CLINICAL TRIALS: FASTER, EASIER, PROACTIVE

Planning While in Preclinical

Why wait? We begin planning your clinical study while your preclinical safety assessment is ongoing to start your first-in-human trials sooner.

1) PROACTIVE PLANNING & DESIGN

Preparing During Regulatory Review

While your IND/CTA is undergoing review, we're preparing logistics to ensure quick study start-up.

Synchronized Manufacturing

- Clinic-ready formulation for your small molecule drug product
- Dosage or formulation updates handled on the fly

2) PROMPT STUDY START

Starting Clinical Activities Swiftly

We start your first-in-human trials as soon as you receive regulatory approval.

3) FLEXIBLE STUDY CONDUCT

Efficient Recruiting

- 400K+ participant database
- Access to patient and special population from partnership with nearby hospitals
- In-house, multilingual recruiting staff

95% Retention Rate

- Comfortable, modern facilities
- 500+ beds in three units
- Bed-side entertainment and game rooms
- High-speed Wi-Fi
- Catered meals

Timed Bioanalysis

Quick and informed dosing decisions throughout your trial.

Up to 25-40% Time-savings

From preclinical to clinical, and beyond, we save you time and money through proactive planning and parallel processing your drug development.

REDUCED HAND-OFFS AND SEAMLESS TRANSITIONS BETWEEN MILESTONES WITH A SINGLE CRO/CDMO PARTNER