

Service

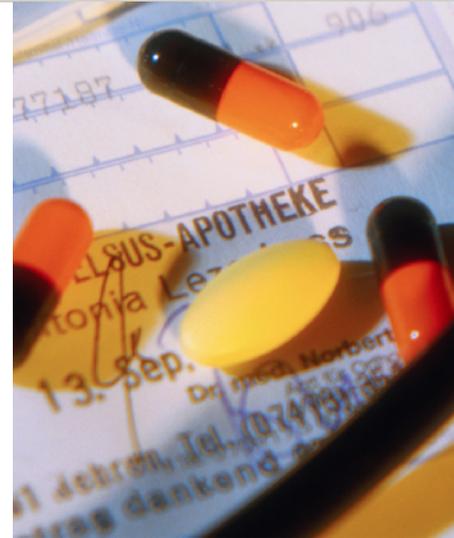
Health Technology Assessment

You'll find the support of our specialist expertise invaluable in evaluating novel drugs, medical technologies and diagnostics for your Health Technology Assessment.

Adhering to all relevant regulatory regimes, our comprehensive approach to statistical support effectively supports the assessment of drugs, medical services and diagnostics – through randomised controlled trials, systematic reviews, meta-analyses and the setting up and running of registers and databases.

Of course this is 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****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.

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.

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.

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

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