MEDPACE

MAKING THE COMPLEX

SEAMLESS®

Clinical development is complex and scientifically demanding. Let Medpace execute your clinical trials seamlessly with a corporate structure that is streamlined for execution, cohesive study teams, integrated services and technology, and global resources to scale with your compound through all stages of clinical development.

QUALITY AT OUR CORE

Built to execute—enabled by organic growth

Accelerate your clinical development by partnering with a CRO that is structured to keep study teams focused on execution. This commitment to quality extends to the projects we engage in – we are selective to ensure we deliver the highest levels of service and partnering. Our unique approach has been preserved by 30+ years of organic growth – delivering consistency and stability while avoiding the disruptions associated with mergers and acquisitions.

TEAM CONTINUITY

Expertise embedded in trials—throughout the project lifecycle

Engage with one cross-functional team that actively participates in your trial or program – from early planning through close-out. Our approach maintains continuity and optimizes performance while building strong team chemistry, minimizing disruptions, and ensuring high-quality results across our full breadth of services. With deep scientific and therapeutic expertise, our teams are poised to lead the industry's most innovative and challenging clinical trials.

INTEGRATED SERVICES

End-to-end partnership—a model of productivity

Streamline even the most complex clinical trials with a full-service, single-vendor outsourcing strategy. Our comprehensive CRO services are supported by our wholly-owned Central Laboratories, Bioanalytical Lab, Imaging Core Lab, ECG Core Lab, and Phase I Unit, as well as a Clinical Trial Management System that ties all study data together in a single platform. With integrated services and systems, we achieve higher levels of efficiency and productivity.

GLOBAL ENGAGEMENT

Navigating cultures and requirements—reaching patients around the world

Leverage our country-specific experience and breadth of in-house expertise to navigate languages, cultures, and clinical and regulatory environments around the globe. Our resources extend across North America, Europe, Latin America, and Asia Pacific—with strong site and Investigator relationships to reach targeted and diverse patient populations. Our global reach and experience across all phases of development and key therapeutic areas can help streamline your clinical trials and reach your development goals faster.