MEDPACE

WOMEN'S HEALTH

Experts. Experience. Execution.

Discover the POWER OF X

The Power of X in Women's Health

Experts:

- Medpace Medical experts are deeply involved throughout your trials and work closely with the study team
- Global staff experienced in women's health studies
- Late Phase expertise in both interventional and non-interventional type studies, including Registries

Experience:

- Managed several trials with over 8000 patients
- Depth of knowledge and experience working with global regulatory authorities
- Broad experience in both drug and device trials
- Phase I-IV trial experience with coverage in over 45 countries

Execution:

- Ongoing relationships with investigative sites and patient advocacy groups with proven track records and years of experience in the execution of clinical trials for women's health disorders
- Full-service approach to drug development (Visit Medpace.com for a full list of comprehensive services.)
- Fully-integrated, web-based decision support system provides IWRS/IVRS, study drug management, data management, laboratory management, and image management needs

Women's Health requires a different viewpoint.

Medpace has experience conducting trials that affect every stage of a woman's life. From pregnancy to menopause, Medpace understands the complexity and unique challenges faced when managing these studies. Supported by physicians specializing in oncology, metabolic disorders, endocrinology and cardiology, Medpace has the medical expertise to support all indications within women's health drug/device development.

List of indications:

- Obstetrics
- Menstrual Disorders
- Endometrial Ablation Therapy
- Osteoporosis
- Hormone Therapy
- Incontinence
- Pelvic Organ Prolapse
- Sexual Disorders
- Breast Cancer
- Gynecologic Cancers

Site Relationships

Site selection can be a challenge for women's health trials. Medpace has developed relationships with several investigators, site networks and hospitals, specializing in women's health disorders. In addition, Medpace works with patient advocacy groups to increase the visibility of the study and generate interest in participation for both investigators and women within the patient population.

Integrated Project Teams

Over the past 13 years, Medpace has been involved in trials focused in women's health. This experience is a direct result of well-established Sponsor relationships, exceptional clinical operations, and committed study teams. Medpace has built a team that includes medical experts and several Clinical Trial Managers who have been involved with the management and execution of women's health trials from the Sponsor/CRO project management role. All team members are involved from project initiation to completion, producing truly seamless drug development.

Real-time System Support

Fully integrated real-time information is supported with ClinTrak®, Medpace's proprietary web-based decision support system. ClinTrak provides IWRS/IVRS, study management, data management, laboratory management, image management, and endpoint adjudication.

Who We Are

Medpace is a global full-service clinical research organization providing Phase I-IV core development services for drug, biologic, and device programs. With medical and regulatory expertise in multiple therapeutic specialties, Medpace has assembled experienced and therapeutically focused teams to execute at every level of the company's operations, providing complete and seamless drug development services.