

# Reliability. Excellence. Velocity.

At the core of every **clinical partnership.**

# About

Building on our history as CBCC Global Research, REV Clinical reflects our conviction and our focus on serving innovators with intent and purpose. Science moves fast, and the industry needs clinical partners who can move with reliability, excellence, and velocity.

As a US-based global clinical CRO, we collaborate with small and mid-sized sponsors across multiple therapeutic areas. With a growing international footprint, REV delivers speed, flexibility, and capital-efficient execution from first-in-human through late-phase studies. Our teams operate as an extension of yours. We are nimble, transparent, and deeply invested in moving your study forward.

## Therapeutic Expertise

■ Oncology

■ Neurology

■ Ophthalmology

■ Rare Diseases

REV Clinical focuses on complex, high-stakes therapeutic areas where precision and speed directly influence outcomes. Our teams bring hands-on experience with immune-modulating therapies, targeted oncology, device-adjacent ophthalmic programs, and hard-to-enroll neurological and rare conditions. We anticipate the scientific and operational hurdles inherent to these areas and design studies that move programs forward with confidence.

## Core Capabilities

Clinical Operations & Site Support

Biometrics & Data Management

Regulatory & Submission Expertise

Trial Rescue Services

REV Clinical delivers the essential capabilities biotechs need to move fast and stay on track. Our clinical operations teams ignite startup, drive enrollment, and keep execution sharp from FIH through Phase III. Regulatory specialists navigate global pathways with precision, shaping phase-appropriate strategies and crafting submissions that stand up to scrutiny. Our biometrics and data management experts power clean, reliable data flows and analysis-ready outputs that accelerate decisions. If existing studies hit turbulence, our rescue specialists diagnose root issues quickly and deploy cross-regional resources to stabilize execution and restore trial momentum.

## **Global Velocity Network™**

REV's cross-regional model integrates operations in the US, India, and Australia under one leadership structure to deliver:

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Faster global submissions

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Rapid site activation

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Scalable monitoring and data teams

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Cross-regional project management aligned to a single standard

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## **Quality & Compliance**

Since 2021: 50+ regulatory inspections with zero material findings

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GCP-aligned processes across all regions

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Dedicated QA/QC & Regulatory Affairs departments

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Robust vendor qualification and oversight frameworks

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## **Scope of Operations**

Clinical operations across the US, India, and Australia

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Phase I-III trial experience

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Proven ability to run U.S. trials with India-level efficiency

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Strength in oncology, immune-modulating therapies, and ophthalmic/device-adjacent studies

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Strong track record navigating accelerated pathways

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## **Why Sponsors Choose REV**

Startup-speed study activation (as fast as two weeks)

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Capital-efficient execution across regions

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Hands-on leadership team that sponsors can access

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Global Velocity Network for scale + speed

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Flexible teams that reduce bureaucracy and keep trials moving

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**REV**  
Clinical™

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