About Medpace

Medpace is a scientifically-driven, global, full-service clinical contract research organization (CRO) providing Phase I-IV clinical development services to the biotechnology, pharmaceutical and medical device industries. Medpace's mission is to accelerate the global development of safe and effective medical therapeutics through its physician-led, high-science, and disciplined operating approach that leverages local regulatory and deep therapeutic expertise across all major areas including oncology, cardiology, metabolic disease, endocrinology, central nervous system and anti-viral and anti-infective. Headquartered in Cincinnati, Ohio, Medpace employs approximately 2,500 people across 35 countries.

A Full-Service Clinical Research Partner

Throughout the development life cycle, Medpace provides medical and regulatory leadership and guidance, with efficient, disciplined operational execution of your studies around the world. Our unique global partnering philosophy emphasizes an uncompromising commitment to clinical research and to 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 partners with best-in-class service.

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

Driven by a full-service CRO model, Medpace provides an accountable, seamless, integrated and efficient platform for executing clinical research – increasing quality and speed while significantly reducing the need for duplicate management oversight. Our disciplined processes, site relationships, and technologies enable us to execute even the most complex global studies.

Explore our full range of integrated services:

- Medical Affairs
- Regulatory Affairs and Medical Writing
- Clinical Monitoring
- Clinical Trial Management
- Biometrics & Data Sciences
- Safety & Pharmacovigilance
- Quality Assurance
- Proprietary CTMS

Global Laboratories

Fully integrated labs facilitate efficiency and collaboration.

- Medpace Central Laboratories: wholly owned and purpose built laboratories around the world
- Medpace Bioanalytical laboratory: experience working with small molecules, biologics, and biomarkers across a wide variety of technologies and therapeutic areas
- Imaging Core Lab and Cardiovascular Core Lab further support studies with imaging and safety requirements

Physician Led - Therapeutically Focused

Increasingly complex clinical research demands that you engage a team of medical experts to navigate the challenges. Medpace is unique in its physician-led approach to clinical research. The Medpace model gives you the advantage of early and ongoing insight and guidance from therapeutic experts (MDs and PhDs) throughout trial design and execution. Your project team will be led by medical, regulatory and operational experts with deep therapeutic experience who are fully engaged throughout your study, proving guidance and averting potential roadblocks by staying close to the project.

Therapeutic Areas:

- Cardiovascular
- Endocrine & Metabolic
- Hematology & Oncology
- Infectious Diseases and Vaccines
- Neurology & Psychiatry
- Nephrology

Specialty Areas:

- Pediatrics
- Rare and Orphan Disease
- Advanced Therapies

Accelerate your drug, biologic, or medical device clinical development with a full-service, highly disciplined and therapeutically-focused model.

For more information email info@medpace.com

MEDPACE

Therapeutically Specialized Clinical Development

North America Latin America Europe Asia Pacific Middle East Africa