

Accelerate & Diversify Your Vaccine Program

Count on Alliance Clinical to accelerate your vaccine program—even as we diversify your participant profile. With a robust database of prescreened participants across seven fully integrated sites, we stand ready to meet your enrollment timelines and key study milestones.



50 vaccine studies
in 5 years



700+ prescreens
per month



280,000+ database
of consented,
prescreened
participants



85% of database
from
underrepresented
populations



>20% of database
65+

Proven Expertise in a Broad Range of Vaccine Areas

COVID-19

Respiratory Syncytial
Virus (RSV)

Influenza

Human
Papillomavirus
(HPV)

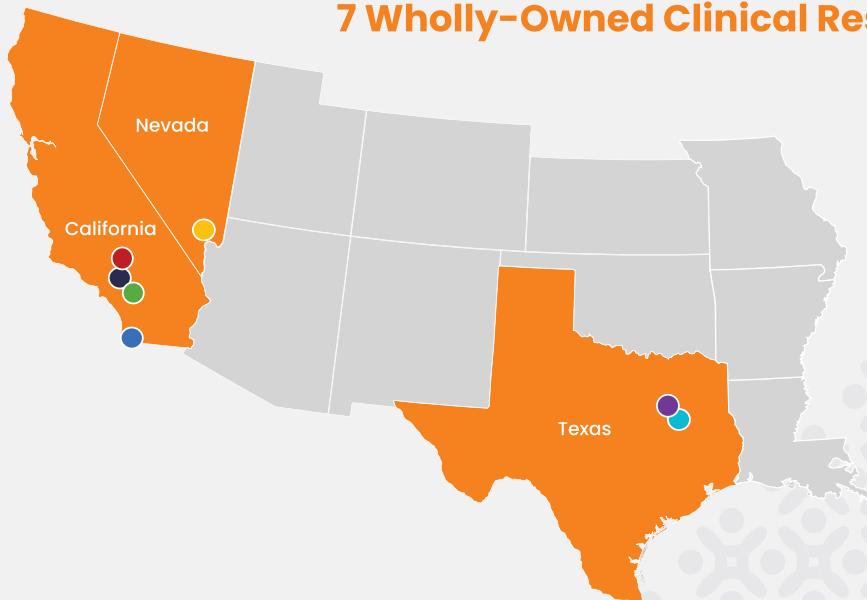
Human
Immunodeficiency
Virus (HIV)

Herpes Simplex
Virus - 2

Cytomegalovirus
(CMV)

Urinary Tract
Infections (UTI)

7 Wholly-Owned Clinical Research Sites



- **Alliance Clinical West Hills**
West Hills, CA
- **Alliance Clinical Canoga Park**
Canoga Park, CA
- **Alliance Clinical Los Angeles**
Inglewood, CA
- **Alliance Clinical San Diego**
San Diego, CA
- **Alliance Clinical Las Vegas**
Las Vegas, NV
- **Alliance Clinical Lewisville**
Lewisville, TX
- **Alliance Clinical Dallas**
Dallas, TX

Three Keys to a Successful Vaccine Study

Rapid Study Startup. Audit-Ready Data. Diverse Participant Population.



Rapid Study Startup Enables Brisk Enrollment

We begin with a proprietary database of 280,000+ prescreened participants. Harnessing technology, we then quickly identify and contact appropriate candidates across our sites, proactively scheduling them in anticipation of site activation. This accelerates time to FPI.

In a recent vaccine study, we contributed 1,506 participants in just 2 months, making Alliance Clinical the third-highest enroller globally; all participants were from just one of our eight sites.



Comprehensive Focus on Quality Produces Audit-Ready Data

We use a streamlined visit management process that allows us to efficiently consent and treat participants with minimal downtime and maximum quality. We bolster that with:

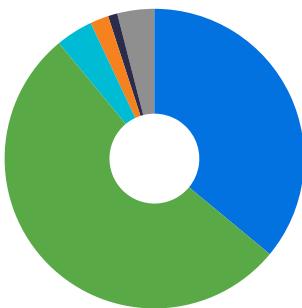
- Overnight quality control on 100% of visits
- Automated eSource exception reporting
- Ongoing third-party internal audits

These tactics—implemented across all projects—enable us to catch and correct issues in near real time. The result: conclusive, audit-ready data for your clinical study.



Intentional Design Delivers Diverse Participant—and Staff—Population

Establishing purpose-built study sites in communities and cities with underrepresented residents, Alliance Clinical places a deliberate focus on helping ensure that innovative new therapies are effective across populations.



Breakdown by Race

Caucasian: 36%
Black/AA: 53%
Asian: 4%
American Indian & Alaska Native: 2%
Native Hawaiian & Pacific Islander: 1%
Other: 4%

Breakdown by Ethnicity

Hispanic/
Latino: 38%

Not Hispanic/
Latino: 62%

About Alliance Clinical Network

Alliance Clinical's fully integrated, wholly owned and operated clinical research sites help advance human health by delivering exceptional clinical trial data gathered from a proprietary database of 280,000+ consented participants. Our centralized management harmonizes processes and systems to deliver consistent study execution across our seven sites—a solid foundation from which to meet your study needs.

To learn more, please contact us.



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