



CLINICAL TRIALS OF TEXAS, INC.

Promoting Health Through Research

Curriculum Vitae

Jose Manuel Ruiz III, MD

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EDUCATION

- 1986-1990 Residency in OB/GYN, Saint Luke's Roosevelt Hospital Women's Hospital, New York City, New York, Dr. Robert Neuwirth, Chairman
1986 Doctor of Medicine Degree, Louisiana State University School of Medicine, New Orleans, Louisiana
1982 Bachelors of Science Degree in Zoology, Louisiana State University, Baton Rouge, Louisiana

UNIVERSITY APPOINTMENT

- 1996-1998 Interviewer Admissions Committee, University of Arizona Health Science Center
1994-1998 Clinical Assistant Professor - Dept. of OB/GYN, University Medical Center, Tucson, Arizona

LICENSE

- 1998 Texas Medical Board, # K4993

CERTIFICATION(S)

- 1992 The American College of Obstetrics and Gynecology

PROFESSIONAL EXPERIENCE

- 1998-Present Institute for Women's Health, San Antonio, Texas
1990-1998 El Rio OB/GYN Associates, Tucson, Arizona

PROFESSIONAL MEMBERSHIPS

Diplomat American Board of OB/GYN
Fellow of the American College of OB/GYN
Pima County Medical Society
Southwest Society of OB/GYN
Tucson OB/GYN Society
Chairman of OB/GYN Dept, St. Mary's Hospital 1994-1996
Bexar County Medical Society

HOSPITAL AFFILIATIONS

St. Lukes Baptist Hospital, San Antonio, Texas

RECENT PAPERS PRESENTED AND PUBLISHED

"The Incidence of Complications Associated with the Use of Hyskon During Hysteroscopy: Experience in 1,703 consecutive patients", Journal of Gynecologic Surgery

RESEARCH AFFILIATIONS

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

RESEARCH EXPERIENCE

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

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 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: 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

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

2007-2009: 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

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

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"

2007-2008: 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

2007-2008: 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

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

2000: Ortho Evra- Birth Control Study

1997: Ditropan- Urge Incontinence Study

1997: Climara Hormone Patch Study

1996: Vagifem For Contraception

1994: Cox II Inhibitor (Celebrex) For Post-Op Pain In GYN Patients

1992-1998: Additional Contraceptive Study With Various Agent

Signature and Date