QPS Clinical Development Services

Proven scientific expertise and collaborative solutions from a nimble, specialty CRO to help you navigate the entire clinical development process

Building a Trusting Partnership with You Throughout the Clinical Development Process

Beginning with our industry-leading Phase I capabilities, our Clinical Development Team combines superior scientific expertise with a passion for building lasting relationships upon open communication and trust. From First-in-Man (FIM) and Proof of Concept (POC) through multicenter, multinational studies, you will gain peace of mind knowing we are meticulously managing every aspect of your drug program, providing you with high enrollment rates, industry-best timelines, high-quality data, and a competitive price.

Industry-Leading Capabilities to Maximize Your Clinical Success

When you partner with QPS, you gain access to industry-leading Phase I capabilities, featuring more than 500 beds on three continents. Our six strategically located Phase I facilities are staffed by expert clinical pharmacology teams that routinely conduct hundreds of Phase I/IIa studies annually. We also have scientific expertise in the design of all types of Phase I studies, as well as the interpretation of study data.

Additionally, each of our Phase I sites have unique capabilities that compliment each other to achieve maximum efficiency without overlap, ensuring your drug candidate moves seamlessly through clinical development. Taking advantage of our full portfolio of services enables your staff to focus more time on what they do best: developing new drugs.

QPS provides you with a full range of early stage clinical drug development services, including:

See how QPS' proven scientific expertise and flexible, collaborative solutions can help your therapy reach its maximum potential. Call to schedule your tour at one of QPS' worldwide locations today.

- Review of Preclinical Data
- Study Design and Protocol Writing
- Clinical Conduct
- Bioanalysis/Biomarker Assays
- Data Management
- Biostatistics
- PK/PD Analysis
- Provision of CDISC-compliant Clinical Study Reports
- First-in-Man Programs (SAD + MAD + FE + CYP450 Interaction)
- Clinical PK/PD Studies
- Bioavailability Studies
- Bioequivalence Studies
- Drug Interaction Studies
- 505(b)(2) NDA Studies
- Human Mass Balance Studies
- Microdosing Studies
- Imaging (PET, fMRI) Studies
- Vaccine Studies
- Thorough QT/QTc Studies

Specialty/Patient Study Experience to Meet Your Unique Needs

Our Phase I sites have access to large numbers of patients for all types of clinical studies across a wide variety of indications. We work independently or together with local university and general hospitals to complete POC studies in patient or specialty populations. In addition, we develop pharmacodynamic endpoints and laboratory biomarker assays that are critical for the evaluation of your early stage compounds.

Customized Clinical Development Solutions Provide Peace of Mind

When you continue your trusted relationship with QPS through further clinical development, we will work with you to create customized solutions that meet your unique needs. You will also have peace of mind knowing that QPS acts as a collaborative partner to address potential challenges, leading to decreased necessary oversight and increased cost savings.

Our experienced Clinical Development Teams operate from strategically located sites throughout Europe, the United States, and Asia Pacific. In every location, we focus on selecting the investigators who will best fit your study. Our staff undergoes regular, intensive training to stay up-to-date on the latest clinical and safety procedures and adheres to the highest quality clinical conduct according to ICH-GCP.

We offer a full range of clinical development services for the evaluation of compounds targeting a wide variety of therapeutic areas. Our services include:

- Program Management
- Project Management
- Site Management & Monitoring
- Data Management
- Biostatistics
- Medical Writing
- Temporary Staffing & Flexible Sourcing Solutions
- Patient Recruitment Services

Proven Scientific Expertise and Guidance with Open, Responsive Communication

At QPS, we work hard to ensure optimal communications. Each 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 remains your single point of contact throughout the project's duration. Clinical Project Managers are readily available to provide insight and guidance, and all QPS sites are connected through a global data network to simplify management of your study.

QPS helps you navigate

Clinical Development Centers

QPS helps you navigate

Company Values: Integrity - We make and keep promises and build trust through honesty, proactive communication and reliability - Quality - We bring the highest level of technical expertise and judgement to our work - Customer Focus - We tap our global resources to provide service that is fast, flexible and integrated - Commitment - We work hard to solve problems and deliver results - People - We treat people with dignity, respect and fairness, and embrace our differences - Culture - We are friendly and fun. We provide opportunities to grow, we value loyalty and teamwork, and we recognize and reward performance.

Time is of the essence in drug development, so contact a member of the QPS Business Development Team today and find out what QPS can do for you.

GLOBAL BD Office

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