

Service

Preclinical and Clinical Trials

We'll work with you from the start to finish of your trials, ensuring that the statistical aspects of your project are in accordance with ICH/GCP and internal SOPs as well as complying fully with MHRA regulations.

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

- Research strategy development
- Preclinical animal studies selection, study design and modelling – and interpretation of resulting data
- Developing the protocol's statistical and design aspects
- Working closely with our data management team to ensure that we collect all the necessary and precisely the right data
- Writing the statistical analysis plan and related information for the Data Monitoring Committee and Trial Steering Committee as required
- Writing reports for publication or submission to regulatory agencies.

Of course Preclinical and Clinical services are just one of many ways in which we can compliment your own 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 whom 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****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.

Data Management and Database Developments

We will 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 to ensure you get all the data you need in the format your study requires.

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?

Email: medstats-kch@kcl.ac.uk
Internal email: KCL Electronical Mail Directory
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