

Service

## Data Management and Database Development

We'll work with you and your team on the management of your data from start to finish of the trial, adhering to ICH and all other regulatory standards

Typically, the service might include any or all of the following:

- Working with your database and clinical teams to develop the tools (often eCRF) to ensure collected data's in the right format for your study's question – and that you get all the data you need
- Design, development, implementation and hosting of databases for ethics-approved non-CTIMP studies and external CTIMP studies
- Design, development, implementation and hosting of bespoke data management software and web applications (including specialist expertise with statistical algorithm-based servers).

Far less typically, but vitally, we also offer the following services in case of serious adverse event notification:

- Continuous discrepancy management: that is, ensuring your data is of the highest standard – a process that will call for extra resources
- Documenting all data management/laboratory data management activities and plans and so on
- SAS dataset preparation.

Data Management and Database Development is just one of the services we offer to compliment your team's very specific strengths and specialisms. Equally importantly, we find our research partners come to value the peace of mind to be found in working closely with a team for who co-operation in a common purpose is a way of life – and the only way we work.

**Please don't hesitate to get in touch, talk through the possibilities, and ask us to provide a quote for this or one of our other statistical services.**



## Other services

### **Preclinical and Clinical Trials**

*From developing research strategy and the protocol's statistical and design aspects to ethical approvals and submissions for publication or regulatory approval.*

### **Observational Studies and Electronic Patient Records**

*From case series to hypothesis generation and beyond, we'll work with you to validate your epidemiological ideas and make the best possible use of your data.*

### **Medical Devices**

*Working with you from start to finish to ensure your project is in accordance with all relevant regulations, including development of research strategy, ethics approvals and writing the statistical analysis plan and related information.*

### **Health Technology Assessment**

*Specialist expertise in the evaluation of novel drugs, medical technologies and diagnostics to support your HTA submissions and assist in achieving reimbursement.*

## Have you got a project that might benefit from our help?

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