



**JEFFREY E. ATKINSON, MD**  
BOARD CERTIFIED SPECIALIST  
INTERNAL MEDICINE

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THE HORIZONS  
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**EMPLOYMENT**

**July 1992 – October 2003**

Potomac Physicians, PA  
Baltimore, Maryland

**February 2004 – Present**

Jeffrey E. Atkinson, MD, Self-Employed  
Pasadena, Maryland

**EDUCATION AND TRAINING**

**ABIM**

1997 – 2007  
2008 – 2018

**Post Graduate**

Internal Medicine Residency  
Allegheny General Hospital  
Medical College of Pennsylvania  
Pittsburgh, Pennsylvania  
July 1989 – July 1992

**Graduate**

M.D., Medical College of Virginia  
Richmond, Virginia  
August 1985 – May 1989

M.S., Biology – Cellular/Molecular  
Virginia Commonwealth University  
Richmond, Virginia

**Undergraduate**

B.S., Biology  
Virginia Commonwealth University  
Richmond, Virginia

**High School**

Severna Park Senior High School  
Severna Park, Maryland

**PROFESSIONAL LICENSURE**

State of Maryland

**CURRENT ORGANIZATIONS**

American Medical Association  
Medical and Chirurgical Faculty of Maryland  
Anne Arundel County Medical Society

**RESEARCH AFFILIATION**

June 2004 - Present  
 Principal Investigator/Sub-Investigator  
 Chesapeake Research Group, LLC  
 8030B Ritchie Highway  
 Pasadena, Maryland 21122

**CLINICAL RESEARCH****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 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 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 ELI-200-003-2014 (2015)**

A Multi-Center, Randomized, Multiple-Dose, Double-Blind, Placebo-Controlled, Parallel Group, Phase 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

**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-NM-002 (2013)**

A Phase 2, Randomized, Controlled, Double Blind Multi-Center Study to Evaluate Safety and Immunogenicity of (the study vaccine) compared with Menactra

**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 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 (2011-2012)**

A Phase 3 Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Evaluation of the Safety and Analgesic Efficacy of COV795 ER Tablets 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 M12-807 (2011)**

A Phase 2, Randomized Withdrawal Study of the Analgesic Efficacy and Safety of (the study drug) Compared to Placebo in Subjects with Chronic Low Back Pain

**Protocol Q8003-022 (2011)**

A Randomized, Double-Blind, Multicenter, Repeat-Dose Comparison of the Effects of Q8003 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 F1J-US-HMGL (2009-2010)**

A Randomized, Placebo-Controlled Trial of (the study drug) Added to Nonsteroidal Anti-inflammatory Drugs in Patients with Knee Pain due to Osteoarthritis who have had Suboptimal Response to Nonsteroidal Anti-inflammatory Drug Treatment

**Protocol Q8003-008 (2009-2010)**

A Randomized, Double-blind, Multicenter, Repeat-dose Comparison of Analgesic Efficacy and Safety of Q8003 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 Opioid Combination Q8003 to each of the Individual Milligram Components (Oxycodone and Morphine) in the Management of Acute Moderate to Severe Pain Following Bunionectomy Surgery

**\*\* Protocol M10-277 (2009)**

A Phase 3, Open-Label Period Followed by a Randomize, Double-Blind, Placebo-Controlled Study of the Analgesic Efficacy of (the study drug) Compared to Placebo in Subjects with Chronic Low Back Pain

**Protocol DFC-005 (2008)**

A randomized, double-blind, active- and placebo-controlled study of the analgesic efficacy and safety of repeated dosing of DIC075V relative to parenteral ketorolac and placebo in patients with acute moderate to severe post-surgical pain following mixed elective general orthopedic surgery

**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 SKY2028-1-003 (2008)**

A Randomized, Double-Blind, Placebo- and Active-Controlled, Parallel Group, 6-Week Study to Evaluate the Effect of Multiple Doses of (the study drug) twice daily, Prednisone and Placebo on the Hypothalamic-Pituitary-Adrenal Axis in Adult Subjects with Mild to Moderate Asthma

**Protocol KET-017 (2008)**

A randomized, multiple-center, double-blind, placebo-controlled study of the safety and analgesic efficacy of repeated dosing of PMI-150 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 Q8003 in Patients with Acute Moderate to Severe Pain

**Protocol 2005-008 (2006)**

A Phase 2, Randomized, Double-Blind, Parallel Group, Multicenter Lock-out Determination Study of (the study drug) Administered by a Patient-Controlled Analgesia (PCA) Pump for the Treatment of Postoperative Pain Following Elective Bunionectomy Surgery

**Protocol Q8003-007 (2008)**

A Placebo-Controlled, Randomized, Double-Blind Study of the Safety and Efficacy of Q8003 in the Management of Post-Bunionectomy Pain

**\*\* Protocol M06-850 (2008)**

A Global, Multicenter, Randomized, Double-Blind, Placebo-Controlled Study Comparing the Safety and Efficacy of ABT-894, 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 H7U-MC-IDAW (2006-2008)**

A Phase 3, Open-Label, Parallel Group Treatment Concordance Study to Compare Insulin Use and Its Effect on Glycemic Control in Patients with Type 2 Diabetes Mellitus: Two Populations with Different Insulin Treatment Options

**\*\* Protocol M04-697 (2007)**

A Phase 3, Randomized, Multicenter, Double-Blind Study Comparing the Analgesic Efficacy of (the study drug) to Placebo in Subjects With Osteoarthritis

**Protocol R331333PAI3003 (2006-2007)**

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 XP21L-301 (2006)**

A Multicenter, Randomized, Double-Blind, Parallel Group, Placebo-Controlled Study of the Analgesic Efficacy of (the study drug) in Subjects with Pain Following Bunionectomy Surgery

**Protocol EN3269-301 (2006)**

A Randomized, Double Blind, Placebo Controlled Parallel Group Phase III Study of the Efficacy, Tolerability and Safety of (study drug) in the Treatment of Pain Associated With Grade 1 or Grade 2 Ankle Sprain or Strain

**Protocol VGF3554g (2006)**

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

**Protocol 3000-0523 (2006)**

A Phase 3 Open-Label, Single Arm Study to Assess the Safety of (study drug) For Minimal-To- Moderate Sedation in Patients Undergoing Minor Surgical Procedures

**\*\* Protocol M05-790 (2006)**

A Phase 3, Open-Label Period Followed by a Randomized, Double-Blind, Placebo-Controlled Study of the Analgesic Efficacy and Safety of (the study drug) Compared to Placebo in Subjects with Chronic Low Back Pain

**\*\* Protocol HKT-500-US05 (2006)**

A Randomized, Multicenter, Double-Blind, Placebo-Controlled, Two-Week Study to Assess the Safety and Efficacy of (the study drug) in Subjects with Pain from Moderate Lateral Epicondylitis

**Protocol SB-767905/014 (2005-2006)**

A Randomized, Double-Blind, Placebo-Controlled, Multi-Center Phase 3 Study to Evaluate the Long-Term Safety of (the study drug) for 12 Months in the Treatment of Opioid-Induced Bowel Dysfunction in Adults taking Opioid Therapy for Persistent Non-Cancer Pain

**\*\* Protocol HKT-500-US06 (2005)**

An Open Label Safety Study with Intermittent Use of (the study drug) in Subjects with Lower Back Pain, Pain from Osteoarthritis of the Knee, Shoulder Pain or Lateral Epicondylitis Pain

**\*\* Protocol M03-666 (2005)**

An Open Label Study Evaluating the Safety and Tolerability of Long Term Administration of (the study drug) in Subjects with Moderate to Severe Chronic Non-Malignant Pain

**Protocol 3000-0412 (2004-2005)**

A Phase III, Randomized, Open-Label Study to Assess the Safety and Efficacy of (the study drug) versus Midazolam HCl for Sedation in Patients Undergoing Minor Surgical Procedures

**\*\* Protocol M03-643 (2004-2005)**

A Randomized, Multi-Center Double-Blind Study Comparing the Analgesic Efficacy of (the study drug) and Placebo in Subjects with Osteoarthritis

**Protocol 4975-2-003-1 (2004-2005)**

Randomized, Double-Blind, Placebo-Controlled Dose Ranging Trial of (the study drug) in Subjects Undergoing Bunionectomy with First Metatarsal Osteotomy Surgery

**\*\* Principal Investigator**