

## CURRICULM VITAE

**Nancy Ann Bedrick Harris, MD**

Precision Trials  
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Phoenix, AZ 85032  
602-931-4507

### **Education:**

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| 1978-1982 | Brown University, Providence, RI<br>ScB – Neuroscience                     |
| 1982-1984 | University of Cincinnati College of Medicine<br>Cincinnati, OH             |
| 1984-1986 | Case Western Reserve University School of Medicine<br>Cleveland, OH – M.D. |

### **Postgraduate Training:**

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| 1986-1990 | Residency – Mt. Sinai Medical Center<br>Cleveland, OH<br>Obstetrics & Gynecology |
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### **Professional Experience:**

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| 2006-present | Investigator<br>Precision Trials<br>Phoenix, AZ   |
| 1998-present | Obstetrician & Gynecologist/ Investigator<br>Arizona Wellness Center for Women, P.C.<br>Phoenix, AZ   |
| 1995-1998    | Obstetrician & Gynecologist/ Investigator<br>Obstetrics & Gynecology Consultants, P.C.<br>Phoenix, AZ |

**Nancy Harris, MD**  
**Curriculum Vitae**  
**Page 2**

**Professional  
Experience  
Continued:**

1994-1995	Gynecologist – Private Practice Scottsdale, AZ
1994-1995	Moved to Scottsdale, AZ, on leave from employment to be with children
1990-1994	Obstetrician & Gynecologist Department of OB/GYN University MEDNET Cleveland, OH

**Professional  
Memberships:**

American College of Obstetrics & Gynecology

**Hospitals:**

Paradise Valley Hospital  
Scottsdale Healthcare

**Licenses/Certifications:**

Arizona # 22215

**Board Certification:**

American Board of Obstetrics & Gynecology  
1994 – Certificate # 29687

**Research Experience:**

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

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

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 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 2, 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.

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 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.

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

**Nancy Harris, MD**  
**Curriculum Vitae**  
**Page 4**

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

A Multi-center, 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, Multi-center, 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

**Nancy Harris, MD**  
**Curriculum Vitae**  
**Page 5**

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

A Randomized, Multi-Center, 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 Multi-center, 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”

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

**Nancy Harris, MD**  
**Curriculum Vitae**  
**Page 6**

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

A phase II, Multi-center, Randomized, Placebo Controlled, Double-Blinded Study of XXXX in Menopausal Women with Symptomatic Leiomyomata

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

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

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 2, 18 week, double blind, placebo controlled, multi-center study evaluating the safety and efficacy of XXXXX combination cream with XXXXX and Placebo in the treatment of Vulvar Vestibulitis Syndrome.

An open label study of the contraceptive efficacy of an extended regime of XXXXX.

Open-label study of the efficacy of a new low dose oral contraceptive containing XXXXX and XXXXX

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.

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 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

**Nancy Harris, MD**  
**Curriculum Vitae**  
**Page 7**

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

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.

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