

# Safety and performance of high-risk medical devices: the role of real-world data

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Chapter 2 – Quality and utility of European cardiovascular and orthopaedic registries for the regulatory evaluation of medical device safety and performance across the implant lifecycle: a systematic review



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#### Abstract

Background: The European Union Medical Device Regulation (MDR) requires manufacturers to undertake post-market clinical follow-up (PMCF) to assess the safety and performance of their devices following approval and Conformité Européenne (CE) marking. The quality and reliability of device registries for this Regulation have not been reported. As part of the Coordinating Research and Evidence for Medical Devices (CORE-MD) project, we identified and reviewed European cardiovascular and orthopedic registries to assess their structures, methods, and suitability as data sources for regulatory purposes.

Methods: Regional, national and multi-country European cardiovascular (coronary stents and valve repair/replacement) and orthopedic (hip/knee prostheses) registries were identified using a systematic literature search. Annual reports, peer-reviewed publications, and websites were reviewed to extract publicly available information for 33 items related to structure and methodology in six domains and also for reported outcomes.

Results: Of the 20 cardiovascular and 26 orthopedic registries fulfilling eligibility criteria, a median of 33% (IQR: 14%-71%) items for cardiovascular and 60% (IQR: 28%-100%) items for orthopedic registries were reported, with large variation across domains. For instance, no cardiovascular and 16 (62%) orthopedic registries reported patient/procedure-level completeness. No cardiovascular and 5 (19%) orthopedic registries reported outlier performances of devices, but each with a different outlier definition. There was large heterogeneity in reporting on items, outcomes, definitions of outcomes, and follow-up durations.

Conclusion: European cardiovascular and orthopedic device registries could improve their potential as data sources for regulatory purposes by reaching consensus on standardised reporting of structural and methodological characteristics to judge the quality of the evidence as well as outcomes.

## Background

A vital mechanism for assuring safety and performance of high-risk medical devices in patients is that they are subject to systematic post-market surveillance, which includes the collection of high-quality clinical data by registries. For regulatory purposes, such post-market clinical follow-up (PMCF) is mandatory for cardiovascular devices like stents and valves and for orthopedic devices like hip and knee implants.

The International Medical Device Regulators Forum (IMDRF) defines a medical device registry as "an organized system with a primary aim to increase the knowledge on medical devices contributing to improve the quality of patient care that continuously collects relevant data, evaluates meaningful outcomes and comprehensively covers the population defined by exposure to particular device(s) at a reasonably generalizable scale (e.g. international, national, regional, and health system)." A medical device registry is thus an unselected population-based health information system collecting large numbers of real-world data regarding safety and performance of specific devices over time, with the aim to improve the quality of patient care,<sup>1-4</sup> and therefore well suited to provide clinical evidence on PMCF of devices for regulatory purposes.

The European Medical Device Regulation (MDR) requires manufacturers to plan and conduct surveillance of their devices (Article 83 of (EU) 2017/7455), but the list of sources of available information that can be used for this purpose includes "relevant specialist or technical literature, databases and/or registers" and "information, including feedbacks and complaints, provided by users, distributors and importers" (Annex III, clause 1.1(a)).<sup>5</sup> Real-world data collected by medical device registries are particularly useful as they enable continuous benchmarking across longer follow-up in many more patients than enrolled in clinical trials.<sup>6-10</sup>

The utility of medical device registries organised by medical professional associations is exemplified by the case of the "Metal on Metal" (MoM) hip implants. Originally developed as a more durable alternative to implants with ceramic or polyethylene components, mid-term follow-up registry data of patients with MoM showed far higher revision rates when compared with other implants. The Australian Orthopaedic Association National Joint Replacement Registry identified these implants as having an outlier performance, three years before their withdrawal from the market in 2010. Performance, three years before their withdrawal from the market in 2010. Performance or cardiovascular diseases, device registries have provided important insights on the safety of coronary stents, by documenting increased rates of low-frequency events such as stent thrombosis with specific stent platforms. Performance or sentence of the platforms.

Principles have been proposed by regulators to evaluate whether the quality of clinical data on medical devices meets the scientific standards to be used for PMCF. They include coverage (i.e. extent of participation in data collection), completeness (i.e. data used in analyses are consistently captured), accuracy (i.e. data recorded is an accurate reflection of the healthcare event), consistency (i.e. uniformity in following the same procedures for data capture), integrity (i.e. consistent recording of unique identification of medical devices), and reliability (i.e. reproducibility of data elements).¹ Specific criteria have not been proposed, however, and it is therefore unknown if existing medical device registries in Europe would allow manufacturers to meet the MDR requirements to an acceptable standard. As part of the Coordinating Research and Evidence for Medical Devices (CORE-MD) project, this systematic review therefore aims to: i) identify current European cardiovascular and orthopedic medical device registries, and ii) review these registries by 33 items that related to their structures, methodologies, and quality of data.

#### Methods

This systematic review was conducted according to the Preferred Reporting Items for Systematic Reviews and MetaAnalyses (PRISMA) 2020 guidelines,<sup>17</sup> and it was registered in the Center for Open Science in October 2021 (https://osf.io/7yuwx/) prior to data collection.

## Search strategy

A previous study identified European registries on implantable medical devices<sup>18</sup> from which we adapted and updated its search strategy in order to identify new registries and expand the list of registries for this systematic review. Eight literature libraries (Centre for Reviews and Dissemination York, Cochrane library, Embase, Emcare, Google Scholar, Medline, PubMed, and Web of Science) were searched for publications between January 1, 2013, and July 7, 2021, using a systematic search strategy (Supplementary File 1 – available online) created by a librarian (JWS). References were imported to EndNote (Version X9, Clarivate Analytics, Philadelphia, the USA) which was used to remove duplicate publications, and subsequently exported to the web application Rayyan (Doha, Qatar)<sup>19</sup> which was used for study selection.

#### Study selection

Two reviewers (LAH and THG) independently screened titles and abstracts and then independently assessed eligibility of full texts. Discrepancies were resolved by discussion. If consensus could not be reached, the senior researcher (PJMvdM) was consulted for a decisive vote. Studies were included firstly if they described a European regional, national, or multi-country cardiovascular medical device registry in which data were captured on coronary stents and/or on percutaneous or surgical valve repair or replacement. We focused on coronary artery stents as they are commonly used high-risk devices with a low frequency of adverse events so that a large number of patients is needed to detect safety issues, and on valve prostheses because there are many new devices for which guidance is needed on benchmarking

safety and performance. Secondly, we also included European registries capturing data on hip and/or knee prostheses since they are the most common orthopedic high-risk devices. By applying these criteria and by excluding multicentre studies, we complied with the IMDRF definition of a registry, which is particularly relevant to evaluate implant performance in the entire population receiving such a device in daily practice, rather than in selected (high performing) centres, Additional inclusion criteria were: i) an active/accessible website at the time of study collection; or ii) at least one publication and/or annual report containing registries' data between 2013 and 2021. We defined an "active registry" as a registry that published at least one annual report and/or peer-reviewed paper containing registries' data, during or later than 2018. The reason for making a distinction between "active" and "non-active" registries is to give a better estimate regarding the number of registries able to contribute evidence for regulatory purposes in practice. In addition, "active" registries may also report the structural and methodological characteristics determining the quality of the data more consistently. No language restriction was applied. Data were extracted from any peer-reviewed publication(s) that described the registries' structure and methodology, and combined with data from the most recent published annual report(s) (if available) and/or registries' website (if available). To identify any more registries that were not vet included in this review, the references in publications and annual reports were checked, and clinical experts were consulted (five for the cardiovascular and eight for the orthopedic field). For orthopedic registries, we also checked the list on the EFORT —Network of Orthopedic Registries of Europe (NORE) — website (https://efortnet.efort.org/nore-map/#/nore/map-all).

#### Data extraction and analysis

the literature including a study reporting recommendations,20 LAH and PIMvdM developed a list of items that could be used to assess registries' structures and methodological characteristics, reflecting the previously mentioned principles<sup>1</sup> and therefore relevant to judge the quality of registry data for regulatory purposes as required by the MDR. These were sent to 13 experts in the cardiovascular (n=7) and/or orthopedic (n=6) fields, for feedback and suggestions of relevant additional items. Consensus was reached on a total of 33 quality items covering six domains: i) Identification (six items) to understand which population the registry intends to describe; ii) Maturity (three items) to contextualise the numbers of procedures and extent to which longer-term outcomes may already be captured; iii) Governance (five items) to enable assessment of the integrity of data; iv) Coverage, design & organisation (eight items) to reflect the aforementioned principles of coverage and consistency; v) Data quality & completeness (four items) to reflect the aforementioned principles of completeness and accuracy, and vi) Safety & performance (seven items) to capture reliability of data in using standard definitions to assess safety; details of each item are given in Box 1. Data were also collected on: i) the number of peer-reviewed publications since foundation of the registry, as an indicator of scientific utility; ii) the number of included manufacturers and the total number of patients/ procedures, to indicate the average experience with a specific device, that would potentially be relevant when assessing the performance based on a minimum sample size to obtain reliable

estimates, and iii) reported outcomes, including definitions and durations of followup.

Using a prespecified format, publicly available data were extracted independently by LAH and THG for each registry and each item. Otherwise, items were recorded as "Not reported" (N/R). Median values (given the skewed distributions) and interquartile ranges (IQRs) were calculated for the percentage of items reported per domain and across all domains, for both cardiovascular and orthopedic registries. Analyses were performed using Microsoft Excel (Excel version 2012, Microsoft, Redmond, the USA).

## Results

#### Literature search

The searches identified 4,538 cardiovascular and 4,485 orthopedic publications, of which 1,727 cardiovascular and 1,360 orthopedic publications remained after removing duplicates. Title and abstract screening identified a total of 81 cardiovascular and 27 orthopedic registries, mentioned in publications from January 2013 to July 2021 (Figure 1). Twelve cardiovascular registries were excluded because they focused on other cardiovascular devices (e.g. pacemakers) (n=11) or no devices (n=1) and a further 51 cardiovascular and 7 orthopedic registries were excluded during full-text screening, mostly because of reporting on a single or multicentre study, or due to registry mergers (Figure 1). Manual search identified two additional cardiovascular<sup>21,25</sup> and six orthopedic registries,<sup>47,51,53,57,60,66</sup> that did not publish any peer-reviewed papers and therefore were not found in the literature search. Thus, a total of 20 cardiovascular<sup>21,40</sup> and 26 orthopedic registries<sup>41-66</sup> were selected for data extraction.

# Overall findings

Across all domains, a median of 33% (IQR 14%-71%) of the predefined 33 quality items were reported by cardiovascular registries and 60% (IQR 28%-100%) by orthopedic registries. The highest median value was reached for the domain 'Identification' since almost all registries reported information on e.g. the type of registry: 75% (IQR 69%-100%) for cardiovascular and 100% (IQR 100%-100%) for orthopedic registries (Figure 2). The lowest percentages were observed for the domains 'Data quality & completeness' and 'Safety & performance'; for cardiovascular registries these were respectively 25% (IQR 0%-25%) and 0% (IQR 0%-4%) and for orthopedic registries they were 38% (IQR 0%-69%) and 50% (IQR 0%-71%) (Figure 2).

# Domains "Identification" and "Maturity"

The majority of included registries (41 out of 46; 89%) were national registries,<sup>21-26,28-48,51,53,54,56-66</sup> with only three (7%) regional registries<sup>27,52,55</sup> and two (4%) multicountry registries<sup>49,50</sup> (Table S1A and S1B, Supplementary Files 2 and 3 – online

available). The first cardiovascular registry was founded in 1978<sup>23</sup> and the two most recent in 2013,<sup>35,37</sup> while the first orthopedic registry was established in 1975<sup>65</sup> and the most recent in 2019.<sup>53</sup> Initial motivations to set up a registry were mostly reported (by 60% of cardiovascular<sup>21,23,25-27,29,33,35-37,39,40</sup> and 92% of orthopedic registries<sup>42-44,46-66</sup>) and often involved ensuring patients' safety. More orthopedic than cardiovascular registries publish annual reports (77% *versus* 30%), although for some registries (35%) data were last reported more than four years ago and therefore labelled as "non-active" (Table 1). Of the active registries (65%), a median of 43% (IQR 25%-80%) of the 33 quality items were reported by cardiovascular registries and 75% (IQR 41%-100%) by orthopedic registries (Figure 3).

Domains "Governance" and "Coverage, Design & Organisation"

Mandatory enrolment of eligible patients was implemented in 8 (40%) cardiovascular<sup>22,24,27,29,30,37,39,40</sup> and 12 (46%)orthopedic registries42,43,46,48,50,51,55,56,59,60,62,64 (Table S2A and S2B - online available). Few cardiovascular<sup>21,24,27,29,35-37,39,40</sup> and orthopedic<sup>42-44,46,53,54,61-63,65</sup> registries have reported on their funding and few report on the patient informed consent process<sup>24,25,27,29,31,33-37,39,40,42,44,46,48,50, 54,60,63,64</sup> (Table S3A and S3B – online available). The number of participating hospitals per registry varied largely, with a median of 28 (IQR 17-89) hospitals for cardiovascular registries and 71 (IQR 42-116) hospitals for orthopedic registries (Table S4A and S4B - online available). The proportion of all eligible hospitals that participated in the registry (i.e. hospital-level coverage) was only reported by six (30%) cardiovascular registries, <sup>24,26-28,31,34</sup> with a median hospital-level coverage of 100% (IQR 98%-100%) and by nine (35%) orthopedic registries, 44-46,48,52,54,60,64,65 also with a median hospital-level coverage of 100% (IOR 95%-100%) (Table S4A and S4B - online available).

In general, cardiovascular registries report on studies for which selected patient groups are included, so data on the total number of patients receiving an implant were reported by only four (20%) registries.<sup>21,25,29,34</sup> The median for stents was 12,395 (IOR 3,985-201,647) and the median for valves was 2,325 (IOR 861-10,479) (Table S4A and S4B - online available). Given the regular publication of annual reports, the total and annual volume of implant procedures in orthopedic registries was mostly reported; details were on both items was not available for seven (27%) registries. 41,45,47,49,53,54,61 Overall, orthopedic registries reported on a median of 120,408 (IQR 52,391-218,445) hip implants and a median of 102,649 (IQR 51,700-194,076) knee implants (Table S4A and S4B – online available). Data linkage with other sources—mostly national clinical databases—was reported by 8 (40%) cardiovascular<sup>21,24,27,29,34,36,37,39</sup> orthopedic registries.42,44and 14 (54%)46,48,50,52,54,55,60,62-65

Information was mostly provided on hospital and/or device-level, while in some cases also surgeon-level information was provided. There were more different types of implants in orthopedic than in cardiovascular registries, shown by totals of 37 different manufacturers for knee implants and 63 for hip implants compared with 13 different manufacturers of valves and 11 of stents (Table S5A and S5B – online available).

## Domain "Data quality & completeness"

None of the cardiovascular registries reported patient/procedure-level data completeness (Table S6A and S6B – online available). Techniques to handle missing data were described in only one cardiovascular registry (5%), $^{21}$  which applied a data completeness threshold (i.e. a certain variable will only be analysed if its completeness is  $\geq 95\%$ ). Most (55%) cardiovascular registries $^{21,23,26,27,29,30,34-37,40}$  reported on procedures to check the quality of their data, such as checking on the range and consistency of entries, and verification by audits or an external electronic tool.

Patient/procedure-level completeness was reported by 16 (62%) orthopedic registries, 42-46,48,50,52-55,60,62-65 which varied from 19% for hip prostheses in the Irish National Orthopaedic Register to 98%-99% for knee prostheses in the Danish Knee Arthroplasty Register. Both registries used data linkage with national patient databases to determine patient/procedure-level completeness (Table S6A and S6B – online available). Techniques to handle missing data were clearly described by only one orthopedic registry (4%),50 which sent requests for missing data to each orthopedic department once every three months. Almost half (46%) of the orthopedic registries, 42,43,46,50,52-55,60,63-65 reported that they implemented techniques for quality assurance of the data, which in the majority consisted of comparing registry data with national patient databases or implant databases.

# Reported Outcomes, Definitions, and Duration of Follow-up

The number of peer-reviewed publications per registry in the period January 2013 – July 2021 varied, with a median of 11 (IQR 3-33) published articles among cardiovascular registries and 9 (IQR 2-45) among orthopedic registries.

A wide variety of outcomes as well as their definitions and durations of follow-up were reported by both cardiovascular and orthopedic registries (Table S7A and S7B – online available).

The most frequently reported outcome in cardiovascular registries was mortality; reported by 18 (90%) registries.<sup>21-24,26-37,39,40</sup> Mortality was reported using 70 different time-points, from in-hospital mortality to mortality at 21-years, the majority of registries (80%) reported on 30-day mortality.<sup>21,22, 24,27-37,39,40</sup> Major

cardiovascular events (MACE) were reported as combined end-points by eight (40%) registries,<sup>21,27-29,32,36,37,40</sup> but with seven different combinations of complications included in this endpoint and seven different time intervals with most (50%) registries reporting on 1-year MACE.<sup>28,29,36,40</sup> Reporting on other single outcomes also showed large variability, ranging from 3 to 40 outcome variables per registry (Table S7A and S7B – online available).

In orthopedic registries, revision surgery (for any cause) was the most frequently reported outcome, reported by 20 (77%) registries. 42-44,46,48,50-60,62,63,65 It was mostly reported as the revision rate or cumulative revision risk but at 30 different time-points up to 25-years, with the most common endpoint being the 1year revision rate which was reported by ten registries (38%). 42,43,46,50-52,56,59,60,66 Specific reasons for revision were reported by 19 (73%) registries. 42-44,46,48,50-57,59,60,62,63,65,66 but these reasons for revision varied between registries (e.g. infection, loosening, component failure, etc). Patient reported outcome measurements (PROMs) were reported by five (19%) orthopedic registries, 44,46,48,63,65 with a total of 8 different scores for knee surgery patients and 11 scores for hip surgery patients. All registries measuring PROMs reported pre-operative PROMs, but post-operative PROMs were measured at different time-points up to 10-years postoperatively. Other outcomes (e.g. renal failure, hip dislocation, deep venous thrombosis, etc) were inconsistently reported by 13 (50%) registries, 44,46,48,50,51,54-56,58,60,62,63,65 the majority (77%) reported on mortality<sup>44,50,51,55,56,58,60,62,63,65</sup> (Table S7A and S7B online available).

## Domain "Safety & performance"

Public reporting on how feedback on e.g. devices, hospitals, and surgeons is provided was reported by three (15%) cardiovascular registries<sup>21,29,36</sup> (Table S8A and S8B – online available). Managerial procedures to detect individual hospitals or specific devices using an outlier performance analysis based on benchmark thresholds was reported by one (5%) cardiovascular registry, the British Cardiovascular Intervention Society registry (BCIS). The outlier was defined using funnel plots, with 2 and 3 standard deviations. Outlier results regarding the timing of treatment (to assess any delay before treatment is delivered) compared between hospitals, as well as adverse outcomes per hospital, were publicly available. However, outlier reports on patients' survival data per hospital were only disclosed confidentially to each hospital. No outlier reports for specific implants were reported by cardiovascular registries.

Public reporting on the frequency of feedback provided was reported by 14 (54%) orthopedic registries.<sup>42-44,46,48, 50,53,55,58,60,62,63,65,66</sup> Most registries report that they provide annual feedback, while two registries (the Irish National Orthopaedic Register and the Swiss national registry for hip and knee replacement) do so both

annually and quarterly. The majority provided feedback both at the hospital-level and for individual devices. Details of outlier procedures including statistical testing were reported by eight (31%) registries, of which three reported solely on outlier devices, <sup>59,60,66</sup> two solely on outlier hospitals, <sup>58,62</sup> one on outlier devices and hospitals, <sup>65</sup> and two on outlier devices, hospitals, and surgeons. <sup>50,63</sup> Outlier procedures were mostly publicly available. No registries shared the same definition of an outlier (e.g. above the 95% control limit in the funnel plot *versus* revision rates of more than twice compared to the relevant group). Overall, in all annual reports, a total of 95 total hip (TH) component combinations, 3 TH cups, 2 TH stems, and 24 total knee (TK) implants were identified by these eight registries as outlier implants. Overall, registries all identified different outlier implants, with only one outlier implant (a TH component combination) identified by more than one registry.

#### Discussion

In this systematic review we have evaluated structural and methodological characteristics as well as the data quality of 46 European cardiovascular and orthopedic medical device registries, in an attempt to gain insight into the usability of these data sources for regulatory purposes. Medical device registries are potentially well suited for post-market surveillance as they may collect data from unselected patient populations and monitor safety and performance throughout the lifetime of specific devices. However, we found heterogeneity and incomplete transparency in quality items related to their structure and methodology, implying that it would be difficult currently for registries to agree upon common principles, to report the information needed by regulators to judge the quality of their data, and to collect and report comparable information across Europe.

The European Union (EU) has regulatory requirements relating to the PMCF of medical devices. 67-69 As stated by the MDR in Article 83, manufacturers have to set up, document, maintain, and update a post-market surveillance system for each device, in which relevant data on the quality, performance, and safety of an implant are evaluated, directly after Conformité Européenne (CE) approval and throughout the entire expected lifetime of a device. 68 To allow for lifetime evaluation and benchmarking of implants, registries need clearly defined methods to detect outliers and to report safety concerns for specific implants, but these were reported by only 5% of the cardiovascular and 31% of the orthopedic registries that were included in this systematic review. Even more, none of the registries used the same definition, making it difficult for manufacturers, regulators, but also patients to assess whether the device performs worse in all or only in some settings. Furthermore, four orthopedic registries identified >100 components and combinations of implants as outliers, with only one outlier implant identified by more than one registry, which may partly result from the different definitions used

from the fact that and that not all implants are used in all countries and/or regions and thereby included in the registry.

Another way to enable benchmarking of implants across registries is to implement objective performance classification systems such as the Orthopaedic Data Evaluation Panel (ODEP). The ODEP rating provides benchmarks for orthopedic prostheses (hip, knee, and shoulder implants) based on the number of years for which the product has been monitored and on the strength of the evidence provided by different data sources, including registry data, randomised controlled trials, peer-reviewed publications, podium presentations, and manufacturers' inhouse data sources.<sup>70,71</sup> The ODEP rating can be considered as an absolute benchmark to identify if implants meet the benchmark criteria, whereas others have suggested relative benchmark approaches within a given registry e.g. comparing with the best implant construct<sup>72-75</sup> or with all other similar implants.<sup>8</sup>

The MDR in Article 108 states that registries need to establish common principles, so that they can collect comparable information and thereby contribute to the independent evaluation of the long-term safety and performance of devices.<sup>69</sup> They need to capture the same outcomes, based on the same definitions and the same durations of follow-up, before they can be used to benchmark devices and pool data for early detection of safety concerns. Current European device registries do not meet these recommended principles, however, since our systematic review showed large heterogeneity between recorded outcomes, definitions of outcome variables, and time-points for follow-up. Comparable findings were reported by a recent study of the quality of cardiac registries across all subspecialties of cardiac care, in which several registries gave explicit definitions for only a low percentage of variables.<sup>76</sup> Similar findings were also observed for orthopedic registries, with considerable heterogeneity in captured outcomes and definitions used for revision procedures.<sup>77-79</sup> Another aspect to consider before outcomes across registries can be pooled, is whether registries use the same implant library to classify implants by relevant device characteristics.80 The European Medical Device nomenclature is a generic classification intended for this purpose, but more detailed libraries are used by registries to capture their specialty-specific characteristics as well. For orthopedic devices for instance, the International Society of Arthroplasty Registers (ISAR) has proposed a global registry library in 2019 to ensure the same classification of orthopedic devices across registries.80 Also, this problem of using different implant libraries can be solved if registries document the unique device identifier for each implant.

In combination, these findings highlight the importance of international agreement on definitions of data and outcomes, as well as time-points used for measuring outcomes within registries. This might be reached by developing

consensus frameworks to achieve common datasets that must be captured by registries<sup>81</sup> such as the clinical outcome endpoints in heart failure trials created by the European Society of Cardiology Heart Failure Association, the common dataset for acute coronary syndromes and percutaneous coronary interventions created by the EuroHeart data science group, the benchmarking document for hip and knee arthroplasties by the ISAR, and the common dataset for demographics and implant survival following THA by the Nordic Arthroplasty Register Association.<sup>82-85</sup>

In addition to these common data specifications, the IMDRF states that registries should include at least 95% of all patients receiving a device, to have sufficiently robust high quality data to inform regulatory decisions. As shown in our systematic review, patient/procedure-level completeness was not reported publicly by any of the cardiovascular registries, but it was available for the majority (65%) of orthopedic registries. Of the latter only 11 of 13 orthopedic registries reported recent data (2018 and beyond) that reached a patient/procedure-level completeness of 95% or above. Similar findings were shown for European TH and TK registries by Lübbeke et al, with 67% reporting patient-level completeness, and for cardiovascular registries, of which the majority had data completeness below 50% or not available.

Making it mandatory to enrol all patients in a registry would help to increase patient/procedure-level completeness. In this systematic review, however, none of the mandatory cardiovascular registries and only 75% of the mandatory orthopedic registries reported patient/procedure-level completeness. Since completeness of patients is often checked against electronic medical records, it could also help to automatically populate certain data fields regarding patient and implant characteristics from the electronic medical records, so that less information needs to be entered by medical professionals, thereby preventing data loss as well as double data entry. However, rather than considering single items that on their own will contribute to higher quality data, the quality of the evidence provided by registry data is ultimately determined by the combination of multiple factors.

The strength of this systematic review is its' comprehensiveness. We updated the search strategy used by Niederländer et al,<sup>18</sup> and expanded it with support from an experienced librarian. In addition, experts in the field (cardiologists and orthopedic surgeons) were consulted, resulting in the addition of two cardiovascular registries. Furthermore, European orthopedic registries listed on the EFORT – NORE-website were checked for their eligibility, resulting in an additional six orthopedic registries and the completeness of included European cardiovascular registries as well as orthopedic registries was checked by experts in the relevant field. Thus, the likelihood of missing relevant registries is very low. However, some limitations remain. Firstly, we relied on publicly available information regarding

registries' structure and methodological characteristics as well as outcomes, which means that some items that we did not find may have been available if we had approached each registry directly. Therefore, the regulatory utility of the data generated by some registries may be higher than that found by this analysis. Secondly, this systematic review only focuses on cardiovascular and orthopedic registries, because they represent the most commonly used high-risk medical devices aiming to reduce patients' mortality and morbidity.<sup>87</sup> However, the items used to determine the regulatory utility of these registries would also be applicable to other (high-risk) medical device registries.

An overview of publicly available information, as summarised in this systematic review, demonstrates the transparency of European cardiovascular and orthopedic medical device registries and what information could already be available for regulators. We have proposed characteristics that can be used to interpret whether the data provided by registries are of sufficient quality, and we have identified registries that had an active/accessible website at the time of study selection and/or that published at least one paper or annual report between 2013 and 2021. No data were collected since 2018 were available for 35% of these registries (shown in Table), and so there is a chance that some are no longer active and thereby would not be able to contribute evidence for regulatory purposes. However, the cut-off point to define an active registry was arbitrary and we therefore highlighted that the median of items reported across all domains among active registries was higher than items reported across all registries combined (i.e. both "active" and "in-active" labelled registries).

## Conclusion

This systematic review showed large heterogeneity and incomplete public transparency related to structure and methodological characteristics of the registries that were reviewed, which implies that it would be difficult to combine and judge the regulatory utility of data reported by registries. Effort is needed from registries to agree upon a minimum set of quality criteria that all registries should publicly report to provide information needed by regulators to judge the quality of registry data and use them for medical device safety surveillance. Developing comprehensive and trustworthy medical device registries will be tremendously valuable, not only for manufacturers to meet the requirements of the MDR for PMCF of their devices, but also for healthcare professionals and patients to support evidence-based choices of devices and contribute to their long-term safety and efficacy.

# **Figures and Tables**

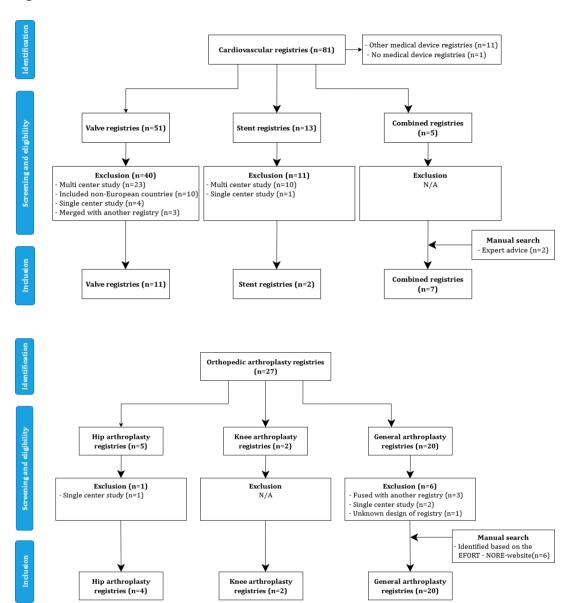
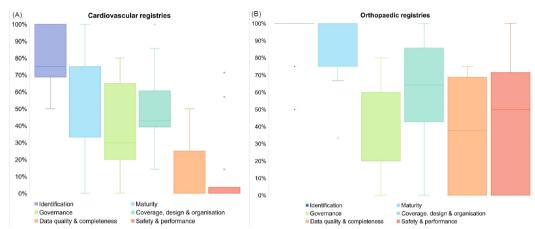
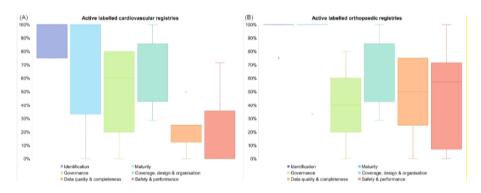


Figure 1. PRISMA Flowchart – (A) Cardiovascular and (B) Orthopedic registries. Abbreviation: PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses



**Figure 2.** Reported items by cardiovascular (A) and orthopedic (B) registries in each domain indicating the variation in reporting across registries (with the lower end of the boxes representing the first quartile and the higher upper end the third quartile; the solid lines in the boxes representing the median values (if not visible the solid lines are at the same level as the first or third quartile); the T-shaped whiskers the maximum or minimum values (without outliers); the individual points representing outlier values)



**Figure 3.** Reported items by the active labelled cardiovascular (A) and orthopedic (B) registries in each domain indicating the variation in reporting across registries (with the lower end of the boxes representing the first quartile and the higher upper end the third quartile; the solid lines in the boxes representing the median values (if not visible the solid lines are at the same level as the first or third quartile); the T-shaped whiskers the maximum or minimum values (without outliers); the individual points representing outlier values)

#### Identification

- 1. Class of device (cardiovascular registries stents/cardiovascular registries valves/cardiovascular registries combined)/(orthopedic arthroplasty registries hips/orthopedic arthroplasty registries knees)
- 2. Name of registry
- 3. Initial motivation/goal to set up the registry
- 4. Country (country or countries in which the registry is conducted)
- 5. Design (regional/national/multi-country)
- 6. Website (available yes/no)

#### Maturity

- 7. Starting year (year of first patient/procedure included)
- 8. First annual report (year of publication)
- 9. Most recent (or last, if registry no longer active) annual report (year of publication)

#### Governance

- 10. Mandatory (if mandatory for surgeons/hospitals to submit to the registry; yes/no)
- 11. Patients' consent (patients' consent required before entering their data to the registry; required/not-required)
- 12. Funding (public/private/both)
- 13. Who can access the data and see results?
- 14. Privacy regulation for patients' identifiable information (privacy regulation reported as implemented: yes/no? And if yes: how?)

## Coverage, design & organisation

- 15. Number of participating hospitals and % of hospital-level coverage (defined as number of participating hospitals relative to the total number of eligible hospitals)
- 16. Number of patients/procedures (cumulative total in registry)
- 17. Number of selected patients/procedures in study population (if cumulative total in registry is not reported)
- 18. Annual number of patients/procedures in registry
- 19. Data capture and collection method (e.g. electronic/manual/barcodes-industry/surgeon-reported)

- 20. Method of access to registry for users/members (e.g. dashboard/real-time/secure server)
- 21. Level of information provided (data is reported at hospital/medical device/surgeon-level)
- 22. Data linkage with other sources (e.g. registry data is linked to hospital statistics/manufacturer vigilance data/national competent authority on medical devices)

# Data quality & completeness

- 23. Quality assurance system defined/quality check of data (e.g. data verification)
- 24. Missing data for patients' characteristics reported (%)(e.g. body mass index, ASA classification, gender)
- 25. Methods for handling missing data described
- 26. Data completeness reported at patient/procedure-level (%)

## Safety & performance

- 27. Frequency of feedback provided to surgeons/hospitals (e.g. annually/quarterly)
- 28. Level of feedback information provided (e.g. hospital/medical device/surgeon-level)
- 29. Feedback time period (the duration of observation before assessment of performance is possible)
- 30. Outlier reports procedures (the type of outlier reports or procedures a registry has established and published methods to define outlier performance)
- 31. Accessibility of outlier results (e.g. publicly available or only accessible for individual hospitals/surgeons/members)
- 32. Definition of an outlier (e.g. using funnel plots)
- 33. Number of outliers identified (has this registry identified and published details of any specific hospitals/medical devices/surgeons with outlier performance?)

**Box 1.** Description of the items in each domain that were extracted for each registry

	Published paper(s) containing registries' data (2018 and beyond)	Published annual report(s) containing registries' data (2018 and beyond)	Active registry	
Cardiovascular registries - combined			5 out of 7 (71%)	
British Cardiovascular Intervention Society	No	Yes	Yes	
East Denmark Heart Registry	No	No	No	
German Society for Thoracic and Cardiovascular Surgery	Yes	Yes	Yes	
Polish National Database of Cardiac Surgery Procedures	Yes	No	Yes	
Portuguese National Registry of Intervention Cardiology	No	No	No	
Spanish Cardiac Catheterization and Coronary Intervention Registry	Yes	Yes	Yes	
Western Denmark Heart Registry	Yes	No	Yes	
Cardiovascular registries - stents			2 out of 2 (100%)	
Polish National Percutaneous Coronary Intervention Registry	Yes	No	Yes	
Swedish Coronary Angiography and Angioplasty Registry	Yes	Yes	Yes	
Cardiovascular registries - valves 4 out of 11 (36%				
Quality Assurance Registry on Aortic Valve Replacement	No	No	No	
Austrian-TAVI Registry	No	No	No	
Belgian TAVI Registry	No	No	No	
Czech TAVI Registry	No	No	No	
FinnValve Registry	No	No	No	
FRANCE-TAVI Registry	No	No	No	
German Aortic Valve Registry	Yes	No	Yes	
Polish Registry of Transcatheter Aortic Valve Implantation	Yes	No	Yes	

Spanish Registry of Heart Valves Repair	No	No	No	
Swedish Transcatheter Cardiac Intervention Registry	Yes	Yes	Yes	
Swiss TAVI Registry	Yes	No	Yes	
Orthopedic arthroplasty registries - combined				
Croatian Register of endoprothesis	No	No	No	
German Arthroplasty Register	Yes	Yes	Yes	
Finnish Arthroplasty Register	No	Yes	Yes	
Irish National Orthopaedic Register	No	Yes	Yes	
Lithuanian Arthroplasty Register	Yes	No	Yes	
Dutch Arthroplasty Register	Yes	Yes	Yes	
Hungarian Arthroplasty Register	No	No	No	
Norwegian Arthroplasty Register	Yes	Yes	Yes	
Nordic Arthroplasty Register Association	Yes	No	Yes	
National Joint Registry for England, Wales, Northern Ireland, the Isle of Man, and the States of Guernsey	Yes	Yes	Yes	
Belgian National Arthroplasty Register	No	Yes	Yes	
Catalan Arthroplasty Register	No	No	No	
National Arthroplasty Registry of Slovenia	No	Yes	Yes	
Italian Arthroplasty Registry	No	Yes	Yes	
Emilia-Romagna Region Arthroplasty Register	Yes	Yes	Yes	
Romanian National Arthroplasty Register	No	No	No	
Portuguese National Arthroplasty Register	No	No	No	
Scottish Arthroplasty Project Joint Registry	No	Yes	Yes	
Slovakian National Arthroplasty Register	No	No	No	

Swiss Arthroplasty Register	No	Yes	Yes		
Orthopedic arthroplasty registries – hips 3 out of 4 (75%)					
Czech Republic Arthroplasty Register	No	No	No		
French Arthroplasty Register	No	Yes	Yes		
Danish Hip Arthroplasty Register	Yes	Yes	Yes		
Swedish Hip Arthroplasty Register	Yes	Yes	Yes		
Orthopedic arthroplasty registries - knees 2 out of 2 (100					
Danish Knee Arthroplasty Register	Yes	No	Yes		
Swedish Knee Arthroplasty Register	Yes	Yes	Yes		

 Table 1. Recent activity of included registries

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- 29. Swedish Coronary Angiography and Angioplasty Registry
- 30. Quality Assurance Registry on Aortic Valve Replacement
- 31. Austrian-TAVI Registry
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- 40. Swiss TAVI Registry
- 41. Croatian Register of endoprothesis
- 42. German Arthroplasty Register
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- 51. Belgian National Arthroplasty Register
- 52. Catalan Arthroplasty Register
- 53. National Arthroplasty Registry of Slovenia
- 54. Italian Arthroplasty Registry

- 55. Emilia-Romagna Region Arthroplasty Register
- 56. Romanian National Arthroplasty Register
- 57. Portuguese National Arthroplasty Register
- 58. Scottish Arthroplasty Project Joint Registry
- 59. Slovakian National Arthroplasty Register
- 60. Swiss Arthroplasty Register
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