

## CURRICULUM VITAE

**Dhiren R. Thakker**

**RESIDENCE:** 317 Dalton Drive  
Raleigh, NC 27615  
(919) 606 6688 (Cell)  
(919) 966-1122 (Office)  
dhiren\_thakker@unc.edu

**EDUCATION:** Ph.D. (Biochemistry) University of Kansas, Lawrence, KS, 1975  
M.S. (Pharmaceutical Chem.) Columbia University, NY, 1972  
B.S. (Pharmacy) Bombay University, Bombay, India, 1970

### PROFESSIONAL EXPERIENCE:

**2018 – Current**

John A. and Margaret P. McNeill, Sr. Distinguished Professor

**2017 – current**

Interim Dean, UNC Eshelman School of Pharmacy  
Interim Director, Eshelman Institute for Innovation

**1996 - current**

Howard Q Ferguson Distinguished Professor of Pharmaceutical Sciences (1996-2018)  
Division of Pharmacotherapy and Experimental Therapeutics (since 2009)  
Division of Molecular Pharmaceutics (formerly Division of Drug Delivery and Disposition)  
(1996-2009)  
UNC Eshelman School of Pharmacy (formerly School of Pharmacy)  
The University of North Carolina at Chapel Hill  
Chapel Hill, NC 27599

**2008 – 2016**

Associate Dean, Entrepreneurial Development and Global Engagement (EDGE)  
(Previously referred as Economic Development and International Partnerships)  
UNC Eshelman School of Pharmacy

**2015 – current**

Director, Bill and Karen Campbell Faculty Mentoring Program  
UNC Eshelman School of Pharmacy

**1998 - 2008**

Associate Dean, Research and Graduate Education  
School of Pharmacy (now UNC Eshelman School of Pharmacy)

**1996 - 2005**

*Joint appointment:* Pharmacology  
University of North Carolina at Chapel Hill  
Chapel Hill, NC 27599

**1995 - 1996**

Visiting Professor  
Pharmaceutics Division, School of Pharmacy  
University of North Carolina at Chapel Hill,  
Chapel Hill, NC 27599

**1992 - 1995**

Director, Drug Metabolism Department, Glaxo Research Institute,  
Research Triangle Park, NC 27709

**1990 - 1992**

Department Head, Drug Metabolism Department, Glaxo Research Institute,  
Research Triangle Park, NC 27709

**1987 - 1990**

Section Head, Bioorganic Mechanisms Section, Drug Metabolism Department,  
Glaxo Inc., Research Triangle Park, NC 27709

**1993 - 1996**

Adjunct Professor, Department of Pharmacology, School of Medicine,  
University of North Carolina, Chapel Hill, NC 27514

**1990 - 1996**

Adjunct Professor, Department of Medicinal Chemistry,  
School of Pharmacy, University of North Carolina,  
Chapel Hill, NC 27514

**1984 - 1987**

Senior Investigator, Laboratory of Molecular Pharmacology, Division of  
Biochemistry and Biophysics, Center for Drugs and Biologics, Food and Drug  
Administration, Bethesda, MD 20892.

**1978 - 1984**

Senior Staff Fellow/Research Chemist, Laboratory of Bioorganic Chemistry, National  
Institute of Arthritis, Diabetes, and Digestive and Kidney Diseases, National Institutes  
of Health.

**1976 - 1978**

Staff Fellow, Laboratory of Chemistry, National Institute of Arthritis, Metabolism, and Digestive Diseases, National Institutes of Health.

**1975 - 1976**

Visiting Fellow, Laboratory of Chemistry, National Institute of Arthritis, Metabolism and Digestive Diseases, National Institutes of Health.

**MANAGEMENT**

**EXPERIENCE:**

**1. Serves (or has served) on Boards and Scientific Advisory Boards of several Biotech Companies**

**2. Led and Managed the Preclinical Drug Metabolism Department at**

**Glaxo**

- The department consisted of a staff of 36 including 19 Senior Scientists with Ph.D. or equivalent experience.
- The annual operating budget was \$6 million, and the capital budget was approximately \$1.5 million.
- The primary responsibility of the department was to conduct non-clinical metabolism, pharmacokinetic and toxicokinetic studies in support of development candidates, and to participate in the discovery and selection of new drug candidates to achieve optimum metabolic and pharmacokinetic profiles.
- The department also had the responsibility to synthesize radioisotope-labeled compounds for the Research and Development Divisions.

**3. As a member of the Technical Development Team (management team), participated in the coordination of exploratory development programs**

- The Technical Development Team had representation from Toxicology, Drug Metabolism, Pharmaceuticals, Analytical Chemistry, Process Chemistry, Clinical Pharmacology, Regulatory Affairs, and Project Planning.
- The Technical Development Team participated in the formulation of strategies for exploratory development (preclinical through phase I/phase II) of drug candidates and was responsible for the implementation of these strategies.

**4. Interacted with the FDA on the non-clinical ADME issues**

**5. As a member of the Research Management Committees, participated in the management of Discovery Programs in Cancer, Metabolic Diseases, and Inflammation**

**PROFESSIONAL** Editorial Board, J. Pharm. Sci. (1997-current)

**EXPERIENCE** Editorial Board, Medicinal Research Reviews (2000 – current).

UNC Global Advisory Council, UNC Global (2016- )

Chancellor's UNC Global Taskforce (2016- )

Advisory Board, Office of Technology Development, UNC-Chapel Hill

Co-Editor-in-Chief, Medicinal Research Reviews (2003-2006)

Advisory Board, Office of Business and Economic Development, UNC-Chapel Hill

Advisory Board, Carolina Student Biotechnology Network, UNC, Chapel Hill

International Affairs Advisory Council, UNC Global Education Center, UNC, Chapel Hill

IntraHealth Advisory Council, IntraHealth International, Durham, NC

National University of Singapore Department of Pharmacy Visiting Committee (external review committee) 2009

Chair, National University of Singapore Department of Pharmacy Visiting Committee (external review committee) 2014

Editorial Board, Current Drug Metabolism, 1999-2002.

Editorial Board, Drug Metabolism and Disposition, 1994-1997

Board of Directors and Scientific Advisory Board, Qualyst Inc.

Board of Directors, Chesson Labs (current)

Board of Directors, Tergus Pharma (current)

Co-founder and Chair of Scientific Advisory Board, Sphaera Pharma (current)

Drug Development Advisory Board, Scios Inc. (subsidiary of J & J)

Science and Technology Advisory Board, Oread Inc.

Scientific Advisory Board, Navicyte

Awards Committee, Society for Biomolecular Screening (SBS)

AAPS Taskforce on Drug Discovery Interface

Chair, Drug Metabolism Focus Group, AAPS

Founding Member of the Steering Committee, RTP Drug Metabolism Discussion Group, 1999-2002

Member of the External Review Committee  
Graduate Program, School of Pharmacy, University of Washington, Seattle

Basic Pharmacology Advisory Committee  
Pharmaceutical Manufacturers' Association Foundation

Special Emphasis Panel, National Institute of General Medical Sciences, NIH

Organizer - Short Course on "Designing Safe Drugs - Integration of Disposition Studies in Drug Discovery and Development", Residential School on Medicinal Chemistry, Drew University

Organizer and Chair- Short Course on "Prodrug Design – Enhanced and Targeted Delivery of Therapeutic Agents" at the 7<sup>th</sup> North American ISSX Meeting, October 1996, Sand Diego, CA.

Chair - Special Symposium on "*Drug Delivery and Prodrug Technologies*" at the 31st ACS Western Regional Meeting held in October 1995

Chair - Session on "*Integration of Preclinical ADME Studies in the Preclinical and Clinical Safety Assessment*" at the annual meeting of the Drug Information Association, 1994

Chair - Session on "*Delivery and Disposition of Peptides and Oligonucleotides- Current Status and Future Challenges*" at the annual meeting of the International Society for the Study of Xenobiotics held in October 1994

Special reviewer for Geriatric Review Committee, Institute on Aging

Reviewer for Investigational New Drug (IND) applications primarily dealing with Interferons and Monoclonal antibodies

Reviewer for several leading journals including Nature, Cancer Research, Journal of American Chemical Society, Molecular Pharmacology, Carcinogenesis, Chemico-Biological Interactions, Analytical Biochemistry, Toxicology and Applied Pharmacology, and Chemical Research in Toxicology, Drug Metabolism and Disposition, Pharmaceutical Research, Journal of Pharmaceutical Sciences, Journal of Experimental Therapeutics.

Served as a consultant to Amgen, Amylin Pharmaceuticals, BASF Bioscience, Du Pont Pharmaceuticals, Glaxo Wellcome, ICAGEN, Intercardia, Parke Davis, Procter & Gamble, Synaptics, Triangle Pharmaceuticals, Trimeris, Wyeth, Pozen, Ontogen, Chiron, Scios, Medivation, Arete, Virobay, Sanofi-Aventis, USVP venture partners

Served as an Expert Witness for several law firms representing pharmaceutical companies in patent litigation

External Examiner, Ph.D. Examination, Uppsala University, Sweden.

External Examiner, Ph.D. Examination for Huadong Sun, University of Toronto,

Toronto, Ontario, Canada

External Examine, Ph.D. Examination, Nirma University, Gujarat, India

Organizer, Symposium on ADME in Drug Discovery at the Institute of Chemical Technology, Mumbai, India, 2012

**HONORS AND  
AWARDS:**

INVENTOR OF THE YEAR AWARD – The University of North Carolina at Chapel Hill, Chapel Hill, NC, USA, 2015

DISTINGUISHED ALUMNUS AWARD - Academic, UDCT Alumni Association and Chemical Institute of Technology, Mumbai, India, 2014

FELLOW, American Association of Pharmaceutical Scientists

SATO MEMORIAL INTERNATIONAL AWARD presented by the Pharmaceutical Society of Japan in 1987.

PHILLIP NEWMARK AWARD for "Excellence in Biochemical Research" at the University of Kansas in 1974.

DISSERTATION FELLOWSHIP at the University of Kansas in 1975.

GRADUATE HONORS FELLOW at the University of Kansas.

Placed SECOND in Bombay University at the B. Pharm. degree examination.

**RESEARCH:**

- (1) Mechanisms of drug transport across intestinal epithelium
- (2) Novel intestinal absorption mechanism and pharmacology of the antidiabetic drug metformin
- (3) Anticancer effects of metformin
- (4) Regulation and modulation of tight junctions in biological barriers
- (5) Pediatric drug disposition
- (6) Stereoselectivity of cytochrome P450 isoenzymes (past interest)
- (7) Metabolic activation of xenobiotics to mutagens and carcinogens (past interest)

**Ph.D.**

- (1) Development of specific irreversible inhibitors of the enzyme, COMT, which

play an important role in the metabolism of epinephrine, norepinephrine, dopamine, as well as several catecholic endogenous compounds and xenobiotics.

- (2) Study of relationship between physico-chemical properties of 8-hydroxyquinolines and their COMT inhibitory activity (Hansch approach).
- (3) Affinity chromatography of COMT.
- (4) Chemical modification of functional groups on COMT active-site.

### M.S.

Synthesis of imidazole derivatives as potential chymotryptic agents.

**FUNDING:** Over \$4 million in research funding from federal and state agencies as well as from pharmaceutical and biotech companies

### **Current Funding (2016):**

#### **RSG CCE 128826**

01/01/2016 – 12/31/2019

**American Cancer Society** (Contact PI: Bae-Jump, V.) Annual Direct: \$164,963

Role: Co-Investigator

#### **Obesity, Cation-Selective Transporters and Metformin in Endometrial Cancer**

We will evaluate the contribution of a representative metformin transporter to the anti-tumor efficacy of metformin in obese and non-obese orthotopic mouse models, in which tumors are derived from an endometrial cancer cell line with normal or overexpression of a representative transporter. In addition, we will correlate treatment response to metformin in EC patients with (i) expression and genetic variants of the metformin transporters, (ii) modulation of the AMPK-mTOR pathway, and (iii) metabolic factors associated with obesity.

#### **PhRMA Foundation** (PI – Dhiren Thakker; Predoctoral Fellowship for Christine Lee)

Annual Direct: \$20,000, Role: Advisor to Christine Lee

01/01/2017 – 06/30/2018

#### **Use of Human Intestinal Tissue and PBPK Modeling to Predict the Pharmacokinetics of Orally Administered Amoxicillin in Adults and Infants**

### **STUDENTS AND FELLOWS:**

#### **Graduate Students:**

1. Kiho Lee (MCNP) - Ph.D., May 2000
2. Hui Ouyang (MCNP) – Ph.D., Dec 2001
3. Peter Ward (Pharmacology) – Ph.D., May 2001
4. Matt Troutman (DDD) – Ph.D., July 2001
5. Gururaj Rao (DDD) – Ph.D., May 2003
6. Chee Ng (DDD) - (Chair, Dissertation Committee and co-advisor), Ph.D., May 2003
7. Seongwong Hong (MCNP) – Ph.D., May 2004
8. David Bourdet (DDD)- Ph.D., August, 2005
9. Stephanie Faucette (Molecular Pharmaceutics) co-advisor – Ph.D., May 2006
10. Richard Graham (Molecular Pharmaceutics) co-advisor – Ph.D., May 2006

11. Tim Tippin (Molecular Pharmaceutics), Ph.D., Dec 2006
12. Beverly Mowrey (Molecular Pharmaceutics), Ph.D. Aug 2007
13. Ryan Klein (Molecular Pharmaceutics), Ph.D. Dec 2007
14. Xin Ming (Molecular Pharmaceutics), Ph.D. Aug 2008
15. Will Proctor (Molecular Pharmaceutics) Ph.D. May 2010
16. Claudia Generaux (Molecular Pharmaceutics) co-advisor Ph.D. August, 2010
17. Matthew Dufek (Molecular Pharmaceutics) Ph.D. May 2011
18. Souzan Yanni (Katholic U. Leuven, Belgium) Ph.D., June 2010
19. Chester Costales (Molecular Pharmaceutics) Ph.D., May 2013
20. Kevin Han (Molecular Pharmaceutics) Ph.D., May 2013
21. Yunhui Zhang (Ji Lin University, Peoples' Republic of China), Ph.D., 2013
22. Nicole Zane (Pharmacotherapy & Exp. Therapeutics), Ph.D., May 2015
23. Hao Cai (Pharmacotherapy & Exp Therapeutics), Ph.D., May 2016
24. Christine Lee (Pharmacotherapy & Exp Therapeutics), 4th year
25. Lawrence Ku (Pharmacotherapy & Exp Therapeutics), 4th year

**Postdoctoral Fellows:**

1. Dr. Aruljothi Muralidharan (2017- )
2. Dr. Arti Thakkar (2013-2015)
3. Dr. Ravindra Varma Alluri (2011 - 2013)
4. Dr. Wujian Ju (2009 – 2010)
5. Dr. Yan Gao (2006- 2008)
6. Darkhan Utepbergenov (2005-2007)
7. Dr. Lorraine King (2001-1002)
8. Dr. Johne Ansede (2001-2003)
9. Dr. Dongzhou Liu (1996-1998)
10. Dr. Shailesh Desai (1996-1999)
11. Dr. Cuiping Chen (1997-1998)
12. Dr. Sonia Serron (1997-1998)
13. Dr. Pieter Annaert (1998-1999)
14. Dr. Ashwin Patel (1998-2000)

**Research Assistant Professor:**

1. Dr. Ruth Everett (2010 - )
2. Dr. Steven Qian (2004-2005)

**Visiting Scholars:**

1. Dr. Yufeng Xia (2013-2014)
2. Dr. Ruth Everett (2008-2009)
3. Souzan Yanni (2008-2010)
4. Dr. Harish Padh, Vice Chancellor, Sardar Patel University (June – September 2006)
5. Dr. Innocent Ononiwu (2012-2014)

**DISSERTATION  
COMMITTEES:**

1. Cuiping Chen, DDD (Ph.D. 1997)
2. Catherine Booth, DDD (Ph.D. 1997)
3. Xinrong Liu, DDD (Ph.D. 1998)
4. Mark Bush, DDD (Ph.D. 1999)
5. Kay Rittenhaus, DDD (Ph.D. 1999)



6. James Whisnant, DDD (M.S. 1999)
7. Mark Summerville, DDD (Ph.D. 2000)
8. Jennifer(Quin) Dong, DDD (Ph.D. 2000)
9. Claude Degenais, DDD (Ph.D. 2001)
10. Preethi Krishnan, MCNP (Ph.D. 2001)
11. Christopher Lowden, MCNP (Ph.D. 2001)
12. Jessica Smith, DDD (Ph.D., 2001)
13. Jian Zong, DDD (Ph.D., 2002)
14. Ryan Turncliff, DDD (Ph.D. 2004)
15. Chris Matheny, DDD (Ph.D. 2003)
16. Lian Zhou, MCNP (Ph.D., 2002)
17. Carolyne Cyne (Pharmacology - Ph.D. committee)
18. Peter Sazani, Pharmacology (Ph.D., 2002)
19. Craig Lee, Exp. Therapeutics (Chair) (Ph.D., 2006)
20. Scott Barrow, Toxicology Curriculum (Ph.D., 2004)
21. Enzo Palma, Molecular Pharmaceutics (Ph.D. 2006)
22. Dongmei Liu, Molecular Pharmaceutics (Chair) (Ph.D. 2007)
23. Jian Jiang, Curriculum in Applied and Material Sciences (Ph.D. 2007)
24. Xihong Xu, Chemistry, Duke University (Ph.D. 2008)
25. Brandon Swift, Experimental Therapeutics (Ph.D. 2009)
26. Melanie Joy, MCNP (Ph.D. 2009)
27. David Szabo, Toxicology Curriculum (Ph.D. 2011)
28. Michael Cohen-Wolkowicz Experimental Therapeutics (Ph.D. 2012)
29. Christina Won Experimental Therapeutics (Ph.D. 2012)
30. Nathan Pfeifer, Experimental Therapeutics (Ph.D. 2013)
31. Katsuhiko Sueda, Molecular Pharmaceutics (Ph.D. 2014)
32. Brandon Gufford, Experimental Therapeutics (SAC Committee)
33. Kyunghee Yang, Experimental Therapeutics (Ph.D. 2014)
34. Brian Ferslew, Experimental Therapeutics (Ph.D. 2014)
35. James Huckle, Molecular Pharmaceutics (Ph.D. 2014)
36. Jing Fu, Molecular Pharmaceutics (Ph.D. 2016)
37. Kevin Watt, Experimental Therapeutics (Ph.D. 2016)
38. Jason Slizgi, Experimental Therapeutics, (Ph.D. Committee)

**PUBLICATIONS:** Co-author of over 160 publications which include papers in the peer reviewed journals, review articles and book chapters (List Attached), co-editors of 2 books, and co-inventor on 7 patents (issued and pending).

- AFFILIATIONS:**
- (1) American Chemical Society (ACS)
  - (2) American Association for Advancement of Science (AAAS)
  - (3) American Society of Pharmacology and Experimental Therapeutics (ASPET)
  - (4) American Association for Cancer Research (AACR)
  - (5) American Association of Pharmaceutical Scientists (AAPS)
  - (6) International Society for the Study of Xenobiotics (ISSX)
  - (7) Indian Association for Cancer Research

### PUBLICATIONS

1. Borchardt, R.T. and Thakker, D.: Affinity Labeling of Catechol-O-Methyltransferase with N-Iodoacetyl-3,5-dimethoxy-4-hydroxyphenylethylamine. Biochem. Biophys. Res. Commun., 54: 1233-1239, 1973.
2. Borchardt, R.T. and Thakker, D.R.: Catechol-O-methyltransferase. 6. Affinity Labeling with N-Iodoacetyl-3,5-dimethoxy-4-hydroxyphenylalkylamines. J. Med. Chem., 18: 152-158, 1975.
3. Borchardt, R.T. and Thakker, D.R.: Affinity Labeling of Catechol-O-methyltransferase by N-Halonacetyl Derivatives of 3,5-Dimethoxy-4-hydroxyphenylethylamine and 3,4-Dimethoxy-5-hydroxyphenylethylamine. Kinetics of Inactivation. Biochemistry, 14: 4543-4551, 1975.
4. Borchardt, R.T., Cheng, C.G. and Thakker, D.R.: Purification of Catechol-O-methyltransferase. 8. Structure-Activity Relationships for Inhibition by 8-Hydroxyquinolines. J. Med. Chem., 63: 69-77, 1975.
5. Borchardt, R.T., Thakker, D.R., Warner, V.D., Mirth, D.B. and Sane, J.M.: Catechol-O-methyltransferase. 8. Structure-Activity Relationships for Inhibition by 8-Hydroxyquinolines. J. Med. Chem., 19: 558-560, 1976.

6. Brochardt, R.T. Reid, J. R., Thakker, D.R., Liang, Y.O., Wightman, R.W. and Adams, R.N.: Catechol-O-methyltransferase. 9. Mechanisms of Inactivation by 6-Hydroxydopamine. J. Med. Chem., 19: 1201-1209, 1976.
7. Borchardt, R.T. and Thakker, D.R.: Evidence for Sulfhydryl Groups at the Active Site of Catechol-O-methyltransferase. Biochim. Biophys. Acta, 445: 598-609, 1976.
8. Borchardt, R.T. and Thakker, D.R.: Affinity Labeling of Catechol-O-Methyltransferase using N-Haloacetyl Derivatives of 3,5-Dimethoxy-4-hydroxyphenylethylamine and 3,4-Dimethoxy-5-hydroxyphenylethylamine. In Jacoby, W.B. and Wilchek, M. (Ed.): Methods in Enzymology - Affinity Labeling. New York, Academic Press, 1977, Vol. 46, pp. 554-561.
9. Lu, A.Y.H., Levin, W., Vore, M. Conney, A.H., Thakker, D.R., Holder, G. and Jerina, D.M.: Metabolism of Benzo[a]pyrene by Purified Liver Microsomal Cytochrome P-448 and Epoxide Hydrase. In Freudenthal, R.I. and Jones, P.W. (Ed.): Polynuclear Aromatic Hydrocarbons: Chemistry, Metabolism, and Carcinogenesis. New York, Raven Press, 1976, Vol. 1, pp. 115-126.
10. Thakker, D.R., Yagi, H., Lu, A.Y.H., Levin, W., Conney, A.H. and Jerina, D.M.: Metabolism of Benzo[a]pyrene IV. Conversion of (+)-Trans-7,8-dihydroxy-7,8-dihydrobenzo[a]pyrene to the Highly Mutagenic 7,8-Diol-9,10-epoxides, Proc. Natl. Acad. Sci. USA, 73: 3381-3385, 1976.
11. Thakker, D.R., Yagi, H., Akagi, H., Koreeda, M., Lu, A.Y.H., Levin, W., Wood, A. W., Conney, A.M.: Metabolism of Benzo[a]pyrene VI. Stereoselective Metabolism of Benzo[a]pyrene and Benzo[a]pyrene 7,8-Dihydrodiol to Diol Epoxides. Chem.-Biol. Interact., 16: 281-300, 1977.
12. Thakker, D.R., Yagi, H., Levin, W., Lu, A.Y.H., Conney, A.H. and Jerina, D.M.: Stereospecificity of Microsomal and Purified Epoxide Hydrase from Rat Liver: Hydration of Arene Oxides of Polycyclic Hydrocarbons. J. Biol. Chem., 252: 6328-6334, 1977.
13. Yagi, H., Thakker, D.R. Hernandex, O., Koreeda, M. and Jerina, D.M.: Synthesis and Reactions of the Highly Mutagenic 7,8-Diol-9,10-epoxides of the Carcinogen Benzo[a]pyrene. J. Am. Chem. Soc., 99: 1604-1611, 1977.
14. Yagi, H., Thakker, D.R., Mah, H.D., Koreeda, M. and Jerina, D.M.: Absolute Stereochemistry of the Highly Mutagenic 7,8-Diol-9,10-epoxides Derived from the Potent Carcinogen trans-7,8-Dihydroxy-7,8-dihydrobenzo[a]pyrene. J. Am. Chem. Soc., 99: 2358-2359, 1977.
15. Whalen, D.L. Montemarano, J.A., Thakker, D.R., Yagi, H. and Jerina, D.M.: Changes of Mechanism and Product Distributions in the Hydrolysis of benzo[a]pyrene 7,8-Diol-9,10-epoxide metabolites Induced by Changes in pH. J. Am. Chem. Soc., 99: 5522-5524, 1977.
16. Wood, A.W., Chang, R.L., Levin, W., Yagi, H. Thakker, D.R., Jerina, D.M., and Conney, A.H.: Differences in Mutagenicity of the Optical Enantiomers of the Diastereomeric Benzo[a]pyrene 7,8-Diol-9,10-epoxides. Biochem. Biophys. Res. Commun., 77: 1389-1396, 1977.
17. Wood, A.W., Levin, W., Ryan, D., Thomas, P.E., Yagi, H., Mah, H.D., Thakker, D.R., Jerina, D.M., and Conney, A.H.: High Mutagenicity of Metabolically Activated Chrysene 1,2-Dihydrodiol: Evidence for Bay Region Activation of chrysene. Biochem. Biophys. Res. Commun., 78: 847-854. 1977.
18. Levin, W., Wood, A.W., Lu, A.Y.H., Ryan, D., West, S., Thakker, D.R., Yagi, H., Jerina, D.M., and Conney, A.H.: Role of Purified Cytochrome P-448 and Epoxide Hydrase in the Activation and Detoxification of Benzo[a]pyrene. In Jerina, D.M. (Ed.): Drug Metabolism Concepts. Washington, D.C., American Chemical Society, 1977, ACS Symposium Series 44, pp. 99-126.
19. Jerina, D.M. Lehr, R., Shaefer-Ridder, M., Yagi, H., Karle, J.M., Thakker, D.R., Wood, A.W., Lu, A.Y.H., Ryan, D., West, S., Levin, W., and Conney, A.H.: Bay Region Epoxides of Dihydrodiols: A Concept which Explains the

Mutagenic and Carcinogenic Activity of Benzo[a]pyrene and Benzo[a]anthracene. In Hiatt, H., Watson, J.D. and Winsten, I. (Ed.): Origins of Human Cancer. New York, Cold Spring Harbor Laboratories, 1977, PP. 639-658

20. Levin, W., Lu, A.Y.H., Ryan, D., Wood, A.W., Kapitulnik, J., West, S., Huang, M.T., Thakker, D.R., Holder, G., Yagi, H., Jerina, D.M. and Conney, A.H.: Properties of the Liver Microsomal Monooxygenase System and Epoxide Hydrase: Factors Influencing the Metabolism and Mutagenicity of Benzo[a]pyrene. In Hiatt, H., Watson, J.D., and Winsten, I. (Ed.): Origins of Human Cancer. New York, Cold Spring Harbor, Cold Spring Harbor Laboratories, 1977, pp.659-682.
21. Bresnick, E., Stoming, T.A., Vaught, J.B., Thakker, D.R., and Jerina, D.M.: Nuclear Metabolism of Benzo[a]pyrene and of (+)-trans-7,8-Dihydroxy-7,8-dihydrobenzo[a]pyrene. Comparative Chromatographic Analysis of Alkylated DNA. Arch. Biochem. Biophys., **183**: 31-37, 1978.
22. Lehr, R.E., Yagi, H., Thakker, D.R., Levin, W., Wood, A.W., Conney, A.H. and Jerina, D.M.: The Bay Region Theory of Polycyclic Aromatic Hydrocarbon Induced Carcinogenicity. In Freudenthal, R.I. and Jones, P.W. (Ed.): Polynuclear Aromatic Hydrocarbons: Chemistry, Metabolism and Carcinogenesis. New York, Raven Press, Vol. 2., pp.231-241.
23. Thakker, D.R., Levin, W., Stoming, T.A., Conney, A.H., and Jerina, D.M.: Metabolism of 3-Methylcholanthrene by Rat Liver Microsomes and a Highly Purified Monooxygenase System With and Without Epoxide Hydrase. In Freudenthal, R.I., and Jones, P.W.(Ed.): Polynuclear Aromatic Hydrocarbons: Chemistry, Metabolism and Carcinogenesis. New York, Raven Press, 1978, Vol. 2., pp.253-264.
24. Thakker, D.R., Yagi, H., and Jerina, D.M.: Analysis of Polycyclic Aromatic Hydrocarbons and Their Metabolites by High Pressure Liquid Chromatography. In Packer, L. and Fleischer, S. (Ed.): Methods in Enzymology: Biomembranes. New York, Academic Press, 1978, Vol. 51 (part C), pp. 279-296.
25. Thakker, D.R., Levin, W., Wood, A.W., Conney, A.H., Stoming, T.A., and Jerina, D.M.: Metabolic Formation of 1,9,10-Trihydroxy-9,10-dihydro-3-methylcholanthrene: A Potential Proximate Carcinogen from 3-Methylcholanthrene. J. Am. Chem. Soc., **100**: 654-647, 1978.
26. Thakker, D.R., Yagi, H., Lehr, R.E., Levin, W., Lu, A.Y.H., Change, R.L., Wood, A.W., Conney, A.H., and Jerina, D.M.: Metabolism of trans-9,10-Dihydroxy-9,10-dihydrobenzo[a]pyrene Occurs Primarily by Arylhydroxylation Rather than Formation of a Diol Epoxide. Mol. Pharmacol., **14**: 502-513, 1978.
27. Levin, W., Thakker, D.R., Wood, A.W., Chang, R.L., Lehr, R.E., Jerina, D.M. and Conney, A.H.: Evidence that Benzo[a]anthracene 3,4-Diol-1,2-epoxide is an Ultimate Carcinogen on Mouse Skin. Cancer Res., **38**: 1705-1710, 1978.
28. Jerina, D.M., Yagi, H., Thakker, D.R., Karle, J.M., Mah, H.D., Boyd, D.R., Gadaginamath, G., Wood, A.W., Beuning, M., Chang, R.L. Levin, W., and Conney, A.H.: Stereoselective Metabolic Activation of Polycyclic Aromatic Hydrocarbons. In Cohen, Y. (Ed.): Advances in Pharmacology and Therapeutics, Vol. 9, Toxicology. New York, Pergamon Press, 1978, pp. 53-62.
29. Jerina, D.M. Yagi, H., Lehr, R.E., Thakker, D.R., Schaefer-Ridder, M., Karle, J.M., Levin, W., Wood, A.W., Change, R.L., and Conney, A.H.: The bay-Region Theory of Carcinogenesis by Polycyclic Aromatic Hydrocarbons. In Gelboin, H.V. and Ts'o P.O.P. (Ed.): Polycyclic Hydrocarbons and Cancer: Chemistry, Molecular Biology, and Environment. New York, Academic Press, 1978, pp. 173-188.
30. Jerina, D.M., Thakker, D.R., Yagi, H., Levin, W. Wood, A.W., and Conney, A.H.: Carcinogenicity of Benzo[a]pyrene Derivatives: The Bay-Region Theory. In Pure and Applied Chemistry. Vol. 50, Oxford, England, Pergamon Press, 1978, pp. 1033-1044.

31. Buening, M.K., Wislocki, P.G., Levin, W., Yagi, H., Thakker, D.R., Akagi, H., Koreeda, M., Jerina, D.M., Conney, A.H.: Tumorigenicity of the Optical Enantiomers of the Diastereomeric Benzo[a]pyrene 7,8-Diol-9,10-epoxides in Newborn Mice: Exceptional Acitivity of (+)-7,8-Dihydroxy, 9,10-epoxy-7,8,9,10-tetrahydrobenzo[a]pyrene. Proc. Natl. Acad. Sci. USA, 75: 5358-5361, 1978.
32. Wood, A.W., Chang, R.L., Levin, W., Thomas, P.E., Ryan, D., Stoming, T.A., Thakker, D.R., Jerina, D.M., and Conney, A.H.: Metabolic Activation of 3-Methylcholanthrene and Its Metabolites to Mutagenes Towards Bacteria and Mammalian Cells. Cancer Res., 33: 3398-3404, 1978.
33. Kapitulnick, J., Wislocki, P.G., Levin, W., Yagi, H., Thakker, D.R., Akagi, H., Koreeda, M., Jerina, D.M., and Conney, A.H.: Marked Differences in the Carcinogenic Activity of Optically Pure (+) and (-) Trans-7,8-Dihydroxy-7,8-Dihydrobenzo[a]pyrene in Newborn Mice. Cancer Res., 38: 2661-2665, 1978.
34. Jerina, D.M., Yagi, H., Thakker, D.R., Lehr, R.E., Karle, J.M., Shaefer-Ridder, M., Levin, W., Wood, A.W., and Conney, A.H.: The Bay-Region Theory of Polycyclic Aromatic Hydrocarbon-Induced Carcinogenicity. In Bolt, H. (Ed.): Primary Liver Tumors. Lancaster, England, MTP Press, Ltd., 1979, pp.253-260.
35. Thakker, D.R., Yagi, H., Karle, J.M., Lehr, R.E., Levin, W., Ryan, D., Thomas, P.E., Conney, A.H., and Jerina, D.M.: Metabolism of Benzo[a]anthracene by Rat Liver Microsomes and by Purified Cytochrome P-448 with and without Epoxide Hydrase. Mol. Pharmacol., 15: 174-189, 1979.
36. Wood, A.W., Levin, W., Chang, R.L., Yagi, H., Thakker, D.R., Lehr, R.E., Jerina, D.M., and Conney, A.H.: Bay region activation of carcinogenic polycyclic hydrocarbons. In Jones, P. and Leber, P. (Eds.): Polynuclear Biology, Carcinogenesis and Mutagenesis, Vol. 4. Ann Arbor, Michigan, Ann Arbor Science Publishers, Inc. 1979, pp. 531-551.
37. Lehr, R.E., Taylor, C.W., Kumar, S., Levin, W., Chang, R., Wood, A.W., Conney, A.H., Thakker, D.R., Yagi, H., Mah., H.D., and Jerina, D.M.: Differences in Metabolism Provide a Basis for the Low Mutagenicity and Carcinogenicity of Benzo[e]pyrene Compared to Benzo[a]pyrene. In Jones, P.W. and Leber, P. (Eds.): Polynuclear Aromatic Hydrocarbons: Third International Symposium on Chemistry and Mutagenesis Vol.4. Ann Arbor, Michigan, Ann Arbor Science Publishers, Ins., 1979, pp. 37-49.
38. Thakker, D.R., Nordqvist, M., Yagi, H., Levin, W., Ryan, D., Thomas, P., Conney, A.H., and Jerina, D.M.: Comparative Metabolism of a Series of Polycyclic Aromatic Hydrocarbons by Rat Liver Microsomes and Purified Cytochrome P-450. In Jones, P.W. and Leber, P.(Eds.): Polynuclear Biology, Carcinogenesis and Mutagenesis, Vol. 4. Ann Arbor, Michigan, Ann Arbor Science Publishers, Inc., 1979. pp. 455-472.
39. Nordqvist, M., Thakker, D.R., Levin, W., Yagi, H., Ryan, D.E., Thomas, P.E., Conney, A.H., and Jerina, D.M.: The Highly Tumorigenic 3,4-Dihydrodiol is a Principal Metabolite Formed from Dibenzo[a,h]anthracene by Liver Enzymes. Mol. Pharmacol., 16: 634-655, 1979.
40. Levin, W., Buening, M.K., Wood, A.W., Chang, R.L., Thakker, D.R., Jerina, D.M., and Conney, A.H.: Tumorigenic Activity of 3-Methylcholanthrene Metabolites on Mouse Skin and in Newborn Mice. Cancer Res., 39: 3549-3553, 1979.
41. Wislocki, P.G., Buening, M.K., Levin, W., Lehr, R.E., Thakker, D.R., Jerina, D.M., and Conney, A.H.: Tumorigenicity of the Diastereomeric Benzo[a]anthracene 3,4-diol-1,2-epoxides and the (+) and (-)-Enantiomers of Benzo[a]anthracene 3,4-dihydrodiol in Newborn Mice. J. Natl. Cancer Inst., 63: 201-204, 1979.
42. Wood, A.Q., Levin, W., Thakker, D.R., Yagi, H., Chang, R.L., Ryan, D.E., Thomas, P.E., Dansette, P.M., Whittaker, N., Turujman, S., Lehr, R.E., Kumar, S., Jerina, D.M., and Conney, A.H.: Biological Activity of Benzo[e]pyrene: An Assessment Based on Mutagenic Activities and Metabolic Profiles of the Polycyclic Hydrocarbon and its Derivatives. J. Biol. Chem., 254: 4408-4415, 1979.

43. Yagi, H. Thakker, D.R., Lehr, R.E., and Jerina, D.M.: Benzo-ring Diol Epoxides of Benzo[e]pyrene and Triphenylene. J. Org. Chem., 44: 3449-3442, 1979.
44. Whalen, D.L., Ross, A.M., Manlenmarano, J.A., Thakker, D.R., Yagi, H., and Jerina, D.M.: General Acid Catalysis in the Hydrolysis of Benzo[a]pyrene 7,8-diol-9,10-epoxides. J. Am. Chem. Soc., 101: 5086-5088, 1979.
45. Chang, R.L., Wood, Levin, W., Mah, H.D., Thakker, D.R., Jerina, D.M., and Conney, A.H.: Differences in Mutagenicity and Cytotoxicity of (+)- and (-)-Benzo[a]pyrene 4,5-oxide: A Synergistic Interaction of Enantiomers. Proc. Natl. Acad. Sci. USA., 76: 4280-4284, 1979.
46. Thakker, D.R., Levin, W., Yagi, Turujman, S., Kapadia, D., Conney, A.H., and Jerina, D.M.: Absolute Stereochemistry of the Trans Dihydrodiols Formed from Benzo[a]anthracene by Liver Microsomes. Chem.-Biol. Interact. 27: 145-161, 1979.
47. Nordqvist, M., Thakker, D.R. Yagi, H., Lehr, R.E., Wood, A.W., Levin, W., Conney, A.H., and Jerina, D.M.: Evidence in Support of the Bay Region Theory as a Basis for the Carcinogenic Activity of Polycyclic Aromatic Hydrocarbons. In Bhatnagar, R.S. (Ed.): Molecular Basis of Environmental Toxicity. Ann Arbor, Michigan, Ann Arbor Science Publishers, Inc., 1980, pp. 329-357.
48. Lehr, R.E., Kumar, S., Levin, W., Wood, A.W., Chang, R.L., Buening, M.K., Conney, A.H., Whalen, D.L., Thakker, D.R., Yagi, H., and Jerina, D.M.: Benzo[a]pyrene Dihydrodiols and Diol Epoxides: Chemistry, Mutagenicity and Tumorigenicity. In Bjorseth, A. and Dennis, A.J. (Eds.): Polynuclear Aromatic Hydrocarbons: Chemistry and Biological Effects. Columbus, Ohio, Battelle Press, 1980.
49. Jerina, D.M., Yagi, H., Thakker, D.R., Lehr, R.E., Wood, A.W., Levin, W., and Conney, A.H.: Bay-region Activation of Polycyclic Aromatic Hydrocarbons to Ultimate Mutagens and Carcinogens. In Coon, M.J., Conney, A.H., Estabrook, R.W., Gelboin, H.V., Gillette, J.R., and O'Brian, P. (Eds.): Microsomes, Drug Oxidations, and Chemical Carcinogenesis. New York, Academic Press, Inc., 1980, pp. 1041-1051.
50. Thakker, D.R., Levin, W., Yagi, H., Tada, M., Conney, A.H., and Jerina, D.M.: Comparative Metabolism of Dihydrodiols of Polycyclic Aromatic Hydrocarbons to Bay-region Diol Epoxides. In Bjorseth, A. and Dennis, A.J. (Eds.): Polynuclear Aromatic Hydrocarbons: Chemistry and Biological Effects. Columbus, Ohio, Battelle Press, 1980, pp. 267-286.
51. Thakker, D.R., Yagi, H., Levin, W., Wood, A.W., Conney, A.H., and Jerina, D.M.: Metabolic Activation of Polycyclic Aromatic Hydrocarbons at the Bay-region. In Breimer, D.D. (Ed.): Towards Better Safety of Drugs and Pharmaceutical Products. Amsterdam, Elsevier/North-Holland, Biomedical Press, 1980, pp. 207-226.
52. Thakker, D.R., Yagi, H., Whalen, D.L., Levin, W., Wood, A.W., Conney, A.H., and Jerina, D.M.: Metabolic Formation and Reactions of Bay-region Diol Epoxides: Ultimate Carcinogenic Metabolites of Polycyclic Aromatic Hydrocarbons. In McKinney, J.R. (Ed.): Environmental Health Chemistry: The Chemistry of Environmental Agents as Potential Human Hazards. Ann Arbor, Michigan, Ann Arbor Science Publishers, Inc., 1980, pp. 383-401.
53. Levin, Wayne, Buening, Mildred K., Wood, Alexander W., Chang, Richard L., Kedzierski, Bogdan, Thakker, Dhiren R. Boyd, Derek R. Gadaginamath, Guru S., Armstrong, Richard N. Yagi, Haruhiko, Karle, Jeane M., Sloga, Thomas J., Jerina, Donald M., and Conney, Allan H.: An Enantiomeric Interaction in the Metabolism and Tumorigenicity of (+)- and (-)-Benzo[a]pyrene 7,8-oxide. J. Biol. Chem., 255: 9067-9074, 1980.
54. Conney, A.H., Levin, W., Wood, A.W., Yagi, H., Thakker, D.R., Lehr, R.E., and Jerina, D.M.: Metabolism of Polycyclic Aromatic Hydrocarbons to Reactive Intermediates with High Biological Activity. In Coulston, R. and Shubik, P. (Eds.): Human Epidemiology and Animal Laboratory Correlations in Chemical Carcinogenesis. Norwood, NJ, Ablex Publishing Corp., 1980, pp. 153-183.

55. Jerina, D.M., Sayer, J.M., Thakker, D.R., Yagi, H., Levin, W., Wood, A.W., and Conney, A.H.: Carcinogenicity of Polycyclic Aromatic Hydrocarbons: The Bay-region Theory. B. Pullman, P.O.P. Ts'O and H. Gelboin (Eds.). Carcinogenesis: Fundamental Mechanisms and Environmental Effects. Dordrecht, Holland, D. Reidel Publishing Co., 1980, pp. 1-12.
56. Kedzierski, B., Thakker, D.R., Armstrong, R.N., and Jerina, D.M.: Absolute Configuration of the K-region 4,5-Dihydrodiols and 4,5-Oxide of Benzo[a]pyrene. Tetrahedron Letters, 22: 405-408, 1980.
57. Wood, A.W., Change, R.L. Huang, M.-T., Levin, W., Lehr, R.E., Kumar, S., Thakker, D.R., Yagi, H., Jerina, D.M., and Conney, A.H.: Mutagenicity of Benzo[a]pyrene and Triphenylene Tetrahydroepoxides and Diol Epoxides in Bacterial and Mammalian Cells. Cancer Res., 40: 1985-1989, 1980.
58. Thakker, D.R., Levin, W., Buening, M. Yagi, H., Lehr, R.E., Wood, A.W., Conney, A.H., and Jerina, D.M.: Species Specific Enhancement by 7,8-Benzoflavone of Hepatic Microsomal Metabolism of Benzo[a]pyrene 9,10-dihydrodiol to Bay-region Epoxides. Cancer Res., 41: 1938.
59. Nordqvist M., Thakker, D.R., Levin, W., Yagi, H., Conney, A.H., and Jerina, D.M.: Metabolism of Chrysene and Phenanthrene to Bay-region Diol Epoxides by Rat Liver Enzymes. Mol. Pharmacol., 19: 168-178, 1981.
60. Vyas, K.P., Thakker, D.R., Levin, W., Yagi, H., Conney, A.H., and Jerina, D.M.: Stereoselective Metabolism of the Optical Isomers of Trans-1,2-dihydroxy-1,2-dihydrophenanthrene to Bay-region Diol Epoxides by Rat Liver Microsomes. Chem.-Biol. Interact., 38: 203-213, 1982.
61. Yagi, H., Vyas, K.P., Tada, M., Thakker, D.R., and Jerina, D.M.: Synthesis of the Enantiomeric Bay-region Diol Epoxides of Benz[a]anthracene and Chrysene. J. Org. Chem., 47: 1110-1117, 1982.
62. Jerina, D.M., Sayer, J.M., Yagi, H., Croisy-Delcey, M., Ittah, Y., Thakker, D.R., Wood, A.W., Chang, R.L., Levin, W., and Conney, A.H.: Highly Tumorigenic Bay-region Diol Epoxides from the Weak Carcinogen Benzo[c]phenanthrene. Adv. Exp. Med. Biol.: Biological Reactive Intermediates, IIA. Snyder, R., Parke, D.V., Kocsis, J., Jollow, D.J., and Gibson, G.G.m (Eds.): Plenum Publishing Co., New York, 1982, pp. 501-524.
63. Thakker, D.R., Levin, W., Yagi, H., Conney, A.H., and Jerina, D.M.: Regio-and Stereoselectivity of Hepatic Cytochrome P-450 toward Polycyclic Aromatic Hydrocarbon Substrates. Adv. Exp. Med. Biol.: Biological Reactive Intermediates, IIA. Snyder, R., Parke, D.V., Kocsis, J., Jollow, D.J., and Gibson, G.G., (Eds.): Plenum Publishing Co., New York. 1982, pp. 525-539.
64. Thakker, D.R., Levin, W., Yagi, H., Tada, M., Ryan, D.E., Thomas, P.E., Conney, A.H., and Jerina, D.M.: Stereoselective Metabolism of the (+)-and (-)-Enantiomers of Trans-3,4-dihydroxy-3,4-dihydrobenzo[a]anthracene by Rat Liver Microsomes and by a Purified and Reconstituted Cytochrome P-450 System. J. Biol. Chem., 257: 5103-5110, 1982.
65. Van Bladeren, P.J., Armstrong, R.N., Cobb, D., Thakker, D.R., Ryan, D.M.: Stereoselective Formation of Benz[a]anthracene (+)-(5S,6R)-oxide and (+)-(8R,9S)-oxide by a Highly Purified and Reconstituted System Containing Cytochrome P-450c. Biochem. Biophys. Res. Commun., 106: 602-609, 1982.
66. Thakker, D.R., Yagi, H., Nordqvist, M., Lehr, R.E., Levin, W., Wood, A.W., Chang, R.L., Conney, A.H., and Jerina, D.M.: Polycyclic Aromatic Hydrocarbons and Carcinogenesis: The Bay-region Theory. In Arcos, J.C., Woo, Y.-T., and Argus, M.F. (Eds.): Chemical Induction of Cancer. New York, Academic Press, Inc., 1982, pp. 727-746.
67. Levin, W., Wood, A., Chang, R., Ryan, D., Thomas, P., Yagi, H., Thakker, D., Vyas, K., Boyd, C., Chu, S.-Y., Conney, A., and Jerina, D.: Oxidative Metabolism of Polycyclic Aromatic Hydrocarbons to Ultimate Carcinogens. Drug Metabolism Reviews. 13: 555-580. 1982.

68. Jerina, D.M., Michaud, D.P., Feldmann, R.J., Armstrong, R.N., Vyas, K.P., Thakker, D.R., Yagi, H., Thomas, P.E., Ryan, D.E., and Levin, W.: Stereochemical Modeling of the Catalytic Site of Cytochrome P-450c. Fifth International Symposium on Microsomes and Drug Oxidations, (Microsomes, Drug Oxidations, and Drug Toxicity). SATO, R. and KATO, R. (Eds.), Japan Scientific Societies Press, Tokyo, 1982, pp. 195-201.
69. Yagi, H., Thakker, D.R., Vyas, K.P., Chang, R.L., Wood, A.W., Levin, W., Conney, A.H., and Jerina, D.M.: The Role of Relative and Absolute Stereochemistry in the Expression of Mutagenic and Carcinogenic Activity of Metabolites from Polycyclic Aromatic Hydrocarbons. Fifth International Symposium on Microsomes and Drug Oxidations (Microsomes, Drug Oxidations and Drug Toxicity). SATO, R. and KATO, R. (Eds.), Japan Scientific Societies Press, Tokyo, 1982, pp. 539-540.
70. Lehr, R.E., Wood, A.W., Levin, W., Conney, A.H., Thakker, D.R., Yagi, H., and Jerina, D.M.: The bay region theory: History and Current Perspectives. Polycyclic Aromatic Hydrocarbons: Physical and Biological Chemistry. Sixth International Symposium, Dennis, A.J. and Cooke, W.M. (Eds.), Battelle Press, Columbus, Ohio, 1982, pp. 859-971.
71. Vyas, K.P., Yagi, H., Thakker, D.R., Chang, R.L., Wood, A.W., Levin, W., Conney, A.H., and Jerina, D.M.: Stereoselectivity in the Metabolism, Mutagenicity and Tumorigenicity of the Polycyclic Aromatic Hydrocarbon Chrysene. Polycyclic Aromatic Hydrocarbons: Physical and Biological Chemistry. Sixth International Symposium, Dennis, A.J. and Cooke, W.M. (Eds.), Battelle Press, Columbus, Ohio, 1982, pp. 859-871.
72. Thakker, D.R., Kirk, K.L., and Creveling, C.R.: Enzymatic O-methylation of Norephrine: Studies on the Site of Methylation by High Pressure Liquid Chromatography. The Biochemistry of S-Adenosylmethionine and Related Compounds, Usdin, E., Borchardt, R., and Creveling, C.R. (Eds.), MacMillan Press, London, 1982, pp. 473-477.
73. Vyas, K.P., Levin, W., Yagi, H., Thakker, D.R., Ryan, D.E., Thomas, P.E., Conney, A.H., and Jerina, D.M.: Stereoselective Metabolism of the (+)-and (-)-Enantiomers of Trans-1,2-dihydroxy-1,2-dihydrochrysene to Bay-region 1,2-Diol-3,4-epoxide Diastereomers by Rat Liver Enzymes. Mol. Pharmacol., 22: 182-189, 1982.
74. Buhler, D.R., Unlu, F., Thakker, D.R., Slaga, T.J., Newman, M.S., Levin, W., Conney, A.H., and Jerina, D.M.: Metabolism and Tumorigenicity of 7-, 8-, 9-, and 10-Fluorobenzo[a]pyrenes. Cancer Res., 42: 4779-4783, 1982.
75. Buhler, D.R., Unlu, F., Thakker, D.R., Slaga, T.J., Conney, A.H., Wood, A.W., Chang, R.L., Levin, W., and Jerina, D.M.: Effect of a 6-Fluoro Substituent on the Metabolism and Biological Activity of Benzo[a]pyrenes. Cancer Res., 43: 1541-1549, 1983.
76. Ittah, Y., Thakker, D.R., Levin, W., Croisy-Delcey, M., Ryan, D.E., Thomas, P.E., Conney, A.H., and Jerina, D.M.: Metabolism of Benzo[c]phenanthrene by Rat Liver Microsomes and by a Purified Monooxygenase System Reconstituted with Different Forms of Cytochrome P-450. Chem.-Biol. Interact., 45: 15-28, 1983.
77. Vyas, K.P., Thakker, D.R., Yagi, H., Sayer, J.M., Levin, W., and Jerina, D.M.: Regioselectivity and Stereoselectivity in the Metabolism of Trans-1,2-dihydroxy-1,2-dihydrobenz[a]anthracene. Mol. Pharmacol., 24: 115-123, 1983.
78. Yagi, H., Thakker, D.R., Ittah, Y., Croisy-Delcey, M., and Jerina, D. M.: Synthesis and Assignment of Absolute Configuration to the Trans-3,4-dihydrodiols and 3,4-Diol-1,2-epoxides of Benzo[c]phenanthrene. Tetrahedron Lett., 24: 1349-1352, 1983.
79. Wood, A.W., Chang, R.L., Levin, W., Yagi, H., Thakker, D.R., van Bladeren, P.J., Jerina, D.M., and Conney, A.H.: Mutagenicity of the Enantiomers of the Diastereomeric Bay-region Benz[a]anthracene 3,4-Diol-1,2-epoxides in Bacterial and Mammalian Cells. Cancer Res., 43: 5821-5825, 1983.
80. Levin, W., Chang, R.L., Wood, A.W., Yagi, H., Thakker, D.R., Jerina, D.M., and Conney, A.H.: High Stereoselectivity Among the Optical Isomers of the Diastereomeric Bay-region Epoxides of Benz[a]anthracene in the Expression of Tumorigenic Activity. Cancer Res., 44: 929-933, 1984.



81. Levin, W., Wood, A.W., Chang, R.L., Newman, M.S., Thakker, D.R., Conney, A.H., and Jerina, D.M.: The Effect of Steric Strain in the Bay-region of Polycyclic Aromatic Hydrocarbons: Tumorigenicity of Akyl-substituted Benz[a]anthracenes. Cancer Lett., **20**: 139-146, 1984.
82. Wood, A.W., Change, R.L., Levin, W., Thakker, D.R., Yagi, H., Jerian, D.M., and Conney, A.H.: Mutagenicity of the Enantiomers of the Diastereomeric Bay-Region Benzo[c]phenanthrene 3,4-Diol-1,2-epoxides in Bacterial and Mammalian Cells. Cancer Res., **44**: 2320-2324, 1984.
83. Jerina, D.M., Yagi, H., Thakker, D.R., Sayer, J.M., Van Bladeren, P.J., Lehr, R.E., Whalen, D.L., Levin, W., Wood, A.W., and Conney, A.H. Identification of the Ultimate Carcinogenic Metabolites of the Polycyclic Aromatic Hydrocarbons: Bay-region (R,S)-Diol-(S,R)-epoxides. In Caldwell J. and Paulson, G. D. (Editors): Foreign Compound Metabolism. Taylor and Francis Ltd., London, 1984, pp. 257-266.
84. Thakker, D.R., Yagi, H., Sayer, J.M., Kapur, U., Levin, W., Change, R.L., Wood, A.W., Conney, A.H., and Jerina, D.M.: Effects of a 6-Fluoro Substituent on the Metabolism of Benzo[a]pyrene 7,8-Dihydrodiol to Bay-region Diol Epoxides by Rat Liver Enzymes. J. Biol. Chem., **259**: 11249-11256, 1984.
85. Thakker, D.R., Yagi, H., Levin, W., Wood, A.W., Conney, A.H., and Jerina, D.M.: Metabolic Activation of Polycyclic Aromatic Hydrocarbons to Ultimate Carcinogens. In Anders M.W., (Eds.): Bioactivation of Foreign Compounds. Academic Press, New York, 1985, pp. 177-241.
86. Jerina, D.M., Sayer, J.M., Yagi, H., Van Bladeren, P.J., Thakker, D.R., Levin, W., Chang, R.L., Wood, A.W., and Conney, A.H.: Stereoselective Metabolism of Polycyclic Aromatic Hydrocarbons to Carcinogenic Metabolites. In Boobis, A.R., Caldwell, J., deMattis, S., and Elcombe, C.R., (Eds.): Microsomes and Drug Oxidations. Proceedings of the 16th International Symposium. Taylor and Francis, Ltd. London, 1985, pp. 310-319.
87. Thakker, D.R., Boehlert, C., Kirk, K.L., Antkowiak, R., and C.R. Creveling: Regioselectivity of Catechol O-Methyltransferase: The Effect of pH on the Site of O-Methylation of Fluorinated Norepinephrines. J. Biol. Chem., **261**: 178-184, 1986.
88. Thakker, D.R., Levin, W., Yagi, H., Yeh, H.J.C., Ryan, D.E., Thomas, P. E., Conney, A.H. and Jerina, D.M.: Stereoselective Metabolism of (+)-(S,S)- and (-)-(R,R)-Enantiomers of Trans-3,4-dihydroxy-3,4-dihydrobenzo[c]phenanthrene by Rat and Mouse liver Microsomes and by a Purified and Reconstituted Cytochrome P-450 System. J. Biol. Chem., **261**: 5404-5413, 1986.
89. Levin, W., Chang, R.L., Wood, A.W., Thakker, D.R., Yagi, H., Jerina, D.M. and Conney, A.H.: Tumorigenicity of Optical Isomers of the Bay-region 3,4-Diol-1,2-epoxides of Benzo[c]phenanthrene in Murine Tumor Models. Cancer Res., **46**: 2257-2261, 1986.
90. Boyd D.R., Kennedy, D.A., Malone, J.F., O'Kane, G., Thakker, D.R., Yagi, H. and Jerina, D.M.: Synthesis and Absolute Configuration of The Arene 1,2-Oxide and Trans-1,2-Dihydrodiol Metabolites of Triphenylene. Crystal Structure of (-)-(1R,2R)-Trans-2-bromo-1-menthyloxy-1,2,3,4-tetrahydrotriphenylene. J. Chem. Soc. Perkin Transactions I, 369-375, 1987.
91. Van Bladeren, P.J., Balani, S.K., Sayer, J.M., Thakker, D.R., Boyd, D.R., Ryan, D.E., Thomas, P.E., Levin, W. and Jerina, D.M.: Stereoselective Formation of Benzo[c]phenanthrene (+)-(3S,4R)- and (+)-(5S,6R)-Oxides by Cytochrome P-450c in a Highly Purified and Reconstituted System. Biochem. Biophys. Res. Commun., **145**: 160-167, 1987.
92. Yagi, H., Sayer, J.M., Thakker, D.R., Levin, W. and Jerina, D.M.: Novel Bay-region Diol Epoxides from 6-Fluorobenzo[a]pyrene. J. Am. Chem. Soc., **109**: 838-846, 1987.

93. Thakker, D.R., Levin, W., Wood, A.W., Conney, A.H., Yagi, H. and Jerina, D.M. Stereoselective Biotransformation of Polycyclic Aromatic Hydrocarbons to Ultimate Carcinogens. In Waisser, I.W. and Drayer, D.E.(Eds.): Drug Stereochemistry: Analytical Methods and Pharmacology. Marcel Dekker, New York, 1988, pp. 271-296.
94. Thakker, D.R., Boehlert, C., Mirsadeghi, S., Levin, W., Ryan, D.E., Thomas, P.E., Yagi, H., Pannell, L., Sayer, J.M., and Jerina, D.M.: Differential Stereoselectivity on Metabolism of Triphenylene by Cytochromes P-450 in Liver Microsomes from 3-Methylcholanthrene and Phenobarbital-treated Rats. J. Biol. Chem., 263: 98-105, 1988.
95. Thakker, D.R., Boehlert, C., Kirk, K.L., and Creveling, C.R.: Interaction of Fluorinated Catecholamines with Catechol O-Methyltransferase. In Dahlstrom A., Sandler, M. and Belmaker, R. (Eds.): Progress in Catecholamine Research, Part A: Basic Aspects and Peripheral Mechanisms. Alan R. Liss, New York, 1988.
96. Chang, R.L., Wood, A.W., Conney, A.H., Yagi, H., Sayer, J.M., Thakker, D.R., Jerina, D.M., and Levin, W.: Role of Diaxial versus Diequatorial Hydroxyl Groups in the Tumorigenic Activity of a Benzo[a]pyrene Bay-region Diol Epoxides. Proc. Natl. Acad. Sci. USA., 84: 8633-8636, 1988.
97. Prasad, G.K.B., Mirsadeghi, S., Boehlert, C., Byrd, A., and Thakker, D.R.: Oxidative Metabolism of the Carcinogen 6-Fluorobenzo[c]phenanthrene. Effect of a K-region Fluoro Substituent on the Regioselectivity of Cytochromes P-450 in Liver Microsomes from Control and Induced Rats. J. Biol. Chem., 263: 3676-3683, 1988.
98. Mirsadeghi, S., Prasad, G. K. B., Whittaker, N., and Thakker, D.R.: Synthesis of K-region Monofluoro- and Difluorobenzo[c]phenanthrenes. J. Org. Chem., 54: 3091-3096, 1989.
99. Thakker, D.R.: Perspective: Polycyclic Aromatic Hydrocarbon-DNA Adducts and Chemical Carcinogenesis, Drug Metabolism News Letter, 18: 1-2, 1988.
100. Thakker, D.R. and Creveling, C.R.: O-Methylation. In Mulder, G. (Ed.) Conjugation Reactions in Drug Metabolism, An Integrated Approach. Taylor and Francis, London, pp. 193-232, 1989.
101. Iyer, R.P., Phillips, L.R., Biddle, J.A., Thakker, D.R., and Egan, W.: Synthesis of Acyloxyalkyl Acylphosphonates as Potential Prodrugs of the Antiviral, Trisodium Phosphonoformate (Foscarnet Sodium). Tetrahedron Lett., 30: 7141-7144, 1989.
102. Patrick, M.A., Sethi, S., Unger, S., Mirsadeghi, S., Ribeiro, A., and Thakker, D.R.: A Novel Reaction of a Polycyclic Aromatic Hydrocarbon  $\alpha$ -Quinone with Acetone. J. Org. Chem., 56: 888-891, 1991.
103. Thakker, D.R., Boehlert, C., Levin, W., Ryan, D.E., Thomas, P.E., and Jerina, D.M.: Novel Stereoselectivity of Rat Liver cytochromes P450 Toward Enantiomers of Trans-1,2-dihydrodiol of Triphenylene. Arch. Biochem. Biophys., 288: 54-63, 1991.
104. Gan, L.-S., Hsyu, P.-H., Pritchard, J.F., and Thakker, D.R.: Mechanism of Intestinal Absorption of Ranitidine and Ondansetron Transport across Caco-2 Cell Monolayers. Pharmaceutical Res. 10: 1722-1725, 1993.
105. Augustijn, P., Gan, L.-S., Bradshaw, T., Hendren, W., and Thakker, D.R.: Evidence for the Presence of Multidrug Resistance Protein on the Apical side of the Caco-2 Cell Monolayers, Biochem. Biophys. Res. Commun., 197: 360-365 (1993).
106. Gan, L.-S., Niederer, T., Eads, C., and Thakker, D.R.: Evidence for Predominantly Paracellular Transport of Thyrotropin-Releasing Hormone (TRH) Across Caco-2 Cell Monolayers. Biochem. Biophys. Res. Commun., 197: 771-777 (1993).
107. Thakker, D.R. and St. Claire, R.L. Book Review - Chromatography Today, J. Am. Chem. Soc., 115: 8890 (1993).

108. Creveling, C.R. and Thakker, D.R.: Conjugation -DeConjugation Reactions. In Kauffman, S, ed. Drug Metabolism and Toxicity: Handbook of Experimental Pharmacology, vol 112, pp. 189-219, 1994.
109. Gan, L.-S., Eads, C., Niederer, T., Bridgers, A., Yanni, S., Hsyu, P.-H., Pritchard, F.J., and Thakker, D.R. Use of Caco-2 Cells as an In Vitro Intestinal Absorption and Metabolism Model. Drug Development and Industrial Pharmacy, 20: 615-631, 1994.
110. Prakash, S.R., Correa, I.D., Gan, L.-S., Chism, JP., Arrendale, R.F., Miwa, G.T., and Thakker, D.R. Tumor Targeting by Reductive Activation of Nitroimidazole Carbamate. In Allen, J (Ed.). In Synthesis and Application of Isotopically Labelled Compounds, John Wiley, London, pp. 573-576, 1995.
111. Sinhababu, A., Gan, L.-S., Lee, F, Boehlert, C.C., and Gan, L.-S., Yanni, S.B., Thakker, D.R.: High Performance Liquid Chromatographic Purification, Optimization of the Assay and Properties of Ribonucleoside Diphosphate Reductase from Rabbit Bone Marrow. Arch. Biochem. Biophys., 317: 285-291, 1995.
112. Gan, L.-S., Moseley, M.A., Khosla, B., Augustijns, P.F., Bradshaw, T.P., Hendren, R.W., and Thakker, D.R.: CYP3A-Like Cytochrome P450-Mediated Metabolism and Polarized Efflux of Cyclosporin A in Caco-2 Cells: Interaction between the Two Biochemical Barriers to Intestinal Transport, Drug Metabolism and Disposition, 24: 344-349, 1996.
113. Sinhababu, A. and Thakker, D.R.: Prodrugs of Anticancer Agents. Advanced Drug Delivery Reviews, 19, 241-273, 1996.
114. Gan, L.-S. and Thakker, D.R.: Applications of the Caco-2 Model in the Design and Development of Orally Active Drugs: Elucidation of Biochemical and Physical Barriers Pose by the Intestinal Epithelium. Advanced Drug Delivery Reviews, 23, 77-98, 1997.
115. Kang, S.H., Sinhababu, A.K., Cory, J.G., Mitchell, B.S., Thakker, D.R., and Cho, M.J.: Cellular Delivery of Nucleoside Diphosphates: A prodrug Approach. Pharmaceutical Res., 14, 706-712, 1997.
116. Gan, L.-S., Yanni, S., and Thakker, D.R.: Modulation of the Tight Junctions of the Caco-2 Cell Monolayers by H<sub>2</sub>-antagonists. Pharmaceutical Res., 15, 54-58, 1998.
117. Lee, K. and Thakker, D. R.: Saturable Transport of H<sub>2</sub>-Antagonists Ranitidine and Famotidine Across Caco-2 Cell Monolayers. J. Pharm. Sci., 88, 680-687, 1999.
118. Liu, D. Z., LeCluyse, E. L., and Thakker, D. R.: Dodecylphosphocholine (DPC)-mediated Enhancement of Paracellular Permeability and Cytotoxicity in Caco-2 Cell Monolayers. J. Pharm. Sci., 88, 1161-1168, 1999.
119. Liu, D. Z., Morris-Natschke, S. L., Kucera, L. S., Ishaq, K. S., and Thakker, D. R.: Structure-Activity Relationships for Enhancement of Paracellular Permeability by 2-Alkoxy-3-alkylamidopropylphosphocholines across Caco-2 Cell Monolayers. J. Pharm. Sci., 88, 1169-1174, 1999.
120. Ward, P. D., Tippin, T. K., Thakker, D. R.: Enhancement of Paracellular Permeability by Modulation of Epithelial Tight Junctions, Pharmaceutical Science and Technology Today, 3, 346-358, 2000.
121. Troutman, M. D., Luo, G., Gan, S.-S., and Thakker, D. R.: The Role of P-Glycoprotein in Drug Disposition – Significance to Drug Development. In Drug-Drug Interactions: From Basic Pharmacokinetic Concepts to Marketing Issues (Rodrigues, D., Ed.), Marcel Dekker, 2001, pp. 295-357.
122. Alfrangis, L. H., Thakker, D. R., Pope, M., Christensen, I. T., Hovgaard, L., Frokjaer, S. Structure-absorption Relationships for Tetrapeptides in MDCK Cells. Eur. J. Pharm. Sci., in press, 2002.

123. Annaert, P. P., Ryan, T., Booth, C., Thakker, D. R., and Brouwer, K. P-glycoprotein-mediated *In Vitro* Biliary Excretion of Rhodamine 123 in Sandwich-cultured Rat Hepatocytes. Drug Metab. Disp., 29, 1277-1283, 2001.
124. Chen C., and Thakker, D. R.: The Fallacy of Using Adrenochrome Reaction for Measurement of Reactive Oxygen Species formed During Cytochrome P450-mediated Metabolism of Xenobiotics. J. Pharmacol. Exp. Therap., 300, 417-420, 2002.
125. Ouyang, H., Morris-Natschke, S. L., Ishaq, K. S., Ward, P., Liu, D. Z., Leonard, S., and Thakker, D. R.: Structure-activity Relationship for Enhancement of Paracellular Permeability across Caco-2 Cell Monolayers by 3-Alkylamido-2-alkoxypropylphosphocholines, J. Med. Chem. 45, 2857-2866, 2002.
126. Lee, K., Ng, C., Brouwer, K.L.R., and Thakker, D. R.: Role of a Basolateral Transporter and P-glycoprotein in the Secretory Transport of Ranitidine and Famotidine across Caco-2 Cell Monolayers. J. Pharmacol. Exp. Ther., 303, 1-7, 2002.
127. Ward, P. D., Troutman, M. D., Desai, S. and Thakker, D. R.: Phospholipase C- $\gamma$  Regulates Tight Junction Permeability of Epithelial Barrier by Phosphorylation of the Tight Junction-associated Protein ZO-2. J. Biol. Chem., 277, 35760-35765, 2002..
128. Zhou, L., Voyksner, R. D., Hall, J. E., Thakker, D. R., Srephens, C. E., Anbazhagan, M., Boykin, D. W., Tidwell, R. R.: Characterizing the Fragmentations of 2,5-Bis(4-amidinophenyl)furan-bis-O-methylamidoxime and Selected Metabolites Using Ion Trap Mass Spectrometry. Rapid Commun. Mass Spect., 16: 1078-85, 2002.
129. Zhou, L., Lee, K., Thakker, D. R., Boykin, D., Tidwell, R. R., Hall, J. E.: Transport of 2,5-Bis(4-amidinophenyl)furan and its Novel Orally Active O-methylamidoxime Prodrug Across Caco-2 Cell Monolayers. Pharm. Res., 19: 1689-1695, 2002.
130. Ward, P. D., Ouyang, H., and Thakker, D. R. : Role of Phospholipase C- $\beta$  in the Modulation of Epithelial Tight Junction Permeability. J. Pharmacol. Exp. Ther., 304: 1-10, 2003.
131. Troutman, M. D. and Thakker, D. R.: Rhodamine 123 Requires Carrier-Mediated Influx for its Activity as a P-glycoprotein Substrate in Caco-2 Cells. Pharm. Res., 20, 1192-1199, 2003
132. Troutman, M. D. and Thakker, D. R.: The Efflux Ratio Cannot Assess P-glycoprotein-mediated Attenuation of Absorptive and Secretory Transport across Caco-2 Cell Monolayers. Pharm. Res., 20, 1200-1209, 2003
133. Troutman, M. D. and Thakker, D. R.: Novel Experimental Parameters to Quantify the Modulation of Absorptive and Secretory Transport of Compounds by P-glycoprotein in Cell Culture Models of Intestinal Epithelium. Pharm. Res., 20, 1210-1226, 2003
134. Ansele, J. H. and Thakker, D. R.: In Vitro High Throughput Screening for Inhibition and Metabolic Stability of Compounds Towards Cytochrome P450-mediated Oxidative Metabolism. J. Pharm. Sci., 93, 239-255, 2004
135. Bourdet, D. L., Lee, K., and Thakker, D. R.: Photoaffinity Labeling of the Anionic Sites in Caco-2 Cells Mediating Saturable Transport of Hydrophilic Cations Ranitidine and Famotidine. J. Med. Chem., 47, 2935-2938, 2004.
136. Thakker, D. R.: Strategic Use of Preclinical Pharmacokinetic Studies and In Vitro Models in Optimizing ADME Properties of Lead Compounds. In: Optimizing Drug-Like Properties During Discovery Lead Optimization. (Ed. Borchardt RT, Hageman, MJ, Kerns EH, Stevens, JL, and Thakker, DR), Arlington, VA, AAPS Press, 2006, pp 1-24.

137. Bourdet, D. L., Pritchard, J. B. and Thakker, D. R.: **Differential Substrate and Inhibitory Activities of Ranitidine and Famotidine toward hOCT1 (SLC22A1), hOCT2 (SLC22A2), and hOCT3 (SLC22A3).** J. Pharmacol. Exp. Ther., 315, 1288-1297, 2005.
138. Bourdet, D. L., and Thakker, D. R.: Saturable Absorptive Transport of Ranitidine in Caco-2 Cells: Role of pH-dependent Organic Cation Uptake System and P-glycoprotein. Pharm. Res., 23, 1265-1177, 2006.
139. Bourdet, D. L., Pollack, G. L., and Thakker, D. R.: Intestinal Transport of Hydrophilic Cations: A Kinetic Modeling Approach to Elucidate the Role of Uptake and Efflux Transporters and Paracellular vs. Transcellular Transport in the Absorptive Transport of Ranitidine. Pharm. Res., 23, 1178-1187, 2006.
140. Yanni, S. and Thakker, D. R.: Prodrugs: Absorption, Distribution, Metabolism, and Excretion (ADME) Issues. In Prodrugs: Challenges and Rewards (Ed Stella VJ, Borchardt RT, Hageman MJ, Oliyai R, Jilley JW, and Maag H.), Arlington VA, AAPS Press, 2006, pp 313-352.
141. Knight, B., Troutman, M, and Thakker, D. R.: Deconvoluting P-glycoprotein's Effect on Intestinal SYP3A: A Major Challenge. Current Opinions in Pharmacology, 6, 1-5, 2006.
142. Tippin, T.K., Thakker D.R.: Biorelevant Refinement of the Caco-2 Cell Culture Model to Assess Efficacy of Paracellular Permeability Enhancers. J. Pharm. Sci., 97, 1977-1993 2008.
143. Troutman, M., Luo, G., Knight, B. M., Thakker, D., and Gan, L. S.: The Role of P-Glycoprotein in Drug Disposition: Significance to Drug Development in *Drug-drug Interactions* (Rodrigues A. D., ed.), Marcel and Dekker, New York, NY, 2008, pp. 359-434.
144. Yanni, S.B., Annaert, P. Augustijns, P., Bridges, A.A., Gao, Y., Benjamin, Jr., D.K., and Thakker, D.R. Role of Flavin-containing Monooxygenase in Oxidative Metabolism of Voriconazole by Human Liver Microsomes. Drug Met. Disp., 36, 1119-1125, 2008.
145. Proctor, W.R., Bourdet, D.L., and Thakker, D.R. Mechanisms Underlying Saturable Intestinal Absorption of Metformin. Drug Met. Disp., 36, 1650-1658, 2008.
146. Ming X., Ju, W., Wu, H., Tidwell, R.D., Hall, J.E., and Thakker, D.R. Transport of Dicationic Drugs Pentamidine and Furamidine by Human Organic Cation Transporters. Drug Met. Disp., 37, 424-430, 2009.
147. Proctor, W.R., Ming, X., and Thakker, D.R.: In Vitro Techniques to Study Drug-Drug Interactions Involving Transport: Caco-2 Model for Study of P-glycoprotein and Other Transporters in *Enzymatic- and Transporter-Based Drug-Drug Interactions* (Peng, S., Rodrigues A.D., and Peter, R.), Springer, New York, NY, 2010, pp 257-282.
148. Ming X. and Thakker, D.R.: MRP4 Mediates Basolateral Efflux of Adefovir Formed in Caco-2 Cells from its Prdrug – Adefovir Dipivoxil, Biochem. Pharmacol, 79, 455-462, 2010 (Epub 2009 Sep 6).
149. Yanni, S.B., Annaert, P.P., Augustijns, P, Ibrahim, J.G., Benjamin, Jr., D.K., and Thakker, D.R.: *In Vitro* Hepatic Metabolism Explains Higher Clearance of Voriconazole in Children versus Adults: Role of CYP2C19 and Flavin-containing Monooxygenase-3. Drug Met. Disp., 38, 25-31, 2010.

150. Yanni S.B., Benjamin D.K., Augustijns P, Brouwer, K.L., Thakker D.R., and Annaert P.P. *In vitro* investigation of the hepatobiliary disposition mechanism of the antifungal agent micafungin in humans and rats. *Drug Met. Disp.*, 38, 1848-1856, 2010.
151. Yanni S.B., Smith P.B., Benjamin D.K., Augustijns P., Thakker D.R., and Annaert PP. Annaert. Age-dependent disposition of micafungin in humans: Effect of plasma binding. *Biopharm. Drug Disp.*, 32, 222-232, 2011.
152. Klein R.R., Bourdon D.M., Costales C.L., Wagner C.D., White W.L., Williams J.D., Hicks S.N., Sondek J., and Thakker D.R., Direct Activation of Human Phospholipase C $\beta$ 3 by its Well Known Inhibitor U73122, *J. Biol. Chem.*, 286, 12407-12416, 2011, (e-pub Jan 25 2011).
153. Ming X, Knight, B.M., and Thakker, D.R. Vectorial Transport of Fexofenadine across Caco-2 Cells: Involvement of Apical Uptake and Basolateral Efflux Transporters, *Mol. Pharmaceutics*, 8, 1677-1686, 2011 (e-pub Aug 5 2011).
154. Knight, B.M., Proctor, W.R., Ming, X., Bridges, A., and Thakker, D.R. P-glycoprotein Attenuates Oxidative Metabolism of Terfenadine during Absorptive Transport across Mouse Intestinal Tissue but not across Caco-2 Cell Monolayers, submitted to *Biochemi. Pharmacol.*, 2011.
155. Dufek, M.B., Knight B.M., Bridges, A.S., and Thakker, D.R., P-glycoprotein (P-gp) Reduces First-pass Intestinal Metabolism and Enhances Portal Bioavailability of Loperamide in Mouse, *Drug Met. Disp.*, 41, 642-650, 2013
156. Generaux, C.N., Ainslie, G.R., Bridgers, A.S., Ismail, M.A., Boykin, D.W., Tidwell, R.R., Thakker, D.R., and Pain, M.F. Compartmental and Enzyme Kinetic Modeling to Elucidate the Biotransformation Pathway of a Centrally Acting Antitripanosomal Drug, *Drug Met. Disp.*, 41, 518-528, 2013
157. Swift, B., Nebot, N., Lee, J.K., Han, T., Proctor, W.R., Thakker, D.R., Lang, D., and Radtke, M., Gnoth, M.J., and Brouwer, K.L. Sorafenib Hepatobiliary Disposition: Mechanisms of Hepatic Uptake and Disposition of Generated Metabolites. *Drug Met. Disp.*, E-Pub, 2013
158. Han, Tianxiang, Everett R, Proctor W.R., Ng, C.M., Costales, C.L., Brouwer, K.L.R., and Thakker D.R. "Apical Localization of Organic Cation Transporter 1 in Caco-2 Cell Monolayers and in Mouse and Human Intestinal Epithelium." *Mol Pharmacol*, 84, 182-189, 2013
159. Dufek, M.B., Bridges, A.S., and Thakker, D.R., Intestinal First-pass Metabolism by Cytochrome P450 and not P-gp is the Major Barrier to Amrpenavir, *Drug Met. Disp.*, 41, 1695-1702, 2013
160. Varma Alluri R., Ward P., Kunta, J., Ferslew, B.C., Thakker, D.R., and Dallas, S. "In Vitro Characterization of Intestinal and Hepatic Transporters: MRP2" in *Optimization in Drug Discovery In Vitro Methods, 2<sup>nd</sup> edition* (Casey D.C.), Humana Press, New York, NY, 2013, pp
161. Zane N and Thakker D.R., "'A Physiologically Based Pharmacokinetic Model for Voriconazole Disposition Predicts Intestinal First-pass Metabolism in Children". *Clin Pharmacokinet*, 2014, 53, 1171-1182, 2014
162. Han T.K., Proctor W.R., Costales C.L., Cai H., Everett R.S., Thakker D.R., "Four Cation-Selective Transporters Contribute to Apical Uptake and Accumulation of Metformin in Caco-2 Cell Monolayers". *J. Pharmacol. Exp. Ther.* 352, 519-528, 2015

163. Zane, N.R. and Thakker, D.R., A Letter to the Editor, Response to Prediction of Non-Linear Pharmacokinetics Using a Pediatric Physiologically Based Pharmacokinetic Modelling Approach, Clin. Pharmacokin. DOI 10, 1907/s40262-4015-4054-6, 2015.
164. Ohutsu, Y, Thakker D.R., Gibbons, J.A., Tsubota, K, Otsuka, S, and Arai, H. Determination of the Androgen Receptor Inhibitor Enzalutamide and Its Metabolites in Animal Plasma and Brain Homogenates Using LC/MS-MS and Its Application to Pharmacokinetic Studies. Chromatography, 36, 115-122, 2015 (published in 2015 but was not listed last year)
165. Cai, H., Zhang U., Han, T.K., Everett R., and Thakker, D.R. "Cation-selective Transporters are Critical to the AMPK-mediated Antiproliferative Effects of Metformin in Human Breast Cancer Cells." Int J Cancer. 138(9):2281-92, 2016.
166. Proctor W.R., Ming X, Bourdet D, Han T.K., Everett R.S., Thakker D.R. Why Does the Intestine Lack Basolateral Efflux Transporters for Cationic Compounds? A Provocative Hypothesis. J Pharm Sci. 105(2):484-96, 2016.
167. Chen Y, Zane NR, Thakker DR, and Wang MZ. Quantification of Flavin-containing Monooxygenases 1, 3 and 5 in Human Liver Microsomes by UPLC-MRM-based Targeted Quantitative Proteomics and Its Application to the Study of Ontogeny. Drug Metab Disp. 44(7):975-983, 2016
168. Hao Cai, Muhammad Wahajuddin, Nicole Zane, Ruth S. Everett, and Dhiren R. Thakker. Cation-selective Transporters Drive the Antitumor Efficacy of Metformin against Breast Cancer: An Unequivocal Evidence. Int J Cancer, under review.
169. Han, T.K. and Thakker, D.R., Role of Serotonin Reuptake Transporter (SERT) in the Intestinal Absorption and Adverse Effects of Metformin. In Preparation for submission in 2016. Target publication Science
170. Proctor, W.R., Van Itallie, C.M., Wang, S., Anderson, J.M., and Thakker, D.R. "Vitamin D<sub>3</sub> enhances paracellular transport of metformin across Caco-2 cell monolayers via induction of claudin-2", Manuscript in Preparation for submission in 2016. Target publication Mol Pharmacol.
171. Costales, C. and Thakker, D.R. "Cation-selective transporters involved in apical uptake and accumulation of metformin in mouse intestinal epithelium." Manuscript in preparation for submission in 2016. Target publication: Drug Met Disp.
172. Costales, C., Mackiovic, B., and Thakker, D.R. "Transporters involved in the Apical Uptake and Efflux of Metformin in Mouse Intestinal Epithelium", Manuscript in preparation for submission in 2016. Target publication: *Drug Met Disp*.
173. Costales, C.L., Alluri, R.V., and Thakker, D.R., "Metformin Intestinal Absorption in Mice is Enhanced by Cycling between Enterocytes and Intestinal Lumen: A Novel Oral Absorption Mechanism for the Antidiabetic Drug", Manuscript in preparation for submission in 2016. Target publication: Mol Pharmacol.
174. Alluri, R.V., Costales, C.L., and Thakker, D.R., "Interaction between Commonly Used Anesthetic Agents Ketamine/Xylazine and Organic Cation Drug Metformin at the Intestinal Epithelium", Manuscript in preparation for submission in 2016. Target publication: Drug Met Disp

175. Proctor, W.R., Han, T., Van Itallie, C.M., Holmes, J., Wang, S., Anderson, J.M., and Thakker, D.R. "Cation-selective claudins facilitate paracellular transport of small organic cations across epithelial monolayers." Manuscript in preparation for submission in 2016. Target publication J Cell Sci.
176. Zane NR, Chen Y, Wang MZ, and Thakker DR. Age-Dependent Differences in Expression and Functional Activity of CYP and FMO Families. Manuscript in preparation for submission in 2016. Target publication Ped Research.
177. Zane NR, Bridges AS, and Thakker DR. Improving In Vitro Techniques to Predict Intestinal First-pass Metabolism in Children. Manuscript in preparation for submission in 2016. Target publication DMD.
178. Zane NR, Laughon MM and Thakker DR. A Systems Pharmacology Approach to Predicting Clinical Exposure of Sildenafil in Premature Neonates. Manuscript in preparation for submission in 2016. Target publication CPT.

## BOOKS

1. Pharmaceutical Profiling in Drug Discovery for Lead Selection. Borchardt, R. T., Kerns, E. H., Lipinski, C. A. Thakker, D. R., and Wang, B. (Editors). AAPS Press, Arlington, VA, 2004.
2. Optimizing Drug-Like Properties during Discovery Lead Optimization. Borchardt R. T, Hageman, M. J., Kerns E. H., Stevens, J. L., and Thakker, D. R. (Editors) AAPS Press, Arlington, VA, 2006.





## PATENTS

1. Method of Screening Candidate Compounds for Susceptibility to Oxidative Metabolism, US 6,312,917.
2. PLC-based Absorption Enhancers, US 7,713,949
3. Pharmacokinetic Tool and Method for Predicting Metabolism of a compound in a Mammal, EP 1386274 (A2) WO 02/10746
4. High Throughput Reactive Oxygen Species-Based Cytochrome P450 Inhibition Assay. Patent pending US 2006/0223135 (abandoned).
5. Methods and Kits for Determining Metabolic Stability of Compounds. Patent pending US 2006/0046278 (abandoned).
6. Direct Activation of Human Phospholipase C $\beta$ 3 (HPLC $\beta$ 3) by U73122 in Dodecylmaltocide (DDM) Mixed Micelles via Alkylation at Cysteine Residues, US Patent issued

## ABSTRACTS

1. Borchardt, R.T. and Thakker, D.R. "Affinity Labeling of Catechol-O-methyltransferase," 9th Midwest American Chemical Society Meeting, 1973, Lawrence, Kansas.
2. Borchardt, R.T. and Thakker, D.R.: "Affinity Labeling of Catechol-O-methyltransferase," 10th Midwest American Chemical Society Meeting, 1974, Iowa City, Iowa.
3. Borchardt, R.T., Thakker, D.R. and Cheng, C.F.: "Purification of Catechol-O-methyltransferase by Affinity Chromatography," 10th Midwest American Chemical Society Meeting, " 1974, Iowa City, Iowa.
4. Borchardt, R.T. and Thakker, D.R.: "Affinity Labeling of Catechol-O-methyltransferase," 6th International Congress of Pharmacology, 1975, Helsinki, Finland.
5. Jerina, D.M., Yagi, H., Thakker, D.R., Holder, G., Hernandez, O., Dansette, P.M., Cheng, P., Wislocki, P., Lu, A.Y.H., Wood, A.W., Levin, W. and Conney, A.H.: "A Metabolism oriented Approach to the Carcinogenicity of Benzo[a]pyrene," Alfred Benzo Symposium X, 1976, Copenhagen.
6. Thakker, D.R., Lu, A.Y.H., Levin, W., Conney, A.H. and Jerina, D.M.: "Stereospecificity of Rat Liver Cytochrome P-450 Monooxygenase and Epoxide Hydrase," Federation of American Societies of Experimental Biology, 1977, Chicago, Illinois.
7. Bresnick, E., Vaught, J.B., Thakker, D.R., Jerina, D.M. and Stoming, T.A.: Nuclear Metabolism of Benzo[a]pyrene and of (+)-Trans-7,8-dihydroxy-7,8-dihydrobenzo[a]pyrene," Proceedings of American Association for Cancer Research, 19, 1977.
8. Karle, J.M., Lehr, R.E., Yagi, H., Thakker, D.R., Jerina, D.M., Wood, A.W., Levin, W. and Conney, A.H.: "Carcinogenesis by Polycyclic Aromatic Hydrocarbons: The Bay-Region Theory," Proceeding of American Association for Advancement of Sciences, 1978.
9. Thakker, D.R., Levin, W., Wood, A.W., Conney, A.H., Stoming, T.A. and Jerina, D.M.: "9,10-dihydrodiols of 1-hydroxy-3-methylcholanthrene: Potential Proximate Carcinogens Derived from 3-methylcholanthrene," Proceedings of American Association for Cancer Research, 19, 200, 1978.
10. Thakker, D.R., Yagi, H., Lehr, R.E., Levin, W., Lu, A.Y.H., Chang, R.L., Wood, A.W., Buening, M., Conney, A.H. and Jerina, D.M.: "Metabolism of Benzo[a]pyrene 9,10-Dihydrodiol," Federation Proceedings, 37, 749, 1978.
11. Thakker, D.R., Levin, W., Turujman, S., Kopadia, D., Conney, A.H. and Jerina, D.M.: "Absolute Stereochemistry of the Trans Dihydrodiols Formed from Benzo[a]anthracene by Rat Liver Microsomes," Proceedings of American Association for Cancer Research, 20, 1979.
12. Thakker, D.R., Levin, W., Yagi, H., Tada, M., Conney, A.H. and Jerina, D.M.: "Comparative Metabolism of Dihydrodiols of polycyclic aromatic hydrocarbons to bay-region diol epoxides," The Fourth International Symposium on Polynuclear Aromatic Hydrocarbons, Columbus, Ohio, 1979.
13. Thakker, D.R., Yagi, H., Levin, W., Wood, A.W., Conney, A.H. and Jerina, D.M.: "Metabolic formation and reactions of diol epoxide: Ultimate Carcinogenic Metabolites of the polycyclic aromatic hydrocarbons," 178th National ACS Meeting, American Chemical Society Division of Environmental Chemistry, Abstract #81, Washington, DC.
14. Thakker, D.R., Levin, W., Wood, A.W., Lehr, R.E., Kumar, S., Conney, A.H. and Jerina, D.M.: "Metabolism and Metabolic Activation of Benzo[e]pyrene 9,10-dihydrodiol by Hepatic Microsomes from Several Species," Proceedings of the American Association for Cancer Research, 21, 89, 1980.

15. Thakker, D.R., Levin, W., Yagi, H., Tada, M., Ryan, D.G., Thomas, P.E., Conney, A.H. and Jerina, D.M.: "Stereoselective Metabolism of (+)- and (-)-Benzo[a]anthracene 3,4-dihydrodiols by Rat Liver Microsomes and a Reconstituted Cytochrome P-450 System," *Proceeding of the American Association for Cancer Research*, 22, 98, 1981.
16. Thakker, D.R., Ittah, Y., Levin, W., Crosy-Delcey, M., Conney, A.H. and Jerina, D.M.: "Metabolism of Benzo[c]phenanthrene by Rat Liver Microsomes and by a Highly Purified and Reconstituted Cytochrome P-450 System," *13th International Cancer Congress, Seattle, WS, 1982.*
17. Vyas, K.P., Yagi, H., Tada, M., Thakker, D.R., Ryan, D.E., Thomas, P.E., Conney, A.H., Levin, W., and Jerina, D.M.: "Stereoselective Metabolism of the (+)- and (-)-Enantiomers of Trans-1,2-dihydroxy-1,2-dihydrochrysene to Bay-region 1,2-Diol-3,4-epoxides by Rat Liver Enzymes," *Federal Proceedings*, 41,1222, 1982.
18. Thakker, D.R., Bubler, D., Yagi, H., Unlu, F., Kapur, U., Slaga, T.J., Conney, A.H., Change, R.L., Wood, A.W., Levin, W., and Jerina, D.M.: "Metabolism of 6-Fluorobenzo[a]pyrene (FBP) to Bay-region Diol Epoxides by Rat Liver Microsomes," *Proceedings of the American Association for Cancer Research*, 24, 75, 1983.
19. Van Bladeren, P., Thakker, D.R., Vyas, K., Yagi, H., Ryan, D., Thomas, P., Levin, W., and Jerina, D.M.: "Mapping the Active Site of Cytochrome P-450c with (+)- and (-)-Trans-1,2-dihydroxy-1,2-dihydrobenz[a]anthracene and Anthracene," *Proceedings of the American Association for Cancer Research*, 24, 75, 1983.
20. Wood, A., Chang, R.L., Levin, W., Thakker, D.R., Yagi, H., Jerina, D.M., and Conney, A.H.: "Mutagenicity of Enantiomers of the Diastereomeric Bay-region Benzo[c]phenanthrene 3,4-diol-1,2-epoxides," *Proceeding of American Association for Cancer Research*, 24, 95, 1983.
21. Thakker, D.R., Yagi, H., Levin, W., Ryan, D., Thomas, P.E., Conney, A.H., and Jerina, D.M.: "Regio- and Stereoselective Metabolism of (+)-and (-)-Benzo[c]phenanthrene 3,4-dihydrodiols to Bay-region Diol Epoxides, Bisdihydrodiols and Phenolic Dihydrodiols." *Proceedings of the American Association for Cancer Research*, 25, 118, 1984.
22. Wood, A.W., Change, R.L., Levin, W., Yagi, H., Thakker, D.R., Sayer, J.M., Jerina, D.M. and Conney, A.H.: "Effects of a 6-Fluoro Substituent on the Biological Activity of Bay-region Diol Epoxides of Benzo[a]pyrene." *Proceedings of the American Association for Cancer Research*, 25, 97, 1984.
23. Chang, R.L., Levin, W., Wood, A.W., Thakker, D.R., Yagi, H. Jerina, D.M., and Conney, A.H.: "Tumorigenicity of the Enantiomers of the Diastereomeric Bay-region Benzo[c]phenanthrene 3,4-diol-1,2-epoxides (B[c]ph Diol Epoxides) on Mouse Skin and in Newborn Mice." *Proceedings of the American Association for Cancer Research*, 25, 97, 1984.
24. Thakker, D.R., Yagi, H., Levin, W., Sayer, J.M., Change, R.L., Wood, A.W., Conney, A.H., and Jerina, D.M.: "Effects of a 6-Fluoro Substituent on the Metabolic Formation and Mutagenic Activity of the Bay-region 7,8-Diol-9,10-epoxides of Benzo[a]pyrene." *Proceeding of the 6th International Symposium on Microsomes and Drug Oxidations*, 1984.
25. Thakker, D.R., Shirai, N., Levin, W., Ryan, D.E., Thomas, P.E., Lehr, R.E., Conney, A.H., and Jerina, D.M.: "Metabolism of Dibenz[c,h]acridine by Rat Liver Microsomes and by Cytochrome P-450c With and Without Epoxide Hydrolase." *Proceedings of the American Association for Cancer Research*, 26, 114, 1985.
26. Thakker, D.R., Boehlert, C., Levin, W., Yagi, H., and Jerina, D.M.: "Metabolism of Triphenylene by Liver Microsomes from Control, Phenobarbital (PB)-treated and 3-Methylcholanthrene (MC)-treated Rats." *Proceedings of the American Association of Cancer Research*, 26, 113, 1985.
27. Thakker, D.R., Boehlert, C., Kirk, K.L., and Creveling, C.R.: "Site-selective O-Methylation of Fluorinated Norepinephrines by Catechol O-Methyltransferase." *Proceedings of the International Symposium on the Biochemistry of S-Adenosylmethionine as a Basis for Drug Design*, 1985.

28. Wood, A.W., Levin, W., Chang, R.L., Lehr, R.E., Thakker, D.R., Yagi, H., Sayer, J.M., Jerina, D.M. and Conney, A.H.: "Stereochemical and Electronic Determinants of the Mutagenicity and Tumorigenicity of Polycyclic Hydrocarbon Bay-region Diol Epoxides." Proceeding of UCLA Symposium on Biochemical and Molecular Epidemiology of Cancer, 1985.
29. Thakker, D.R., Levin, W., Yagi, H., Yeh, H.J.C., Ryan, D.E., Thomas, P. E., Conney, A.H. and Jerina, D.M.: "Stereoselectivity in the Metabolism of (+)-(S,S)- and (-)-(R,R)-Benzo[c]phenanthrene 3,4-dihydrodiols by Rat and Mouse Liver Microsomes and by Cytochrome P-450c." Proceedings of the American Association of Cancer Research, 27, 116, 1986.
30. Thakker, D.R., Boehlert, C., Levin, W., Conney, A.H., Ryan, D.E., Thomas, P.E., Yagi, H. and Jerina, D.M.: "Novel Stereoselectivity in the Metabolism of the Enantiomeric Trans 1,2-Dihydrodiols of Triphenylene to Diol Epoxides by Hepatic Cytochromes P-450." Proceedings of the American Association for Cancer Research, 27, 116, 1986.
31. Prasad, G., Boehlert, C. and Thakker, D.R.: "Metabolism of Carcinogenic 6-Fluorobenzo[c]phenanthrene by Hepatic Cytochromes P-450 from Control and Induced Rats." Proceedings of the American Association for Cancer Research, 28, 124, 1987.
32. Schaffer, V. Mpanza, Z. and Thakker, D.R.: "Stereoselective Metabolism of (+)-(1S,2S)- and (-)-(1R,2R)-Dihydrodiols of Naphthalene by Hepatic Cytochromes P-450 from Control and Induced Rats." Federation Proceedings, 46, 865, 1987.
33. Ayubi, N., Boehlert, C., Creveling, C.R. and Thakker, D.R.: "Purification of Catechol O-Methyltransferase by High Performance Liquid Chromatography." Federation Proceedings 46, 2277, 1987.
34. Mirsadeghi, S., Prasad, G. and Thakker, D.R.: "Synthesis of Fluorinated Benzo[c]phenanthrenes." Proceedings of the Pacific regional meeting of the American Chemical Society, 1987.
35. Thakker, D.R.: "Stereochemical Considerations in the Metabolic Activation of Polycyclic Aromatic Hydrocarbons to Ultimate Carcinogens." Proceedings of the Pharmaceutical Society of Japan, 48-50, 1987.
36. Prasad, G., Mirsadeghi, S., Boehlert, C., and Thakker, D.R.: "Influence of a Fluoro Substituent on the Regio- and Stereoselectivity of Rat Liver Cytochrome(s) P-450 in the Metabolism of Benzo[c]phenanthrenes." Proceedings of the 2nd Annual Meeting of International Society for the Study of Xenobiotics, 1988.
37. Patrick, M.A., Sethi, S., Unger, S., Rebeiro, A., Mirsadeghi, S., and Thakker, D.R.: "A Novel Reaction of K-Region  $\alpha$ -Quinones of Polycyclic Aromatic Hydrocarbons With Acetone." Proceedings of the 199th Annual Meeting of American Chemical Society, 1990.
38. Patrick, M., Levin, W., Wood, A.W., Mirsadeghi, S., and Thakker, D.R.: "Metabolic Basis for Low Tumorigenicity of 6,7-Difluorobenzo[c]phenanthrene (6,7-F<sub>2</sub>BcPh)." Proceedings of the American Association for Cancer Research, 31, 112, 1990.
40. Boehlert, C., Lee, F., Gan, L.-S., Levesque, D., Sinhababu, A.K., and Thakker, D.R.: "Rapid and Reproducible Purification of Ribonucleoside Diphosphate Reductase from Rabbit Bone Marrow by HPLC Gel Filtration and Kinetics of Its Inactivation." Proceedings of the American Association for Cancer Research, 31, 11, 1990.
41. Prakash, S.R., Correa, I.D., Unger, S., Davis, I., Miwa, G.T., and Thakker, D.R.: "Study of the Solvolysis of Nitrogen Mustards by <sup>1</sup>H NMR." Proceedings of the American Association for Cancer Research, 31, 332, 1990.
42. Sinhababu, A. K., Boehlert, C., Levesque, D., Gan, L.-S., and Thakker, D. R.: "Mechanism of Inhibition of Ribonucleoside Diphosphate Reductase from Rabbit Bone Marrow by 2'azido-2'-deoxycytidine-5'diphosphate." Proceedings of the American Association for Cancer Research, 32, 12, 1991.

43. Lahey, A. P., Niederer, T., Tung, T., St. Claire III, R. L., Gan, L.-S., and Thakker, D. R.: "Transport and Metabolism of Dipeptide L-Alanyl-L-alanine in a Cell culture System for the Intestinal Epithelium." *The FASEB Journal*, A762, 1991.
44. Thakker, D. R., Patrick, M. A., Prasad, K. B., Boehlert, C., and Mirsadeghi, S.: "Effect of the Fluoro Substituent on the Regio- and Stereoselectivity of Cytochromes P-450-catalyzed Oxidation of Polycyclic Aromatic Hydrocarbons." *Proceedings of the Third International Meeting of the International Society for the Study of Xenobiotics (ISSX)*, 302, 1991.
45. Gan, L.-S., Hsyu, P.-H, Pritchard, J.F., and Thakker, D.R.: "Study of Intestinal Absorption of Ranitidine and Ondansetron Using the In Vitro Caco-2 Cell Culture System." *Pharmaceutical Research*, 8, S217, 1991.
46. Sinhababu, A.K., Levesque, D.L., Ghate, J., and Thakker, D.R.: "Inactivation of Ribonucleoside Diphosphate Reductase from Rabbit Bone Marrow by Nucleotide 2',3'-Dialdehydes." *Proceedings of the American Association for Cancer Research*, 33, 1992.
47. Sinhababu, A.K., Boehlert, C.C., Levesque, D.L., Yanni, S.B., Gan, L.-S., and Thakker, D.R.: "Inactivation of Ribonucleoside Diphosphate Reductase from Rabbit Bone Marrow by 2'-Azido-2'-deoxyCDP." *Proceedings of the American Association for Cancer Research*, 33, 29, 1992.
48. Levesque, L.L., Sinhababu, A.K., Keefer, L.K., Saavedra, J.E. and Thakker, D. R.: "Inhibition of Ribonucleoside Diphosphate Reductase from Rabbit Bone Marrow by Nitric Oxide/Nucleophile Complexes." *American Association of Cancer Research*, 1993.
49. Gan, L.-S., Eads, C., Niederer, T., and Thakker, D.R.: Study of Intestinal Transport of Glycylsarcosine and Thyrotropin-Releasing Hormone Using the In Vitro Caco-2 Cell Culture System. *FASEB Journal*, 7, A578, 1993.
50. Bradshaw, T.P., Brouwer, K.L.R., and Thakker, D.R.: The study of the Hepatic Extraction of Thyrotropin Releasing Hormone and Its Free Acid Analogue Using Isolated Rat Hepatocytes and Isolated Perfused Rat Liver Model. *ISSX Proceedings*, 4, 221, 1993.
51. Augustijns, P., Bradshaw, T., Gan, L.-S., Hendren, W. and Thakker, D. Drug Transport Modulation by a Polarized Efflux System in Caco-2 Cells. *Journal de Pharmacie de Belgique*, 1994.
52. Bridgers, A., Russell, D., Gan, L.-S., Brown, F., Profeta Jr., S., and Thakker, D.: "The Influence of Molecular Structure on Drug Transport across Intestinal Epithelial Caco-2 Cells." *The FASEB Journal*, 8, A3, 1994.
53. Gan, L.-S., Boehlert, C., Thakker, D., and Serabjit-Singh, C.: "Inhibition of CYP1A, CYP2B, and CYP3A Activities by 5HT<sub>3</sub> Antagonists." *The FASEB Journal*, 8, A1248, 1994.
54. Yanni, S., Gan, L.S., and Thakker, D.R.: "Modulation of Tight Junctions of Intestinal Epithelia by H<sub>2</sub>-Antagonists." *Pharmaceutical Research*, 11, S250, 1994.
55. Prakash, S., Correa, I., Gan, L.-S., Chism, J., Arrendale, R., Miwa, G., and Thakker, D.: "Tumor Targeting by Reductive Activation of Nitroimidazole Carbamates." *Proceedings of 5th International Symposium on the Synthesis and Applications of Isotopically Labelled Compounds*, 1994.
56. Gan, L.-S., Moseley, A., Khosla, B., Augustijns, P., Bradshaw, T., Hendren, W., and Thakker, D.: "Formation and Polarized Efflux of Cyclosporin A Metabolites in the In Vitro Caco-2 Cell Culture System." *The FASEB Journal*, 9, A367, 1995.
57. Thakker, D.R. and Sinhababu, A.K.: "Prodrug Design: Opportunities for Cancer Chemotherapy." *Proceedings of the Western Biotech Conference, San Diego, CA*, p.172, 1995.

58. Gan, L.-S. and Thakker, D.R.: "Attenuation of Polarized Efflux of Matrix Metalloprotease (MMP) Inhibitors across Caco-2 Cell Monolayers via Prodrug Approaches." *Pharmaceutical Research*, 13, S240, 1996.
59. Desai, S.R., Liu, D.Z., and Thakker, D.R.: "The Unusual Effect of Pyridyl Group on the Transcellular Transport of Compounds Across the Caco-2 Cell Monolayers." *Pharmaceutical Research*, 14, S21, 1997.
60. Liu, D.Z. and Thakker, D.R.: "Modulation of Caco-2 Cell Tight Junctions by Dodecylphosphorylcholine (DPC)." *Pharmaceutical Research*, 14, S23, 1997.
61. Lee, K. and Thakker, D.R.: "Modulation of the Tight Junctions in Caco-2 Cell Monolayers by H<sub>2</sub>-Antagonists." *Pharmaceutical Research*, 14, S23, 1997.
62. Lee, K. and Thakker, D. R.: A Novel Saturable Mechanism for the Paracellular Drug Transport across Caco-2 cell Monolayers, *PharmSci*, 1, S-452, 1998.
63. Lee, K. and Thakker, D. R.: Photoaffinity Labeling of the Anionic Cellular Components Mediating Paracellular Saturable Transport of Cations across Caco-2 Cell Monolayers, *PharmSci*, 1, S-452, 1998.
64. Lee, K. and Thakker, D. R.: Potential for Drug-Drug Interactions in the Absorption of Cationic Compounds across Intestinal Epithelium, *PharmSci*, 1, S-453, 1998.
65. Desai, S. R., Lee, K. and Thakker, D. R.: Use of the Nonionic Surfactant Polysorbate 80 in the Measurement of Intrinsic Transport Rate of Highly Lipophilic Compounds across Caco-2 Cell Monolayers, *PharmSci*, 1, S-456, 1998.
66. Chen, C., Ramirez, S., Kinder, D. S., Geysen, H. M., and Thakker, D. R.: Determination of Permeability of Peptoids from a Combinatorial Library with Caco-2 Assay System, *PharmSci*, 1, S-139, 1998.
67. Chen, C., Ward, P. D., and Thakker, D. R.: Inactivation of Cytochrome P450 3A4 (CYP3A4) by Epinephrine and Adrenochrome, *PharmSci*, 1, S-37, 1998.
68. Chen, C. and Thakker, D. R.: The Fallacy of the Use of Adrenochrome Reaction for Measurement of Cytochrome P450-mediated Reactive Oxygen Species (ROS) Formation. *PharmSci*, 1, S-600, 1998.
69. Liu, D.Z., Morris-Natschke, S. L., Ishaq, K.S., and Thakker, D. R.: Structural Requirements for Synthetic Phospholipids that Modulate Tight Junction Permeability across Caco-2 Cells. *PharmSci*, 1, S-453, 1998.
70. Annaert, P.P. Laethem, R.M., Thakker, D.R., LeCluyse, E.L., Brouwer, K.L.R.: *In Vitro* Biliary Excretion of the P-glycoprotein (P-gp) Substrate Rhodamine-123 (rh123) by Sandwich-Cultured Rat Hepatocytes. *PharmSci*.1999.
71. Lee, K., Thakker, D.R.: Carrier-Mediated Secretory Transport of H<sub>2</sub>-Antagonists Ranitidine and Famotidine across Caco-2 Cell Monolayers. *PharmSci*.1999.
72. Troutman, M., Thakker, D.R.: Structure-Activity Relationship for P-glycoprotein (P-gp) mediated Efflux of Steroids during Transport Across Caco-2 Cells. *PharmSci*.1999.
73. Ward, P.D., Thakker, D.R.: Hexadecylphosphocholine (HPC) is a Highly Potent Enhancer of Paracellular Permeability across MDCK and Caco-2 Cell Monolayers. *PharmSci*.1999.
74. Desai, S.R., Thakker, D.R.: The Solid Phase Synthesis of Intestinal ACAT Inhibitor RP64477. 218th ACS National Meeting, New Orleans, LA (1999).
75. Rao, G.A., Rabinowitz, J., Samulski, R.J, and Thakker D.R.: A Novel Method For Dual Isotope (3H/14C) Labeling Of Plasmid DNA. *PharmSci*, 1, S-629 1999.

76. Liu, D.Z., Ward, P.D., and Thakker, D.R.: Redistribution and Hyperphosphorylation of Epithelial Tight Junctional Components by Dodecylphosphocholine. *PharmSci*. 2000.
77. Ward, P.D., Ouyang, H., and Thakker, D.R.: Phospholipase C Inhibition is Associated with the Increase in Paracellular Permeability across MDCK Cell Monolayers. *PharmSci*. 2000.
78. Troutman M.D., Carson S.W., Hill-Zabala C.E., and Thakker D.R.: Activation and Inhibition of P-glycoprotein (P-gp) Mediated Efflux of Digoxin by St. John's Wort Extract. *PharmSci*. 2000.
79. Troutman M.D. and Thakker D.R.: Permeability of a 6-Aza, 17-Carboxylate Steroid across Caco-2 cell monolayers is affected by basolaterally and apically directed transporters other than P-glycoprotein(P-gp). *PharmSci*. 2000.
80. Ward, P.D., Troutman, M.D., Desai, S., and Thakker, D.R.: 3-Nitrocoumarin Increases Paracellular Permeability Across MDCK Monolayers by Inhibition of Phospholipase C And Hyperphosphorylation of Tight Junction Protein, ZO-2. *PharmSci*. 2001.
81. Rao, G.A., Rabinowitz, J., Samulski, R.J., and Thakker D.R.: A Novel Method For Site-Specific Dual Isotope (3H/14C) Labeling Of Plasmid DNA. American Society of Gene Therapy 4th Annual Meeting, Seattle, WA, 2001.
82. Troutman M.D. and Thakker D.R.: Efflux Ratio does not predict the role of P-glycoprotein (P-gp) in attenuating the intestinal absorption of its substrates. *PharmSci*. 2001.
83. Troutman M.D. and Thakker D.R.: P-glycoprotein (P-gp) is not responsible for the inability of rhodamine 123 to enter Caco-2 cells across apical membrane during absorptive transport. *PharmSci*. 2001.
84. Hong, S. and Thakker, D.R.: A fluorescent derivative of famotidine provides evidence that absorptive transport of famotidine and ranitidine across Caco-2 cell monolayers occurs primarily via paracellular route. *PharmSci*. 2001.
85. Bourdet, D. L. and Thakker, D. R., Evidence For Carrier-Mediated Apical Uptake of Ranitidine in Caco-2 Cells, AAPS, 2003.
86. Hong, S., Balimane, P., Chong, S., Morrison, R. A., and Thakker, D. R.: Use of Saturable Transport Mechanism of Ranitidine across Caco-2 Cell Monolayers to Screen Hydrophilic Compounds with Good Oral Absorption *In Vivo*, AAPS, 2003.
87. Hong, S. and Thakker, D. R.: Evaluation of Paracellular and Transcellular Contribution to Absorptive Transport of Ranitidine and Famotidine across Caco-2 Cell Monolayers, AAPS, 2003.
88. Klein, R. R., Yanni, S., Hong, S., and Thakker, D. R.: Role of Phospholipase C in the Regulation of Tight Junction Permeability across Caco-2 Cell Monolayers, AAPS, 2003.
89. Tippin, T. K., Morgan, D. G., and Thakker, D. R.: GI Contents but Not Serum Impact Cell Association and Efficacy of Amphiphilic Paracellular Permeability Enhancers, AAPS, 2003.
90. Ansele, J. H. and Thakker, D. R.: Development of a High-Throughput Assay for Measuring the Metabolic Stability of Compounds Towards Cytochrome P450 Oxidative Metabolism, ISSX, 2003.
91. Bourdet<sup>1</sup>, D.L., Pritchard, J.B., and Thakker, D.R.: Interaction Of H<sub>2</sub>-Antagonists With Human Organic Cation Transporters Hoct1 (Slc22a1), Hoct2 (Slc22a2), And Hoct3 (SLC22A3), AAPS, 2004.
92. Bourdet, D.L., Hong, S., and Thakker, D.R.: Absorptive Transport And Apical Uptake Of Quaternary Ammonium Organic Cations In Caco-2 Cells: Differential Uptake And Efflux Mechanisms For Mpp<sup>+</sup> And Tea, AAPS, 2004.



93. Bourdet, D.L., Hong, S and Thakker, D.R.: Absorptive Transport And Uptake Of Quaternary Ammonium Organic Cations In Caco-2 Cells: Differential Uptake And Efflux Mechanisms For Mpp<sup>+</sup> And Tea, AAPS, 2004
94. Klein, R.R., Hong, S., Zhao, B, and Thakker, D.R.: Role For Phospholipase C-Beta In The Regulation Of Tight Junction Permeability Across Caco-2 Cells, AAPS, 2004.
95. Hong, S., and Thakker, D.R.: A Nove Method to Assess Paracellular Vs. Transcellular Contribution To Absorptive Transport Across Caco-2 Cell Monolayers, AAPS, 2004.
96. Tippin, T.K., and Thakker, D. R.: Modeling GI Motility And Its Impact On Efficacy And Flux Of Paracellular Permeability Enhancers In Caco-2 Cell Monolayers, AAPS, 2004.
97. Ansede, J.H., Brouwer, K.R., and Thakker, D.R.: Development of A High Throughput Reactive Oxygen Species-Based Cytochrome P450 3a4 Inhibition Assay (ACCURATE™-IA), ISSX, 2004.
98. Bourdet, D.L., Pritchard, J.B., and Thakker, D.R.: Interaction of H<sub>2</sub>-Antagonists With Human Organic Cation Transporters hOCIT1 (Slc22a1), hOCT2 (Slc22a2), And hOCT3 (SLC22A3), ISSX, 2004.
99. Bourdet, D.L., Ming, X. and Thakker, D.R.: Do Organic Cation Transporters (Slc22a1-5) Have A Role In The Intestinal Absorption Of Hydrophilic Cationic Drugs?, AAPS Workshop On Drug Transporters – From Bench To Bedside, 2005.
100. Boudet, D. L. and Thakker, D. R., Intestinal Transport of Hydrophilic Cations: A Kinetic Modeling Approach to Elucidate the Role of Uptake and Efflux Transporters and Paracellular vs. Transcellular Transport in the Absorptive Transport of Ranitidine, AAPS, 2005.
101. Bourdet, D. L. and Thakker, D. R., Intestinal Transport of Hydrophilic Cations: A Kinetic Modeling Approach to Elucidate the Role of Uptake and Efflux Transporters and Paracellular vs. Transcellular Transport in the Absorptive Transport of Ranitidine, AAPS, 2005.
102. Klein R, Boudon, D., and Thakker, D. R. Paracellular Permeability Enhancers U73122 and Alkylphosphocholines Inhibit Phospholipase C □ Isozymes in A Cell-Free System, AAPS, 2005.
103. Tippin T. K., Cummings, C. A., and Thakker, D. R. A Cell Culture Model to Evaluate Local and Systemic Toxicity of Intestinal Paracellular Permeability, AAPS, 2005.
104. Ming, X., Hong, S., and Thakker, D. R. Evidence for Cell Membrane Damage by EGTA During Ca<sup>+2</sup>-switch Experiment in Caco-2 Cells to assess Paracellular Transport of Compounds, AAPS, 2005.
105. Ming, X., Bourdet, D. L., and Thakker, D. R. Gene Expression Profile of Human Organic Cation Transporters along the Gastrointestinal Tract and in Caco-2 Cells, AAPS, 2005.
106. Ming, X., and Thakker, D. R. Expression and Functional Activity of the Heteromeric Organic Solute Transporter alpha-beta in Caco-2 Cells, AAPS, 2006
107. Proctor, W., Bourdet, D. L., and Thakker, D. R. A Kinetic Modeling Approach to Determine the Role of Uptake Transporters and Relative Contribution of Paracellular vs. Transcellular Transport in the Absorptive Transport of Metformin in Caco-2 Cells, AAPS, 2006.
108. Mowrey, B. and Thakker, D.R. CYP3A-mediated Metabolism of Terfenadine During Absorptive Transport across Intestinal Tissue from P-gp-deficient and P-gp-competent Mice., AAPS , 2006

109. Ming, X., Knight, B., and Thakker, D. R. Involvement of Multidrug Resistance-associated Proteins (MRPs) in Intestinal Transport of the Anti-allergic Drug Fexofenadine, AAPS Transporter Meeting, 2007.
110. Proctor, W., and Thakker, D. R. Saturable Absorptive Transport of Metformin across Caco-2 Cell Monolayers Occurs Predominantly via a Novel Paracellular Transport Mechanism, AAPS Transporter Meeting, 2007
111. Knight, B., and Thakker, D. R. Transport/Metabolism Interplay for Loperamide and Terfenadine in the CYP3A-Expressing Caco-2 Cell Model, AAPS Transporter Meeting, 2007.
112. Klein, R. and Thakker, D. R. A role for phospholipase C $\beta$  in the regulation of paracellular permeability in a human intestinal epithelial cell line. *FASEB J.* 21:729.8 2007.
113. Klein, R.R., Bourdon, D.M., Wagner, C.D., White, W.L., Williams, J.D., and Thakker, D.R. Direct activation of human phospholipase C $\beta$ 3 (hPLC $\beta$ 3) by U73122 in dodecylmaltoside (DDM) mixed micelles via alkylation at cysteine residues. *FASEB J.* 21:729.9 2007.
114. Yanni, S.B., Annaert, P.P., Augustijns, P, Benjamin Jr. D.K., and Thakker, D.R. Higher Clearance of the Antifungal Agent Voriconazole in Children Compared to Adults: Potential Role of FMO and CYP2C19. ISSX, 2008.
115. Proctor, W.R. and Thakker, D.R. Mechanisms Underlying Saturable Intestinal Absorption of Metformin. Podium Session, Globalization of Pharmaceutics Education Network (GPEN) Biannual Meeting, Leuven, Belgium. September 2008.
116. Proctor, W.R. and Thakker, D.R. The Mechanisms of Saturable Paracellular Transport of Hydrophilic Cations across Epithelial Cell Monolayers. AAPS, 2008
117. Ming X. and Thakker, D.R. MRP4 Mediates Basolateral Efflux of Adefovir Formed in Caco-2 Cells from its Prodrug - Adefovir Dipivoxil. AAPS, 2008.
118. Ming X. and Thakker, D.R. Vectorial Transport of Fexofenadine across Caco-2 Cells: Involvement of Apical Uptake and Basolateral Efflux Transporters. AAPS, 2008.
119. Yanni SB, Benjamin DK, Augustijns P, Thakker DR, and Annaert PP. *In vitro* investigation of the hepatobiliary disposition mechanism of the antifungal agent micafungin in humans and rats. Drug Metabolism and Disposition, ISSX meeting, Baltimore MD, 2009
120. Proctor, W.R., Everett, R.S., and Thakker, D.R. A Novel Mechanism for Intestinal Absorption of the Type II Diabetes Drug Metformin: Potential Role in Metformin Pharmacology and Adverse Effects. Podium Presentation, Asian Symposium on ADMET Profiling in Drug Discovery, Singapore, September 2009.
121. Claudia N. Generaux, Scott J. Brantley, Richard R. Tidwell, James E. Hall, Mary F. Paine, Dhiren. R. Thakker. *Trypanosomal Infection Alters the Pharmacokinetics of Furamidine and its Prodrug Pafuramidine in Rats.* AAPS 2009
122. Dufek MB, Knight BM, Thakker DR. Effect of P-glycoprotein (P-gp) on CYP3A mediated metabolism of dual substrates during absorptive transport across stripped intestinal epithelium from P-gp competent and P-gp deficient mice. American Association of Pharmaceutical Scientists (AAPS), Los Angeles, CA. November 2009.
123. Proctor, W.R., Van Itallie, C.M., Holmes, J., Anderson, J.M., and Thakker, D.R. Paracellular Transport of Organic Cations is Facilitated by Claudin-2. Poster Session, American Association of Pharmaceutical Sciences (AAPS) Annual Meeting, Los Angeles CA, November 2009.

124. Proctor, W.R., Van Itallie, C.M., Holmes, J., Anderson, J.M., and Thakker, D.R. Vitamin D<sub>3</sub> Treatment Increases Absorptive Transport of Metformin Across Caco-2 Cell Monolayers. Poster Session, American Association of Pharmaceutical Sciences (AAPS) Annual Meeting, Los Angeles CA, November 2009.
125. Proctor, W. R., Han, T., Van Itallie, C. M., Anderson, J. M., Thakker, D. R. Saturable Paracellular Transport of Hydrophilic Organic Cations is Facilitated by Tight-Junction Protein Family, Claudins: Implications on Metformin Intestinal Absorption. AAPS 2010
126. Costales C.L., Proctor W.R, Dufek M.B., Everett R.S, Thakker D.R. Apically Localized Mouse Intestinal Cation-selective Transporters Play a Role in the Oral Absorption and Pharmacology of Metformin. Globalization of Pharmaceutics Education Network: Eighth Meeting, Chapel Hill, NC, November 2010.
127. Dufek M.B., Bridges A, Zhao L, Knight BM, Thakker DR. P-glycoprotein (P-gp) Enhances Intestinal Bioavailability of Loperamide in Mouse by Reducing First-pass Intestinal Metabolism. AAPS, New Orleans, LA. November 2010.
128. Dufek M.B., Zhao L, Thakker DR. The Effect of P-glycoprotein (P-gp) on Metabolism of Loperamide in Human and Mouse Intestinal Epithelium. AAPS, New Orleans, LA. November 2010.
129. Dufek M.B., Bridges, A, Zhao L, Knight B, Thakker DR. P-glycoprotein (P-gp) Enhances the Portal Bioavailability of Loperamide in Mouse by Reducing First-pass Intestinal Metabolism. Global Pharmaceutical Education Network (GPEN) Biannual Conference, Chapel Hill, NC. November 2010.
130. Dufek M.B., Knight BM, Bridges AS, Overby DW, Farrel TM and Thakker DR. Can P-glycoprotein (P-gp) Increase Oral Bioavailability of Dual P-gp and Cytochrome P450 3A (CYP3A) Substrates in Human? AAPS Workshop on Drug Transporters in ADME: From the Bench to the Bedside. Bethesda, MD. March 2011.
131. Costales C.L., Proctor W.R., Dufek M.B., Everett R.S., Overby D, Farrell T, Thakker D.R. Metformin Absorption is mediated by Apical Cation-selective Transporters in Mouse and Human Intestinal Tissue. AAPS Workshop on Drug Transporters in ADME: From the Bench to the Bedside, Bethesda, MD, March 2011.
132. Everett, R., Han, T., Proctor, W.R., Ng, C., Costales, C., and Thakker D.R. The Apical Localization of the Organic Cation Transporter 1 (OCT1) in the Caco-2 cell Monolayer. AAPS Workshop on Drug Transporters, Bethesda, MD, March 2011.
133. Han, T., Proctor, W.R., Costales, C., Everett, R., and Thakker, D.R. The Role of the Organic Cation Transporter 1 (OCT1) and Plasma Membrane Monoamine Transporter (PMAT) in Metformin Apical Uptake in Caco-2 cells. AAPS Workshop on Drug Transporters, Bethesda, MD, March 2011.
134. Alluri, R.V., Dufek, M.B., Proctor, W.R., Costales, C.L., Everett, R.S., Thakker, D.R. The Role of Intestinal Organic Cation Transporter 1 (OCT1) in Influencing the Intestinal Absorption (Portal Concentrations) and Overall Pharmacokinetics of Metformin in Male Diabetic db/db Mice. AAPS, Washington D.C., October 2011
135. Costales C.L., Proctor W.R., Thakker, D.R. Identification and Characterization of Mouse Cation-Selective Transporters Involved in the Intestinal Absorption of Metformin. AAPS, Washington D.C., October 2011
136. Han, T., Thakker, D.R. The Interaction between Serotonin Reuptake Transporter (SERT) and Metformin: Mediating the Mechanisms of Absorption and Adverse Effects of Metformin. AAPS, Washington D.C., October 2011
137. Zane, N., Thakker D.R. *In Vitro* Data Suggest Decreased Metabolic Clearance of Voriconazole in Pediatric CYP2C19-Poor Metabolizers with Normal CYP3A4 Activity. AAPS, Washington D.C., October 2011
138. Zhang, Y., Everett, R.S., Thakker, D.R. Are MCF-7 Cells a Relevant *In Vitro* Model for Evaluating Metformin Anticancer Efficacy in Breast Cancer? AAPS, Washington D.C., October 2011

141. Han, T.H., Everett, R.S., Proctor, W.R., Ng, C.M., Costales, C.L., and Thakker, D.R., Intestinal Cation-Selective Transporters Involved in the Apical Uptake, Accumulation and Absorption of Metformin. Globalization of Pharmaceutics Education Network (GPEN) Meeting, Australia, 2012
142. Han, T.H., Proctor, W.R., Costales, C.L., Everett, R.S., and Thakker, D.R., Cation-Selective Transporters Involved in the Apical Uptake and Accumulation of Metformin in Caco-2 Cell Monolayers. AAPS– Chicago, IL, 2012
143. Han, T.H., Everett, R.S., Proctor, W.R., Ng, C.M., Costales, C.L., and Thakker, D.R., Apical Localization of the Organic Cation Transporter (OCT) 1 in Caco-2 Cell Monolayers, and Human and Mouse Intestine. AAPS– Chicago, IL, 2012
144. Romo-Fewell, O., Han, T.H., Alluri R.V., and Thakker, D.R., Evidence for “Metformin-Like” Cation Transporter-Assisted Paracellular Absorptive Transport of the Dietary Hydrophilic Organic Cation Choline across Caco-2 Cell Monolayers. AAPS– Chicago, IL, 2012
145. Costales C, Alluri R, and Thakker D.R., Metformin Intestinal Absorption and Accumulation in Mouse is Inhibited by Tricyclic Anti-depressant Desipramine: Insights into a Novel Absorption Mechanism of Metformin. AAPS– Chicago, IL, 2012
146. Alluri R, Costales C, and Thakker D. Interactions of Commonly Used Anesthetics on Intestinal Absorption and Systemic Disposition of Metformin. AAPS– Chicago, IL, 2012.
147. Alluri R, Costales C, Everett R, Asokan A and Thakker D. Development of an Adeno-Associated Virus (AAV) Vector-mediated Transporter Knockdown Mouse Model to Study the Contribution of Mouse Organic Cation Transporter 1 (mOct1) in the Intestinal Absorption of Metformin. AAPS– Chicago, IL, 2012
148. Han T, Proctor W, Costales C, Everett R, and Thakker D. Cation-Selective Transporters Involved in the Apical Uptake and Accumulation of Metformin in Caco-2 Cell Monolayers. AAPS– Chicago, IL, 2012
149. Han T, Everett R, Proctor W, Ng C, Costales C, and Thakker D. Organic Cation Transporter 1 (OCT1/Oct1) is Localized to the Apical Membrane in Caco-2 Cell Monolayers and Intestinal Epithelium of Mouse and Human. AAPS– Chicago, IL, October 2012
150. Costales C, Alluri R, and Thakker D. Insights into a Novel Absorption Mechanism of the Antidiabetic Drug Metformin. Globalization of Pharmaceutics Education Network: Ninth Meeting – Melbourne, AU, November 2012.
151. Costales C, Alluri R, and Thakker D. Apical Uptake and Efflux Transporters Enhance the Intestinal Absorption of Metformin in Mouse. AAPS Workshop on Drug Transporters in ADME: From the Bench to the Bedside – Bethesda, MD, 2013.
152. Mackowiak B, Costales C, Alluri R, Han T, Everett R, and Thakker D. Apical Membrane Transporters in Human and Mouse Intestinal Epithelia Mediate Metformin Efflux. AAPS Workshop on Drug Transporters in ADME: From the Bench to the Bedside – Bethesda, MD, 2013.
153. Han T, Everett R, Proctor W, Ng C, Costales C, Brouwer K, and Thakker D. Organic Cation Transporter 1 (OCT1/Oct1) is Localized in the Apical Membrane of Caco-2 Cell Monolayers and Enterocytes. AAPS Workshop on Drug Transporters in ADME: From the Bench to the Bedside – Bethesda, MD, 2013.
154. Han, T.H., Everett, R.S., Proctor, W.R., Ng, C.M., Costales, C.L., and Thakker, D.R., Apical Localization of the Organic Cation Transporter (OCT) 1 in Caco-2 Cell Monolayers, and Human and Mouse Intestine. AAPS Workshop on Drug Transporters in ADME: From the Bench to the Bedside – Bethesda, MD, 2013.

155. Han, T.H., Everett, R.S., and Thakker, D.R., The Interaction Between Serotonin Reuptake Transporter (SERT) and Metformin: A Novel Mechanism of Oral Absorption and Intestinal Adverse Effects of Metformin. AAPS Workshop on Drug Transporters in ADME: From the Bench to the Bedside – Bethesda, MD, 2013.
156. Everett, R.S., Zhang, Y, Onoliwu, I., Bae-Jump, V.L., and Thakker, D.R. Multiple Cation-selective Transporters Contribute to the Anti-proliferative Effect of Metformin in Ovarian Cancer Cell Lines, AACR, Washington, D.C., 2013.
157. Damery, E.F., Zhang, Y., Ononiwu, I.M., Everett, R.S., Bae-Jump, V.L., and Thakker, D.R. Cation-selective Transporters are Critical to the Anti-cancer Efficacy of Metformin in Endometrial Cancer. The Society of Gynecologic Oncology 44th Annual Meeting on Women's Cancer, Los Angeles, CA, 2013.
158. Zane, N.R. and Thakker D.R. A Physiologically Based Pharmacokinetic Model Incorporating Extrahepatic Metabolism Explains Voriconazole Pediatric Bioavailability Differences. The American Society for Pharmacology and Experimental Therapeutics (ASPET), Boston, MA, 2013.
159. Zane NR and Thakker DR. Age-Dependent Differential Gene Expression Explains Higher Voriconazole Clearance in Children. American Association of Pharmaceutical Sciences (AAPS), San Antonio, Tx. November 2013
160. Cai, H., Wahajuddin, M., Alluri, R.V., Everett R.S., and Thakker, D.R. Cation-selective Transporters are Critical Determinants of the Anti-proliferative Effects of Metformin in Breast Cancer Cells. AAPS Annual Meeting, San Antonio, TX, 2013.
161. Zane NR and Thakker DR. Integrating In Vitro Sildenafil Metabolism into a Physiologically-Based Pharmacokinetic (PBPK) Model: Validating a Bottom-Up Modeling Approach. American Association of Pharmaceutical Sciences (AAPS), San Diego, CA. November 2014
162. Zane NR and Thakker DR. In Vitro Techniques to Predict Intestinal First-pass Metabolism in Children and Adults. American Association of Pharmaceutical Sciences (AAPS), San Diego, CA. November 2014
163. Zane NR, Chen Y, Wang M, and Thakker DR. Higher CYP2C19 Functional Activity in Children is Not Entirely Explained by Higher Gene or Protein Expression. International Society for the Study of Xenobiotics (ISSX), San Francisco, CA. October 2014
164. Zane NR, Chen Y, Wang M, and Thakker DR. Evidence for a New Testosterone Metabolite Uniquely formed by CYP3A7: A Potential Probe for CYP 3A7-mediated Metabolism in the Newborn. International Society for the Study of Xenobiotics (ISSX), San Francisco, CA. October 2014
165. Cai, H., Wahajuddin, M., Everett R.S., and Thakker, D.R. Cation-Selective Transporters Enhance the Antitumor Efficacy of Metformin in Xenograft Mouse Models of Breast Cancer. AAPS Annual Meeting, San Diego, CA, 2014.
166. Cai, H., Wahajuddin, M., Everett R.S., and Thakker, D.R. Do cation-selective transporters help or hurt the antitumor efficacy of metformin in breast cancer? AACR Annual Meeting, San Diego, CA, 2014.
167. Thakker, A.R., and Thakker, D.R. Lower Pediatric Oral Bioavailability of Voriconazole is Not Due to Lower Intestinal Bile Salt Concentration in Children. AAPS Annual Meeting, San Diego, CA, 2014.
168. Thakker, A.R., Alluri, R.V., Everett, R.S., and Thakker, D.R. Adeno-associated Viral Vector-mediated shRNA Strategies to Generate Intestinal and Hepatic Metformin Transporters Knockdown Mouse Models. AAPS Annual Meeting, San Diego, CA, 2014.
169. Xia, Y., Thakker, A.R., and Thakker, D.R. A Novel Mechanism for Glucose Lowering Effect of Metformin Involving the Intestine. AAPS Annual Meeting, San Diego, CA, 2014.

170. Hao Cai, Ruth S. Everett, and Dhiren R. Thakker. Cation-Selective Transporters Are Central to the Antitumor Efficacy of Metformin against Breast Cancer. American Association of Pharmaceutical Scientists (AAPS) Annual Meeting and Exposition, Orlando, FL, 2015.
171. Hao Cai, Ruth S. Everett, and Dhiren R. Thakker. Interaction of the Insulin-Dependent and AMPK-Dependent Pathways in Human Breast Cancer Cells Improves the Antiproliferative Efficacy of Metformin. AAPS Annual Meeting and Exposition, Orlando, FL, 2015.
172. Lee, C.M., Zane, N.R., Thakker, D.R. Use of Physiologically-Based Pharmacokinetic Model to Elucidate the Role of Metabolism and Transport in Vincristine Disposition. AAPS Annual Meeting and Exposition; Orlando, FL, 2015
173. Lee, C.M., Thakkar, A., Ward, P.D., Sepassi, K., Thakker, D.R. Use of Human Intestinal Tissue and Modeling to Replace Preclinical Species in the Evaluation of Oral Formulations. AAPS Annual Meeting and Exposition; Orlando, FL. 2015
174. Arti R. Thakkar, Ravindra V Alluri, Chester Costales, Garrett Berry, Aravind Asokan, Ruth S. Everett and Dhiren R. Thakker. Strategies to Generate Intestinal and Hepatic Metformin Transporter Knockdown Mouse Models: Delivery of Transporter-specific Multiple microRNA by Adeno-associated Viral Vector. 7th International Symposium on DMPK, NIPER, Mohali, India, 2015.
175. Staley AS, Roque DR, Schuler KM, Rambally BS, Sampey B, Everett R, Thakker D, Gehrig PA, O'Connor S, Makowski L, Bae-Jump VL. Molecular and metabolic differences of treatment responders versus non-responders in a phase 0 clinical trial of metformin in endometrial cancer. Society of Gynecologic Oncology 47th Annual Meeting on Women's Cancer, San Diego, CA, 2016.
176. Emily Meichun Ko, Stephanie Sullivan, Brooke Rambally, Siobhan O'Connor, Ruth Everett, Dhiren Thakker, Dominic T. Moore, John Byron, Victoria Lin BaeJump. Metformin for the Treatment of Endometrial Hyperplasia. American Society of Clinical Oncology Annual Meeting, Chicago, IL 2016.
179. Christine M Lee, Nicols Zane, and Dhiren R Thakker. A Physiologically-Based Pharmacokinetic Model Suggests that Intracellular Binding, in Addition to Metabolism and Efflux Transport, Defines the Disposition of Vincristine in Adult and Pediatric Populations. AAPS Annual Meeting and Exposition, Denver, CO, 2016
180. Christine M Lee, Hao Cai, and Dhiren R Thakker. Elucidation of the Intestinal Absorption Mechanism of Amoxicillin Using the Caco-2 Cell Monolayer Model and Human Intestinal Tissue. AAPS Annual Meeting and Exposition, Denver, CO, 2016.
181. Hao Cai, Ruth S Everett, and Dhiren R Thakker. Higher Sensitivity of Human Breast Cancer Stem Cells Compared to Non-stem Cancer Cells to Antiproliferative Effect of Metformin Results from Higher Metformin Transporter Expression. AAPS Annual Meeting and Exposition, Denver, CO, 2016.
182. Hao Cai, Ruth S Everett, and Dhiren R Thakker. A Higher Dose of Metformin is Necessary for Treatment of Breast Cancer than the Doses Used for Treatment of Diabetes. AAPS Annual Meeting and Exposition, Denver, CO, 2016.
183. Hao Cai, Ruth S Everett, and Dhiren R Thakker. Inhibition of the Insulin Pathway Sensitizes Breast Tumors to the Anticancer Efficacy of Metformin. AAPS Annual Meeting and Exposition, Denver, CO, 2016.
184. Ben M. Clements, Ruth S Everett, and Dhiren R Thakker. The Critical Nutrient Choline is Absorbed across the Intestinal Epithelium by a Mechanism Similar to the Enterocyte-luminal Cycling Absorptive Transport of Metformin. AAPS Annual Meeting and Exposition, Denver, CO, 2016.
185. Derek L. Bolhuis, Erin F. Damery, Innocent M. Onoiwu, Aakash Patel, Hao Cai, Christine M. Lee, Ruth S. Everett, Victoria L. Bae-Jump, Dhiren R. Thakker. Cation-Selective Transporters are Central to AMPK-mediated Anti-

proliferative Efficacy of Metformin in Endometrial Cancer Cells. AAPS Annual Meeting and Exposition, Denver, CO, 2016.

**186.** Lawrence C. Ku, P. Brian Smith, Michael Cohen-Wolkowicz, and Dhiren R. Thakker. Interaction between Formula Feedings and Caffeine Pharmacokinetics in Premature Infants. AAPS Annual Meeting and Exposition, Denver, CO, 2016.

**187.** Dhiren R. Thakker. Strategic Need and Optimal Use of Preclinical Determinants of ADME Properties In Conjunctions with Toxicological Endpoints. AAPS Annual Meeting and Exposition, Denver, CO, 2016.

## INVITED LECTURES

1. Clinical Society of Washington, Rockville, MD., 1976.
2. Department of Pharmacology, George Washington University, Washington, D.C., 1977.
3. Department of Chemistry, George Washington University, Washington, D.C., 1977.
4. Syntex Pharmaceutical Co., Palo Alto, CA., 1978.
5. Departments of Medicinal Chemistry and Biochemistry, University of Kansas, Lawrence, KS, 1978.
6. Environmental Health Chemistry Symposium, American Chemical Society, Washington, DC., 1979.
7. 39th International Congress of Pharmaceutical Sciences, Brighton, U.K., 1979.
8. Department of Chemical Technology, Bombay University, Bombay, India, 1980.
9. Cancer Research Institute, Bombay, India, 1980.
10. Bristol Myers Laboratories, Syracuse, NY., 1980.
11. Laboratory of Developmental Pharmacology, National Institute of Child Health and Human Development, NIH, Bethesda, MD., 1981.
12. Laboratory of Biochemistry and Metabolism, NIADDK, NIH, Bethesda, MD., 1981.
13. International Symposium on Biological Reactive Intermediates, Gilford, Surrey, U.K., 1981.
14. Interx-Merck Pharmaceutical Co., Lawrence, KS., 1982.
15. Smith Kline Beckman, Philadelphia, PA., 1982.
16. Indian Pharmaceutical Association, Bombay, India, 1983.
17. Cancer Research Institute, Bombay, India, 1983.
18. National Center for Drugs and Biologics, FDA, Bethesda, MD., 1983.
19. Department of Pharmaceutical Chemistry, University of Kansas, Lawrence, KS., 1984.
20. Smith Kline Beckman, Philadelphia, PA., 1985.
21. Cancer Research Institute, Bombay, India, 1986.
22. Central Drug Research Institute, Lucknow, India, 1986.
23. Bhabha Atomic Research Center, Bombay, India, 1986.
24. Hoechst Pharmaceuticals, Bombay, India, 1986.
25. 6th International Catecholamine Symposium, Jerusalem, Israel, 1987.
26. SATO Memorial International Award Lecture at the 107th Annual Meeting of the Pharmaceutical Society of Japan,



Kyoto, Japan, 1987.

27. Nagoya City University, Faculty of Pharmaceutical Sciences, Nagoya, Japan, 1987.
28. Tohoku University, Research Institute for Tuberculosis and Cancer Sendai, Japan, 1987.
29. Keio University School of Medicine Tokyo, Japan, 1987.
30. Tokyo College of Pharmacy, Tokyo, Japan, 1987.
31. National Institute of Hygienic Sciences, Tokyo, Japan, 1987.
32. Department of Chemical Technology, Bombay University, Bombay, India, 1987.
33. Laboratory of Bioorganic Chemistry, NIDDK, Bethesda, MD., 1987.
34. Carcinogen Risk Assessment, Banbury Center, Cold Spring Harbar, 1987.
35. Pharmaceutical Institute, School of Medicine, Keio University, Tokyo, Japan, 1988.
36. Tokyo College of Pharmacy, Tokyo, Japan, 1988.
37. Dept. of Pharmaceutical Sciences, Nagoya City University, Nagoya, Japan, 1988
38. Dept. of Medicinal Chemistry, University of North Carolina, 1989.
39. Dept. of Medicinal Chemistry, University of North Carolina, 1990 (presented 3 lectures in a Course on Drug Metabolism).
40. Dept. of Medicinal Chemistry, University of North Carolina, 1991 (presented 3 lectures in a Course on Drug Metabolism).
41. Dept. of Pharmacology, University of North Carolina, 1991.
42. Annual Meeting of the PMA Drug Metabolism Section, New Orleans, 1991.
43. 25th Annual Higuchi Symposium, Lake of the Ozarks, 1992.
44. Enz Lecture, Dept. of Pharmaceutical Chem., University of Kansas, 1992
45. Dept. of Medicinal Chemistry, University of North Carolina, 1992 (presented 3 lectures in a course on Drug Metabolism).
46. Dept. of Medicinal Chemistry, University of North Carolina, 1993 (presented 3 lectures in a course on Drug Metabolism).
47. Dept. of Chemistry, Catholic University, Washington, D.C., 1993.
48. Lab of Bioorganic Chemistry, NIDDK, NIH, 1993.
49. Chaired a session on the "Integration of Preclinical ADME studies in the Preclinical and Clinical Safety Assessment" at the National Meeting of Drug Information Association, at Washington, D.C., 1994.
50. Chaired a session on "Delivery and Disposition of Peptides and Oligonucleotides- Current Status and Future

Challenges” at the annual meeting of the International Society for the Study of Xenobiotics (ISSX), Raleigh, NC, October 1994.

51. Chaired a Special Symposium on “Drug Delivery and Prodrug Technologies” at the 31st ACS Western Regional Meeting, San Diego, CA, October 1995.
52. Conference on “Lead Generation and Optimization”, Princeton, NJ, September, 1996.
53. Organized a short course presented a talk on Prodrugs at the 7th North American ISSX Meeting, San Diego, October 1996.
54. Winter Conference on “Medicinal and Bioorganic Chemistry”, Steamboat Springs, Colorado, January, 1997.
55. Third International Symposium on Innovations in Pharmaceutical Sciences and Technology”, Ahmadabad, India, February 1997.
56. S.N.D.T. College of Pharmacy, Bombay University, Bombay, India, February, 1997.
57. Tata Institute for Cancer Research, Bombay, India, February, 1997.
58. Amylin Pharmaceuticals, San Diego, CA, April, 1997
59. Merck Research, West Point, PA, April, 1997.
60. Proctor & Gamble, Cincinnati, OH, June, 1997.
61. AAPS Southeast Regional Meeting, Research Triangle Park, NC, June, 1997.
62. Drew University, (faculty), Princeton, NJ, July, 1997.
63. DuPont Merck, Newark, DE, July 1997.
64. Formulations and Drug Delivery, ACS Conference, San Diego, CA, October 1997.
65. Glaxo Wellcome, Inc. Pharmaceuticals Division, RTP, NC, January, 1998.
66. Swiss Chemical Society – Minisymposium on Oral Drug Delivery, Basel Switzerland, May, 1998.
67. University of Leuven, Workshop on Industry – Academic Collaborations, Leuven, Belgium, May 1998.
68. Glaxo Wellcome, Inc., Biomet Division, Ware, U.K., May, 1998.
69. AAPS Western Regional Meeting, San Francisco, CA, May, 1998.
70. AAPS Southeastern Regional Meeting, RTP, NC, June, 1998.
71. 28<sup>th</sup> Annual Gordon Research Conference on Drug Metabolism, Plymouth, NH, July, 1998.
72. Drew University, (faculty), Princeton, NJ, July, 1998.
73. Pfizer, Inc., Groton, CT, July, 1998.
74. Bristol-Meyer Squibb, Inc., Wallingford, CT, July, 1998.

75. School of Pharmacy, University of Michigan, Ann Arbor, MI, September, 1998.
76. Higuchi Research Seminar, Lake of Ozarks, MO, March, 1999.
77. Glaxo Wellcome, Pharmaceuticals Dept., RTP, NC, March, 1999.
78. BASF, Boston, MA, June, 1999.
79. Parke Davis, Inc., Ann Arbor, MI, July, 1999.
80. Novartis, East Hanover, NJ, September, 1999.
81. Institute for Innovative Research, (faculty), San Diego, CA, December, 1999.
82. Agouron Pharmaceuticals, San Diego, CA, December 1999.
83. Glaxo Wellcome (Pharmaceutics), RTP, NC, March 2000.
84. Laboratory of Biorganic Chemistry, NIDDK, NIH, Bethesda, MD, May 2000
85. Pratt Fellowship Program, NIH, Bethesda, MD, May 2000
86. Wyeth-Ayerst, Pearl River, N.Y., June 2000
87. Drew University (faculty), Princeton, NJ, June 2000
88. Chiron, CA, 2001.
89. Roche, Palo Alto, CA, 2001
90. Guilford Pharmaceuticals, Baltimore, MD, 2001
91. Drew University (faculty), Princeton, NJ, 2001
92. Drew University (faculty), Course Organizer, Princeton, NJ, 2001
93. Transform Pharmaceuticals, Boston, MA, 2001
94. Bristol-Myers Squibb, Princeton, NJ, 2001
95. Albany Biolomolecules, Albany, NY, 2001
96. Lilly Pharmaceuticals, Indianapolis, IN, 2001
97. 3-D Pharmaceuticals, Philadelphia, PA, 2001
98. Pfizer, Ann Arbor, MI, 2002
99. Gilford Pharmaceuticals, Baltimore, MD, 2002
100. Pfizer (Agouron), San Diego, CA, 2002
101. Soreno Pharmaceuticals, Soreno, Switzerland, 2002

102. Avantis, Frankfurt, Germany, 2002
103. Bristol Myers Squibb, Lawrenceville, NJ, 2002
104. Avantis, Pearl River, NJ, 2002
105. Roche, Nutley, NJ, 2002
106. Pfizer, Groton, CT, 2002
107. GPEN, Detroit, MI, 2002
108. GlaxoSmithKline, Philadelphia, PA, 2002
109. Arena Pharmaceuticals, San Diego, CA, 2002
110. Drew University, Newark, NJ, 2003
111. Novartis, Jolla CA, 2003
112. AstraZeneca, Boston, MA, 2003
113. Fibrogen, San Francisco, CA 2003
114. ADMET-1, San Diego, CA 2004
115. AstraZeneca, San Diego, CA 2004
116. LabFusion, Boston, Ma 2004
117. Millennium, Cambridge, MA 2004
118. Schering-Plough, Kenilworth, NJ 2004
119. Neurocrine, San Diego, CA 2004
120. Abbott, Abbott Park, IL 2004
121. Pfizer, Groton, CT, 2005
122. ICOS, Bothell, WA, 2005
123. SCIOS, Inc. San Francisco, CA, 2005
124. Biogen, Cambridge, MA 2005
125. Genzyme, Framingham, MA 2005
126. Sunesis, San Francisco, CA 2005
127. Drew University, Newark, NJ 2005
128. Amgen, Boston, MA, 2005

“Probing the Intestinal Epithelial Barrier: A Tortuous Journey”

129. Pediatric Pharmacology Research Unit Network Meeting, Washington, D. C., 2005  
“Implementing In Vitro Drug Disposition Studies to Improve Pediatric Therapy – Challenges and Opportunities”
130. Cambridge Healthcare Institute – Mastering Medicinal Chemistry/ADME In-depth, San Francisco, CA, 2006  
“ADME In-depth: A Medicinal Chemistry Perspective” (Lead Lecture)
131. Scitech Center, Mumbai, India, 2006  
“Integration of ADME in Drug Discovery: Past, Present, and Future”
132. Indian Drug Manufacturers’ Association Meeting – Bioavailability and Bioequivalence, Mumbai, India, 2006.  
“Integration of ADME in Drug Discovery: Past, Present, and Future”
133. PERD Center, Ahmedabad, India, 2006  
“Integration of ADME in Drug Discovery: Past, Present, and Future”  
  
“Probing the Intestinal Epithelial Barrier: A Tortuous Journey”
134. Drew University, Madison, NJ, 2006.  
“Use of Preclinical Pharmacokinetics and In Vitro Models to Assess Problems of Poor Oral Bioavailability” (with Kim Brouwer)  
  
“Role of Intestinal and Hepatic Metabolism in Drug Clearance – Chemical Strategies to Optimize Metabolic Clearance”
135. Genentech, San Francisco, CA, 2006.  
“Use of Preclinical Pharmacokinetics and In Vitro Models to Assess Problems of Poor Oral Bioavailability” (with Kim Brouwer)  
  
“Role of Intestinal and Hepatic Metabolism in Drug Clearance – Chemical Strategies to Optimize Metabolic Clearance”
136. New Jersey Drug Metabolism Discussion Group, Newark, NJ, 2006.  
Intestinal Drug Transport: A Black Box or a Target for Rational Design of Orally Active Drugs?
137. Bristol Myers Squibb, NJ, 2006.  
Interplay of Transport and Metabolism in the Intestinal Epithelium
138. Nektar Pharmaceuticals, Mobile, Alabama/San Diego, CA, 2007  
“Use of Preclinical Pharmacokinetics and In Vitro Models to Assess Problems of Poor Oral Bioavailability” (with Kim Brouwer)  
  
“Role of Intestinal and Hepatic Metabolism in Drug Clearance – Chemical Strategies to Optimize Metabolic Clearance”
139. Molecular Medicine Tri-Conference – ADMET short course, San Francisco, CA, February, 2007.  
Lead Speaker - *In Vitro* ADME Assays: Strategic Application to Drug Discovery
140. New England Drug Metabolism Discussion Group Symposium, Boston, MA, April 2007  
“Drug Metabolism-Pharmacokinetics-guided (DM/PK-guided) Lead Optimization: A Historical Perspective of a Paradigm Shift”
141. Bayer Pharmaceuticals, Dusseldorf, Germany, January 2008

- “Use of Preclinical Pharmacokinetics and In Vitro Models to Assess Problems of Poor Oral Bioavailability” (with Kim Brouwer)
- “Role of Intestinal and Hepatic Metabolism in Drug Clearance – Chemical Strategies to Optimize Metabolic Clearance”
142. Molecular Medicine Tri-Conference – ADMET short course, San Francisco, CA, March, 2008.  
Lead Speaker – “*In Vitro* ADME Assays: Strategic Application to Drug Discovery”
143. Sepracor Short Course, Boston, MA, July 2008  
“Use of Preclinical Pharmacokinetics and In Vitro Models to Assess Problems of Poor Oral Bioavailability- Part 2
144. University of Toronto, Canada, August 2008  
“Orally Active Hydrophilic Cationic Drugs – Design or Serendipity?”
145. National Institute of Pharmaceutical Education and Research (NIPER) 1<sup>st</sup> International Symposium on Metabolism and Pharmacokinetics, Chandigarh, India, 2009  
“Introduction to Drug Transporters”  
“Pharmacogenomics of Drug Transporters”
146. Wolfe Biopharma , Conference – Filling Biopharma’s Pipeline Boston, MA, April 2009  
”ADMET: Strategic Use of In Vitro and Preclinical Studies to De-risk Clinical Candidates”
147. GSK, Research Triangle Park, NC, August 2009  
“A Novel Mechanism for Intestinal Absorption of Metformin: Possible Role in Metformin Pharmacology”
148. Asian Symposium on ADMET Profiling for Drug Discovery: Metabolism and Medication Safety, September 2009  
“Role of ADME in Adverse Effects of Drugs” – Keynote Address
149. J & J, La Jolla, CA, November 2009  
“Overcoming the Intestinal Barrier: Challenges and Opportunities for Oral Drug Delivery”
150. Institute of Pharmaceutical Education and Research (NIPER) 2<sup>nd</sup> International Symposium on Metabolism and Pharmacokinetics, Chandigarh, India, 2010  
“Understanding the Role of Transporters in Efficacy and Adverse Effects of Therapeutic Agents: Past, Present, and Future”
151. Advinus Therapeutics, Poone, India, March 2010.  
“Transporters in Efficacy and Adverse Effects of Therapeutic Agents”
152. Scitech, Mumbai, Bombay, India, March, 2010.  
“Transporters: An Overview of the Role of Transporters in Drug Development”
153. Medicinal Chemistry Symposium, National University of Singapore, Singapore, September 2010: “Challenging Landscape of Drug Discovery in the 21<sup>st</sup> Century”
154. Institute of Pharmaceutical Education and Research (NIPER) 3<sup>rd</sup> International Symposium on Metabolism and Pharmacokinetics, Chandigarh, India, February, 2011: “Transporters in Drug Efficacy and Toxicity: Revisiting History and Looking to the Future”
155. Indian Institute of Technology –Bombay, India, February, 2011: “Absorption of Therapeutic Agents across the Intestinal Epithelial Barrier: A Tortuous Journey”
156. ISF College of Pharmacy, Moga, India, February, 2011: “Strategies to Design Orally Active Therapeutic Agents”

157. Ili Lilly & Co., April, 2011: “Interactions of Metformin and Loperamide with the Intestinal Epithelial Barrier: A New Look at Old Drugs”
158. New Jersey Drug Metabolism Discussion Group, October 2011: “Interplay of Drug Metabolizing Enzymes and Transporters”.
159. Bristol Myer Squibb, New Jersey, October 2011: “Interplay of Drug Metabolizing Enzymes and Transporters”.
160. American Association of Indian Pharmaceutical Scientists – Dr. R. S. Baichwal Seminar – Department of Pharmaceutical Sciences and Technology, Institute of Chemical Technology, Bombay, India, February 2012: Chair, Symposium on “ADME (DM/PK) as a Success Factor in Drug Discovery”
161. American Association of Indian Pharmaceutical Scientists – Dr. R. S. Baichwal Seminar – Department of Pharmaceutical Sciences and Technology, Institute of Chemical Technology, Bombay, India, February 2012: Symposium on “ADME (DM/PK) as a Success Factor in Drug Discovery  
“Science and Application of Absorption, Distribution, Metabolism, Excretion (ADME) in Therapy: Past, Present, and Future”
162. Dow Chemical, Mumbai, India, February 2012:  
“Activation of Phospholipase C: A Serendipitous Discovery in Search of Applications”
163. Indian Institute of Technology, Bombay, India, February 2012:  
“Transporters in Drug Efficacy and Toxicity: Past Present, and Future”
164. Institute of Pharmaceutical Education and Research (NIPER) 4<sup>th</sup> International Symposium on Drug Metabolism and Pharmacokinetics, Chandigarh, India, February 2012:  
“Intestinal Transport-Metabolism Interactions”
165. The Delaware Valley Drug Metabolism Discussion Group, Langhorne, PA, September 2012:  
“Complex Interplay of Intestinal Apical Uptake/Efflux and Basolateral Efflux Transporters in Affecting Oral Absorption of Hydrophilic Ionic Compounds: Implications for Drug-drug Interactions”
166. The joint conference of the Australasian Pharmaceutical Science Association – Australasian Society of Clinical and Experimental Pharmacologists and Toxicologists, December 2012: Role of organic cation transporters in mediating oral drug absorption”
167. Applied Pharmaceutical Analysis-India, Annual Meeting, Hyderabad, India, March, 2013:  
“Intestinal Transporters: A Gateway to the Body but Also a First Line of Defense”
168. National Institute of Pharmaceutical Education and Research (NIPER) 5<sup>th</sup> International Symposium on Drug Metabolism and Pharmacokinetics, Chandigarh, India, March 2013:  
A Keynote Address - “DMPK in the 21<sup>st</sup> Century”
169. Scitech Center, Mumbai, Bombay, India, March 2013:  
“DMPK in the 21<sup>st</sup> Century”
170. Parul Institutes - College of Pharmacy, Baroda, India, March 2013:  
“Complex Interplay of Intestinal Apical and Basolateral Transporters Affecting Oral Absorption of Hydrophilic Ionic compounds: Implications for DDI”
171. University of Maryland School of Pharmacy, November 2013:  
“The Metformin Story: How Multiple Transporters Affect Its Oral Absorption, Efficacy in Type II Diabetes and Cancer, and Adverse Effects”

172. Applied Pharmaceutical Analysis-India, Annual Meeting, Ahmedabad, India, February 2014:  
“Transporter-driven Vectorial Transport of Cationic, Anionic, and Zwitterionic Drugs in the Intestine”
173. National Institute of Pharmaceutical Education and Research (NIPER) 6<sup>th</sup> International Symposium on Drug Metabolism and Pharmacokinetics, Chandigarh, India, February28-March 2 2014:  
“The Journey of Metformin in the Intestine and Tumors: How Transporters Affect Metformin Pharmacology in Diabetes and Cancer”
174. Institute of Chemical Technology, Department of Pharmaceutical Sciences and Technology, Mumbai, India, March 2014:  
Organizer of Dr. R. S. Baichwal Seminar: “Intellectual Property in an Academic Institution: Conception to Commercialization”
175. Institute of Chemical Technology, Department of Pharmaceutical Sciences and Technology, Mumbai, India, March 2014:  
Dr. R. S. Baichwal Seminar: Intellectual Property in an Academic Institution: Conception to Commercialization:  
“Creation of Intellectual Property and Entrepreneurism: An Integral Part of Academic Pursuit in 21<sup>st</sup> Century”
176. Piramel Enterprises Ltd., Mumbai, India, March 2014:  
“How Transporters Affect Metformin Pharmacology in Diabetes and Cancer”
177. Pharmaceutical Sciences World Congress, Melbourne, Australia, April 2014:  
Debate on “Carrier Mediated Transport through Membranes: The Exception or The Rule”
178. Pharmaceutical Sciences World Congress, Melbourne, Australia, April 2014:  
“Interplay of Drug Metabolizing Enzymes and Transporters” in a Symposium entitled “Barrier Mechanisms Team Up: Interplay between Transporters, Enzymes, and Tight Junctions”
179. Nicolae Testemitanu State University of Medicine and Pharmacy of the Republic of Moldova, Chisinau, Moldova, April 2014:  
“Inside UNC Eshelman School of Pharmacy: Highlighting the Research Enterprise, Educational Outreach, and Health care Impact”
180. North Dakota State University, Fargo, North Dakota, May 2014:  
“Creation of Intellectual Property and Entrepreneurism to Translate Discoveries for Societal Benefit: An Integral Part of Academic Pursuit in 21<sup>st</sup> Century” in a Symposium entitled “Frontiers in Biomedical Research”
181. National Institute of Pharmaceutical Education and Research (NIPER) 6<sup>th</sup> International Symposium on Drug Metabolism and Pharmacokinetics, Chandigarh, India, February28-March 2 2015:  
“Transporters in Breast Cancer Cells Play a Critical Role in Anticancer Efficacy of Metformin”.
182. Applied Pharmaceutical Analysis-India, Annual Meeting, Mumbai, India, February 2015:  
“Four Cation-selective Transporters Contribute to Intestinal Absorption and Pharmacology of The Type II Diabetes Drug Metformin”.
183. JSS University, School of Pharmacy, Mysore, India, February 2015:  
“Inside UNC Eshelman School of Pharmacy: Highlighting the Research Enterprise, Educational Outreach, and Health care Impact”
184. AAPS Annual meeting (Sunrise Session), Denver, CO, November 2016  
“Strategic Need and Optimal Use of preclinical determination ADME of ADME properties in conjunction with Toxicological endpoints”