A FLEXIBLE APPROACH TO EARLY PHASE CLINICAL TRIALS

WITH FOUR PHASE I SITES ON THREE CONTINENTS, QPS OFFERS comprehensive clinical trial services across all phases of development and a wide variety of therapeutic indications.

QPS EARLY PHASE CLINICAL TRIALS OVERVIEW

QPS is well known for its success in first-in-man (FIM) clinical trials. All QPS Phase I sites are staffed by expert clinical pharmacology teams that routinely conduct hundreds of Phase I/IIa trials annually. We also have scientific expertise in the design of all types of early phase clinical trials, including dose response, single ascending dose (SAD), multiple ascending dose (MAD), drug-drug interaction studies and more.

Professional and Experienced Staff

Global Flexibility & Capacity

Customer Focus

Preferred Provider Relationships

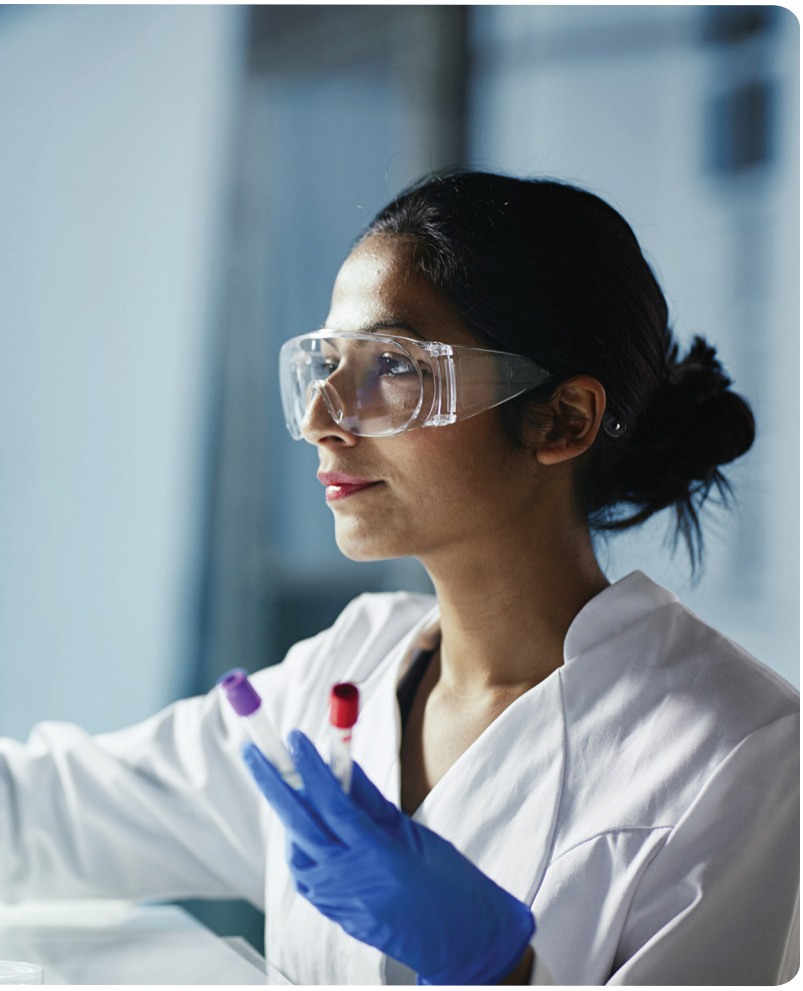
PROFESSIONAL AND EXPERIENCED STAFF

Each Phase I/II trial will have a dedicated client team, headed by an experienced Clinical Project Manager, who has the overall responsibility for the conduct of your trial and is your single point of contact throughout the trial.

As a full service CRO, QPS provides clinical research services across the entire early phase drug development process, including: 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 and Biomarker Assays.

FULL RANGE OF EARLY PHASE TRIALS

- First-in-Man & Proof of Concept
- Clinical PK/PD studies
- Bioavailability & Bioequivalence
- Drug-Drug Interaction
- Human Mass Balance
- Imaging (PET, fMRI)
- Thorough QT/QTc
- Microdosing
- Specialty Populations



GLOBAL FLEXIBILITY AND CAPACITY

At QPS, we realize that you face many challenges with strict timelines, patient enrollment, demanding clinical data collection and reporting requirements. With four state-of-the-art and strategically located global Phase I facilities, we are in an excellent position to meet these needs. QPS has access to large numbers of study subjects for many types of clinical trials.

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 always have complete visibility into your project's status. You can rest assured that your deadlines will be met.

GLOBAL CAPABILITIES

QPS' early phase clinical trial sites have unique and complimentary core capabilities with expertise that remains focused in order to gain maximum efficiency. The QPS Clinical Pharmacology Teams work to ensure each study is completed in line with all appropriate global standards.

All QPS sites are connected through a global data network to simplify study management. This ensures fully integrated information during the entire course of the drug development process.

SCIENTIFIC LEADERSHIP AND PROVEN RESULTS

Our dedicated, experienced team ensures that all clinical trials conducted in QPS facilities meet timelines and regulatory requirements. QPS provides high quality data along with direct access to our technical staff, regularly scheduled updates in a format that works for you, and prompt and courteous answers to your inquiries at a fair and competitive price.

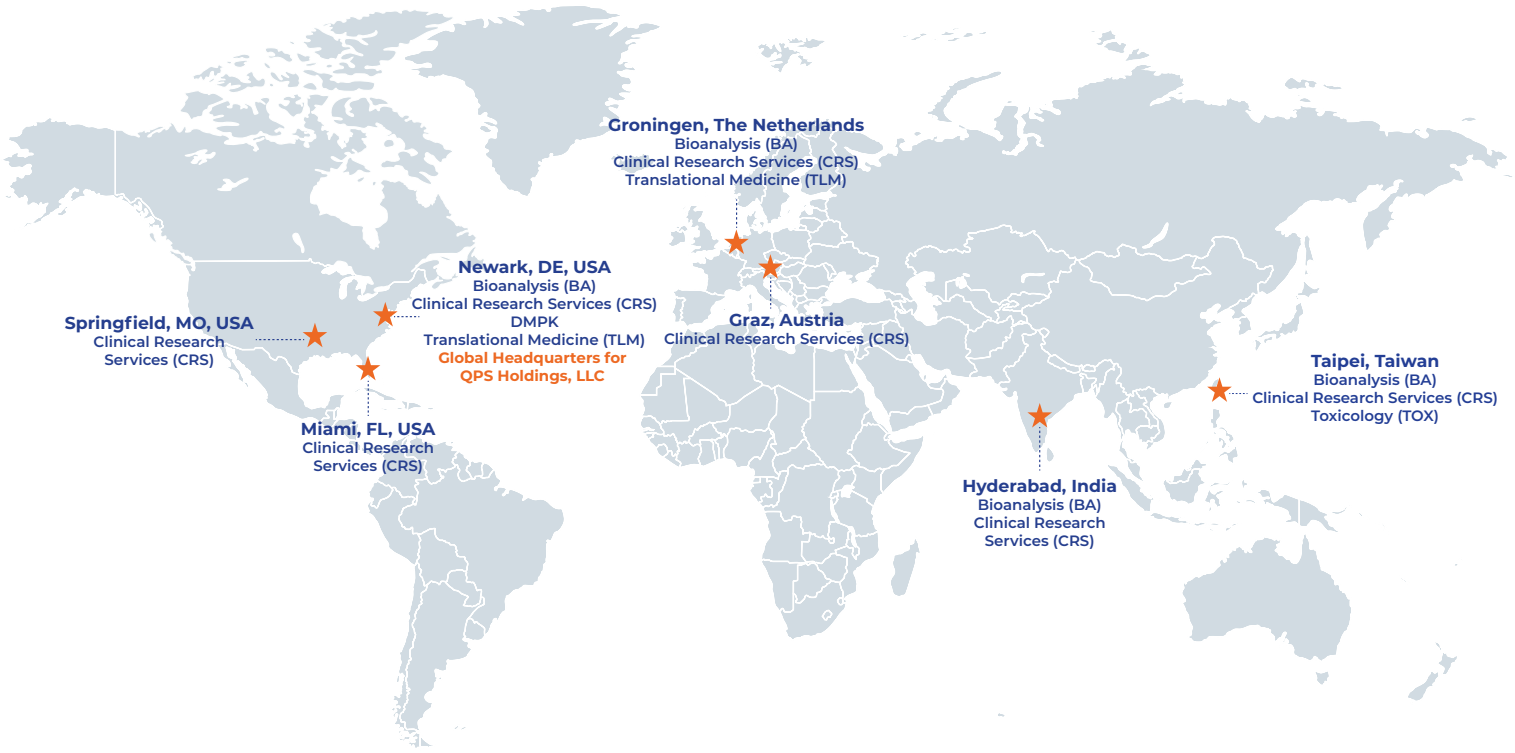
- Phase I
- Phase IIa
- Clinical Research Services
- Data Management

QPS
CUSTOM-BUILT RESEARCH™

QPS IS A GLOBAL CRO WITH LOCATIONS AROUND THE WORLD

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

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.



QPS
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