



A CRO built differently for early phase certainty

We help biotech and pharmaceutical companies meet study timelines through disciplined, early phase execution.

15

Average years of leadership tenure

97 %

Of studies achieve or surpass recruitment goals

1,000+

Clinical trials conducted by our Investigators

100

Bed purpose-built Phase I unit

Specialized Studies

- ✓ First-in-Human
- ✓ Drug-Drug Interaction
- ✓ Early QT/TQT
- ✓ Food Effect
- ✓ PK/PD
- ✓ Renal Impairment
- ✓ Hepatic Impairment
- ✓ Human Abuse Potential
- ✓ Tobacco Harm Reduction

Therapeutic Areas

- ✓ CNS, Psychiatry & Neurology
- ✓ Substance Use Disorders & Addiction
- ✓ Allergy & Immunology
- ✓ Cardiometabolic & Obesity
- ✓ Pain
- ✓ Respiratory
- ✓ Dermatology
- ✓ Infectious Diseases
- ✓ Vaccines



What our sponsors are saying

In our experience, this was the first time that we have ever engaged an organization of their caliber and if we ever have the need for a Phase I CRO we would only think of DVCR.

Executive Director of
Clinical Pharmacology

Full-Service CRO Solutions



Project Management

Experienced Project Managers serve as your dedicated guide and single point of contact from study award through archive.



Data Management

Delivering efficient, quality-driven solutions that transform complex clinical data into trusted information, streamline timelines and maintain data integrity.



Biostatistics & Programming

From study design consultation to submission-ready CDISC packages, our programmers collaborate to deliver peer-reviewed, validated outputs.



PK/PD Analysis

Our experienced PK scientists support your study from start to finish across the full spectrum of clinical pharmacology.



PK Modeling & Simulation

Turn pharmacokinetic and pharmacodynamic data into quantitative evidence for dose selection, study design, go/no-go decisions, labeling and regulatory strategy.



Clinical Monitoring

Experienced CRAs conduct both blinded and unblinded monitoring, operate independently of the study team and are presented for your approval before assignment.



Medical Monitoring

A dedicated Medical Monitor supports your study from protocol synopsis through the CSR, guided by a tailored monitoring plan.



Medical Writing

We prepare key study documents on structured timelines using DVCR or sponsor templates.

Our Phase I Research Campus

PK Processing and Safety Laboratory

Our CLIA certified in-house laboratory performs chemistry, hematology, coagulation, urinalysis and other testing operating under established quality systems, validated testing procedures and applicable quality-control and proficiency-testing standards.

cGMP Pharmacy

A 2,500 square-foot cGMP-compliant compounding pharmacy inside our Phase I unit, DEA-registered to handle and store Schedule I-V controlled substances.

Contact Us to Learn More



+1 913-333-3000

