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DATA HANDLING

Your boost for enhanced data quality

Challenges

The requirements for data quality, integrity and traceability in clinical trials are constantly increasing - as is the effort required for data collection, transfer and query management. For trial centers with limited staffing availabilities, professional data processing is therefore becoming an increasing challenge. These challenges are compounded by complex eCRF systems, specific trial protocol requirements and the need to provide parallel patient care.

- **Limited resources in test center**
  Many sites don't have sufficient staff that ensures qualitative data collection and efficient conduct of the study.
- **Complex eCRF systems & high data integrity requirements**
  Using IT systems and databases alongside strict regulations makes it difficult to capture data that is both complete and error-free.
- **Investigational sites focus on patient care**
  Site teams often have limited experience with working on research data, which can lead to delays and increased efforts spent on correcting data.

Why us

- ✓ Established workflows
- ✓ Selected team extension
- ✓ Seamless documentation
- ✓ Accelerated data management
- ✓ GDPR compliance
- ✓ Enhanced data quality

We care

Camovis Data Services focus on the accurate and compliant collection, cleaning, and integration of study data at your clinical trial site. We ensure data is valid, complete and entered in a timely manner. We resolve queries in accordance with regulations and with maximum efficiency, thereby reducing the workload of your site staff.

Our experienced research professionals are specially trained in clinical trial data management ensuring you study data is reliably processed and quality-assured from start to finish. Following a thorough needs analysis, we will clarify all the important details with you and establish a solid foundation for a clean and smooth process. If desired, we then define customized workflows and prepare the team for your project. Throughout the data collection and processing stages, we ensure careful recording, continuous coordination and strict validation with the ultimate goal of achieving the highest data quality. This provides transparent verified data that fully meets all the requirements of the sponsors and regulatory authorities.

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Service Packages

Data transfer & integration

- Data validation prior to input to detect errors (sources) at an early stage
- Ensure compatibility between source data & eCRF
- Transfer of source data to eCRF and database systems

Query resolution & data cleansing

- Identification and elimination of data inconsistencies
- Tracking and documentation of open queries
- Proactive avoidance of incorrect data entry through data plausibility checks
- Close collaboration with study centers and monitors to clarify issues

Retrospective studies & archive data processing

- Identification & extraction of relevant study data from archive systems
- Data preparation and structured archiving for later analysis
- Harmonization & standardization of data for regulatory requirements

Decentralized Trials (DCT) - Data Handling

- Camovis Data Handling Services support the decentralized conduct of clinical trials. Please refer to the DCT Service Information to find out more about the wide range of available options.

Let`s get to work

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Feel free to contact us — we're here and happy to help! :)