Keykavous Parang

Associate Dean of Research, Graduate Studies and Global Affairs, Full Professor School of Pharmacy, Chapman University

Address

Chapman University 9401 Jeronimo Road Irvine, California 92816

E-Mail: parang@chapman.edu

Phone: 714-516-5489 Fax: 714-516-5481

EDUCATION

Sept. 1992-Aug. 1997 Ph.D. in Medicinal Chemistry, Faculty of Pharmacy, University of

Alberta (Drs. L. I. Wiebe, E. E. Knaus, Supervisors)

Jan. 1984-Jan. 1989 Pharm.D., Faculty of Pharmacy, University of Tehran

(Dr. A. Shafiee, Supervisor)

RESEARCH EXPERIENCE

July 2013-Present Associate Dean of Research, Graduate Studies and Global

Affairs, Full Professor, School of Pharmacy, Chapman University

Sept. 2013-Present Member, Chao Family Comprehensive Cancer Center, University of

California, Irvine

Nov. 2012-Present Adjunct Faculty, Department of Molecular Pharmacology,

Physiology and Biotechnology, Brown University

July 2013-Present Research Professor, Department of Biomedical and

Pharmaceutical Sciences, College of Pharmacy, University of

Rhode Island

July 2008-June 2013 Full Professor, Department of Biomedical and Pharmaceutical

Sciences, College of Pharmacy, University of Rhode Island

March 2013-Present Adjunct Faculty, International Center for Chemical and Biological

Sciences, University of Karachi

May 2012-June 2013 Program Coordinator, Rhode Island IDeA Network of Biomedical

Research Excellence (INBRE) sponsored by NIH

January 2010-Dec. 2010 Sabbatical Research, Department of Molecular Pharmacology,

Physiology and Biotechnology, Brown University

Nov. 1, 2000-Present Visiting scientist (Scholar), Department of Pharmacology and

Molecular Sciences, School of Medicine, Johns Hopkins University

April 2006-July 2008 Associate professor, Department of Biomedical and

Pharmaceutical Sciences, College of Pharmacy, University of

Rhode Island

Oct. 2000-April 2006 Assistant professor, Department of Biomedical and

Pharmaceutical Sciences, College of Pharmacy, University of

Rhode Island

1. Designing self-assembled peptide nanomaterials for drug delivery and tissue engineering.

- 2. Designing protein kinase inhibitors through target-based strategies using molecular modeling, synthetic, and biochemical tools.
- 3. Novel approaches in designing bifunctional anti-infective agents. Currently our focus is to design prodrugs of antifungal agents and HIV microbicides.
- 4. Novel strategies in the solid-phase synthesis.
- 5. Polymer-based drug delivery
- 6. Synthesis and evaluation of modified nucleotides and nucleic acids
- 7. Anticancer drug design

July 1999-Oct. 2000 Postdoctoral fellow, Department of Pharmacology and Molecular

Sciences, School of Medicine, Johns Hopkins University

(Burrough Welcome Fund) (Dr. Philip A. Cole)

Chemical cross-linking agents for identification of substrates for protein kinases, Bisubstrate ATP-peptide inhibitors of tyrosine kinases.

Jan. 1999-July 1999 Postdoctoral fellow, Department of Bioorganic Chemistry,

Rockefeller University (Dr. Philip A. Cole)

Tyrosine analogs as alternative substrates for protein kinase Csk.

Sept. 1997-Jan. 1999 Postdoctoral fellow, Department of Chemistry, University of

Alberta (Dr. Ole Hindsgaul)

Sulfation and phosphorylation of carbohydrates and nucleosides using solid phase organic synthesis.

Sept. 1992-Aug. 1997 Ph.D., Faculty of Pharmacy, University of Alberta

(Drs. L. I. Wiebe, E. E. Knaus, Supervisors)

- 1. Synthesis, biodistribution, pharmacokinetic, and biological evaluation of 5'-O-myristoyl derivatives of 3'-azido-3'-deoxythymidine (AZT) and 3'-fluoro-3'-deoxythymidine (FLT) as prodrugs for the treatment of AIDS and hepatitis B.
- Synthesis, antifungal and antiviral activities of myristic acid analogs.
 This research project involved the design and synthesis of double-barreled myristoyl, and heteroatom-modified myristoyl, derivatives of nucleoside analogs for the treatment of AIDS and Hepatitis B.
- 3. Synthesis of derivatives of 2-fluoromethyl and 3-fluoromethyl tyrosine.
- 4. Synthesis of 4-hydroxymethyl-2-nitro-1-trityl-imidazole.
- 5. Synthesis of 1'- and 2,2'-deuterated nucleosides.

Jan. 1992-Sept. 1992 Sabbatical leave, Faculty of Pharmacy, University of Alberta (Drs. L. I. Wiebe, E. E. Knaus, Supervisors)

Jan. 1989-Dec. 1991 Ph.D. student, Faculty of Pharmacy, University of Tehran (Dr. A. Shafiee, Supervisor)

Synthesis of derivatives of 1-phenylethyl substituted 1,2,4-triazoles and imidazoles as spermicides.

Jan. 1984-Jan. 1989 Pharm.D. thesis, Faculty of Pharmacy, University of Tehran (Dr. A. Shafiee, Supervisor)

Synthesis of substituted 2-(1-methyl-5-nitro-2-imidazolyl)quinolines, effective drugs against tropical diseases.

MEMBERSHIP IN PROFESSIONAL SOCIETIES

- 1. American Chemical Society (1996-Present)
- 2. American Association for the Advancement of Science (1999-present)
- 3. American Association of Pharmaceutical Scientists (AAPS) (1993-1995) (2004-Present)
- 4. The Johns Hopkins Medical and Surgical Association (2002-Present)
- 5. American Diabetes Association (2004-Present)
- 6. American Association of Colleges of Pharmacy (February 2001-2008)
- 7. SIGMA XI (The Scientific Research Society) (November 2000-November 2001)
- 8. Distinguished member of PIRE Advisory Board, NSF, University of Rhode Island Partnership in International Research and Education (PIRE) (2008)

SCIENTIFIC JOURNAL EDITOR OR EDITORIAL ADVISORY BOARD

- 1. The Executive Guest Editor of "Current Pharmaceutical Design" (2001-Present).
- 2. Editorial Advisory Board "Current Medicinal Chemistry, Central Nervous System Agents" (June 2004-Present).

- 3. Editorial Advisory Board of "Recent Patent Reviews on CNS Drug Discovery" (March 2005-Present).
- 4. Editorial Board member of "Perspectives in Medicinal Chemistry" (January 2007-Present).
- 5. Editorial Board Member of "Open Biochemistry Journal" (June 2007-Present).
- 6. Editorial Board Member of "Daru Journal of Pharmaceutical Sciences" (Aug. 2008-Present).
- 7. Editorial Board Member of Associate Editors, "The Beilstein Journal of Organic Chemistry" (December 2008-Present).
- 8. Editorial Board Member, "Journal of Herbal Drugs" (Aug. 2009-Present).
- 9. Editorial Advisory Board, "Chemistry Biology Interface" (September 2011-Present).
- 10. Honorary Editorial Board, "Research and Reports in Transdermal Drug Delivery" (Dec. 2011-Present)
- 11. Editorial Board Member of "Journal of Pharmacological & Biomedical Analysis" (June 2012-Present)

PEER REVIEW ACTIVITY AND GRANT REVIEW ACTIVITY

- Reviewer for Organic Letters, Bioorganic Medicinal Chemistry Letters, Bioorganic Medicinal Chemistry, Journal of Combinatorial Chemistry, Current Pharmaceutical Design, Tetrahedron Letters, Journal of Organic Chemistry, Chemistry and Biology, and Journal of Medicinal Chemistry, ChemMedChem.
- Center For Scientific Review, NIH, Microbicidal Preclinical Development Program, ZRG1 AARR-1, August 7-9, 2001
- Center For Scientific Review, NIH, *Microbicidal Preclinical Development Program* ZRG1 AARR-1, December 9, 2002.
- Center For Scientific Review, NIH, *Microbicidal Preclinical Development Program*, ZRG1 AARR-1, July 9, 2003.
- Center For Scientific Review, NIH, *Drug Delivery & Drug Discovery SBIR/STTR Panel*, ZRG1 SSS-L (10), June 30, 2003- July 01, 2003.
- Center For Scientific Review, NIH, *Microbicidal Preclinical Development Program*, ZRG1 AARR-A 51, December 8-9, 2003.

- Center For Scientific Review, NIH, Microbicide Innovation R21/R33 program, ZAI1 BLG-A (S1), May 23-24, 2006.
- Ad Hoc Reviewer, INSF, Development of COX-2 Inhibitors, December 2006.
- Reviewer, 2006 Assessment of Research Doctorate Programs, National Research Council (NRC), December 2006.
- The Estonian Science Foundation, Evaluation of proposals, July 2008.
- Organic and Macromolecular Chemistry CAREER Panel, National Science Foundation (NSF), October 6-7, 2008.
- Center For Scientific Review, NIH, Microbicide Innovation R21/R33 program (MIP IV), ZAI1 RB-A (J1) 2, November 20-21, 2008.
- Center For Scientific Review, NIH, *Microbicide Innovation R21/R33 program (MIP VI)*, ZAI1 RB-A (J1), December 2-3, 2010.
- Cancer Drug Discovery Peer Review Committee, American Cancer Society, June 23-24, 2011.
- Cancer Drug Discovery Peer Review Committee, American Cancer Society, January 19-20, 2012.
- Cancer Drug Discovery Peer Review Committee, American Cancer Society, June 14-16, 2012.
- City University of New York, Cuny Collaborative Research Grant Proposal Review, April 6, 2013.
- **Permanent member**, Cancer Drug Discovery Peer Review Committee, American Cancer Society, August 1, 2012-Present.

AWARDS AND HONORS

Year	<u>Award</u>	<u>Awarding</u>	Nature of Award
		Agency/Institution	
2014	Outstanding Intellectual	University of Rhode	Efficient Synthesis of
	Property	Island	CN2097 and RC7 and
	Development		Their Analogs
2014	Outstanding Intellectual	University of Rhode	Synthesis of Maplexin J
	Property	Island	
	Development		
2014	Outstanding Intellectual	University of Rhode	Phenolic Compounds
	Property	Island	with Antioxidant and
	Development		Anti-Cancer Properties
2014	Outstanding Intellectual	University of Rhode	Antibacterial Peptides
	Property	Island	Containing Arginine and
	Development		Tryptophan Residues
2013	Outstanding Intellectual	University of Rhode	Solid-phase synthesis of
	Property	Island	modified
	Development		oligonucleotides
			containing
			diphosphodiester inter-
			nucleotide linkages.
2013	Outstanding Intellectual	University of Rhode	Preparation of quebecol
	Property	Island	and its analogs as anti-
	Development		cancer agents.
2012	Research Excellence Award	College of	Recognition
		Pharmacy	
		University of Rhode	
		Island	
2012	M2012 Travel Scholarship	Microbicides: From	Travel scholarship and
		Discovery to Delivery	invitation for giving an
			oral presentation
2011	Outstanding Researcher	University of Rhode Island	Recognition
2011	The Rho Chi Society	University of Rhode	Honorary Member
2011	The Academic Honor Society in	Island	Tionorary McMibel
	Pharmacy Beta Pi Chapter	isianu	
2010	M2010 Travel Scholarship	Microbicide: Building	Travel scholarship and
2010	Wizo to Travel Scholarship	Bridges in HIV	invitation for giving an
		Prevention	oral presentation
		Organizing	oral prosentation
		Committee	
		Committee	

2009	Harry and Elsa Jiler—American Cancer Society Professors	American Cancer Society	Travel scholarship and invitation for giving an
	Meeting Travel Scholarship	Coolety	oral presentation
2009	Honorary Fellow of the Indian Society Chemists and Biologists (ISCB)	Indian Society Chemists and Biologists	Honorary Fellow
2004	Outstanding Intellectual Property Development	University of Rhode Island	Research: Recognition for Discovery of the Docking Site of Tyrosine Kinases
2004	Outstanding Intellectual Property Development	University of Rhode Island	Research: Recognition for Discovery of Bisubstrate Inhibitors of Protein Tyrosine Kinases as Anticancer Agents
2004	Outstanding Intellectual Property Development	University of Rhode Island	Research: Recognition for Discovery of Novel Bisubstrate Antifungal Derivatives
2003	Listed in Marquis Who's Who in America	Marquis Who's Who	Honor
2001	Commercial Innovation Award	Slater Center For Biomedical Technology	Research: Industrial Innovation Research
1998	American Chemical Society Travel Grant	American Chemical Society	Research Travel Award
1997- 1999	Alberta Heritage Foundation for Medical Research Postdoctoral Fellowship	Alberta Heritage Foundation for Medical Research	Postdoctoral Fellowship Scholarship
1997	The Most Positive Influence Award	University of Alberta	Service Honorary Award
1996	Golden Bulb Light Award	University of Alberta	Service
1994-	Alberta Heritage Foundation for	Alberta Heritage	Ph.D. Student
1996	Medical Research Studentship Scholarship	Foundation for Medical Research	Scholarship
1995	J. Gordon Graduate Student Award	University of Alberta	Research
1994	Dr. Wu Hong Fund Poster Prize	University of Alberta	Research, Poster Prize
1993	Mike Wolowyk Graduate Scholarship	University of Alberta	Research
1993	Myer Horowitz Graduate Scholarship	University of Alberta	Research

PROFESSIONAL AND TECHNICAL EXPERIENCE

- 1. Medicinal chemistry.
- 2. Nano-based drug design and drug delivery tools.
- 3. Peptide synthesis and application as enzyme inhibitors and drug delivery tools.
- 4. Biomaterials characterization using Transmission Electron Microscopy (TEM), Scanning Electron Microscopy (SEM), and Dynamic Light Scattering (DLS)
- 5. Cell-based assays and imaging techniques (flow cytometry, fluorescence microscopy, confocal microscopy).
- 6. Solid-phase organic synthesis and combinatorial chemistry.
- 7. Synthetic organic chemistry of lipids, peptides, nucleosides, nucleic acids, carbohydrates, and heteroaromatics.
- 8. Analytical and spectroscopy methods like HPLC, NMR, IR, UV and MS.
- 9. Enzymatic reactions.
- 10. Molecular modeling and quantitative structure-activity relationship (QSAR) using Insight II, Biosym, Hyperchem, Rasmol, ViewerLite, ChemPro. Software.
- 11. *In vivo* biodistribution and pharmacokinetic studies.
- 12. Techniques in molecular biology (electrophoresis, cloning of DNA, hybridization, gene transfection).
- 13. Enzyme kinetic studies.
- 14. Using radioactive compounds for biological assays like enzymatic studies.

PUBLICATIONS

- 1. Shafiee, A., Pirouzzadeh, B., Ghasemian, F., <u>Parang, K.</u> Synthesis of 2-acetyl-1-methyl -5-nitroimidazole. *J. Heter. Chem.* (1992) 29, 1021-1023 and *Cheminform* (1992) 23, 50.
- 2. Shafiee, A., <u>Parang, K.</u>, Khazen, M., Ghasemian, F. Synthesis of substituted 2-(1-methyl-5-nitro-2-imidazolyl)quinolines. *J. Heter. Chem.* (1992) 29, 1859-1861.
- 3. Sharifan, A., <u>Parang, K.</u>, Zorrieh-Amirian, H., Nazarinia, M., Shafiee, A. Synthesis of 1-substituted 1,2,4-triazoles, imidazoles and benzimidazoles. *J. Heter. Chem.* (1994) 31, 1421-1423.
- 4. **Parang, K.**, Knaus, E. E., Wiebe, L. I., Sardari, S., Daneshtalab, M., Csizmadia, F. Synthesis and antifungal activities of myristic acid analogs. *Arch. Pharm.-Pharm. Med. Chem.* **(1996)** 329, 475-482.
- 5. **Parang, K.**, Wiebe, L. I., Knaus, E. E., Huang, J. S., Tyrrell, D. L., Csizmadia, F. *In vitro* antiviral activities of myristic acid analogs against human immunodeficiency and hepatitis viruses. *Antiviral Research* **(1997)** *34*, 75-90.
- 6. **Parang, K.**, Wiebe, L. I., Knaus, E. E. Syntheses and biological evaluation of 5'-O-myristoyl derivatives of thymidine against human immunodeficiency virus (HIV-1). *Antiviral. Chem. Chemother.* **(1997)** *8*, 417-427.
- 7. **Parang, K.**, Knaus, E. E., Wiebe, L. I. Synthesis, *in vitro* anti-HIV structure-activity relationships and stability of 5'-O-myristoyl analogue derivatives of 3'-azido-2',3'-dideoxythymidine as potential prodrugs of 3'-azido-2',3'-dideoxythymidine (AZT). *Antiviral. Chem. Chemother.* **(1998)** *9*, 311-323.
- 8. **Parang, K.**, Wiebe, L. I., Knaus, E. E. *In vivo* pharmacokinetic parameters, liver and brain uptake of (±)-3'-azido-2',3'-dideoxy-5'-O-(2-bromomyristoyl)thymidine as potential prodrug of 3'-azido-3'-deoxythymidine. *J. Pharm. Pharmacol.* **(1998)** *50*, 989-996.
- 9. **Parang, K.**, Knaus, E. E., Wiebe, L. I. Synthesis, *in vitro* anti-HIV activity, and biological stability of 5'-O-myristoyl analogue derivatives of 3'-fluoro-2',3'-dideoxythymidine (FLT) as potential prodrugs of FLT. *Nucleosides & Nucleotides* (1998) 17, 987-1008.
- 10. Parang, K., Wiebe, L. I., Knaus, E. E., Huang, J. S., Tyrrell, D. L. *In vitro* anti-hepatitis B virus activities of 5'-O-myristoyl analogue derivatives of 3'-fluoro-2',3'-dideoxythymidine (FLT) and 3'-azido-2',3'-dideoxythymidine (AZT). *J. Pharm. Pharmaceut. Sci.* (1998) 1, 107-113.
- 11. Ablooglu, A. J., Till, J. K., Kim, K., <u>Parang, K.,</u> Cole, P. A., Hubbard, S. R., Kohanski, R. A. Probing the catalytic mechanism of the insulin receptor kinase with a tetrafluorotyrosine-containing peptide substrate. *J. Biol. Chem.* **(2000)** *275*, 30394-30398.

- 12. Kim, K., <u>Parang, K.,</u> Lau, O. D., Cole, P. A. Tyrosine analogs as alternative substrates for protein tyrosine kinase Csk: insights into substrate selectivity and catalytic mechanism. *Bioorg. Med. Chem.* **(2000)** *8*, 1263-1268.
- 13. Parang, K., Wiebe, L. I., Knaus, E. E. Novel approaches in designing prodrugs of AZT. *Current Med. Chem.* (2000) 7, 995-1039.
- 14. Parang, K., Till, J. H., Ablooglu, A. J., Kohanski, R. A., Hubbard, S. R., Cole, P. A. Mechanism-based design of a protein kinase inhibitor. *Nature Structural Biology* (2001) 8, 37-41.
- 15. <u>Parang, K.</u>, Fournier, E. J.-L., Hindsgaul, O. A solid phase reagent for the capture phosphorylation of carbohydrate and nucleosides. *Org. Lett.* **(2001)** *3*, 307-309.
- 16. **Parang, K.**, Preface, Anti-HIV design, Bentham Science Publishers, Hilversum, *Current Pharmaceutical Design* **(2002)** *8*, 8.
- 17. <u>Parang, K.,</u> Miri, R. Review of Organic Chemistry Principals, Kushamehr Publications, First Edition, Shiraz, **2002**, Persian.
- 18. **Parang, K.**, Kohn, J. A., Saldahna, A., Cole, P. A. Development of photo-crosslinking reagents for protein kinase-substrate interactions. *Febs Letters* **(2002)** *520*, 156-160.
- 19. **Parang, K.** Polymer-supported reagents for methylphosphorylation and phosphorylation of Carbohydrates. *Bioorg. Med. Chem. Lett.* **(2002)** *12*, 1863-1866.
- 20. **Parang, K.**, Cole, P. A. Bisubstrate inhibitors of protein kinases. *Pharmacology and Therapeutics* **(2002)** 93, 145-157.
- 21. Nam, N. H., <u>Parang, K.</u> Current Drug Targets for Anticancer Drug Discovery. *Current Drug Targets* (2003), *4*, 159-179.
- 22. <u>Parang, K.</u>, Preface, Anti-HIV design, Bentham Science Publishers, Hilversum, *Current Pharmaceutical Design* **(2003)** *9*, 22.
- 23. Nam, N. H., Sardari, S., <u>Parang, K.</u> Reactions of solid supported reagents and solid supports with alcohols and phenols through their hydroxyl functional group. *J. Comb. Chem.* (2003) *5*, 479-546 and *ChemInform* (2003), *34*, 49.
- 24. Sardari, S., Pourmorad, F., Tiemo, A., Nam, H., <u>Parang, K.</u> Protein kinases and their modulation in the central nervous system. *Current Medicinal Chemistry-CNS* (2003) 3, 341-364.
- 25. Lee, S., Lin, X., Nam, N. H., <u>Parang, K.,</u> Sun G. Determination of the substrate-docking site of protein tyrosine kinase Csk. *Proc. Nat. Acad. Sci. U.S.A.* **(2003)** *100*, 14707-14712.

- 26. Force, T., Kuida, K., <u>Parang, K.,</u> Kyriakis, J. M. Inhibitors of protein kinase signaling pathways: emerging therapies for cardiovascular disease. *Circulation* **(2004)** *109*, 1196-1205.
- 27. Carballeira, N. M., Ortiz, D., <u>Parang, K.,</u> Sardari, S. Total synthesis and *in vitro* antifungal activity of 2-methoxytetradecanoic acid. *Arch. Pharm.-Pharm. Med. Chem.* (2004) 337, 152-155.
- 28. Nam, N.-H., Ye, G., Sun, G., <u>Parang, K.</u> Conformationally constrained peptide analogues of pTyr-Glu-Glu-Ile as inhibitors of the Src SH2 domain binding. *J. Med. Chem.* **(2004)** *47*, 3131-3141.
- 29. Nam, N. H., Pitts, R., Sun, G., Sardari, S., Tiemo, A., Xie, M., Yan, B., <u>Parang, K.</u> Design of tetrapeptide ligands as inhibitors of the Src SH2 domain. *Bioorg. Med. Chem.* (2004) 12, 779-787.
- 30. Schmidt, B., Jiricek, J., Titz, A., Ye, G., <u>Parang, K.</u> Copper dipicolinates as peptidomimetic ligands for the Src SH2 domain. *Bioorg. Med. Chem. Lett.* **(2004)** *14*, 4203-4206.
- 31. Nam, N. H., Sardari, S., Selecky, M., <u>Parang, K.</u> Carboxylic acids and phosphate ester derivatives of fluconazole: synthesis and antifungal activities. *Bioorg. Med. Chem.* (2004) *12*, 6255-6269.
- 32. Nam, N. H., Lee, S., Ye, G., Sun, G., <u>Parang, K.</u> ATP-phosphopeptide conjugates as inhibitors of Src tyrosine kinases. *Bioorg. Med. Chem.* (2004) *12*, 5753-5766.
- 33. **Parang, K.**, Sun, G. Design strategies for protein kinases inhibitors. *Current Opinions In Drug Discovery* **(2004)** *7*, 630-638.
- 34. Ahmadibeni, Y., <u>Parang, K.</u> Solid-phase reagents for selective monophosphorylation of carbohydrates and nucleosides. *J. Org. Chem.* **(2005)** *70*, 1100-1103.
- 35. **Parang, K.**, Sun, G. Protein kinase inhibitors in drug discovery. *Drug Discovery Handbook*, **2005**, Wiley-Interscience, New Jersey, Ed. Gad, S. C. 1191-1257.
- 36. Lin, X., Ayrapetov, M. K., Lee, S., <u>Parang, K.,</u> Sun G. Probing the communication between the regulatory and catalytic domains of a protein tyrosine kinase, Csk. *Biochemistry* **(2005)** *44*, 1561-1567.
- 37. Hines, A. C., <u>Parang, K.,</u> Kohanski, R. A., Hubbard, S. R., Cole, P. A. Bisubstrate analog probes for the insulin receptor protein tyrosine kinase: Molecular yardsticks for analyzing catalytic mechanism and inhibitor design. *Bioorg. Chem.* **(2005)** 33, 285-297.
- 38. Carballeira, N. M., O'Neill, R., <u>Parang K.</u> Racemic and optically active 2-methoxy-4-oxatetradecanoic acids: Novel synthetic fatty acids with selective antifungal properties. *Chem. Phys. Lipids* (2005) *136*, 47-54.

- 39. Ahmadibeni, Y., <u>Parang, K.</u> Polymer-bound oxathiaphospholane: A solid-phase reagent for regioselective monothiophosphorylation and monophosphorylation of unprotected nucleosides and carbohydrates. *Org. Lett.* **(2005)** *7*, 1955-1958.
- 40. Ayrapetov, M. K., Nam, N. H., Ye, G. Kumar, A., <u>Parang, K.,</u> Sun, G. Functional diversity of Csk, Chk, and Src SH2 domains due to a single residue variation. *J. Biol. Chem.* (2005) 280, 25780-25787.
- 41. Carballeira, N. M., O'Neill, R., <u>Parang, K.</u> Total synthesis and further scrutiny of the in vitro antifungal activity of 6-nonadecynoic acid. *Arch. Pharm.-Pharm. Med. Chem.* (2005) 338, 441-443.
- 42. Ye, G., Ayrapetov, M., Nam, N. H., Sun, G., <u>Parang, K.</u> Solid-phase binding assays of peptides using EGFP-Src SH2 domain fusion protein and biotinylated Src SH2 domain. *Bioorg. Med. Chem. Lett.* **(2005)** *15*, 4994-4997.
- 43. **Parang, K.**, Sun, G. Recent advances in the discovery of Src kinases inhibitors. *Expert Opin. Ther. Patents* **(2005)** *15*, 1183-1207.
- 44. Ahmadibeni, Y., <u>Parang, K.</u> Selective diphosphorylation, dithiodiphosphorylation, triphosphorylation, and trithiotriphosphorylation of unprotected carbohydrates and nucleosides. *Org. Lett.* **(2005)** *7*, 5589-5592.
- 45. **Parang, K.** Editor. Novel approaches in designing anti-HIV microbicides and anti-HIV Agents. *Curr. Pharm. Des.* **(2005)** *11*, 123 pp.
- 46. Lin, X., Wang, Y., Ahmadibeni, Y., <u>Parang, K.</u>, Sun, G. Structural basis for domain-domain communication in a protein tyrosine kinase, Csk. *J. Mol. Biol.* **(2006)** *357*, 1263-1273.
- 47. Lee, S., Ayrapetov, M. K., Kemble, D., <u>Parang, K.</u>, Sun, G. Docking-based substrate recognition by the catalytic domain of a protein tyrosine kinase, the C-terminal Src kinase. *J. Biol. Chem.* **(2006)** *281*, 8183-8189.
- 48. Ahmadibeni, Y., <u>Parang, K.</u> Solid-phase synthesis of dinucleoside and nucleoside-carbohydrate phosphodiesters and thiophosphodiesters. *J. Org. Chem.* **(2006)** *71*, 6693-6696.
- 49. Ahmadibeni, Y., **Parang, K.** Application of a solid-phase β-triphosphitylating reagent in the synthesis of nucleoside β-triphosphates. *J. Org. Chem.* **(2006)** *71*, 5837-5839.
- 50. Kumar, A., Ye, G., Wang, Y., Lin, X., Sun, G., <u>Parang, K.</u> Synthesis and structure-activity relationships of linear and conformationally constrained peptide analogs of CIYKYY as Src tyrosine kinase inhibitors. *J. Med. Chem.* (2006) *49*, 3395-3401.

- 51. Carballeira, N. M., Sanabria, D., Cruz, C., <u>Parang, K.</u>, Wan, B., Franzblau, S. 2,6-Hexadecadiynoic acid and 2,6-nondecadiynoic acid Novel synthesized acetylenic fatty acids as potent antifungal agents. *Lipids* (2006) *41*, 507-511.
- 52. Ayrapetov, M. K., Wang, Y.-H., Xiaofeng, L., Gu X., <u>Parang, K.</u>, Sun G. Conformational basis for SH2-pTYR527 binding in SRC inactivation. *J. Biol. Chem.* **(2006)** *281*, 23776-23784.
- 53. Kumar, A., Ye, G., Ahmadibeni, Y., <u>Parang, K.</u> Synthesis of polymer-bound 4-acetoxy-3-phenylbenzaldehyde derivatives: Applications in solid-phase organic synthesis. *J. Org. Chem.* **(2006)** *71*, 7915-7918.
- 54. Gu, X., Wang, Y., Kumar, A., Ye, G., <u>Parang, K.</u>, Sun, G. Design and evaluation of hydroxamate derivatives as metal-mediated inhibitors of a protein tyrosine kinase. *J. Med. Chem.* **(2006)** *49*, 7532-7539.
- 55. Agarwal, H. K., <u>Parang, K.</u> Application of solid-phase chemistry for the synthesis of 3'-fluoro-3'-deoxythymidine. *Nucleosides, Nucleotides& Nucleic Acids* (2007) 26, 317-322.
- 56. Ahmadibeni, Y., <u>Parang, K.</u>, Synthesis and evaluation of oligodeoxynucleotides containing diphosphodiester internucleotide linkages. *Angew. Chem. Int. Ed.* **(2007)** *46*, 4739-4743.
- 57. Kumar, A., Wang, Y., Lin, X., Sun, G., <u>Parang, K.</u> Synthesis and evaluation of 3-phenylpyrazolopyrimidine-peptide conjugates as Src tyrosine kinase inhibitors. *ChemMedChem* (2007) 2, 1346-1360.
- 58. Carballeira, N. M., O'Neil, R., <u>Parang, K.</u> Synthesis and antifungal properties of alphamethoxy and alpha-hydroxyl substituted 4-thiatetradecanoic acids. *Chem. Phys. Lipids* (2007) *150*, 82-88.
- 59. Ahmadibeni, Y., Hanley, M., White, M., Ayrapetov, M., Lin, X., Sun, G., <u>Parang, K.</u> Metal-binding properties of a dicysteine-containing motif in protein tyrosine kinases. *ChemBioChem* (2007) *8*, 1592-1605.
- 60. Ye, G., Nam, N. H., Saleh, A., Kumar, A., Sun, G., Shenoy, D. B., Amiji, M. M., **Parang**, **K.** Synthesis and evaluation of tripodal peptide analogues for cellular delivery of phosphopeptides. *J. Med. Chem.* **(2007)** *50*, 3604-3617.
- 61. Chimalakonda, C., Agarwal, H., Kumar, A., <u>Parang, K.</u>, Mehvar, R. Synthesis, analysis, in vitro characterization, and in vivo disposition of a lamivudine-dextran conjugate for selective antiviral delivery to the liver. *Bioconj. Chem.* **(2007)** *18*, 2097-2108.
- 62. Bhandari, R., Saiardi, A., Ahmadibeni, Y., Snowman, A. M., Resnick, A. C., Kristiansen, T. Z., Molina, H., Pandey, A., Werner, Jr. J. K., Juluri, K. R., Xu, Y., Prestwich, G. D., Parang, K., Snyder, S. H. Protein pyrophosphorylation by inositol pyrophosphates is a posttranslational event. *Proc. Nat. Acad. Sci. U.S.A.* (2007) 104, 15305-15310.

- 63. Ahmadibeni, Y., <u>Parang, K.</u> Solid-phase synthesis of symmetrical 5',5'-dinucleoside mono-, di-, tri-, and tetraphosphodiesters. *Org. Lett.* **(2007)** *9*, 4483-4486.
- 64. Penugonda, S., Kumar, A., Agarwal, H. K., <u>Parang, K.</u>, Mehvar, R. Synthesis and *in vitro* characterization of novel dextran-methylprednisolone conjugates with peptide linkers: Effects of linker length on hydrolytic and enzymatic release of methylprednisolone and its peptidyl intermediates. *J. Pharm. Sci.* (2008) *97*, 2649-2664.
- 65. Ahmadibeni, Y., <u>Parang, K.</u> Solid-supported diphosphitylating and triphosphitylating reagents for nucleoside modification. *Current Protocols in Nucleic Acid Chemistry* (2008) Chapter 13:Unit13.8.1-13.8.29.
- 66. Ye, G., Tiwari, R., <u>Parang K.</u> Development of Src tyrosine kinase substrate binding site inhibitors. *Current Opinions Investigational Drugs* **(2008)** *9*, 605-613.
- 67. Agarwal, H. K., Doncel, G., <u>Parang, K.</u> Synthesis and anti-HIV activities of phosphate triester derivatives of 3'-fluoro-2',3'-dideoxythymidine and 3'-azido-2',3'-dideoxythymidine. *Tetrahedron Lett.* (2008) 49, 4905-4907.
- 68. Ahmadibeni, Y., <u>Parang K.</u> Symmetrical dinucleosides. *Synfacts* (2008) 2, 0207 (highlight publication).
- 69. Ahmadibeni, Y., <u>Parang, K.</u> Solid-supported reagents for synthesis of nucleoside monothiophosphates, dithiodiphosphates, and trithiotriphosphates. *Current Protocols in Nucleic Acid Chemistry* (2009) Chapter 13:Unit13.9.
- 70. Ahmadibeni, Y., Tiwari, R., Sun, G., <u>Parang, K.</u> Synthesis of nucleoside mono-, di-, and triphosphoramidates from solid-phase cycloSaligenyl phosphitylating reagents. *Org. Lett.* **(2009)** *11*, 2157-2160.
- 71. Ye, G., Schuler, A., Ahmadibeni, Y., Morgan, J. R., Faruqui, A., Huang, K., Sun, G., Zebala, J. A., **Parang, K.** Synthesis and evaluation of peptides containing iminodiacetate groups as binding ligands of the Src SH2 domain. *Bioorg. Chem.* (2009) 37, 133-142.
- 72. Carballeira, N. M., Miranda, C., <u>Parang, K.</u> The first total synthesis of the (±)-4-methoxydecanoic acid: a novel antifungal fatty acid. *Tetrahedron Lett.* **(2009)** *50*, 5699-5700.
- 73. Tiwari, R., <u>Parang K.</u> Protein conjugates of SH3 domain ligands and ATP-competitive inhibitors as bivalent inhibitors of protein kinases. *ChemBioChem* (2009) *10*, 2445-2448.
- 74. Ahmadibeni, Y., Dash, C., Grice, S. F., <u>Parang, K.</u> 5'-O-β,γ-Methylenetriphosphate derivatives of nucleoside. *Synfacts* **(2010)** *8*, 0961 (highlight publication).

- 75. Penugonda, S., Agarwal, H. K., <u>Parang, K.</u>, Mehvar, R. Plasma pharmacokinetics and tissue disposition of novel dextran-methylprednisolone conjugates with peptide linkers in rats. *J. Pharmaceutical Sci.* **(2010)**, *99*, 1627-1637.
- 76. Ye, G., Gupta, A., DeLuca, R., <u>Parang, K.</u>, Bothun, G. D. Bilayer disruption and liposome restructuring by a homologous series of small Arg. Rich synthetic peptides. *Colloids and Surfaces B.: Biointerfaces* (2010) 76, 76-81.
- 77. Ahmadibeni, Y., Dash, C., Hanley, M. J., Le Grice, S. F. J., Agarwal, H. K., <u>Parang K.</u> Synthesis of nucleoside 5'-O-α,β-methylene-β-triphosphates and evaluation of their potency towards inhibition of HIV-1 reverse transcriptase. *Org. Biomol. Chem.* (2010) 8, 1271-1274.
- 78. Tiwari, R., Brown, A., Narramaneni, S., Sun, G., <u>Parang, K.</u> Synthesis and evaluation of conformationally constrained peptide analogues as the Src SH3 domain binding ligands. *Biochimie* (2010) *92*, 1153-1163.
- 79. Ahmadibeni, Y., Dash, C., Le Grice, S. F. J., <u>Parang K.</u> Solid-phase synthesis of 5'-*O*-β,γ-methylenetriphosphate derivatives of nucleosides and evaluation of their inhibitory activity against HIV-1 reverse transcriptase. *Tetrahedron Lett.* **(2010)** *51*, 3010-3013.
- 80. Sharma, D., Bhatia, S., Sharma, R. K., Tiwari, R., Olsen, C. E., Mandal, D., Lehmann, J., <u>Parang, K.</u>, Parmar, V. S., Prasad, A. S. Synthesis, Src kinase inhibitory and anticancer activities of 1-substituted 3-(*N*-alkyl-*N*-phenylamino)propane-2-ols. *Biochimie* (2010) 92, 1164-1172.
- 81. Chhikara, B. S., Mandal, D., <u>Parang, K.</u> Synthesis and evaluation of fatty acyl ester derivatives of cytarabine as anti-leukemia agents. *Eur. J. Med. Chem.* **(2010)** *45*, 4601-4608.
- 82. Agarwal, H. K., Kumar, A., Doncel, G. F., <u>Parang K.</u> Synthesis, antiviral and contraceptive activities of nucleoside-sodium cellulose sulfate acetate and succinate conjugates. *Bioorg. Med. Chem. Lett.* (2010) 20, 6993-6997.
- 83. Chhikara, B. S., <u>Parang, K.</u> Development of cytarabine prodrugs and delivery systems for leukemia treatment. *Expert Opin. Drug Del.* **(2010)** 7, 1399-1414.
- 84. **Parang, K.**, Sun, G. Protein kinase inhibitors in drug discovery. *Pharmaceutical Sciences Encyclopedia: Drug Discovery, Development, and Manufacturing* **(2010)** John Wiley & Sons, Inc.
- 85. Kumar, D., Buchi Reddy, V., Kumar, A., Mandal, D., Tiwari, R., <u>Parang, K.</u> Click chemistry inspired one-pot synthesis of 1,4-disubstituted 1,2,3-triazoles and their Src kinase inhibitory activity. *Bioorg. Med. Chem. Lett.* (2011) 2, 449-452.
- 86. Kumar, A., Ahmad, I., Chhikara, B. S., Tiwari, R., Mandal, D., <u>Parang, K.</u> Synthesis of 3-phenylpyrazolopyrimidine-1,2,3-triazole conjugates and evaluation of their Src kinase

- inhibitory and anticancer activities. Bioorg. Med. Chem. Lett. (2011) 21, 1342-1346.
- 87. Ahmadibeni, Y., Tiwari, R., Swepson, C., Pandhare, J., Dash, C., Doncel, G. F., Parang, K. Synthesis and anti-HIV activities of bis-(cycloSaligenyl) pronucleotides derivatives of 3'-fluoro-3'-deoxythymidine and 3'-azido-3'-deoxythymidine. *Tetrahedron Lett.* (2011) *52*, 802-805.
- 88. Agarwal, H. K., Loethan, K., Mandal, D., Doncel, G. F., <u>Parang, K.</u> Synthesis and anti-HIV activities of fatty acyl ester derivatives of 2',3'-didehydro-2',3'-dideoxythymidine. *Bioorg. Med. Chem. Lett.* **(2011)** *21*, 1917-1921.
- 89. Fallah-Tafti, A., Tiwari, R., Shirazi, A. N., Akbarzadeh, T., Mandal, D., Shafiee, A., Parang, K., Foroumadi, A. 4-Aryl-4*H*-chromene-3-carbonitrile derivatives: Evaluation of Src kinase inhibitory and anticancer activities. *Med. Chem.* (2011) 7, 466-472.
- 90. Chhikara, B. S., St. Jean, N., Mandal, D., Kumar, A., <u>Parang, K.</u> Fatty-acyl amide derivatives of doxorubicin: Synthesis and *in vitro* anticancer activities. *Eur. J. Med. Chem.* (2011) 46, 2037-2042.
- 91. Gupta, A., Mandal, D., Ahmadibeni, Y., <u>Parang, K.</u>, Bothun, G. Hydrophobicity drives the non-specific cellular uptake of short cationic peptide ligands. *Eur. Biophysics J.* (2011) *40*, 727-736.
- 92. Rao, V. K., Chhikara, B. S., Nasrolahi Shirazi, A. N., Tiwari, R., **Parang, K.**, Kumar, A. 3-Substitued indoles: One-pot synthesis and evaluation of anticancer and Src kinase inhibitory activities. *Bioorg. Med. Chem. Lett.* **(2011)** *21*, 3511-3514.
- 93. Dash, C., Ahmadibeni, Y., Hanley, M. J., Pandhare, J., Gotte, M., Le Grice, S. F. J., Parang, K. Inhibition of multi-drug resistant HIV-1 reverse transcriptase by nucleoside β-triphosphates. *Bioorg. Med. Chem. Lett.* (2011) 21, 3519-3522.
- 94. Kathuria, A., Jalal, S., Tiwari, R., Nasrolahi Shirazi, A., Gupta, S., Kumar, S., <u>Parang, K.</u>, Harma, S. K. Substituted coumarin derivatives: Synthesis and evaluation of antiproliferative and Src kinase inhibitory activities. *Chemistry Biology Interface* (2011) *1* (2), 279-296.
- 95. Fallah-Tafti, A., Foroumadi, A., Tiwari, R., Shirazi, A. N., Hangauer, D. G., Bu, Y., Akbarzadeh, T., <u>Parang, K.</u>, Shafiee, A. Thiazolyl N-benzyl-substituted acetamide derivatives: Synthesis, Src kinase inhibitory and anticancer activities. *Eur. J. Med. Chem.* (2011) *46*, 4853-4858.
- 96. Mandal, D., Nasrolahi Shirazi, A., <u>Parang, K.</u> Cell-penetrating homochiral cyclic peptides as nuclear-targeting molecular transporters. *Angew. Chem. Int. Ed.* **(2011)** *50*, 9633-9637.
- 97. Rao, M. S., Chhikara, B. S., Tiwari, R., Nasrolahi Shirazi, A., <u>Parang, K.</u>, Kumar, A. Greener synthesis of 2-aminochromenes in ionic liquid and evaluation of their

- antiproliferative activities. Chemistry & Biology Interface (2012) 2, 362-372.
- 98. Rafinejad, A., Fallah-Taftia, A., Tiwari, R., Nasrolahi Shirazi, A., Mandal, D., Shafiee, A., Parang, K., Foroumadi, A., Akbarzadeh, T. 4-Aryl-4H-naphthopyrans derivatives: Onepot synthesis, Evaluation of Src kinase inhibitory and anti-proliferative activities. *DARU* Journal of Pharmaceutical Sciences, 2012, 20, 100.
- 99. Muthyala, M. J., Chhikara, B. S., <u>Parang, K.</u>, Kumar, A. Ionic-liquid-supported 1,5,7-triazabicyclo[4.4.0]dec-5-ene— An efficient and recyclable organocatalyst for Michael addition to α,β-unsaturated ketones. *Can. J. Chem.* **(2012)** *90*, 290-297.
- 100. Agarwal, H. K., Doncel, G. F., <u>Parang K.</u> Synthesis and anti-HIV activities of Suramin conjugates of 3'-fluoro-2',3'-dideoxythymidine and 3'-azido-2',3'-dideoxythymidine. *Med. Chem.* **2012**, *8*, 193-197.
- 101. Rao, M. S., Chhikara, B. S., Tiwari, R., Shirazi, A. N., **Parang, K.**, Kumar, A. Microwave-assisted and scandium triflate catalyzed synthesis of tetrahydrobenzo[α]xanthen-11-ones. *Monatsh. Chem.* **(2012)** *143*, 263-268.
- 102. Rao, V. K., Chhikara, B.S., Tiwari, R., Nasrolahi Shirazi, A., <u>Parang, K.</u>, Kumar, A. One-pot regioselective synthesis of tetrahydroindazolones and evaluation of their anti-proliferative and Src kinase inhibitory activities. *Bioorg. Med. Chem. Lett.* (2012) 22, 410-414.
- 103. Munthayala, M., Chhikara, B., <u>Parang, K.</u>, Kumar, A. Ionic liquid-supported synthesis of sulfonamides and carboxamides. *ACS Combinatorial Science* **(2012)** *14*, 60-65.
- 104. Chhikara, B. S., Mandal, D., <u>Parang, K.</u> Synthesis, anticancer activities, and cellular uptake studies of lipophilic derivatives of doxorubicin succinate. *J. Med. Chem.* **(2012)** *55*, 1500-1510.
- 105. Agarwal, H. K., Chhikara, B. S., Quiterio, M., Doncel, G. F., <u>Parang, K.</u> Synthesis and anti-HIV activities of glutamate and peptide conjugates of nucleoside reverse transcriptase inhibitors. *J. Med. Chem.* (2012) *55*, 2672-2687.
- 106. Agarwal, H. K., Chhikara, B. S., Hanley, M. J., Ye, G., Doncel, G. F., <u>Parang, K.</u> Synthesis and biological evaluation of fatty acyl ester derivatives of (-)-2',3'-dideoxy-3'-thiacytidine. *J. Med. Chem.* **(2012)** *55*, 4861-4871.
- 107. Agarwal, H. K., Buckheit, K. W., Buckheit, R. W. Jr, <u>Parang, K.</u> Synthesis and anti-HIV activities of symmetrical dicarboxylate esters of dinucleoside reverse transcriptase inhibitors. *Bioorg. Med. Chem. Lett.* (2012) *22*, 5451-5454.
- 108. Shaik, I. H., Agarwal, H. K., <u>Parang, K.</u>, Mehvar, R. Hepatic immunosuppressive effects of systemically administered novel dextran-methylprednisolone prodrugs with peptide linkers in rats. *J. Pharm. Sci.* **(2012)** *101*, 4003-4012.

- 109. Tiwari R. K., <u>Parang, K.</u> Conformationally constrained peptides as protein tyrosine kinase inhibitors. *Curr. Pharm. Des.* (2012) *18*, 2852-2866.
- 110. Chhikara, B. S., Tiwari, R., <u>Parang, K.</u> *N*-Myristoylglutamic acid derivative of 3'-fluoro-3'-deoxythymidine as an organogel. *Tetrahedron Lett.* **(2012)** *53*, 5335-5337.
- 111. Sharma, R. K., Singh, S., Tiwari, R., Mandal, D., Olsen, C. E., Parmar, V. S., <u>Parang</u>, <u>K.</u>, Prasad, A. K. O-Aryl α,β-D-ribofuranoside: Synthesis & highly efficient biocatalytic separation of anomers and evaluation of their Src kinase inhibitory activity. *Bioorg. Med. Chem.* (2012) *20*, 6821-6830.
- 112. Kumar, A., Muthyala, M. K., Choudhary, S., Tiwari, R. K., <u>Parang, K.</u> Ionic liquid as soluble support for synthesis of 1,2,3-thiadiazoles and 1,2,3-selenadiazoles. *J. Org. Chem.* (2012) 77, 9391-9396.
- 113. Nasrolahi Shirazi, A., Mandal, D., Tiwari, R. K., Guo, L., Lu, W., <u>Parang, K.</u> Cyclic peptide-capped gold nanoparticles as drug delivery systems. *Molecular Pharmaceutics* (2013), *10*, 500-511.
- 114. Agarwal, H. K., Chhikara, B. S., Bhavaraju, S., Mandal, D., Doncel, G. F., <u>Parang, K.</u> Emtricitabine prodrugs with improved anti-HIV activity and cellular uptake. *Molecular Pharmaceutics* (2013) *10*, 467-476.
- 115. Nasrolahi Shirazi, A., Tiwari, R. K., Chhikara, B. S., Mandal, D., <u>Parang, K.</u> Design and evaluation of cell-penetrating peptide-doxorubicin conjugates as prodrugs. *Molecular Pharmaceutics* (2013) *10*, 488-499.
- 116. Chand, K., Nasrolahi Shirazi, A., Yadav, P., Tiwari, R. K., Kumari, M., <u>Parang K.</u>, Sharma, S. K. Synthesis, antiproliferative and c-Src kinase inhibitory activities of cinnamoyl- and pyranochromen-2-one derivatives. *Can. J. Chem.* (2013) *91*, 741-754.
- 117. Cao, C., Rioult-Pedotti, M. S., Migani, P., Yu, C. J., Tiwari, R., <u>Parang, K.</u>, Spaller, M. R., Goebel, D. J., Marshall, J., Impairment of TrkB-PSD-95 signaling in Angelman syndrome, *PLoS Biology*, **(2013)** *11*, e1001478.
- 118. Nasrolahi Shirazi, A., Tiwari, R. K., Oh, D, Banerjee, A, Yadav, A, <u>Parang, K.</u> Efficient delivery of cell impermeable phosphopeptides by a cyclic peptide amphiphile containing tryptophan and arginine. *Mol. Pharm.* **(2013)** *10*, 2008-2020.
- 119. Nasrolahi Shirazi, A., Tiwari, R. K., Brown, A., Mandal, D., Sun, G., <u>Parang, K.</u> Cyclic peptides containing tryptophan and arginine as Src kinase inhibitors. *Bioorg. Med. Chem. Lett.* **(2013)** *23*, 3230-3234.

- 120. Kameshwara Rao V., Shelke, G. M., Tiwari, R., <u>Parang, K.</u>, Kumar A. A simple and efficient synthesis of 2,3-diarynaphthofurans using sequential hydroarylation/Heck oxyarylation. *Org. Lett.* **(2013)** *15*, 2190-2193.
- 121. Kumar, A., Ye, G., Gu, X., Wang, Y., Sun, G., <u>Parang K.</u> Synthesis of pyrazolo[3,4-d]pyrimidine derivatives and evaluation of their Src kinase inhibitory activities. *Chemistry & Biology Interface* (2013) 3, 264-269.
- 122. Chand, K., Tiwari, R. K., Kumar, S., Shirazi, A. N., Sharma, S., Eycken, E. V. V. D, Parmar, V. S., <u>Parang, K.</u>, Sharma, S. K. Synthesis, antiproliferative, and c-Src kinase inhibitory activities of chromone derivatives. *J. Heterocyclic Chemistry* (2013) DOI 10.1002/jhet.2106.
- 123. Chand, K.; Shirazi, A. N.; Yadav, P.; Tiwari, R. K.; Kumari, M.; <u>Parang, K.</u>; Sharma, Sunil K. *Canadian Journal of Chemistry* (2013) 91, 741-754.
- 124. Kumar, A., Rao, V. K., Tiwari, R., Chhikara, B. S., Shirazi, A. N., <u>Parang, K.</u> Copper triflate-mediated synthesis of 1,3,5-triarylpyrazoles in [bmim][PF6] ionic liquid and evaluation of their anticancer activities. *RSC Adv.* (2013), 3, 15396-15403.
- 125. Nasrolahi Shirazi, *A.,* Tiwari, R. K., Mandal, D., <u>Parang, K.</u> Surface decorated gold nanoparticles by linear and cyclic peptides as molecular transporters. *Molecular Pharmaceutics.* **(2013)** *10*, 3137-3151.
- 126. Pericherla, K., Nasrolahi Shirazi, Rao, A. V. K., Tiwari, R., DaSilva, N., Mccaffrey, K. T. Seeram, N., Parang. K., Kumar A., Synthesis and antiproliferative activities of Quebecol and its analogs. *Bioorg. Med. Chem. Lett.* (2013) 23, 5329-5331.
- 127. Mandal, D., Tiwari, R. K., Nasrolahi Shirazi, A., Ye, G., Banerjee, A., Yadav, A., Parang. K.. Self-assembled surfactant cyclic peptide nanostructures as stabilizing agents. Soft Matter. (2013) 9. 9465-9475.
- 128. Pericherla K., Poonam Khedar, P., Khungar, B., <u>Parang, K.,</u> Kumar, A. Copper catalyzed tandem oxidative C–H amination/cyclizations: Direct access to imidazo[1,2-a]pyridines. *RSC Adv.* (2013) 3, 18923-18930.
- 129. Shelke, G. M., Rao, V. K., Tiwari, R., Chhikara, B. S., <u>Parang K.</u>, Kumar A. Bismuth triflate-catalyzed condensation of indoles with acetone. *RSC Adv.* (**2013**) *3*, 22346-22352.

- 130. Suresh, N., Nagesh, H. N., Sekhar, K., V. G. C., Kumar, A., Shirazi, A. N., <u>Parang K.</u> Synthesis of novel ciprofloxacin analogues and evaluation of their anti-proliferative effect on human cancer cell lines. *Bioorg. Med. Chem. Lett.* (2013) 23, 6292-6295.
- 131. Nasrolahi Shirazi, A., Oh, D., Tiwari, R. K., Sullivan, B., Gupta, A., Bothun, G. D., Parang, K. Peptide amphiphile containing arginine and fatty acyl chains as molecular transporters. *Mol. Pharmaceutics* (2013) 10, 4717-4727.
- 132. Yoon, Y. K., Ali, M., A., Wei, A. C., Choon, T. S., Osman, H., <u>Parang, K.,</u> Shirazi, A. M. Synthesis and evaluation of novel benzimidazole derivatives as sirtuin inhibitors with antitumor activities. *Bioorg. Med. Chem.* (2014) 22, 703-710.
- 133. El-Sayed, N.S., Shirazi, N. A., El-Meligy, M. G., El-Ziaty, A. K., Nagib, Z. A., <u>Parang, K.</u> Synthesis of 4-aryl-6-indolylpyridine-3-carbonitriles and evaluation of their antiproliferative activity. *Tetrahedron Lett.* **(2014)** *55*, 1154-1158.
- 134. Motavallizadeh, S., Fallah-Tafti, A., Maleki, S., Nasrolahi Shirazi, A., Pordeli, M., Safavi, M., Kabudanian Ardestani, S., Asd, S., Tiwari, R., Oh, D., Shafiee, A., Foroumadi, A., Parang, K., Akbarzadeh T., Synthesis and evaluation of cytotoxic activity of substituted N-(9-oxo-9H-xanthen-4-yl)benzenesulfonamides. *Tet. Lett.* (2014) 55, 373-375.
- 135. Ghasemi Pirbalouti, A, Sajjadi, S. E., <u>Parang, K.</u> A Review (Research and Patents) on jasmonic acid and its derivatives. *Arch Pharm (Weinheim).* (2014) *347*, 229-239.
- Pemmaraju, B., Agarwal, H. K., Oh, D., Buckheit, K. W., Buckheit Jr. R. W., Tiwari, R., <u>Parang, K.</u> Synthesis and biological evaluation of 5'-O-dicarboxylic fatty acyl monoester derivatives of anti-HIV nucleoside reverse transcriptase inhibitors. *Tetrahedron Lett.* (2014) 55, 1983-1986.
- 137. Chand, K., Prasad, S., Tiwari, R. K., Shirazi, A. N., Kumar, S., Parang, K., Sharma, S. K. Synthesis and Evaluation of c-Src Kinase Inhibitory Activity of Pyridin-2(1*H*)-one Derivatives, *Bioorganic Chem.* **(2014)** *53*, 75-82.
- 138. Verma, A. K., Patel, M., Joshi, M., Likhar, P. R., Tiwari, R. K., <u>Parang, K.</u> Base-mediated chemo- and stereoselective addition of 5-aminoindole/tryptamine and histamines onto alkynes. *J. Org. Chem.* **(2014)** *79*, 172-186.
- 139. Mandal, D., Nasrolahi Shirazi, A., Parang, K. Self-assembly of peptides to nanostructures. *Org. Biomol. Chem.* **2014**, **DOI:** 10.1039/C4OB00447G.

PATENTS

- 1. Courtney A., Cole, P, A., <u>Parang, K.,</u> Ablooglu, A., Kohanski, R. Bisubstrate inhibitors of kinases. *PCT Int. Appl.* **2001**, WO 20010170770, *US Patent* 7,045,617 B2, May **2006**.
- 2. Parang, K., Nam, N. H. Sardari, S. Preparation of azole monosaccharide as antifungal agents. *PCT Int. Appl.* **2005**, WO 2005006860, *U.S. Patent* 7,351,731, April **2008**.
- 3. Parang, K., Nam, N. H., Anil, K., Sun, G. Bisubstrate inhibitors of protein tyrosine kinases as therapeutic agents. *PCT Int. Appl.* **2005**, WO 2005117932, *U.S. Patent* 7,799,753 September **2010**.
- Parang, K., Ahmadibeni, Y. Solid-phase synthesis of modified oligonucleotides containing diphosphodiester inter-nucleotide linkages. PCT Int. Appl. 2007, WO 2007030227; Polymer-Bound Phosphitylating Reagents for the Synthesis of Organophosphorus Compounds, US patent 8,193,384 B2, June 5, 2012.
- 5. **Parang, K.**, Doncel, G., Agarwal, H. K. Substituted nucleoside derivatives with antiviral and antimicrobial properties. *PCT Int. Appl.* **2009** WO 2009009625.
- 6. Seeram, N. P., Barbeau, J., Beland, G., <u>Parang, K.</u>, Preparation of quebecol and its analogs as anti-cancer agents. *PCT Int. Appl.* **2012**, WO 2012167364.

PRESENTATIONS AND PUBLISHED MEETING ABSTRACTS

- 1. Shafiee, A., Pirouzzadeh, B., Ghasemian, F., <u>Parang, K.</u> A convenient synthesis of 2-acetyl-1-methyl-5-nitroimidazole. *Abstracts of papers of the American Chemical Society Meeting*, 202:200-ORGN, Part 2 Aug. **1991**.
- 2. <u>Parang, K.</u>, Knaus, E. E., Wiebe, L. I. Synthesis and biological evaluation of 5'-O-myristoyl derivatives of 3'-azido-3'-deoxythymidine (AZT) and 3'-fluoro-3'-deoxythymidine (FLT) as potential antiviral agents against human immunodeficiency virus (HIV). Page 62, *International Symposium on Lipophilicity in Drug Research and Toxicology*, University of Lausanne, Switzerland, March 21-24, **1995**.
- 3. <u>Parang, K.</u>, Knaus, E. E., Wiebe, L. I., Sardari, S., Daneshtalab, M. Synthesis, antifungal and antiviral activities of myristic acid analogs. Page 46, *Pharmaceutical Research in the 21st Century*, University of Alberta, November 3, **1995**.
- 4. <u>Parang, K.</u>, Wiebe, L. I., Knaus, E. E. Synthesis and antiviral activities of myristic acid analogs and 5'-O-myristoyl derivatives of 3'-azido-3'-deoxythymidine (AZT) and 3'-fluoro-3'-deoxythymidine (FLT). *American Association of Pharmaceutical Scientists (AAPS) Annual Meeting*, Seattle, Washington, September **1996** (*Pharmaceutical Research*, 13, 141, **1996**).
- Parang, K., Knaus, E. E., Wiebe, L. I., Parang, K., Knaus, E. E., Wiebe, L. I. Comparative pharmacokinetics of 3'-azido-3'-deoxythymidine (AZT) and 2', 3'-dideoxy-5'-O-(2-bromomyristoyl)-3'-azidothymidine in mice. *American Association of Pharmaceutical Scientists (AAPS) Annual Meeting*, Boston, 1253, November 1997 (*Pharmaceutical Research*, 14, 79, 1997).
- Parang, K., Knaus, E. E., Wiebe, L. I. Hydrolysis of 5'-O-myristoylesters of 3'-azidodeoxythymidine (AZT) and 3'-fluorodeoxythymidine (FLT) by rat brain homogenate, plasma and porcine liver esterase and their relation to anti-HIV 1 activity. American Association of Pharmaceutical Scientists (AAPS) Annual Meeting, Boston, 2538, November 1997 (Pharmaceutical Research, 14, 389, 1997).
- 7. Parang, K., Wiebe, L. I., Knaus, E. E. Myristic acid derivatives of thymidine targeting HIV-1. American Association of Pharmaceutical Scientists (AAPS) Annual Meeting, 2537, Boston, November 1997 (Pharmaceutical Research, 14, 388, 1997).
- 8. <u>Parang, K.</u>, Hindsgaul, O. New methodology for carbohydrate sulfation. *216th American Chemical Society Meeting*, 216: 242-MEDI, Part 2, Boston, August **1998**.
- 9. <u>Parang, K.</u>, Wiebe, L. I., Knaus, E. E. *In vitro* anti-hepatitis B virus activities of 5'-O-myristoyl analogue derivatives of 3'-fluoro-2', 3'-dideoxythymidine (FLT) and 3'-azido-2',3'-dideoxythymidine (AZT). *216th American Chemical Society Meeting*, 216: 166-MEDI, Part 2, Boston, August **1998**.

- Parang, K., Kim, K., Lau, O. D., Cole, P. A. Substrate selectivity of unnatural tyrosine derivatives for protein tyrosine kinase Csk. 219th American Chemical Society Meeting, 358318, San Francisco, 219: 78-BIOL Part 1, 2000; (Biochemistry, 2000, 39, 78).
- 11. <u>Parang, K.</u>, Fournier, E., Hindsgaul, O. Solid-phase strategy in the monophosphorylation of carbohydrates and nucleosides. *219th American Chemical Society Meeting*, ORGN-170, March 26-30, **2000**, San Francisco.
- 12. <u>Parang, K.</u>, Ablooglu, A. J., Till, J. K., Kim, K., Cole, P. A., Hubbard, S. R., Kohanski, R. A. Probing the catalytic mechanism of the insulin receptor kinase with a tetrafluorotyrosine-containing peptide substrate. *Gordon Conference on Bioorganic Chemistry*, New Hampshire, June 18-23, **2000**.
- 13. <u>Parang, K.</u>, Till, J. H., Ablooglu, A. J., Kohanski, R. A., Hubbard, S. R., Cole, P. A., Mechanism-based design of a protein kinase inhibitor. *New England Pharmacologists 30th Annual Meeting*, Rhode Island, Jan 26-27, **2001**.
- 14. <u>Parang, K.</u>, Till, J. H., Ablooglu, A. J., Kohanski, R. A., Hubbard, S. R., Cole, P. A. Peptide-ATP bisubstrate inhibitors of protein kinases. *Gordon Research Conference on Bioorganic Chemistry*, New Hampshire, June 17-21, **2001**.
- 15. <u>Parang, K.</u> Designing photo-crosslinking reagents for studying protein kinase-substrate interactions. *Gordon Research Conference on Bioorganic Chemistry*, New Hampshire, June 9-14, **2002**.
- 16. Pitts, R., <u>Parang, K.</u>, Sun, G. Design and synthesis of peptide analogues as inhibitors of Src tyrosine kinases. Abstract of Papers, *224th ACS National meeting, Boston*, MA, August 18-22, **2002**, MEDI-107.
- 17. Sardari, S., **Parang, K.** Modeling of activity for biological samples using artificial neural network. Abstracts of Papers, *226th ACS National Meeting*, New York, NY, United States, September 7-11, **2003**, MEDI-352.
- 18. Selecky, M., Nam, N. H., Sardari, S., <u>Parang, K.</u> Novel bisubstrate antifungal agents 1st Annual RI-BRIN Summer Undergraduate Research Day, University of Rhode Island, Kingston, RI, August 19, **2003**.
- Ayrapetov, M., Nam, N. H., <u>Parang, K.,</u> Sun, G. Feud in the Family: Biochemical studies of two protein tyrosine kinases in the Csk Family. *NERMCAP XXVIII*, June 13-14, **2003**, University of Rhode Island.
- 20. Ye, G., <u>Parang, K.</u> Design and solid-phase assay of the Src SH2 domain inhibitors. *NERMCAP XXVIII*, June 13-14, **2003**, University of Rhode Island.
- 21. Nam, H. N., Pitts, R., Sun, G., Sardari, S., Tiemo, A., Xie, M., Yan, B., <u>Parang, K.</u> Design of Tetrapeptide Analogs as Inhibitors of Src SH2 Domain. *Gordon Research Conference on Bioorganic Chemistry*, New Hampshire, June 15-20, **2003**.

- 22. Nam, N. H., Pitts, R., Sardari, S., Tiemo, M., Xie, M., <u>Parang, K.</u> Design of tetrapeptide ligands as inhibitors of the Src SH2 domain. *BRIN Second Annual Retreat*, University of Rhode Island, July 11, **2003**.
- 23. Carballeira, N. M., Oritz, D., <u>Parang, K.</u>, Sardari, S., Goldblum, A., The marine 2-methoxylated fatty acids as new antifungal agents. *44th Annual Meeting of the American Society of Pharmacognosy*, Chapel Hill, NC, July 12-16, **2003**.
- 24. <u>Parang, K.</u>, Ye, G., Sun, G., Avrapetov, M. Design and Solid-phase Assay of Src SH2 Domain Inhibitors. W4086, *Drug Design and Discovery Section of AAPS Annual Meeting*, Baltimore, November 7-11, **2004**.
- 25. <u>Parang, K.</u>, Nam, N.-H., Ye, G., Sun G., Design and Evaluation of Artificial Receptors Mimicking Src SH2 Domain. W4107, *Drug Design and Discovery Section of AAPS Annual Meeting*, Baltimore, November 7-11, **2004**.
- 26. Ahamdibeni, Y., <u>Parang, K.</u> Studying the interactions of toxic metals with protein tyrosine kinases. Joint Toxicology Symposium, *Rhode Island IDEA Network of Biomedical Research Excellence*, W. Alton Jones Campus, June 17, **2005**.
- 27. White, M., Ahmadibeni, Y., <u>Parang, K.</u> Toxic heavy metal interactions with peptides and proteins. *Summer Undergraduate Research Program*, RI-INBRE, University of Rhode Island, August 17, **2005**.
- 28. Hanley, M., Agarwal, H., <u>Parang, K.</u> The design, synthesis, and evaluation of fatty acid prodrugs of lamivudine as anti-HIV-1 microbicides. *Summer Undergraduate Research Program*, RI-INBRE University of Rhode Island, August 17, **2005**.
- 29. Loethen, K, Agarwal, H., <u>Parang, K.</u> Designing prodrugs of antiviral nucleosides for applications as microbicides and targeted hepatic delivery. *Summer Undergraduate Research Program*, RI-INBRE, University of Rhode Island, August 17, **2005**.
- 30. Ahmadibeni, Y., <u>Parang, K.</u> Regioselective solid-phase diphosphorylation and diphosphodithioation of unprotected nucleosides and carbohydrates. *230th ACS National Meeting*, Washington, DC, United States, Aug. 28-Sept. 1, **2005**, ORGN-589.
- 31. Ahmadibeni, Y., <u>Parang, K.</u> Regioselective solid-phase synthesis of dinucleoside and nucleoside-carbohydrate phosphodiesters and thiophosphodiesters using polymer-bound oxathiaphospholanes. *230th ACS National Meeting*, Washington, DC, United States, Aug. 28-Sept. 1, **2005**, ORGN-588.
- 32. Ahmadibeni, Y., <u>Parang, K.</u> Regioselective solid-phase synthesis of nucleosides and carbohydrates triphosphates and triphosphotrithioates. *230th ACS National Meeting*, Washington, DC, United States, Aug. 28-Sept. 1, **2005**, ORGN-587.

- 33. Kumar, A., Wang, Y., Sun, G., <u>Parang, K.</u> Converting a weak peptide inhibitor of Src kinase to potent peptide inhibitors by systemic structural modification. *230th ACS National Meeting*, Washington, DC, United States, Aug. 28-Sept. 1, **2005**, MEDI-405.
- 34. Kumar, A., Wang, Y., Sun, G., <u>Parang, K.</u> N-Heteroaromatic-peptide conjugates as Src kinase inhibitors. *230th ACS National Meeting*, Washington, DC, United States, Aug. 28-Sept. 1, **2005**, MEDI-404.
- 35. Ye, G., Kumar, A., Wang, Y., Sun, G., <u>Parang, K.</u> Conformationally constrained peptide analogs of CIYKYY as inhibitors of Src tyrosine kinase. *230th ACS National Meeting*, Washington, DC, United States, Aug. 28-Sept. 1, **2005**, MEDI-403.
- 36. Gu, X., Wang, Y., Kumar, A., Sun, G., <u>Parang, K.</u> Synthesis and evaluation of hydroxamate derivatives as metal-mediated inhibitors of Csk. *230th ACS National Meeting*, Washington, DC, United States, Aug. 28-Sept. 1, **2005**, MEDI-378.
- 37. Ahmadibeni, Y., <u>Parang, K.</u>, White, M., Sun, G. Studying the interactions of toxic metals with protein tyrosine kinases. *230th ACS National Meeting*, Washington, DC, United States, Aug. 28-Sept. 1, **2005**, MEDI-219.
- 38. <u>Parang, K.</u> Antimycotic drug development. *Rhode Island Showcase: Cutting-edge Medical Technologies, Biotech Research & Discovery*, Providence, RI, February 16, **2006**.
- 39. Hanley, M., **Parang, K.** Solid-phase synthesis of nucleoside 5'-β-triphosphates. *Summer Undergraduate Research Day*, University of Rhode Island, August 10, 2006.
- 40. Kumar, A., Ye, G.; <u>Parang, K.</u> Application of two safety-catch linkers in solid-phase synthesis of sulfonamides. *232nd ACS National Meeting*, San Francisco, CA, United States, Sept. 10-14, **2006**, ORGN-221.
- 41. Ahmadibeni, Y., <u>Parang, K.</u> Synthesis of modified oligonucleotides containing diphosphate diester bridges. *232nd ACS National Meeting*, San Francisco, CA, United States, Sept. 10-14, **2006**, ORGN-218.
- 42. Ahmadibeni, Y., <u>Parang, K.</u> Selective solid-phase β-triphosphorylation of unprotected nucleosides. *232nd ACS National Meeting*, San Francisco, CA, United States, Sept. 10-14, **2006**, ORGN-217.
- 43. Ye, G., Nam, N. H., Kumar, A., <u>Parang, K.</u> Cellular delivery of phosphopeptides by positively-charged tripodal peptides. *232nd ACS National Meeting*, San Francisco, CA, United States, Sept. 10-14, **2006**, MEDI-561.
- 44. Carballeira, N. M., Sanabria, D. <u>Parang, K.</u> 2,6-Hexadecadiynoic acid and 2,6-nonadecadiynoic acid: Novel synthesized acetylenic fatty acids as potent antifungal agents. *232nd ACS National Meeting*, San Francisco, CA, United States, Sept. 10-14, **2006**, MEDI-336.

- 45. Agarwal, H. K.; <u>Parang, K.</u> Solid-phase synthesis of 3'-fluoro-3'-deoxythymidine. *232nd ACS National Meeting*, San Francisco, CA, United States, Sept. 10-14, **2006**, CARB-005.
- 46. Chimalakonda, C., Kumar, A., <u>Parang, K.,</u> Mehvar, R. Simultaneous HPLC analysis of Lamivudine and its succinate in rat plasma. *Annual Meeting of the American Association of Pharmaceutical Scientists*, San Antonio, Texas, October 29-November 2, **2006**.
- 47. Kumar, A., Wang, Y., Sun, G., <u>Parang, K.</u> Bivalent ligand inhibitors of Src kinases. 2nd International Conference on Heterocyclic Chemistry, Jaipur, India December 16-19, **2006**.
- 48. Kumar, A., Ye, G., Wang, Y., Sun, G., <u>Parang, K.</u> Synthesis of linear and conformational constrained peptide inhibitors for Src tyrosine kinase by systematic structural modification. 9th National Symposium in Chemistry, New Delhi, India, February 1-4, **2007**.
- 49. Kumar, A., Ye, G., Wang, Y., Sun, G., <u>Parang, K.</u> Linear and conformational constrained peptide inhibitors for Src tyrosine kinase inhibitors.11th International Conference on Advances in Drug Discovery Research, Aurangabad, India, February 24-26, **2007**.
- 50. Ahmadibeni, Y., Hanley, M., Agarwal, H. K., <u>Parang, K.</u> Synthesis of nucleoside 5'-*O*-β-triphosphates containing alpha-beta methylene bridge. *233rd ACS National Meeting,* Chicago, IL, United States, March 25-29, **2007**, ORGN-641.
- 51. Ahmadibeni, Y., <u>Parang, K.</u> Solid-phase synthesis of symmetrical dinucleoside monophosphodiesters, diphosphodiesters, triphosphodiesters, and tetraphosphodiesters. *233rd ACS National Meeting*, Chicago, IL, United States, March 25-29, **2007**, ORGN-640.
- 52. Ahmadibeni, Y., <u>Parang, K.</u> Solid-phase synthesis of 5'-*O*-(α,β-methylene)triphosphate and 5'-*O*-(β,γ-methylene)triphosphate derivatives of nucleosides. *233rd ACS National Meeting*, Chicago, IL, United States, March 25-29, **2007**, ORGN-639.
- 53. Nam, N. H., Sardari, S., <u>Parang, K.</u> Synthesis and antifungal activities of modified fluconazole derivatives. *233rd ACS National Meeting*, Chicago, IL, United States, March 25-29, **2007**, MEDI-060.
- 54. Ahmadibeni, Y., <u>Parang, K.</u> Interactions of transition metals with cysteine-containing peptides. Abstracts of Papers, *233rd ACS National Meeting*, Chicago, IL, United States, March 25-29, **2007**, INOR-256.
- 55. Agarwal, H. K., Kumar, A., Mehvar, R., <u>Parang, K.</u> Synthesis of nucleoside-succinate-dextran conjugates. *233rd ACS National Meeting*, Chicago, IL, United States, March 25-29, **2007**, CARB-094.
- 56. Kumar, A., Nam, N. H., Agarwal, H. K., Mehvar, R., <u>Parang, K.</u> Synthesis of methylprednisolone-peptide-dextran conjugates. *233rd ACS National Meeting*, Chicago, IL, United States, March 25-29, **2007**, CARB-093.

- 57. Williams, H., Ye, G., <u>Parang, K.</u> Design and synthesis of cyclic octapeptides as nanotube drug delivery system. 2007 Summer Undergraduate Research Fellows Meeting & Faculty Retreat, August 10, **2007**.
- 58. Ahmadibeni, Y., Bhandari, R., Snyder, S. H., <u>Parang, K.</u> Synthesis of pyrophosphopeptides and methylphosphopeptides. *234th ACS National Meeting*, Boston, MA, United States, August 19-23, **2007**, ORGN-903.
- 59. Ahmadibeni, Y., <u>Parang, K.</u> Circular dichroism, thermal denaturation, and nuclease stability studies of modified oligodeoxynucleotides containing mixed sequences of four nucleosides attached through diphosphodiester bridges. *234th ACS National Meeting*, Boston, MA, United States, August 19-23, **2007**, ORGN-151.
- 60. Ye, G., Nam, N. H., Kumar, A., <u>Parang, K.</u> Noncovalent cellular delivery of phosphopeptides by amphipathic peptides. *234th ACS National Meeting*, Boston, MA, United States, August 19-23, **2007**, MEDI-112.
- 61. Agarwal, H. K., Doncel, G. F., <u>Parang, K.</u> Synthesis and evaluation of bis(fatty acylglycol)phosphate triester derivatives of 3'-fluoro-2',3'-dideoxythymidine. *234th ACS National Meeting*, Boston, MA, United States, August 19-23, **2007**, MEDI-097.
- 62. Agarwal, H. K., Hanley, M., Doncel, G. F., <u>Parang, K.</u> Synthesis and anti-HIV activities of fatty acyl derivatives of 2',3'-dideoxy-3'-thiacytidine. *234th ACS National Meeting*, Boston, MA, United States, August 19-23, **2007**, MEDI-096.
- 63. Penugonda, S., Kumar, A., Agarwal, H., <u>Parang, K.</u>, Mehvar, R. Simultaneous HPLC analysis of methylprednisolone, its succinate, and five of its peptidyl derivatives in rat liver lysosomes. *Annual Meeting of the American Association of Pharmaceutical Scientists*, San Diego, California, November 11-15, **2007**.
- 64. Chimalakonda, C., Agarwal, H., Kumar, A., <u>Parang, K.</u>, Mehvar, R. Pharmacokinetics of a liver-targeted dextran prodrug of the antiviral drug Lamivudine in rats. *Annual Meeting of the American Association of Pharmaceutical Scientists*, San Diego, California, November 11-15, **2007**.
- 65. Penugonda, S., Kumar, A., Agarwal, H., <u>Parang, K.</u>, Mehvar, R. In vitro release characteristics of methylprednisolone and its peptidyl intermediates from novel dextranmethylprednisolone prodrugs with peptide linkers of various lengths. *Annual Meeting of the American Association of Pharmaceutical Scientists*, San Diego, California, November 11-15, **2007**.
- 66. Penugonda, S., Naik, B., Kumar, A., Agarwal, H., <u>Parang, K.</u>, Mehvar, R. Effects of linker length on the plasma and tissue disposition of novel dextran-peptide-methylprednisolone prodrugs in rats. *Annual Meeting of the American Association of Pharmaceutical Scientists*, San Diego, California, November 11-15, **2007**.

- 67. Ye, G., <u>Parang, K</u>. Synthesis and evaluation of pyrazolo[3,4-d]pyrimidine derivatives as Src kinase inhibitors. *Annual Meeting of the American Association of Pharmaceutical Scientists*, San Diego, California, November 11-15, **2007**.
- 68. Ahmadibeni, Y., <u>Parang, K.</u> Synthesis and evaluation of dendrimer modified oligodeoxynucleotides containing β-triphosphotriester internucleotide bridges. *236th ACS National Meeting*, Philadelphia, PA, United States, August 17-21, **2008**, ORGN-274.
- 69. Ahmadibeni, Y., Tiwari, R., <u>Parang, K.</u> Application of polymer-bound salicyl alcohol for the synthesis of nucleoside 5'-O-monophosphates. *236th ACS National Meeting*, Philadelphia, PA, United States, August 17-21, **2008**, ORGN-660.
- 70. Agarwal, H. K., Kumar, A., Doncel, G, F., <u>Parang, K.</u> Synthesis of cellulose sulfate-nucleoside conjugates as bifunctional anti-HIV agents. 236th ACS National Meeting, Philadelphia, PA, United States, August 17-21, **2008**, MEDI-442.
- 71. Tiwari, R., Gu, X., Wang, Y., Narramaneni, S., <u>Parang, K.</u>, Sun, G. Development of a potent and specific ligand for the Src SH3 domain. *236th ACS National Meeting*, Philadelphia, PA, United States, August 17-21, **2008**, MEDI-367.
- 72. Agarwal, H. K., Loethan, K. A., Doncel, G. F., <u>Parang, K.</u> Synthesis and anti-HIV activity of fatty acyl derivatives of Stavudine. *236th ACS National Meeting*, Philadelphia, PA, United States, August 17-21, **2008**, MEDI-325.
- 73. Lehmann, J., Kumar, A., <u>Parang K.</u> Synthesis and Evaluation of modified nucleoside oligomers containing 1,2,3-triazole internucleoside linkages as antisense agents. *Summer Undergraduate Research Fellows Conference*, University of Rhode Island, August 4, **2008**.
- 74. Ye, G., Schuler, A. D., Ahmadibeni, Y., Morgan, J. R., Faruqui, A., Zebala, J., <u>Parang, K.</u> Synthesis and evaluation of peptides containing iminodiacetate groups as binding ligands of the Src SH2 domain. *236th ACS National Meeting*, Philadelphia, PA, United States, August 17-21, **2008**, MEDI-043.
- 75. Ahmadibeni, Y., <u>Parang, K</u>. Laboratory teaching of solid-phase reactions, *60th Southeast Regional Meeting of the American Chemical Society*, Nashville, TN, United States, November 12-15, **2008**, SERM-552.
- 76. Ahmadibeni, Y., Tiwari, R., <u>Parang, K.</u> Application of polymer-bound salicyl alcohol for the synthesis of nucleoside mono-, di-, and triphosphoramidates. *238th ACS National Meeting*, Washington, DC, United States, August 16-20, **2009**, ORGN-424.
- 77. Ahmadibeni, Y., Tiwari, R., <u>Parang, K.</u> Using polymer-bound bi(cycloSaligenyl) phosphitylating reagents for the synthesis of nucleoside mono-, di-, and triphosphates. *238th ACS National Meeting*, Washington, DC, United States, August 16-20, **2009**, ORGN-524.

- 78. Mandal, D., Ye, G., Gupta, A., Deluca, R., Bothun, G., <u>Parang, K.</u> Amphipathic peptides containing arginine residues and hydrophobic linkers: Synthesis, cellular uptake, and interactions with phospholipid bilayer membranes. *238th ACS National Meeting*, Washington, DC, United States, August 16-20, **2009**, MEDI-190.
- 79. Ahmadibeni, Y., Tiwari, R., Doncel, G. F., <u>Parang, K.</u> Bis(cycloSaligenyl) derivatives of nucleosides: Synthesis and anti-HIV activities. *238th ACS National Meeting*, Washington, DC, United States, August 16-20, **2009**, MEDI-114.
- 80. Agarawal, H. K., Doncel, G. F., <u>Parang K.</u> Synthesis and anti-HIV activity of glutamate-nucleoside conjugates. *238th ACS National Meeting*, Washington, DC, United States, August 16-20, **2009**, MEDI-113.
- 81. Ahmadibeni, Y., Tiwari, R., <u>Parang, K.</u> Tris(2-(hydroxymethyl)phenol) derivatives: Synthesis and application as acid/base indicators. *238th ACS National Meeting*, Washington, DC, United States, August 16-20, **2009**, ANYL-117.
- 82. Gupta, A., Bothun, G., Deluca, R., Ye, G., Mandal, D., Parang, K. Effects of synthetic linear and cyclic peptide analogs on liposome phase behavior, transport, and morphology. *AIChE Annual meeting*, Nashville, **2009**, 502a.
- 83. Carroll, B., Ahmadibeni, Y., <u>Parang, K.</u> Design of lipophilic bifunctional masked nucleoside monophosphates as anti-HIV agents. *2009 Rhode Island Summer Undergraduate Research Fellowship Conference*, University of Rhode Island, August, 3, 2009.
- 84. <u>Parang, K</u> Synthesis and evaluation of pyrazolo[3,4-d]pyrimidine derivatives as Src kinase inhibitors. *Harry and Elsa Jiler American Cancer Society Professors Meeting*, Naples, Florida, November 11-4, **2009**.
- 85. <u>Parang, K.</u> Design and evaluation of bifunctional anti-HIV-1 agents as microbicides, *Trends in Microbicide Formulation*, Arlington, Virginia, January **2010**.
- 86. Gupta, A., Deluca, R., Ye, G., Mandal, D., <u>Parang, K.</u>, Bothun, G. Membrane activity of novel cationic, amphiphilic, and lipophilic cell penetrating peptides. *Dr. David M. Dooley Inaugural Research and Scholarship Poster Presentation*, University of Rhode Island, April 7, 2010.
- 87. Gagnon, A., K., Tiwari, R., <u>Parang, K.</u> Design and evaluation of self-assembled compounds as drug delivery vehicles. *2010 Rhode Island Summer Undergraduate Research Fellowship Conference*, University of Rhode Island, July 30, 2010.
- 88. Ahmadibeni, Y., Tiwari, R., Doncel, G. F., <u>Parang, K.</u> Synthesis of bis(cycloSaligenyl) pronucleotides containing two nucleosides from bis(methoxymethyl)benzene-1,4-diol. *240th ACS National Meeting*, Boston, MA, United States, August 22-26, **2010**, ORGN-1075.

- 89. Ahmadibeni, Y., Carroll, B., Doncel, G. F., <u>Parang, K.</u> Synthesis of lipophilic dodecyl bis(acylSaligenyl) nucleoside monophosphate triesters. *240th ACS National Meeting*, Boston, MA, United States, August 22-26, **2010**, ORGN-1019.
- 90. Chhikara, B. S., Mandal D., <u>Parang, K.</u> Synthesis and anticancer activity of fatty acid derivatives of cytarabine, *240th ACS National Meeting*, Boston, MA, United States, August 22-26, **2010**, MEMDI-49.
- 91. Mandal, D., <u>Parang, K.</u> Self-assembly of cyclic peptides and potential application as molecular transporters. *240th ACS National Meeting*, Boston, MA, United States, August 22-26, **2010**, ORG-107.
- 92. Tiwari, R., Brown, A., Narramaneni, S., Sun, G., <u>Parang, K.</u> Conformationally constrained peptides as the Src SH3 domain binding ligands. *240th ACS National Meeting*, Boston, MA, United States, August 22-26, **2010**, MEDI-230.
- 93. Tamami, B., **Parang, K.**, Shirazi, A.N. Polystyrene supported aluminum chloride as an efficient and reusable catalyst for one-pot preparation of α-aminophosphonates via three component coupling reaction of aldehydes, amines, and diethyl phosphite. *240th ACS National Meeting*, Boston, MA, United States, August 22-26, **2010**, ORGN-968.
- 94. <u>Parang, K.</u>, Mandal, D., Nasrolahi Shirazi, A. Gold nanoparticle-capped peptides: Design, characterization, and application in drug delivery. *RI Nanotechnology Showcase*, Providence, RI, April 7, 2011.
- 95. Nasrolahi Shirazi, A., Mandal, D., Tiwari, R., <u>Parang K.</u> Peptide-capped gold nanoparticles: Design, characterization, and their application in drug delivery. *42nd Middle Atlantic Regional Meeting of the American Chemical Society*, College Park, MD, United States, May 21-24, **2011**, MARM-221.
- 96. Muthyala, M. K., Chhikara, B. S., <u>Parang, K.</u>, Kumar, A. [TBDbmim]Br: An efficient and recyclable organocatalyst for Michael addition to α,β-unsaturated ketones, *National symposium on organic synthesis*, IIS University, Jaipur, February 18-19, **2011**, PP-2.
- 97. <u>Parang K.,</u> Mandal, D., Nasrolahi Shirazi, A., Cell-penetrating homochiral peptides as nuclear targeting molecular transporters. *Nature Chemical Biology Symposium*, Oct 20-22, **2011**, Cambridge, MA.
- 98. Priya, B., Singh, B., Malekar, S., Doncel, G., Tiwari, R., Worthen, D., <u>Parang, K.</u> Design, synthesis, antiviral activity, and pre-formulation evaluation of multifunctional poly-L-arginine-fatty acyl derivatives of antiviral nucleosides. T2053, **2011** *AAPS Annual Meeting and Exposition*. Washington, DC.
- 99. Ahmadibeni, Y., Williams, T., Seeram, N., <u>Parang, K.</u> Triphenylmethanol Derivatives: Synthesis and Application as Acid-Base Indicators for Acid-Base Titrations. *63rd Southeast Regional Meeting of the American Chemical Society*, Richmond, VA, United States, October 26-29, **2011**, SERM-239.

- 100. Nasrolahi Shirazi, A. N.; Tiwari, R.; Mandal, D.; <u>Parang, K.</u> Enhanced drug delivery by linear and cyclic peptides containing trypotophan and lysine through capping of gold nanoparticles. *63rd Southeast Regional Meeting of the American Chemical Society*, Richmond, VA, United States, October 26-29, **2011**, SERM-23.
- 101. Northup, K., Oh, D., <u>Parang, K.</u> Cyclic polyarginine peptide-fatty acid conjugates: Synthesis and comparative cellular uptake studies. *2012 Rhode Island Summer Undergraduate Research Fellowship Conference*, University of Rhode Island, July 27, **2012**.
- 102. Nasrolahi Shirazi, A. N.; Tiwari, R.; Oh, D.; Ye, G.; <u>Parang, K.</u> Amphiphilc cyclic peptide [WR]₄ as an efficient transporter of negatively charged phosphopeptides. *43rd Middle Atlantic Regional Meeting of the American Chemical Society*, Baltimore, MD, United States, May 31-June 2, **2012**, MARM-343.
- 103. Nasrolahi Shirazi, A. N.; Tiwari, R.; Mandal, D.; <u>Parang, K.</u> Cyclic and linear homochiral decapeptides containing tryptophan and arginine/lysine residues as Src kinase inhibitors. *43rd Middle Atlantic Regional Meeting of the American Chemical Society*, Baltimore, MD, United States, May 31-June 2, **2012**, MARM-341.
- 104. Nahhas, A. F.; Tiwari, R. K.; <u>Parang, K.</u> Application of self-assembled amphiphilc peptides containing tryptophan, arginine, and glutamic acid for generation of gold nanoparticles. *43rd Middle Atlantic Regional Meeting of the American Chemical Society*, Baltimore, MD, United States, May 31-June 2, **2012**, MARM-264.
- 105. Oh, D.; LaPlante, K. L.; Bobcock, K. M.; <u>Parang, K.</u> Acylated cyclic polyarginine cell-penetrating peptides as molecular transporters and antibacterial agents. *244th ACS National Meeting & Exposition*, Philadelphia, PA, United States, August 19-23, **2012**, MEDI-356.
- 106. Tiwari, R. K.; Brown, A.; Shirazi, A. N.; Sun, G.; <u>Parang, K.</u> Dasatinib-fatty acid conjugates: Synthesis and evaluation of tyrosine kinase inhibitory and anticancer activities. *244th ACS National Meeting & Exposition, Philadelphia*, PA, United States, August 19-23, **2012**, MEDI-326.
- 107. Tiwari, R. K.; Brown, A.; Shirazi, A. N.; Sun, G.; <u>Parang, K.</u> Synthesis and evaluation of dasatinib amino acid derivatives for their anticancer and protein tyrosine kinase activity. *244th ACS National Meeting & Exposition*, Philadelphia, PA, United States, August 19-23, **2012**, MEDI-254.
- 108. Shirazi, A. N.; Tiwari, R.; Chhikara, B. S.; Mandal, D.; <u>Parang, K.</u> Synthesis and evaluation of cell-penetrating peptide-doxorubicin conjugates. *244th ACS National Meeting & Exposition*, Philadelphia, PA, United States, August 19-23, **2012**, MEDI-66.

- 109. Nahhas, A. F.; Tiwari, R. K.; <u>Parang, K.</u> Peptides containing tryptophan and arginine as Src kinase inhibitors. *244th ACS National Meeting & Exposition*, Philadelphia, PA, United States, August 19-23, **2012**, MEDI-54.
- 110. Darwish, S., Tiwari, R., Oh, D., <u>Parang, K.</u> An amphiphillic bicyclic peptide as a cellular delivery agent of phosphopeptides. *1st International Conference on Frontiers in Pharmaceutical Sciences: Global Perspectives*, Kingston, RI, Sept. 28-30, **2012**, C4.
- 111. Oh, D., Northup, K., <u>Parang, K.</u> Cyclic polyarginine peptide-fatty acid conjugates as cell-penetrating molecular transporters. *1st International Conference on Frontiers in Pharmaceutical Sciences: Global Perspectives*, Kingston, RI, Sept. 28-30, **2012**, C18.
- 112. Nasrolahi Shirazi, A., <u>Parang, K.</u> amphiphilic cyclic peptide-selenium nanoparticles: Synthesis and antiproliferative activity. *1st International Conference on Frontiers in Pharmaceutical Sciences: Global Perspectives*, Kingston, RI, Sept. 28-30, **2012**, C26.
- 113. Tiwari, R. K., Brown, A., Nasrolahi Shirazi, A., Boltan, J., Sun, G., <u>Parang, K.</u> Dasatinib-amino acid and dasatinib-fatty acid conjugates: Synthesis, comparative tyrosine kinase inhibition, and antiproliferative activity. *1st International Conference on Frontiers in Pharmaceutical Sciences: Global Perspectives*, Kingston, RI, Sept. 28-30, **2012**, C31.
- 114. Nasrolahi Shirazi, A., Mandal, D., Tiwari, R., Neira, K., Howlett, N., <u>Parang, K.</u> Synthesis of a new generation of cyclic peptides and their applications in biomedical sciences. *The Institute for Molecular and Nanoscale Innovation (IMNI)*, Brown University, November 9, **2012**.
- 115. Beni, Y. A.; Tiwari, R.; Parang, K. Synthesis of tris(cycloSaligenyl) pronucleotides containing three nucleosides from 4,4',4"-methanetriyltris(2-(hydroxymethyl)phenol). 245th ACS National Meeting & Exposition, New Orleans, LA, United States, April 7-11, **2013**, ORGN-808.
- 116. Nasrolahi Shirazi, A.; Neira, K.; Mandal, D.; Howlett, N.; Parang, K. Cyclic peptide-capped gold nanoparticles for enhanced siRNA delivery. 245th ACS National Meeting & Exposition, New Orleans, LA, United States, April 7-11, **2013**, MEDI-352.
- 117. Nasrolahi Shirazi, A., Parang, K. Peptide amphiphiles as new transporters for the delivery of phosphopeptides. 245th ACS National Meeting & Exposition, New Orleans, LA, United States, April 7-11, **2013**, MEDI-118.
- 118. Nasrolahi Shirazi, A., Parang, K. Cyclic peptide-capped selenium nanoparticles as molecular transporters. 246th ACS National Meeting & Exposition, Indianapolis, IN, United States, September 8-12, **2013**, BIOL-147.

- 119. Parang, K, Peptide nanostructures as nolecular transporters of anticancer agents. 16th Annual Chao Family Comprehensive Cancer Center Scientific Retreat, Palm Springs, CA, November 15-16, **2013**.
- 120. Beni, Y. A.; Parang, K. Synthesis and thermal stability evaluation of six-membered cyclic triphosphates containing three nucleotides. 65th Southeast Regional Meeting of the American Chemical Society, Atlanta, GA, United States, November 13-16, **2013**, SERM-204.
- 121. Shirazi, A., Tiwari, R., Parang, K. Cyclic peptide-capped selenium nanoparticles as nano drug delivery systems. Gordon Conference: Chemistry and Biology of Peptides, Ventura, CA, United States, February 23-28, **2014**.
- 122. Parang, K, Peptide nanostructures as nolecular transporters of anticancer agents. Gordon Conference: Chemistry and Biology of Peptides, Ventura, CA, United States, February 23-28, **2014**.

INVITED ORAL PRESENTATIONS

- 1. **Parang, K.** Chemistry and Biology of Phosphorylation, College of Pharmacy, *Long Island University*, February 2000.
- 2. **Parang, K.** Chemistry and Biology of Phosphorylation, College of Pharmacy, *Idaho State University*, April 2000.
- 3. **Parang K.** Chemistry and Biology of Phosphorylation, College of Pharmacy, *University of Illinois at Chicago*, Sept. 8, 2001.
- 4. **Parang, K. Keynote Speaker,** Implementing Mechanism Based Approaches to Protein Kinase Inhibitor Design and Functional Analysis, *Discovery Tech 2001*, November 28, 2001, Palm Beach, Florida.
- 5. **Parang, K.** Novel bifunctional Antifungal Agents, Intellectual Property Committee, *University of Rhode Island*, April 24, 2003.
- 6. **Parang, K. Keynote Speaker,** Designing Bisubstrate Analog Inhibitors for Protein Kinases. *Cambridge Healthtech Institute's Protein Kinase Targets*, Strategies for Drug Development, June 10, 2003, Boston, Massachusetts.
- 7. **Parang, K.** Bisubstrate Inhibitors of Protein Tyrosine Kinases as Anticancer Agents. Intellectual Property Committee, *University of Rhode Island*, August 6, 2003.
- 8. **Parang, K. Keynote Speaker,** Novel Approaches in Designing Protein Kinase Inhibitors. *Genentech*, October 3, 2003, San Francisco, California.
- 9. **Parang, K.** Opportunities in Biology and Chemistry of Phosphorylation for Drug Discovery, *University of Alberta*, Canada, June 10, 2005.
- 10. **Parang, K.** Chirality in Drug-Receptor Interactions, *University of Alberta*, Canada, June 10, 2005.
- 11. **Parang, K.** Chemistry and Biology of Phosphorylation, Department of Chemistry, *University of Rhode Island*, Sept. 12, 2005.
- 12. **Parang, K.** Synthesis of Modified DNA Molecules Containing Diphosphate or Dithiophosphate Bridges Using Novel Diphosphorylating Reagents, Intellectual Property Committee, *University of Rhode Island*, Sept. 16, 2005.
- 13. **Parang, K.** Evaluation of Reactivity and Stability of Peptides and Proteins for Pharmaceutical Formulations, *University of Rhode Island*, November 18, 2005.
- 14. **Parang, K.** Chemistry and Biology of Phosphorylation, Department of Medicinal Chemistry, *University of Utah*, January 19, **2006**.

- 15. **Parang, K.** Chemistry and Biology of Phosphorylation, *Texas Tech School of Pharmacy,* November 20, **2006**.
- 16. Ahmadibeni, Y., Hanley, M., Agarwal, H. K., **Parang, K.** Applications of polymer-bound phosphitylating reagents in the synthesis of organophosphorus compounds. *233rd ACS National Meeting*, Chicago, IL, United States, March 29, **2007**, ORGN-240.
- 17. **Parang, K.** Research Experiences and Opportunities at the Interface of Chemistry and Biology, Research Seminar Series, November 23, **2007**, University of Rhode Island.
- 18. **Parang, K.** Understanding the Regulation, Activation, and Interactions of Protein Tyrosine Kinases for Rational Design of the Next Generation of Inhibitors, *Akron University*, November 27, **2007**.
- 19. **Parang, K.** Research Experiences and Opportunities at the Interface of Chemistry and Biology Moffit Cancer Center, February 6, **2008**.
- 20. **Parang K.**, New Strategies in Designing Protein Kinase Inhibitors. *International Conference on the Interface of Chemistry-Biology in Biomedical Research*, Pilani, India, February 23, **2008** (Invited lecture).
- 21. **Parang, K.** Research Experiences and Opportunities at the Interface of Chemistry and Biology. *The 11th Iranian Pharmaceutical Sciences Conference IPSC2008*, Kerman, August **2008**.
- 22. Gupta, A., Bothun, G. D., Deluca, R., Ye, G., **Parang, K.** Interaction mechanisms between a homologous series of tripodal cationic peptides and lipid bilayer membranes. *American Institute of Chemical Engineers (AIChE)* annual meeting, Philadelphia, November 16, **2008**, 196/1-196/5.
- 23. **Parang, K.** Designing protein tyrosine kinase inhibitors, 13th **ISCBC** International Conference on Interplay of Chemical and Biological Sciences, New Delhi, India, Feb. 28, **2009**.
- 24. **Parang, K.** Design and evaluation of multifunctional anti-HIV agents. Research Seminar Series, University of Rhode Island, July 17, **2009**.
- 25. Ahmadibeni, Y., **Parang, K**. Solid-phase phosphitylating reagents: design and application in the synthesis of phosphorylated biomolecules. 61st Southeast Regional Meeting of the American Chemical Society, San Juan, Puerto Rico, October 21-24, **2009**, SRM-473.
- 26. Gupta, A., Deluca, R., Ye, G., Mandal, D., **Parang, K.**, Bothun, G. D. Lipid vesicles, disks, and interlamellar attachments: Aggregation effects of cell penetrating peptides on zwitterionic-anionic vesicles. 238th ACS National Meeting, Washington, DC, United States, August 16-20, **2009**, COLL-053.

- 27. **Parang, K.** Design and evaluation of multifunctional anti-HIV agents. T3D-2010 International Symposium on Trends in Drug Discovery and Development New Delhi, India, January 5-8, **2010**.
- 28. **Parang, K**. Nucleoside-polymer conjugates as anti-HIV microbicides. Current Trends in Pharmaceutical Research: Focus on Orphan Diseases, Patna, India, January 10, **2010**.
- 29. **Parang K.** Research Experiences and Opportunities at the Interface of Chemistry and Biology, University of Puerto Rico Humacao, April 20, **2010**.
- 30. **Parang K.** Designing Protein Tyrosine Kinase Inhibitors, University of Puerto Rico San Juan, April 21, **2010**.
- 31. **Parang K.**, Agarwal, H., Doncel G. 5'-O-Fatty acyl ester derivatives of 3'-fluoro-2',3'-dideoxythymidine (FLT) as microbicidal agents. M2010 Microbicides, Pittsburgh, May 25, **2010**.
- 32. Gupta, A., Shirazi, A., **Parang K.**, Bothun, G. Self-Assembled Peptide-Amphiphile/Lipid Mixtures Nanoscale Science and Engineering forum session (AIChE 2010 Meeting), November 12, 2010, Salt Lake City, Utah.
- 33. **Parang K.** Peptide nanostructures as molecular transporters of therapeutic Agents. University of Delhi, India, Jan. 31, 2011.
- 34. **Parang K.** Designing protein kinase inhibitors. Delhi Indian Institute of Technology, India, Jan. 31, 2011.
- 35. **Parang K.** Peptide nanostructures as molecular transporters of therapeutic Agents. BITS, Pilani, India, Feb. 2, 2011.
- 36. **Parang K.** Peptide nanostructures as molecular transporters of therapeutic Agents. ISCBC, Rajkot, India, Feb. 5, 2011.
- 37. Nasrolahi Shirazi, A.; Tiwari, R.; Mandal, D.; **Parang, K.** Enhanced drug delivery by linear and cyclic peptides containing trypotophan and lysine through capping of gold nanoparticles. 2011 *Southestern Regional Meeting of the American Chemical Society (SERMACS 2011).* Medicinal Chemistry, *Oct. 26, 2011.*
- 38. Ahmadibeni, Y.; Williams, T.; Seeram, T.; Seeram, N.; **Parang, K.** Triphenylmethanol derivatives: Synthesis and application as acid-base indicators for acid-base titrations. 2011 *Southestern Regional Meeting of the American Chemical Society (SERMACS 2011).* Medicinal Chemistry, *Oct. 27, 2011.*
- 39. **Parang, K.**, Peptide nanostructures as molecular transporters of therapeutic Agents. Rhode Island College. *February 15, 2012*.

- 40. **Parang, K.**, Agarwal, H., Doncel, G. Novel lipophilic fatty acyl derivatives of lamivudine and emtricitabine as potential microbicidal agents. *M2012 International Microbicide Conference*, Sydney, Australia, April 18, 2012.
- 41. **Parang K.** Peptide nanoparticles as molecular transporters of therapeutic agents. Technical University Braunschweig, Germany, May 7, 2012.
- 42. **Parang K.**, Peptide nanostructures as molecular transporters of therapeutic agents. Roger Williams University, February 20, 2013.
- 43. **Parang K.**, Molecular modeling of peptide nano self-assemblies. The Rhode Island Consortium for Nanoscience and Nanotechnology. University of Rhode Island, March 6, 2013.
- 44. **Chikkara, B.**, Parang, K. Fatty acyl and peptidic Adriamycin synthesis and evaluation as new Prodrugs with novel pharmacological profile for cancer treatment*" 3rd Annual National Convention of Association of Pharmacy Professionals' Dharmsinh Desai, University, Nadiad, Gujarat, January 18, 2014.

FUNDED GRANT PROPOSALS

- 1. "Tyrosine Kinase Inhibitors as Anti-Cancer Agents", *Special Research Funding*", University of Rhode Island Research Office, \$3,500 June 2001 for equipment purchase (Role: PI).
- 2. "Investigator, "Bioisosteres of Tyrosine Phosphate as Inhibitors of SH2 Domain of Tyrosine Kinases", *Biomedical Research Infrastructure Network (BRIN) Grant* (P.I.: Dr. Zahir Shaikh), NIH, PR16457-01, 9/30/01-8/31/04 (~\$120,000 for Summer Salary support, travel, student support, supplies).
- 3. "Molecular Modeling Laboratory Initiative", *URI Teaching with Technology Student Assistants Program (TTSA)*, Information and Instructional Technology Services, University of Rhode Island, October 2001, I got support for one student.
- 4. "Designing Prodrugs Against Human Immunodeficiency Virus", *URI Foundation*, \$2,000 for supplies, March 2002 (Role: PI).
- 5. Principal Investigator, "Molecular Modeling Laboratory Initiative", *University of Rhode Island Foundation Competitive Grant Application*, \$4,800, Jan. 1, 2002 through Jan 1, 2003 (Role: PI).
- 6. "Bisubstrate Analogs of Protein Tyrosine Kinases against Cancer", Medical Research Grant, *Rhode Island Foundation*, \$10,000, February 2003-March 2004 (Role: PI).
- 7. "Antimycotic Drug Development", *Slater Center for Biomedical Technology*, May 2001, \$100,101, March 2002-January 2004 (Role: PI).
- 8. "Drug Development against Osteoporosis", *The University of Rhode Island Council for Research*, \$5,910, 08/01/2005-12/31/2005 (Role: PI).
- 9. "Novel Bifunctional Anti-HIV-1 Agents as Microbicides with and without Contraceptive Activity," *American Foundation for Pharmaceutical Education (AFPE)*, \$5,000, 07/01/2005-05/01/2006 (Role: PI).
- 10. "Antimycotic Drug Development", Principal Investigator, *URI Foundation*, \$7,500, December 2005 (Role: PI).
- 11. "Solid-Phase Triphosphorylation of Nucleosides", *Helicos BioSciences Corporation*, \$13,680, September 2006 (Role: PI).
- 12. "Solid-Phase Triphosphitylating Reagents", *Biota Corporation*, \$4,975, September 2006 (Role: PI).
- 13. "Design and Synthesis of Cyclic Octapeptides as Nanotube Drug Delivery System", *Rhode Island Sea Grant and Honors Program*, \$798, 01/8/2008 (Role: PI).
- 14. "Novel Bifunctional Anti-HIV-1 Agents as Microbicides With and Without Spermicide Activity", *Contraceptive Research and Development (CONRAD)*, MSA-03-367, \$751,117,

- 12/01/03-12/31/09 (Role: PI).
- 15. "Synthesis and Development of Novel Neuroprotective Agents", *Brown University/STAC*, \$28,000, 08/01/09-5/31/10 (Role: PI).
- 16. "Synthesis of KP-17", CONRAD, PPA-09-028, \$43,804, 09/01/09-03/31/10 (Role: PI).
- 17. (Co-PI) Establishment of an Animation and 3-D Stereo Visualization Facility, *Champlin Foundations Grant*, Approved Nov. 2003, \$106,000 (PI. Dr. Bongsup Cho).
- 18. Investigator, "Mechanistic Studies of Protein Tyrosine Kinase Activation by Arsenite", INBRE NIH P20 RR016457, \$76,795, 08/1/04-08/01/05.
- 19. (Co-PI, 25% effort, \$178,635), "Mechanistic Studies and Inhibitor Design for Protein Tyrosine Kinases", *American Cancer Society*, RSG CDD-106966 (Total grant \$714,543) 08/01/04-08/01/09 (P.I.: Dr. Gongqin Sun).
- 20. (Co-PI, 15% effort, \$130,000), "Local Immunosuppression for Liver Transplantation", *NIH*, 1R01 GM069869 (Total grant \$866,980) 04/01/05-04/01/09, (PI. Dr. Reza Mehvar).
- 21. (Co-PI, 20% effort, \$172,800), "Mechanism of PTK Substrate Recognition and Specificity", NIH, 1R01CA111687 (Total grant \$864,000) March 2005-December 2009 (PI: Dr. Gongqin Sun).
- 22. (Mentor and Sponsor), "Fatty Acyl Amide Derivatives of Doxorubicin: Synthesis and in vitro Anticancer Activities", Undergraduate Research Initiative (Total grant \$800) 03/01/2010-08/31/2010 (Student Nicole St. Jean).
- 23. (Mentor and Sponsor), "Self-Assembled Cyclic Peptides as Nanostructures", Undergraduate Research Initiative (Total grant \$1,161) 01/01/2010-06/31/2010 (Students Doujia Li, Can Nguyen, Phouthone Malayphone
- 24. (Co-PI, 30% effort, \$31,607), "Controlled Release and Medicated Combination Devices for the Treatment of Bone Fractures and Other Bone Disorders", *IllumiOss Medical, Inc.*, (Total grant \$105,357), 11/01/10-06/01/12 (PI: Dr. David Worthen).
- 25. (Mentor and Sponsor), "Development of Neuroprotective PDZ-Domain Inhibitors for Treatment of Stroke", *American Heart Association*, 11POST5820019 (Total grant \$94,000) 12/31/11-12/31/12 (Postdoc.: Rakesh Tiwari).
- 26. (Mentor and Sponsor), "Designing Cell-Penetrating Cyclic Peptides as Molecular Transporters", Undergraduate Research Initiative (Total grant \$1,200) 03/01/2012-08/31/2012 (Students Brian Sullivan and Kellen McCaffrey).
- 27. "The Tick-Bite Patch", *Rhode Island Science & Technology Advisory Council (STAC)*, RIRA-CA-2011, (Total grant \$199,479) (01/01/2011-12/31/2012) (Pl. Dr. Thomas Mather) (Role: Pl, Multiple Pl grant).

- 28. Treatment of Stroke Using a Novel PDZ Binding Peptidomimetic Drug, NIH, 1R43NS074651-01 (Total grant \$352,596, my portion \$75,723), 09/01/11-08/31/12) (Role: PI, Multiple PI grant).
- 29. "Novel Bifunctional Anti-HIV-1 Agents as Microbicides", CONRAD/USAID, PPA-09-035, \$340,567, 11/01/09-07/01/12 (Role: PI).
- 30. "Development of Src Kinase Inhibitors as Anticancer Agents". *American Cancer Society*, RSG-07-290-01-CDD, \$627,000, 07/01/2007-07/01/2012 (Role: PI).
- 31. "Synthesis of Bicyclic Peptides as Antimicrobial Agents", *Joint Supervision Mission Grant*, Egyptian Cultural Sector- Ministry of Higher Education. \$10,000, 01/01/2012-01/01/2013 (Role: PI).
- 32. "Synthesis and Evaluation of Peptides Synthesized through Click Chemistry", *Programme de bourses Libye-Amérique du Nord*, Academic Manager for the Libyan North American Scholarship, \$24,000 (\$6,000/year), 12/01/2012-12/01/2016 (Role: PI).
- 33. "Design, Green Synthesis And Structure–Activity Relationship Of Novel Indolyl Nicotinonitriles Conjugated To Water Soluble Cellulose Derivative As Tumor Necrosis Factor Alpha (TNF-A) Antagonists For The Treatment of Rheumatoid Arthritis", *Joint Supervision Mission Grant*, Egyptian Cultural Sector- Ministry of Higher Education. \$10,000, 01/15/2013-01/01/2014 (Role: PI).
- 34. (Mentor and Sponsor), "Synthesis and Evaluation of Multifunctional Cyclic Peptide-Doxorubicin-Folic acid Conjugates as Selective Anticancer Agents", Undergraduate Research Initiative (Total grant \$1,200) 01/15/2013-06/31/2013 (Students Brian Sullivan and Kellen McCaffrey).
- 35. "Synthesis of Organophosphorus Compounds Using Solid-Phase Reagents", *NSF*, CHE 0748555, \$300,000, 02/01/2008-02/01/2013 (Role: PI).
- 36. "Synthesis of Cyclic Pyranopterin Monophosphate (cPMP) and the Prodrugs", *Alexion* (Total grant \$139,656) (4/01/2012-10/01/2013) (Role: PI).
- 37. "Molecular Modeling of Peptide Nano Self-Assemblies", RIN^2 , \$8000 (8/31/2012-8/31/2013) (Role: PI).
- 38. "Purification of Cyclic Peptides and Chiral Compounds", *Waters Academic Grant*, \$12,000 (12/15/2012-04/01/2013) (Role: PI).
- 39. "Synthesis of Molybdopterin and its Analogs", \$213,179 Alexion, (8/1/13-10/1/14) (Role: PI).

COLLABORATORS in U.S.

COLLABORATORS AT URI

- 1. Dr. Geoff Bothun, Assistant Professor, Department of Chemical Engineering at University of Rhode Island. Dr. Bothun and I are collaborating in studying the interactions of amphipathic peptides with phospholipid bilayer.
- 2. Dr. Niall Howlett, Associate Professor, Department of Cell and Molecular Biology at University of Rhode Island. Dr. Howlett and I are collaborating on evaluation of peptides as delivery tools for siRNA.
- 3. Dr. Gongqin Sun, Professor, Department of Cell and Molecular Biology at University of Rhode Island: I am a co-PI (25% effort) in a project titled, "Mechanistic Studies and Inhibitor Design for Protein Tyrosine Kinases" funded by *American Cancer Society* and a project titled "Mechanism of PTK substrate recognition and specificity" funded by NIH (1R01CA111687).
- 4. Dr. Wei Lu, Department of Biomedical and Pharmaceutical Sciences at University of Rhode Island. Dr. Lu and I are collaborating on evaluation of cyclic peptides and cyclic peptide-capped gold nanoparticles in xenograft mice model.
- 5. Dr. Navindra Seeram, Department of Biomedical and Pharmaceutical Sciences at University of Rhode Island. Dr. Seeram and I are collaborating on the synthesis of Quebecol.
- 6. Dr. Kerry LaPlante, Department of Biomedical and Pharmaceutical Sciences at University of Rhode Island. Dr. LaPlante and I are collaborating on design and evaluation of amphiphilic peptides as antibacterial agents.
- 7. Dr. David Rowley, Department of Biomedical and Pharmaceutical Sciences at University of Rhode Island. Dr. Rowley and I are working on synthesis and evaluation of antimicrobial peptides.
- 8. Dr. David Worthen, Department of Biomedical and Pharmaceutical Sciences at University of Rhode Island. Dr. Worthen and I are collaborating on determining the stability of a lipophilic anti-HIV agent.
- Dr. Thomas Mather, URI Center for Vector-Borne Disease, University of Rhode Island. Dr. Mather and I are collaborating on design Tick vaccine using peptides synthesized in my laboratory.

COLLABORATORS at OTHER U.S. INSTITUTIONS

- Dr. Gustavo F. Doncel, Professor, Department of Obstetrics and Gynecology, Eastern Virginia Medical School, Norfolk, VA. Dr. Doncel and I are collaborating in design and evaluation of anti-HIV microbicides. Funded by CONRAD.
- 2. Dr. Reza Mehvar, Professor, Texas Tech. College of Pharmacy. Dr. Mehvar (PI) and I are collaborating in a project titled "Local Immunosuppression for Liver Transplantation" funded by NIH (1R01 GM069869).
- Dr. Solomon H. Snyder, Professor, Dept Neuroscience, Johns Hopkins Medical School. Dr. Snyder and I are collaborating in designing pyrophosphate derivatives of IP7 to investigate protein pyrophosphorylation. A manuscript in this subject was published in PNAS (2007, 104, 15305-15310).
- 4. Dr. John Marshall, Department of Molecular Pharmacology, Physiology and Biotechnology, Brown University, "Design and evaluation of peptides as neuroprotective agents".
- 5. Dr. Dennis Goebel, Department of Anatomy and Cell Biology Wayne State University, Dr. Goebel and I are collaborating on design and evaluation of peptides containing polyarginine and cysteine residues as neuroprotective agents.
- 6. Dr. Sujata Bhatia, Bioengineering Labs, Harvard School of Engineering and Applied Sciences. Dr. Bhatia and I are collaborating on designing biomaterials for wound healing and nerve regeneration using self-assembled peptides.
- 7. Dr. Shailendra Giri, Dr. Ramandeep Rattan, Mayo Clinic, Rochester, I am collaborating with Drs. Giri and Rattan in designing compounds for autoimmune diseases.
- 8. Dr. Robert Buckheit, Imquest BioSciences, Inc., I am collaborating with this company in developing anti-HIV microbicides.

INTERNATIONAL COLLABORATORS

- 1. Dr. Ashok Prasad, Department of Chemistry, University of Delhi, India. "Collaboration in the area of the synthesis of anti-cancer agents".
- 2. Dr. Nestor Carballeira, Department of Chemistry, University of Puerto Rico, "Evaluation of modified fatty acids as antifungal agents".
- 3. Dr. Aziz Ghahary, Department of Surgery, University of British Columbia, "Evaluation of self-assembled peptides for tissue engineering".

- 4. Dr. Sunil Sharma, Department of Chemistry, University of Delhi, India. "Collaboration in the area of the synthesis of anti-cancer agents".
- 5. Dr. Akhilesh K. Verma, Department of Chemistry, University of Delhi, India. "Collaboration in the area of the synthesis of anti-cancer agents".
- 6. Dr. Anil Kumar, Department of Chemistry, Birla Institute of Technology, India. Dr. Kumar and I are collaborating with Dr. Seeram for the synthesis of Quebecol, anticancer agents, and ionic liquids.
- 7. Dr. Alireza Foroumadi, Dr. Abbas Shafiee, Dr. Tahmineh Akbarzadeh, College of Pharmacy, Tehran University of Medical Science. Screening of heteroaromatic compounds for kinase inhibition and anticancer activities.
- 8. Dr. Arpita Yadav, University Institute of Engineering & Technology, C.S.J.M. University, Kanpur 208024. Molecular modeling of cyclic peptides and determination of mechanism of self assembly.
- 9. Dr. Dalip Kumar, Department of Chemistry, Birla Institute of Technology, India. Dr. Kumar and I are collaborating in synthesis and evaluation of Src kinase inhibitors.
- 10. Dr. Mohamed Ashraf Ali, Institute for Research in Molecular Medicine, Universiti Sains Malaysia, Malaysia. Dr. Kassim and I are collaborating in the synthesis and evaluation of benzimidazole derivatives.

LEADERSHIP AND MISCELLANEOUS WORK EXPERIENCE

May 2012-July 1, 2013 **Program Coordinator,** Rhode Island IDeA Network

of Biomedical Research Excellence (INBRE)

sponsored by NIH

October 2012-September 30, 2012 International Conference Committee Chair

November 2012-Septemebr 14, 2012 One Day Symposium Committee Member

May 2012 Initiated international collaboration between Technical

University Braunschweig, Germany and the College of Pharmacy after invitation by IEP program. Dean Jordan and Associate Dean Cho accompanied me on this trip. A German delegation was invited to the

International Conference for follow up visit.

February 2011 Initiated and established international collaboration

between Dean of College of Pharmacy and University of Delhi and Indian Birla Institute of Technology and Science. President Dooley and Dean Jordan traveled to India and two MOUs were signed for student and

faculty exchange.

February 2011 Along with Dean Jordan initiated discussion for

establishing research collaboration between the URI College of Pharmacy and the International Center for Chemical and Biological Sciences. One MOU was

signed following our visit.

February 2011 Along with Dean Jordan and Dean Zawia initiated

discussion between the URI College of Pharmacy and

the U.A. E. University

May 2012-present Graduate and Research Committee

January 2011-April 2011 Space committee

September 2010-Jan. 2012 Assessment committee member

September 2010-Jan. 2012 Curriculum committee member

September 2008-Jan. 2010	Scholastic Standing Committee Member
December 2006-June 2007	Serving as a member of evaluation committee for Dean of Pharmacy
January 2006-August 2006	Serving in two search committees for two faculty positions in Department of Biomedical and Pharmaceutical Sciences and Department of Cell and Molecular Biology
March 2004-April 2007	Senator representing the College of Pharmacy in University of Rhode Island Senate
2007	Advisor for Pharmacy professional students, class of 2009
2002-2007	A member of organizing Committee of Annual AAPS- NERDG in Rocky Hill, CT
April 2004-April 2007	Organizing INBRE Seminar Series
Oct. 2000-Presnt	Serving in Graduate Student Thesis Committees: I have been serving in several graduate student thesis committees (totally for 60 graduate students).
May 1996-April 1997	Vice President Operations, HUB Community Association, U. of Alberta
February 1997	Leadership Retreat, University of Alberta
May 1995-April 1997	Sports Unit Manager, University of Alberta
May 1995-April 1997	A member of COSL committee (Council of student life), University of Alberta
August 1996	Leadership Retreat, University of Alberta
September 2-4, 1996	Leadership in Residence, University of Alberta
May 1995-April 1996	Vice President Programming, HUB Community Association, U. of Alberta

Sept. 1995- Dec. 1995 Teaching assistant in Radioisotope Tracer Methodology

(Pharm. 601, Faculty of Pharmacy, University of

Alberta)

1994 Volunteer, HUB community association, University of

Alberta

1994 Leadership Program (Speechcraft)

1993-1996 Vice President Finance (Treasurer), Pharmacy

Graduate Student Association Faculty of Pharmacy,

University of Alberta

1989-1991 Teaching in Organic Chemistry, General Chemistry,

Instrumental Analysis and Medicinal Chemistry (Faculty

of Pharmacy, University of Tehran)

Organizing Conference and Symposiums

Program	Date	Title of symposium
Fifth Annual AAPS-Northeast	April 22nd,	Impact of Genome Therapeutics
Regional Discussion Group	2002	on Drug Delivery
planning committee		
Sixth Annual AAPS-Northeast	April 28,	Overcoming Physical, Chemical,
Regional Discussion Group	2003	and Biological Barriers to Drug
planning committee		Delivery
Seventh Annual AAPS-Northeast	April 26,	Cellular Targeting and Drug
Regional Discussion Group	2004	Delivery
Planning Committee		
Organizing INBRE Seminar	April 2004- April 2007	Related Topics to Toxicology, Biology, and Drug Discovery
Eighth Annual AAPS-Northeast	April 22,	Successful Prediction of In Vivo
Regional Discussion Group	2005	Performance of Oral Dosage
Planning Committee		Forms from In Vitro/ In Silico
Ninth Annual AAPS-Northeast	April 28,	Studies Drugs from Discovery to the
Regional Discussion Group	2006	Clinic
Planning Committee	2000	Giii ii G
Tenth Annual AAPS-Northeast	April 20,	Methods and Applications for the
Regional Discussion Group	2007	Optimization of Pre-clinical Oral
Planning Committee		Dosage Formulations
12 th International Conference on	February	Scientific Advisory Board
The Interface of Chemistry-	22-24,	Colonano Advisory Board
Biology in Biomedical Research	2008	
(ISCBC)		
Drug Therapy in the 21st	September	A member of organizing
Century: Discovery and Clinical	14, 2012	committee
Use		
1st International Conference on	September	Conference Chair
Frontiers in Pharmaceutical	28-30,	
Sciences: Global Perspectives	2012	

MAJOR PROFESSOR FOR UNDERGRADUATE STUDENTS, GRADUATE STUDENTS, AND POSTDOCTORAL FELLOWS

Student	Degree	Thesis or
		Non-Thesis Option
Ali Saleh	B.Sc.	Summer Student
Amie Tiemo	B.Sc.	Summer Student
Meredith Selecky	B.Sc.	Summer Student
Amelia Lyman	B.Sc.	Summer Student
Carrie Funk	B.Sc.	Summer Student
Megrose Quiterio	B.Sc.	Academic year
Michael Zompa	B.Sc.	Summer Student
Elana Viola	B.Sc.	Academic year
Cang Nguyen	B.Sc.	Summer Student
Sojin Kim	B.Sc.	Summer Student
Phouthone Malayphone	B.Sc.	Academic year
Thuytien Dang	B.Sc.	Academic year
Daravuth Hav	B.Sc.	Academic year
Julia Robidoux	B.Sc.	Summer Student
Victoria Lomas	B.Sc.	Summer Student
Jared Bolton	B.Sc.	Academic year
Brian Sullivan	B.Sc.	Academic year
Kellen McCafferly	B.Sc.	Academic year
Kevin Northup	B.Sc.	Academic year
Kellye Loethen	Pharm.D.	Summer Student
Michael Hanley	Pharm.D.	Summer Student and
		Academic year
Millie White	Pharm.D.	Summer Student
Hilary Williams	Pharm.D.	Summer Student
Jessica Lehmann	Pharm.D.	Summer Student and
		Academic year
Kate Waggner	Pharm.D.	Summer Student
Benjamin Carrol	Pharm.D.	Academic year
Nicol Patel	Pharm.D.	Academic year
Derek Joseph	Pharm.D.	Academic year
Neal Russell	Pharm.D.	Academic year
Vasudha Gupta	Pharm.D.	Academic year
Nicole St. Jean	Pharm.D.	Academic year
Duojia Li	Pharm.D.	Academic year
Victor Carlu	Pharm.D.	Academic year

	Pharm.D.	Academic year
	Pharm.D.	Academic year
a Young Kim F	Pharm.D.	Academic year
_	Pharm.D.	Academic year
anhe Cui F	reshman	Academic year
a Young Kim F	reshman	Academic year
itaran Bhavaraju F	Ph.D.	Summer Student
guyen Hai Nam F	Ph.D.	Postdoctoral Fellow
oroush Sardari F	Ph.D.	Postdoctoral Fellow
nil Kumar F	Ph.D.	Postdoctoral Fellow
ousef Ahmadibeni F	Ph.D.	Postdoctoral Fellow
ianfeng Gu F	Ph.D.	Postdoctoral Fellow
akesh Kumar Tiwari F	Ph.D.	Postdoctoral Fellow
itaran Bhavaraju F	Ph.D.	Postdoctoral Fellow
eendayal Mandal F	Ph.D.	Postdoctoral Fellow
jay Dixit F	Ph.D.	Postdoctoral Fellow
hupender Singh F	Ph.D.	Postdoctoral Fellow
habban Asd F	Ph.D.	Visitor Scientist
unil Kumar F	Ph.D.	Postdoctoral Fellow
ebecca Pitts N	VI.S.	Thesis
uofeng Ye N	M.S., Ph.D.	Thesis
itesh Agarwal F	Ph.D.	Thesis
onghoo Oh F	Ph.D.	Thesis
nju Gupta F	Ph.D.	Thesis
hanu Priya N	M.S.	Thesis
mir Nasrolahi Shirazi F	Ph.D.	Thesis
asser Sayeh F	Ph.D.	Thesis

SELECTED ALUMNI MENTORED IN MY LABORATORY

- 1. **Dr. Yousef Ahmadibeni**, Assistant Professor, Department of Chemistry, Tennessee State University, USA.
- 2. **Dr. Rakesh Tiwari**, Assistant Professor, Department of Life Sciences, Shiv Nadar University, India.
- 3. **Dr. Deendayal Mandal**, Associate Professor, School of Biotechnology, KIIT University, Bhubaneswar, Orissa, India.
- 4. **Dr. Nguyen Hai Nam**, Associate Professor, Hanoi University of Pharmacy, Vietnam.
- 5. **Dr. Anil Kumar**, Associate Professor, Birla Institute of Technology and Science, India.
- 6. **Dr. Xianfeng Gu**, Assistant Professor, Fudan University, China.
- 7. **Dr. Ajay Kumar Dixit**, R&D Associate Scientist, ITC R&D Centre, Peenya Industrial Are, Bangalore, India.
- 8. **Guofeng Ye**, Scientist II, Agilux Laboratories, Previously in Novartis Institutes for Biomedical Research, NIBR, Cambridge, MA
- 9. Hitesh K. Agarwal, Senior Postdoctoral Researcher, Mount Sinai Medical Center.
- 10. Aimee Gagnon, Research Associate, Ipsen
- 11. Rebecca Pitts, Scientist II, Novartis Institutes for Biomedical Research
- 12. Sitaran Bhavaraju, US Pharmacopea.
- 13. Carrie Funk, Pfizer.
- 14. **Jessica Lehmann**, Resident, Baylor College of Medicine, Dallas
- 15. **Michael Hanley,** Ph.D., Clinical Pharmacology Contractor, Millennium Pharmaceuticals Inc.
- 16. **Benjamin Carroll**, Pharm.D., M.S., Postdoctoral Fellow at Johnson and Johnson.
- 17. Kellye Loethen, Pharm. D., Pharmacy Resident, Naval Medical Center San Diego.
- 18. Victoria Lomas, B.S., AmeriCorps NCCC FEMA Corps.