cliantha
Science Integrity

BIOMETRICS

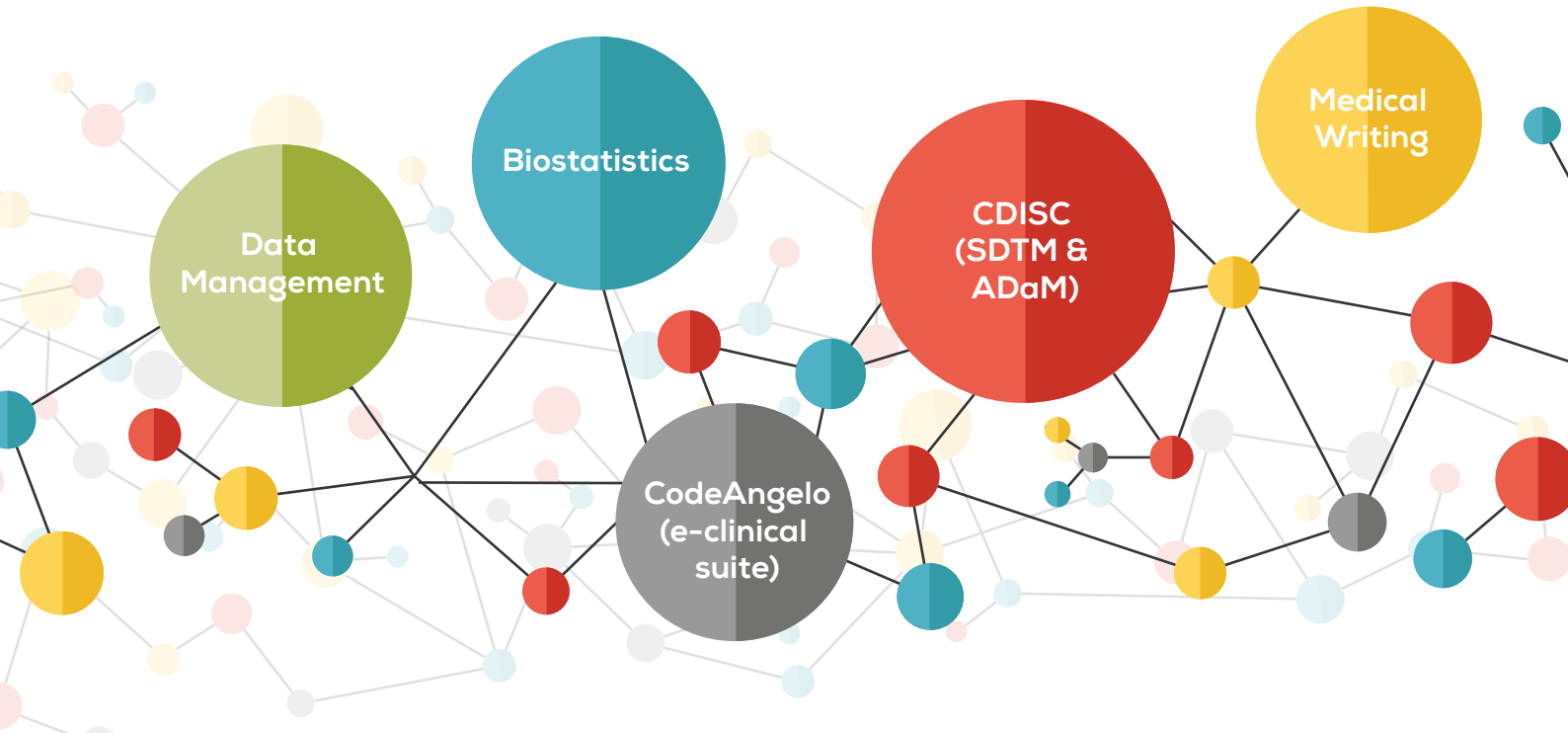
Clinical Trial Data Management Solutions

Cliantha provides integrated services offerings in the areas of Clinical Data Management, Biostatistics, Consumer Research and Third party auditing/monitoring and consulting services.

Our aim is to provide the best services to our sponsors through offering hands-on solutions with the right technology, meticulous planning and seamless implementation. Our Validated Clinical Trial Management System CodeAngelo provides the platform which is easy to deploy, user-friendly and is a next generation, 21 CFR Part 11, integrated EDC + CTMS + eTMF + SDTM Mapping Tool with a Clinical Data Repository.

Our services include:

- **Data Management**
- **Biostatistics & SDTM - ADaM Services**
- **Medical Writing**
- **CodeAngelo eClinical Suite**



"Your trusted partner for all your Data Operations needs"

Experience in different Therapeutic Area: (but not limited to this)

- Oncology
- Dermatology
- Diabetes
- Medical device
- Respiratory
- CNS
- Women and Men's health
- Ophthalmology

Clinical Data Management:

- Complete Clinical Data Management services from CRF Design , Deploying the database, Query generation to Database Lock (DBL) for both eCRF & Paper studies.
- Robust IT infrastructure ensures your data security, privacy & uninterrupted services.
- Rich experience of working with industry renowned CDM systems like Rave, Inform, Medrio, CodeAngelo, Acceliant, Omnicomm, SAS PheedIT, ClinTrial, etc.
- Efficient and tested processes with detailed documentation including SOPs, working procedures & study specific documents to comply with ICH standards, national regulations & guidelines.
- Highly skilled talent pool, who have worked on Phase I-IV trials for a variety of therapeutic areas including Biologics and Biosimilars having study duration ranging from 1 month to 5 months and having patient population ranging from 20- 25000.
- Stringent quality procedures including senior review (QC) & independent QA at various stages of documentation & data processing.
- Our primary objective is to deliver high quality data, on time, to achieve client's expectations and to supply a data set worthy of regulatory submission.

"Successful track record of **120+ Paper CRF & 75+ eCRF Studies**"

CDISC

CDISC standards help improve data exchange, increase study efficiency, saves time and speeds up approval and makes standardization possible. We have a comprehensive package that includes STDM and ADaM datasets.