

DANIEL RUTRICK, M.D.

MEDICAL DIRECTOR AND PRINCIPAL INVESTIGATOR AT ADAMS CLINICAL TRIALS
BOARD CERTIFIED IN PSYCHIATRY
521 MOUNT AUBURN STREET, SUITES 107, 109, 209 • WATERTOWN, MASSACHUSETTS 02472
T: (617)744-8542 • F: (617)744-8544 • DRUTRICK@ADAMSCLINICAL.COM

EDUCATION/SCHOLARSHIP

HARVARD MEDICAL SCHOOL, Cambridge, MA

Resident in Psychiatry, June 1975

Chief Resident: Clinical Fellow of Harvard Medical School; Assistant Clinical Director of In-Patient Psychiatry, Mount Auburn Hospital, Cambridge, MA.

SUNY UPSTATE MEDICAL UNIVERSITY, Syracuse, NY

Resident in Psychiatry, May 1974

Doctor of Medicine, May 1971

CITY COLLEGE OF NEW YORK, New York, NY

Bachelor of Science, May 1967

BRONX HIGH SCHOOL OF SCIENCE, Bronx, NY

PSYCHIATRIC CLINICAL RESEARCH EXPERIENCE

ADAMS CLINICAL TRIALS, Watertown, MA

December 2010 - Present

Medical Director: Lead research team at dedicated psychiatric clinical research site with a focus on testing treatments for mood disorders, eating disorders, and substance use disorders for all age groups in Phase II and III trials. Staff consists of eight full-time study coordinators, five physicians, and two psychologists.

Principal Investigator on the 38 clinical trials listed below:

NCT03107026

2017

A Study to Evaluate a Drug (Dasotraline) on the Safety, Effectiveness and How Well the Body Tolerates it, in Adults With Moderate to Severe Binge Eating Disorder

NCT02951988

2017

A Study of Rapastinel as Adjunctive Therapy in the Prevention of Relapse in Patients With Major Depressive Disorder

NCT03002077

2017

Long-term Safety Study of Rapastinel as Adjunctive Therapy in Patients With Major Depressive Disorder

NCT02431806

2017

Safety and Efficacy of Levomilnacipran ER in Adolescent Patients With Major Depressive Disorder

NCT02932943

2016

A Study of Rapastinel as Adjunctive Therapy in Major Depressive Disorder

NCT02497287

2016

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Chief Resident: Clinical Fellow of Harvard Medical School; Assistant Clinical Director of In-Patient Psychiatry, Mount Auburn Hospital, Cambridge, MA.

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NCT02436239	2016
Safety of Vilazodone in Pediatric Patients with Major Depressive Disorder	
NCT01878292	2016
Safety and Efficacy of Vilazodone in Pediatric Patients with Major Depressive Disorder	
NCT02684279	2016
An Open-label, Flexibly-dosed, Multicenter, Extension Study of Dasotraline to Evaluate Long-term Safety and Tolerability in Adults with Binge-eating Disorder	
NCT02564588	2016
A 12-week, Randomized, Double-blind, Parallel-group, Placebo Controlled, Flexibly Dosed, Multicenter Study to Evaluate the Efficacy, Safety, and Tolerability of Dasotraline in Adults with Moderate to Severe Binge Eating Disorder	
NCT02510014	2015
Safety and Tolerability Study of Depot Buprenorphine in Treatment Seeking Subjects With Opioid Use Disorder	
NCT02357901	2015
A Phase 3, A Randomized, Double-Blind, Placebo-Controlled, Multicenter Study To Assess the Efficacy, Safety, and Tolerability of Multiple Subcutaneous Injections of Depot Buprenorphine (RBP-6000) Over 24 Weeks in Treatment-Seeking Subjects with Opioid Use Disorder	
NCT01998633	2015
A Phase 3, Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel-group Trial with an Open-label Extension Phase to Evaluate the Efficacy and Safety of Subcutaneously Administered Bremelanotide in Premenopausal Women with Hypoactive Sexual Desire Disorder (HSDD) (with or without Decreased Arousal)	
NCT01878292	2015
A Phase 3 Multicenter, Double-blind, Placebo- and Active-Controlled Parallel-Group Evaluation of the Safety and Efficacy of Vilazodone in Pediatric Patients With Major Depressive Disorder	
NCT02288325	2014
A Phase 4 Multicenter, Randomized, Double-blind, Placebo-Controlled, Relapse Prevention Study With Levomilnacipran ER in Patients With Major Depressive Disorder	
NCT02141399	2014
A Phase 3 Multicenter Study of the Long-Term Safety and Tolerability of ALKS 5461 for the Adjunctive Treatment of Depression in Pediatric Patients with Major Depressive Disorder	
NCT01878292	2016
Safety and Efficacy of Vilazodone in Pediatric Patients with Major Depressive Disorder	
NCT02684279	2016
An Open-label, Flexibly-dosed, Multicenter, Extension Study of Dasotraline to Evaluate Long-term Safety and Tolerability in Adults with Binge-eating Disorder	
NCT02564588	2016
A 12-week, Randomized, Double-blind, Parallel-group, Placebo Controlled, Flexibly Dosed, Multicenter Study to Evaluate the Efficacy, Safety, and Tolerability of Dasotraline in Adults with Moderate to Severe Binge Eating Disorder	

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NCT02009163	2014
A Phase 3, Multicenter, Double-blind, Placebo-controlled, Randomized-withdrawal Study to Evaluate the Maintenance of Efficacy of SPD489 in Adults Aged 18-55 Years With Moderate to Severe Binge Eating Disorder.	
NCT01312909	2013
A Twelve-Week, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Dose-Ranging Study With Follow-Up Evaluating The Safety and Efficacy Of Varenicline For Smoking Cessation in Healthy Adolescent Smokers	
NCT01944969	2013
Interventional, Open-label, Long-term Extension Study to Evaluate the Safety and Tolerability of Brexpiprazole as Adjunctive Treatment in Patients with Major Depressive Disorder	
NCT01837797	2013
Interventional, Randomized, Double-blind, Parallel-group, Placebo-controlled, Fixed-dose Study to Evaluate the Efficacy and Safety of Brexpiprazole (1 and 3mg/Day) as Adjunctive Treatment in Elderly Patients with Major Depressive Disorder With an Inadequate Response to Antidepressant Treatment	
NCT01813019	2013
A Randomized, Double-blind, Placebo-controlled, Parallel-group Proof of Concept Study to Evaluate the Effect of AFQ056 in Obsessive Compulsive Disorder (OCD) Patients Resistant to Selective Serotonin Reuptake Inhibitor (SSRI) Therapy	
NCT01878292	2013
A Double-blind, Placebo-Controlled Evaluation of the Safety and Efficacy of Vilazodone in Adolescent Patients with Major Depressive Disorder.	
NCT01457677	2013
A Multi Center, Randomized, Double-blind, Placebo Controlled, Parallel-group Study to Investigate the Efficacy and Safety of RO4995819 Versus Placebo, as Adjunctive Therapy in Patients With Major Depressive Disorder Having Inadequate Response to Ongoing Antidepressant Treatment.	
NCT01360866	2013
A Long-term, Phase 3, Multicenter, Open-label Trial to Evaluate the Safety and Tolerability of Oral OPC-34712 as Adjunctive Therapy in Adults With Major Depressive Disorder, the Orion Trial.	
NCT01727726	2013
Interventional, Multicenter, Double-blind, Placebo-controlled, Randomized-withdrawal Study to Evaluate the Maintenance of Efficacy of SPD489 in Adults Aged 18-55 Years With Moderate to Severe Binge Eating Disorder.	
NCT01312909	2013
A Twelve-Week, Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, Dose-Ranging Study With Follow-Up Evaluating The Safety and Efficacy Of Varenicline For Smoking Cessation in Healthy Adolescent Smokers	
NCT01944969	2013
Interventional, Open-label, Long-term Extension Study to Evaluate the Safety and Tolerability of Brexpiprazole as Adjunctive Treatment in Patients with Major Depressive Disorder	
NCT01837797	2013

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NCT01657019

2012

A Phase 3, Multicenter, Open-Label, 12-Month Extension Safety and Tolerability Study of SPD489 in the Treatment of Adults with Binge Eating Disorder.

NCT01718509

2012

The SPD489-344, Phase 3, Multicenter, Randomized, Double-blind, Parallel-group, Placebo-controlled, Dose-optimization Study to Evaluate the Efficacy, Safety, and Tolerability of SPD489 in Adults Aged 18-55 Years With Moderate to Severe Binge Eating Disorder.

NCT01436175

2011

A Phase 3, Open-label, Multicenter, 12-month Extension Safety and Tolerability Study of SPD489 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.

NCT01436149

2011

The SPD489-322 Phase 3, Multicenter, Randomized, Double-blind, Parallel-group, Placebo-controlled, Flexible Dose Titration, Efficacy and Safety Study of SPD489 in Combination With an Antidepressant in the Treatment of Adults With Major Depressive Disorder With Inadequate Response to Prospective Treatment With an Antidepressant.

NCT01110902

2011

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

NCT01395147

2011

A Phase 3, Long-Term, Open-Label, Flexible-Dose, Extension Study Evaluating the Safety and Tolerability of Lu AA21004 (15 and 20 mg) in Subjects With Major Depressive Disorder.

NCT01179516

2011

A Phase 3, Randomized, Double-Blind, Parallel-Group, Placebo-Controlled, Duloxetine-Referenced, Fixed-Dose Study Comparing the Efficacy and Safety of 2 Doses (10 and 15 mg) of Lu AA21004 in Acute Treatment of Adults With Major Depressive Disorder.

NCT01121536

2011

A 6-Month, Open-Label, Flexible-Dosage (150 to 200 mg/Day) Extension Study of the Safety and Efficacy of Armodafinil Treatment as Adjunctive Therapy in Adults With Major Depression Associated with Bipolar I Disorder, Multicenter, Open-Label, 12-Month Extension Safety and Tolerability Study of SPD489 in the Treatment of Adults with Binge Eating Disorder.

NCT01718509

2012

The SPD489-344, Phase 3, Multicenter, Randomized, Double-blind, Parallel-group, Placebo-controlled, Dose-optimization Study to Evaluate the Efficacy, Safety, and Tolerability of SPD489 in Adults Aged 18-55 Years With Moderate to Severe Binge Eating Disorder.

NCT01436175

2011

A Phase 3, Open-label, Multicenter, 12-month Extension Safety and Tolerability Study of SPD489 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.

NCT01436149

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NCT01056289

2010

A Randomized, Double-Blind, Parallel Group Study To Compare Discontinuation Symptoms In Abrupt Discontinuation Versus A 1-Week Tapering Regimen In Subjects With MDD Treated For 24 Weeks With Open-Label 50 mg DVS SR Formulation.

NCT00887224

2010

A Multicenter, Double-Blind, Placebo-Controlled, Randomized Withdrawal, Parallel Group Study To Evaluate The Efficacy And Safety Of 50 mg/Day Of DVS SR In Adult Outpatients With Major Depressive Disorder

ATLANTIC CLINICAL RESEARCH, LLC, Watertown, MA

2004 - 2010

Principal Investigator on the 4 clinical trials listed below:

NCT00824291

2009

A Multicenter, Double-Blind, Randomized, Placebo-Controlled Study To Evaluate Functional Outcome In Outpatients With Major Depressive Disorder Treated With Desvenlafaxine Succinate Sustained Release.

NCT00483548

2007

A Six-Week, Double-Blind, Multicenter, Placebo Controlled Study Evaluating The Efficacy And Safety Of Flexible Doses Of Oral Ziprasidone As Add-On, Adjunctive Therapy With Lithium, Valproate Or Lamotrigine In Bipolar I Depression.

NCT00235508

2005

The Efficacy of Eszopiclone 3 mg as Adjunctive Therapy in Subjects With Insomnia Related to Generalized Anxiety Disorder.

NCT00368030

2004

Depression Response to Eszopiclone in Adults With Major Depressive Disorder (DREAMDD): A Randomized, Double-Blind, Placebo-Controlled, Parallel-Group, 8-Week, Safety & Efficacy Study of Eszopiclone 3 mg Compared to Placebo in Subjects With Insomnia Related to MDD Acronym: DREAMDD.

OTHER CLINICAL RESEARCH EXPERIENCE

ATLANTIC CLINICAL RESEARCH, LLC, Watertown, MA

Sub-Investigator on the 8 clinical trials listed below:

NCT01156415

2011

A 52-week, Multi-center, Open-label Study of the Safety and Tolerability of Agomelatine Sublingual Tablets in Patients With Major Depressive Disorder (MDD).

~~NCT01056289, Double-Blind, Parallel Group Study To Compare Discontinuation Symptoms In Abrupt Discontinuation Versus A 1-Week Tapering Regimen In Subjects With MDD Treated For 24 Weeks With Open-Label 50 mg DVS SR Formulation.~~

NCT00887224

2010

A Multicenter, Double-Blind, Placebo-Controlled, Randomized Withdrawal, Parallel Group Study To Evaluate The Efficacy And Safety Of 50 mg/Day Of DVS SR In Adult Outpatients With Major Depressive Disorder

ATLANTIC CLINICAL RESEARCH, LLC, Watertown, MA

2004 - 2010

Principal Investigator on the 4 clinical trials listed below:

NCT00824291

2009

A Multicenter, Double-Blind, Randomized, Placebo-Controlled Study To Evaluate Functional Outcome In Outpatients With Major Depressive Disorder Treated With Desvenlafaxine Succinate Sustained Release

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NCT01350141

2010

A Phase 2, Double-Blind, Placebo-Controlled, Randomized Study To Assess The Efficacy, Safety, And Tolerability Of PF-04950615 (RN316) Following Multiple Intravenous Doses In Hypercholesterolemic Subjects On High Doses Of Atorvastatin, Rosuvastatin Or Simvastatin

NCT01289990

2010

A Phase III Double-blind, Extension, Placebo-controlled Parallel Group Safety and Efficacy Trial of BI 10773 (10 and 25mg Once Daily) and Sitagliptin (100mg Once Daily) Given for Minimum 76 Weeks (Incl. 24 Weeks of Preceding Trial) as Monotherapy or With Different Back-ground Therapies in Patients With Type 2 Diabetes Mellitus Previously Completing Trial 1245.19, 1245.20 or 1245.23

NCT01131676

2010

BI 10773 add-on to Usual Care Compared With Usual Care Alone in Patients With Type 2 Diabetes Mellitus at High Cardiovascular Risk.

NCT01194830

2009

A Phase IIIB, 24-week, randomized, placebo-controlled, double-blinded, efficacy and safety study of linagliptin (BI 1356) in Black/African American patients with type 2 diabetes with a MTT sub-study.

NCT01042769

2009

A Safety and Efficacy Study to Evaluate the Potential of Aleglitazar to Reduce Cardiovascular Risk in Coronary Heart Disease (CHD) Patients With a Recent Acute Coronary Syndrome (ACS) Event and Type 2 Diabetes Mellitus (T2D).

NCT00788944

2008

An Open-Label Study To Evaluate The Prevalence Of Phenotypic Poor Metabolizers At CYP2D6 Among Venlafaxine-Treated Outpatients With Depression.

PROFESSIONAL CLINICAL EXPERIENCE

PRIVATE PSYCHIATRIC PRACTICE, Everett, MA, and Watertown, MA

July 1975 - Present

Opioid Dependency Treatment: Prescribe Suboxone for the treatment of 80 patients suffering from opioid dependency with a special waiver from the Drug Enforcement Administration under the Drug Addiction Treatment Act of 2000. Treatment consists of a comprehensive out-patient program with both individual and group therapy. Medication management is closely monitored and combined with frequent urine toxicology screens. (January 2006 - present).

Everett Chronic Pain Program: Direct out-patient program for chronic intractable pain syndrome patients.

Everett Chronic Pain Program: Direct out-patient program for chronic intractable pain syndrome patients. Focus on the use of non-pharmaceutical, non-invasive modalities such as acupuncture, yoga, and Tai Chi. Tolerability Of PF-04950615 (RN316) Following Multiple Intravenous Doses In Hypercholesterolemic Subjects On High Doses Of Atorvastatin, Rosuvastatin Or Simvastatin

NCT01289990

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practitioners. Manage group therapies for disability; grief; mid-life crisis; panic disorder; chronic pain; and social anxiety disorder. (July 1975 - present).

BOSTON PAIN CENTER, SPAULDING REHABILITATION HOSPITAL, Boston, MA 1976 - 1988

Consultant: Advised multi-disciplinary in-patient treatment program focusing on chronic intractable low back pain and headaches. Program consisted of behavioral and dynamic psychotherapy in association with physical therapy, and required psychopharmacology, milieu management, and crisis intervention when patients suffered from depressive and other psychiatrically diagnosable reactions to chronic pain.

MOUNT AUBURN HOSPITAL, Watertown, MA September 1974 - June 1975

Assistant Director of In-Patient Psychiatry: Hired staff, developed orientation program, determined admissions, and managed a sixteen bed voluntary, adult, in-patient unit. Diagnosed and managed treatment for patients suffering from bipolar disorder, major depression with suicidal behavior, personality disorders, and psychotic states. Treatment consisted individual, group, and psychopharmacological therapy.

HOSPITAL AFFILIATIONS

LAWRENCE MEMORIAL HOSPITAL OF MEDFORD, Medford, MA	2001 - 2014
MELROSE-WAKEFIELD HOSPITAL, Melrose, MA	2001 - 2014
ST. ELIZABETH'S MEDICAL CENTER, Brighton, MA	1976 - Present
MOUNT AUBURN HOSPITAL, Cambridge, MA	1975 - Present

PUBLICATIONS

Combined Psychotherapy of Chronic Pain Patients on a Pain Unit. *Hospital Practice*. September, 1983.
Psychiatrist's View of Chronic Pain. *International Anesthesiology Clinics*. Volume 21, No. 4, Winter, 1983.
Psychodynamic Psychotherapy of Chronic Pain Patient. With Gerald Aronoff, M.D., 1985.
Mavoglurant in Obsessive-Compulsive Disorder Resistant to Selective Serotonin Reuptake Inhibitor: A Proof-of-Concept, Randomized, Placebo-Controlled, Phase 2 Study. *Advances in Therapy*. With Dan. J Stein, FRCPC, PhD, Ganesan Subramanian, PhD, Brian Smith, PhD, Maurizio Fava, MD, Gregor Hasler, MD, Jang-Ho Cha, MD, PhD, Toni Donchev, MD, PhD, Magdalena Ocwieja, PhD, and Baltazar GomezMancilla, M.D., Ph.D., June, 2016.

PROFESSIONAL MEMBERSHIPS

American Psychiatric Association; Massachusetts Psychiatric Association; American Society of Clinical Psychopharmacology; Association of Clinical Research Professionals.

BOSTON PAIN CENTER, SPAULDING REHABILITATION HOSPITAL, Boston, MA 1976 - 1988

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