

RAYMOND GEORGE BOOTH

CURRICULUM VITAE

RESEARCH SYNOPSIS:

Drug Discovery and Development for Neuropsychiatric Disorders

The laboratory is on track to an Investigational New Drug (IND) Application regarding novel drug(s) that target brain serotonin receptors for treatment of neuropsychiatric disorders, including, autism spectrum disorder, obsessive-compulsive disorders (binge eating, attention deficit-hyperactivity disorder [ADHD], addiction) and diseases involving cognition impairment (schizophrenia, dementias, psychoses). Development will proceed under the auspices of the National Institutes of Health (NIH) and Boston-based pharmaceutical industry partners. In addition to structure-based design and synthesis of new chemical entities, laboratory drug discovery and development technology includes computational chemistry and molecular modeling, molecular neuropharmacology, pharmacokinetics, and preclinical in vivo behavioral methodologies for development of drug candidates to treat neuropsychiatric disorders.



EDUCATION

1988-90 NIH Postdoctoral Fellow (Psychiatry and Neuroscience)
HARVARD MEDICAL SCHOOL and MCLEAN HOSPITAL
Boston, Massachusetts

1983-88 Ph.D. (Medicinal/Pharmaceutical Chemistry)
UNIVERSITY OF CALIFORNIA at SAN FRANCISCO
San Francisco, California

1978-83 B.S. (Pharmacy)
NORTHEASTERN UNIVERSITY
Boston, Massachusetts

PROFESSIONAL EXPERIENCE

2012- Professor, Department of Pharmaceutical Sciences (interim Chair, 2017-current), Center for Drug Discovery (Associate Director), Department of Chemistry & Chemical Biology, Northeastern University, Boston.

2007-2013 Professor, Department of Medicinal Chemistry (College of Pharmacy), and, Department of Pharmacology & Therapeutics (College of Medicine), University of Florida
Current: courtesy appointment in the Department of Medicinal Chemistry

2005-2007 Associate Professor, Department of Medicinal Chemistry, College of Pharmacy, University of Florida

1997-2005 Associate Professor, Division of Medicinal Chemistry (School of Pharmacy), and, Division of Toxicology (School of Medicine), University of North Carolina at Chapel Hill

1990-1997 Assistant Professor, Division of Medicinal Chemistry, School of Pharmacy, University of North Carolina at Chapel Hill

RESEARCH

Active funding

Source: Department of Defense Congressionally Directed Medical Research Programs Peer Reviewed Medical Research Program
 Number: PR160095
 Project Name: *Development of Novel Drugs Targeting Serotonin Receptors to Treat Motor, Social, Cognitive, and Sensory Domains of Autism Spectrum Disorder Using Mouse Models*
 Role: MPI with Dr. Clint Canal
 Funding Period: 09/01/2017-08/31/2019
 Total costs: \$591,890

Source: Department of Defense Congressionally Directed Medical Research Programs Peer Reviewed Medical Research Program
 Number: PR141869
 Project Name: *Translation of Novel Serotonin 5-HT₇ Agonist Drug Candidates in Rodent Models of Fragile X Syndrome*
 Role: MPI with Dr. Clint Canal
 Funding Period: 09/01/2015-08/31/2017
 Total Costs: \$309,000

Source: NIH (National Institute on Drug Abuse, NIDA)
 Number: RO1 DA030989
 Project Name: *Functionally-Selective Serotonin 5HT₂ Drugs for Amphetamines Abuse/Disorders*
 Role: MPI (contact)
 Funding Period: 09/01/12-08/31/17
 Total Costs: \$1,802,016

Completed funding

Source: NIH (National Institute on Drug Abuse, NIDA)
 Number: T32 DA07312
 Project Name: Training Program in Medications Development for Drugs of Abuse
 Role: co-Principal Investigator (Makriyannis, PI)
 Funding Period: 07/01/1999 - 06/30/2014
 Total Costs: \$2,396,693.00

Source: NIH (National Institute on Mental Health, NIMH)
 Number: R01MH081193
 Project Name: *Serotonin 5HT_{2C} Agonist Ligands with 5HT_{2A/B} Antagonist Activity*
 Role: Principal Investigator
 Funding Period: 04/01/08-02/28/14
 Total Costs: \$1,841,323

Source: NIH (Rapid Access to Interventional Development, RAID)
 Number: R01MH081193-W1
 Project Name: *Novel Anxiolytic Agents Targeting Serotonin 5HT_{2A/2C} Receptors*
 Role: Principal Investigator
 Funding Period: 10/01/09-02/28/14
 Total Costs: \$73,250

Source: NIH (NIDA)
 Number: R01DA023928
 Project Name: *Novel 5HT2C Agonist Drugs with 5HT2A Antagonist Activity for Cocaine Addiction*
 Role: Principal Investigator
 Funding Period: 09/15/07-09/14/12
 Total Costs: \$1,475,130

Source: NIH (NIDA)
 Number: R01DA023928-03S109
 Project Name: *International Collaborative Research to Develop Cocaine Abuse Pharmacotherapy*
 Role: Principal Investigator
 Funding Period: 08/01/09-09/14/12
 Total Costs: \$91,563

Source: University of Florida Research Foundation Commercialization Fund (Project 100721)
 Project Name: *Commercialization of Drug Candidate for Schizophrenia*
 Role: Principal Investigator
 Funding Period: 03/01/12-09/01/12
 Total Costs: \$25,311

Source/Number: NIH (Mental Health, NIMH) R01MH068655
 Project Name: *Functional Probes for Brain Histamine H₁ Receptors*
 Role: Principal Investigator
 Funding Period: 04/01/04-03/31/09
 Total Costs: \$1,055,113

Source/Number: UF Opportunity Fund 65651
 Project Name: *Preclinical Development of Drugs for Obesity and Cocaine Addiction*
 Role: Principal Investigator
 Funding Period: 07/01/07-06/30/09
 Total Costs: \$81,550

Source/Number: NIH (Neurological Disorders and Stroke, NINDS) R-29-NS35216
 Project Name: *Novel Sigma Ligands in Neurodegeneration*
 Role: Principal Investigator
 Funding Period: 04/01/97-03/31/02
 Total Costs: \$502,855

Source/Number: Pharmacy Foundation of North Carolina 6-68379-4501
 Project Name: *Evaluation of NMDA receptor-active compounds using primary glial cell cultures*
 Role: Principal Investigator
 Funding Period: 07/01/00-06/30/02
 Total Costs: \$20,000

Source/Number: Otsuka Pharmaceuticals 6-68347-4281
 Project Name: *Biochemical Effects of Novel Quinoline Compounds at Brain Sigma Receptors*
 Role: Co-Principal Investigator (Richard B. Mailman, Co-PI)
 Funding Period: 09/01/98-12/31/01
 Total Costs: \$97,0000

Source/Number: Environmental Protection Agency
Project Name: *Neurotoxicity of Polychlorinated Biphenyls*
Role: Principal Investigator
Funding Period: 08/01/97-09/31/99
Total Costs: \$45,000

Source/Number: NIMH RO1 MH34006
Project Name: *Pharmacology of Dopamine Receptors in CNS*
Role: Investigator (Ross Baldessarini, PI)
Funding Period: 07/01/95-06/30/99
Total Costs: \$1,200,000

Source/Number: Pharmacy Foundation of North Carolina
Project Name: *Putative Sigma-3 Receptors in Mammalian Brain*
Role: Principal Investigator
Funding Period: 01/01/95-12/31/97
Total Costs: \$10,000

Source/Number: Environmental Protection Agency 4D-1882
Project Name: *Polychlorinated Biphenyls Effects on Brain Dopamine Synthesis*
Role: Principal Investigator
Funding Period: 08/01/94-07/31/96
Total Costs: \$5,000

Source/Number: UNC-CH Faculty Research Grant 5-44786
Project Name: *Adenosine and Sigma Receptor-Mediated Regulation Dopamine Synthesis*
Role: Principal Investigator
Funding Period: 07/01/93-06/30/95
Total Costs: \$2,970

Source/Number: Otsuka Pharmaceuticals 6-68347
Project Name: *Novel Quinoline Inhibitors of Tyrosine Hydroxylase*
Role: Co-Principal Investigator (Richard B. Mailman, Co-PI)
Funding Period: 07/01/93-06/30/97
Total Costs: \$24,000

Source/Number: NIMH RO1 MH40537
Project Name: *A Novel Molecular Site for Antidopaminergic Action*
Role: Investigator (Richard B. Mailman, PI)
Funding Period: 04/01/93-03/31/97
Total Costs: \$1,291,981

Source/Number: UNC-CH Faculty Research Grant 5-44339
Project Name: *Development of Autoreceptor Agonists*
Role: Principal Investigator
Funding Period: 12/01/90-06/30/93
Total Costs: \$2,550

Source/Number: UNC-CH Junior Faculty Development Award-6-69410
Project Name: *Characterization of Autoreceptors in Mammalian Forebrain*
Role: Principal Investigator
Funding Period: 01/01/91-12/31/92
Total Costs: \$3,000

Source/Number: NIH BSRG RR05967
Funding Period: 10/01/91
Total Costs: \$5,800 (instrument purchase)

Source/Number: NIMH MH14275-15
Project Name: *Neuropharmacology of Presynaptic Dopamine Agonists*
Role: Investigator (W.H. Morse, PI)
Funding Period: 11/01/88-08/01/90
Total Costs: \$39,700

TEACHING (Northeastern University)

GRADUATE CURRICULUM

CHEM 5676 Bio-organic and Medicinal Chemistry

3 semester hours, elective; 15 students
2014-current (Course Coordinator)

PHSC 6300 Seminar in Drug Discovery

1 semester hour, required; 12 students
2014-current (Course Coordinator)

PHSC 5100 Concepts in Pharmaceutical Science

Drug Metabolism and Drug Design and Development
2 semester hours, required; 75 students
2013-current

PMCL 6262 Receptor Pharmacology

2 semester hours, required; 25 students
2014-current

CHEM 5620 Protein Chemistry

3 semester hours, elective; 25 students
2016-current

PHSC 6222: Chemistry and Biology of Drugs of Abuse

2 semester hours, elective; 20 students
2013-current

PHSC 5934 Research in Pharmaceutical Science (MS level lab course)

2 semester hours, elective, 5 students
2013-current

CHEM 4456 Organic Chemistry III

2 semester hour, elective; 20 students
2015 – 2016 (3 lecture hours)

PROFESSIONAL (PHARM.D.) AND UNDERGRADUATE CURRICULA

HLTH 1555 Honors Special Topics in Healthcare: Drug Discovery and Delivery
4 semester hours, 12 students; elective
2012-2014, 2015 (course coordinator and lecturer)

HONR 3310-12: Approaches to Drug Therapy in the US.
1 semester hour, elective; 15 students
2014 – current

GRADUATE STUDENT TRAINEES**CURRENT GRADUATE STUDENT TRAINEES**

Austen Casey, Ph.D., Medicinal Chemistry, 2020 (expected)

Kirin Gada, Ph.D., Pharmacology, 2020 (expected)

Charles Perry, Ph.D., Medicinal Chemistry, 2018 (expected)

FORMER GRADUATE STUDENT TRAINEES

Sisy Hu, M.S., Pharmaceutical Sciences 2017

Yiming Chen, MS., Pharmaceutical Sciences 2017

Hima Patel M.S., Pharmaceutical Sciences 2017

Eliza Miller, M.S., Chemistry and Chemical Biology, 2016

Yajun Lin, M.S., Pharmaceutical Sciences 2016

Yan Zhou, M.S., Pharmaceutical Sciences 2016

Laura Purcell, M.S., Pharmaceutical Sciences 2016

Daoyang Chen, M.S., Pharmaceutical Sciences 2016

Ngyn Tran, M.S., Pharmaceutical Sciences, 2015

Rinkal Soni, M.S., Pharmaceutical Sciences, 2015

Bryce Suchomel, M.S., Pharmaceutical Sciences, 2015

Wanying Zhu, M.S., Pharmaceutical Sciences, 2014

Daniel Felsing, Ph.D, Medicinal Chemistry, 2016.

Thesis title: Drug discovery targeting serotonin G protein-coupled receptors to treat neuropsychiatric disorders.

Current position: Postdoctoral research fellow, Virginia Commonwealth University, Department of Biochemistry and Biophysics

Krishnakanth Kondabolu, Ph.D, Medicinal Chemistry 2013.

Thesis title: In vitro and in vivo pharmacology of novel phenylaminotetralin (PAT) analogs at serotonin 5-HT₂ receptors: Development of drugs for neuropsychiatric disorders.

Current position: Postdoctoral Research Fellow, Dept. Pharmacology, Boston University

Sean Travers, Ph.D., Medicinal Chemistry, 2011

Thesis title: Characterization of the molecular determinants for class A G protein-coupled receptors: Drug discovery targeting the histamine H₁ receptor

Current position: Technical Research Advisor Rigaku Raman Technologies

Zhuming Sun, Ph.D., Medicinal Chemistry, 2010.

Thesis title: "Novel phenylaminotetralin (PAT) analogs: Multifunctional serotonin 5-HT₂ receptor drugs for neuropsychiatric disorders".

Current position: Research Scientist, Novartis, Shanghai, China

Dawn Covington, M.S. Medicinal Chemistry (non-thesis), 2007.

Ola Maher Ghoneim, Ph.D., Medicinal Chemistry, 2006

Thesis title: "Synthesis, analytical, and molecular modeling studies of novel aminophenyltetralin ligands to characterize human histamine and serotonin receptor signaling"

Current position: Assistant Professor (teaching, research, service), Department of Medicinal Chemistry, St. Joseph College, Hartford, Connecticut

Nader Moniri, Ph.D., Medicinal Chemistry, 2004

Thesis Title: "Histamine H₁ receptor multifunctional signaling characterized using novel tetrahydro-(naphthalene and benzocycloheptane) ligands"

Current position: Associate Professor (teaching, research, service), Medicinal Chemistry, Mercer University, Atlanta, GA

Jacqueline Legere, Ph.D., Medicinal Chemistry, 2004

Thesis Title: "Synthesis and pharmacological activity of 2-dimethylamino-5-(6)-phenyl-1,2,3,4-tetrahydronaphthalenes as novel ligands for the human histamine H₁ receptor"

Current Position: Director of Biology Research Core, Genzyme, Framingham, MA

Neepa Choksi, Ph.D., Medicinal Chemistry, 1999

Thesis Title: "Neuropharmacological Characterization of Brain Receptors Recognized by 1-Phenyl-3-aminotetralins"

Current position: Senior Research Scientist, Integrated Laboratory Systems Inc., Research Triangle Park, NC

Constance E. Owens, Ph.D., Medicinal Chemistry, 1997

Thesis title: "A Novel Brain Receptor System Linked to Modulation of Catecholamine Synthesis"

Current position: Senior Research Scientist, Endacea, Research Triangle Park, North Carolina

Anwar Hussain, M.S., Medicinal Chemistry, 1995

Thesis Title: "The Role of Adenosine A₂ Receptors in Stimulation of Brain Dopamine Synthesis"

Last known position: Senior Research Scientist, Bristol-Myers-Squibb, New Brunswick, New Jersey.

POSTDOCTORAL & RESEARCH ASSISTANT PROFESSOR TRAINEES

Munmun Mukherjee, Ph.D. (2017-current)

Shan Zhu, Ph.D. (2014-2016)

Yue Liu, Ph.D. (2013-2016); current position is research scientist at Pfizer Inc., Cambridge, MA

Rajender Vemula, Ph.D. (2012-2016); current position is research scientist at Harvard University Department of Chemistry, Cambridge, MA.

Tania Cordova-Sintjago, Ph.D. (2010-2016); current position is faculty (teaching and research) at Santa Fe College, Gainesville FL

Dario Ambrosini, Ph.D. (2012-2015); current position is postdoctoral researcher at University of Milan Department of Chemistry, Milan, Italy.

Clinton Canal, Ph.D. (Postdoctoral: 2010-2012); current position is Research Assistant Professor at Northeastern University Department of Pharmaceutical Sciences and Center for Drug Discovery, Boston, MA

Myong Sang Kim, Ph.D. (Postdoctoral: 2009-2012); current position is research scientist at Firebird Biomolecular Sciences, Alachua, FL 32615-9465, USA.

Nancy Villa, Ph.D. (2009-2012); current position is research scientist at University of Florida Department of Oncology, Gainesville, FL

Jean-Claude Nzimulinda, Ph.D. (2010-2011); current position is staff pharmacist in Louisville, KY.

Rajeev Sakhujia, Ph.D. (2009-2011); current position is faculty (teaching, research) at Birla Institute of Technology & Science, Pilani, Rajasthan 333031, India.

Adam Vincek, Ph.D (2008-2011); current position is research scientist in Center for Drug Discover at Mount Sinai School of Medicine, New York, NY 10029, USA.

Li Fang, Ph.D. (2006-2009); current position is research scientist at University of Florida College of Medicine Department of Nephrology

Andrzej Wilczynski, Ph.D (2006-2008); current position is research scientist at Perkin Elmer, Boston, MA

Sashi Sivendran, Ph.D. (2006-2008); current position is Dow Chemicals, Andover, MA

Yingsu Huang, Ph.D. (2004-2005)

Aaron Meng, M.D. (1999-2001)

Bonita L. Blake, D.V.M., Ph.D. (1997-1999)

UNDERGRADUATE and MS RESEARCH TRAINEES

George (Jia Xing) Guo, B.S., Biochemistry (expected, 2018)

Christopher Chang, B.S., Chemistry and Chemical Biology (expected, 2018)

Jessica Mecklosky, B.S. Neuroscience, 2016.

Daniel Felsing, B.S., Chemistry, 2011

Sean Wimberly, Pharm.D., 2011

Roberto Campillo, Pharm.D., 2007

Russell Moore, Pharm.D. 2006

Christopher Smelick, B.S. 2006

Tammy Bristow, B.S., Chemistry, 2004

Alexandra Calves; B.S., Pharmacy, 1998

Kathrinn Fitzpatrick, B.S., Pharmacy, 1997

Brenda Aske, B.S., Pharmacy, 1996

Kelly Hendricks; B.S., Pharmacy, 1996

R. Donald Harvey; Pharm.D., 1994

Jonathan Ducar; B.S., Pharmacy, 1992

THESIS DISSERTATION COMMITTEES

Northeastern University

Chandrashekar Honrao Ph.D., Pharmaceutical Sciences (Drug Discovery/Medicinal Chemistry), 2016 (expected); Committee Member

Anisha Korde, Ph.D., Pharmaceutical Sciences (Medicinal Chemistry), 2016 (expected), Committee Member

Shashank Kulkarni, Ph.D., Pharmaceutical Sciences (Medicinal Chemistry), 2016 (expected), Committee Member

Pooja Naresh Sabhachandani, Ph.D., Pharmaceutical Sciences (Pharmaceutics) 2016 (expected), Committee Member

Abhijit Kulkarni, Ph.D., Pharmaceutical Sciences (Medicinal Chemistry), 2016 (expected), Committee Member

Mohammed Baradwan, Ph.D., Pharmaceutical Sciences (Drug Discovery/Medicinal Chemistry), 2018 (expected); Committee Member.

Michael Johnson, Ph.D., Pharmaceutical Sciences (Drug Discovery/Medicinal Chemistry), 2014; Committee Member.

University of Florida

Ebrahim Hossein Ghazvini Zadeh, Ph.D., Chemistry, 2015; Committee Member

Harald Messer, Ph.D., Interdisciplinary Program in Biomedical Science, 2015 (expected), Committee Member

Zheng Zheng, Ph.D., Chemistry, 2014 (expected), Committee Member

James Kasper, Ph.D., Pharmacodynamics, 2012 (expected), Committee Member

Wenjen Li, Ph.D, Medicinal Chemistry, 2011, Committee Member

Jongwoo Park, Ph.D., Chemistry, 2011, Committee Member

Bin Song, Ph.D., Computer Information Science & Engineering, 2010, Committee Member

Jason Kwan, Ph.D., Medicinal Chemistry, 2010, Committee Member

Bettina Proneth, Ph.D., Medicinal Chemistry, 2007

James Sacco, Ph.D., Medicinal Chemistry, 2006

Joshua Thomas, Ph.D., Medicinal Chemistry, 2006

Susruta Majumdar, Ph.D, Medicinal Chemistry, 2006

University of North Carolina at Chapel Hill

Richard Durham, Ph.D., Medicinal Chemistry, 2005

Erin Heinzen, Ph.D., Pharmaceutics Division: Drug Delivery and Disposition, 2004

Yolanda Taylor, M.S., 2001 (Biochemistry)

Erhen Bucholtz, Ph.D., Medicinal Chemistry, 1998
Brian Hoffman, Ph.D., Medicinal Chemistry, 1998
Daphne Da, M.S., Pharmaceutics Division: Drug Delivery and Disposition, 1998
Merril Miller, Ph.D., Medicinal Chemistry, 1996
Andrew Myers, Ph.D., Medicinal Chemistry, 1995
Jin Sung Cho, Ph.D., Medicinal Chemistry, 1995
Ke Chen, Ph.D., 1995
Lynne Canne, Ph.D. Medicinal Chemistry, 1995
Leping Lee, Ph.D., Medicinal Chemistry, 1994
Bruce Burnham, Ph.D., Medicinal Chemistry, 1994
Debra Minor, Ph.D., Medicinal Chemistry, 1994
Christopher Waller, Ph.D., Medicinal Chemistry, 1993

TEACHING AWARDS

- 2006:** BEST PROFESSOR AWARD, University of North Carolina at Chapel Hill
School of Pharmacy, Class of 2006
- 2005:** ACADEMIC EXCELLENCE AWARD IN TEACHING
Nominated by Chair of Medicinal Chemistry – award to be announced July 2005
- 2003:** BEST PROFESSOR AWARD, University of North Carolina at Chapel Hill
School of Pharmacy, Class of 2005
- 2002:** BEST PROFESSOR AWARD, University of North Carolina at Chapel Hill
School of Pharmacy, Class of 2004
- 2001:** BEST PROFESSOR AWARD, University of North Carolina at Chapel Hill
School of Pharmacy Class, of 2003
- 1999:** BEST PROFESSOR AWARD, University of North Carolina at Chapel Hill
School of Pharmacy, Class of 2000
- 1998:** DISTINGUISHED TEACHING AWARD FOR POST-BACCALAUREATE
INSTRUCTION, *Nominee*, University of North Carolina at Chapel Hill
- 1997:** BEST PROFESSOR AWARD, *Honorable Mention*, University of North Carolina at Chapel Hill
School of Pharmacy, Class of 1999

SERVICE

National Institutes of Health

NIDA Board of Scientific Counselors (2014 - current).

K25 Award Mentor for Dr. Anna Sromek's project titled, Dextromorphan Medications for Treating Opioid Addiction (2014-current)

ETTN-M 11 B: Drug Discovery for Aging, Neuropsychiatric and Neurologic Disorders (2012 - current)

EMNR-R (56): PAR Panel NIDDK Translational Research *Ad hoc* (2014)
 Emerging Technologies and Training in Neurosciences C-11 (2009 - 2012)
 NIA Translational Research in Aging ZAG1 ZIJ-1 B *Ad hoc* (December, 2011)
 NIDA Medications Development Program Projects for Substance-Related Disorders (October, 2011)
 NIDA New Molecular Entities to Treat Substance Use Disorders, Chairman (May, 2011)
 NIDA Clinical Research Training Program *Ad hoc* (March: 2009, 2010, 2011, 2012; 2013; 2014)
 Molecular, Cellular, and Developmental Neuroscience MDCN-F *Ad hoc* (February, 2010)
 NIDA Development of Therapeutic Agents Special Emphasis Panel *Ad hoc* (December, 2008)
 NIMH Conference to Advance Mental Health Research Special Emphasis Panel *Ad hoc* (October, 2008)
 Brain Disorders and Clinical Neuroscience Study Section F-11 (2005-2008)
 Brain Disorders and Clinical Neuroscience Study Section K-15 *Ad hoc* (March, 2006)
 Fundamental Neurosciences Study Section, *Ad hoc* (1997)
 Brain Disorders and Clinical Neuroscience Study Section K-15 (1996-2000)
 Technology and Applied Sciences Study Section (1992-1996)

Environmental Protection Agency

Chemical Mixtures in Environmental Health Panel (1997-2000)
 Exploratory Research on Environmental Neurotoxicants Panel (1997-2000)

Editorial Board

Obesity Insights (2008-current)
 Medicinal Chemistry: Current Research (2010-current)
 Medicinal Chemistry (open access) (2012-current)
 (ISRN) Pharmacology (2010-current)
 Journal of Addiction and Prevention (2013-current)

Northeastern University

Pharmaceutical Sciences Interim Chair (2017-current)
 Pharmaceutical Sciences Graduate Committee (2016-current)
 Pharmaceutical Sciences Senior Hires (2) Pharmacology Recruitment Committee 2013-2017 (Chair)
 Pharmaceutical Sciences Chair Recruitment Committee 2014-2016 (Chair)
 University Grievance Committee 2014-15 (member)
 Chemistry and Chemical Biology Computational Chemistry Senior Hire Recruitment 2013-14 (Member)
 Center for Drug Discovery Senior Hires Recruitment Committee 2012- (co-Chair)
 School of Pharmacy Assessment Committee 2012 – (Member)
 Bouvé College of Health Sciences Dean's Leadership Team 2013 – (Member)
 Northeastern University Committee for Research Policy Oversight 2013- (Member)
 Northeastern University Research Awards Administration Advisory Committee 2013- (Member)

CONSULTING ACTIVITY

Sterne, Kessler, Goldstein, Fox, LLP; Washington DC (2007-2013)

Johnson and Johnson Pharmaceutical Research and Development (2008-2010)

Patterson, Belknap, Webb & Tyler, LLP; NYC (2004-2005)

Pitts, Hugenschmidt & Devereux, PA; Asheville, NC (1998-2000)

GlaxoSmithKline, Research Triangle Park, NC (1992-99)

CURRENT LICENSURE Registered Pharmacist: Massachusetts #19004; California #PL-038684

INVITED ORAL PRESENTATIONS

D₂-Autoreceptor-Mediated Inhibition of Dopamine Synthesis in Brain. Department of Pharmacology and Experimental Therapeutics, School of Medicine, Boston University, October 16, 1990

Presynaptic Modulation of Brain Dopamine Synthesis. Department of Medicinal Chemistry, School of Pharmacy, University of North Carolina at Chapel Hill, November 1, 1990.

Stimulation of Brain Dopamine Synthesis by Sigma Ligands. Department of Medicinal Chemistry, School of Pharmacy, Medical College of Virginia, Richmond, VA, February 15, 1991

Effects of Novel Dopamine Agonists on Striatal Dopamine Synthesis. Biological Sciences Research Center, School of Medicine University of North Carolina at Chapel Hill, March 15, 1991.

Regulation of Brain Catecholamine Synthesis. Department of Neuropharmacology, Molecular Devices Corporation Menlo Park, CA, Sept. 11, 1992.

Mechanisms of Dopamine Cell Death. Neuroscience Department, Amgen Inc. Thousand Oaks, CA, October 2, 1992

Introduction to Pharmacology. Department of Medicinal Chemistry, Glaxo Pharmaceutical Research, Research Triangle Park, NC, June 2-11, 1992.

Receptor-Mediated Regulation of Dopamine Synthesis. Division of Pharmaceutics, School of Pharmacy, University of North Carolina at Chapel Hill, Nov. 12, 1992.

Principles of Pharmacology with Applications for Medicinal Chemistry. Department of Medicinal Chemistry, Glaxo Pharmaceutical Research, Research Triangle Park, NC, April 19-30, 1993.

A Novel Sigma Receptor Modulates Brain Dopamine Synthesis. Program in Neurobiology, School of Medicine, University of North Carolina at Chapel Hill, March 31, 1994.

Discovery of a Novel Sigma Receptor in Mammalian Brain. Division of Pharmaceutics, School of Pharmacy, University of North Carolina at Chapel Hill, February 1, 1996

Characterization of a Novel Brain Sigma Receptor. University of the Pacific, School of Pharmacy, Stockton, CA, March 22, 1996.

Neuromodulatory Adenosine A₂ receptors. Curriculum in Neurobiology, School of Medicine, University of North Carolina at Chapel Hill, February, 19, 1997.

Environmental Agents Linked to Neurotoxicity and Parkinson's Disease. Environmental Protection Agency, Research Triangle Park, North Carolina, December 11, 1997

Polychlorinated Biphenyl Compounds Inhibit Brain Dopamine Synthesis. Environmental Protection Agency, Research Triangle Park, North Carolina, November 5, 1998

Phenylaminotetralins as Novel Histamine H₁ Ligands. Department of Pharmacochimistry, Vrije (Free) University, Amsterdam, The Netherlands; March 9, 1999.

Novel Phenylaminotetralins Activate Neuromodulatory H₁ Receptors to Stimulate Brain Dopamine Synthesis. Department of Pharmacochimistry, Vrije (Free) University, Amsterdam, The Netherlands; April 11, 2000.

Molecular Mechanisms Regulating Brain Neurotransmitter Synthesis. Cogent Neurosciences Corporation, Durham, North Carolina, September 8, 2000.

Medicinal Chemical Probes Useful to Characterize H₁ Dimer Expression. Department of Pharmacochimistry, Vrije (Free) University, Amsterdam, The Netherlands; March 29, 2001.

Functional Heterogeneity of Histamine H₁ Receptors. Presented at the European Medicinal Chemistry Society Meeting, Noordwijkerhout, The Netherlands; May 7-11, 2003.

Ligand-directed multifunctional signaling of histamine H₁ and related G protein-coupled receptors. Histamine Research Society, Cologne, Germany, April 27-May 2, 2004.

Ligand Functional Selectivity at G Protein-Coupled Receptors: Impact on Compound Selection in Drug Discovery. Gordon Research Conference on Molecular Pharmacology, Barga, Italy, May 8-13, 2005.

Novel Ligands Stabilize Stereo-Selective Conformations of the H₁ Receptor: Functionally-Selective Modulation of Intracellular Signaling Pathways. Histamine Research Society, Delphi, Greece, May 10-14, 2006.

Molecular Determinants of Ligand-Directed G_q versus G_s Signaling for the Histamine H₁ Receptor. Histamine Research Society, Florence, Italy, May 9-13, 2007.

GPCR Conformations that Activate Phospholipase C vs Adenylyl Cyclase: Stereoselective Stabilization of Histamine H₁ Signaling by Novel Ligands. University of Camerino Medicinal Chemistry Symposium, Camerino, Italy, September 9-13, 2007.

Histamine H₁ Receptor Mutagenesis Studies to Characterize Molecular Determinants for Ligand Activation. Department of Pharmacochimistry, Vrije (Free) University, Amsterdam, The Netherlands; March 13, 2008

New Drug Candidates for Treating Obesity and Neuropsychiatric Disorders. Health Care Ventures. Cambridge, MA, March 20, 2008

Novel Drugs Targeting Serotonin 5HT_{2A/C} Receptors: Antipsychotics Devoid of Weight-Gain Effects. Department of Psychiatry, Harvard Medical School and McLean Psychiatric Hospital, Belmont, MA, April 14, 2008.

Drug Discovery Targeting Brain Serotonin 5HT₂ G Protein-Coupled Receptors: Antiobesity and Antipsychotic Pharmacotherapy. Conference on New Methods in Drug Research, Limassol, Cyprus, May 11 – 14, 2008.

Transmembrane Helix Position 5.48 Involvement in the Dimer Interface of Domain-Swapped GPCRs. 27th Symposium on Trends in Drug Research, Noordwijkerhout, The Netherlands, May 3-7, 2009.

Novel serotonin 5HT_{2C} agonist drugs with 5HT_{2A/2B} inverse agonism for drug addiction. National Institute of Drug Abuse Binational Research Collaboration on Drug Abuse and Addiction. Washington, DC. October 22-23, 2009.

Functionally-Selective Serotonin 5-HT₂ GPCR Drugs: 5-HT_{2C} agonists with 5-HT_{2A/2B} inverse agonist activity for neuropsychiatric disorders. 28th Camerino-Cyprus-Noordwijkerhout International Medicinal Chemistry Symposium. Camerino, Italy, May 16-20, 2010.

Drug Development Targeting Serotonin 5HT_{2A/2C} Receptors: Pharmacotherapy for Neuropsychiatric Disorders Without Weight-Gain. University of Florida, College of Medicine, Department of Pharmacology and Therapeutics, October 19, 2010.

Drug Development Targeting Serotonin 5HT_{2A/2C} Receptors: Pharmacotherapy for Neuropsychiatric Disorders Without Weight-Gain. Northeastern University Department of Pharmaceutical Sciences, April 21, 2011.

Novel Serotonin 5-HT₂ Receptor Modulators Demonstrate In Vivo Efficacy for Psychoses. 29th Camerino-Cyprus-Noordwijkerhout International Medicinal Chemistry Symposium., Limassol, Cyprus, October 2-5, 2011.

Novel serotonin 5HT₂ drugs for cocaine addiction. National Institute of Drug Abuse Binational Research Collaboration on Drug Abuse and Addiction. The Hague, Netherlands, October 11-14, 2011.

Development of Serotonin 5HT₂ Modulators to Treat Psychoses and Addiction. University of Florida Department of Psychiatry, November 17, 2011.

Integrating Medicinal Chemistry, Pharmacology, and Behavioral Research Toward Novel Serotonin 5-HT₂-based Pharmacotherapy for Neuropsychiatric Disorders. Lowenthal Symposium Keynote Speaker. Virginia Commonwealth University/Medical College of Virginia School of Pharmacy, May 24, 2012.

Novel orally-active 5-HT_{2c} agonists with 5-HT_{2A/2B} antagonist activity demonstrate efficacy to treat psychoses without weight gain. 30th Camerino-Cyprus-Noordwijkerhout International Medicinal Chemistry Symposium., Camerino Italy, May 19-24, 2013.

Serotonin Receptor Type-2 (5-HT₂) Drugs for Obesity and Neuropsychiatric Disorders: What you should know before putting your patient to sleep. New England Assembly of Nurse Anesthetists Annual Meeting, Burlington, MA April 4-6, 2014

Serotonin 5-HT₂ Receptor Structure-Based Drug Design Leads to Clinical Candidates for Neuropsychiatric Disorders. The 32nd Cyprus-Noordwijkerhout-Camerino Symposium in Trends in Drug Research, sponsored by the European Federation of Medicinal Chemistry, Cyprus, May 18-22, 2014.

PUBLICATIONS

Peer-reviewed research journal papers

1. Booth, R.G., Selassie, C.D., Hansch, C., and Santi, D.V. Quantitative structure-activity relationship of triazine-antifolate inhibition of *Leishmania* dihydrofolate reductase and cell growth. *Journal of Medicinal Chemistry* 30: 1218-1224 (1987).
2. Ramsay, R.R., McKeown, K.A., Johnson, E.A., Booth, R.G., and Singer, T.P. Inhibition of NADH oxidation by pyridine derivatives. *Biochemical and Biophysical Research Communications* 146: 53-60 (1987).
3. Booth, R.G., Rollema, H., and Castagnoli, N. *In vivo* dopaminergic neurotoxicity of the 2-b-methyl-carbolinium ion, a potential endogenous MPP⁺ analog. *European Journal of Pharmacology* 153: 131-134 (1988).
4. Sirawaraporn, W., Sertsrivanich, R., Booth, R.G., Hansch, C., Neal, R.A., and Santi, D.V. Inhibition of *Leishmania* dihydrofolate reductase and *Leishmania* growth by 5-benzyl-2,4-diaminopyrimidines. *Molecular Biochemical Parasitology* 31: 79-86 (1988).
5. Booth, R.G., Trevor, A.J., Singer, T.P., and Castagnoli, N. Studies on semi-rigid tricyclic analogs of the nigrostriatal toxin 1-methyl-4-phenyl-1,2,3,6-tetrahydropyridine (MPTP). *Journal of Medicinal Chemistry* 32: 473-477 (1989).
6. Booth, R.G., Castagnoli, N., and Rollema, H. Intracerebral microdialysis neurotoxicity studies of quinoline and isoquinoline derivatives related to MPTP/MPP⁺. *Neuroscience Letters* 100: 306-312 (1989).

7. Johnson, E.A., Wu, E.Y., Rollema, H., Booth, R.G., Trevor, A.J., and Castagnoli, N. MPP⁺ Analogs: *In vivo* neurotoxicity and inhibition of striatal synaptosomal dopamine uptake. *European Journal of Pharmacology* 166: 65-74 (1989).
8. Booth, R.G., Baldessarini, R. J., Kula, N.S., Zong, R., Gao, Y., and Neumeyer, J.L. Presynaptic inhibition of dopamine synthesis in rat striatal tissue by enantiomeric mono- and dihydroxyaporphines. *Molecular Pharmacology* 38: 92-101 (1990).
9. Rollema, H., Booth, R.G., Caldera, P., Johnson, E.A., Lampen, P., Youngster, S.K., Trevor, A.J., Naiman, N., and Castagnoli, N. *In vivo* intracerebral microdialysis studies in rats of MPP⁺ analogs and related charged species. *Journal of Medicinal Chemistry* 33: 2221-2230 (1990).
10. Booth, R.G., and Baldessarini, R.J. Adenosine A₂ stimulation of tyrosine hydroxylase activity in rat striatal minces is reversed by dopamine D₂ autoreceptor activation. *European Journal of Pharmacology* 185: 217-221 (1990).
11. Booth, R.G., Baldessarini, R.J., Kula, N., and Neumeyer, J.L. Stereochemical effects of mono- and dihydroxyaporphines on presynaptic inhibition of tyrosine hydroxylase *in vitro*. *Annals of New York Academy of Science* 604: 592-595 (1990).
12. Baldessarini, R.J., Booth, R.G., Campbell, A., and Neumeyer, J.L. S(+)-Aporphines as potential limbic-selective antipsychotic agents. *Schizophrenia Research* 4: 311-312 (1991).
13. Booth, R.G., Baldessarini, R.J, and Campbell, A. Inhibition of dopamine synthesis in rat striatal minces: Evidence of autoreceptor supersensitivity to S(+) but not to R(-)-N-n-propylnorapomorphine after repeated pretreatment with fluphenazine. *Biochemical Pharmacology* 41: 2040-2043 (1991).
14. Booth, R.G. and Baldessarini, R.J. (+)-Benzomorphan sigma ligands stimulate dopamine synthesis in rat corpus striatum tissue. *Brain Research* 557: 349-352 (1991).
15. Teicher, M.H., Gallitano, A.L., Gelbard, H.A., Evans, H.K., Marsh, E.R., Booth, R.G., and Baldessarini, R.J. Dopamine D₁ autoreceptor function: Possible expression in developing rat prefrontal cortex and striatum. *Developmental Brain Research* 63: 229-235 (1992).
16. Wyrick, S.D., Booth, R.G., Myers, A.M., Kula, N.S., and Baldessarini, R.J. Synthesis of [*N*-C³H₃]-racemic-*trans*-1-phenyl-3-dimethylamino-6-chloro-7-hydroxy-1,2,3,4-tetrahydronaphthalene (PAT-6). *Journal of Labeled Compounds and Radiopharmaceuticals* 31:871-874 (1992).
17. Wyrick, S.D., Booth, R.G., Myers, A.M., Owens, C.E., Kula, N.S., Baldessarini, R.J, Mailman, R.B., Synthesis and pharmacological evaluation of 1-phenyl-3-amino-1,2,3,4-tetrahydronaphthalenes as ligands for a novel receptor with sigma-like neuromodulatory activity. *Journal of Medicinal Chemistry* 36: 2542-2551 (1993).
18. Booth, R.G., Wyrick, S.D., Baldessarini, R.J., Kula, N.S., Myers, A.M., and Mailman, R.B. A new sigma-like receptor recognized by novel phenylaminotetralins: Ligand binding and functional studies *Molecular Pharmacology* 44: 1232-1239 (1993).
19. Wyrick, S.D., Myers, A.M., Booth, R.G., Kula, N.S., Baldessarini, R.J., and Mailman, R.B. Synthesis of [*N*-C³H₃]-*trans*-(1*R*,3*S*)-(-)-1-phenyl-3-*N,N*-dimethylamino-1,2,3,4-tetrahydronaphthalene (H₂-PAT). *Journal of Labeled Compounds and Radiopharmaceuticals* 34: 131-134 (1994).

20. Booth, R.G., Baldessarini, R.J., Owens, C.E., and Marsh, E. Actions of 7-hydroxy-*N,N*-di-*n*-propyl-2-aminotetralin (7-OH-DPAT) on dopamine synthesis in limbic and extrapyramidal regions of rat brain. *Brain Research* 662:283-288 (1994).
21. Booth, R.G., and Wyrick, S.D. Development of phenylaminotetralin ligands for a novel sigma (σ_3) receptor in brain. *Medicinal Chemistry Research* 4:225-237 (1994).
22. Myers, A.M., Charifson, P.S., Owens, C.E., Kula, N.S., Baldessarini, R.J., McPhail, A.T., Booth, R.G., and Wyrick, S.D. Conformational analyses, pharmacophore identification, and comparative, molecular field analyses of ligands for the neuromodulatory σ_3 receptor. *Journal of Medicinal Chemistry* 37:4109-4117 (1995).
23. Wyrick, S.D. and Booth, R.G. Progress in sigma receptor research. *Drugs of the Future* 20:1033-1044 (1995).
24. Wyrick, S.D., Booth, R.G., Myers, A.M., Owens, C.E., Bucholtz, E.C., Hooper, P.C., Kula, N.S., Baldessarini, R.J., and Mailman, R.B. 1-Phenyl-3-amino-1,2,3,4-tetrahydronaphthalenes and related derivatives as ligands for the neuromodulatory σ_3 receptor: Further structure-activity relationships. *Journal of Medicinal Chemistry* 38:3857-3864 (1995).
25. Choksi, N.Y., Hussain, A., and Booth, R.G. 2-Phenylaminoadenosine stimulates dopamine synthesis in rat forebrain *in vitro* and *in vivo* via adenosine A_2 receptors. *Brain Research* 761:151-155 (1997).
26. Choksi, N.Y., Kodavanti, P.R.S., Tilson, H.A., and Booth, R.G. Effects of Polychlorinated Biphenyls (PCBs) on Brain Tyrosine Hydroxylase Activity and Dopamine Synthesis in Rats. *Fundamentals of Applied Toxicology* 39:76-80 (1997)
27. Bucholtz, E.C., Wyrick, S.D., Owens, C.E., and Booth, R.G. 1-Phenyl-3-dimethylaminotetralins (PATs): Effect of stereochemistry on binding and function at brain histamine receptors. *Medicinal Chemistry Research* 8:322-332 (1998).
28. Bucholtz, E.C., Brown, R.L., Tropsha, A., Booth, R.G, and Wyrick, S.D. Synthesis, Evaluation and Comparative Molecular Field Analysis of 1-Phenyl-3-amino-1,2,3,4-tetrahydronaphthalenes as Ligands for Histamine H_1 Receptors. *Journal of Medicinal Chemistry* 42:3041-3054(1999).
29. Booth, R.G., Owens, C.E., Brown, R.L., Bucholtz, E.C., Lawler, C.P., and Wyrick, S.D. Putative σ_3 sites in mammalian brain have histamine H_1 receptor properties: Evidence from ligand binding and distribution studies with the novel H_1 radioligand [3H]-(-)-*trans*-1-phenyl-3-aminotetralin (PAT). *Brain Research* 837:95-105 (1999).
30. Choksi, N.Y., Nix, William B., Wyrick, S.D., and Booth, R.G. A novel phenylaminotetralin recognizes histamine H_1 receptors and stimulates dopamine synthesis *in vivo* in rat brain. *Brain Research* 852:151-160 (2000).
31. Mottola, D., Kilts, J., Lewis, M., Smith, H., Walker, Q.D., Jones, S., Booth, R.G., Hyslop, D., Piercey, M., Wightman, M., Lawler, C., Nichols, D.E., and Mailman, R.B. Functional selectivity of dihydrexidine: I. Selective activation of post-synaptic dopamine D_2 receptors linked to adenylate cyclase. *Journal of Pharmacology and Experimental Therapeutics* 301:1166-1178 (2002).
32. Booth RG, Moniri NH, Bakker RA, Choksi NY, Timmerman H, and Leurs R. A novel phenylaminotetralin radioligand reveals a sub-population of histamine H_1 receptors. *Journal of Pharmacology and Experimental Therapeutics* 302:328-336 (2002).

33. Moniri NH, Booth RG. Functional heterogeneity of histamine H₁ receptors. *Inflammation Research* 53:71-73 (2004)
34. Bakker RA, Dees G, Carrillo JJ, Booth RG, López-Gimenez JF, Graeme Milligan G, Strange PG, Leurs R. Domain swapping in the human histamine H₁ receptor. *Journal of Pharmacology and Experimental Therapeutics* 311:131-138 (2004).
35. Moniri NH, Covington-Strachan D, Booth RG. Ligand-directed functional heterogeneity of histamine H₁ receptors: Novel dual-function ligands selectively activate and block H₁-mediated phospholipase C and adenylyl cyclase signaling. *Journal of Pharmacology and Experimental Therapeutics* 311:274-281 (2004).
36. Heinzen EL, Booth RG, Pollack GM. Neuronal nitric oxide modulates morphine antinociceptive tolerance by enhancing constitutive activity of the μ -opioid receptor. *Biochemical Pharmacology* 69:679-688 (2005).
37. Booth RG, Moniri NH. Ligand-directed multifunctional signaling of histamine H₁ receptors *Inflammation Research* 54: S44-45 (2005).
38. Ghoneim OM, Legere JA, Glibraikh A, Tropsha A, Booth RG. Novel ligands for the human histamine H₁ receptor: Synthesis, pharmacology, and comparative molecular field analysis studies of 2-dimethylamino-5-(6)-phenyl-1,2,3,4-tetrahydronaphthalenes. *Bioorganic and Medicinal Chemistry* 14:6640-6658 (2006).
39. Moniri NH, Booth RG. Role of PKA and PKC in Histamine H₁ Receptor-Mediated Activation of Tyrosine Hydroxylase and Catecholamine Synthesis in Mammalian Brain and Adrenal Tissues. *Neuroscience*, 407:249-253 (2006).
40. Booth RG, Moniri NH. Novel Ligands Stabilize Stereo-Selective Conformations of the Histamine H₁ Receptor to Activate Catecholamine Synthesis. *Inflammation Research* 56:1-12 (2007).
41. Sansuk K, Balog CI, van der Does AM, Booth R, de Grip WJ, Deelder AM, Bakker RA, Leurs R, Hensbergen PJ. GPCR Proteomics: Mass Spectrometric and Functional Analysis of Histamine H₁ Receptor after Baculovirus-Driven and in Vitro Cell Free Expression. *Journal of Proteome Research*, 7:621-629 (2008).
42. Booth RG, Fang L, Wilczynski A, Sivendren S, Sun Z, Travers S, Bruysters M, Sansuk K, Leurs R. Molecular determinants of ligand-directed signaling for the histamine H(1) receptor. *Inflammation Research*, 57:S40-44 (2008).
43. Rowland N, Crump E, Nguyen N, Robertson K, Booth RG. Effect of (-)-Trans-PAT, a novel 5-HT_{2C} receptor agonist, on intake of palatable food in mice. *Pharmacology, Biochemistry and Behavior* 91:176-180 (2008).
44. Booth RG, Rowland N, and Gingrich JA. A Novel Serotonin 5-HT_{2C} agonist with 5-HT_{2A}/5-HT_{2B} inverse agonist activity demonstrates antipsychotic efficacy without weight-gain liability. *Drugs of the Future* 33 (Suppl. A): 68-69 (2008).
45. Booth RG, Fang L, Huang Y, Wilczynski A, Sivendran S. (1R, 3S)-(-)-Trans-PAT: A novel full-efficacy serotonin 5-HT_{2C} receptor agonist with 5-HT_{2A} and 5-HT_{2B} receptor inverse agonist/antagonist activity. *European Journal of Pharmacology* 615: 1-9 (2009).
46. Vincek AS, Booth RG. Title: New Approach to 4-Phenyl- β -aminotetralin from 4-(3-Halophenyl)tetralen-2-ol Phenylacetate. *Tetrahedron Letters* 50:5107-5108 (2009).

47. Sansuk K, Balog CI, van der Does AM, Booth RG, de Grip WJ, Deelder AM, Bakker RA, Leurs R, Hensbergen PJ. GPCR proteomics: mass spectrometric and functional analysis of histamine H1 receptor after baculovirus-driven and in vitro cell free expression. *J Proteome Res.* 2008 Feb;7(2):621-9. doi: 10.1021/pr7005654. PMID:18177001
48. Canal CE, Cordova-Sijntjago T, Villa N, Fang L, Booth RG. Drug discovery targeting human 5-HT_{2C} receptors: Residues S3.36 and Y7.43 impact ligand-binding pocket structure via hydrogen bond formation. *European Journal of Pharmacology* 2011; 673:1-12.
49. Ghoneim OM, Ibrahim DA, El-Deeb IM, Lee SH, Booth RG. A novel potential therapeutic avenue for autism: Design, synthesis and pharmacophore generation of SSRIs with dual action. *Bioorganic and Medicinal Chemistry* 21:6714-6723 (2011).
50. Córdova-Sintjago T, Villa N, Canal CE, Booth RG. Human serotonin 5-HT_{2C} G protein-coupled receptor homology model from the β_2 adrenoceptor structure: Ligand docking and mutagenesis studies. *Int. Journal of Quantum Chemistry* 2011; 112:140-149.
51. Córdova-Sintjago T, Sakhuja R, Kondabolu K, Canal CE, Booth RG. Molecular determinants for ligand binding at serotonin 5-HT_{2A} and 5-HT_{2C} GPCRs: Experimental affinity results analyzed by molecular modeling and ligand docking studies. *Int. Journal Quantum Chemistry* 2012; 112:3807-3814.
52. Córdova-Sintjago, TC, Fang, L, Bruysters M, Leurs, R, Booth RG. Molecular determinants of ligand binding at the human histamine H1 receptor: Site-directed mutagenesis results analyzed with ligand docking and molecular dynamics studies at H1 homology and crystal structure models. *Journal of Chemical and Pharmaceutical Research* 2012; 4:2937-2951.
53. Morgan D, Canal CE, Kondabolu K, Sakhuja R, Robertson K, Rowland NE, Booth RG. A Novel Serotonin-2 (5-HT₂) Modulator as a Candidate Drug to Treat Impulsive Behavioral Disorders and Psychoses without Weight Gain as a Side Effect. *Neuropsychopharmacology* 2012;38:S104-105.
54. Canal CE, Booth RG, Morgan D. Support for 5-HT_{2C} receptor functional selectivity in vivo utilizing structurally diverse, selective 5-HT_{2C} receptor ligands and the 2,5-dimethoxy-4-iodoamphetamine elicited head-twitch response model. *Neuropharmacology* 2013;70C:112-121.
55. Morgan D, Kondabolu K, Kuipers A, Sakhuja R, Robertson KL, Rowland NE, Booth RG. Molecular and behavioral pharmacology of two novel orally-active 5HT₂ modulators: potential utility as antipsychotic medications. *Neuropharmacology.* 2013;72:274-281.
56. Kasper J, Tikamdas R, Kim MS, MacFadyen K, Aramini R, Ladd J, Sarah Bisceglia S, Booth RG, Peris J. The serotonin-2 receptor modulator, (-)-trans-PAT, decreases voluntary ethanol consumption in rats. *European Journal of Pharmacology* 2013 718:98-104
57. Canal CE, Cordova-Sintjago T, Liu Y, Kim MS, Morgan D, Booth RG. Molecular pharmacology and ligand docking studies reveal a single amino acid difference between mouse and human serotonin 5-HT_{2A} receptors that impacts behavioral translation of novel phenylaminotetralin ligands. *Journal of Pharmacology and Experimental Therapeutics* 2013 347:705-716
58. Canal CE, Morgan D, Felsing D, Kondabolu K, Rowland NE, Robertson KL, Sakhuja R, Booth RG. A novel aminotetralin-type serotonin (5-HT)_{2C} receptor-specific agonist and 5-HT_{2A} competitive antagonist/5-HT_{2B} inverse agonist with preclinical efficacy for psychoses. *Journal of Pharmacology and Experimental Therapeutics* 2014 349:1-9.

59. Córdova-Sintjago T, Villa, N, Fang, L, Booth RG. Aromatic interactions impact ligand binding and function at serotonin 5-HT_{2C} G protein-coupled receptors: receptor homology modelling, ligand docking, and molecular dynamics results validated by experimental studies, *Molecular Physics* 2014 112: 398-407. PMID:24729635
60. Kasper J, Booth RG, Peris J. Serotonin-2C Receptor Agonists Decrease Potassium-Stimulated GABA Release In the Nucleus Accumbens. *Synapse* 2014 69:78-85. PMID: 25382408
61. Córdova-Sintjago T, Liu T, Booth RG. Molecular interactions of agonist and inverse agonist ligands at serotonin 5-HT_{2C} G protein-coupled receptors: computational ligand docking and molecular dynamics studies validated by experimental mutagenesis results. *Molecular Physics* 2015 113:348-358.
62. Sakhuja R, Kondabolu K, Córdova-Sintjago T, Travers S, Vincek AS, Kim MS, Abboud KA, Fang L, Sun Z, Canal CE, Booth RG. Novel 4-substituted-N,N-dimethyltetrahydronaphthalen-2-amines: synthesis, affinity, and in silico docking studies at serotonin 5-HT₂-type and histamine H₁ G protein-coupled receptors. *Bioorg Med Chem.* 2015; 23:1588-600. PMID: 25703249
PMCID: PMC4363177
63. Morgan D, Canal CE, Orza PC, Rose JL, Kim MS, Booth RG. A novel 5HT_{2C}-specific agonist/5HT_{2A-2B} antagonist attenuates psychomotor behaviors induced by methamphetamine, oxycodone, and their combination. *Drug and Alcohol Dependence* 146 2015; 146:e46.
64. Canal CE, Felsing DE, Liu Y, Zhu W, Wood JT, Perry CK, Vemula R, Booth RG. An Orally-Active Phenylaminotetralin-Chemotype Serotonin 5-HT₇ and 5-HT_{1A} Receptor Partial Agonist that Corrects Motor Stereotypy in Mouse Models. *ACS Chem Neurosci.* 2015; 6:1259-1270. PMID:26011730.
65. Liu Y, Canal CE, Cordova-Sintjago TC, Zhu W, Booth RG. Mutagenesis analysis reveals distinct amino acids of the human serotonin 5-HT_{2C} receptor underlie the pharmacology of distinct ligands. *ACS Chem Neurosci.* 2017; 8:28-39. PMID: 27580242.
66. Mongeau E, Yuan G, Minden Z, Waldron S, Booth RG, Felsing D, Ondrechen MJ, Jones GB. Homology Modeling Inspired Synthesis of 5-HT_{2A} Receptor Inhibitors: A Diazepine Analogue of the Atypical Antipsychotic JL13. *Cent Nerv Syst Agents Med Chem.* 2017 Apr 26. doi: 10.2174/1871524917666170426123607. [Epub ahead of print]

Book Chapters

67. Castagnoli, N., Trevor, A.J., Singer, T.P., Sparatore, A., Leung, L., Shinka, T., Wu., E.Y., and Booth, R.G. Metabolic studies on the nigrostriatal toxin MPTP. In *Progress in Catecholamine Research*, Alan R. Liss, Inc., New York, 93-100 (1988).
68. Neumeyer, J.L and Booth, R.G. Chapter 12: Neuroleptics and anxiolytic agents. In *Principles of Medicinal Chemistry 4th. Edition* (William O. Foye, ed.) Lea Febiger, Philadelphia, 199-231 (1996).
69. Neumeyer, J.L and Booth, R.G. Chapter 13: Drugs used to treat neuromuscular disorders. In *Principles of Medicinal Chemistry 4th. Edition* (William O. Foye, ed.) Lea Febiger, Philadelphia, 232-246 (1996).
70. Neumeyer, J.L and Booth, R.G. Chapter 42: Pesticides. In *Principles of Medicinal Chemistry 4th. Edition* (William O. Foye, ed.) Lea Febiger, Philadelphia, 908-926 (1996).

71. Nickell, W. Ward, H.E., and Booth, R.G. Antianxiety agents. In Burger's Medicinal Chemistry and Drug Discovery, 5th Edition, Volume 5 (Manfred E. Wolff, ed.) John Wiley and Sons, New York, 153-194 (1997).
72. Booth, R.G. and Neumeyer, J.L. Psychotherapeutic Drugs: Chapter 17: Antipsychotic and Anxiolytic Agents. In *Foye's Principals of Medicinal Chemistry 5th Edition* (Lemke et al., eds) Williams and Wilkins, Baltimore; pp. 408-434 (2002).
73. Booth, R.G. and Neumeyer, J.L. Chapter 20: Drugs used to treat neuromuscular disorders: Antiparkinsonian and Spasmolytic Agents. In *Foye's Principals of Medicinal Chemistry 5th Edition* (Lemke et al., eds) Williams and Wilkins, Baltimore; pp. 480-497 (2002).
74. Neumeyer, J.L. Baldessarini, R.J. Booth, R.G. Chapter 12, Therapeutic and diagnostic agents for Parkinson's disease. In: *Burgers' Medicinal Chemistry and Drug Discovery, Sixth Edition*, Donald J. Abraham (Ed), John Wiley and Sons, New York, pp 711-741, 2003.
75. Booth, R.G. Psychotherapeutic Drugs: Chapter 22: Antipsychotic and Anxiolytic Agents. In *Foye's Principals of Medicinal Chemistry 6th Edition* (Lemke et al., eds) Williams and Wilkins, Baltimore; pp. 601-630 (2007).
76. Booth, R.G. Chapter 25: Drugs used to treat neuromuscular disorders: Antiparkinsonian and Spasmolytic Agents. In *Foye's Principals of Medicinal Chemistry 6th Edition* (Lemke et al., eds) Lippincott Williams and Wilkins, Baltimore; pp. 679-697 (2007).
77. Booth R.G, Neumeyer, J.L. Baldessarini, R.J. Therapeutic and diagnostic agents for Parkinson's disease. Chapter 15 in Volume 9: Nervous System Therapeutics: In: *Burgers' Medicinal Chemistry and Drug Discovery, Seventh Edition*, Donald J. Abraham (Ed), John Wiley and Sons, New York, pp 529-568 (2010).
78. Booth, R.G. Chapter 13: Drugs used to treat neuromuscular disorders: Antiparkinsonian and Spasmolytic Agents. In *Foye's Principals of Medicinal Chemistry 7th Edition* (Lemke et al., eds) Lippincott Williams and Wilkins, Baltimore; pp. 419-447 (2013).
79. Booth, R.G. Chapter 14: Antipsychotic and Anxiolytic Agents. In *Foye's Principals of Medicinal Chemistry 7th Edition* (Lemke et al., eds) Williams and Wilkins, Baltimore; pp. 448-484 (2013).

Published International Patents Pending for New Chemical Entities and Therapeutic Use

PCT/US15/31523

"Serotonin Receptor Targeting Compounds and Methods"

Abstract: This invention relates to, in part, compositions and methods that are useful for the treatment of various diseases, including those linked to serotonin receptor binding, including, for example, neuropsychiatric diseases or disorders. Accordingly, the present invention provides for compositions and methods that agonize or antagonize one or more serotonin receptors and which find use in the treatment of various neuropsychiatric diseases or disorders including, without limitation, autism spectrum disorder (ASD) or associated symptoms.

Filed: May 19, 2015

Assignee: Northeastern University

Inventor: Raymond G Booth

Patent number: 9024071

"Therapeutic compounds"

Abstract: The invention relates to protein binding interacting/binding compounds and methods of identifying and using them. The invention further relates to pharmaceutical compositions and methods for treating serotonin and/or RSK disorders, including diseases and disorders mediated by GPCRs and/or RSKs.

Type: Grant

Filed: May 5, 2010

Issued: May 5, 2015

Assignee: University of Florida Research Foundation, Inc.

Inventor: Raymond G. Booth

Application number: 20140155490

“Therapeutic compounds”

Abstract: The invention relates to protein binding interacting/binding compounds and methods of identifying and using them. The invention further relates to pharmaceutical compositions and methods for treating serotonin disorders, including diseases and disorders mediated by GPCRs.

Type: Application

Filed: November 19, 2013

Issued: June 5, 2014

Assignee: University of Florida Research Foundation, Inc.

Inventor: Raymond G. Booth

Application number: 20120149693

“Therapeutic compounds”

Abstract: The invention relates to protein binding interacting/binding compounds and methods of identifying and using them. The invention further relates to pharmaceutical compositions and methods for treating serotonin and/or RSK disorders, including diseases and disorders mediated by GPCRs and/or RSKs.

Type: Application

Filed: May 5, 2010

Issued: June 14, 2012

Assignee: University of Florida Research Foundation, Inc.

Inventor: Raymond G. Booth

Application number: 20100305212

“Therapeutic compounds”

Abstract: The invention relates to protein binding interacting/binding compounds and methods of identifying and using them. The invention further relates to pharmaceutical compositions and methods for treating serotonin disorders, including diseases and disorders mediated by GPCRs.

Type: Application

Filed: June 13, 2008

Issued: December 2, 2010

Assignee: University of Florida Research Foundation, Inc.

Inventor: Raymond G. Booth

Therapeutic compounds and methods of use

Application number: 20100227934

Abstract: The invention relates to protein binding interacting/binding compounds and methods of identifying and using them. The invention further relates to pharmaceutical compositions and methods for treating serotonin disorders, including diseases and disorders mediated by GPCRs.

Type: Application

Filed: June 13, 2008

Issued: September 9, 2010

Assignee: University of Florida Research Foundation, Inc

Inventor: Raymond G. Booth