



# CLINICAL TRIALS OF TEXAS, INC.

*Promoting Health Through Research*

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## Curriculum Vitae

### Timothy Stewart Orihel, MD

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#### EDUCATION

1980-1983 Internship in Anesthesiology, Wilford Hall USAF Medical Center, Lackland AFB, Texas  
1976-1980 Doctor of Medicine, Tulane Medical School, New Orleans, Louisiana  
1972-1976 Bachelors Degree, Louisiana State University, Baton Rouge, Louisiana

#### LICENSE(S)

2009 Department of Public Safety Narcotic Service  
2008 Drug Enforcement Administration  
1983 Texas State Board of Medical Examiners, # G4667

#### CERTIFICATION(S)

1998 American Board of Pain Management, # 12573  
1985 American Board of Anesthesiology, # 12573

#### PROFESSIONAL EXPERIENCE

1994-Present Private Practice, Tejas Anesthesia, P.A., San Antonio, Texas

#### HOSPITAL AFFILIATION(S)

Christus Santa Rosa Health Care, San Antonio, Texas  
Methodist Healthcare System, San Antonio, Texas  
Baptist Health System, San Antonio, Texas

#### RESEARCH AFFILIATION(S)

2004-Present Investigator, Clinical Trials of Texas, Inc., San Antonio, Texas

#### RESEARCH EXPERIENCE

2009-Present: A One-Year, Randomized, Open-Label, Parallel-Group, Multiple-Dose Long-Term Safety Study With Controlled Adjustment Of Dose Of "Study Drug" And "Study Drug" In Subjects With Chronic, Painful Diabetic Peripheral Neuropathy (DPN)

2009-Present: A Phase IIa, Double-Blind, Randomized, Parallel-Group, Multi-Centre Study To Evaluate The Analgesic Efficacy Of 28 Days Oral Administration Of "Study Drug" With One-Day Escalation Compared To Placebo In Peripheral Neuropathic Pain Patients With Mechanical Hypersensitivity

2008-Present: An Open-Label Study To Evaluate The Long-Term Safety Of Subcutaneous "Study Drug" For Treatment Of Opioid-Induced Constipation In Subjects With Nonmalignant Pain

2008–2009: Study For The Validation Of The BF-Diary For Assessing Opioid-Induced Constipation

2007-2009: A Phase II Randomized, Double-Blind, Placebo-Controlled, Multicenter, Parallel Group, Proof Of Concept Study Of The Analgesic Effects Of “Study Drug” In Adult Patients With Post-Herpetic Neuralgia

2007-2008: A Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study Of Subcutaneous “Study Drug” For The Treatment Of Opioid-Induced Constipation In Subjects With Chronic Non-Malignant Pain

2007-2008: A Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study Of Oral “Study Drug” For The Treatment Of Opioid-Induced Bowel Dysfunction In Subjects With Chronic Non-Malignant Pain

2006-2008: A Randomized, Double-Blind, Placebo-Controlled, Parallel-Group With A Crossover Confirmation Period Study Of “Study Drug” For The Treatment Of Postherpetic Neuralgia

2006-2007: An Open-Label Extension Trial Assessing The Safety And Tolerability Of “Study Drug” In Patients With Postherpetic Neuralgia

2006-2007: An Observational Study To Characterize The Burden Of Illness Associated With Laxative Use In Subjects Using Opioids For The Management Of Persistent Pain

2005-2006: A 16-Week, Randomized, Double-Blind, Placebo And Pregabalin Controlled, Multi-Center Trial Of “Study Drug” In Patients With Postherpetic Neuralgia

2004-2006: A Randomized, Double-Blind, Placebo-Controlled, Multicenter Phase 3 Study To Evaluate The Efficacy And Safety Of “Study Drug” 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

2004-2006: A Randomized, Double-Blind, Placebo-Controlled, Multicenter Phase 3 Study To Evaluate The Long-Term Safety Of “Study Drug” Twice Daily For 12 Months For The Treatment Of Opioid-Induced Bowel Dysfunction In Adults Taking Opioid Therapy For Persistent Non-Cancer Pain

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Signature and Date