QPS
CUSTOM-BUILT RESEARCH™

# A GLOBAL LEADER IN **BIOANALYSIS**, **PRECLINICAL** AND **CLINICAL RESEARCH SERVICES**

**QPS is a Global Full Service CRO across USA, Europe and Asia offering Phase I-IV Drug Development Services**

## FULL SERVICE GLOBAL CRO

- Pharmacology
- Toxicology
- DMPK
- Bioanalysis
- Translational Medicine
- Clinical Trial Conduct
- Clinical Research Services
- Cell Therapy and PBMC
- Clinical Trial Kits

## 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

## THERAPEUTIC AREAS OF EXPERTISE

- Dermatology
- Endocrinology
- Neuropsychiatry
- Obesity
- Oncology
- Rheumatology
- NASH/NAFLD
- Vaccines
- Women's Health

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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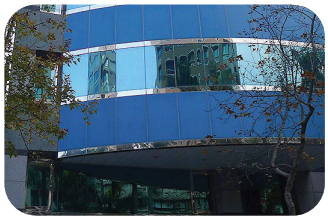
# EARLY PHASE CLINICAL SITES

## Miami

- 2 Clinical Units with 106 Phase I beds
- Over 3,000 studies completed, including POC, FIM, SAD, MAD, and specialty populations
- On-site board-certified physicians
- Database of over 35,000 potential study patients
- Excellent subject retention rates of over 95%
- Extensive experience in vaccines, diabetes, obesity, cardiology, RA, NASH/NAFLD, and more
- Capabilities for over 65 consecutive inpatient nights
- 15 FDA inspections
- Certified PBMC lab and technicians onsite

## Missouri

- 6 Clinical Units with 240 Phase I beds
- Over 2,000 studies completed including POC, FIM, SAD, MAD, and specialty populations
- On-site board-certified physicians
- Database of 46,000 potential study subjects
- 31 FDA inspections
- On-site pharmacy, negative pressure room, and safety & central labs
- Certified PBMC lab and technicians on-site
- Leukopak and cell therapy facilities
- Clinical trial kit production and central lab kit logistics



## Taiwan

- Full-service CRO offerings, serving global clients across Phase I-IV studies
- Clinical trial conduct, IND packages, site selection and monitoring, project management, data management, medical and regulatory affairs, biostatistics, and report writing
- Over 365 studies completed, including POC, FIM, SAD, MAD, and specialty populations
- Extensive experience in oncology, diabetes, RA, cardiology, and more
- 3 FDA inspections
- Toxicology and Pharmacology studies
- Bioanalysis and Sample Analysis

## India

- 4 clinical units with 138 actively monitored beds
- Over 2,000 Phase I-IV studies
- Database of over 23,000 potential study subjects
- On-site sample collection/processing, pharmacy and safety lab
- BA/BE & PK/PD studies in healthy volunteers (DDI, dermal, inhalation, PM women, biosimilars and long acting injectables)
- PK/PD & clinical endpoint studies in patients (oncology, CNS, endocrinology, pulmonology, ophthalmology)
- 15 FDA, 1 ANVISA, 3 WHO, 2 MHRA, 1 EMA, I NPRA & 2 GCC successful inspections
- Bioanalytical MD/MV and Sample Analysis

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