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Albuquerque, NM 87109

**CONTACT INFORMATION:**

Site Selection and Information:  
Bobbie Theodore, Alliance Director  
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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  
Constipation • Dementia • Depression • Diabetes (Type II) • Fibromyalgia • Influenza • Insomnia  
Migraine • Post Herpetic Neuralgia • 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 – September 2016  
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 – July 2015  
Christian Counseling Professionals, Albuquerque, NM

*Senior Psychiatric Inpatient/Outpatient*, March 2011 – January 2019  
Artesia General Hospital, Albuquerque, NM

**PROFESSIONAL EXPERIENCE (*continued*):**

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

*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-Controlled Study of XXX in Subjects with Alzheimer's Disease

A Phase II/III, Randomized, Double-blind, Placebo-controlled, Multicenter Studies to Evaluate the Safety and Efficacy of XXX as a Treatment for Subjects With Mild Alzheimer's Disease and Impaired Glucose Tolerance

A Phase II, Randomized, Double-Blind, Placebo Controlled, 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

**CLINICAL TRIAL EXPERIENCE (continued):**

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

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 Acetylcholinesterase 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

**CLINICAL TRIAL EXPERIENCE (*continued*):**

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

***Anxiety***

A Phase III, Multicenter, Randomized, Double-Blind, Placebo Controlled Trial of XXX in Generalized Anxiety Disorder

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

**CLINICAL TRIAL EXPERIENCE (*continued*):**

***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)

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 I 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

**CLINICAL TRIAL EXPERIENCE (continued):**

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

***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***

A Phase IIa, Randomized, Double-blind, Placebo-controlled Proof of Concept Study to Evaluate the Effects of Oral XXX Versus Placebo in Subjects With Major Depressive Disorder

A Phase III, Randomized, Double-blind, Placebo-controlled Study of the Efficacy and Safety of XXX with a Fixed, Repeated Treatment Regimen on Relapse Prevention in Adults with Major Depressive Disorder

A Phase III, Open-label, 1-year Study of the Safety, Tolerability, and Need for Re-treatment with XXX in Adult Subjects with Major Depressive Disorder

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



**CLINICAL TRIAL EXPERIENCE (continued):**

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

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

**CLINICAL TRIAL EXPERIENCE (*continued*):**

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

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

**CLINICAL TRIAL EXPERIENCE (*continued*):**

A Phase IIb, 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

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)

**CLINICAL TRIAL EXPERIENCE (*continued*):**

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

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

***Insomnia***

A Phase III Multicenter, Open-Label Study to Evaluate the Substitution of XXX with XXX for Monotherapy of Insomnia

A Randomized, Double-blind, Placebo-controlled Study of the Safety, Tolerability, and Efficacy of XXX Compared to Placebo in Adult Subjects with Comorbid Major Depressive Disorder and Insomnia

**CLINICAL TRIAL EXPERIENCE (*continued*):**

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

***Migraine***

A Randomized, Double-Blind, Parallel Group, Placebo-Controlled, Multicenter Study to Evaluate the Efficacy, Safety, and Tolerability of Single Doses of XXX Nasal Powder in the Acute Treatment of 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

**CLINICAL TRIAL EXPERIENCE (*continued*):**

***Schizophrenia***

A Phase II Randomized, Double-blinded, Placebo-controlled Parallel Group Trial to Examine the Efficacy and Safety of XXX Once Daily With Adjunctive Computerized Cognitive Training Over 12 Week Treatment Period in Patients With Schizophrenia

A Phase IV Post-XXX Study Interviews to obtain feedback on the digital therapeutic used in the XXX trial as well as new ideas for a future version

A Phase II, Randomized, Sham-Controlled Study of XXX as an adjunct to standard-of-care treatment for schizophrenia

A Phase IIa, Randomized, Double-Blind, Placebo-Controlled, Parallel-group Study to Assess the Safety and Efficacy of XXX as Add-on Treatment for Cognitive Impairment in Subjects with Schizophrenia on Stable Doses of Antipsychotic Medication

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

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

**CLINICAL TRIAL EXPERIENCE (*continued*):**

***Other Indications***

A Phase IV Freestyle XXX Glucose Monitoring System Post Approval Study for Pediatric Patients

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

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,<br>American Medical Association.  | 1976-Present |
| Elected to Waring Society, University of Colorado, School of Medicine.<br>Paper presented: "Penitentes: Neoplatonists of New Mexico"                   | 1966-1968    |
| Robert H. Felix Metropolitan Mental Health Award for Best Freshman<br>Psychiatric Paper at University of Colorado: "The Spanish-American as a Patient" | 1965         |

**APPOINTMENTS:**

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|--|-----------|
| <i>Medical Director</i> , Memorial Psychiatric Hospital, Albuquerque, NM   | 1992-2002 |
| <i>Assistant Professor</i> , Department of Psychiatry, University of Kentucky<br>School of Medicine, Lexington, Kentucky   | 1976-1978 |
| <i>Associate Chief</i> , Psychiatry Service, V.A. Hospital, Lexington, Kentucky  | 1976-1978 |
| <i>Clinical Instructor, Psychiatry</i> , Columbia College Physicians & Surgeons,<br>Presbyterian Hospital, NYC             | 1974-1976 |
| <i>Research Psychiatrist I</i> , New York State Psychiatric Institute, NYC, NY   | 1974-1976 |
| <i>Chief of Medical Staff</i> , Embudo Presbyterian Hospital, Embudo, NM   | 1970-1971 |
| <i>Staff Physician</i> , Embudo Presbyterian Hospital, Embudo, NM  | 1969-1971 |
| <i>Research Assistant</i> , Lovelace Foundation for Medical Research, Department<br>of Physiology, Albuquerque, New Mexico | 1963-1964 |
| <i>Graduate Assistant</i> , 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"



**PRESENTATIONS (*continued*):**

**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

**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

**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

***Bibliography available upon request***