



# Dallas Clinical Research Unit

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The Labcorp Clinical Research Unit (CRU) in Dallas is a top-tier facility with more than 55,000 square feet of space. The CRU occupies over four floors and includes a cGMP pharmacy, dedicated ophthalmology procedure rooms, plus dedicated screening and outpatient visit areas. The cGMP pharmacy includes three sterile and two non-sterile manufacturing suites, which can manage and handle high-potency and hazardous drugs. An advanced telemetry system enables the team to perform continuous high quality data monitoring, enabling staff to better monitor study participants and quickly make necessary adjustments to treatment regimens.

With more than 25 years of research experience, the Dallas site has conducted over 700 inpatient and outpatient Phase I-IV clinical trials of investigational medications. In addition to healthy normal populations, we have experience in special populations such as type 1 and 2 diabetes, healthy elderly, high cholesterol, obese and postmenopausal women.

As the ninth largest city in the U.S. and the fourth largest metropolitan area, Dallas offers a tremendous opportunity for recruiting study participants and experienced staff. Our extensive database, which contains more than 56,000 active volunteers, has a unique racial distribution: 30 percent Caucasian, 30 percent African-American and 30 percent Hispanic.

## Site Features

- Total capacity – 100 beds
- Participant database – 56,802
- Longest continuous in-house stay – 38 days
- Emergency response time – < 5 minutes
- Fibrosan 530 and TransAir 3 PFT System capable of dynamic static spirometry, DLCO, lung volumes via nitrogen recovery, and more
- Dedicated ophthalmology suite
- cGMP pharmacy, including three sterile and two non-sterile manufacturing suites



### Phase I cGMP manufacturing potential

**25-50%**

drug production  
costs savings

significant when total  
manufacturing costs for  
drug development are

**20-40%**

## A GMP Pharmacy Case Study

A mid-sized drug manufacturer was in Phase I clinical trials for a molecule designed to treat adenoviral infection. Their contract manufacturing organization (CMO) predicted a delay in drug production leading to a three- to six-month delay in study start-up. The client inquired if Labcorp could manufacture the drug on time and manage the clinical trial with their supplied active pharmaceutical ingredient (API). This sponsor needed fast turnaround to get the study on track and on budget, with cGMP quality plus flexibility to adjust the formulation if needed.

### Understanding the Challenge

- cGMP quality for patient safety
- Fast turnaround to meet study timelines
- Flexibility for adaptive trial design
- Control over costs and decision-making

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### Game-Changing Solution: cGMP Manufacturing and Clinical Trial on One Campus

Labcorp proposed that our cGMP Pharmacy manufacture a small quantity of the client's formulation for the Phase I clinical trial. Their auditors were assured of quality and oversight by reviewing our documented cGMP process. On-site manufacturing ensured drug quality and patient safety plus fast delivery – a concern when some formulations may be stable for only minutes or hours.

With the pharmacy and trial site located on one campus, our team could quickly deliver doses from manufacture to clinic. A large CMO would add distance and shipping time between manufacture of the product and delivery to the clinic. Labcorp removed the time and distance gap between pharmacy and trial site for fast turnaround.

The client also gained flexibility to adjust their formulation on the fly based on observed trial results. A CMO could only deliver one formulation and one dosage, with the expectation of a large run. Our adaptive trial design provided for formula or dosage adjustment in response to trial indications – with less waste of API.

With cGMP manufacturing, the client gained control and cost savings during the trial. Instead of one large CMO run of one formulation, Labcorp delivered cost-efficiency for small runs, with the ability to change or halt the process without delay or material loss.

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**The client met their study timeline, with quality cGMP drugs produced in small runs for flexible formulation, expedient regulatory review, controlled costs and informed decision-making.**