



VALOS

Statistical solutions
for your clinical trial

Data Management

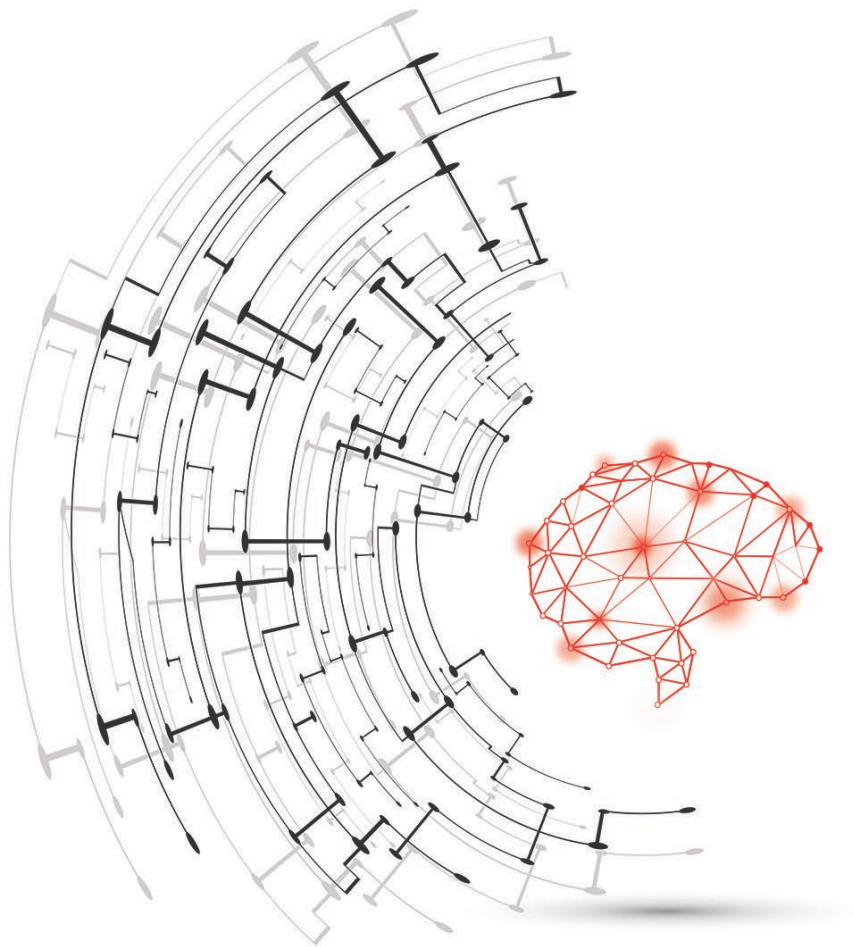
Statistical Analysis

Statistical Programming

Regulatory Submission

CDISC Specialists

Strong Education



www.valos.it

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VALOS provides comprehensive **data management, biostatistics, and statistical programming services** for Phase I-IV studies and retrospective analyses.

If you need to outsource a part of your clinical trial or drug development project, **VALOS** is the partner you can count on.

VALOS's professionals will communicate effectively and will meet project milestones using processes that match the Sponsor's systems.

VALOS ensures best quality and increases the success of your clinical trial.

Statistical Programming

- Statistical Programming in SAS® and R languages
- Data analysis, including designing analysis data sets, statistics, interim analysis, and customized DSMBs (also posted via a secure Web site)
- Reporting of results, production of analysis summaries: production of Tables Listings Figures (TLF) according to Client's needs
- Exploratory analyses for publications, abstracts, and marketing
- Double independent validation
- CDISC data conversion (SDTM, ADaM)
- Graphical patient profiles
- Collaboration on ad hoc analyses, regulatory interactions, applications (IND) and annual safety report, reconciliation of safety data
- Pharmacokinetics
- Programming support for Data Cleaning and Discrepancy Management
- Creation, validation, and auditing of clinical database and data warehouse

Our people

At VALOS all 80+ of our professionals hold a university degree in Mathematics, Statistics, Genomic Statistics or Informatics and have 10+ years' experience in clinical trials. 8 programmers have more than 10+ of experience in SAS and R programming for Clinical Trials.

Focus Areas

- Oncology
- Neurology
- Cardiovascular
- Gastrointestinal
- Respiratory
- Allergology
- Immunology
- Antibiotics
- Ophthalmology
- Epidemiology
- Medical Devices



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