

**GLOBALIZED CENTRAL LABORATORIES DELIVER PRECISE  
RESULTS SUPPORTING SUCCESSFUL DRUG DEVELOPMENT**

The global reach and capability to conduct studies, assist with regulatory requirements, and deliver custom solutions, specific to your needs, on six continents.



# MEDPACE REFERENCE LABORATORIES

**MEDPACE**

THE ADVANTAGE OF FOCUS

# MEDPACE REFERENCE LABORATORIES



## A global network of laboratories

Medpace Reference Laboratories (MRL) is a global leader in providing customized, high-quality laboratory services to the pharmaceutical and biotech clinical development industries. A full-service central laboratory with locations in Cincinnati, Ohio; Leuven, Belgium; Mumbai, India; and Beijing, China, MRL combines a unique partnering philosophy with state-of-the-art infrastructure. Our team of medical and technical experts has extensive experience in all areas of central laboratory services – including cardiovascular and metabolic diseases, and oncology.

MRL has the global reach and the local presence to conduct studies and assist with regulatory requirements on six continents. Our dedicated experts provide comprehensive, flexible laboratory services that maintain uniform instrumentation, reagents, calibration, and standard operating procedures (SOPs), regardless of location. Identical reference ranges throughout a global trial are maintained guaranteeing that all results generated, in all laboratories, are the same and require no manipulation or conversion.

With the infrastructure needed to support patient participation and accessibility as well as longstanding partnerships with the best investigative sites around the world, only Medpace has the expertise to deliver high-quality results, efficiently and accurately. And, our state-of-the-art proprietary Laboratory Information Management System (LIMS), ClinTrak® Lab, gives you the power to compare customized results from subjects in Prague to subjects in Mumbai in near real-time.

## A customized approach

Drawing upon one of the most experienced teams in the central laboratory field, we listen to and analyze your needs to create a customized approach designed specifically to meet your project objectives. Collaborating with the therapeutic and regulatory experts of Medpace, MRL senior management are involved in every step of the drug development process ensuring you receive the personalized service, professional oversight, and expertise you deserve.

### Complete control for powerful solutions

As a provider of central lab services for different landmark trials, standardization programs, and NDAs, MRL understands how to add value to every project. MRL controls its operations at all locations – leveraging reference methodology, the highest levels of national and international standardization, and attention to even the smallest detail – to ensure the highest quality of specimens during collection and transport. From logistics to project management to sample analysis, this complete control delivers truly identical results on a global basis from start to finish.

Advancing your drug from development plan to regulatory approval in the most efficient and cost-effective manner demands a concentrated effort every step of the way. That's why MRL is committed to science and service, not marketing and volume.

## A broad range of capabilities

Key benefits of MRL include:

- Experienced, service-oriented, expert leadership and professional support staff with a proven track record in central lab services
- Certified pathologists and experienced scientists available for consultation during all phases of the study
- Wholly-owned state-of-the-art facilities, instrumentation, and methodology (identical in all Medpace labs) with global coverage
- Industry-appropriate quality assurance and accreditation, including CDC Part III (gold standard) lipid standardization, NGSP (Level I) certification, and CAP accreditation
- Dedicated project management as a single point-of-contact for Sponsors and sites
- Key Performance Indicators and Financial Performance Indicators for optimal trial monitoring and planning
- Intelligent supply chain management, including full lot and expiry date tracking, for production and on-time distribution of customized sample collection kits
- Innovative solutions for controlled sample shipments across the world
- In-process data cleaning
- Customized reporting and delivery options to accommodate international and study-specific requirements
- Access to near real-time data via our fully customizable, web-based Laboratory Information Management System, ClinTrak Lab
- Long-term, secure, on-site archival storage of specimens (-20°C to -180°C)
- Quality-assurance compliant with all applicable regulatory requirements (GLP, GCP, CLIA'88, CAP, FDA 21 CFR Part 11)

## 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.

### Medpace group of companies

**Medpace**

**Imagepace**

**Medpace Bioanalytical Laboratories**

**Medpace Clinical Pharmacology**

**Medpace Reference Laboratories**