Shana Hart, BSN, RN

Education:

December 2012 – Anne Arundel Community College – Arnold, MD
Associates of Arts in Nursing
August 2016 – Walden University – Minneapolis, MN
Bachelor of Science in Nursing

Certifications:

CITI – Coordinators Basic Course – 2015
CITI – Coordinators Refresher Course – 2016-present
CITI – Good Clinical Practice Basic Course – 2015
CITI – GCP Refresher Course – 2017-present
CITI – Health Information Privacy & Security – 2015-pres
IATA Certification 2017-present
CPR/Basic Life Support

Professional Experience:

Feb 2015 – present	**Clinical Research Nurse** **Chesapeake Research Group** Pasadena, MD 21122 410-761-0118
Oct 2010 – present	**Intensive Individual Support Services Instructor** **Trellis Services** Sparks, MD
Aug 2003 – Feb 2015	**Pets Product Manager** Petsmart Pasadena, MD

Clinical Trial Experience:

Protocol VX18-150-104 (2018)
A Phase 2B Randomized, Double-blind, Placebo-controlled, Dose-ranging, Parallel-design Study of the Efficacy and Safety of (study drug) for Acute Pain Following Bunionectomy

Protocol TLC590A1002 (2020)
A Phase 2, Randomized, Double-blind, Comparator- and Placebo-controlled Study to Evaluate the Safety, Pharmacokinetics, and Efficacy of (the study drug) for Postsurgical Pain Management Following Bunionectomy

Protocol CPL-01_AB-001 (2019)
Phase 2a, Randomized, Double-blind, Placebo-controlled study to Evluate the Safety and Pharmacokinetic Profile of (the study drug) in the Management of Acute Postoperative Pain After Mini-abdominoplasty Surgery

Protocol AFT-MXIV-11 (2019)
A Phase 3, Open-Label, Multiple-Dose, Single-Arm Exposure Study of (the study drug) in Patients with Acute Pain Following Orthopedic, General or Plastic Surgery

Protocol D-FR-52120-237 (2019)
A Multiple-Dose, Double-Blind, Randomised, Placebo-Controlled Study to Evaluate the Efficacy and Safey of (the study drug) for the Treatment of Pain Associated with Hallux Abducto Valgus

Protocol CA-PS-201 (2018)
A Phase 2, Randomized, Double-blind, Placebo-controlled Safety, Pharmacokinetics and Efficacy Study of (study drug) in Patients Undergoing Bunionectomy

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-07 (2016)
A Phase 3, Randomized, Double-Blind, Multiple-Dose, Parallel-Group and Placebo-Controlled Study of (the study drug) 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 INS005-15-062 (2016)
A Phase 3, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Evaluation of the Efficacy and Safety of (the study drug) Following Bunionectomy

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 (the 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 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

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