QPS
CUSTOM-BUILT RESEARCH™

A flexible approach to

IND-enabling Preclinical Studies

QPS IS A GLOBAL CRO WITH DIVERSE CAPABILITES THAT OFFERS END TO END DRUG DEVELOPMENT SERIVCES. Partnering with QPS for a well-conceived and executed ind-enabling preclinical program will provide you with a detailed assessment of your drug candidate and the most agile, flexible and timely pathway to filing an IND.

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

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

Why perform your IND-enabling preclinical studies at QPS?

Before we begin executing your IND-enabling preclinical program, you will receive strategic review and advice on the design and execution of your ADME and pharmacology-toxicology studies:

- Your proposed non-clinical plan including proof of concept studies, pharmacology and ADME studies, and toxicology/safety program will be analyzed in depth.
- Your proposed non-clinical plan will be reviewed to identify deficiencies and potential roadblocks and hurdles and whenever possible solutions identified.
- Timelines for preclinical development of your overall and individual programs will be mapped out and preclinical development objectives and crucial milestones will be confirmed.

During execution of your IND-enabling program you will benefit from QPS's operational strengths, strong scientific/regulatory pre-IND/IND support, and drug development experience.

Operational Strengths

- ADME scientists and toxicologists with extensive industry and CRO experience allow for optimal planning and execution of ADME and pharmacology-toxicology studies.
- State-of-the-art ADME, toxicology and bio-analytical facilities.
- Rapid execution and completion of all pre-clinical studies required for IND submission.
- All studies will be carefully monitored and every phase of the studies critically assessed for scientific rigor and quality.
- Fast turnaround on high-quality non-clinical study reports.
- Extensive experience in the preparation of ADME and pharmacology-toxicology sections of IND submissions.
- An experienced program manager will be assigned to ensure rigorous program oversight.

Scientific/Regulatory Pre-IND/IND Support

- ADME scientists and toxicologists with extensive industry and CRO experience allow for optimal planning and execution of ADME and pharmacology-toxicology studies.
- Review and gap analysis of available data & pre-clinical development plans.
- Advice on the design and timing of ADME, safety pharmacology, and toxicology studies.
- Provide expert advice on ADME and pharmacology-toxicology issues associated with a broad range of therapeutic areas.
- Rapid completion of the ADME and pharmacology-toxicology sections of the IND to enable client to file the IND in a timely manner.

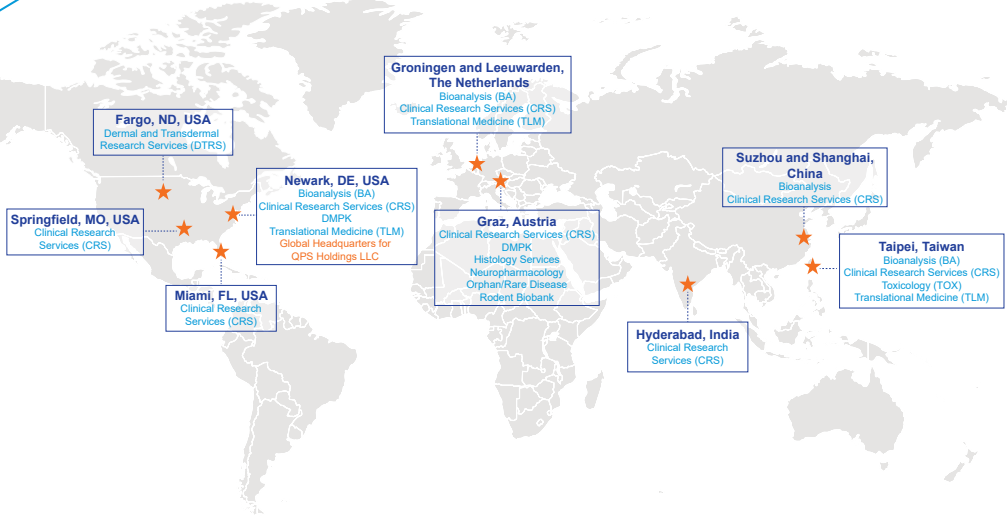
QPS
CUSTOM-BUILT RESEARCH™

IND-enabling Preclinical Studies

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

QPS is a Global CRO

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



WWW.QPS.COM