Meagan Downey, CCRC

Curriculum Vitae

PROFESSIONAL EXPERIENCE:

February 2012 – present
Supervisor of Clinical Operations

Chesapeake Research Group, LLC
8030 B Governor Ritchie Highway
Pasadena, Maryland 21122

Responsibilities:

- Review and become familiar with sponsor protocols.
- Recruit, screen and conduct study visit procedures per protocol.
- Completing protocol related activities, including, but not limited to: subject education and administration of pain scales, phlebotomy and laboratory processing, obtaining EKG's.
- Maintaining accurate records of patient care, condition, progress and concerns.
- Administering medication, within limitations, and under supervision.

EDUCATION AND CERTIFICATIONS:

Certified Clinical Research Coordinator (ACRP)- since 2015
CitiTraining: HIPAA & GCP's – since 2012
BLS for Healthcare Providers (CPR & AED) – since 2013
IATA- since 2013

TRIAL EXPERIENCE:

Protocol HTX-011-301 (2017)
A Phase 3, Randomized, Double-Blind, Saline Placebo- and Active-Controlled, Multicenter Study of (the study drug) via Local Administration for Postoperative Analgesia and Decreased Opioid Use Following Unilateral Simple Bunionectomy

Protocol AVE-901-102 (2017)
A Phase 3, Multicenter, Randomized, Double Blind, Three-Arm Study to Evaluate the Efficacy and Safety of (the study drug) versus Placebo in the Management of Postoperative Pain Following Bunionectomy

Protocol CLCT-018 (2017)
A Randomized, Double-Blind, Placebo- and Active-Controlled Study to Determine the Efficacy and Safety of (the study drug) as a Treatment for Moderate-to-Severe Acute Pain and the Prevention of Opioid-Induced Nausea and Vomiting (OINV) Following Orthopedic Surgery

Protocol MOB015B-IV (2017)
A Multi-Centre, Double-Blind, Randomized, Vehicle-Controlled Study of Efficacy and Safety of (the study drug) in the Treatment of Mild to Moderate Distal Subungual Onychomycosis

Protocol ESTEVE-SUSA-301 (2017)
A Randomized, Double-blind, Active- and Placebo-controlled, Parallel Groups, Phase 3 Clinical Trial to Establish the Efficacy of (the study drug) for the Management of Moderate to Severe Post-surgical Pain after Bunionectomy

Protocol AFT-MXIV-703 (2016)
A Phase 3, Randomized, Double-Blind, Multiple-Dose,
Parallel-Group and Placebo-Controlled Study of IV (study drug), IV acetaminophen and IV ibuprofen for the Treatment of Acute Postoperative Pain after Bunionectomy

Protocol CP130-3002 (2016)
A Phase 3, Multicenter, Randomized, Double-Blind, Placebo- and Active-Controlled Study of (the study drug) for the Treatment of Moderate to Severe Acute Pain After Abdominoplasty

Protocol CP130-3001 (2016)
A Phase 3, Multicenter, Randomized, Double-Blind, Placebo- and Active-Controlled Study of (the study drug) for the Treatment of Moderate to Severe Acute Pain After Bunionectomy

Protocol HTX-011-C2015-203 (2016)
A Phase 2, Randomized, Controlled Evaluation of the Efficacy and Safety of (the study drug) for Post-Operative Analgesia Following Abdominoplasty Surgery

Protocol HTX-011-C2016-208 (2016)
A Phase 2, Randomized, Controlled, Multicenter, Evaluation of the Efficacy and Safety of Locally Administered (study drug) for Postoperative Analgesia Following Bunionectomy

Protocol REC-15-016 (2016)
A Phase 3, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Evaluation of the Efficacy and Safety of (the study drug) Following Bunionectomy

Protocol INS005-16-062 (2016)
A Phase 3, Randomized, Double Blind, Multiple Dose, Parallel Group, Placebo Controlled Study of (the study drug) for the Treatment of Moderate to Severe Pain

Protocol 4975-MN-203 (2015)
An Open-Label, Multiple-Dose Extension Study to Evaluate the Safety and Efficacy of (the study drug) in Subjects with Painful Intermetatarsal Neuroma (Morton's Neuroma)

Protocol TV46763-CNS-30031 (2015)
A Multicenter, Randomized, Double-blind, Placebo-controlled, Phase 3 Study to Evaluate the Analgesic Efficacy and Safety of (study drug) Every 4 to 6 Hours in Patients with Moderate to Severe Pain Following Bunionectomy

Protocol REC-15-014 (2015)
A Phase 2, Single-Center, Randomized, Double-Blind, Placebo-Controlled, Evaluation of the Safety, Efficacy, and Pharmacokinetics of (the study drug) Following Bunionectomy

Protocol ELI-200-003-2014 (2015)
A Multi-Center, Randomized, Multiple-Dose, Double-Blind, Placebo-Controlled, Parallel Group, Phase 3 Study to Evaluate the Efficacy and Safety of (the study drug) for the Treatment of Adults with Moderate to Severe Pain Following Bunionectomy Surgery

Protocol INS-14-026 (2015)
A Phase 3, Randomized, Double-Blind, Multiple-Dose, Parallel-Group, Placebo-Controlled Study of (the study drug) for the Treatment of Moderate to Severe Pain

Protocol 4975-MN-202 (2014)
A Randomized, Double-Blind, Placebo-Controlled, Dose-Ranging Study to Evaluate the Efficacy and Safety of CNTX-4975 in Subjects with Painful Intermetatarsal Neuroma (Morton's Neuroma)

Protocol CP130-2001 (2014)
A Phase 2, Randomized, Controlled, Double-Blind, Multiple-Dose, Adaptive, -Placebo and Active-controlled Study of TRV130 for the treatment of Acute Postoperative Pain After Bunionectomy

Protocol MN-201 (2014)
An Open label, Ascending Dose Study to Evaluate the Safety and Tolerability of CNTX-4975 in Subjects with Painful Intertmetatarsal Neuroma (Morton's Neuroma)

Protocol JN-MN-002 (2013)
A Phase 2, Randomized, Controlled, Double Blind, Multicenter Study to evaluate the Safety and Immunogenicity of an Investigational Vaccine compared to an FDA approved Vaccine

Protocol TCO2-2012-01 (2012)
A Prospective, Randomized, Double-Blind Multicenter Study Comparing (the study device) standard Moist Wound Therapy (MWT) in the Treatment of Diabetic Foot Ulcers

Protocol COV15010232-US (2012)
A Phase 3 Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Evaluation of the Safety and Analgesic Efficacy of COV155 Tablets in Moderate to Severe Post-Operative Bunionectomy Pain Followed by an Open-Label Extension

Protocol RBI.2012.003.P.2 (2012)
A Multicenter, Randomized, Sham-Controlled, Double-Blinded Study to Evaluate the Analgesic Efficacy and Safety of Pulsed Radiofrequency Energy (PRFE) in Bunionectomy Surgery for the Treatment of Post-Operative Pain

Protocol IND3-10-06 (2012)
A Phase 3, Randomized, Double-Blind, Multiple-Dose, Parallel-Group, Placebo-Controlled Study for the Treatment of Acute Postoperative Pain After Bunionectomy

Protocol IND3-08-04b (2011-2012)
A Phase 3, Randomized, Double-Blind, Multiple-Dose, Parallel-Group, Active- and Placebo-Controlled Study of Capsules for the Treatment of Acute Postoperative Pain After Bunionectomy.

Protocol COV795 (2011-2012)
A Phase 3 Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Evaluation of the Safety and Analgesic Efficacy of COV795 (study drug) ER Tablets in Moderate to Severe Post-Operative Bunionectomy Pain Followed by an Open-Label Extension

Protocol DIC3-08-04 (2011-2012)
A Phase 3, Randomized, Double-Blind, Multiple-Dose, Parallel-Group, Active- and Placebo-Controlled Study of Capsules for the Treatment of Acute Postoperative Pain After Bunionectomy