



A FLEXIBLE APPROACH TO PHASE II-IV CLINICAL SERVICES

QPS IS A GLOBAL CRO WITH DIVERSE CAPABILITIES

that offers end to end services to pharma and biopharma companies, enabling them to bring products to market in a faster and more compliant manner. QPS is ideally positioned to address the key global product development requirements of quality, compliance, and time to market.





PHASE II-IV CLINICAL SERVICES

QPS Services:

- ▶ Biostatistics
- ▶ Data Management
- ▶ Medical Writing
- ▶ Monitoring
- ▶ Pharmacogenomics (RNA/DNA)
- ▶ Pharmacovigilance
- ▶ PK/PD evaluation
- ▶ Program, Project and Study Management
- ▶ Regulatory Affairs
- ▶ Medical Affairs

Therapeutic Area Focus:

- ▶ CNS – including Parkinson's disease and Alzheimer's disease
- ▶ Endocrinology – including Diabetes and Obesity
- ▶ Oncology
- ▶ Pulmonology/Respiratory – including Asthma, Emphysema and COPD
- ▶ Dermatology – Dermal and Transdermal delivery systems
- ▶ Vaccines – Flu, RSV, Hepatitis, HPV, Ebola, Zika, and others

Why Choose QPS for Phase II-IV Services?

- ▶ Full service CRO providing the highest level of custom-built research
- ▶ Global footprint with 7 locations in Asia, Europe and North America
- ▶ Low turn-over of staff
- ▶ Monitoring efficiency increased by strategic site and study support
- ▶ Access to a large network of investigators across therapeutic areas
- ▶ Experienced personnel with medical, regulatory and cultural knowledge
- ▶ Direct access to scientists

QPS GLOBAL CLINICAL TRIAL FOOTPRINT

QPS India

- ▶ Extensive experience with generic and biosimilar studies
- ▶ Fast recruitment of patients

QPS Missouri

- ▶ All 5 units available for Phase II-IV trials
- ▶ PI Experience: Family practice, Oncology (dermal), Hepatitis C and Ophthalmology

QPS Netherlands

- ▶ Clinical Research Operations Services across European countries, including the UK and Scandinavia

QPS Taiwan

- ▶ Fast governmental regulatory approval to start studies
- ▶ Clinical Research Coordinators who can increase study efficiency

QPS Austria

- ▶ Clinical studies have involved over 15,000 subjects
- ▶ In-depth knowledge and understanding of European and local regulations

QPS Miami

- ▶ Over 1,200 clinical trials conducted across Phase I, Phase II-IV and Vaccines
- ▶ A robust database of 30,000 subjects/patients





QPS 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 bioanalytical services to support drug development needs from discovery, through clinical development and regulatory filing.



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