



CLINICAL TRIALS OF TEXAS, INC.

Promoting Health Through Research

Curriculum Vitae

Brian Waid Harle, MD, FACOG

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EDUCATION

1994 Fellow – American Board of Obstetrics and Gynecology
1989-1992 Residency – Department of OB-GYN, University of Texas Medical Branch, Galveston, Texas
1988-1989 Intern – Department of OB-GYN, University of Texas Medical Branch, Galveston, Texas
1986-1988 Texas Tech University School of Medicine, Lubbock, Texas
1984-1986 Howard University College of Medicine, Washington, D.C.
1980-1984 Texas A & M University, College Station, Texas

LICENSE

1988 Texas Medical Board, # H6274

CERTIFICATIONS

1994 Diplomate, American Board of OB-GYN

PROFESSIONAL EXPERIENCE

1992-Present Private Practice, San Antonio, Texas
1996-Present Affiliated with Seven Oaks Women's Center, San Antonio, Texas

PROFESSIONAL MEMBERSHIPS

American College of Obstetrics and Gynecology
Willard R. Cooke Obstetrics and Gynecology Society
Texas Association of Obstetrics and Gynecology
Texas Medical Association
Bexar County Medical Society

HOSPITAL AFFILIATIONS

1992-Present Southwest Texas Methodist Hospital, San Antonio, Texas

RESEARCH AFFILIATIONS

2001-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 Sexual Desire Distress Questionnaire[®] In Pre- And Post-Menopausal Women With And Without Hypoactive Sexual Desire Disorder (HSDD)

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-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"

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

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

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: 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: An Open Label Study Of The Contraceptive Efficacy Of An Extended Regimen Of “Study Drug” And “Study Drug”

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

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 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

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 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

2006-2008: 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-Comtrolled, Parallel-Group Study Comparing Oral “Study Drug” 30 mg And 60 mg Daily Doses With Placebo

2006-2007: A Prospective, Multicenter, Open-Label Study To Evaluate The Safety And Efficacy Of The 28-Day Oral Contraceptive

2006-2007: A Double-Blind, Randomized, Placebo-Controlled Study To Evaluate The Safety And Efficacy Of “Study Drug” On The Reduction Of Symptoms Associated With Endometriosis During Treatment And Post Treatment In Reproductive-Aged Women

2005-2007: An 18-Month Study To Assess The Efficacy, Safety, And Tolerability Of “Study Drug” In Obese Patients

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: A Multi-Center, Randomized, Blinded, Placebo Controlled, Cross-Over Study To Investigate The Safety And Tolerability Of “Study Drug” In Patients With Iron Deficiency Anemia

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 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

2001-2003: A Study Of The Safety And Efficacy Of "Study Drug" In The Treatment Of Vaginal Atrophy In Postmenopausal Women

2001-2003: A Randomized, Open-Label, 16-Week Study Comparing Breakthrough Bleeding Profiles Of Women On "Study Drug" 1/5 Or "Study Drug"

2001-2003: Female Sexual Dysfunction (2 Separate Protocols: Hypo-Arousal And Hypo-Desire)

Signature

Date