JENNY NGUYEN, D.P.M

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EDUCATION

07 / 2003 – 06 / 2006	Cooper University Hospital: Camden, New Jersey PSR 24+ Podiatric Surgical Resident: Chief Resident Graduation Date: June 2006
09 / 1999 – 05 / 2003	Temple University School of Podiatric Medicine Doctor of Podiatric Medicine: May 2003
09 / 1994 – 05 / 1999	University of Maryland at College Park Bachelor of Science in Neurology and Physiology May 1999 – Cum Laude Honor Graduate

AWARDS

09 / 1999 – 05 / 2000	Temple University School of Podiatric Medicine Admissions Scholarship
05 / 1999	Cum Laude Honor Graduate
09 / 1995 – 05 / 1999	Who's Who among American College Students National Dean's List all Semesters Life Time Golden Key National Honor Society Phi Delta Sigma Honor Society

EXPERIENCE

08 / 2010 – Present	Board Certified, American Board of Podiatric Surgery.
10 / 2011 – Present	Podiatric Physician at Foot and Ankle Specialists of the Mid-Atlantic

07 / 2006 – 2011	Podiatric Physician at Chesapeake Foot and Ankle Center.
07 / 2006 – Present	Sub-Investigator at Chesapeake Research Group, LLC 8028 Ritchie Hwy. Suites 100-106, Pasadena, Maryland 21122
07 / 2003 – 06 / 2006	Three Years of Extensive Training in Podiatric Medicine and Surgery, PSR 24 +, in Fore-foot, Rear-foot, Ankle, Trauma Surgery, Diabetic Wound Care, and Limb Salvage at Cooper University Hospital, NJ
07 / 2006 - Present	Foot Screenings at Multiple Community Health Fairs.
12 / 2001 – 06 / 2002	Temple University School of Podicatric Medicine Gait Study Center

CREDENTIALS

08 / 2010 – Present	Board Certified, American Board of Podiatric Surgery
01 / 2006 - Present	Maryland Podiatric License
07 / 2003 – Present	Certified BLS.

PROFESSIONAL MEMBERSHIP

08 / 2010 - Present	American College Foot and Ankle Surgeon Member
09 / 1998 – Present	American Podiatric Medical Association Member
06 / 2006 - Present	Maryland Podiatric Medical Association Member
06 / 2006 - Present	Podiatry Management, Podiatry Today Member

LANGUAGES

Bilingual: Fluent in English and Vietnamese

RESEARCH

07 / 2006 – Present	Chesapeake Research Group, LLC 8030B Ritchie Highway Pasadena, Maryland 21122
06 / 2000 – 05 / 2002	Gait Analysis using Musgrave, Vicon 370TM, Kistler Force Plate.

CLINICAL 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 MOB015-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 (DSO)

Protocol ESTEVE-SUSA-301
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 INS005-16-062 (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 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, Multicenter, Randomized, Double-Blind, Multiple-Dose, Adaptive, Placebo-and Active-controlled Study of (the study drug) for the Treatment of Acute Post-Operative Pain After Bunionectomy

Protocol MN-201 (2014)
An Open Label, Ascending Dose Study to Evaluate the Safety and Tolerability of (the study drug) in Subjects with Painful Intermetatarsal Neuroma (Morton's Neuroma)

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 (the study drug) 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 (the study device) 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 of (the study drug) for the Treatment of Acute Postoperative Pain After Bunionectomy

Protocol 15000182-US (2012)
A Phase 3 Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Evaluation of the Safety and Analgesic Efficacy of (the study drug) in Moderate to Severe Post-Operative Bunionectomy Pain followed by an Open Label Extension

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

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

Protocol Q8003-022 (2011)
A Randomized, Double-Blind, Multicenter, Repeat-Dose Comparison of the Effects of (the study drug) to the Morphine-Equivalent Doses of Oxycodone and of Morphine on the Opioid-Related Adverse Events of Moderate to Severe Nausea, Emesis, and Dizziness in Subjects with Acute Moderate-to-Severe Postoperative Pain Following Bunionectomy

Protocol KD-PV01 (2010)
A multi-Center, Randomized, Double-Blinded, Pivotal Study of the Safety, Local Tolerability and Efficacy of (the study drug) for the Treatment of Onychomycosis

Protocol BK15 (2009)
Evaluation of a Medical Food for Chronic Wounds

Protocol Q8003-008 (2009)
A Randomized, Double-Blind, Multicenter, Repeat-dose Comparison of Analgesic Efficacy and Safety of (the study drug) with Oxycodone and Morphine for the Management of Acute Moderate to Severe Postoperative Pain Following Bunionectomy Surgery

Protocol Q8003-021 (2009)
A Double-Blind, Randomized, Multi-Center, Repeat Dose, Comparison of the Analgesic Efficacy and Safety of the (the study drug) to each of the Individual Milligram Components (Oxycodone and Morphine) in the Management of Acute Moderate to Severe Pain Following Bunionectomy Surgery

Protocol DFC-005 (2008)
A randomized, double-blind, active- and placebo-controlled study of the analgesic efficacy and safety of repeated dosing of (the study drug) relative to parenteral ketorolac and placebo in patients with acute moderate to severe post-surgical pain following mixed elective general orthopedic surgery

Protocol INN-TOP-002 (2008)
A Phase II, Randomized, Parallel, Double-blind, Placebo-controlled Study to Assess Prevention of Infection Using (the study drug) in Diabetic Patients with Uninfected Lower Extremity Skin Ulcers

Protocol INN-TOP-001 (2008)
A Randomized, Controlled, Open-Label Study to Investigate the Safety and Efficacy of (the study drug) Compared to Levofloxacin in Diabetic Patients with a Mild Infection of a lower Extremity Skin Ulcer

Protocol M10-421 (2008)
A Randomized, Multicenter, double-blind Study comparing the Analgesic Efficacy and Safety of (the study drug) to Placebo in Subjects with Acute Pain Following Bunionectomy

Protocol HKT-500-US10 (2008)
A Randomized, Multicenter, Double-Blind, Placebo-Controlled Study to Evaluate the Efficacy and Safety of (the study drug) in the Treatment of Pain Associated with Grade I or Grade II Ankle Sprain

Protocol KET-017 (2008)
A randomized, multiple-center, double-blind, placebo-controlled study of the safety and analgesic efficacy of repeated dosing of (the study drug) to treat acute post-operative pain following orthopedic trauma, injury or surgery

Protocol R331333PAI3018 (2008)
A Randomized, Double-Blind, Active- and Placebo-Controlled, Parallel-Group, Multicenter Study to Evaluate the Efficacy and Safety of (the study drug) in the Treatment of Acute Pain From Bunionectomy

Protocol Q8003-010 (2007)
A Double-Blind, Multi-Center Extension Study to Evaluate the Safety and Efficacy of (the study drug) in Patients with Acute Moderate to Severe Pain

Protocol Q8003-007 (2007)
A Placebo-Controlled, Randomized, Double-Blind Study of the Safety and Efficacy of (the study drug) in the Management of Post-Bunionectomy Pain

Protocol M06-850 (2007)
A Global, Multicenter, Randomized, Double-Blind, Placebo-Controlled Study Comparing the Safety and Efficacy of (the study drug), Duloxetine and Placebo in Subjects with Diabetic Neuropathic Pain

Protocol KF5503/37 (2007)
A Phase 3, Randomized, Double-Blind, Parallel-Group, Multi-Center, Active- and Placebo-Controlled Trial to Evaluate the Analgesic Efficacy and Safety of Multiple Doses of (the study drug) for Postoperative Pain Following Bunionectomy

Protocol 2005-005 (2006)
A Phase III, Randomized, Double-Blind, Placebo-Controlled Study Evaluating the Safety and Efficacy of (the study drug) for Sedation During Monitored Anesthesia Care

Protocol R331333PAI3003 (2006)
A Randomized, Double-Blind, Active- and Placebo-Controlled, Parallel-Group, Multicenter Study to Evaluate the Efficacy and Safety of Multiple Doses of (the study drug) Immediate Release Formulation in the Treatment of Acute Pain From Bunionectomy Followed by a Voluntary Open-Label Extension

Protocol VGF3554g (2006)
A Phase II, Double-Blind, Randomized, Placebo-Controlled Study to Assess the Effect of (the study drug) for Induction of Healing of Diabetic Foot Ulcers

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