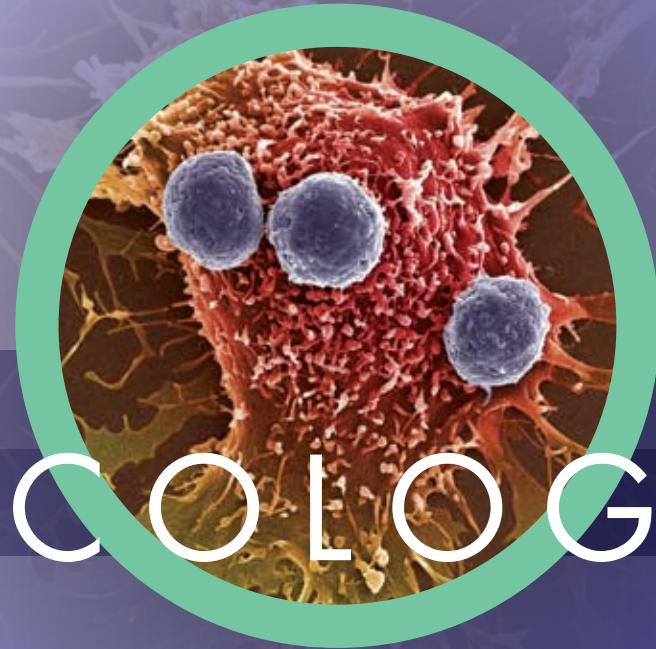


**THERAPEUTIC EXPERTISE IN ONCOLOGY CLINICAL  
RESEARCH TO SUPPORT PRODUCT APPROVAL**

The medical and regulatory expertise to deliver effective recruitment and accelerate accurate results in trials of varied scope on six continents.

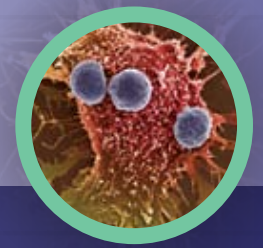


# FOCUS: ONCOLOGY

**MEDPACE**

THE ADVANTAGE OF **FOCUS**

# FOCUS: ONCOLOGY



## Research with a sense of urgency

At Medpace, a unified team of oncology experts and highly knowledgeable clinical operations staff are dedicated to meeting the distinct challenges of oncology trials and quickly providing partners with the efficacy and safety data needed to support product approval worldwide.

With experience in trials from early through late phase development (Phase I-IV), including classic multi-cytotoxic agent and targeted and / or biologic therapy trials, the oncologic experts at Medpace deliver rapid and accurate results in trials of varied scope – from small, focused trials to large studies.

There is an urgent need to develop new oncologic therapies. The complexity of oncology development demands the right kind of research partner. Powerful solutions demand the thought leaders at Medpace.

## Overcoming recruitment challenges

Medpace takes a proactive approach to the challenges of oncology trial recruitment. Our experts have conducted oncology trials throughout the world and have medical and regulatory expertise in Africa, Asia / Pacific, Australia, China, Central and Eastern Europe, India, Latin America, the Middle East, North America, and Western Europe. This experience, coupled with longstanding relationships with oncologists and clinical sites around the world, allows us to target the areas where recruitment will be most effective. Leveraging our unique partnering philosophy, Medpace consults with our partners, Sponsors, and investigative sites, to ensure every protocol is focused correctly for maximum results.

## The advantage of oncology focus

The Medpace oncology medical team provides clients with:

- Medical team leadership for oncology
- Experience in oncology product development planning, trial design, and regulatory filings
- Project teams with experience in oncology-specific grading systems
- Oncology expertise at every level of clinical operations, safety / pharmacovigilance, biostatistics, and data management
- Established oncology investigator relationships
- EDC and DSMB planning and support

## The Medpace difference

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 our 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 execution to product approval. With clinical trial experience in over 40 countries, Medpace has the global reach and capability to successfully conduct studies and navigate regulatory requirements worldwide.

The Medpace approach creates therapeutically-focused project teams and increases the quality of every service, including:

- Medical Expertise
- Regulatory Affairs
- Safety / Pharmacovigilance
- Clinical Operations
- Data Management
- Biostatistics
- Quality Assurance
- Medical Writing
- Pharmacokinetics
- Bioanalytics
- Central Imaging

## Medpace group of companies

- Medpace**
- Imagepace**
- Medpace Bioanalytical Laboratories**
- Medpace Clinical Pharmacology**
- Medpace Reference Laboratories**