QPS
CUSTOM-BUILT RESEARCH™

A flexible approach to

Early Phase Clinical Trials

WITH EIGHT PHASE I SITES ON THREE CONTINENTS, QPS offers comprehensive services from small-scale, complex and high-capacity studies in healthy volunteers, to Proof of Concept (POC) studies in patient or specialty populations for a wide variety of therapeutic indications.

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

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Professional and Experienced Staff

Each Phase I study will have a client dedicated team, headed by an experienced Clinical Project Manager, who has the overall responsibility for the conduct of your study and is your single point of contact throughout your Phase I study. As a full service unit, QPS can assist you with the entire early phase drug development process:

- Review of Preclinical Data
- Study Design and Protocol Writing
- Clinical Conduct
- Provision of CDISC-compliant Clinical Study Reports
- Data Management
- PK/PD Analysis
- Biostatistics
- Bioanalysis Assays
- Biomarker Assays

Global Flexibility and Capacity

At QPS, we realize that you face many challenges in strict time lines, patient enrollment, demanding clinical data collection and reporting requirements, and budgetary constraints. With our eight state-of-the-art and strategically located global Phase I facilities, we are in an excellent position to meet these needs. QPS provides high quality data, client-specified time lines and competitive pricing. Our sites also have access to large numbers of healthy and patient volunteers for all types of clinical studies.

Customer Focus

QPS focuses on meeting the needs of each client. We make certain all studies are recruited fully and completed on time. We also ensure optimal communications so you have always complete visibility into your project's status. You can rest assured that your deadlines will be met and your budgets will not be exceeded.

Preferred Provider Relationship

QPS' Phase I sites have unique and complimentary core capabilities with expertise that remains focused in order to gain maximal efficiency. This translates to improved quality and time management which in turn translates to enhanced cost effectiveness. QPS enables maximum benefit by leveraging the distinct features of our global Phase I sites to keep your overall early phase clinical development programs on track. The QPS Clinical Pharmacology Teams in the Netherlands, Taiwan, India and US will work to ensure your study is completed in line with all appropriate global standards. All QPS sites are connected through a global data network to simplify study management for sponsors, providing integrated information and perspective during the entire course of the drug development process.

QPS offers a full range of Phase 1 services including:

- First-in-Man Programs (SAD + MAD + FE + CYP450 Interaction)
- Clinical PK/PD studies
- Bioavailability studies
- Bioequivalence studies
- Drug Interaction studies
- Human Mass Balance studies
- Imaging (PET, fMRI) studies
- Thorough QT/QTc studies
- Microdosing studies
- Proof of Concept (PoC) across multiple disease indications
- Specialty Populations

QPS is a Global CRO with locations around the world