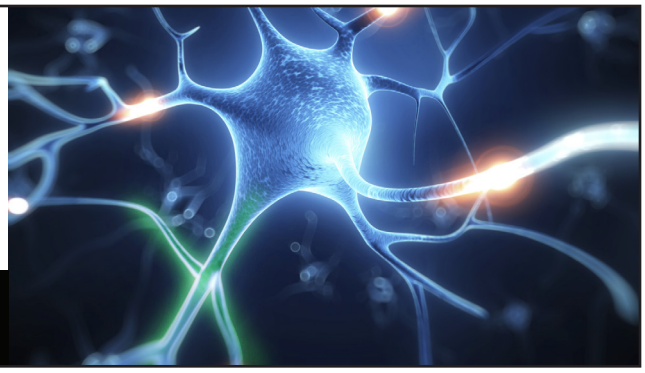
MEDPACE

PSYCHIATRY

Experts. Experience. Execution.

Discover the POWER OF X®



The Power of X in Psychiatry

Experts:

- Medical, regulatory and clinical project teams with experience designing and conducting psychiatry clinical trials
- Led by a team of medical doctors with extensive Phase I-IV experience in Psychiatry and Neurology drug development
- Skilled, proactive operational staff experienced with the unique challenges of psychiatric studies

Experience:

- Broad experience in Psychiatric drug development, including depression, bipolar disorder, anxiety disorders, and schizophrenia
- Spans adult and pediatric populations, with additional expertise in geriatric populations
- Includes drug, diagnostic and device

Execution:

- Full-service approach to drug development - visit medpace.com for a list of comprehensive services
- Strong relationships with key industry investigative sites expedites study start-up, patient recruitment, and query resolution
- Coordination of rater training and certification including subjective assessments, rater reliability, and scale validation
- Powerful, web-based Clinical Trial Management technology platform

Peace of Mind - Partner with a Specialty Team for Your Psychiatric Clinical Development

Developing effective treatments for psychiatric disorders such as depression, bipolar disorder, anxiety disorders, and schizophrenia require unique expertise and capabilities. To streamline your clinical development, Medpace offers medical and regulatory experts, therapeutic experience, and the disciplined processes, site relationships, and technologies required to execute even the most difficult studies

Full-service Advantage

Medpace supports our sponsors through specialized expertise in the design and management of psychiatric programs that maximize a drug candidate's probability of success. Utilizing a one-stop approach to drug development, Medpace keeps your programs on course. We combine efficient clinical trial management, comprehensive regulatory consulting, and innovative technologies to execute at the highest level.

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Key differentiators include:

- A unique scientifically-driven approach to clinical development puts our team of neuroscience medical doctors at the center of your trials, providing strategic direction for study design and planning, training operational staff, working with primary investigators, providing medical monitoring, and meeting with regulatory agencies
- Full-time, in-house psychiatrist with 14+ years in the pharmaceutical industry designing and managing trials in psychiatry and neurology
- Experienced in key indications including depression, bipolar disorder, anxiety disorders, and schizophrenia
- With a reputation for managing some of the industry's most complex and challenging trials, Medpace has a unique concentration of study experience in rare disease and orphan indications
- Our physicians and professional staff have in-depth knowledge of screening tools and rater scales and will provide oversight for rater services including subjective assessments, inter-rater reliability, and scale validation
- Broad experience conducting Phase I-IV clinical trials spanning adult and pediatric populations
- Long-term relationships with successful, experienced sites, networks, and key opinion leaders (KOLs)
- Extensive early phase and transitional development expertise supported by Medpace's Phase I Clinical Pharmacology Unit
- Integrated imaging and central laboratory services, ensuring seamless logistics, review and testing
- Global footprint with operations in 50 countries in North America, Latin America, Europe, Asia-Pacific and Africa

Medpace Services

- Comprehensive clinical research support
- Medical and regulatory consulting
- Clinical development plan preparation
- Protocol development
- Case report form design
- Regulatory document preparation
- Clinical monitoring
- Study management
- Medical monitoring
- Web-based technologies for study tracking
- Safety surveillance and reporting
- Data management
- Statistical analysis
- Medical writing
- Good Clinical Practice compliance audits
- US and International regulatory submissions
- Real world evidence generation
- Health economics and outcomes research
- Central laboratory services
- Bioanalytical laboratory services
- Imaging
- Pharmacology
- medical device

Who We Are

Medpace is a scientifically-driven, global, full-service clinical contract research organization (CRO) providing Phase I-IV clinical development services to the biotechnology, pharmaceutical and medical device industries. Medpace's mission is to accelerate the global development of safe and effective medical therapeutics through its high-science and disciplined operating approach that leverages local regulatory and deep therapeutic expertise across all major areas including oncology, cardiology, metabolic disease, endocrinology, central nervous system and anti-viral and anti-infective. Headquartered in Cincinnati, Ohio, Medpace employs approximately 2,500 people across 35 countries.