

A FLEXIBLE APPROACH TO ALZHEIMER'S DISEASE CLINICAL TRIALS

AND TO THE STATE OF THE STATE O

WHEN YOUR FOCUS IS CLINICAL TRIALS,



QPS EXPERTISE IN ALZHEIMERS RESEARCH

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

OPS IS A GLOBAL CRO WITH LOCATIONS AROUND THE WORLD

BENEFIT FROM THE RESOURCES OF A GLOBAL CRO



Whether your focus is small molecules, protein biotherapeutics, vaccines, gene therapy or cell therapy, QPS provides a full range of clinical trial services to support all drug development needs from discovery, through clinical development and regulatory filing.



IETWORK USA / Austria / Australia / Argentina / Belgium / Bosnia / Brazil / Bulgaria / Chile / China / Croatia / Czech Republic / Denmark/ France / Finland / Germany / OCATIONS Hungary / India / Italy / Lithuania / Netherlands / Poland / Romania / Serbia / Slovakia / South Africa / South Korea / Spain / Sweden / Taiwan / United Kingdom



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

Call +1 512 350 2827 Email infobd@qps.com