

## CURRICULM VITAE

**Steven Winger, M.D. CPI**

Precision Trials  
3815 East Bell Road, Suite 4500  
Phoenix, AZ 85032  
602-931-4507

### **Education:**

- 1980 Oberlin College/Conservatory of Music  
Bachelor Degree in Music
- 1984 Post Graduate Studies  
University of Massachusetts at Amherst
- 1989 Tufts University School of Medicine  
Medical Doctor  
Boston, MA

### **Postgraduate Training:**

- 1989-1990 Internship, Pennsylvania Hospital  
Philadelphia, PA
- 1990-1993 Resident in OB/GYN Pennsylvania Hospital  
Philadelphia, PA

### **Professional Experience:**

- 2006-present Principal Investigator/ Physician/ Medical Director  
Precision Trials (formerly Arizona Wellness Center for Women  
Research Department)  
Phoenix, AZ
- 1998-present Obstetrics & Gynecology/Investigator/Medical Director  
Arizona Wellness Center for Women  
Phoenix, AZ
- 1995-1998 Obstetrics & Gynecology/Investigator  
Obstetrics and Gynecology Consultants, Ltd  
Phoenix, AZ

**Steven Winger, MD CPI**  
**Curriculum Vitae**  
**Page 2**

**Professional  
Experience  
Continued:**

1993-1995                      Obstetrics & Gynecology  
   Seaton Obstetrics and Gynecology  
   Boston, MA

**Academic Affiliations:**

Assistant Professor  
Tufts University School of Medicine  
Boston, MA

Assistant Instructor in Obstetrics & Gynecology  
University of Pennsylvania School of Medicine  
Philadelphia, PA

Department Chairman in OB GYN  
Paradise Valley Hospital, Phoenix, AZ

**Professional  
Memberships:**

American Associates of Gynecologic Laparoscopists  
American College of Obstetrics & Gynecology  
Massachusetts Medical Society  
Maricopa Medical Society

**Hospital Appointments:**

2004 Board of Directors  
Paradise Valley Hospital  
Phoenix, AZ

**Hospitals:**

Paradise Valley Hospital  
Scottsdale Healthcare

**Steven Winger, MD CPI**  
**Curriculum Vitae**  
**Page 3**

**Licenses/Certifications:**

Arizona #23378  
Massachusetts # 77483  
Pennsylvania # MD044992L

**Certification:**

American Board of Obstetrics & Gynecology 1997  
Certified Clinical Research Investigator (CCTI) ACRP 2005

**Honors/Awards:**

1988 – McGraw Hill Award for Outstanding Contribution and Academic Achievement  
1993 - Resident Training Award, University of Pennsylvania School of Medicine  
1993 - CREOG Award, Pennsylvania School of Medicine

**Publications:**

Prenatal Diagnosis Vol. 14:839-843 (1994)  
Syndromes identified with fetuses with prenatally diagnosed cephaloceles

K. Candiotti, N. Singla, S. Winger, H. Minkowitz, J. Breitmeyer  
Poster 2008. Study Results of Intravenous Acetaminophen over 48 hours for the  
treatment of Postoperative Pain after Gynecologic Surgery

K. Candiotti, N Singla, S. Winger, H. Minkowitz, M Royal  
Poster 2008. Study Results of Intravenous Acetaminophen over 48 hours for the  
treatment of Postoperative Pain after Gynecologic Surgery

Fertility and Sterility Vol. No. 2008.  
J. Simon, MD, K Reape, MD., S. Winger, MD., H. Hait MS.  
Clinical Article on Synthetic Conjugates Estrogens for the treatment of Vulvovaginal  
Atrophy in healthy Postmenopausal Women.

**Research Experience:**

A Phase III Open-Label, Single Arm Study to Assess the Safety of XXXXX Injection for  
Minimal-to-Moderate Sedation in patients undergoing Minor Surgical Procedures.

**Steven Wininger, MD CPI**  
**Curriculum Vitae**  
**Page 4**

A Phase III Randomized, Double-Blind, Placebo-Controlled, Multi-Center, Parallel-Group, Multiple-Dose study of the XXXXX Efficacy and Safety of Intravenous XXXXX versus Placebo over 48 hours for the Treatment of Postoperative Pain After Gynecologic Surgery

A Phase II, Multi-centered, Randomized, Placebo-controlled, Double-blind study of the XXXXX in pre-menopausal women with symptomatic leiomyomata

A twenty-four week, randomized, double-blind, placebo-controlled, safety and efficacy trial of XXXXX daily and, with XXXXX daily in Premenopausal women with hypoactive sexual desire disorder.

A Randomized, Multi-center, Double-Blind, Placebo-Controlled Trial, to demonstrate the safety and efficacy of Daily XXXXX and XXXXX daily for the treatment of vasomotor symptoms in postmenopausal women.

A 12 month, Open Label, Multi-center Study to Evaluate the safety of XXXXX Formulation administered three times daily for women with heavy menstrual bleeding associated with menorrhagia.

A multi-center, randomized, double-blind phase III study of the efficacy and safety of XXXXX compared to XXXXX for women requiring cervical ripening and induction of labor.

A phase 3, randomized, double blind, dose-controlled study to assess the efficacy and safety of XXXXX injection for minimal-to-moderate sedation in patients undergoing flexible Bronchoscopy.

A multi-center, double-blind, randomized, placebo-controlled study of XXXXX in the treatment of High-Grade cervical intraepithelial lesions of the uterine cervix.

A Phase II, multi-center, double-blind, randomized, placebo-controlled study to evaluate induction of Withdrawal bleeding after administration of XXXXX in women with secondary amenorrhea.

A phase II b multi-centered, double-blind, placebo-controlled, parallel-group, dose-ranging study evaluating the efficacy and safety of XXXXX for the treatment of moderate to severe vasomotor symptoms associated with menopause.

A Randomized, multi-center, double-blind, placebo-controlled trial to compare the effects of 12 weeks of treatment with XXXXX vs. XXXXX on vulvovaginal atrophy in healthy post menopausal women.

**Steven Winger, MD CPI**  
**Curriculum Vitae**  
**Page 5**

A phase II, multi-center, double-blind, randomized, placebo-controlled study to evaluate two doses of XXXXX for the management of moderate to severe endometriosis related non-menstrual pelvic pain.

A Pilot clinical study to evaluate the efficacy of XXXXX in the treatment of the signs and symptoms of atrophic Vaginitis.

Evaluating the Safety and Efficacy of XXXX in the treatment of Vaginal Atrophy in Postmenopausal Women

Evaluating the Safety & Efficacy of XXXX, vaginal cream compared with Metro Gel vaginal cream for the treatment of Bacterial Vaginosis

Evaluating the Safety & Efficacy of XXX, combination pack in bedtime versus daytime administration

Evaluating the Safety & Efficacy of XXXX, vaginal ointment, plus XXXX external vulvar cream of XXX, vaginal cream in the Treatment of Vulvovaginal candidiasis.

Evaluate the Treatment of Women with Menorrhagia with Intrauterine XXXX Ablation

Evaluate the dosing optimization study of XXXX for Endometriosis

Evaluate the Safety & Efficacy of Triphasic Oral Contraception

Evaluating the Safety & Efficacy in a double-blinded study of XXXX versus Placebo in Subjects with Endometriosis

Evaluating the Safety & Efficacy of XXXX estrogen combination in Postmenopausal Women

Evaluating XXXX for Contraceptive use

Evaluating XXX in Women using XXXX to Depo-Provera for Contraception

Evaluate the Safety & Efficacy of XXXX in the ovulatory, endocrine and metabolic abnormalities of Women with Polycystic Ovary Syndrome

Evaluation of XXXX vs hysterectomy in Safety & Efficacy trial for Permanent Female Sterilization

**Steven Wininger, MD CPI**  
**Curriculum Vitae**  
**Page 6**

A Multicenter, Randomized, Double-Blind, Study to Evaluate the Safety And Efficacy of XXXX versus XXXX versus Placebo in Women Diagnosed with Overactive Bladder Who Have Symptoms of Predominant or Pure Urge Incontinence

A Randomized, Multicenter, Double-Blind, Placebo-Controlled Trial to Demonstrate the Safety and Efficacy of XXXX for the Treatment of Vasomotor Symptoms in Postmenopausal Women

A Randomized, Multicenter, Double-Blind, Placebo-Controlled Trial to Compare the Efficacy and Safety of XXXX vs. XXXX Vulvovaginal Atrophy in Healthy Postmenopausal Women

A Randomized, Multicenter, Double-Blind, Placebo Controlled Trial to Evaluate the Efficacy and Safety of XXXX for the Treatment of Vulvovaginal Atrophy in Healthy Postmenopausal Women

A Phase II, Multicenter, Double-Blind, Randomized, Placebo Controlled Study to Evaluate Induction of Withdrawal Bleeding After Administration of XXXX in Women with Secondary Amenorrhea

A Randomized, MultiCenter, Double-Blind Study to Evaluate the Efficacy of XXXX Compared to Conventional Oral Contraceptive Therapy for the Treatment of Cyclic Pelvic Pain in Women

A Multicenter, Open-Label Study to Evaluate the Efficacy and Safety of an XXXX for the Prevention of Pregnancy in Women

A Multicenter, Randomized, Double Blind, Placebo-controlled, Dose-ranging, Parallel Group Phase II Study to Evaluate the Safety, Efficacy, and Pharmacokinetics of XXXX When Administered with XXXX for the Prevention of Post-operative Nausea and Vomiting (PONV) and Post-discharge Nausea and Vomiting (PDNV) in Female Subjects with Known Risk Factors for PONV Who are Undergoing Surgical Procedures Associated with an Increased Emetogenic Risk

A Randomized, Double-Blind, Multicenter, Parallel Group, Balanced, Stratified, Phase 3 Study to Evaluate the Efficacy and the Safety of XXXX Following Elective Abdominal or Gynecological Laparoscopic Surgery

Comparison of the Safety and Efficacy of XXXX in the Treatment of Iron Deficiency Anemia Secondary to Heavy Uterine Bleeding”

**Steven Winger, MD CPI**  
**Curriculum Vitae**  
**Page 7**

Comparison of the Safety and Efficacy of XXXX in Subjects who Display Postpartum Anemia

A Phase II, Randomized, Double-Blind, Placebo-Controlled Study of XXXXX in Endometriosis

A Phase II, Randomized, Double-Blind, Placebo-Controlled Twice-Daily Dosing Study of XXXX in Endometriosis

A Phase III, Multicenter, Randomized, Double-Blind, Placebo-Controlled Study of XXXX Breast Disease in Otherwise Healthy, Premenopausal Women

A 12-Month, Open Label, Multicenter Study to Evaluate the Safety of XXXX During the Menstrual Cycle in Women with Heavy Menstrual Bleeding Associated with Menorrhagia

A Double Blind, Randomized, Placebo Controlled, Efficacy and Safety of XXXX for the Treatment of Vasomotor Symptoms Associated with Menopause.

An Open-Label Study of the Safety and Efficacy of a New Low Dose Oral Contraceptive Containing XXXXX.

An Open Label Study of the Contraceptive Efficacy of an Extended Regimen of XXXXX and XXXXX.

A Multicenter, Open-Label Study to Evaluate the Efficacy and Safety of an Extended Cycle, Low Dose, Combination Oral Contraceptive Regimen XXXXX During the Seven Day Interval Between Each 84-Day Cycle of Combination Therapy For the Prevention of Pregnancy in Women

A Double-Blind, Randomized, Placebo-Controlled, Efficacy and Safety Study of XXXXX Combinations for Treatment of Vasomotor Symptoms associated with Menopause

Whole Blood Collection Protocol for XXXXX Clinical Trial in Women with Ovarian Tumors

A Phase II, 18-Week, Double-Blind, Placebo-Controlled, Multicenter Study Evaluating the Safety and Efficacy of XXXXX Combination Cream Compared with XXXXX Placebo Creams in the Treatment of Vulvar Vestibulitis Syndrome

**Steven Winger, MD CPI**  
**Curriculum Vitae**  
**Page 8**

A Phase II, Randomized, Double-Blind, Active-Controlled Study to Assess the Safety and Efficacy of XXXXX in Subjects with Endometriosis

A twelve month, open-label, safety trial of XXXXX daily in women with Hypoactive Sexual Desire Disorder.

A randomized, double-blind, placebo-controlled, multi-center, 52 week study to evaluate the endometrial safety of XXXXX in naturally postmenopausal women with hypoactive sexual desire disorder.

A Phase III Randomized, Double-Blind, Placebo-Controlled, Multi-Center, Parallel-Group, Multiple-Dose Study of the Analgesic Efficacy and Safety of Intravenous XXXXX Versus Placebo over 48 hours for the Treatment of Postoperative pain after Gynecologic Surgery

A Phase III Randomized, Double-Blind, Placebo-Controlled, Multi-Center, Parallel-Group, Repeated-Dose Study of the Analgesic Efficacy and Safety of Intravenous Acetaminophen versus Placebo for the Treatment of Postoperative Pain after Laparoscopic Surgery

A multi-center, open-label, randomized study to assess the safety and contraceptive efficacy of two doses of the ultra low dose XXXXX contraceptive intrauterine system for a maximum of 3 years in women 18 to 35 years of age

A multi-center, randomized, controlled study to investigate the safety and tolerability of XXXXX vs. standard medical care in treating Iron Deficiency Anemia in Heavy Uterine Bleeding and Post Partum patients.

Multi-center clinical study to determine the effectiveness and safety of HEDA in the screening of women for breast cancer

A Phase II single dose, blinded prospective study to investigate the efficacy, safety and Pharmacokinetics profile of the XXXXX in women following abdominal hysterectomy.

A Phase II clinical study evaluating the safety and efficacy of two regimens of XXXXX administered intravaginally for three months in women with moderate to severe pain associated with endometriosis.

Randomized, double-blind, parallel-group study evaluating the safety and efficacy of XXXXX in the treatment of mixed bacterial vaginosis/vulvovaginal candidiasis infections.

**Steven Wininger, MD CPI**  
**Curriculum Vitae**  
**Page 8**

A Phase III, Randomized, Double-blind, Placebo-controlled, Multi-center, Efficacy and Safety Study of XXXXX creams in the treatment of External Genital Warts.

Evaluation of the XXXXX HPV Test and the Linear Array High Risk HPV Genotyping Test for the Detection of High-Grade Cervical disease in Women Undergoing Routine Cervical Cancer Screening Using Cervical Samples Prepared With the Cobas x 421 Instrument

Updated: July 2008