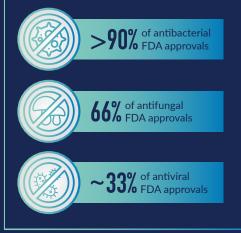


INFECTIOUS DISEASES & VACCINES A COMPETITIVE EDGE FOR CLINICAL DEVELOPMENT

DIFFERENTIATORS & BENEFITS

- Collaborative and cross-functional teams of seasoned medical, regulatory, and clinical project teams with extensive experience designing and conducting infectious diseases and vaccine clinical trials
- Led by noted experts, Medpace medical team is deeply embedded in your trials from protocol design through submission and approval
- Expertise to secure government and non-dilutive funding such as BARDA, NIH, CARB-X, NAMD, NIAID, and Wellcome Trust
- A specialized Data Integrity Unit dedicated to collaborating with the clinical team and ensuring the protocol is maximally operationalized to collect appropriate microbiological, diagnostic, virology and biomarker data to evaluate study outcomes
- Integrated global central laboratory services to support global infectious diseases trials and coordinate the complex logistics with specialty testing labs

In the past 7 years, Medpace ID&V experts have contributed to over:



GLOBAL, PHASE I-IV CLINICAL DEVELOPMENT EXPERIENCE

Broad experience in anti-infective drug development, including antivirals, antibiotics, antifungals, biologics, and vaccines across indications spanning adult and pediatric patients, as well as special at-risk populations.

Antivirals

- Nucleos(t)ide analogs
- RTIs
- Entry inhibitors
- Protease inhibitors and boosters
- Allogeneic and stem cell therapies
- RNAi
- Monoclonal antibodies

Antifungals

- Polyenes
- Echinocandins
- Triazoles

Vaccines/Biologics

- Protein/polysaccharide
- DNA
- Live attenuated

Anti-bacterial

- Aminoglycosides
- Oxazolidinones
- Macrolides/ketolides
- Fluoroquinolones
- Fab-Inhibitors
- Carbapenems
- Beta-lactams
- Beta-lactamase inhibitors
- Pleuromutilins
- Phage therapy
- Microbiome
- Monoclonal antibodies

MEDPACE

INTEGRATED EFFICIENCY

A Model Of Collaboration – An End To End Partner



- Central Labs
- Bioanalytical Lab
- Core Labs
- Phase I Unit

THERAPEUTIC FOCUS

Scientific Expertise Embedded In Trials-Hands-On & Specialized



SEAMLESS

ORGANIC GROWTH

Stable And Disciplined— Preserving A Culture Of Quality



| What's The Difference? | Why It Matters |
|-------------------------------------|---|
| 30+ year history o | Stability in leadership and deep institutional experience |
| Embedded physician OO leadership | Therapeutic expertise through trial lifecycle |
| Strategic regulatory OO affairs | Early planning for long term success |
| Scientific reputation o | Access to KOLs and sites |
| One-of-a-kind O | Built-in efficiency |
| ClinTrak [®] o | Technology for proactive study management |
| Budget management OO model | Operations focused on execution |
| Rigorous training o | Consistent quality around the world |
| Office-based model O | Facilitates collaboration and communication |

GLOBAL REACH

Operational Around The World—Wherever Research Is Happening

