MEDPACE

MAKING THE COMPLEX

SEAMLESS™

WE CAN'T SIMPLIFY CLINICAL DEVELOPMENT — BUT WE CAN EXECUTE IT SEAMLESSLY

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.

THERAPEUTIC FOCUS

Scientific expertise embedded in trials — hands-on and specialized

Translating medical, operational and regulatory knowledge into execution while leveraging established KOL and site relationships to drive successful research.

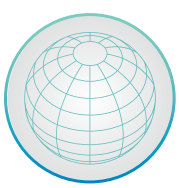
Medpace is unique in its scientifically-driven approach to clinical research. The Medpace model gives you the advantage of early and ongoing insight and guidance from therapeutic experts throughout trial design and execution. Our therapeutically-focused culture facilitates cross-collaboration across specialties to incorporate various medical perspectives and considerations. All project teams are led by medical, regulatory and operational experts with deep therapeutic experience who are fully engaged throughout every study, providing guidance and averting potential roadblocks by staying close to the project.

INTEGRATED EFFICIENCY

A model of collaboration — an end to end partner

The built-in collaboration and efficiencies of working with a single vendor facilitates a streamlined strategy for executing even the most complex global studies.

Integrating core clinical trial services delivers efficient and streamlined execution. Medpace offers comprehensive and fully integrated laboratory services including global central laboratories, bioanalytical laboratories, imaging and ECG core laboratories, as well as a clinical pharmacology unit.



GLOBAL REACH

Operational around the world — wherever research is happening

With resources around the globe, Medpace skillfully navigates local languages, cultures and processes to avoid delays and missteps — delivering seamless execution amid the complex landscape of global clinical development.

As a global CRO with an operational footprint across 40 countries, Medpace has broad experience designing and conducting Phase I-IV clinical trials around the globe. From feasibility, to patient recruitment, to study start-up, Medpace has the resources to advance your medical therapeutic in any region. Global reach also means that our medical and operational specialists have country-specific expertise to deliver faster enrollment and obtain access to country-specific patient populations, while our regulatory experts can plan and coordinate each aspect of regulatory strategy and engagement—locally and globally.

ORGANIC GROWTH

Stable and disciplined — preserving a culture of quality

Our 29 year history with purposeful, organic growth provides consistency in leadership, deep institutional experience, and incomparable efficiencies as a top 10 CRO.

While the CRO industry has grown primarily through mergers and acquisitions, Medpace has expanded through disciplined organic growth—expanding globally to operations in 40 countries and 4,100 employees. Over decades, we've systematically added specialized medical, regulatory and operational experts, and refined and enhanced custom-built technologies and processes to best serve the needs of our clients. The result is a culture built on quality that has not been disrupted by acquisitions, and that delivers ongoing efficiencies and stability.