

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. With operations in India, USA, Europe and Taiwan, QPS is ideally positioned to address the key global generic product development requirements of quality, compliance, and time to market.

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

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



Services Offered

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

Therapeutic Area Focus

- CNS - including Parkinson's Disease and Alzheimer's Disease
- Endocrinology - including Diabetes and Obesity
- Oncology
- Pulmonary/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?

- Boutique style CRO providing the highest level of customized service.
- Global footprint with 11 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

The Global Footprint Of QPS

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, Skin Cancer & Hepatitis C Specialists, Ophthalmology

QPS Netherlands

- Capable of monitoring studies in several countries (Belgium, Northern Germany, NL, UK)
- Cardiology studies are well promoted in NL, with good recruitment rates

QPS Taiwan

- Fast governmental regulatory approval to start studies
- Clinical Research Coordinators with UNITIX, an SMO, 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 1, Late Phase and Vaccines
- A robust database of 30,000 subjects/patients

FLEXIBILITY. AGILITY. SPEED.

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with locations around the world
to serve the evolving needs of the
Pharmaceutical and Biotech industries

