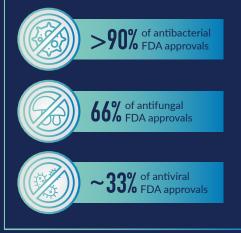


# **INFECTIOUS DISEASES & VACCINES** A COMPETITIVE EDGE FOR CLINICAL DEVELOPMENT

# 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 <sup>®</sup> 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

