

# CURRICULUM VITAE

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## 1. EDUCATION

<u>Year</u>	<u>Degree</u>	<u>Institution</u>
1990-1993	B.Sc. (Chemistry)	University of Mumbai, India
1993-1995	M.Sc. (Chemistry)	Indian Institute of Technology (IIT), Mumbai, India
1995-2000	Ph.D. (Organic Chemistry)	Institute of Chemical Technology, Mumbai, India.

## 2. POSTDOCTORAL TRAINING

07/2000- 08/2003 Postdoctoral Fellow, Department of Pharmaceutical Sciences, School of Pharmacy, University of Connecticut, Storrs, CT.  
Supervisor: Dr. Alexandros Makriyannis

## 3. UNIVERSITY APPOINTMENTS

09/2003-10/2004 Assistant Research Professor, Department of Pharmaceutical Sciences, School of Pharmacy, University of Connecticut, Storrs, CT.  
11/2004-06/2005 Senior Research Scientist, Center for Drug Discovery, NEU, Boston, MA, USA  
07/2005- 08/2010 Assistant Research Professor, Center for Drug Discovery, NEU, Boston, MA, USA.  
09/2010- 06/2016 Assistant Professor (Tenure-track), Department of Pharmaceutical Sciences, Northeastern University, Boston, MA, USA.  
07/2016 onwards- Associate Professor (Tenured), Department of Pharmaceutical Sciences, Northeastern University, Boston, MA, USA.

## 4. AWARDS AND HONORS

1995 Recipient of Fellowship of University Grant Commission (UGC) as a Junior Research Fellow  
1998 Recipient of Fellowship of University Grant Commission (UGC) as a Senior Research Fellow  
1998 Recipient of "Best Paper Presentation Award" in XIV<sup>th</sup> Carbohydrate Conference Conducted by Indian Institute of Technology (IIT) in collaboration with Association of Carbohydrate Chemists & Technologists, India.  
2009 Early Career Development Award in Chemistry of Drug Abuse and Addiction (ECHEM) from NIDA.  
2011 Excellence in Teaching Award, School of Pharmacy, Northeastern University, Boston, MA 02115  
2014 The Schumacher Award for Excellence in Research, Northeastern University, Boston MA 02115  
2014 NIH Study Section, Drug Discovery for the Nervous System, Ad Hoc reviewer, June 2014  
2014 NIH Study Section, Special Emphasis Panel, Molecular Probes, Ad Hoc Reviewer, June 2014  
2014 NIH Study Section, Drug Discovery for the Nervous System, Ad Hoc reviewer, October 2014  
2014 NIH Study Section, Special Emphasis Panel, Molecular Probes, Ad Hoc Reviewer, October 2014  
2015 RISE (Research, Innovation, Scholarship and Expo) Award in Health Sciences, NEU, Boston, 2015  
2015 Nominated for Northeastern University's University-wide "Excellence in Teaching Award"  
2015 Nominated for "Bouve College Distinguished Educator Award" for year 2015.  
2015 Guest Speaker, Rho Chi Society's Annual Lecture Series, 2015.  
2016 RRD8 Study Section, Career Development Award Review Committee, US Department of Veteran Affairs, Ad Hoc Reviewer, Feb.2016

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**SCOPUS ANALYSIS:** (as of 06/25/2016): **H-index = 19**; Total Citation = **1220**; Total publications = **58**;  
Total patents = **6**.

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#### **4. IN NEWS/MEDIA:**

- 1) “ Last Dance with Mary Jane” <http://www.northeastern.edu/news/2014/03/thakur/>
- 2) “ A Promising Alternative to Medical Marijuana:  
<http://www.northeastern.edu/news/2014/09/promising-alternative-medical-marijuana/>
- 3) “Smarten Up” <http://www.northeastern.edu/news/2014/03/smarten-up/>
- 4) “Local researchers work to eliminate high from medical marijuana”  
<http://boston.cbslocal.com/tag/ganesh-thakur/>
- 5) “Hope the prescription can mimic marijuana’s benefits”  
<https://www.bostonglobe.com/lifestyle/health-wellness/2014/11/17/some-parents-children-who-have-seizures-are-hoping-that-prescription-drug-will-able-mimic-marijuana-benefits/KoGizuhG99EO3w0RkyOrcO/story.html>

#### **5. MAJOR COMMITTEE ASSIGNMENTS**

##### **Department of Pharmaceutical Sciences, College of Pharmacy, NEU**

- 2011- 2013 Member, Faculty Research Development and Mentoring.  
2012- Present Scholarship and Award Committee.  
2012 John Neumeyer Award Committee.  
2013-Present Assessment Committee.  
2013-Present Faculty Search Committee/Bouve College.

##### **NIH**

- 2014 Drug Discovery for the Nervous System Study Section (DDNS), *Ad hoc* Reviewer  
2014 NIH Study Section, Special Emphasis Panel, Molecular Probes, *Ad Hoc* Reviewer

##### **Reviewer**

- 1) Chemical Biology and Drug Design
- 2) Expert Opinion on Therapeutic Patents
- 3) Bio-organic Medicinal Chemistry
- 4) ChemBioChem (Wiley VCH-de)
- 5) Recent Patents on CNS Drug Discovery
- 6) Life Sciences
- 7) Journal of Pharmacology and Experimental Therapeutics (JPET)
- 8) Neuropsychopharmacology, Nature Publishing Group (NGP)
- 9) Bio-organic Medicinal Chemistry Letters
- 10) Drugs of the Future
- 11) ACS Chemical Neuroscience

#### **6. MEMBERSHIP PROFESSIONAL SOCIETIES**

- 1999-current Association of Carbohydrate Chemists and Technologists  
2000-current American Chemical Society  
2003-current International Cannabinoid Research Society (ICRS)  
2008-current Member of NIDA Networking Project (NNP)  
2010-current Member of American Association of Pharmaceutical Scientists (AAPS)  
2011-current Member of Rho Chi Honor Society  
2015-current Member of Society of Neuroscience

## **7. RESEARCH GRANTS**

1. **1R01 EY024717-01/NIH-NEI**  
Title: A Novel Pharmacotherapy for Glaucoma  
Role: PI  
Date: 09/01/2014- 08/30/2019
2. **1R01CA206028-01/NIH/NCI**  
Title: Mitigation of Chemotherapy Induced Peripheral Neuropathy  
Role: PI on the subcontract with VCU  
Date: 04/15/2016- 04/14/2017
3. **1R43MH103936-01 NIH/NIMH**  
Title: Novel treatment of posttraumatic stress disorder  
Role: PI on the subcontract  
Date: 08/01/2014-07/31/2016
4. **Instruments and Supplies:** Received from Pharma Industries (estimated cost = ~120K).

## **Recently completed grants:**

1. **RO3 DA027113/NIH-NIDA**  
Early Career Development Award in Chemistry of Drug Abuse and Addiction (ECHEM)  
Title: Allosteric Modulators of CB1 Cannabinoid Receptor  
Role: PI  
Requested Start Date: 08/01/2009-07/31/2013
2. **Indo-US Singh Obama Post-doctoral Fellowship Program**  
Title: Development of peripherally acting CB2-selective ligands.  
(Dr. Ganesh Chaturbhuj, post-doctoral fellow)  
Role: Mentor  
Date: 10/01/2013- 09/30/2014
3. **1R01DA026795/NIH-NIDA**  
Title: Novel Medications for Cannabis Dependence  
Role: Co-I  
Date: 07/15/2009- 05/31/2014
4. **Davis Foundation Post-doc Fellowship Program in Eating Disorder Research**  
Title: Positive Allosteric Modulators of CB1 Cannabinoid Receptor for treatment of Anorexia Nervosa. (Dr. Pushkar Kulkarni, postdoctoral fellow)  
Role: Mentor  
Date: 07/01/2012- 06/30/2015

## **8. MAJOR TEACHING EXPERIENCE**

2006-2010: Assistant Research Professor (Northeastern University)  
2010-Present: Assistant Professor (tenure-track; Northeastern University)

### **Graduate Courses**

2011-Present : Drug Discovery Journal Club (PHSC6300-003)- **Course Coordinator**  
2011 : Chemistry and Biology of Drugs of Abuse (PHSC6222)  
2013- Present: Pharmacokinetics and Drug Metabolism (PMST6252)

## Undergraduate Courses

2006-Present : Pharmacology and Medicinal Chemistry I (PHSC4501); Principal instructor for teaching medicinal chemistry

2007-Present : Pharmacology and Medicinal Chemistry II (PHSC4502); Principal instructor for teaching medicinal chemistry (**Course Coordinator**)

2014-Present : Anti-infective Agents (PHSC5360)

**9. RESEARCH MENTORING:** (\* indicate students for whom my role is mentor as well as advisor)

### UNDERGRADUATE MENTORING:

#	Year	Student's name	Degree	Current Position
1	08/2011-12/2011	Shivan Acharya	Pharm.D.	Graduated from NEU
2.	01/2012-04/2012	Monica Tiang	Pharm D	Graduated from NEU
3.	02/2014- 09/2014	Olatokunbo Onabanjo	Pharm.D.	Student at NEU (P4)
4.	02/2014- 08/2014	Lucia Zhu	Pharm.D.	Student at NEU (P4)

### PH.D./M.S. THESIS MENTORING:

1.	1999-2001	Alok Singh <sup>+</sup>	M.S. Med. Chem.	Novartis (NIBR), MA.
2.	2003-2006	Jin Zhang <sup>+</sup>	M.S. Med. Chem.	Harvard Medical School, MA.
3.	01/2007-08/2007	Vince Abeyta <sup>+</sup>	M.S. Med. Chem.	Boehringer Ingelheim, CT.
4.	2007-2011	Heidi Teng <sup>+</sup>	Ph.D. Med. Chem.	Aldrich, MA.
5.	2010-09/2011	Abhijit Kulkarni*	M.S. Med. Chem.	Pursuing Ph.D. NEU
6.	09/2011-04/2012	Khushbu Shah*	M.S. Med. Chem.	Pursuing Ph.D. Duquesne Univ.
7.	09/2012-09/2014	Prisca Mungalachetty*	M.S. Med Chem.	Novartis (NIBR), Boston
8.	09/2011-12/2012	Ameya Ranade*	M.S. Med. Chem.	Pursuing Ph.D. in Canada
9.	10/2011-11/2013	Vasanth Duggirla*	M.S. Med Chem.	
10.	2007-06/2012	Marsh D'Souza <sup>+</sup>	Ph.D. Med. Chem.	Post-doc at Scripps
11.	2006-06/2012	Rishi Sharma <sup>+</sup>	Ph.D. Med. Chem.	Scientist at MicroConstants Inc.
12.	2007-08/2012	Shama Bajaj**	Ph.D. Med. Chem.	MIT, Cambridge, MA.
13.	10/2008-10/2012	Ritesh Tichkule**	Ph.D. Med Chem.	Novartis (NIBR), Boston.
14.	09/2011-contd	Abhijit Kulkarni*	Ph.D. Med. Chem.	Pursuing Ph.D. NEU
15.	02/2014-08/2015	Siddhi Honavar*	M.S. Med. Chem.	GSK, Philadelphia
16.	09/2014-contd	Sharvik Shirodkar*	M.S. Med Chem.	Working on his M.S. Degree
17.	09/2014-contd	Ninad Dixit*	M.S. Med Chem.	Working on his M.S. Degree
18.	01/01/2016-	Peter Schaffer	M.S. Med Chem.	Working on his M.S. Degree
19.	01/01/2016-	Lucas Cantwell	M.S. Med Chem	Working on his M.S. Degree

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\* indicates my role as advisor; \*\* indicates my role as co-advisor; + indicates my role as mentor

### POST-DOCTORAL FELLOW MENTORING:

1.	2005-2008	Dr. Vidyanand Shukla+	---	Aldrich, Natick, MA.
2.	2007-2009	Dr. Shaine Cararas+	---	--
3.	02/2011-08/2015	Dr. Pushkar Kulkarni --- (Davis Foundation Fellowship)		Barnett Institute/Department of Pharm. Sci. NEU
4.	10/2013-09/2014	Dr. Ganesh Chaturbuj --- (Indo-US project; Raman Fellowship)		Department of Pharm. Sci. NEU
5.	01/2015-present	Dr. Sumanta Garai --- (NIH Post-doc Fellow)	--	Department of Pharm. Sci. NEU
6.	05/2015-present	Dr. Gopalkrushna Waghule --- (NIH Post-doc Fellow)		Department of Pharm. Sci. NEU

## GRADUATE STUDENT THESIS COMMITTEE

1.	2007-2012	Shama Bajaj, Ph.D. (Medicinal Chemistry)
2.	2007-2012	Ritesh Tichkule, Ph.D. (Medicinal Chemistry)
3.	2011- 2013	Namita Dodwadkar, Ph.D. (Pharmaceutics)
4.	2011-2013	Vasantha Duggirala, M.S. (Pharm. Sci.)
5.	2011-2013	Ameya Ranade, M.S. (Pharm. Sci.)
6.	2012-2014	Madhura Deshpande, Ph.D. (Pharmaceutics)
7.	2012-2014	Prisca Mungalachetty, M.S. (Pharm. Sci.)
8.	2013-2015	Siddhi Honavar, M.S. (Medicinal Chemistry)
9.	2012-Current	Pranali Deshpande, Ph.D. (Pharmaceutics)
10.	2012- Current	Abhijit Kulkarni, Ph.D. (Medicinal Chemistry)
11.	2014- Current	Ninad Dixit, M.S. (Medicinal Chemistry)
12.	2014- Current	Sharvik Shirodkar (Medicinal Chemistry)
13.	2012- Current	Aditi Jhaveri (Pharmaceutics)

## 10. RESEARCH INTERESTS

My research interests are focused on developing novel medications for the treatment of diseases that represent unmet medical need such as: **drug addiction, glaucoma and associated neurodegeneration, chronic pain, neuropathic pain, eating disorders** and **cognition** and **memory deficits** associated with **Alzheimer's disease** and **Schizophrenia** (with emphasis on auditory gating). The drug targets include currently being pursued in my lab are: CB1/CB2 receptors,  $\alpha 7$  nAChR and nNOS-PSD95.

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### **Targeting Cannabinoid Receptors for the Treatment of Pain, Glaucoma, PTSD, Huntington Disease and GI Disorders/Molecular Characterization of CB1 orthosteric/allosteric site(s)**

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$\Delta^9$ -Tetrahydrocannabinol ( $\Delta^9$ -THC), the major psychoactive constituent of marijuana (*Cannabis Sativa L.*) produces its physiological effects by interaction with cannabinoid receptors (CB1 and CB2). Cannabinoid receptors influence diverse cellular signal-transduction pathways. CB1 receptors are mainly located in the central nervous system, whereas the CB2 receptors are found mainly in peripheral tissues, such as spleen, tonsils and immunocytes. The adverse CNS side effects of  $\Delta^9$ -THC are mainly mediated by brain CB1 receptors. The CB2 subtype is of particular interest since it has been identified as a potential target for immune- modulator therapies. Several CB selective and CB1/CB2 mixed agonists have demonstrated preclinically potential clinical utility in treating pain, glaucoma, depression and GI disorders. CB2 selective agonists have potential utility in the treatment of peripheral and neuropathic pain and inflammation without any undesirable CB1 mediated CNS side effects which include dizziness, dry mouth, tiredness/fatigue, muscle pain and palpitations.

#### **A. Development of sub-type selective orthosteric ligands of CB receptors:**

- 1) Development of CB1 selective and water-soluble cannabinoids
- 2) Development of CB2 selective cannabinoids
- 3) Development of peripherally-restricted cannabinoids

## **B. Mapping the Binding-Sites of Cannabinoid Receptors (both Orthosteric and Allosteric sites):**

In order to improve the selectivity of CB-receptor ligands with an eye towards their therapeutic use, the three-dimensional structure of CB receptors (and virtually all other GPCRs) needs to be characterized. In the absence of experimentally determined structures of cannabinoid receptors, information about their ligand-binding sites may best be obtained through the use of affinity probes coupled with mass spectrometric analysis of labeled protein. This project involves design and synthesis of:

- 1) Covalent and electrophilic probes
- 2) “Click-chemistry” probes

Currently, my laboratory has developed key covalent probes for CB1 allosteric sites that have been characterized in functional assays are evaluated in receptor labeling studies. **One key probe, GAT100 is the FIRST EVER PROBE that has been shown to label the CB1 receptor allosteric site covalently and represents a pure NAM that lacks inverse agonist activity.** Two papers on this work are currently under review.

## **C. Allosteric Modulators of CB1 Cannabinoid Receptors**

Recent discovery of an allosteric binding site on the cannabinoid CB1 receptor invites new approaches to potential drugs that modulate cannabinoid signaling for therapeutic benefit. Several GPCRs have been shown to contain allosteric binding sites for endogenous/synthetic ligands which are discrete from the agonist binding (orthosteric) site. The binding of allosteric modulators leads to a conformational change which affects the affinity and/or efficacy of the orthosteric (endogenous) ligands, thereby fine-tuning its actions. Putative advantages of CB1-receptor allosteric inhibitors, and very recently discovered functional enhancers, include greater receptor subtype selectivity and reduced side effects. My research work, which is funded by NIH, is focused on developing novel CB1 selective allosteric modulators which include both **Positive Allosteric Modulators** (PAMs) and **Negative Allosteric Modulators** (NAMs). We have developed some potent and functionally selective allosteric modulators which are well-characterized *in vitro* and *in vivo*. Our current finding shows that CB1 PAMs may provide all or most of the beneficial effects of CB1 activation but are devoid of CB1 agonist related side effects. We hope that this work together with the allosteric covalent probe/LAPS studies will contribute significantly towards the understanding of structural requirements of the CB1 receptor’s allosteric binding site and facilitate the development of preclinical candidates. The most potent CB1 PAMs have already established their utilities for treatment of Glaucoma, PTSD, Neuropathic Pain Huntington’s Disease and Anorexia Nervosa in animal models and the results received so far are extremely promising.

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## **Development of Inhibitors of nNOS-PSD95 protein-protein interaction:**

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In 2012, we initiated our efforts on developing novel inhibitors of nNOS-PSD95 protein-protein interactions for the treatment of neuropathic pain, chemotherapy-induced neuropathy, TBI and post-traumatic stress disorder. We have recently reported (publication in *neuropharmacology*, 2015, 97; 464-75; doi:

10.1016/j.neuropharm.2015.05.038) that such novel inhibitors are very effective in treating pain and chemotherapy induced neuropathy but are devoid of CNS side effects commonly seen with current medication on market for treating neuropathic pain. Two key ligands GAT404 and GAT405 have been shown to be more potent than literature established tools and are being currently further evaluated. These novel inhibitors are also effective in treating PTSD, a disorder with an unmet medical need. This work has been currently supported by a 2-year Phase-I, SBIR grant from NIMH/NIH.

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## Development of Selective and Potent Allosteric Modulators of $\alpha 7$ Nicotinic Acetylcholine Receptors/ Molecular Characterization of $\alpha 7$ allosteric site(s)

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Nicotinic acetylcholine receptors (nAChR), members of Cys-loop superfamily of cationic ligand-gated ion channels are involved in the physiological responses to the neurotransmitter acetylcholine (ACh) and are distributed throughout the central and peripheral nervous systems. The  $\alpha 7$  nAChRs are expressed at high levels in areas involved with learning and memory and play pivotal roles in modulating neurotransmission in these area. It has been considered a promising target for improving cognitive impairments in diseases such as Alzheimer's (AD) and schizophrenia as well as for treatment of inflammation and neuropathic pain. So far our laboratory has developed potent  $\alpha 7$  nAChR PAMs and ago-PAMs and the lead molecule GAT107 shown good potency in *improving memory and cognition* in animal models as well as reducing pain in animal models of *neuropathic and chronic pain*. We show that these compounds are orally bioavailable, non-cytotoxic and very selective for  $\alpha 7$  nAChR subtype. More than 100 compounds have been synthesized so far in my laboratory to identify structural requirement for ago-PAM activity. A series of publications that involves medical chemistry as well as structural biology have been communicated recently and are under review.

### **11. PEER-REVIEWED PUBLICATIONS (IN CHRONOLOGICAL ORDER)**

1. Rao, P. S., **Thakur, G.A.**, Lahiri, G. K., Synthesis, characterization and redox properties of ruthenium (II) phenolato Schiff base mixed ligand complexes. *Indian J. Chem., Sect. A: Inorg., Bio-inorg., Phys., Theor., Anal. Chem.*, **1996**, 35A (11), 946-51.
2. Santra, B. K., **Thakur G. A.**, Ghosh, P., Pramanik, A., Lahiri, G. K., A novel example of metal mediated aromatic thiolation in ruthenium complex: crystal structure of RuII (SC<sub>6</sub>H<sub>4</sub>N:NC<sub>5</sub>H<sub>4</sub>N)<sub>2</sub>. *Inorg. Chem.*, **1996**, 35(10), 3050-2.
3. **Thakur, G. A.**, Narayanswami, K., Lahiri, G. K., Synthesis, characterization and redox properties of ruthenium (II) dithiocarbonato complexes having 2,2'-bipyridine co-ligands. *Indian J. Chem., Sect. A: Inorg., Bio-inorg., Phys., Theor., Anal. Chem.*, **1996**, 35A (5), 379-84.
4. Nikas, S. P., **Thakur, G. A.**, Makriyannis, A., A convenient and effective synthesis of 3-(3,5-Dimethoxyphenyl)propanal. *Synth. Comm.*, **2002**, 32(11), 1751-1756.

5. Palmer, S., **Thakur, G. A.**, Makriyannis, A., Cannabinergic ligands. *Chem. and Phys. of Lipids*, **2002**, 121(1-2), 3-19.
6. Nikas, S. P., **Thakur, G. A.**, Makriyannis, A., Synthesis of side chain specifically deuterated(-)- $\Delta^9$ -tetrahydrocannabinoids. *J. Label Compd. Radipharm.*, **2002**, 45, 1-12.
7. Nikas, S. P., **Thakur, G. A.**, Makriyannis, A., Regiospecifically deuterated (-)- $\Delta^9$ -tetrahydrocannabinoids. *J. Chem. Soc. Perkin. Trans.1*, **2002**, 22, 2544-2548.
8. **Thakur, G. A.**, Palmer, S., Harrington, P. E., Stergiades, I. A., Tius, M. A., Makriyannis, A., Enantiomeric resolution of a novel chiral cannabinoid receptor ligand. *J. Biochem. Biophys. Methods*, **2002**, 54(1-3), 415-422.
9. Lu, D., Meng, Z., **Thakur, G. A.**, Fan, P., Steed, J., Tartal, C. L., Hurst, D. P., Reggio, P. H., Deschamps, J. R., Parrish, D. A., George, C., Jarbe, T. U., Lamb, R. J., Makriyannis, A., Adamantyl Cannabinoids: A novel class of cannabinergic ligands. *J. Med. Chem.*, **2005**, 48 (14), 4576-4585.
10. McLaughlin, P. J., Lu, D., Winston, K. M., **Thakur, G. A.**, Swezey, L. A., Makriyannis, A., Salamone, J. D., Behavioral effects of the novel cannabinoid CB1 agonist AM411. *Pharmacol. Biochem. Behav.*, **2005**, 81(1), 78-88.
11. McLaughlin, P. J., Brown, C. M., Winston, K. M., **Thakur, G. A.**, Lu, D., Makriyannis, A., Salamone, J. D., The novel cannabinoid agonist AM 411 produces a biphasic effect on accuracy in a visual target detection task in rats. *Behav. Pharmacol.*, **2005**, 16(5-6), 477-486.
12. Picone, R. P., Khanolkar, A. D., Xu, W., Ayotte L. A., **Thakur, G. A.**, Hurst, D. P., Abood, M. E., Reggio, P. H., Fourier, D. J., Makriyannis, A., (-)-7'-Isothiocyanato-11-hydroxy-1',1'-dimethylheptylhexahydrocannabinol (AM841), a high-affinity electrophilic ligand, interacts covalently with a cysteine in helix six and activates the CB1 cannabinoid receptor. *Mol. Pharmacol.*, **2005**, 68 (6), 1623-1635.
13. **Thakur, G. A.**, Nikas, S. P., Makriyannis, A., CB1 Cannabinoid receptor ligands. *Mini Reviews in Medicinal Chemistry*, **2005**, 5(7), 631-640.
14. **Thakur, G. A.**, Duclos, R.,I.,Jr., Makriyannis, A., Natural cannabinoids: templates for drug discovery. *Life Sci.*, **2005**, 78(5), 454-466.
15. Pavlopoulos, S., **Thakur, G. A.**, Nikas, S. P., Makriyannis, A., Cannabinoid receptors as therapeutic targets. *Curr. Pharm. Design*, **2006**, 12(14), 1751-1769.
16. Kapur, A., Hurst, D. P., Fleischer, D., Whitnell, R., **Thakur, G. A.**, Makriyannis, A., Reggio, P. H., Abood, M. E., Mutation studies of Ser7.39 and Ser2.60 in the human CB<sub>1</sub> cannabinoid receptor: evidence for a serine induced bend in CB<sub>1</sub> transmembrane helix 7. *Mol. Pharmacol.*, **2007**, 71(6), 1512-1524.
17. Nikas, S. P., **Thakur, G. A.**, Parrish, D. A., Alapafuja, S. O., Huestis, M., Makriyannis, A., A concise methodology for the synthesis of (-)- $\Delta^9$ -tetrahydrocannabinol and (-)- $\Delta^9$ -tetrahydrocannabinol metabolites and their regiospecifically deuterated analogs. *Tetrahedron*, **2007**, 63, 8112-8123.
18. Khanolkar, A. D., Lu, D., Ibrahim, M., Duclos, R.I. Jr., **Thakur, G. A.**, Malan, T. P. Jr., Porreca, F., Veerappan, V., Tian, X., George, C., Parrish, D. A., Papahatjis, D. P., Makriyannis, A., Cannabilactones: a novel class of CB2 selective agonists with peripheral analgesic activity. *J. Med. Chem.*, **2007**, 50 (26), 6493-6500.
19. Sink, K.S., McLaughlin, P.J., Wood, J.A., Brown, C., Fan, P., Vemuri, V.K., Peng, Y., Olzewska, T., **Thakur, G.A.**, Makriyannis, A., Parker, L.A., Salamone, J.D. (2008), The novel cannabinoid CB1 receptor neutral antagonist AM4113 suppresses food intake and food-reinforced behavior but does not induce signs of nausea in rats. *Neuropsychopharmacology*, 33(4), 946-955.
20. Kapur, A., Samaniego, P., **Thakur, G. A.**, Makriyannis, A., Abood, M. E., Mapping the structural requirements in the CB1 cannabinoid receptor transmembrane helix II for signal transduction. *JPET*, **2008**, 325 (1), 341-8.
21. Bergman, J., Delatte, M. S., Paronis, C. A., Vemuri, V. K., **Thakur, G. A.**, Makriyannis, A., Some effects of CB1 antagonists with inverse agonist and neutral biochemical properties. *Physiol. Behav.*, **2008**, 93(4-5), 666-70.



22. Rahn, E. J., Zvonok, A. M., **Thakur, G. A.**, Khanolkar, A. D., Makriyannis, A., Hohmann, A. G., Selective activation of cannabinoid CB<sub>2</sub> receptors suppresses neuropathic nociception induced by treatment with the chemotherapeutic agent paclitaxel in rats. *JPET.*, **2008**, 327(2), 584-91.
23. Pei, Y., Mercier, R. W., Anday, J. K., **Thakur, G. A.**, Zvonok, A. M., Reggio, P. H., Janero, D. R., Makriyannis, A., Ligand-binding architecture of human CB<sub>2</sub> cannabinoid receptor: evidence for a receptor subtype-specific binding motif and modeling GPCR activation. *Chem. Biol.*, **2008**, 15(11), 1207-19.
24. Sink K. S., Segovia K. N., Nunes E. J., Collins L. E., Vemuri V. K., **Thakur G. A.**, Makriyannis A., Salamone J. D. Intracerebroventricular administration of cannabinoid CB<sub>1</sub> receptor antagonists AM251 and AM4113 fails to alter food-reinforced behavior in rats. *Psychopharmacology (Berl.)*, **2009**, 206(2), 223-232.
25. **Thakur, G. A.**,\*Tichkule, R., Bajaj, S., Makriyannis, A., Recent advances in cannabinoid receptor agonist. *Expert Opinion on Therapeutic Patents.* **2009**, 19 (12), 1647-73.
26. Dixon, D. D., Sethumadhavan, D., Benneche, T., Banaag, A. R., Tius, M. A., **Thakur, G. A.**, Bowman, A., Wood, J., Makriyannis, A. Heteroadamantyl Cannabinoids. *J. Med. Chem.*, **2010**, 53(15), 5656-66.
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## 12. BOOK CHAPTERS

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2. “*Cannabinoid Receptors as Therapeutic Targets*” given at Bombay College of Pharmacy (BCP), Mumbai, India (2011).
3. “*Targeting Obesity with Negative Allosteric Modulators of CB1 Cannabinoid Receptor*” in Discovery of Target meeting, (2012).
4. “*Tuning the Endocannabinoid System: Allosteric Modulators of CB1 Cannabinoid Receptor*” given in the department of Pharmaceutical Sciences, St. John University, (2013).
5. “*Allosteric Modulators of CB1 Cannabinoid Receptor*” given in the department of pharmacology and therapeutics, University of Florida, (2013).
6. “*Tuning Endocannabinoid System for Therapeutic Gain*” given at ICT, India, on 28<sup>th</sup> March 2014.
7. “*Novel Therapeutic Opportunities via Allosteric Modulation of CB1 Cannabinoid Receptor*” given at Indian Institute of Science Education and Research (IISER), Pune, India, (2014).
8. “*Tuning the Endocannabinoid System: Allosteric Modulators of CB1 Cannabinoid Receptor*” given at Indian Institute of Technology (IIT), Bombay, India, (2014).
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12. “*Tuning the Endocannabinoid System: Allosteric Modulators of CB1 Cannabinoid Receptors*” given at the *Department of Pharmaceutical Sciences, Eugene Applebaum College of Pharmacy and Health Sciences (EACPHS), Wayne State University, Detroit, MI*, (2014).
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