

PERSONAL DATA

Name: Nicholas Joseph Schork
Address: Department of Molecular and Experimental Medicine
The Scripps Research Institute
10550 N. Torrey Pines Rd
La Jolla, CA 92037
Mail Drop: MEM 275A
Phone: (858) 784-2308
Administrative: (858) 822-5788
Fax: (858) 822-2113
e-mail: nschork@scripps.edu; nschork@ucsd.edu
Web site: <http://psychiatry.ucsd.edu/>
<http://polymorphism.scripps.edu/>

EDUCATION

1991–1994 University of Michigan, Ann Arbor, Michigan; Ph.D. (Epidemiology)
1988–1991 University of Michigan, Ann Arbor, Michigan; M.A. (Applied Statistics)
1986–1992 University of Michigan, Ann Arbor, Michigan; M.A. (Philosophy)
1982–1985 University of Michigan, Ann Arbor, Michigan; B.A. (Philosophy)
1980–1982 Kalamazoo College, Kalamazoo, Michigan

APPOINTMENTS

Scripps Health

2007-Present Director of Research, Scripps Genomic Medicine

The Scripps Research Institute

2007-Present Professor, Molecular and Experimental Medicine
2003-2007 Professor (adjunct), Department of Molecular and Experimental Medicine

University of California at San Diego

Faculty

2007-Present Professor of Psychiatry and Biostatistics (adjunct)
2004-2007 Professor of Biostatistics, Department of Family & Preventive Medicine
2001-2007 Professor of Psychiatry (with tenure), Department of Psychiatry
2001 Visiting Professor, Department of Psychiatry

Academic

2006-Present Deputy Director of Research, Stein Institute for Research on Aging (SIRA)
2005-Present Director, Scientific Advisory Board, Stein Institute for Research on Aging
2004-Present Co-Director, Center for Human Genetics and Genomics
2004-Present Professor, Biomedical Sciences Graduate Program
2002-Present Member, UCSD Cancer Center
2001-Present Founding Professor, Bioinformatics Graduate Program
2006-2007 Director, Biomedical Informatics, UCSD Moores Cancer Center
2006-2007 Co-director, Cancer Genetics Program, UCSD Moores Cancer Center

Singapore

2007-Present Member, NUS-GIS Center for Molecular Epidemiology, International Advisory Panel

2007-Present Member, The Singapore Consortium of Cohort Studies, International Advisory Panel

San Diego State University

2004-Present Adjunct Clinical Professor in the Graduate School of Public Health

Case Western Reserve University

1997–2001 Associate Professor, Department of Epidemiology & Biostatistics
 1995–1997 Assistant Professor, Department of Epidemiology & Biostatistics
 1995–1998 Assistant Professor (secondary faculty), Department of Genetics
 1994–1995 Assistant Professor, Department of Genetics

Harvard University School of Public Health

1998–2001 Associate Professor (adjunct), Department of Biostatistics
 1995–1997 Assistant Professor (adjunct), Department of Biostatistics

Jackson Laboratories

1994– 2002 Associate Research Scientist (adjunct), The Jackson Laboratory, Bar Harbor, Maine.

Cleveland Clinic Foundation

1996–2001 Assistant Scientist (adjunct staff), Department of Biostatistics and Epidemiology

The Genset Corporation

1999–2000 Vice President, Statistical Genomics, La Jolla, California (sponsored and coordinated leave of absence from CWRU)

CONSULTING POSITIONS

2006-Present DermTech International (San Diego, California): consultant
 2006-Present Ligand Pharmaceuticals (San Diego, California): consultant
 2001-2005 First Genetic Trust (Chicago, Illinois): founding scientific advisory board member
 2001-2005 Celera Diagnostics (Oakland, California): consultant
 1999-Present Genizon (prev. Galileo) Genomics (Montreal, Quebec): member, scientific advisory board
 2001-2003 Merck (Newark, New Jersey): member, pharmacogenetics advisory board
 2001-2002 Affymetrix (Santa Clara, California): consultant
 2001 Xenon Genetics (Vancouver, Canada): consultant
 1997–1999 GENSET (Paris, France): scientific advisor
 1998 Roche Biosciences (Basel, Switzerland): occasional consultant
 1998 Roche Molecular Systems (Alameda, CA): occasional consultant
 1998 Roche Discovery (Welwyn, England): occasional consultant
 1996–1999 Gemini Research (Cambridge, England): member, scientific advisory board
 1996–1999 DNAX Institute of Molecular and Cellular Biology (Palo Alto, CA): occasional consultant
 1996–1999 SmithKline Beecham (King of Prussia, PA): occasional consultant
 1996–1998 Algene Biotechnologies (Montreal, Canada): member, scientific advisory board
 1996–2000 Abbott Laboratories (Chicago, Illinois): member, pharmacogenetics advisory board
 1994–1997 Millennium Pharmaceuticals, Inc. (Cambridge, MA): scientific advisor
 1995–1996 Marion Merrell Dow (Cincinnati, Ohio); occasional consultant.
 1993–1996 Warner-Lambert (Ann Arbor, Michigan): project consultant
 1991–1992 Cleveland Clinic Foundation (Cleveland, Ohio): project consultant
 1989–1990 District Health Department No. 3 (Charlevoix, Michigan): project consultant

PATENTS

Schork NJ, Essioux L, Cohen-Akenin A, Blumenfeld M, Cohen D. “Methods, Software and Apparati for Identifying Genomic Regions Harboring a Gene Associated with a Detectable Trait”. (Issued: 6,29,182 B1)

Schork NJ, Fallin, D, Lisseguere S. “Methods of DNA Marker-based genetic analysis using estimated haplotype frequencies and uses thereof” (Patent Pending; 10/296,867).

Jacob HJ, Schork NJ, Tonellato P, Cowley, AW “Physiological Profiling” (Patent Pending; 09/960,234).

Lanchbury J, Hall M, Schork NJ. “Methodology for identifying genomic regions that affect a biological phenotype” (Application Number: PCTUS02/21,377; United Kingdom Patent Application Submitted).

SCIENTIFIC ACTIVITIES***Editorial Boards***

2000-Present	Associate editor, <i>American Journal of Pharmacogenetics</i>
2002-Present	Associate editor (Statistical Genetics), <i>American Journal of Psychiatry</i>
2000-2003	Scientific advisor, <i>Encyclopedia of Human Genetics</i>
2002-Present	Scientific advisor, <i>Encyclopedia of Diagnostic Genomics and Proteomics</i>
2000-2003	Associate editor, <i>Physiological Genomics</i>
1999-2001	Associate editor, <i>American Journal of Human Genetics</i>
1998-2002	Section editor, <i>Current Hypertension Reports</i>
1995-1999	Associate editor, <i>Gene-COMBIS</i>

Study Sections and Grant Review

2007	National Academy of Sciences, “Nutrigenomics and Beyond,” reviewer
2007	National Health Research Institute of Taiwan, External reviewer
2005	Institute for Mental Health Research (Pheonix, AZ) grant reviewer
2005	Tourette’s Syndrome Association grant reviewer
2004	NIH special contract proposal reviewer, NIDA
2004	NIH, CEBRA reviewer, NIDA
2002	Health Research Board of Ireland, review panel member
2001	European Commission on Bioinformatics, review panel member
2000	MRC of Great Britain, ad hoc review panel member
1999	NIH, special study section member, Renal and Cardiovascular Disease
1999	NIH, special study section member, NIDDK
1998	NIH, special study section member, NIDDK
1998	NIH, special study section member, BSS
1998	NIH, Quantitative Genetics Study Section member, NIMH
1997	NIH, ad hoc review panel member, NHLBI
1997	MRC of Great Britain, ad hoc review panel member
1996	MRC of Great Britain, ad hoc review panel member
1996	MRC of Canada, site visit team member
1996	NIH, Centers of Excellence in Molecular Genetics, review panel member, NCHGR
1995	NIH, Family Heart Study, review panel member, NHLBI
1992	NIH, site visit team member, RCMI

ACTIVE GRANT SUPPORT

Active

- 07/01/05-06/30/06 National Institutes of Health (2 U01 HL064777-07), Family Blood Pressure Program, Nicholas Schork, P.I., 10% effort, annual direct costs: \$96,519
- 06/01/05-5/31/2010 National Institutes of Health (2 P01 HL58120-06A1), Sympathetic Neuroeffector Junctions and Blood Pressure – B: Informatics and Statistical Genetics, Daniel O'Connor, M.D., P.I.; Nicholas Schork, P.I. of Core B, 5% effort, annual direct costs: \$1,527,400
- 04/04/03-02/29/08 National Institutes of Health (5 R01 MH065571-02), The Genetics of Endophenotypes in Schizophrenia, David Braff, M.D., P.I.; Nicholas Schork, director of statistical genetics, 19% effort, annual direct costs: \$493,194
- 09/26/03-08/31/07 National Institutes of Health (1 R01 HL074730-02), Hypertension Pharmacogenetics, Julie A. Johnson, P.I.; (subcontract from the University of Florida, Gainesville), Nicholas Schork, co-investigator, 4% effort, annual direct costs: \$35,198
- 12/01/03-11/30/07 National Institutes of Health (2 R01 MH059567-05A2) Collaborative Genomic Study of Bipolar Disorder, John R. Kelsoe, P.I.; Nicholas Schork, co-investigator, 4% effort, total direct costs \$250,000
- 04/01/04-01/31/09 National Institutes of Health (1 R01 MH068503-01A1) Genomic Studies of Bipolar Disorder and Chromosome 22, John R. Kelsoe, P.I.; Nicholas Schork, co-investigator, 8% effort, total direct costs \$489,650
- 04/15/04-02/28/09 National Institutes of Health (1 R01 HL070137-01A1) Genetics of the Sodium-Lithium Countertransport, Abraham Aviv, P.I.; (subcontract from the University of Medicine and Dentistry New Jersey), Nicholas Schork, co-investigator, 10% effort, total direct costs \$44,501
- 04/01/04-03/31/07 National Institutes of Health (1 R01 HL071123-01A2) Genetic Variants in Circadian Rhythm Sleep Disorders, Daniel Kripke, M.D., P.I.; Nicholas Schork, co-investigator, 4% effort, annual direct costs \$226,081
- 09/30/04-06/30/09 National Institutes of Health (1 U19 AG023122-01A1) A Consortium to Study the Genetics of Longevity, Steven R. Cummings, M.D., P.I.; Nicholas Schork, co-investigator, 15% effort, annual direct costs \$107,453
- 10/01/04-09/30/08 Donald W. Reynolds Foundation, Novel Statistical Approaches to Cardiovascular Genetics, Helen H. Hobbs, M.D.; P.I., Nicholas Schork, co-investigator, 1% effort, annual direct costs \$25,000
- 07/15/05-06-30-09 National Institutes of Health, Human Telomere Genetics, Abraham Aviv, P.I.; (subcontract from the University of Medicine and Dentistry, New Jersey), Nicholas Schork, co-investigator, 1% effort, annual direct costs \$39,970.
- 02/15/05-01/31/10 National Institutes of Health, An Affected SibPair Study of ADHD in Costa Rica. (1R01 NS48376-01A1) Carol A. Mathews, M.D., P.I.; Nicholas Schork co-investigator, 4% effort, annual direct costs \$211,777

HONORS AND AWARDS

- 1997 Burroughs-Wellcome Researcher on Infectious Disease (with Jim Kazura)
- 1992 Student Travel Award Recipient, Spring Biometric Society (ENAR), Cincinnati, OH (Stephen Lagakos, Harvard University, organizer)
- 1991 Student Travel Award Recipient, The 23rd Symposium on the Interface of Computing Science and Statistics, Seattle, WA (Jon R. Kettenring, Bellcore, organizer).

TEACHING ACTIVITIES***Students***

2007-Present Hyun Min Kang, UCSD (Bioinformatics), doctoral thesis committee member
 2007-Present Vikas Bansal, UCSD(Bioinformatics), doctoral thesis committee member
 2006-Present Ali Torkamani, UCSD (Biomedical Sciences), doctoral thesis advisor
 2006-Present Alex Joyner, UCSD (Biomedical Sciences), doctoral thesis advisor
 2006-Present Noah Zaitlen, UCSD (Bioinformatics), doctoral thesis committee member
 2006-Present Jimmie Ye, UCSD (Bioinformatics), doctoral thesis committee member
 2006-Present Ali Bashir, UCSD (Bioinformatics), doctoral thesis committee member
 2005-Present Tobias Gerhard, University of Florida, doctoral thesis committee member
 2004-Present Sean O'Rourke, UCSD (Computer Science), doctoral thesis committee member
 2004-Present Matt Zapala, UCSD (MSTP), doctoral thesis advisor (MD/Ph.D. program)
 2004-2006 Coleman Mosley, MD, UCSD, doctoral thesis advisor
 2004-Present Anne Valle, UCSD, doctoral thesis committee member
 2004-2005 Guillermo Moratorio, UCSD, master's thesis committee member
 2003-Present Michele Day, UCSD, doctoral thesis committee member
 2003-Present Sherri Liang, UCSD, doctoral thesis committee member
 2003-2006 James Rodgers, SDSU, master's thesis committee member
 2004-Present Ryan Fries, UCSD, doctoral thesis committee member
 2003-2004 Ryan Fries, UCSD, master's thesis committee member
 2003-Present Rany Salem, UCSD, doctoral studies research advisor
 2003-2006 Jennifer Wessel, UCSD, doctoral thesis advisor
 2002-Present Martin Jirout, MD, UCSD, doctoral thesis advisor
 2001-2004 Lynn Evans, UCSD, doctoral thesis committee member
 1999-2001 Nadia Tahri, University of Paris XI, doctoral thesis advisor
 1998-2003 Fabrice Larribe, University of Montreal, doctoral thesis co-advisor
 1998-2001 Li-Lian Kim, CWRU, doctoral thesis committee member
 1998-2000 Danielle Fallin, CWRU, doctoral thesis advisor
 1998 Markus Perola, University of Helsinki, doctoral committee member
 1998 Cixuang Zheng, CWRU, doctoral thesis committee member
 1997 Stacy Hirth, master's thesis advisor
 1995-1998 Bonnie Thiel, CWRU, master's thesis advisor
 1995-1997 Wade Najem, CWRU, doctoral thesis advisor
 1995-1996 Eric Puffenberger, CWRU, doctoral thesis committee member
 1994-1998 Tim Nui, Harvard University, doctoral thesis committee member
 1994-1996 Audrey Lynn, CWRU, doctoral thesis committee member

Lab Rotations

2006 Stephanie Nissan, UCSD, doctoral student
 2006-Present Nzali Campbell, UCSD PREP Awardee mentor
 2006 Trygve Bakken, UCSD, doctoral student
 2006- Pamela Chayavichitsilp, UCSD medical school ISP rotation project committee
 2006 Caroline McGruther, UCSD, M.D./Ph.D. student
 2006 Nina Haste, UCSD, doctoral student, School of Pharmacy
 2006 Oanh Nguyen, UCSD medical school ISP project co-advisor
 2006 Ali Torkamani, UCSD doctoral student
 2006 Alex Joyner, UCSD doctoral student
 2005-2007 Jan McClure, UCSD/SDSU, student research rotation advisor
 2005 Ohn Chou, UCSD doctoral student
 2004-2005 Chileshe Mabula, UCSD SIRA undergraduate student investigator, advisor
 2004-2005 Matt Bengard, UCSD, medical school ISP project co-advisor
 2004-2005 Carie Nguyen, UCSD, medical school ISP project co-advisor
 2004 Ashley Ebert, UCSB undergraduate, lab rotation advisor
 2003-2004 Samuel Payne, UCSD, doctoral student, Biomedical Science

2002-2003 Ashwin Chandra, UCSD undergraduate student
 2002-2005 Brad Kohlenberg, UCSD undergraduate research mentor
 1996-1998 Joshua Pinkus, The Jackson Laboratory, undergraduate student intern co-mentor
 1995 Michael Decker, CWRU, undergraduate research mentor

Post-Doctoral Mentor

2007-Present Jennifer Wessel, Ph.D., post-doctoral mentor
 2006-Present Nathalie Malo, Ph.D., post-doctoral mentor
 2002-2007 Tiffany Greenwood, Ph.D., project scientist mentor
 2002-Present Caroline Nievergelt, Ph.D., project scientist mentor
 2001-2007 Brinda K. Rana, Ph.D., project scientist mentor
 2002-2004 Miguel T. Robinson, Ph.D., project scientist mentor
 1999-2000 Swapan Nath, Ph.D., post-doctoral mentor
 1997-2000 Mark Seielstad, Ph.D., post-doctoral co-mentor
 1997-1998 Zanquin Liu, Ph.D., post-doctoral co-mentor
 1997 Vincent Carrier, Ph.D., post-doctoral mentor
 1996 John Rogus, Ph.D., post-doctoral co-mentor
 1995-1998 Pamela St. Jean, Ph.D., post-doctoral mentor

Career Development Mentor (K-Awardees , etc.)

2006-Present Steven Glatt, Ph.D., UCSD (K-Award)
 2005-Present Eleazar Eskin, Ph.D., UCSD (K-Award)
 2001-2006 Carol Mathews, M.D., UCSD, informal career mentor
 2002-2005 Sally Bao, M.D., Ph.D., UCSD (K-Award)
 2001-Present Tom Barrett, Ph.D., UCSD (K-Award)
 2000-2005 Jordon Smoller, M.D., Harvard University (K-Award)
 2002-2004 Murray Stein, M.D., UCSD, formal mentor for Johns Hopkins MPH degree program
 1997-1998 Tom Ferraro, Ph.D., University of Pennsylvania

Visiting Scientist/Mini-Sabbatical Mentor

2006 Alan B. Weder, M.D., University of Michigan
 2003 Maud Artaud, M.S., Université de Poitiers
 2003 Mélanie Langlois, M.S., Université de Poitiers
 2003 Richard Prebish, M.S., University of Michigan, Ann Arbor
 1997 Heidi Stirnadel, Ph.D., Swiss Tropical Institute
 1997 Niall Anderson, Ph.D., University of Glasgow
 1996 Robert Zee, Ph.D., Harvard University
 1996 David Allison, Ph.D., St. Luke's Hospital/Columbia University

International Teaching Lectures

1996 Lectures on the genetic analysis of complex traits; University of Oulu Course Short Course on "Molecular Genetics and Diseases," University of Oulu, Finland (2 x 1.5 hour lectures)
 1996 Lectures on "Linkage disequilibrium mapping of human quantitative traits;" University of Helsinki Medical School, University of Helsinki, Finland (1 2 hour lecture, workshop participation).

National Teaching Lectures

1996 Lecture on "Linkage Mapping of Traits Exhibiting Marked Age-Trends," Workshop on Statistical Genetics, Institute of Mathematics and its Applications, University of Minnesota, MN (1 hour lecture).

- 1996 Lecture on “Sibpairs vs. Pedigrees: What are the Advantages?” American Diabetes Association Workshop on the Genetics of Diabetes Susceptibility, Aspen, Colorado (2 x 1 hour lectures)
- 1996 Lecture on the “Potential Contribution of Molecular Genetics to Speech and Language Research,” American Speech and Hearing Association Workshop on the Genetics of Speech and Language Disorders, Seattle, Washington (1 hour lecture)
- 1998 Center for Environmental Genetics Workshop organizer and lecturer: Marker-Based Human Genetic Epidemiologic Analysis (University of Cincinnati) 1 8-hour workshop.
- 1998 Abbott Laboratories one-day workshop/lecture series on the “Modern Marker-Based Genetic Analysis,” 6 1-hour lectures. Abbott Park, Illinois.
- 1998 Roche/Jackson Laboratories Workshop on Functional Genomics and Gene Expression
- 2000 “Modern Genetic Epidemiology” Epidemiology Research Institute Short Course in Epidemiologic Methods, Boston, MA July 24-28,2000

University of California, San Diego

- 2002 Lectures: BIOM 243 Human Genetics, four lectures on linkage and association mapping; Tony Wynshaw-Boris, organizer; spring quarter
- 2002 Lecture: Chem 92 Undergraduate Pharmacology Seminar, lecture on pharmacogenetics; Bob Tukey, organizer; spring quarter
- 2002 Lecture: BILD 94 Professional Issues in Bioinformatics, lecture on statistical genetics; Suresh Subramani, organizer; spring quarter
- 2002 Lecture: SoM 218 Epidemiology/Biostatistics, lecture on genetic epidemiology
- 2002 Lecturer: Epidemiology II (2 lectures on genetic epidemiology); UCSD CREST program in graduate epidemiology, October 30 and November 6
- 2002 Mentor, undergraduate research presentations at the annual undergraduate research symposium.
- 2003 Lecturer: SoM Pharm/Biom 235/Med 230 Pharmacogenomics, two 1-1/2 hour lectures on statistical genetics; Daniel O’Connor, organizer; April 1 and April 10
- 2003 Lecture: Chem 92 Undergraduate Pharmacology Seminar, lecture on pharmacogenetics; Bob Tukey, organizer; April 25
- 2003 Lecture: BICD 180 Genetics of Model Organisms, lecture on Homo sapiens: human and medical genetics; Randall Johnson, instructor; May 5
- 2003 Mentor, undergraduate research presentations at the annual undergraduate research symposium.
- 2003 Lecture: Math 283 Bioinformatics, lecture on statistical issues in genetic mopping; John O’Quigley, instructor; June 2
- 2003 Lectures: CREST Program, Epidemiology II (two lectures on genetic epidemiology); November 19 and December 10
- 2004 Organizer and lecturer: Math 283 Statistical Methods in Bioinformatics, spring quarter

- 2004 Lecture: Chem 92 Undergraduate Pharmacology Seminar, lecture on pharamcogenetics, Bob Tukey, organizer; April 24
- 2005 Organizer and lecturer, SPPS 216: “Human Genetics and Genomics,” (Palmer Taylor, overall director), January 27
- 2005 Lecturer, FPM 237: “Microarray applications in the health sciences,” (Sonia Jain, Mike Cleary, Gary Hardiman, organizers), March 3.
- 2005 Lecture: Chem 92 Undergraduate Pharmacology Seminar, lecture on pharamcogenetics, Bob Tukey, organizer; April 27
- 2005 Organizer and lecturer, SIRA workshop on Genetics and Imaging. “Modern Human Genetics and Genomics,” (Lisa Eycler, overall organizer), June 8.
- 2005 Lecture: (2): BIOM 200, “Genetic Dissection of Complex Traits”, “Evolutionary Genetics/Genomics” (Bruce Hamilton, organizer), September 26 and October 4.
- 2005 Lecture: SOM 218 “Statistical Genetics: Concepts in Genetic Mapping”, (Reena Deutsch, organizer), November 19.
- 2005 Fall term independent studies course (15 meetings) on the Genetics of Infectious Disease for Lorne Walker, MSTP student. Fall Term.
- 2005 Lecture: SPPS 216 “Statistical Genetics and Pharmacogenetics”, (Palmer Taylor, overall course director), January 31.
- 2005 Moderator, BGGN 219, Classic Papers in Human Genetics (Tony Wynshaw-Boris, organizer), March 8.
- 2006 Lecture: SPPS 216 “Introduction to Statistical Genetics and Pharmacogenetics,” (Palmer Taylor, overall course director), January 28.
- 2006 Lecture: FPM 243, “Clinical Genomics” (Lisa Madlensky, organizer), February 23.
- 2006 Lecture: BIOM 252 “Linkage Disequilibrium” (Tony Wynshaw-Boris, organizer), February 27.
- 2006 Lecture: FPM 237, “Applications of Microarrays to the Health Sciences,” (Sonia Jain, organizer), March 9.
- 2006 Lecture: GeroPsychiatry Fellowship Lecture Series, “Modern Genetic Epidemiology,” (Lori Montross, organizer), April 5, 2006.
- 2006 Chemistry 92, “Pharmacogenetics and Pharmacogenetic Inquiry” (Robert Tukey, organizer), April 21.
- 2006 Lecture: FPM 258, “Large-scale genetic association analysis” (Kristen Deveraux, organizer), May 23.
- 2006 Organizer, Biomedical Science Student Orientation, University of California, San Diego, September, 2006.
- 2006 Lecture: (2): BIOM 200, “Genetic Dissection of Complex Traits”, “Evolutionary Genetics/Genomics” (Steve Dowdy and Alexandra Newton, organizers), September 25 and October 5.

- 2007 Lecture: (2), BGGN 219 “Classic Papers in Human Genetics: Human Population Genetics”, January 8 and January 10.
- 2007 Lecture: Bioinformatics Graduate Program Faculty Lecture: “Statistical Genetics,” January 15.
- 2007 Lecture: SPPS 216 “Introduction to Statistical Genetics and Pharmacogenetics,” (Palmer Taylor, overall course director), February 6.
- 2007 Lecture, Graduate Fellowship in Biological Psychiatry and Neuroscience Seminar Series on “Whole Genome Association Studies,” (Eric Turner, fellowship coordinator), March 1.
- 2007 Lecture: FPM 237, “Applications of Microarrays to the Health Sciences,” (Sonia Jain, organizer), March 6.

University of Alabama, Birmingham

- 2003 Lecture series: Department of Biostatistics, School of Public Health, five one-hour seminars in the Section on Statistical Genetics, August 18-22
- 2003 Visiting professor, Department of Biostatistics, School of Public Health

The Jackson Laboratory

- 1995 Lectures on complex trait analysis and the origins of complex traits, and computer laboratory organizer for gene mapping; The Jackson Laboratory Short Course on "Experimental Genetics of the Laboratory Mouse," Bar Harbor, ME (2 x 2 hour lectures; 3 x 3 hour workshops)
- 1996 Lectures on complex trait analysis and the origins of complex traits, and computer laboratory organizer for gene mapping; The Jackson Laboratory Short Course on "Experimental Genetics of the Laboratory Mouse," Bar Harbor, ME (2 x 2 hour lectures; 3 x 3 hour workshops)
- 1996 Lectures on complex trait analysis and the origins of complex traits, and computer laboratory organizer for gene mapping; The 37th Annual Joint Jackson Laboratory/Johns Hopkins University Short Course on "Human and Experimental Genetics," Bar Harbor, ME (2 x 2 hour lectures; 3 x 3 hour workshops)
- 1996 Lectures on complex trait analysis and the origins of complex traits, and computer laboratory organizer for gene mapping; The Joint Jackson Laboratory/NHLBI/ American Type Culture Short Course on the Genetics of Heart, Lung, and Blood Diseases, Bar Harbor, ME (2 x 2 hour lectures; 1 2 hour workshops)
- 1997 Lectures on complex trait analysis and the origins of complex traits, and computer laboratory organizer for gene mapping; The Jackson Laboratory Short Course on "Experimental Genetics of the Laboratory Mouse," Bar Harbor, ME (1 x 3 hour lectures; 3 x 1.5 hour workshops)
- 1997 Lectures on complex trait analysis and the origins of complex traits, and computer laboratory organizer for gene mapping; The Joint Jackson Laboratory/NHLBI/ American Type Culture Short Course on the Genetics of Heart, Lung, and Blood Diseases, Bar Harbor, ME (2 x 2 hour lectures; 1 2-hour workshops)

- 1998 Lectures on complex trait analysis and the origins of complex traits, The Jackson Laboratory Short Course on "Experimental Genetics of the Laboratory Mouse," Bar Harbor, ME (1 2-hour lecture)
- 1999 Lectures on complex trait analysis and the origins of complex traits, and computer laboratory organizer for gene mapping; The Joint Jackson Laboratory/NHLBI/ American Type Culture Short Course on the Genetics of Heart, Lung, and Blood Diseases, Bar Harbor, ME (1 2-hour lecture)
- 1999 Lectures on complex trait analysis and the origins of complex traits, The Jackson Laboratory Short Course on "Experimental Genetics of the Laboratory Mouse," Bar Harbor, ME (1 2-hour lecture)
- 2000 Lectures on complex trait analysis and the origins of complete traits, and computer laboratory organizer for gene mapping; The Joint Jackson Laboratory/NHLBI/American Type Culture Short Course on the Genetics of the Heart, Lung, and Blood Diseases, Bar Harbor, ME (1 2-hour lecture)
- 2000 Co-organizer and lecturer, "Mathematical Approaches to the Analysis of Complex Traits and Diseases," The Jackson Laboratory, ME (2 1-hour lectures)
- 2002 Short course on Mathematical Approaches to the Analysis of Complex Phenotypes, "Issues in Contemporary Human Genetic Analysis: Haplotype Maps, Bioinformatics Tools and Novel Statistical Methods," Bar Harbor, ME (1 2-hour lecture)
- 2003 Short course on Mathematical Approaches to the Analysis of Complex Phenotypes, "Issues in contemporary human genetic analysis: haplotype maps, bioinformatics tools, and novel statistical methods," Bar Harbor, ME (1 2-hour lecture)

Harvard University

- 1996 EHS 287: Introduction to Genetic Epidemiology, course developer (3 x 2 hour lecture; 50 students)
- 1999 Short Course: Variance Components Models in Modern Genetic Epidemiology (5 x 2 hour lectures; 100 students), February, 1999.

Case Western Reserve University

- 1994 Medical Student Preclinical Survey Course, Department of Medicine (2 hour lecture)
- 1994–1997 Gene 504: Survey Course, Department of Genetics (3 x 2 hour lectures)
- 1995 EPBI 451: Introduction to Genetic Epidemiology (2 hour lecture)
- 1995-1999 EPBI 494: Infectious Disease, Department of Epidemiology and Biostatistics (2 hour lecture)
- 1996 EPBI 701: Population Genetics in Genetic Epidemiology, Department of Epidemiology and Biostatistics, Course Developer and Lecturer
- 1998 EPBI 454: Evolution and Population Genetics in Genetic Epidemiology. Course organizer and teacher.
- 1999-2000 EPBI 454: Evolution and Population Genetics in Genetic Epidemiology. Course organizer and leader.

EXTRAMURAL INVITED PRESENTATIONS

Published Lectures

2007.1 "Evolutionary Genetic Epidemiology," Henry Stewart Talks (www.hstalks.com); Series on Evolutionary Medicine (Randolph Nesse, lecture series organizer).

Past Speaking Engagements

1993.1 "Detecting Genes That Influence Traits Exhibiting Marked Postnatal Developmental Trends," invited seminar, Department of Genetics, Harvard University, Massachusetts General Hospital, November, 1993.

1993.2 "Detecting Genes That Influence Traits Exhibiting Marked Postnatal Developmental Trends," invited seminar, University of Florida, Center for Mammalian Genetics, December, 1993.

1994.1 "Extended Multipoint Identity-by-Descent Mapping of Complex Quantitative Phenotypes," invited seminar, Harvard University, Department of Biostatistics, January, 1994.

1994.2 "Issues in the Analysis of Multifactorial Traits," invited seminar, Center d'Etude du Polymorphisme Humain, Paris, France, February, 1994.

1994.3 "Linkage Analysis of Complex Traits," invited seminar, University of Washington, Department of Biostatistics, February, 1994.

1994.4 "Linkage Mapping of Traits Exhibiting Marked Age-Trends: Applications Using Twins," Workshop on Methodology for Genetic Studies of Twins and Families, University of Colorado, February, 1994.

1994.5 "Linkage Analysis of Complex Traits," invited seminar, National Center for Human Genome Research, NIH, Bethesda, MD, March 1994.

1994.6 "Linkage Mapping of Traits Exhibiting Marked Age-Trends," and "A Biometrical Genome Scan Search Reveals the Multilocus Basis of Blood Pressure Regulation," invited seminars, Medical College of Wisconsin, Department of Physiology, Madison, WI, March, 1994.

1994.7 "Linkage Analysis of Complex Traits," invited seminar, Marquette University, Department of Mathematics and Statistics, March, 1994.

1994.8 "A Biometrical Genome Scan Search Reveals the Multilocus Basis of Blood Pressure Regulation," invited seminar, The Jackson Laboratories, June, 1994.

1994.9 "Issues in the Analysis of Complex Traits," invited speaker, Alder Foundation Symposium: Alzheimer's Disease: A cutting edge discussion of genetic epidemiology and the role of APO-E, Duke University, NC, August 1994.

1994.10 "Linkage Analysis of Complex Genetic Traits in Mice," invited speaker, Symposium on Mouse Molecular Neurogenetics, The Jackson Laboratories, Bar Harbor, Maine, August, 1994.

1994.11 "Models and Methods for Genetic Analysis of Complex Traits," invited seminar, Harvard University, Department of Biostatistics, December, 1994.

1995.1 "Methods in the Dissection of Complex Genetic Traits," invited seminar, Sergievsky Center, Columbia University, NY, January, 1995.

1995.2 "The Y Chromosome and Blood Pressure Variation in Inbred Rat Strains," invited seminar, The University of Michigan, Department of Medicine, Ann Arbor, Michigan, March, 1995.

- 1995.3 “A Biometrical Genome Scan Search Reveals the Multilocus Basis of Blood Pressure Regulation,” invited seminar, National Institute of Environmental Health Sciences, North Carolina, May, 1995.
- 1993.4 “Extensions to QTL Mapping,” invited speaker, Gordon Research Conference on the Genetics of Disease, Salve Regina College, Rhode Island, August, 1995.
- 1995.5 “On the Use of Multiple Phenotypes in Mapping Complex Trait Genes,” invited speaker, NIH Workshop on “Phenotypic Issues in Studies of Asthma Genetics,” August, 1995.
- 1995.6 “Mapping Multigenic Traits,” invited seminar, Harvard University, Department of Biostatistics, Boston, MA, October, 1995.
- 1996.1 “On the Use of Multiple Phenotypes in Mapping Complex Trait Genes,” invited speaker, Banbury Center, Cold Spring Harbor Laboratory, conference on "Looking to the Next Generation of Genetic Mapping," New York, November, 1996.
- 1996.2 “Novel Approaches to the Analysis of Complex Traits,” invited seminar, The University of Michigan, Department of Human Genetics, Ann Arbor, Michigan, May, 1996.
- 1996.3 “Novel Approaches to the Analysis of Complex Traits,” invited seminar, Ohio State University Symposium on Biostatistical Methodology, Columbus, Ohio, May, 1996.
- 1996.4 “Linkage Mapping of Multigenic Diseases,” invited speaker, Fu Wai Cardiovascular Hospital, Chinese Academy of Medical Sciences, Beijing, China, May, 1996.
- 1996.5 “Extensions of QTL Mapping,” invited speaker, International Society of Hypertension Satellite Symposium on Genetics, Glasgow, Scotland, June, 1996.
- 1996.6 “Issues in Assessing Statistical Significance in QTL Mapping Studies,” invited speaker, NIAAA workshop on QTL mapping, Bethesda, MD, August, 1996.
- 1996.7 “Promising Study Designs” and “Overall Research Strategies for the Genetics of Aging,” invited speaker, NIA Panel on the Epidemiology and Genetics of Longevity, Bethesda, MD, August, 1996.
- 1996.8 “The Genetic Dissection of Complex Traits,” invited speaker, American Hypertension Association Annual Meeting, Chicago, IL, August, 1996.
- 1996.9 “Novel Methods for Mapping Complex Traits,” invited seminar, University of Pittsburgh, Department of Human Genetics, Pittsburgh, PA, November, 1996.
- 1996.10 “Novel Methods for Mapping Complex Traits,” invited seminar, University of Helsinki, Department of Medical Genetics, Helsinki, Finland, December, 1996.
- 1997.1 “The Genetics of Complex Diseases: Issues and Strategies,” invited keynote speaker, American Association for Asthma Research, Boca Raton, Florida, January, 1997.
- 1997.2 “Novel Methods for Mapping Complex Genetic Traits,” invited speaker, Gordon Research Conference on Quantitative Genetics and Biotechnology, Ventura, CA, February, 1997.
- 1997.3 “Issues in Assessing Associations and Linkages with Addictive Behaviors,” invited speaker, UCLA Conference on Molecular Genetics of Alcoholism and Other Addictive/Compulsive Disorders, Los Angeles, CA, February, 1997.
- 1997.4 “Multigenic Diseases,” invited speaker, Human Genome Meeting Sponsored by the Human Genome Organization (HUGO), Toronto, Canada, March, 1997.

- 1997.5 “Allele Sharing Methods for the Analysis of Twin Data,” invited speaker, NordicTwin Registry Workshop, Helsinki, Finland, March, 1997.
- 1997.6 “Novel Methods for Mapping Complex Genetic Traits,” invited speaker, Max Delbruck Center for Molecular Medicine, Berlin, Germany, March, 1997.
- 1997.7 “Novel Methods for Mapping Complex Genetic Traits,” invited speaker, distinguished lecture series, University of Medicine and Dentistry of New Jersey, Newark, New Jersey, April 8, 1997.
- 1997.8 “Genetic Mapping Strategies for Complex Traits Based on the Short-term Evolution of Chromosomes,” invited speaker, 9th International Congress on Genes, Gene Families, and Isozymes, San Antonio, TX, April, 1997.
- 1997.9 “Multivariate Gene Mapping Methods,” invited speaker, New York Obesity Research Meeting, New York, July, 1997.
- 1997.10 “Novel Methods for Mapping Complex Genetic Traits,” invited speaker, American Statistical Association Annual Meeting, Anaheim, CA, August, 1997.
- 1997.11 “Novel Methods in the Meiotic Mapping of Complex Traits,” invited speaker, Sequana Therapeutics, La Jolla, CA, August, 1997.
- 1997.12 “Novel Methods for Mapping Complex Genetic Traits,” invited speaker, Harvard University, Department of Medicine, Boston, MA, September, 1997.
- 1997.13 “New Directions in the Analysis of Complex Genetic Traits,” invited speaker, Harvard University School of Public Health, Boston, MA, September, 1997.
- 1997.14 “Issues and Strategies in the Assessment of Genetic Function and Complexity,” invited speaker, The Jackson Laboratory, Bar Harbor, Maine, October 6, 1997.
- 1997.15 “Issues in the Analysis of Gene Expression Data,” organizer, Roche/Jackson Laboratories workshop on Functional Genomics, Roche Biosciences, Palo Alto, CA, November 6, 1997.
- 1997.16 “Novel Approaches to Investigating Complex Traits in Humans: A Two Village Study in China ,” invited speaker, University of Cincinnati, Department of Medicine, November 13, 1997.
- 1997.17 “Statistical Methods for Assessing the Genetic Basis of Complex Traits ,” invited speaker, Department of Biostatistics, University of Cincinnati, November 13, 1997.
- 1997.18 “Novel Approaches to Investigating Complex Traits in Humans: Population Genetics and Large-Scale Epidemiology ,” invited speaker, StrataGene Corporation, La Jolla, CA, November 20, 1997.
- 1998.1 “Issues in Marker-Based Analysis of Complex Genetic Traits,” invited speaker, Marshfield Clinic, Marshfield, Wisconsin, February 14, 1998.
- 1998.2 “Weighing the Odds in Genetic Analysis: Approaches,” invited speaker, Gordon Research Conference, Ventura, CA, February 17, 1998.
- 1998.3 “Issues in Marker-Based Analysis of Complex Genetic Traits,” invited speaker, Algene Biotechnologies, Montreal, Canada, February 26, 1998.
- 1998.4 “Issues in Marker-based Genetic Analysis of Disease,” invited speaker, Smith-Kline Beecham, London, England, March 25, 1998.
- 1998.5 “Use of Twins in QTL Mapping,” invited speaker, First International Workshop on Genetic Epidemiology of Complex Traits using Twins and Sibpairs, Cambridge, England, March 26, 1998.

- 1998.6 “Analysis of Target Gene Variations in Complex Disease,” invited speaker, Cold Spring Harbor Laboratory meeting on the Molecular Basis of Asthma, March 30-April-2, 1998.
- 1998.7 “Issues in the Marker-based Analysis of Complex Traits,” invited speaker, Lunenfeld Research Center, University of Toronto, Canada, April 7, 1998.
- 1998.8 “Marker-based Analysis of Complex Traits,” invited speaker, Department of Human Genetics, UCLA, April 15, 1998.
- 1998.9 “Issues in the Marker-based Analysis of Complex Traits,” invited speaker, 50th anniversary of the National Heart, Lung, and Blood Institute, San Francisco, CA, April 19, 1998.
- 1998.10 “Issues in the Marker-based Analysis of Complex Traits,” invited speaker, O’Brien Kidney Center, University of Iowa, May 7, 1998.
- 1998.11 “The Future of Polymorphism Analysis,” invited speaker and organizer, Roche/Jackson Laboratories Workshop on Polymorphisms and Drug Discovery, Roche Discovery, Welwyn, England, May 19, 1998.
- 1998.12 “Disease models and functional genomics,” invited speaker, Roche/Jackson Laboratories research de-briefing, The Jackson Laboratory, May 26, 1998.
- 1998.13 “Statistical Methods Used in Genome Scans,” and “Issues in Association Studies,” National Institute of Diabetes and Digestive and Kidney Diseases Conference on “Strategies for Identification of Nephropathy Susceptibility Genes,” Bethesda, MD, June 28-29, 1998.
- 1998.14 “Approaches to the Analysis of Complex Genetic Traits,” invited speaker, Department of Biostatistics, University of Southern California, August 7, 1998.
- 1998.15 “Genetic Analysis and SNPs,” co-invited speaker, Swedish conference on SNPs and Genetic Analysis, Stockholm, Sweden, August 30, 1998.
- 1998.16 “Novel Approaches to the Genetic Analysis of Complex Traits,” invited speaker, Department of Epidemiology and Biostatistics seminar series, Case Western Reserve University, Cleveland, Ohio, September 23, 1998.
- 1998.17 “Issues in the Genetic Analysis of Complex Traits,” invited speaker, MetroHealth Medical Center, Cleveland, Ohio, September 29, 1998.
- 1998.18 “The Genetics of Hypertension,” invited speaker, Division of Hypertension, Department of Medicine, University of Michigan, October 5, 1998.
- 1998.19 “Disease Models and Functional Genomics” The Jackson Laboratory Symposium on the Mouse and Drug Discovery, Asticou, ME, October 20-21, 1998.
- 1998.20 “Statistical Issues in Gene Mapping,” invited speaker, Roche/Jackson Laboratories workshop on Bioinformatics. Basel, Switzerland, November 1-2, 1998.
- 1998.21 “Whole Genome Association Studies,” co-invited speaker, Banbury Center meeting on large-scale discovery and genetic applications of SNPs, Cold Spring Harbor, NY, November 10-13, 1998.
- 1999.1 “Population Genetic Analysis of Social and Geographical Isolates”, invited speaker, VIIth Annual CEPH Conference on Human Genetics, May 27-28, 1999
- 1999.2 “New Approaches to Therapeutic Target Discovery – Overview Lecture”, invited speaker, Fifth International Symposium on the Immunotherapy of the Rheumatic Diseases, Cyprus, May 26-30, 1999

- 1999.3 “Genetic Analysis of Single Nucleotide Polymorphisms,” invited lecture, Roche Molecular Systems, Alameda, CA, July 15, 1999.
- 1999.4 “The Future of Genetic Analysis,” invited speaker, Hoffman-La Roche Symposium on the future of genetics research, Chantilly, WV, September 19, 1999.
- 1999.5 “Linkage Disequilibrium Strength as a Function of Social and Demographic Factors,” slide presentation, American Society of Human Genetics Annual Meeting, San Francisco, CA, October 22, 1999.
- 1999.6 “The Future of Genetic Case-Control Studies,” invited lecture, Roche Molecular Systems Symposium on the future of genetic analysis, Alameda, CA, October 25, 1999.
- 1999.7 “Quantitative Trait Loci,” invited speaker, Association for Molecular Pathology annual meeting, St. Louis, MO, November 5, 1999.
- 1999.8 “The Future of Genetic Case-Control Studies,” invited speaker, Bristol-Myers Squibb, Philadelphia, PA, December 1, 1999.
- 1999.9 “Analysis with Genetic Markers,” invited speaker, Association of Physiological Research, Paris, France, December 9-11, 1999.
- 2000.1 “Modern Genetic Case/Control Studies”, invited speaker, University of California, San Diego, CA, February 2, 2000
- 2000.2 “Optimal Design of SNP Studies,” invited speaker, NHGRI and SNP Consortium meeting, Bethesda, MD, March 7-8, 2000
- 2000.3 “Genetic Association Studies,” invited speaker, ASPET meeting, Boston, MA, June 4, 2000.
- 2000.4 “Genetic Case/Control Studies,” invited speaker, Pharmacogenetics 2000, San Diego, CA, June 14, 2000.
- 2000.5 “Whole Genome Association Studies,” invited speaker, 3rd Annual Conference on Single Nucleotide Polymorphisms and the Analysis of Complex Disease, Taos, NM, September 8-11, 2000.
- 2000.6 “Modern Genetic Analysis,” invited speaker, International Society of Nephrology Research meeting, Toronto, Canada, October 7, 2000.
- 2000.7 “The Future of Genetic Case/Control Studies,” invited speaker, Roswell Park Cancer Institute, Roswell Park, NY, October, 30, 2000.
- 2000.8 “Whole Genome Association Study of Calcium Entry into the Cell,” slide presentation, American Society of Human Genetics Annual Meeting, October 3-9, 2000
- 2000.9 “Linkage Disequilibrium and Whole Genome Association Studies: Theory and Applications,” invited speaker, The Curagen Corporation, New Haven, CT, November 7, 2000
- 2000.10 “Linkage Disequilibrium and Whole Genome Association Studies: Theory and Applications,” invited speaker, University of California, San Diego, Department of Mathematics, November 8, 2000
- 2000.11 “Linkage Disequilibrium and Whole Genome Association Studies: Theory and Applications,” invited speaker, University of California, San Diego, Department of Psychiatry, November 15, 2000
- 2000.12 “Linkage Disequilibrium and Whole Genome Association Studies: Theory and Applications,” invited speaker, Department of Statistics, Case Western Reserve University, 17, 2000
- 2000.13 “Genetic Epidemiology of SNPs,” invited presentation, Atherothrombosis, Lisbon, Spain, December 11, 2000.

- 2000.14 “Linkage Disequilibrium and Whole Genome Association Studies: Theory and Applications,” invited speaker, Department of Biostatistics, University of Michigan, Ann Arbor, Michigan, December 19, 2000.
- 2001.1 “Statistical Analysis Tools for Modern SNP-Based Genetic Analysis.” invited speaker, IBC USA 3rd Annual Pharmacogenetics, SNPs, and Genetic Patenting Meeting. San Diego, CA, February 12-14, 2001.
- 2001.2 “The Future of Association Studies of Complex Traits,” invited speaker, Medical College of Wisconsin, Department of Medicine, Milwaukee, Wisconsin, February 19, 2001.
- 2001.3 “Whole-genome association studies of complex traits,” invited speaker, . Program in Human Genetics, University of California, San Francisco, March 13, 2001.
- 2001.4 “Novel analytical approaches in analyzing the complex trait, asthma,” David G Marsh Invited Lectureship, The American Academy of Allergy, Asthma, and Immunology, 57th annual meeting, New Orleans, LA, March 16-21, 2001.
- 2001.5 “Linkage Disequilibrium Analysis,” invited speaker, First Genetic Trust, New York, NY, April 2, 2001.
- 2001.6 “The Future of Association Studies of Complex Traits” and “A Comparison of Mean Effects and Variance Components Models for Group Comparisons,” invited speaker, University of California, San Diego, Departments of Psychiatry and Mathematics, April 11-13, 2001.
- 2001.7 “Applied Population Genetics,” invited speaker, Short Course in Statistical Genetics, University of Alabama, Birmingham, May 15-18, 2001.
- 2001.8 “Applied Population Genetics: Concepts, Methods, and Relevance to Disease Gene Mapping in Isolates,” invited speaker, American Society of Human Genetics symposium on Genetic Studies in Isolated Populations, San Diego, CA, October 15, 2001.
- 2001.9 “Modern Genetic Analysis Tools,” invited speaker, IBC Meeting on Functional Genomics, Seattle, WA, October 16-17, 2001.
- 2002.1 “Power calculations for genetic associations studies using empirically-derived parameter distributions,” invited speaker, Human Genetics Journal Club, University of California, San Diego, January 22, 2002.
- 2002.2 “Power calculations for genetic associations studies using empirically-derived parameter distributions,” invited speaker, Department of Biostatistics, Stanford University, Palo Alto, CA, January 31, 2002.
- 2002.3 “Power calculations for genetic associations studies using empirically-derived parameter distributions,” invited speaker, Celera Diagnostics, Alameda, CA, February 1, 2002.
- 2002.4 “Improving Case-Control Studies: Haplotypes and Homogeneity Assessment,” invited speaker, Second Longevity Consortium Meeting, Napa, CA, February 4, 2002.
- 2002.5 “SNP-Based Genetic Analysis Tools,” invited speaker, Cambridge Healthtech Institute’s Tri-Genome Conference 2002, Santa Clara, CA, February 25-27, 2002.
- 2002.6 “SNP-Based Genetic Analysis Tools,” invited speaker, Roche Molecular Systems, Alameda, CA, February 27, 2002.
- 2002.7 “Future Directions in Statistical Genetics,” invited keynote speaker and “Haplotype Analysis,” invited speaker, Third Annual Conference on the Genetic Analysis of Complex Traits, Cambridge, United Kingdom, April 1-3, 2002.
- 2002.8 “Novel DNA-marker Based Analyses: Parametric IBD Linkage Analysis and Phenotype-Genotype Distance Analysis,” invited speaker, Family Blood Pressure Program, Baltimore, MD, July 14-16, 2002

- 2002.9 "Identifying Drug Response Genes," invited speaker, New York Academy of Medicine Pharmacogenetics Symposium, New York, NY, October 3-4, 2002.
- 2002.10 "Bioinformatics: Applied Population Genetics," invited speaker, Annual Meeting of the American College of Clinical Pharmacy, Albuquerque, NM, October 21-23, 2002.
- 2002.11 "Large-scale Genetic Association Studies," invited speaker, Celera Diagnostics, Oakland, CA, November 1, 2002.
- 2002.12 "Large-scale Genetic Association Studies," invited speaker, University of Florida, School of Pharmacy, Gainesville, FL, November 11-13, 2002.
- 2002.13 "Haplotype Analysis with applications to Alzheimer's disease," invited speaker, National Academy of Sciences' Fourteenth Annual Beckman Frontiers of Science Symposium, Irvine, CA, November 14-16, 2002.
- 2002.14 "Genetic Epidemiology," invited speaker, Department of Biostatistics, University of California, San Diego, La Jolla, CA, December 11, 2002.
- 2002.15 "Statistical Issues in Cancer Genetics Research," invited speaker, National Cancer Institute, Cancer Susceptibility Genes in Human and Model Organisms, University of California at Irvine, Newport Beach, CA, December 11-13, 2002.
- 2003.1 "The Haplotype Map Initiative and Whole Genome Association Studies," invited speaker, Department of Biostatistics, University of Southern California, Los Angeles, CA, January 30, 2003.
- 2003.2 "The Haplotype Map Initiative and Whole Genome Association Studies," invited speaker, Contemporary Issues in Human Genetics Seminar Series, University of California, San Diego, La Jolla, CA, February 20, 2003.
- 2003.3 "The Haplotype Map Initiative and Association Studies," invited speaker, NIH Pharmacogenetics Network Annual Meeting, Memphis, TN, March 4, 2003.
- 2003.4 "Complex trait genetics: Insights for hypertension and renal disease," invited speaker, Renal Conference, UCSD School of Medicine continuing medical education Category 1 activity, Veterans Affairs Medical Center, La Jolla, CA, March 18, 2003.
- 2003.5 "Race and Modern Population and Epidemiological Genetics," invited speaker, American Society for Clinical Pharmacy and Therapeutics, Washington, DC, April 4, 2003.
- 2003.6 "The Haplotype Map Initiative and Whole Genome Association Studies," invited speaker, Department of Medicine, University of Medicine and Dentistry of New Jersey, Newark, NJ, April 7, 2003.
- 2003.7 "Methodological Issues in the Genetic Analysis of Complex Traits," invited speaker, McKusick-Nathans Center for Human Genetics, Johns Hopkins University, Baltimore, MD, April 8, 2003
- 2003.8 "Systems Biology and Genomics," invited speaker, GeneGo, Inc. symposium on systems reconstruction, San Diego Supercomputer Center, University of California, San Diego, La Jolla, CA, April 30, 2003
- 2003.9 "Methods for Assessing Cryptic Population Substructure," invited speaker, The Donald W. Reynolds Symposium: Understanding the Complexity of Heart Disease, Stanford University, Stanford, CA, May 19, 2003.
- 2003.10 "Issues in the use of the haplotype map resource," invited speaker, Scripps Research Institute, La Jolla, CA, June 17, 2003.
- 2003.11 "Genetics and Race," invited speaker, Collaboration in the Health and Social Sciences: A Workshop, University of California, San Diego, La Jolla, CA, June 21, 2003.

- 2003.12 “Issues in the Genetic Epidemiologic Analysis of Complex Traits,” invited speaker, University of Texas Southwestern Medical Center, Dallas, TX, July 17, 2003.
- 2003.13 “Issues in Large-Scale Genetic Epidemiology Studies,” invited speaker, The Donald W. Reynolds Cardiovascular Clinical Research Center Training Seminar, Stanford School of Medicine, Stanford, CA July 24, 2003.
- 2003.14 “Analyses of population substructure,” invited speaker, Roche Palo Alto, University of California, San Francisco, October 24, 2003.
- 2003.15 “Using Population Data to Find Genes for Complex Diseases,” invited speaker, American Society of Nephrology Annual Meeting, postgraduate education course, “Clinical Science Course – Genetics Issues in Kidney Disease: Platforms for Investigation.” San Diego, CA, November 13, 2003.
- 2004.1 “Novel Multivariate Analysis Methods for Genomic Analysis,” invited speaker, Mathematical Sciences Research Institute, workshop, “Genetics of Complex Disease.” University of California, Berkeley, February 9, 2004.
- 2004.2 “Issues in Statistical Genetics and Statistical Genomics,” invited speaker, San Diego Supercomputer Center, University of California, San Diego, La Jolla, California, March 22, 2004.
- 2004.3 “Issues and Methodology in Statistical Genomics,” invited speaker, San Diego Supercomputer Center, University of California, San Diego, La Jolla, California, May 10, 2004.
- 2004.4 “Integrated Approaches to Genome-wide Association Studies,” invited speaker, American Diabetes Association 64th Scientific Sessions, Orlando, Florida, June 6, 2004
- 2004.5 “Modern Human Genetic Association Studies: Statistics, Biology, or Both?” grand rounds, University of California, San Diego, Stein Institute for Research on Aging, August 23, 2004
- 2004.6 “Novel Multivariate Analysis Methods for Genomics,” invited speaker, Department of Biostatistics, UCSD, September 6, 2004.
- 2004.7 “Genomics for Dummies,” invited speaker, Advances in Psychopharmacology, San Diego, California, October 8, 2004
- 2005.1 “The Genetics of Complex Disease,” invited speaker, Kawasaki Disease Symposium, San Diego, California, February 18, 2005.
- 2005.2 “Human Genetics and Genomics,” invited speaker, West Coast College of Biological Psychiatry, Pasadena, California, April 1, 2005.
- 2005.3 “Whole Genome and Genetic Background Analysis,” invited speaker, USC symposium on Whole Genome Association Studies (Duncan Thomas, organizer), USC Campus, Los Angeles, California, April 14, 2005.
- 2005.4 “Genetic Analysis in the Consortium on the Genetics of Schizophrenia (COGS),” invited speaker, Biological Psychiatry Meeting, Atlanta, Georgia, May 21.
- 2005.5 “Genomic Profiling,” invited speaker, VIth FAP Symposium, La Jolla, CA, August 24-26.
- 2005.6 “Clinical and Molecular Genomic Profiling for Gene Discovery and Phenotypic Characterization,” invited speaker, Cancer Center, La Jolla, CA, August 24.
- 2005.7 “Clinical and Molecular Genomic Profiling for Gene Discovery and Phenotypic Characterization,” invited speaker, LOCI Seminar, La Jolla, CA, September 7.

- 2005.8 “DNA sequencing and microarrays: An introduction to the field of modern statistical genomics and bioinformatics,” invited speaker, San Diego Chapter of the American Statistical Association, La Jolla, CA, September 10.
- 2005.9 “Clinical and Molecular Genomic Profiling for Gene Discovery and Phenotypic Characterization,” invited speaker, Translational Genomics Research Institute (TGen), Phoenix, AZ, September 19.
- 2005.10 “Clinical and Molecular Genomic Profiling for Gene Discovery and Phenotypic Characterization,” invited speaker, Venter Institute’s Meeting on “genomes, medicine, and the environment.” Hilton Head, Virginia, October 17-19.
- 2005.11 “Some New Directions for Data Analysis,” invited speaker and section organizer, 4th Annual Meeting of the Longevity Consortium, San Diego, CA, November 30.
- 2005.12 “Genomics and Proteomics Discovery Strategies: Integrative Genomics and Quantitative Human Clinical Studies,” Workshop Canadian Consulate for Biotechnology, San Diego, CA, November 30.
- 2005.13 “Clinical and Molecular Genomic Profiling for Gene Discovery and Phenotypic Characterization,” invited speaker, University of Colorado, December 2.
- 2005.14 “Molecular, Clinical, and Population Profiling in Similarity Analysis,” invited speaker, 6th Annual Meeting of the ADHD Molecular Genetics Network, Miami, Florida, November 20, 2005.
- 2006.1 “Multivariate Profiling in Imaging and Genetics” invited speaker, UCI Workshop on Imaging and Genetics, University of California, Irvine, January 16, 2006.
- 2006.2 “Molecular, Clinical, and Population Profiling in Similarity Analysis,” invited speaker, University of Washington, February 2, 2006.
- 2006.3 “Tutorial on Applied and Human Population Genetics,” invited speaker and organizer, Inaugural Symposium on Information Theory and Applications, University of California, San Diego, Feb. 6 – 10, 2006.
- 2006.4 “Genetic, Molecular, and Clinical Profiling via Similarity Analysis,” invited speaker, Ligand Pharmaceutical, La Jolla, CA, February 14, 2006.
- 2006.5 “Genetic Epidemiology and Modern Human Genetics and Genomics,” guest speaker, UCSD T32 Fellows lecture series, San Diego VA Hospital, April 15, 2006.
- 2006.6 “Similarity Analysis: Motivation and Applications,” invited speaker, UCLA psychiatric genetics group, Los Angeles, CA, May 3, 2006.
- 2006.7 “Modern Statistics Genetics Resources,” invited speaker, Annual Meeting of the Society for Biological Psychiatry, Toronto, Canada, May 19, 2006.
- 2006.8 “Computational Science in Modern Human Genomics,” invited speaker, Alliance for Equity in Higher Education, San Diego, June 27, 2006
- 2006.9 “Pharmacogenetics and Pharmacogenomics,” invited speaker, Ligand Pharmaceuticals Board of Directors meeting, San Diego, July 17, 18, 2006.
- 2006.10 “Evolution, Humanity and Genomic Diversity: Deviations vs. Variations,” invited speaker, UCSD Medical Scientist Graduate Training Program Annual Retreat. La Jolla, California, August 27, 2006.
- 2006.11 “New Directions for Genetic Association Studies: Whole Genome and Sequence-Based Analyses,” invited speaker, Venter Institute, Rockville, Maryland, September 22, 2006.

- 2006.12 “A Comparison of Statistical Methods for Genome-Wide Association Studies,” invited speaker, Genetic Epidemiology Research Institute, University of California, Irvine, October 6, 2006.
- 2006.13 “Novel Analysis Methods for Whole Genome Association Studies,” invited speaker, Biostatistics and Bioinformatics Seminar Series, University of California, San Diego, October 18, 2006.
- 2006.14 “New Directions for Genetic Association Studies,” invited speaker, 10th Scientific Advisory Board Meeting, Genome Institute of Singapore, Singapore, October 30 – November 1, 2006.
- 2006.15 “Novel Analysis Methods for Whole Genome Association Studies,” invited speaker, Department of Preventive Medicine Biostatistics Seminar Series, University of Southern California, Los Angeles, November 8, 2006.
- 2006.16 “Novel Approaches to Genetic Analysis,” invited participant, Genome Quebec and Moores UCSD Cancer Center collaborative discussions, La Jolla, November 16, 2006.
- 2006.17 “Similarity Analysis: Applications in Genetics and Genomics,” invited speaker, San Diego Chapter American Statistical Association, University of California, San Diego, November 17, 2006.
- 2007.1 “Multivariate Analysis of Combined Imaging and Genomic Data,” invited speaker, UCI Workshop in Imaging and Genetics, University of Irvine, January 15, 2007.
- 2007.2 “The Analysis of Multivariate Similarity: Theory and Applications,” invited speaker, UCSD Artificial Intelligence Seminar Series. March 5, 2007.

Invited and Pending Presentations

- 2007.3 “Large-scale data analysis in bioinformatics and genomics,” invited speaker, Annual Meeting of the Biological Psychiatry Organization. San Diego, California, May 17, 2007.

PUBLISHED INTERVIEWS

- 1995.1 Sue Goetinck, “Pressure Lookers – Scientists search for genetic causes behind hypertension,” *The Dallas Morning News*, D6-D8, Monday, September 4, 1995
- 1996.1 Virginia Morell, “Manic Depression Findings Spark Polarized Debate,” *Science*, 272:31-32; 1996.
- 1999.1 Malorye A. Branca, Kenneth Rubenstein, “Single Nucleotide Polymorphisms: Commercial and Scientific Prospects,” Cambridge, Cambridge Health Tech Institute, 1999.
- 1999.2 “Applying the Discoveries from the Genome Project,” *Genetic Engineering News*, September 1, 1999, Volume 19 (15)
- 1999.3 “SNPs and SNP Analysis,” Cold Spring Harbor Laboratory WWW interview series, (Jan Witkowski, organizer), October, 1999.
- 2000.1 Leslie Roberts, “SNP Mappers Confront Reality and Find It Daunting,” *Science* 287; 1898-1988; 2000.
- 2000.2 Anthony R. Porcari and Ronald J. Shebuski, “Meeting report of the ASPET Colloquium: Functional genomics,” *Clinical and Experimental Therapeutics*.
- 2002.1 Lucy J. Sannes, “Predictive Pharmacogenomics: Revolutionizing Health Care,” Cambridge: Cambridge HealthTech Institute
- 2002.2 Jean McCann, “Color of skin no way to prescribe drugs,” *Drug Topics*, May 19, 2003

2006.1 Nancy Jenkins, "Movers," Nature 441:904, 2006

COMMITTEE AND ADMINISTRATIVE SERVICE

National

1994	NIA-appointed member, Alzheimer's Research Planning Committee, Bethesda, MD.
1995-2002	Steering committee member: The Family Blood Pressure Program, NIH-NHLBI.
1995	Co-organizer: Symposium on the Biology and Genetics of Complex Traits, The Jackson Laboratory, Bar Harbor, ME, (September 13-17).
1996-2003	NIA-selected working group member: The Genetics of Longevity.
1996-2001	Associate director, Harvard Program for Population Genetics.
1997	NIH-appointed working group member: Research on the Molecular Genetics of Speech and Language.
1997	Organizer: Roche/Jackson Laboratory Symposium on Function Genomics, Roche Biosciences, Palo Alto, CA (October 6).
1998	Organizer: Roche/Jackson Laboratory Symposium on Polymorphisms and Drug Discovery, Roche Discovery, Welwyn, England (May 19).
1998	Organizer: Astra/Jackson Laboratory Workshop on the Mouse in Genetics Research, Sweden (August 30).
1998-2000	Member, scientific advisory committee, "Kidney Development and Cytogenesis," NIH program project grant awarded to Harvard University (Amin Arnaout, PI).
1998-2001	Member, Data Safety and Monitoring Board, The Family Heart Study, NIH-NHLBI
1999	Review team member, Wellcome Trust, Oxford Centre for Gene Function (December)
1998-2001	Advisor on the grant, "The Genetics of Alcoholism," University of Pittsburgh, (5 R01 AA05909, Shirley Hill, PI).
2000	Advisory board member, Harvard Medical School, Department of Medicine, SCOR Grant (Gordon Williams, PI).
2000	Co-chair, association studies advisory board, National Association on Aging
2000-2006	Member of the Committee on Twins Studies, Institute of Medicine, the National Academies
2001	Review panel member, NIDDK, Phoenix, AZ (Clifton Borgardus, P.I.)
2002-2003	Co-director, Committee on Biostatistics and Population Genetics, NIH PharmGKB initiative
2002-Present	Member of the Internal Scientific Review Committee for Opportunity Funds, National Institute on Aging, the Longevity Consortium
2003-2005	External advisory board member, Mt. Sinai School of Medicine, Children's Oncology Group Acute Lymphoblastic Leukemia SPORE grant (William Carroll, P.I.)

- 2003-Present Member of the Food and Nutrition Board, Institute of Medicine, the National Academies
- 2003 NIH-appointed review panel member, Laboratory of Epidemiology, Demography, and Biometry
- 2003-Present Member of the Scientific Advisory Board, NIH-funded consortium grant on ADHD
- 2003-2005 External steering committee member, Johns Hopkins Pepper Center in Aging Research.
- 2004 Reviewer, *Identifying and Assessing Unintended Effects of Genetically Engineered Foods on Human Health*; Division on Earth and Life Studies, National Research Council
- 2004 Reviewer, *Safety of Genetically Engineered Foods: Approaches to Assessing Unintended Health Effects*. Institute of Medicine, Food and Nutrition Board
- 2004 Member, NCI-sponsored “Susceptibility and Resistance to Cancer Think Tank.”
- 2005-Present Organizer and Chair, “Nutrigenomics and Beyond: Informing the Future.” National Academy of Sciences, Washington, D.C., June 1-2.

State of California

- 2006- Present Committee Member, Joint Committee on interactions between the state judiciary and the University of California system.

University of California, San Diego

- 2001-Present Co-organizer, Contemporary Issues in Human Genetics seminar series
- 2001-2006 Member, Department of Psychiatry, Faculty Ad Hoc Review Committees (3)
- 2002-2005 Member, School of Medicine, Conflict of Interest Management Committee
- 2002-Present Scientific review board member, Center for Medicinal Cannabis Research
- 2003-Present Member, UCSD Health Sciences Faculty Research Council
- 2003-2003 Member, San Diego Supercomputer Center Biosciences Advisory Council
- 2004-2007 Member, Department of Psychiatry K Award Committee
- 2004-2006 Member, Campus-wide Shared Resources Committee
- 2004-2005 Member, Burroughs-Wellcome Fellowship Program Committee
- 2004-2005 Member, Systems Engineering and Medicine Committee
- 2005-2007 Organizer, Interactions with the Venter Institute of Rockville, Maryland.
- 2005-2007 Chairman, School of Medicine, Conflict of Interest Management Committee
- 2006 Co-organizer, “Information Theory and Applications”, Inaugural Workshop,
- 2006-Present Member, Working Group on the Interface of Medicine and Engineering (Dean’s office)
- 2006-Present Member, CTSA Steering Committee

2006-2007	Medical Scientist Training Program Admissions Committee Member
2006-Present	Member, Senate-Administration Task Force on Multidisciplinary Joint Faculty Appointments
2006	Steering committee, Cancer Center Retreat
2007-Present	Member, San Diego Supercomputer Center Oversight and Executive Committee

University of California, Irvine

2004-Present	Member, Genetic Epidemiology Research Institute Advisory Board
--------------	--

Case Western Reserve University

1994-1995	Seminar series co-organizer, Department of Genetics
-----------	---

BIBLIOGRAPHY

Completed Publications in Scientific Journals

Peer Reviewed Publications

1. Hinderliter AL, Fitzpatrick MA, Schork NJ, Julius S: Research utility of noninvasive methods for measurement of cardiac output. *Clin Pharmacol Ther* 1987; 41:419-25.
2. Weder AB, Sekkarie MA, Takiyyuddin M, Schork NJ, Julius S: Antihypertensive and hypotensive effects of atrial natriuretic factor in men. *Hypertension* 1987; 10:582-9.
3. Johnson EH, Schork NJ, Spielberger CD: Emotional and familial determinants of elevated blood pressure in black and white adolescent females. *J Psychosom Res* 1987; 31:731-41.
4. Egan BM, Panis R, Hinderliter A, Schork N, Julius S: Mechanism of increased alpha-adrenergic vasoconstriction in human essential hypertension. *J Clin Invest* 1987; 80:812-7.
5. Schork, NJ and Schork MA: Skewness and mixtures of normal distributions. *Communications in Statistics - Theory and Methods* 1988; 17:3951-69.
6. Egan BM, Schork N, Panis R, Hinderliter A: Vascular structure enhances regional resistance responses in mild essential hypertension. *J Hypertens* 1988; 6:41-8.
7. Egan BM, Conlon ME, Campbell R, Schork N, Zweifler A, Julius S: Effects of ketanserin on blood pressure and platelet aggregation in elderly men with mild hypertension. *Am J Hypertens* 1988; 1:324S-30S.
8. Julius S, Schork N and Schork A: Sympathetic hyperactivity in early stages of hypertension: the Ann Arbor data set. *J Cardiovasc Pharmacol* 1988; 12:S121-9.
9. Weder AB and Schork NJ: Mixture analysis of erythrocyte lithium-sodium countertransport and blood pressure. *Hypertension* 1988; 13:145-50.
10. Gupta MA, Gupta AK, Kirkby S, Weiner HK, Mace TM, Schork NJ, Johnson EH, Ellis CN, Voorhees JJ: Pruritus in psoriasis: A prospective study of some psychiatric and dermatologic correlates. *Arch Dermatol* 1988; 124:1052-7.
11. Gupta MA, Gupta AK, Kirkby S, Schork NJ, Gorr SK, Ellis CN, Voorhees JJ: A psychocutaneous profile of psoriasis patients who are stress reactors: A study of 127 patients. *Gen Hosp Psychiatry* 1989; 11:166-73.

12. Schork NJ and Schork MA: Testing separate families of segregation hypotheses: bootstrap methods. *Am J Hum Genet* 1989; 45:803-13.
13. Gupta MA, Gupta AK, Kirkby S, Schork NJ, Weiner HK, Ellis CN, Voorhees JJ: Pruritus associated with nocturnal awakenings: organic or psychogenic? *J Am Acad Dermatol* 1989; 21:479-84.
14. Egan BM, Schork NJ, Weder AB: Regional hemodynamic abnormalities in overweight men: focus on alpha-adrenergic vascular responses. *Am J Hypertens* 1989; 2:428-34.
15. Kjeldsen SE, Schork NJ, Leren P, Eide IK: Arterial plasma norepinephrine correlates to blood pressure in middle-aged men with sustained essential hypertension. *Am Heart J* 1989; 118:775-81.
16. Mejia AD, Egan BM, Schork NJ, Zweifler AJ: Artefacts in measurement of blood pressure and lack of target organ involvement in the assessment of patients with treatment-resistant hypertension. *Ann Intern Med* 1990; 112:270-7.
17. Schork NJ, Weder AB, Schork MA, Bassett DR, Julius S: Disease entities, mixed multi-normal distributions, and the role of the hyperkinetic state in the pathogenesis of hypertension. *Stat Med* 1990; 9:301-14.
18. Schork NJ, Weder AB, Schork MA: On the asymmetry of biological frequency distributions. *Genet Epidemiol* 1990; 7:427-46.
19. Mejia AD, Julius S, Jones KA, Schork NJ, Kneisley J: The Tecumseh Blood Pressure Study: normative data on blood pressure self-determination. *Arch Intern Med* 1990; 150:1209-13.
20. Kneisley J, Schork N, Julius S: Predictors of blood pressure and hypertension in Tecumseh, Michigan. *Clin Exp Hypertens A* 1990; 12:693-708.
21. Julius S, Jamerson K, Mejia A, Krause L, Schork N, Jones K: The association of borderline hypertension with target organ changes and higher coronary risk: Tecumseh Blood Pressure Study. *JAMA* 1990; 264:354-8.
22. Gupta MA, Gupta AK, Schork NJ, Ellis CN, Voorhees JJ: The aging face: A psychocutaneous perspective. *J Dermatol Surg Oncol* 1990; 16:902-4.
23. Gupta MA, Gupta AK, Schork NJ, Ellis CN, Voorhees JJ: Psychiatric aspects of the treatment of mild to moderate acne: some preliminary observations. *Int J Dermatol* 1990; 29:719-21.
24. Julius S, Mejia A, Jones K, Krause L, Schork N, van de Ven C, Johnson E, Petrin J, Sekkarie MA, Kjeldsen SE, Schmouder R, Gupta R, Ferraro J, Nazzaro P and Weissfeld J: "White coat" versus "sustained" borderline hypertension in Tecumseh, Michigan. *Hypertension* 1990; 16:617-23.
25. Schork NJ and Schork MA: Histograms: multimodal or poorly constructed? (Letter to the Editor), *Am J Hum Genet* 1990; 46:396-7.
26. Julius S, Jones K, Schork N, Johnson E, Krause L, Nazzaro P, Zemva A: Independence of pressure reactivity from blood pressure levels in Tecumseh, Michigan. *Hypertension* 1991; 17(Supplement III):12-21.
27. Schork NJ: Efficient computation of patterned covariance matrix mixed models in quantitative segregation analysis. *Genet Epidemiol* 1991; 8:29-46.
28. Weder AB, Schork NJ, Krause L, Julius S: Red blood cell lithium-sodium countertransport in the Tecumseh blood pressure study. *Hypertension* 1991; 17:652-60.
29. Julius S, Krause L, Schork N, Mejia AD, Jones KA, van de Ven C, Johnson EH, Sekkarie MA, Kjeldsen SE, Petrin J, Schmouder R, Gupta R, Ferraro J, Nazzaro P and Weissfeld J: Hyperkinetic borderline hypertension in Tecumseh, Michigan. *J Hypertens* 1991; 9:77-84.

30. Weder AB, Schork NJ, Julius S: Linkage of MN locus and erythrocyte lithium-sodium countertransport in Tecumseh, Michigan. *Hypertension* 1991; 17:977-81.
31. Gupta MA, Goldfarb MT, Schork NJ, Weiss JS, Gupta AK, Ellis CN and Voorhees JJ: Treatment of mildly to moderately photoaged skin with topical tretinoin has a favorable psychosocial effect: a prospective study. *J Am Acad Dermatol* 1991; 24:780-1.
32. Schork NJ: Extended pedigree patterned covariance matrix mixed models for quantitative phenotype analysis. *Genet Epidemiol* 1992; 9:73-86.
33. Schork NJ: Detection of genetic heterogeneity for complex quantitative phenotypes. *Genet Epidemiol* 1992; 9:207-23.
34. Julius S, Jamerson K, Gudbrandsson T, Schork N: White coat hypertension: a follow-up. *Clin Exp Hypertens A*. 1992; 14:45-53.
35. Gudbrandsson T, Julius S, Krause L, Jamerson K, Randall OS, Schork N, Weder A: Correlates of the estimated arterial compliance in the population of Tecumseh, Michigan. *Blood Press* 1992; 1:27-34.
36. Jamerson KA, Schork NJ, Julius S: Effect of home blood pressure and gender on estimates of the familial aggregation of blood pressure: The Tecumseh Blood Pressure Study. *Hypertension* 1992; 20:314-8.
37. Bretz WA, Krahn DD, Drury M, Schork N, Loesche WJ: Effects of fluoxetine on the oral environment of bulimics. *Oral Microbiol Immunol* 1993; 8:62-4.
38. Gupta MA, Schork NJ, and Dhaliwal JS: Stature, drive for thinness and body dissatisfaction: a study of males and females from a non clinical sample. *Can J Psychiatry* 1993; 38:59-61.
39. Gupta MA, Schork NJ, Gupta AK, Kirkby S, Ellis CN: Suicidal ideation in psoriasis. *Int J Dermatol* 1993; 32:188-90.
40. Gupta MA, Schork NJ, Gupta AK and Ellis CN: Alcohol intake and treatment responsiveness of psoriasis: a prospective study. *J Am Acad Dermatol* 1993; 28:730-2.
41. Schork NJ, Schork MA: The relative efficiency and power of small-pedigree studies of the heritability of a quantitative trait. *Hum Hered* 1993; 43: 1-11.
42. Schork NJ: The design and use of variance component models in the analysis of human quantitative pedigree data. *Biometrical J* 1993; 35:387-405.
43. Schork NJ and Schork MA: The effect of pairwise correlated observations on likelihood ratio tests for the difference between two means. *Communications in Statistics—Theory and Methods* 1993; 22:2609-18.
44. Bretz WA, Eklund SA, Radicchi R, Schork MA, Schork N, Schottenfeld D, Lopatin DE, Loesche WJ: The use of a rapid enzymatic assay in the field for the detection of infections associated with adult periodontitis. *J Public Health Dent* 1993; 53:235-40.
45. Topol EJ, Ellis SG, Cosgrove DM, Bates ER, Muller DW, Schork NJ, Schork MA, Loop FD: Analysis of coronary angioplasty practice in the United States with an insurance-claims data base. *Circulation* 1993; 87: 1489-97.
46. Schork NJ: Combining Monte Carlo and Cox tests of non-nested hypotheses. *Communications in Statistics—Simulation and Computation* 1993; 22:939-54.
47. Schork NJ, Boehnke M, Terwilliger J, Ott J: Two-trait-locus linkage analysis: a powerful strategy for mapping complex genetic traits. *Am J Hum Genet* 1993; 53:1127-36.

48. Schork NJ: Extended multipoint identity-by-descent analysis of human quantitative traits: efficiency, power, and modeling considerations. *Am J Hum Genet* 1993; 53:1306-19.
49. Gupta MA, Schork NJ: Aging-related concerns and body image: possible future implications for eating disorders. *Int J Eat Disord* 1993; 14:481-6.
50. Schork NJ, Guo SW: Pedigree models for complex human traits involving the mitochondrial genome. *Am J Hum Genet* 1993; 53:1320-37.
51. Gupta MA, Schork NJ, Ellis CN: Psychosocial correlates of the treatment of photodamaged skin with topical retinoic acid: a prospective controlled study. *J Am Acad Dermatol* 1994; 30:969-72.
52. Schork NJ, Weder AB, Trevisan M, Laurenzi M: The contribution of pleiotropy to blood pressure and body-mass index variation: the Gubbio Study. *Am J Hum Genet* 1994; 54:361-73.
53. Weder AB, Schork NJ: Adaptation, Allometry, and Hypertension. *Hypertension* 1994; 24:145-56.
54. Gupta MA, Gupta AK, Schork NJ, Ellis CN: Depression modulates pruritus perception: a study of pruritus in psoriasis, atopic dermatitis, and chronic idiopathic urticaria. *Psychosom Med* 1994; 56:36-40.
55. Gudbrandsson T, Julius S, Jamerson K, Smith S, Krause L, Schork N: Recreational exercise and cardiovascular status in the rural community of Tecumseh, Michigan. *Blood Pres* 1994; 3:178-84.
56. Schork NJ, Jokelainen P, Grant EJ, Schork MA, Weder AB: Relationship of growth and blood pressure in inbred rats. *Am J Psychiol* 1994; 266:R702-8.
57. Lander ES, Schork NJ: Genetic dissection of complex traits. *Science* 1994; 265:2037-48.
58. Marcus R, Krause L, Weder AB, Dominguez-Meja A, Schork NJ and Julius S: Sex-specific determinants of increased left ventricular mass in the Tecumseh Blood Pressure Study. *Circulation* 1994; 90:928-36.
59. Schork NJ, Boehnke M, Terwilliger JD, Ott J. Reply to Sham et al. (Letter to the Editor), *Am J Hum Genet* 1994; 5:856-8.
60. Schork NJ: Sampling guidelines for testing secondary attack rates associated with short-latency infectious diseases. *Stat Med* 1994; 13:1563-73.
61. Smith S, Julius S, Jamerson K, Amerena J, Schork N: Hematocrit levels and physiologic factors in relationship to cardiovascular risk in Tecumseh, Michigan. *J Hypertens* 1994; 12:455-62.
62. Gupta MA, Gupta AK, Schork NJ: Psychosomatic study of self-excoriative behavior among male acne patients: preliminary observations. *Int J Dermatol* 1994; 33:846-8.
63. Gupta MA, Gupta AK, Schork NJ, Watteel GN. Perceived touch deprivation and body image: some observations among eating disordered and non-clinical subjects. *J Psychosom Res* 1995; 39:459-64.
64. Gupta MA, Schork NJ. Touch deprivation has an adverse effect on body image: some preliminary observations. *Int J Eat Disord* 1995; 17:185-9.
65. Davidson AO, Schork NJ, Jacques BC, Kelman AW, Sutcliffe RG, Reid JL, Dominiczak AF. Blood pressure in genetically hypertensive rats: Influence of the Y chromosome. *Hypertension* 1995; 26:452-9.
66. McAleer MA, Reifsnnyder P, Palmer SM, Prochazka M, Love JM, Copeman JB, Powell EE, Rodrigues NR, Prins JB, Serreze DV, DeLarato NH, Wicker LS, Peterson LB, Schork NJ, Todd JA, Leiter EH. Crosses of NOD mice with the related NON strain: A polygenic model for type I diabetes. *Diabetes* 1995; 44:1186-95.
67. Thibbonier M, Schork NJ. The genetics of hypertension. *Curr Opin Genet Dev* 1995; 5:362-70.

68. Schork NJ, Krieger JE, Trollet MR, Franchini KG, Koike G, Krieger EM, Lander ES, Dzau VJ, Jacob HJ. A biometrical genome search in rats reveals the multigenic basis of blood pressure variation. *Genome Res* 1995; 5:164-72.
69. Letts VA, Schork NJ, Copp AJ, Bernfield M, Frankel WN. A curly-tail modifier locus, *mct1*, on mouse chromosome 17. *Genomics* 1995; 29:719-24.
70. Ghosh S, Schork NJ: Genetic analysis of NIDDM. The study of quantitative traits. *Diabetes* 1996; 45:1-14.
71. Schork NJ, Allison DB, Thiel B: Mixture Distributions in Human Genetics Research. *Stat Methods Med Res* 1996; 5:155-78.
72. Wolf NG, Abdul-Karim FW, Schork NJ, Schwartz S: Origins of heterogenous ovarian carcinomas: A molecular cytogenetic analysis of histologically benign, low malignant potential, and fully malignant components. *Am J Pathol* 1996; 149: 511-20.
73. Witte JS, Elston RC, Schork NJ: Genetic dissection of complex traits. (Letter to the Editor). *Nat Genet* 1996; 12:355-6.
74. Schork NJ, Weder AB: The use of genetic information in large-scale clinical trials: Applications to Alzheimer research. *Alzheimer Dis Assoc Disord* 1996; 10 Suppl 1:22-6.
75. Krause BR, Schork NJ, Kieft KA, Smith MP, Maciejko JJ: High correlation but lack of agreement between direct high-performance gel chromatography analysis and conventional indirect methods for determining lipoprotein cholesterol. *Clin Chem* 1996; 42:1996-2001.
76. Eicher EM, Washburn LL, Schork NJ, Lee BK, Shown EP, Xu X, Dredge RD, Pringle MJ, Page DC: Sex-determining genes on mouse autosomes identified by linkage analysis of C57BL/6J-YPOS sex reversal. *Nat Genet* 1996; 14:206-9.
77. Schork NJ, Nath SP, Lindpaintner K, Jacob HJ: Extensions to quantitative trait locus mapping in experimental organisms. *Hypertension* 1996; 28:1104-11.
78. Gupta MA, Gupta AK, Schork NJ. Psychological factors affecting self-excoriative behavior in women with mild-to-moderate facial acne vulgaris. *Psychosomatics* 1996; 37:127-30.
79. Frankel WN, Schork NJ: Who's afraid of epistasis? *Nat Genet* 1996; 14:371-3.
80. Schork NJ: Genetically complex cardiovascular traits. Origins, problems, and potential solutions. *Hypertension* 1997; 29:145-9.
81. Salvador SL, Grisi MF, Romanelli RG, Silva Netto CR, Schork NJ, Bretz WA: Similarities of periodontal clinical and microbiological parameters in mother-child pairs. *Braz Dent J* 1997; 8:99-104.
82. Cassidy SB, Forsythe M, Heeger S, Nicholls RD, Schork NJ, Benn P, Schwartz S: Comparison of phenotype between patients with Prader-Willi Syndrome due to deletion 15q and uniparental disomy 15. *Am J Med Genet* 1997; 68:433-40.
83. Ferraro TN, Golden GT, Smith GG, Schork NJ, St Jean P, Ballas C, Choi H, Berrettini WH: Mapping murine loci for seizure response to kainic acid. *Mamm Genome* 1997; 8:200-8.
84. Strohl KP, Thomas AJ, St Jean P, Schlenker EH, Koletsky RJ, Schork NJ: Ventilation and metabolism among rat strains. *J Appl Physiol* 1997; 82:317-23.
85. Beebe AM, Mauze S, Schork NJ, Coffman RL: Serial backcross mapping of multiple loci associated with resistance to *Leishmania major* in mice. *Immunity* 1997; 6:551-7.

86. Allison DB, Schork NJ: Selected methodological issues in meiotic mapping of obesity genes in humans: issues of power and efficiency. *Behav Genet* 1997; 27:401-21.
87. Schork NJ, Xu X: Sibpairs vs. Pedigrees: what are the advantages? *Diabetes Reviews* 1997; 5:116-22.
88. Xu X, Schork NJ: Linking genes and environmental exposure: why China presents special opportunities. *Cancer Causes Control* 1997; 8:518-23.
89. Schork NJ: Genetics of complex disease: approaches, problems, and solutions. *Am J Respir Crit Care Med* 1997; 156:S103-9.
90. Allison DB, Heo M, Schork NJ, Wong SL, Elston RC. Extreme selection strategies in gene mapping studies of oligogenic quantitative traits do not always increase power. *Hum Hered* 1998; 48:97-107.
91. Schork NJ, Schork CM: Issues and Strategies in the Genetic Analysis of Alcoholism and Related Addictive Behaviors. *Alcohol* 1998; 16:71-83.
92. Schork NJ, Thiel B, St Jean P: Linkage analysis, kinship, and the short-term evolution of chromosomes. *J Exp Zool* 1998; 282:133-49.
93. Schork NJ, Cardon LR, Xu X. The future of genetic epidemiology. *Trends Genet* 1998; 14:266-72.
94. Allison DB, Thiel B, St Jean P, Elston RC, Infante MC, Schork NJ: Multiple phenotype modeling in gene-mapping studies of quantitative traits: ower advantages. *Am J Hum Genet* 1998; 63:1190-201.
95. Herrera VL, Xie HX, Lopez, LV, Schork NJ, Ruiz-Opazo N: The alpha1 Na,K-ATPase gene is a susceptibility hypertension gene in the Dahl salt-sensitive HSD rat. *J Clin Invest* 1998; 102:1102-11.
96. Ginns EI, St. Jean P, Philibert RA, Galdzicka M, Damschroder-Williams P, Long R, Ingraham L, Dalwaldi H, Murray M, Ehlert M, Paul S, Remortel B, Patel A, Anderson M, Dymarskaia I, Martin B, Stubblefield B, Falls K, Keith T, Fann C, Thiel B, Lacy LG, Allen C, Hostetter AM, Ott J, Elston RC, Schork NJ, Egeland JA, Paul SM. A genome-wide search for chromosomal loci linked to mental health wellness in relatives at high risk for bipolar disorder among the old order Amish. *Proceedings of the National Academy of Sciences* 1998; 95:15531-36.
97. Faust, C, Lawson KA, Schork NJ, Thiel B, Magnuson T. The Polycomb-group gene *eed* is required for normal morphogenetic movements during gastrulation in the mouse embryo. *Development* 1998; 125: 4495-506.
98. Anderson NH, Devlin AM, Graham D, Morton JJ, Hamilton CA, Reid JL, Schork NJ, Dominiczak AF: Telemetry for cardiovascular monitoring in a pharmacological study: new approaches to data analysis. *Hypertension* 1999; 33:248-55
99. Bix M, Wang ZE, Thiel B, Schork NJ, Locksley RM: Genetic regulation of commitment to interleukin 4 production by a CD4(+) T cell-intrinsic mechanism. *J Exp Med* 1998; 188:2289-99.
100. Xu X, Rogus JJ, Terwedow HA, Yang J, Wang Z, Chen C, Niu T, Wang B, Xu H, Weiss S, Schork NJ, Fang Z. An Extreme-Sib-Pair Genome Scan for Genes Regulating Blood Pressure. *Am J Hum Genet* 1999; 64:1694-701.
101. Schork NJ: Review of 'Angiotensin Genotype, Sodium Reduction, Weight Loss, and Prevention of Hypertension: Trials of Hypertension Prevention, Phase II'. *Curr Hypertens Rep* 1999; 1:13-14
102. Niu T, Chen C, Cordell H, Yang J, Wang B, Wang Z, Fang Z, Schork NJ, Rosen CJ, Xu X. A Genome-Wide Scan for Loci Linked to Forearm Bone Mineral Density. *Hum Genet* 1994; 104:226-33.

103. Niu T, Chen C, Yang J, Wang B, Wang Z, Schork N, Fang Z, Xu X: Blood pressure and the T174M and M235T polymorphisms of the angiotensinogen gene. *Ann Epidemiol* 1999; 9:245-53.
104. Allison DB, Neale MC, Zannolli R, Schork NJ, Amos CI, Blangero J: Testing the robustness of the likelihood-ratio test in a variance-component quantitative-trait-loci mapping procedure. *Am J Hum Genet* 1999; 65:531-44.
105. Drazen JM, Yandava CN, Dube L, Szczerback N, Hippensteel R, Pillari A, Israel E, Schork NJ, Silverman ES, Katz DA, Drajesk J: Pharmacogenetic association between ALOX5 promoter genotype and the response to anti-asthma treatment. *Nat Genet* 1999; 22:168-70.
106. Ferraro TN, Golden GT, Smith GG, St Jean P, Schork NJ, Mulholland N, Ballas C, Schill J, Buono RJ, Berrettini WH: Mapping loci for pentylenetetrazol-Induced seizure susceptibility in mice. *J Neurosci* 1999; 19:6733-39.
107. Xu X, Yang J, Rogus J, Chen C, Schork NJ, Xu X. Mapping of a blood pressure quantitative trait locus to chromosome 15q in a Chinese population. *Hum Mol Genet* 1999; 8:2551-5
108. MacGregor AJ, Snieder H, Schork NJ, Spector TD: Twins – Novel uses to study complex traits and genetic diseases. *Trends Genet* 2000; 16:131-4.
109. Bihain BE, Schork N, Bougueleret L, Blumenfeld M, Chumakov I, Yen F, Cohen D, *Génomique promesses et réalités. Médecine Sciences*, 2000;16(1): 17-21
110. Barlassina C, Schork NJ, Manunta P, Citterio L, Sciarrone M, Lanella G, Bianchi G, and Cusi D. Synergistic effect of alpha-adducin and ACE genes causes blood pressure changes with body sodium and volume expansion. *Kidney Int* 2000; 57:1083-90
111. Schork NJ: Xenobiotics, Dietary Interventions, and Genetically-Mediated Therapies. *Curr Hypertens Rep* 2000; 2:11-12.
112. Jeanclos E, Schork NJ, Kyvik KO, Kimura M, Skurnick JH, Aviv A: Telomere length inversely correlates with pulse pressure and is highly familial. *Hypertension* 2000; 36:195-200.
113. Fallin D, Schork NJ: Accuracy of Haplotype Frequency Estimation for Biallelic Loci, via the Expectation-Maximization Algorithm for Unphased Diploid Genotype Data. *Am J Hum Genet* 2000; 67:947-59.
114. Lanchbury JS, Schork NJ: Peaks and troughs in linkage mapping for the rheumatic diseases. *Rheumatology* 2000; 39:453-6.
115. Cowley AW Jr, Stoll M, Greene AS, Kaldunski ML, Roman RJ, Tonellato PJ, Schork NJ, Dumas P, and Jacob HJ: Genetically defined risk of salt-sensitivity in an intercross of Brown Norway and Dahl S rats. *Physiol Genomics* 2000; 2:107-15.
116. Schork NJ, Fallin D, Lanchbury JS: Single nucleotide polymorphisms and the future of genetic epidemiology. *Clin Genet* 2000; 58:250-64.
117. Schork NJ, Nath SK, Fallin D, Chakravarti A: Linkage Disequilibrium Analysis of Bi-allelic DNA Markers, Human Quantitative Trait Loci, and Threshold-Defined Cases and Controls. *Am J Hum Genet* 2000; 67:1208-18.
118. LeStunff C, Fallin D, Schork NJ, Bougnères P: The insulin gene VNTR is associated with fasting insulin levels and development of juvenile obesity. *Nat Genet* 2000; 26:444-6.
119. Schork NJ, Chakravarti A, Thiel B, Fornage M, Jacob HJ, Cai R, Rotimi CN, Cooper RS, Weder AB: Lack of Association Between a Biallelic Polymorphism in the Adducin Gene and Blood Pressure in Whites and African Americans. *Am J Hypertens* 2000; 13:693-8.

120. Nair RP, Stuart P, Henseler T, Jenisch S, Chia NV, Westphal E, Schork NJ, Kim J, Lim HW, Christophers E, Voorhees JJ, Elder JT: Localization of Psoriasis-Susceptibility Locus PSORS1 to a 60-kb Interval Telomeric to HLA-C. *Am J. Hum Genet* 2000; 66:1833-44.
121. LeStunff C, Le Bihan C, Schork NJ, Bougnères P: A common promoter variant of the leptin gene is associated with changes in the relationship between serum leptin and fat mass in obese girls. *Diabetes* 2000; 49:2196-200.
122. Paigen B, Schork NJ, Svenson KL, Cheah YC, Mu JL, Lammert F, Wang DQ, Bouchard G, Carey MC: Quantitative trait loci mapping for cholesterol gallstones in AKR/J and C57L/J strains of mice. *Physiol Genomics* 2000; 4:59-65.
123. Hadley EC, Albers SM, Bailey-Wilson J, Baron J, Cawthon R, Christian JC, Corder EH, Franceschi C, Kestenbaum B, Kruglyak L, Lauderdale DS, Lubitz J, Martin GM, McClearn GE, McGue M, Miles T, Mineau G, Ouellette G, Pedersen NL, Preston SH, Page WF, Province M, Schachter F, Schork NJ, Vaupel JW, Vijg J, Wallace R, Wang E, Wijsman EM: Genetic Epidemiologic Studies on Age-Specified Traits. *Am J Epidemiol* 2000; 152:1003-8
124. Fallin D, Cohen A, Essioux L, Chumakov I, Blumenfeld M, Cohen D, Schork NJ: Genetic Analysis of Case/Control Data Using Estimated Haplotype Frequencies: Application to APOE Locus Variation and Alzheimer's disease. *Genome Res* 2001; 11:143-51.
125. Stoll M, Cowley AW Jr., Tonellato PJ, Greene AS, Kaldunski ML, Roman RJ, Dumas P, Schork NJ, Wang Z, Jacob HJ: A Genomic-Systems Biology Map for Cardiovascular Function. *Science* 2001; 294:1723-26.
126. Nath SK, Chen CH, Schork NJ: Two-trait-locus linkage analyses of asthma susceptibility. *Genet Epidemiol* 2001; 21 Suppl: S278-83.
127. Schork NJ: Genome Partitioning and Whole-Genome Analysis. *Adv Genet* 2001 42:299-322
128. Schork NJ, Fallin D, Thiel B, Xu X, Broeckel U, Jacob HJ, and Cohen D: The Future of Genetic Case-Control Studies. *Adv Genet* 2001; 42:191-212
129. Morrison SJ, Qian D, Jerabek L, Thiel BA, Park IK, Ford PS, Kiel MJ, Schork NJ, Weissman IL, Clarke MF: A genetic determinant that specifically regulates the frequency of hematopoietic stem cells. *J Immunol* 2002; 168: 635-42.
130. Shaw SH, Hutchison D, Saiz R, Abel K, Delisi LE, Schork NJ, Sherrington R: Evaluation of linkage disequilibrium between chromosome 22q11 single nucleotide polymorphisms in a large outbred population. *Am J Med Genet* 2002; 114:205-13.
131. Nath SK, Chakravarti A, Chen CH, Cooper R, Weder A, Schork NJ: Segregation analysis of blood pressure and body mass index in a rural US community. *Hum Biol* 2002; 74:11-23.
132. Hall MA, Norman PJ, Thiel B, Tiwari H, Peiffer A, Vaughan RW, Prescott S, Leppert M, Schork NJ, Lanchbury JS: Quantitative-trait loci on chromosomes 1, 2, 3, 4, 8, 9, 11, 12, and 18 control variation in levels of T and B lymphocyte subpopulations. *Am J Hum Genet* 2002; 70:1172-82.
133. Schork NJ: Power Calculations for Genetic Association Studies Using Estimated Probability Distributions. *Am J. Hum Genet* 2002; 70:1480-9.
134. The Family Blood Pressure Program Investigators. Multi-Center Genetic Study of Hypertension: the Family Blood Pressure Program (FBPP). *Hypertension* 2002; 39:3-9.
135. Larribe F, Lessard S, Schork NJ: Gene mapping via the ancestral recombination graph. *Theor Popul Biol* 2002; 62:215-29.

136. Schork NJ, Gardner JP, Zhang L, Fallin D, Thiel B, Jakubowski H, Aviv A: Genomic association/linkage of sodium lithium countertransport in CEPH pedigrees. *Hypertension* 2002; 40:619-28.
137. Kotchen TA, Broeckel U, Grim CE, Hamet P, Jacob H, Kaldunski ML, Kotchen JM, Schork NJ, Tonellato PJ, Cowley AW Jr.: Identification of hypertension-related QTLs in African American sib pairs. *Hypertension* 2002; 40:634-9.
138. Ralph Knöll, Masahiko Hoshijima, Hal M.Hoffman, Veronika Person, Ilka Lorenzen-Schmidt, Marie-Louise Bang, Takeharu Hayashi, Nobuyuki Shiga, Hideo Yasukawa, Hideo Yasukawa, Wolfgang Schaper, William McKenna, Mitsuhiro Yokoyama, Nicholas J Schork, Jeffrey H Omens, Andrew D McCulloch, Akinori Kimura, Carol C. Gregorio, Wolfgang Poller, Jutta Schaper, Heinz P Schultheiss, and Kenneth R Chien: The Cardiac Mechanical Stretch Sensor Machinery Involves a Z Disc Complex that is Defective in a Subset of Human Dilated Cardiomyopathy. *Cell* 2002; 111:943-55.
139. Lee A, Rana BK, Schiffer HH, Schork NJ, Brann MR, Insel PA, Weiner DM: Distribution analysis of nonsynonymous polymorphisms within the G-protein-coupled receptor gene family. *Genomics* 2003; 81:245-8.
140. Thiel BA, Chakravarti A, Cooper RS, Luke A, Lewis S, Lynn A, Tiwari H, Schork NJ, Weder AB: A genome-wide linkage analysis investigating the determinants of blood pressure in Whites and African Americans. *Am J Hypertens* 2003; 16:151-3.
141. Jirout M, Krenova D, Kren V, Breen L, Pravenec M, Schork NJ, Printz MP: A New Framework Marker- Based Linkage Map and SDPs for the Rat HXB/BXH Strain Set. *Mamm Genome* 2003; 14(8):537-46
142. Broeckel U, Schork NJ: Identifying Genes and Genetic Variation Underlying Human Diseases and Complex Phenotypes via Recombination Mapping. *J Physiol.* 2004 Jan 1;554:40-5
143. Gold DA, Baek SH, Schork NJ, Rose DW, Larsen DD, Sachs BD, Rosenfeld MG, Hamilton BA: RORalpha coordinates reciprocal signaling in cerebellar development through sonic hedgehog and calcium-dependent pathways. *Nuron* 2003 Dec 18, 40(6):1119-31
144. Schork NJ, Greenwood TA: Inherent bias toward the null hypothesis in conventional multipoint nonparametric linkage analysis. *Am J Human Genet* 2004 Feb, 74(2):306-16
145. Conti LH, Jirout M, Breen L, Vanella JJ, Schork NJ, Printz MP: Identificaion of quantitative trait Loci for anxiety and locomotion phenotypes in rat recombinant inbred strains. *Behav Genet* 2004 Jan, 34(1):93-103
146. Wen G, Mahata SK, Cadman P, Mahata M, Ghosh S, Mahapatra NR, Rao F, Stridsberg M, Smith DW, Mahboubi P, Schork NJ, O'Connor DT, Hamilton BA: Both rare and common polymorphisms contribute functional variation at CHGA, a regulator of catecholamine physiology. *Am J Hum Genet* 2004 Feb, 74(2):197-207
147. Fries RS, Mahboubi P, Mahapatra R, Mahata SK, Schork NJ, Schmid-Schoenbein GW, O'Connor DT: Neuroendocrine Transcriptome in Genetic Hypertension: Multiple Changes in Diverse Adrenal Physiological Systems. *Hypertension* 2004 Jun;43(6):1301-11
148. Greenwood TA, Cadman PE, Stridsberg M, Nguyen S, Taupenot L, Schork NJ, O'Connor DT: Genome-wide linkage analysis of chromogranin B expression in the CEPH pedigrees: Implications for exocytotic sympathochromaffin secretion in humans. *Physiol Genomics* 2004 Jun 17;18(1):119-27
149. Nievergelt CM, Smith DW, Kohlenberg JB, Schork NJ: Large-scale integration of human genetic and physical maps. *Genome Res* 2004 Jun;14(6):1199-205
150. Greenwood TA, Rana BK, Schork NJ: Human haplotype block sizes are negatively correlated with recombination rates. *Genome Res.* 2004 July;14(7):1358-61

151. Pascal Le Corre, Robert J. Parmer, Mala T. Kailasam, Brian P. Kennedy, Todd Skaar, Herbert Ho, Roger Leverage, Douglas W. Smith, Michael G. Ziegler, Paul A. Insel, Nicholas J. Schork, David Flockhart, and Daniel T. O'Connor: Human sympathetic activation by α_2 -adrenergic blockade with yohimbine: Bimodal, epistatic influence of cytochrome P450-mediated drug metabolism. *Clinical Pharmacol Ther.* 2004 Aug; 76(2):139-53.
152. Stein MB, Schork NJ, and Gelernter J: A polymorphism of the beta-1-adrenergic receptor is associated with low extraversion. *Biol Psychiatry* 2004 Aug 15;56(4):217-24.
153. Cory A. Ogden, Michael E. Rich, Nicholas J. Schork, Martin P. Paulus, Mark A. Geyer, James B. Lohr, Ronald Kuczenski, and Alexander B. Niculescu: Candidate genes, pathways and mechanisms for bipolar (manic-depressive) and related disorders: An expanded convergent functional genomics approach. *Mol Psychiatry.* 2004 Nov;9(11):1007-29.
154. Zhang L, Fangwen Rao, Jennifer Wessel, Brian P. Kennedy, Brinda K. Rana, Laurent Taupenot, Elizabeth O. Lillie, Nicholas J. Schork, Michael G. Ziegler, and Daniel T. O'Connor: Functional allelic heterogeneity and pleiotropy of a repeat polymorphism in tyrosine hydroxylase: Prediction of catecholamines and response to stress in twins. *Physiological Genomics.* 2004 Nov 17;19(3):277-91.
155. Salem R, Wessel J, Schork NJ: A Comprehensive Review of Haplotyping Software and Methods for Use with Unrelated Individuals. *Human Genomics* 2005;2(1):39-66
156. Nievergelt CN, Schork NJ: Admixture mapping as a gene discovery approach for common complex human traits and diseases. *Current Hypertension Reports* 2005 Feb;7(1):31-7.
157. Jorgenson E, Tang H, Gadde M, Province M, Leppert M, Kardia S, Schork N, Cooper R, Rao, Boerwinkle E, Risch N (2004). Ethnicity and human genetic linkage maps. *American Journal of Human Genetics.* 2005 Feb;76(2):276-90.
158. Tang H, Quertermous T, Rodriguez B, Kardia SL, Zhu X, Brown A, Pankow JS, Province MA, Hunt SC, Boerwinkle E, Schork NJ, Risch NJ (2004). Genetic Structure, Self-Identified Race/Ethnicity, and Confounding in Case-Control Association Studies. *Am J Hum Genet.* 2005 Feb;76(2):268-75.
159. Reiner AP, Ziv E, Lind DL, Nievergelt CM, Schork NJ, Cummings SR, Phong A, Burchard EG, Harris TB, Psaty BM, Kwok PY (2005): Population structure, admixture, and aging-related phenotypes in African-American adults: The Cardiovascular Health Study. *American Journal of Human Genetics.* 2005 Mar;76(3):463-77.
160. Murthy K, Mahboubi VS, Santiago A, Barragan M, O'Connor DT, Schork NJ, Rana BK (2005). Fidelity of Multiple Displacement Amplification to Generate a Renewable Source of Genomic DNA for Population Genetic Studies. *Human Mutation.* 2005 August;26(2):145-52.
161. Mustanski BS, Dupree MG, Nievergelt CM, Bocklandt S, Schork NJ, Hamer DH. A genomewide scan of male sexual orientation. *Human Genetics.* 2005 Mar;116(4):272-8.
162. Soares ML, Coelho T, Sousa A, Batalov S, Conceicao I, Sales-Luis ML, Ritchie MD, Williams SM, Nievergelt CM, Schork NJ, Saraiva MJ, Buxbaum JN. Susceptibility and modifier genes in Portuguese transthyretin V30M amyloid polyneuropathy: complexity in a single gene disease. *Human Molecular Genetics.* 2005 Feb 15;14(4):543-53.
163. An P, Freedman BI, Hanis CL, Chen YI, Weder AB, Schork NJ, Boerwinkle E, Province MA, Hsiung CA, Wu X, Quertermous T, Rao DC: Genome-wide Linkage Scans for Glucose, Insulin, and Insulin Resistance in the National Heart, Lung, and Blood Institute Family Blood Pressure Program: Evidence of Linkages to Chromosome 7q36 and 19q13 from Meta-Analysis. *Diabetes.* 2005 Mar;54(3):909-14.
164. Friese RS, Mahboubi P, Mahapatra NR, Mahata SK, Schork NJ, Schmid-Schonbein GW, O'Connor DT. Common genetic mechanisms of blood pressure elevation in two independent rodent models of human essential hypertension. *Am J Hypertens.* 2005 May;18(5 Pt 1):633-52.

165. King D, Etzel JP, Chopra S, Smith J, Cadman PE, Rao F, Funk SD, Rana BK, Schork NJ, Insel PA, O'Connor DT. Human response to alpha2-adrenergic agonist stimulation studied in an isolated vascular bed in vivo: Biphasic influence of dose, age, gender, and receptor genotype. *Clin Pharmacol Ther.* 2005 May;77(5):388-403.
166. Bretz WA, Corby PM, Hart TC, Costa S, Coelho MQ, Weyant RJ, Robinson M, Schork NJ. Dental caries and microbial acid production in twins. *Caries Res.* 2005 May-Jun;39(3):168-72.
167. Etzel JP, Rana BK, Wen G, Parmer RJ, Schork NJ, O'Connor DT, Insel PA. Genetic variation at the human alpha2B-adrenergic receptor locus: role in blood pressure variation and yohimbine response. *Hypertension.* 2005 Jun;45(6):1207-13.
168. Stein MB, Fallin MD, Schork NJ, Gelernter J. COMT Polymorphisms and Anxiety-Related Personality Traits. *Neuropsychopharmacology.* 2005 Nov;30(11):2092-102.
169. Murthy KK, Mahboubi VS, Santiago A, Barragan MT, Knoll R, Schultheiss HP, O'connor DT, Schork NJ, Rana BK. Assessment of multiple displacement amplification for polymorphism discovery and haplotype determination at a highly polymorphic locus, MC1R. *Hum Mutat.* 2005 Aug;26(2):145-52.
170. Bao X, Paul J. Mills, Brinda K. Rana, Joel E. Dimsdale, Nicholas J. Schork, Douglas W. Smith, Fangwen Rao, Daniel T. O'Connor, Milos Milic. Interactive Effects of Common Beta2-Adrenoceptor Haplotypes and Age on Susceptibility to Hypertension and Receptor Function. *Hypertension.* 2005 Aug;46(2):301-7.
171. Schwimmer J, Behling R, Newbury R, Deutsch R, Nievergelt C, Schork N, Lavine J. The Histopathology of Pediatric Nonalcoholic Fatty Liver Disease. *Hepatology* 2005 Sept.;42(3):641-9.
172. Greenwood TA, Nicholas J. Schork, Eleazar Eskin, John R. Kelsoe. Identification of additional variants within the human dopamine transporter gene provides further evidence for an association with bipolar disorder in two independent samples. *Molecular Psychiatry.* 2005 Nov; 11(2):125-33.
173. Bretz WA, Corby PM, Schork NJ, Robinson MT, Coelho M, Costa S, MeloFilho MR, Weyant RJ, Hart TC. Longitudinal analysis of heritability for dental caries traits. *J Dent Res.* 2005 Nov;84(11):1047-51.
174. Mathews CA, , Kerry L. Jang, Luis Diego Herrera, Thomas Lowe, Cathy Budman, Gerald Erenberg, Allan Naarden, Ruth Bruun, Nicholas J. Schork, Nelson B Freimer, Victor I. Reus. Tic Symptom Profiles in Subjects with Tourette Syndrome from two Genetically Isolated Populations. *Neuropsychiatric Genetics* 2005 (in press).
175. Nievergelt CM, Daniel F. Kripke, Thomas B. Barrett, Elyssa Burg, Ronald A. Remick, A. Dessa Sadovnik, Susan L. McElroy, Paul E. Keck Jr., Nicholas J. Schork, John R. Kelsoe. Suggestive evidence for association of the circadian genes *PERIOD3* and *ARNTL* with bipolar disorder. *Molecular Psychiatry* 2006, 141:234-241.
176. Chang YP, Kim JD, Schwander K, Rao DC, Miller MB, Weder AB, Cooper RS, Schork NJ, Province MA, Morrison AC, Kardina SL, Quertermous T, Chakravarti A. Related Articles, Links. The impact of data quality on the identification of complex disease genes: experience from the Family Blood Pressure Program. *Eur J Hum Genet.* 2006, 14:469-477.
177. Libiger, O, Schork, NJ. Simulation-based homozygosity mapping with the GAW14 COGA dataset on alcoholism. *BMC Genetics.* 2005 Dec; Supplement 1: 535-540.
178. Luo X, Kranzler HR, Zuo L, Wang S, Schork NJ, and Gelernter, J. Diplotype trend regression (DTR) analysis of the *ADH* gene cluster and *ALDH2* gene: Multiple significant associations for alcohol dependence. *American Journal of Human Genetics* 2006; 78:973-987.
179. Seasholtz TM, Wessel J, Rao F, Rana BK, Khandrika S, Kennedy BP, Lillie EO, Ziegler MG, Smith DW, Schork NJ, Brown JH, O'connor DT. Rho Kinase Polymorphism Influences Blood Pressure and Systemic Vascular Resistance in Human Twins. Role of Heredity. *Hypertension.* 2006 May; 47(5):937-47.

180. Greenwood TA, Rao F, Stridsberg M, Mahapatra NR, Mahata M, Lillie EO, Mahata SK, Taupenot L, Schork NJ, O'Connor DT. Pleiotropic effects of novel trans-acting loci influencing human sympathochromaffin secretion. *Physiol Genomics*. 2006 25(3):470-9.
181. Tang H, Jorgenson E, Gadde M, Kardia, S, Rao DC, Zhu X, Schork N, Hanis C, Risch N. Racial admixture and its impact on BMI and blood pressure in African and Mexican Americans. *Human Genetics* 2006 Jul;119(6):624-33.
182. Niculescu AB, Lulow L, Ogden C, Salomon DR, Schork NJ, Caligiuri M, and Lohr JB: PhenoChipping of psychotic disorders: a novel approach for deconstructing and quantitating psychiatric phenotypes. *Molecular Psychiatry* 2006; 141(6):653-62.
183. Mathews CA, Nievergelt CM, Azzam A, Garrido H, Chavira DA, Wessel J, Bagnarello M, Reus VI, Schork NJ. Heritability and Clinical Features of Multigenerational Families With Obsessive-Compulsive Disorder and Hoarding. *American Journal of Medical Genetics: Neuropsychiatric Genetics* 2007 144(2):174-82..
184. Bretz WA, Corby PM, Melo MR, Coelho MQ, Costa SM, Robinson M, Schork NJ, Drewnowski A, Hart TC. Heritability estimates for dental caries and sucrose sweetness preference. *Archives of Oral Biology*. 2006; 51(12):1156-60.
185. Radant AD, Dobie DJ, Calkins ME, Olincy A, Braff DL, Cadenhead KS, Freedman R, Green MF, Greenwood TA, Gur RE, Light GA, Meichle SP, Mintz J, Nuechterlein KH, Schork NJ, Seidman LJ, Siever LJ, Silverman JM, Stone WS, Swerdlow NR, Tsuang MT, Turetsky BI, Tsuang DW. Successful multi-site measurement of antisaccade performance deficits in schizophrenia. *Schizophr Res*. 200; 89(1-3):320-9.
186. Wessel J, Schork NJ. Generalized genomic distance-based regression methodology for multilocus association analysis. *Am J Hum Genet*. 2006 Nov;79(5):792-806.
187. Calkins ME, Dobie DJ, Cadenhead KS, Olincy A, Freedman R, Green MF, Greenwood TA, Gur RE, Gur RC, Light GA, Mintz J, Nuechterlein KH, Radant AD, Schork NJ, Seidman LJ, Siever LJ, Silverman JM, Stone WS, Swerdlow NR, Tsuang DW, Tsuang MT, Turetsky BI, Braff DL. The Consortium on the Genetics of Endophenotypes in Schizophrenia: Model Recruitment, Assessment, and Endophenotyping Methods for a Multisite Collaboration. *Schizophr Bull*. 2007 33(1):33-48.
188. Schork NJ, Greenwood TA, Braff DL. Statistical Genetics Concepts and Approaches in Schizophrenia and Related Neuropsychiatric Research. *Schizophr Bull*. 2007; 33(1):95-104.
189. Braff DL, Freedman R, Schork NJ, Gottesman II. Deconstructing Schizophrenia: An Overview of the Use of Endophenotypes in Order to Understand a Complex Disorder. *Schizophr Bull*. 2007; 33(1):21-32.
190. Rao F, Wen G, Mahata SK, Kennedy BP, Gayen JR, Rana BK, Mahata M, Salem RM, Stridsberg M, Abel K, Smith DW, Eskin E, Schork NJ, Hamilton BA, Ziegler MG, and O'Connor DT. The Catecholamine Release Inhibitory Peptide Catestatin (Chromogranin A [CHGA]352-372): A Naturally Occurring Amino Acid Variant (Gly364Ser) Causes Profound Changes In Human Autonomic Activity and Alters Risk for Hypertension. *Circulation* (in press).
191. Rana BK, Insel PA, Payne S, Abel K, Beutler E, Ziegler MG, Schork NJ, O'Connor DT. A large population-based sample reveals the contribution of gene-gender interactions in blood pressure in Caucasian-Americans. *Hypertension*; 2007 49(1):96-106.
192. Zapala M, Schork NJ. Multivariate regression analysis of distance matrices for testing associations between gene expression patterns and related variables. *Proceedings of the National Academy of Sciences* 2006; 103(51):19430-5.
193. Glatt SJ., Chayavichitsilp P., Depp C, Schork NJ, Jeste DV. Successful Aging: From Phenotype to Genotype. *Biological Psychiatry* 2007 (Epub Ahead of Print).

194. Gong Y, Beitelshes AL, Wessel J, Langae TY, Schork NJ, Johnson JA. SNP discovery and haplotype analysis of bk channel beta 1 subunit. *American Journal of Pharmacogenomics* (in press).
195. Wessel J, Moratorio G, Rao F, Mahata M, Zhang L, Greene W, Rana BK, Kennedy BP, Khandrika S, Lillie EO, Smith DW, Ziegler MG, Witztum J, Schork NJ, Schmid-Schönbein GW, O'Connor DT. C reactive protein, an "intermediate phenotype" for inflammation: human twin studies reveal heritability, association with blood pressure and the metabolic syndrome, and the influence of common polymorphism at catecholaminergic/□-adrenergic pathway loci *Journal of Hypertension* 2007 Feb;25(2):329-343.
196. Greenwood TA, Ondrej Libiger, Sharon Kardia, Craig Hanis, C. Charles Gu, Michael Miller, Alan B. Weder, Schork NJ, Comprehensive Linkage and Linkage Heterogeneity Analysis of 4344 Sibling Pairs Affected with Hypertension from the Family Blood Pressure Program. *Genetic Epidemiology* (in press).
197. Luo X, Kranzler HR, Zuo L, Wang S Schork NJ, Gelernter J (2006). Multiple ADH and ALDH genes modulate risk for drug dependence in both African- and European-Americans. *Human Molecular Genetics* 2007; 16(4):380-90.
198. Swerdlow NR, Sprock J, Light GA, Cadenhead K, Calkins ME, Dobie DJ, Freedman RR, Green MF, Greenwood TA, Gur RE, Mintz J, Olincy A, Nuechterlein KH, Radant AD, Schork NJ, Seidman LJ, Siever LJ, Silverman JM, Stone WS, Tsuang DW, Tsuang MT, Turetsky BI, Braff DL (2007). Multi-site studies of acoustic startle and prepulse inhibition in humans: Initial experience and methodological considerations based on studies by the Consortium on the Genetics of Schizophrenia. *Schizophrenia Research* (in press).
199. Beitelshes AL, Gong Y, Wang D, Schork NJ, Cooper-DeHoff RM, Langae TY, Shriver MD, Sadee W, Pepine CJ, Johnson JA for the INVEST Investigators. KCNMB1 genotype influences response to verapamil SR and adverse outcomes in the INternational VErapamil SR/Trandolapril STudy (INVEST). *Pharmacogenetics and Genomics* (in press).
200. Hovatta I, Zapala MA, Broide RS, Schadt EE, Libiger O, Schork NJ, Lockhart DJ, and Barlow C. DNA variation and brain-region specific expression profiles show different relationships between inbred mouse strains: implications for eQTL mapping studies. *Genome Biology* (in press).
201. Neivergelt CM, Libiger O, Schork NJ. Generalized molecular analysis of variance. *PLOS Genetics* (in press).
202. Schork NJ, Wessel J, Malo N. DNA Sequence-Based Phenotypic Association Analysis. *Advance in Genetics: Genetic Dissection of Complex Traits*, second edition (DC Rao, editor; in press).
203. Braff D, Schork NJ, Gottesman II. Endophenotyping Schizophrenia. *Schizophrenia Bulletin* (in press).
204. Stein MB, Schork NJ, Gelernter J. Gene-by-Environment (Serotonin Transporter and Childhood Maltreatment) Interaction for Anxiety Sensitivity, an Intermediate Phenotype for Anxiety Disorders. *Neuropsychopharmacology* (in press).
205. Gu C, Hunt SC, Kardia S, Turner S, Chakravarti A, Schork NJ, Olshen R, Curb D, Jacquish C., Rao DC. An Investigation of Genome-wide Associations of Hypertension with Microsatellite Markers in the Family Blood Pressure Program (FBPP). *Human Genetics* 2007; (in press; electronic pre-publication)
206. Rao F, Wessel J, Wen G, Zhang L, Rana BK, Kennedy BP, Greenwood TA, Salem RM, Chen Y, Khandrika S, Hamilton BA, Smith DW, Holstein-Rathlou NH, Ziegler MG, Schork NJ, O'connor DT. Renal Albumin Excretion. Twin Studies Identify Influences of Heredity, Environment, and Adrenergic Pathway Polymorphism. *Hypertension*. 2007 Mar 12 (in press).
207. Torkamanini A, Schork NJ. Prediction of Disease Causing Nonsynonymous Polymorphisms within the Human Kinase Gene Family. *Genomics* (in press).
208. Wessel J, Zapala MA, Schork NJ. Accommodating Pathway Information in Expression Quantitative Trait locus ("eQTL") Analysis. *Genomics* (in press).

209. Greenwood TA, Braff DL, Cadenhead KS, Calkins ME, Dobie DJ, Freedman R, Green MF, Gur RE, Gur RC, Light GA, Mintz J, Nuechterlein KH, Olincy A, Radant AD, Seidman LJ, Siever LJ, Silverman JM, Stone WS, Swerdlow NR, Tsuang DW, Tsuang MT, Turetsky BI, Schork NJ. The Consortium on the Genetics of Schizophrenia (COGS): Preliminary Heritability Analyses of Endophenotypic Measures for Schizophrenia Archives of General Psychiatry (in press).

Articles Submitted for Publication

1. Barrett, TB, John E. Emberton, Caroline M. Nievergelt, Sherri G. Liang, Eleazar Eskin, Nicholas J. Schork, John R. Kelsoe. Further evidence for association of GRK3 to bipolar disorder suggests a second disease mutation. *Psychiatric Genetics* (in review).
2. Bretz WA, Patricia M. A. Corby, Mario R. Melo Filho, Simone M. Costa, Robert J. Weyant, Jennifer Wessel, Nicholas J. Schork, Adam Drewnowski, Thomas C. Hart. Inheritance of Occlusal Tomography. *Pediatric Dentistry* (in review).
3. Libiger O, Schork NJ. An Extended Gene-Dropping Approach to the Probabilistic Evaluation of Chromosome Segment Sharing in Studies Exploiting Genealogical Information. *European Journal of Human Genetics* (in review).
4. Wessel J, Libiger O, Schork NJ. Whole Genome Association Studies Using Window-Based Multivariate distance Matrix Regression Analysis. *Genetic Epidemiology* (in review).
5. Wessel J, Schork AJ, Tiwari HK, Schork NJ. Powerful Designs for Molecular Genetic Association Studies that consider Twins and Sibling Pairs with Discordant Genotypes. *Genetic Epidemiology* (in review).
6. Gerhard T, Gong Y, Beitelshes AL, Lobmeyer MT, Cooper-DeHoff RM, Langae TY, Schork NJ, Pepine CJ, Johnson JA for the INVEST investigators. Association between Cardiovascular Outcomes, Diuretic Therapy and the α -adducin Polymorphism: Results from the International Verapamil SR-trandolapril Study Genetic Substudy (INVEST-GENES). *Circulation* (in review).
7. Ye C, Zapala MA, Kang HM, Wessel J, Eskin E, Schork NJ. High-Density QTL Mapping to Identify Phenotypes and Loci Influencing Gene Expression Patterns in Entire Biochemical Pathways. *BMC Genetics* (in review).
8. Mosley C, Taupenot L, Taulane JP, Biswas N, Vaingankar S, Wen G, Schork NJ, Ziegler MG, Mahata SK, O'Connor DT. Biogenesis of the secretory granule: Chromogranin A (CHGA) coiled-coil structure predicts unusual physical properties and suggests a mechanism for granule core condensation. *Journal of Molecular Biology* (in review).
9. Lillie EO, Mahata M, Khandrika S, Rao F, Bunday RA, Wen G, Vaingankar S, Biswas N, Taupenot L, Hamilton BA, Rana BK, Smith DW, Mahata SK, Ziegler MG, Cockburn M, Schork NJ, O'Connor DT. Heredity of endothelin secretion: Human twin studies reveal the influence of polymorphism at the chromogranin A locus, a novel determinant of endothelial function. *Circulation* (in review).
10. Bhatnagar V, O'Connor DT, Schork NJ, Rana BK, Salem RM, Smith DW, Lipkowitz MK and the AASK Study Investigators. Polymorphisms at the angiotensin converting (ace) enzyme locus predict the time-course of blood pressure response to ace inhibition in the AASK trial. *Hypertension* (in review).
11. Evans LM, Akiskal HS, Greenwood TA, Nievergelt CM, Keck Jr. PE, McElroy SL, Sadovnick AD, Remick AD, Schork NJ, Kelsoe JR. Suggestive Linkage of Different Chromosomal Loci to Cyclothymic and Anxious Temperaments in Bipolar Disorder. *American Journal of Medical Genetics: Psychiatric Genetics* (in review).
12. Wen G, Wessel J, Zhou W, Ehret GB, Rao F, Stridsberg M, Mahata SK, Gent PM, Das M, Cooper RS, Chakravarti A, Zhou H, Schork NJ, O'Connor DT, Hamilton BA. An ancestral SCG2 variant confers

regulation by PHOX2 transcription factors and increased risk of hypertension. *Journal of Clinical Investigation* (in review).

13. Zhang L, Rao F, Zhang K, Khandrika S, Das M, Vaingankar SM, Kennedy BP, Rana BK, Smith DW, Wessel J, Salem RM, Rodriguez JL, Mahata SK, Schork NJ, Ziegler MG, O'Connor DT. GTP cyclohydrolase (GCH1), a rate-limiting enzyme in neurotransmitter biosynthesis: Discovery of common human genetic variants governing nitric oxide, autonomic activity, and cardiovascular risk in twin pairs. *Journal of Clinical Investigation* (in review).
14. Ursin G, Lillie EO, Cockburn M, Cozen W, Parisky YP, Hamilton AS, Schork NJ, Astrahan MA, Mack T. A Revised Estimate of the Heritability of Mammographic Density. *Journal of the National Cancer Institute* (submitted).
15. Rao F, Zhang L, Kennedy BP, Wessel J, Taupenot L, Wen G, Zhang K, Rana BK, Smith DW, Lillie EO, Cadman PE, Salem RM, Schork NJ, Ziegler MG, O'Connor DT. Tyrosine hydroxylase (TH), the rate-limiting enzyme in catecholamine biosynthesis: Discovery of common human genetic variants governing transcription, autonomic activity and blood pressure in vivo. *Circulation* (in review).
16. Greenhall JA, Zapala MA, Cáceres M, Libiger O, Schork NJ, Barlow C, Lockhart DJ. Detecting Genetic Variation in Microarray Expression Data. *Genome Research* (in review).

Chapters in Books

1. Schork NJ and Hardwick J: Supercomputer-intensive multivariable randomization tests. In: *Proceedings of the 22nd Symposium on the Interface of Computing Science and Statistics*. New York: Springer Verlag, 1992: 509-13.
2. Schork NJ: The parallel computation of pedigree likelihoods. In: *Proceedings of the 23rd Symposium on the Interface of Computing Science and Statistics*. Fairfax Station, VA: The Interface Foundation, 1992: 262-5.
3. Schork NJ: Bootstrapping likelihood ratios in quantitative genetics. In: LePage R, Billard L, eds. *Exploring the Limits of Bootstrap*. New York: John Wiley & Sons, 1992:389-96.
4. Schork NJ, Chakravarti A: A non-mathematical overview of modern gene mapping method techniques used in humans. In, S. Mockrin, ed. *Molecular Genetics and Gene Therapy of Cardiovascular Diseases*. New York: Marcel Dekker 1996:79-109.
5. St. Jean P, Schork NJ: Genetic epidemiologic approaches to finding genes that influence susceptibility to cancer. *Inherited Susceptibility to Cancer*, WD Foulkes and SV Hodgson, eds. Cambridge: Cambridge University Press 1998:109-33.
6. Schork NJ: Mixed models in segregation analysis. *Encyclopedia of Biostatistics*. Armitage, Peter and Theodore Colton, editors-in-chief. John Wiley & Sons, Ltd, 1998: v.4:2662-4.
7. Schork NJ, Kashkoush S, Xu X: Hypertension as a complex trait amenable to genetic analysis: Basic strategies and integrative approaches. In, Dominiczak A.F., Connell J. M. C., Soubrier F. eds. *Molecular Genetics of Hypertension*. Oxford: BIOS Scientific 1999:1-30.
8. Iyengar SK, Weder AB, Koike G, Jokelainen P, Jacob HJ, Schork NJ: The developmental evolution of hypertension: evidence, genetic models, and mechanisms. In, McCarty R, Blizzard DA, Chevalier RL eds. *Handbook of Hypertension. Development of the Hypertensive Phenotype: Basic and Clinical Studies*. New York: Elsevier 1999; Volume 19:77-102.
9. Xu X, Chen C, Niu T, Wang B, Rogus J, Schork NJ, Wang A: Twin and sibpair studies in developing countries. In, Spector T, Sneider H, and MacGregor A eds. *Advances in Twin and Sib Pair Analysis*. London: Greenwich Medical Media 2000:91-101.

10. Schork NJ, Xu X: The use of twins in QTL mapping. In, Spector T, Sneider H, and MacGregor A eds. *Advances in Twin and Sib Pair Analysis*. London: Greenwich Medical Media 2000:189-202.
11. Schork NJ, Fallin D, Tiwari HK, Schork MA: Pharmacogenetics. *Handbook of Statistical Genetics* In Balding D, Bishop M, and Cannings C.eds. Chichester 2001:741-64. (First and Second editions only).
12. Schork NJ: Basic principles of genetics. In Weder A, ed. *Hypertension Primer* 2003, Ch A73:212-5.
13. Rana BK, Schork NJ: Genetic epidemiology of complex traits. *Encyclopedia of the Human Genome*. Native Publishing Group 2005 (www.ehgonline.net).
14. Rana BK, Smith D, Schork NJ: Polymorphism and sequence assembly. *Encyclopedia of Genetics and Genomics*. Proteomics, and Bioinformatics. New York: John Wiley and Sons 2005.
15. Baker DG, Risbrough, V, Schork NJ: Post traumatic stress disorder: genetic and environmental risk factors. *Biobehavioral Resilience to Stress*, forthcoming.

Abstracts, Preliminary Communications, Panel Discussions (Available upon request)