



Clinical Trials of Texas, Inc.

Dedicated to the advancement of new medical therapies.

Curriculum Vitae

Ronald Anthony Valdez, MD

7940 Floyd Curl Drive, Suite 700
San Antonio, Texas 78229

7940 Floyd Curl Drive, Suite 900
San Antonio, Texas 78229

EDUCATION

- 1984-1988 Residency in Obstetrics and Gynecology, Texas Tech School of Medicine, El Paso, Texas
1983-1984 Rotating Internship, Texas Tech School of Medicine, El Paso, Texas
1983 Doctrine of Medicine Degree, University of Texas Medical Branch, Galveston, Texas
1979 Bachelors of Arts Degree: Graduated Magna Cum Laude, St. Mary's University, San Antonio, Texas

LICENSURE

Texas Medical Board, # G5722

CERTIFICATIONS

- 1993 Fellow of the American College of Obstetrics and Gynecology
1991 The American Board of Obstetrics and Gynecology

PROFESSIONAL ORGANIZATIONS

Bexar County Medical Society
Texas Medical Association
American College of OB/GYN

HOSPITAL APPOINTMENTS

- 1999-2001 Chairman – Department of Obstetrics and Gynecology, Baptist Hospital System
1997-Present Member, Audit Policy Committee, Baptist Hospital System
1997-1998 Chief of Services - Department of Gynecology, St. Luke's Hospital
1996-Present Board of Directors, Health Corp
1996 Founding Partner, Institute For Women's Health
1995 President, Mexican American Physicians Association
1994-1996 Medical Care Advisory Committee, Department of Human Services
1994-1996 Admissions Committee, University of Texas Health Science Center, San Antonio
1994-1996 Obstetrics and Gynecology Advisory Committee, Pacificare of Texas
1991-1994 Executive Committee, Mexican American Physician Association
1991-1994 Chairman-Scholarship Committee, Mexican American Physician Association
1991-1999 Chairman, Utilization Management Committee Select OB/GYN Associates (SOGA)

RESEARCH AFFILIATION

2006-Present Investigator, Clinical Trials of Texas, Inc., San Antonio, Texas

PROFESSIONAL EXPERIENCE

1996-Present Institute for Women's Health, San Antonio, Texas
1990-1996 Live Oak OB/GYN, San Antonio, Texas
1988-1990 Private Practice, Beeville, Texas

RESEARCH EXPERIENCE

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

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

2008-Present: 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-Present: 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" 60 MG Daily Dose With Placebo In Postmenopausal Women

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

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

2007-2008: Long-Term Safety Of 30 MG And 60 MG Oral Daily Doses Of "Study Drug" In The Treatment Of Vulvar And Vaginal Atrophy (VVA) In Postmenopausal Women With An Intact Uterus: A 40-Week Randomized, Double-Blind, Placebo-Controlled, Follow-Up To "Protocol"

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


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

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

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

2006-2008: 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: 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


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