



GLOBAL REACH

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

FOCUS: LATIN AMERICA

MEDPACE

THE ADVANTAGE OF **FOCUS**

Latin America Partnerships

Overview

Latin America is emerging as a key global region for clinical trials because of its large population, highly-qualified medical personnel, similarities to Western medicine, and established regulations. With a staff average of 10 or more years of clinical research experience, Medpace continues to open offices and work with local partners to support clinical research in the region. Medpace has its Latin American headquarters in Sao Paulo, Brazil.

Recruitment

Medpace has developed consistent, longstanding, and positive relationships with local Latin American regulatory authorities, leading investigators, and research sites, resulting in expedited study start-up and rapid enrollment. The Latin American population exceeds 500 million, with 70% concentrated in urban areas and a large percentage of patients qualifying as treatment-naïve. The region's expanding number of clinical trials in breast and lung cancer, cardiovascular studies, and diabetes complement Medpace therapeutic expertise in oncology, cardiology, and metabolism. Strong physician-patient relationships foster high retention rates. In Latin America, typical seasonal illnesses that impede patient enrollment occur six months out of phase with North America, permitting Medpace to take advantage of year-round recruiting.

Data

Regionally, the standard of medical training is high, and knowledge of the guidelines established by the International Conference on Harmonisation and Good Clinical Practice (GCP) is extensive. The strong relationships Medpace has formed with regulatory authorities, leading investigators, and research sites promote better data quality.

Regulatory

Medpace has clinical trial submission expertise in Latin America, along with current knowledge of regulations in the region. In addition, Medpace understands the strict and complex regulatory importation processes. Medpace knowledge of the medical and cultural issues in the region is essential to its success in Latin America.

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