QPS
CUSTOM-BUILT RESEARCH™

EARLY-PHASE ONCOLOGY DEVELOPMENT WITH TRANSLATIONAL AND BIOANALYTICAL EXPERTISE

QPS is a leading full-service global CRO with an experienced and dedicated global team providing comprehensive services from preclinical studies to bioanalysis and all phases of clinical trials. QPS supports oncology sponsors from first-in-human through proof-of-concept by combining early-phase clinical expertise, bioanalytical support, and biomarker and PK/PD-driven development.

WHY CHOOSE QPS FOR ONCOLOGY STUDIES?

1. Oncology Drug Development is Complex

- First-in-human and dose escalation requires careful safety assessment and dose selection
- Biomarker strategies are critical for patient selection and demonstrating mechanism of action
- PK/PD and translational endpoints are essential for understanding exposure/response relationships and guiding decisions
- Recruitment of targeted patient populations can impact study timelines

2. Early-Phase Oncology Studies

- First-in-human (FIH)
- Dose escalation and dose expansion
- PK/PD and safety evaluation
- Patient-based studies and healthy volunteer studies

3. Global Bioanalytical Laboratories

- PK/PD analysis
- Biomarker integration

4. Flexible Access to Oncology Expertise

- A global network of investigators and key opinion leaders
- Clinical and scientific teams supported by experienced oncology specialists

5. Global Study Execution

- Flexible model tailored to study and sponsor needs
- Clinical operations across North America, Europe, and Asia
- Coordinated execution from early-phase through later-stage studies
- Experience supporting multinational oncology studies across regions

6. Oncology Experience

- Studies across solid tumours and other malignancies
- Small molecules, biologics, and ATMPs
- Support for early- and late-phase clinical trials

CLINICAL RESEARCH OPERATIONS

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

Since 1995, QPS has provided discovery, preclinical, and clinical drug development services. An award winning leader focused on bioanalytics and clinical trials, QPS is known for proven quality standards, technical expertise, a flexible approach to research, client satisfaction and turnkey laboratories and facilities. For more information on QPS, visit www.qps.com, or email infobd@qps.com.

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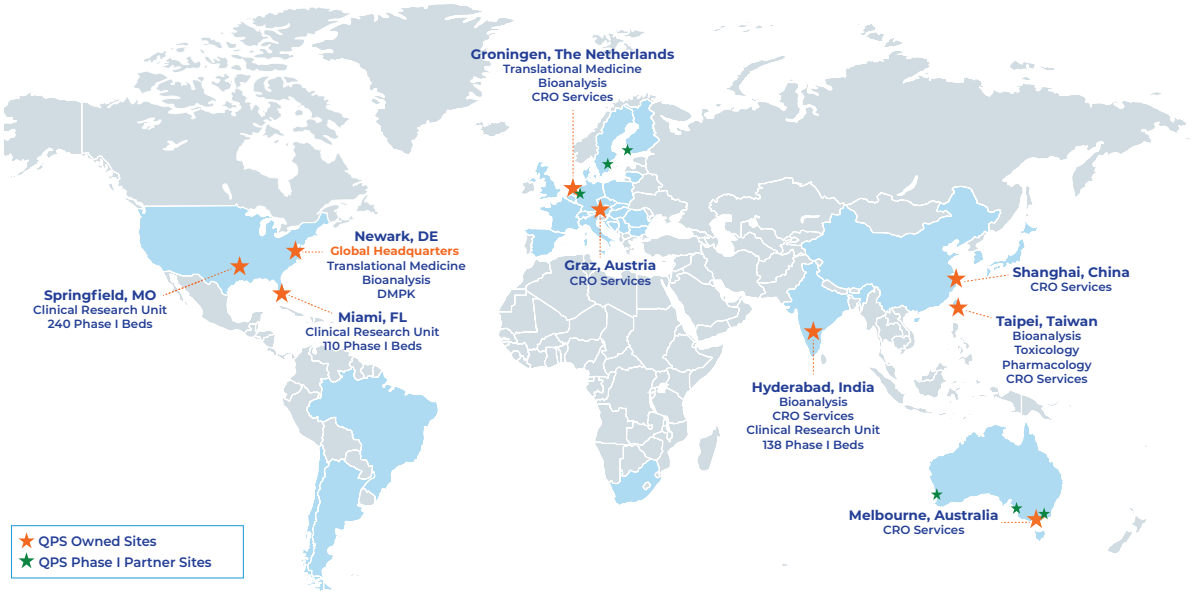
ONCOLOGY KEY OPINION LEADER (KOL) NETWORK

Europe
- Dr. Mindaugas Jievaltas, Urologist | University Hospital Kaunas/Clinic of Modern Oncology | Lithuania
- Dr. Vincas Urbonas, Medical Oncologist | Early Phase Clinical Trials Lead | National Cancer Institute | Lithuania
- Christina Junvik, Masters Biopharmaceutical Sciences | Early Drug Development in Oncology | Sweden

USA
- Dr. Bruno Fang, Oncologist | Director of Clinical Research at Astera Cancer Care | New Jersey
- Dr. Gordon Brown, Uro-Oncologist | Summit Health/New Jersey Urology | New Jersey
- Dr. Raoul Concepcion, Uro-Oncologist | US Urology Chief Medical Officer | Tennessee

Taiwan
- Dr. Yen-Hwa Chang, Uro-Oncologist | Taipei Veterans General Hospital | Taiwan
- Dr. Peng-Hui Wang Director, Obstetrics and Gynecology | Taipei Veterans General Hospital | Taiwan
- Dr. See-Tong Pang, Uro-Oncologist | Chang Gung Memorial Hospital (CGMH) | Taiwan

ONCOLOGY CLINICAL TRIALS SITE 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

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