A FLEXIBLE APPROACH TO PHASE II-IV CLINICAL TRIALS

QPS SUPPORTS DRUG DEVELOPMENT IN EVERY CLINICAL STAGE WITH A STRONG FOCUS on Phase II-IV clinical trials and medical device studies. QPS' Clinical teams bring years of experience catering to the unique needs of virtual, small, mid-size and large pharma, biotech and medical device companies.

QPS PHASE II-IV CLINICAL TRIALS OVERVIEW

With its site management & monitoring teams operating from 30 locations on three continents (Asia/Pacific, USA and Europe), QPS has become a new strong player in the space of late phase clinical research services.

Professional and Experienced Staff

Global Flexibility & Capacity

Customer Focus

Preferred Provider Relationships

KEY DIFFERENTIATORS

We combine scientific leadership with disciplined execution, global access to patients, and local and global regulatory expertise to accelerate your compound to approval.

- Full-time physicians provide strategic direction for study design and planning, train operations staff, work with primary investigators, provide medical monitoring, and meet with regulatory agencies as needed.
- Long-standing interactions with global key academic and industry opinion leaders.
- Close collaborations and ongoing programs with high recruiting sites at academic and independent centers.
- Our expertise as a full service late phase CRO provides us instrumental insight into the needs and challenges of the sites allowing to proactively manage and support them.
- Our unique operational platform allows us to assign the most experienced and therapeutically focused team members to your project.
- A proven study initiation process that supports and encourages team work, transparent communication, risk mitigation insight and individual ownership of expectations.

PROFESSIONAL AND EXPERIENCED STAFF

With QPS you get immediate access to our Global Clinical Development Teams who are capable of meticulously managing the conduct of your trial while working collaboratively with your development team.

From the first interaction we are there to provide insight, consultative services, protocol review, site feasibility and recommendations. We will evaluate your protocol to determine the right project team mix and experience necessary to successfully perform your trial within the designated time lines and within budget.

We have the regional coverage, regulatory prowess and medical support to ensure a smooth start-up and selection of the right sites. With offices throughout the world we are in constant contact with thought leaders and regulatory authorities to help ensure the success of your trial.

Our project management team offers a diverse therapeutic expertise as well as years of management experience. When a therapeutic need arises that is beyond our in-house expertise, we will proactively offer experienced outside consultants.

We believe you come to CROs for their experience, operational depth, in-house consultation, and geographic reach. We work hard to ensure that our clients are impressed with our teams on all levels which helps establish long lasting relationships.

THERAPEUTIC EXPERTISE

QPS has extensive experience across multiple Therapeutic Areas, including: Neurology, Pain, Inflammation, Vaccines, Dermatology, Gynecology, Urology, Diabetes, Metabolic Disorders, Infectious Disease, CNS, Respiratory, and Rare and Orphan Diseases.

QPS
CUSTOM•BUILT RESEARCH™

QPS IS A GLOBAL CRO WITH LOCATIONS AROUND THE WORLD

BENEFIT FROM THE WORLDWIDE RESOURCES THAT A GLOBAL BIOANALYTICAL CONTRACT RESEARCH ORGANIZATION BRINGS

QPS maintains four advanced bioanalytical facilities in the USA, Netherlands, Taiwan, and India, offering strategic solutions to companies with sites or trials, both in the US and overseas and/or wishing to complete studies in Asia and/or India.

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

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