


*Curriculum Vitae*  
Cynthia L. Mann, RN

Signature:   
Date Signed: 1/30/08

**Business Address:**

CAMC Health Education and  
Research Institute  
Clinical Trials Center  
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**EDUCATION:**

1993-1996                      WVIT / Nursing Major / Spirit of Nursing Award  
Montgomery, WV

1988                              Windfield High School / Diploma

**PROFESSIONAL EXPERIENCE:**

2008 – Present                Contract Clinical Research Coordinator (Clinical Trials Center)

2001 - 2007                    Clinical Research Nurse Coordinator II (Clinical Trials Center)  
CAMC Health Education and Research Institute  
Charleston, WV

1998 – 2001                    CAMC Memorial Division Cardiac Cath Lab.  
Charleston, WV

1996 – 1997                    CAMC Memorial Division – Open Heart Recovery Unit  
Charleston, WV

1990 – 1993                    St. Francis West Healthcare, Inc. / Office of Adla Adi, MD  
Charleston, WV

**PROFESSIONAL AFFILIATIONS:**

Association of Clinical Research Professionals

**CERTIFICATIONS:**

Basic Life Support – 2006  
American Stroke Association NIH Stroke Scale Certification – 2006

Certified Clinical Research Coordinator – 2005  
CITI – Course in The Protection of Human Research Subjects – 2005

**CLINICAL RESEARCH EXPERIENCE:**

6 Years 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.

**Current Responsibilities:**  
***Co-Coordinator***

HS# 06-09-1855: *MEND CABG II*- Protocol Number 06004: A Randomized, Double-Blind, Placebo-Controlled, Multi-Center Study to Evaluate the Cardio Protective Effects of MC-1 in Patients Undergoing High Risk Coronary Artery Bypass Graft (CABG) Surgery, *Medicare International, Inc. c/o Clinical Development Research Institute, (CDRI) Inc*

**Past Responsibilities:**  
***Lead Coordinator***

HS# 01-09-1302: *PARODI* - The Parodi Anti-Emboli System (PAES) as an adjuvant cerebral protection device during Carotid Stent-supported Angioplasty with the Boston Scientific Carotid Wallstent Monorail Endoprosthesis (Wallstent), *Arteria Medical Science, Inc.*

HS # 01-09-1301: *SHELTER*- Stenting of High Risk Patients: Extracranial Lesion Trial with Emboli Removal, *Boston Scientific Corporation*

HS# 02-09-1407: *CREST*- Carotid Revascularization Endarterectomy vs. Stent Trial, *Acknowledged Cardiovascular Systems, Inc. – Guidant Corporation*

HS# 04-10-1641: *CAPTURE EV* - A Post Approval Study of the Guidant Carotid Stent Systems and Embolic Protection Systems (Carotid RX Acculink/ Accunet Post-Approval Trial to Uncover Unanticipated or Rare Events, *Guidant Corporation*

HS# 05-11-1737: *EXACT* – Protocol Number 640-0063-01: Emboshield (R) and Xact (R) Post Approval Carotid Stent Trial Using the Emboshield BarWire(TM) Rapid Exchange Embolic Protections System and Xact (R) Rapid Exchange Carotid Stent System, *Abbott Vascular Devices*

HS# 05-07-1704: *VIVA* - The VIVEXX Carotid Revascularization Trial for High Surgical Risk Patients with Extracranial Carotid Artery Stenosis using the Bard VIVEXX Carotid Stent and Emboshield Bare Wire Rapid Exchange Embolic Protection System, *Bard Peripheral Vascular, Inc*

HS# 04-04-1591: *CAPTURE EP*- Complete Automatic Pacing Threshold Utilization Recorded by EnPulse, *Medtronic, Inc*

HS# 04-03-1582: *PROPATEN* - Comparison of Primary Patency Between GORE-TEX(R) PROPATEN Vascular Grafts and Thin Walled GORE-TEX(R) Stretch Vascular Grafts, *W. L. Gore, & Associates, Inc*

HS# 04-01-1556: *VALOR* - Evaluation of Medtronic Vascular Talent Thoracic Stent Graft System for the Treatment of Thoracic Aortic Aneurysms, *Medtronic Vascular*

HS# 03-11-1537: *RENAISSANCE* - A Prospective, Multi-Center, Single-Arm Study Evaluating The Express(TM) Renal Premounted Stent System In The Treatment of Atherosclerotic Lesions in the Aortorenal Ostium, *Boston Scientific Corp.*

### ***Co-Coordinator***

HS# 02-09-1406: *MOBILE* - A Prospective, Phase III, Randomized Trial to Evaluate the Safety and Effectiveness of the Novoste™ Corona™ System for the Treatment of In-Stent Restenosis of Native Superficial Femoral Arteries (SFA) and Popliteal Arteries When Used Immediately After Successful Percutaneous Intervention (*MOBILE Study*), *Novoste Corporation*

HS# 04-06-1609: *D-Stat Dry*: A Randomized, Placebo Controlled Trial of the D-Stat Dry Bandage in The Diagnostic Patient, *Vascular Solutions, Inc*

HS# 03-03-1468: *D-STAT* - Prepectoral Pocket Investigation (Utilization of D-Stat Flowable Hemostat in a Prepectoral Pocket, *Vascular Solutions*

HS# 02-05-1366: *SYMBIOT III* - A Prospective Randomized Trial Evaluating the Symbiot III Covered Stent System in Saphenous Vein Grafts, *Boston Scientific Corporation*

+++ HS# is Internal study Identifier