



# CLINICAL TRIALS OF TEXAS, INC.

*Promoting Health Through Research*

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## Curriculum Vitae

### **Parke Joseph Hedges, MD, FACOG**

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San Antonio, Texas 78229

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San Antonio, Texas 78229

#### **EDUCATION**

1983-1986 Residency, University of Texas Health Science Center at San Antonio, San Antonio, Texas  
1982-1983 Internship, University of Texas Health Science Center at San Antonio, San Antonio, Texas  
1982 Doctor of Medicine, University of Texas Health Science Center, San Antonio, Texas  
1978 Texas A & M University, College Station, Texas

#### **LICENSE**

1983 Texas Medical Board, # G3450

#### **CERTIFICATION(S)**

1988 American Board of Obstetrics and Gynecology

#### **PROFESSIONAL EXPERIENCE**

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

#### **PROFESSIONAL MEMBERSHIPS**

American College of Obstetrics and Gynecology, 1988  
Texas Medical Association  
Bexar County Medical Society  
San Antonio OB & GYN Society

#### **HOSPITAL AFFILIATION(S)**

Southwest Texas Methodist

#### **RESEARCH AFFILIATION(S)**

2003-Present Investigator, Clinical Trials of Texas, Inc., San Antonio, Texas

#### **RESEARCH EXPERIENCE**

2009-Present: A Twenty-Four Week, Randomized, Double-Blind, Placebo Controlled, Safety And Efficacy Trial Of "Study Drug" (100 Milligrams) Administered Orally Once Daily In Premenopausal Women With Hypoactive Sexual Desire Disorder In The United States

2009-Present: A Twenty-Four Week, Randomized, Double-Blind, Placebo-Controlled, Safety And Efficacy Trial Of "Study Drug" (100 Milligrams) Administered Orally Once Daily In Naturally Postmenopausal Women With Hypoactive Sexual Desire Disorder In The United States

2009-Present: A Multi-Center, Randomized, Active Controlled Study To Investigate The Efficacy And Safety Of Intravenous "Study Drug" In Patients With Iron Deficiency Anemia (IDA)

2009-Present: A Phase 2, Randomized, Double-Blind, Placebo-Controlled Study To Assess The Efficacy And Safety Of "Study Drug" In Subjects With Endometriosis

2009-2009: Psychometric Evaluation Of The "Study Questionnaire" In Pre- And Post-Menopausal Women With And Without Hypoactive Sexual Desire Disorder (HSDD)

2008-Present: A Multicenter, Randomized, Double-Blind, Parallel Group Study To Evaluate The Efficacy And Safety Of Two Doses Of "Study Drug" Versus Placebo In Women With Overactive Bladder

2008-Present: A Multi-Center, Randomized, Controlled Study To Investigate The Safety And Tolerability Of Intravenous "Study Drug" Vs. Standard Medical Care In Treating Iron Deficiency Anemia

2008-Present: A Multi-Center, Randomized, Controlled Study To Investigate The Safety And Tolerability Of A Single Dose Of Intravenous "Study Drug" Vs. Standard Medical Care In Treating Iron Deficiency Anemia In Subjects Who Are Not Dialysis Dependent

2008-2009: A Phase II, Randomized, Double-Blind, Placebo-Controlled Study To Assess The Efficacy And Safety Of "Study Drug" In Subjects With Endometriosis

2008-2009: Efficacy And Safety Of "Study Drug" In The Treatment Of Moderate To Severe Vaginal Dryness And Vaginal Pain Associated With Sexual Activity, Symptoms Of Vulvar And Vaginal Atrophy (VVA), Associated With Menopause: A 12-Week, Randomized, Double-Blind, Placebo-Controlled, Parallel Group Study Comparing Oral "Study Drug" Daily Dose With Placebo In Postmenopausal Women

2008-2009: Open-Label Study Of "Study Drug" Oral Daily Dose, In The Treatment Of Moderate To Severe Vaginal Dryness And Pain Associated With Sexual Activity, Symptoms Of Vulvar And Vaginal Atrophy (VVA) Associated With Menopause: A Follow-Up To "Protocol"

2008-2009: A Multi-Center, Placebo Controlled, Safety And Efficacy Study Of The Selective "Study Drug" In Anemic, Pre-Menopausal Women With Symptomatic Uterine Fibroids Requiring Hysterectomy

2008-2009: A Phase III, Three-Arm, Parallel Design, Placebo-Controlled, Randomized, Double-Blind, Multicenter Study Evaluating The Safety And Efficacy Of "Study Drug" In The Treatment Of Premenopausal Women With Symptomatic Uterine Fibroids

2007-2009: A Multi-Center, Randomized, Controlled Study To Investigate The Safety And Tolerability Of Intravenous "Study Drug" Vs. Standard Medical Care In Treating Iron Deficiency Anemia In Heavy Uterine Bleeding And Postpartum Patients

2007-2009: A Randomized, Double-Blind, Placebo-Controlled, Multicenter, 52-Week Study To Evaluate The Endometrial Safety Of Transdermal "Study Drug" (300 Mcg/Day) In Naturally Postmenopausal Women With Hypoactive Sexual Desire Disorder

2007-2009: An Open Label Study Of The Contraceptive Efficacy Of An Extended Regimen Of "Study Drug" And "Study Drug"

2007-2008: A Randomized, Placebo-Controlled Phase II Study Of Multiple Dosing Regimens Of Intravaginally Administered "Study Drug" Gel For The Treatment Of Cervical High Risk HPV Infection

2007-2008: Long-Term Safety Of "Study Drug" 60 mg Oral Daily Dose For The Treatment Of Vulvar And Vaginal Atrophy (VVA) In Postmenopausal Women Without A Uterus: A 52-Week Open-Label Follow-Up To "Protocol"

2007-2008: A Randomized, Double-Blind, Placebo-Controlled, Dose-Ranging Multicenter Evaluation Of The Use Of Topically Administrated "Study Drug" Versus Placebo In Subjects With Pain Associated With Fibrocystic Breast Disease

2007-2008: A Phase II, Three-Arm, Parallel Design, Dose-Ranging Placebo-Controlled, Randomized, Double-Blind, Multicenter Study Evaluating The Safety And Efficacy Of The “Study Drug” In The Treatment Of Premenopausal Women With Symptomatic Endometriosis

2006-2009: A Twelve Month, Open-Label, Safety Trial Of “Study Drug” 50 Milligrams To 100 Milligrams Daily In Women With Hypoactive Sexual Desire Disorder

2006-2008: A 24-Week, Randomized, Double-Blind, Placebo Controlled, Safety And Efficacy Trial Of “Study Drug” 50 And 100 Milligrams Each Evening In Premenopausal Women With Hypoactive Sexual Desire Disorder

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 “Study Drug” Compared To “Study Drug” Oral Contraceptive Regimen

2006-2007: Efficacy And Safety Of “Study Drug” 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 “Study Drug” 30 Mg And 60 Mg Daily Doses With Placebo

2006-2007: A Double-Blind, Randomized, Placebo- And Active-Controlled Efficacy And Safety Study Of “Study Drug” 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 “Study Drug” 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 “Study Drug” Vaginal Ring For The Management Of Moderate To Severe Endometriosis-Related Nonmenstrual Pelvic Pain

2005-2006: Comparison Of The Safety And Efficacy Of A “Study Drug” 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 “Study Drug” 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 “Study Drug” 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 “Study Drug” For The Treatment Of Vasomotor Symptoms Of Menopause In Postmenopausal Women

2003-2004: A Study Of The Safety And Efficacy Of “Study Drug” In The Treatment Of Vaginal Atrophy In Postmenopausal Women

2002-2004: A Study To Demonstrate Immunogenicity And Tolerability Of The “Study Drug” Vaccine In Pre-Adolescents And Adolescents

2001-2002: A Study Of The Safety And Efficacy Of “Study Drug” In The Treatment Of Vaginal Atrophy In Postmenopausal Women

2001-2004: A Double-Blind Placebo-Controlled, Parallel Group Design Study Of Two Doses Of “Study Drug” 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 "Study Drug" 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 "Study Drug" 1/5 Or "Study Drug"

1998-2001: A Randomized, Multi-Center, Open-Label, Dose-Ranging Study Comparing The Safety And Efficacy Of "Study Drug" To "Study Drug" 3.75 Mg In Women With Endometriosis-Associated Pain

The Treatment Of Vasomotor Symptoms Associated With Menopause

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Signature and Date