



# CLINICAL TRIALS OF TEXAS, INC.

*Promoting Health Through Research*

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## Curriculum Vitae

### **Paula Bilica, DO**

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San Antonio, Texas 78229

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#### **EDUCATION**

2004-2008 OB-GYN Residency, Jersey Shore University Medical Center, Neptune, New Jersey  
2004 Doctor of Osteopathic Medicine, Kirksville College of Osteopathic Medicine, Kirksville, Missouri  
1999 Bachelors of Science in Psychology, Southwest Texas State University, San Marcos, Texas  
1995 Austin Community College, Austin, Texas

#### **LICENSE**

2008 Texas Medical Board, # M8535

#### **CERTIFICATIONS**

ACLS Certified  
Neonatal Resuscitation Program (NRP) Certified

#### **PROFESSIONAL EXPERIENCE**

2008-Present Group Practice, Institute for Women's Health, San Antonio, Texas  
2000-2000 Office Manager, Kelly Hawkins Physical Therapy Clinic  
1995-1999 Medical Assistant, Dr. T.A. Castoldi and Associate, San Marcos, Texas

#### **PROFESSIONAL MEMBERSHIPS**

ACOG Junior Fellow  
American Society for Colposcopy and Cervical Pathology

#### **HOSPITAL AFFILIATIONS**

Northcentral Baptist Hospital, San Antonio, Texas

#### **RESEARCH AFFLIATION(S)**

2009-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 II, Randomized, Double-Blind, Placebo-Controlled Study To Assess The Efficacy And Safety Of "Study Drug" In Subjects With Endometriosis

2009-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

2009-2009: 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

2009-2009: 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

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

2009-2009: 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-Extension Study

2009-2008: 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

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Signature and Date