



# Clinical Trials of Texas, Inc.

Dedicated to the advancement of new medical therapies.

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## Curriculum Vitae

### **Karen Maria Carcamo, MD, MPH**

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#### **EDUCATION**

- 1992-1996      Residency- Obstetrics and Gynecology, University of Texas Medical Branch,  
Galveston, Texas
- 1992            Doctor of Medicine, University of California at San Francisco, San Francisco, California
- 1988            Masters of Public Health, University of California at Los Angeles, Los Angeles,  
California
- 1985            Bachelors of Arts in Chemistry and Spanish, University of New Mexico, Albuquerque,  
New Mexico

#### **HONORS**

- 1988      UCLA SPH Alumni Association Outstanding Graduate Award
- 1988      UCLA SPH Health Career Opportunities Program Student of the Year Award
- 1985      Phi Beta Kappa

#### **PROFESSIONAL ORGANIZATIONS**

American College of Obstetrics and Gynecology  
Diplomat American Board of OB/GYN  
Bexar County Medical Society  
Fellow of the American College of OB/GYN

#### **LICENSE**

Texas Medical Board, # K0663  
Pennsylvania Medical License, # 0457796-L

#### **RESEARCH AFFILIATION**

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

#### **PROFESSIONAL EXPERIENCE**

- 1999-Present    Institute for Women's Health, San Antonio, Texas
- 1997-1999       Delos Rive Walk OB/GYN Associates, San Antonio, Texas
- 1996-1997       MacGregor Medical Association, Houston, 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



1/30/09