



CLINICAL TRIALS OF TEXAS, INC.

Promoting Health Through Research

Curriculum Vitae

Christine Marie De La Garza, MD, FACOG

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EDUCATION

- 1988-1992 Internship and Residency: Obstetrics & Gynecology, Bexar County Hospital District, San Antonio, Texas
- 1988 Doctor of Medicine, The University of Texas Health Science Center, San Antonio, Texas
- 1984 Masters of Science in Nursing: Medical Surgical Nursing, The University of Texas Health Science Center, San Antonio, Texas
- 1983 The University of Texas at San Antonio, San Antonio, Texas
- 1980 Incarnate Word College, San Antonio, Texas
- 1979 University of Illinois Circle Campus, Chicago, Illinois
- 1979 Bachelors of Science in Nursing: with distinctions, University of Minnesota, Minneapolis, Minnesota

LICENSE

- 1989 Texas Medical Board, H6156

CERTIFICATIONS

- 1994 American Board of Obstetrics & Gynecology

PROFESSIONAL EXPERIENCE

- 1996-Present Private Practice, Seven Oaks Women's Center, San Antonio, Texas
- 1992-1996 Private Practice in Obstetrics and Gynecology, San Antonio, Texas

Staff Nurse: Neonatal Intensive Care Unit, Bexar County Hospital District, San Antonio, Texas

PROFESSIONAL MEMBERSHIPS

American College of Obstetrics & Gynecology
Texas Medical Association
Bexar County Medical Society

HOSPITAL AFFILIATIONS

Southwest Texas Methodist

RESEARCH AFFILIATIONS

- 2004-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" 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" 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 "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 "Study Drug" 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

2006-2008: 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

2006-2008: 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

2006-2007: 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-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

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

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

2005-2007: The Safety And Efficacy Of A “Study Drug” In The Treatment Of Postpartum 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

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

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