





Pharmacy Capabilities

At Worldwide Clinical Trials (Worldwide), we are committed to delivering excellence in clinical research through our state-of-the-art pharmacy services. Our cGMP Phase I on-site pharmacy, coupled with Investigation Drug Services located within our Clinical Pharmacology Unit, is designed to meet the highest industry and regulatory standards, ensuring the utmost security and confidentiality of your study trial and investigational products. With a team of highly experienced clinical pharmacists and certified pharmacy technicians, we provide a comprehensive range of services tailored to support your clinical trials. Our on-site pharmacy is equipped with advanced compounding equipment and sophisticated environmental monitoring systems, guaranteeing precise and reliable dose preparation.













State-of-the-Art Facilities

-  **On-site cGMP Phase I Compounding Pharmacy**
Fully dedicated to clinical trials with established processes and procedures.
-  **Dedicated Dose Preparation Areas**
Over 3,100 sq. ft. with HEPA filtration, segregated IP storage, and secure access.
-  **Sterile and Non-Sterile Compounding Suite**
ISO Class 7 clean room and ISO Class 5 laminar flow unit for sterile compounding.
-  **Radiolabel Compounding Suite**
Complies with USP Chapter (800).



Comprehensive Services

-  **Oral Dose Dispensation**
-  **Non-Sterile Dose Compounding**
Per USP Chapter (795).
-  **Sterile Dose Compounding**
Per USP Chapter (797).
-  **Radiologicals**
-  **Retention Sampling**
-  **Handling of Controlled Substances**
Experience with Schedule I-V substances following FDA, cGMP, GCP and DEA Guidelines.
-  **Temperature-Controlled Storage**
 -  Room Temperature: 20°C to 25°C
 -  Refrigeration: 2°C to 8°C
 -  Frozen: -20°C to -80°C



Commitment to Excellence

- ✓ **FDA Compliance**
Adherence to cGMP standards for Phase I investigational drugs since 2010.
- ✓ **Secure Access**
Security monitoring with cameras, limited access card ID system, and locked drug storage areas.
- ✓ **Comprehensive Training**
Includes didactic, observation, mentoring, and QC oversight for all personnel.
- ✓ **Mock Batch Procedures**
Ensures detailed and properly sequenced processes before finalizing the Batch Record Template.
- ✓ **Detailed SOPs and Manuals**
Provides meticulous protocols for equipment and processes.
- ✓ **In-Depth and Consistent Documentation**
Guarantees quality study documentation through well-defined procedures.
- ✓ **Methodological Rigor**
Dedicated to upholding the highest standards of quality data and subject safety.
- ✓ **Environmental Monitoring**



Why Partner with Worldwide Clinical Trials?

At Worldwide, we have an unwavering commitment to methodological rigor, quality data, subject safety, and personalized attention. With a proven track record spanning more than 17 years, we have successfully conducted a wide range of clinical pharmacology studies. Our integrated services fully support early and late-stage development under one set of SOPs, ensuring seamless and efficient operations. Additionally, we leverage advanced technology, including a fully validated eSource data collection system, to provide efficient and accurate data management, making us a trusted partner for your clinical research needs.



Worldwide
Clinical Trials

About Worldwide Clinical Trials

Worldwide Clinical Trials (Worldwide) is a global CRO serving development-driven biopharmaceutical companies, with more than 4,400 professionals operating across more than 70 countries. The company delivers therapeutically dedicated expertise in neuroscience, oncology, rare disease, and internal medicine, with comprehensive support across every development phase – from early-stage and first-in-human studies through Phase III registration trials.

The company's flexible service model – spanning full-service trial management to functional service partnerships through Worldwide Flex – is powered by a people-first, partnership-driven outsourcing approach that strengthens collaboration, enhances data transparency, and supports more informed decision-making. This provides sponsors with tailored, adaptable solutions that keep pace with the evolving demands of clinical research.

Learn more at www.worldwide.com