

GLOBAL REACH

Medpace has the global reach and capability to conduct studies and assist with regulatory requirements throughout the world.



FOCUS: INDIA

MEDPACE

THE ADVANTAGE OF **FOCUS**

India Partnerships

Overview

India provides significant advantages for the conduct of clinical research trials. The country offers a large patient pool, a low per-patient cost, and a highly-qualified professional medical community. These advantages are complemented by India's world class medical centers and clinical research facilities. With highly-developed information technology systems in place and no language barrier, India has become an important country for global clinical trials. Medpace has responded positively to these incentives by opening its Indian headquarters in Mumbai.

Recruitment

India excels in patient recruitment because of a substantial heterogeneous population pool of over 1 billion, a well-trained medical community, and an offering of incentives from the government for clinical research activities. High incidences of cancer and diabetes coincide with the Medpace areas of specialization in oncology, cardiology, and metabolism. Large numbers of cardiovascular events and stroke contribute to shorter recruitment timelines for critical endpoint trials. Medpace continues to form regional partnerships and has developed positive relationships with research sites, investigators, and key opinion leaders.

Data

India has technologically-sophisticated data capabilities supporting the use of electronic data capture by investigators and research facilities. Medpace ensures data quality through comprehensive site selection, global training, and certification processes. Investigators and coordinators are trained in Good Clinical Practice (GCP) and maintain excellent quality control for data collection.

Regulatory

India has one of the most rapid and predictable clinical trial regulatory timelines in the world. Medpace has clinical trial submission expertise along with complete knowledge of regulations in the region. Medpace understands the procedures covering regulatory policy and importation and exportation processes which is essential to its success in India.

Reaching a Global Population

Medpace Worldwide

Patient recruitment is the greatest obstacle in clinical trials today. Finding the right patient populations has fueled the rapid international growth of Medpace. With operations in nearly 40 countries, Medpace is well-positioned to meet changing client needs. Medpace offices are located in Africa, Asia/Pacific, Australia, China, Eastern Europe, India, Latin America, the Middle East, North America, and Western Europe.

Medpace is focused on making every global study cost-effective and well-organized. Start-up and efficiency metrics are maintained for each country where trials are conducted. Medpace assists sponsors in prioritizing international sites, not only by patient populations, but also by start-up speed and relative cost.

Clinical Research Associates (CRAs) are regionally-based and have in-depth knowledge of the local culture, language, and regulatory environment where they work. Medpace CRAs typically are assigned just two protocols each. They gain a thorough understanding of the therapeutic area and protocol, which allows them to communicate knowledgeably with investigators.

Therapeutic Areas of Expertise

Oncology

Oncology clinical trial development is longer and more intricate than trials in any other therapeutic area. The complexity of this development requires the right kind of research partner.

At Medpace, a unified team of oncology experts and highly knowledgeable clinical operations staff are prepared to quickly provide clients with the efficacy and safety data needed to support product approval worldwide.

Medpace has experience in classic multi-cytotoxic agent and targeted and/or biologic therapy trials and has conducted oncology trials throughout the world.

Cardiology and Metabolism

Medpace is home to renowned regulatory and therapeutic experts who are committed to the complex and intertwined therapeutic areas of cardiology and metabolism.

The medical team includes international therapeutic leaders who have helped pioneer many of the preventative cardiovascular and metabolic compounds introduced in the last 15 years.

Medpace regulatory experts are former government regulatory officials who have invaluable insight into the specific areas of cardiovascular and metabolic drug approval.

The Medpace Difference

The Advantage of Focus

Medpace is a global, research-based drug development company led by top therapeutic and regulatory experts who are driven to further the advancement of pharmaceutical agents for use in cardiology, metabolism, and oncology. Medpace has assembled the industry's most experienced and therapeutically-focused team to execute at all levels of the company's operations, providing complete and seamless drug development services.

Through specialized regulatory expertise and therapeutically focused clinical operations, Medpace creates strategic partnerships with pharmaceutical and biotechnology companies to provide the most efficient and cost-effective path to drug development – from program planning and executing to product approval.

With clinical trial experience in over 40 countries, Medpace has the global reach and capacity to conduct studies and navigate regulatory requirements worldwide. The Medpace approach creates therapeutically-focused project teams and increases the quality of every service offered.

Medpace group of companies

Medpace

Imagepace

Medpace Bioanalytical Laboratories

Medpace Clinical Pharmacology

Medpace Reference Laboratories