PRN PACIFIC RESEARCH NETWORK
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JILL E. CRUSEY, Ph.D.

EDUCATION

1982 Ph.D: Clinical Northwestern University – University of Chicago, Developmental Psychology & Research, Chicago, Illinois
Degree: Ph.D.

1975 Masters: University of Illinois, Psychology and Recreational Therapy
Champaign, Illinois, Degree: M.A.

1973 Undergraduate: University of Illinois, Psychology and Recreational Therapy
Champaign, Illinois, Degree: B.A.

2007 In Progress Masters University of California, San Diego, and California Western
School of Law – Masters Program in Health Law

TRAINING

1981– 1982 Internship: Clinical Research Associate, CNS Research Division Psychological
Services of San Diego, San Diego, California

1979 – 1980 Internship: Clinical Associate, Tavistock Clinic, England Royal Free Hospital
London, England

MEDICAL EXPERIENCE

1982 – Present Associate Clinical Director, CNS Research Division, Pacific Research Network, San Diego, California

1982 – Present Clinical Psychologist, Private Practice, San Diego, California

1982 – Present Adjunct Professor, Department of Psychiatry, University of California San Diego, La Jolla, California

1984 – 1990 Adjunct Professor, Graduate/Doctoral Courses, California School of Professional Psychology, San Diego, California

1982 Geriatric Consultant, Senior Citizens Community Medical Clinic, San Diego, California

1982 – 1983	Adjunct Professor, Upper Level Division, Department of Psychology Chapman College, San Diego, California
1980 – 1981	Psychological Assistant, Women's Counseling Association of San Diego San Diego, California
1978 – 1979	Counselor for Patients and Families, Oncology Service, Northside Veterans Administration Hospital, Chicago, Illinois
1977 – 1979	Mental Health Specialist, Northwestern Memorial Hospital, Institute of Psychiatry, Chicago, Illinois
1977 – 1979	Program Consultant for Nursing Home and Residential Facilities Chicago, Illinois and Outlying Suburbs
1975	Resource Development Specialist with Focus to Development of Two Year College Training Program in Recreation, University of Illinois, Champaign, Illinois
1974 – 1975	Teaching Assistant, Department of Psychology and Sports Medicine University of Illinois, Champaign, Illinois
1973	Program Director of Camping Services for the Handicapped, Camp Allen, Inc. Bedford, New Hampshire

CERTIFICATION AND LICENSURE

LICENSURE:

State of California
License Number: PSY12840
Expiration Date: December 31, 2013

PRESENTATIONS

Many Conferences With Focus to Dealing with Clinical and Treatment Topics for Specific Age and Gender Related Issues.

PUBLICATIONS

1. *Menopause As A Developmental Process: Aging and Life Cycle Issues,* Edited by Dr. D. Gutmann, University of Chicago Press, Fall 1988.

2. *Dynamic Therapy With a Sixty-Two Year Old Man: Race Against Time,* Edited by Dr. R. Nemiroff and Dr. C. Colarusso, Pleneum Publishing, Spring 1985.

3. *Sexual Issues and Alzheimer's Disease*, Caring, Volume 1, No. 9, Janssen Research Foundation, August, 1985.

4. *Menopause and Its Developmental Implications*, an Unpublished Dissertation, Northwestern University, Chicago, IL, 1983.

5. Developing Curriculum for Community College Students*: Competency Based Skills in Working with the Handicapped,* an Unpublished Thesis, University of Illinois, Champaign, IL 1978.

6. *Mid-Life Shifts and The Emergence of the Feminine Self*, Clinical Implications for an Aging Population, Edited by D. Gutmann, Ph.D., University of Chicago Press, in press.

PROFESSIONAL ORGANIZATIONS

Advancement of Professional Psychology
Gerontological Society of America
Southern Caregivers Resource Center, Board of Directors, San Diego, California

CLINICAL RESEARCH EXPERIENCE

1. "A Placebo-Controlled Study of XXXXX and XXXXXX in Patients with Generalized Anxiety Disorder" (2597)

2. "Open-Label Safety Study of XXXXX in Patients with Anxiety Disorders" (2387)

3. "A Randomized, Single-Dosed, Double-Blind, Parallel Study Comparing XXXXX 20 mg, XXXXX 10 mg and Placebo in Preventing Heartburn When Administered Immediately Prior to a Provocative Breakfast Meal" (2666)

4. "A Double-Blind, Placebo-Controlled Study of XXXXX in the Treatment of Behavioral Agitation in Elderly Patients with Dementia" (2637)

5. "A Multicenter, Randomized, Double Blind, Placebo Controlled Flexible Dose Study of XXXXX in the Treatment of Institutionalized Patients with Psychosis Associated with Dementia of the Alzheimer's Type" (2721)

6. "A Multi-center, Double-Blind, Placebo-Controlled, Randomized Fixed Dose Study of XXXXX in the Treatment of Depressed Patients" (2721)

7. "Long-Term Open Label Treatment with XXXXX for Evaluation and Safety" (1743)

8. "A Double-Blind, Placebo-Controlled Dose Finding Study Evaluating the Safety and Efficacy of XXXXX 1.5, 6 and 24 mg/day (0.5, 2, 8 mg tid) in the Treatment of Major Depressive Disorder" (2553)

9. "Safety and Efficacy of Long-Term Administration of XXXXX in the Treatment of Major Depressive Disorder: A 4 Month Double-Blind Extension to Study XXXXX" (2553x1)

10. "Safety of Open-Label Standard Antidepressant Therapy in the Treatment of Major Depressive Disorder: A 1 Month Follow-Up After Termination of Study XXXXX" (2553x2)

11. “A Double-Blind, Placebo-Controlled, Parallel-Group Comparison of XXXXX Extended-Release Capsules and XXXXX in Outpatients with Generalized Social Anxiety Disorder” (2672)

12. “Long-Term Safety and Efficacy of XXXXX in the Treatment of Alzheimer’s Disease” (1138)

13. “Long-Term Safety of XXXXX in the Treatment of Alzheimer’s Disease” (2485)

14. “Open-Label use of Synthetic XXXXX in the Treatment of Alzheimer’s Disease” (1138x1)

15. “XXXXX For the Treatment of Mild Cognitive Impairment and Prevention of Conversion to Alzheimer’s Disease” (1160)

16. “The Safety and Efficacy of XXXXX in Slowing the Progression of the Symptoms of Alzheimer’s Disease” (2486)

17. “The Safety and Efficacy of XXXXX 25mg in Delaying the Progression of the Symptoms of Alzheimer’s Disease in Patients with Probable AD” (2855)

18. “Randomized, Double-Blind, Placebo-Controlled Multi-center Trial to Demonstrate the Clinical Efficacy and Safety of Two Different Doses of XXXXX in Patients Suffering From Dementia of the Alzheimer’s Type According to DSM-IV and NINCDS/ADRDA Criteria” (1250)

19. “A Randomized, Double-Blind, Placebo-Controlled Trial to Evaluate the Safety and Efficacy of XXXXX Sodium Therapy for Agitation in Nursing Home Residents with Probable or Possible Alzheimer’s Disease” (2860)

20. “A Placebo-Controlled Study of XXXXXX in patients With Social Phobia.” (2257A1)

21. “A 10-Week, Randomized, Double-Blind, Placebo-Controlled Study of XXXXXX and XXXXXX in Patients With Social Phobia.” (3315)

22. “A Randomized, Double-Blind, Placebo-Controlled Trial to Evaluate the Safety and Efficacy of XXXXXX Sodium Therapy for Agitation in Nursing Home Residents with Probable or Possible Alzheimer’s Disease.” (2860)

23. “XXXXXX: Double-Blind, Placebo-Controlled, Dose-Response Study of Tolerability, Safety, and Efficacy in Patients with Early Parkinson’s Disease” (3401)

24. “XXXXXX: Open-Label, Long Term, Flexible Dose Study of Safety, Tolerability, and Therapeutic Response in Patients with Parkinson’s Disease.” (3429)

25. “Efficacy and Safety of a Flexible Dose of XXXXXX Versus Placebo in the Treatment of
Psychosis of Alzheimer’s Disease.” (2956)

26. “A Dose-Ranging, Double-Blind, Placebo-Controlled, Safety, Tolerability, and Efficacy Study
of XXXXXX in Patients with Major Depressive Disorder By DSM-IV Criteria.” (3348)

27. “Placebo Controlled Evaluation of XXXXXX in the Treatment of Alzheimer’s Disease: Safety and Efficacy of Controlled Release Formulation.” (2918)

28. “A Randomized Double-Blind Placebo-Controlled Trial to Evaluate the Efficacy and Safety of XXXXXX in Subjects with Mild Cognitive Impairment (MCI) Clinically at Risk for Development of Clinically Probable Alzheimer’s Disease.” (2919)

29. “A Well-Controlled Safety and Efficacy Study of XXXXXX In Subjects with Mild to Moderate
Alzheimer’s Disease.” (3261)

30. “An 8-Week, Double-Blind, Randomized, Multicenter, Flexible-Dose, Placebo-Controlled
Study of XXXXXX in Patients with Panic Disorder.” (3789)

31. “A Randomized, 26-Week, Double-Blind, Placebo-Controlled Trial to Evaluate the Safety and
Efficacy of XXXXXX in the Treatment of Dementia Secondary to Cerebrovascular Disease. (3666)

32. “A 6-Week, Double-Blind, Randomized, Multicenter, Fixed-Dose, Placebo-Controlled Study of
XXXXXX Dosed Twice a Day in Patients with Generalized Anxiety Disorder.” (3979)

33. “An Open-Label Extension Trial to Assess the Long-Term Safety of a Controlled Release
Formulation of XXXXXX HBr in the Treatment of Alzheimer’s Dementia.” (2918x1)

34. “A Randomized, Double-Blind, Placebo-Controlled Trial to Evaluate the Efficacy and Safety
of XXXXXX in Subjects with Mild Cognitive Impairment (MCI) Clinically at Risk for
Development of Clinically Probable Alzheimer’s Disease.”(2920)

35. “A Randomized, Double-Blind, Placebo-Controlled Evaluation of the Safety and Efficacy of
XXXXXX in Patients with Mild to Moderate Dementia of the Alzheimer’s Type. (3811)

36. “A Long-Term, Open-Label Study of XXXXXX in Subjects with Mild to Moderate Alzheimer’s

Disease." (3261x1)

37. "A Long-Term Extension Study Evaluating the Safety and Tolerability of BID and QD
Administration of XXXXXX in Patients with Mild to Moderate Dementia of the Alzheimer's
Type." (3811X1)

38. "An Eight-Week, Double-Blind, Placebo-Controlled Multicenter Study to Evaluate the Safety and Efficacy of 2 Doses of XXXXXX (1.5mg and 3.0mg) and XXXXXX in Subjects with
Major Depressive Disorder." (3495)

39. "A Randomized, Double-Blind, Placebo-Controlled Evaluation of the Safety and Efficacy of
XXXXXX in Patients with Mild to Moderate Dementia of the Alzheimer's Type."
IND#: 33,392 (4441)

40. "An Open-Label Extension Trial to Assess the Safety of XXXXXX HBr in the Treatment of
Vascular Dementia." (4556)

41. "A Double-Blind, Placebo-Controlled Dose-Finding Study Evaluating the Safety and Efficacy of XXXXXX, 80 mg bid, and 20 and 80 mg QD in the Treatment of Mild to Moderate Alzheimer's Disease. (4807)

42. A Long-term, Open-label, Flexible-dose Study of the Efficacy and Safety of XXXX in Patients with Idiopathic Restless Legs Syndrome. (4382)

43. A Randomized, Double-Blind, Placebo-Controlled Evaluation of the Safety and Efficacy of XXXX in Patients with Moderate to Severe Dementia of the Alzheimer's Type. (4887)

44. An Open-Label Extension Study to Assess the Long-Term Safety and Tolerability of XXXX in the Treatment of Mild Cognitive Impairment. (5075)

45. A Randomized, Double-Blind, Placebo-Controlled Evaluation of the Safety and Efficacy of XXXX in Patients with Moderate to Severe Dementia of the Alzheimer's Type. (4988)

46. Pharmacogenomics Blood Sampling Protocol to Obtain DNA in a Reference Population of Patients Diagnosed with Restless Legs Syndrome. (5152)

47. An 8-Week, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Flexible-Dosage Study to Evaluate the Efficacy and Safety of XXXX, at Dosages up to 16mg/day, in the Treatment of Generalized Anxiety Disorder in Adults. (202359)

48. A 12-Month, Open-Label, Flexible-Dosage Study to Evaluate the Safety of XXXX, at Dosages up to 16mg/day in Adults with Generalized Anxiety Disorder. (202976)

49. A Randomized, Multicenter, Double-Blind, Placebo-Controlled, 18-Month Study of the Efficacy of XXXX in Patients with Mild-To-Moderate Dementia of the Alzheimer's Type. (202793)

50. A Randomized, Double-Blind, Placebo-Controlled Trial of XXXX to Attenuate the Progression of Alzheimer's Disease. (202530)

51. A Randomized, Double-Blind, Placebo-Controlled, Parallel Group, Multicenter, Fixed-Dose, Polysomnographic Study of XXXX in Patients with Primary Insomnia. (202700)

52. A Multi-center, Randomized, Open-Label Study Evaluating the Effects of XXXX, 80 mg b.i.d., vs. XXXX, 5 or 10 mg, on Adrenal Function in Patients with Mild Alzheimer's Disease. (202538)

53. A One year, Multi-center, Randomized, Double-Blind, Placebo-Controlled Evaluation of the Efficacy and Safety of XXXX in Subjects with Mild Cognitive Impairement. (202493)

54. Long-Term Safety and Efficacy of Open-Label XXXX, 80 mg b.i.d. in the Treatment of Probable Alzheimer's Disease: A 6-Month Follow-Up After Completion of the Study XXXX. (202859)

55. A Randomized, Double-Blind, Placebo-Controlled Trial to Evaluate the Safety and Efficacy of XXXX 25 mg in Slowing the Progression of Alzheimer's Disease. (202531)

56. An Evaluation of the Long-Term Safety and Efficacy of XXXX in Patients with Moderate to Severe Dementia of the Alzheimer's Type. (202891)

57. A Multicenter, Randomized, Double-Blind, Placebo-Controlled, Phase II, Study of Efficacy and Safety of XXXX in Subjects with Mild to Moderate Alzheimer's Disease. (00012)

58. A Study To Define the Non-Restorative Sleep Population. (00010)

59. A Randomized, Doulbe-Blind, Placebo-Controlled Trial of XXXX to Attenuate the Progression of Alzheimer's Disease. (00026)

60. A 6-Month Safety Follow-Up Study of Select Patients Previously Enrolled and Randomized to XXXX in Studies XXXX, XXXX, or XXXX. (00028)

61. Evaluation of the Long-Term Efficacy and Safety of XXXX 12.5mg Compared to Placebo, When Both Are Administered Over a Long-Term Period "as needed", in Patients with Chronic Primary Insomnia. (A

Randomized, Double-Blind, Placebo-Controlled, Parallel Group, Multicenter, Phase IIIb Clinical Study). (00007)

62. A Randomized, Double-Blind, Placebo-Controlled, Crossover Study to Assess the Efficacy and Safety of XXXX in Patients with Restless Legs Syndrome. (203812PRN)

63. A Randomized, Double-Blind, Placebo-Controlled, 4-Period-Cross-Over Pilot Study of the Safety and Efficacy of Multiple Doses of XXXX in Subjects with Alzheimer's Disease. (202856)

64. A 12-Week, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study to Evaluate the Efficacy and Safety of XXXX (150 and 250 mg/day) as Treatment for Adults with Excessive Sleepiness Associated with Narcolepsy. (203231)

65. A 12-Week, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study to Evaluate the Efficacy and Safety of XXXX (150 and 250 mg/day) as Treatment for Adults with Residual Excessive Sleepiness Associated with Obstructive Sleep Apnea/Hypopnea Syndrome. (203233)

66. A Phase III Study of the Efficacy and Safety of XXXX in Patients with Mild to Moderate Alzheimer's Disease. (203077PRN)

67. A 12-Month, Open-Label, Flexible-Dosage (100-250 mg/day) Extension Study of the Safety and Efficacy of XXXX in the Treatment of Patients with Excessive Sleepiness Associated with Narcolepsy, Obstructive Sleep Apnea/Hypopnea Syndrome, or Chronic Shift Work Sleep Disorder. (203903)

68. A Phase IIa, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Multiple Ascending Dose, Safety, Tolerability, Pharmacokinetic, Pharmacodynamic, and Immunogenicity Trial of XXXX in Patients with Mild to Moderate Alzheimer's Disease. (00025)

69. A Randomized, Double-Blind, Placebo-Controlled Study to Assess the Safety and Efficacy of Once Daily XXXX in Patients with Restless Legs Syndrome. (00036)

70. A Double-Blind, Randomized, Placebo-Controlled, Phase IIa, Multiple Dose, Multicenter Study in Patients with Mild to Moderate Dementia of the Alzheimer's Type to Evaluate the Safety and Tolerability of Two 10-Week Dose Regimens of Orally-Administered XXXX. (00063)

71. A Phase III, Randomized, Double-Blind, Placebo-Controlled, Polysomnographic Study to Assess the Efficacy and Safety of a Modified Release Formulation of XXXX in Primary Insomnia Patients with Sleep Maintenance Difficulties. (00047)

72. A Double Blind, Placebo Controlled Study of XXXX for the Treatment of Mild-To-Moderate Alzheimer's Disease. (00065)

73. The Efficacy of XXXX 3mg as Adjunctive Therapy in Subjects with Insomnia Related to Generalized Anxiety Disorder (GAD). (00057)

74. Phase 3 Multicenter, Randomized, Double-Blind, Placebo Controlled Study of the Effect of Daily Treatment with XXXX on Measures of Cognition, Activities of Daily Living and Global Function in Subjects with Mild Dementia of the Alzheimer's Type. (00023)

75. A 12 Week, Double-Blind, Placebo Controlled, Parallel Group Study to Assess the Efficacy and Safety of XXXX (Extended Release) in Patients with Restless Legs Syndrome.

76. A Multi-Center, Randomized, Double-Blind, Placebo-Controlled, Five-Arm Parallel-Group Trial to Investigate the Efficacy and Safety of Four Different Transdermal Doses of XXXX in Subjects with Idiopathic Restless Legs Syndrome. (00046)

77. A Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study of the Efficacy, Safety and Tolerability of XXXX in Patients with Generalized Anxiety Disorder. (00054)

78. A Randomized, Double-Blind, Placebo-Controlled, Multi-Center, Parallel-Group Study to Evaluate the Efficacy and Safety of XXXX in Migraine Prophylaxis. (00082)

79. A 52-Week, Open-Label Study to Assess the Long-Term Safety of XXXX Extended Release (XR) in Patients with Restless Legs Syndrome (RLS). (00075)

80. A Randomized, Double-Blind Comparison of 5 mg of XXXX, 15 mg of XXXX, and Placebo in the Treatment of Patients with Primary Insomnia. (00076)

81. A Randomized, Double-Blind, Placebo Controlled, Cross-Over Study to Evaluate the Effects of the XXXX in Patients with Insomnia. (00077)

82. A Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Effects of XXXX on Cognitive Functions in the Elderly. (00058)

83. An Open-Label Extension Trial to Investigate the Safety and Tolerability of Long-Term Treatment with Transdermal XXXX in Subjects with Idiopathic Restless Legs Syndrome. (00093)

84. A Randomized, Double-Blind, Placebo-Controlled, Cross-Over Study to Evaluate the Effects of XXXX (2.5, 10 and 30 mg) On Polysomnographic Sleep Recordings, Subjective Sleep Assessment, and Daytime Cognitive Function in Elderly and Nonelderly Subjects with Primary Insomnia. (00088)

85. A Randomized, Double-Blind, Placebo-Controlled Study to Assess the Efficacy and Safety of XXXX in Patients with Restless Legs Syndrome. (00056)

86. An Open-Label Extension of the Phase III Study XXXX with XXXX in Patients with Alzheimer's Disease. (00107)

87. A 28-Week Open Label Extension Study Evaluating the Safety and Tolerability of XXXX in Subjects with Mild Cognitive Impairment. (00104)

88. A Randomized, Double-Blind, Placebo and Active-Controlled, Multicenter, Proof of Concept Trial of XXXX in Subjects with Non-Restorative Sleep. (00095)

89. Efficacy and Safety of XXXX 5mg/day on Sleep Maintenance Insomnia: A 6-Week, Multicenter, Randomized, Double-Blind, Placebo-Controlled Study. (00103)

90. A Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel Study to Investigate the Efficacy and Safety of a Single Oral Dose of XXXX (20, 50, and 100 mg) and Matching Placebo in Healthy Male and Female Subjects with Induced Transient Insomnia. (00107)

91. An Open-Label, 52-Week Extension Study Assessing XXXX Safety and Efficacy in Patients with Restless Legs Syndrome. (00114)

92. A 54-Week, Double-Blind, Randomized, Placebo-Controlled, Parallel-Group Study to Investigate the Effects of XXXX (extended release tablets) as Adjunctive Therapy to Acetylcholinesterase Inhibitors on Cognition and Overall Clinical Response in APOE E4-Stratified Subjects with Mild to Moderate Alzheimer's Disease. (00123)

93. A Multi-Center, Double-Blind, Parallel Group, Fixed Dose, 4-Arm, Placebo and Paroxetine Controlled 8-Week Efficacy Study of 2 Oral Doses of XXXX (175 mg or 350 mg, bid) in Adult Outpatients with Major Depressive Disorder. (00022)

94. Phase 3 Multinational, Randomized, Double-Blind, Placebo-Controlled Study of the Effect of Daily Treatment with XXXX on Measures of Cognition, Activities of Daily Living and Global Function in Subjects with Mild Dementia of the Alzheimer's Type. (00128)

95. A Randomized, Double-Blind, Placebo-Controlled, Dose-Response Study to Assess the Efficacy, Safety and Pharmacokinetics of XXXX in Patients with Restless Legs Syndrome. (00141)

96. Open Label Study of the Effect of Daily Treatment with XXXX in Subjects with Dementia of the Alzheimer's Type.

97. A Randomized, Double-Blind, Placebo-Controlled, Parallel Group Study to Demonstrate the Subjective Treatment Effects of XXXX 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. (00145)

98. An 8-Week, Randomized, Double-Blind, Fixed-Dosage, Placebo-Controlled, Parallel-Group, Multicenter Study of the Efficacy, Safety and Tolerability of XXXX 25 mg and 50 mg in the Treatment of Major Depressive Disorder (MMD) Followed by a 52-Week, Open-Label Extension (XXXX). (00125)

99. A Double-Blind, Randomized, Placebo-Controlled Study of the Efficacy, Safety and Tolerability of 8 Week Treatment of XXXX 8 mg (QHS) in Sleep Disturbed, Mild to Moderately Severe Alzheimer's Disease Subjects. (00090)

100. Effect of XXXX in Slowing the Progression of Alzheimer's Disease. (00134)

101. A Phase 2, Double-Blind, Placebo-Controlled Study to Evaluate the Safety, Tolerability, and Effect on Cognitive Function of XXXX After 12 Weeks of Intranasal Administration in Subjects with Mild Cognitive Impairment. (00163)

102. A Six-Week, Double-Blind, Randomized, Placebo-Controlled, Parallel Group, Efficacy and Safety, Sleep Lab Trial with XXXX in Patients with Chronic Primary Insomnia. (00118)

103. A Randomized, Double-Blind, Placebo-Controlled, Parallel, Proof of Concept Study to Evaluate the Effectiveness of XXXX To Advance the Timing of Sleep in Individuals with Delayed Sleep Phase Syndrome (DSPS). (00198)

104. A Randomized, Double-Blind, Placebo-Controlled, Phase IIa Study to Assess the Short-Term Effects of XXXX Alone and in Combination with Donepezil in Subjects with Mild Alzheimer's Disease. (00144)

105. Efficacy and Safety of 2 mg/Day XXXX on Sleep Maintenance Insomnia: A 6-Week, Multicenter, Randomized, Double-Blind, Placebo-Controlled Polysomnographic Study. (00192)

106. A Phase II, Double-Blind, Randomized, Placebo-Controlled, Parallel-Group, Dose-Ranging Study Assessing the Efficacy and Safety of XXXX Tablets Twice Daily in Adults with Mild to Moderate Alzheimer's Disease. (00214)

107. Fifty-Two Weeks, Open-Label Extension Trial To Evaluate Safety and Efficacy of XXXX in Outpatients with Chronic Primary Insomnia Who Completed Clinical Trial Protocol 176001 or 176002. (00217)

108. A Randomized, Double-Blind, Active- and Placebo-Controlled, Parallel Group Safety Study Assessing Simulated Driving Performance in XXXX – Treated Patients with Restless Legs Syndrome. (00227)

109. XXXX Dose-Ranging Trial: A Randomized, Double-Blind, Placebo-Controlled, 5-Way Crossover, Multicenter Polysomnography Trial of XXXX in Adults with Primary Insomnia. (00284)

110. A 52-Week Open-Label Extension Study of the Long-Term Safety and Efficacy of XXXX Extended-Release (RSGXR) as Adjunctive Therapy to Acetylcholinesterase Inhibitors in Subjects with Mild-to-Moderate Alzheimer's Disease (AVA102675, REFLECT-4). (00220)

111. A Phase 3, Randomized, Double-Blind, Placebo Controlled, Parallel Group, 10-Week Study Evaluating the Efficacy and Safety of XXXX for the Treatment of Generalized Anxiety Disorder. (00275)

112. A Double-Blind, Randomized, Placebo-Controlled Study Examining, the Safety, Efficacy, and Tolerability of XXXX in Subjects with Major Depressive Disorder (including Atypical and Melancholic Features). (00230)

113. A Randomized, Long-Term Safety Study of XXXX in Elderly Outpatients with Chronic Primary Insomnia Examining the Effects of 1.5 mg or 3.0 mg of XXXX. (00226)

114. A Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Phase 3 Study of XXXX in Adult Migraineurs for a Single Migraine Followed by Open-Label Extensions to 26/52 Weeks. (00219)

115. A Randomized, Double-Blind, Placebo-Controlled Subjective Study to Assess the Efficacy of XXXX in Patients with Primary Insomnia Characterized by Difficulty Maintaining Sleep. (00324)

116. A Randomized, Double-Blind, Placebo-Controlled, Parallel-Group Study to Evaluate the Efficacy and Safety of XXXX at a Target Dosage of 200 mg/Day as Treatment for Adults with Excessive Sleepiness Associated with Obstructive Sleep Apnea/Hypopnea Syndrome With Comorbid Major Depressive Disorder or Dysthymic Disorder. (00323)

117. XXXX Dose-Ranging Trial: A Randomized, Double-Blind, Parallel Group, Placebo-Controlled, Multicenter Outpatient Trial of XXXX in Adults with Non-Restorative Sleep. (00321)

118. A Phase 2a Multi-Center, Randomized, Double Blind, Placebo-Controlled Study to Investigate the Efficacy and Safety of XXXX in Patients with Mild to Moderate Alzheimer's Disease. (00318)

119. A Double-Blind, Placebo-Controlled, Randomized, Multicenter Study Evaluating the Efficacy and Safety of Eighteen Months of Treatment with XXXX (XXXX) in Participants with Mild-To-Moderate Alzheimer's Disease. (00355)

120. A Polysomnography Study of XXXX (XXXX) Extended Release Tablets Versus Placebo in the Treatment of Restless Legs Syndrome (RLS) and Associated Sleep Disturbance. (00338)

121. Effect of y-Secretase Inhibition on the Progression of Alzheimer's Disease: XXXX Versus Placebo. (00294)

122. A Randomized, Double-Blind, Placebo-Controlled, Ascending-Dose Phase 1b Safety Study of Three Different Doses of an Alpha-7 Nicotinic Acetylcholine Receptor Agonist (XXXX) or Placebo in Patients with Mild to Moderate Probable Alzheimer's Disease. (00343)

123. A 52 Week Open Label Safety Study of XXXX in Subjects with Generalized Anxiety Disorder. (00305)

124. A Phase 2 Study of XXXX Compared with Placebo for the Treatment of Alcohol Dependence. (00347)

125. Efficacy and Safety of XXXX 5mg/day in Insomnia Characterized by Sleep Maintenance Difficulties: A 6-Week, Randomized, Double-Blind, Placebo-Controlled, Polysomnography Study. (00350)

126. A Phase 3 Multicenter, Randomized, Placebo-Controlled, Twelve-Month Safety and Efficacy Study Evaluating XXXX in Patients with Mild-to Moderate Alzheimer's Disease on Donepezil. (00404)

127. A Double-Blind Placebo-Controlled Preliminary Study of the Efficacy, Safety and Tolerability of XXXX in the Treatment of Alzheimer's Disease. (00382)

128. Randomized, Double-Blind, 12-Month Study of XXXX in Subjects with Restless Legs Syndrome. (00418)

129. A Multicenter, Randomized, Double-Blind, Placebo-Controlled Study of the Safety, Tolerability, Pharmacodynamic and Pharmacokinetic Effects of XXXX in the Treatment of Patients with Mild to Moderate Alzheimer's Disease. (00341)

130. A Multicenter, Randomized, Double-Blind, Placebo-Controlled Study of the Safety, Tolerability, Pharmacodynamic and Pharmacokinetic Effects of XXXX in the Treatment of Patients with Prodromal Alzheimer's Disease. (00412)

131. A Double-Blind Placebo-Controlled Preliminary Study of the Efficacy, Safety and Tolerability of XXXX in the Treatment of Alzheimer's Disease in Subjects Concurrently Receiving Donepezil (Aricept®). (00403)

132. An Open Label Study of the Safety and Tolerability of XXXX in Subjects with Alzheimer's Disease. (00446)

133. A Randomized, Double-Blind, Placebo-Controlled, Functional Neuroimaging Study of the Effects of XXXX (200 mg/day) Treatment on Prefrontal Cortical Activation in Patients with Residual Excessive Sleepiness Associated with Obstructive Sleep Apnea/Hypopnea Syndrome. (00407)

134. A 12-Week, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Fixed-Dosage Study to Evaluate the Efficacy and Safety of XXXX (50, 150 and 250 mg/day) as Treatment for Patients with Excessive Sleepiness Associated with Mild or Moderate Closed Traumatic Brain Injury. (00397)

135. A Phase 2 Multicenter, Double Blind, Placebo Controlled, Parallel Group Study of XXXX in Subjects with Mild to Moderate Alzheimer's Disease. (00440)

136. A Randomized, Double-Blind, Placebo-Controlled, 3-Way Crossover, Multicenter Polysomnography Study of XXXX and XXXX in Adults with Restless Legs Syndrome. (00457)

137. A Phase III, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Long-Term Safety Study of XXXX in Patients with Primary Insomnia. (00448)

138. Open-Label Extension for Alzheimer's Disease Patients Who Complete One of Two XXXX Phase 3 Double-Blind Studies (H6l-MC-LFAN or H6L-MC-LFBC). (00294x1)

139. A 12-Month, Open-Label Study to Evaluate the Safety, Tolerability, and Efficacy of XXXX (150 and 250 mg/Day) as Treatment for Patients with Excessive Sleepiness Associated with Mild or Moderate Closed Traumatic Brain Injury. (00506)

140. Randomized, Double-Blind, 12-Week Study of XXXX in Subjects with Restless Legs Syndrome. (00515)

141. A Worldwide, Randomized, Double Blind, Placebo Controlled, Parallel Group Clinical Trial to Evaluate the Safety and Efficacy of XXXX for the Acute Treatment of Migraine in Children and Adolescents. (00502)

142. A Worldwide, Open Label, Clinical Trial to Examine the Long Term Safety and Tolerability of XXXX in Pediatric Migraineurs for the Treatment of Migraine With or Without Aura. (00469)

143. Effect of Passive Immunization on the Progression of Alzheimer's Disease: XXXX (XXXX) Versus Placebo. (00503)

144. A Phase III, Multicenter, Randomized, Double-Blind, Placebo-Controlled, Parallel Group, Study to Evaluate the Safety and Efficacy of XXXX in Patients with Primary Insomnia – Study A. (00514)

145. An Open-Label Extension of the CONCERT Protocol (DIM18) Evaluating XXXX (XXXX) in Patients with Alzheimer's Disease. (00529)

146. A Multiple Dose Study of the Safety, Tolerability, Pharmacokinetic and Pharmacodynamic Effects of XXXX Administered Once Daily to Subjects with an Excessive Daytime Sleepiness Disorder. (00541)

147. Continued Efficacy and Safety Monitoring of XXXX, an Anti-Amyloid β Antibody in Patients with Alzheimer's Disease. (00565)

148. A Randomized, Double-Blind, Placebo-Controlled, Combined Single Ascending Dose and Multiple Ascending Dose Study to Assess Safety, Tolerability, Immunogenicity, Pharmacodynamic Response, and Pharmacokinetics of Intravenous Infusions of XXXX in Subjects with Mild to Moderate Alzheimer's Disease. (00582)

149. A Phase 2, 24-Month, Multi-Center, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Amyloid-Imaging Positron Emission Tomography (PET) and Safety Study of XXXX and XXXX Adjuvant in Subjects with Mild to Moderate Alzheimer's Disease. (00591)

150. A 4-Week, Randomized, Double-Blind, Parallel Group, Placebo Controlled Study to Investigate the Safety and Efficacy of XXXX as Monotherapy in Patients with Treatment-Resistant Major Depression. (00585)

151. A Multicenter, Randomized, Double-Blind, Active-Controlled Study of the Efficacy and Safety of Flexibly-Dosed XXXX in Patients with Treatment Resistant Major Depression. (00571)

152. A Phase 2, Multicenter, Open-Label Study to Assess the Safety and Tolerability of Oral XXXX as Adjunctive Therapy in Adult Patients with Major Depressive Disorder. (00416)

153. A Randomized, Double-Blind, Placebo-Controlled Efficacy and Safety Study of XXXX in Subjects with Spasticity Due to Multiple Sclerosis. (00595)
154. An Open Label, 26-Week Study Assessing XXXX Safety and Efficacy in Subjects with Spasticity Associated with Multiple Sclerosis. (00614)

155. A Phase 1b, Multicenter, Double-Blind, Randomized, Placebo-Controlled, Sequential Dose Study of the Safety and Tolerability of XXXX in Subjects with Alzheimer's Disease on Donepezil. (00618)

156. A Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Multicenter, Phase II Study to Evaluate the Efficacy and Safety of XXXX in Patients with Mild to Moderate Alzheimer's Disease. (00579)

157. A Multicenter, Double-Blind, 58-Week Rollover Study to Assess the Safety and Tolerability of XXXX in Patients with Treatment Resistant Major Depression. (00583)

158. A Study to Evaluate the Effects of XXXX in Patients with Chronic Obstructive Pulmonary Disease. (00619)

159. A Double-Blind, Parallel Group, Randomized, Placebo Controlled Sleep Laboratory Study of Efficacy and Safety of XXXX in Insomnia Patients Aged 18-80. (00635)

160. A Phase 3 Randomized, Double-Blind, Placebo-Controlled Study of the Safety and Effectiveness of XXXX (Human), 10% Solution (XXXX, 10%) for the Treatment of Mild to Moderate Alzheimer's Disease (AD). (00626)

161. A Phase IIa Safety and Tolerability Study to Investigate the Effect on Sleep of 3 Doses of XXXX and Placebo in Patients with Mild Alzheimer's Disease and Mild Cognitive Impairment During 4 Weeks of Treatment, Designed as a Randomized, Double-Blind, Multi-Center, Parallel Group, Placebo-Controlled Study. (00658)

162. Effect of XXXX in Obese Subjects with Moderate or Severe Obstructive Sleep Apnoea A 32-Week Randomized, Double-Blind, Placebo-Controlled, Parallel Group, Multi-Center and Multinational Trial. (00696)

163. A Randomized, Double-Blinded, Placebo-Controlled Multiple Dose Study to Assess the Safety, Tolerability, Pharmacokinetics and Pharmacodynamics of XXXX in Subjects with Prodromal or Mild Alzheimer's Disease. (00700)

164. A Phase 2, Randomized, Double-Dummy, Double-Blind, Placebo-Controlled Study to Assess the Efficacy, Safety and Tolerability of XXXX (XXXX/XXXX) for the Treatment of Symptoms of Agitation in Patients with Alzheimer's Disease. (00697)

165. A Multicenter, Open-Label, Long-Term Safety Extension of Phase II Studies ABE4869g and ABE4955g in Patients with Mild to Moderate Alzheimer's Disease. (00729)

166. Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, 18-Month Safety and Efficacy Study of XXXX in Subjects with Mild Alzheimer's Disease. (00722)

167. A Twelve-Week, Double-Blind, Placebo-Controlled, Randomized, Parallel-Group, Multi-Center Study of the Safety and Efficacy of XXXX in the Treatment of Excessive Daytime Sleepiness in Subjects with Narcolepsy. (00733)

168. A 26-Week, Double-Blind, Randomized, Placebo-Controlled, Parallel-Group Study to Investigate the Effects of Daily Administration of XXXX in Participants with Mild to Moderate Alzheimer's Disease. (00731)

169. Effect of Passive Immunization on the Progression of Mild Alzheimer's Disease: XXXX (XXXX) Versus Placebo. (00743)