8.6)
Legion	2,185	15	68	1.9 (1.4-2.6)	6.9 (5.8-8.2)	9.2 (7.9-10.8)	14.2 (11.9-16.8)	
NK flex	1,841	41	70	1.3 (0.8-1.9)	4.0 (3.2-5.0)	5.1 (4.1-6.2)	6.7 (5.5-8.0)	7.1 (5.9-8.5)

Table 5.9 Part one

Long term evaluation: Failure rates of primary total knee arthroplasty systems (all diagnoses, all component fixations)

Time since operation, 2012–2024. Please note that if reported system involves multiple sub-variants, it is possible that the long-term performance of these sub-variants may be significantly different from their combined performance.

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects.

A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Younger mean age signifies that the case mix is less “usual” and potentially biased towards higher revision risk.

their own results thoroughly and switch to a brand more frequently used in Switzerland. Mobile bearings did perform worse in all knee systems compared to other subtypes of TKA of the same brand, except for those from Medacta. Interestingly, their system with mobile bearing was associated with lower long-term revision rates compared to the other knee systems in their portfolio. Medial pivot (MP) had an elevated revision rate in an old system but within the limits of the Swiss average (rate at 12 years: 9.1%, CI 7.6-11.0%). The market leader MP system from Medacta had a better rate of 7.8% (CI 6.9-8.8%) at 12 years than the national average of 8.4% (CI 8.2-8.6%). Even more

promising was a newer system (Persona) with a significantly lower failure rate after five years of 4.5% (CI 4.0-5.1%) for the CR-MC and 3.4% (CI 3.0-3.85) for the CR-UC version compared to the average of 5.5% (CI 5.4-5.6%). Persona CPS missed significance due to wider confidence intervals (Table 5.9).

Some problematic brands lack ten-year results so far. They had an acceptable revision rate at one year, but then significantly elevated revision rates up to seven years (Table 5.9). Both problematic systems were identified as potential outliers. None of the knee systems used in Switzerland was classified as definitive outlier anymore.

System	Total N	CCS*	Mean age**	1 year (95% CI)	3 years (95% CI)	5 years (95% CI)	10 years (95% CI)	12 years (95% CI)
Nexgen CR/LPS-Flex	2,320	14	69	1.5 (1.1-2.1)	3.6 (2.8-4.4)	4.6 (3.7-5.6)	6.6 (5.5-8.0)	6.9 (5.7-8.3)
Nexgen LCCK	514	15	71	1.6 (0.8-3.2)	3.2 (2.0-5.3)	4.8 (3.1-7.5)	6.1 (4.0-9.2)	
Origin PS	2,794	23	69	1.9 (1.5-2.6)	4.4 (3.6-5.5)	6.1 (4.8-7.7)		
Persona CPS	1,188	15	72	1.2 (0.7-2.1)	2.7 (1.8-3.9)	3.7 (2.6-5.4)		
Persona CR-MC	10,556	9	69	1.3 (1.1-1.6)	3.6 (3.2-4.1)	4.5 (4.0-5.1)		
Persona CR-UC	11,195	37	70	0.9 (0.8-1.1)	2.5 (2.2-2.9)	3.4 (3.0-3.8)	4.9 (4.2-5.6)	
Persona PS	7,672	11	70	1.7 (1.5-2.1)	4.0 (3.6-4.6)	5.2 (4.7-5.8)	7.9 (7.0-9.0)	
RT-plus	1,383	14	77	3.0 (2.2-4.0)	4.5 (3.5-5.9)	5.1 (4.0-6.6)	5.6 (4.3-7.2)	5.6 (4.3-7.2)
Sigma CR-FB	4,829	28	71	0.9 (0.6-1.2)	2.4 (2.0-2.9)	3.4 (2.9-3.9)	4.3 (3.7-5.0)	4.5 (3.8-5.2)
Sigma CR-RP	2,256	40	68	2.5 (1.9-3.2)	5.8 (4.9-6.8)	6.6 (5.6-7.7)	8.3 (7.1-9.7)	9.1 (7.8-10.6)
Sigma PS-FB	1,334	59	72	1.1 (0.6-1.8)	3.3 (2.4-4.4)	4.1 (3.1-5.4)	5.6 (4.3-7.2)	6.2 (4.7-8.2)
Sigma PS-RP	1,661	11	70	1.6 (1.1-2.4)	3.9 (3.1-5.0)	4.7 (3.8-5.9)	6.2 (5.1-7.5)	6.7 (5.5-8.2)
TC-plus primary FB	2,766	32	69	1.5 (1.1-2.0)	3.8 (3.1-4.6)	5.0 (4.2-5.9)	7.4 (6.2-8.8)	8.5 (7.0-10.3)
TC-plus primary RP	2,077	31	70	1.2 (0.8-1.8)	3.7 (3.0-4.7)	5.1 (4.2-6.2)	8.8 (7.3-10.6)	10.1 (8.1-12.7)
Triathlon CR/CS	2,164	32	69	2.2 (1.6-2.9)	5.3 (4.4-6.5)	6.5 (5.4-7.8)	8.8 (7.4-10.6)	11.0 (8.6-14.1)
Triathlon PS	3,852	34	69	2.0 (1.6-2.6)	5.5 (4.7-6.4)	6.6 (5.7-7.8)	9.1 (7.4-11.1)	9.1 (7.4-11.1)
Unity	661	36	69	1.1 (0.5-2.4)	3.0 (1.8-4.9)	3.3 (2.0-5.3)		
Vanguard CR	1,240	32	67	1.5 (0.9-2.3)	4.4 (3.3-5.7)	5.7 (4.5-7.3)	7.6 (6.1-9.6)	8.1 (6.3-10.3)
Vanguard PS	1,072	57	68	2.0 (1.3-3.0)	5.1 (3.9-6.6)	6.7 (5.4-8.4)	9.1 (7.3-11.3)	9.1 (7.3-11.3)
Other systems	8,606		71	2.0 (1.7-2.4)	5.3 (4.8-5.9)	7.0 (6.4-7.7)	10.4 (9.5-11.4)	11.4 (10.2-12.7)
CH average for group				1.6 (1.5-1.6)	4.3 (4.2-4.4)	5.5 (5.4-5.6)	7.7 (7.5-7.9)	8.4 (8.2-8.6)

Table 5.9 Part two

Long term evaluation: Failure rates of primary total knee arthroplasty systems (all diagnoses, all component fixations)

Time since operation, 2012–2024. Please note that if reported system involves multiple sub-variants, it is possible that the long-term performance of these sub-variants may be significantly different from their combined performance.

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects.

A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Younger mean age signifies that the case mix is less “usual” and potentially biased towards higher revision risk.

One older (Sigma CR-FB) and one of the newer TKA systems (Persona CR-UC) performed significantly better at long term than the average (Figure 5.22). For Sigma CR-FB, this does not speak by itself for a better performance, as older systems were often used in older patients, inherently more reluctant to revision than younger and more active patients.

Two newer systems from one manufacturer (Journey II and Legion, from Smith&Nephew) performed significantly worse than the Swiss average, but still below

long-term outlier limits. The effect started early after primary TKA, and the revision rates remained elevated up to 12 years of follow-up (Figure 5.23).

With one exception, the newer systems did not lead to improved revision rates at medium and long term. The remaining brands of TKA had revision risks within the margins of the lower and upper limits at 66% and 150% of the group average, respectively (Figures 5.22 and 5.23).

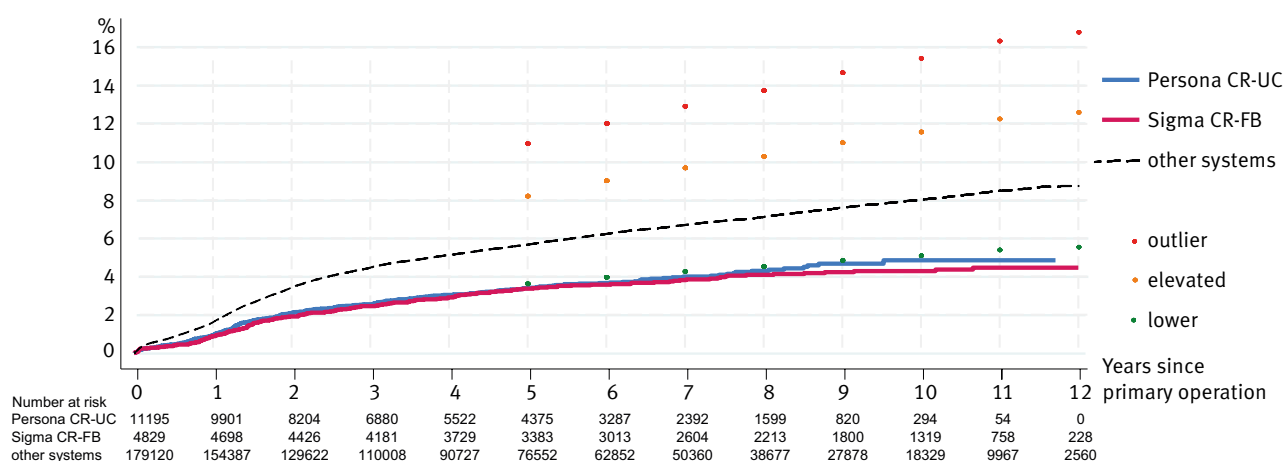


Figure 5.22

Implant combinations with below-average long-term revision rates (all TKA)

Below-average was defined as an 9-year/10-year revision rate of up to 66% of the group average (and upper bounds of the 95% confidence interval staying below the lower bound of the group average; and at least 25 cases at risk at 9 years/10-years). The dots indicate upper and lower limits.

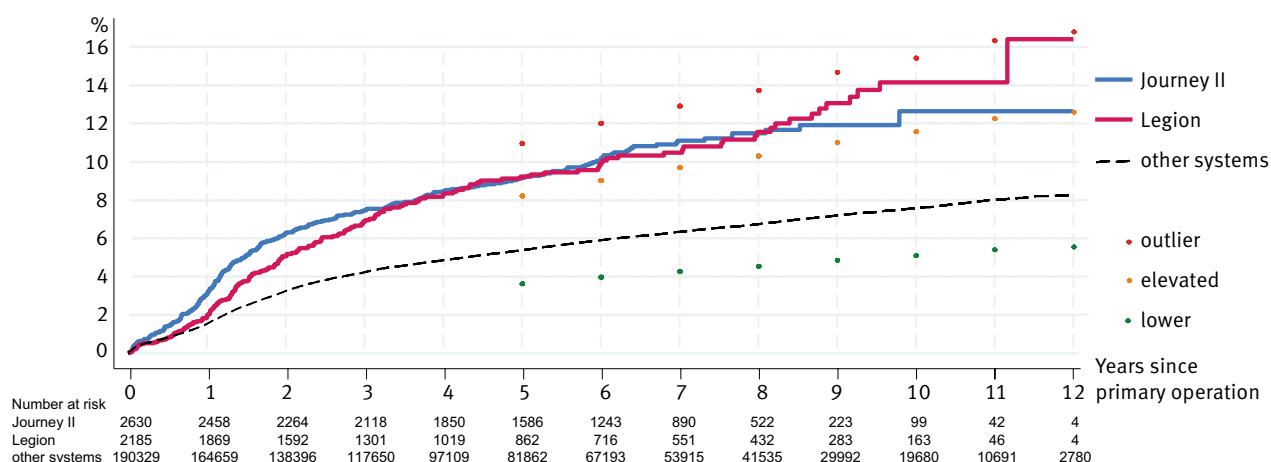


Figure 5.23

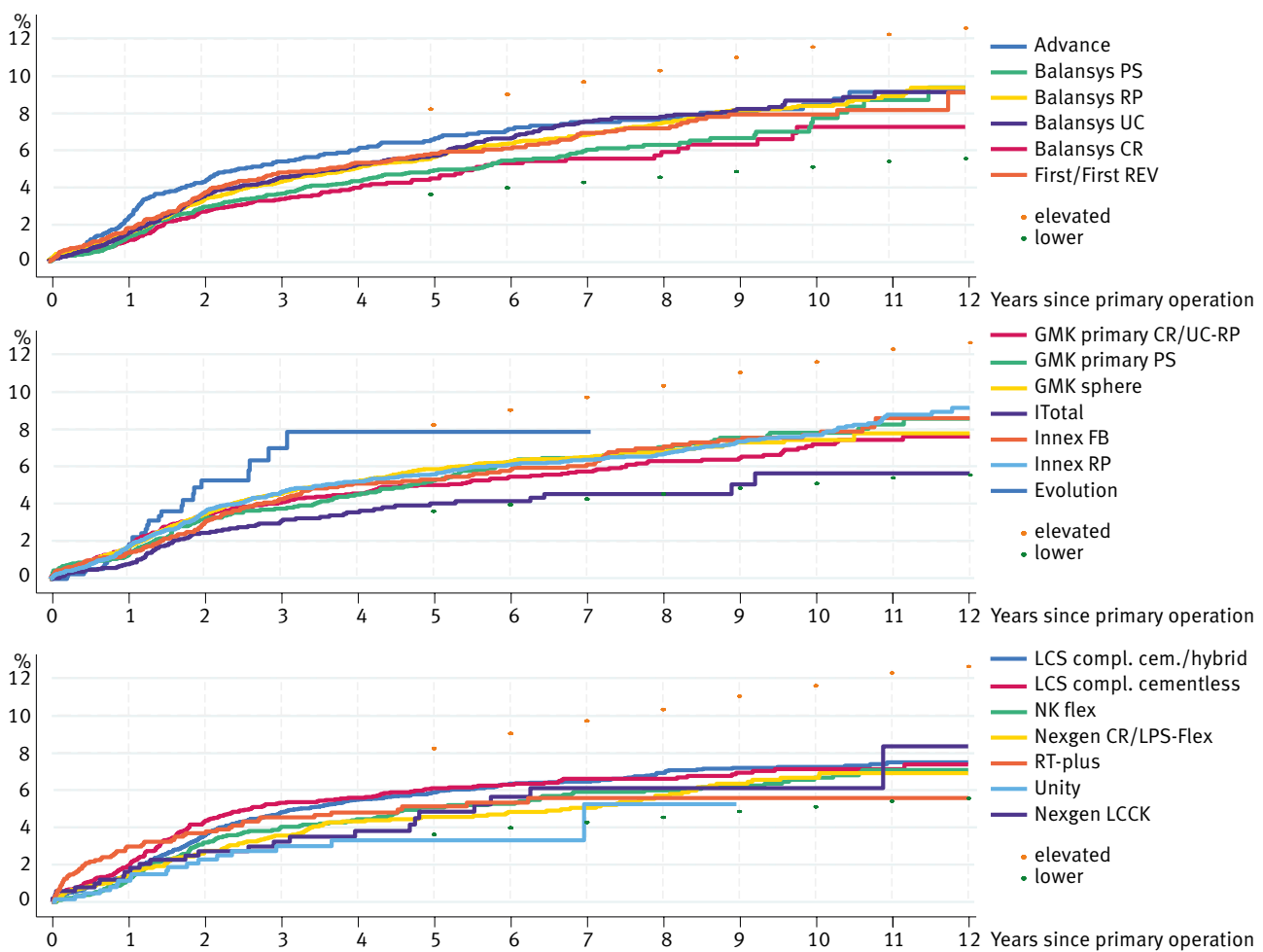
Implant combinations with elevated long-term revision rates (all TKA)

An elevated revision rate was defined as a deviation of at least 50% above the group average at any time between year 5 and year 12 (and lower bounds of the 95% confidence interval exceeding the upper bound of the group average; and at least 50 cases at risk at 5 years). The dots indicate upper and lower limits.

5.5.4 Brands and subsystems

Figures 5.24 show performance of the remaining TKA brands, including subsystems, up to twelve years after primary arthroplasty. The upper and lower limits of the Swiss average are illustrated by green (lower) and orange (upper margin) dots. The goal of this differentiation is to reduce camouflage effects. It could be demonstrated some years ago by other registers, such as the British NJR, that subdividing subsystems – for which the average performance of the whole brand was within the countywide range or even better – some subtypes performed significantly worse or

were even outliers, the effect though hidden by the rest of the well performing variants. In Switzerland, such an example is the well-known Sigma TKA (Figures 5.24), of which the subsystems show differing results. While Sigma CR-FB had excellent results (Figure 5.22), the other variants (CR-RP, PS-FB, PS-RP) showed a spread of results in the long-term evaluation. All subsystems grouped together had revision rates within the expected range, even with a tendency toward decent 12-year results. However, PS-RP cases from 2019 to 2022 had outlier status already in the 2-year evaluation (Figures 5.25). It could be assumed in this case that not only the knee system,



Figures 5.24 Part one

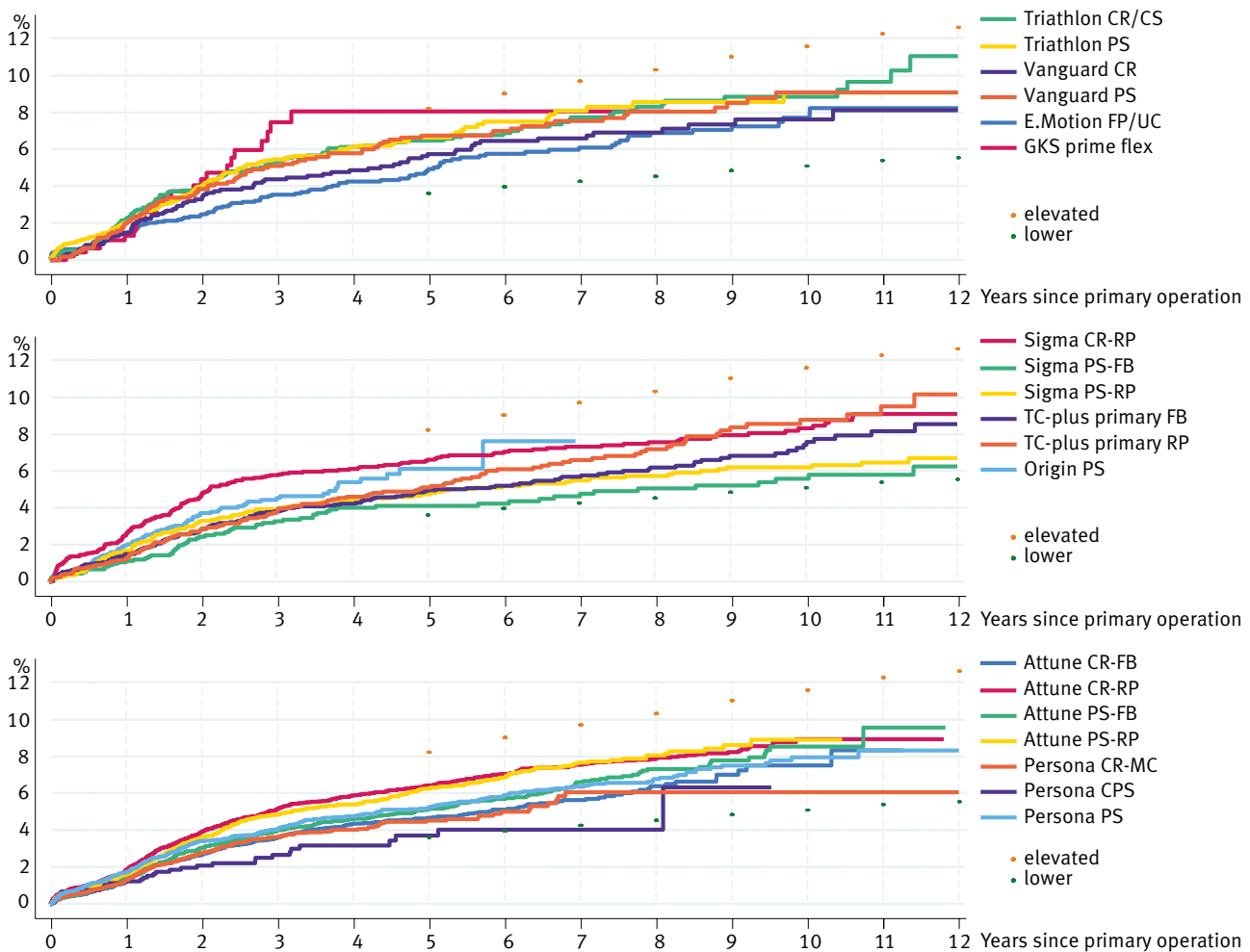
All remaining implant combinations with average revision risks (all TKA)

Also showing upper and lower limits (corresponding to elevated and below-average version risk at 150% and 66% of the group average respectively). upper and lower limits.

but the surgeons influenced the results significantly, particularly due to this temporal specificity and as the 12-year revision rates of this system are excellent. The PS-RP variation was rarely used from 2021 onwards, the brand Sigma completely disappeared from the Swiss marked in 2025.

All remaining knee systems showed long-term results close to the Swiss average (Figures 5.24). Total and Unity tended to perform better than Swiss average, the latter with a follow up to 9 years only. Interest-

ingly, a rotating hinge implant (RT-plus) had elevated revision rates up to three years after surgery but was performing at the lower margin of Swiss average in long-term. Two older subsystems of a brand (Sigma PS-FB and PS-RP) tended to perform better than average as well as two newer subsystems (Persona CR-MC and CPS). Persona CPS however has only 9 years of follow-up available and had a remarkable step to worse at 8 years. This may however best be explained by small numbers remaining at risk at this time point.



Figures 5.24 Part two

All remaining implant combinations with average revision risks (all TKA)

Also showing upper and lower limits (corresponding to elevated and below-average version risk at 150% and 66% of the group average respectively). upper and lower limits.

System	CCS*	Mean age	at risk N**	Revised N	% (95% CI)***
Advance	24	68	708	31	4.4 (3.1-6.3)
Anatomic	28	69	128	4	3.2 (1.2-8.3)
Attune CR-FB	14	69	4,459	111	2.5 (2.1-3.0)
Attune CR-RP	9	69	5,534	212	3.9 (3.4-4.4)
Attune CRS	11	69	127	6	4.8 (2.2-10.4)
Attune PS-FB	15	70	2,184	54	2.5 (1.9-3.3)
Attune PS-RP	15	70	3,381	125	3.7 (3.2-4.4)
Balansys CR	19	70	2,101	72	3.5 (2.8-4.4)
Balansys PS	50	70	2,457	84	3.5 (2.8-4.3)
Balansys RP	18	71	1,597	60	3.8 (3.0-4.9)
Balansys UC	22	71	1,784	74	4.2 (3.4-5.2)
E.Motion FP/UC	100	70	602	15	2.5 (1.5-4.2)
E.Motion PS	100	73	56	1	1.8 (0.3-12.0)
Endo-Model SL	20	77	137	5	3.8 (1.6-8.8)
Endo-Model rotation	50	80	102	2	2.0 (0.5-7.9)
Enduro	40	77	91	3	3.4 (1.1-10.1)
Evolution	46	68	265	15	5.7 (3.5-9.3)
First/First REV	30	71	717	34	4.8 (3.5-6.7)
GKS prime flex	26	69	296	12	4.1 (2.3-7.1)
GMK hinge	14	76	156	4	2.6 (1.0-6.8)
GMK primary CR/UC-FB	48	71	80	3	3.8 (1.2-11.4)
GMK primary CR/UC-RP	27	70	347	11	3.2 (1.8-5.7)
GMK primary PS	26	71	303	19	6.5 (4.2-9.9)
GMK sphere	12	69	9,684	345	3.6 (3.2-4.0)
Gemini SL	64	67	109	2	1.9 (0.5-7.2)
Genus	99	72	123	8	6.5 (3.3-12.6)
ITotal	29	69	1,082	31	2.9 (2.0-4.1)
Innex FB	90	72	246	4	1.6 (0.6-4.3)
Innex RP	32	71	633	21	3.4 (2.2-5.1)
Journey II	28	68	909	36	4.0 (2.9-5.5)
LCS complete cemented/hybrid	34	70	1,935	78	4.1 (3.3-5.1)
LCS complete cementless	40	68	577	31	5.4 (3.8-7.6)
Legion	24	68	814	38	4.7 (3.5-6.4)
NK flex	45	70	141	5	3.6 (1.5-8.4)
Nexgen CR/LPS-Flex	23	68	397	14	3.6 (2.1-6.0)

Figures 5.25 Part one

2-year evaluation: Revision rates of primary total knee arthroplasty systems within 24 months

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

*** Rates adjusted for effects of mortality and emigration.

System	CSS*	Mean age	at risk N**	Revised	
				N	% (95% CI)***
Nexgen LCCK	22	72	156	3	2.0 (0.6-6.0)
Nexgen RHK	28	77	128	3	2.4 (0.8-7.1)
Optetrak logic	86	67	84	1	1.2 (0.2-8.2)
Origin CR	56	68	52	0	0.0 (-.)
Origin PS	21	69	1,532	57	3.7 (2.9-4.8)
Persona CPS	16	72	568	9	1.6 (0.8-3.0)
Persona CR-MC	8	69	4,780	131	2.8 (2.3-3.3)
Persona CR-UC	39	70	4,673	91	2.0 (1.6-2.4)
Persona PS	13	70	2,798	98	3.5 (2.9-4.3)
RT-plus	22	78	511	19	3.8 (2.5-5.9)
Score / Score II	84	69	166	5	3.0 (1.3-7.2)
Sigma CR-FB	31	72	953	24	2.5 (1.7-3.8)
Sigma CR-RP	52	68	461	27	5.9 (4.1-8.5)
Sigma PS-FB	90	72	337	11	3.3 (1.8-5.9)
Sigma PS-RP	42	71	91	8	8.9 (4.5-16.9)
TC-plus primary FB	42	70	754	28	3.8 (2.6-5.4)
TC-plus primary RP	32	71	595	16	2.7 (1.7-4.4)
Triathlon CR/CS	39	69	583	19	3.3 (2.1-5.1)
Triathlon PS	38	68	1,609	70	4.4 (3.5-5.5)
U2	93	70	162	8	4.9 (2.5-9.6)
Unity	33	69	339	10	3.0 (1.6-5.5)
Vanguard CR	51	67	263	7	2.7 (1.3-5.5)
Vanguard PS	42	68	198	6	3.0 (1.4-6.7)
Other systems		70	501	23	4.7 (3.1-7.0)
CH average for group					3.4 (3.3-3.6)

Figures 5.25 Part two

2-year evaluation: Revision rates of primary total knee arthroplasty systems within 24 months

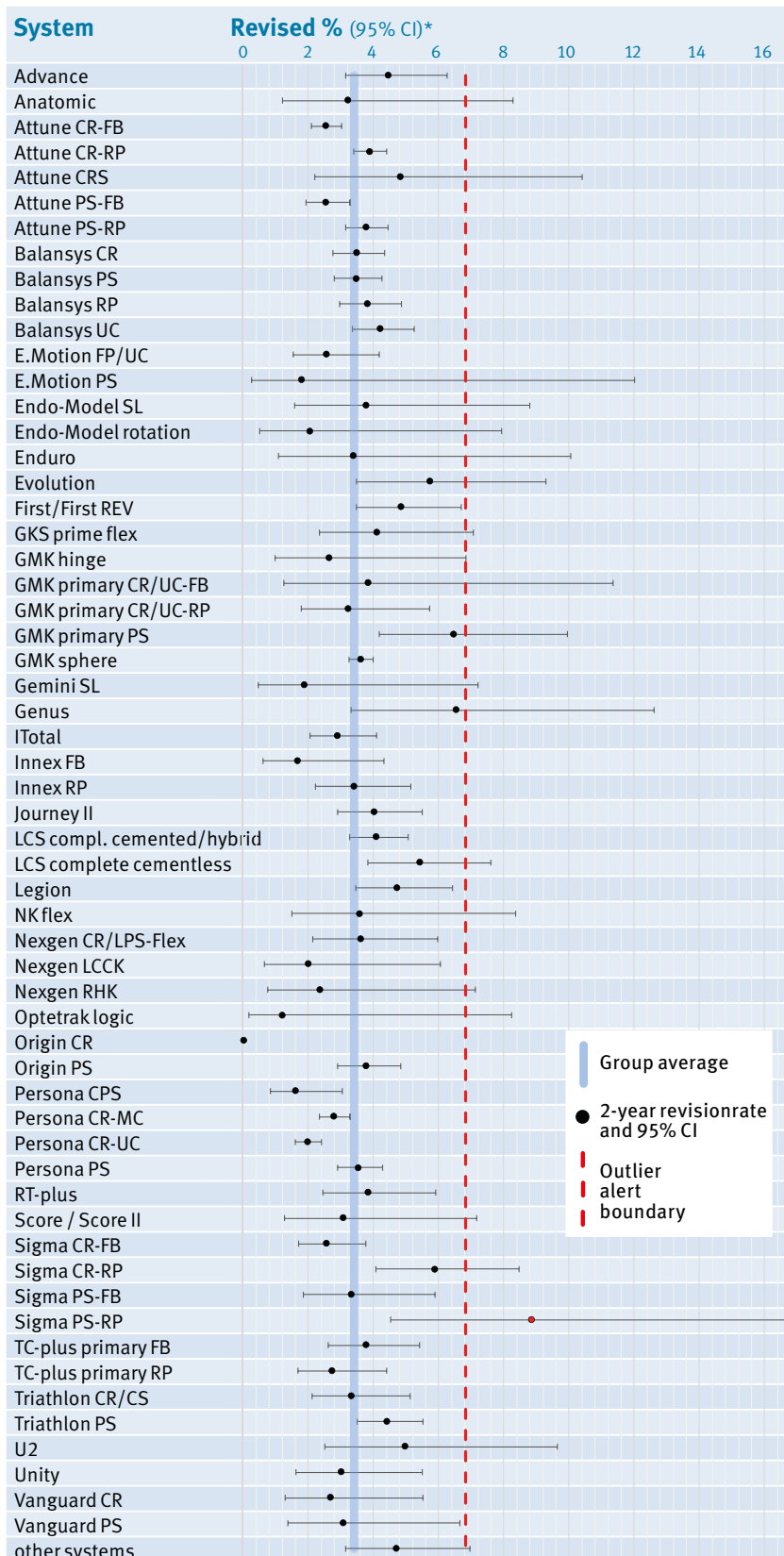
4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

- * Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.
- ** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).
- *** Rates adjusted for effects of mortality and emigration.

Important information on the use of the implant performance tables below

- Estimated revision rate exceeds the alert boundary, but we do not identify this implant combination as an outlier because the 95% confidence interval overlaps the confidence zone of the reference group.
- Identified as potential outliers. Please note the statistical confidence intervals. The outlier status comes with varying degrees of statistical probability. We consider the potential outlier status „highly likely“ when both the estimated revision rate and the complete confidence interval exceed the outlier alert boundary.

Please be aware that relatively rare implant combinations are frequently used in only a small number or indeed only in one hospital in Switzerland. Observed revision rates may be determined by local factors and performance may differ significantly between locations. Manufacturers of detected outlier implants and the hospitals where they were used (and revisions occurred) have been informed by SIRIS.



Figures 5.25 Part three

2-year evaluation: Revision rates of primary total knee arthroplasty systems within 24 months

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

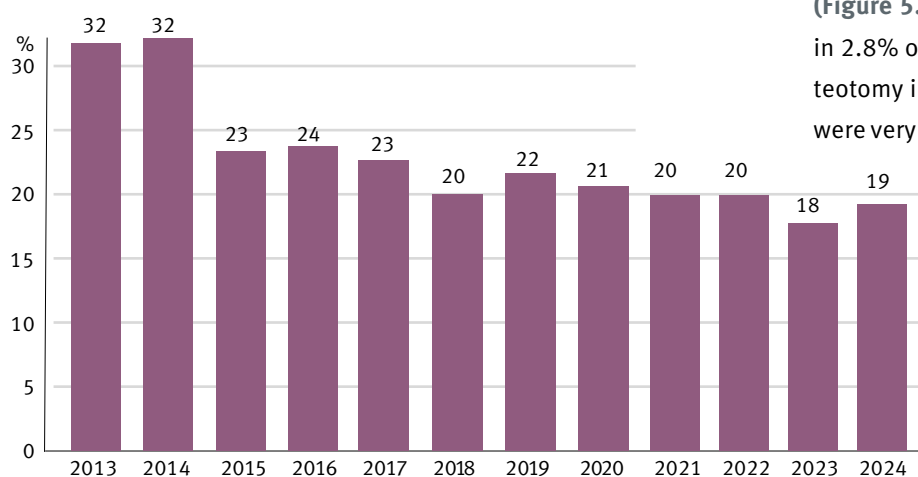
* Rates adjusted for effects of mortality and emigration.



Partial knee arthroplasty

5.6 Primary partial knee arthroplasty

Since 2012, a total of 35,908 primary PKA were registered (Table 3.10). The proportion of PKA was 15.4% over the past 12 years. In 2024, 21,568 TKA and 3,725 PKA had been performed, resulting in a share of PKA of 14.7%. For further details regarding incidence and demography please consult also Chapter 3.3.3.



Figures 5.26

Share of partial knee patients who had knee arthroscopy prior to arthroplasty (%)

Previous surgery	N	%	Intervention	N	%
None	12,980	64.8	Unicompartment medial	16,769	83.7
Knee arthroscopy	3,951	19.7	Unicompartment lateral	1,146	5.7
Meniscectomy	4,771	23.8	Femoropatellar	1,336	6.7
ACL reconstruction	401	2.0	Other (including combinations)	251	1.3
Osteotomy tibia close to knee	212	1.1	Other (type unknown)*	535	2.7
Osteosynthesis tibia close to knee	100	0.5	Conventional (including minimally invasive)	17,346	86.6
Surgery for patella stabilization	215	1.1			
Synovectomy	122	0.6	Technology	N	%
Osteotomy femur close to knee	33	0.2	Computer assisted / navigation	340	1.7
Osteosynthesis femur close to knee	35	0.2	Patient specific instrumentation	1,398	7.0
Surgery for treating infection	9	0.0	Robotic-assisted (v2021)	890	4.4
Surgery for tumor	6	0.0	Other	371	1.9
Other	339	1.7			

Table 5.10

Primary partial knee arthroplasty: Surgery characteristics

All diagnoses, all component fixations (2019–2023).

* In those cases TKA categories were chosen on the data entry form but partial knee systems registered. We consider implant registration more reliable than form entry and therefore recognise them as partial knee procedures.

5.6.1 Previous surgery

Over the past six years, no previous operations were registered in 64.8% of the PKA, whereas 23.8% had a meniscectomy, and 19.7% an arthroscopy. As both these options could be selected simultaneously, one may expect that an earlier open meniscectomy was performed in 4.1%. Prior meniscal surgery declined steadily since 2012 (32%), plateauing around 19% (Figure 5.26). Former ACL reconstruction was noted in 2.8% of the cases, followed next by high tibial osteotomy in 1.1%. All other previous types of surgery were very rare (Table 5.10).

5.6.2 Type of PKA and fixation

Medial PKA represented 83.7% of all the PKA cases, lateral PKA 5.7%, and PFJ replacement 6.7%. Others, including combinations, were rare (1.3%). In 2.7%, the type was incorrectly classified as a TKA (mentioned as “other, type unknown”), but the implant data identified them as PKA (Table 5.10). This underscores the importance of checking properly the options available at case registration.

Most of the PKA were fully cemented during the period from 2019 to 2024. The share of cemented components continuously increased and reached 87.2% in 2024. Correspondingly, the share of uncemented PKA continuously decreased over the same period, hybrid fixation playing no relevant role (Figure 5.27).

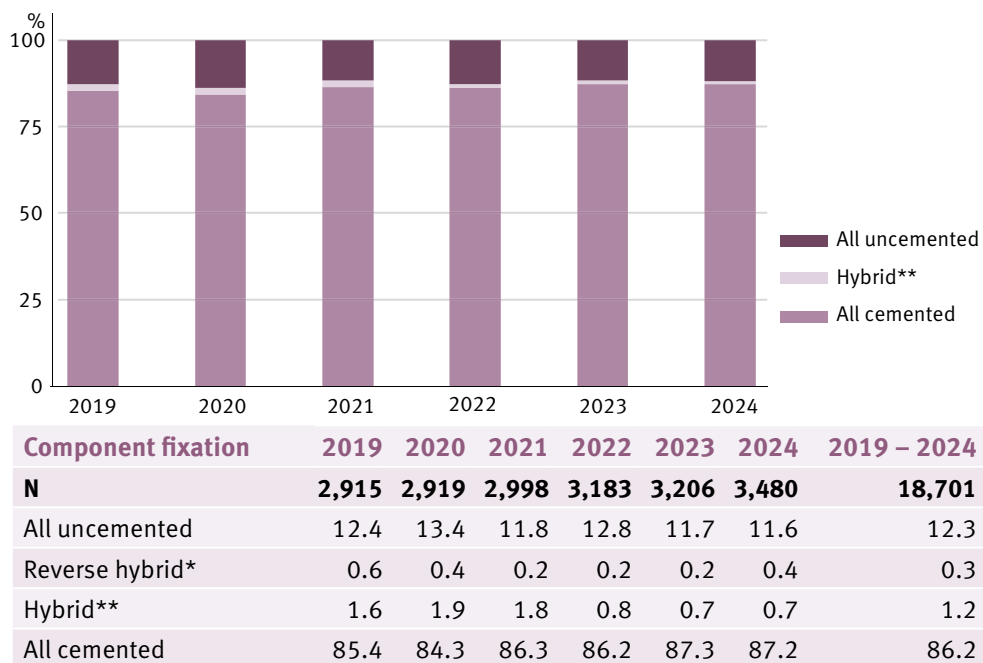


Figure 5.27

Primary partial knee arthroplasty: Component fixation (%)

All diagnoses

* femur cemented, tibia uncemented

** femur uncemented, tibia cemented

5.6.3 Technology

Between 2019 and 2024, PKA were implanted conventionally in 86.6% of the cases. Technical support was still rarely used during PKA in Switzerland, accounting for 13.1% of all cases during the same period (Table 5.10), although a slight increase could be observed since 2018 mainly for robotic assistance (Figure 5.28). Conventional computer navigation

was used in 1.7%, PSI in 7.0% and robotics in 4.4% (Table 5.10). It would have been expected that technical support would be more frequently used in PKA than in TKA to reduce the risk of surgical errors, mainly on the tibial side, mostly responsible for elevated early revisions after PKA. Obviously, Swiss surgeons rely far less on technological support than surgeons in other countries like Australia.

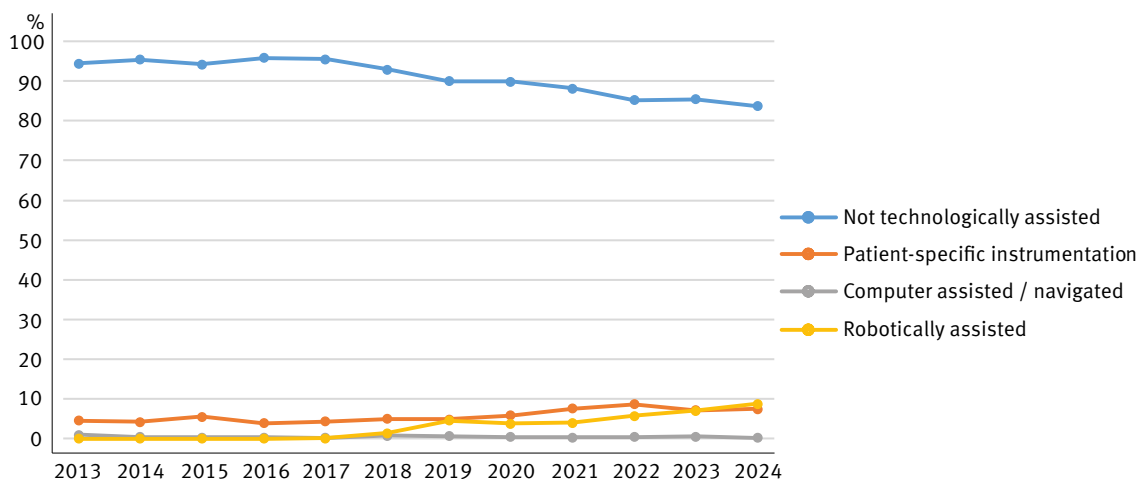


Figure 5.28

Partial knee arthroplasty: technological assistance over time (%)

NB: robotically assisted cases before v2021 were derived from free text entries. “Other” responses were coded as “Not tech. assisted” unless they specifically mentioned robotic, PSI oder navigation”.

There was a recognisable overlap of computer navigated and other responses, indicating that those cases were actually robotic-assisted.

5.7 First revision within two years after primary partial knee arthroplasty

First revisions are revisions linked to primary PKA registered in SIRIS and occurring for the first time. Of the 35,908 PKA documented since 2012 (Table 3.10), 12,874 had been performed between 01.01.2019 and 31.12.2022 and were at risk for revision in the 4-year moving window used to calculate the current 2-year revision rate. Of the implants in this cohort, 633 had been revised, accounting for a 2-year revision rate of 5.0% (CI 4.6 – 5.3%) (Table 5.11). Please refer also to Chapter 3.3.5 for incidence and demography.

	N at risk*	Revised		95% CI	
		N	%**	lower	upper
Overall	12,874	633	5.0	4.6	5.3
Gender					
Women	5,341	276	5.2	4.6	5.8
Men	6,159	266	4.4	3.9	4.9
Age group					
<55	1,709	119	7.0	5.9	8.3
55-64	3,989	207	5.2	4.6	6.0
65-74	3,635	146	4.1	3.5	4.7
75-84	1,915	58	3.1	2.4	3.9
85+	252	12	4.8	2.8	8.4

Table 5.11
First revision of primary partial knee arthroplasty within 24 months overall and according to baseline characteristics

All diagnoses, all component fixations.

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

** Rates adjusted for effects of mortality and emigration.

5.7.1 Reasons for revision

The main reason for revision of PKA was loosening of the tibial component (30.0%), followed by progression of OA (17.2%), loosening of the femoral component (13.3%), femorotibial instability (9.8%), pain of unclear origin (7.1%) and patella problems (7.1%). Infections were registered in 9.6% and periprosthetic tibial fractures in (7.9%). Malposition of the tibial component was registered in 5.2% and in 3.0% of the femoral component (Table 5.12).

Excluding infections, most of the other reasons for revision speak for technical errors or mistakes in indication when the revision had to be performed within less than two years after primary PKA.

Reason for early first revision	N revised	%
Loosening tibia	190	30.0
Progression of unicomp. OA	109	17.2
Loosening femur	84	13.3
Femorotibial instability	62	9.8
Infection	61	9.6
Periprosthetic fracture tibia	50	7.9
Patella problems	45	7.1
Pain (of unclear origin)*	45	7.1
Component malposition tibia	33	5.2
Component malposition femur	19	3.0
Joint stiffness/arthrofibrosis	16	2.5
Wear of inlay	12	1.9
Loosening patella	8	1.3
Sizing femoral component	7	1.1
Sizing tibial component	5	0.8
Patellar instability	3	0.5
Periprosthetic fracture patella	2	0.3
Periprosthetic fracture femur	0	0.0
Other	67	10.6
Total Revisions	633	

Table 5.12
Reason for early first revision of primary partial knee arthroplasty

All diagnoses, all component fixations. 4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

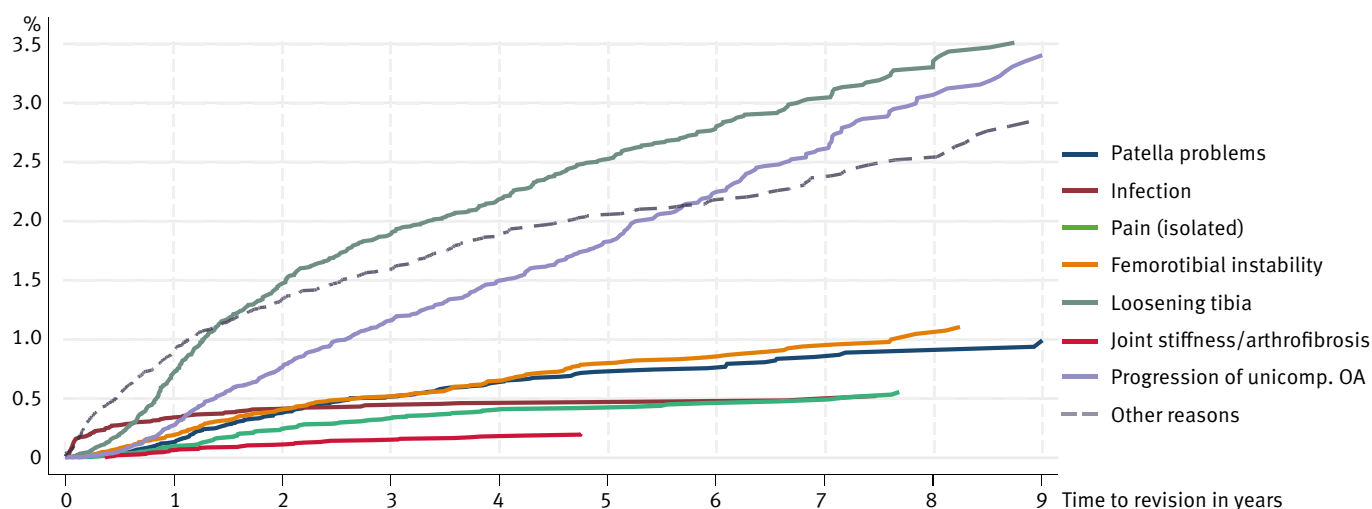
Early first revisions are those occurring within 2 years of the primary arthroplasty. Multiple responses possible (percentages do not sum to 100)

* Pain was frequently reported alongside other reasons. The proportion of “isolated pain” was 7%.

Taking in account cumulative incidence rates for revision between 2015 and 2024, loosening of the tibial component was most prominent, followed by patella problems and “other reasons”. The curve for all the other revision reasons had a flatter pattern (Figure 5.29). Similarly to TKA, only periprosthetic joint infections (PJI) were revised early after the index surgery, most of the other reasons leading to a revision peak 11.8 to 14.6 months after the primary PKA. Revision for early tibial loosening had its peak 11.8 months after PKA. Early revision because of isolated pain of unknown origin was performed later at median 16.0 months (Figure 5.30).

Fixation and revision

Uncemented PKA had a higher revision rate than cemented ones until 12 years of follow-up (Figure 5.31). A higher revision rate in uncemented PKA can be expected especially early after surgery, as uncemented implants require sufficient primary stability for osteointegration, a critical issue in some cases, particularly regarding the tibial component. After the initial failures had manifested, the revision curve of the uncemented implants remained parallel to the one observed for cemented implants. Astonishingly, the difference was getting even more pronounced from the 10th year onward (Figure 5.31). Of note, patients with uncemented PKA were younger than those with cemented implants. This selection bias influences the revision rate unfavourably.

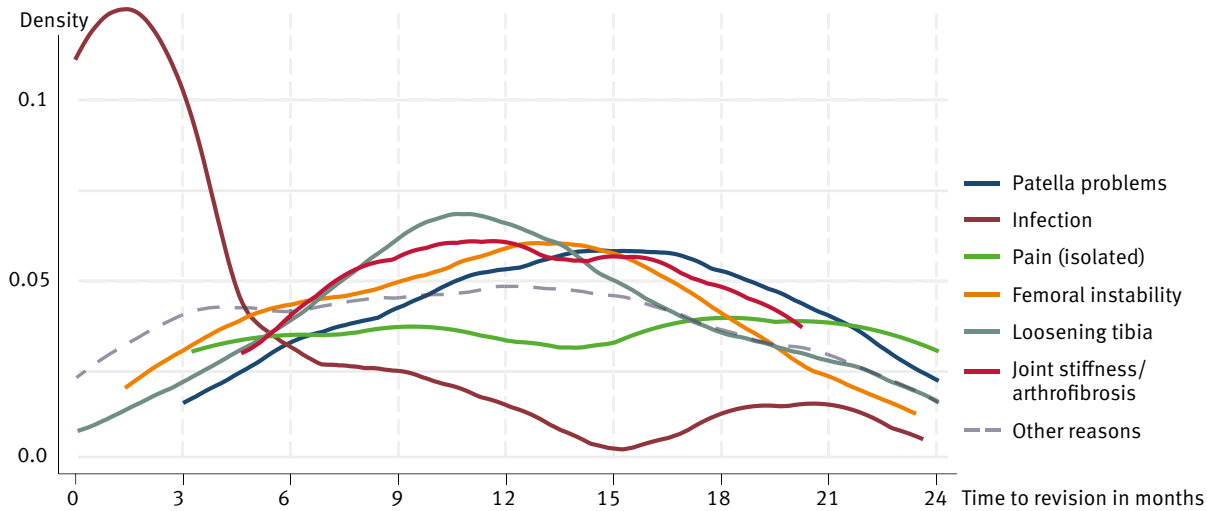


Diagnosis for revision	1 year	2 years	3 years	5 years	6 years	8 years	9 years
Patella problems	0.1 (0.1-0.2)	0.4 (0.3-0.5)	0.5 (0.4-0.6)	0.7 (0.6-0.9)	0.8 (0.6-0.9)	0.9 (0.8-1.1)	1.0 (0.8-1.2)
Infection	0.3 (0.3-0.4)	0.4 (0.3-0.5)	0.4 (0.4-0.5)	0.5 (0.4-0.6)	0.5 (0.4-0.6)	0.5 (0.4-0.6)	0.5 (0.4-0.6)
Pain (isolated)	0.1 (0.1-0.1)	0.2 (0.2-0.3)	0.3 (0.3-0.4)	0.4 (0.3-0.5)	0.5 (0.4-0.6)	0.6 (0.4-0.7)	0.6 (0.4-0.7)
Femorotibial instability	0.2 (0.1-0.2)	0.4 (0.3-0.5)	0.5 (0.4-0.6)	0.8 (0.7-0.9)	0.9 (0.7-1.0)	1.0 (0.9-1.2)	1.1 (0.9-1.3)
Loosening TI	0.7 (0.6-0.8)	1.5 (1.3-1.6)	1.9 (1.7-2.1)	2.5 (2.3-2.7)	2.8 (2.6-3.1)	3.4 (3.1-3.7)	3.5 (3.2-3.9)
Joint stiffness / arthrofibrosis	0.1 (0.0-0.1)	0.1 (0.1-0.2)	0.1 (0.1-0.2)	0.2 (0.2-0.3)	0.2 (0.2-0.3)	0.2 (0.2-0.3)	0.2 (0.2-0.3)
Progression of unicomp. OA	0.3 (0.2-0.3)	0.8 (0.7-0.9)	1.2 (1.0-1.3)	1.8 (1.6-2.0)	2.3 (2.0-2.5)	3.1 (2.8-3.4)	3.4 (3.0-3.8)
Other reasons	0.9 (0.8-1.0)	1.3 (1.2-1.5)	1.6 (1.4-1.8)	2.1 (1.9-2.3)	2.2 (2.0-2.4)	2.5 (2.3-2.8)	2.9 (2.5-3.2)

Figure 5.29

Cumulative incidence rates for different revision diagnosis of partial knee arthroplasty

Time since operation, 2015–2024, % of implants revised. Detailed reasons for revisions available since 2015.

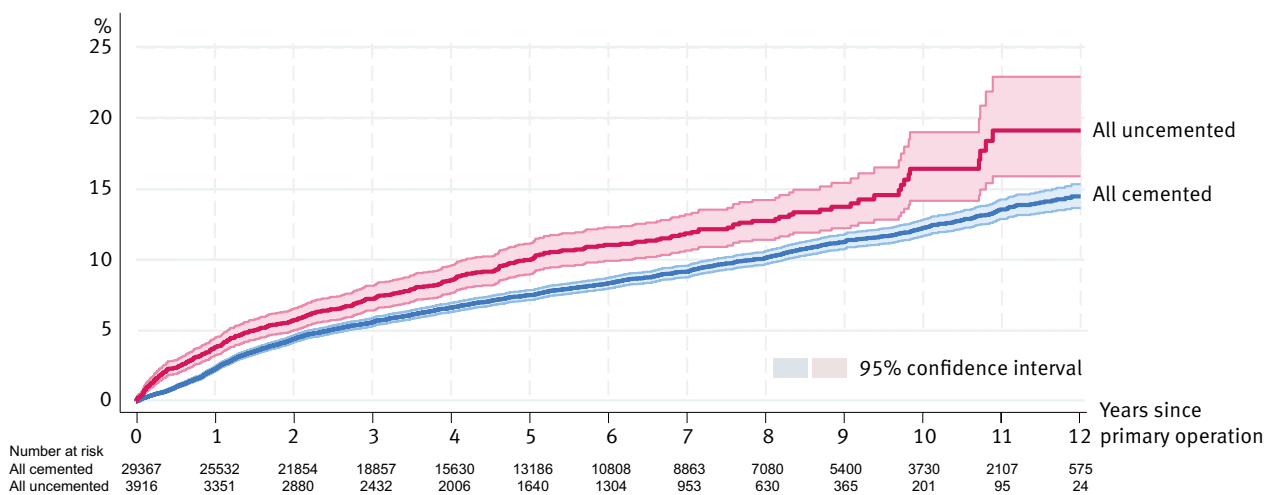


Reason for revision	N	Median	IQR 25%	IQR 75%
Patella problems	45	14.6	9.7	18.1
Infection	61	1.4	0.8	6.5
Pain (isolated)	15	16.0	5.1	21.1
Femoral instability	62	12.3	6.9	15.5
Loosening tibia	190	11.8	8.3	16.6
Joint stiffness/arthrofibrosis	16	12.2	8.8	17.5
Other	316	11.1	6.0	16.5

Figure 5.30

Time interval between primary partial knee arthroplasty and first revision by reason

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024). Early first revisions are those occurring within 2 years of the primary arthroplasty. All diagnoses.



Component fixation	1 year	2 years	3 years	5 years	6 years	8 years	10 years	12 years
All cemented	2.3 (2.1-2.5)	4.4 (4.2-4.7)	5.6 (5.3-5.9)	7.5 (7.2-7.9)	8.4 (8.0-8.7)	10.2 (9.7-10.6)	12.3 (11.7-12.8)	14.5 (13.7-15.3)
All uncemented	3.8 (3.2-4.5)	5.7 (5.0-6.6)	7.3 (6.4-8.2)	10.0 (9.0-11.1)	11.1 (9.9-12.3)	12.8 (11.4-14.2)	16.4 (14.2-19.0)	19.1 (15.9-22.9)

Figure 5.31

Estimated failure rates of primary partial knee arthroplasty for main types of component fixation

Time since operation, 2012–2024, all diagnoses.

Type of PKA and revision

Estimated failure rates of PKA since 2012 were equal for medial and lateral PKA. PFJ performed much worse regarding revision rates, diverging already between one to two years after surgery. At 4 years of follow-up, revision rates were already twice as high as for medial or lateral PKA, maintaining a revision rate about twice as high until 12 years after primary PKA (Figure 5.32).

Technical support and revision

Comparing technical support during primary PKA, PSI did not perform better than conventional technique (Figure 5.33), as did conventional computer navigation (Figure 5.34). In contrast, robotically assisted PKA was associated with less revisions from early on, although the effect seemed to flatten out from the fifth year after primary PKA (Figure 5.34). At 6 years,

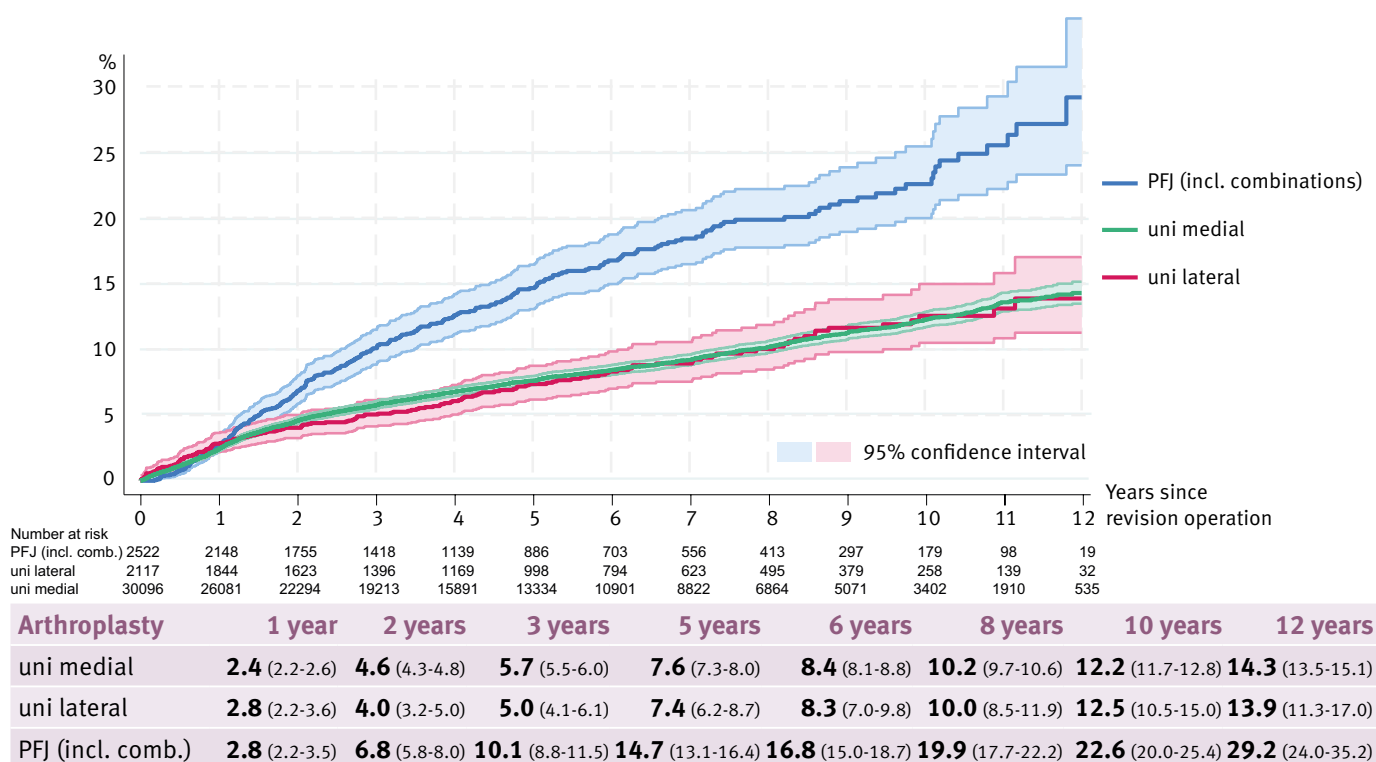


Figure 5.32

Estimated failure rates of primary partial knee arthroplasty: type of arthroplasty

Time since operation, 2012–2024, all diagnoses.

robotically assisted PKA had a revision rate of 6.8% (CI 4.1–11.3%) compared to conventional technique with 9.0% (CI 8.7–9.4%). The larger confidence intervals of robotic PKA due to smaller numbers rendered

the difference non-significant. More robotic systems with varying value in supporting surgery and more surgeons with less experience could deteriorate initially promising results, as had been observed in Australia.

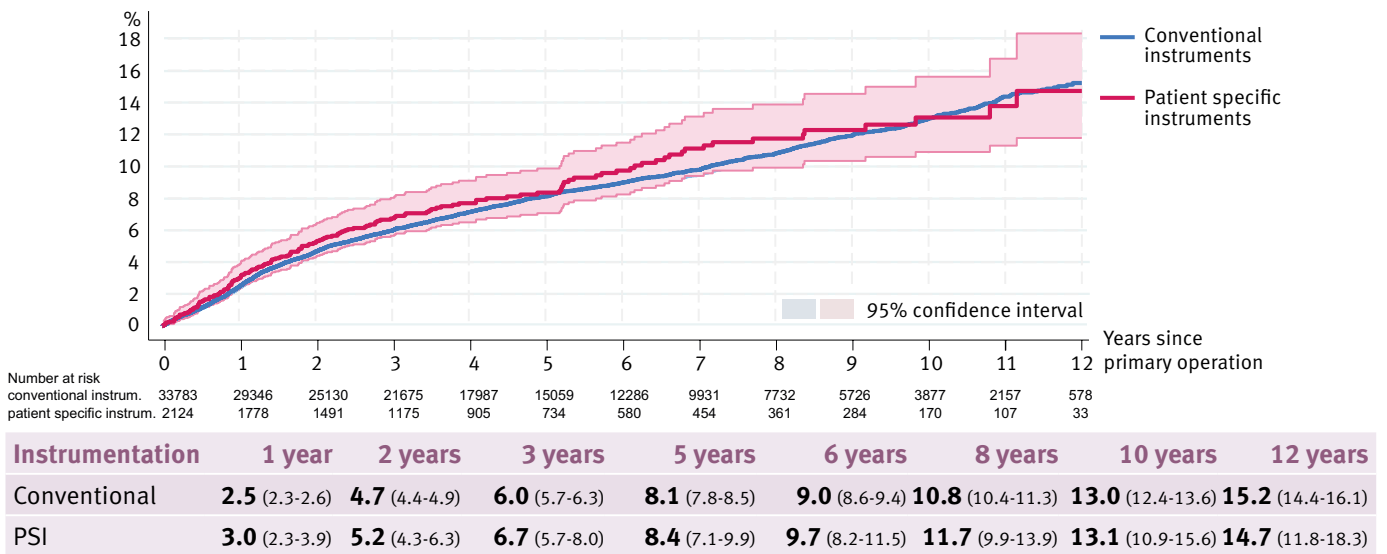


Figure 5.33
Estimated failure rates of primary partial knee arthroplasty: conventional vs. patient specific instrumentation
 Time since operation, 2012–2024, all diagnoses.

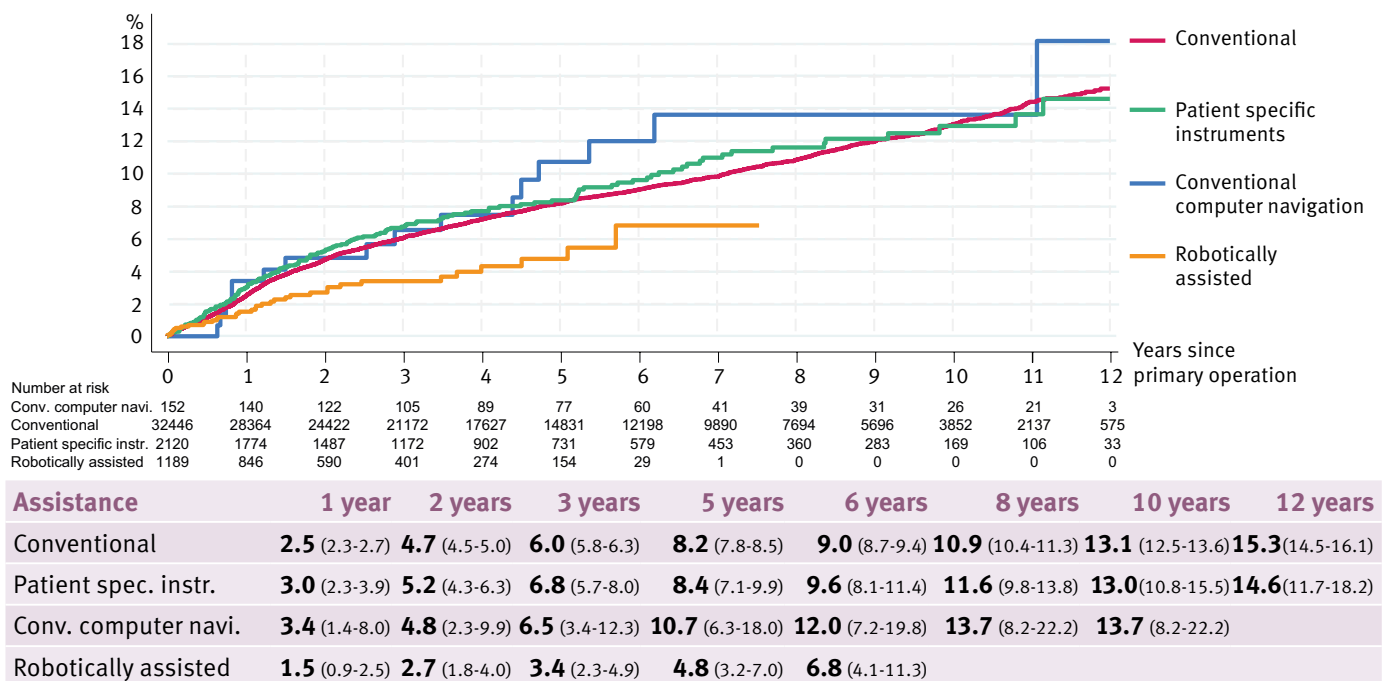


Figure 5.34
Estimated failure rates of primary partial knee arthroplasty: technological assistance
 Time since operation, 2012–2024, all diagnoses.

Type of early first revision	N revised	%
Conversion from unicom. to total prosthesis*	443	70.0
Exchange of PE liner	106	16.7
Tibial revision	25	3.9
Complete revision*	15	2.4
Femoral revision	12	1.9
Subsequent partial prosthesis, second compartment	12	1.9
Patella revision	11	1.7
Component removal with spacer implantation	5	0.8
Component removal without spacer implantation	2	0.3
Subsequent patella prosthesis	1	0.2
Reimplantation of prosthesis	1	0.2
Subsequent patella prosthesis with exchange of PE liner	0	0.0
Other	0	0.0
Total Revisions	633	100

Table 5.13

Type of early first revision of primary partial knee arthroplasty

All diagnoses, all component fixations. 4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

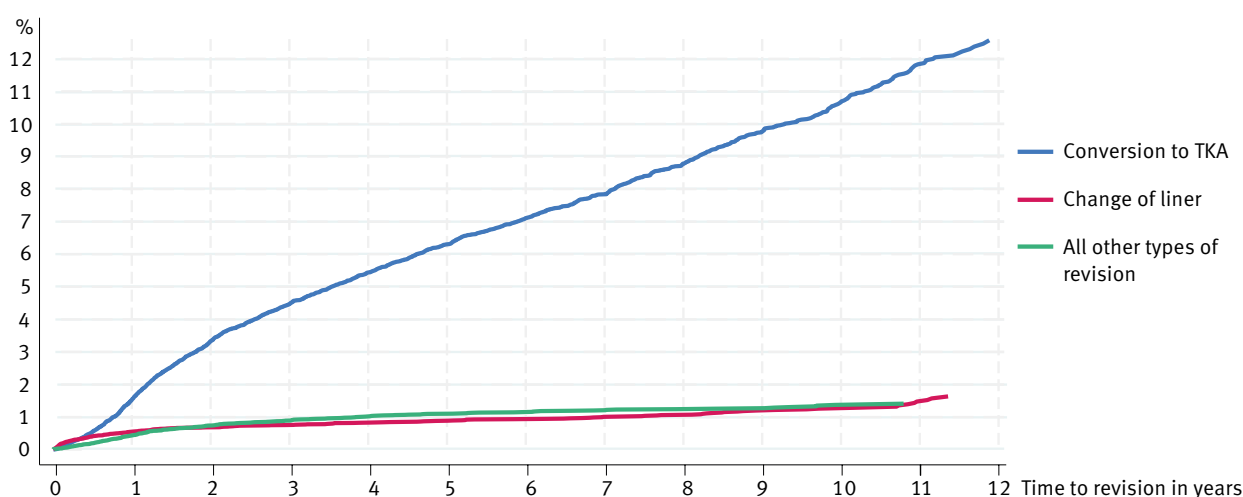
Early first revisions are those occurring within 2 years of the primary arthroplasty

* A large share of conversions is entered locally as “complete revisions”.

Such responses have been recoded as conversions if TKA components were registered or if a TKA was indicated on the revision proforma.

5.7.2 Type of early PKA revision

A total of 70.0% of the PKA revised were converted to TKA. Isolated liner exchange was performed in 16.7% of revisions, followed by an isolated tibial revision in 3.9% and isolated femoral revision in 1.9% (Table 5.13). Complete revision was performed in 2.4%, meaning conversion to TKA in many cases. These responses were reclassified as conversion if TKA components were registered at the revision. The high number of conversions to TKA in case of failed PKA is even more obvious in Figure 5.35 depicting cumulative incidence rates of all cases since 2012. Subsequent PKA in another compartment was rare (1.9%). Component removal with spacer due to PJI was performed in only 2 of the 443 revisions performed between 01.01.2019 and 31.12.2022. For re-revision surgery after primary PKA please consult Chapter 5.4.2.



Type of revision	1 year	2 years	3 years	5 years	6 years	8 years	10 years	12 years
Conversion to TKA	1.5 (1.4-1.7)	3.3 (3.1-3.5)	4.5 (4.2-4.7)	6.3 (6.0-6.6)	7.1 (6.8-7.4)	8.8 (8.4-9.2)	10.7 (10.2-11.2)	12.6 (11.8-13.3)
Change of liner	0.5 (0.5-0.6)	0.7 (0.6-0.8)	0.8 (0.7-0.9)	0.9 (0.8-1.0)	0.9 (0.8-1.1)	1.1 (0.9-1.2)	1.3 (1.1-1.4)	1.6 (1.4-2.0)
All other types	0.4 (0.4-0.5)	0.7 (0.6-0.8)	0.9 (0.8-1.0)	1.1 (1.0-1.2)	1.2 (1.0-1.3)	1.2 (1.1-1.4)	1.4 (1.2-1.6)	1.4 (1.2-1.6)

Figure 5.35

Cumulative incidence rates for different types of revisions of partial knee arthroplasty

Time since operation, 2012–2024, % of implants revised

5.8 Results of implants in unicondylar partial knee arthroplasty

The analysis is performed separately for unicondylar PKA (chapter 5.8) and for PFJ PKA (chapter 5.9), as the revision rates observed largely differed.

Table 5.14 shows Switzerland's current top ten of the most frequently used unicondylar PKA systems, representing 94% (17,588) of the PKA performed between 2019 and 2024. During this period, 1,050 implants (5.6%) belonged to the less common systems, grouped together as "other systems". The share of these is decreasing continuously over the years. Only 62 implants (0.3%) could not be classified.

System	2019	2020	2021	2022	2023	2024	2019 – 2024
Balansys uni	354	298	352	420	333	337	2,094
GMK uni	222	205	159	155	116	95	952
Journey uni	90	89	74	47	52	48	400
Moto	31	66	125	188	150	237	797
Oxford cemented/hybrid	314	270	254	210	199	271	1,518
Oxford cementless	319	354	321	362	344	362	2,062
Persona partial knee	427	412	445	517	592	603	2,996
Physica ZUK	251	329	333	403	469	468	2,253
Restoris MCK	128	110	112	178	227	290	1,045
Sigma partial knee	505	600	615	507	573	671	3,471
Other systems	245	175	198	182	140	110	1,050
Total	2,886	2,908	2,988	3,169	3,195	3,492	18,638

Table 5.14

Top 10 (94%) of primary partial knee arthroplasty systems (all diagnoses, all component fixations) 2019 – 2024

Important information on the use of the implant performance tables below

- Estimated revision rate exceeds the alert boundary, but we do not identify this implant combination as an outlier because the 95% confidence interval overlaps the confidence zone of the reference group.
- Identified as potential outliers. Please note the statistical confidence intervals. The outlier status comes with varying degrees of statistical probability. We consider the potential outlier status „highly likely“ when both the estimated revision rate and the complete confidence interval exceed the outlier alert boundary.

Please be aware that relatively rare implant combinations are frequently used in only a small number or indeed only in one hospital in Switzerland. Observed revision rates may be determined by local factors and performance may differ significantly between locations. Manufacturers of detected outlier implants and the hospitals where they were used (and revisions occurred) have been informed by SIRIS.

5.8.1 Two-year revision rates of unicondylar partial knee arthroplasty

The two-year revision rates of the unicondylar PKA from the current moving window is shown in **Figure 5.36**, reflecting how the implants used between 01.01.2019 and 31.12.2022, performed at a two-year follow-up by 31.12.2024. Of the 13 implant combinations used most frequently (the rest being summarised under “other systems”), none was identified as an outlier. The system reaching a definitive outlier

status at the third year (Journey Uni) after PKA had only a slightly elevated revision rate at two years, remaining within the boundaries of the average. This is reflective of recent improvements with this implant. In contrast, the summary group “other systems” had the highest two-year revision rate and remained only just below the outlier-boundary (Figure 5.36). Therefore, it is advisable for surgeons using less commonly used implants to carefully review their outcomes and consider switching to brands more frequently used in Switzerland if there is any uncertainty.

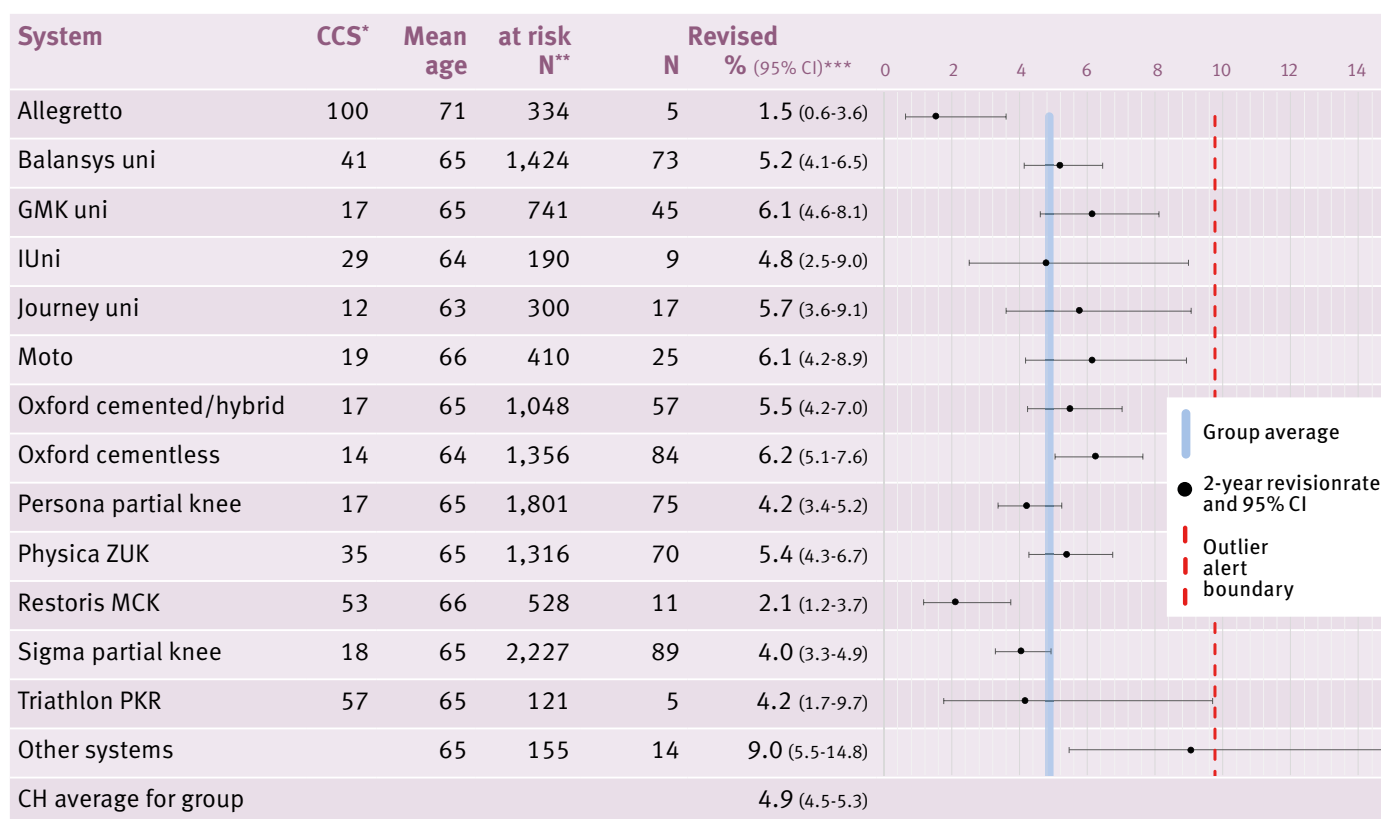


Figure 5.36

2-year evaluation: Revision rates of primary partial knee arthroplasty systems within 24 months

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

*** Rates adjusted for effects of mortality and emigration.

5.8.2 Long term survival of unicondylar partial knee arthroplasty

The long-term evaluation for all systems registered since 2012, all diagnoses, and all fixation methods is depicted in **Table 5.15**, showing results up to 12 years after surgery. The revision rate after 12 years for all systems was 14.5% (CI 13.7-15.3%), considerably higher than after TKA, where an average 12-year revision rate of 8.4% (CI 8.2–8.6%) was observed (**Table 5.9**).

Primary PKA subsystems (such as cemented or cementless) were analysed separately if numbers were sufficient and differed relevantly regarding revision rates. Different implants performed rather differently on the short, medium, and long term (**Table 5.15**).

The revision rate after 12 years varied from 5.9% for the best to 25.6% for the worst system. Wider confidence intervals reflect higher variability due to small numbers in the subgroups. Please take note of the case concentration score (CCS), indicating the share of the largest providing hospital. A higher value indicates an increased likelihood of bias due to local effects, i.e. individual providers may influence results of systems not widely used. For instance, Allegretto was implanted by just one surgeon, resulting in a CCS of 100. For some brands, only five-year results were available. Until then, they performed better than the Swiss average, with one system (Restoris MCK) performing even statistically significantly better (**Table 5.15**).

System	Total N	CCS*	Mean age**	1 year (95% CI)	3 years (95% CI)	5 years (95% CI)	10 years (95% CI)	12 years (95% CI)
Allegretto	1,074	100	70	0.5 (0.2-1.1)	1.7 (1.1-2.7)	3.2 (2.3-4.5)	5.9 (4.3-8.0)	5.9 (4.3-8.0)
Balansys uni	4,064	47	65	2.2 (1.8-2.7)	5.6 (4.8-6.4)	7.0 (6.1-7.9)	11.0 (9.7-12.5)	12.2 (10.5-14.1)
GMK uni	1,852	21	66	3.3 (2.6-4.2)	7.5 (6.4-8.9)	9.4 (8.0-10.9)	14.0 (11.8-16.5)	15.7 (12.7-19.4)
Journey uni	1,128	13	63	3.5 (2.6-4.8)	8.9 (7.4-10.8)	15.8 (13.6-18.3)	24.5 (21.3-28.2)	25.6 (22.1-29.5)
Moto	818	15	65	3.4 (2.3-5.1)	5.8 (4.2-8.0)	6.9 (4.9-9.7)		
Oxford cemented/hybrid	4,590	19	66	2.5 (2.1-3.1)	5.6 (5.0-6.4)	7.5 (6.7-8.4)	12.8 (11.6-14.1)	16.0 (14.2-18.1)
Oxford cementless	3,434	10	64	3.8 (3.2-4.5)	6.7 (5.9-7.7)	9.2 (8.2-10.4)	14.6 (12.3-17.3)	16.9 (13.3-21.3)
Persona partial knee	3,441	14	65	2.0 (1.6-2.6)	4.8 (4.1-5.7)	6.6 (5.5-7.8)		
Physica ZUK	4,477	20	65	1.9 (1.5-2.4)	5.5 (4.8-6.3)	7.0 (6.2-7.9)	11.8 (10.6-13.2)	14.7 (12.8-16.8)
Restoris MCK	1,080	48	65	1.2 (0.7-2.2)	3.0 (2.0-4.5)	4.2 (2.7-6.4)		
Sigma partial knee	6,013	18	65	2.2 (1.8-2.6)	5.4 (4.8-6.0)	6.9 (6.2-7.6)	9.7 (8.6-10.9)	11.2 (9.7-12.9)
Other systems	1,436		64	3.5 (2.7-4.6)	8.0 (6.6-9.6)	10.7 (9.0-12.7)	19.0 (15.8-22.8)	19.7 (16.3-23.7)
CH average for group				2.5 (2.3-2.6)	5.7 (5.5-6.0)	7.7 (7.4-8.1)	12.4 (11.9-13.0)	14.5 (13.7-15.3)

Table 5.15

Long term evaluation: Failure rates of primary partial knee arthroplasty systems (all diagnoses, all component fixations)

Time since operation, 2012–2024. Please note that if reported system involves multiple sub-variants, it is possible that the long-term performance of these sub-variants may be significantly different from their combined performance.

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects.

A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Younger mean age signifies that the case mix is less “usual” and potentially biased towards higher revision risk.

PKA systems grouped under the label “other systems” because of small numbers, accounting all together for 5.6% of the PKA, had an average revision rate at 12 years of 19.7% (CI 16.3 – 23.7%). This means that none of the less commonly used systems would reach a place in the midfield at twelve years of follow-up. One system (Journey Uni) (Figure 5.37) even had to be classified as a long-term outlier at 12 years follow-up. Interestingly, Journey Uni is from the same company as the potential outlier system

and the one with “elevated” long-term revision risk identified in TKA. As usual, the potential and definitive outlier identification will result in a detailed outlier report investigating the reasons for the observed deviations from the national average. As in former years, one PKA system (Allegretto) had a significantly better survival curve than the Swiss average, but it must be underscored that it was used in just one surgical unit (Figure 5.38).

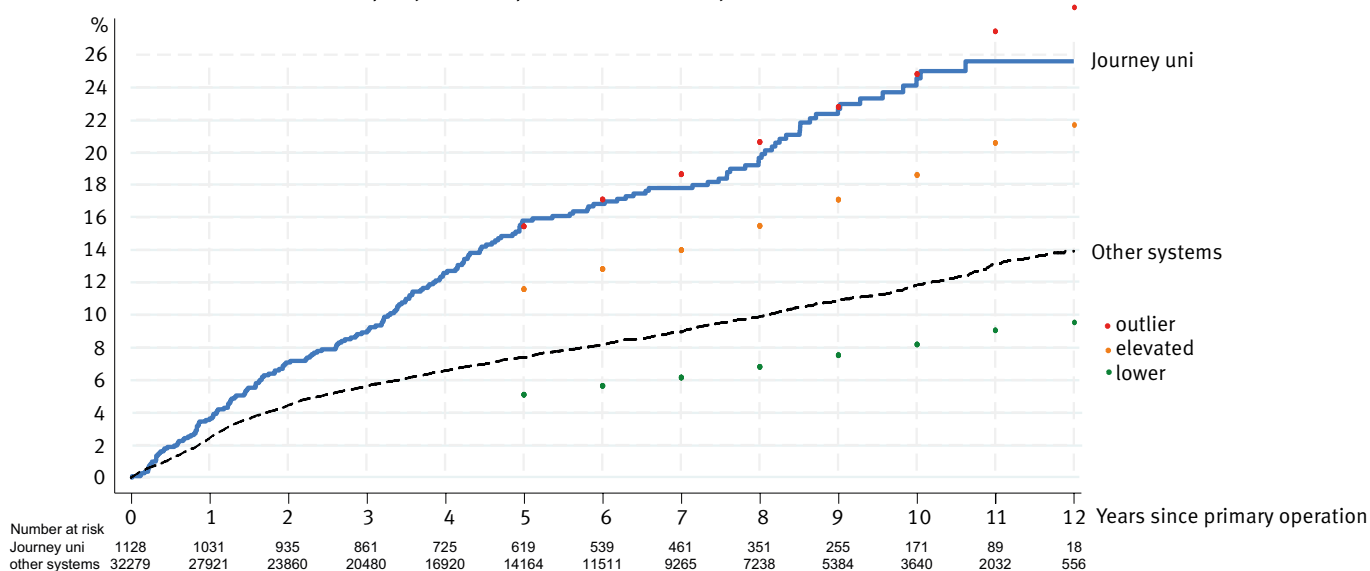


Figure 5.37

Implant combinations with long-term evaluation outlier status (all unicompartmental PKA)

Outlier status was defined as a revision rate of twice the group average at any time between year 5 and year 12 (and lower bounds of the 95% confidence interval exceeding the upper bound of the group average; and at least 50 cases at risk at 5 years). The dots indicate upper and lower limits.

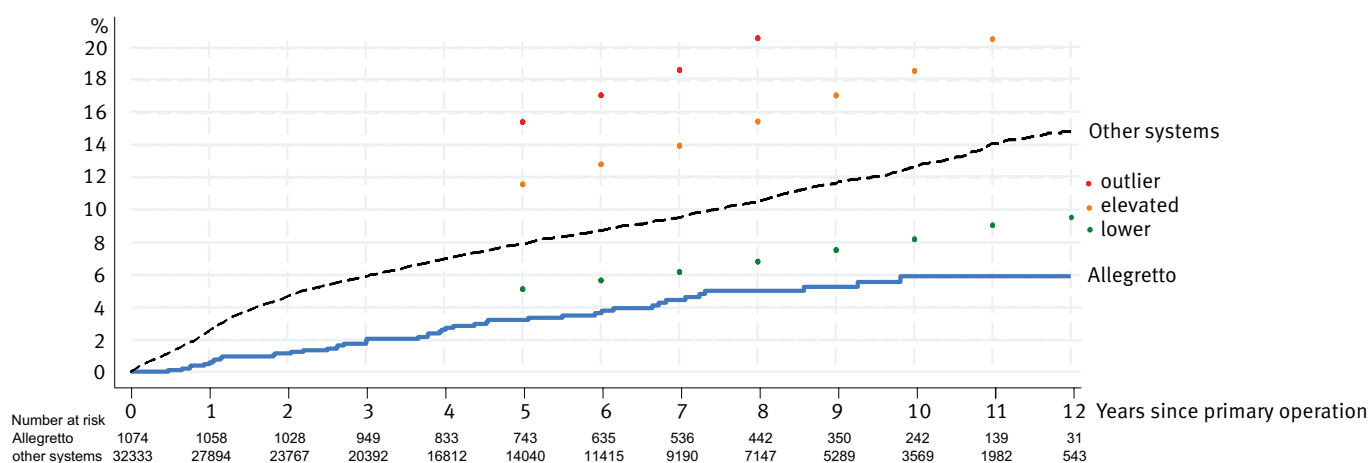


Figure 5.38

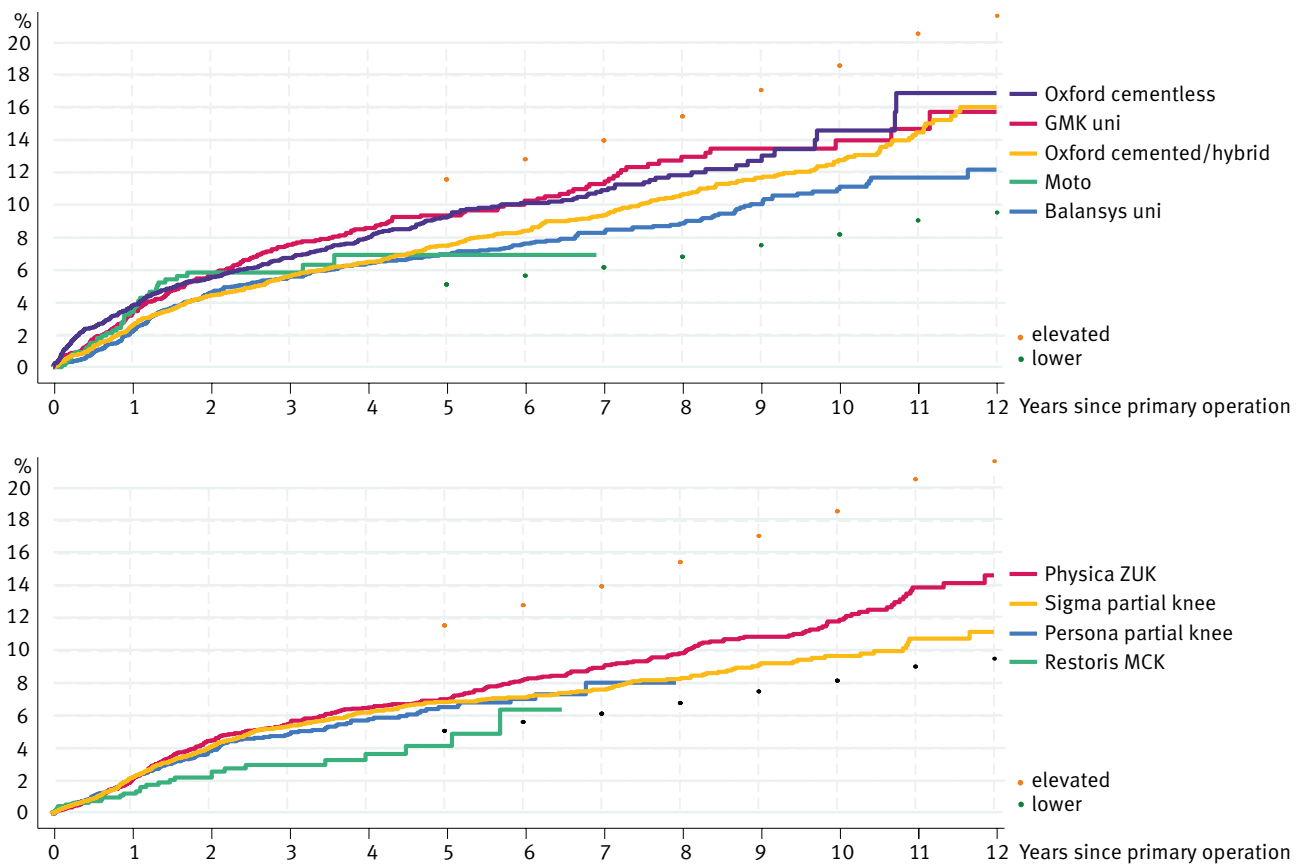
Implant combinations with below-average long-term revision rates (unicompartmental PKA)

Below-average was defined as an 9-year/10-year revision rate of up to 66% of the group average (and upper bounds of the 95% confidence interval staying below the lower bound of the group average; and at least 25 cases at risk at 9 years/10-years). The dots indicate upper and lower limits.

These excellent results are most probably to be explained by the outstanding expertise of the surgeon than by the system itself. With numbers decreasing rapidly in recent years, it is expected that this system will not be used anymore in the future, either because the surgeon changed implant or is retiring. The remaining brands of PKA had revision rates within the margins of the lower and upper limits at 66% and 150% of the group average, respectively (Figures 5.39).

5.8.3 Brands of unicondylar PKA

Figures 5.39 show the performance of the different PKA brands up to twelve years of follow-up, with upper and lower limits of the Swiss average illustrated in dotted lines. All the systems evaluated performed at the level of the Swiss average. One older and one newer brand performed better than average, as did a system which is implanted with image-based robotic assistance, but results are limited to 6 years of follow-up. Visible steps to the worse from the fourth year after primary surgery are again related to limited numbers (Figures 5.39).



Figures 5.39

All remaining implant combinations with average revision risks (all PKA)

Also showing upper and lower limits (corresponding to elevated and below-average version risk at 150% and 66% of the group average respectively).

5.9 Results of implants in patellofemoral joint partial knee arthroplasty

Table 5.16 shows Switzerland's current top five of the most frequently used patellofemoral PKA systems, representing 98% (1,513) of the PFJ performed between 2019 and 2024. Only 32 implants (2.1%) in this period belonged to the less common systems, grouped as "other systems". Quite a significant share of 61 implants (3.9%) could not be identified, which is more than the 32 patellofemoral PKA summarised under "other systems".

5.9.1 Two-year revision rate of patellofemoral PKA

The two-year revision rates of the individual brands are shown in Figure 5.40, reflecting the PKA performed between 01.01.2019 and 31.12.2022, with a completed two-year follow-up by 31.12.2024. Of the five implant systems registered, including the rest summarised under "other systems", none was a definitive outlier, although one system (IBalance PFJ) was above the outlier boundary but was not classified as such due to statistical uncertainty. One system implanted with image-based robotics (Restoris) was above the outlier boundary but was not classified as such due to statistical uncertainty. One system implanted with image-based robotics (Restoris)

System	2019	2020	2021	2022	2023	2024	2019 – 2024
Gender PFJ	109	162	110	167	154	142	844
Hemicap PF classic/wave (PFJ)	24	33	40	48	28	29	202
IBalance PFJ	17	24	17	30	29	15	132
Journey PFJ	18	20	20	9	19	15	101
Restoris MCK PFJ	24	25	28	32	58	35	202
Other systems	0	6	13	5	4	4	32
Total	192	270	228	291	292	240	1,513

Table 5.16
Top 5 (98%) of primary patellofemoral joint systems (all diagnoses, all component fixations) 2019 – 2024

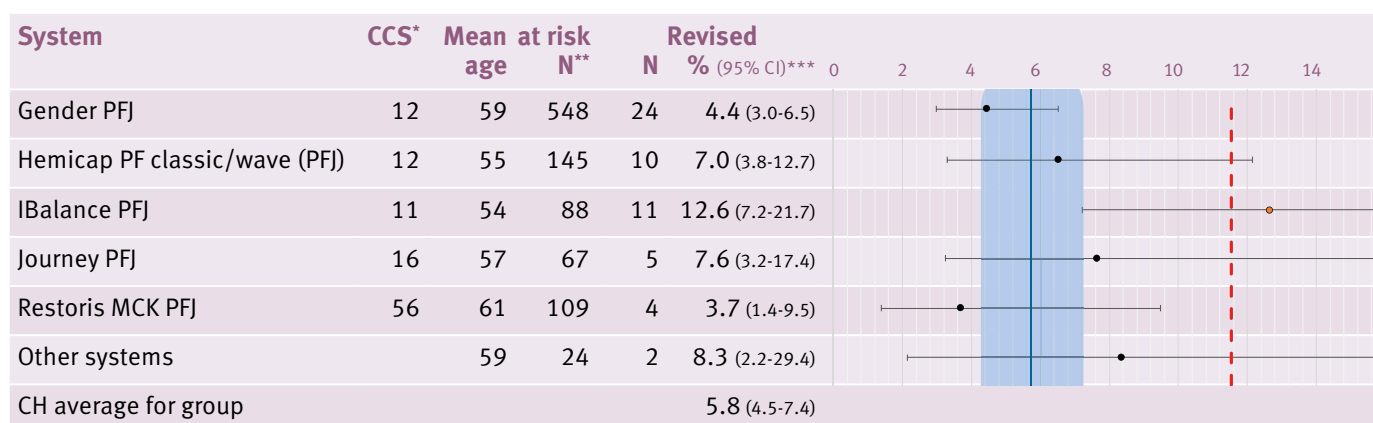


Figure 5.40
2-year evaluation: Revision rates of primary patellofemoral joint systems within 24 months

4-year moving average covering implants between 01.01.2019 and 31.12.2022, with two years follow-up (31.12.2024).

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects. A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Number of patients with at least two years follow-up (i.e. primary prosthesis in moving average).

*** Rates adjusted for effects of mortality and emigration.

Group average
● 2-year revision rate and 95% CI
Outlier alert boundary

MCK PFJ) and the most frequently used PFJ system (Gender PFJ) had better results than the average. This is of special interest as Gender PFJ had a low case concentration score (CCS) of 12% and therefore multiple providers. This would rather be a risk for elevated revision rates due to small individual caseloads. Therefore, the results speak in favour of a reliable system. In contrast, Restoris MCK PFJ system had a CCS of 56%, thus concentrated in the hand of fewer surgeons with potentially higher expertise beside robotic assistance.

5.9.2 Long term survival of patellofemoral PKA

Numbers are too small to differentiate each of the five brands. Therefore, comparison was reduced to the brand used most commonly and all the rest of the implants, summarised under “other systems” (Table 5.17). The singular system performed significantly better than all the other systems together from the first year on, though results beyond 10 years of follow-up were not yet available. It is essential to realise that revision rates of PFJ replacements are more than three times higher than those of unicondylar PKA eleven years after surgery. At 7 years, the most frequently used and best performing PFJ system had a revision rate more than double of the unicondylar PKA average (Table 5.17, Figure 5.30).

System	Total N	CCS*	Mean age**	1 year (95% CI)	3 years (95% CI)	5 years (95% CI)	10 years (95% CI)	12 years (95% CI)
Gender PFJ	1,238	8	59	2.2 (1.5-3.2)	7.9 (6.3-9.8)	11.8 (9.8-14.3)	19.7 (15.6-24.7)	
Other systems	1,212		57	3.4 (2.5-4.6)	12.1 (10.2-14.3)	17.1 (14.7-19.8)	26.1 (22.5-30.2)	34.4 (27.0-43.2)
CH average for group				2.8 (2.2-3.5)	10.0 (8.7-11.4)	14.5 (12.9-16.3)	23.1 (20.3-26.1)	30.1 (24.4-36.8)

Table 5.17

Long term evaluation: Failure rates of primary patellofemoral joint systems (all diagnoses, all component fixations)

Time since operation, 2012–2024. Please note that if reported system involves multiple sub-variants, it is possible that the long-term performance of these sub-variants may be significantly different from their combined performance.

* Share of implants accounted for by main user hospital service. A higher share signifies an increased likelihood of biased figures due to local effects.

A share of 50%+ would suggest that reported results are likely determined by one hospital service.

** Younger mean age signifies that the case mix is less “usual” and potentially biased towards higher revision risk.



Patient reported outcome measures
PROMs

6. Patient reported outcome measures (PROMs)

Since 2019, the core components of the new Patient-Reported Outcome Measures (PROMs) instrument have been implemented in the Canton of Zürich and selected hospitals, ahead of the nationwide registration within SIRIS scheduled for late 2025. For this report, we analysed complete pre- and post-operative data for 9,065 total hip arthroplasties (THA), 7,087 total knee arthroplasties (TKA), and 1,351 partial knee arthroplasties (PKA) performed for primary osteoarthritis. Additionally, we included 974 THA and 991 TKA cases for secondary osteoarthritis. PROMs were assessed using the Swiss Orthopaedics Minimal Dataset (SO-MDS), which comprises joint-specific pain and satisfaction evaluations via a numeric rating scale (NRS), as well as general quality of life assessment using the EQ-5D-5L score. Further details on the project and coverage rates are provided in the methods chapter. All analyses adopt an intention-to-treat (ITT) approach, meaning revised implants are included unless stated otherwise.

6.1 Joint-specific pain

Pain reduction is a central outcome measure for arthroplasty. Preoperative pain levels were generally similar across groups, with the exception of secondary hip osteoarthritis, which showed notably higher values. Joint-specific pain was assessed using a 0–10 numeric rating scale (NRS). Mean preoperative pain scores for primary osteoarthritis were 6.60 for total hip arthroplasty (THA), 6.64 for total knee arthroplasty (TKA), and 6.38 for partial knee arthroplasty (PKA). For secondary osteoarthritis, mean scores were 7.06 for THA and 6.42 for TKA. As illustrated in **Figure 6.1**, the distribution of preoperative pain scores was relatively consistent across most groups, with the majority of patients reporting values of seven or eight. Patients undergoing PKA experienced slightly less pain than those undergoing TKA, particularly at the higher end of the scale, while patients with secondary hip osteoarthritis reported the highest proportion of severe pain (scores of 9 or 10). Postoperatively, mean pain scores decreased substantially: to 0.86 for THA, 1.64 for TKA, and 1.73 for PKA in primary osteoarthritis, and to 1.09 for THA and 1.68 for TKA in secondary osteoarthritis. The postoperative distribution of pain scores was effectively reversed, with a significantly larger proportion of THA patients achieving complete pain remission (**Figure 6.2**).

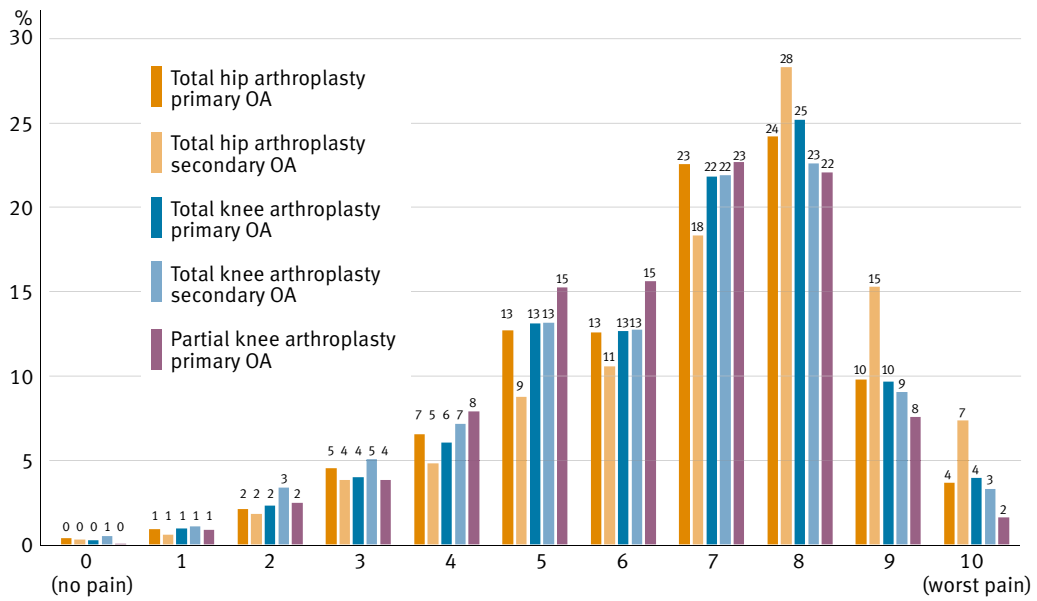


Figure 6.1
Preoperative pain

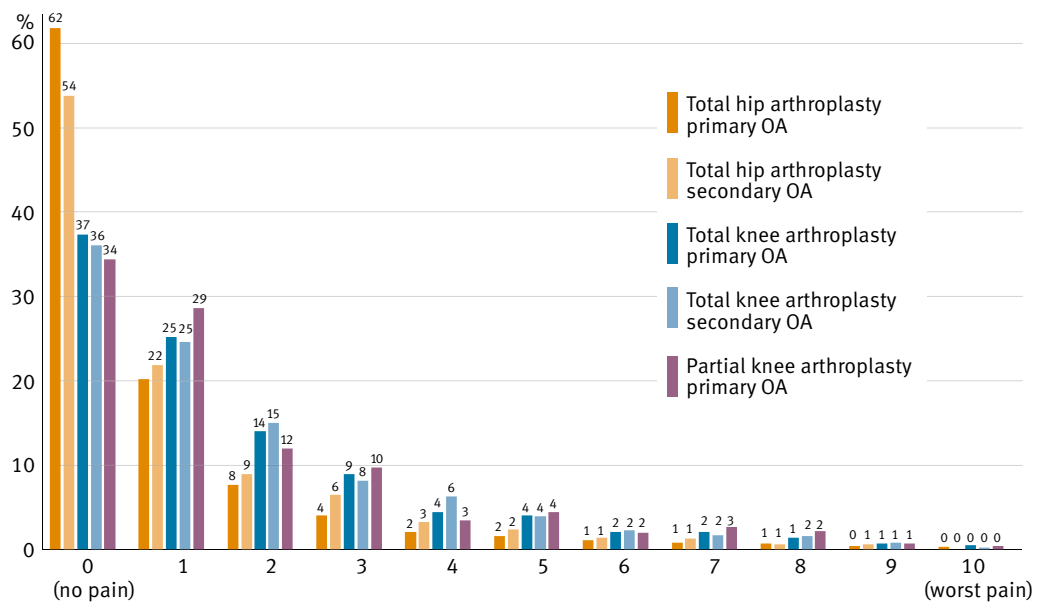


Figure 6.2
Postoperative pain
Follow-up 1 year (9 to 18 months allowed), intention-to-treat perspective

6.2 Quality of life

The EQ5D-5L instrument was used to assess health-related quality of life before and after treatment across five dimensions: pain, mobility, self-care, ability to perform usual activities, and anxiety. Among these, mobility stands out as the undisputed core dimension in elective arthroplasty.

As illustrated in **Figure 6.3**, hip (THA) and knee (TKA) patients exhibited remarkably similar mobility restrictions prior to treatment. The majority reported at least moderate problems, with over a third of both THA and TKA patients experiencing severe mobility limitations. Partial knee arthroplasty (PKA) patients, on average, reported slightly fewer restrictions, while THA patients with secondary arthritis were the most likely to report severe mobility issues. Notably, a small but significant proportion of patients in each group—ranging from 4.1% to 9.8%—reported no mobility problems at all before treatment. Post-treatment, the situation improved markedly, though outcomes varied by procedure (**Figure 6.4**). 78.5% of THA patients reported no further mobility problems, compared to 60.7% of TKA and 64.0% of PKA patients. Despite TKA and PKA patients starting from a better baseline, THA patients achieved the most substantial gains in mobility following treatment.

Radar charts are highly effective for visualising multi-dimensional concepts, such as those assessed by the EQ5D-5L. In this instrument, all items are measured on a consistent 5-point ordinal scale, ranging from “no problem” to “severe problem.” For the purposes of these visualisations, the scale was recoded so that the worst possible quality of life state is represented as 0 and the best as 4. **Figure 6.5a** presents the pre- and postoperative mean values for each EQ5D-5L dimension among THA patients with primary osteoarthritis (prim OA). At a glance, it is clear that mobility, pain/discomfort, and usual activities were the most affected areas before surgery. For instance, the mean mobility score was approximately 2.2, corresponding to the modal category of 41.9% of THA (prim OA) patients reporting moderate problems. In contrast, self-care and anxiety/depression were rarely problematic prior to treatment. Postoperatively, substantial improvements were observed not only in the three main problem areas but also, to a lesser extent, in self-care and anxiety/depression. THA patients with secondary osteoarthritis (sec OA) exhibited worse baseline

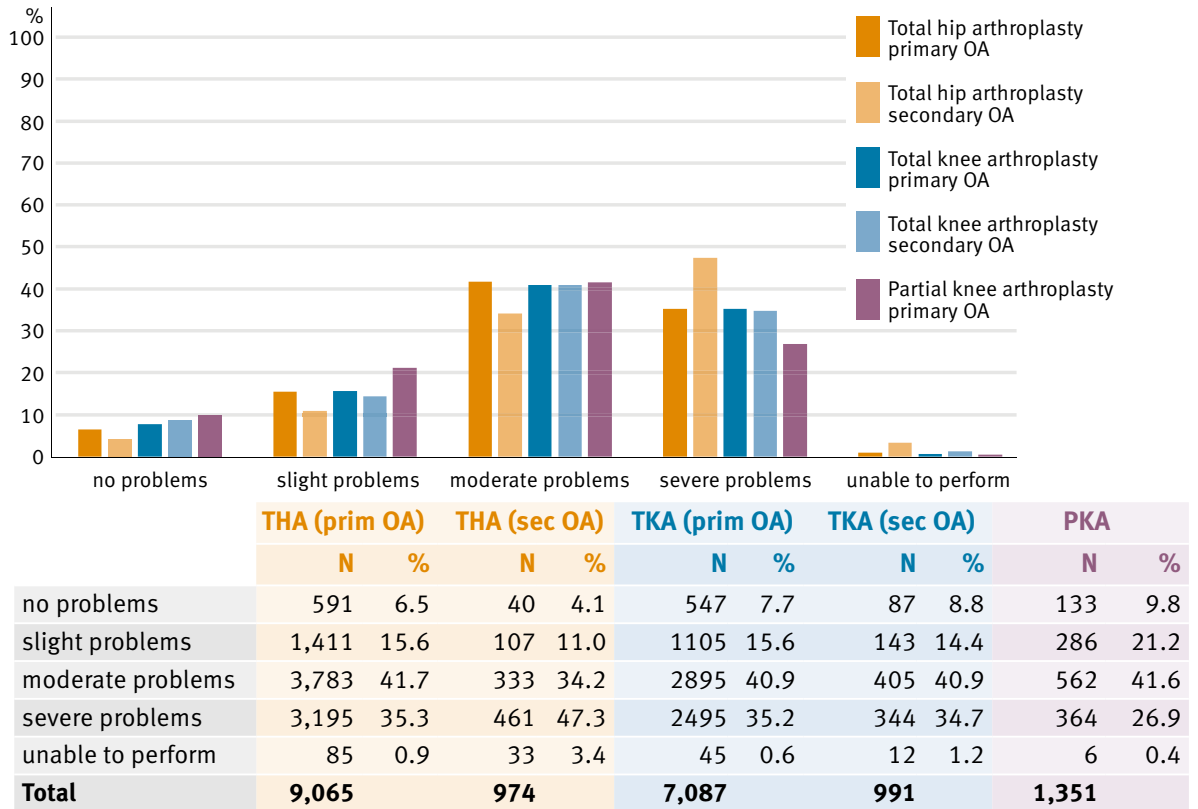


Figure 6.3

Preoperative quality of life (EQ5D): mobility

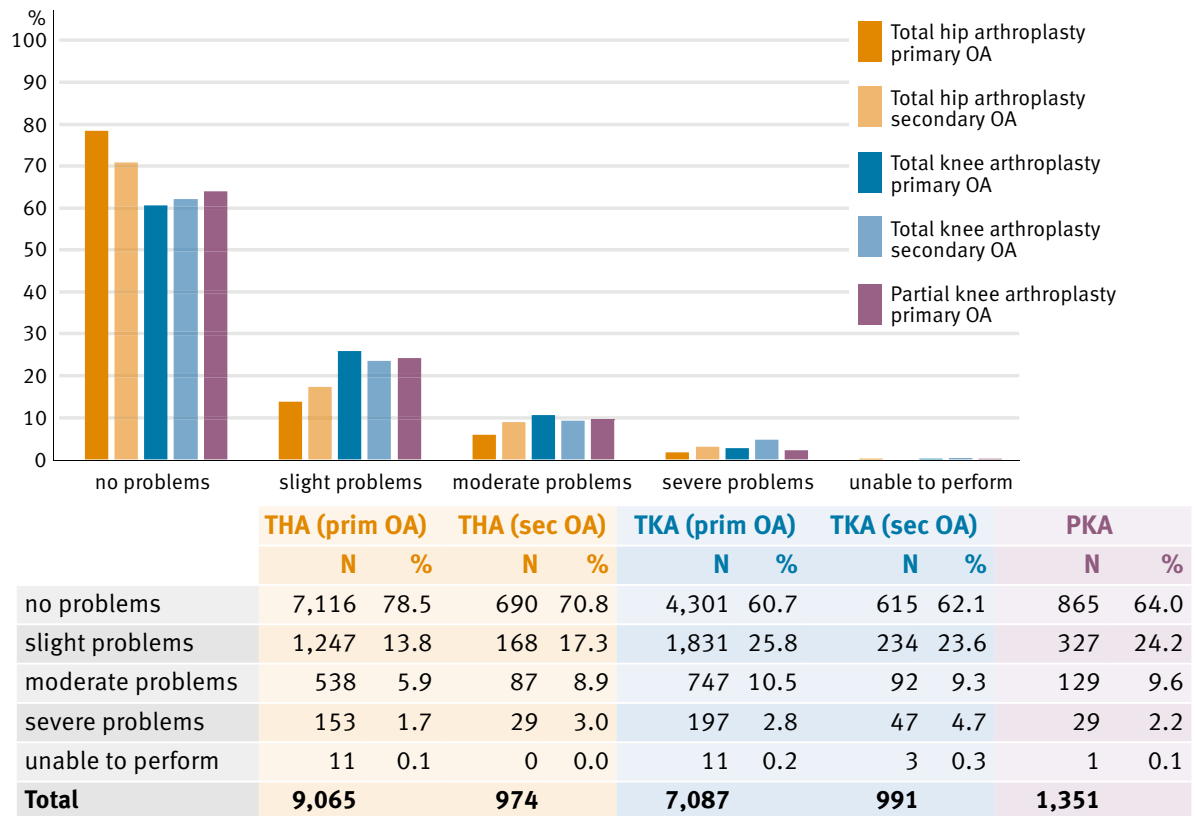


Figure 6.4

Postoperative quality of life (EQ5D): mobility

Follow-up 1 year (up to 18 months allowed), intention-to-treat perspective

values and, on average, achieved less favourable outcomes (Figure 6.5b). Figures 6.5c, d, and e depict similar patterns for knee patients (TKA and PKA), though the magnitude of improvement across all five dimensions was consistently less pronounced compared to THA patients with primary osteoarthritis.

6.3 Joint-specific satisfaction

Comparisons of preoperative and postoperative satisfaction ratings further corroborate the previous findings. Before surgery, hip and knee patients reported similarly high levels of dissatisfaction, with 72.5% to 84.7% stating they were very dissatisfied with their situation. One year after the operation, this picture was largely reversed. However, a clear satisfaction gap persisted between hip and knee patients.

While 81.2% of hip patients with primary osteoarthritis reported being very satisfied, the corresponding figures for knee implant recipients ranged from only 60.3% to 63.8% (Figures 6.6 and 6.7). Although a larger proportion of knee patients expressed dissatisfaction with their outcomes, the primary difference lay not in outright dissatisfaction but rather in the degree of satisfaction—many knee patients fell into intermediate or less conclusive categories rather than extreme dissatisfaction.

6.4 Treatment effects

Treatment success can be directly assessed by calculating the treatment effect. Figure 6.8 compares the distribution of pain reduction outcomes across the five patient groups, excluding the small minority who reported no preoperative pain.

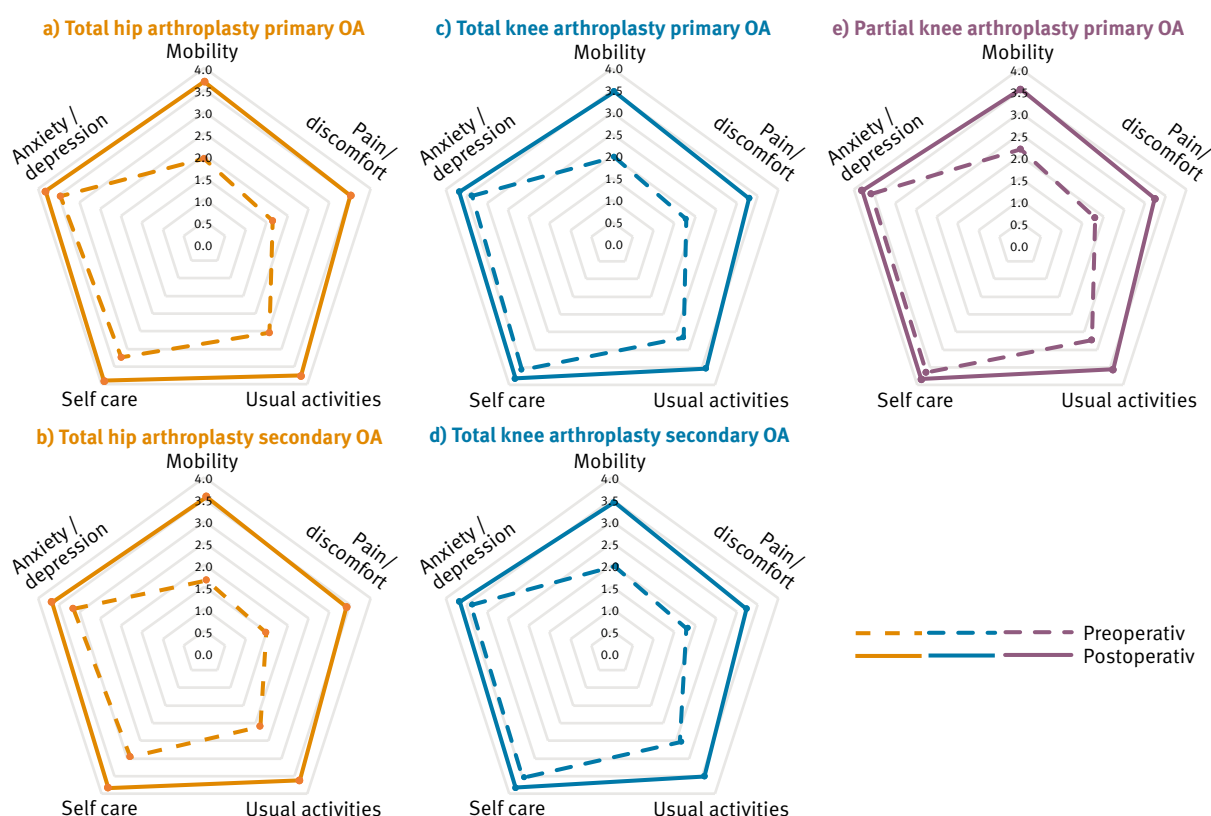


Figure 6.5

Quality of life (EQ-5D-5L): overview

Follow-up 1 year (up to 18 months allowed), intention-to-treat perspective

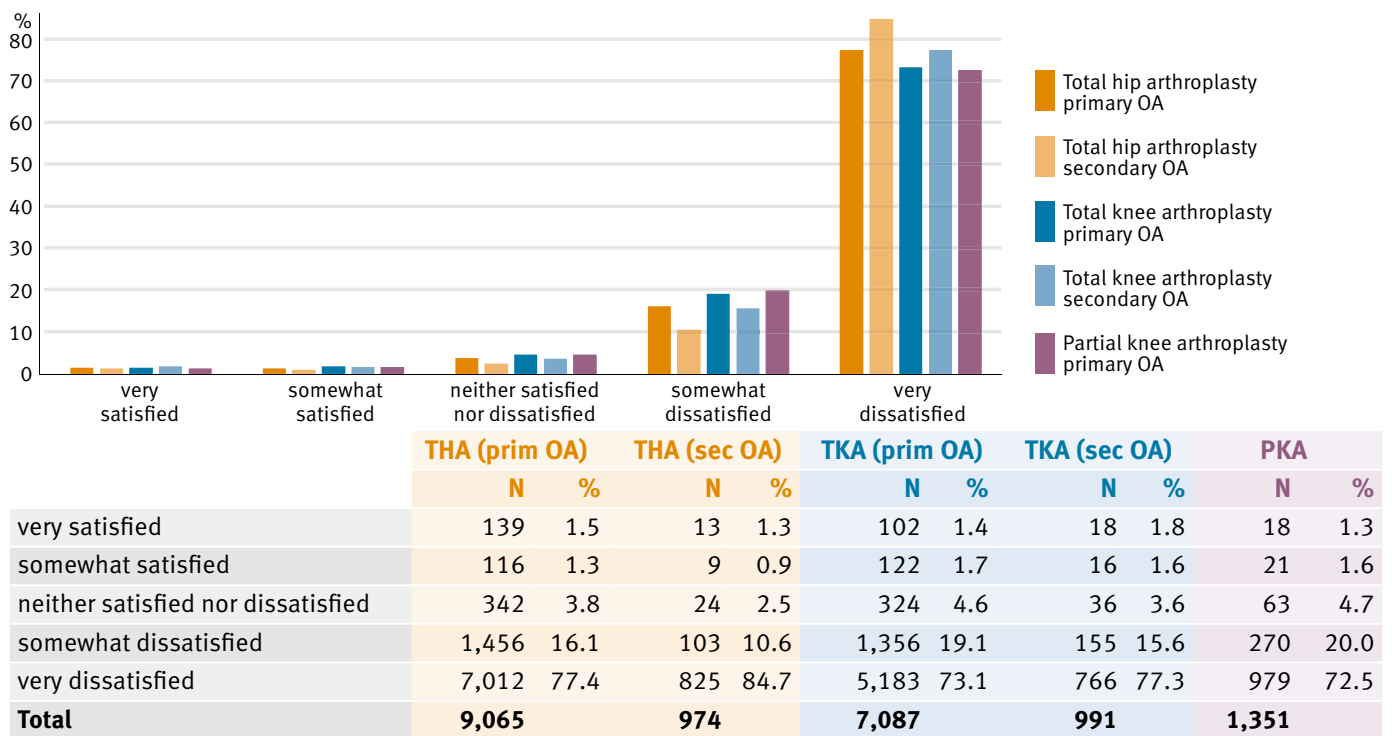


Figure 6.6
Preoperative satisfaction with current situation

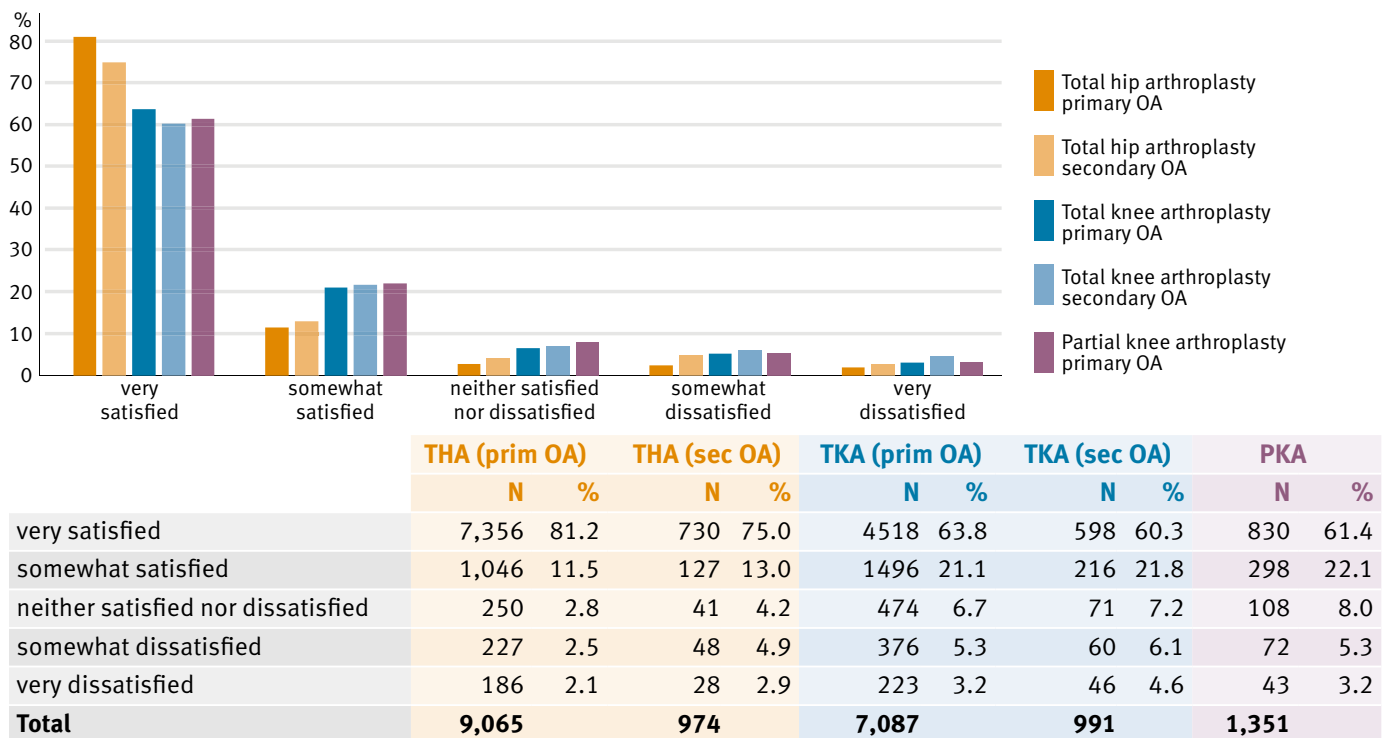


Figure 6.7
Postoperative satisfaction with current situation

Follow-up 1 year (9 to 18 months allowed), intention-to-treat perspective

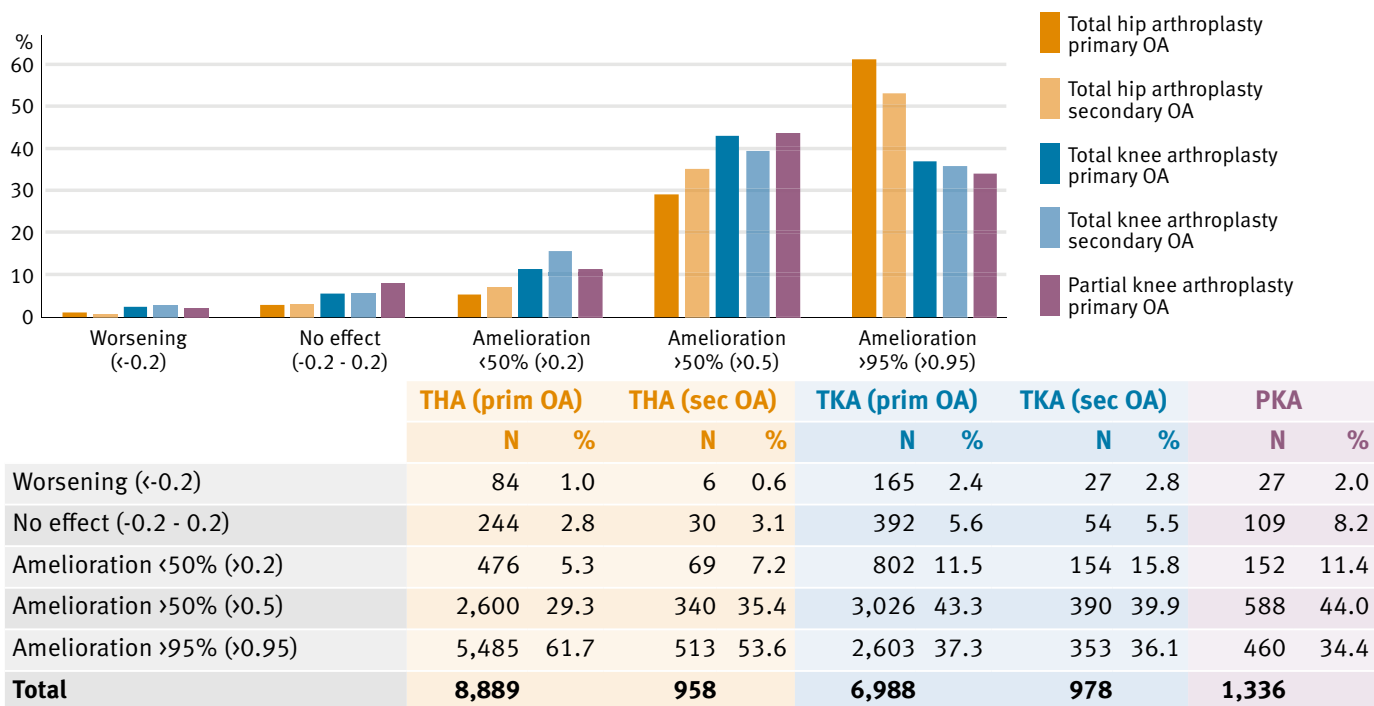


Figure 6.8

Treatment effect pain

Follow-up 1 year (9 to 18 months allowed), intention-to-treat perspective . Patients without reported pain excluded as calculation of TE not possible

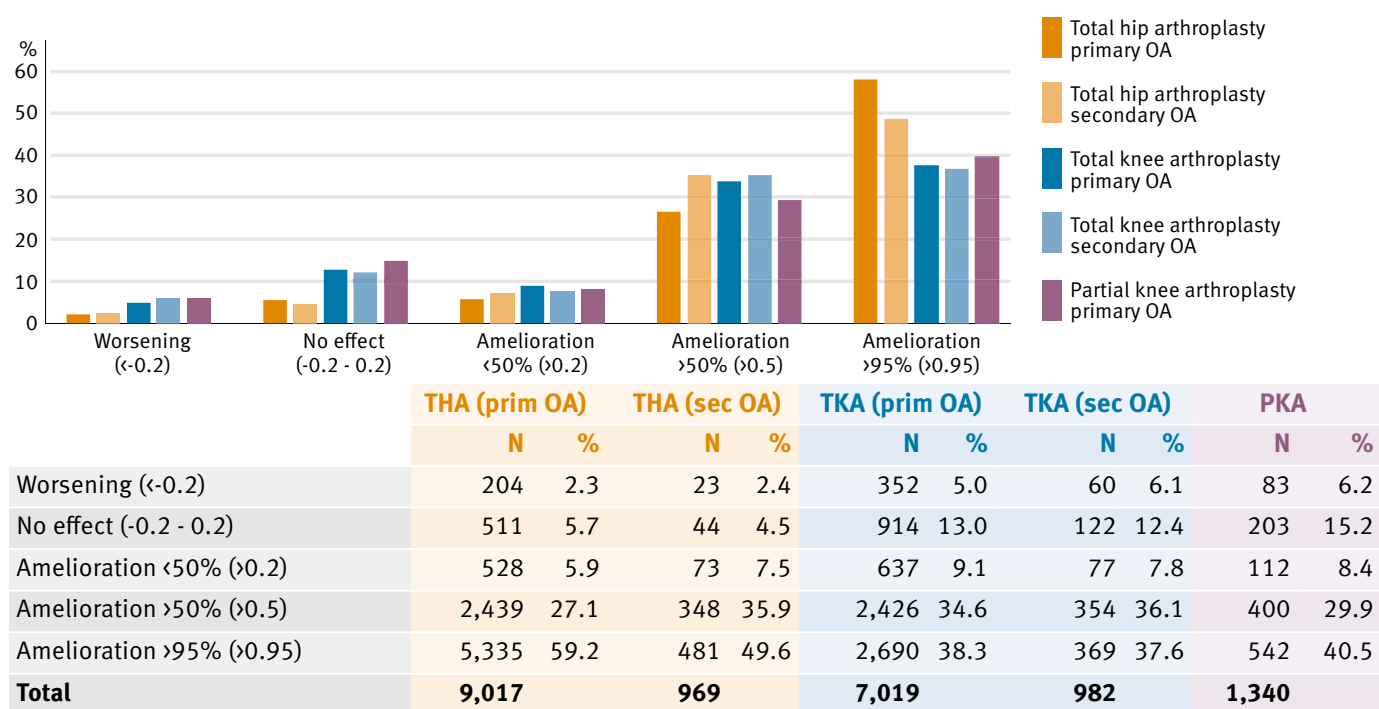


Figure 6.9

Treatment effect quality of life (EQ5D)

Follow-up 1 year (9 to 18 months allowed), intention-to-treat perspective . Patients without reported pain excluded as calculation of TE not possible

THA patients with primary osteoarthritis achieved the most favourable results: 61.7% experienced a symptom reduction of at least 95%, effectively becoming pain-free after treatment. An additional 29.3% reported a pain reduction of more than 50%, classified as a good outcome. For example, a patient with a pre-operative pain score of 8 typically reported a postoperative score of no higher than 3. Only a small fraction of THA patients had less favourable outcomes: 2.8% reported no improvement, and 1.0% experienced increased pain one year after surgery. Notably, these outcomes were observed across the full range of pre-operative pain levels. For THA patients with secondary osteoarthritis, the rate of pain-free outcomes was lower (53.6%), but the proportion of patients who did not benefit at all was not higher than in the primary osteoarthritis group.

Knee patients, however, had significantly less favourable outcomes: only 34.4% to 37.3% achieved a perfect, pain-free result. While over 40% experienced a pain reduction of more than 50%, a substantial minority – 8.0% to 10.2% – reported no improvement in pain. The differences between TKA and PKA outcomes were not statistically significant, despite PKA patients generally being younger and more active, with higher expectations for both pain reduction and functional restoration.

The treatment effect on the EQ5D quality of life measure – which encompasses functional limitations – mirrored the patterns observed for pain outcomes. This analysis was based on the EQ5D-5L summary score. Overall, hip patients achieved somewhat better outcomes than knee patients, while results for TKA and PKA were largely comparable within the limits of statistical precision. However, PKA outcomes exhibited greater variability than those for TKA. Notably, the higher proportion of poor TKA results, relative to THA, was statistically significant. Furthermore, the share of patients who did not experience an improvement in overall quality of life was substantially larger than the proportion who saw no benefit in pain reduction alone. This was true for both hip and knee patients. This discrepancy may partly reflect the broader, less joint-specific nature of the EQ5D measure. Nevertheless, the proportion of poor outcomes among hip patients remained less than half that observed in knee patients (Figure 6.9).

6.5 Conclusion

In summary, patient-reported outcome measures (PROMs) offer a more comprehensive assessment of the clinical effectiveness of hip and knee arthroplasties, complementing revision rates, which capture only a single endpoint of treatment outcomes. However, the current geographical coverage of these PROMs remains limited, raising concerns about the generalisability of the findings. The data are drawn predominantly from the Canton of Zurich, and factors such as implant selection, surgical practices, and cultural influences may not be fully representative of the broader Swiss context. The planned nationwide integration of PROMs into the Swiss Implant Registry, scheduled for late 2025, is expected to address these limitations and provide a more complete and representative overview of arthroplasty outcomes across Switzerland.

SIRIS outlier watch list – hip implants

Implant or implant combination	Risk-adjusted hazard ratios for either 2-year revision risk or overall revision risk (long-term evaluation)							
	Detected as outlier in report	Adjusted for age and sex			Adjusted for age, sex, BMI, ASA and Charnley Class (from 2015, if available)			
	HR	lb95%	ub95%	HR	lb95%	ub95%		
Uncemented stem/cup combinations (primary osteoarthritis)								
Active outlier implants				Summary				
CLS Spotorno + Allofit	2025	1.38	1.05	1.80	1.55	1.08	2.23	The CLS Spotorno + Allofit implant combination demonstrated unremarkable revision performance between 2012 and 2019. However, an increase in early revision risk was observed starting in 2020, with relatively high rates persisting through 2021 and 2022. Given the limited number of main users during this period, the possibility of an adverse local effect cannot be ruled out. Preliminary data for 2023—following the involvement of a new main user—show no cause for concern. Recommended course of action: Investigate the underlying reasons for the rise in revision risk after 2019.
SPS evolution + April poly	2025	1.69	1.00	2.86	1.26	0.41	3.91	The SPS evolution + April poly implant combination has previously avoided being classified as a potential outlier because of low statistical precision. While it exhibited unremarkable revision performance between 2012 and 2019, its usage has since declined. More recently, revision rates have risen to outlier levels. Currently, this implant combination is only registered in one remaining hospital and very infrequently.
Twinsys + RM pressfit	2025	1.97	1.22	3.17	2.04	1.02	4.09	The Twinsys + RM pressfit implant combination has previously avoided being classified as a potential outlier due to low statistical precision. While it exhibited unremarkable revision performance between 2012 and 2018, its usage has since declined. More recently, revision rates have risen to outlier levels and are now very close to definitive outlier status. Currently, this implant combination is only relatively infrequently registered in two remaining hospitals.
Nanos + R3	2024	2.25	1.28	3.97	2.76	1.24	6.15	Nanos + R3 has been used in small numbers in only a few hospitals and revision performance varies between hospitals, which raises the possibility of local effects. The statistical precision of the potential outlier status is low. The R3 cup is widely used and has an excellent performance record. The Nanos stem, on the other hand, is mostly used in this particular combination. It is noteworthy that half of early revisions were due to dislocation. Recommended course of action: investigate reasons for revision, especially dislocations, and observe future performance.
	2025	2.16	1.23	3.82	2.58	1.16	5.76	
Accolade II + Trident II	2023	2.96	1.54	5.69	2.60	1.08	6.26	The Accolade II + Trident II implant combination was first registered in 2018, with significant usage beginning in 2019. Currently, the risk-adjusted 2-year revision risk for this combination exceeds the critical threshold of 2. However, statistical precision remains limited at this stage. Notably, implants from the early years (2018 and 2019) exhibited a particularly high 2-year revision risk. In contrast, data for implants from 2020 onwards are, as yet, inconclusive. Infection accounted for 40% of early revisions—an unusually high proportion—suggesting that the early revision performance may have been adversely affected by random events or specific circumstances. Based on the current trajectory, this implant combination is expected to lose its outlier status in the 2026 annual report, as the poorly performing implants from 2019 will no longer be included in the observation window.
	2024	2.89	1.55	5.38	2.79	1.25	6.21	
	2025	2.66	1.43	4.94	2.82	1.26	6.28	
Symbol + Symbol DMHA/DS evol.	2023	2.62	1.31	5.23	2.40	1.19	4.81	The Symbol + Symbol DMHA implant combination was initially registered in 2019, with a notable increase in registrations occurring in 2020. Importantly, only the original Symbol DMHA cups were recorded in this combination; none of the equivalent DS evolution cups were included. During the most recent evaluation period, the risk-adjusted 2-year revision risk for this combination has fallen below the critical threshold of 2. However, statistical precision remains very low. It is worth noting that infections accounted for half of all early revisions—a proportion that is unusually high. This raises concerns that the observed performance may have been adversely affected by random events, particularly at one specific hospital. To date, the performance of implants used in 2023 and 2024 has been unremarkable. Should this positive trend continue, the implant combination may no longer be classified as an outlier in the 2026 annual report.
	2024	1.91	1.03	3.56	1.80	0.97	3.36	
	2025	1.72	1.02	2.91	1.48	0.86	2.56	
SPS evolution + April ceramic	2020	2.22	1.72	2.88	3.67	2.47	5.47	The SPS Evolution + APRIL Ceramic implant combination has been identified as a definitive outlier based on its overall performance over multiple years, both as a combination and for its individual components, across several hospitals. The risk-adjusted hazard ratio significantly exceeds the critical value of two, including its confidence interval. This combination, which remains in active clinical use, surpasses the outlier threshold at both the 2-year and long-term (beyond 5-year) follow-up intervals. While recent data indicate some improvement in early revision performance, it remains uncertain whether these changes will be sufficient to mitigate its outlier status. Recommended course of action: Investigate the specific causes of above-average revision rates. Continue to monitor future performance closely.
	2021	2.33	1.84	2.96	3.50	2.42	5.06	
	2022	2.50	2.01	3.11	3.50	2.51	4.88	
	2023	2.44	1.97	3.02	3.46	2.53	4.74	
	2024	2.49	2.02	3.05	3.34	2.44	4.57	
	2025	2.48	2.03	3.03	3.26	2.39	4.44	
SL-plus MIA + HI (long-term evaluation)	2025	1.76	1.36	2.28	2.39	1.52	3.75	The SL-plus MIA + HI implant combination was first detected as a potential long-term outlier in the 2025 report because the 11-year revision rate exceeded the upper boundary. First analyses of reasons for revisions indicate relatively frequent problems with implant position/orientation as well as acetabular loosening. The SL-plus MIA stem is widely used and has an excellent performance record, whilst the HI cup is more associated with the long-term upward separation of revision rates as shown in the KM curves in this report. Note that the AR2025 reported risk-adjusted hazard ratios refer to all implants with full follow-up, as the implant is now only classified as a potential long-term evaluation outlier.

SIRIS outlier watch list – hip implants

Implant or implant combination	Risk-adjusted hazard ratios for either 2-year revision risk or overall revision risk (long-term evaluation)							
	Detected as outlier in report	Adjusted for age and sex			Adjusted for age, sex, BMI, ASA and Charnley Class (from 2015, if available)			
	HR	lb95%	ub95%	HR	lb95%	ub95%		
Uncemented stem/cup combinations (primary osteoarthritis)								
Former outlier implants (improved)				Summary				
SPS HA + April ceramic	2021	2.61	1.44	4.73	2.85	1.18	6.87	No longer identified as a potential outlier. Consult previous reports for details.
	2022	2.61	1.44	4.72	2.84	1.18	6.85	
Former outlier implants (not in use)				Summary				
Exception + Avantage	2025	1.62	1.21	2.18	2.14	1.27	3.63	Although detected for the first time as a potential outlier in the 2025 annual report, the last recorded use of this combination was in 2022. Revision rates were unproblematic for most of its use time, but high early revision risk of the last exemplars used prompted the outlier status in the 2019 to 2022 observation period.
Polarstem + EP-fit	2020	1.93	1.30	2.86	2.52	1.42	4.45	
	2021	1.89	1.30	2.74	2.31	1.39	3.84	
	2022	1.92	1.36	2.71	2.14	1.38	3.33	
	2023	1.94	1.40	2.68	2.11	1.41	3.16	
	2024	1.87	1.35	2.58	1.99	1.33	2.97	
GTS + G7 bi-spherical	2019							
	2020	5.27	3.22	8.62	3.39	1.52	7.57	
	2021	5.15	3.24	8.19	3.84	1.92	7.71	
	2022	5.15	3.28	8.09	3.96	2.06	7.63	
	2023	5.15	3.28	8.09	3.96	2.05	7.62	
Alloclassic + Fitmore	2022	1.49	1.02	2.17	1.26	0.60	2.64	
	2023	1.55	1.07	2.24	1.24	0.59	2.61	
(AMiStem + Versafitcup DM)	2020	2.14	1.02	4.51	2.30	1.03	5.15	
	2021	2.00	0.95	4.21	2.18	0.98	4.88	
Amistem-H prox coating + Versafitcup DM	2022	3.11	1.29	7.49	3.17	1.31	7.62	These combinations are not currently in active use. Consult previous reports for details.
	2023	3.14	1.30	7.54	3.22	1.34	7.76	
(Harmony + Gyracup)	2020	3.97	1.98	7.94	3.55	1.76	7.13	
Harmony + Symbol DMHA/DS evol.	2022	3.67	1.83	7.35	3.20	1.60	6.42	
	2023	3.66	1.83	7.33	3.20	1.60	6.42	
SPS modular + April ceramic	2019							
	2020	2.95	1.94	4.49	1.61	0.23	11.50	
	2021	2.90	1.91	4.41	1.59	0.22	11.32	
Stelia-stem + Ana.nova hybrid	2019							
	2020	2.65	1.71	4.12	2.30	1.26	4.22	
	2021	2.60	1.68	4.04	2.20	1.20	4.01	
Twinsys + Selexys PC	2020	1.96	0.98	3.93	4.93	1.58	15.34	
Exception + Exceed	2020	1.53	0.69	3.40	1.30	0.33	5.22	
	2021	1.59	0.76	3.33	1.48	0.48	4.61	
(AMiStem-H + Mpact)	2019							
GTS + Exceed	2019							
Corail + Delta motion	2019							

SIRIS outlier watch list – hip implants

Implant or implant combination	Risk-adjusted hazard ratios for either 2-year revision risk or overall revision risk (long-term evaluation)						
	Detected as outlier in report	Adjusted for age and sex			Adjusted for age, sex, BMI, ASA and Charnley Class (from 2015, if available)		
		HR	lb95%	ub95%	HR	lb95%	ub95%

Hybrid fixation stem-cup combinations (primary osteoarthritis)

Active outlier implants

Summary

There are currently no active outliers in this implant category

Former outlier implants (improved)

Implant combination	Year	HR	lb95%	ub95%	HR	lb95%	ub95%	Notes
CCA +	2020	1.83	0.75	4.45	1.91	0.60	6.07	This combination is no longer identified as potential outlier. Please consult previous report for details.
RM Pressfit vitamys	2021	2.05	0.91	4.63	1.86	0.59	5.91	
	2023	1.94	0.97	3.88	1.62	0.61	4.33	
Twinsys (cem) + RM pressfit	2019							

Former outlier implants (not in use)

Implant combination	Year	HR	lb95%	ub95%	HR	lb95%	ub95%	Notes
PF + Fitmore	2020	0.84	1.22	3.17	2.04	1.02	4.09	These combinations are not currently in active use. Consult previous reports for details.
Weber + Alloclassic	2019							
	2020	2.91	1.20	7.05	3.48	1.10	11.02	

Uncemented stem-cup combinations (fractures)

Active outlier implants

Summary

There are currently no active outliers in this implant category

Former outlier implants (improved)

Implant combination	Year	HR	lb95%	ub95%	HR	lb95%	ub95%	Notes
CLS Spotorno + Allofit	2022	2.18	1.20	3.95	2.12	0.94	4.81	This combination is no longer identified as potential outlier. Please consult previous report for details.
	2023	2.09	1.15	3.79	1.97	0.87	4.45	
Fitmore + Allofit	2022	1.37	0.77	2.43	1.87	0.88	3.98	

Cemented stem-head combinations (fractures)

Active outlier implants

Summary

There are currently no active outliers in this implant category

Former outlier implants (not in use)

Implant combination	Year	HR	lb95%	ub95%	HR	lb95%	ub95%	Notes
Harmony (cemented) + OHST bipolar head	2025	3.15	1.63	6.08	2.88	1.43	5.82	Although detected for the first time as a potential outlier in the 2025 annual report, the last recorded use of this combination was in 2023.
Harmony (cemented) + Symbios bibop	2024 2025	1.75	1.14	2.68	1.53	0.79	2.98	Although detected for the first time as a potential outlier in the 2024 annual report, the last recorded use of this combination was in 2020.
Quadra-C + Medacta bipolar head	2024	1.49	0.82	2.70	1.61	0.52	5.02	This combination is not currently in active use. Consult previous reports for details.

SIRIS outlier watch list – knee implants

Implant or implant combination	Risk-adjusted hazard ratios for either 2-year revision risk or overall revision risk (long-term evaluation)							Summary
	Detected as outlier in report	Adjusted for age and sex			Adjusted for age, sex, BMI, ASA and Charnley Class (from 2015, if available)			
	HR	lb95%	ub95%	HR	lb95%	ub95%		
Total knee systems								
Active outlier implants				Summary				
There are currently no active outliers in this implant category								
Former outlier implants (improved)								
Genius	2024	1.93	1.00	3.71	2.10	1.00	4.41	This combination is no longer identified as a potential outlier. Please consult previous report for details. Update 2025: 2-year revision rate has fallen just short of the outlier alert boundary.
Legion	2024	1.60	1.36	1.87	1.35	1.05	1.73	This combination is no longer identified as a potential outlier. Please consult previous report for details. Update 2025: 10-year revision performance just fell short of the critical threshold. Legion is now classified as showing elevated long term revision rates.
Journey II	2019							
	2020	2.17	1.81	2.61	2.10	1.69	2.61	This combination is no longer identified as a potential outlier. Please consult previous report for details.
	2021	2.06	1.74	2.46	2.00	1.63	2.45	
	2022	1.93	1.64	2.29	1.81	1.48	2.20	
Former outlier implants (not in use)								
Sigma PS-RP	2024	1.00	0.77	1.31	2.42	1.54	3.80	These combinations are not currently in active use. Consult previous reports for details.
	2025	1.00	0.77	1.31	2.41	1.53	3.78	
Physica KR	2019							
	2020	3.97	2.13	7.38	3.20	1.20	8.54	
	2021	3.80	2.04	7.07	3.06	1.14	8.17	
Physica PS	2019							
	2020	3.32	1.96	5.61	3.06	1.73	5.41	
	2021	3.11	1.84	5.25	2.91	1.65	5.51	
Physica KR/PS	2022	3.25	2.17	4.85	2.83	1.73	4.63	
E.Motion PS	2019							
Partial knee systems								
Active outlier implants				Summary				
Journey Uni (long-term evaluation)	2020	1.82	1.38	2.39	1.56	0.96	2.53	It is likely that Journey Uni was a problematic knee system at least between 2015 and 2019, but there were signs of improvement in 2020. The statistical precision within the report's main timeframe of interest (2-year revision rate) is relatively low. Whilst the system actually fell below the outlier boundary in 2022 and has so far stayed there, the development of the revision risk beyond two years follow-up strongly suggests an unusual pattern. The system was identified as an outlier in the first round of long-term-evaluation (from 5 years) in 2022 and this was confirmed in 2023 and 2024. It should also be noted that the better short-term revision position since 2022 was also due to the inclusion of poorly performing "other systems" in the evaluation and thus a right-shift of the outlier boundary. Recommended course of action: investigate reasons for revisions and observe future performance. Note that the AR2025 reported risk-adjusted hazard ratios refer to all implants with full follow-up, as the implant is now only classified as a potential long-term evaluation outlier.
	2021	1.81	1.39	2.35	1.68	1.10	2.58	
	2022	1.61	1.25	2.08	1.51	1.02	2.23	
	2023	1.57	1.23	2.00	1.40	0.96	2.03	
	2024	2.00	1.71	2.33	1.79	1.39	2.30	
	2025	1.94	1.67	2.24	1.74	1.37	2.22	

List of manufacturers and distributors

List of companies with implants registered in the SIRIS registry 2024

Company	Headquarters Switzerland	Corporate domicile
Adler Ortho	-	Italy
Amplitude Switzerland	Genf	France
Argomedical AG	Cham	Switzerland
Arthrex Swiss AG	Belp	Germany
Arthrosurface	-	USA
ATF	-	France
B. Braun Medical AG	Sempach	Germany
CeramTec	-	Germany
Conformis	-	Germany
Corin GSA GmbH	Solothurn	United Kingdom
Dedienne Santé	-	France
DePuy Synthes Johnson&Johnson	Zuchwil/Zug	USA
enovis-surgical (includes Mathys and Lima)	Bettlach	Switzerland
Exactech International Operation AG	-	USA
Heraeus Medical Schweiz AG	Zürich	Germany
Implantcast Suisse SA	Basel	Germany
Link Implants AG	Bern	Germany
Medacta International SA	Frauenfeld	Switzerland
OHST Medizintechnik AG	-	Germany
Permedica ORTHOPAEDICS (I)	Scairolo di Collina d'Oro	Italy
Peter Brehm GmbH (Schweiz)	Dietikon	Germany
PLUSOrtho Prothetik GmbH	Oftringen	Switzerland
Smith&Nephew Orthopaedics AG	Baar	United Kingdom
Stemcup Medical Products AG	Zürich	Switzerland
Stryker Osteonics SA	Biberist	USA
Swiss Synergy AG	Baar	Switzerland
Symbios Orthopédie SA	Yverdon-les-Bains	Switzerland
United Orthopedic Corporation Suisse SA	Yverdon-les-Bains	Switzerland
Zimmer Biomet	Winterthur	USA

Definitions

Acetabular component The part of a hip prosthesis that is implanted into the acetabulum – the socket part of a ball and socket joint.

Arthrodesis A procedure in which a natural joint is fused together.

Arthrofibrosis Rigidity of the joint as a consequence of connective tissue adhesion.

Arthrotomy The opening of a joint during surgery.

Articulation The two surfaces that move together (articulate) in a total joint replacement.

ASA score The scoring system of the American Society of Anaesthesiologists (ASA) for grading the overall physical condition of the patient, as follows: I: fit and healthy; II: mild disease, not incapacitating; III: incapacitating systemic disease; IV: life-threatening disease.

Benchmark Comparing the performances at a specific hospital to the mean performances of hospitals throughout Switzerland.

Bilateral Replacing the same joint on both sides of the body (typically both hips or knees) by means of a prosthesis (here meaning the replacement on both sides in one session).

Body Mass Index. . Is obtained by dividing body weight in kilograms by height in meters squared. Interpretation: <18.5: underweight; 18.5–24.9: normal weight; 25–29.9: overweight; 30–34.9: obese class I; 35–39.9: obese class II; >40: obese class III.

Case mix Term used to describe variation in the population, relating to factors such as diagnosis, patient age, gender and health condition.

Cement Material (polymethyl methacrylate) used to fix joint replacements to bone.

Charnley score Clinical classification system – A: one joint affected; B1: both joints affected; B2: contralateral joint with a prosthesis; C: several joints affected or a chronic disease that affects quality of life.

Competing risks survival analysis Method to calculate survival taking into account various outcomes, in this case revision and death.

Cumulative incidence Overall incidences over a specific period of an event (such as the revision of a prosthesis or death of a patient).

Cumulative revision percentage Overall revision percentage over a specific period.

Femoral component Part of a hip or knee prosthesis that is implanted into the femur (thigh bone) of the patient.

Girdlestone Hip revision procedure in which the hip joint or hip prosthesis is removed and no new prosthesis is implanted (usually because of a bacterial infection).

Hybrid fixation Fixation of a prosthesis in which one of the two parts of a prosthesis is cemented and the other one uncemented.

Head component Part of a hip prosthesis that is implanted on top of the femoral component of a hip prosthesis and moves inside the acetabular component of the hip joint.

Hospital service volumes In the tables depicting the total number arthroplasty procedures per year. Four categories of hospital service volume were used (<100, 100–199, 200–299, 300+ procedures per year). The calculation of the annual volume was performed separately for hip and knee surgeries, using the average of all (primary and revision) procedures recorded in each hospital service in 2013–2021.

Acetabular inlay (insert) Intermediate component (inner layer), made usually of polyethylene (but also other materials), which is placed in the acetabular component.

Kaplan-Meier survival analysis Method to calculate survival, in which only one end point is possible, in this case revision.

Kernel density plot A variation of a histogram that uses kernel smoothing to plot values. The underlying kernel is usually Gaussian distribution. One advantage of density plots over histograms is that they are not stepped depending of the number of bins used (histogram bars), but are always smooth lines. The second advantage is that several lines can be plotted over each other and still be visible, which could be difficult with more than two overlaying histograms.

Liner (Knee) named alternatively inlay or insert. Intermediate component of the knee prosthesis. It is made of polyethylene and placed between the femoral and tibial component, fixed somehow to the latter.

Lateral collateral ligament Lateral (outer) knee ligament.

Malalignment Malpositioning of prosthetic components significantly deviating from physiological norms.

Meniscectomy Meniscus removal.

Metallosis Deposition of metal debris in soft tissues of the body, usually around the prosthesis.

Osteoarthritis Disease of the joint in which the cartilage is damaged/destroyed, and the underlying bone altered

Osteochondral bone defect Defect of the joint surface in which both cartilage and the underlying bone are affected

Osteonecrosis Cellular death of bone tissue.

Osteosynthesis Securing broken bone parts together with plates, pins and/or screws.

Osteotomy Cut of the bone with a saw or chisel in order to correct its position, to shorten or lengthen it.

Patellar component Part of a knee prosthesis that is implanted on the inner side of the knee cap.

Patellofemoral prosthesis Two-piece knee prosthesis that provides a prosthetic (knee) articulation surface between the patella and trochlea (furrow) of the thigh bone (femur).

Primary prosthesis The first time replacement of the original joint with a prosthesis.

PROMs Patient Reported Outcome Measures.

Resurfacing hip arthroplasty Hip prosthesis in which the cup (acetabulum) is replaced and a metal cap is implanted on top of the femoral head.

Reverse hybrid fixation hip prosthesis Fixation of a hip or knee prosthesis in which one component is cemented and the other uncemented.

Revision A revision procedure is a secondary surgical procedure of a patient's hip or knee joint whereby the complete primary implant or parts thereof are replaced by new components.

Reoperation All secondary procedures, where no components of the primary implantation are removed.

Revision burden The ratio of revision procedures to all primary and arthroplasty procedures.

Sarcopenia The degenerative loss of skeletal muscle mass and strength associated with aging.

Synovectomy Removal of inflamed mucosa in a joint.

Tibial component Part of a knee prosthesis that is inserted in the tibia (shin bone) of a patient.

Total joint arthroplasty Arthroplasty in which the entire joint of a patient is replaced.

Unicompartmental knee arthroplasty Replacement of half the knee (either inner or outer side) by a prosthesis.

Abbreviations

ASA	American Society of Anaesthesiologists
AVN	Avascular Necrosis
BCR	Bicruciate retaining
BMI	Body Mass Index
CCK	Constrained condylar knee
CI	Confidence Interval
CR	Cruciate retaining
CS	Cruciate sacrificing
CRF	Case Report Form
HA	Hemiarthroplasty of the hip
HR	Hazard ratio
IQR	Interquartile range
KLM	Kaplan Meier estimate
lb/ub	Lower, upper bound (of a confidential ratio)
MCL	Medical Collateral (Inner Knee) Ligament
OA	Osteoarthritis
PCR	Posterior cruciate retaining
PROMs	Patient Reported Outcome Measures
PS	Posterior stabilised
SD	Standard Deviation
SHR	Subhazard ratio
Sig	Significance
TE	Treatment effect
THA	Total Hip Arthroplasty
TKA	Total Knee Arthroplasty
UCOR	Ultra congruent rotating
UKA	Unicompartmental Knee Arthroplasty

Participating hospitals 2024 (144)

Group	Clinic
AG	Kantonsspital Aarau
AG	Kantonsspital Baden
AG	Spital Muri
AG	Spital Zofingen
AG	Asana Gruppe
AG	Spital Leuggern
AG	Asana Gruppe
AG	Spital Menziken
AG	Gesundheitszentrum Fricktal
AG	Spital Rheinfelden
AG	Hirslanden Gruppe
AG	Klinik Aarau
AG	Swiss Medical Network
AG	Privatklinik Villa im Park
AR	Berit Klinik AG
AR	Hirslanden Gruppe
AR	Klinik Am Rosenberg AG
AR	Spitalverbund Appenzell (AR)
AR	Spital Herisau
BE	Klinik Hohmad
BE	Spitalzentrum Biel
BE	Hirslanden Gruppe
BE	Klinik Linde AG
BE	Hirslanden Gruppe
BE	Salem-Spital
BE	Hirslanden Gruppe
BE	Klinik Permanence
BE	Swiss Medical Network SA
BE	Réseau de l'Arc
BE	Hôpital de Saint-Imier
BE	Swiss Medical Network SA
BE	Réseau de l'Arc
BE	Hôpital de Moutier
BE	Insel Gruppe
BE	Spital Aarberg
BE	Insel Gruppe
BE	Inselspital, Unispital Bern
BE	Insel Gruppe
BE	Spital Riggisberg
BE	Lindenhofgruppe
BE	Lindenhofspital
BE	Lindenhofgruppe
BE	Sonnenhofspital
BE	Spital Emmental AG
BE	Spital Burgdorf
BE	Spital Emmental AG
BE	Spital Langnau
BE	Spitäler fmi
BE	Spital Frutigen
BE	Spitäler fmi
BE	Spital Interlaken
BE	Spital Region Oberaargau SRO
BE	Spital Langenthal
BE	Spital STS
BE	Spital Thun
BE	Spital STS
BE	Spital Zweisimmen
BE	Swiss Medical Network
BE	Privatklinik Siloah
BS	Merian Iselin Klinik für Orthopädie und Chirurgie
BS	Universitätsspital Basel
BS	Standort Bethesda Spital AG
BS	Universitätsspital Basel
BS	Standort Uni-Spital
BL	Praxisklinik Rennbahn
BL	Hirslanden Gruppe
BL	Klinik Birshof
BL	Kantonsspital Baselland
BL	Bruderholz
BL	Ergolz Klinik

Group	Clinic
FL	Liechtensteinisches Landesspital
FR	Hôpital fribourgeois HFR
FR	HFR Hôpital cantonal
FR	Swiss Medical Network
FR	Clinique Générale Ste-Anne
GE	Hôpital de La Tour
GE	Hôpitaux universitaires de Genève HUG
GE	Hirslanden Gruppe
GE	Clinique La Colline SA
GE	Hirslanden Gruppe
GE	Clinique des Grangettes SA
GE	Swiss Medical Network
GE	Clinique Générale-Beaulieu
GL	Kantonsspital Glarus
GR	Flury Stiftung Spital Schiers
GR	Gesundheitszentrum Unterengadin
GR	Kantonsspital Graubünden
GR	Regionalspital Surselva AG
GR	Spital Davos
GR	Spital Oberengadin
GR	Spital Thusis
GR	Klinik Gut
GR	Standort Fläsch
GR	Klinik Gut
GR	Standort St. Moritz
JU	Hôpital du Jura
JU	Site de Delémont
LU	Hirslanden Gruppe
LU	Klinik St. Anna AG
LU	Hirslanden Gruppe
LU	St. Anna in Meggen
LU	Luzerner Kantonsspital LUKS
LU	Luzern
LU	Luzerner Kantonsspital LUKS
LU	Sursee
LU	Luzerner Kantonsspital LUKS
LU	Wolhusen
LU	Schweizerisches Paraplegiker-Zentrum
NE	Réseau hospitalier neuchâtelais
NE	La Chaux-de-Fonds
NE	Réseau hospitalier neuchâtelais
NE	Pourtalès
NE	Swiss Medical Network
NE	Clinique Montbrillant
NE	Swiss Medical Network
NE	Hôpital de la Providence
NE	Clinique Volta SA
NW	Spital Nidwalden AG
OW	Kantonsspital Obwalden

Group	Clinic
SG	Spital Linth
SG	Hirslanden Gruppe
SG	Spitalregion Fürstenland Toggenburg
SG	Spitalregion Rheintal Werdenberg Sarganserland
SG	Spitalregion Rheintal Werdenberg Sarganserland
SG	Kantonsspital Graubünden
SG	Kantonsspital St. Gallen
SG	Swiss Medical Network
SH	Spitäler Schaffhausen
SH	Swiss Medical Network
SO	Solothurner Spitäler AG
SO	Solothurner Spitäler AG
SO	Solothurner Spitäler AG
SO	Swiss Medical Network
SZ	Spital Lachen
SZ	Spital Schwyz
SZ	AMEOS
TG	Klinik Seeschau
TG	Spital Thurgau AG
TG	Spital Thurgau AG
TI	Gruppo Ospedaliero Moncucco
TI	Gruppo Ospedaliero Moncucco
TI	Ente Ospedaliero Cantonale
TI	Ente Ospedaliero Cantonale
TI	Ente Ospedaliero Cantonale
TI	Ente Ospedaliero Cantonale
TI	Swiss Medical Network
UR	Kantonsspital Uri
VD	CHUV Centre hospitalier universitaire vaudois
VD	Clinique de la Source
VD	Clinique La Prairie
VD	Clinique CIC Suisse SA
VD	Ensemble Hospitalier de la Côte EHC

Group	Clinic
VD	Etablissements Hospitaliers du Nord Vaudois eHnv
VD	Etablissements Hospitaliers du Nord Vaudois eHnv
VD	Groupement Hospitalier de l'Ouest Lémanique (GHOL)
VD	Hirslanden Gruppe
VD	Hôpital intercantonal de la Broye HIB
VD	Hôpital Riviera-Chablais HRC
VD	Réseau Santé Balcon du Jura RSBJ
VD	Swiss Medical Network
VD	Swiss Medical Network
VS	Clinique CIC Valais
VS	Hôpital du Valais - Spital Wallis
VS	Hôpital du Valais - Spital Wallis
VS	Hôpital du Valais - Spital Wallis
VS	Hôpital du Valais - Spital Wallis
VS	Swiss Medical Network
ZG	Zuger Kantonsspital
ZG	Hirslanden Gruppe
ZH	Kantonsspital Winterthur
ZH	Swiss Medical Network
ZH	Schulthess Klinik
ZH	Spital Bülach
ZH	Spital Limmattal
ZH	Spital Männedorf
ZH	Spital Uster
ZH	Spital Zollikerberg
ZH	Universitätsspital Zürich
ZH	Universitätsklinik Balgrist
ZH	GZO
ZH	Hirslanden Gruppe
ZH	Hirslanden Gruppe
ZH	See-Spital
ZH	Stadtspital Zürich
ZH	Stadtspital Zürich
ZH	Swiss Medical Network
ZH	Swiss Medical Network

List of tables and figures

1. Introduction

Table 1.1	Variables collected by the SIRIS registry	12
-----------	---	----

2. Methods

Table 2.1	Retrospective coverage analysis 2017–2022 based on National Office of Public Health figures (BAG)	16
Figure 2.1	Coverage rates for primary hip prostheses in the SIRIS registry 2023	17
Figure 2.2	Coverage rates for primary knee prostheses in the SIRIS registry 2023	17

3. Demography and Epidemiology

Table 3.1	Total and partial hip arthroplasty (THA and HA), primary and revisions/reoperations	22
Figure 3.1	Incidence of primary total hip arthroplasties registered in SIRIS	23
Figure 3.2	Growth rates of primary hip arthroplasties by canton: 2019 – 2024	23
Figure 3.3	Kaplan Meier estimate of cumulative postoperative revision risk after primary hip arthroplasty	24
Figure 3.4	Kaplan Meier estimate of cumulative postoperative revision risk after THA by time period	25
Table 3.2	Primary total hip arthroplasty: Baseline patient characteristics by year	26
Figure 3.5	Age distribution at surgery of primary total hip arthroplasty and hemiarthroplasty	27
Figure 3.6	Primary total hip arthroplasty: Mean age at primary arthroplasty depending on BMI class	27
Table 3.3	Primary total hip arthroplasty: Baseline patient characteristics by main diagnostic group	28
Figure 3.8	Relative share of total hip arthroplasty procedures using 36 mm heads by canton (2019 – 2024)	30
Figure 3.9	Relative share of total hip arthroplasty procedures using dual mobility cups by canton (2019 – 2024)	30
Table 3.4	Number of participating hospital services and median procedures per unit per year	32
Table 3.5	Number of hospital services and number of primary total hip arthroplasties according to hospital volume	32
Figure 3.10	What share of selected procedures is performed in hospital services with different service volumes?	33
Figure 3.11	Cases per hospital service 2024: Total hip arthroplasty and hemiarthroplasty	33
Table 3.6	Primary total hip arthroplasty: Baseline patient characteristics of primary total hip arthroplasty by hospital service volume	34
Figure 3.12	2-year revision rate of primary total hip arthroplasty by service	35
Table 3.7	Fracture of the hip: Baseline patient characteristics by year	36
Table 3.8	Fracture of the hip: Baseline patient characteristics by type of treatment	37
Table 3.9	Fracture of the hip: Baseline patient characteristics by hospital service volume	38
Figure 3.13	2-year revision rate of primary hemiarthroplasty by service	39
Table 3.10	Total and partial knee arthroplasty (TKA, PKA), primary and revisions/reoperations	40
Figure 3.14	Incidence of all primary knee arthroplasties registered in SIRIS	41
Figure 3.15	Growth rates of primary knee arthroplasties by Canton: 2019 – 2024	41
Table 3.11	Primary total knee arthroplasty: Baseline patient characteristics by year	42
Figure 3.16	Age distribution at surgery of primary total and partial knee arthroplasty	43
Figure 3.17	Primary total knee arthroplasty: Mean age at primary arthroplasty depending on BMI class	43
Table 3.12	Primary total knee arthroplasty: Baseline patient characteristics by main diagnostic group	44
Table 3.13	Baseline patient characteristics of primary partial knee arthroplasty by hospital service volume	45
Table 3.14	Primary partial knee arthroplasty: Baseline patient characteristics by year	47
Figure 3.18	Risk of revision within 5 years and 10 years after primary TKA in percent	48
Figure 3.19	Kaplan Meier estimate of cumulative postoperative revision risk after TKA by time period	49

Figure 3.20	Risk of revision within 5 years after primary PKA in percent	50
Figure 3.21	Kaplan Meier estimate of cumulative postoperative revision risk after primary knee arthroplasty	50
Figure 3.22	2-year revision rate of primary total knee arthroplasty by service	51
Figure 3.23	2-year revision rate of primary total knee arthroplasty by service, without isolated secondary patella resurfacing	52
Figure 3.24	2-year revision rate of partial knee arthroplasty by service	52
Figure 3.25	Relative share of TKA procedures using PS or CS/CR systems by Canton: 2019 – 2024	53
Figure 3.26	Primary total knee arthroplasty: Type of bearing	54
Figure 3.27	Relative share of TKA procedures using mobile bearing by Canton: 2019 – 2024	54
Figure 3.28	Primary total knee arthroplasty: Patellar component	55
Figure 3.29	Relative share of TKA procedures using primary patellar replacement by Canton: 2019 – 2024	55
Table 3.15	Number of participating hospital services and median procedures per unit per year	56
Table 3.16	Number of hospital services and number of primary total knee arthroplasties according to hospital volume	56
Figure 3.30	What share of selected procedures is performed in hospital services with different service volumes?	57
Figure 3.31	Cases per hospital service 2024: Total and partial knee arthroplasty	57
Table 3.17	Primary total knee arthroplasty: Baseline patient characteristics by hospital service volume	58
Figure 3.32	Distributions of different diagnoses in 36 hospitals >200 cases (2024)	59

4. Hip arthroplasty

Table 4.1	Primary total hip arthroplasty: Surgery characteristics by main diagnostic group	63 and 64
Figure 4.1	Share of dual-mobility cups over time by main pathology	64
Table 4.2	Primary total hip arthroplasty: Component fixation methods by diagnostic group by year	65
Table 4.3	Surgical approach in total hip arthroplasty for primary osteoarthritis by year	66
Figure 4.2	Relative share of total hip arthroplasty procedures using different surgical approaches by Canton (2019 – 2024)	66
Table 4.4	Primary total hip arthroplasty: bearing surface in primary osteoarthritis by year (in %)	67
Figure 4.3	Relative shares of bearing surfaces in total hip arthroplasty over time	67
Table 4.5	Primary total hip arthroplasty: bearing surface in primary osteoarthritis by age	68
Figure 4.4	Primary total hip arthroplasty: bearing surface in primary osteoarthritis by age	68
Table 4.6	Primary total hip arthroplasty: fixation methods in primary osteoarthritis by age	69
Table 4.7	Primary total hip arthroplasty: fixation methods in primary osteoarthritis by sex	69
Figure 4.5	Primary total hip arthroplasty: technologies used in primary OA	70
Table 4.9	First revision of primary total hip arthroplasty within 24 months overall and according to baseline characteristics	71
Table 4.10	Reason for early first revision of primary total hip arthroplasty (primary OA)	72
Table 4.11	Type of early first revision of total hip arthroplasty (primary OA)	72
Table 4.12	Hip early first revision of primary OA THA: main components used by age at revision (2019 – 2024)	74
Table 4.13	Hip early first revision of primary OA THA: main brands used (2019 – 2024)	73
Table 4.14	First revision of primary total hip arthroplasty within 24 months according to stem fixation, articulation and approach	75
Figures 4.6	Reason for early first revision by time interval since primary total hip arthroplasty	76
Figures 4.7	Cumulative incidence rates for different revision diagnoses (primary OA THA)	77 and 78
Figure 4.8	Estimated failure rates of primary total hip arthroplasty for different bearing surfaces	79
Figure 4.9	Estimated failure rates of primary total hip arthroplasty for different types of head sizes (standard cups: all uncemented fixation)	80
Table 4.15	Estimated failure rates of primary total hip arthroplasty for different types of head sizes by bearing surface (standard cups: all uncemented fixation)	81
Figure 4.10	Estimated failure rates of primary total hip arthroplasty for different fixation methods	82
Figure 4.11	Estimated failure rates of primary total hip arthroplasty for different fixation methods	82

Figure 4.12	Estimated failure rates of primary total hip arthroplasty for different BMI	83
Figure 4.13	Estimated failure rates of primary total hip arthroplasty for different BMI	84
Figure 4.14	Estimated failure rates of primary total hip arthroplasty for different types of cups (all uncemented fixation)	85
Figure 4.15	Estimated failure rates of primary total hip arthroplasty for different types of cups (hybrid fixation)	85
Figure 4.16	Estimated failure rates of primary total hip arthroplasty for different types of dual mobility cups (all uncemented fixation)	86
Figure 4.17	Estimated failure rates of primary total hip arthroplasty for different types of short stems (all uncemented fixation)	87
Figure 4.18	Estimated failure rates after revision of total hip arthroplasty: primary and secondary OA	88
Figure 4.19	Risk of revision within 5 years and 10 years after primary THA (primary OA or sec OA) in percent	88
Figure 4.20	Cumulative incidence rates for different re-revision diagnoses after main component changes	89
Figure 4.21	Estimated failure rates after revision of total hip arthroplasty (primary and secondary OA): types of procedures	90
Figure 4.22	Cumulative incidence rates for different re-revision diagnoses after head/inlay changes	91
Figure 4.23	Estimated failure rates after revision of total hip arthroplasty (primary and secondary OA): types of stems used in revision	92
Figure 4.24	Estimated failure rates after revision of total hip arthroplasty (primary and secondary OA): types of cups used in revision	93
Table 4.16	Top 75% of primary total hip arthroplasty uncemented combinations (primary OA) 2019 – 2024	95
Table 4.17	Long term evaluation: Failure rates of primary total hip arthroplasty uncemented combinations (primary OA)	96
Figure 4.25	Implant combinations with elevated long-term revision rates (primary OA, uncemented THA)	97
Figure 4.26	Implant combinations with long-term evaluation outlier status (primary OA, uncemented THA)	97
Figure 4.27	Implant combinations with below-average long-term revision rates (primary OA, uncemented THA)	98
Figure 4.28	All remaining implant combinations with average revision risks (primary OA, uncemented THA)	99 and 100
Table 4.18	Long term evaluation: Failure rates of primary total hip arthroplasty uncemented combinations and different bearing surfaces (primary OA)	101 and 102
Table 4.19	Top 75% of primary total hip arthroplasty hybrid combinations (primary OA) 2019 – 2024	103
Table 4.20	Long term evaluation: Failure rates of primary total hip arthroplasty hybrid combinations (primary OA)	104
Figure 4.29	Implant combinations with below-average long-term revision rates (primary OA, hybrid THA)	105
Figures 4.30	All remaining implant combinations with average revision risks (primary OA, hybrid fixation THA)	105
Table 4.21	Top 75% of primary total hip arthroplasty uncemented combinations (secondary OA) 2019– 2024	106
Table 4.22	Long term evaluation: Failure rates of primary total hip arthroplasty uncemented combinations (secondary OA)	107
Figure 4.31	Implant combinations with elevated long-term revision rates (secondary OA, uncemented THA)	107
Figure 4.32	Implant combinations with below-average long-term revision rates (secondary OA, uncemented THA)	108
Figures 4.33	All remaining implant combinations with average revision risks (secondary OA, uncemented THA)	108
Table 4.23	2-year evaluation: Revision rates of uncemented primary total hip arthroplasty combinations within 24 months (primary OA)	110, 111, 112
Figure 4.34	2-year evaluation: Revision rates of uncemented primary total hip arthroplasty combinations within 24 months (primary OA)	113
Figure 4.35	2-year evaluation: Revision rates of hybrid primary total hip arthroplasty combinations within 24 months (primary OA)	114
Figure 4.36	2-year evaluation: Revision rates of uncemented primary total hip arthroplasty combinations within 24 months (secondary OA)	115
Table 4.24	Fractures of the hip: Surgery characteristics by treatment group	117
Tables 4.25 and Figures 4.37	Fracture of the hip: Component fixation methods by type of treatment by year	118
Tables 4.26 and Figures 4.38	Fracture of the hip: Surgical approach by type of treatment by year	119
Table 4.27	Fracture of the hip: First revisions within 24 months overall and according to baseline characteristics	120
Table 4.28	Fracture of the hip: First revisions according to stem fixation and approach	121
Table 4.29	Fracture of the hip: Reasons for early first revisions	121
Table 4.30	Fracture of the hip: Reasons for early first revisions (unipolar vs. bipolar hemi heads; cemented stems only)	122

Figure 4.39	Fracture of the hip: Estimated failure rates of hemiarthroplasty of the hip: unipolar heads versus bipolar heads	122
Figures 4.40	Fracture of the hip: cumulative incidence rates for different revision diagnoses, THA and HA	123
Table 4.31	Fracture of the hip: Type of revisions by primary treatment modality, THA versus HA	124
Table 4.32	Top 75% of primary total hip arthroplasty uncemented combinations to treat fractures 2019 – 2024	125
Figure 4.41	2-year evaluation: Revision rates within 24 months of uncemented primary total hip arthroplasty comb. to treat fractures	126
Table 4.33	Top 75% of primary total hip arthroplasty hybrid combinations to treat fractures 2019 – 2024	127
Figure 4.42	2-year evaluation: Revision rates within 24 months of hybrid primary total hip arthroplasty combinations to treat fractures	128
Table 4.34	Fracture of the hip: top 75% stem/head combinations used in hemiarthroplasty (HA) 2019 – 2024	129
Figure 4.43	2-year evaluation: Revision rates within 24 months of hybrid primary total hip arthroplasty combinations to treat fractures	130 and 131

5. Knee arthroplasty

Table 5.1	Primary total knee arthroplasty: Surgery characteristics	135
Figure 5.1	Share of TKA patients who had knee arthroscopy prior to arthroplasty	136
Figure 5.2	Primary total knee arthroplasty: Component fixation	137
Figure 5.3	Use of stems by BMI and type of knee system	138
Figure 5.4	Use of stems as a percentage of primary TKAs with cemented tibial components	138
Figure 5.5	Primary total knee arthroplasty: technologies used	139
Table 5.3	First revision of primary total knee arthroplasty within 24 months overall and according to baseline characteristics	140
Table 5.4	Reason for revision of primary total knee arthroplasty	141
Figure 5.6	Cumulative incidence rates for different revision diagnosis of primary total knee arthroplasty	142
Figure 5.7	Time interval between primary total knee arthroplasty and first revision by reason	143
Table 5.5	Median time interval between primary total knee arthroplasty and early first revision (in months) according to reason	143
Table 5.6	First revision of primary total knee arthroplasty within 24 months overall and according to component fixation	144
Figure 5.8	Estimated failure rates of primary total knee arthroplasty for different fixation methods	144
Figure 5.9	Estimated failure rates of primary total knee arthroplasty for different implant types	145
Figure 5.10	Estimated failure rates of primary total knee arthroplasty: use of stems (CR/CS/PS/MP)	146
Figure 5.11	Estimated failure rates of primary total knee arthroplasty: use of stems (Hinged, SC/CCK)	147
Figure 5.12	Estimated failure rates of primary total knee arthroplasty: status of patella after primary operation	148
Table 5.7	Early first revision of primary total knee arthroplasty: Surgery characteristics	149
Figure 5.14	Early first revision of primary total knee arthroplasty: Component fixation	150
Figure 5.15	Selected additional components used in early first revisions	151
Figure 5.16	Cumulative incidence rates for different re-revision diagnosis of primary total knee arthroplasty	152
Figure 5.17	Estimated failure rates after revision of total knee arthroplasty: types of revisions	153
Figure 5.18	Estimated failure rates after revision of total knee arthroplasty: secondary patella replacement	154
Figure 5.19	Cumulative incidence rates for different re-revision diagnosis after secondary patella replacements (TKA)	155
Figure 5.20	Estimated failure rates after conversion from partial knee to total knee arthroplasty	156
Figure 5.21	Cumulative incidence rates for different re-revision diagnoses of conversions to primary total knee arthroplasty	156
Table 5.8	Top 75% of primary total knee arthroplasty systems (all diagnoses, all component fixations) 2019 – 2024	157
Table 5.9	Long term evaluation: Failure rates of primary total knee arthroplasty systems (all diagnoses, all component fixations)	158 and 159
Figure 5.22	Implant combinations with below-average long-term revision rates (all TKA)	160
Figure 5.23	Implant combinations with elevated long-term revision rates (all TKA)	160
Figures 5.24	All remaining implant combinations with average revision risks (all TKA)	161 and 162
Figures 5.25	2-year evaluation: Revision rates of primary total knee arthroplasty systems within 24 months	163, 164, 165
Figures 5.26	Share of partial knee patients who had knee arthroscopy prior to arthroplasty	167

Table 5.10	Primary partial knee arthroplasty: Surgery characteristics	167
Figure 5.27	Primary partial knee arthroplasty: Component fixation	168
Figure 5.28	Partial knee arthroplasty: technological assistance over time	169
Table 5.11	First revision of primary partial knee arthroplasty within 24 months overall and according to baseline characteristics	170
Table 5.12	Reason for early first revision of primary partial knee arthroplasty	170
Figure 5.29	Cumulative incidence rates for different revision diagnosis of partial knee arthroplasty	171
Figure 5.30	Time interval between primary partial knee arthroplasty and first revision by reason	172
Figure 5.31	Estimated failure rates of primary partial knee arthroplasty for main types of component fixation	172
Figure 5.32	Estimated failure rates of primary partial knee arthroplasty: type of arthroplasty	173
Figure 5.33	Estimated failure rates of primary partial knee arthroplasty: conventional vs. patient specific instrumentation	174
Figure 5.34	Estimated failure rates of primary partial knee arthroplasty: technological assistance	174
Table 5.13	Type of early first revision of primary partial knee arthroplasty	175
Figure 5.35	Cumulative incidence rates for different types of revisions of partial knee arthroplasty	175
Table 5.14	Top 10 (94%) of primary partial knee arthroplasty systems (all diagnoses, all component fixations) 2019 – 2024	176
Figure 5.36	2-year evaluation: Revision rates of primary partial knee arthroplasty systems within 24 months	177
Table 5.15	Long term evaluation: Failure rates of primary partial knee arthroplasty systems (all diagnoses, all component fixations)	178
Figure 5.37	Implant combinations with long-term evaluation outlier status (all unicompartmental PKA)	179
Figure 5.38	Implant combinations with below-average long-term revision rates (unicompartmental PKA)	179
Figures 5.39	All remaining implant combinations with average revision risks (all PKA)	180
Table 5.16	Top 5 (98%) of primary patellofemoral joint systems (all diagnoses, all component fixations) 2019 – 2024	181
Figure 5.40	2-year evaluation: Revision rates of primary patellofemoral joint systems within 24 months	181
Table 5.17	Long term evaluation: Failure rates of primary patellofemoral joint systems (all diagnoses, all component fixations)	182

6. Patient reported outcome measures (PROMs)

Figure 6.1	Preoperative pain	185
Figure 6.2	Postoperative pain	185
Figure 6.3	Preoperative quality of life (EQ5D): mobility	187
Figure 6.4	Postoperative quality of life (EQ5D): mobility	187
Figure 6.5	Quality of life (EQ-5D-5L): overview	188
Figure 6.6	Preoperative satisfaction with current situation	189
Figure 6.7	Postoperative satisfaction with current situation	189
Figure 6.8	Treatment effect pain	190
Figure 6.9	Treatment effect quality of life (EQ5D)	190

Foundation for quality assurance
in implant surgery – SIRIS

c/o Core Treuhand AG, 3007 Bern
info@siris-implant.ch, www.siris-implant.ch