Celerion is the largest global early clinical research provider, with three facilities and over 600 beds across North America and Europe. Our state-of-the-art clinic in Phoenix, Arizona offers comprehensive clinical capabilities and a full clinical laboratory. With 315 beds (including 65 intensive care beds) the clinic has the largest early phase clinical-bed capacities in the Western United States.

Global Clinical Research – Phoenix, AZ

Cardiovascular Safety Services
Extensive experience and innovative solutions enable Celerion to deliver high-quality and cost-effective QT assessment.

- Celerion is a market leader in cardiovascular safety services, with over 60 Thorough QT (TQT) and over 400 ECG intense studies conducted over the last five years, in group sizes of up to 65 participants
- Optimized cardiac safety data collection resulting in lower variability—participant QTc standard deviation of 6msec versus industry average of approximately 10msec, enabling smaller group sizes and therefore, lower cost
- The first CRO with a Highly Automated ECG Core Lab system utilizing Bluetooth-enabled Holter monitors and highly automated ECG review providing clients faster access to high-quality data at a lower cost

Platelet Aggregation
Celerion has achieved a major milestone of generating over 25,000 platelet aggregation sample results. Platelet function assays are inherently variable, so a high level of experience is critical to produce reliable and accurate platelet aggregation results.

- Broad experience with several agonists including thrombin, TRAP, ADP (in multiple concentrations), arachidonic acid, collagen and epinephrine
- Clients benefit from higher quality data, reduced timelines and lower costs

EXPERIENCE & FLEXIBILITY
Celerion’s facility in Phoenix has been in operation over 30 years. The senior leadership team average over 10 years of clinical research experience, while dedicated investigators average 12 years.

This extensive knowledge and experience has resulted in the development of creative solutions for client programs to optimize success.

- More than 300 studies completed in the last five years: greater than 50 percent were PK, bioavailability, drug-drug interaction and First-in-Human (FIH)
- Case Report Forms data entry within 48 hours of collection: using ClinQuick®, paper and EDC systems
CULTURE OF QUALITY

Celerion’s quality and compliance systems, processes and personnel have instilled a culture of quality inherent in every program, with real-time quality control to enable delivery of high-quality, accurate data. Global Standard Operating Procedures (SOPs) and processes ensure consistency for all studies conducted across all sites.

Ophthalmic Services

Proven record and experience in conducting ophthalmic studies, involving over 800 participants, enables Celerion to efficiently run studies that save time and cost.

Our on-site ophthalmic suite is equipped for collecting refraction, biomicroscopy, fundus examination, Intra-ocular Pressure (IOP) (Goldmann applanation and airpuff), pupil size assessment, best-corrected visual acuity, macroscopic hyperemia, PK tear collection data, visual field and pachymetry.

Technology for Data You Can Trust

ClinQuick®, used in all Celerion clinics, is a proprietary Phase I clinical study management system with electronic data acquisition as its core function. ClinQuick assures consistency of clinical operations and data collection across sites, giving access to accurate high-quality information. This information enables faster go/no-go decisions based on accurate data and reduces time to database lock. Preprogrammed alerts ensure all events occur according to protocol, thereby reducing potential error and increasing participant safety.