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EDUCATION:

1998 Doctor of Medicine
University of Texas Southwestern Medical Center at Dallas, TX

1994 Bachelor of Science
Texas A&M University, College Station, TX

INTERNSHIPS AND RESIDENCIES:

2005-2006 Fellowship, Sleep Medicine
Tulane University Health Sciences Center at New Orleans, LA

2002-2005Fellowship, Pulmonary/ Critical Care Medicine
Tulane University Health Sciences Center at New Orleans, LA

2001-2002 Chief Resident, Internal Medicine
Tulane University Health Sciences Center at New Orleans, LA

INTERNSHIPS AND RESIDENCIES (*continued*):

2001 Residency, Internal Medicine
Tulane University Health Sciences Center at New Orleans, LA

1999 Internship, Internal Medicine
Tulane University Health Sciences Center at New Orleans, LA

CERTIFICATIONS:

Board Eligible: Critical Care Medicine
ABIM-Diplomate: Sleep Medicine (2011)
Diplomate-American Board of Sleep Medicine (2006)
ABIM-Diplomate: Pulmonary Diseases (2004)
ABIM-Diplomate: Internal Medicine (2001)

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MEMBERSHIPS:

American College of Chest Physicians
American Thoracic Society
American Academy of Sleep Medicine
Travis County Medical Society

PROFESSIONAL EXPERIENCE:

Sub-Investigator, 2009-present
FutureSearch Trials of Neurology, LP, Austin, TX

Research Assistant, 2004-2005
Tulane University Health Sciences Center, New Orleans, LA

Research Assistant, 2002-2003
Tulane University Health Sciences Center, New Orleans, LA

CLINICAL RESEARCH EXPERIENCE:

Asthma • Alzheimer's Disease • Cognition • Chronic Idiopathic Constipation • Chronic Pain COPD • Diabetic Peripheral Neuropathy • Fibromyalgia • Insomnia • Irritable Bowel Syndrome - Constipation • Migraine • Narcolepsy • Neuropathic Pain • Phase I • Postherpetic Neuralgia Osteoarthritis • Restless Legs Syndrome • Sleep Disorders • Tinnitus • Tourette's

CLINICAL TRIAL EXPERIENCE (Sub-I):

Phase I Insomnia

A Phase I study designed to obtain consumer preference feedback on an integrated continuous positive airway pressure (CPAP)/mask system and the marketing materials created for this product

A Multicenter, Open-label Study to Determine the Effects of XXX on Sleep in Healthy Subjects

Alzheimer's Disease

An Open-Label, Extension Study of the Effects of XXX in Subjects with Alzheimer's Disease or Behavioral Variant Frontotemporal Dementia

Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, 12-Month Trial of XXX in Subjects with Mild to Moderate Alzheimer's Disease (2 studies: Study A: Multinational, Randomized, Double blind, Placebo controlled, Parallel Group, 18 month, Study of XXX in patients with Mild Alzheimer's Disease Study B: Randomized Double blind, Placebo-controlled, Parallel group, 52 week study of 2 dose levels of XXX in pts w/treated or untreated Mild to Moderate Alzheimer's Disease)

Cognition

A Phase II, Randomized, Multicenter, Double-Blind, Placebo-Controlled, Parallel-Group Study Comparing XXX with Placebo in Subjects with Age-Associated Memory Impairment (AAMI)

Epilepsy

Pilot Study to Evaluate a Baseline Prototype Device Designed to Collect ECG (Electrocardiogram) and Accelerometer Data

A Prospective Randomized 12-week Controlled Study of Visual Field Change in Subjects with Partial Seizures Receiving XXX or Placebo

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

A Phase III, 12-month, Open-label Study Evaluating the Safety and Tolerability of Flexible Doses of XXX as Adjunctive Therapy in Pediatric Patients Ages 1 month to 16 years with Partial Onset Seizures and Pediatric and Adult Patients Ages 5 to 65 years with Primary Generalized Tonic-Clonic Seizures

A Randomized, Double-Blind, Placebo-Controlled, Parallel Group, Multi-Center Trial of XXX Controlled Release Formulation as Adjunctive Therapy in Adult Subjects with Partial onset Seizures

A Multi-center, Open-Label Study to Assess the Efficacy of as First Add-on Treatment in Adult Patients (17-65) with Partial-Onset Seizures. This Phase III b/IV Study will Assess Two Groups of Epilepsy Patients: Patients who are Not Responding to Monotherapy Anti-Epileptic Drug (AED) Treatment (Diagnosis ≤ 12 Months); OR Patients who Have Not Responded to More Than Two Treatments (Diagnosis ≥ 5 Years)

A Randomized, Double-Blind, Parallel-Group, Multi-center Study to Evaluate the Retention Rate, Efficacy, Safety, and Tolerability of XXX, XXX, and XXX as Adjunctive Therapy in Subjects with Partial Onset Seizures

A 9-11 Week Multi-center, Randomized, Double-Blind, Placebo-Controlled, Parallel Group Study to Determine the Effects of Adjunctive XXX on the Sleep Architecture of Adult Subject (18 – 45 Years of Age) with Partial Onset Epilepsy Receiving a First Generation Anti-Epileptic Drug

Fibromyalgia

An Open-label Extension Study of XXX for 52 weeks in Pain Associated with Fibromyalgia

A randomized, double-blind, double-dummy, placebo- and active-controlled, multi-center study of XXX in subjects with pain associated with fibromyalgia

A Phase IIIb Multicenter, Double-blind, Randomized, Placebo-controlled, 2-way Crossover Study of XXX in the Treatment of Fibromyalgia with Concurrent Antidepressant Therapy for Comorbid Depression

A Comprehensive Evaluation of Impacts and Possible Outcome Assessments

A Phase III Double-Blind, Randomized, Placebo-Controlled, Safety and Efficacy Study of Once Daily Controlled Release XXX in the Treatment of Patients with Fibromyalgia

A Phase II, Multicenter, Open-label, 52-Week Extension Study to Evaluate the Safety and Efficacy of XXX in Pediatric Patients With Primary Fibromyalgia

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

A Phase II, Multicenter, Randomized, Double-blind, Placebo-Controlled Withdrawal Study to Evaluate the Safety, Tolerability, and Efficacy of XXX in Pediatric Patients With Primary Fibromyalgia

A Safety and Tolerability Study Comparing XXX Given as an Oral Solution to a Single-Blinded Combination of Oral Tablets Plus Oral Solution in Subjects with Fibromyalgia

Insomnia

A Phase II, Randomized, Placebo-Controlled, DoubleBlind, Fixed-Dose, Multiple Cohort, Multiple Crossover, Dose-Finding Study of Oral XXX in Adults with Idiopathic Hypersomnia or Narcolepsy Type 2

A Phase II Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Bayesian Adaptive Randomization Design, Dose Response Study of the Efficacy of XXX in Adults and Elderly Subjects with Chronic Insomnia

A12-Month, Open-Label Study to Evaluate the Safety, Tolerability, and Efficacy of XXX as Treatment for Patients With Excessive Sleepiness Associated With Mild or Moderate Closed Tramatic Brain Injury

A Phase III, Multi-center, Randomized, Double-Blind, Placebo-Controlled, Parallel Group, Polysomnography Study to Evaluate the Safety and Efficacy of XXX in [adults and] Elderly Patients with Primary Insomnia

A Randomized, Double-Blind, Parallel Group, Placebo-Controlled, Multi-center Outpatient Trial of XXX in Adults with Nonrestorative Sleep

Narcolepsy and Excessive Daytime Sleepiness

A twelve-week, double-blind, placebo-controlled, randomized, parallel-group, multi-center study of the safety and efficacy of XXX in the treatment of excessive daytime sleepiness in subjects with narcolepsy

Migraine

A Phase III, Multicenter, Randomized, Double-Blind, Placebo Controlled, Efficacy, Tolerability, and Safety Study of XXX in Episodic Migraine With or Without Aura

A Multicenter, Randomized, Open-Label Extension Study to Evaluate the Long-Term Safety and Tolerability of Oral XXX in the Acute Treatment of Migraine With or Without Aura

A Phase III, Multicenter, Randomized, Double-Blind, Placebo-Controlled Single Attack Study to Evaluate the Efficacy, Safety and Tolerability of Oral XXX in the Acute Treatment of Migraine

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

A Phase III, Open-label, LonG-term, Safety Study of XXX (100 mg and 200 mg) in the Acute Treatment Of MigRaine

A Phase III, Study of Three Doses of XXX (50 mg, 100 mg and 200 mg) Compared to Placebo in the Acute Treatment of Migraine: A Randomized, Double-blind, Placebo-controlled Parallel Group Study

An Observational Research Study: Prospective Cohort Study to Describe Patient-Reported Outcomes in Subjects with Migraine Eligible for Prophylaxis

A Phase III, Multicenter, Randomized, Double-Blind, Parallel-Group Study Evaluating the Long-Term Safety, Tolerability, and Efficacy of Subcutaneous Administration of XXX for the Preventive Treatment of Migraine

A Phase III, Multicenter, Randomized, Double-Blind, Double-Dummy, Placebo-Controlled, Parallel-Group Study Comparing the Efficacy and Safety of 2 dose regimens of subcutaneous administration of XXX versus Placebo for the Preventive Treatment of Episodic Migraine

A Phase III, Multicenter, Randomized, Double-Blind, Double-Dummy, Placebo-Controlled, Parallel-Group Study Comparing the Efficacy and Safety of 2 dose regimens of subcutaneous administration of XXX versus Placebo for the Preventive Treatment of Chronic Migraine

A Phase III, Randomized, Double-Blind, Placebo-Controlled Study of XXX in Patients with Chronic Migraine – the XXX Study

A Phase III, Randomized, Double-Blind, Placebo-Controlled Study of XXX in Patients with Episodic Migraine – the XXX Study

A Phase III, Randomized, Double-blind, Placebo-controlled Study to Evaluate the Efficacy and Safety of XXX in Migraine Prevention

An Open-label Extension (OLE) Study to Assess the Long-term Safety and Efficacy of XXX in Chronic Migraine Prevention

A Phase II, Randomized, Double-blind, Placebo-controlled Study to Evaluate the Efficacy and Safety of XXX in Chronic Migraine Prevention

A Randomized, Multicenter, Double-Blind, Placebo Controlled, Two-Arm Study Evaluating Efficacy of a Single Dose XXX (10 mg vs. Placebo) in Patients with Acute Migraine Headache With or Without Aura

A Phase IIb, Randomized, Doub-Blind, Placebo-Controlled Study of XXX in Patients with Episodic Migraine

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

A Multicenter, Double-Blind, Placebo-Controlled, Parallel Group, Multi-dose Study to Compare the Efficacy and Safety of Subcutaneous XXX with Placebo for the Preventive Treatment of Chronic Migraine

A Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Study Comparing the Efficacy and Safety of Two Doses of Subcutaneous XXX with Placebo for the Preventive Treatment of High Frequency Episodic Migraine (HFEM)

A Phase II, Randomized, Double-blind, Placebo-controlled Study to Evaluate the Efficacy and Safety of XXX in Episodic Migraine Prevention, Dosed monthly by subcutaneous (SC) injection.

A Phase II, Randomized, Double-Blind, Placebo-Controlled Study of XXX in Patients with Migraine

A Phase IIb Double blind Randomized Placebo controlled, Dose-ranging Trial of XXX for the Acute Treatment of Migraine

Pain

A Phase II, Randomized, Double-Blind, Placebo-Controlled, Titration-to-Effect Study of Orally Administered XXX in Patients with Osteoarthritis of the Hip or Knee

A Phase III, Open-Label, Long-Term Safety and Tolerability Study of XXX in Patients with Irritable Bowel Syndrome with Constipation (IBS-C)

An Open-label Phase III Trial to Evaluate the Safety and Tolerability of XXX Tablet, in Patients with Moderate-to-Severe Chronic Noncancer Pain

A Second Phase III, Randomized, 12-Week, Double-Blind, Placebo-Controlled Study of the Safety and Efficacy of XXX in Patients with Irritable Bowel Syndrome with Constipation (IBS-C)

A Phase III, Open-Label Extension (OLE), Long-term Safety and Tolerability Study of XXX in Patients with Chronic Idiopathic Constipation (CIC)

A Phase III, Randomized, Double Blind, Placebo and Active-Controlled, Multicenter, Parallel-Group Study of the Analgesic Efficacy and Safety of XXX in Adult Patients with Chronic Low Back Pain

A Phase II, Double-blind, Placebo-controlled, Randomised Dose Ranging Trial to determine the safety and efficacy of two dose levels of XXX (angiotensin II type 2 receptor antagonist) administered orally in patients with postherpetic neuralgia

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

A Phase III Multicenter, Open-Label, 52-week study to Evaluate the Long Term Safety and Tolerability of XXX in Subjects with Moderate to Severe Chronic Noncancer Pain

A Phase III Double-Blind, Placebo-Controlled Study to Assess the Efficacy, Safety, and Tolerability of XXX in Opioid-Naïve Subjects with Moderate to Severe Chronic Low Back Pain

A Safety and Efficacy Evaluation of XXX Laxative in Constipated Adults

A Phase IIa, Randomized, Double-Blind, Multicenter, Placebo and Active-Controlled Study to Assess Analgesic Efficacy and Safety of XXX in Subjects with Painful Diabetic Peripheral Neuropathy

A Phase II, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Safety and Efficacy of Topically Applied XXX Ointment in Patients with Postherpetic Neuralgia

A Randomized Double-blind, Placebo-controlled, Parallel-group, Multicenter, Phase III Study to Evaluate the Long-term Safety of XXX for the Treatment of Opioid-induced Constipation in Subjects with Non-malignant Chronic Pain Receiving Opioid Therapy

A Double-blind, Randomized, Vehicle-controlled, Parallel-group Evaluation of XXX and Vehicle Nasal Sprays in the Treatment of Postherpetic Neuralgia of the Trigeminal Nerve

A National, Randomized, 12-Week, Double-Blind, Placebo-Controlled Study to Assess the Safety and Efficacy of XXX (3.0 and 6.0 mg) in Patients with Chronic Idiopathic Constipation

A Multicenter, randomized, double-blind, placebo-controlled trial to assess the safety and efficacy of XXX in Subjects with Type 2 Diabetes and Diabetic Peripheral Neuropathy

A Phase III, Randomized, Double-blind, Placebo-controlled, Parallel-group Study of XXX in the Treatment of Opioid-induced Constipation in Subjects with Non-malignant Pain Receiving Opioid Therapy

A Phase II, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study to Evaluate the Efficacy and Safety of XXX in Subjects with Postherpetic Neuralgia

A Phase II Multicenter, Randomized, Double-Blinded, Placebo-Controlled Study Using a Bayesian Adaptive Design to Assess the Efficacy, Safety, Tolerability, and Serum Exposure of Multiple Doses of XXX in Subjects with Painful Lumbar Radiculopathy

An Phase III open-label extension study of up to 52 weeks to assess the safety, tolerability, and analgesic efficacy of XXX in the management of moderate to severe chronic pain requiring ATC opioid analgesia for an extended period of time

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

A Phase III, 6-Month, Open-Label, Extension Study to Evaluate the Safety of XXX at 15 to 90 mg Every 12 Hours for Relief of Moderate to Severe Pain in Patients With Chronic Low Back Pain Who Require Opioid Treatment for an Extended Period of Time

A Phase III, 12-Week, Randomized, Double-Blind, Placebo-Controlled, Randomized-Withdrawal Study to Evaluate the Efficacy and Safety of XXX at 30 to 90 mg Every 12 Hours for Relief of Moderate to Severe Pain in Opioid-Experienced Patients With Chronic Low Back Pain Who Require Opioid Treatment for an Extended Period of Time

A Randomized, Double-Blind, Placebo- and Active-Controlled Study to Evaluate the Safety and Efficacy of XXX in Patients with Moderate to Severe Chronic Pain Due to Osteoarthritis of the Knee

A Phase II, Randomized, Double-blind, Placebo- and Active-controlled, Parallel-group, Multicenter Study Evaluating the Analgesic Efficacy and Safety of XXX in Subjects With Moderate to Severe Chronic Pain Due to Postherpetic Neuralgia (PHN) pursuant to Protocol XXX

A Phase III, Double-blind, Placebo-controlled, Multicenter, Randomized Withdrawal Study to Evaluate the Analgesic Efficacy, Safety, and Tolerability of XXX in Opioid-Experienced Subjects with Moderate to Severe Chronic Low Back Pain Requiring Around-the-clock Opioid Analgesia for an extended period of time

A Phase III, Multi-Center, Randomized, Double-Blind, Placebo-Controlled, Safety, Tolerability, and Efficacy Study of XXX Versus Placebo in Opioid-Experienced Subjects with Moderate to Severe Chronic Low Back Pain

A Phase III Open Label Safety Study of XXX in Subjects with Osteoarthritis or Chronic Low Back Pain

A Multicenter, Randomized, Double-blind, Placebo-controlled, Phase III Study to Evaluate the Efficacy and Safety of XXX for the Treatment of Opioid-Induced Constipation in Adults taking Opioid Therapy for Chronic Non-Cancer Pain AND A Multicenter, Randomized, Double-blind, Placebo-controlled, Phase III Study to Evaluate the Long-Term Safety and Tolerability of XXX for the Treatment of Opioid-Induced Constipation in Adults taking Opioid Therapy for Chronic Non-Cancer Pain

A Phase III, Double-blind, Placebo-controlled, Multicenter, Randomized withdrawal Study to Evaluate the Analgesic Efficacy, Safety, and Tolerability of XXX in Opioid-naïve Subjects with Moderate to Severe Chronic Low Back Pain Requiring around-the-clock Opioid Analgesia for an Extended Period of Time

A Phase II Study, Assessing the Content Validity of XXX in Patients with Chronic Opioid-Induced Constipation

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

An Open-label Evaluation of XXX Extended Release (ER) in Subjects with Moderate to Severe Chronic Pain Following Conversion from XXX

A Multi-center, 12-week, Double-blind, Placebo-controlled, Randomized Withdrawal Study to Determine the Efficacy and Safety of XXX Extended-release Capsules in Subjects with Moderate to Severe Chronic Low Back Pain

A Phase IV, Randomized, Double-Blind, Parallel-Arm Clinical Trial to Compare the Clinical Effectiveness of XXX vs. XXX in Subjects with Moderate to Severe Low Back Pain

A Phase IIa, Multicenter, Double-Blind, Placebo-Controlled, Parallel-Group Clinical Trial to Evaluate the Safety and Efficacy of XXX in Patients with Painful Diabetic Neuropathy

A Randomized, Double-Blind, Placebo and Active Comparator-Controlled Study of XXX for Treatment of Neuropathic Pain Associated with Diabetic Peripheral Neuropathy

A Phase II, Multinational, Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel-group Study of Efficacy and Safety of XXX 20MG and 120MG Twice Daily for 4 Weeks in Patients with Chronic Peripheral Neuropathic Pain

A Randomized, Double-blind, Double-dummy, Placebo-controlled, Active-controlled, Parallel-group, Multicenter Trial of XXX controlled-release Tablets to Assess the Analgesic Efficacy (Compared to Placebo) and the Management of Opioid-induced Constipation (Compared to XXX) in Opioid-experienced Subjects with Controlled Moderate to Severe Chronic Low Back Pain and a History of Opioid-induced Constipation who Require Around-the-clock Opioid Therapy

A Randomized, Double-blind, Double-dummy, Placebo-controlled, Active-controlled, Parallel-group, Multicenter Trial of XXX controlled-release Tablets to Assess the Analgesic Efficacy (Compared to Placebo) and the Management of Opioid-induced Constipation (Compared to XXX) in Opioid-experienced Subjects with Uncontrolled Moderate to Severe Chronic Low Back Pain and a History of Opioid-induced Constipation who Require Around-the-clock Opioid Therapy

An Open Label, Safety and Tolerability Study of Multi-Layer, Extended-Release Tablets of XXX in the Treatment of Patients with Moderate to Severe Acute Pain of Up to 30 Days, of Non-Malignant Origin

A Phase IV, Open Label, Study of Safety and Effectiveness of XXX Tablets in the Treatment of Patients with Post herpetic Neuralgia in Clinical Practice

A Multicenter, Randomized, Double-blind, Placebo-controlled Study With an Open-label Run-in to Assess the Efficacy and Safety of XXX Tablets Once-daily in Subjects with Moderate to Severe Chronic Low Back Pain

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

A Randomized, Double-blind, Placebo-controlled, Parallel-group Study of XXX for the Treatment of Opioid-induced Constipation (OIC) in Subjects with Non-malignant Chronic Pain Receiving Opioid Therapy

Qualitative Study to Assess the Content Validity of the Modified Insomnia Severity Index (ISI) in Chronic Pain Patients

A Open-label, Multicenter Study to Assess the Long -term Safety of XXX Tablets 20 to 120 mg Once-daily in Subjects with Moderate to Severe Chronic Nonmalignant and Non-neuropathic Pain

A Randomized, Double-blind, Placebo-controlled, Multicenter Trial with an Enriched Study Design to Assess the Efficacy and Safety of XXX Controlled-release Tablets Compared to Placebo in Opioid-experienced Subjects with Moderate to Severe Pain due to Chronic Low Back Pain who Require Around-the-clock Opioid Therapy

Validation of Short Treatment Satisfaction Questionnaire for Use in Neuropathic Pain Patients Treated with XXX

A Phase II, 52-week, Open-label, Long-term Treatment Evaluation of the Safety and Efficacy of XXX in subjects with moderate to severe Chronic Pain

A Multicenter, Double-blind, Placebo-controlled, Cross-over Study of the Safety and Efficacy of XXX in Patients with Post-herpetic Neuralgia (PHN)

A Phase IV Multi-center, Primary Care-Based, Open-Label Study to Assess the Success of Converting Opioid-Experienced Patients, with Chronic, Moderate to Severe Pain, to XXX Using a Standardized Conversion Guide, and to Identify Behaviors Related to Prescription Opioid Abuse, Misuse, and Diversion

A Phase III, Multicenter, 12-Month, Open-Label, Single-Arm, Safety Study of XXX and XXX Extended-Release Capsules in Subjects With Moderate to Severe Chronic Non-cancer Pain

A 12-week placebo-controlled, double-blind, randomized withdrawal study to evaluate the efficacy and safety of XXX in subjects with moderate to severe chronic low back pain

A Multi-center, Double-Blind, Randomized, Placebo-controlled, Repeat Treatment (two cycle) Study of the Safety and Efficacy of XXX in Patients with Postherpetic Neuralgia

A Phase IIa, Randomized, Double-Blind, Placebo-Controlled, Two-Treatment, Two-Period Cross-Over Study to Evaluate the Safety Tolerability, Preliminary Efficacy and Systemic Exposure of XXX in Patients with Post Herpetic Neuralgia

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

A Phase IIa, Randomized, Double-Blind, Placebo-Controlled, Two-Treatment, Two-Period Cross-Over Study to Evaluate the Safety Tolerability, Preliminary Efficacy and Systemic Exposure of Topical XXXin Patients with Post Herpetic Neuralgia

A Randomized, Double-Blind, Placebo-Controlled, Multi-center Trial with an Enriched Study Design to Assess the Efficacy and Safety of XXXCompared to Placebo in Opioid-experienced Subjects with Moderate to Severe Chronic Low Back Pain

A Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Single-Dose Study of the Safety and Efficacy of Subcutaneously Administered XXX in Patients with Osteoarthritis of the Knee

A Multi-center, Double-Blind, Randomized, Placebo-Controlled, Repeat Treatment (two cycle) Study of the Safety and Efficacy of XXX in Patients with Post-herpetic Neuralgia

A Randomized, Double-Blind, Placebo-Controlled with Open-label Run-in Assessing Efficacy, Tolerability, Safety of XXX Compared to Placebo in Opioid-Naïve Subjects with Moderate to Severe, Chronic Pain Due to OA of the Knee

A Phase III Randomized, Double-Blind, Placebo-Controlled, Multi-center Study of the Analgesic Efficacy and Safety of the Subcutaneous Administration of XXX in Patients with Osteoarthritis of the Knee

A 12-Week, Randomized, Double-Blind, Placebo-Controlled, Parallel Group Study of the Efficacy and Safety of XXX in Patients with Chronic Low Back Pain

A Phase IIb, Randomized, Double-Blind, Two-Arm, Multi-center, Placebo-Controlled, Study to Assess the Efficacy and Safety of XXX in Subjects with Moderate to Severe Chronic Low Back Pain (CLBP)

A Phase IIa, Randomized, Blinded, Placebo- and Active-controlled, 2-Period Crossover Study to Assess the Analgesic Efficacy, Safety, and Tolerability of XXX in Subjects with Post herpetic Neuralgia

A Randomized, center, Double-Blind, Parallel-Group Trial with Controlled Adjustment of Dose Assessing the Analgesic Efficacy and Safety of a New Analgesic Compared with Placebo in Subjects with Painful Diabetic Peripheral Neuropathy

A Phase II Randomized, Double-Blind, Multi-Dose, Active- and Placebo-Controlled, Multi-center, Parallel Group Study of the Analgesic Effect of XXX in Adult Patients with Chronic Low Back Pain

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

Respiratory

A Phase III, 24 week treatment, multicenter, randomized, double blinded, double dummy, parallel-group, clinical trial evaluating the efficacy and safety of XXX fixed-dose combination BID compared with each monotherapy

A Phase III, Randomized, Double-blind, Placebo-controlled Study to Assess the Efficacy and Safety of XXX in patients with Uncontrolled Asthma who are on Inhaled Corticosteroids and a Second Controller Medication

A Phase IV Safety and Efficacy Study of Inhaled XXX Combination versus Inhaled XXX in the Treatment of Adolescent and Adult Subjects with Asthma

Restless Legs Syndrome

A Randomized, Double-Blind, Placebo-Controlled, Fixed-Dose, Parallel-Group Study to Compare the Efficacy, Tolerability, and Safety of 3 Doses of XXX With Placebo in the Treatment of Subjects With Moderate-to-Severe Primary Restless Legs Syndrome

A Phase IIIb, Double-blind, Randomized, Placebo-controlled Study of XXX and it's Effect on All-day Fuctioning and Quality of Life in Subjects with Moderate to Severe Idiopathic Restless Legs Syndrome

A Phase IIIb multi-center, randomized, double-blind, placebo-controlled, parallel-group, polysomnography (PSG) study to investigate safety and efficacy of the XXX in subjects with restless legs syndrome and end-stage renal disease requiring hemodialysis

A Randomized, Double-blind, Placebo-controlled, Safety and Efficacy Study of XXX in the Treatment of Moderate to Severe Restless Legs Syndrome (RLS)

A Multicenter, Open-label, 2-group, Dose Escalation Study of Monotherapy Administration of XXX in Pediatric Subjects with Idiopathic Restless Legs Syndrome

A Phase II, Open-label, Long-term, Follow-up Study to determine the Safety, Tolerability and Efficacy of XXX as Monotherapy in Adolescents with Restless Legs Syndrome

A Fixed Dose Randomized, Double-Blind, 12-week Study of XXX Subjects with Moderate to Severe Idiopathic Restless Legs Syndrome

A Randomized, Double-Blind, Placebo-Controlled, 3-Way Crossover, Multi-center Polysomnography Study of XXX and XXX in Adults with Restless Legs SyndromeA Fixed-dose, Randomized, 12-Week Placebo-Controlled, 52-week Comparator-Controlled, Double-Blind Study to Assess the Rates of Augmentation, Efficacy and Safety of XXX and XXX in Subjects with Moderate to Severe Idiopathic RLS

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

Sleep Disorders

A Phase III, Long-Term, Open-Label Safety and Maintenance of Efficacy Study of XXX in the Treatment of Excessive Sleepiness in Subjects with Narcolepsy or Obstructive Sleep Apnea

A Phase III, Twelve-Week, Double-Blind, Placebo-Controlled, Randomized, Parallel-Group, Multicenter Study of the Safety and Efficacy of XXX in the Treatment of Excessive Sleepiness in Subjects with Obstructive Sleep Apnea (OSA)

A Phase III, Twelve-week, Double-blind, Placebo-controlled, Randomized, Parallel-group, Multicenter Study of the Safety and Efficacy of XXX in the Treatment of Excessive Sleepiness in Subjects with Narcolepsy

A Phase II/III Double-Blind, Placebo-Controlled, Randomized-Withdrawal, Multicenter Study of the Efficacy and Safety of XXX with an Open-Label Pharmacokinetic Evaluation and Safety Extension in Pediatric Subjects with Narcolepsy with Cataplexy

A twelve-week, double-blind, placebo-controlled, randomized, parallel-group, multi-center study of the safety and efficacy of XXX in the treatment of excessive daytime sleepiness in subjects with narcolepsy

A Four-week, Double-blind, Placebo-controlled, Randomized, Cross-over Study of the Safety and Efficacy of XXX in the Treatment of Excessive Daytime Sleepiness

Other Indications

A Double-blind, Randomized, Placebo-controlled, Parallel-group, Phase IV Study to Evaluate the Effect of XXX on Long-term Cardiovascular Safety and COPD Exacerbations in Patients with Moderate to Very Severe COPD

A Phase III, 28-Week, Multi-Center, Randomized, Double-Blind, Parallel-Group, Active-Controlled Safety Extension Study to Evaluate the Safety and Efficacy of XXX, XXX and XXX in Subjects With Moderate to Very Severe COPD, With XXX as an Active Control

A Phase III, Randomized, Double-Blind (Test Products and Placebo), Chronic Dosing (24 Weeks), Placebo-Controlled, Parallel Group, Multi-Center Study to Assess the Efficacy and Safety of XXX, XXX , and XXX in Subjects With Moderate to Very Severe COPD, Compared With Placebo and XXX as an Active Control

A Phase III, Clinical Evaluation Of The Safety Of XXX When Administered Once Daily To Subjects With Spasticity Due To Multiple Sclerosis (MS): An Open Label, Long Term, Safety Trial

CLINICAL TRIAL EXPERIENCE (Sub-I) (*continued*):

A Phase III Multicenter, Randomized, Double-blind, Placebo-controlled Study Evaluating the Safety and Efficacy of Fixed-dose Once-daily XXX in Children and Adolescents with Tourette's Disorder

An Phase III Open-Label, Multicenter Study Evaluating the Safety and Tolerability of Once-daily XXX in Children and Adolescents with Tourette's Disorder

A Phase III Study to Assess Repeat Treatment Efficacy and Safety of XXX in Subjects with Irritable Bowel Syndrome with Diarrhea (IBS-D)

A Placebo-Controlled Randomized Withdrawal Evaluation Of The Efficacy And Safety of XXX In Subjects With Spasticity Due To Multiple Sclerosis

Open Label Long Term Safety Study: An Open-Label Study to Assess the Long-Term Safety of XXX in Patients with Opioid-Induced Constipation (OIC)

A Phase III Randomized, Double-Blind, Placebo-Controlled Study to Assess the Efficacy and Safety of XXX in Patients with Non-Cancer-Related Pain and Opioid-Induced Constipation (OIC)

A Phase III, Six-month Randomized, Active Comparator, Open label, Multi-center Study to Evaluate Patient Outcomes, Safety and Tolerability of XXX in patients with Relapsing Remitting MS who are candidates for MS Therapy Change from previous Disease Modifying Therapy

A Multicenter, Randomized, Placebo-controlled, Double-blinded Study of the Efficacy and Safety of XXX in Subjects with Opioid-Induced Bowel Dysfunction

A Randomized, Double-blind, Placebo-Controlled Study Evaluating the Efficacy, Safety, and Tolerability of 2 Doses of XXX Compared With Placebo for 12 Weeks in Patients with Moderate to Severe, Stable Chronic Obstructive Pulmonary Disease Followed by a 40-Week Evaluation of the 2 XXX Doses

A Randomized, Double-Blind, Placebo-Controlled, Clinical Evaluation of the Efficacy, Safety and Tolerability of XXX in Patients with Subjective Tinnitus AND an Open-Label, Long-Term Treatment Study to Assess the Long-Term Safety and Tolerability and Efficacy of XXX in Patients with Subjective Tinnitus

AWARDS:

- Healthgrades Honor Roll
- Top 100 Physicians, 2012
- Top Physician in Austin, TX, 2012
- 1999 Intern of the Year Teaching Award