QPS
CUSTOM-BUILT RESEARCH™

QPS OBESITY 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 OBESITY STUDIES?

1. Expert Global Team

- **Specialized Expertise:** Our team includes board-certified obesity medicine physicians, experienced clinical research coordinators, project managers, and skilled data analysts specializing in obesity.
- **Patient-Centric Approach:** We prioritize patient comfort and compliance, ensuring high retention rates and reliable data collection.

2. Phase I State-of-the-Art Facilities

- **Over 500 Phase I beds:** Multiple clinics across 3 countries accommodate short- and long-stay trials.
- **On-Site Pharmacy:** Facilitates seamless medication management and ensures adherence to study protocols.

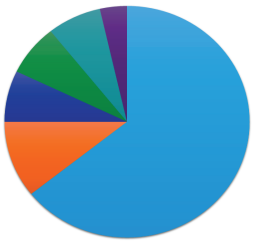
3. Comprehensive Services

- **Advanced Research Centers:** Equipped with cutting-edge technology for precise diagnostics and monitoring.
- **End-to-End Support:** From protocol development to regulatory submissions, QPS handles every aspect of your obesity study.
- **Customized Solutions:** Tailored study designs to meet the specific needs of obesity research, including biomarker analysis and nutritionists on staff.

4. Proven Track Record

- **Successful Trials:** Over 28 successful obesity trials since 2015, including multiple GLP-1, GIP, and GIP/GLP-1 studies.

QPS CLINICAL EXPERIENCE IN OBESITY BY CLASS OF DRUG



QPS Has Conducted 28 GLP-1 Studies

- GLP-1/GLP-1R/GIP/GIPR Agonists (18=64%)
- GPR40/GPR119 Agonists (3=11%)
- Peptide YY (PYY) Analog (2=7%)
- Triple agonist for GIP/GLP-1/GCG (2=7%)
- Non Peptide/Small Molecule GLP-1R Agonist (2=7%)
- GIPR, GLP-1R, and GcgR Activators (1=4%)

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

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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A GLOBAL SERVICE CRO

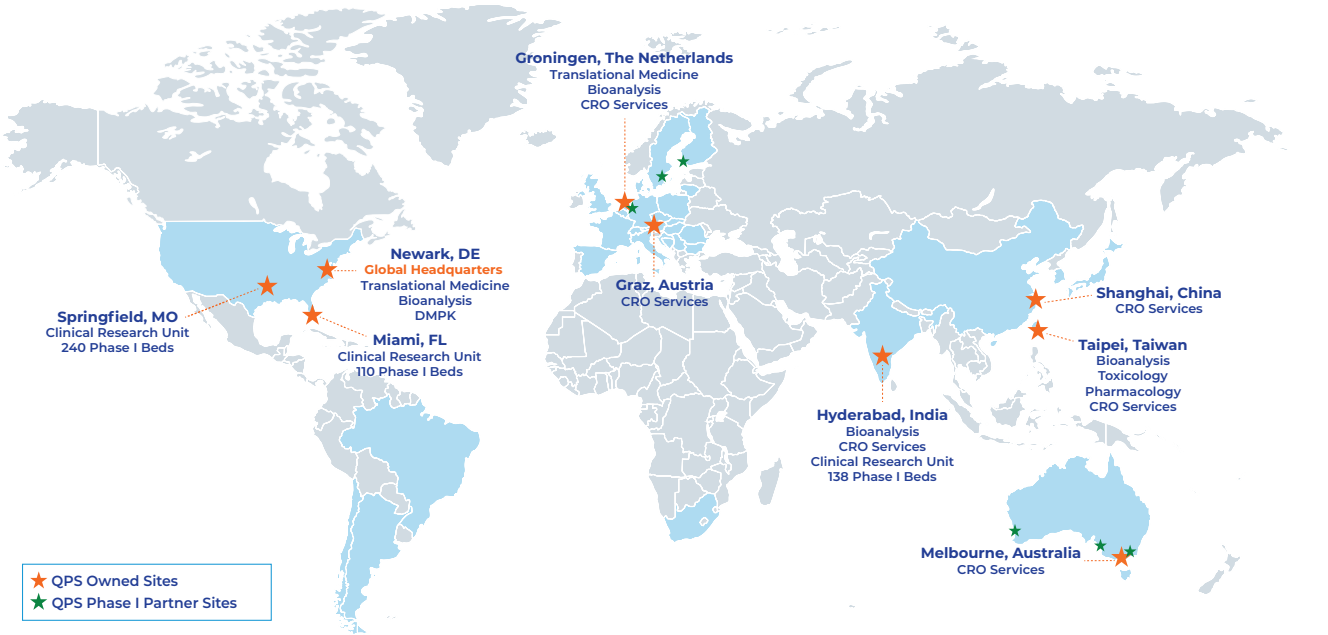
Drug Development Services

- Pharmacology
- Toxicology
- DMPK
- Bioanalysis
- Translational Medicine
- Clinical Trial Units
- Clinical Research Services
- Clinical Trial Kits
- Central Lab Services
- PBMC and DEXA Scanning Services

Clinical Research Services

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

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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