

Curriculum Vitae
Krystal R. McConihay

Signature: 
Date Signed: 18 June 2008

Business Address:

CAMC Health Education and
Research Institute
Clinical Trials Center
3100 MacCorkle Avenue, SE
Suite 306
Charleston, WV 25304
Phone: (304) 388-9984
Fax: (304) 388-9935
Email: krystal.mcconihay@camc.org

EDUCATION:

2000 Bachelor of Science
Major Sports Medicine/ Athletic Training
University of Charleston
Charleston, WV

PROFESSIONAL EXPERIENCE:

2008 - Present Clinical Research Assistant (Clinical Trials Center)
CAMC Health Education and Research Institute
Charleston, WV

2007 - 2008 Exercise Technologist / Industrial Athletic Trainer
Holzer Clinic Health & Rehabilitation Center
So. Charleston, WV

2005 - 2007 Customer Service / Marketing Representative
Holzer Cardiovascular Institute
Gallipolis, OH

2003 - 2005 Exercise Technologist/ Industrial Athletic Trainer
Holzer Clinic Health & Rehabilitation Center
So. Charleston, WV

2002 - 2003 Rehabilitation Coordinator
Smith & Jones Chiropractic
Dunbar, WV

CERTIFICATIONS:

Basic Life Support – 2007

AED Training – 2007

CITI – Course in The Protection of Human Research Subjects – 2008

Safe-T Pak Training (Shipping of Dangerous Goods) - 2008

CAMC HIPPA Awareness and Guidelines – 2008

Various Other CAMC training inc. blood-born pathogens, infectious diseases, patient confidentiality, and Biohazard Waste – 2008

CLINICAL RESEARCH EXPERIENCE:

1 Year of Research Experience to date

Clinical Research Coordinator: the primary role of the Research Coordinator is to screen and assess the patients in accordance with the protocol for eligibility, baseline, and follow-up procedures. The research coordinator is also responsible for assisting in the informed consent process, completing all case report forms, collecting the data, and maintaining documents and correspondence with the sponsor. The coordinator also completes all internal and external regulatory requirements.

HS# 07-01-1893: *PEDS # 2 6-17* - Protocol Number CVAL489K2302: A multicenter, randomized, double-blind, parallel-group, evaluation of 12 weeks of valsartan compared to enalapril on sitting systolic blood pressure in children 6 to 17 years of age with hypertension, *Novartis*

HS# 07-01-1894: *PEDS # 2 6-17 Extension* – Protocol Number CVAL489K2302 E01: A 14 week Extension to a randomized, double-blind, multicenter, parallel-group, active-controlled study to evaluate the long-term safety, tolerability and efficacy of valsartan and enalapril combined and alone in children 6 to 17 years of age with hypertension, *Novartis*

HS# 07-10-1979: *VIVUS 302* - A Phase III Randomized, Double-Blind, Placebo-Controlled Multicenter Study to Determine the Safety and Efficacy of VI-0521 in the Treatment of Obesity in Adult Population with BMI ≥ 35 (OB-302), *VIVUS, Inc.*

HS# 07-10-1980: *VIVUS 303* - A Phase III Randomized, Double-Blind, Placebo-Controlled Multicenter Study to Determine the Safety and Efficacy of VI-0521 in the Treatment of Obesity in Adults with Obesity-Related Co-Morbid Conditions (OB-303), *VIVUS Inc.*

HS # 07-06-1954 – *SNUG II* - Study of Glypromate[®] in patients undergoing Cardiopulmonary Bypass (CPB) surgery. Studying Neurons Using Glypromate[®], *Nueren Pharmaceuticals Ltd.*

HS# 08-04-2049: *ZESCA* - Zyban as an Effective Smoking Cessation Aid for Patients Following an Acute Coronary Syndrome, *Canadian Institutes of Health Research*

HS# *CANYON* - A Comparative Clinical and Pharmacoeconomic Study Comparing Argatroban IV vs. Desirudin SC for Patients with Suspected Heparin-Induced Thrombocytopenia (HIT) without Clinically Evident Thrombosis, *Canyon Pharmaceuticals Inc*

HS# *ExOS* - Exenatide Observational Study in the U.S., *Elli Lilly and Company, Inc*

HS# *DVT*- Protocol B0661001 - CV185056: A Safety and Efficacy Trial Evaluating the Use of Apixaban in the Treatment of Symptomatic Deep Vein Thrombosis and Pulmonary Embolism, *Bristol-Meyers Squibb Company*

+++ HS# is Internal study Identifier