

A flexible approach to

Alzheimer's Disease Clinical Trials



CUSTOM-BUILT RESEARCH™

TIME IS OF THE ESSENCE IN DRUG DEVELOPMENT.
CONTACT THE QPS BUSINESS DEVELOPMENT TEAM TODAY!

CALL +1 512 350 2827 EMAIL infobd@qps.com



QPS has excellent connections with leading AD researchers, and can therefore support site selection that guarantees the highest quality investigators and efficient patient recruitment and retention.

QPS has a longstanding history of conducting trials in neurodegenerative disorders such as Alzheimer's disease (AD). Our clinical team is specialized and focused on AD treatment development. The team consists of medical, regulatory and operational experts with significant experience in AD study planning and conduct. Our clinicians are aware of the latest diagnostic criteria, and the most recent developments in study design, to meet the requirements of modern AD research. We have performed trials in Phase I, Phase II and Phase III, and we have nearly 3,000 potential AD study subjects in our database.

QPS Regulatory Affairs successfully supports study start up, while offering strategic advice on regulatory pathways and possibilities. In addition, we provide direct access to, and organize/chair meetings with relevant contact persons at Regulatory Agencies. Whether you are planning a clinical trial of small molecules, biologics, immunotherapies or medical devices, QPS has the experience and resources to handle your study and deliver high quality data on time and within the planned budget.



QPS' Experience in **AD Clinical Development**

TRIAL EXPERIENCE

Study Design of **16** Trials

Regulatory Affairs in **21** Trials

Monitoring in **26** Trials

Safety in **15** Trials

Data Management in **18** Trials

Project Management in **21** Trials

Site Selection in **21** Trials

Medical Writing in **21** Trials

Statistics in **21** Trials



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QPS is a Global CRO
with locations around the world
to serve the evolving needs of the
Pharmaceutical and Biotech industries

