

MEDPACE[®]

MAKING THE COMPLEX
SEAMLESS[®]



**WE CAN'T SIMPLIFY CLINICAL DEVELOPMENT –
BUT WE CAN EXECUTE IT SEAMLESSLY.**

PHILOSOPHY FOR FULL-SERVICE OUTSOURCING

A LETTER FROM THE CEO

To our current and future Sponsors,

Over our 30+ year history, Medpace has steadfastly held to a model of providing full-service clinical development services to biopharmaceutical and device Sponsors. Even as the industry explored various outsourcing/insourcing models — functional, clinical staffing and hybrids — Medpace chose to drive success for Sponsors through full-service outsourcing.

We know from our long-standing relationships with sponsors that the full-service outsourcing model ultimately delivers higher quality results. When we can fully engage with our medical, regulatory and operational teams and work under our SOPs, we can perform at the highest levels to deliver quality results in the most timely and efficient manner. Competence and empowerment to coordinate all services under one roof provides an accountable, seamless, integrated and efficient platform — increasing quality and speed while significantly reducing a Sponsor's need for duplicate management oversight.

Investing in extraordinary talent produces exceptional results.

- Therapeutically strong clinical development teams for superior execution
- Global regulatory experts who can provide local knowledge and support
- Strong investigative site — key opinion leader relationships
- Leading technology platform — ClinTrak® — for total study decision support

It is this investment in our talented teams and systems that bring superior value to our Sponsors.

I share my philosophy on full-service outsourcing with you because it has stood the test of time. I truly believe it is the best path for Medpace and for our Sponsors.



August Troendle, MD
Medpace Founder and CEO



OUR MISSION: ACCELERATING THE GLOBAL DEVELOPMENT OF SAFE AND EFFECTIVE MEDICAL THERAPEUTICS

Clinical development is more complex and scientifically-demanding than ever before. Successful clinical trials require expertise, experience, and a disciplined team collaborating in unison. Operating under a full-service model, Medpace provides a therapeutically focused, integrated, global approach for seamless execution and quality results.



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.



TEAM CONTINUITY

EXPERTISE EMBEDDED IN TRIALS THROUGHOUT THE PROJECT LIFECYCLE

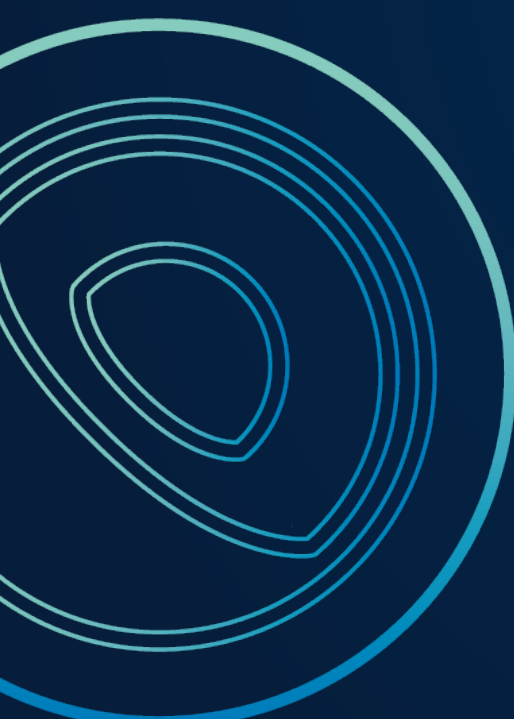
Engage with one cross-functional team that actively participates in your trial or program – from early planning through close-out. Our approach maintains continuity and optimizes performance while building strong team chemistry, minimizing disruptions, and ensuring high-quality results across our full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



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 varied 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.



QUALITY AT OUR CORE

BUILT TO EXECUTE—ENABLED BY ORGANIC GROWTH

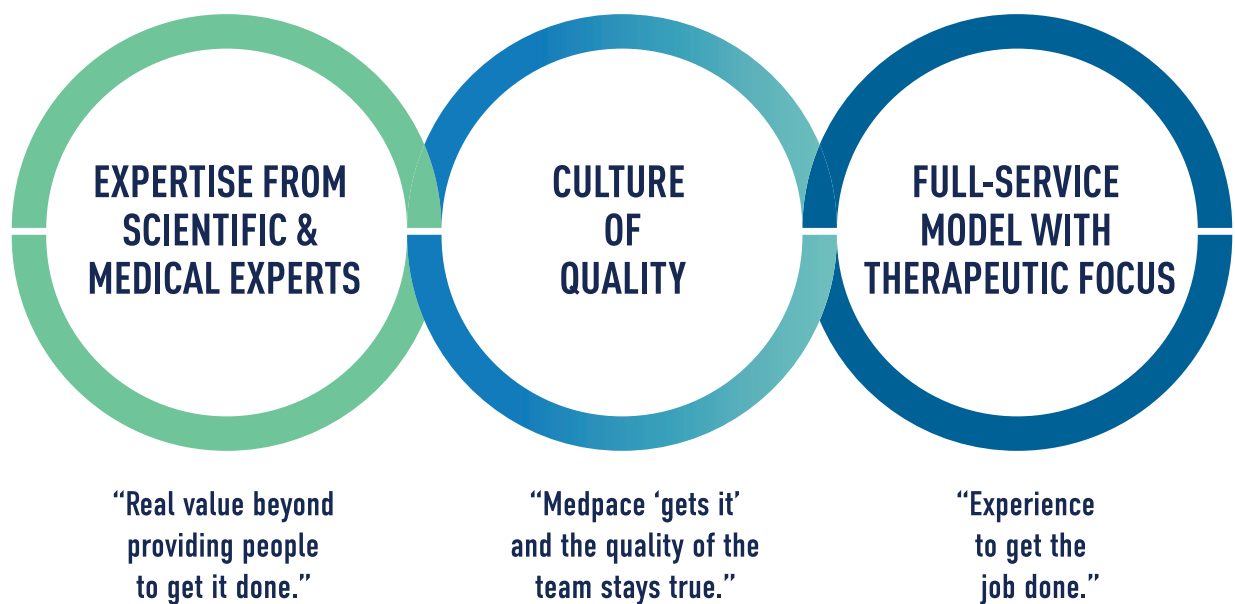
Accelerate your clinical development by partnering with a CRO that is structured to keep study teams focused on execution. This commitment to quality extends to the projects we engage in – we are selective to ensure we deliver the highest levels of service and partnering. Our unique approach has been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

YOUR PARTNER IN CLINICAL DEVELOPMENT

AN EXTENSION OF YOUR TEAM THAT SHARES YOUR COMMITMENT

Medpace's dedicated teams serve as an extension of your team. We engage quickly and provide strategic thinking to ensure quicker start-up times, superior quality, and efficient delivery for any phase of development. Our therapeutic and regulatory experts are committed to streamlining the path to approval so every partnership is designed to create research solutions focused on your critical needs.

Our philosophy emphasizes an uncompromising commitment to clinical research with the highest level of ethical standards and performance in our jobs. We are selective about the projects we engage in because we are devoted to quality and providing our Sponsors with best-in-class service.

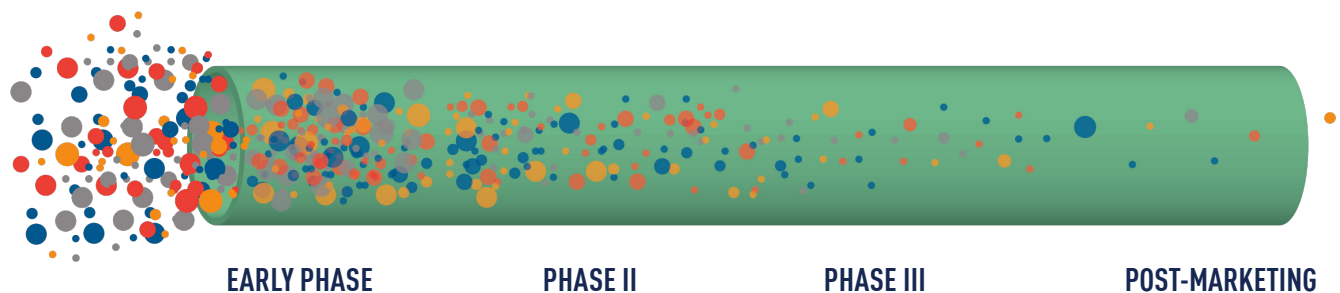


***"We wanted to work with a company that was as entrepreneurial in spirit, as innovative, and adaptable as we were. And I think that is what we ultimately gained with Medpace as our partner."
- CEO, small- mid-size biotech***

FULL-SERVICE CAPABILITIES AND EXPERTISE

COMPREHENSIVE SERVICES FOR ACCELERATED DEVELOPMENT

Throughout the clinical development life cycle, Medpace provides medical and regulatory leadership guidance and efficient, disciplined operational execution of studies around the world. Our comprehensive capabilities, resources and global footprint ensure quality and timely development across all phases of research.



CORE SERVICES

- Clinical trial management
- Clinical monitoring
- Clinical trial submissions
- Clinical packaging and supplies
- Study start-up
- Vendor management
- Patient recruitment & retention
- Medical monitoring
- Safety and pharmacovigilance
- Endpoint adjudication
- Independent assessment committees
- Data management
- ePRO technology
- IRT
- Biostatistics
- Medical writing
- Quality assurance
- Regulatory support

INTEGRATED LABS

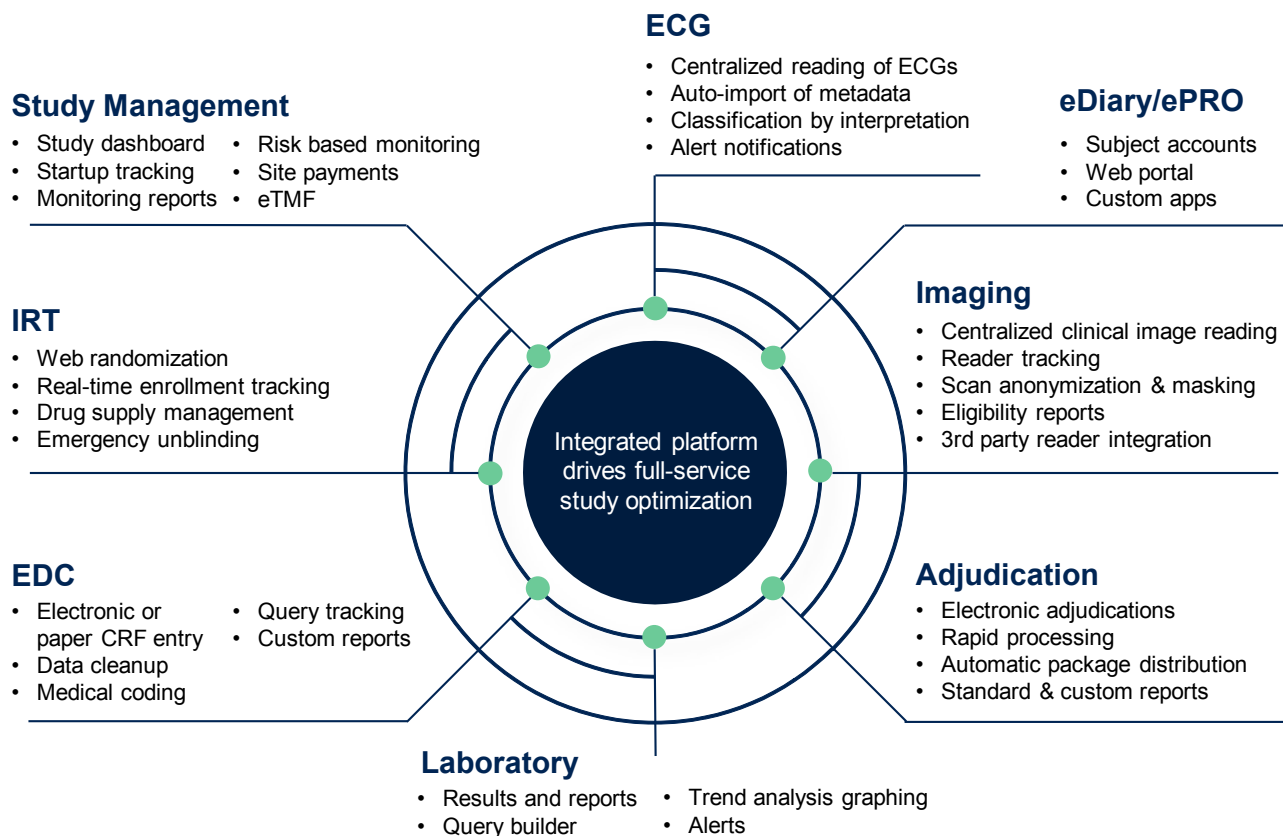
Our wholly-owned laboratories offer full-service scalable support for all phases of development worldwide. This “single-source” approach drives significant efficiencies throughout the study while simplifying vendor management for our Sponsors.

- Central Labs and Biorepository
- Bioanalytical Labs
- Imaging and ECG Core Labs

ClinTrak®

CLINICAL TRIAL MANAGEMENT SYSTEM

Medpace offers an innovative suite of proprietary technology with our ClinTrak clinical suite. ClinTrak is a study management system built for cross-team coordination and decision support for Sponsors and sites while ensuring global teams are organized for maximum efficiencies. ClinTrak uses a common data platform and infrastructure allowing for study optimization and real-time access with a single login for critical study data, tracking, interpreting, and communicating information in a timely, secure, and cost-effective manner.



***One platform. One sign-on.
All aspects of the clinical development process at your fingertips.***



MEDPACE®