CHESAPEAKE RESEARCH GROUP, LLC

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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 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 Evaluate 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 (study drug) in Patients with Acute Pain Following Orthopedic, General or Plastic Surgery

Protocol D-FR-52120-237 (2019)
A Multiple-dose, Double-blind, Randomized, Placebo-controlled Study to Evaluate the Efficacy and Safety of (study drug) for the Treatment of Pain Associated with Hallux Abducto Valgus

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 CA-PS-201 (2018)
A Phase 2, Randomized, Double-Blind, Placebo-Controlled Safety, Pharmacokinetics and Efficacy Study of (the 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 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 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

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 Extenstion 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 1or 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**

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Document Name: CV_Atkinson
Document ID: 182
No. Pages: 6

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