



Patient Registries: An Underused Resource for Medicines Evaluation

Operational proposals for increasing the use of patient registries in regulatory assessments

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Abstract

Introduction Patient registries, ‘organised systems that use observational methods to collect uniform data on a population defined by a particular disease, condition, or exposure, and that is followed over time’, are potentially valuable sources of data for supporting regulatory decision-making, especially for products to treat rare diseases. Nevertheless, patient registries are greatly underused in regulatory assessments. Reasons include heterogeneity in registry design and in the data collected, even across registries for the same disease, as well as unreliable data quality and data sharing impediments. The Patient Registries Initiative was established by the European Medicines Agency in 2015 to support registries in collecting data suitable to contribute to regulatory assessments, especially post-authorisation safety and effectiveness studies.

Methods We conducted a qualitative synthesis of the published observations and recommendations from an initiative-led multi-stakeholder consultation and four disease-specific patient registry workshops. We identified the primary factors facilitating the use of registry data in regulatory assessments. We generated proposals on operational measures needed from stakeholders including registry holders, patients, healthcare professionals, regulators, marketing authorisation applicants and holders, and health technology assessment bodies for implementing these.

Results Ten factors were identified as facilitating registry use for supporting regulatory assessments of medicinal products. Proposals on operational measures needed for implementation were categorised according to three themes: (1) nature of the data collected and registry quality assurance processes; (2) registry governance, informed consent, data protection and sharing; and (3) stakeholder communication and planning of benefit-risk assessments.

Conclusions These are the first explicit proposals, from a regulatory perspective, on operational methods for increasing the use of patient registries in medicines regulation. They apply to registry holders, patients, regulators, marketing authorisation holders/applicants and healthcare stakeholders broadly, and their implementation would greatly facilitate the use of these valuable data sources in regulatory decision-making.

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1 Introduction

Health-related real world data provide crucial support for regulatory decision-making, especially in post-authorisation assessments of medicinal products [1]. There are multiple sources including patient (disease) registries, electronic health records, insurance claims databases, health surveys, and prescription dispensing databases [2]. Patient registries, ‘organised systems that use observational methods to collect uniform data on a population defined by a particular disease, condition, or exposure, and that is followed over time’, are a potentially rich source of data, especially for evaluating the course of rare diseases and effects of new treatments [3–5]. Despite this, they are greatly underused in regulatory

Key Points

Patient registries are potentially valuable sources of data for supporting regulatory decision-making on medicines, but they are greatly underused owing to heterogeneity in registry design, the data collected and its quality, as well as to data sharing impediments.

The European Medicines Agency's Patient Registries Initiative aims to support registries in collecting data suitable to contribute to regulatory assessments, especially post-authorisation safety and effectiveness studies.

We have generated operational proposals on patient registry data, quality assurance processes, governance and stakeholder communication that will help to increase the use of these valuable resources in regulatory benefit-risk assessments of medicines.

assessments of medicines. There are many reasons, including heterogeneity in registry design within individual disease areas, unreliable data quality and data sharing barriers, all amplified by limited national and international collaboration [5, 6].

1.1 Use of Patient Registries for Supporting Regulatory Assessments

In pivotal studies supporting marketing authorisation of medicinal products, randomised controlled trial (RCT) data are preferred by regulators. However, in situations where RCT data are limited or where RCTs are not ethical or are not feasible, as with many rare diseases, patient registry data may provide crucial support for regulatory decision-making. For example, in the case of haemophilia, the updated guideline on Factor VIII products removes the obligation to perform clinical trials in previously untreated patients but requires post-authorisation studies based on a set of core data elements to be collected in patient registries [7]. For products granted conditional marketing approval, registry studies may provide post-authorisation data to fulfil regulator-imposed specific obligations to confirm safety and/or effectiveness, as is the case with the recently authorised chimeric antigen receptor (CAR) T-cell products, tiagenlecleucel and axicabtagene ciloleucel [8, 9]. Some registries may be of particular value in terms of the patient population size and representativeness, the duration of follow-up data for treatment-exposed patients and availability of information not collected in other real world repositories.

Notwithstanding such potential, the under-use of patient registries in the regulatory context is striking [10, 11]. Of

335 products recommended for approval between 2005 and 2013 by the main scientific committee of the European Medicines Agency (EMA), the Committee for Medicinal Products for Human Use (CHMP), 31 registries were requested to fulfil a condition of the marketing authorisation, but by December 2017, just ten had been completed [11]. Potentially, registries could provide data permitting comparisons of outcomes from different treatments across different countries and healthcare settings over time as well as assessment of the impact of measures taken to minimise risks of medicinal products [12, 13]. Their limitations for such assessments are well-described, but a description of the features needed to encourage increased use is lacking [4, 12–14]. Our aim with this work was to provide proposals, from a regulatory perspective, on how registry stakeholders could fill this gap.

2 Methods

We conducted a qualitative synthesis of the observations and recommendations published in five reports arising from a multi-stakeholder consultation and four disease-specific workshops conducted as part of EMA's Patient Registries Initiative.

From the consultation report, we identified the primary factors considered by stakeholders as facilitating the use of registry data for supporting medicines regulation.

From each of the four disease-specific workshop reports, we abstracted the participant observations and recommendations on utilising patient registry data in regulatory assessments along with the measures needed from stakeholders in order to implement them. We then generated operational proposals, applicable to patient registries and stakeholder groups broadly, for implementing these measures.

3 Patient Registries Initiative

In 2015, EMA established a Patient Registries Initiative to support a systematic and standardised approach for registry contribution to medicines assessment, especially for post-authorisation safety studies (PASS) and post-authorisation effectiveness studies (PAES) [6]. The initiative aims to create a registry framework with collaboration between registry coordinators, including healthcare professionals' and patients' associations, academic institutions and national agencies responsible for overseeing healthcare services, and potential users of registry data, such as medicines regulators, reimbursement bodies, and pharmaceutical companies. Key elements of its strategy include facilitating the use of existing patient registries within the current legal and regulatory framework for medicinal products and providing methodological support for the establishment of new registries [15].

4 Multi-Stakeholder Consultation and Disease-Specific Registry Workshops

At a consultation in October 2016, 122 expert stakeholders, including registry holders, patients, healthcare professionals, regulators, marketing authorisation holders and applicants (MAHs/MAAs), and health technology assessment (HTA) and reimbursement bodies, and European Commission representatives shared their views on barriers to and facilitators of registry use and on optimising the use of registries for regulatory assessments [16]. The discussions provided the groundwork for four disease-specific registry workshops held during 2017 and 2018 that explored the use of registry data for supporting regulatory assessments in four areas of active product development where new products had recently been approved or were undergoing assessment (Appendix Table 1, see the electronic supplementary material): cystic fibrosis [17], multiple sclerosis [18], CAR T-cell therapies [19], and haemophilia therapies [20].

The four workshops together included 266 participants representing all of the stakeholder groups in each case. The individual workshop reports providing participants' observations and recommendations, along with the report of the multi-stakeholder consultation, are published on the EMA patient registries webpage [6]. The objectives and methods of the individual patient registry workshops are described in Appendix Table 2 (see the electronic supplementary material). This work did not require ethics approval.

5 Results

5.1 Factors Supporting the Use of Registries

Synthesis of the participant observations from the multi-stakeholder consultation report generated a list of factors that facilitated the use of patient registries for regulatory assessments. They included the use of common core data sets, common coding terminologies, complete data collection, especially on medications, facility for data access and sharing, data linkage capacity, quality assurance processes and governance, early consideration of registries in the regulatory process, stakeholder communication, registry sustainability and the availability of a registry framework. Their value in facilitating registry use is described in Table 1. In each case, absence or incompleteness greatly impeded registry use.

5.2 Proposals on Operational Measures to Increase Registry Use in Regulatory Assessments

Three themes, generalisable to patient registries broadly, emerged from the published observations and

recommendations made by the participants in each of the four disease-specific patient registry workshops:

1. Nature of the data collected and registry quality assurance processes
2. Registry governance, informed consent, data protection and sharing
3. Stakeholder communication and planning of benefit-risk assessments.

5.2.1 Nature of the Data and Registry Quality Assurance Processes

The need for registries in a given disease area to collect core common data elements, commonly defined, was acknowledged by all stakeholders as essential for ensuring that data from multiple registries in a given disease area could be combined to enhance both the generalisability and the power of studies that could be conducted using the data (Table 2, Box 1). EMA scientific advice may assist in clarifying the suitability of individual registries for defined purposes [23]. Knowledge of data quality is fundamental for regulatory assessments. Quality may be judged according to three components: consistency, accuracy, and completeness. Table 2 defines each quality component, summarises potential indicators of quality that could be applied in registries, and describes the systems or solutions needed to facilitate these in operational terms.

5.2.2 Registry Governance, Informed Consent, Data Protection and Sharing

Proposals are summarised in Table 3 for measures needed on registry governance, informed consents, and data sharing and protection in order to ensure that data are accessible for regulatory assessments and may be shared in the context of the applicable legal and governance frameworks.

Examples of recommendations from the individual workshop reports are quoted in Box 2. Registry sustainability measures were not a focus of the disease-specific workshop discussions given the regulatory context, but were acknowledged by all stakeholders as crucial for registry stability and development.

5.2.3 Communication with Stakeholders

Acknowledged areas of improvements needed to support the use of registry data in regulatory assessments include communication between stakeholders early in the marketing authorisation process in order to plan for post-authorisation studies (Fig. 1, Box 3). The studies are needed so that marketing authorisation, if granted, may be followed-up with timely evidence on the benefit-risk balance of new products

Table 1 Factors facilitating registry use for supporting regulatory assessments

Factor	Value in supporting registry use for regulatory assessments
Use of common core data sets	Collecting a common core set of data items with agreed definitions and data dictionaries increases the capacity to combine or pool data across patients or registries for regulatory assessments. Ideally, data items match regulatory needs. Capacity to collect additional data elements, even for a limited period, may be beneficial.
Common data coding terminologies	The availability of coding terminologies such as the Medical Dictionary for Regulatory Activities (MedDRA [®]) that could be used by all registries helps in facilitating the conduct of studies using data from multiple registries [21].
Complete information collection	Complete information on critical disease variables is necessary. Medication information is often limited; primary disorder medication information is essential and should include the start and stop (where applicable) dates. Most registries do not record other medications, but some information is desirable. PROs are of increasing interest to stakeholders, but are not collected in most registries.
Data access and sharing	Clear consent specifications on data use facilitate sharing of registry data with third parties including regulators and MAAs/MAHs. Data sharing and access are further determined by relevant national and European data protection legislation.
Data linkage capacity	Linkages to external databases, for example, prescription dispensing, employment, or death registries add to the value of registry data, but linkages may be variable across member state.
Registry reporting, and quality assurance processes and governance	Most registries have processes in place for annual reporting and for quality assurance including source data verification. While these are heterogeneous currently, they represent good baselines for further development in individual registries.
Timeliness of consideration	Consideration of registry data in the authorisation process generally occurs when risk management plans and post-authorisation data needs are being discussed. Planning early in the authorisation process for registry use facilitates data access by reducing timelines for data upload from treating centres and for registry quality assurance processes.
Direct communication	To best fulfil regulator-requested or regulator-imposed studies, regulators, MAAs/MAHs and registry holders need to communicate directly.
Sustainability	Registry funding and support may be limited, causing difficulties in maintaining database systems, reliable quality assurance processes, and staff training. Data entry is often done on a voluntary basis and manually by clinical staff either directly or by importing information from electronic health records. Registry sustainability is crucial for long-term development.
Availability of a regulatory framework	EMA guidelines and procedures for PASS and PAES provide a structure for stakeholder dialogue on registry use [22]. EMA scientific advice can support CHMP qualification opinions or advice on the suitability of a registry for undertaking pharmacoepidemiological studies [23].

CHMP Committee for Medicinal Products for Human Use, *EMA* European Medicines Agency, *MAAs/MAHs* marketing authorisation applicants/holders, *PAES* post-authorisation efficacy studies, *PASS* post-authorisation safety studies, *PROs* patient reported outcomes

for patient and public health, most especially for products with specific obligations where delays in study completion are common [25–27]. Opportunities arise early in the marketing authorisation process to pro-actively identify likely data needs, especially for post-authorisation studies. These opportunities are illustrated in Fig. 1. Where registry data could potentially contribute, there should be three-way contact involving regulators, MAAs/MAHs and registry holders to explore data availability.

6 Discussion

6.1 Priorities for Implementation of Proposals

Our proposals have been synthesised from the published reports of a consultative discussion and four disease-specific workshops that together included almost 400 specialist stakeholders with patient registry expertise. In doing so, we have

leveraged the deep knowledge of participants in each disease area to generate proposals that apply to registries broadly. Implementation of the proposals by registry stakeholders collectively would help to establish a harmonised patient registry environment within many other individual disease areas, thereby increasing the suitability of registry data for regulatory assessments of related medicinal products.

From a regulatory perspective, the priorities for the development of a European Union-wide framework on patient registries are:

- Availability of core common data sets, specific for individual disease areas, with commonly defined data elements across registry networks
- Registry operational procedures for MAAs/MAHs and regulators to access data in accordance with national regulation and European General Data Protection Regulation [20]
- Transparent quality assurance processes in registries.

Table 2 Proposals on data elements and data quality attributes necessary in patient registries and on the operational measures required for implementation

Topic	Proposals	Operational measures required
Core common data elements	Core common data elements to be collected by all contributing registries in a specific disease area	Agree on the core common data elements to be included in specific disease area registries, including the associated definitions and data dictionaries
	Harmonise data element definitions across registries	Provide data element definition information or source to stakeholders
	Agree on core PROs that could feasibly be collected systematically	All stakeholders to collaborate on defining PROs (appropriate as necessary for patient age, capacity, language, and for caregivers)
Data quality	Indicators on data consistency, accuracy and completeness to be implemented and reported	Registries to publish at agreed intervals reports or audits of data quality
Quality components	Indicators of quality	Operational measures required
Consistency: Uniformity of the data over time (e.g. laboratory data routinely entered)	Proportion of data fields changed over time	Audits and centre level data checks
	Proportion of fields missing over time	Standard terminology and coding Standard operating procedures Registry data entry dashboard
Accuracy: Accuracy of data entry—no errors, contradictions or impossibilities in the data Absence of duplicates	Change in value of data filed by <i>x</i> % creates alerts	Drop down menus, alerts, text prompts
	Variability of data values across common fields	Validate registry data sample (e.g. 10%) against source data Software checks Staff training Help screens/desks Funding for data managers
Completeness: Proportion of data missing Absence of core variables	Agreed % of fields completed in audit procedures (e.g. >90%)	Audits Mandatory fields
	Proportion of patients lost to follow-up/attrition rates	Agreement on entry of 'not done' or 'null' values
	Minimum agreed core common data elements reported All treated patients reported, not selected patients only	Engagement with patients and HCPs Agreed list of data elements and definitions Cross-check patient numbers with numbers of products used at treating centres (applicable for some advanced therapies)

HCPs healthcare professionals, PROs patient reported outcomes

Box 1 Nature of the data collected and registry quality assurance processes

Workshop participants' recommendations included:

'Agree on standards for data quality indicators, terminologies/coding and reporting requirements to apply to national registries and to the ECF-SPR' (European Cystic Fibrosis Society Patient Registry) [17].

'Agreement on the data elements to be collected in MS (multiple sclerosis) registries would facilitate treatment evaluations and comparisons of safety and effectiveness outcomes between different MS populations and across multiple countries' [18].

'Established quality standards should be in place and adequate for all registry studies; a dedicated data control and follow-up system should be introduced only for very specific studies or where the existing system is not [yet] adequate' [19].

'Definitions for the data elements required by the FVIII Guideline need to be agreed and applied across treating centres and registries; the associated data dictionaries need to be established and maintained' [20].

Therefore, in addressing the factors described in Table 1, the proposals on data elements, quality assurance, governance, patient consents, and data protection summarised in Tables 2 and 3, together with communication and planning early in the authorisation process, are critical if the potential of patient registry data for regulatory assessments of medicinal products is to be realised.

6.2 EMA Actions to Increase Registry Use

EMA has provided scientific advice to support CHMP qualification opinions on two patient registries regarding their suitability for supporting regulatory assessments of medicines, the European Cystic Fibrosis Society Patient Registry and the cellular therapy module of the European Blood and

Table 3 Proposals on measures required for registry governance, informed consent, data protection and sharing

Topic	Proposals for measures needed from stakeholders
Registry governance	<p>Regulators and/or MAAs/MAHs to identify early in the authorisation process whether a potentially relevant registry exists and identify data elements needed, especially for post-authorisation assessments likely to be requested or imposed, and to agree on a common study protocol.</p> <p>Regulators and MAAs/MAHs to be aware of the data elements that can feasibly be collected systematically by relevant registries and to inform registries on their data needs.</p> <p>Registry holders to establish a centralised data application <i>process</i> (with a standard template) for stakeholders to request and obtain data.</p> <p>Communicate to patients and the public the benefits and uses of patient registry data and the value of high levels of patient inclusion in registries.</p>
Informed consent	<p>Registry holders to ensure clinical/treating centres confirm that registry patients have provided consent and review whether current patient consent is broad enough for possible future situations taking into account European GDPR [24].</p>
Data sharing and data protection	<p>Registry holders to develop a <i>policy</i> on data analysis and sharing summary, pseudo-anonymised, and individual patient data that aligns with national regulation and European GDPR.</p>

GDPR General Data Protection Regulation, MAA/MAH marketing authorisation applicant/holder

Box 2 Registry governance, informed consent, data protection and sharing

Workshop participants' recommendations included:

Registry holders need to optimise communications with patients, MAHs, and regulators by: informing patients on the benefits and uses of patient registry data including appropriate sharing with relevant stakeholders and by informing MAHs and regulators of the type and detail of registry data that may feasibly be shared within consent and governance parameters' [17].

Standing agreements between MAHs and registry holders could facilitate provision of data for regulatory procedures, either routine (e.g., periodic safety update reports (PSURs), or exceptional (e.g., during a referral procedure) [18].

Data analysis should preferably be performed by the registry owner or by a third-party (e.g. academic centre, contract research organisation) rather than by MAHs/MAAs. If data analysis is conducted by the registry holder or a third party, results of product-specific data analysis should be shared with regulators and the concerned MAHs/MAAs in line with provisions of the study protocol' [19].

Prior to commencing imposed studies, transparent arrangements should be in place for sharing and publishing data and results' [19].

Registries should take a central role in working with their affiliated treating centres to harmonise patient consents ensuring they are aligned with the GDPR as well as with national requirements allowing sharing of aggregated and anonymised patient-level data for research or regulatory purposes' [20].

Specific protocols need to be sufficiently detailed as to allow registries to assess whether they can participate (in terms of data availability and quality)' [20].

Marrow Transplant registry [19]. In both cases, the opinions describe contexts of use for which the registry data are considered suitable by CHMP for undertaking pharmacoepidemiology studies. The possibility for registries to obtain scientific advice to support a qualification opinion may go some considerable way in assuring stakeholders that registries so qualified are satisfactory for regulatory studies.

Internally, EMA has instituted measures to identify products during pre-submission stages of authorisation processes (pre-submission meetings, scientific advice and priority medicines [PRIME] discussions [28]) (Figure 1) where registry or other real world data may be needed for post-authorisation follow-up if marketing authorisation is granted. This action anticipates and permits pro-active planning for post-authorisation assessments and reflects the regulatory policy of benefit-risk assessment throughout

the product lifecycle. To assist stakeholders in identifying potentially relevant registries, a publicly available inventory of patient registries is hosted on the European Network of Centres for Pharmacoepidemiology and Pharmacovigilance (ENCePP) resources database [29].

EMA should not duplicate other initiatives aiming to enhance registry use in healthcare. Therefore, its patient registries strategy aligns with the European Commission policy framework on rare diseases as well as with priorities of the European Research Networks for rare diseases, the Horizon 2020 programme and Joint Action initiatives such as the European Network for Health Technology Assessment (EUnetHTA), and takes into account national endeavours such as those underway in the Netherlands and Sweden [30–35]. The European Platform on Rare Diseases Registration (EU RD Platform) has developed a 'Set of common data

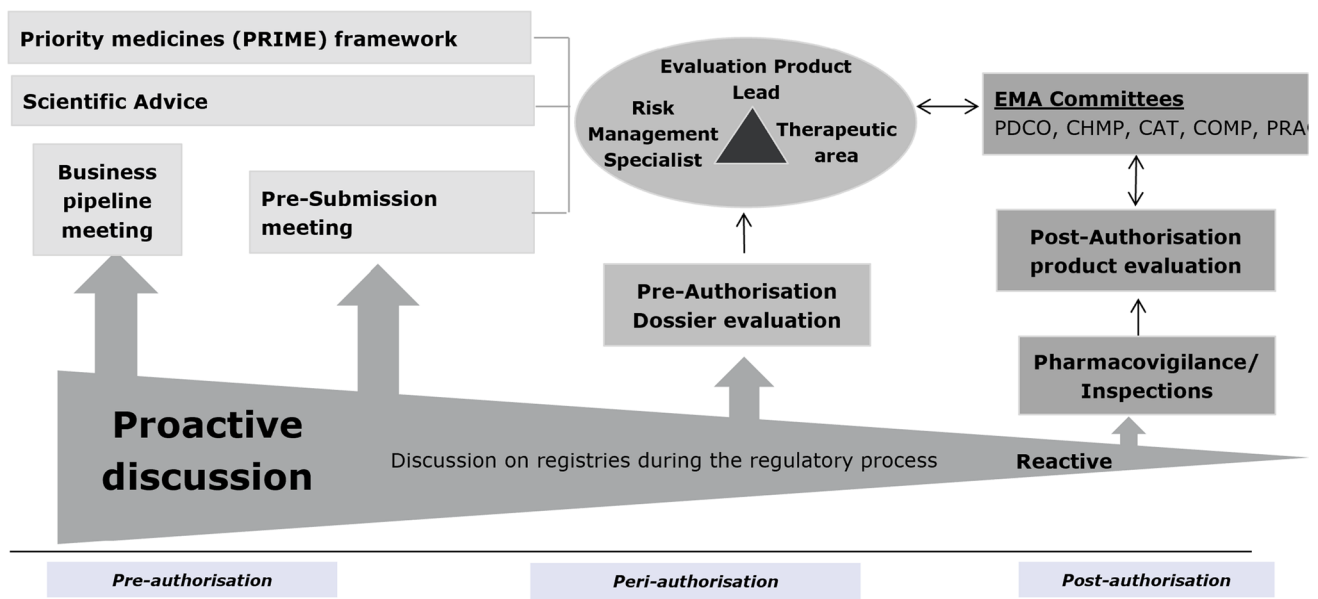


Fig. 1 Opportunities during the regulatory cycle to identify where registry data may be needed for post-authorisation follow-up. Source: Clin Pharmacol Ther 2019 <https://doi.org/10.1002/cpt.1414>

Box 3 Stakeholder communication and planning of benefit-risk assessments

Workshop participants' recommendations included:

Communicate the value of registries, their limitations, and the importance of consistent data quality to all participating healthcare professionals and to those using the data including MAHs, regulators, HTA and reimbursement bodies' [17].

MAHs, regulators and registry holders, plus other stakeholders where relevant (for example, reimbursement bodies), should engage in discussions early during the regulatory processes for approval of new treatments to consider data needs and scientific / study protocols and to understand the range and nature of data that registries could provide, especially for post-authorisation studies' [18].

MAHs / MAAs need to 'commence planning for post-authorisation data collection early in product development' and 'develop a preliminary study protocol and explore with the registry holder/s and regulators if the registry could fulfil the data needs, for example, through a scientific advice procedure' [19].

MAAs must '[i]nitiate discussions with registries and regulators before or at an early stage of a marketing authorisation application on the relevance and adequacy of one or several existing disease registries for the long-term monitoring of their specific product' [20].

elements for RD Registration' [36]. It is aimed at the European Reference Network's existing registries and registries under development; other rare disease registries at national, regional, and local levels in EU Member States; researchers and patient organisations.

From a regulatory perspective, the ultimate requirement of patient registries is that they permit the conduct of high-quality studies that evaluate the safety and effectiveness of medicines. Recognising the benefits and challenges inherent in using observational data for medicines assessments, in 2019, EMA will publish methodological and operational advice on handling registry data in post-authorisation studies, taking into account responses to its open consultation on a preliminary discussion paper [6]. Application of the

advice will be underpinned by clear understanding of the differences between a registry and a registry study (Table 4).

7 Conclusions

This is the first time that the factors necessary for patient registry data to adequately support regulatory assessments, together with operational proposals required for their implementation, have been set out explicitly from a regulatory perspective. In explaining what is needed, taking account of the current legal and regulatory framework for medicinal products, the proposals empower stakeholders seeking to capitalise on the potential of patient registries broadly to

Table 4 Methodological aspects: distinguishing a registry from a registry study

Characteristic	Registry	Registry study
Nature	Data collection system	Investigation of a research question or hypothesis
Timelines	Long-term, open-ended	Defined by the study objective and described in the study protocol
Patient enrolment	Exhaustive within the boundaries of the purpose of the registry (e.g. all patients diagnosed with a disease in a hospital, region or country)	Defined by research objective and described in the study protocol—it may be a subset of the registry population
Data collection	Wide range of data may be collected depending on the purpose of the registry	Restricted to what is needed by the research question including data on potential confounders and effect modifiers—additional data collection may be required
Analysis plan	Routine periodical data analysis; additional ad-hoc analyses	Statistical analysis plan separate from the study protocol in line with the objectives
Collection and reporting of suspected adverse reactions	National requirements as regards the management of safety data apply. Any active data collection with involvement of a MAH must follow the regulatory framework for PASS	National requirements may apply. Regulatory requirements to MAHs differ between studies with primary or secondary data collection
Data quality control	Applied routinely to all data and processes	Additional quality assurance may be needed
Regulatory status	Non-interventional	Non-interventional or interventional

MAH marketing authorisation holder, PASS post-authorisation safety study

support and contribute to regulatory decision-making on medicines.

Actions by all stakeholders, registry owners as well as MAAs/MAHs, regulators, patients/their representatives, healthcare professionals, and HTA and medicines reimbursement bodies, are required to implement the proposals and thereby consolidate registry value in patient and public health.

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Compliance with Ethical Standards

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