Glenn Michael Dempsey, M.D.
Albuquerque Neuroscience, Inc.
101 Hospital Loop NE, Suite 209
Albuquerque, NM 87109

CONTACT INFORMATION:

Site Selection and Information:
Bobbie Theodore, Alliance Director
Tel. (916) 939-6696
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Email: clinicaltrials@alliancesites.com

EDUCATION:

1964-1968 Doctor of Medicine
University of Colorado School of Medicine, Aurora, Colorado

1962-1968 Master of Science
University of New Mexico, Albuquerque, New Mexico

1958-1963 Bachelor of Science
University of New Mexico, Albuquerque, New Mexico

ADDITIONAL TRAINING:

Feb. 2000 ACLS Certificate
Advanced Cardiac Life Support, Albuquerque, New Mexico

Nov. 1974 Post Graduate Nuclear Medicine course in Psychiatric Research
AEC Associated University, Oak Ridge, Tennessee

Various Conferences and CME activities for 20 or more CME Category I credits per year.

INTERNSHIP & RESIDENCE:

1968-1969 Internship
University of Maryland, Medical Center Hospital, Baltimore, Maryland

1971-1974 Residency in Psychiatry
University of Iowa, Psychopathic Hospital, Iowa City, Iowa

CERTIFICATIONS & LICENCES:

Board Certification, American Board of Psychiatry and Neurology	June, 1976 - Present
Colorado State Board of Medical Examiners, inactive status lic. 16105	June, 1971- Present
New Mexico Board of Medical Examiners, lic. #69126	Nov. 17, 1969 - Present
Colorado State Board of Medical Examiners, active status lic.# 16105	June 30, 1969-1971
National Board of Medical Examiners	July 1, 1969

MEMBERSHIPS:

American Psychiatric Association
Psychiatric Medical Association of New Mexico, Past President, 1985-1986
American Medical Association
Greater Albuquerque Medical Association
Academy of Clinical Psychiatry

INVESTIGATOR EXPERIENCE:

ADHD • Alzheimer's Disease • Anxiety • Autism Spectrum Disorder • Bipolar Disorder
Dementia • Depression • Diabetes (Type II) • Fibromyalgia • Insomnia • Post Herpetic Neuralgia
Influenza • Women's Health • Panic Disorder • Schizophrenia • Sexual Dysfunction

PROFESSIONAL EXPERIENCE:

President and Principal Investigator, Jan. 1985 - Present
Outpatient, Specializing in CNS field clinical trials
Albuquerque Neuroscience, Inc., Albuquerque, NM

Attending Physician, Inpatient, May 2014 - Present
Haven Behavioral Health Hospital, Albuquerque, NM

Locum Tenens, Feb. 2014 - Present
University of New Mexico, Albuquerque, NM

Active Medical Staff, Feb. 2014 - Present
UNM Sandoval Regional Medical Center, Albuquerque, NM

Attending Physician, Outpatient, Dec. 2012- Present
Christian Counseling Professionals, Albuquerque, NM

Senior Psychiatric Inpatient/Outpatient, March 2011 - Present
Artesia General Hospital, Albuquerque, NM

Attending Physician, Affiliate, July 1978 - Present
Presbyterian Hospital, Albuquerque, NM

RECENT PROFESSIONAL EXPERIENCE:

Attending Physician, Psychiatric in-patient, 2002-2013
Lovelace Downtown Medical Center, Albuquerque, New Mexico

Attending Physician, 2011-2012
Socorro Mental Health, Socorro, NM

Medical Director, 1992-2002
Memorial Psychiatric Hospital (Hospital sold 2002), Albuquerque, New Mexico

Attending Physician, 1984-2002
Memorial Psychiatric Hospital, (Hospital sold 2002), Albuquerque, New Mexico

Attending Physician, 1978-2002
St. Joseph Hospital, Albuquerque, New Mexico

Outpatient private psychiatric practice, 1978-2000
Albuquerque, New Mexico

Director, 1989-1991
Memorial Hospital Evaluation and Testing Institute

Attending Physician, 1979-1985
Española Hospital, Española, New Mexico

Adjunct faculty member, 1979-1982
University of New Mexico School of Medicine, Albuquerque, NM

RECENT TEACHING EXPERIENCE:

St. Francis University, Albuquerque, NM: Supervised Physician Assistant Students 2006-2014

Memorial Psychiatric Hospital. Freshman Director of Continuing Medical Education 1987-1991

CLINICAL TRIAL EXPERIENCE:

ADHD

A Phase III, Double-blind, Randomized, Multi-center, Placebo-controlled, Dose-optimization Study Evaluating the Safety, Efficacy, and Tolerability of Once-daily Dosing with Extended-release XXX in Adolescents Aged 13-17 years Diagnosed With Attention-deficit/Hyperactivity Disorder (ADHD)

A Phase IV, Randomized, Double-blind, Multicenter, Parallel-group, Active-controlled, Forced-dose Titration, Safety and Efficacy Study of XXX Compared with XXX with a Placebo Reference Arm, in Adolescents Aged 13-17 Years with Attention-deficit/Hyperactivity Disorder (ADHD)

Alzheimer's Disease

A Phase II, Randomized, Double-Blind, Placebo Contolled, parallel group study to evaluate the efficacy and safety of XXX in participants at risk for the onset of clinical symptoms of AD

A Phase III, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Efficacy and Safety study of XXX in Patient's with Prodromal to Mild Alzheimer's Disease

A Phase II, 26-Week, Double-blind, Randomized, Placebo-controlled, Parallel-group Study to Investigate the Effects of Daily Administration of XXX in Participants with Mild to Moderate Alzheimer's Disease (AD) with an Optional 26-Week Open-label Extension

A Phase III Safety and Efficacy Study of XXX in Subjects with Evidence of Early Alzheimer's Disease

A Phase III, Randomized, Double-Blind, Placebo Controlled, Multi-Center Registration Trial to Evaluate the Efficacy and Safety of XXX in Patients with Mild Alzheimer’s Disease Receiving XXX Inhibitors and/or XXX

A Phase III, 26-Week Extension Study of the Safety and Clinical Effects of XXX in Subjects with Alzheimer's Disease Currently or Previously Receiving an XXX Inhibitor Medication

A Randomized, Double-Blind, Placebo- Controlled, Parallel-Group, 26-Week, Phase III Study of Two Doses of XXX or Placebo in Subjects With Mild to Moderate Alzheimer's Disease Currently or Previously Receiving an XXX Inhibitor Medication

A Phase III, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, 18-Month Safety and Efficacy Study of XXX in Subjects with Mild Alzheimer's Disease

A Phase III, Effect of Passive Immunization on the Progression of Mild Alzheimer's Disease: XXX Versus Placebo

CLINICAL TRIAL EXPERIENCE (*continued*):

A Phase II, Multicenter, Randomized, Double-Blind, Parallel-Group, Placebo-Controlled Study to Investigate the Efficacy and the Safety of XXX Added to the Background Therapy of the Acetylcholinestrase Inhibitors XXX or XXX in Patients with Moderate Severity Alzheimer's Disease

A Phase III, Continued Efficacy and Safety Monitoring of XXX, an Anti-Amyloid 13 Antibody in Patients with Alzheimer's Disease

A Long-Term Follow-Up Study of Oral XXX in Subjects With Alzheimer's Disease

A Phase III, Extension, Multicenter, Double-Blind, Long Term Safety and Tolerability Treatment Trial of XXX in Subjects with Alzheimer's Disease who Participated in Study XXX or in Study XXX

An Open Label Extension to Evaluate The Long Term Safety And Tolerability Of XXX in Patients With Alzheimer's Disease

Multicenter, Randomized, Double-Blind, Placebo-Controlled, Phase III Study Comparing XXX and Placebo for 18 Months in Approximately 1000 Patients with Mild to Moderate Alzheimer's Disease

A Phase III, multi-center, randomized, double-blind placebo-controlled study to evaluate the safety and tolerability of XXX for up to 26-weeks in patients with mild to moderate Alzheimer's disease

A Phase II, Randomized, Double-Blind, Placebo-Controlled, Dose-Ranging, Safety and Efficacy Study of Oral XXX in Alzheimer's Disease

A Phase III, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Efficacy and Safety Trial of XXX in Patients With Mild to Moderate Alzheimer's Disease Who Are Apolipoprotein E4 Non- Carriers

A Phase III, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Efficacy and Safety Trial of XXX in Patients With Mild to Moderate Alzheimer's Disease Who Are Apolipoprotein E4 Carriers

A Phase III, Double-Blind Placebo-Controlled Study of XXX For the Treatment of Mild-to-Moderate Alzheimer's Disease

A Phase II, Double-Blind, Randomized, Placebo-Controlled Study of the Efficacy, Safety and Tolerability of 8 Week Treatment of XXX in Sleep Disturbed, Mild to Moderately Severe Alzheimer's Disease Subjects

CLINICAL TRIAL EXPERIENCE (*continued*):

Anxiety

A Phase III, Double-Blind, Placebo-Controlled, Flexible-Dose Study of XXX in Patients with Generalized Anxiety Disorder

A Phase III, Randomized, Double-Blind, Parallel-Group, Placebo-Controlled, Active-Referenced, Fixed-Dose Study Comparing the Efficacy and Safety of 3 Doses of XXX in Acute Treatment of Adults With Generalized Anxiety Disorder

A Phase III, Randomized, Double-Blind, Parallel Group, 10-Week Placebo Controlled Fixed Dose Study Of XXX And XXX Evaluating The Efficacy And Safety Of XXX For The Treatment Of Generalized Anxiety Disorder AND A 52-Week Open-Label Safety Study of XXX in Subjects With Generalized Anxiety Disorder

An 8-Week, Double-Blind, Placebo-Controlled, Phase III Trial of XXX (150-600 mg/Day) in the Adjunctive Treatment of Patients With Generalized Anxiety Disorder (GAD) Who Have Not Optimally Responded to Existing Therapies (GAD)

An Eight-week, Multicenter, Double-blind, Placebo- and XXX-controlled Study Evaluating the Efficacy and Tolerability of Two Fixed Doses of XXX in Outpatients With Generalized Anxiety Disorder

A Phase III, Multicenter, Randomized, Double-Blind, Parallel-Group, Placebo-Controlled, Active-Controlled Study of the Efficacy and Safety of Sustained-Release XXX Compared With Placebo in the Treatment of Generalized Anxiety Disorder

Generalized Anxiety, Adjunctive Comparison of Sub-Optimally Responsive to Standard Psychotherapy

A Phase III, Comparison of XXX, XXX Extended Release, and Placebo in the Treatment of Generalized Anxiety Disorder

Autism

A Phase II, Open-Label Study of the Safety and Tolerability of XXX in Pediatric Patients with Autism, Asperger's Disorder, or Pervasive Developmental Disorder Not Otherwise Specified (PDD-NOS) and To Identify Responders for Participation In The Follow-Up Randomized Withdrawal Study

A Phase II, Open-Label Extension Study of the Safety and Tolerability of XXX in Pediatric Patients with Autism, Asperger's Disorder or Pervasive Developmental Disorder Not Otherwise Specified (PDD-NOS)

CLINICAL TRIAL EXPERIENCE (*continued*):

A Phase II, Double-Blind, Placebo-Controlled , Randomized Withdrawal Study of the Safety and Efficacy of XXX in Pediatric Patients with Autism Asperger's Disorder, or Pervasive Developmental Disorder Not Otherwise Specified (PDD-NOS) Previously Treated with XXX

Bipolar Disorder

A Randomized, Double-Blind, Placebo-Controlled, Phase III Study To Evaluate the Efficacy and Safety of Once a Day, XXX 0.1, 0.4, and 0.8mg as an Adjunctive Therapy to Treatment as-Usual in the Maintenance Treatment of Bipolar I Disorder in Adult Subjects

A Randomized, Double-Blind, Placebo-Controlled, Phase III Study to Evaluate the Efficacy and Safety of Once a Day, XXX Tablet for Sublingual Administration XXX 0.1, 0.4, and 0.8 mg as an Adjunctive Therapy in the Treatment of Acute Depressive Episodes Associated with Bipolar 1 Disorder in Adult Subjects

A Phase II, Double-Blind, Placebo-Controlled Evaluation of the Safety and Efficacy of XXX in Patients With Bipolar Depression

A Phase III, Evaluation of XXX as an Add-on Treatment for Bipolar I Disorder in Children and Adolescents, 10 to 17 Years of Age

A Phase III, 24-Week, Flexible-Dose, Open-Label Extension Study of XXX for the Treatment of Bipolar I Depression

A Phase III, Randomized, 6-Week, Double-Blind, Placebo-Controlled, Flexible-Dose, Parallel-Group Study of XXX for the Treatment of Bipolar I Depression

A Phase III, Randomized, 6-Week, Double-Blind, Placebo-Controlled, Flexible-Dose, Parallel-Group Study of XXX Adjunctive to XXX or XXX for the Treatment of Bipolar I Depression

A Four Week, Double-Blind, Placebo Controlled Phase III Trial Evaluating The Efficacy, Safety And Pharmacokinetics Of Flexible Doses Of Oral XXX In Children And Adolescents With Bipolar I Disorder (Manic Or Mixed) AND 26-Week Open-Label Extension Study Evaluating The Safety And Tolerability Of Flexible Doses Of Oral XXX In Children And Adolescents With Bipolar I Disorder (Manic Or Mixed)

A Phase III, Multicenter, Randomized, Placebo-Controlled, Parallel-Group, Double-Blind, Phase III Study to Compare the Efficacy and Safety of XXX Versus Placebo as Adjunct Therapy With Mood Stabilizers (XXX or XXX) for the Treatment of Alcohol Dependence in Patients With Bipolar I Disorder

CLINICAL TRIAL EXPERIENCE (*continued*):

Dementia

An Open-Label, Multicentre, One Year Extension of the Evaluation of the Safety of XXX in Patients With Dementia Associated With Cerebrovascular Disease

A Phase III, 24-Week, Multi-Center, Randomized, Double-Blind, Placebo-Controlled Evaluation of the Efficacy, Safety and Tolerability of XXX in Patients With Dementia Associated With Cerebrovascular Disease

Depression

An Open-label, Long-term Safety Study of XXX as Adjunctive Therapy in Patients with Major Depressive Disorder

A Randomized, Double-blind, Placebo-controlled, Multicenter Study of XXX as Adjunctive Therapy in the Prevention of Relapse in Patients with Major Depressive Disorder

A Randomized, Double-blind, Placebo-controlled, Multicenter Study of XXX as Adjunctive Therapy in Major Depressive Disorder

A Phase III, Multicenter Study of the Long-term Safety and Tolerability of XXX for the Adjunctive Treatment of Major Depressive Disorder in Adults who Have an Inadequate Response to Antidepressant Therapy

A Phase III, Efficacy and Safety Study of XXX for the Adjunctive Treatment of Major Depressive Disorder

A Phase II, Double-Blind, Placebo-Controlled, Randomized Add-On Study of XXX For Patients With Major Depressive Disorder (MDD) Who have had an Inadequate Response to Current Antidepressant Therapy

A Phase III, Double-Blind, Placebo-Controlled Evaluation of the Safety and Efficacy of XXX in Adolescent Patients With Major Depressive Disorder

A Phase II, 8-week, Randomized, Double-blind, Placebo-controlled, Parallel-group, Multi-center Study of the Efficacy and Safety of XXX Administered Once Daily in Patients with Major Depressive Disorder

A Phase III, Long-term, Open-label Study of Safety and Tolerability of XXX as Adjunctive Therapy in Major Depressive Disorder

A Phase III, Double-Blind, Placebo-Controlled Study of XXX As Adjunctive Therapy In Major Depressive Disorder

CLINICAL TRIAL EXPERIENCE (*continued*):

A Phase III, Open-label, Multicenter, 12 month Extension Safety and Tolerability Study of XXX in Combination with an Antidepressant in the Treatment of Adults with Major Depressive Disorder with Residual Symptoms or Inadequate Response Following Treatment with an Antidepressant

A Phase II, Multicenter, Double-blind, Parallel-group, Randomized, Placebo-controlled, Forced-dose Titration, Dose-ranging Efficacy and Safety Study of XXX in Combination with an Antidepressant in the Treatment of Adults with Major Depressive Disorder with Inadequate Response to Prospective Treatment with an Antidepressant

A Phase III, Long-Term, Open-Label, Flexible-Dose, Extension Study Evaluating the Safety and Tolerability of XXX (15 and 20 mg) in Subjects With Major Depressive Disorder

A Phase III, 6-Month, Open-Label, Multi-Center, Flexible-Dose Extension Study to the XXX Study to Evaluate the Safety, Tolerability AND Efficacy of XXX Sustained-release (SR) Tablets in the Treatment of Children and Adolescent Outpatients with Major Depressive Disorder

A Phase III, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study to Evaluate the Efficacy, Safety and Tolerability of XXX Sustained-Release (SR) in the Treatment of Children and Adolescent Outpatients with Major Depressive Disorder

A Phase II, Multicenter, Double-Blind, 58-week Rollover Study to assess the Safety and Tolerability of XXX in Patients with Treatment Resistant Major Depression

A Phase II, Multicenter, Randomized, Double-blind, Active-Controlled, Comparative, Fixed-Dose, Dose Response Study of the Efficacy and Safety of XXX in Patients with Treatment Resistant Major Depression

A Phase III, Randomized, Double-Blind, Parallel-Group, Placebo-Controlled, Fixed-Dose Study Comparing the Efficacy and Safety of 2 Doses (10 and 15 mg) of XXX in Acute Treatment of Adults with Major Depressive Disorder

A Phase III, Randomized, Double-Blind, Parallel-Group, Active-Controlled, Flexible-Dose Study Evaluating the Effect of XXX vs XXX on Sexual Functioning in Adults With Well-Treated Major Depressive Disorder Experiencing Selective Serotonin Reuptake Inhibitor—Induced Sexual Dysfunction

Phase III, XXX Compared to Placebo as Adjunctive Therapy to SSRI in the Prevention of Symptom Re-emergence in Major Depressive Disorder

A Phase III, 6-Month, Open-Label, Flexible-Dosage (150-200 mg/day) Extension Study of the Safety and Efficacy of XXX Treatment as Adjunctive Therapy in Adults With Major Depression Associated With Bipolar I Disorder

CLINICAL TRIAL EXPERIENCE (*continued*):

A Phase III, Double-Blind, Placebo-Controlled, Parallel-Group, Fixed Study to Evaluate the Efficacy and Safety of XXX Treatment (150 and 200 mg/day) as Adjunctive Therapy in Adults With Major Depression Associated With Bipolar I Disorder

A Phase III, Multicenter, 52-week, Open-label Study to Assess the Safety and Tolerability of an Oral XXX Combination Therapy in Patients with Major Depressive Disorder

A Phase III, Multicenter, Randomized, Double-blind Study to Evaluate the Efficacy, Safety and Tolerability of an Oral XXX Combination Therapy in Patients With Major Depressive Disorder

A Phase III, Long-Term, Open-Label, Safety Study of XXX 12 to 18 mg Once Daily as Adjunctive Treatment for Patients with Major Depressive Disorder Who are Partial Responders to Selective Serotonin Reuptake Inhibitor Treatment

A Phase III, 52-week, multi-center, open-label study of the safety and tolerability of XXX sublingual tablets in patients with Major Depressive Disorder (MDD)

A Phase III, 8-week, Randomized, Double-blind, Placebo-controlled, Parallel-group, Multi-center Study of the Efficacy and Safety of XXX 0.5 mg and 1 mg Sublingual Tablets Administered Once Daily in Patients With Major Depressive Disorder (MDD)

A Phase III, Double-Blind, Efficacy and Safety Study of XXX versus Placebo in the Treatment Of Children and Adolescents with Major Depressive Disorder

A Phase Ilb, Multicenter, Randomized, Double-blind, Parallel Group, Placebo-controlled Efficacy and Safety Study of Adjunctive XXX in Subjects with Severe Major Depressive Disorder (MDD) and a History of Poor Response to Antidepressants

A Phase II/ III, Randomized, Double-Blind Comparison of XXX and Placebo and Long Term Treatment With XXX in Adult Patients With Major Depressive Disorder

A Phase II, Double-Blind, Placebo-Controlled Study Examining The Safety, Efficacy, and Tolerability of XXX in Subjects With Major Depressive Disorder (Including Atypical and Melancholic Features)

A Phase II, Six-Week, Multicenter, Randomized, Double Blind, Placebo-Controlled, Parallel-Group Study Evaluating the Efficacy, Safety, and Tolerability of XXX compared to Placebo in Female Subjects, Diagnosed with Major Depressive Disorder

A Phase III, 52-week, Randomized, Double-blind, Placebo-controlled, Multi-center, Parallel-group Study of the Long-term Efficacy, Tolerability and Safety of XXX 25 and 50 mg in the Prevention of Relapse of Major Depressive Disorder (MDD) Following Open-label Treatment of 16-24 Weeks

CLINICAL TRIAL EXPERIENCE (*continued*):

A Phase II, Multicenter, Randomized, Double-Blind, Placebo and XXX Controlled Trial of the Safety and Efficacy of XXX in the Treatment of Outpatients With Major Depressive Disorder

An 8-week, Randomized, Double-blind, Fixed Dosage, Placebo-controlled, Parallel-group, Multi-center Study of the Efficacy, Safety and Tolerability of XXX 25 mg and 50 mg in the Treatment of Major Depressive Disorder (MDD) Followed by a 52-week, Open-label Extension

An Eight-Week, Double-Blind, Placebo-Controlled, Multicenter Study With XXX as Positive Control, Evaluating the Efficacy, Safety, Tolerability of Two Fixed Doses of XXX in Outpatients With MDD

A Multicenter, Double-Blind, Randomized, Parallel-Group, Placebo-Controlled and Active-Controlled Phase III Study of the Efficacy and Safety of XXX Sustained-Release as Monotherapy in the Treatment of Patients With MDD

Depression, Double-Blind, Safety & Efficacy with Sub-Optimally Responsive to Standard Anti-Depressant Therapy

A Phase III, Randomized, Double-Blind, Placebo-Controlled, Parallel Group Study of the Safety and Efficacy of Three Dose Levels of XXX Plus an Antidepressant vs. Placebo Plus an Antidepressant in the Treatment of Psychotic Symptoms in Patients With Major Depressive Disorder With Psychotic Features (PMD)

A Phase III, Dose Escalation, Double-Blind Treatment With XXX Hydrochloride Once Daily Dosing for Evaluation of Safety in Major Depression

A Phase III, XXX Versus Placebo in the Treatment of Elderly Patients With Major Depressive Disorder

Additional 10+ Phase I-IV depression studies conducted including Psychotic Depression, Child Depression, Treatment Resistant Depression, and Case Comparison Study.

Fibromyalgia

An open-label extension of XXX for 52 weeks in pain associated with Fibromyalgia

A Phase III, Randomized, Double-Blind, Placebo- and Active-Controlled Study of XXX in Subjects with Pain Associated with Fibromyalgia

A Phase III, Effect of XXX 30/60 mg Once Daily versus Placebo in Adolescents with Juvenile Primary Fibromyalgia Syndrome

A Phase II, multicenter, randomized, double-blind controlled withdrawal study to evaluate the safety, tolerability, and efficacy of XXX in pediatric patients with primary fibromyalgia

CLINICAL TRIAL EXPERIENCE (*continued*):

A Phase II, multicenter, open-label, 52 week extension study to evaluate the safety and efficacy of XXX in pediatric patients with primary fibromyalgia

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 Long-Term, Open-Label, Safety and Efficacy Extension Study of XXX in Subjects With Fibromyalgia

A Phase III, Randomized, Double-Blind, Placebo-Controlled, Safety and Efficacy Study of XXX in Subjects With Fibromyalgia

Migraine

A Multicenter, Open Label, Long-Term Safety Study of XXX in Patients with Acute Migraines

A Phase III, Double-Blind, Randomized, Placebo Controlled, Safety, Efficacy, Trial of XXX for the Acute Treatment of Migraine

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

A Multicenter, Randomized, Open-Label, Extension study to evaluate the Long-term Safety and Tolerability of XXX in the Acute Treatment of Migraine with or without Aura

A Multicenter, Randomized, Open-Label Extension Study to Evaluate Long-Term Safety and Tolerability of Oral XXX in the Acute Treatment of Migraine with Aura

A Phase III, Long term, Open-Label Safety Study of XXX in Patients with Migraine

Panic Disorder

Double-Blind Study of XXX in Adults With Panic Disorder

Double-Blind Study of XXX in Adolescents With Panic Disorder

High dose XXX Withdrawal in Panic Disorder

Schizophrenia

A Phase III, 52-Week, Open-Label, Extension Study of XXX for the Adjunctive Treatment of Schizophrenia

CLINICAL TRIAL EXPERIENCE (*continued*):

A Phase II, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Efficacy and Safety of XXX as Adjunctive Treatment for the Negative Symptoms of Schizophrenia

A Phase III Study to Evaluate Weight Gain of XXX Compared to XXX in Adults with Schizophrenia

A Phase III, Study to Evaluate Weight Gain of XXX Compared to XXX in Adults with Schizophrenia

A Phase III, Multicenter 26-Week Extension Study to Evaluate the Safety and Clinical Effects of Prolonged Exposure to 1 and 2 mg Doses of XXX, an Alpha-7 Nicotinic Acetylcholine Receptor Agonist, as an Adjunctive Pro-cognitive Treatment in Subjects with Schizophrenia on Chronic Stable Atypical Antipsychotic Therapy

A Randomized, Double-blind, Placebo-controlled, Parallel, 12-Week, Phase III Study of 2 Doses of an Alpha-7 Nicotinic Acetylcholine Receptor Agonist XXX or Placebo as an Adjunctive Pro-cognitive Treatment in Schizophrenia Subjects on Chronic Stable Atypical Antipsychotic Therapy

A Phase IV, 12-week, Randomized, Multi-center, Open-Label, XXX, (12-24 mg/Day), Flexible Dose Study Assessing Efficacy, Safety and Tolerability of Two Switch Approaches in Schizophrenia Patients Currently Receiving XXX, XXX, or XXX

A Phase III, Population Pharmacokinetic Study in Adolescent Patients With Schizophrenia or Bipolar I Disorder Treated With XXX

Sleep

A Long-Term Multicenter, Randomized, Double-Blind, Controlled, Parallel-Group Study of the Safety and Efficacy of XXX in Subjects With Insomnia Disorder

A Phase III, Efficacy and Safety of 2 mg/Day of XXX on Sleep Maintenance Insomnia With a Sub-study of the Effect of XXX on Stable Type II Diabetes Mellitus: a 12-week, Multi-center, Randomized, Double-blind, Placebo-controlled Study

A Phase IV, Randomized, Double Blind, Placebo-Controlled, Parallel Group Study to Demonstrate the Subjective Treatment Effects of XXX on Sleep Using a Post Sleep Questionnaire - Interactive Voice Response System (PSQ-IVRS) in an "At-Home Setting" in an Adult Population With Chronic Insomnia

Other Indications

A Phase II, A Randomized, Double-blind, Parallel Group, Multicenter, Placebo-controlled, Dose-ranging Study to Evaluate the Glycemic Effects, Safety, and Tolerability of XXX Delayed-Release in Subjects with Type 2 Diabetes Mellitus

CLINICAL TRIAL EXPERIENCE (*continued*):

A prospective, multicenter, randomized, double-blind, Sham-controlled study to assess the efficacy and safety of the XXX capsule administered 5 times per week, XXX Chronic Idiopathic Constipation

A Safety & Efficiency Study of the XXX Capsule in Aiding Patients with Functional Constipation

A multinational, multicenter, prospective Double-Blind, sham controlled, randomized study to assess the performance, efficacy and safety of Vibrating Capsule medical device in aiding relieving Constipated Individuals

A Phase II, Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel-group Study of the Safety and Efficacy of a Single Treatment of XXX in Patients With Postherpetic Neuralgia

A Phase III, Multicenter, Randomized, Double-blind, Placebo Controlled Study to Evaluate the Efficacy and Safety of Intravenous XXX in Subjects With Uncomplicated Influenza

A Phase III, Safety of XXX Versus Placebo in Women Taking a Selective Serotonin Reuptake Inhibitor or Norepinephrine Serotonin Reuptake Inhibitor With Decreased Sexual Desire

Double-blind Study of XXX in the Treatment of Weight-gain

Double-blind Study of XXX Effects on Sexual Functioning

AWARDS:

Physician's Recognition Award in Continuing Medical Education,
American Medical Association. 1976-Present

Elected to Waring Society, University of Colorado, School of Medicine.
Paper presented: "Penitentes: Neoplatonists of New Mexico" 1966-1968

Robert H. Felix Metropolitan Mental Health Award for Best Freshman
Psychiatric Paper at University of Colorado: "The Spanish-American as a Patient" 1965

APPOINTMENTS:

Medical Director, Memorial Psychiatric Hospital, Albuquerque, NM 1992-2002
Assistant Professor, Department of Psychiatry, University of Kentucky
School of Medicine, Lexington, Kentucky 1976-1978
Associate Chief, Psychiatry Service, V.A. Hospital, Lexington, Kentucky 1976-1978
Clinical Instructor, Psychiatry, Columbia College Physicians & Surgeons,
Presbyterian Hospital, NYC 1974-1976

APPOINTMENTS (*continued*):

Research Psychiatrist I, New York State Psychiatric Institute, NYC, NY	1974-1976
Chief of Medical Staff, Embudo Presbyterian Hospital, Embudo, NM	1970-1971
Staff Physician, Embudo Presbyterian Hospital, Ernbudo, NM	1969-1971
Research Assistant, Lovelace Foundation for Medical Research, Department of Physiology, Albuquerque, New Mexico	1963-1964
Graduate Assistant, University of New Mexico Department of Biology	1962-1963

PRESENTATIONS:

Columbia College of Physicians and Surgeons and the New York State Psychiatric Institute

Review of Cation Physiology in Psychiatric Illness
Role of Masked Depression in the Setting of a Medical Clinic

University of Kansas School of Medicine

Schizo-Affective Disorder, Diagnosis and Treatment — October 1985

University of Iowa

Anorexia Nervosa
Use of Lithium in a Patient with Affective Disorder Presenting as Drug Psychosis
Some Side-Effects of Neuroleptic Drugs:
Use of Lithium in Schizoaffective Disorder with Tardive Dyskinesia
Use of Tofranil in Adolescent Depression and Behavior Problems
Medical Grand Rounds — Depression and Suicide
Two presentations to the Iowa Genetics Club:
"Familial Retardation"
"A Family with Genetic Translocation and Multiple Psychiatric Disorders"

University of Kentucky

Use of Clinical Laboratory in Management of Lithium Patients
Lithium Toxicity
Electrolyte and Other Metabolic Disturbances Associated with Alcoholism

VI World Congress of Psychiatry, Honolulu, Hawaii

A 30 - year Follow-up of Atypical (Schizoaffective) Schizophrenia

University of New Mexico, School of Medicine, Department of Psychiatry Grand Rounds, April 1986

Electroconvulsive Therapy
Psychiatric Manifestations of Temporal Lobe Seizure
Wolverhampton England, Second British Lithium Congress, September 1987 The Concurrent Use of Lithium and Clonazepam

PRESENTATIONS (*continued*):

Albuquerque, New Mexico

Diagnosis and Management of Drug Overdoses and Adverse Reactions to Psychiatric Drugs and Illicit Drugs
Diagnosis and Management of Organic Brain Syndromes and Dementia New Therapeutic Approaches in Psychopharmacology
Diagnosis and Management of Depression, with Emphasis on the Use of Antidepressant Medication
Survey of Medical Psychiatry, presentation to Alliance of Mentally Ill Association, 1991
Biological Aspects of Affective Disorders, presentation to Depressive and Manic Depressive Association
Child and Adolescent Psychopharmacology, presented to Memorial Hospital staff, 1992
Multiple lectures systematically reviewing DSM-III-R & DSM-IV, presented to Memorial Hospital Staff, 1992 to 2002

Bibliography available upon request