QPS
CUSTOM-BUILT RESEARCH™

QPS ONCOLOGY EXPERIENCE
FULL SERVICE CRO SUPPORT

QPS is a leading full-service global CRO with extensive experience conducting research, offering comprehensive services to ensure successful drug development. Our experienced and dedicated global team provides unmatched support, from preclinical studies to bioanalysis and all phases of clinical trials.

WHY CHOOSE QPS FOR ONCOLOGY STUDIES?

1. Expert Global Team

- **Specialized Expertise:** A dedicated framework of oncologists, clinical research coordinators, project managers, and data analysts.
- **Patient-Centric Approach:** Powerful associations with study sites across USA, Europe, and Asia and innovative patient recruitment strategies.

2. State-of-the-Art Facilities

- **Advanced Bioanalysis Laboratories:** Equipped with cutting-edge technology for precise method development and assay execution.

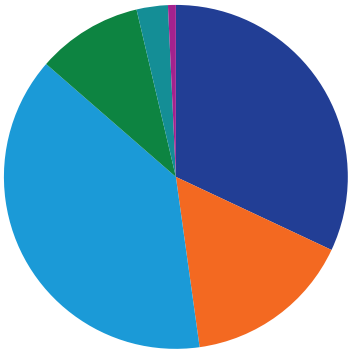
3. Comprehensive Services

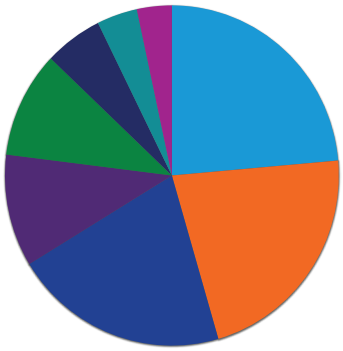
- **End-to-End Support:** From preclinical studies to bioanalysis, and protocol development to regulatory submissions, QPS handles every aspect of your oncology study.

4. Proven Track Record

- **Successful Trials:** QPS has successfully conducted a wide variety of Phase I - IV oncology studies around the world.
- **Broad Range of Cancers:** Including breast, head & neck, lung, and prostate cancer.

QPS CLINICAL EXPERIENCE IN ONCOLOGY BY PHASE AND INDICATION





CLINICAL RESEARCH OPERATIONS

- Project Management
- Patient Recruitment
- Clinical & Medical Monitoring
- Data Management & Biostatistics
- Medical Writing
- Site Selection & Feasability
- Regulatory & Medical Affairs
- Safety & Pharmacovigilance
- Quality Assurance

QPS is your ideal CRO partner for generic product development and can also assist in understanding the regulatory requirements of different countries for generic drug approval. QPS can partner with your product development groups in taking you from study design to the final dossier for regulatory submission. For more information on QPS, visit www.qps.com, or email infobd@qps.com.

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QPS IS A GENERIC DRUG DEVELOPMENT LEADER

QPS has the experience and expertise to work with Sponsors in the global development of a diverse range of products from conventional generics to more complex products including modified release formulations, novel drug delivery systems for NCE-1, ANDA submissions.

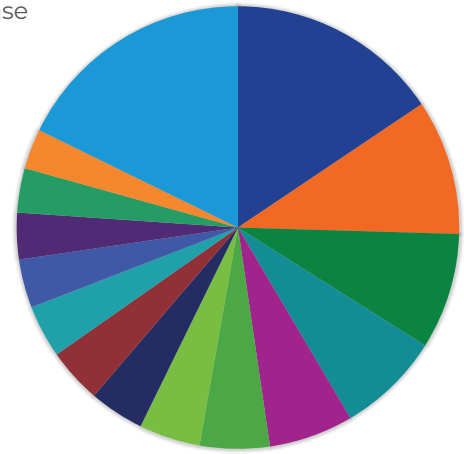
With over 500 beds worldwide, our Clinical Phase I units provide extensive capacity, including 138 beds at QPS India.

Generic drug study design experience

- 138 beds and over 2,000 Phase I-IV studies
- Over 23,000 potential subjects in the database
- BA/BE & PK/PD in healthy volunteers
- BA/BE in patient populations
- Clinical Endpoint Studies
- 505(b)2 NDA submissions

Our comphrehensive services include:

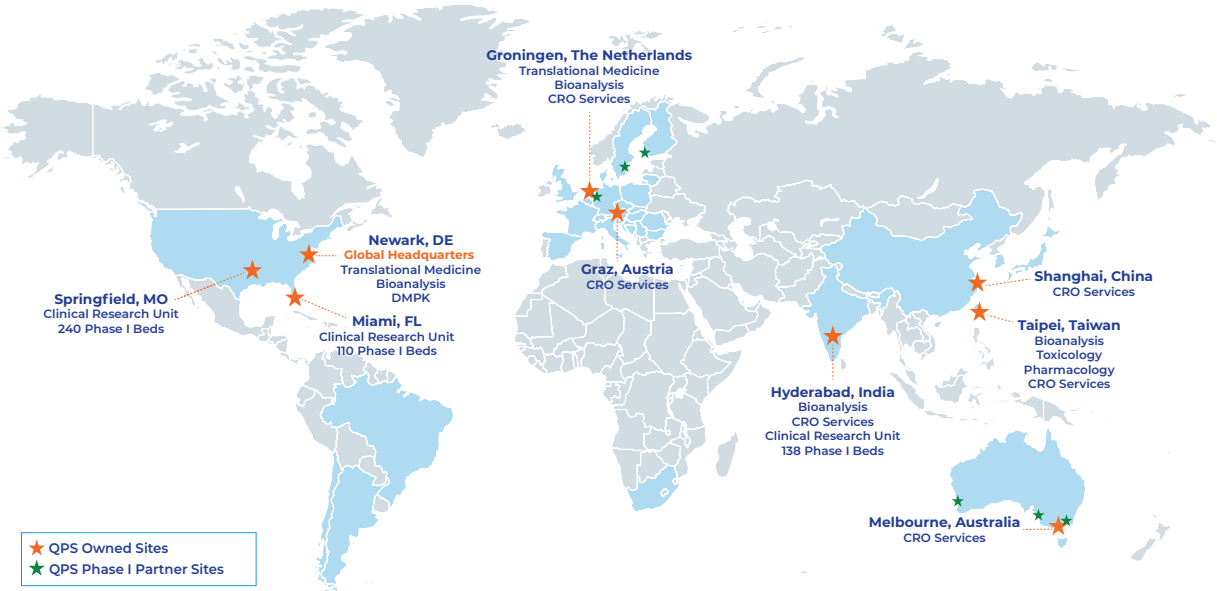
- Study Design
- Medical Writing
- Clinical Study Conduct
- Bioanalysis
- Data Management & CDISC
- PK & Statistical Analysis



Generic Drug Studies

- Antidiabetic (101)
- Tyrosine kinase inhibitor (64)
- Cholinergic (55)
- Antiviral (49)
- Anticonvulsant (40)
- Antihistamines (33)
- Antiretroviral (29)
- Antibiotic (26)
- Antihypertensive (26)
- Proton Pump Inhibitor (25)
- Antidepressant (23)
- Analgesic (22)
- Steriods (21)
- Antipsychotic (19)
- All Other (115)

QPS CLINICAL TRIAL SITES AND NETWORK LOCATIONS



NETWORK LOCATIONS: USA / Austria / Australia / Argentina / Belgium / Bosnia / Brazil / Bulgaria / Chile / China / Croatia / Czech Republic / Denmark/ France / Finland/ Germany / Hungary / India / Italy / Lithuania / Netherlands / Poland / Romania / Serbia / Slovakia / South Africa / South Korea / Spain / Sweden / Taiwan / United Kingdom

Discover how QPS can accelerate your clinical research.

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