



Developing Next-Generation Therapies for Breast Cancer Patients

Breast cancer is a core area of expertise for Worldwide Clinical Trials. We support breast cancer programs from early-phase development through registrational studies across biomarker-defined patient populations and next-generation treatment modalities.

Successfully executing breast cancer trials requires expertise in biomarker-driven patient identification and enrollment, and molecular testing strategies across a highly competitive recruitment landscape. Partnering with an oncology CRO that understands these complexities is critical to accelerating development and study success.

Robust pipeline of breast cancer trials

Global Footprint

- Programs across North America, Europe, Asia-Pacific, Latin America, and the Middle East and North Africa

Study Experience

- 40+ breast cancer studies
- 4,300+ patients enrolled
- Early-phase through registrational development
- Support across HR+, HER2+, HER2-low, and other biomarker-defined patient populations

Site Experience

- 350+ unique sites
- Partner with leading academic centers and community oncology networks worldwide

Next-Generation Drug Modalities

- Antibody-drug conjugates (ADCs)
- Bispecific antibodies
- Cell therapy/CAR T
- Hormone antagonists
- Kinase inhibitors
- Cancer vaccines

**40+**Breast Cancer
Studies**4,300+**Patients
Enrolled**350+**Unique
Sites**Global
Footprint**NA, EU, APAC,
LATAM, MENA

We understand breast cancer trials

Disease Heterogeneity, Complex Sub-populations, and I/E Criteria

- ER+/-, HER2+/-, TNBC, MSI status, etc.

Competitive Landscape

- Highest number of studies of any indication, investment, and rapidly evolving therapies

Global Disparities in Standard of Care (SoC)

- Varying and multiple lines of prior therapy

Drug Class Nuances

- Targeted therapies: biomarker selection/CDx
- Immuno-oncology: combination and sequencing with CPIs

What Sets Us Apart



Biotech Mindset Delivered Worldwide

We are purpose built around a biotech mindset that shapes both our partnership philosophy and delivery model, bringing the agility, responsiveness, and focus needed to meet critical oncology development milestones.



Oncology Expertise and Team Continuity

Oncology expertise is embedded across all functions, ensuring studies are led by professionals who understand the complexities of oncology science and trial execution. With an average of 11+ years of oncology experience across key roles and an industry-leading 90%+ retention rate in the oncology business unit, you can expect a seasoned, specialized team supporting your study end to end.



Flexible solutions
& best-in-class
technologies



Data-centric approach
with visual analytics
& clinical science



Site relationships
with rapid startup

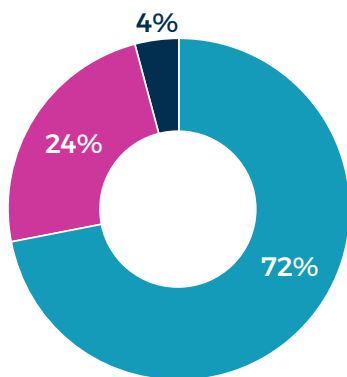


Global footprint:
4,000+ across NA, EU,
LATAM, MENA & APAC

Active Next-Gen Oncology Experience

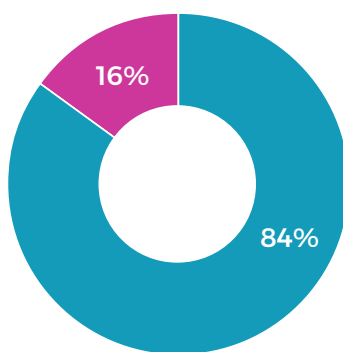
Operating both locally and globally across Phases I through III in a range of solid tumor and hematologic indications, we understand the nuances of complex study designs, novel endpoints, and cutting-edge modalities.

Phase



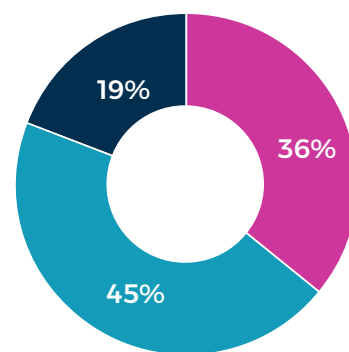
■ Early Phase ■ Late Phase
■ Observational/LTFU

Indication



■ Solid ■ Heme

Drug Class



■ Immuno-Oncology ■ Other
■ Targeted Therapy



Worldwide
Clinical Trials

Oncology at Worldwide

Worldwide features a dedicated oncology business unit within our full-service global contract research organization. We partner with biopharmaceutical sponsors to deliver expert-led, globally scalable oncology trials across every phase of development. We help sponsors move fast with confidence and scale without disruption.

[Contact us](#) to learn how we can support your oncology program from FIH through registration.