

## **CURRICULUM VITAE**

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### **EDUCATION**

- 1986-1987 Undergraduate, University of Akron, Akron, Ohio
- 1987-1991 Undergraduate, University of Toledo, Toledo, Ohio
- 1991 B.S., Pharmacy with Honors in Pharmacology, University of Toledo, Toledo, Ohio
- 1991-1996 Graduate Student, Pharmacology, Vanderbilt University
- 1996 Ph.D., Pharmacology, Vanderbilt University

### **PROFESSIONAL TRAINING AND EXPERIENCE**

- 1996-1998 Postdoctoral Fellow, Pharmacology, Vanderbilt University
- 1998-2003 Senior Fellow, Pharmacology, University of Washington
- 2003-2004 Acting Instructor, Pharmacology, University of Washington
- 2004-2009 Research Assistant Professor, Pharmacology, Vanderbilt University  
Director, Molecular Pharmacology, Vanderbilt Program in Drug Discovery
- 2009-present Research Associate Professor, Pharmacology, Vanderbilt University  
Director, Molecular Pharmacology, Vanderbilt Center for Neuroscience Drug Discovery

## **AWARDS AND HONORS**

- 1991 Valedictorian, College of Pharmacy
- 1991 Phi Kappa Phi Graduate Fellowship
- 1991 AAPS-AFPE Gateway Scholarship
- 1991 The Merck-Sharp and Dohme Award in Medicinal Chemistry
- 1991 The Upjohn Award in Pharmacology
- 1991 The SmithKline Beechem Award in Clinical Pharmacy
- 1993 First Place, Vanderbilt University Graduate Student Research Day
- 1994 First Place, Vanderbilt University Graduate Student Research Day
- 1995 Grass Foundation Fellowship to attend “Neurobiology of Human Neurological Disease: Mechanisms of Neurodegeneration”, Cold Spring Harbor Laboratory
- 2000 Travel award for abstract submitted to “Obesity and the Regulation of Energy Homeostasis”, Keystone Symposium, Taos, NM

## **PROFESSIONAL MEMBERSHIPS**

- 1991-present Phi Kappa Phi
- 1991-1998,  
2006-present Society for Neuroscience
- 2007-present American Society for Pharmacology and Experimental Therapeutics

## **RESEARCH SUPPORT-CURRENT**

**1R01 MH108498-01 (Niswender, CM/Lindsley CW) 12/10/2015-11/30/2018 25% effort**  
**NIMH \$1,498,134, Total Project**

**Development of an mGlu<sub>2/4</sub> heterodimer-selective allosteric modulator.** We propose that heterodimerization may underlie a divergence in *in vivo* pharmacological profiles that we have observed with various mGlu<sub>4</sub> positive allosteric modulators that either do or do not potentiate the activity of mGlu<sub>2/4</sub> heterodimers. We will chemically optimize small molecules with selectivity for mGlu<sub>2/4</sub> heteromers over mGlu<sub>2</sub> or mGlu<sub>4</sub> homomeric receptors.

**1R01MH110389-01 (Niswender) 07/01/2016-06/30/209 15% effort**  
**NIMH \$1,714,670 Total Project**

**Optimization of a Metabotropic Glutamate Receptor 7 Positive Allosteric Modulator.**

We propose to perform a chemical optimization campaign to develop mGlu<sub>7</sub> PAMs with enhanced specificity and pharmacokinetic properties and to use deficits in Mecp2-deficient mice to profile native tissue and *in vivo* activity of optimized compounds, with the goal of developing a highly selective mGlu<sub>7</sub> PAM to support continued exploration of the biology and therapeutic potential of this important receptor.

**PR160102 (Niswender, CM) 08/01/2017-07/31/2020 15% effort**  
**Department of Defense \$1,875,949, Total Project**

**The role of metabotropic glutamate receptor 7 in the etiology and treatment of Rett syndrome.** This grant will examine the role of mGlu<sub>7</sub> in Rett syndrome by profiling mGlu<sub>7</sub> knockout mice, crossing mice with various mGlu<sub>7</sub> expression levels with Mecp2-deficient animals, and developing new mGlu<sub>7</sub> positive allosteric modulator probes.

**3503 rettsyndrome.org (Niswender, CM) 04/01/2017-03/31/2019 10% effort**  
**Rettsyndrome.org \$150,000, Total Project**

**Exploration of metabotropic glutamate receptor 3 as a target for MeCP2-related disorders.** This grant proposes to determine if mGlu<sub>3</sub> is a new target that can be modulated with small molecules for therapeutic benefit in models of MeCP2-related disorders.

**8766 Autism Speaks (Niswender, CM) 03/01/2014-08/28/2017 15% effort**  
**Autism Speaks in NCE \$449,996, Total Project**

**Temporal divergence of hypoconnectivity and excitotoxicity in Rett Syndrome.**

The goals of these studies are to build upon our preliminary data showing that mGlu<sub>5</sub> PAMs reduce symptom severity and phenotypes in RS mice when administered in advanced disease states. We will also assess the liability for PAMs to elicit adverse consequences when administered early in disease due to temporal changes in glutamatergic transmission. Furthermore, we will determine whether mGlu<sub>5</sub> NAMs can temper excitotoxicity and prevent synapse loss when administered to young RS mice, thereby reducing the impact of developmental regression on RS phenotypes.

## **PENDING**

**rettsyndrome.org (Niswender, CM)**      **01/01/2018-12/31/2019**  
**Rettsyndrome.org**                                **\$150,000, Total Project**  
Tailoring gene replacement therapy for MECP2-related disorders

## **COMPLETED**

### **Predoctoral/Postdoctoral Awards**

1994-1996	Pharmaceutical Research and Manufacturers of America Foundation Predoctoral Award in Pharmacology
1997-1998	Pharmaceutical Research and Manufacturers of America Foundation Postdoctoral Award in Pharmacology
1999-2000	Fellow, Neurobiology and Behavior Training Grant, University of Washington
2000-2003	Fellow, National Service Research Award, NIDDK
03/03/03-02/28/04	University of Washington Diabetes and Endocrinology Research Center Pilot and Feasibility Award

### **1 R03 MH076398-01 (Niswender, C.M.) 08/1/05-07/31/06**

**NIH/NIMH**                                        \$3,000 plus access to screening  
Measurement of GPCR-mediated thallium flux through GIRK. Access grant to Molecular Libraries Screening Center Network.

### **1R21NS053536-01/1R21NS053536-01S1 (Niswender, C.M.) 09/30/05-02/28/08**

**NIH/NIMH**                                        Direct Costs: \$125,000 plus \$25,000 supplement.  
A Direct Assay for HTS of Gi/o-linked GPCRs: mGluR7 as the Prototype.

### **1 X01 MH077607-01 (Niswender, C.M.) 02/01/06-01/31/07**

Discovery of novel allosteric agonists of the M4 muscarinic receptor. Access grant to Molecular Libraries Screening Center Network.

**1R01 (Niswender, C.M.)**                            **2/01/04-11/30/09**  
**NIH/NINDS**                                        **1,165,000 direct costs current funding period**  
Metabotropic Glutamate Receptors in Basal Ganglia. Grant transferred from P. Jeffrey Conn in final year of funding.

**LEAPS Award (Conn, P.J.)**                            **11/01/07 – 10/30/11**  
**Michael J. Fox Foundation**                            **\$3,609,000 Direct Costs**  
Discovery of novel allosteric modulators of mGluR4 for treatment of Parkinson's disease.  
Biology Project Team Leader

**Basic Research Award (Niswender, C.M.) 01/01/12-12/31/13**

**International Rett Syndrome Foundation. \$100,000, Total Project**

Metabotropic glutamate receptor 7: a novel therapeutic candidate for Rett Syndrome. This grant examined the regulation of mGlu<sub>7</sub> levels in the context of MeCP2 loss and provided the basis for new grants to develop novel allosteric modulators for MeCP2-based disorders.

**R21 NS078262 (Niswender, C.M.) 04/01/12-03/13/14**

**NIH/NINDS \$420,809, Total Project**

Metabotropic glutamate receptors in the basal ganglia. The goals of this grant were to test the hypothesis that metabotropic glutamate receptor 8 is a novel target in Parkinson's disease. The grant involved a combination of molecular pharmacology, electrophysiology, and in vivo behavioral profiling.

**1UH2 NS099066-01 (Niswender, CM) 09/01/2016-09/29/2021**

**NIH \$124,768 Total Project**

**Development of VU0652957 for the treatment of Parkinson's disease (PD)**

We propose to utilize the Blueprint Neurotherapeutics Network (BPN) to advance VU0652957 through Investigational New Drug (IND)-enabling studies in preparation for clinical studies in PD patients. This is a development grant to advance an mGlu<sub>4</sub> positive allosteric modulator into clinical development for Parkinson's disease. *Project ended due to lack of exposure required for efficacy in a nonhuman primate model of Parkinson's disease.*

**5R21 MH102548-02 (Niswender, CM) 08/01/2014-07/31/2017**

**NIMH \$431,476, Total Project**

**Metabotropic Glutamate Receptor Regulation in MeCP2-Related Disorders.**

Based on dramatic decreases in mGlu<sub>7</sub> levels seen in mice that model Rett syndrome, we propose to test the hypothesis that mGlu<sub>7</sub> levels are conversely increased in an *MECP2* Duplication mouse model, further substantiating a role for MeCP2 in mGlu<sub>7</sub> gene regulation. Additionally, we will test the hypotheses that modulation of mGlu<sub>7</sub> activity using pharmacological tools will normalize synaptic plasticity and behavioral deficits in mice under- and overexpressing MeCP2.

## **INTRAMURAL AND EXTRAMURAL ACTIVITIES**

### **2006**

- Grant reviewer, Melanoma Research Foundation
- Manuscript reviewer, Journal of Neurochemistry
- Paper discussion leader for Interdisciplinary Graduate Program Core Course. Kim et al., 2001,  
JBC. The role of phosphorylation in D1 dopamine receptor desensitization.

### **2007**

- Manuscript reviewer, Molecular Pharmacology
- Lecturer in Neuroscience 345 Course. Neuromodulation and metabotropic glutamate receptors.
- Paper discussion leader in Neuroscience 345 Course. Sansig et al., 2001, J. Neuroscience.  
Increased seizure susceptibility in mice lacking metabotropic glutamate receptor 7.
- Paper discussion leader for Interdisciplinary Graduate Program Core Course. Ross et al., 1978  
JBC. Reconstitution of hormone-sensitive adenylate cyclase activity with resolved components of the enzyme.

### **2008**

- Manuscript reviewer, Neuropharmacology
- Paper discussion leader for Interdisciplinary Graduate Program Core Course. Kim et al., 2001,  
JBC. The role of phosphorylation in D1 dopamine receptor desensitization.
- Lecturer in Neuroscience 345 Course. Neuromodulation and metabotropic glutamate receptors.
- Paper discussion leader in Neuroscience 345 Course. Volk et al., 2007, J. Neuroscience.  
Multiple Gq coupled receptors converge on a common protein synthesis-dependent long term-depression that is affected in fragile X mental retardation.

### **2009**

- Grant reviewer, Michael J. Fox Foundation
- Manuscript reviewer, Molecular Pharmacology
- Manuscript reviewer, ACS Chemical Neuroscience
- Manuscript reviewer, Journal of Neuroscience
- Manuscript reviewer, J Pharmacol Exp Therapeutics
- Manuscript reviewer, British Journal of Pharmacology

### **2010**

- Manuscript reviewer, British Journal of Pharmacology
- Manuscript reviewer, Molecular Pharmacology
- Manuscript reviewer, Neuropharmacology
- Manuscript reviewer, Neuroscience

### **2011**

- Manuscript reviewer, Journal of Biological Chemistry

Manuscript reviewer, The International Journal of Neuropsychopharmacology  
Manuscript reviewer, Nature Protocols  
Manuscript reviewer, Proceedings of the National Academy of Sciences

2012

Grant reviewer, Multiple Sclerosis Fast Forward Foundation  
Manuscript reviewer, Expert Opinion on Drug Discovery  
Manuscript reviewer, Neuropharmacology  
Manuscript reviewer, Pharmacology, Biochemistry, and Behavior  
Manuscript reviewer, Molecular Pharmacology  
Manuscript reviewer, ACS Chemical Neuroscience  
Guest lecturer, Drug Discovery Course. “mGlu<sub>4</sub> positive allosteric modulators for the treatment of CNS disorders”. Vanderbilt University, October 17, 2012.

2013

-Manuscript reviewer, Journal of Neurochemistry  
-Manuscript reviewer, Neuropharmacology  
-Grant reviewer, Michael J. Fox Foundation  
-Grant reviewer, Pilot and Feasibility Awards program, Vanderbilt University for Obesity and Metabolism  
-Grant reviewer, NIH EUREKA program  
-Special Guest Editor, Neuropharmacology, “The Synaptic Basis of Neurodegeneration”  
-Manuscript reviewer, Journal of Neurochemistry  
-Guest lecturer, Neuroscience 235 Course. “Drug Discovery in Academia: mGlu<sub>4</sub> positive allosteric modulators for the treatment of CNS disorders”. Vanderbilt University, March 14, 2013.  
-Guest lecturer, Drug Discovery Course. “mGlu<sub>4</sub> positive allosteric modulators for the treatment of CNS disorders”. Vanderbilt University, November 6, 2013.  
-Co-organizer, Academic Drug Discovery Conference, Nashville TN, October 9-11, 2013.

2014

-Manuscript reviewer, Molecular Pharmacology  
-Manuscript reviewer, Neuropharmacology  
-Manuscript reviewer, Journal of Pharmacology and Experimental Therapeutics  
-Grant reviewer, Trailblazer Award, Autism Speaks  
-Guest lecturer, Neuroscience 235 Course. “Drug Discovery in Academia: mGlu<sub>4</sub> positive allosteric modulators for the treatment of CNS disorders”. Vanderbilt University, March 11, 2014.  
-Grant reviewer, IWT  
-Grant reviews, Meixner Postdoctoral Fellowship Program, Autism Speaks  
-Guest lecturer, Drug Discovery Course. “mGlu<sub>4</sub> positive allosteric modulators for the treatment of CNS disorders”. Vanderbilt University, October, 2014.

2015

-Manuscript reviewer, Neuropharmacology

- Guest lecturer, Neuroscience 235 Course. "Drug Discovery in Academia: mGlu4 positive allosteric modulators for the treatment of CNS disorders". Vanderbilt University, February 24, 2015.
- Grant reviewer, Meixner Postdoctoral Fellowship Program, Autism Speaks
- Guest lecturer, Drug Discovery Course. "mGlu<sub>4</sub> positive allosteric modulators for the treatment of CNS disorders". Vanderbilt University, October, 2015.

2016

- Manuscript reviewer, Neuropharmacology
- Manuscript reviewer, Frontiers in Neural Circuits
- Guest lecturer, Neuroscience 235 Course. "Drug Discovery in Academia: mGlu4 positive allosteric modulators for the treatment of CNS disorders". Vanderbilt University, February 24, 2015.
- Guest lecturer, Modern Drug Discovery, "mGlu4 positive allosteric modulators for the
- Grant reviewer, Meixner Postdoctoral Fellowship Program, Autism Speaks
- Member, VICTR "Drug Repurposing Think Tank"

2017

- Manuscript reviewer, Neuropharmacology
- Introduction to the Vanderbilt Center for Neuroscience Drug Discovery: Case Study in Rett Syndrome, MSTP Second Look Day, Vanderbilt University, April 7, 2017.
- Co-chair of Autism Spectrum Disorders Speaking Session, 9<sup>th</sup> International Meeting on Metabotropic Glutamate Receptors

## **STUDENTS/POSTDOCS DIRECTLY SUPERVISED/THESIS COMMITTEES**

- |                      |  |
|----------------------|--|
| Caroline Kim         | VUMC Emphasis Program for Medical Students, 2007-2008.<br>‘Investigation of type III metabotropic glutamate receptor potentiators using a novel HTS assay using thallium flux through GIRK channels’. Current position, General Surgeon at Houston Methodist, San Jacinto.                                 |
| Alexander Kane       | Vanderbilt University undergraduate. “The <i>in vivo</i> characterization of TBPB, a novel allosteric agonist of M1 muscarinic receptors: Implications for the role of M1 muscarinic receptors in treatment of schizophrenia”. Honors thesis, May, 2008.   |
| Shen Yin             | Vanderbilt University pharmacology graduate student. Co-mentor, graduated September, 2013. “Allosteric modulation of metabotropic glutamate receptors”. Current position, Associate Clinical Scientist, Genentech.   |
| Olivia Veatch        | Vanderbilt University genetics graduate student. Thesis committee member, graduated August, 2013. “Identifying biological pathways implicated in defined subgroups of phenotypic expression for autism spectrum disorders.” Current position, Research Instructor in Neurology, Vanderbilt Kennedy Center. |
| Frank Byers          | Vanderbilt University Masters in Laboratory Investigation student, thesis committee, graduated May 2014.   |
| Rebecca Klar Senter  | Vanderbilt University pharmacology graduate student, Co-mentor. Awarded Weatherstone predoctoral fellowship from Autism Speaks 7/1/2014, graduation 7/14/2015. Current position, Scientist, Flexion Therapeutics.  |
| Rachel Crouch        | Vanderbilt University pharmacology graduate student, co-chair of thesis committee. Graduation 11/2016. Current position, Postdoctoral fellow, VCNDD.   |
| Thomas Utley         | Vanderbilt University pharmacology postdoc. Current position, Licensing Agent, Vanderbilt Technology Transfer office.  |
| Julie Roper-Field    | Vanderbilt University pharmacology postdoc. Current position, Project Manager, VICTR and CTSA Consortium Coordinating Center, Vanderbilt University Medical Center.  |
| Nidhi Jalan-Sakrikar | Vanderbilt University pharmacology postdoc (2012-2014), Current position, postdoctoral fellow at Mayo Clinic.  |

Rocco Gagliotti	Vanderbilt University pharmacology postdoc, joined lab in 2012, awarded postdoctoral fellowship from International Rett Syndrome Foundation, 2014. Awarded BBRF fellowship in 2016 (current).
Nicole Fisher	Vanderbilt University Interdisciplinary Graduate Student, joined lab Spring of 2015, awarded Pharmacology Training grant slot, 2015 (current).
Branden Stansley	Vanderbilt University Pharmacology Postdoctoral Fellow, joined lab Spring of 2015, awarded T32 postdoctoral training award and an F32 NRSA, June 2016 (current).
Sheryl Vermudez	Vanderbilt University Interdisciplinary Graduate Student, joined lab Spring of 2017 (current).
Annah Moore	Vanderbilt University Interdisciplinary Graduate Student, joined Sweatt Lab in Spring of 2017, co-mentor (current).
Christian Marks	Vanderbilt University Molecular Physiology and Biophysics Graduate Student, thesis committee (current).
Sean Moran	Vanderbilt Neuroscience Graduate Student, thesis committee (current).
Nick Harris	Vanderbilt University Molecular Physiology and Biophysics MD/PhD Student, thesis committee (current).

## **PRESENTATIONS**

### 1998

-Identification and Functional Characterization of Edited Human Serotonin 2c Receptor Isoforms. 4<sup>th</sup> IUPHAR Satellite Meeting on Serotonin. October 1998.

### 2006

-High Throughput Screening and Medicinal Chemistry at Vanderbilt in Support of Drug Discovery for Muscarinic Receptor Ligands. Vanderbilt Institute for Chemical Biology Retreat.

-Allosteric modulation of metabotropic glutamate receptor 5, M1, and M4 muscarinic receptors: potential therapeutic directions for schizophrenia. 27<sup>th</sup> Annual Meeting of the Southeastern Pharmacology Society. November 2006.

### 2007

-High Throughput Screening and Medicinal Chemistry at Vanderbilt: The search for Allosteric Ligands of the M1 and M4 Muscarinic Receptors. HTS Users Group Meeting, Vanderbilt University. March, 2007

### 2008

-Allosteric modulation of mGluR4 as a novel therapeutic direction for the treatment of Parkinson's disease. American Society for Biochemistry and Molecular Biology, Experimental Biology, San Diego, CA. April, 2008.

-Allosteric modulation of GCPs as a novel therapeutic direction for the treatment of CNS disorders. American Society of Pharmacology and Experimental Therapeutics, Experimental Biology, San Diego, CA, April. 2008.

-Allosteric modulation of mGluR4 as a novel therapeutic direction for the treatment of Parkinson's disease. Keystone Symposia; G Protein Coupled Receptors: New Insights in Functional Regulation and Clinical Application, Killarney, Ireland. May, 2008.

-Allosteric modulation of mGluR4: a novel therapeutic direction for the treatment of Parkinson's disease. 6<sup>th</sup> International Meeting on Metabotropic Glutamate Receptors, Taormina, Sicily, Italy. September, 2008.

### 2009

-The development of positive allosteric modulators of mGluR4 for the treatment of Parkinson's disease. WFN World Congress on Parkinson's Disease and Related Disorders. Miami, FL. December, 2009.

## 2010

-The development of positive allosteric modulators of mGluR4 for the treatment of Parkinson's disease—Parkinson's Disease Case Study. 4th Annual Drug Discovery for Neurodegeneration Conference. Houston, TX. February, 2010.

-HTS results from academia: discovery of allosteric modulators of mGluR4 and 5. Metabotropic Glutamate Receptors: Translation from Discovery to Clinical Trials. The New York Academy of Sciences. New York, NY. February, 2010.

-Functionally selective and context dependent pharmacology of GPCR allosteric modulators. Pharmacology drive assays for GPCRs and ion channels. Ninth Annual World Pharmaceutical Congress. Philadelphia, PA, June, 2010.

-Novel ion channel-based assays: detecting and characterizing 7TM receptor modulators. Pharmacology drive assays for GPCRs and ion channels. Ninth Annual World Pharmaceutical Congress. Philadelphia, PA, June, 2010.

-The Development of Positive Allosteric Modulators of Metabotropic Glutamate Receptor 4. Allosteric Modulator Drug Discovery Congress, San Diego, CA, November, 2010.

## 2011

-mGlu4 receptor positive allosteric modulator development for the treatment of CNS disorders. 7<sup>th</sup> International Meeting on Metabotropic Glutamate Receptors, Taormina, Italy. October, 2011.

## 2012

-mGlu4 positive allosteric modulators for the treatment of CNS disorders. 45<sup>th</sup> Annual Winter Conference on Brain Research. Snowbird, Utah, January.

## 2013

-mGlu4 positive allosteric modulators for the treatment of CNS disorders. Guest presentation to Vanderbilt Parkinson's Disease Advisory Board, Vanderbilt University, April 19, 2013.

-Metabotropic glutamate receptor 7: a novel therapeutic target for MeCP2-related disorders. Vanderbilt University Rett Syndrome Symposium, October 16, 2013.

## 2014

-Metabotropic glutamate receptor potentiation as a therapeutic direction in Rett syndrome. Invited talk, International Rett Syndrome Foundation meeting, June 25, 2014.

-Metabotropic glutamate receptor 7 (mGlu<sub>7</sub>): a novel target for the treatment of Rett syndrome. Invited talk, 8<sup>th</sup> International Meeting on Metabotropic Glutamate Receptors, October 2, 2014.

-Metabotropic glutamate receptor 7 (mGlu<sub>7</sub>): a novel target for the treatment of Rett syndrome. Invited talk, Multimodal Interventions in IDD from Drug Discovery to Clinical Trials, Vanderbilt Kennedy Center, October 22, 2014.

2015

-Metabotropic glutamate receptor 7: a new therapeutic target for both Rett and *MECP2* Duplication syndromes. Invited talk, MECP2 Duplication Conference, Houston TX, Sept 2015.

2016

-Metabotropic Glutamate Receptor 4 Positive Allosteric Modulators for Parkinson's Disease: Impact of Receptor Heterodimerization. CNS Diseases World Summit. Boston, MA, Sept 12, 2016.

2017

-Therapeutic Potential of mGlu7 in Neurodevelopmental Disorders. Invited Speaker, 9<sup>th</sup> International Meeting on Metabotropic Glutamate receptors. Taormina, Italy, October 2017.

## **ISSUED PATENTS**

1. Benzamide mGluR5 Positive Allosteric Modulators and Methods of Making and Using Same. Patent 8,853,392. Filed 6/3/2008. Issued patent, USA.
2. Benzamide mGluR5 Positive Allosteric Modulators and Methods of Making and Using Same. Patent 5622568. Filed 12/2/2009. Issued patent, Japan.
3. Substituted Dioxopiperidines and Dioxopyrrolidines as mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction. Patent 8,759,377. Filed 11/23/2010. Issued patent, USA.
4. Alkyl-3-((2-Amidoethyl)Amino)-8-Azabicyclo[3.2.1]Octane-8-Carboxylate Analogs as M1 Allosteric Agonists and Methods of Making and Using Same. Patent 8,697,691. Filed 12/21/2010. Issued patent, USA.
5. Substituted 1,1,3,1-Tetraoxidobenzo{D} {1,3,2}Dithiazoles as mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction. Patent 8,658,650. Filed 1/28/2011. Issued patent, USA.
6. Pyrazolopyridine, Pyrazolopyrazine, Pyrazolopyrimidine, Pyrazolothiophene and Pyrazolothiazole Compounds as mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction. Patent zl 201180017221.4. Filed 2/11/2011. Issued patent, China.
7. Pyrazolopyridine, Pyrazolopyrazine, Pyrazolopyrimidine, Pyrazolothiophene and Pyrazolothiazole Compounds as mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction. Patent 2012/06416. Filed 2/11/2011. Issued patent, South Africa.
8. Pyrazolopyridine, Pyrazolopyrazine, Pyrazolopyrimidine, Pyrazolothiophene and Pyrazolothiazole Compounds as mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction. Patent 9,163,015. Filed 2/11/2011. Issued patent, USA.
9. Pyrazolopyridine, Pyrazolopyrazine, Pyrazolopyrimidine, Pyrazolothiophene and Pyrazolothiazole Compounds as mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction. Patent 2011215638. Filed 2/11/2011. Issued patent, Australia.
10. Pyrazolopyridine, Pyrazolopyrazine, Pyrazolothiophene and Pyrazolothiazole Compounds as mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction. Patent 9,108,963. Filed 8/13/2012. Issued patent, USA.
11. mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction. Filed 4/23/2012. Patent 9,180,192. Issued patent, USA.

12. mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction. Patent 8,779,157. Filed 6/12/2012. Issued patent, USA.
13. Aryl and Heteroaryl Sulfones as mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction. Patent 8,912,336. Filed 6/12/2012. Issued patent, USA.
14. Benzisoxazoles and Azabenzisoxazoles as mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction, Patent 8,916,584, Filed 8/13/2012. Issued patent, USA.
15. Heterocyclic Sulfone mGluR4 Allosteric Potentiators, Compositions, and Methods of Treating Neurological Dysfunction. Patent 9,192,603, Filed 11/13/2012. Issued patent, USA.

## **ABSTRACTS**

1. **Burns, CM**, Obermiller PS, Emeson RB. RNA editing of AMPA receptor subunit (GluR-B) mRNA. , Society for Neuroscience Abstracts, 1993.
2. **Burns CM**, Rueter SM, Emeson RB. Molecular mechanisms mediating the editing of glutamate receptor (GluR-B) RNA transcripts. Albany Conference, RNA Editing, 1994.
3. Rueter, SM, **Burns CM** and Emeson RB. *In vitro* editing of glutamate receptor (GluR-B) RNA transcripts. RNA Processing: Cold Spring Harbor Laboratory, 1995.
4. **Burns CM**, Chu H, Rueter SM, Sanders-Bush E, Emeson RB. Identification and characterization of RNA editing within the serotonin 2C receptor. Society for Neuroscience Abstracts, 1996.
5. Chu H, **Burns C**, Canton H, Emeson RB, Sanders-Bush E. Functional characterization of rat serotonin 5-HT<sub>2C</sub> receptor editing isoforms. Society for Neuroscience Abstracts, 1996.
6. Rueter SM, **Burns CM**, Emeson RB. Characterization of AMPA receptor subunit (GluR-B) RNA editing in rat brain nuclear extracts. Society for Neuroscience Abstracts, 1996.
7. **Burns CM**, Chu H, Rueter SM, Sanders-Bush E, Emeson RB. RNA editing of transcripts encoding the serotonin 2C receptor. EMBO workshop on RNA Editing, 1996.
8. **Burns CM**, Chu H, Rueter SM, Hutchinson LK, Canton H, Sanders-Bush E, Emeson RB. RNA editing of serotonin 2C receptor transcripts. RNA Society Meeting, 1997.
9. **Niswender CM**, Copeland S, Emeson RB and Sanders-Bush E. Identification and Functional Characterization of Edited Human Serotonin 2c Receptor Isoforms. 4<sup>th</sup> IUPHAR Satellite Meeting on Serotonin, 1998.
10. **Niswender CM**, Copeland SC, Dilley G., Meltzer HY, Overholser JC, Stockmeier CA, Emeson RB, Sanders-Bush E. RNA editing of human serotonin 2C receptor transcripts. Society for Neuroscience Abstracts, 1998.
11. Berg KA, Cropper JD, **Niswender CM**, Sanders-Bush E, Emeson RB Clarke WP. Activation of PLC and PLA<sub>2</sub> by h5-HT<sub>2C</sub> receptor RNA-edited isoforms. Society for Neuroscience Abstracts, 1999.
12. **Niswender CM**, McKnight GS. Cre recombinase-mediated expression of a constitutively active form of protein kinase A. Keystone Conference, Obesity and the Regulation of Energy Homeostasis, Taos, NM, 2001.
13. Willis BS, **Niswender CM**, McKnight GS. Cre recombinase-mediated expression of dominant-negative and constitutively active protein kinase A subunits. Society for Neuroscience Abstracts, 2001.

14. **Niswender CM**, Willis BS, Sweet IR, Wallen A, Thompson B, Wu C, Lange AJ, McKnight GS. Expression of a constitutively active PKA holoenzyme in the liver results in glucose intolerance and impaired glucose-stimulated insulin secretion in mice. Keystone Conference, Toward Understanding Islet Biology, Keystone, CO, 2003.
15. **Niswender CM**, Myers KA, Banko JL, Rodriguez AL, Edl J, Zhang Y, Shirey JK, Saleh SA, Weaver CD, Conn PJ. Identification of novel allosteric modulators of group III mGluRs: New tools for the study of synaptic transmission. 5<sup>th</sup> Meeting on Metabotropic Glutamate Receptors, Taormina, Italy, 2005.
16. Edl J, Rodriguez AL, Tamagnan G, Alagille D, Johnson RL, **Niswender C**, Myers K, Conn PJ. Characterization of structural analogs of a Group III mGluR agonist and mGluR4/5 allosteric modulators. Society for Neuroscience Abstracts, 2005.
17. Myers KA, **Niswender CM**, Williams R, Edl J, Saleh S, Jones CK, Weaver CD, Orton D, Conn PJ. Characterization of novel allosteric antagonists of metabotropic glutamate receptor subtype 7. Society for Neuroscience Abstracts, 2006.
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\*\*\*Best poster winner at respective conference

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