MEDIAL DEVICE & DIAGNOSTICS

A COMPETITIVE EDGE FOR CLINICAL DEVELOPMENT

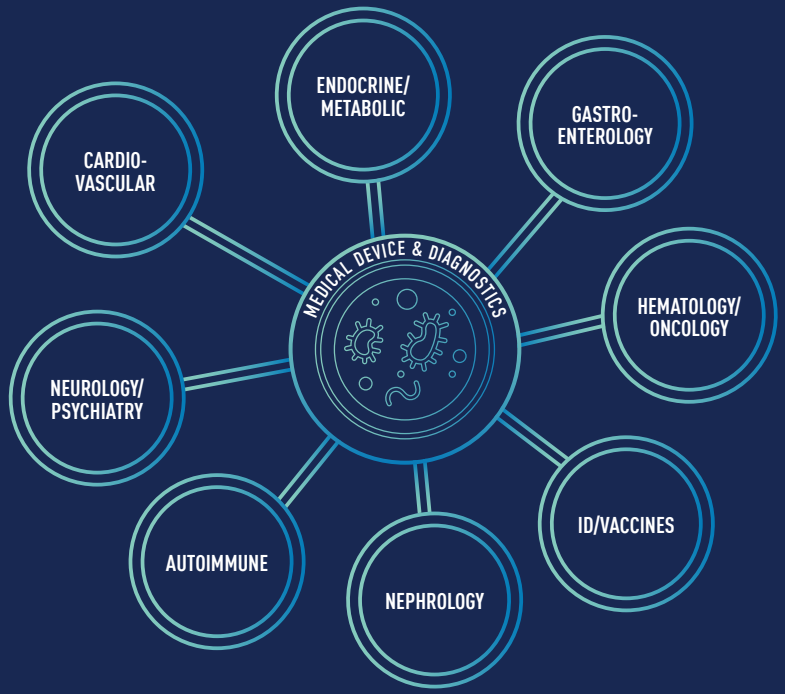
GLOBAL AND FULL-SERVICE

Clinical and Regulatory Services

- Strategies and Trial Design
- Early Feasibility Studies
- Clinical Trial Execution
- Reporting and Regulatory Approval
- Post-Market Support

THERAPEUTIC FOCUS

Scientific Expertise Embedded In Trials—Hands-On & Specialized



GLOBAL DEVICE CLINICAL DEVELOPMENT EXPERIENCE

Medpace is a global, full-service CRO with a scientifically driven approach to clinical research. Our dedicated medical device and diagnostics team brings deep operational expertise to every stage of development—from first-in-human and feasibility studies to pivotal trials and large-scale post-market evaluations. With experience across a broad range of therapeutic areas and technologies, we partner with Sponsors to efficiently design and execute clinical investigations and performance studies that meet global regulatory requirements.

DIFFERENTIATORS & BENEFITS

- Cross-functional teams of medical, operational, and regulatory experts that are actively involved throughout the lifecycle of the project
- Specialized medical and device expertise in key areas of diagnostic development including hematology and oncology, infectious diseases, and cell and gene therapies
- Integrated imaging, ECG management, and central lab to provide seamless logistics, review and testing
- Experience with varied medical settings including bed-side, laboratory or other healthcare professional settings
- Strategic regulatory affairs leadership provides early guidance and execution for developing diagnostics
- Our purpose-built CTMS, ClinTrak®, delivers on-demand access to all your clinical study data in an intuitive interface – unlocking deeper insights and empowering objective, data-driven decision making

MEDPACE®

MEDPACE®

MAKING THE COMPLEX SEAMLESS®