



# Clinical Trials of Texas, Inc.

Dedicated to the advancement of new medical therapies.

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## Curriculum Vitae

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#### EDUCATION

1978 Texas A & M University, College Station, Texas  
1982 University of Texas Health Science Center, San Antonio, Texas  
1982-1983 Internship – University of Texas Health Science Center at San Antonio, San Antonio, Texas  
1983-1986 Residency – University of Texas Health Science Center at San Antonio, San Antonio, Texas  
1988 Fellow – American Board of Obstetrics and Gynecology

#### HONORS

Residency – Administrative Chief Resident  
Medical School – Alpha Omega Alpha Honor Society

#### LICENSE

1983 Texas Medical Board, # G3450

#### CERTIFICATIONS AND TRAINING

1988 American Board of Obstetrics and Gynecology  
2007 Good Clinical Practices: Practical Application and Implementation

#### PROFESSIONAL ORGANIZATIONS

American College of Obstetrics and Gynecology  
Texas Medical Association  
Bexar County Medical Society  
San Antonio OB & GYN Society

#### PROFESSIONAL EXPERIENCE

1986-1987 Corpus Christi, Texas  
1987-Present San Antonio, Texas  
1996-Present Affiliated with Seven Oaks Women's Center, San Antonio, Texas  
1998-2000 Chairman, Department of Obstetrics and Gynecology, Southwest Texas Methodist Hospital

#### RESEARCH EXPERIENCE

2008-Present; "A Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study Evaluating the Safety and Efficacy of XXXXX Vaginal Cream in the Treatment of Mixed Bacterial Vaginosis/Vulvovaginal Candidiasis Infections"

2008- Present; "A Multicenter, Randomized, Double-Blind, Parallel Group Study to Evaluate the Efficacy and Safety of Two Doses of XXXXX Versus Placebo in Women with Overactive Bladder"

2007-Present; "A Randomized, Placebo-Controlled Phase II Study of Multiple Dosing Regimens of Intravaginally Administered XXXXX Gel for the Treatment of Cervical High Risk HPV Infection"

2007-Present; "A Multi-Center, Randomized, Controlled Study to Investigate the Safety and Tolerability of Intravenous XXXXX vs. Standard Medical Care in Treating Iron Deficiency Anemia in Heavy Uterine Bleeding and Postpartum Patients"

2007-Present; "A Randomized, Double-Blind, Placebo-Controlled, Multicenter, 52-week Study to Evaluate the Endometrial Safety of Transdermal XXXXX (300 mcg/day) in Naturally Postmenopausal Women with Hypoactive Sexual Desire Disorder"

2007-Present; "A Randomized, Double-Blind, Placebo-Controlled, Dose-Ranging Multicenter Evaluation of the Use of Topically Administrated XXXXX versus Placebo in Subjects with Pain Associated with Fibrocystic Breast Disease"

2007-Present; "A Phase II, Three-Arm, Parallel Design, Dose-Ranging Placebo-Controlled, Randomized, Double-Blind, Multicenter Study Evaluating the Safety and Efficacy of the XXXXX in the Treatment of Premenopausal Women with Symptomatic Endometriosis"

2007-Present; "An Open Label Study of the Contraceptive Efficacy of an Extended Regimen of XXXXX and XXXXX"

2006-2007; "A Prospective, Multicenter, Double-Blinded, Randomized Study to Evaluate Bleeding Patterns in Women Using One of Three Different Ascending EE Dose Extended Cycle (91-Day) Oral Contraceptive Regimens XXXXX Compared to Seasonale® Oral Contraceptive Regimen"

2006-Present; "A 24-Week, Randomized, Double-Blind, Placebo Controlled, Safety and Efficacy Trial of XXXXX 50 and 100 Milligrams Each Evening in Premenopausal Women With Hypoactive Sexual Desire Disorder"

2006-Present; "A Twelve Month, Open-Label, Safety Trial of XXXXX 50 milligrams to 100 milligrams Daily in Women with Hypoactive Sexual Desire Disorder"

2006-2007; "Efficacy and Safety of XXXXX in the Treatment of Vulvar and Vaginal Atrophy (VVA) in Postmenopausal Women: A 12-Week, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study Comparing Oral XXXXX 30 mg and 60 mg Daily Doses with Placebo"

2006-Present; "A Double-Blind, Randomized, Placebo- and Active-Controlled Efficacy and Safety Study of XXXXX Combinations for Prevention of Endometrial Hyperplasia and Prevention of Osteoporosis in Postmenopausal Women"

2006-2007; "A Prospective, Multicenter, Open-Label Study to Evaluate the Safety and Efficacy of the 28-Day Oral Contraceptive"

2006-2007; "Double-Blind, Randomized, Placebo-Controlled Study to Evaluate the Safety and Efficacy of 75 mg and 150 mg of XXXXX on the Reduction of Symptoms Associated with Endometriosis During Treatment and Post Treatment in Reproductive-Aged Women"

2005-2007; "A Phase 2, Multicenter, Double-Blind, Randomized, Placebo-Controlled Study to Evaluate Two Doses of a XXXXX Vaginal Ring for the Management of Moderate to Severe Endometriosis-Related Nonmenstrual Pelvic Pain"

2005-2006; "Comparison of the Safety and Efficacy of a XXXXX versus Oral Iron in the Treatment of Iron Deficiency Anemia Secondary to Heavy Uterine Bleeding"

2005-2006; "Comparison of the Safety and Efficacy of a XXXXX versus Oral Iron in Subjects Who Display Postpartum Anemia"

2005-2006; "A Multicenter, Double-Phase, Randomized, Double-Blind, Placebo Controlled (12-Week Double-Blind Followed By 12-Week Open-Label) Study Evaluating the Effect of XXXXX on Urgency Urinary Incontinence (UUI), Urgency, Frequency, Sexual Quality Of Life And Sexual Function In Women With Overactive Bladder"

2004-2006; "A Multicenter, Double-Blind, Placebo-Controlled, Parallel-Group Study Evaluating the Efficacy and Safety of XXXXX for the Treatment of Vasomotor Symptoms of Menopause in Postmenopausal Women"

2003-2004; "A Study of the Safety and Efficacy of XXXXX in the Treatment of Vaginal Atrophy in Postmenopausal Women"

2002-2004; "A Study to Demonstrate Immunogenicity and Tolerability of the XXXXX Vaccine in Pre-Adolescents and Adolescents"

2001-2002; "A Study of the Safety and Efficacy of XXXXX in the Treatment of Vaginal Atrophy in Postmenopausal Women"

2001-2004; "A Double-Blind Placebo-Controlled, Parallel Group Design Study of Two Doses of XXXXX vs. Placebo for the Treatment of Sexual Dysfunction (Hypoactive Desire) in Postmenopausal Women"

2001-2004; "A Double-Blind Placebo-Controlled, Parallel Group Design Study of Two Doses of XXXXX vs. Placebo for the Treatment of Sexual Dysfunction (Arousal Disorder) in Postmenopausal Women"

2001-2004; "A Randomized, Open-Label 16-Week Study Comparing Breakthrough Bleeding Profiles of Women on XXXXX 1/5 or XXXXX"

1998-2001; "A Randomized, Multi-Center, Open-Label, Dose-Ranging Study Comparing the Safety and Efficacy of XXXXX to XXXXX 3.75 mg in Women with Endometriosis-Associated Pain"

The Treatment of Vasomotor Symptoms Associated with Menopause