




CURRICULUM VITAE

SIGNATURE:		DATE:	27 Jan 2025
NAME:	Roy C. Brownlow, MD	DATE	Jan 2025
TITLE:	Sub Investigator	UPDATED:	

RESEARCH SITE ADDRESS & PHONE:

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EDUCATION/TRAINING:

INSTITUTION AND LOCATION	YEAR(s)	FIELD OF STUDY
East Carolina University, Greenville, NC	1974-1978	Bachelors of Science
East Carolina University, Greenville, NC	1978-1982	Masters of Education
Wake Forest University, Winston-Salem, NC	1982-1987	Doctor of Medicine
North Carolina Baptist Hospital -Winston Salem	1987-1988	General Surgery Internship
North Carolina Baptist Hospital -Winston Salem	1988-1991	Anesthesia Residency
NIDA Clinical Trials Network	2022	Good Clinical Practice

POSITIONS AND EMPLOYMENT:

Jan 2025 - Present	Advanced Clinical Research Center, dba TriWest Research Associates, LLC, Chula Vista, CA
May 2024 - Present	Investigator, BioSolutions Clinical Research Center, LLC, La Mesa, CA
Aug 2008 - Present	Investigator, Better Health Clinical Research Inc, Newnan, GA
Sept 2023- Present	Owner/America's Pain, Spine Center, Interventional Pain Management Specialist, Escondido, La Mesa Jolla, CA

Aug 2022 - Sept 2023	Equity Owner/Pain Management Specialist/Stockbridge, Newnan and Griffin, GA
Jul 2013 - Aug 2022	Owner/Anesthesiologist, White Oak Surgery Center, Newnan, GA
May 2008 - Aug 2022	Owner/Georgia Pain and Spine Care, Newnan, GA
2013 - 2020	Owner, Care Plus Pharmacy, Newnan, GA
2015 - 2020	Owner, Newnan Plaza Pharmacy, Newnan, GA
Dec 2005 - Apr 2008	Sub-Investigator, Drug Studies America, Marietta, GA
May 1999 - Apr 2008	Pain Management Specialist, Pain Solutions Treatment Centers, Marietta, GA
Nov 1998 - Mar 1999	Pain Management Specialist, Coast Pain Management, San Diego, CA
Jul 1996 - Nov 1998	Staff Anesthesiologist & Pain Management Specialist Northside Hospital, Atlanta, GA
Feb 1996 - Jun 1996	Staff Anesthesiologist & Pain Management Specialist Kaiser Permanente, San Diego, CA
Jul 1994 - Feb 1996	Anesthesiologist & Pain Management Specialist JLR Anesthesia Group, Orlando, FL
1991-1994	Anesthesiologist & Pain Management Specialist, U.S. Naval Medical Center, San Diego, CA

CURRENT LICENSES & CERTIFICATIONS:

Georgia License #042196
 California License #G72363
 Certified Principal Investigator (CPI), Association of Clinical Research Professionals
 Diplomate, American Board of Anesthesiology
 ABA Certificate of Added Qualifications in Pain Management
 Diplomate, American Academy of Pain Management
 Diplomate, American Academy of Pain Medicine
 Fellow in Interventional Pain Practice, WIP & ASIPP
 Certified I American Acupuncture

PROCEDURAL SKILLS:

Spinal Cord Stimulation
 DRG Stimulator
 IT Pumps
 Verti flex/MILD
 Radiofrequency Ablation
 Epidural Steroid Injections
 Transforaminal Epidural Steroid injections
 Ultrasound Diagnostics
 Ultrasound Guided Injections
 Platelet Rich Plasma Injections
 Peripheral Nerve Blocks

Discography: Cervical, Thoracic and Lumbar
Joint Injections
Peripheral Nerve Stimulation
Kyphoplasty

CLINICAL RESEARCH EXPERIENCE:

"A Study of the Efficacy and Safety of XXXX Extended Release (XXER) Compared to Placebo in Patients with Chronic Pain." (Sub Investigator)

"A Randomized, Double-Blind, Two-Period Crossover Study Comparing the Efficacy, Safety, and Tolerability of XXXX (XXXX, controlled release) and XXXX (XXXX, controlled release) in Cancer Patients Who Require Opioid Treatments." (Sub- Investigator)

"An Open-Label Extension study to Evaluate the Long-Term Safety, Tolerability, and Analgesic Efficacy of XXXX (XXXX, controlled release) in Subjects with Cancer Pain or Chronic Lower Back Pain." (Sub-Investigator)

"A prospective, Open-Label, Multicenter Study of the Effectiveness and Safety of XXXX as Add-on Treatment in Patients with Post-herpetic Neuralgia, Diabetic Neuropathy, or Low Back Pain." (Sub-Investigator)

"A Prospective, Multicenter, Open-Label Study of the Effectiveness and Safety of XXXX and Add-on or Mono-therapy in Patients with Pain from Osteoarthritis in One or Both Knees." (Sub-Investigator)

"A Study of the Efficacy and Safety of XXXX (XXXX, extended release) Compared to Placebo in Subjects with Persistent Pain." (Sub-Investigator)

"A phase 3, Randomized, Multicenter, Double-Blind Study Comparing the Analgesic Efficacy of Extended Release XXXX Tablets (XXXX) to Placebo in Subjects with Osteoarthritis." (Principal Investigator)

"A Multi-Center, Double-Blind, Randomized, Placebo-Controlled, Multiple Dose, Parallel Design, Dose Ranging Study of the Safety and Efficacy of XX.XX in Painful Diabetic Peripheral Neuropathy." {Sub-Investigator)

"XX.XX Phase III Clinical Trial: A Twenty Four Week, Randomized Double-Blind, Placebo Controlled, Safety and Efficacy Trial of XXXX 50 and 100 Milligrams Each Evening in Premenopausal Women With Hypoactive Sexual Desire Disorder." {Sub- Investigator)

"An Open Label, 12-Month Study to Evaluate the Safety, Tolerability, and Efficacy of :XXXX for the Management of Breakthrough Pain in Opioid Tolerant Patients with Chronic Non-cancer Pain." (Sub-Investigator)

"A Double-Blind, Randomized, Placebo-Controlled Study to Evaluate the Efficacy and Safety of XXXX for the Management of Breakthrough Pain in Opioid-Tolerant Patients with Chronic Neuropathic Pain." (Sub-Investigator)

"A Double-Blind, Randomized, Placebo-Controlled Study to Evaluate the Efficacy and Safety of XXXX for the Management of Breakthrough Pain in Opioid-Tolerant Patients with Chronic Low Back Pain." (Sub-Investigator)

"A 12-Week, Open-Label Study With 3 Within-Patient Double-Blind Placebo- Controlled Periods to Evaluate the Efficacy and Safety of XXXX Treatment for the Management of Breakthrough Pain in Opioid-Tolerant Patients With Non-Cancer- Related Chronic Pain." (Sub-Investigator)

"A Multi-Center, Double-Blind, Placebo-Controlled Randomized Study of XXXX 200 mg, XXXX 300 mg, and XXXX 400 mg in the Treatment of Chronic Low Back Pain." (Sub-Investigator)

"A Multi-Center, Standard of Care-Controlled Study to Evaluate the Long-Tenn Safety of XX.XX for the Treatment of Chronic Low Back Pain." {Sub-Investigator)

"A Double-Blind, Randomized, Placebo-Controlled, Multicenter Study to Evaluate the Efficacy and Safety of XXXX for Treatment of Breakthrough Pain In Opioid Tolerant Cancer Patients Followed by an up to 12 -Month, Non-Randomized, Open-Label Extension to Assess Long-Tenn Safety." (Sub-Investigator)

"A randomized, multi-center, double blind, parallel-group study assessing the analgesic efficacy and safety of different dosages of: XXXX *bid* compared to active comparator *bid* and placebo *bid* in subjects with chronic knee-joint osteoarthritis." (Sub-Investigator)

A Randomized, Double-Blind, Placebo-Controlled, Multicenter Phase 3 Study to Evaluate the Efficacy and Safety of XX.XX 0.5mg Once Daily and 0.5mg Twice Daily for 12 Weeks for the Treatment of Opioid-Induced Bowel Dysfunction in Adults taking Opioid Therapy for Persistent Non-Cancer Pain." (Sub-Investigator)

"A Randomized, Double-Blind, Placebo-Controlled, Multicenter Phase 3 Study to Evaluate the Long-Tenn Safety of XXXX 0.5mg Twice Daily for 12 Months for the Treatment of Opioid-Induced Bowel Dysfunction in Adults taking Opioid Therapy for Persistent Non-Cancer Pain." (Sub-Investigator)

"An observational study to characterize the burden of illness associated with laxative use in subjects using opioids for the management of persistent pain." (Sub-Investigator)

"A Randomized, Double-Blind, Active-Control, Parallel Group, 90-Day Safety Study of XXXX Immediate Release or XXXX Immediate Release in Subjects With Chronic Pain From Low Back Pain or Osteoarthritis of the Hip or Knee." (Sub-Investigator)

"A Double-Blind, Randomized, Placebo-Controlled Study of XXXX and XXXX in Patients with Excessive Daytime Sleepiness Due to Opioid Therapy." (Sub-Investigator)

"A Randomized, Double-Blind, Controlled Study of XXXX for the Treatment of Postherpetic Neuralgia." (Sub-Investigator)