

Benefit from Kayentis' data management culture for all your clinical studies

and consider data management at each step of your study



1 Get ready before study launch:

WE MAKE SURE EVERYONE HAS THE SAME EXPECTATIONS FOR THE DATA

We assign a Data Manager to your study who:

- Joins from the very start (during the Kickoff Meeting)
- Reviews the protocol



2 Set-up phase: WE THINK OF ALL THE "WHAT IFs!"

The Data Manager:

- Drafts study specific documentation
- Defines which data can be updated & which profiles are allowed to update data
- Ensures programming & validation of all edit checks



3 Throughout the study: WE ENSURE THE DELIVERY OF CLEANED DATA TO SAVE TIME AT THE STUDY CLOSE

- Edit checks automatically raise flags on our Clin'form WebPortal
- Reminders for query management are sent to the investigator/site staff
- Kayentis Data Managers monitor all queries on our Clin'form WebPortal
- Data changes are quality controlled and audit trailed
- Data transfer files are generated automatically by the validated export module of the Clin'form WebPortal
- Monthly data transfers can be ramped up as the study close approaches



4 Closing the study: WE GUARANTEE THE QUALITY OF THE FINAL DATABASE, DELIVERED QUICKLY TO SITES AND CLINICAL TEAMS

- A Quality Check Form is completed for each device used to collect patient data
- A global quality control is performed before the final database lock
- The database lock is completed within 3 working days maximum



Ask us for a demo at sales@kayentis.com