MD STRATEGIES RESEARCH CENTERS

CURRICULUM VITAE

SIGNATURE:		**DATE:**	
NAME: **TITLE:**	Kofi D. Sefa-Boakye, MD, FACOG Sub-Investigator	**DATE UPDATED:**	Aug 2017

RESEARCH SITE NAME/ ADDRESS:

Main Site MD Strategies Research Centers 6280 Jackson Drive, Suite 8 San Diego, California 92119	Office: (619) 464-1607 Fax: (760) 687-2825 Email: drsefa-boakye@mdsrsandiego.com

EDUCATION/TRAINING

INSTITUTION AND LOCATION	DEGREE *(if applicable)*	YEAR(s)	FIELD OF STUDY
National Institute of Health (NIH) Web-based Training	Certificate	2017	Protecting Human Subject Research Participants
Collaborative Institutional Training Initiative (CITI)	Certificate	2017	Good Clinical Practices
American College of Obstetrics & Gynecology		1994-present	Fellowship
St. Joseph's Hospital Phoenix, AZ		1985-1989	Residency
Harbor/UCLA Medical Center Torrance, CA		1984-1985	OB/GYN Internship
University of California Los Angeles(UCLA) Los Angeles, CA	M.D.	1980-1984	Medicine
University of Southern California (USC) Los Angeles, CA	BS	1978-1980	Biology

BOARD CERTIFIED:

BOARD CERTIFIED/ELIGIBLE	YEAR(s)	SPECIALTY
Board Ce1tified (California)	1994	Obstetrics & Gynecology
Board Certified (Arizona)	1986	Obstetrics & Gynecology

POSITIONS AND EMPLOYMENT:

2016 - Present	Sub-Investigator, MD Strategies Research Centers, San Diego, CA
1991-Present	Owner and Medical Director and Sub-Investigator, a Lady's Doctor Excellence in OB/GYN Medical Clinic, San Diego, CA
1981 – 1982	Research Assistant, Sickle Cell Research Program, Martin Luther King Hospital, Los Angeles, CA
2008	Department Co-Chair OB/GYN Scripps Mercy Hospital, Chula Vista, CA
1998	Department Co-Chair OB/GYN Coronado Hospital, CA

CLINICAL RESEARCH EXPERIENCE:

Sub-Investigator: A Phase 3 Study to Evaluate the Safety and Efficacy of XXXXXX in Combination with XXXXXXXXXX Acetate in Subjects with Moderate to Severe Endometriosis-Associated Pain

Sub-Investigator: A Double-Blind, Placebo and Comparator Controlled, Single Dose Parallel Study of the Analgesic Efficacy and safety of XXXXX in Female Patients with Moderate to Severe Post Abdominal or Pelvic Surgical Pain.

Sub-Investigator: A Phase III, Multi-National, Randomized, Double-Blind, Parallel Group, Placebo-Controlled Study to Evaluate the Efficacy and Safety of XXXX in Naturally Menopausal Women with Hypoactive Sexual Disorder on Concurrent Oral Hormone Replacement Therapy

Sub-Investigator: A Phase III, Multi-National, Randomized, Double-blind, Parallel Group, Placebo-Controlled 24 week Study to Evaluate the Safety and Efficacy of XXXX in Women with Hypoactive Sexual Disorder on Concurrent Estrogen Replacement Therapy Who Have Undergone Hysterectomy and Bilateral Oophorectomy

Sub-Investigator: An Open Label, Single Arm Multi-Center Study to Evaluate the Safety and Efficacy of XXXX in the Episodic and Suppressive Treatment of Recurrent Genital Herpes, 2003-2005

Sub-Investigator: A Multi-Center, Double-Blind, Placebo-Controlled, Randomized Study of XXXX for the Treatment of Post-Operative Pain Following Vaginal Hysterectomy

Sub-Investigator: A Phase III Clinical Study Assessing the Efficacy and Safety of XXX in the Treatment of Moderate to Severe Vaginal Dryness and Vaginal Pain Associated with Sexual Activity

Sub-investigator: Symptoms of Vulvar and Vaginal Atrophy (VVA) Associated with Menopause: A 12-Week, Randomized, Double Blind, Placebo-Controlled, Parallel-Group Study Comparing Oral XXX with Placebo in Postmenopausal Women