

Survey questions, with collected feedbacks sorted by participating sites

Site Mapping

Sites-ID	Site Name
1	-
2	-
3	-
4	-
5	-
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7	-
8	-
9	-
10	-

Question 1: On average, how many data requests or local data use projects are handled at your site DIZ per quarter?

Sites-ID	Feedbacks
1	20 (We expect significantly more requests in the following quarter, as our data catalog is still in the publication mechanism. The 20 therefore refer to requests that have already been generated even though the data catalog has not yet been published.)
2	Our UAC processes approximately 3 data use requests per quarter.
3	approx. 15
4	An average of 4 requests per quarter
5	approx 2 per quarter
6	3 (estimation)
7	Q1/2022: 4 projects Quarterly very irregular Total since project start 35
8	Officially 0, as not yet approved by data protection
9	approx. 6 in Q1/2022 n=7
10	One data request per quarter on average

Question 2: Which contents are mostly in focus in your local data use projects (e.g.: care evaluation in transfusion medicine, etc.), and which data repositories are most frequently queried in this context (i2b2, OMOP, FHIR, ...)?

Sites-ID	Feedbacks
1	<ul style="list-style-type: none"> clinical research Self-research by physicians Currently, the clinical data repository is the most queried. FHIR is in the process of implementation.
2	Internal research queries, quality ensuring and reporting are mainly performed using the DWH. The i2b2/OMOP/FHIR repositories are mainly used for MI-I/MIRACUM specific requests.

3	Qualifying research questions (doctoral dissertations etc.), quality assessment, proof of qualification, where so far mostly the mirror system of ORBIS serves as data repository; the mentioned i2b2/OMOP/FHIR mostly play a role only for MI-I-/MIRACUM-specific queries.
4	Lab values and diagnosis within specialties: Neurology, Urology, Pneumology, Internal Medicine. i2b2, omop, cdr (internal projects), fhir the analysis is performed on OPAL/DataSHIELD
5	Clinical research questions e.g. number and context data on splenectomies, context data on urological sepsis. Target repository is i2b2 and FHIR
6	Query only possible directly at Data Integration Center (DIC); DIC extracts the data and makes it available for use. The most frequently requested data items are stored in the local research repository CentraXX. This is followed by requests for FHIR data from the national projects.
7	Case numbers for diagnoses/treatment procedures Requests by all specialties i2b2 (with Apache Superset as interface since 2022) fhir-server
8	n/a
9	Department- and unit-specific clinical questions e.g., prediction of departmental sepsis and associations with specific treatment procedures/ICD diagnoses. Other example: patient case-based analysis of multiple clinical complications associated with specific clinical and demographic characteristics. Most queries through the cDWH and i2b2 repo.
10	Mainly retrospective data analysis in pulmonology. Analyzing is performed via DataSHIELD, therefore no direct query in data repositories. (Indirect i2b2)

Question 3: How are data use project-specific data quality (DQ) requirements collected from the perspective of data requesters at their DIC?

Sites-ID	Feedbacks
1	During the data request, we advise that the requested data should be described as fine-grained and exact as possible. If the data provided does not match the request, a "post-processing" process will be initiated.
2	In general, the heads of the projects contact the transfer office/UAC office and clarify which data can be extracted and which variables are useful for a scientific evaluation and what should be considered (specific conventions/documentation)
3	I am not sure exactly how the question is meant. In any case, the requested data are usually discussed at least once with the requester and quality-reducing aspects are worked out together, e.g. free text information, documentation practice in the respective data-providing institution (usually the requester comes from the same institution and knows it very well).
4	Is not collected.
5	In interactive discussion with the researchers Environment at the moment still too heterogeneous for a standardized approach
6	In personal conversation during consultation.

7	manual explorations of the data with requesters and providers 100% correct data quality is assumed
8	n/a
9	Project-related data quality requirements are gathered using a Feasibility Request (FR) form completed by the data requester & internal data request administrator. Documentation of the intended cohort property (in terms of expected minimum cohort size,...) and project specifications (e.g. inclusion and exclusion criteria) takes place there
10	These are additionally described in the project proposal under the item "Data description".

Question 4: In addition to the current MIRACUM DQA tool, what tools or technical approaches do you employ for data use project-specific data quality assessment?

Sites-ID	Feedbacks
1	An initial concept of completeness of data elements is under review and will be implemented in Q3 2022
2	For project-specific validation, comparison of hit ratio from different systems created by an independent person: e.g. separate i2b2 SQL queries compared to FHIR/staging area/DWH queries, etc. Before data delivery/provision, mutual control (DIC internal as well as with clinicians) and official release of results by the head of the transfer office.
3	Mutual control before issue/provision (4-eyes principle), an MDR-supported DQA tool is under development
4	Simple site-specific count comparison of identical SQL content on CDR and source DB is established.
5	Resource-specific tracking of the datapath based on a unified system of FHIR business identifiers.
6	no further tools in addition to MIRACUM DQA-Tool.
7	n/a
8	n/a
9	4-eyes principle: Content validation of the queries by a second data scientist (possibly also with a separate query), so that it is ensured that the query actually does what it is supposed to do. Content-related plausibility control of the results from the query through medical colleagues.
10	Formless communication to the transfer office of the DIC

Question 5: What measures are taken at your location to communicate with data requesting sites about the quality of data provided for the specific purpose of the data use project, so that data requesters have opportunities to estimate the fitness of the data for the intended project?

Sites-ID	Feedbacks
1	Creation of a transfer office. The transfer office communicates with the data requesting offices. After data provision, the transfer office inquires about the satisfaction / suitability of the data with the data requesting office. After checking with the data requester site, this consults with the transfer office. If there are deficiencies in the quality of the data, the transfer office forwards this to the architects of the data. They contact the data requesting office directly in order to work out solutions together.

2	Conduct feasibility study Communicate mid-term results
3	This is done in direct dialog with the requester. (see also 3)
4	The DIC advises the data requesters individually. So far, there are only a few projects in which the DIC was not scientifically represented.
5	Overview dashboard in the self-developed data integration portal
6	Not relevant yet
7	Scope of the core data set vs. expectations in the context of a consultation. Feasibility queries Comparison with known data from the hospital vs. data set together with requesters Provision of a data dashboard for own queries by requesters
8	n/a
9	Delivery of the data with involvement of the data requesters First, the feasibility request determines to what extent the number of patients suitable for the planned project is available in sufficient amount Then the data are delivered by the data request administrator, who goes through the data to be delivered together with the data requester. In case of change requests/incorrect quality in the data, the data selection queries are adjusted and validated and documented again via the 4-eyes principle This results in the feedback cycle: data requestor => data request administrator => internal data scientists => data request administrator => data requestor Only in case of a complete match (from the data requestor's perspective) the final data delivery takes place.
10	Plausibility check of the provided data together with researchers (physicians) before using the data for the analysis. Use of the uniform data dictionary (metadata). Verification of the data format or type, the number of variables via DataSHIELD before the analyses. If it detects inconsistencies in the research data, it will cross-check them with the source system and identify problems

Question 6: What would be their expectations/requirements for a fitness-for-use cross-site DQ framework that you could adopt in the future to measure DQ related to their data use projects?

Sites-ID	Feedbacks
1	Implementation of a dashboard
2	Flexible organization of the DQ system Locally assessed DQ compared to sites Integrate project-specific data plausibility Understandability for the clinician and data scientist/statistician Fitness-for-use dashboard
3	Generally enough that it can be used in every DIZ and for every request. It should be pragmatic and easy to understand, so that it can always be used as a basic tool and its benefits are seen equally by all parties (data provider, data supplier, data requester). In

	the short term, it is limited to the essentials to be able to use it and gain experience. In the long term, it may even be possible to modularize it and thus use it only in parts.
4	Graphical representation over time (gaps, leaps in values).Graphical representation over time (gaps, leaps in values).
5	Integration of the already used resource-specific tracking of the datapath based on a unified system of FHIR business identifiers into the DQ system.
6	These cannot yet be definitively determined
7	Complete non-interactive integration of the DQ process as an operation within the data pipelines for complete monitoring of the mapping of source and target systems with automatic machine-readable report generation (no PDF) Automated comparison of previous reports (in the context of performed developments or updates)
8	Completeness Plausibility Currentness
9	Provision of a uniform template for documenting DQ and possibly also data requestor feedbacks in the context of project-related data deliveries across the DIZs Mapping and automation of DQ checks based on the specific data quality metrics <ul style="list-style-type: none"> • Data completeness: are there enough patients at the DIZ site to carry out the planned projects • Data plausibility: formulation & automation of general-transferable plausibility checks (e.g., no readmission after a death, ...) that could affect the outcomes of most DRs • Data conformity: uniform mapping and verification of conformity of ICD , OPS, LOINC codes, and adequate reporting in the systematics Structured Provenance Documentation: <ul style="list-style-type: none"> • where did the data come from, • what processing steps were performed on the data up to the time of data delivery, • Are there changes to the data that may represent a potential impact on the planned data use project? FHIR as a single target repository for the data requests (also needs to be coordinated across DIZ). Inclusion of i2b2 and OMOP as additional repositories depending on whether a specific repo is preferred/specified by the data request.
10	Uniform FHIR profiles across MIRACUM partners. Standardization of LOINC mapping Uniform measurement units