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CURRICULUM VITAE

NAME: M. Ilyas Kamboh
CITIZENSHIP: USA
BUSINESS ADDRESS: Department of Human Genetics
Graduate School of Public Health
University of Pittsburgh
A311 Crabtree Hall
130 DeSoto Street
Pittsburgh, PA 15261
Telephone(412) 624-3021
Telefacsimile: (412) 624-3020
E-mail: kamboh@pitt.edu



EDUCATION AND TRAINING

Undergraduate

University of Punjab B.Sc., 1976 MAJORS: Biology
Forman Christian College
Lahore, Pakistan

Graduate

University of Punjab M.Sc., 1979 MAJORS: Human &
Government College,
Department of Zoology,
Lahore, Pakistan

Post Graduate

The Australian National Ph.D., 1984 Human Genetics
University, John Curtin School
of Medical Research,
Department of Human Biology
Canberra, Australia Advisor: Dr. R. L. Kirk

APPOINTMENTS AND POSITIONS

Academic

October 2008- October 2013	Department of Human Genetics University of Pittsburgh	Chairman
January 2005- October 2008	Department of Human Genetics University of Pittsburgh	Interim Chair
July 2004- December 2004	Department of Human Genetics University of Pittsburgh	Vice Chair
July 2010- Present	Department of Epidemiology University of Pittsburgh	Professor (Secondary Appointment)
July 1998- Present	Department of Psychiatry University of Pittsburgh	Professor (Secondary Appointment)
January 1997- Present	Department of Human Genetics University of Pittsburgh	Professor (Tenured)
July 1991- December 1996	Department of Human Genetics University of Pittsburgh	Associate Professor (Tenured in 1993)
1993-Present	Pittsburgh Cancer Institute	Member
February 1987- June 1991	Human Genetics Division University of Pittsburgh	Assistant Professor
April 1985- January 1987	Department of Biostatistics Human Genetics Division University of Pittsburgh	Research Associate
December 1984- March 1985	Institute of Experimental Medicine Lahore, Pakistan	Senior Research Officer
1984	Department of Human Biology John Curtin School of Medical Research Australian National University Canberra, Australia	Research Assistant
1980 -1981	Government Imamia Degree College Sahiwal, Pakistan	Lecturer
1979 -1980	Department of Zoology Government College Lahore, Pakistan	Demonstrator

MEMBERSHIP IN PROFESSIONAL AND SCIENTIFIC SOCIETIES

Years Inclusive	Organization
1981-1984	Human Genetics Society of Australasia
1985-present	American Society of Human Genetics
1987-present	American Association of Physical Anthropologists
1990-1993	International Electrophoresis Society
1990-present	American Association for the Advancement of Science
1990-present	American Heart Association <ul style="list-style-type: none"> • Fellow-Arteriosclerosis, Thrombosis and Vascular Biology Council • Fellow-Epidemiology Council
1990-present	International Atherosclerosis Society
1991-1992	The New York Academy of Sciences
1999-present	International Genetic Epidemiology Society
1999-present	American College of Rheumatology
2015-present	Alzheimer's Association International Society to Advance Alzheimer's Research and Treatment (ISTAART)

HONORS

Year	Title of Award	Awarding Association
1976-1978	Academic Merit Scholarship on the basis of B.Sc. examination results	University of the Punjab
1977	Merit Scholarship	Fauji Foundation
1979	First position with distinction in M.Sc. examination	Government College, Lahore
1979	Roll of Honor in M.Sc.	Government College, Lahore
1980	First position by merit in the selection of Lectureship	Punjab Public Service Commission, Lahore
1981-1984	Ph.D. Merit Scholarship	The Australian National University
1981-1983	Ahmadia Open Merit Scholarship	Ahmadia Education Foundation
1985	Selected for INSERM Fellowship, France	INSERM

1994-present	Who's Who in the East	Marquis
1994-present	American Men and Women of Science	R.R. Bowker
2000-present	Who's Who in America	Marquis Who's Who
2000	Delta Omega Omicron, Public Health National Honor Society	Delta Omega National Honor Society for Schools of Public Health by the Omicron Chapter of Delta Omega, Graduate School of Public Health, University of Pittsburgh
2001- present	Fellow of the American Heart Association	American Heart Association
2003 – present	Who's Who in the World	Marquis Who's Who
2003	Excellence in Science Award	Muslim Physician Society of Greater Pittsburgh

PROFESSIONAL ACTIVITIES

1. Teaching

a. Courses Taught

Number	Title	Hours of Lecture	Enrollment	Credits
HUGEN 2025	1991-1993: "Human Genetics Seminar"	1	~ 50	1
HUGEN 2017	1991-2000: "Human Genetics" (Co-Instructor)	3	7 – 26	3
EPIDEM 2600	1994: "Molecular Epidemiology" (Guest Lecturer)	3	20	3
HUGEN 2034	2001-2019: "Biochemical and Molecular Genetics of Complex Diseases" (Co-Instructor; Instructor from 2005)	3	20 – 30	3
HUGEN 2090	2020-2021: "Genetics of Complex Disease I" (Instructor)	3	30-52	2
HUGEN 2091	2022-Present: "Genetics of Complex Disease II" (Instructor)	3	52	1

b. Other Teaching

Type of Teaching	Title	Date(s)
Demonstrator	Teaching practical and experimental	1979-1980

Department of Zoology Government College Lahore, Pakistan	aspects in biological sciences.	
Lecturer Government Imamia Degree College Sahiwal, Pakistan	Teaching various courses in biological sciences, including Biochemistry, Genetics, Evolution, Paleontology, Embryology, Ecology.	1980-1981
Teaching Advanced Topics in:	Human Genetics (Bios 224), Human Genetics Seminars (Bios 225), Guest lecturer in Human Biochemical and Molecular Genetics (Bios 234); helping graduate students in designing experiments and evaluating results of their research projects.	1985-1990

c. Supervision of Graduate Student Essays, Theses, and/or Dissertations

Name of Student	Degree Awarded, Year	Type of Document and Title
George Wang	Ph.D., 1993-1999	Genetic and Functional Analyses of Apolipoprotein A-IV in Relation to the Quantitative Risk Profile For Coronary Heart Disease.
Hamid Razzaghi	Ph.D., 1994-1999	Population Screening of the Lipoprotein Lipase Gene for Mutations Associated With High Triglyceride/Low HDL-Cholesterol Phenotype and Structure-Function Analysis of the Mutations Using Molecular Modeling.
Xiaoyan Wang	Ph.D., 1995-1999	Molecular Genetic Analysis of the α_1 - Antichymotrypsin Gene in Relation to the Risk of Alzheimer's Disease.
Purnima Desai	Ph.D., 1996-2001	Molecular Genetics and Immunohistochemical Analyses of Apolipoprotein D in Relation to Alzheimer's Disease.
Erin Luedeking-Zimmer	Ph.D., 1997-2001	Association of Candidate Genes on Chromosome 12 in Late-Onset Alzheimer's Disease.
Qi Chen	Ph.D., 1998-2002	Genetic Factors in Coronary Heart Disease in Women.
Ayla Ozturk	Ph.D., 2002-2005	Investigation of the Relationship of Candidate Genes on Chromosome 10 with the Risk and Age-at-Onset of Alzheimer's Disease
Pei-an Betty Shih	Ph.D., 2004-2007	Evidence for Haplotype-based Association in SLE at the C-reactive protein Locus: Population-based and Family-based Association Studies
Sangita Suresh	Ph.D., 2003-2008	Functional Characterization of Apolipoprotein H

		Promoter Polymorphisms and Their Relation With Systemic Lupus Erythematosus
Sudeshna Dasgupta	Ph.D., 2005-2008	Association of Paraoxonase-2 Genetic Variation With Serum Paraoxonase Activity and Systemic Lupus Erythematosus
Vipavee Niemsiri	Ph.D., 2010-2016	Genetic Influence of Sequence Variants in <i>SCARB1</i> and <i>ABCA1</i> Genes on Major Lipid Traits: A Candidate Gene Association Study
Dilek Pirim	Ph.D., 2011- 2014	Variants Discovery in the <i>LPL</i> and <i>CETP</i> Genes and Their Associations With Plasma Lipid and Apolipoprotein Levels
Zaheda Radwan	Ph.D., 2011-2013	A Comprehensive Association Study of Apolipoprotein <i>E-C1-C4-C2</i> Gene Cluster on Chromosome 19 with Plasma Lipoprotein Traits
Samantha Rosenthal	Ph.D., 2011-2016	Whole-exome and targeted sequencing of Alzheimer's disease: an integrative approach
Vibha Acharya	Ph.D., 2018-Present	Genetics of Cognitive Decline in Older Adults
Lily Francis	Ph.D., 2018-Present	Genetic Studies on Alzheimer's disease Varinats
Ruyu Shi	Ph.D., 2020-Present	Genetics of the Natural History of Alzheimer's disease
Corintha Goble	M.S., 1989-1991	The Role of the Apolipoprotein B Allelic Variation in Determining Lipoprotein/ Lipid Levels in a Biethnic Population in the San Luis Valley, Colorado.
Shabnam Ali	M.S., 1994-1996	Association of Two Polymorphisms in the Apolipoprotein(a) Gene With Serum Lipoprotein(a) Levels Among African Blacks.
Megan Harris	M.S. Genetic Counseling; 1994-1996	The Role of the Low Density Lipoprotein Receptor-Related Protein (LRP) Gene Polymorphism in Lipid Metabolism.
Jennifer Anderson	M.S. Genetic Counseling; 1994-1996	Relationship of Two Apolipoprotein B Polymorphisms With Plasma Lipoprotein- Lipid Levels in African Blacks.
Cara Nestlerode	M.S., 1994-1997	Genetic Contribution of Apolipoprotein J in Affecting Serum Lipoprotein-Lipid and Apolipoprotein Levels.
Linda Chiu	M.S. Genetic Counseling; 1997-1999	Apolipoprotein(a) Polymorphisms and Plasma Lipoprotein(a) Levels in Hispanics and Non- Hispanic Whites.
Tejal J. Bhojak	M.S., 1998-1999	Polymorphisms in the Cathepsin D and Interleukin-6

		Genes and their Possible Correlations With the Risk of Alzheimer's Disease.
Shannon Lindsay	M.S. Genetic Counseling; 1998-1999	Promoter Polymorphisms in the Apolipoprotein E Gene and the Risk of Alzheimer's Disease.
Janel Markey	M.S., 1996-1998	The Role of Paraoxonase 1/Codon 192 Polymorphism and its Relationship With Coronary Heart Disease.
Kristen DePrince	M.S. Genetic Counseling; 1997-1998	Apolipoprotein(a) Polymorphisms and Plasma Lipoprotein(a) Concentrations in Samoans.
Erin Jacobs	M.S. Genetic Counseling; 2003-2005	Association of Single Nucleotide Polymorphisms in the Promoter of Apolipoprotein H in Systemic Lupus Erythematosus
Laura Tripi	M.S. Genetic Counseling; 2003-2005	Genetic Variation in the Paraoxonase-1 Gene and Association With Systemic Lupus Erythematosus
Leia Corthell	M.S. Genetic Counseling; 2006-2007	Association of Single Nucleotide Polymorphisms in Apolipoprotein H with the Risk of Systemic Lupus Erythematosus
Jessica Figgins	M.S. Genetic Counseling; 2006-2008	Association of 22 Candidate Genes with Late-Onset Alzheimer's Disease
Sally Hollister	M.S. Genetic Counseling; 2006-2008	Sequence Variation in the APOA2 Gene and its Relationship with Plasma HDL-Cholesterol Levels
Sarah Hill	M.S. Genetic Counseling; 2007-2009	Sequence Variation in the APOA1 and APOA4 Genes and Their Relationship with Plasma HDL-Cholesterol Levels
Kate Sleutz	M.S. Genetic Counseling; 2008-2010	Sequence variation in the CD36 gene and its Relationship with plasma HDL cholesterol levels
Lauren Burns	M.S. Genetic Counseling; 2008-2010	A Candidate Gene Study of Late-Onset Alzheimer's Disease
Zaheda Radwan	M.S., 2009-2010	Effects of Common and rare Genetic Variants of Apolipoprotein C4 on HDL-Cholesterol Level
Dilek Pirim	M.S., 2009-2010	Lipoprotein Lipase Gene Sequencing and Lipid Profile
Sharna Tingle	M.P.H., 2010-2011	IL-10 Genetic Variation and Risk of Systemic Lupus Erythematosus
Jordan Harper	M.S., 2020-2021	Genome-Wide Association Study of Dementia in a Community-Based Sample of Oldest-Old
Courtney M. Kasturiarachi	M.P.H., 2020-2022	Genetic Variation in PCSK9 and Plasma Lipid Profile

d. Co-Supervision of International PhD Students

Name of Student	Degree, Year	Foreign Institute
Syed Fazal Jalil	PhD, 2011-2013	National University of Science and Technology, Islamabad, Pakistan
Mamoonah Chaudhry	PhD, 2013-2015	University of Punjab, Lahore, Pakistan
Asima Zia	PhD, 2013-2015	National University of Science and Technology, Islamabad, Pakistan
Aysha Kiani	PhD, 2013-2015	National University of Science and Technology, Islamabad, Pakistan
Asma Cheema	PhD, 2013-2016	National University of Science and Technology, Islamabad, Pakistan
Muaaz Aslam	PhD, 2017-2021	National University of Science and Technology, Islamabad, Pakistan
Sabiha Bibi	PhD, 2017-Present	National University of Science and Technology, Islamabad, Pakistan

e. Service on Masters and Doctoral Committees

Position	Committee Name	Student's Name	Dates Served
Member	Ph.D.	Bahman Sepehrnia	1986-1988
Member	Ph.D.	Shelly Cole	1988-1990
Member	M.S.	Susan Slaugenhaus	1988
Member	Ph.D.	Sudha Iyengar	1990-1992
Member	M.S.	Joanne Oleck	1990
Member	Ph.D.	John Law	1990-1991
Member	M.S.	Yvette Perry	1992
Member	M.P.H.	Kathleen McHugh	1993-1995
Member	Ph.D.	Weiching Wang	1994-1996

Member	M.S.	Junbum Kim	1997-1998	
Member	Ph.D.	Janet Johnson	1997-1998	
Member	M.S.	Manisha Balwani	1999	
Member	Ph.D.	William Swaney	2001	
Committee Chair	Ph.D.	Xiaojing Ye	2005	
Member	Ph.D.	Pattarana Sue-Chew	2005	
Member	Ph.D.	Andrea Robinson	2009-2012	
Member	Ph.D.	Janelle Zacherl	2010-2013	
Member	Ph.D.	Sharanya Kumar	2010-2013	
Member	Ph.D.	Ferdouse Begum	2012-2013	
Committee Chair/Member	M.S.	Laura Smith	2013	
M.S.	Committed Chair/Member	Laura SmM.S.	Shazina Saeed 2013	2014
Committee Chair/Member	Ph.D	Janelle Zacherl	2013-2014	
Member	M.S.	Amanda D. Matchette	2014	
Member	M.S.	Avani Ahuja	2019	

f. Supervision of Post-Doctoral Students, Residents, and Fellows

Name of Student	Position	Dates Supervised
Young I. Ahn, Ph.D.	Research Associate	1991-1993
Dharambir K. Sanghera, Ph.D.	Research Associate	1993-1995
Hyun Sup Kim, Ph.D.	Research Fellow, Kongjun National University, Korea	1996
Chew-Kiat Heng, Ph.D.	Research Fellow, National University of Singapore	1997
Hamid Razzaghi, Ph.D.	Post-doctoral Fellow	1999-2002
Ousama Khalifa, M.D.	Post-doctoral Fellow	2000-2001

Purnima Desai, Ph.D.	Research Associate	2001-2004
Erin Luedeking-Zimmer, Ph.D.	Post-Doctoral Fellow	2002
Qi Chen, Ph.D.	Research Associate	2003-2006
Li-Xia Yang, MD., Ph.D.	Research Associate	2003-2006
F. Yesim K. Demirci, M.D.	Research Associate	2005-2006
Xingbin Wang, Ph.D.	Post-doctoral Fellow	2011-2013
M. Muaaz Aslam Ph.D.	Post-doctoral Fellow	2021-Present

g. Faculty Support

Name of Faculty	Position of Faculty	Dates
Dharambir K. Sanghera, Ph.D.	Assistant Professor	1995-1999
Haider Mehdi, Ph.D.	Assistant Professor	1996-2001
Dharambir K. Sanghera, Ph.D.	Visiting Faculty	2000
F. Yesim K. Demirci, M.D.	Research Assistant Professor	2006-2009
Xingbin Wang, Ph.D.	Research Assistant Professor	2013-2016
Kang-Hsien Fan, Ph.D.	Research Assistant Professor	2017-Present

h. Mentoring Students in Field Placements

Name of Student	Degree/Program	Description	Field Site(s)
Kendall A. Norman-Cooper 1991-1992	Undergraduate	Biology	University of Pittsburgh –“QUEST”, Jean Hamilton Walls and Minority Research Apprenticeship Program (MRAP)
Nicole McCrief 1992-1993	Undergraduate	Biology	University of Pittsburgh –“QUEST”, Jean Hamilton Walls and Minority Research Apprenticeship Program (MRAP)
Carmel F. Portugal 1992	Undergraduate	Biology	University of Pittsburgh –“QUEST”, Jean Hamilton Walls and Minority Research Apprenticeship Program (MRAP)

Krista Suggs 1992-1993	Undergraduate	Biology	University of Pittsburgh –“QUEST”, Jean Hamilton Walls and Minority Research Apprenticeship Program (MRAP)
Creighton Moorehead 1993-1994	Undergraduate	Biology	University of Pittsburgh –“QUEST”, Jean Hamilton Walls and Minority Research Apprenticeship Program (MRAP)
Melisa Bennett 1994	Undergraduate	Biology	University of Pittsburgh –“QUEST”, Jean Hamilton Walls and Minority Research Apprenticeship Program (MRAP)
Toral Surti 1994	Undergraduate	Biology	Public Health Careers Opportunity Program (PHCOP) Students
William M. Chaparro 1994-1995	Undergraduate	Biology	University of Pittsburgh –“QUEST”, Jean Hamilton Walls and Minority Research Apprenticeship Program (MRAP)
Omari Humphries 1997	Undergraduate	Biology	University of Pittsburgh –“QUEST”, Jean Hamilton Walls and Minority Research Apprenticeship Program (MRAP)
Asia Jordan 2001	Undergraduate	Biology	University of Pittsburgh –“QUEST”, Jean Hamilton Walls and Minority Research Apprenticeship Program (MRAP)
Nida Jawed 2004	Medical student	Summer internship	Agha Khan Medical College, Karachi, Pakistan
Fazia Ahmad Mir 2005	Medical student	Summer internship	Agha Khan Medical College, Karachi, Pakistan

2. Research and Training

a. *Grants and Contracts Received (Active Funding)*

Years Inclusive	Grant or Contract Number and Title	Source	Role	Amount
4/2021 – 3/2025	R01 AG064877: Genetic Architecture of Alzheimer’s Disease Proteinopathies	NIH/NIA	Principal Investigator	Total cost: \$8,377,181

8/2019 – 4/2021	RF56 AG064877: Genetic Architecture of Alzheimer's Disease Proteinopathies	NIH/NIA	Principal Investigator	Total cost: \$2,626,839
4/2016 – 3/2026	R01AG023651/R37:Mild Cognitive Impairment: a Prospective Community Study	NIH/NIA	Co-Investigator	Total cost: \$2,021,320 Kamboh's budget
9/2020 – 8/2025	U19AG068054 Alzheimer's Biomarkers Consortium-Down Syndrome (ABC-DS):	NIH/NIA	MPI, Project 2: Genomics	\$1,511,144 Kamboh's budget
5/2018 – 4/2023	R01AG058549: Risk factors for MCI and Dementia in a Diverse Senior Cohort	NIH/NIA	Co-Investigator	Total cost: \$117,141 for Kamboh's Genetics studies
5/2000- 4/2025	P50 AG05133/P30 AG066468: University of Pittsburgh Alzheimer's Disease Research Center	NIH/NIA	Director, Biomarker and Neurogenetics Core	Total cost: \$2,649,241 for Kamboh's Genetics Core
1/2021 – 12/2025	R01 AG069912: Genetics and Molecular Correlates of White Matter Pathology in Alzheimer's disease	NIH/NIA	Co-Investigator	Total cost: \$383,597 Kamboh's budget
9/2021 – 8/2022	RF56 AG074467: Neurocognitive Outcomes and Precursors of Alzheimer Disease Related Dementias after SARS-CoV-2 Infection: an Ultrahigh Field (7T) MRI Study in a Diverse, Multinational Cohort	NIH/NIA	Co-Investigator	Total cost: \$70,511 Kamboh's budget

b. Grants and Contracts Received (Previous Funding)

Years Inclusive	Grant or Contract Number and Title	Source	Role	Amount
2/2009 – 5/2022	R01/R02 AG030653: Search for the Alzheimer's Disease Genes	NIH/NIA	Principal Investigator	Total cost: \$7,720,857
5/2012 – 4/2022	R01 AG027224: Prediction of Psychosis in Alzheimer's Disease	NIH/NIA	Co-Investigator	Total cost: \$360,854 for Kamboh's budget

9/2020 – 8/2021	R01 AG053952-01: Investigating Gains in Neurocognition in an Intervention Trial of Exercise	NIH/NIA	Co-Investigator	Total cost: \$62,260 Kamboh's budget
5/2016 – 4/2020	U01 AG051197: Connectomics of Brain Aging and Dementia	NIH/NIA	Co-Investigator	Total cost: \$250,306 Kamboh's budget
9/2015 - 4/2020	U01 AG051406: Neurodegeneration in Aging Down Syndrome (NiAD): A Longitudinal Study of Cognition and Biomarkers of Alzheimer's Disease	NIH/NIA	Co-Investigator	Total cost: \$358,897 Kamboh's Genetics studies
9/2012- 8/2019	R01 AG041718: Deep Resequencing of Candidate Gene Regions in Late-onset Alzheimer's Disease	NIH/NIA	Principal Investigator	Total cost: \$3,066,289
12/2009- 11/2014	R01 AR057028: Molecular Genetics of BAT Genes and SLE Risk	NIH/NIAMS	Co-investigator	Total cost: \$1,780,461
12/2008- 11/2012	R01 HL092397: Immune/Inflammation Genomics and the Risk of SLE and CHD	NIH/NHLBI	Principal Investigator	Total Cost: \$2,735,942
5/2007- 4/2012	R01 HL088648: Prothrombin Gene Variation and the Risk of SLE and CHD	NIH/NHLBI	Principal Investigator	Total cost: \$1,963,545
2/2007- 1/2012	R01 HL084613: Genetic Determinants of HDL-Cholesterol	NIH/NHLBI	Principal Investigator	Total cost: \$2,901,017
9/2009- 8/2012	R01 DK082766: Genome-Wide Association Scan to Identify Risk Genes for Type 2 Diabetes in Asian Indian Sikhs	NIH/NIDDK	Co-investigator	Total cost: \$27,372
7/2005- 6/2010	Mild Cognitive Impairment: A Prospective Community Study	NIH/NIA	Co-investigator	Total cost: \$292,694 Kamboh's budget
7/2003- 6/2007	R01 HL074165: Genetic & Structure-Function Studies: Paraoxonase in SLE	NIH/NHLBI	Principal Investigator	Total cost: \$1,646,430

1/2003-12/2007	R01 HL 070169: Functional Characterization of Lipoprotein Lipase Mutations	NIH/NHLBI	Principal Investigator	Total cost: \$1,373,798
5/1996-4/2007	R01/R02 AG13672: Risk Genes in Alzheimer's Disease	NIH/NIA	Principal Investigator	Total cost: \$2,817,625
1/1997-5/2007	R01/R02 HL54900: Molecular Genetics Analysis of Apolipoprotein H in SLE	NIH/NHLBI	Principal Investigator	Total cost: \$3,593,092
3/2001-2/2003	Prothrombin Gene Variation and the Risk of SLE and Thrombosis	Lupus Foundation of America	Principal Investigator	Direct cost: \$60,000
4/1995-3/2000	R01 HL52611: Genetics of CVD Risk Factors in Samoans	NIH/NHLBI	Principal Investigator	Total cost: \$1,556,924
5/1995-4/2000	P50 AG05133: Genetic Epidemiology of Alzheimer's Disease (sub-project of Program Project "University of Pittsburgh Alzheimer's Disease Research Center")	NIH/NIA	Co-Principal Investigator	Direct cost: \$529,534
4/1994-3/1999	R01 HL49074: Genetic Epidemiology of CHD Risk Factors in Blacks	NIH/NHLBI	Principal Investigator	Total cost: \$973,764
4/1995-3/1999	R01 AG09202: Indo-US Cross National Dementia Epidemiology Study	NIH/NIA	Co-investigator	Direct cost: \$200,000
1/1990-12/1995	Genetic Variation in Genes of Lipid Metabolism as Determinants of Lipoprotein/Lipid Levels in Man	National Dairy Promotion & Research Board	Principal Investigator	Direct cost: \$467,271
1/1993-12/1995	R01 HL45778: Genetic Epidemiology of Blood Lipids and Obesity: NGHS"	NIH/NHLBI	Co-investigator	Direct cost: \$90,630
4/1994-3/1995	R01 AG07562-06: Administrative Supplement to "Epidemiology of Dementia: A Prospective Community Study aka the Monongahela Valley Independent Elders Study"	NIH/NHLBI	Co-investigator	Direct cost: \$17,506
1/1994-12/1994	β_2 -Glycoprotein I Variability Among SLE Patients	Lupus Foundation of America	Principal Investigator	Direct cost: \$15,000

				Total cost:
5/1991- 6/2002	R01/R02 HL44672: Genetic Epidemiology of Lipoprotein-Lipid Levels	NIH/NHLBI	Principal Investigator	\$1,955,087
7/1990- 6/1991	Genetic Contribution in Determining Cholesterol Levels	Central Development Fund Award, University of Pittsburgh	Principal Investigator	Direct cost: \$7,195
6/1990- 5/1991	2 S07 RR05451-29: Genetic typing of LP(a)	BRSG, Univ. of Pittsburgh	Principal Investigator	Direct cost: \$18,432
9/1989- 8/1990	Genetics of Hemostatic Factors	Samuel and Emma Winters Foundation Research Grant	Principal Investigator	Direct cost: \$10,050
8/1988- 7/1989	2S07 RR05451-27: Direct Assessment of LDL-Receptor Mutations	BRSG, Univ. of Pittsburgh	Principal Investigator	\$7,500
12/1985- 1/1986	2S07 RR05451: Development of New Markers for the Development of Human Genetic Maps	BRSG, Univ. of Pittsburgh	Principal Investigator	\$5,596

PUBLICATIONS

1. Refereed Articles

1. Shami SA, Tahir AM, **Kamboh MI**: The study of natural selection on human head forms in a population of Punjab (Pakistan). *Biologia* 1978; 24:255-261.
2. Shami SA, **Kamboh MI**: Variation of ABO and Rh blood group gene frequencies in the populations of Punjab (Pakistan) I. *Biologia* 1978; 25:71-84.
3. Shami SA, **Kamboh, MI**: Geographic and age related variations in the gene frequencies of ABO and Rh blood groups in the populations of Punjab (Pakistan) II. *Biologia* 1979; 25:85-96.
4. **Kamboh MI**, Kirk RL: Distribution of transferrin (Tf) subtypes in Asian, Pacific and Australian Aboriginal populations: Evidence for the existence of a new subtype TfC6. *Hum Hered* 1983; 33:237-243.
5. **Kamboh MI**, Kirk RL, Clark P: Alpha-1-antitrypsin (PI) types in Asian, Pacific and Australian Aboriginal populations. *Dis Markers* 1983; 1:33-42.
6. **Kamboh MI**, Kirk RL: Investigation of PGM1 3, PGM1 6, PGM1 7 variants by isoelectric focusing. Evidence

- for new subtypes of the PGM1 3 and PGM1 7 alleles. *Hum Genet* 1983; 64:58-60.
7. **Kamboh MI:** Population genetic studies of PI, Tf, Gc and PGM1 subtypes among various caste groups in North India. *Acta Anthroponogenet* 1984; 8:159-179.
 8. **Kamboh MI,** Kirk RL: Genetic studies of PGM1 subtypes: Population data from the Asian-Pacific area. *Ann Hum Biol* 1984; 11:211-219.
 9. **Kamboh MI,** Kirwood C: Genetic polymorphism of thyroxin-binding globulin (TBG) in the Pacific area. *Am J Hum Genet* 1984; 36:646-654.
 10. **Kamboh MI,** Board PG, Kirwood C: Reevaluation of the proposed interrelationship between thyroxine-binding globulin (TBG) and alpha-1-antitrypsin (PI). *Clin Chim Acta* 1984; 139:65-73.
 11. **Kamboh MI,** Ranford PR, Kirk RL: Population genetics of the vitamin D-binding protein (GC) subtypes in the Asian-Pacific area. Description of new alleles at the GC locus. *Hum Genet* 1984; 67:378-384.
 12. **Kamboh MI:** Heterogeneity of factor XIIIIB: A new method for the determination of factor XIIIIB phenotypes by isoelectric focusing in 6 M urea. *Electrophoresis*. 1985; 6:185-186.
 13. **Kamboh MI:** Biochemical and genetic aspects of human serum alpha-1-proteinase inhibitor protein. *Dis Markers* 1985; 3:135-154.
 14. Musk AW, Zilko PJ, Manners P, Kay P, **Kamboh MI:** Genetic studies in familial fibrosing alveolitis – Possible linkage with immunoglobulin allotypes (Gm). *Chest* 1986; 89:206-210.
 15. **Kamboh MI,** Ferrell RE: A sensitive immunoblotting technique to identify thyroxin-binding globulin protein heterogeneity after isoelectric focusing. *Biochem Genet* 1986; 24:273-280.
 16. **Kamboh MI,** Ferrell RE: Ethnic variation in vitamin D-binding protein (GC): A review of isoelectric focusing studies in human populations. *Hum Genet* 1986; 72:281-293.
 17. **KambohMI,** Ferrell RE: Genetic studies of low abundance human plasma proteins. I. Microheterogeneity of zinc alpha 2-glycoprotein in biological fluids. *Biochem Genet* 1986; 24:849-857.
 18. **Kamboh MI,** Ferrell RE: Genetic studies of low abundance human plasma proteins. II. Population genetics of coagulation factor XIIIIB. *Am J Hum Genet* 1986; 39:817-825.
 19. **Kamboh MI,** Ferrell RE: Genetic studies of low abundance human plasma proteins. III. Polymorphism of the C1R subcomponent of the first complement component. *Am J Hum Genet* 1986; 39:826-831.
 20. **Kamboh MI,** Ferrell RE: Human transferrin polymorphism. *Hum Hered* 1987; 37:65-81.
 21. **Kamboh MI,** Ferrell RE: Analysis of the reported relationship between thyroxin-binding globulin and alpha-1-antitrypsin. *Biochem Genet* 1987; 25:175-179.
 22. **Kamboh MI,** Ferrell RE: Genetic studies of human apolipoproteins. I. Polymorphism of apolipoprotein A-IV. *Am J Hum Genet* 1987; 41:119-127.
 23. Escallon MH, Ferrell RE, **Kamboh MI:** Genetic studies of low abundance human plasma proteins. IV. Improved typing of alpha-1-acid glycoprotein (orosomucoid) by isoelectric focusing and immunoblotting.

Hum Hered 1987; 37:294-299.

24. Escallon MH, Ferrell RE, **Kamboh MI**: Genetic studies of low abundance human plasma proteins. V. Evidence for a second orosomucoid structural locus (ORM2) expressed in plasma. *Am J Hum Genet* 1987; 41:418-427.
25. **Kamboh MI**, Ferrell RE: Genetic studies of low abundance human plasma proteins.VI. Polymorphism of hemopexin. *Am J Hum Genet* 1987; 41:645-653.
26. **Kamboh MI**, Ferrell RE, Sepehrnia B: Genetic studies of human apolipoproteins. II. A rapid one-dimensional isoelectric focusing technique to characterize apolipoproteins A-I, A-II, A-IV and C-II of unfractionated plasma. *Electrophoresis* 1987; 8:355-358.
27. **Kamboh MI**, Ferrell RE: Genetic studies of low abundance human plasma proteins. VII. Heterogeneity of the C1S subcomponent of the first complement component. *J Immunogenet* 1987; 14:231-238.
28. **Kamboh MI**, Ferrell RE: Genetics studies of low abundance human plasma proteins. VIII. Inherited structural variation in antithrombin III. *Ann Hum Genet* 1988; 52:17-24.
29. DeCroo S, **Kamboh MI**, Leppert M, Ferrell RE: Isoelectric focusing studies of superoxide dismutase. Report of the unique SOD*2 allele in a U.S. white population. *Hum Hered* 1988; 38:1-7.
30. Sepehrnia B, **Kamboh MI**, Ferrell RE: Genetic studies of human apolipoproteins. III. Polymorphism of apolipoprotein C-II. *Hum Hered* 1988; 38:136-143.
31. **Kamboh MI**, Ferrell RE, Sepehrnia B: Genetic studies of human apolipoproteins. IV. Structural heterogeneity of apolipoprotein H (β_2 -glycoprotein-I). *Am J Hum Genet* 1988; 42:452-457.
32. **Kamboh MI**, Ferrell RE, Kottke B: Genetic studies of human apolipoproteins. V. A rapid novel procedure to screen apolipoprotein E polymorphism. *J Lipid Res* 1988; 29:1535-1543.
33. Sepehrnia B, **Kamboh MI**, Adams-Campbell L, Nwankwo M, Ferrell RE: Genetic studies of human apolipoproteins. VII. Population distribution of polymorphisms of apolipoproteins A-I, A-II, A-IV, C-II, E and H in Nigeria. *Am J Hum Genet* 1988; 43:847-853.
34. **Kamboh MI**, Lyons L, Ferrell RE: Genetic studies of low-abundance human plasma proteins. IX. A new allele at the complement subcomponent ClR structural locus. *Hum Genet* 1988; 81:93-94.
35. Ferrell RE, **Kamboh MI**, Sepehrnia B, Adams-Campbell LL, Weiss KM: Genetic variation in apolipoproteins C-II and C-III. *Adv Exp Med Biol* 1988; 243:81-85.
36. **Kamboh MI**, Ferrell RE: Genetic studies of low-abundance human plasma proteins. X. Coagulation factor XIIIIB variants in Blacks. *Electrophoresis* 1989; 10:53-57.
37. **Kamboh MI**, Sepehrnia B, Ferrell RE: Genetic studies of human apolipoproteins. VI. Common polymorphism of apolipoprotein E in Blacks. *Dis Markers* 1989; 7:49-55.
38. Lyons L, **Kamboh MI**, Ferrell RE: Genetic studies of low-abundance human plasma proteins. XI. Linkage analysis and population genetics of the C1S subcomponent of the first complement component. *Complement Inflamm* 1989; 6:81-87.
39. Kahl LE, **Kamboh MI**, Decroo S, Ferrell RE: Alpha-1-antitrypsin (PI) and vitamin D-binding globulin (GC) phenotypes in rheumatoid arthritis: Absence of an association. *Dis Markers* 1989; 7:71-78.

40. Eichner JE, Kuller LH, **Kamboh MI**, Ferrell RE: Phenotypic effects of apolipoproteins structure variation on lipid profiles. I. APO H and quantitative lipid measures in the Healthy Women Study. *Genet Epidemiol* 1989; 6:311-318.
41. Nakamura Y, Lathrop M, O'Connell P, Leppert M, **Kamboh MI**, Lalouel J-M, White R: Frequent recombination is observed in the distal end of the long arm of chromosome 14. *Genomics* 1989; 4:76-81.
42. Eichner JE, **Kamboh MI**, Cook T, Ferrell RE: Genetic studies of low-abundance human plasma proteins. XII. A new variant of corticosteroid binding globulin detected by IEF/immunoblotting. *Hum Hered* 1989; 39:170-173.
43. Sepehrnia B, **Kamboh MI**, Adams-Campbell LL, Bunker CH, Nwankwo M, Majumder PP, Ferrell RE: Genetic studies of human apolipoproteins. VIII. Role of the apolipoprotein H polymorphism in relation to serum lipoprotein concentrations. *Hum Genet* 1989; 82:118-122.
44. **Kamboh MI**, Lyons LA, Ferrell RE: Genetic studies of low-abundance human plasma proteins. XIII. Population genetics of C1R complement subcomponent and description of new variants. *Am J Hum Genet* 1989; 44:148-153.
45. **Kamboh MI**, Albers JJ, Majumder PP, Ferrell RE: Genetic studies of human apolipoproteins. IX. Apolipoprotein D polymorphism and its relation to serum lipoprotein lipid levels. *Am J Hum Genet* 1989; 45:147-154.
46. Eichner JE, Kuller LH, **Kamboh MI**, Ferrell RE: Phenotypic effects of apolipoproteins structure variation on lipid profiles. II. APO A-IV and quantitative lipid measures in the Healthy Women Study. *Genet Epidemiol* 1989; 6:493-499.
47. Sepehrnia B, **Kamboh MI**, Adams-Campbell LL, Bunker CH, Nwankwo M, Majumder PP, Ferrell RE: Genetic studies of human apolipoproteins. X. The effect of the apolipoprotein E polymorphism on quantitative levels of lipoproteins in Nigerian blacks. *Am J Hum Genet* 1989; 45:586-591.
48. Sepehrnia B, **Kamboh MI**, Adams-Campbell LL, Bunker CH, Nwankwo M, Majumder PP, Ferrell RE: Genetic studies of human apolipoproteins. XI. The effect of the apolipoprotein C-II polymorphism on lipoprotein levels in Nigerian Blacks. *J Lipid Res* 1989; 30:1349-1355.
49. Eichner JE, Kuller LH, Ferrell RE, **Kamboh MI**: Phenotypic effects of apolipoprotein structural variation on lipid profiles. IV. Apolipoprotein polymorphisms in a small group of black women from the Healthy Women Study. *Genet Epidemiol* 1989; 6:681-689.
50. **Kamboh MI**, Ferrell RE: Three F XIIIa gene loci? *Hum Genet* 1989; 84:102.
51. Ferrell RE, **Kamboh MI**, Majumder PP, Valdez R, Weiss KM: Genetic studies of human apolipoproteins. XIII. Quantitative polymorphism of apolipoprotein C-III in the Mayans of Yucatan Peninsula. *Hum Hered* 1990; 40:127-135.
52. Eichner JE, Friedrich CA, Cauley JA, **Kamboh MI**, Gutai JP, Kuller LH, Ferrell RE: Phenotypic effects of alpha 2-HS glycoprotein on quantitative hormone and bone measures in postmenopausal women. *Calcified Tissue Internatl* 1990; 47:345-349.
53. **Kamboh MI**, Ferrell RE: An improved method for complement subcomponent C1R typing. *J Forensic Sci*

1990; 35:190-192.

54. Eichner JE, Kuller LH, Ferrell RE, Meilahn EN, **Kamboh MI**: Phenotypic effects of apolipoproteins structural variation on lipid profiles. III. The contribution of APO E to prediction of total cholesterol, APO B and LDL-C in the Healthy Women Study. *Arteriosclerosis* 1990; 10:379-385.
55. **Kamboh MI**, Bhatia KK, Ferrell RE: Genetic studies of human apolipoproteins. XII. Population genetics of apolipoproteins in Papua New Guinea. *Am J Hum Biol* 1990; 2:17-23.
56. Ferrell RE, Sepehrnia B, **Kamboh MI**, VandeBerg JL: Highly polymorphic apolipoprotein A-IV locus in the baboons. *J Lipid Res* 1990; 31:131-135.
57. **Kamboh MI**, Kelly LJ, Ferrell RE: Genetic studies of human apolipoproteins. XIV. A simple agarose isoelectric focusing gel method for apolipoprotein E phenotyping. *Electrophoresis* 1990; 11:314-318.
58. **Kamboh MI**, Ferrell RE: Genetic studies of human apolipoproteins. XV. An overview of IEF-immunoblotting methods to screen apolipoprotein polymorphisms. *Hum Hered* 1990; 40:193-207.
59. **Kamboh MI**, Weiss KM, Ferrell RE: Genetic studies of human apolipoproteins. XVI. Apo E polymorphism and cholesterol levels in the Mayans of the Yucatan Peninsula, Mexico. *Clin Genet* 1991; 39:26-32.
60. Crews DE, **Kamboh MI**, Bindon JR, Ferrell RE: Genetic studies of human apolipoproteins. XVII. Population genetics of apolipoprotein polymorphisms in American Samoa. *Am J Phys Anthropol* 1991; 84:165-170.
61. **Kamboh MI**, Serjeantson SW, Ferrell RE: Genetic studies of human apolipoproteins. XVIII. Apolipoprotein polymorphisms in Australian Aborigines. *Hum Biol* 1991; 63:179-186.
62. Kelly LJ, **Kamboh MI**, Ferrell RE: Genetic studies of human apolipoproteins. XIX. Apolipoproteins E and A-IV phenotyping from whole blood and blood stains. *Applied Theoret Electrophoresis* 1991; 2:53-57.
63. Kaprio J, Ferrell RE, Kottke BA, **Kamboh MI**, Sing CF: Effects of polymorphisms in apolipoproteins E, A-IV and H on quantitative traits related to risk for cardiovascular disease. *Arterioscler Thromb* 1991; 11:1330-1348.
64. DeCroo S, **Kamboh MI**, Ferrell RE: Population genetics of alpha-1 antitrypsin polymorphism in US Whites, US Blacks and African Blacks. *Hum Hered* 1991; 41:215-221.
65. Eichner JE, Kuller LH, Ferrell RE, **Kamboh MI**: A simplified method to screen apolipoprotein E polymorphism. *Hum Hered* 1991; 41:61-64.
66. **Kamboh MI**, Ferrell RE: Apolipoprotein H polymorphism and its role in lipid metabolism. *Adv Lipid Res* 1991; 1:9-18.
67. Chakraborty R, **Kamboh MI**, Ferrell RE: "Unique" alleles in admixed populations: A strategy for determining "hereditary" population differences of disease frequencies. *Ethnicity Dis* 1991; 1:245-256.
68. **Kamboh MI**, Ferrell RE, Kottke B: Expressed hypervariable polymorphisms in apolipoprotein (a). *Am J Hum Genet* 1991; 49:1063-1074.
69. **Kamboh MI**, Harmony JAK, Sepehrnia B, Nwankwo M, Ferrell RE: Genetic studies of human apolipoproteins. XX. Genetic polymorphism of apolipoprotein J and its impact on quantitative lipid traits in normolipidemic subjects. *Am J Hum Genet* 1991; 49:1145-1154. [Published with an editorial; 49:1139-

1144].

70. **Kamboh MI**, Hamman RF, Iyengar S, Aston CE, Ferrell RE: Apolipoprotein A-IV polymorphism and its role in determining variation in lipoprotein-lipid, glucose and insulin levels in normal and non-insulin dependent diabetic individuals. *Atherosclerosis* 1991; 91:25-34.
71. Reilly SL, Ferrell RE, Kottke B, **Kamboh MI**, Sing CF: The gender specific apolipoprotein E genotype influence on the distribution of lipids and apolipoproteins in the population of Rochester, MN. I. Pleiotropic effects on means and variances. *Am J Hum Genet* 1991; 49:1155-1166.
72. Saha N, **Kamboh MI**, Kelly LJ, Ferrell RE, Tay JSH: Apolipoprotein H (β_2 -glycoprotein I) polymorphism in Asians. *Hum Biol* 1992; 64:617-621.
73. **Kamboh MI**, Iyengar S, Aston CE, Hamman RF, Ferrell RE: Apolipoprotein A-IV genetic polymorphism and its impact on quantitative traits in normoglycemic and non-insulin dependent diabetic Hispanics from the San Luis Valley, Colorado. *Hum Biol* 1992; 64:605-616.
74. Eichner JE, Ferrell RE, **Kamboh MI**, Kuller LH, Becker DJ, Drash AL, Stein EA, Orchard TJ: The impact of the apolipoprotein E polymorphism on the lipoprotein profile in insulin dependent diabetes. The Pittsburgh Epidemiology of Diabetes Complication Study. IX. *Metabolism* 1992; 41:347-351.
75. Chakraborty R, **Kamboh MI**, Nwankwo M, Ferrell RE: Caucasian genes in the American Blacks: New data. *Am J Hum Genet* 1992; 50:145-150.
76. Mailly F, Moll PP, Kottke BA, **Kamboh MI**, Humphries SE, Ferrell RE: Estimation of the frequency of isoform-genotype discrepancies at the apolipoprotein E locus in heterozygotes for the isoforms. *Genet Epidemiol* 1992; 9:239-248.
77. Ahn YI, **Kamboh MI**, Ferrell RE: Two new alleles in the tetranucleotide repeat polymorphism at the lipoprotein lipase (LPL) locus. *Hum Genet* 1992; 90:184.
78. **Kamboh MI**, Hamman RF, Ferrell RE: Two common polymorphisms in the APOA-IV coding gene: Their evolution and linkage disequilibrium. *Genet Epidemiol* 1992; 9:305-315.
79. Cleve H, Vogt U, **Kamboh MI**: Genetic polymorphism of apolipoprotein H (β_2 -glycoprotein I) in African Blacks from the Ivory Coast. *Electrophoresis* 1992; 13:849-851.
80. **Kamboh MI**, Williams ER, Law J, Aston CE, Bunker CH, Ferrell RE, Pollitzer WS: Molecular basis of a unique African variant (A-IV 5) of human apolipoprotein A-IV and its significance in lipid metabolism. *Genet Epidemiol* 1992; 9:379-388.
81. Ferrell RE, **Kamboh MI**: Interaction of apolipoprotein E genotype and dietary cholesterol in determining plasma cholesterol levels. *Am J Hum Genet* 1992; 50:236.
82. Crews DE, Bindon JR, **Kamboh MI**: Apolipoprotein polymorphisms and phenotypic variability in American Samoans: Preliminary data. *Am J Hum Biol* 1993; 5:39-48.
83. Crews DE, **Kamboh MI**, Mancilha-Carvalho JJ, Kottke B: Population genetics of apolipoproteins A-4, E and H polymorphisms in Yanomami Indians of northwestern Brazil: Associations with lipids, lipoproteins and carbohydrate metabolism. *Hum Biol* 1993; 65:211-224.

84. **Kamboh MI**, Aston CE, Ferrell RE, Hamman RF: Impact of the apolipoprotein E polymorphism in determining interindividual variation in total cholesterol and low density lipoprotein cholesterol in Hispanics and non-Hispanic Whites. *Atherosclerosis* 1993; 98:201-211.
85. Cauley JA, Eichner JE, **Kamboh MI**, Ferrell RE, Kuller LH, Black DM: Apo E allele frequencies in younger (age 42-50) vs. older (age 65-90) women. *Genet Epidemiol* 1993; 10:27-34.
86. Ahn YI, **Kamboh MI**, Hamman RF, Cole SE, Ferrell RE: Two DNA polymorphisms in the lipoprotein lipase gene and their association with factors related to cardiovascular disease. *J Lipid Res* 1993; 34:421-428.
87. **Kamboh MI**, Bunker CH, Nwankwo MU, Ferrell RE: Hemopexin: A unique polymorphism in populations of African ancestry. *Hum Biol* 1993; 65:655-660.
88. Saha N, **Kamboh MI**, Ahn YI, Tay JSH, Ferrell RE: Influence of the apolipoprotein H polymorphism on serum lipoprotein and apolipoprotein levels in two Asian populations. *Ethnicity Dis* 1993; 3:250-254.
89. Ahn YI, Ferrell RE, Hamman RF, **Kamboh MI**: Association of lipoprotein lipase gene variation with the physiological components of the Insulin Resistance Syndrome in the population of the San Luis Valley, Co. *Diabetes Care* 1993; 16:1502-1506.
90. Craig WY, Poulin SE, **Kamboh MI**: Apolipoprotein (a): A comparison of variants identified by SDS-PAGE or by SDS-agarose gel electrophoresis. *Electrophoresis* 1993; 14:1038-1041.
91. **Kamboh MI**, Svitko CM, Williams ER, Ferrell RE, Pollitzer WS: Hypervariable polymorphism of APO(a) in Blacks and Whites as reflected by phenotyping. *Chem Physics Lipids* 1994; 67/68:283-292.
92. **Kamboh MI**, Kelly LJ, Ahn YI, Ferrell RE: Genetic polymorphism of apolipoprotein A-IV in the chimpanzee – A common deletion of a conserved 12 nucleotides tandem repeat. *Human Biol* 1994; 66:625-638.
93. Ahn YI, **Kamboh MI**, Aston CE, Ferrell RE, Hamman RF: Role of common genetic polymorphisms in the LDL receptor gene in affecting plasma cholesterol levels in the general population. *Arterioscler Thromb* 1994; 14:663-670.
94. **Kamboh MI**, Friedlaender JS, Ahn YI, Ferrell RE: A common deletion polymorphism in the apolipoprotein A4 gene and its significance in lipid metabolism. *Arterioscler Thromb* 1994; 14:656-662.
95. Deka R, McGarvey ST, Ferrell RE, **Kamboh MI**, Yu LM, Aston CE, Jin L, Chakraborty R: Genetic characterization of American and Western Samoans. *Hum Biol* 1994; 66:805-822.
96. Islam S, Guitin B, Smith C, Treiber F, **Kamboh MI**: Association of apolipoprotein(a) phenotypes in children with family history of premature coronary artery disease. *Arterioscler Thromb* 1994; 14:1609-1616.
97. Rewers M, **Kamboh MI**, Hoag S, Shetterly SM, Ferrell RE, Hamman RF: Apolipoprotein A-IV polymorphism associated with myocardial infarction in obese NIDDM patients. The San Luis Valley Diabetes Study. *Diabetes* 1994; 43:1485-1489.
98. **Kamboh MI**, Aston CE, Hamman RF: The relationship of APOE polymorphism and cholesterol levels in normoglycemic and diabetic subjects in a biethnic population from the San Luis Valley, Colorado. *Atherosclerosis* 1995; 112:145-159.
99. **Kamboh MI**: Apolipoprotein E polymorphism and susceptibility to Alzheimer's disease. *Hum Biol* 1995;

67:195-215.

100. **Kamboh MI**, Wegenknecht DR, McIntyre JA: Heterogeneity of the apolipoprotein *H*3* allele and its role in affecting the binding of apolipoprotein H (β_2 -glycoprotein I) to anionic phospholipids. *Hum Genet* 1995; 95:385-388.
101. Ganguli M, Cauley JA, DeKosky ST, **Kamboh MI**: Dementia among elderly APOE 4/4 homozygotes: a prospective study. *Genet Epidemiol* 1995; 12:309-311.
102. **Kamboh MI**, Sanghera DK, Ferrell RE, DeKosky ST: *APOE*4*-associated Alzheimer's disease risk is modified by α_1 -antichymotrypsin polymorphism. *Nature Genet* 1995; 10:486-488.
103. **Kamboh MI**, Evans RW, Aston CE: Genetic effect of apolipoprotein[a] and apolipoprotein E polymorphisms on plasma quantitative risk factors for coronary heart disease in American Black Women. *Atherosclerosis* 1995; 117:73-81.
104. **Kamboh MI**, DeKosky ST: Apolipoprotein E genotyping in the diagnosis of Alzheimer's disease. *Ann Neurol* 1995; 38:967-969.
105. **Kamboh MI**, Crawford MH, Aston CE, Leonard WR: Population distributions of APOE, APOH and APOA4 polymorphisms and their relationship with quantitative plasma lipid levels among the Evenki Herders of Siberia. *Hum Biol* 1996; 68:231-243.
106. Marshall JA, **Kamboh MI**, Bessesen DH, Hoag S, Hamman RF, Ferrell RE: Association between dietary factors and serum lipids by apolipoprotein E polymorphism. *Am J Clin Nutr* 1996; 63:87-95.
107. Harris M, Sanghera DK, **Kamboh MI**: Two new alleles in the tetranucleotide polymorphism in the LDL-receptor protein (LRP) gene. *Clin Genet* 1996; 50:54-55.
108. DeKosky ST, Aston CE, **Kamboh MI**: Polygenic determinants of Alzheimer's disease: Modulation of the risk of Alzheimer's disease by alpha 1-antichymotrypsin. *Ann New York Acad Sci* 1996; 802:27-34.
109. Sanghera DK, Ferrell RE, Aston CE, McAllister AE, **Kamboh MI**, Kimm SYS: Quantitative effects of the apolipoprotein E polymorphism in a biracial sample of 9-10 year old girls. *Atherosclerosis* 1996; 126: 35-42.
110. Evans RW, Sankey SS, Hauth BA, Sutton-Tyrrell K, **Kamboh MI**, Kuller LH: Effect of sample storage on quantitation of lipoprotein(a) by an enzyme-linked immunosorbent assay. *Lipids* 1996; 31:1197-1203.
111. **Kamboh MI**, Aston CE, Svitko CM, McAllister AE, Hamman RF: Haplotype analysis of two APOA1/*MspI* polymorphisms in relation to plasma levels of apoA-I and HDL-cholesterol. *Atherosclerosis* 1996; 127:255-262.
112. Merriwether DA, Houston S, Iyengar S, Hamman R, Norris JM, Shetterly SM, **Kamboh MI**, Ferrell RE: Mitochondria versus nuclear admixture estimates demonstrate a past history of directional mating. *Am J Phys Anthropol* 1997; 102:153-159.
113. Sanghera DK, Saha N, Aston CE, **Kamboh MI**: Genetic polymorphism of paraoxonase and the risk of coronary heart disease. *Arterioscler Thromb Vas Biol* 1997; 17:1067-1073.
114. **Kamboh MI**, Sanghera DK, Aston CE, Bunker CH, Hamman RF, Ferrell RE, DeKosky ST: Gender-specific nonrandom association between the α_1 -antichymotrypsin and apolipoprotein E polymorphisms in the general population and its implication for the risk of Alzheimer's disease. *Genet Epidemiol* 1997; 14:169-

180.

115. Sanghera DK, Wagenknecht DR, McIntyre JA, **Kamboh MI**: Identification of structural mutations in the fifth domain of apolipoprotein H (β_2 -glycoprotein I) which affect phospholipid binding. *Hum Mol Genet* 1997; 6:311-316.
116. Saha N, Wang G, Vashist S, **Kamboh MI**: Influence of two APOA4 polymorphisms at codons 347 and 360 on non-fasting plasma lipids and apolipoproteins in Asian Indians. *Atherosclerosis* 1997; 131:249-255.
117. Sanghera DK, Kristensen T, Hamman RF, **Kamboh MI**: Molecular basis of the apolipoprotein H (β_2 -glycoprotein I) protein polymorphism. *Hum Genet* 1997; 100:57-62.
118. Anderson JL, Bunker CH, Aston CE, **Kamboh MI**: Relationship of two apolipoprotein B polymorphisms with serum lipoprotein and lipid levels in African Blacks. *Hum Biol* 1997; 69:793-807.
119. Lopez OL, **Kamboh MI**, Becker JT, Kaumfer DI, DeKosky ST: The apolipoprotein E4 allele is not associated with psychiatric symptoms or extrapyramidal signs in probable Alzheimer's disease. *Neurology* 1997; 49:794-797.
120. **Kamboh MI**, Rewers M, Aston CE, Hamman RF: Plasma apolipoprotein A-I, apolipoprotein B and lipoprotein(a) concentrations in normoglycemic Hispanics and non-Hispanic Whites from the San Luis Valley, Colorado. *Am J Epidemiol*; 1997; 146:1011-1018.
121. **Kamboh MI**, Aston CE, Ferrell RE, DeKosky ST: Genetic effect of α_1 -antichymotrypsin on the risk of Alzheimer's disease. *Genomics* 1997; 41:382-384.
122. Sanghera DK, Saha N, **Kamboh MI**: The codon 55 polymorphism in the paraoxonase 1 gene is not associated with the risk of coronary heart disease in Asian Indians and Chinese. *Atherosclerosis* 1998; 136:217-223.
123. Harris MR, Bunker CH, Hamman RF, Sanghera DK, Aston CE, **Kamboh MI**: Racial differences in the distribution of a low density lipoprotein receptor related protein (LRP) polymorphism and its association with serum lipid and apolipoprotein levels. *Atherosclerosis* 1998; 137:187-195.
124. Ali S, Bunker CH, Aston CE, Ukoli FA, **Kamboh MI**: Apolipoprotein (a) kringle 4 polymorphism and serum lipoprotein (a) concentrations in African Blacks. *Hum Biol* 1998; 70:477-490.
125. Sanghera DK, Aston CE, Saha N, **Kamboh MI**: DNA polymorphisms in two paraoxonase genes (PON1 and PON2) are associated with the risk of coronary heart disease. *Am J Hum Genet* 1998; 62:36-44. [Published with an editorial; 62:20-24].
126. Montoya SE, Aston CE, DeKosky ST, **Kamboh MI**, Lazo JS, Ferrell RE: Bleomycin hydrolase is associated with risk of Alzheimer's disease. *Nature Genet* 1998; 18:211-212.
127. **Kamboh MI**, Ferrell RE, DeKosky ST: Genetic association studies between Alzheimer's disease and two polymorphisms in the low density lipoprotein receptor-related protein gene. *Neurosci Lett* 1998; 244:85-88.
128. Lopez OL, Lopez-Pousa S, **Kamboh MI**, Adroer R, Oliva R, Becker JT, DeKosky ST: Apolipoprotein E polymorphism in Alzheimer's disease: A comparative study of two research populations from Spain and the United States. *Euro Neurol* 1998; 39:229-233.
129. Styren ST, **Kamboh MI**, DeKosky ST: Differential expression of immune factors in temporal cortex and cerebellum: The role of alpha-1-antichymotrypsin, apolipoprotein E and reactive glia in the progression of

Alzheimer's disease. *J Compar Neurol* 1998; 396:511-520.

130. Wang X, DeKosky ST, Wisniewski S, Aston CE, **Kamboh MI**: Genetic association of two chromosome 14 genes (presenilin 1 and α_1 -antichymotrypsin) with Alzheimer's disease. *Ann Neurol* 1998; 44:387-390.
131. **Kamboh MI**, Mehdi H: Genetics of apolipoprotein H (β_2 -glycoprotein I) and anionic phospholipid binding. *Lupus* 1998; 7 (suppl. 2): S10-S13.
132. Sweet RA, Nimgaonkar VL, **Kamboh MI**, Lopez OL, Zhang F, DeKosky ST: Dopamine receptor genetic variation, psychosis, and aggression in Alzheimer's disease. *Archiv Neurol* 1998; 55:1335-1340.
133. Kim HS, **Kamboh MI**: Genetic polymorphisms of apolipoproteins A-IV, E and H in Koreans. *Hum Hered* 1998; 48:313-317.
134. **Kamboh MI**, Aston CE, DeKosky ST: Association between ACT polymorphism and Alzheimer's disease. *Neurology* 1998; 50:574-576.
135. Okkels H, Rasmussen TE, Sanghera DK, **Kamboh MI**, Kristensen T: Structure of the human β_2 -glycoprotein I (apolipoprotein H) gene. *Eur J Biochem* 1999; 259:435-440.
136. **Kamboh MI**, Bunker CH, Aston CE, Nestlerode CS, McAllister AE, Ukoli FA: Genetic association of five apolipoprotein polymorphisms with serum lipoprotein-lipid levels in African blacks. *Genet Epidemiol* 1999; 16:205-222.
137. Nestlerode CS, Bunker CH, Sanghera DK, Aston CE, Ukoli FA, **Kamboh MI**: Apolipoprotein J polymorphisms and serum HDL-cholesterol levels in African blacks. *Hum Biol* 1999; 71:197-218.
138. Luedeking EK, Ganguli M, DeKosky ST, **Kamboh MI**: Genetic polymorphism in the persyn (γ -synuclein gene and the risk of Alzheimer's disease. *Neurosci Lett* 1999; 261:186-188.
139. Islam S, Gutin B, Treiber F, Hobbs G, **Kamboh MI**, M. Lopes-Virella: Association of apolipoprotein A phenotypes and oxidized low density lipoprotein-immune complexes in children. *Arch Pediatr Adolesc Med* 1999; 153:57-62.
140. **Kamboh MI**, Aston CE, Perez-Tur J, Kokmen E, Ferrell RE, Hardy J, DeKosky ST: A novel mutation in the apolipoprotein E gene (*APOE*4 Pittsburgh*) is associated with the risk of late-onset Alzheimer's disease. *Neurosci Lett* 1999; 263:129-132.
141. Mehdi H, Aston CE, Sanghera DK, Hamman RF, **Kamboh MI**: Genetic variation in the apolipoprotein H (β_2 -glycoprotein I) gene affects plasma apolipoprotein H concentrations. *Hum Genet* 1999; 105:63-71.
142. Kimm SYS, Pasagian-Macauley A, Aston CE, McAllister AE, Glynn NW, **Kamboh MI**, Ferrell RE: Correlates of lipoprotein(a) levels in a biracial cohort of young girls: The NHLBI Growth and Health Study. *J Pediatr* 1999; 135:169-176.
143. **Kamboh MI**, Manzi S, Mehdi H, Fitzgerald S, Sanghera DK, Kuller LH, Aston CE: Genetic variation in the apolipoprotein H (β_2 -glycoprotein I) affects the occurrence of antiphospholipid antibodies and apolipoprotein H concentrations in systemic lupus erythematosus. *Lupus* 1999; 8:742-750.
144. Saha N, Sanghera DK, **Kamboh MI**: The p22-phox polymorphism (C242T) is not associated with CHD risk in Asian Indians and Chinese. *Eur J Clin Invest* 1999; 29:999-1002.
145. Kerr ME, Kraus M, Marion D, **Kamboh MI**: Evaluation of apolipoprotein E genotypes on cerebral blood

- flow and metabolism following traumatic brain injury. *Adv Exp Med Biol* 1999; 471:117-124.
146. Mehdi H, Naqvi A, **Kamboh MI**: An hydrophobic sequence at positions 313-316 (Leu-Ala-Phe-Trp) in the fifth domain of apolipoprotein H (β_2 -glycoprotein I) is crucial for cardiolipin binding. *Eur J Biochem* 2000; 267:1770-1776.
 147. **Kamboh MI**, McGarvey ST, Aston CE, Ferrell RE, Bausserman L: Plasma lipoprotein(a) distribution and its correlates among samoans. *Hum Biol* 2000; 72:321-336.
 148. **Kamboh MI**, Aston CE, Hamman RF: DNA sequence variation in human apolipoprotein C4 gene and its effect on plasma lipid profile. *Atherosclerosis* 2000; 152:193-201.
 149. Ganguli M, Chandra V, **Kamboh MI**, Johnston JM, Dodge HH, Thelma BK, Juyal RC, Pandav R, Belle SH, DeKosky ST: *APOE* polymorphism and Alzheimer's disease: the Indo-US cross-national dementia study. *Archiv Neurol* 2000; 57:824-830.
 150. Johnston JM, Nazar-Stewart V, Kelsey SF, **Kamboh MI**, Ganguli M: Relationships between cerebrovascular events, *APOE* polymorphism, and Alzheimer's disease in a community sample. *Neuroepidemiol* 2000; 19:320-326.
 151. Chiu L, Hamman RF, **Kamboh MI**: Apolipoprotein(a) polymorphisms and plasma lipoprotein(a) concentrations in non-Hispanic whites and Hispanics. *Hum Biol* 2000; 72:821-835.
 152. Luedeking EL, DeKosky ST, Mehdi H, Ganguli M, **Kamboh MI**: Analysis of genetic polymorphisms in the transforming growth factor- β 1 gene and the risk of Alzheimer's disease. *Hum Genet* 2000; 106:565-569.
 153. Bhojak TJ, DeKosky ST, Ganguli M, **Kamboh MI**: Genetic polymorphisms in the cathepsin D and interleukin-6 genes and the risk of Alzheimer's disease. *Neurosci Lett* 2000; 288:21-24.
 154. Razzaghi H, Aston CE, Hamman RF, **Kamboh MI**: Genetic Screening of the lipoprotein lipase gene for mutations associated with high triglyceride/low HDL-Cholesterol levels. *Hum Genet* 2000; 107:257-267.
 155. Minster RL, DeKosky ST, Ganguli M, Belle S, **Kamboh MI**: Genetic association studies of interleukin-1 (IL-1A and IL-1B) and interleukin-1 receptor antagonist (IL-1RN) genes and the risk of Alzheimer's disease. *Ann Neurol* 2000; 48:817-818.
 156. Saha N, Aston CE, Low PS, **Kamboh MI**: Racial and genetic determinants of plasma factor XIII activity. *Genet Epidemiol* 2000; 19:440-455.
 157. Lopez OL, Becker JT, Klunk W, Saxton J, Hamilton RL, Kaufer DI, Sweet R, Cidis Meltzer C, Wisniewski S, **Kamboh MI**, DeKosky ST: Research evaluation and diagnosis of probable Alzheimer's disease over the last two decades I. *Neurology* 2000; 55:1854-1862.
 158. Lopez OL, Becker JT, Klunk W, Saxton J, Hamilton RL, Kaufer DI, Sweet R, Cidis Meltzer C, Wisniewski S, **Kamboh MI**, DeKosky ST: Research evaluation and diagnosis of possible Alzheimer's disease over the last two decades II. *Neurology* 2000; 55: 1863-1869.
 159. Pfaff CL, Parra EJ, Bonilla C, Heister K, McKeigue PM, **Kamboh MI**, Hutchinson RG, Ferrell RE, Boerwinkle E, Shriver MD: Population structure in admixed populations: Effects of admixture dynamics on the pattern of linkage disequilibrium. *Am J Hum Genet* 2001; 68:198-207.

160. DePrince KM, McGarvey ST, McAllister AE, Bausserman L, Aston CE, Ferrell RE, **Kamboh MI**: Genetic effect of two Apo(a) repeat polymorphisms (kringle 4 and pentanucleotide repeats) on plasma Lp(a) levels in American Samoans. *Hum Biol* 2001; 73:91-104.
161. Parra EJ, Kittles RA, Argyropoulos G, Pfaff CL, Hiester K, Bonilla C, Sylvester N, Parrish-Gause D, Garvey WT, Jin L, McKeigue PM, **Kamboh MI**, Ferrell RE, Pollitzer WS, Shriner MD: Ancestral proportions and admixture dynamics in geographically defined African Americans living in South Carolina. *Am J Phys Anthropol* 2001; 114:18-29.
162. Wang X, Lueddecking EK, Minster RL, Ganguli M, DeKosky ST, **Kamboh MI**: Lack of association between α 2-macroglobulin polymorphisms and Alzheimer's disease. *Hum Genet* 2001; 108:105-108.
163. Evans RW, Shpilberg O, Shaten BJ, Ali S, **Kamboh MI**, Kuller LH: Prospective association of lipoprotein(a) concentrations and apo(a) size with coronary heart disease among men in the Multiple Risk Factors Intervention Trial. *J Clin Epidemiol* 2001; 54:51-57.
164. Nebes RD, Vora IJ, Meltzer CC, Fukui MB, Williams RL, **Kamboh MI**, Saxton, Houck PR, DeKosky ST, Reynolds CF, III: The relation of deep white matter hyperintensities and non-depressed older adults. *Am J Psychiatry* 2001; 158:878-884.
165. Sanghera DK, Nestlerode CS, Ferrell RE, **Kamboh MI**: Chimpanzee apolipoprotein H (β 2-glycoprotein I): Report on the gene structure, a common polymorphism and a high prevalence of antiphospholipid antibodies. *Hum Genet* 2001; 109:63-72.
166. Razzaghi H, **Kamboh MI**: VGT-SSCP: A highly sensitive and non-radioactive mutation detection method based on vertical temperature SSCP. *Electrophoresis* 2001; 22:2665-2669.
167. Bhojak TJ, DeKosky ST, Ganguli M, **Kamboh MI**: Genetic polymorphism in the cathepsin G gene and the risk of Alzheimer's disease. *Neurosci Lett* 2001; 309:138-140.
168. Razzaghi H, Day BW, McClure RJ, **Kamboh MI**: Structure-function analysis of D9N and N291S mutations in human apolipoprotein lipase using molecular modeling. *J Mol Graphics Modell* 2001; 19:487-494.
169. Sweet RA, **Kamboh MI**, Wisniewski SR, Lopez OL, Klunk WE, Kaufer DI, DeKosky ST: APOE and ACT genotypes do not predict time to psychosis in Alzheimer's disease. *J Geriatr Psychiatry Neurol* 2002; 15:24-30.
170. Wang X, DeKosky ST, Ikonomovic M, **Kamboh MI**: Distribution of plasma α 1-antichymotrypsin levels in Alzheimer's disease patients and their genetic controls. *Neurobiol Aging* 2002; 23:377-382.
171. Romero LJ, Schuyler M, **Kamboh MI**, Quals C, LaRue A, Liang HC, Rhyne R: The APO E4 allele and cognition in New Mexico Hispanic Elderly. *Ethnicity Dis* 2002; 12:235-241.
172. Desai PP, Bunker CH, Ukoli FAM, **Kamboh MI**: Genetic variation in the apolipoprotein D gene among African blacks and its significance in lipid metabolism. *Atherosclerosis* 2002; 163:329-338.
173. Wang X, DeKosky ST, Lueddecking-Zimmer E, Ganguli M, **Kamboh MI**: Genetic variation in α 1-antichymotrypsin and its association with Alzheimer's disease. *Hum Genet* 2002; 110:356-365.
174. Desai PP, DeKosky ST, **Kamboh MI**: Genetic variation in the cholesterol 24-hydroxylase (CYP46) gene and the risk of Alzheimer's disease. *Neurosci Lett* 2002; 328:9-12.

175. Luedeking-Zimmer E, DeKosky ST, Chen Q, Barmada MM, **Kamboh MI**: Investigation of oxidized LDL-receptor 1 (OLR1) as the candidate gene for Alzheimer's disease on chromosome 12. *Hum Genet* 2002; 111:443-451.
176. Johson JM, Nazar-Stewart V, Kelsey SF, **Kamboh MI**, Ganguli M: Relationship between cerebrovascular events, APOE polymorphism and Alzheimer's disease in a community sample. *Dementia Rev J* 2002; 4: 8-9.
177. Desai PP, Hendrie HC, Evans RM, Murrell JR, DeKosky ST, **Kamboh MI**: Genetic variation in apolipoprotein D affects the risk of Alzheimer's disease in African Americans. *Am J Med Genet (Neuropsychiatr Genet)* 2003; 116B:98-101.
178. DeKosky ST, Ikonomovic MD, Wang X, Farlow M, Wisniewski S, Lopez OL, Becker JT, Saxton J, Klunk WE, Sweet R, Kaufer DI, **Kamboh MI**: Plasma and cerebrospinal fluid α 1-antichymotrypsin levels in Alzheimer's disease: Correlation with cognitive impairment. *Ann Neurol* 2003; 53:81-90.
179. Luedeking-Zimmer E, DeKosky ST, Nebes R, **Kamboh MI**: Association of the 3'UTR transcription factor LBP-1c/CP2/LSF polymorphism with late-onset Alzheimer's disease. *Am J Med Genet (Neuropsychiatr Genet)* 2003; 117B:114-117.
180. Chen Q, Reis SE, Kammerer CM, McNamara DM, Holubkov R, Sharaf BL, Sopko G, Pauley DF, Merz CNB, **Kamboh MI**, for the WISE study group: Association between the severity of angiographic coronary artery disease and paraoxonase gene polymorphisms in the NHLBI-sponsored Women's Ischemia Syndrome Evaluation (WISE) study. *Am J Hum Genet* 2003; 72:13-22.
181. Mehdi H, Manzi S, Desai P, Chen Q, Nestlerode C, Bontempo F, Strom SC, Zarnegar R, **Kamboh MI**: A functional polymorphism at the transcriptional initiation site in β 2-glycoprotein I (apolipoprotein H) associated with reduced expression and lower plasma levels of β 2-glycoprotein I. *Eur J Biochem* 2003; 270:230-238.
182. Wang GQ, DiPetro M, Roeder K, Heng C-K, Bunker CH, Hamman RF, **Kamboh MI**: Cladistic analysis of human apolipoprotein A4 polymorphisms in relation to quantitative plasma lipid risk factors of coronary heart disease. *Ann Hum Genet* 2003; 67:107-124.
183. Lambert J-C, Luedeking-Zimmer, E, Merrot S, Hayes A, Thaker U, Desai P, Houzet A, Hermant X, Cotte D, Pritchard A, Iwatsubo T, Pasquier F, Frigard B, Conneally PM, Chartier-Harlin M-C, DeKosky ST, Lendon C, Mann D, **Kamboh MI**, Amouyel P: Association of 3'-UTR polymorphisms of the oxidized LDL-receptor 1 (OLR1) gene with Alzheimer's disease. *J Med Genet* 2003; 40:424-430.
184. Chen, Q, Reis SE, Kammerer C, Craig WY, LaPierre SE, Zimmer EL, McNamara DM, Pauly DF, Sharaf B, Holubkov R, Merz CNB, Sopko G, Bontempo F, **Kamboh MI**: Genetic variation in the lectin-like oxidized LDL receptor (LOX1) gene and the risk of coronary artery disease. *Circulation* 2003; 107:3146-3151.
185. Chen Q, Reis SE, Kammerer CM, McNamara DM, Holubkov R, Sharaf BL, Sopko G, Pauley DF, Merz CNB, **Kamboh MI**, for the WISE study group: APOE polymorphism in angiographic coronary artery disease severity in the Women's Ischemia Syndrome Evaluation (WISE) study. *Atherosclerosis* 2003; 169:159-167.

186. Hokanson JE, **Kamboh MI**, Scarboro S, Eckel RH, Hamman RF: Effect of the hepatic lipase gene on coronary heart disease risk. *Am J Epidemiol* 2003; 158:836-843.
187. Kerr ME, **Kamboh MI**, Yookyoung K, Kraus MF, Puccio AM, DeKosky ST, Marion DW: Relationship between apoE4 allele and excitatory amino acid levels after traumatic brain injury. *Crit Care Med* 2003; 31: 2371-2379.
188. Hope C, Mettenburg J, Gonias SL, DeKosky ST, **Kamboh MI**, Chu CT: Functional analysis of plasma α 2-macroglobulin from Alzheimer's disease patients with *A2M* intronic deletion. *Neurobiol Dis* 2003; 14: 504-512.
189. Butters MA, Sweet RA, Mulsant BH, **Kamboh MI**, Pollock BG, Begley AE, Reynolds III CF, DeKosky ST: APOE is associated with age-of-onset, but not cognitive functioning, in late-life depression. *Int J Geriatr Psychiatry* 2003; 18:1075-1081.
190. Crews DE, Fitton LJ, Kottke BA, **Kamboh MI**: Population genetics of apolipoproteins A-IV, E, H and the angiotensin converting enzyme (ACE): Association with lipids, and apolipoprotein levels in American Samoans. *Am J Phys Anthropol* 2004; 124:364-372.
191. Heng C-K, Lal S, Saha N, Low P-S, **Kamboh MI**: The impact of factor XIIIa V34L polymorphism on plasma factor XIII activity in the Chinese and Asian Indians from Singapore. *Hum Genet* 2004; 114: 186-
192. **Kamboh MI**, Sanghera DK, Mehdi H, Nestlerode CS, Chen Q, Khalifa O, Naqvi A, Bunker CH: Single nucleotide polymorphisms in the coding region of the apolipoprotein H (β 2-glycoprotein I) gene and their correlation with the protein polymorphism, anti- β 2 glycoprotein I antibodies and cardiolipin binding: Description of novel haplotypes and their evolution. *Ann Hum Genet* 2004; 68:285-299.
193. **Kamboh MI**: Molecular genetics of late-onset Alzheimer's disease. *Ann Hum Genet* 2004; 68:381-404.
194. Sanghera DK, Manzi S, Bontempo F, Nestlerode C, **Kamboh MI**: Role of an intronic polymorphism in the *PDCD1* gene with the risk of sporadic systemic lupus erythematosus and the occurrence of antiphospholipid antibodies. *Hum Genet* 2004; 115: 393-398.
195. Ozturk A, Desai P, Minster RL, DeKosky ST, **Kamboh MI**: Three SNPs in the GSTO1, GSTO2 and PRS11 genes on chromosome 10 are not associated with age-at-onset of Alzheimer's disease. *Neurobiol Aging* 2005; 26:1161-1165.
196. Tsuang D, Wilson R, Lopez OL, Luedeking-Zimmer EK, Leverenz JB, DeKosky ST, **Kamboh MI**, Hamilton RL: Genetic Association between *APOE*4* allele and Lewy bodies in Alzheimer's disease. *Neurology* 2005; 64: 509-513
197. Desai P, Nebes R, DeKosky ST, **Kamboh MI**: Investigation of the effect of brain-derived neurotrophic factor (BDNF) polymorphism on the risk of late-onset Alzheimer's disease (AD) and quantitative measures of AD progression. *Neurosci Lett* 2005; 379: 229-234.
198. Omalu BI, DeKosky ST, Minster RL, **Kamboh MI**, Hamilton RL, Wecht CH: Chronic traumatic encephalopathy (CTE) in a National Football League player. *Neurosurgery* 2005; 57: 128-134.

199. Desai PP, Ikonomovic MD, Abrahamson EE, Hamilton RL, Isanski BA, Hope CE, Klunk WE, DeKosky ST, **Kamboh MI**: Apolipoprotein D is a component of compact but not diffuse amyloid-beta plaques in Alzheimer's disease temporal cortex. *Neurobiol Dis* 2005; 20: 574-582.
200. Chen Q, **Kamboh MI**: Complete DNA sequence variation in the apolipoprotein H (β 2-glycoprotein I) gene and identification of informative SNPs. *Ann Hum Genet* 2005; 70: 1-11.
201. **Kamboh MI**, Minster RL, Kenney M, Ozturk A, Desai PP, Kammerer CM, DeKosky ST: Alpha-1-antichymotrypsin (ACT or SERPINA3) polymorphism may affect age-at-onset and disease duration of Alzheimer's disease. *Neurobiol Aging* 2006; 27:1435-1439.
202. Ozturk A, DeKosky ST, **Kamboh MI**: Genetic variation in the choline acetyltransferase (CHAT) gene may be associated with the risk of Alzheimer's disease. *Neurobiol Aging* 2006; 27:1440-1444.
203. **Kamboh MI**, Minster RL, Feingold E, DeKosky ST: Genetic association of ubiquilin with Alzheimer's disease and related quantitative measures. *Mol Psychiatry* 2006; 11:273-279.
204. Sanghera DK, Bhatti JS, Bhatti GK, Ralhan SK, Wander GS, Singh JR, Bunker CH, Weeks DE, **Kamboh MI**, Ferrell RE: The Khatri Sikh Diabetes Study (SDS): Study design, methodology, sample collection and initial results. *Hum Biol* 2006; 78: 43-63.
205. Tripi LM, Manzi S, Chen Q, Kenney M, Shaw P, Kao A, Bontempo F, Kammerer C, **Kamboh MI**: Relationship of serum paraoxonase (PON1) activity and PON1 genotype with the risk of systemic lupus erythematosus. *Arthritis Rheum* 2006; 54:1928-1939.
206. Ozturk A, DeKosky ST, **Kamboh MI**: Lack of association of 5 SNPs in the vicinity of the insulin-degrading enzyme (IDE) gene with late-onset Alzheimer's disease. *Neurosci Lett* 2006; 406:265-269.
207. Minster RL, DeKosky ST, **Kamboh MI**: Lack of association of two chromosomes 10q24 SNPs with Alzheimer's disease. *Neurosci Lett* 2006; 408: 170-172.
208. Kerr ME, **Kamboh MI**, Kong Y, Alexander S, Yonas H: Apolipoprotein E genotype and CBF in traumatic brain injured patients. *Adv Exp Med Bio*. 2006; 578: 291-296.
209. Demirci FYK, Manzi S, Ramsey-Goldman R, Minster RL, Kenney M, Shaw PS, Dunlop-Thomas CM, Kao AH, Rhew E, Bontempo F, Kammerer C, **Kamboh MI**: Association of a common interferon regulatory factor 5 (IRF5) variant with increased risk of systemic lupus erythematosus (SLE). *Ann Hum Genet* 2006; 71: 308-311.
210. Omalu BI, DeKosky ST, Hamilton RL, Minster RL, **Kamboh MI**, Shakir AM, Wecht CH. Chronic traumatic encephalopathy in a National Football League player: part II. *Neurosurgery* 2006; 59: 1086-1093.
211. Sundar PD, Feingold E, Minster R, DeKosky ST, **Kamboh MI**: Gender-specific association of ATP-binding cassette transporter 1 (ABCA1) polymorphisms with the risk of late-onset Alzheimer's disease. *Neurobiol Aging* 2007; 28:856-862.
212. Ozturk A, Minster RL, DeKosky ST, **Kamboh MI**: Association of tagSNPs in the urokinase-plasminogen activator (PLAU) gene with Alzheimer's disease and associated quantitative traits. *Am J Med Genet Part B* 2007; 144: 79-82.

213. Lambert JC, Ferreira S, Gussekloo J, Christiansen L, Brysbaert G, Slagboom PE, Cottel D, Petit T, Hauw JC, DeKosky ST, Richard F, Berr C, Lendon CL, **Kamboh MI**, Mann D, Christensen K, Westendorp R, Amouyel P: Evidence for the association of the S100 beta gene with low cognitive performance and dementia in the elderly. *Mol Psychiatry* 2007; 12: 870-880.
214. Alexander S, Kerr ME, Kim Y, **Kamboh MI**, Beers SR, Conley YP: Apolipoprotein E4 allele presence and functional outcome after severe traumatic brain injury. *J Neurotrauma* 2007; 24: 790-797.
215. Demirci FYK, Manzi S, Ramsey-Goldman R, Kenney M, Shaw PS, Dunlop-Thomas CM, Kao A, Rhew EY, Bontempo F, Kammerer C, **Kamboh MI**: Association study of toll-like receptor 5 (TLR5) and toll-like receptor 9 (TLR9) polymorphisms in systemic lupus erythematosus (SLE). *J Rheumatology* 2007; 34: 1708-1711.
216. Minster RL, DeKosky ST, **Kamboh MI**: No association of dynamin binding protein (DNMBP) gene SNPs and Alzheimer's disease. *Neurobiol Aging* 2008; 29:1602-1604.
217. Sanghera DK, Manzi S, Minster RL, Shaw P, Kao A, Bontempo F, **Kamboh MI**: Genetic variation in the paraoxonase-3 (PON3) gene is associated with serum activity. *Ann Hum Genet* 2008; 72: 72-81.
218. **Kamboh MI**: Genome-wide association studies in genomic medicine – Are we there yet? *J Dow Univ Hlth Sci* 2008; 2: 88-90.
219. Chen Q, Razzaghi H, Demirci FY, **Kamboh MI**: Functional significance of lipoprotein lipase *HindIII* polymorphism associated with the risk of coronary artery disease. *Atherosclerosis* 2008; 200: 102-108.
220. Hom G, Graham RR, Modrek B, Ortman W, Taylor KE, Garnier S, Lee AT, Chung SA, Ferreira R, Pant PVK, Ballinger DG, Kosoy R, Demirci FY, **Kamboh MI**, Kao AH, Tian C, Gunnarsson I, Bengtsson AA, Rantapaa-Dahlqvist S, Petri M, Manzi M, Seldin MF, Ronnblom L, Syvanene A-C, Criswell LA, Gregersen PK, Behrens TW: Association of systemic lupus erythematosus with *C8orf13/BLK* and *ITGAM-ITGAX*. *N Engl J Med* 2008; 358: 900-909.
221. Mehdi H, Naqvi A, **Kamboh MI**: Recombinant hepatitis B surface antigen and anionic phospholipids share a binding region in the fifth domain of β 2-glycoprotein I (apolipoprotein H). *Biochim Biophys Acta – Mol Basis Dis* 2008; 782: 163-168.
222. Sanghera DK, Nath SK, Ortega L, Gambarelli M, Kim-Howard X, Singh J, Ralhan SK, Wander GS, Mehra NK, Mulhivill JJ, **Kamboh MI**: TCFL2 polymorphism are associated with type 2 diabetes in Khatri Sikhs from north India: Genetic variation affects lipid levels. *Ann Hum Genet* 2008; 72: 499-509.
223. Sanghera DK, Ortega LM, Han S, Singh JR, Ralhan SK, Wander GS, Mehra NK, Mulvihill JJ, Ferrell RE, Nath SK, **Kamboh MI**: Impact of nine common type 2 diabetes risk polymorphisms in Asian Indian Sikhs: PPARG2 (Pro12Ala), IGF2BP2, TCF7L2 and FTO variants confer a significant risk. *BMC Med Genet* 2008; 9: 59.
224. Taylor KE, Remmers EF, Lee AT, Ortmann WA, Plenge RM, Tian C, Chung SA, Nititham J, Hom G, Kao AH, Demirci FY, **Kamboh MI**, Petri M, Manzi S, Kastner DL, Seldin MF, Gregersen PK, Behrens TW, Criswell LA: Specificity of the *STAT4* genetic association with severe disease manifestations of systemic lupus erythematosus. *PLoS Genet* 2008; 4(5): e1000084.

225. Minster RL, DeKosky ST, **Kamboh MI**: No association of SORL1 SNPs with Alzheimer's disease. *Neurosci Lett* 2008; 440: 190-192.
226. Musone SL, Taylor KE, Lu T, Nititham J, Ferreira RC, Ortmann W, Shifrin N, Petri MA, **Kamboh MI**, Manzi S, Seldin MF, Gregersen PK, Behrens TW, Ma A, Kwok P-Y, Criswell LA: Multiple polymorphisms in the TNFAIP3 region are independently associated with systemic lupus erythematosus. *Nature Genet*; 2008; 40: 1062-1064.
227. Shih PB, Manzi S, Shaw P, Kenny M, Kao AH, Bontempo F, Barmada MM, Kammerer C, **Kamboh MI**: Genetic variation in the C-reactive protein (CRP) gene may be associated with the risk of systemic lupus erythematosus and CRP levels. *J Rheumatology* 2008; 35: 2171-2178.
228. Minster RL, DeKosky ST, **Kamboh MI**: No association of DAPK1 and ABCA2 SNPs on chromosome 9 with Alzheimer's disease. *Neurobiol Aging* 2009; 30:1890-1891.
229. Figgins JA, Minster RL, Demirci FY, DeKosky ST, **Kamboh MI**: Association studies of 22 candidate SNPs with late-onset Alzheimer's disease. *Am J Med Genet Part B* 2009; 150: 520-526.
230. Suresh S, Demirci FY, Jacobs E, Kao AH, Rhew ER, Sanghera SK, Seltzer F, Sutton-Tyrrell K, McPherson D, Bontempo FA, Kammerer CM, Ramsey-Goldman R, Manzi S, **Kamboh MI**: APOH polymorphisms in relation to lupus and lupus-related phenotypes. *J Rheumatology* 2009; 36: 315-322.
231. Conley YP, Mukherjee A, Kammerer C, DeKosky ST, **Kamboh MI**, Finegold DN, Ferrell RE: Evidence supporting a role for the calcium sensing receptor in Alzheimer disease. *Am J Med Genet B (Neuropsychiatr Genet)* 2009; 150B: 703-709.
232. Chapuis J, Hansmannel F, Kerdraon O, Ferreira S, Hubans C, Maurage CA, Huot L, Bensemain F, Laumet G, Ayral AM, Fievet N, Hauw JJ, DeKosky ST, Lemoine Y, Iwatsubo T, Wavrant-Devrieze F, Dartigues JF, Tzourio C, Buee L, Pasquier F, Berr C, Mann D, Lendon C, Alperovitch A, **Kamboh MI**, Amouyel P, Lambert JC: Transcriptomic and genetic studies identify IL33 as a candidate gene for Alzheimer's disease. *Mol Psychiatry* 2009; 14: 1004-1016.
233. Minster RL, Demirci FY, DeKosky ST, **Kamboh MI**: No association between CALHM1 variation and risk of Alzheimer disease. *Hum Mutation* 2009; 30: E566-E569.
234. Yang L, Razzaghi H, Hokanson JE, **Kamboh MI**: Identification and characterization of a novel 5 bp deletion in a putative insulin response element in the lipoprotein lipase gene. *Biochim Biophys Acta - Molecular and Cell Biology of Lipids* 2009; 1791: 1057-1065.
235. Demirci FY, Dressen AS, Hamman RF, Bunker CH, Kammerer CM, **Kamboh MI**: Association of a common G6PC2 variant with fasting plasma glucose levels in non-diabetic individuals. *Ann Nutr Metab* 2009; 56: 59-64.
236. Heng C-K, He X, Saha N, Low PS, Demirci FY, **Kamboh MI**: Association of three lipoprotein lipase polymorphisms with coronary artery disease in Chinese and Asian Indians. *Int J Cardiol* 2010; 144: 412-413.
237. Fukumoto N, Fujii T, Combarros O, **Kamboh MI**, Tsai S-J, Matsushi S, Naemias B, Comings DE, Arboleda H, Ingelsson M, Hyman BT, Akatsu H, Grupe A, Nishimura AL, Zatz M, Mattila KM, Rinne J, Goto Y, Asada T, Nakamura S, Kunugi H: Sexually dimorphic effect of BDNF on susceptibility to Alzheimer's

disease: New data and meta analysis. *Am J Med Genet Part B* 2010; 153B: 235-242.

238. D'Angelo GM, **Kamboh MI**, Feingold E: A likelihood-based approach for missing genotype data. *Hum Hered* 2010; 69: 171-183.
239. Sanghera DK, Demirci FY, Been L, Ortega L, Ralhan SK, Wander GS, Mehra NK, Singh J, Aston CE, Mulvihill JJ, **Kamboh MI**: PPARG and ADIPOQ gene polymorphisms increase type 2 diabetes risk in Asian Indian Sikhs. Pro12Ala still remains as the strongest predictor. *Metabolism* 2010; 59: 492-450.
240. Suresh S, Demirci FY, Lefterov I, Kammerer C, Ramsey-Goldman R, Manzi S, **Kamboh MI**: Functional and genetic characterization of the promoter region of apolipoprotein H (β 2 glycoprotein-I). *FEBS Journal* 2010; 277: 941-963.
241. Omalu BI, Hamilton RL, **Kamboh MI**, DeKosky ST, Bailes J: Chronic traumatic encephalopathy (CTE) in a National Football League Player: Case report and emerging medicolegal practice questions. *J Forensic Nurs* 2010; 6: 40-46.
242. Hansmannel F, Sillaire A, **Kamboh MI**, Lendon C, Pasquier F, Hannequin D, Laumet G, Mounier A Ayral A-M, DeKosky ST, Hauw J-J, Berr C, Mann D, Amouyel P, Campion D, Lambert J-C: Is the urea cycle involved in Alzheimer's disease? *J Alzheimer's Dis* 2010; 21:1013-1021. [PMID: 20693631 - PMCID: PMC2945690]
243. Lambert J-C, Sleegers K, González-Pérez A, Ingelsson M, Gary W Beecham GW, Hiltunen M, Combarros O, Bullido MJ, Brouwers N, Bettens K, Berr C, Pasquier F, Richard F, DeKosky ST, Hannequin D, Haines JL, Tognoni G, Fiévet N, Dartigues J-F, Tzourio C, Engelborghs S, Arosio B, Coto E, Deyn PD, Zompo MD, Mateo I, Boada M, Antunez C, Lopez-Arrieta J, Epelbaum J, Schjeide BM, Frank-Garcia A, Giedraitis V, Helisalmi S, Porcellini E, Pilotto A, Forti P, Ferri R, Delepine M, Zelenika D, Lathrop M, Scarpini E, Siciliano G, Solfrizzi V, Sorbi S, Spalletta G, Ravaglia G, Valdivieso F, Vepsäläinen S, Alvarez V, Bosco P, Mancuso M, Panza F, Naemias B, Bossù P, Hanon O, Piccardi P, Annoni G, Mann D, Marambaud P, Seripa D, Galimberti D, Tanzi RE, Bertram L, Lendon C, Lannfelt L, Licastro F, Campion D, Pericak-Vance MA, Soininen H, Van Broeckhoven C, Alpérovitch A, Ruiz A, **Kamboh MI**, Amouyel P: The CALHM1 P86L polymorphism is a genetic modifier of age at onset in Alzheimer's disease : a meta-analysis study. *J Alzheimer's Dis* 2010; 22:247-255. [PMID: 20847397 - PMCID: PMC2964875]
244. Reynolds CF III, Butters MA, Lopez O, Pollock BG, Dew MA, Mulsant BH, Lenze EJ, Holm M, Rogers JC, Mazumder S, Houck PR, Begley A, Anderson S, Karp JF, Miller MD, Whyte EM, Stack J, Gildengers A, Szanto K, Bensasi S, Kaufer DI, **Kamboh MI**, DeKosky ST: Maintenance treatment of depression in old age: Efficacy and adverse effects of donepezil combined with antidepressant pharmacotherapy. *Arch Gen Psychiatry* 2011; 8:51-60
245. Miller MA, Conley Y, Scanlon JM, Ren DR, **Kamboh MI**, Niyonkuru C, Wagner, AK: APOE Genetic Associations with Seizure Development after Severe Traumatic Brain Injury. *Brain Injury* 2010; 24:1468-1477.
246. **Kamboh MI**, Minster RL, Demirci FY, Ganguli M, DeKosky ST, Lopez OL, Barmada MM: Association of CLU and PICALM variants with Alzheimer's disease. *Neurobiol Aging* 2012; 33:518-521 [Epub June 8, 2010].

247. Rietz C, Cheng R, Rogaeva E, Lee JH, Tokuhiro S, Bettens K, Sleegers K, Tan EK, Kimura R, Shibata N, **Kamboh MI**, Prince JA, Maier W, Riemenschneider M, Owen M, Harold D, Hollingworth P, Cellini E, Takeda M, Pericak-Vance MA, Younkin S, Williams J, van Broeckhoven C, Farrer LA, St.George-Hyslop P, Mayeux R: A meta-analysis of the association between genetic variants in SORL1 and Alzheimer's disease. *Arch Neurol* 2011; 68: 99-106.
248. Omalu BI, Bailes J, Hamilton RL, **Kamboh MI**, Hammers J, Case M, Fitzsimmons R: Emerging histomorphology subtypes of chronic traumatic encephalopathy (CTE) in American athletes. *Neurosurgery* 2011; 69:173-183.
249. Demirci FYK, Dressen AS, Kammerer CM, Barmada MM, Kao AH, Ramsey-Goldman R, Manzi S, **Kamboh MI**: Functional polymorphisms of the coagulation factor II gene (F2) and susceptibility to systemic lupus erythematosus (SLE). *J Rheumatology* 2011; 38: 652-657.
250. Dodge HH, Chang C-CH, **Kamboh MI**, Ganguli M: Risk of Alzheimer disease incidence attributable to vascular disease in the population. *Alzheimers Dement* 2011; 7:356-360.
251. Dasgupta S, Demirci FY, Dressen AS, Kao AH, Rhew EY, Ramsey-Goldman R, Manzi S, Kammerer CH, **Kamboh MI**: Association analysis of PON2 genetic variants with serum paraoxonase activity and systemic lupus erythematosus. *BMC Med Genet* 2011; 12:7.
252. Chen Q, Reis SE, Kammerer C, Craig W, McNamara DM, Holubkov R, Sharaf BL, Sopko G, Pauly DF, C. Merz NB, **Kamboh MI**, for the WISE study group: Association of anti-oxidized LDL and candidate genes with severity of coronary stenosis in the WISE study. *J Lipid Res* 2011; 52: 801-807.
253. Taylor KE, Chung SA, Graham RR, Ortmann WA, Lee AT, Langefeld CD, Jacob CO, **Kamboh MI**, Alarcón-Riquelme ME, Tsao BP, Moser KL, Gaffney PM, Harley JB, Petri M, Manzi S, Gregersen PK, Behrens TW, Criswell LA. Risk alleles for systemic lupus erythematosus in a large case-control collection and associations with clinical subphenotypes. *PLoS Genet* 2011; 7: e1001311.
254. Chung SA, Taylor KE, Graham RR, Nititham J, Lee AT, Ortmann WA, Jacob CO, Alarcon-Riquelme ME, Tsao BP, Harley JB, Gaffney PM, Moser KL, SLEGEN, Petri M, Demirci FY, **Kamboh MI**, Manzi S, Gregersen PK, Langefeld CD, Behrens TW, Criswell LA: Differential genetic associations for systemic lupus erythematosus based on anti-dsDNA autoantibody production. *PLoS Genet* 2011; 7: 21001323.
255. Naj AC, Jun G, Beecham GW, Wang L-S, Vardarajan BN, Buros J, Gallins PJ, Buxbaum JD, Jarvik GP, Crane PK, Larson EB, Bird TD, Boeve BF, Graff-Radford NR, De Jager PL, Evans D, Schneider JA, Carrasquillo MM, Ertekin-Taner N, Younkin SG, Cruchaga C, Kauwe JSK, Nowotny P, Kramer P, Hardy J, Huentelman MJ, Myers AJ, Barmada MM, Demirci FY, Baldwin CT, Green RC, Rogaeva E, George-Hyslop PS, Alzheimer Disease Genetics Consortium (100 authors), Cantwell LB, Dombroski BA, Beekly D, Lunetta KL, Martin ER, **Kamboh MI**, Saykin AJ, Eric M Reiman, David A Bennett, John C Morris, Thomas J Montine, Goate AM, Blacker D, Tsuang DW, Hakonarson H, Kukull WA, Foroud TM, Haines JL, Mayeux R, Pericak-Vance MA, Farrer LA, Schellenberg GD: Common variants in *MS4A4/MS4A6E*, *CD2AP*, *CD33* and *EPHA1* are associated with late-onset Alzheimer's disease. *Nature Genet* 2011; 43: 429-435.
256. Burns LC, Minster RL, Demirci FY, Barmada MM, Ganguli M, Lopez OL, DeKosky ST, **Kamboh MI**: Replication study of genome-wide associated SNPs with late-onset Alzheimer's disease. *Am J Med Genet Part B* 2011; 156:507-512.

257. Genin E, Hannequin D, Wallon D, Sleegers K, Hiltunen M, Combarros O, Bullido MJ, Engelborghs S, De Deyn P, Berr C, Pasquier F, Dubois B, Tognoni G, Fiévet N, Brouwers N, Bettens K, Arosio B, Coto E, Del Zompo M, Mateo I, Epelbaum J, Frank-Garcia A, Helisalmi S, Porcellini E, Pilotto A, Forti P, Ferri R, Scarpini E, Siciliano G, Solfrizzi V, Sorbi S, Spalletta G, Valdivieso F, Vepsäläinen S, Alvarez V, Bosco P, Mancuso M, Panza F, Nacmias B, Bossù P, Hanon O, Piccardi P, Annoni G, Seripa D, Galimberti D, Licastro F, Soininen H, Dartigues J-F, **Kamboh MI**, Van Broeckhoven C, Lambert JC, Amouyel P, Campion D: *APOE* and Alzheimer disease: A major gene with semi-dominant inheritance. *Mol Psychiatry* 2011; 16:903-907.
258. Sanghera DK, Been LF, Ralhan S, Wander GS, Mehra NK, Singh JR, Ferrell RE, **Kamboh MI**, Aston CE: Genome-wide linkage scan to identify loci associated with type 2 diabetes and blood lipid phenotypes in the Sikh Diabetes Study. *PLoS ONE* 2011; 6(6) 221188.
259. Chu SH, Roeder K, Ferrell RE, Devlin B, DeMichele-Sweet MAA, **Kamboh, MI**, Lopez OL: TOMM40 poly-T repeat lengths, age of onset and psychosis risk in Alzheimer disease. *Neurobiol Aging* 2011; 32:2328.e1-9. PMCID: PMC3192304
260. Omalu B, Hammers J, Bailes J, Hamilton RL, **Kamboh MI**, Webster G, Fitzsimmons R. Chronic traumatic encephalopathy [CTE] in an Iraqi war veteran who was diagnosed with post traumatic stress disorder [PTSD] and committed suicide. *Neurosurg Focus* 2011; 31(5): E3.
261. **Kamboh MI**, Barmada MM, Demirci FY, Minster RL, Carrasquillo MM, Pankratz VS, Younkin SG, Saykin AJ, The Alzheimer's Disease Neuroimaging Initiative, Sweet RA, Feingold E, DeKosky ST, Lopez OL. Genome-wide association analysis of age-at-onset in Alzheimer's disease. *Mol Psychiatry* 2012; 17: 1340-1346.
262. Hollingsworth P, Sweet R, Sims R, Harold D, Russ G, Abraham R, Stretton A, Jone N, Gerrish A, Chapman J, Ivanov D, Noskina V, Lovestone S, Priotsi P, Lupton M, Brayne C, Gill M, Lawlor B, Lynch A, David C, McGuiness B, Johnston J, Holmes C, Livingston G, Bass NJ, Gurling H, McQuillin A, GERD Consortium, The National Institute on Aging Late-Onset Alzheimer's Disease Family Study Group, Holmans P, Jones L, Devlin B, Klei L, Barmada MM, Demirci FY, DeKosky ST, Lopez OL, Passmore P, Owen MJ, O'Donovan MC, Mayeux R, **Kamboh MI**, Williams J: Genome-wide association study of Alzheimer's disease with psychotic symptoms. *Mol Psychiatry* 2012; 17:1316-1327.
263. Achkar J-P, Klei L, de Bakker PIW, Bellone G, Robert N, Scott R, Lu Y, Regueiro M, Brzezinsky A, **Kamboh MI**, Fiocchi C, Devlin B, Trucco M, Ringquist S, Roeder K, Duerr RH: Amino acid position 11 of HLA-DRB1 is a major determinant of chromosome 6p association with ulcerative colitis. *Genes Immun* 2012; 13:245-252.
264. Razzaghi H, Santorico SA, **Kamboh MI**: Population-based resequencing of LIPG and ZNF202 genes in subjects with extreme HDL levels. *Front Genet* 2012; 3, 89 doi: 10.3389/fgene.2012.00089.
265. **Kamboh MI**, Demirci FY, Wang X, Minster RL, Carrasquillo MM, Pankratz VS, Younkin SG, Saklin AJ, for the Alzheimer's Neuroimaging Initiative, Jun G, Baldwin C, Logue MW, Buros J, Farrer L, Pericak-Vance MA, Haines JL, Sweet RA, Ganguli M, Feingold E, DeKosky ST, Lopez OL, Barmada MM: Genome-wide association study of Alzheimer's disease.; *Transl Psychiatry* 2012; 2, e117, doi:10.1038/tp.2012.45. PMCID number PMC3365264
266. Rosenthal S, Wang X, Demirci FY, Barmada MM, Ganguli M, Lopez OL, **Kamboh MI**: Beta-amyloid toxicity modifier genes and the risk of Alzheimer's disease. *Am J Neurodegener Dis* 2012; 191-198.

267. Whitcomb DC, Larusch J, Krasinskas AM, Klei L, Smith JP, Brand RE, Neoptolemos JP, Lerch MM, Tector M, Sandhu BS, Guda NM, Orlichenko L; Alzheimer's Disease Genetics Consortium, Albert MS, Albin RL, Apostolova LG, Arnold SE, Baldwin CT, Barber R, Barnes LL, Beach TG, Beecham GW, Beekly D, Bennett DA, Bigio EH, Bird TD, Blacker D, Boxer A, Burke JR, Buxbaum JD, Cairns NJ, Cantwell LB, Cao C, Carney RM, Carroll SL, Chui HC, Clark DG, Cribbs DH, Crocco EA, Cruchaga C, Decarli C, Demirci FY, Dick M, Dickson DW, Duara R, Ertekin-Taner N, Faber KM, Fallon KB, Farlow MR, Ferris S, Foroud TM, Frosch MP, Galasko DR, Ganguli M, Gearing M, Geschwind DH, Ghetti B, Gilbert JR, Gilman S, Glass JD, Goate AM, Graff-Radford NR, Green RC, Growdon JH, Hakonarson H, Hamilton-Nelson KL, Hamilton RL, Harrell LE, Head E, Honig LS, Hulette CM, Hyman BT, Jicha GA, Jin LW, Jun G, **Kamboh MI**, Karydas A, Kaye JA, Kim R, Koo EH, Kowall NW, Kramer JH, Kramer P, Kukull WA, Laferla FM, Lah JJ, Leverenz JB, Levey AI, Li G, Lin CF, Lieberman AP, Lopez OL, Lunetta KL, Lyketsos CG, Mack WJ, Marson DC, Martin ER, Martiniuk F, Mash DC, Masliah E, McKee AC, Mesulam M, Miller BL, Miller CA, Miller JW, Montine TJ, Morris JC, Murrell JR, Naj AC, Olichney JM, Parisi JE, Peskind E, Petersen RC, Pierce A, Poon WW, Potter H, Quinn JF, Raj A, Raskind M, Reiman EM, Reisberg B, Reitz C, Ringman JM, Roberson ED, Rosen HJ, Rosenberg RN, Sano M, Saykin AJ, Schneider JA, Schneider LS, Seeley WW, Smith AG, Sonnen JA, Spina S, Stern RA, Tanzi RE, Trojanowski JQ, Troncoso JC, Tsuang DW, Valladares O, Van Deerlin VM, Van Eldik LJ, Vardarajan BN, Vinters HV, Vonsattel JP, Wang LS, Weintraub S, Welsh-Bohmer KA, Williamson J, Woltjer RL, Wright CB, Younkin SG, Yu CE, Yu L, Alkaade S, Amann ST, Anderson MA, Baillie J, Banks PA, Conwell D, Coté GA, Cotton PB, Disario J, Farrer LA, Forsmark CE, Johnstone M, Gardner TB, Gelrud A, Greenhalf W, Haines JL, Hartman DJ, Hawes RA, Lawrence C, Lewis M, Mayerle J, Mayeux R, Melhem NM, Money ME, Muniraj T, Papachristou GI, Pericak-Vance MA, Romagnuolo J, Schellenberg GD, Sherman S, Simon P, Singh VP, Slivka A, Stoltz D, Sutton R, Weiss FU, Wilcox CM, Zarnescu NO, Wisniewski SR, O'Connell MR, Kienholz ML, Roeder K, Barmada MM, Yadav D, Devlin B: Common genetic variants in the CLDN2 and PRSS1-PRSS2 loci alter risk for alcohol-related and sporadic pancreatitis. *Nature Genet* 2012; 44:1349-1354.
268. Glas J, Seiderer J, Czamara D, Paschütz G, Diegelmann J, Wetzké M, Olszak T, Wolf C, Müller-Myhsok B, Balschun T, Achkar JP, **Kamboh MI**, Franke A, Duerr RH, Brand S: PTGER4 expression-modulating polymorphisms in the 5p13.1 region predispose to Crohn's disease and affect NF-κB and XBP1 binding sites. *PLoS One* 2012; 7(12):252873.
269. Allen M, Zou F, Chai HS, Younkin CS, Crook J, Pankratz VS, Carrasquillo MM, Rowley CN, Nair AA, Middha S, Maharjan S, Nguyen T, Ma L, Malphrus KG, Palusak R, Lincoln S, Bisceglie G, Georgescu C, Schultz D, Rakhshan F, Kolbert CP, Jen J, Haines JL, Mayeux R, Pericak-Vance MA, Farrer LA, Schellenberg GD, Petersen RC, Graff-Radford NR, Dickson DW, Younkin SG, Ertekin-Taner N; Alzheimer's Disease Genetics Consortium (ADGC), Apostolova LG, Arnold SE, Baldwin CT, Barber R, Barmada MM, Beach T, Beecham GW, Beekly D, Bennett DA, Bigio EH, Bird TD, Blacker D, Boeve BF, Bowen JD, Boxer A, Burke JR, Buros J, Buxbaum JD, Cairns NJ, Cantwell LB, Cao C, Carlson CS, Carney RM, Carroll SL, Chui HC, Clark DG, Corneveaux J, Cotman CW, Crane PK, Cruchaga C, Cummings JL, De Jager PL, DeCarli C, DeKosky ST, Demirci FY, Diaz-Arrastia R, Dick M, Dombroski BA, Duara R, Ellis WD, Evans D, Faber KM, Fallon KB, Farlow MR, Ferris S, Foroud TM, Frosch M, Galasko DR, Gallins PJ, Ganguli M, Gearing M, Geschwind DH, Ghetti B, Gilbert JR, Gilman S, Giordani B, Glass JD, Goate AM, Green RC, Growdon JH, Hakonarson H, Hamilton RL, Hardy J, Harrell LE, Head E, Honig LS, Huentelman MJ, Hulette CM, Hyman BT, Jarvik GP, Jicha GA, Jin LW, Jun G, **Kamboh MI**, Karlawish J, Karydas A, Kauwe JS, Kaye JA, Kennedy N, Kim R, Koo EH, Kowall NW, Kramer P, Kukull WA, Lah JJ, Larson EB, Levey AI, Lieberman AP, Lopez OL, Lunetta KL, Mack WJ, Marson DC, Martin ER, Martiniuk F, Mash DC, Masliah E, McCormick WC, McCurry SM, McDavid AN, McKee AC, Mesulam M, Miller BL, Miller CA, Miller JW, Montine TJ, Morris JC, Myers AJ, Naj AC, Nowotny P, Parisi JE, Perl DP, Peskind E, Poon WW, Potter H, Quinn JF, Raj A,

- Rajbhandary RA, Raskind M, Reiman EM, Reisberg B, Reitz C, Ringman JM, Roberson ED, Rogaeva E, Rosenberg RN, Sano M, Saykin AJ, Schneider JA, Schneider LS, Seeley W, Shelanski ML, Slifer MA, Smith CD, Sonnen JA, Spina S, St George-Hyslop P, Stern RA, Tanzi RE, Trojanowski JQ, Troncoso JC, Tsuang DW, Van Deerlin VM, Vardarajan BN, Vinters HV, Vonsattel JP, Wang LS, Weintraub S, Welsh-Bohmer KA, Williamson J, Woltjer RL. Novel late-onset Alzheimer disease loci variants associate with brain gene expression. *Neurology* 2012; 79:221-228.
270. Mathis CA, Kuller LH, Klunk WE, Snitz BE, Price JC, Weisfeld LA, Rosario BL, Lopresti BJ, Saxton JA, Aizenstein HJ, McDade EM, **Kamboh MI**, DeKosky ST, Lopez OL: In vivo assessment of amyloid deposition in non-demented very elderly subjects. *Ann Neurol* 2013; 73:751-761.
271. Elbers CC, Guo Y, Tragante V, van Iperen EP, Lanktree MB, Castillo BA, Chen F, Yanek LR, Wojczynski MK, Li YR, Ferwerda B, Ballantyne CM, Buxbaum SG, Chen YD, Chen WM, Cupples LA, Cushman M, Duan Y, Duggan D, Evans MK, Fernandes JK, Fornage M, Garcia M, Garvey WT, Glazer N, Gomez F, Harris TB, Halder I, Howard VJ, Keller MF, **Kamboh MI**, Kooperberg C, Kritchevsky SB, Lacroix A, Liu K, Liu Y, Musunuru K, Newman AB, Onland-Moret NC, Ordovas J, Peter I, Post W, Redline S, Reis SE, Saxena R, Schreiner PJ, Volcik KA, Wang X, Yusuf S, Zonderland AB, Anand SS, Becker DM, Psaty B, Rader DJ, Reiner AP, Rich SS, Rotter JI, Sale MM, Tsai MY, Borecki IB, Hegele RA, Kathiresan S, Nalls MA, Taylor HA Jr, Hakonarson H, Sivapalaratnam S, Asselbergs FW, Drenos F, Wilson JG, Keating BJ: Gene-centric meta-analysis of lipid traits in african, East asian and Hispanic populations. *PLoS One* 2012;7(12):e50198. doi: 10.1371/journal.pone.0050198. Epub 2012 Dec 7.
272. Saxena R, Saleheen D, Been L, Garavito M, Braun T, Bjonne A, Young R, Ho W, Rasheed A, Frossard P, Sim X, Hassanali N, Radha V, Chidambaram M, Liju S, Rees S, Peng-Keat ND, Wong T-Y, Yamauchi T, Hara K, Tanaka Y, Hirose H, McCarthy M, Morris A, Basit A, Barnett A, Katulanda P, Matthews D, Mohan V, Wander G, Singh J, Mehra N, Ralhan S, **Kamboh MI**, Mulvihill J, Maegawa H, Tobe K, Maeda S, Shin C, Tai E, Kelly M, Chambers J, Kooner J, Kadawaki T, Deloukas P, Rader D, Danesh J, Sanghera D: Genome-wide association study identifies a novel locus contributing to type 2 diabetes susceptibility in Sikhs of Punjabi origin from India. *Diabetes* 2013; 62:1746-1755.
273. Jalil SF, Bhatti A, Demirci FY, Wang X, Ahmed I, Ahmed M, Barmada MM, Malik JM, John P, **Kamboh MI**: Replication of European rheumatoid arthritis loci in the Pakistani population. *J Rheumatol* 2013; 40: 401-407
274. Christiel D, Shofer J, Millard SP, Li E, DeMichele-Sweet MA, Weamer EA, **Kamboh MI**, Lopez OL, Sweet RL, Tsuang D: Genetic association between APOE*4 and neuropsychiatric symptoms in patients with probable Alzheimer's disease is dependent on the psychosis phenotype. *Behavior Brain Funct* 2012; 8: 62
275. **Kamboh MI**, Wang X, Kao AH, Barmada M, Clarke A, Ramsey-Goldman R, Manzi S, Demirci FY: Genome-wide association study of antiphospholipid antibodies. *Autoimmune Dis* 2013; 2013:761046. doi: 10.1155/2013/761046. Epub 2013 Feb 24
276. Bryant EK, Dressen AS, Bunker CH, Hokanson JE, Hamman RF, **Kamboh MI**, Demirci FY: A multiethnic replication study of plasma lipoprotein levels-associated SNPs identified in recent GWAS. *PLOS ONE* 2013; 8 (5) e63469
277. Reitz C, Jun G, Naj A, Rajbhandary R, Vardarajan BN, Wang L-S, Valladares O, Lin C-F, Larson EB, Graff-Radford NR, Evans D, De Jager PL, Crane PK, Buxbaum JD, Murrell JR, Raj T, Ertekin-Taner N, Logue M, Baldwin CT, Green RC, Barnes LL, Cantwell LB, Fallin MD, Go RCP, Griffith P, Obisesan TO, Manly JJ, Lunetta KL, **Kamboh MI**, Lopez OL, Bennett DA, Hendrie H, Hall KS, Goate AM, Byrd GS, Kukull WA, Foroud TM, Haines JL, Farrer LA, Pericak-Vance MA, Schellenberg GD, Mayeux R: Variants in the ATP-binding cassette transporter (ABCA7), Apolipoprotein E ε4, and the risk of late-onset

Alzheimer's disease in African Americans. *JAMA* 2013; 309:1483-1492.

278. Holton P, Ryten M, Nalls M, Trabzuni D, Weale ME, Hernandez D, Crehan H, Gibbs JR, Mayeux R, Haines JL, Farrer LA, Pericak-Vance MA, Schellenberg GD; Alzheimer's Disease Genetics Consortium, Ramirez-Restrepo M, Engel A, Myers AJ, Corneveaux JJ, Huentelman MJ, Dillman A, Cookson MR, Reiman EM, Singleton A, Hardy J, Guerreiro R, Apostolova LG, Arnold SE, Baldwin CT, Barber R, Barmada MM, Beach TG, Beecham GW, Bickley D, Bennett DA, Bigio EH, Bird TD, Blacker D, Boeve BF, Bowen JD, Boxer A, Burke JR, Buros J, Buxbaum JD, Cairns NJ, Cantwell LB, Cao C, Carlson CS, Carney RM, Carrasquillo MM, Carroll SL, Chui HC, Clark DG, Cotman CW, Crane PK, Crocco EA, Cruchaga C, Cummings JL, De Jager PL, DeCarli C, DeKosky ST, Demirci FY, Diaz-Arrastia R, Dick M, Dickson DW, Duara R, Ellis WG, Ertekin-Taner N, Evans D, Faber KM, Fallon KB, Farlow MR, Ferris S, Foroud TM, Frosch MP, Galasko DR, Ganguli M, Gearing M, Geschwind DH, Ghetti B, Gilbert JR, Gilman S, Giordani B, Glass JD, Goate AM, Graff-Radford NR, Green RC, Growdon JH, Hakonarson H, Hamilton RL, Harrell LE, Head E, Honig LS, Hulette CM, Hyman BT, Jarvik GP, Jicha GA, Jin LW, Jun G, **Kamboh MI**, Karlawish J, Karydas A, Kauwe JS, Kaye JA, Kim R, Koo EH, Kowall NW, Kramer P, Kukull WA, Lah JJ, Larson EB, Levey AI, Lieberman AP, Lopez OL, Lunetta KL, Mack WJ, Marson DC, Martin ER, Martiniuk F, Mash DC, Masliah E, McCormick WC, McCurry SM, McDavid AN, McKee AC, Mesulam M, Miller BL, Miller CA, Miller JW, Montine TJ, Morris JC, Naj AC, Nowotny P, Parisi JE, Peskind E, Petersen RC, Poon WW, Potter H, Quinn JF, Raj A, Rajbhandary RA, Raskind M, Reisberg B, Reitz C, Ringman JM, Roberson ED, Rogaeva E, Rosenberg RN, Sano M, Saykin AJ, Schneider JA, Schneider LS, Seeley WW, Shelanski ML, Smith CD, Sonnen JA, Spina S, St George-Hyslop P, Stern RA, Tanzi RE, Trojanowski JQ, Troncoso JC, Tsuang DW, Valladares O, Van Deerlin VM, Vardarajan BN, Vinters HV, Vonsattel JP, Wang LS, Weintraub S, Welsh-Bohmer KA, Williamson J, Woltjer RL, Wright CB, Younkin SG: Initial assessment of the pathogenic mechanisms of the recently identified Alzheimer risk Loci. *Ann Hum Genet* 2013;77: 85-105.
279. Nebes RD, Snitz BE, Cohen AD, Aizenstein HJ, Saxton JA, Halligan EM, Mathis CA, Price JC, **Kamboh MI**, Weissfeld LA, Klunk WE: Cognitive aging in persons with minimal amyloid- β and white matter hyperintensities. *Neuropsychologia* 2013; 51: 2202-2209.
280. Lopez OL, Becker JT, Chang Y-F, Sweet RA, Aizenstein H, Snitz B, Saxton J, McDade E, **Kamboh MI**, DeKosky ST, Reynolds CF, Klunk WE: The long-term effects of typical and atypical antipsychotics in patients with Probable Alzheimer's disease. *Am J Psychiatry* 2013; 170:1051-1058.
281. Heim AF, Coyne MJ, **Kamboh MI**, Ryan C, Jennings JR: The catechol-o methyltransferaseVal158Met polymorphism modulates organization of regional cerebral blood flow response to working memory in adults. *Int J Psychophysiol*. 2013; 90:149-156.
282. Lambert JC, Ibrahim-Verbaas CA, Harold D, Naj AC, Sims R, Bellenguez C, Gyungah J, DeStefano A, Bis JC, Beecham GW, Grenier-Boley B, Russo G, Thornton-Wells TA, Jones N, Smith AV, Chouraki V, Thomas C, Ikram MA, Zelenika D, Vardarajan BN, Kamatani Y, Lin CF, Gerrish A, Schmidt H, Kunkle B, Dunstan ML, Ruiz A, Bihoreau MT, Cho SH, Reitz C, Pasquier F, Hollingworth P, Ramirez A, Hanon O, Fitzpatrick AL, Buxbaum JD, Campion D, Crane PK, Becker T, Gudnason V, Cruchaga C, Craig D, Amin N, Berr C, Lopez OL, De Jager PL, Deramecourt V, Johnston JA, Evans D, Lovestone S, Letteneur L, Moron FJ, Rubinsztein DC, Eiriksdottir G, Sleegers K, Goate AM, Fievet N, Huentelman MJ, Gill M, Brown K, **Kamboh MI**, Keller L, Barberger-Gateau P, McGuinness B, Larson EB, Myers AJ, Dufouil C, Todd S, Wallon D, Love S, Rogaeva E, Gallacher J, St George-Hyslop P, Clarimon J, Lle A, Bayer A, Tsuang DW, Yu L, Tsolaki M, Bossu P, Spalletta G, Proitsi P, Collinge J, Sorbi S, Garcia FS, Fox N,

Hardy J, Naranjo MCD, Bosco P, Clarke R, Brayne C, Galimberti D, Mancuso M, Moebus S, Mecocci P, del Zompo M, Maier W, Hampel H, Pilotto A, Bullido M, Panza F, Caffarra P, Naemias B, Gilbert JR, Mayhaus M, Lannfelt L, Hakonarson H, Pichler S, Carrasquillo MM, Ingelsson M, Beekly D, Alvarez V, Zou F, Valladares O, Younkin SG, Coto E, Hamilton-Nelson KL, Mateo I, Owen MJ, Faber KM, Jonsson PV, Combarros O, O'Donovan MC, Cantwell LB, Soininen H, Blacker D, Mead S, Mosley TH, Bennett DA, Harris TB, Fratiglioni L, Holmes C, de Brujin R, Passmore P, Montine TJ, Bettens K, Rotter JI, Brice A, Morgan K, Foroud TM, Kukull WA, Hannequin D, Powell JF, Nalls MA, Ritchie K, Lunetta KL, Kauwe JSK, Boerwinkle E, Riemenschneider M, Boada M, Hiltunen M, Martin ER, Schmidt R, Rujescu D, Wang LS, Dartigues JF, Mayeux R, Tzourio C, Hofman A, Nothen MM, Graff C, Psaty BM, Jones L, Haines JL, Holman PA, Lathrop M, Pericak-Vance MA, Launer LJ, Farrer LA, van Duijn CM, Van Broekhoven C, Moskvina V, Seshadri S, Williams J, Schellenberg GD, Amouyel P: Extended meta-analysis of 74,538 individuals identifies 11 new susceptibility loci for Alzheimer's disease. *Nature Genet* 2013; 45:152-1458.

283. Hughes TM, Lopez OL, Evans RW, **Kamboh MI**, Williamson JD, Klunk WE, Mathis C, Price JC, Cohen A, Snitz B, DeKosky S, Kuller LK: Markers of cholesterol transport are associated with amyloid deposition in the brain. *Neurobiol Aging* 2014; 35: 820-807.
284. Cruchaga C, Karch CM, Jin SC, Benitez BA, Guerreiro R, Harari O, Norton J, Budde J, Bertelsen S, Jeng AT, Cooper B, Cai Y, Skorupa T, Carrell D, Levitch D, Hsu S, Choi J, Ryten M, SassiC, Bras J, Gibbs RJ, Hernandez DG, Lupton MK, PowellJ, Forabosco P, Ridge PG, Corcoran CD, Tschanz JT, Norton MC, Munger RG, Schmutz C, Leary M, Demirci FY, Bamne MN, Wang X, Lopez OL, Ganguli M, Medway C, Turton J, Lord J, Braae A, Barber I, Brown K, The Alzheimer's Research UK (ARUK) Consortium, Pastor P, Lorenzo-Betancor O, Brkanac Z, Scot E, Topol E, Morgan K, Rogaeva E, Singleton A, Hardy J, **Kamboh MI**, St George-Hyslop P, Cairns N, Morris JC, Kauwe JSK, Goate AM: Rare coding variants in phospholipase D3 confer risk for Alzheimer's disease. *Nature* 2014; 505:550-554.
285. Wang M-S, Fiocchi C, Zhu X, Ripke S, **Kamboh MI**, Robert N, Duerr RH, Achkar J-P. Gene-gene and gene-environment interactions in ulcerative colitis. *Hum Genet* 2014; 133: 547-558.
286. Pirim D, Wang X, Radwan ZH, Niemsiri V, Hokanson JE, Hamman RF, Barmada MM, Demirci FY, **Kamboh MI**: Lipoprotein lipase gene sequencing and plasma lipid profile. *J Lipid Res* 2014; 55:85-93.
287. Bamne MN, Demirci FY, Berman S, Snitz BE, Rosenthal SL, Wang X, Lopez OL. **Kamboh MI**: Investigation of an APP protective mutation (A673T) in a North American case-control sample of late-onset Alzheimer's disease. *Neurobiol Aging* 2014; 1779.e15-1779.e16.
288. Rosenthal SL, **Kamboh MI**: Late-onset Alzheimer's disease genes and the potentially implicated pathways. *Curr Genet Med Rep* 2014; 2:85-101.
289. Rosenthal SL, Barmada MM, Wang X, Demirci FY, **Kamboh MI**: Connecting the dots: Potential of data integration to identify regulatory SNPs in late-onset Alzheimer's disease GWAS findings. *PLoS One* 2014; 9(4):e95152.
290. Stokes ME, Barmada MM, **Kamboh MI**, Visweswaran S: The application of network label propagation to rank biomarkers in genome-wide Alzheimer's data. *BMC Genomics* 2014; 15: 282.
291. Chung SA, Brown EE, Williams AH, Ramos PS, Berthier CC, Bhangale T, Alarcon-Riquelme M, Behrens TW, Criswell LA, Graham DC, Demirci FY, Edberg JC, Gaffney PM, Harley JB, Jacob CO, **Kamboh MI**, Kelly JA, Manzi S, Moser-Sivils KL, Russell LP, Petri M, Tsao BP, Vyse TJ, Zidovetzki R, Kretzler M, Kimberly RP, Freedman BI, Graham RR, Langefeld CD. Lupus nephritis susceptibility loci in women

with systemic lupus erythematosus. *J Am Soc Nephrol* 2014; 25:2859-2870.

292. Escott-Price V, Bellenguez C, Wang LS, Choi SH, Harold D, Jones L, Holmans P, Gerrish A, Vedernikov A, Richards A, DeStefano AL, Lambert JC, Ibrahim-Verbaas CA, Naj AC, Sims R, Jun G, Bis JC, Beecham GW, Grenier-Boley B, Russo G, Thornton-Wells TA, Jones N, Smith AV, Chouraki V, Thomas C, Ikram MA, Zelenika D, Vardarajan BN, Kamatani Y, Lin CF, Schmidt H, Kunkle B, Dunstan ML, Vronskaya M, The UK Brain Expression Consortium, Johnson AD, Ruiz A, Bihoreau MT, Reitz C, Pasquier F, Hollingworth P, Hanon O, Fitzpatrick A, Buxbaum JD, Campion D, Crane PK, Baldwin CT, Becker T, Gudnason V, Cruchaga C, Craig D, Amin N, Berr C, Lopez O, De Jager PL, Deramecourt V, Johnston JA, Evans D, Lovestone SL, Letenneur L, Hernandez I, Rubinsztein DC, Eiriksdottir G, Sleegers K, Goate AM, Fievet N, Huentelman MJ, Gill M, Brown K, **Kamboh MI**, Keller L, Barberger-Gateau P, McGuinness B, Larson EB, Myers AJ, Dufouil C, Todd S, Wallon D, Love S, Rogeava E, Gallacher J, St. George-Hyslop P, Clarimon J, Lleo A, Bayer A, Tsuang DW, Yu L, Tsolaki M, Bossu P, Spalletta G, Priotsi P, Collinge J, Sorbi S, Sanchez Garcia F, Fox N, Hardy J, Naranjo MCD, Bosco P, Clarke R, Brayne C, Galimberti D, Scarpini E, Bonuccelli U, Mancuso M, Siciliano G, Moebus S, Mecocci P, Del Zompo M, Maier W, Hampel H, Pilotto A, Bullido M, Frank-Garcia A, Panza F, Solfrizzi V, Caffarra P, Nacmias B, Perry W, Mayhaus M, Lannfelt L, Hakonarson H, Pichler S, Carrasquillo MM, Ingesson M, Beekly D, Alvarez V, Zou F, Valladares O, Younkin SG, Coto E, Hamilton-Nelson KL, Gu W, Razquin C, Pastor P, Mateo I, Owen MJ, Faber KM, Jonsson PV, Combarros O, O'Donovan MC, Soininen H, Blacker D, Mead S, Mosley Jr TH, Bennett DA, Harris TB, Fratiglioni L, Holmes C, FAG de Brujin R, Passmore P, Montine TJ, Bettens K, Rotter JI, Brice A, Morgan K, Foroud TM, Kukull WA, Hannequin D, Powell J, Nalls MA, Ritchie K, Lunetta KL, Kauwe John SK, Boerwinkle E, Riemschneider M, Boada M, Hiltunen M, Martin ER, Schmidt R, Rujescu D, Dartigues JF, Mayeux R, Tzourio C, Hofman A, Nothen MM, Graff C, Psaty BM, Haines JL, Lathrop M, Pericak-Vance MA, Launer LJ, Van Broeckhoven C, Farrer LA, van Duijn CM, Ramirez A, Seshadri S, Schellenberg GD, Amouyel P, Williams J. Gene-wide analysis detects two new susceptibility genes for Alzheimer's disease. *PLoS One* 2014; 9(6): 294661.
293. Chaudhry M, Hasnain S, Snitz B, Wang X, Rosenthal S, Demirci FY, **Kamboh MI**: Association of *APOE* polymorphisms and stressful life events with dementia in a Pakistani population. *Neurosci Letts* 2014; 570: 42046.
294. Naj AC, Jun G, Reitz C, Kunkle BW, Perry W, Park YS, Beecham GW, Rajbhandary RA, Hamilton-Nelson KL, Wang L-S, Kauwe JSK, Huentelman MJ, Myers AJ, Bird TD, Boeve BF, Baldwin CT, Jarvik GP, Crane PK, Rogeava E, Barmada MM, Demirci FY, Cruchaga C, Kramer P, Alzheimer's Disease Genetics Consortium*, Ertekin-Taner N, Hardy J, Graff-Radford NR, Green RC, Larson EB, George-Hyslop PS, Buxbaum JD, Evans D, Schneider JA, Lunetta KL, **Kamboh MI**, Saykin AJ, Reiman EM, De Jager PL, Bennett DA, Morris JC, Montine TJ, Goate AM, Blacker D, Tsuang DW, Hakonarson H, Kukull WA, Foroud TM, Martin ER, Haines JL, Mayeux R, Farrer LA, Schellenberg GD, Pericak-Vance MA: Effects of multiple genetic loci on age-at-onset in late onset Alzheimer disease: A genome-wide association study. *JAMA Neurol* 2014; 71:1394-404.
295. Logue MW, Schu M, Vardarajan BN, Farrell J, Bennett DA, Buxbaum JD, Byrd GS, Ertekin-Taner N, Evans D, Foroud T, Goate A, Graff-Radford NR, **Kamboh MI**, Kukull WA, Manly JJ, Alzheimer Disease Genetics Consortium, Haines JL, Mayeux R, Pericak-Vance MA, Schellenberg GD, Lunetta KL, Baldwin CT, Fallin MD, Farrer LA. Two Rare *AKAP9* missense variants are associated with Alzheimer disease in African Americans. *Alzheimers Dement* 2014; 10 609-618.

296. Wang X, Lopez OL, Sweet RA, Becker JT, DeKosky ST, Barmada MM, Demirci FY, **Kamboh MI**. Genetic determinants of disease progression in Alzheimer's disease. *J Alzheimers Dis* 2015; 43:649-655.
297. Beecham G, Hamilton K, Naj A, Martin E, Huentelman M, Myers A, Corneveaux J, Hardy J, Vonsattel J-P, Younkin S, Bennett D, De Jager P, Larson E, Crane P, **Kamboh MI**, Kofler JK, Mash D, Duque L, Gilbert J, Gwirtsman H, Buxbaum J, Kramer P, Dickson D, Farrer L, Frosch M, Ghetti B, Haines J, Hyman B, Kukull W, Mayeux R, Pericak-Vance M, Schneider J, Trojanowski J, Reiman E, Schellenberg G, Montine TJ. Genome-wide association meta-analysis of neuropathologic features of Alzheimer's disease and related dementias. *PLoS Genet* 2014; 10(9): e21004606.
298. Niemsiri V, Wang X, Pirim D, Radwan ZH, Hokanson JE, Hamman RF, Barmada MM, Demirci FY, **Kamboh MI**. Impact of genetic variants in human scavenger receptor class B type I (*SCARB1*) on plasma lipid traits. *Cir Cardiovasc Genet* 2014; 7:838-847. [Published with an editorial: Human scavenger receptor class B type I variants, lipid traits, and cardiovascular disease. *Cir Cardiovasc Genet* 2014; 7:735-737].
299. Zheng X, Demirci FY, Barmada MM, Richardson GA, Lopez OL, Sweet RA, **Kamboh MI**, Feingold E. A rare duplication on chromosome 16p11.2 is identified in patients with psychosis in Alzheimer's disease. *PLoS One* 2014; 9(11):e111462.
300. Floudas CS, Um N, **Kamboh MI**, Barmada MM, Visweswaran S. Identifying genetic interactions associated with late-onset Alzheimer's disease. *BioData Min* 2014; 19: 7(1): 35.
301. Radwan ZH, Wang X, Waqar F, Pirim D, Niemsiri V, Hokanson JE, Hamman RF, Bunker CH, Barmada MM, Demirci FY, **Kamboh MI**. Comprehensive evaluation of the association of APOE genetic variation with plasma lipoprotein traits in U.S. whites and African blacks. *PLoS One* 2014; 9(12):e114618.
302. Chaudhry M, Wang X, Bamne MN, Hasnain S, Demirci FY, Lopez OL, **Kamboh MI**. Genetic variation in imprinted genes is associated with risk of late-onset Alzheimer's Disease. *J Alzheimers Dis* 2015; 44:989-994.
303. Pirim D, Wang X, Radwan ZH, Niemsiri V, Bunker CH, Barmada MM, Demirci FY, **Kamboh MI**. Resequencing of LPL in African Blacks and associations with lipoprotein-lipid levels. *Eur J Hum Genet* 2015; 23:1244-1253.
304. Chaudhry M, Hasnain S, Wang X, Snitz BE, Bamne MN, Rosenthal S, Demirci FY, **Kamboh MI**. Association analysis of 23 susceptibility loci with risk of dementia in a Pakistani population. *Psychiatry Res* 2015; 225:223-224.
305. Jones L, Bellenguez C, Wang LS, Choi SH, Harold D, Vedernikov A, Escott-Price V, Stone T, Richards A, Lambert JC, Ibrahim-Verbaas CA, Naj AC, Sims R, Gerrish A, Jun G, DeStefano AL, Bis JC, Beecham GW, Grenier-Boley B, Russo G, Thornton-Wells TA, Jones N, Smith AV, Chouraki V, Thomas C, Ikram MA, Zelenika D, Vardarajan BN, Kamatani Y, Lin CF, Schmidt H, Kunkle B, Dunstan ML, Ruiz A, Bihoreau MT, Reitz C, Pasquier F, Hollingworth P, Hanon O, Fitzpatrick Al, Buxbaum JD, Campion D, Crane PK, Becker T, Gudnason V, Cruchaga C, Craig D, Amin N, Berr C, Lopez OL, de Jager PL, Deramecourt V, Johnston JA, Evans D, Lovestone S, Letteneur L, Morón FJ, Rubinsztein DC, Eiriksdottir G, Sleegers K, Goate AM, Fiévet N, Huentelman MJ, Gill M, Brown K, **Kamboh MI**, Keller L, Barberger-Gateau P, McGuinness B, Larson EB, Myers AJ, Dufouil C, Todd S, Wallon D, Love S, Rogaeva E, Gallacher J, St George-Hyslop P, Clarimon J, Lleo A, Bayer A, Tsuang DW, Yu L, Tsolaki M, Bossù P, Spalletta G, Proitsi P, Collinge J, Sorbi S, Garcia FS, Fox N, Hardy J, Naranjo MCD, Razquin C, Bosco P, Clarke R, Brayne C, Galimberti D, Mancuso M, MRC CFAS, Moebus S, Mecocci P,

del Zompo M, Maier W, Hampel H, Pilotto A, Bullido M, Panza F, Caffarra P, Nacmias B, Gilbert JR, Mayhaus M, Jessen F, Dichgans M, Lannfelt L, Hakonarson H, Pichler S, Carrasquillo MM, Ingelsson M, Beekly D, Alavarez V, Zou F, Valladares O, Younkin SG, Coto E, Hamilton-Nelson KL, Mateo I, Owen MJ, Faber KM, Jonsson PV, Combarros O, O'Donovan MC, Cantwell LB, Soininen H, Blacker D, Mead S, Mosley TH, Bennett DA, Harris TB, Fratiglioni L, Holmes C, de Brujin RFAG, Passmore P, Montine TJ, Bettens K, Rotter JI, Brice A, Morgan K, Foroud TM, Kukull WA, Hannequin D, Powell JF, Nalls MA, Ritchie K, Lunetta KL, Kauwe JSK, Boerwinkle E, Riemenschneider M, Boada M, Hiltunen M, Martin ER, Pastor P, Schmidt R, Rujescu D, Dartigues JF, Mayeux R, Tzourio C, Hofman A, Nöthen MM, Graff C, Psaty BM, Haines JL, Lathrop M, Pericak-Vance MA, Launer LJ, Farrer LA, van Duijn CM, Van Broekhoven C, Ramirez A, Schellenberg GD, Seshadri S, Philippe Amouyel P, Williams J, Holmans PA. Convergent genetic and expression data implicate immunity in Alzheimer's disease. *Alzheimers Dement* 2015; 11:658-671.

306. Hohman TJ, Cooke-Bailey JN, Reitz C, Jun G, Naj A, Beecham GW, Liu Z, Carney RM, Vance JM, Cuccaro ML, Rajbhandary R, Vardarajan BN, Wang L-S, Valladares O, Lin CF, Larson EB, Graff-Radford NR, Evans D, Jager PL, Crane PK, Buxbaum JD, Murrell JR, Raj T, Ertekin-Taner N, Logue MW, Baldwin CT, Green RC, Barnes LL, Cantwell LB, Fallin MD, Go RCP, Griffith P, Obisesan TO, Manly JJ, Lunetta KL, **Kamboh MI**, Lopez OL, Bennett DA, Hardy J, Hendrie HC, Hall KS, Gaote AM, Lang R, Byrd GS, Kukull WA, Foroud TM, Farrer LA, Martin ER, Pericak-Vance MA, Schellenberg GD, Mayeux R, Haines JL, Thornton-Wells TA, for the Alzheimer Disease Genetics Consortium. Global and local ancestry in African Americans: Implications for Alzheimer's Disease risk. *Alzheimers Dement* 2016; 12:233-243.
307. Wang X, Lopez OL, Sweet RA, Becker JT, DeKosky ST, Barmada MM, Feingold E, Demirci FY, **Kamboh MI**. Genetic determinants of survival in patients with Alzheimer's disease. *J Alzheimers Dis* 2015; 45:651-658.
308. Jun G, Ibrahim-Verbaas CA, Vronskaya M, Lambert J-C, Chung J, Naj AC, Kunkle BW, Wang L-S, Bis JC, Bellenguez C, Harold D, Lunetta KL, DeStefano AL, Grenier-Boley B, Sims R, Beecham GW, Smith AV, Chouraki V, Hamilton-Nelson KL, Ikram MA, Fievet N, Denning N, Martin ER, Schmidt H, Kamatani Y, Dunstan ML, Valladares O, Laza AR, Zelenika D, Ramirez A, Foroud TM, Choi S-H, Boland A, Becker T, Kukull WA, van der Lee SJ, Pasquier F, Cruchaga C, Beekly D, Fitzpatrick AL, Hanon O, Gill M, Barber R, Gudnason V, Campion D, Love S, Bennett DA, Amin N, Berr C, Tsolaki M, Buxbaum JD, Lopez OL, Deramecourt V, Fox NC, Cantwell LB, Tarraga L, Dufouil C, Hardy J, Crane PK, Eiriksdottir G, Hannequin D, Clarke R, Evans D, Mosley TH, Letenneur L, Brayne C, Maier W, De Jager P, Emilsson V, Dartigues J-F, Hampel H, **Kamboh MI**, De Brujin RFAG, Tzourio C, Pastor P, Larson EB, Rotter JI, O'Donovan MC, Montine TJ, Nalls MA, Mead S, Reiman EM, Johnson PV, Holmes C, St George-Hyslop PH, Boada M, Passmore P, Wendland JR, Schmidt R, Morgan K, Winslow AR, Powel JF, Carrasquillo M, Younkin SG, Jakobsdottir J, Kauwe JSK, Wilhelmsen KC, Rujescu D, Nothen MM, Hofman A, Jones L, IGAP Consortium, Haines J, Psaty BM, Van Broekhoven C, Holmans P, Launer L, Mayeux R, Lathrop M, Goate A, Escott-Price V, Seshadri S, Pericak-Vance MA, Amouyel P, Williams J, van Duijn CM, Schellenberg GD, Farrer L. A novel Alzheimer disease locus located near the gene encoding Tau protein. *Mol Psychiatry* 2016; 21:108-117. doi: 10.1038/mp.2015.23. Epub 2015 Mar 17
309. Kiani AK, John P, Bhati A, Zia A, Shahid G, Akhtar P, Wang X, Demirci FY, **Kamboh MI**. Association of 32 type 1 diabetes risk loci in Pakistani patients. *Diabetes Res Clin Prac* 2015; 108:137-142.
310. Zheng X, Demirci FY, Barmada MM, Richardson GA, Lopez OL, Sweet RA, **Kamboh MI**, Feingold E. Genome-wide copy number variation study of psychosis in Alzheimer's disease. *Transl Psychiatry* 2015; 5:e574.

311. Cheema AN, Bhatti A, Wang X, Ali J, Bamne MN, Demirci FY, **Kamboh MI**. *APOE* gene polymorphism and risk of coronary stenosis in Pakistani population. *BioMed Res International* 2015; 2015:587465.
312. Kiani AK, Jahangir S, John P, Bhatti A, Zia A, Wang X, Demirci FY, **Kamboh MI**. Genetic link of type 1 diabetes loci with rheumatoid arthritis in Pakistani patients. *Immonogenetics* 2015; 67:277-282.
313. Zia A, Bhatti A, Jalil F, Wang X, John P, Kiani AK, Zafar J, **Kamboh MI**. Prevalence of type 2 diabetes-associated complications in Pakistan. *Int J Diabetes Dev Ctries* 2015; April 15 DOI 10.1007/s13410-015-0380-6 [Epub ahead of print]
314. Rosenthal S, Bamne MN, Wang X, Berman S, Snitz BE, Klunk WE, Sweet RA, Demirci FY, Lopex OL, **Kamboh MI**. More evidence for association of a rare TREM2 mutation (R47H) with Alzheimer's disease risk. *Neurobiol Aging* 2015; 36:2443.e21-e26.
315. Wang LS, Naj AC, Graham RR, Crane PK, Kunkle BW, Cruchaga C, Murcia JD, Cannon-Albright L, Baldwin CT, Zetterberg H, Blennow K, Kukull WA, Faber KM, Schupf N, Norton MC, Tschanz JT, Munger RG, Corcoran CD, Rogaeva E; Alzheimer's Disease Genetics Consortium, Lin CF, Dombroski BA, Cantwell LB, Partch A, Valladares O, Hakonarson H, St George-Hyslop P, Green RC, Goate AM, Foroud TM, Carney RM, Larson EB, Behrens TW, Kauwe JS, Haines JL, Farrer LA, Pericak-Vance MA, Mayeux R, Schellenberg GD; National Institute on Aging-Late-Onset Alzheimer's Disease (NIA-LOAD) Family Study, Albert MS, Albin RL, Apostolova LG, Arnold SE, Barber R, Barmada M, Barnes LL, Beach TG, Becker JT, Beecham GW, Beekly D, Bennett DA, Bigio EH, Bird TD, Blacker D, Boeve BF, Bowen JD, Boxer A, Burke JR, Buxbaum JD, Cairns NJ, Cao C, Carlson CS, Carroll SL, Chui HC, Clark DG, Cribbs DH, Crocco EA, DeCarli C, DeKosky ST, Demirci FY, Dick M, Dickson DW, Duara R, Ertekin-Taner N, Fallon KB, Farlow MR, Ferris S, Frosch MP, Galasko DR, Ganguli M, Gearing M, Geschwind DH, Ghetti B, Gilbert JR, Glass JD, Graff-Radford NR, Growdon JH, Hamilton RL, Hamilton-Nelson KL, Harrell LE, Head E, Honig LS, Hulette CM, Hyman BT, Jarvik GP, Jicha GA, Jin LW, Jun G, Jun G, **Kamboh MI**, Karydas A, Kaye JA, Kim R, Koo EH, Kowall NW, Kramer JH, LaFerla FM, Lah JJ, Leverenz JB, Levey AI, Li G, Lieberman AP, Lopez OL, Lunetta KL, Lyketsos CG, Mack WJ, Marson DC, Martin ER, Martiniuk F, Mash DC, Masliah E, McCormick WC, McCurry SM, McDavid AN, McKee AC, Mesulam WM, Miller BL, Miller CA, Miller JW, Montine TJ, Morris JC, Murrell JR, Olichney JM, Parisi JE, Perry W, Peskind E, Petersen RC, Pierce A, Poon WW, Potter H, Quinn JF, Raj A, Raskind M, Reiman EM, Reisberg B, Reitz C, Ringman JM, Roberson ED, Rosen HJ, Rosenberg RN, Sano M, Saykin AJ, Schneider JA, Schneider LS, Seeley WW, Smith AG, Sonnen JA, Spina S, Stern RA, Tanzi RE, Thornton-Wells TA, Trojanowski JQ, Troncoso JC, Tsuang DW, Van Deerlin VM, Van Eldik LJ, Vardarajan BN, Vinters HV, Vonsattel JP, Weintraub S, Welsh-Bohmer KA, Williamson J, Wishnek S, Woltjer RL, Wright CB, Younkin SG, Yu CE, Yu L. Rarity of the Alzheimer disease-protective APP A673T variant in the United States. *JAMA Neurol* 2015; 72:209-216.
316. Ghani M, Reitz C, Cheng R, Vardarajan BN, Jun G, Sato C, Naj A, Rajbhandary R, Wang L-S, Valladares O, Lin C-F, Larson EB, Graff-Radford NR, Evans D, De Jager PL, Crane PK, Buxbaum JD, Murrell JR, Raj T, Ertekin-Taner N, Logue M, Baldwin CT, Green RC, Barnes LL, Cantwell LB, Fallin MD, Go RCP, Griffith P, Obesisan TO, Manly JJ, Lunetta KL, **Kamboh MI**, Lopez OL, Bennerr DA, Hendrie H, Hall KS, Goate AM, Byrd GS, Kukull WA, Foroud TM, Haines JL, Farrer LA, Perical-Vance MA, Lee JH, Schellenberg GD, St George-Hyslop P, Mayeux R, Rogaeva E, for the Alzheimer Disease Genetics Consortium. Association of long runs of homozygosity with Alzheimer's disease among African Americans. *JAMA Neurol* 2015; 72: 1313-1323.
317. Gregg NM, Kim AE, Gurol ME, Lopez OL, Aizenstein HJ, Price JC, Mathis CA, James JA, Snitz BE, Cohen AD, **Kamboh MI**, Minhas D, Weissfeld LA, Tamburo EL, Klunk WE. Incidental cerebral microbleeds and cerebral blood flow in elderly individuals. *JAMA Neurol* 2015; 72:1021-1028.

318. Demirci FY, Wang X, Kelly JA, Morris DL, Barmada B, Feingold E, Kao AH, Sivils KL, Bernatsky S, Pineau C, Clarke A, Ramsey-Goldman R, Vyse TJ, Gaffney PM, Manzi S, **Kamboh MI**. Identification of a new susceptibility locus for systemic lupus erythematosus on chromosome 12 in individuals of European ancestry. *Arthritis Rheumatol* 2016; 68: 174-183.
319. Barral S, Vardarajan BN, Faber KM, Bird TD, Tsuang D, Bennett DA, Rosenberg R, Boeve BF, Graff-Radford NR, Goate AM, Farlow M, Lantigua R, Medrano MZ, Wang X, **Kamboh MI**, Barmada MM, Schaid DJ, Foroud TM, Weamer EA, Ottman R, Sweet RA, Mayeux R; NIA-LOAD/NCRAD Family Study Group. Genetic variants associated with susceptibility to psychosis in late-onset Alzheimer's disease families. *Neurobiol Aging* 2015; 36: 3116.e9-e16.
320. Zia A, Wang X, Bhatti A, Demirci FY, Zhao W, Rasheed A, Samuel M, Kiani AK, Ismail M, Zafar J, John P, Saleheen D, **Kamboh MI**. A replication study of 49 type 2 diabetes risk variants in a Punjabi Pakistani population. *Diabet Med* 2016; 33:1112-1117.
321. Ombrello MJ, Remmers EF, Tachmazidou I, Grom A, Foell D, Hass J-P, Martini A, Gattorno M, Ozen S, Prahalad S, Zeft AS, Bhonsak JF, Mellins ED, Ilowite NT, Russo R, Len C, Hillario MOE, Oliveria S, Yeung RSM, Rosenberg A, Wedderburn LR, Anton J, Schwarz T, Hinks A, Bilginer Y, Park J, Cobb J, Satorius C, Han B, Baskin E, Signa S, Duerr R, Achkar JP, **Kamboh MI**, Kaufman K, Kotyan LC, Pinto D, Scherer SW, Alarcon-Riquelme ME, Decampo E, Estivill X, Gul A, British Society of Pediatric and Adolescent Rheumatology (BSPAR) study group, Childhood Arthritis Prospective Study (CAPS) group, Randomized Placebo Phase Study of Rilonacept in sJIA (RAPRORT) investigators, Sparks-Childhood Arthritis Response to Medication Study (CHARMS) group, Biologically Based Outcome Predictor in JIA (BBOP), de Bakker OIW, Raychaudhuri S, Longfeld CD, Thompson SD, Zeggini E, Thomson W, Kastner DL, Woo P, on behalf of the International Childhood Arthritis Genetics (INXHARGE) Consortium. *HLA-DRB1*11* and variants of the MHC class II locus are strong risk factors for systemic juvenile idiopathic arthritis. *Proc Natl Acad Sci USA* 2015; 112: 15970-15975.
322. Niemsiri V, Wang X, Pirim D, Radwan ZH, Bunker CH, Barmada MM, **Kamboh MI**, Demirci FY. Genetic contribution of *SCARB1* variants to lipid traits in African Blacks: a candidate gene association study. *BMC Med Genet* 2015; 16(1):106. doi: 10.1186/s12881-015-0250-6.
323. Prim D, Wang X, Niemsiri V, Radwan ZH, Bunker CH, Hokanson J, Hamman RF, Barmada MM, Demirci FY, **Kamboh MI**. Resequencing of the CETP gene in American whites and African blacks: Association of rare and common variants with HDL-cholesterol levels. *Metabolism* 2016; 65:36-47.
324. Mez J, Chung J, Jun G, Kriegel J, Bourlas AP, Sherva R, Logue MW, Barnes LL, Bennett DA, Buxbaum JD, Byrd GS, Crane PK, Ertekin-Taner N, Evans D, Fallin MD, Foroud T, Goate A, Graff-Radford NR, Hall KS, **Kamboh MI**, Kukull WA, Larson EB, Manly JJ; Alzheimer Disease Genetics Consortium., Haines JL, Mayeux R, Pericak-Vance MA, Schellenberg GD, Lunetta KL, Farrer LA. Two novel loci, *COBL* and *SLC10A2*, for Alzheimer's disease in African Americans. *Alzheimers Dement* 2017; 13:119-129.
325. Ombrello MJ, Arthur VL, Remmers EF, Hinks A, Tachmazidou I, Grom A, Foell D, Martini A, Gattorno M, Ozen S, Prahalad S, Zeft AS, Bhonsak JF, Ilowite NT, Mellins ED, Russo R, Len C, Hillario MOE, Oliveria S, Yeung RSM, Rosenberg A, Wedderburn LR, Anton J, Hass J-P, Rosen-Wolff A, Minden K, Tenbrook K, Demirkaya E, Cobb J, Baskin E, Signa S, Shuldiner E, Duerr RH, Achkar JP, **Kamboh MI**, Kaufman KM, Kotyan LC, Pinto D, Scherer SW, Alarcon-Riquelme ME, Decampo E, Estivill X, Gul A, British Society of Pediatric and Adolescent Rheumatology (BSPAR) study group, Inception Cohort of Newly Diagnosed Patients with Juvenile Idiopathic Arthritis (ICON-JIA), Childhood Arthritis Prospective Study (CAPS) group, Randomized Placebo Phase Study of Rilonacept in sJIA (RAPRORT) investigators, Sparks-Childhood Arthritis Response to Medication Study (CHARMS) group, Biologically Based

Outcome Predictor in JIA (BBOP), Langefled CD, Thompson S, Zeggini E, Kastner DL, Woo P, Thomson W. Genetic architecture distinguishes system juvenile idiopathic arthritis from other forms of juvenile idiopathic arthritis: Clinical and therapeutic implications. *Ann Rheum Dis* 2017; 76:906-913.

326. Demirci FY, Wang X, Morris DI, Feingold E, Bernatsky S, Pineau C, Clarke A, Ramsey-Goldman R, Manzi S, Vyse T, **Kamboh MI**. Multiple signals at the extended 8p23 locus are associated with susceptibility to systemic lupus erythematosus (SLE). *J Med Genet* 2017; 54:381-389.
327. Jun GR, Chung J, Mez J, Barber R, Beecham GW, Bennett DA, Buxbaum JD, Byrd GS, Carrasquillo MM, Crane PK, Cruchaga C, De Jager P, Ertekin-Taner N, Evans D, Fallin MD, Foroud T, Friedland RR, Goate A, Graff-Radford NR, Hendrie H, Hall KS, Hamilton-Nelson KL, Inzelberg R, **Kamboh MI**, Kauwe JSK, Kukull WA, Kunkle BW, Kuwano R, Larson EB, Logue MW, Manly JJ, Martin ER, Montine TJ, Mukerjee S, Naj A, Reiman EM, Reitz C, Sherva R, St George-Hyslop P, Thorton T, Younkin SG, Vardarajan BN, Wang L-S, Wendlund JR, Winslow AR, Alzheimer Disease Genetics Consortium., Haines JL, Mayeux R, Pericak-Vance MA, Schellenberg GD, Lunetta KL, Farrer LA. Transthectic genome-wide scan identifies novel Alzheimer disease loci. *Alzheimers Dement* 2017; 13: 727-738.
328. Sims R, van der Lee SJ, Naj AC, Bellenguez C, Badarinarayanan N, Jakobsdottir J, Kunkle BW, Boland A, Raybould R, Bis JC, Martin ER, Grenier-Boley B, Heilmann-Heimbach S, Chouraki V, Kuzma AB, Sleegers K, Vronskaya M, Ruiz A, Graham RR, Olaso R, Hoffmann P, Grove ML, Vardarajan BN, Hiltunen M, Nöthen MM, White CC, Hamilton-Nelson KL, Epelbaum J, Maier W, Choi SH, Beecham GW, Dulary C, Herms S, Smith AV, Funk CC, Derbois C, Forstner AJ, Ahmad S, Li H, Bacq D, Harold D, Satizabal CL, Valladares O, Squassina A, Thomas R, Brody JA, Qu L, Sánchez-Juan P, Morgan T, Wolters FJ, Zhao Y, Garcia FS, Denning N, Fornage M, Malamon J, Naranjo MCD, Majounie E, Mosley TH, Dombroski B, Wallon D, Lupton MK, Dupuis J, Whitehead P, Fratiglioni L, Medway C, Jian X, Mukherjee S, Keller L, Brown K, Lin H, Cantwell LB, Panza F, McGuinness B, Moreno-Grau S, Burgess JD, Solfrizzi V, Proitsi P, Adams HH, Allen M, Seripa D, Pastor P, Cupples LA, Price ND, Hannequin D, Frank-García A, Levy D, Chakrabarty P, Caffarra P, Giegling I, Beiser AS, Giedraitis V, Hampel H, Garcia ME, Wang X, Lannfelt L, Mecocci P, Eiriksdottir G, Crane PK, Pasquier F, Boccardi V, Henández I, Barber RC, Scherer M, Tarraga L, Adams PM, Leber M, Chen Y, Albert MS, Riedel-Heller S, Emilsson V, Beekly D, Braae A, Schmidt R, Blacker D, Masullo C, Schmidt H, Doody RS, Spalletta G, Jr WTL, Fairchild TJ, Bossù P, Lopez OL, Frosch MP, Sacchinelli E, Ghetti B, Yang Q, Huebinger RM, Jessen F, Li S, **Kamboh MI**, et al. [Multiple additional authors]. Rare coding variants in *PLCG2*, *ABI3* and *TREM2* implicate microglia-mediated innate immunity in Alzheimer's disease. *Nat Genet* 2017; 49: 1372-1384.
329. DeMichele-Sweet MA, Weamer EA, Klei L, Vrana DT, Hollingshead DJ, Seltman HJ, Sims R, Foroud T, Hernandez I, Moreno-Grau S, Tarraga L, Boada M, Ruiz A, Williams J, Mayeux R, Lopez OL, Sibile EL, **Kamboh MI**, Devlin B, Sweet RA. Genetic risk for schizophrenia and psychosis in Alzheimer disease. *Mol Psychiatry* 2018; 23: 963-972.
330. Zhao Y, Tudorascu DL, Lopez OL Cohen AD, Mathis CA, Aizenstein HJ, Price JC, Kuller LH, **Kamboh MI**, DeKosky ST, Klunk WE, Snitz BE. Amyloid-beta deposition and suspected non-Alzheimer pathophysiology show different cognitive decline patterns over 12 years in oldest-old without dementia. *JAMA Neurol* 2018; 75: 88-96.
331. Cheema AN, Rosenthal SL, **Kamboh MI**. Proficiency of data interpretation: identification of signaling SNPs/specific loci for coronary artery disease. *Database* 2017; 2017, 1 January 2017, bax078, <https://doi.org/10.1093/database/bax078>.
332. Snitz BE, Wang T, Cloonan YK, Jacobsen E, Chang CH, Hughes TF, **Kamboh MI**, Ganguli M. Risk of

progression from subjective cognitive decline to mild cognitive impairment: The role of study setting. *Alzheimers Dement* 2018; 14:734-742.

333. Lopez OL, Becker JT, Chang Y, Klunk WE, Mathis C, Price J, Aizenstein HJ, Snitz B, Cohen AD, DeKosky ST, Ikonomovic M, **Kamboh MI**, Kuller LH. Amyloid deposition and brain structure as long-term predictors of MCI, dementia, and mortality. *Neurology* 2018; 90: e1920-e1928.
334. Lyons CE, Tudorascu D, Snitz BE, Price J, Aizenstein H, Lopez O, Lopresti B, Laymon C, Minhas D, **Kamboh MI**, Mathis C, Klunk W, Cohen AD. The relationship of current cognitive activity to brain amyloid burden and glucose metabolism. *Am J Geriatr Psychiatry* 2018; 26:977-984.
335. The Brainstorm Consortium, Anttila V, Bulik-Sullivan B, Finucane HK, Walters RK, Bras J, Duncan L, Escott-Price V, Falcone GJ, Gormley P, Malik R, Patsopoulos NA, Ripke S, Wei Z, Yu D, Lee PH, Turley P, Grenier-Boley B, Chouraki V, Kamatani Y, Berr C, Letenneur L, Hannequin D, Amouyel P, Boland A, Deleuze JF, Duron E, Vardarajan BN, Reitz C, Goate AM, Huentelman MJ, **Kamboh MI**, et al [multiple additional authors]. Analysis of shared heritability in common disorders of the brain. *Science* 2018; 360 (6395): eaap8757. doi:10.1126/science.aap8757
336. Yan Q, Nho K, Del-Aguila JL, Wang X, Risacher SL, Fan KH, Snitz BE, Aizenstein HJ, Mathis CA, Lopez OL, Demirci FY, Feingold E, Klunk WE, Saykin AJ, Cruchaga C, **Kamboh MI**. Genome-wide association study of brain amyloid deposition as measured by Pittsburgh Compound-B (PiB)-PET imaging. *Mol Psychiatry* 2021; 26:309-321. Epub Oct 25, 2018 [*Paper selected for the Research Highlights section in Nat Rev Neurol 2018; 14:689 'Genetics influences development of AD pathology'*]
337. Ganguli M, Jia Y, Hughes TF, Snitz BE, Chang CH, Berman SB, Sullivan KJ, **Kamboh MI**. Mild cognitive impairment that does not progress to dementia: A population-based study. *J Am Geriatr Soc* 2018; 67:232-238.
338. Hu Z, Wang L, Ma S, Kirisci, L, Feng Z, Xue Y, Klunk WE, **Kamboh MI**, Sweet, RA, Becker J, Lv Q, Lopez OL, Xie X-U. Synergism of antihypertensives and cholinesterase inhibitors in Alzheimer's disease. *Alzheimers Dement (N Y)* 2018; 4:542-555.
339. Mukherjee S, Mezz J, Tritschuh E, Saylin A, Gibbons L, Fardo D, Wessels M, Bauman J, Moore M, Choi SE, Gross A, Rich J, Louden D, Saunders R, Grabowski T, Bird T, McCurry S, Snotz B, **Kamboh MI**, Lopez O, De Jager P, Bennett D, Keene C, Larson E, Crane P. Genetic data and cognitively-defined late-onset Alzheimer's disease subgroups. *Mol Psychiatry* 2020; 25:2942-2951.
340. **Kamboh MI**. A brief synopsis on the genetics of Alzheimer's disease. *Curr Genet Med Rep* 2018; 6:133-135.
341. Kunkle BW, Grenier-Boley B, Sims R, Bis JC, Damotte V, Naj AC, Boland A, Vronskaya M, van der Lee SJ, Amlie-Wolf A, Bellenguez C, Frizatti A, Chouraki V, Martin ER, Sleegers K, Badarinarayanan N, Jakobsdottir J, Hamilton-Nelson KL, Moreno-Grau S, Olaso R, Raybould R, Chen Y, Kuzma AB, Hiltunen M, Morgan T, Ahmad S, Vardarajan BN, Epelbaum J, Hoffmann P, Boada M, Beecham GW, Garnier JG, Harold D, Fitzpatrick AL, Valladares O, Moutet ML, Gerrish A, Smith AV, Qu L, Bacq D, Denning N, Jian X, Zhao Y, Del Zompo M, Fox NC, Choi SH, Mateo I, Hughes JT, Adams HH, Malamon J, Sanchez-Garcia F, Patel Y, Brody JA, Dombroski BA, Naranjo MCD, Daniilidou M, Eiriksdottir G, Mukherjee S, Wallon D, Uphill J, Aspelund T, Cantwell LB, Garzia F, Galimberti D, Hofer E, Butkiewicz M, Fin B, Scarpini E, Sarnowski C, Bush WS, Meslage S, Kornhuber J, White CC, Song Y, Barber RC, Engelborghs S, Sordon S, Voijnovic D, Adams PM, Vandenberghe R, Mayhaus M, Cupples LA, Albert MS, De Deyn PP, Gu W, Himali JJ, Beekly D, Squassina A, Hartmann AM, Orellana A, Blacker D, Rodriguez-Rodriguez E, Lovestone S, Garcia ME, Doody RS, Munoz-Fernandez C, Sussams

- R, Lin H, Fairchild TJ, Benito YA, Holmes C, Karamujić-Čomić H, Frosch MP, Thonberg H, Maier W, Roschupkin G, Ghetti B, Giedraitis V, Kawalia A, Li S, Huebinger RM, Kilander L, Moebus S, Hernández I, **Kamboh MI**, Brundin R, Turton J, et al. Genetic meta-analysis of diagnosed Alzheimer's disease identifies new risk loci and implicates A β , tau, immunity and lipid processing. *Nat Genet* 2019; 51:414-430.
342. Wu M, Thurston RC, Tudorascu DL, Karim HT, Mathis CA, Lopresti BJ, **Kamboh MI**, Cohen AD, Snitz BE, Klunk WE, Aizenstein HJ. Amyloid deposition is associated with different patterns of hippocampal connectivity in men versus women. *Neurobiol Aging* 2019; 76:141-150.
343. Pirim D, Radwan Z, Wang X, Niemsiri V, Hokanson JE, Hamman RF, Feingold E, Bunker CH, Demirci FY, **Kamboh MI**. Apolipoprotein E-C1-C4-C2 gene cluster region and inter-individual variation in plasma lipoprotein levels: a comprehensive genetic association study in two ethnic groups. *PLoS One* 2019; 14(3):e0214060.
344. **Kamboh MI**, Fan K, Yan Q, Beer JC, Wang X, Demirci FY, Snitz BE, Chang CH, Feingold E, Ganguli M. Population-based genome-wide association study of cognitive decline in older adults free of dementia: identification of a novel locus for the attention domain. *Neurobiol Aging* 2019; 84:239.e15-239e24.
345. Aslam MM, John P, Bhatti A, Jahangir S, **Kamboh MI**. Vitamin D a principal factor in mediating rheumatoid arthritis-derived immune system. *BioMed Res International* 2019; 2019: 3494937.
346. Rajakumar K, Yan Q, Khalid A, Feingold E, Vallejo A, Demirci FY, **Kamboh MI**. Gene expression and cardiometabolic phenotypes of vitamin D-deficient overweight and obese black children. *Nutrients* 2019; 11:2016; <http://dx.doi.org/10.3390/nu11092016>
347. Aslam MM, John P, Fan K-H, Bhatti A, Jahangir S, Feingold E, Demirci FY, **Kamboh MI**. Exploration of shared genetic susceptibility loci between type 1 diabetes and rheumatoid arthritis in the Pakistani population. *BMC Res Notes* 2019; 12:544.
348. Erickson KI, Grove GA, Burns JM, Hillman CH, Kramer AF, McAuley E, Vidoni ED, Becker JT, Butters MA, Gray K, Huang H, Jakicic JM, **Kamboh MI**, Kang C, Klunk WE, Lee P, Marsland AL, Mettenburg J, Rogers RJ, Stillman CM, Sutton BP, Szabo-Reed A, Verstynen TD, Watt JC, Weinstein AM, Wollam ME. Investigating gains in neurocognition in an Intervention Trial of Exercise (IGNITE): Protocol. *Contemp Clin Trials* 2019; 85:105832.
349. Reiman EM, Arboleda-Velasquez JF, Quiroz YT, Huentelman MJ, Beach TG, Caselli RJ, Chen Y, Su Y, Myers AJ, Hardy J, Vonsattel JP, Younkin SG, Bennett DA, De Jager PL, Larson EB, Crane PK, Keene D, **Kamboh MI**, Kofler JK, Duque L, Gilbert J, Gwirtsman H, Buxbaum J, Dickson D, Frosch M, Ghetti B, Lunetta K, Wang Li-San, Hyman B, Kukull W, Foroud T, Haines J, Mayeux R, Pericak-Vance M, Schneider J, Trojanowski J, Farrer L, Schellenberg G, Beecham G, Montine T, Jun G. Exceptionally low likelihood of Alzheimer's dementia in APOE2 homozygotes from a 5,000-person neuropathological study. *Nature Commun* 2020; 11:667.
350. Snitz BE, Chang Y, Tudorascu D, Lopez OL, Leprosto B, DeKosky ST, Carlson MC, Cohen AD, **Kamboh MI**, Aizenstein HJ, Klunk WE, Kuller LH. Predicting resistance to amyloid-beta deposition and cognitive resilience in the oldest-old. *Neurology* 2020; 95:e984-e994.
351. Sullivan KJ, Liu A, Chang C-C H, Cohen AD, Lopresti BJ, Minhas DS, Laumon CM, Klunk WE, Aizenstein

- H, Nadkarni NK, Loewenstein DL, **Kamboh MI**, Ganguli M, Snitz BE. Alzheimer's disease pathology in a community-based sample of older adults without dementia: The MYHAT Neuroimaging Study. *Brain Imag Behav* 2020; Aug 3. doi: 10.1007/s11682-020-00334-2. Online ahead of print.
352. Fan KH, Feingold E, Rosenthal SL, Demirci FY, Ganguli M, Lopez OL, **Kamboh MI**. Whole-exome sequencing (WES) analysis of Alzheimer's disease in non-*APOE**4 carriers. *J Alzheimers Dis*; 2020; 76:1553-1565.
353. Cheema AN, Pirim D, Wang X, Ali J, Bhatti A, John P, Feingold E, Demirci FY, **Kamboh MI**. Association study of coronary artery disease-associated genome-wide significant SNPs with coronary stenosis in Pakistani population. *Dis Markers* 2020; 2020:9738567.
354. Jia Y, Chang CCH, Hughes TF, Jacobsen E, Wang S, Berman SB, **Kamboh MI**, Ganguli M. Predictors of dementia in the oldest old: a novel machine learning approach. *Alzheimer Dis Assoc Disord* 2020; 34: 325-332.
355. Lopez O Kofler J Chang YF, Berman S, Becker J, Sweet R, Nadkarni NK, Patira R, **Kamboh MI**, Cohen A, Snitz B, Kuller L, Klunk W. Hippocampal sclerosis, TDP-43, and the duration of the symptoms of dementia of AD patients. *Ann Clin Transl Neurol* 2020; 7:1546-1556.
356. Aslam MM, John P, Fan K-H, Bhatti A, Aziz W, Ahmed B, Feingold E, Demirci FY, **Kamboh MI**. Investigating the GWAS-implicated loci for rheumatoid arthritis in the Pakistani population. *Dis Markers* 2020; 2020:1910215.
357. Aslam MM, Jalil F, John P, Fan KH, Bhatti A, Feingold E, Demirci FY, **Kamboh MI**. A sequencing study of *CTLA4* in Pakistani rheumatoid arthritis cases. *PLoS One* 2020; 15(9):e0239426.
358. Aslam MM, John P, Fan K-H, Bhatti A, Feingold E, Demirci FY, **Kamboh MI**. Association of *VPREB1* gene copy number variation and rheumatoid arthritis susceptibility. *Dis Markers* 2020; 2020:7189626.
359. Pirim D, Bunker CH, Hokanson JE, Hamman RF, Demirci FY, **Kamboh MI**. Hepatic lipase (LIPC) sequencing in individuals with extremely high and low high-density lipoprotein cholesterol levels. *PLoS One* 2020; 15(12): e0243919.
360. Sabiha B, Bhatti A, Fan, K-H, John P, Aslam MM, Ali J, Feingold E, Demirci FY, **Kamboh MI**. Assessment of genetic risk of type 2 diabetes among Pakistanis based on GWAS-implicated loci. *Gene* 2021; 783:145563.
361. Kunkle BW, Schmidt M, Klein HU, Naj AC, Hamilton-Nelson KL, Larson EB, Evans DA, De Jager PL, Crane PK, Buxbaum JD, Ertekin-Taner N, Barnes LL, Fallin MD, Manly JJ, Go RCP, Obisesan TO, **Kamboh MI**, Bennett DA, Hall KS, Goate AM, Foroud TM, Martin ER, Wang LS, Byrd GS, Farrer LA, Haines JL, Schellenberg GD, Mayeux R, Pericak-Vance MA, Reitz C; Writing Group for the Alzheimer's Disease Genetics Consortium (ADGC). Novel Alzheimer disease risk loci and pathways in African American individuals using the African Genome Resources Panel: A meta-analysis. *JAMA Neurol* 2021;17:102-113
362. Khan MJ, Desaire H, Lopez OL, **Kamboh MI**, Robinson RAS. Why inclusion matters for Alzheimer's disease biomarker discovery in plasma. *J Alzheimers Dis*; 2021; 79:1327-1344.
363. Khan MJ, Desaire H, Lopez OL, **Kamboh MI**, Robinson RAS. Dataset of why inclusion matters for Alzheimer's disease biomarker discovery in plasma. *Data in Brief* 2021; 35:106923.
364. Kim T, Kim S-Y, Agarwal V, Cohen A, Roush R, Chang Y-F, Cheng Y, Snitz B, Huppert TJ, Basic A,

Kamboh MI, Doman J, Becker JT. Cardiac-induced cerebral pulsatility, brain structure, and cognition in older adults. *Neuroimage* 2021; 233:117956.

365. DeMichele-Sweet MAA, Klei L, Creese B, Weamer EA, McClain L, Sims R, Hernandez I, Moreno-Grau S, Tárraga L, Boada M, Alarcón-Martín M, Valero S, NIA-LOAD Family Based Study Consortium, Alzheimer's Disease Genetics Consortium (ADGC), Liu Y, Hooli B, Aarsland D, Selbaek G, Bergh S, Rongve A, Saltvedt I, Skjellegrend HK, Engdahl B, Stordal E, Andreassen OA, Djurovic S, Athanasiu L, Seripa D, Borroni B, Albani D, Forloni G, Meccoci P, Serretti A, De Ronchi D, Politis A, Paroni G, AddNeuroMed Consortium, Williams J, Mayeux R, Foroud T, Ruiz A, Ballard C, Lopez OL, **Kamboh MI**, Devlin B, Sweet RA. Genome-wide association identifies the first risk loci for psychosis in Alzheimer disease. *Mol Psychiatry* 2021; 2021; 26:5797-5811.
366. Fitz NF, Wang J, **Kamboh MI**, Koldanova R, Lefterov I. Small nucleolar RNAs in plasma extracellular vesicles and their discriminatory power as diagnostic biomarkers of Alzheimer's disease. *Neurobiol Dis* 2021 Aug 16:105481. doi: 10.1016/j.nbd.2021.105481. Online ahead of print.
367. **Kamboh MI**. Genomics and functional genomics of Alzheimer's disease. *Neurotherapeutics* 2022; 19:152-172.
368. Khan MJ, Chung NA, Hansen S, Dumitrescu L, Hohman TJ, **Kamboh MI**, Lopez OL, Robinson RAS. Targeted lipidomics to understand health disparities in Alzheimer's disease. *Analytic Chem* 2022; 94:4165-4174.
369. Harper JD, Fan KH, Aslam MM, Snitz BE, DeKosky ST, Lopez OL, Feingold E, **Kamboh MI**. Genome-wide association study of incident dementia in a community-based sample of older subjects. *J Alzheimer Dis* 2022; Jun 9. doi: 10.3233/JAD-220293. Online ahead of print.

2. Additional Publications Listed in PubMed in Secondary/Collaborator Authors List as Part of Consortia

1. Snitz BE, O'Meara ES, Carlson MC, Arnold AM, Ives DG, Rapp SR, Saxton J, Lopez OL, Dunn LO, Sink KM, DeKosky ST; **Ginkgo Evaluation of Memory (GEM) Study Investigators**. Ginkgo biloba for preventing cognitive decline in older adults: a randomized trial. *JAMA* 2009; 302:2663-2670.
2. Brinkley TE, Lovato JF, Arnold AM, Furberg CD, Kuller LH, Burke GL, Nahin RL, Lopez OL, Yasar S, Williamson JD; **Ginkgo Evaluation of Memory (GEM) Study Investigators**. Effect of Ginkgo biloba on blood pressure and incidence of hypertension in elderly men and women. *Am J Hypertens* 2010; 23:528-533.
3. Kauwe JS, Cruchaga C, Karch CM, Sadler B, Lee M, Mayo K, Latu W, Su'a M, Fagan AM, Holtzman DM, Morris JC; **Alzheimer's Disease Neuroimaging Initiative**, Goate AM. Fine mapping of genetic variants in BIN1, CLU, CR1 and PICALM for association with cerebrospinal fluid biomarkers for Alzheimer's disease. *PLoS One* 2011; 6(2):e15918.
4. Tosun D, Schuff N, Mathis CA, Jagust W, Weiner MW; **Alzheimer's Disease NeuroImaging Initiative**. Spatial patterns of brain amyloid-beta burden and atrophy rate associations in mild cognitive impairment. *Brain* 2011; 134:1077-1088.
5. McEvoy LK, Holland D, Hagler DJ Jr, Fennema-Notestine C, Brewer JB, Dale AM; **Alzheimer's Disease Neuroimaging Initiative**. Mild cognitive impairment: baseline and longitudinal structural MR imaging measures improve predictive prognosis. *Radiology* 2011; 259:834-43.
6. Eskildsen SF, Coupé P, Fonov V, Manjón JV, Leung KK, Guizard N, Wassef SN, Østergaard LR, Collins DL; **Alzheimer's Disease Neuroimaging Initiative**. BEaST: brain extraction based on nonlocal segmentation technique.

Neuroimage 2012; 59:2362-2373.

7. Cho Y, Seong JK, Jeong Y, Shin SY; **Alzheimer's Disease Neuroimaging Initiative**. Individual subject classification for Alzheimer's disease based on incremental learning using a spatial frequency representation of cortical thickness data. *Neuroimage* 2012; 59:2217-2230.
8. Shen KK, Fripp J, Mériudeau F, Chételat G, Salvado O, Bourgeat P; **Alzheimer's Disease Neuroimaging Initiative**. Detecting global and local hippocampal shape changes in Alzheimer's disease using statistical shape models. *Neuroimage* 2012; 59:2155-2166.
9. Yasar S, Lin FM, Fried LP, Kawas CH, Sink KM, DeKosky ST, Carlson MC; **Ginkgo Evaluation of Memory (GEM) Study Investigators**. Diuretic use is associated with better learning and memory in older adults in the Ginkgo Evaluation of Memory Study. *Alzheimers Dement* 2012; 8:188-195.
10. Zou F, Chai HS, Younkin CS, Allen M, Crook J, Pankratz VS, Carrasquillo MM, Rowley CN, Nair AA, Middha S, Maharjan S, Nguyen T, Ma L, Malphrus KG, Palusak R, Lincoln S, Bisceglie G, Georgescu C, Kouri N, Kolbert CP, Jen J, Haines JL, Mayeux R, Pericak-Vance MA, Farrer LA, Schellenberg GD; **Alzheimer's Disease Genetics Consortium**, Petersen RC, Graff-Radford NR, Dickson DW, Younkin SG, Ertekin-Taner N. Brain expression genome-wide association study (eGWAS) identifies human disease-associated variants. *PLoS Genet* 2012; 8(6):e1002707.
11. Chapman B, Duberstein P, Tindle HA, Sink KM, Robbins J, Tancredi DJ, Franks P; **Ginkgo Evaluation of Memory Study Investigators**. Personality predicts cognitive function over 7 years in older persons. *Am J Geriatr Psychiatry* 2012; 20:612-21.
12. Jun G, Vardarajan BN, Buros J, Yu CE, Hawk MV, Dombroski BA, Crane PK, Larson EB; **Alzheimer's Disease Genetics Consortium**, Mayeux R, Haines JL, Lunetta KL, Pericak-Vance MA, Schellenberg GD, Farrer LA. Comprehensive search for Alzheimer disease susceptibility loci in the APOE region. *Arch Neurol* 2012; 69:1270-1279.
13. Zou F, Chai HS, Younkin CS, Allen M, Crook J, Pankratz VS, Carrasquillo MM, Rowley CN, Nair AA, Middha S, Maharjan S, Nguyen T, Ma L, Malphrus KG, Palusak R, Lincoln S, Bisceglie G, Georgescu C, Kouri N, Kolbert CP, Jen J, Haines JL, Mayeux R, Pericak-Vance MA, Farrer LA, Schellenberg GD; **Alzheimer's Disease Genetics Consortium**, Petersen RC, Graff-Radford NR, Dickson DW, Younkin SG, Ertekin-Taner N. Brain expression genome-wide association study (eGWAS) identifies human disease-associated variants. *PLoS Genet* 2012; 8(6):e1002707
14. Jun G, Vardarajan BN, Buros J, Yu CE, Hawk MV, Dombroski BA, Crane PK, Larson EB; **Alzheimer's Disease Genetics Consortium**, Mayeux R, Haines JL, Lunetta KL, Pericak-Vance MA, Schellenberg GD, Farrer LA. Comprehensive search for Alzheimer disease susceptibility loci in the APOE region. *Arch Neurol* 2012; 69(10):1270-9
15. Miyashita A, Koike A, Jun G, Wang LS, Takahashi S, Matsubara E, Kawarabayashi T, Shoji M, Tomita N, Arai H, Asada T, Harigaya Y, Ikeda M, Amari M, Hanyu H, Higuchi S, Ikeuchi T, Nishizawa M, Suga M, Kawase Y, Akatsu H, Kosaka K, Yamamoto T, Imagawa M, Hamaguchi T, Yamada M, Morihara T, Takeda M, Takao T, Nakata K, Fujisawa Y, Sasaki K, Watanabe K, Nakashima K, Urakami K, Ooya T, Takahashi M, Yuzuriha T, Serikawa K, Yoshimoto S, Nakagawa R, Kim JW, Ki CS, Won HH, Na DL, Seo SW, Mook-Jung I; **Alzheimer Disease Genetics Consortium**, St George-Hyslop P, Mayeux R, Haines JL, Pericak-Vance MA, Yoshida M, Nishida N, Tokunaga K, Yamamoto K, Tsuji S, Kanazawa I, Ihara Y, Schellenberg GD, Farrer LA, Kuwano R. SORL1 is genetically associated with late-onset Alzheimer's disease in Japanese, Koreans and Caucasians. *PLoS One* 2013; 8(4):e58618.
16. Holton P, Ryten M, Nalls M, Trabzuni D, Weale ME, Hernandez D, Crehan H, Gibbs JR, Mayeux R, Haines JL, Farrer LA, Pericak-Vance MA, Schellenberg GD; **Alzheimer's Disease Genetics Consortium**, Ramirez-Restrepo M, Engel A, Myers AJ, Corneveaux JJ, Huettel MJ, Dillman A, Cookson MR, Reiman EM, Singleton A, Hardy J, Guerreiro R. Initial assessment of the pathogenic mechanisms of the recently identified Alzheimer risk Loci. *Ann Hum Genet* 2013; 77(2):85-105
17. Reitz C, Mayeux R; **Alzheimer's Disease Genetics Consortium**. TREM2 and neurodegenerative disease. *N Engl J Med* 2013; 369(16):1564-5.

18. Ridge PG, Mukherjee S, Crane PK, Kauwe JS; **Alzheimer's Disease Genetics Consortium**. Alzheimer's disease: analyzing the missing heritability. *PLoS One* 2013; 8(11):e79771.
19. Tate JA, Snitz BE, Alvarez KA, Nahin RL, Weissfeld LA, Lopez O, Angus DC, Shah F, Ives DG, Fitzpatrick AL, Williamson JD, Arnold AM, DeKosky ST, Yende S; Infection hospitalization increases risk of dementia in the elderly. **GEM Study Investigators**. *Crit Care Med* 2014; 42:1037-1046.
20. Østergaard SD, Mukherjee S, Sharp SJ, Proitsi P, Lotta LA, Day F, Perry JR, Boehme KL, Walter S, Kauwe JS, Gibbons LE; **Alzheimer's Disease Genetics Consortium**; GERAD1 Consortium; EPIC-InterAct Consortium, Larson EB, Powell JF, Langenberg C, Crane PK, Wareham NJ, Scott RA. Associations between potentially modifiable risk factors and Alzheimer Disease: A Mendelian randomization study. *PLoS Med* 2015; 12(6):e1001841.
21. Ridge PG, Hoyt KB, Boehme K, Mukherjee S, Crane PK, Haines JL, Mayeux R, Farrer LA, Pericak-Vance MA, Schellenberg GD, Kauwe JS; **Alzheimer's Disease Genetics Consortium (ADGC)**. Assessment of genetic variance of late-onset Alzheimer's disease. *Neurobiol Aging* 2016; 41: 200.e13-20.
22. Peloso GM, van der Lee SJ; **International Genomics of Alzheimer's Project (IGAP)**, Destefano AL, Seshardi S. Genetically elevated high-density lipoprotein cholesterol through the cholesteryl ester transfer protein gene does not associate with risk of Alzheimer's disease. *Alzheimers Dement (N Y)* 2018;4:542-555.
23. Chauhan G, Adams HHH, Satizabal CL, Bis JC, Teumer A, Sargurupremraj M, Hofer E, Trompet S, Hilal S, Smith AV, Jian X, Malik R, Traylor M, Pulit SL, Amouyel P, Mazoyer B, Zhu YC, Kaffashian S, Schilling S, Beecham GW, Montine TJ, Schellenberg GD, Kjartansson O, Guðnason V, Knopman DS, Griswold ME, Windham BG, Gottesman RF, Mosley TH, Schmidt R, Saba Y, Schmidt H, Takeuchi F, Yamaguchi S, Nabika T, Kato N, Rajan KB, Aggarwal NT, De Jager PL, Evans.....; Stroke Genetics Network (SiGN), the International Stroke Genetics Consortium (ISGC), METASTROKE, **Alzheimer's Disease Genetics Consortium (ADGC)**, and the Neurology Working Group of the Cohorts for Heart and Aging Research in Genomic Epidemiology (CHARGE) Consortium. *Neurology* 2019; Jan 16. pii: 10.1212/WNL.0000000000006851.
24. Bellenguez C, Küçükali F, Jansen IE, Kleineidam L... GR@ACE; DEGESCO; EADI; GERAD; Demgene; FinnGen; **ADGC**; CHARGE,et al. New insights into the genetic etiology of Alzheimer's disease and related dementias. *Nat Genet* 2022; 54:412-436.
25. Heath L, Earls JC, Magis AT, Kornilov SA, Lovejoy JC, Funk CC, Rappaport N, Logsdon BA, Mangravite LM, Kunkle BW, Martin ER, Naj AC, Ertekin-Taner N, Golde TE, Hood L, Price ND; **Alzheimer's Disease Genetics Consortium**. Manifestations of Alzheimer's disease genetic risk in the blood are evident in a multiomic analysis in healthy adults aged 18 to 90. *Sci Rep* 2022;12(1):6117.
26. Queder N, Phelan MJ, Taylor L, Tustison N, Doran E, Hom C, Nguyen D, Lai F, Pulsifer M, Price J, Kreisl WC, Rosas HD, Krinsky-McHale S, Brickman AM, Yassa MA, Schupf N, Silverman W, Lott IT, Head E, Mapstone M, Keator DB; **Alzheimer's Biomarkers Consortium**. Joint-label fusion brain atlases for dementia research in Down syndrome. *Alzheimers Dement (Amst)*. 2022 May 25;14(1):e12324.

3. Book Chapters; Book Reviews; Letters to the Editor

1. Kirk RL, **Kamboh MI**: Pacific peoples: origin and genetic relationships. In: *Genetic Microdifferentiation in Human and Other Animal Populations*. (eds: Ahuja YR, Neel JV). Indian Anthropologist, Occasional Papers in Anthropology University of Delhi, Delhi, India.1985; No. 1, pp. 62-79.
2. Chakraborty R, **Kamboh MI**, Ferrell RE: Response to: Issues in estimating Caucasian admixture in American blacks (letter to the editor). *Am J Hum Genet* 1992; 51:680-681.

3. **Kamboh MI:** Genetic analysis at the phenotypic level. *Encyclopedia of Immunology* 1992; pp. 608-611.
4. **Kamboh MI:** A Laboratory Manual for Human Blood Analysis by Bhagat MK, Chahal SMS (Book Review). *Am J Phys Anthropol* 1998; 105:555-556.
5. **Kamboh MI:** PCR3: PCR *In Situ* Hybridization, edited by Herrington CS, O'Leary JJ (Book Review). *Am J Hum Biol* 1999; 11: 694.
6. **Kamboh MI:** Decoding Darkness: The Search for Genetic Causes of Alzheimer's Disease by Tanzi RE, Parson AB (Book Review). *American Scientist* 2001; 89:83-84.
7. **Kamboh MI:** Genetics of Alzheimer's Disease. In: *Handbook of Medical Psychiatry* (eds: Soares JC, Gershon S). Marcel Dekker, Inc., New York, 2003, pp. 521-536.
8. Omalu BI, DeKosky ST, Minster RL, **Kamboh MI**, Hamilton RL, Wecht CH: Reply to Casson et al. Chronic traumatic encephalopathy (CTE) in the National Football League (NFL) player. *Neurosurgery* 2006; 58: E1003.

4. Published Abstracts and Presentations in Scientific Meetings

1. **Kamboh MI:** Alpha-1-antitrypsin (PI) and transferrin (TF) subtyping in Asian, Pacific and Australian Aboriginal populations. *Pathology* 1983; 15:106.
2. **Kamboh MI:** Isoelectric focusing of PGM1 and GC polymorphisms in different ethnic groups from the Asia, Pacific and Australian area. *Pathology* 1984; 16:103.
3. **Kamboh MI:** Genetic polymorphism of thyroxine-binding globulin in the pacific area and reassessment of its proposed association with α_1 -antitrypsin (PI). *Aust Pediatr J* 1985; 21:313.
4. **Kamboh MI**, Ferrell RE: Heterogeneity of TBG: A new polymorphic genetic variant in Eskimos determined by an immunoblotting technique. *Am J Hum Genet* 1985; 37:A198.
5. Eichner JE, Ferrell RE, **Kamboh MI:** Isoelectric focusing studies of steroid binding proteins. *Am J Hum Genet* 1986; 39:A8.
6. Escallon MH, Ferrell RE, **Kamboh MI:** Genotyping of alpha-1-acid glycoprotein (orosomucoid) by an isoelectric focusing immunoblotting technique. *Am J Hum Genet* 1986; 39:A9.
7. Ferrell RE, **Kamboh MI:** Genetic diversity in the human coagulation factor XIIIIB plasma protein. *Am J Hum Genet* 1986; 39:A234.
8. **Kamboh MI**, Ferrell RE: Immunobiochemical detection of new polymorphisms by IEF-protein blotting methods. *Am J Hum Genet* 1986; 39:A238.
9. **Kamboh MI**, Ferrell RE, Sepehrnia B: Genetics of human apolipoproteins. *Am J Hum Genet* 1987; 41:A8.
10. Sepehrnia B, Ferrell RE, Adams L, **Kamboh MI:** Sex-specific effect of apolipoprotein C-II genotype on lipoprotein levels in Nigerian Blacks. *Am J Hum Genet* 1987; 41:A82.
11. Lyons LA, **Kamboh MI**, Ferrell RE: Population genetics of the ClR and ClS subcomponents of the first

- complement component C1. *Am J Hum Genet* 1987; 41:A258.
12. Leppert M, Ferrell RE, **Kamboh MI**, Beasley J, O'Connell P, Lathrop M, Lalouel JM, White R: Linkage of the polymorphic protein markers F13B, CIS, CIR and blood group antigen Kidd in the CEPH reference families. *Cytogenet Cell Genet* 1987; 46:647.
 13. Ferrell RE, **Kamboh MI**, Freidrich CF: Paternity diagnosis in chimpanzees: Results from the N.I.H. Chimpanzee Breeding and Research Program. *Am J Phys Anthropol* 1988; 75:208.
 14. **Kamboh MI**, Ferrell RE: The anthropological significance of new protein polymorphisms. *Am J Phys Anthropol* 1988; 75:228.
 15. Ferrell RE, **Kamboh MI**, Sepehrnia B, Weiss KM: Genetic variation in apolipoproteins C-II and C-III. *4th Int. Colloq. On Atherosclerosis*, Brussels. 1988; p. 34.
 16. **Kamboh MI**, Sepehrnia B, Ferrell RE: Apolipoprotein E: methodology, polymorphism and, allelic effects on quantitative levels of lipids and lipoproteins in Blacks. *Am J Hum Genet* 1988; 43:A9.
 17. Sepehrnia B, **Kamboh MI**, Ferrell RE: Investigation of the apolipoprotein H polymorphism in relation to plasma lipoprotein concentrations. *Am J Hum Genet* 1988; 43:A222.
 18. Ferrell RE, Sepehrnia B, **Kamboh MI**, Majumder PP: Genetic variation in apolipoprotein C-II and its impact on lipoprotein levels. *Am J Hum Genet* 1988; 43:A48.
 19. Cole SA, **Kamboh MI**, Stern MP, Ferrell RE: Genetic variation at apolipoprotein loci in the Mexican-American population of San Antonio. *Am J Hum Genet* 1988; 43:A212.
 20. Lyons LA, **Kamboh MI**, Ferrell RE: Linkage and population genetics studies of CIR and CIS complement components. *Am J Hum Genet* 1988; 43:A150.
 21. **Kamboh MI**, Ferrell RE: The significance of apolipoprotein D polymorphism in anthropogenetics and lipid metabolism. *Am J Phys Anthropol* 1989; 78:249-250.
 22. Deka R, **Kamboh MI**, Ferrell RE, Smouse PE: Gene and gene product variation at the transferrin locus in Melanesians of New Guinea. *Am J Phys Anthropol* 1989; 78:211-212.
 23. Ferrell RE, **Kamboh MI**, Majumder PP, Valdez R, Weiss KM: Quantitative genetic variation in apolipoprotein C-III and its impact on cholesterol metabolism in Mayans of Yucatan Peninsula, Mexico. *Am J Phys Anthropol* 1989; 78:220.
 24. Eichner JE, Ferrell RE, **Kamboh MI**: Apolipoprotein E phenotypes and cholesterol in IDDM. *Diabetes* 1989; 38(Suppl. II):211A.
 25. Eichner JE, Kuller LH, Ferrell RE, **Kamboh MI**: The influence of the apolipoprotein E polymorphism in perimenopausal women. *2nd International Conference on Preventive Cardiology*. Washington, D.C. 1989; June 18-22, p. 46.
 26. **Kamboh MI**, Weiss KM, Ferrell RE: Apolipoprotein E polymorphism and its impact on cholesterol levels in the Mayans of Yucatan Peninsula, Mexico. *Am J Hum Genet* 1989; 45:241.
 27. Ferrell RE, **Kamboh MI**, Bhatia KK: Population genetics of apolipoprotein polymorphisms in Papua New

- Guinea. *Am J Hum Genet* 1989; 45:A239.
28. Eichner JE, Kuller LH, Ferrell RE, **Kamboh MI**: Phenotypic effects of the apolipoprotein E locus in White and Black perimenopausal healthy women. *Am J Hum Genet* 1989; 45:A237.
 29. Cauley JA, Eichner JE, **Kamboh MI**, Ferrell RE, Kuller LH, Black DM: APO E allele frequencies in younger (age 42-50) vs. older (age 65-90) women. *Circulation*. 1990; 81:725.
 30. **Kamboh MI**, Serjeantson RE, Ferrell RE: Population genetics of apolipoprotein polymorphisms in Australian aborigines. *Am J Phys Anthropol* 1990; 81:246-247.
 31. **Kamboh MI**, Ferrell RE: A high resolution SDS-agarose electrophoresis method to screen apolipoprotein(a) polymorphism. *Arteriosclerosis*. 1990; 10:826a.
 32. **Kamboh MI**, Chakraborty R, Ferrell RE: Caucasian genes in the American Blacks: New data. *Am J Hum Genet* 1990; 47:A318.
 33. Crews DE, Bindon JR, **Kamboh MI**: Apolipoprotein polymorphisms in American Samoa. 1990.
 34. **Kamboh MI**, Ferrell RE: A high resolution SDS-agarose electrophoresis method to screen apolipoprotein(a) polymorphism. *International Meeting of the Electrophoresis Society*, Washington, D.C. 1991; March 19-21.
 35. Iyengar S, Hamman RF, **Kamboh MI**, Marshall JA, Baxter J, Majumder PP, Ferrell RE: Amerindian admixture among the Anglo and Hispanic ethnic groups in the San Luis Valley, Colorado. *Am J Phys Anthropol* 1991; (Suppl12):97-98.
 36. **Kamboh MI**, Hamman RF, Iyengar S, Ferrell RE: Apolipoprotein A-IV polymorphism and its impact on lipoprotein-lipid levels in normal and non-insulin dependent diabetes. *Am J Phys Anthropol* 1991; (Supplement 12):102.
 37. **Kamboh MI**, Kelly LJ, Ferrell RE: Apolipoproteins E and A-IV phenotyping from whole blood and blood stains. *International Meeting of the Electrophoresis Society*, Washington, D.C. 1991; March 19-21.
 38. Eichner JE, Cauley JA, **Kamboh MI**, Ferrell RE, Kuller LH, Black D: Lp(a) concentration in older women: Distribution and association with other risk factors. *Annual Meeting of the Society for Epidemiologic Research*, Buffalo, NY. 1991; June 12-14.
 39. Crews DE, Bindon JR, **Kamboh MI**: Apolipoprotein polymorphisms and body habitus in American Samoans. *Towards the Pacific Century; The Challenge of Change, XVII*. Pacific Science Conference, Meeting of Abstract Volume, Waikiki, Hawaii. 1991; May 26-June 2, pg. 23.
 40. **Kamboh MI**, Aston C, Hamman R, Ferrell RE: The impact of the apolipoprotein E polymorphism on lipoprotein-lipid levels in the Anglo and Hispanic ethnic groups in the San Luis Valley, Colorado. *Am J Human Genet* 1991; 49 (Suppl):472.
 41. Ferrell RE, **Kamboh MI**, Iyengar S, Aston C, Hamman R: Apolipoprotein A- IV and lipoprotein-lipid levels among Hispanics and non-Hispanic Whites in the San Luis Valley, CO. *Am J Hum Genet* 1991; 49 (Suppl):469.
 42. **Kamboh MI**: Current status about the role of APOH genetic polymorphism in lipid metabolism and new perspectives. *Antiphospholipid Antibody/Lupus Anticoagulant Workshop*. NIH, Bethesda, MD. 1991;

September 25.

43. **Kamboh MI**, Aston CE, Ferrell RE, Hamman RF: Apolipoprotein E polymorphism and lipid concentrations in normal and NIDDM subjects. *Am J Phys Anthropol* 1992; (Suppl 14):97.
44. Hamman RF, Ahn YI, Ferrell RE, **Kamboh MI**: A study of association of the HindIII RFLP at the lipoprotein lipase (LPL) locus and levels of plasma lipids and carbohydrates in the Anglo and Hispanic populations in the San Luis Valley, Colorado. *Am J Phys Anthropol* 1992; (Suppl 14):87.
45. Crews DE, **Kamboh MI**, Mancilha-Carvalho JJ, Kottke B: Population genetics of apolipoprotein E, A-IV, H polymorphisms and effects on lipids, lipoproteins and carbohydrate metabolism in Yanomami Indians. *Am J Phys Anthropol* 1992; (Suppl 14):65.
46. Shetterly S, **Kamboh MI**, Marshall J, Ferrell RE: Apolipoprotein E and risk of myocardial infarction, the San Luis Valley Diabetes Study. *Diabetes*. 1992; 41:180A.
47. **Kamboh MI**: APO(a) polymorphism reflected by phenotyping. *Second International Conference on Lipoprotein(a)*, New Orleans, Louisiana. 1992; November 12-14.
48. Ahn YI, **Kamboh MI**, Ferrell RE, Hamman RF: Associations of three DNA polymorphisms in the lipoprotein lipase (LPL) and LDL receptor (LDLR) loci with plasma levels of lipid and carbohydrate traits in Hispanics and non-Hispanic Whites (NHWs). *Am J Hum Genet* 1992; 51(Suppl): A144.
49. **Kamboh MI**, Williams ER, Law J, Aston CE, Bunker CH, Pollitzer WJ, Ferrell RE: Molecular basis of an unique African variant (A-IV 5) of human apolipoprotein A-IV and its significance in lipid metabolism. *Am J Hum Genet* 1992; 51(Suppl):A152.
50. Roube RAS, Buyon JP, Winfield JB, **Kamboh MI**: Phenotypic analysis of β_2 -glycoprotein I in patients with antiphospholipid autoantibodies. *Fifth International Symposium on Antiphospholipid Antibody*, San Antonio. 1992; September 9-12.
51. **Kamboh MI**, Ahn YI, Ferrell RE: Polymorphism and partial DNA sequence of the apolipoprotein A-IV gene in the chimpanzee. *Am J Phys Anthropol* 1993; (Suppl. 16):121.
52. Deka R, **Kamboh MI**, Ferrell RE, McGarvey ST: Genetic characterization of American and Western Samoans. *Am J Phys Anthropol* 1993; (Suppl 16):85.
53. Ahn YI, Ferrell RE, Hamman RF, **Kamboh MI**: An association study of the lipoprotein lipase (LPL) polymorphism with the physiological components of the Insulin Resistance Syndrome in the population of the San Luis Valley, Colorado. *Am J Phys Anthropol* 1993; (Suppl 16):48.
54. Rewers M, **Kamboh MI**, Shetterly SM, Hoag S, Ferrell RE, Hamman RF: Apolipoprotein A-IV phenotype associated with myocardial infarction in diabetes: The San Luis Valley Diabetes Study. *33rd Annual Conference on Cardiovascular Disease Epidemiology*, Santa Fe, New Mexico. 1993; March 17-20.
55. Hamman RF, Ahn YI, Ferrell RE, **Kamboh MI**: Association of lipoprotein lipase (LPL) gene variation with the Insulin Resistance Syndrome: The San Luis Valley Diabetes Study. *Diabetes* 1993; 42 (Suppl 1):212A.
56. Rewers M, **Kamboh MI**, Shetterly SM, Ferrell RE: The increased risk of myocardial infarction in diabetes and its ethnic variation are associated with the APO A-IV Phenotype: The San Luis Valley Diabetes

- Study. *Diabetes* 1993; 42 (Suppl 1):119A.
57. **Kamboh MI**, Evans RW, Ahn YI: Apolipoprotein(a) and E genetic polymorphisms in relation to plasma quantitative levels of lipoprotein(a) and APOB in American Blacks. *Council on Arteriosclerosis*, 1993; p.101.
 58. **Kamboh MI**, Friedlaender JS, Ahn YI, Ferrell RE: A common deletion polymorphism in the apolipoprotein A-IV gene and its association with lipoprotein-lipid levels. *Circulation* 1993; 88 (Suppl): I-267.
 59. Marshall JA, **Kamboh MI**, Hoag S, Hamman RF, Ferrell RE: Association of ApoE polymorphism and dietary cholesterol with serum lipid levels. *Circulation* 1993; 88 (Suppl): I-509.
 60. Rewers M, **Kamboh MI**, Shetterly SM, Hoag S, Ferrell RE, Hamman RF: Association of apolipoprotein H polymorphism with hypertension: The San Luis Valley Diabetes Study. *Circulation* 1993; 88 (Suppl): I-513.
 61. **Kamboh MI**, Aston C, Ferrell RE, Hamman RF: The relationship between APOE polymorphism and cholesterol levels in normoglycemic and diabetic subjects. *Am J Hum Genet* 1993; 53 (Suppl):816.
 62. Ahn YI, **Kamboh MI**, Ferrell RE, Hamman RF: The role of common polymorphisms in the LDL receptor gene in affecting cholesterol levels in the general population. *Am J Hum Genet* 1993; 53 (Suppl):773.
 63. **Kamboh MI**: Genetic markers for studies of coronary heart disease risk factors. Presented in a symposium "Genetics of Quantitative Risk Factors for Coronary Heart Disease" organized by chaired by M.I. Kamboh. The American Association of Physical Anthropologists (AAPA) annual meetings held in Denver, CO. 1994; April 1.
 64. Ferrell RE, Ahn YI, Hamman RF, **Kamboh MI**: Quantitative effects of LDL-receptor locus variation in the population of the San Luis Valley, Colorado. *Am J Phys Anthropol* 1994; (Suppl 18):87.
 65. Marshall JA, **Kamboh MI**, Hoag S, Hamman RF, Ferrell RE: Association of apolipoprotein E polymorphism and dietary cholesterol with serum cholesterol levels. The San Luis Valley Diabetes Study. *Am J Phys Anthropol* 1994; (Suppl 18):137.
 66. Rewers M, **Kamboh MI**, Hamman RF: Association of apolipoprotein A-IV polymorphism with myocardial infarction in non-insulin dependent diabetes mellitus. The San Luis Valley Diabetes Study. *Am J Phys Anthropol* 1994; (Suppl 18):168.
 67. **Kamboh MI**: Genetics of quantitative lipoprotein (a) levels. *Am J Phys Anthropol* 1994; (Suppl 18):118.
 68. Wagenknecht DR, **Kamboh MI**, McIntyre JA: An apolipoprotein H*3 allele product which does not bind anionic phospholipids. 6th International symposium on Antiphospholipid Antibodies, Belgium. 1994; September 14-17.
 69. **Kamboh MI**: Polymorphisms at the apolipoprotein A4 and plsaminogen loci affect plasma lipoprotein (a) levels in American Blacks. *Council on Arteriosclerosis* 1994; p 47.
 70. **Kamboh MI**, Aston CE, Rewers M, Hamman RF: Relationship between the apolipoprotein A4 polymorphism and plasma LP(a) levels in Hispanic men and women. *Council on Arteriosclerosis* 1994; p 54.
 71. **Kamboh MI**, DeKosky ST, Ferrell RE: Over-representation of the *APOE*4* allele in autopsy confirmed

- early- and late-onset sporadic Alzheimer's disease. *Am J Hum Genet* 1994; 55(Suppl):A154.
72. Wang GQ, **Kamboh MI**, Aston CE, Hamman RF: Role of the apolipoprotein A-IV polymorphism at codon 347 in affecting normal plasma lipid and apolipoprotein variation. *Am J Hum Genet* 1994; 55(Suppl):A168.
 73. Sanghera DK, Ferrell RE, Aston CE, **Kamboh MI**, McAllister AE, Kimm SY: Quantitative effects of the apolipoprotein E polymorphism in a biethnic sample of young females. *Am J Hum Genet* 1994; 55(Suppl):A164.
 74. Ferrell RE, **Kamboh MI**, Aston CE: Apolipoprotein (a) polymorphism and serum lipoprotein(a) levels in the chimpanzee. *Am J Phys Anthropol* 1995; (Suppl 20):90.
 75. **Kamboh MI**, Crawford MH, Leonard WR: Apolipoprotein E polymorphism and serum cholesterol levels among the Evenki herders of Siberia. *Am J Phys Anthropol* 1995; (Suppl 20):121.
 76. Friedlaender JS, Merriwether A, **Kamboh MI**: New indicators of Bougainville genetic distinctiveness. *Am J Phys Anthropol* 1995; (Suppl 20):93.
 77. DeKosky ST, Ferrell RE, Becker JT, **Kamboh MI**: Natural history of definite AD as a function of APOE genotypes. *Neurology* 1995; 45(Suppl 4): A373-A374.
 78. McGarvey ST, Kamboh MI, Galanis DJ: ApoE polymorphism and change in lipid and lipoprotein levels in modernizing Samoans. *Am J Hum Biol* 1995; 7:130-131.
 79. DeKosky ST, Sanghera DK, Ferrell RE, **Kamboh MI**: Alpha-1-antichymotrypsin (ACT) genetic polymorphism modifies the risk of Alzheimer's disease conferred by *APOE*4*. *Ann Neurology* 1995; 38:282.
 80. Sanghera DK, Ferrell RE, Aston CE, McAllister AE, **Kamboh MI**, Kimm SYS: Quantitative effects of a common genetic polymorphism in the APOA4 gene in a biracial sample of young females. *Am J Hum Genet* 1995; 57(Suppl): A171.
 81. Aston CE, McAllister AE, Marshall JA, Hoag S, Hamman RF, **Kamboh MI**: Association of LDL-receptor polymorphism and dietary factors with serum lipid levels in Hispanics. *Am J Hum Genet* 1995; 57(Suppl): A159.
 82. Wang GQ, Hamman RF, Aston CE, **Kamboh MI**: Haplotype analysis of two APOA4 polymorphisms in relation to plasma lipoprotein-lipid levels. *Am J Hum Genet* 1995; 57(Suppl): A174.
 83. Harris MR, Hamman RF, Aston CE, **Kamboh MI**: Role of low density lipoprotein-related protein (LRP) gene polymorphism in lipid metabolism. *Am J Hum Genet* 1995; 57(Suppl): A164.
 84. Saha N, Sanghera DK, **Kamboh MI**: Role of common genetic polymorphism in the paraoxonase gene in lipoprotein metabolism. *Am J Hum Genet* 1995; 57(Suppl): A11.
 85. Ali S, Bunker CH, Aston CE, Ukoli FA, **Kamboh MI**: Contribution of the APO(a) size polymorphism in affecting plasma Lp(a) levels in Africans. *Am J Hum Genet* 1995; 57(Suppl): A159.
 86. **Kamboh MI**, Bunker CH, Aston CE, Ukoli FA, Svitko CM, McAllister AE: Relationship of the APOE polymorphism and socioeconomic status in affecting plasma cholesterol levels. *Am J Hum Genet* 1995;

57(Suppl): A165.

87. **Kamboh MI**, Aston CE, McAllister AE, Hamman RF: Haplotype analysis of two APOA1/MsPI polymorphisms in relation to plasma apoA-I and HDL-cholesterol levels. Council on Arteriosclerosis, American Heart Association, 1995, p 85.
88. DeKosky ST, Farlow M, Sanghera DK, Aston CE, **Kamboh MI**: Modification of *APOE**4-mediated risk of Alzheimer's disease by alpha-1-antichymotrypsin in familial AD: Results from the Indiana Alzheimer Center National Cell Repository. *Neurology* 1996; 46:A418.
89. McGarvey ST, **Kamboh MI**: ApoE polymorphism and cross-sectional lipid and lipoprotein levels in adult American Samoans. *Am J Phys Anthropol* 1996; (Suppl 22) 163.
90. Fitton LJ, Crews DE, **Kamboh MI**: Apolipoprotein polymorphisms:Relationship with quantitative lipid and apolipoprotein levels in American Samoans. *Am J Phys Anthropol* 1996; (Suppl 22): 104.
91. Lopez OL, Lopez-Pousa S, **Kamboh MI**, Lozano-Gallego M, Oliva R, Adroer R, DeKosky ST: Apolipoprotein E in Alzheimer's disease:A comparative study of two populations from Spain and the United States. *Neurologia* 1996; 11:363.
92. **Kamboh MI**, Aston CE, Low PS, Saha N: Racial and genetic determinants of plasma Factor XIII activity. *Am J Hum Genet* 1996; 59 (Suppl):A181.
93. Aston CE, Nestlerode CS, McAllister AE, Hamman RF, **Kamboh MI**: Haplotype analysis of two APOA1/*MspI* polymorphisms in relation to plasma levels ApoA-I and HDL-cholesterol. *Am J Hum Genet* 1996; 59 (Suppl):A174.
94. Harris MR, Bunker CH, Sanghera DK, Aston CE, **Kamboh MI**: Association of low density lipoprotein receptor-related protein (LRP) polymorphism with plasma lipid and apolipoprotein levels in African Blacks. *Am J Hum Genet* 1996; 59 (Suppl):A179.
95. Anderson JL, Bunker CH, Aston CE, **Kamboh MI**: Relationship of two APOB polymorphisms with plasma quantitative traits in African Blacks. *Am J Hum Genet* 1996; 59(Suppl):A173.
96. Nestlerode CS, Sanghera DK, Bunker CH, **Kamboh MI**: Molecular basis of apolipoprotein J protein polymorphism and its impact on serum lipid and apolipoprotein levels. *Am J Hum Genet* 1996; 59 (Suppl):A185.
97. Sanghera DK, Saha N, Aston CE, **Kamboh MI**: Genetic polymorphism of paraoxonase and the risk of coronary heart disease. *Am J Hum Genet* 1996; 59 (Suppl):A188.
98. Ali S, Bunker CH, McAllister AE, Aston CE, **Kamboh MI**: A pentanucleotide repeat polymorphism in the APO(a) gene is a significant determinant of serum Lp(a) levels in Blacks. *Am J Hum Genet* 1996; 59 (Suppl):A173.
99. McAllister AE, Bunker CH, Aston CE, **Kamboh MI**: Interaction between BMI and APOE polymorphism influences serum triglyceride levels. *Am J Hum Genet* 1996; 59 (Suppl):A184.
100. Wang GQ, Saha N, Vashist S, **Kamboh MI**: Influence of two APOA4 polymorphisms (Thr 347 Ser and Gln 360 His) on postprandial serum lipid levels. *Am J Hum Genet* 1996; 59 (Suppl):A192.

101. Myung P, McGarvey ST, Steel MS, Bausserman L, **Kamboh MI**, Galanis DJ: Cross-sectional interrelations among diet, lipids and lipoproteins, and ApoE genotype in adult Samoans. *Am J Hum Biol* 1997; 9:137-138.
102. DeKosky ST, Styren SD, **Kamboh MI**: Regional distribution of alpha1-antichymotrypsin immunoreactivity in Alzheimer's disease. *Neurology* 1997; 48:A103-A104.
103. Markey J, Sanghera DK, Hamman RF, **Kamboh MI**: Role of the paraoxonase 1 codon 192 polymorphism in insulin metabolism. *Am J Hum Genet* 1997; 61 (Suppl): A205.
104. **Kamboh MI**, Sanghera DK, Aston CE, Saha N: A common polymorphism in the paraoxonase 2 gene (PON2) is associated with the risk of coronary heart disease. *Am J Hum Genet* 1997; 61 (Suppl): A202.
105. Mehdi H, Sanghera DK, Manzi S, **Kamboh MI**: Genetic variation in the apolipoprotein H gene influence plasma apolipoprotein H concentrations. *Am J Hum Genet* 1997; 61 (Suppl): A206.
106. Nestlerode CS, Sanghera DK, Bunker CH, **Kamboh MI**: The effect of a codon 328 apolipoprotein J polymorphism on serum lipid and apolipoprotein levels in a population from Benin City, Nigeria. *Am J Hum Genet* 1997; 61 (Suppl): A207.
107. Razzaghi H, Day BW, **Kamboh MI**: A three-dimensional molecular model for human lipoprotein lipase and structural analysis of D9N and N291S mutations. *Am J Hum Genet* 1997; 61 (Suppl): A180.
108. Saha N, Low PS, **Kamboh MI**: Association of two polymorphisms in the lipoprotein lipase gene with coronary heart disease. *Am J Hum Genet* 1997; 61 (Suppl): A209.
109. Sanghera DK, Saha N, **Kamboh MI**: Genetic polymorphisms in the paraoxonase 1 (PON1) gene and the risk of coronary heart disease. *Am J Hum Genet* 1997; 61 (Suppl): A210.
110. Wang X, DeKosky ST, Aston CE, **Kamboh MI**: Genetic association of two chromosome 14 genes (PS1 and ACT) with Alzheimer's disease. *Am J Hum Genet* 1997; 61 (Suppl): A214.
111. Sweet RA, Nimgoankar VL, **Kamboh MI**, Lopez OL, Taffe K, DeKosky ST: Genetic variation, psychosis, and aggression in Alzheimer's disease. *Soc Neurosci* 1997; 23:29.
112. **Kamboh MI**, Mehdi H, Sanghera DK, Hamman RF: Genetic variation in the apolipoprotein H gene is a major determinant of plasma apolipoprotein H concentrations. *Circulation* 1998; 98 (Suppl I): I-790.
113. Bhojak T, Bunker CH, **Kamboh MI**: Polymorphism in the Apolipoprotein F gene and its relation to lipid levels in Nigerian Blacks. *Am J Hum Genet* 1998; 63(Suppl): A208.
114. Chiu L, Hamman RF, Aston CE, **Kamboh MI**: Relationship of the kringle 4 polymorphism in the apo(a) gene and plasma Lp(a) levels in non-Hispanic Whites. *Am J Hum Genet* 1998; 63(Suppl): A210.
115. DePrince KM, McGarvey ST, Bausserman L, Aston CE, **Kamboh MI**: Plasma Lp(a) distribution and effect of two apo(a) repeat polymorphisms (kringle 4 and pentanucleotide repeats) on plasma Lp(a) levels in American Samoans. *Am J Hum Genet* 1998; 63(Suppl): A211.
116. Desai PP, Bunker CH, Aston CE, **Kamboh MI**: Identification of three novel missense mutations in the apolipoprotein D gene and their significance in lipid metabolism. *Am J Hum Genet* 1998; 63(Suppl): A324.

117. Heng CK, Lim SL, Saha N, Tong MC, Tan YS, Low PS, **Kamboh MI**: Apo(a) gene polymorphisms of Asian Indians in Singapore: Association with plasma Lp(a) levels and coronary artery disease (CAD). *Am J Hum Genet* 1998; 63(Suppl): A213.
118. **Kamboh MI**, Manzi S, Mehdi H, Sanghera DK: Genetic variation in apolipoprotein H (β_2 -glycoprotein I) affects the occurrence of antiphospholipid antibodies and apolipoprotein H concentrations in lupus patients. *Am J Hum Genet* 1998; 63(Suppl): A330.
119. Luedeking EK, DeKosky ST, **Kamboh MI**: Genetic polymorphisms in transforming growth factor- β 1 and the risk of Alzheimer's disease. *Am J Hum Genet* 1998; 63(Suppl): A216.
120. Razzaghi H, McClure RJ, Day BW, **Kamboh MI**: Molecular modeling of interaction between lipoprotein lipase and its cofactor, apoC-II. *Am J Hum Genet* 1998; 63(Suppl): A190.
121. Saha N, Heng CK, Low PS, Tong MC, **Kamboh MI**: Genetic variations of the promoter region of protein C gene determines plasma protein C antigen levels and CAD risk in Asian Indians. *Am J Hum Genet* 1998; 63(Suppl): A220.
122. Wang X, DeKosky ST, **Kamboh MI**: Characterization of polymorphisms and linkage disequilibrium in the ACT gene and the risk of Alzheimer's disease. *Am J Hum Genet* 1998; 63(Suppl): A313.
123. **Kamboh MI**: Genetics of apolipoprotein H (β_2 -glycoprotein I) and anionic phospholipid binding. Invited Speaker to the 8th International Symposium on Antiphospholipid Antibodies. Sapporo, Japan, October 6-9, 1998.
124. Ikonomovic, MD, O'Malley M, Ciallella JR, Wisniewski S, Aston C, **Kamboh MI**, Farlow M, DeKosky ST: Alpha-1-antichymotrypsin (ACT) serum levels are not affected by ACT signal peptide polymorphism. *Soc Neurosci* 1999; 25:832.
125. **Kamboh MI**, Manzi S, Mehdi H, Fitzgerald S, Sanghera DK, Kuller LH, Aston CE: Genetic variation in apolipoprotein H (β_2 -glycoprotein I) affects the occurrence of antiphospholipid antibodies and apolipoprotein H concentrations in systemic lupus erythematosus. *Arthritis Rheum* 1999; 42(Suppl):S233.
126. Mehdi H, Naqvi A, **Kamboh MI**: An hydrophobic sequence at positions 313-316 (Leu-Ala-Phe-Trp) in the fifth domain of apolipoprotein H (β_2 -glycoprotein I) is critical for cardiolipin binding. *Arthritis Rheum* 1999; 42(Suppl):S233.
127. Bhojak T, DeKosky ST, **Kamboh MI**: The cathepsin D Ala224Val polymorphism is not associated with the risk of Alzheimer's disease. *Am J Hum Genet* 1999; 65(Suppl): A243.
128. Chiu L, Hamman RF, Aston CE, **Kamboh MI**: Apolipoprotein(a) polymorphisms and plasma lipoprotein(a) concentrations in non-Hispanic Whites and Hispanics. *Am J Hum Genet* 1999; 65(Suppl): A199.
129. Chen Q, Markey J, Hamman RF, **Kamboh MI**: Genetic variation in two paraoxonase genes (PON1 and PON2) and the risk of coronary heart disease in Hispanics and non-Hispanic Whites. *Am J Hum Genet* 1999; 65(Suppl): A245.
130. Heng CK, Ho WFK, Saha N, Tong MC, Tan YS, **Kamboh MI**: The relationship of Lp(a) levels with coronary heart disease is genotype dependent for three apo(a) gene polymorphisms in Asian Indians in

Singapore. *Am J Hum Genet* 1999; 65(Suppl): A205.

131. Low PS, Heng CK, Quek SC, Saha N, **Kamboh MI**: Significant influence of the +93 C/T polymorphism in the apolipoprotein(a) gene on Lp(a) concentrations in the Asian Indian neonates from Singapore. *Am J Hum Genet* 1999; 65(Suppl): A210.
132. Luedeking EK, DeKosky ST, Mehdi H, **Kamboh MI**: Genetic polymorphisms in transforming growth factor- β_1 and the risk of Alzheimer's disease. *Am J Hum Genet* 1999; 65(Suppl): A210.
133. Mehdi H, Naqvi A, **Kamboh MI**: Recombinant hepatitis B surface antigen particles and anionic phospholipid bind to identical sites on apolipoprotein H (β_2 -glycoprotein I) as revealed by site-directed mutagenesis. *Am J Hum Genet* 1999; 65(Suppl): A376.
134. Razzaghi H, Aston CE, **Kamboh MI**: Cladistic diplotype: a novel method for organizing genotype data for statistical analysis. *Am J Hum Genet* 1999; 65(Suppl): A395.
135. Saha N, Luedeking EK, **Kamboh MI**: genetic polymorphisms in transforming growth factor- β_1 and the risk of coronary heart disease. *Am J Hum Genet* 1999; 65(Suppl): A397.
136. Wang X, DeKosky ST, **Kamboh MI**: Lack of association between the α_2 -macroglobulin polymorphism and Alzheimer's disease. *Am J Hum Genet* 1999; 65(Suppl): A401.
137. **Kamboh MI**, Aston CE, Hamman RF: DNA sequence variation in human apolipoprotein C4 gene and its effect on plasma lipid profile. *Circulation* 1999; 100(Suppl): I-5.
138. Parra EJ, Hiester K, Argyropoulos G, Garvey WT, Kittles RA, Sylvester N, Parrish-Gause D, **Kamboh I**, Ferrell RE, Shriver MD: High resolution admixture map of African Americans living in South Carolina. *Am J Phys Anthropol* 2000; (Suppl 30):245-246.
139. Ikonomovic MD, Ciallella JR, O'Malley M, Wisniewski S, Aston C, **Kamboh MI**, DeKosky ST: High levels of serum ACT correlate with cognitive impairment in Alzheimer's disease. *Soc Neurosci* 2000; 26:
140. Ikonomovic MD, Desai PP, **Kamboh MI**, Hamilton R, DeKosky ST: Apolipoprotein D distribution in normal and Alzheimer's brains: Implication for its role in the pathogenesis of Alzheimer's disease. *Neurobiol Aging* 2000; 21 (Suppl):S82.
141. **Kamboh MI**: Identification of novel Alzheimer genes using population-based association studies (Invited Speaker by the World Alzheimer Congress). *Neurobiol Aging* 2000; 21(Suppl):S212.
142. Mehdi H, **Kamboh MI**: Identification of two mutations in the third domain of apolipoprotein H (β_2 -glycoprotein I) and their impact on cardiolipin binding. *J Autoimmun* 2000; 15:A34.
143. Razzaghi H, Aston CE, Hamman RF, **Kamboh MI**: Genetic screening of the lipoprotein lipase gene for mutations associated with high triglyceride/low HDL-cholesterol levels. *Am J Hum Genet* 2000; 67(Suppl):212.
144. Sweet RA, **Kamboh MI**, Wisniewski SR, Lopez OL, Klunk WE, Kaufer DI, DeKosky ST: APOE and ACT genotypes do not predict time to psychosis in Alzheimer's disease. *Soc Neurosci* 2000; 26:1828.
145. **Kamboh MI**: Molecular genetics of Alzheimer's disease (Invited Speaker by the Human Biology

- Association). *Am J Hum Biol* 2000; 13:126-127.
146. Sweet RA, Umapathy C, Nimgoankar VL, **Kamboh MI**, Lopez OL, DeKosky ST: Role for Norepinephrine in Treatment of Apathy in Dementia. Proceedings, 13th Annual meeting of the American Association for Geriatric Psychiatry, Miami Beach, FL, March 2000.
 147. Hokanson JE, Hamman RF, **Kamboh MI**: Gene-environmental interaction in the regulation of HDL: The hepatic lipase gene promoter polymorphism limits the physical activity associated increase in HDL. *Circulation* 2001; 103:1352.
 148. Chen Q, Reis SE, McNamara D, Hovubkov R, **Kamboh MI**: Analysis of association between the severity of coronary disease and paraoxonase gene polymorphism in the Women's Ischemia Syndrome Evaluation (WISE) study. *Arterioscler Thromb Vas Biol* 2001; 21:679.
 149. Mehdi H, Desai P, Wang X, Zarnegar R, Strom S, **Kamboh MI**: Reduced expression of β 2-glycoprotein I (apolipoprotein H) associated with a transcriptional initiation site mutation (-1C→A) that also disrupts the binding of hepatic nuclear factors. *Am J Hum Genet* 2001; 69(Suppl):A365.
 150. Chen Q, Reise SE, Pauly D, Sharaf B, McNamara D, Holubkov R, Bairey Merz CN, **Kamboh MI**: Association between the apolipoprotein E polymorphism and the severity of coronary artery disease in the NHLBI-sponsored Women's Ischemia Syndrome Evaluation (WISE) Study. *Am J Hum Genet* 2001; 69(Suppl):A402.
 151. Desai PP, Hendrie HC, Evans RM, Murrell JR, DeKosky ST, **Kamboh MI**: Genetic variation in apolipoprotein D affects the risk of Alzheimer's disease in African Americans. *Am J Hum Genet* 2001; 69(Suppl):A416.
 152. Razzaghi H, **Kamboh MI**: VGT-SSCP: A highly sensitive and non-radioactive mutation detection method based on vertical gradient temperature SSCP. *Am J Hum Genet* 2001; 69(Suppl):A438.
 153. Luedeking-Zimmer E, DeKosky ST, **Kamboh MI**: Candidate genes for late-onset Alzheimer's disease on chromosome 12. *Am J Hum Genet* 2001; 69(Suppl):A518.
 154. Hokanson JE, **Kamboh MI**, Eckel RH, Hamman RF: The hepatic lipase promoter polymorphism is associated with an increase in coronary heart disease. *Circulation* 2001; 104(Suppl):II-808.
 155. Desai PP, Ikonomic MD, Ciallela JR, Wilbur Y, **Kamboh MI**, DeKosky ST: Apolipoprotein D forms plaques in the hippocampus of patients with Alzheimer's disease. *Soc Neurosci* 2001; 27:859.
 156. Butters MA, Sweet RA, Mulsant BH, **Kamboh MI**, Pollock BG, Nebes RD, Begley AE, DeKosky ST, Reynolds CF III: APOE is associated with age-of-onset, but not cognition in late-life depression. 2001 AAGP.
 157. Wilson RK, Luedeking EK, **Kamboh MI**, DeKosky ST, Hamilton RL: Association of the APOE4 allele and Lewy bodies in Alzheimer's disease. *J Neuropathol* 2002; 61:454.
 158. Luedeking-Zimmer E, DeKosky ST, Ganguli M, **Kamboh MI**: Association of transcription factor LBP-1c/CP2/LSF on chromosome 12 with the risk of Alzheimer's disease. *Neurobiol Aging* 2002; 23:S337-338.
 159. **Kamboh MI**, Luedeking-Zimmer E, Ganguli M, DeKosky ST: The oxidized LDL-receptor gene on chromosome 12 with the risk of Alzheimer's disease. *Neurobiol Aging* 2002; 23:S338-339.

160. Ikonomovic MD, Desai PP, **Kamboh MI**, Hamilton RL, Klunk WE, DeKosky ST: Apolipoprotein D and β -amyloid in progression of neuritic plaques. A study of Alzheimer's disease and Lewy body variant. *Neurobiol Aging* 2002; 23:S394.
161. Mehdi H, Manzi S, Desai PP, Chen Q, Wang X, Strom S, Zarnegar R, **Kamboh MI**: A functional $\beta 2$ -glycoprotein-I promoter polymorphism is associated with reduced gene expression and plasma levels. *Lupus* 2002; 11:544.
162. Desai PP, Ikonomovic MD, Hamilton RL, DeKosky ST, **Kamboh MI**: Apolipoprotein D in Alzheimer's disease: Implication for amyloid pathology. *Soc Neurosci* 2002; in press.
163. Chen C, Reis SE, Kammerer C, Luedeking-Zimmer E, McNamara DM, Pauley DF, Sharaf B, Holubkov R, Bairey-Meiz CN, Sopko G, **Kamboh MI**: Genetic variation in the oxidized LDL-receptor-1 (OLR1) gene and the risk of coronary artery disease. *Am J Hum Genet* 2002; 71 (Suppl): 348.
164. Desai PP, DeKosky ST, **Kamboh MI**: Association between the ABCA1/G1051A polymorphism and Alzheimer's disease. *Am J Hum Genet* 2002; 71 (Suppl): 366.
165. Razzaghi H, Hamman RF, **Kamboh MI**: Identification of a novel mutation in the 3' untranslated region (3'UTR) of the lipoprotein lipase gene in Hispanics and its association with the components of glucose homeostasis. *Am J Hum Genet* 2002; 71 (Suppl): 333.
166. Mosher MJ, **Kamboh MI**, Sorenson M, Leonard W: Effect of APO E polymorphisms, gender, and nutrients on plasma LDL in the Buryat. *Am J Hum Biol* 2003; 15: 275-276.
167. **Kamboh MI**, Sanghera DK, Mehdi H, Nestlerode CS, Chen Q, Bunker CH: Evolution of human apolipoprotein H gene haplotypes and their significance in the production of antiphospholipid antibodies and binding to anionic phospholipid. *Am J Hum Genet* 2003; 73(Suppl):376.
168. Sanghera DK, Manzi S, Bontempo F, Nestlerode C, **Kamboh MI**: Association of a regulatory intronic polymorphism in the *PDCD1* gene with the risk of sporadic systemic lupus erythematosus and the occurrence of antiphospholipid antibodies. *Am J Hum Genet* 2003; 73(Suppl):384.
169. Desai PP, DeKosky ST, Hendrie HC, Murrell JR, **Kamboh MI**: The ABCA1/G1051A polymorphism and Alzheimer's disease in Whites and African Americans. *Am J Hum Genet* 2003; 73(Suppl):504.
170. Minster RL, Desai PP, Ozturk A, Chen Q, Ganguli M, Nebes RD, Reynolds III CF, DeKosky ST, **Kamboh MI**: Association of the APOE promoter polymorphisms with late-onset Alzheimer's disease. *Neurobiol Aging* 2004; 25(suppl): S512-S513.
171. Desai PP, Ganguli M, Nebes RD, DeKosky ST, **Kamboh MI**: Lack of association in the brain-derived neurotrophic factor (BDNF) and sporadic late-onset Alzheimer's disease in Caucasians. *Neurobiol Aging* 2004; 25(suppl): S515.
172. Ozturk A, Desai PP, Minster RL, Scott T, Kammerer CM, DeKosky ST, **Kamboh MI**: Alpha-1 antichymotrypsin (ACT or SERPINA3) may affect age-at-onset of Alzheimer's disease. *Neurobiol Aging* 2004; 25(suppl): S496.
173. Conley YP, DeKosky ST, **Kamboh MI**, Finegold DN, Ferrell RE: The calcium sensing receptor in Alzheimer disease susceptibility. *Neurobiol Aging* 2004; 25(suppl): S509.

174. **Kamboh MI**, Jacobs E, Suresh S, Chen Q, Kenney M, Shaw P, Kao A, Kammerer C, Manzi S: Genetic variation in the APOH (β 2GPI) gene is associated with carotid vascular disease in women with Systemic Lupus Erythematosus (SLE). *Arthritis Rheum* 2005; 52 (Suppl) S616.
175. **Kamboh MI**, Tripi L, Chen Q, Kenney M, Shaw P, Kao A, Kammerer C, Bontempo F, Manzi S: Low serum paraoxonase activity is associated with Systemic Lupus Erythematosus (SLE). *Arthritis Rheum* 2005; 52 (Suppl) S615-S616.
176. Shih PB, Manzi S, Shaw P, Kenney M, , Kao A, Kammerer C, Bontempo F, Manzi S, **Kamboh MI**: Association between C-reactive protein polymorphisms and Systemic Lupus Erythematosus . *Arthritis Rheum* 2005; 52 (Suppl) S615.
177. Chen Q, Xu WJ, Razzaghi H, **Kamboh MI**: Functional characterization of the *HindIII* polymorphism in the lipoprotein lipase gene. *Annual American Society of Human Genetics Meeting, Salt Lake City, Utah, October 25-29, 2005*.
178. Minster R, DeKosky ST, **Kamboh MI**: Association of ubiquilin 1 (UBQLN1) SNPs with late-onset Alzheimer's disease in a large case-control sample. *Annual American Society of Human Genetics Meeting, Salt Lake City, Utah, October 25-29, 2005*.
179. Jacobs E, Chen Q, Kenney M, Shaw P, Kao A, Kammerer C, Bontempo F, Manzi S, **Kamboh MI**: Promoter polymorphisms in the apolipoprotein H (APOH) gene and the risk of systemic lupus erythematosus. *Annual American Society of Human Genetics Meeting, Salt Lake City, Utah, October 25-29, 2005*.
180. Ozturk A, DeKosky ST, Kamboh MI: Investigation of the relationship of candidate genes on chromosome 10 with the risk of and age-at-onset of Alzheimer's disease. *Annual American Society of Human Genetics Meeting, Salt Lake City, Utah, October 25-29, 2005*.
181. Tripi L, Chen Q, Kenney M, Shaw P, Kao A, Kammerer C, Bontempo F, Manzi S, **Kamboh MI**: Association of paraoxonase-1 (PON1) polymorphisms with lupus nephritis. *Annual American Society of Human Genetics Meeting, Salt Lake City, Utah, October 25-29, 2005*.
182. Yang L-X, Razzaghi H, **Kamboh MI**: Functional study of a novel 5 bp deletion located in a putative insulin response element in exon 10 of human lipoprotein lipase gene. *Annual American Society of Human Genetics Meeting, Salt Lake City, Utah, October 25-29, 2005*.
183. Sanghera DK, Bhatti JS, Bhatti GK, Ralhan SK, Wander GS, Singh JR, Bunker CH, Weeks DE, **Kamboh MI**, Ferrell RE: The Khatri Sikh Diabetes Study (SDS): Study design, methodology, sample collection and initial results. *Annual American Society of Human Genetics Meeting, Salt Lake City, Utah, October 25-29, 2005*.
184. Chen Q, Kamboh MI: Complete DNA sequence variation in the apolipoprotein H (B2-glycoprotein I) gene and informative SNPs. The 3rd Hong Kong Medical Genetic Conference, April 8-10, 2005.
185. Ikonomovic M, Abrahamson E, **Kamboh MI**, DeKosky S. Glutathione-S-transferase M1-1 protein levels in acute and chronic neurological disorders: Correlation with changes in amyloid-beta peptide. *Neurology* 66:A283, 2006.
186. Shih PB, Manzi S, Shaw P, Kenney M, Kao A, Kammerer C, **Kamboh MI**. C-reactive protein and cardiovascular disease risk in patients with systemic lupus erythematosus. *Arthritis Rheum* 2006;

54(suppl) S295.

187. Dasgupta S, Demirci FY, Minster RL, Kenney M, Shaw P, Kao A, Kammerer C, Bontempo F, Manzi S, **Kamboh MI**: Association of *PON2* variants with PON activity in systemic lupus erythematosus (SLE). *The American Society of Human Genetics Annual Meeting*, October 9-13, 2006. New Orleans, LA
188. Minster RL, DeKosky ST, **Kamboh MI**: Lack of association of two SNPs on chromosome 10 (*rs498055* and *rs4417206*) with Alzheimer's Disease. *The American Society of Human Genetics Annual Meeting*, October 9-13, 2006. New Orleans, LA
189. Sanghera DK, Manzi S, Shaw P, Kao A, Bontempo F, Kammerer C, **Kamboh MI**: Genetic variation in the paraoxonase 3 (PON3) gene is associated with serum PON activity. *The American Society of Human Genetics Annual Meeting*, October 9-13, 2006. New Orleans, LA
190. Suresh S, Minster RL, Demirci FY, Kenney M, Shaw P, Kao A, Kammerer C, Bontempo F, Manzi S, **Kamboh MI**: Effect of *APOH* promoter SNPs on gene expression and plasma levels of APOH. *The American Society of Human Genetics Annual Meeting*, October 9-13, 2006. New Orleans, LA
191. Kodavali V, Sweet RA, **Kamboh MI**, Nimgaonkar VL: Chromosome 6q21: Positional candidate genes for psychosis in Alzheimer's and Schizophrenia. *The American Society of Human Genetics Annual Meeting*, October 9-13, 2006. New Orleans, LA
192. Demirci FY, Manzi S, Minster RL, Bontempo F, Shaw PS, Kao AH, **Kamboh MI**: Association of a common interferon regulatory factor 5 (*IRF5*) variant with increased risk of systemic lupus erythematosus (SLE). *The American Society of Human Genetics Annual Meeting*, October 9-13, 2006. New Orleans, LA
193. Dasgupta S, Demirci FY, Kao A, Bontempo F, Kammerer C, Manzi S, **Kamboh MI**: *PON2* Polymorphisms, PON Activity, and Systemic Lupus Erythematosus (SLE). *The American Society of Human Genetics Annual Meeting*, October 23-27, 2007. San Diego, CA.
194. Minster RL, DeKosky ST, **Kamboh MI**: No association of SORL1 SNPs with Alzheimer's diseases. *The American Society of Human Genetics Annual Meeting*, October 23-27, 2007. San Diego, CA.
195. Figgins JA, Minter RL, DeKosky ST, **Kamboh MI**: No association of selected SNPs in GALP, PCK1, SERPINA13 or TNK1 with Alzheimer's disease. *The American Society of Human Genetics Annual Meeting*, October 23-27, 2007. San Diego, CA.
196. Suresh S, Jacobs E, Manzi S, Sanghera DK, Kao A, Bontempo F, Kammerer C, Demirci FY, **Kamboh MI**: Association of *APOH* promoter polymorphisms with lupus nephritis and cardiovascular disease. *The American Society of Human Genetics Annual Meeting*, October 23-27, 2007. San Diego, CA.
197. Kodavali V, Sweet RA, **Kamboh MI**, Nimgaonkar NL: OSTM1 and NR2E1: Positional candidate genes for psychosis in Alzheimer's and Schizophrenia. *The American Society of Human Genetics Annual Meeting*, October 23-27, 2007. San Diego, CA.
198. Demirci FY, Manzi S, Ramsey-Goldman R, Kao AH, Rhew EY, Bontempo F, Kammerer C, **Kamboh MI**: *TNFSF13(APRIL)* Polymorphisms and systemic lupus erythematosus. *The American Society of Human Genetics Annual Meeting*, October 23-27, 2007. San Diego, CA.
199. Corthell LS, Demirci FY, Kao AH, Rhew EH, Bontempo F, Kammerer C, Ramsey-Goldman R, Manzi S, **Kamboh MI**: Relationship of apolipoprotein H (*APOH*) polymorphisms with susceptibility to

systemic lupus erythematosus. *The American Society of Human Genetics Annual Meeting*, October 23-27, 2007. San Diego, CA.

200. Muscone SL, Taylor KE, Lu T, Nititham J, Ferriera RC, Ortmann W, Shiffrin N, Petri MA, **Kamboh MI**, Manzi S, Seldin MF, Gregersen PK, Behrens TW, Ma A, Kwok P-Y, Criswell LA: Multiple polymorphisms in the TNFAIP3 region are independently associated with systemic lupus erythematosus. *American College of Rheumatology Annual Meeting*, 2008.
201. Suresh S, Demirci FY, Kao AH, Kammerer CM, Manzi S, **Kamboh MI**: Functional characterization of *APOH* promoter and its variants. *The America Society of Human Genetics Annual Meeting* 2008.
202. Dasgupta S, Demirci FY, Dressen A, Kao AH, Kammerer C, Manzi S, Kamboh **MI**: Relation of paraoxonase (*PON2*) polymorphisms with PON activity and systemic lupus erythematosus (SLE). *The America Society of Human Genetics Annual Meeting* 2008.
203. **Kamboh MI**, Hollister SM, Demirci FY, Dressen AS, Bunker CH, Hamman RF, Kammerer CM: *APOA2* genetic variation and its relationship with plasma HDL-cholesterol levels. *The America Society of Human Genetics Annual Meeting* 2008.
204. Waqar F, Demirci FY, Dressen AS, Bunker CH, Hamman RF, Kammerer C M, **Kamboh MI**: *APOM* sequence analysis in relation to plasma HDL-C levels. *The America Society of Human Genetics Annual Meeting* 2008.
205. Demirci FY, Dressen AS, Hamman RF, Kammerer C M, **Kamboh MI**: Association of a common *G6PC2* variant with fasting plasma glucose levels. *The America Society of Human Genetics Annual Meeting* 2008.
206. Figgins J, Minster RL, Demirci FY, DeKosky ST, **Kamboh MI**: Association studies of 22 candidate SNPs with late-onset Alzheimer's disease. *The America Society of Human Genetics Annual Meeting* 2008.
207. Sanghera DK, Demirci FY, Ortega L, Been L, Ralhan SK, Wander GS, Mehra NK, Singh JR, Mulvihill JJ, **Kamboh MI**: Novel polymorphisms in PPARG and ADIPOQ genes increase type 2 diabetes risk in Asian Indian Sikhs: Evidence of gene-gene interaction. *The America Society of Human Genetics Annual Meeting* 2008.
208. Demirci FY, Wang Y, Kao AH, Rhew EY, Ramsey-Goldman R, Manzi S, **Kamboh MI**: Comprehensive resequencing of F2 gene in systemic lupus erythematosus. *The American Society of Human Genetics 2009 Annual Meeting*, October 20-24, 2009. Honolulu, Hawaii.
209. Hill SE, Demirci FY, Dressen AS, Kammerer CM, Bunker CH, Hokanson JE, Hamman RF, **Kamboh MI**: Sequence variation in the *APOA1* and *APOA4* genes and their relationship with plasma HDL-cholesterol levels. *The American Society of Human Genetics 2009 Annual Meeting*, October 20-24, 2009. Honolulu, Hawaii.
210. Burns LC, Minster RL, Demirci FY, DeKosky ST, **Kamboh MI**: Association studies of 12 candidate SNPs with late-onset Alzheimer's disease. *The American Society of Human Genetics 2009 Annual Meeting*, October 20-24, 2009. Honolulu, Hawaii.
211. Sanghera DK, Begum MF, Mukerjee A, Ralhan S, Wander GS, Mehra NK, Singh JR, Ferrell RE, **Kamboh MI**, Weeks DE: A genome-wide linkage scan for blood lipid phenotypes in the Khatri Sikh Diabetes Study (SDS). *The American Society of Human Genetics 2009 Annual Meeting*, October 20-

24, 2009. Honolulu, Hawaii.

212. Bryant EK, Dressen AS, Hokanson JE, Hamman RF, Kammerer CM, **Kamboh MI**, Demirci FY: Replication study of plasma lipoprotein levels: Associated SNPs identified in recent GWAS. *The American Society of Human Genetics 2010 Annual Meeting*, November 2-6, 2010. Washington, DC.
213. Hughes SC, Demirci FY, Dressen AS, Bunker CH, Hokanson JE, Hamman RF, Kammerer CM, **Kamboh MI**: Comprehensive resequencing of the CD36 gene in subjects with extremely low or high plasma HDL-C levels. *The American Society of Human Genetics 2010 Annual Meeting*, November 2-6, 2010. Washington, DC.
214. Miller MA, Conley YP, Scanlon JM, Ren D, Kamboh MI, Niyonkuru C, Wagner AK. APOE genetic associations with seizure development after severe traumatic brain injury. Presented at the *National Neurotrauma Symposium*, June 2010. Las Vegas, NV
215. Barmada MM, Demirci FY, Minster RL, Carrasquillo MM, Pankratz VS, Younkin SG, Saykin AJ, Sweet RA, Feingold E, DeKosky ST, Lopez OL, **Kamboh MI**: Genome-wide association analysis of age-at-onset in Alzheimer's disease. *The International Congress of Human Genetics meeting*, October 11-14, 2011. Montreal, Canada.
216. Radwan ZH, Demirci FY, Hokanson JE, Hamman RF, Bunker CH, Kammerer CM, **Kamboh MI**: Effects of common and rare genetic variants of APOC4 on HDL-cholesterol levels. *The International Congress of Human Genetics meeting*, October 11-14, 2011. Montreal, Canada.
217. Prim D, Demirci FY, Hughes SC, Irfan M, Wang Y, Hokanson JE, Hamman RF, Kammerer CM, **Kamboh MI**: Lipoprotein lipase gene (*LPL*) resequencing and plasma lipid profile. *The International Congress of Human Genetics meeting*, October 11-14, 2011. Montreal, Canada.
218. Chung SA, Brown EE, Williams AH, Bhangale T, Ramos P, Ziegler J, Freedman BI, Kimberly RP, Vyse TJ, Gregersen PK, Jacob CO, Alarcon-Riquelme M, Tsao BP, Harley JB, Behrens TW, Petri M, Demirci FY, **Kamboh MI**, Manzi S, Criswell LA, SLEGEN, Moser KL, Gaffney PM, Graham RR, Langefeld C.: Lupus nephritis susceptibility markers in *PDGRFA-GSX2*, *SLC5A11*, *ID4*, and *HAS2-SNTB1* regions identified from a meta-analysis of genome wide association studies of women with systemic lupus erythematosus. *Annual American College of Rheumatology meeting*, November 4-9, 2011, Chicago.
219. Hollingworth P, Sweet R, Sims R, Harold D, Gerrish M, Devlin B, Klei L, Barmada MM, Demirci F, FF DeKosky ST, Lopez OL, Passmore P, Mayeux R, Kamboh MI, Williams J. Genome wide association study of Alzheimer's disease with psychotic Symptoms. Abstract was presented at the Alzheimer's Association International Conference 2011 (AAIC 2011), Paris, France, July 16-21, 2011.
220. Pirim D, Demirci FY, Wang X, Irfan M, Wang Y, Hokanson JE, R. F. Hamman RF, Kammerer CM, **Kamboh MI**. Resequencing of the lipoprotein lipase (*LPL*) gene in individuals with extreme HDL-cholesterol Levels. *Arteriosclerosis, Thrombosis and Vascular Biology (ATVB) 2012 Scientific Sessions*, April 18-20, 2012. Chicago, Illinois.
221. Hughes TM, Lopez OL, **Kamboh MI**, Klunk WE, Kuller LW: Markers of cholesterol homeostasis are associated with amyloid deposition in the brain. *Gerontological Society of America*, 2012.
222. Pirim D, Demirci FY, Wang X, Irfan M, Wang Y, Hokanson JE, Hamman RF, Kammerer CM, **Kamboh MI**. Resequencing of the lipoprotein lipase gene in individuals with extreme HDL-cholesterol levels. *Arteriosclerosis, Thrombosis, and Vascular Biology* 2012; 32(supplement):A324

223. Demirci FY, Wang X, Kao AH, Clarke A, Ramsey-Goldman R, Manzi S, Barmada MM, **Kamboh MI**. A genome-wide association study of systemic lupus erythematosus in North Americans of European ancestry. *The American Society of Human Genetics 2012 Annual Meeting*, November 6-10, San Francisco.
224. Pirim D, Demirci FY, Wang X, Hokanson JE, Hamman RF, Bunker CH, Kammerer CM, Barmada MM, **Kamboh MI**. Resequencing of the cholesteryl ester transfer protein gene (*CETP*) in U.S. Whites and African Blacks with extreme HDL-C levels. *The American Society of Human Genetics 2012 Annual Meeting*, November 6-10, San Francisco.
225. Radwan ZH, Demirci FY, Wang X, Waqar F, Hokanson JE, Hamman RF, Bunker CH, Kammerer CM, Barmada MM, **Kamboh MI**. Comprehensive evaluation of the effects of *APOE* genetic variation on plasma lipoprotein traits in U.S. Whites and African Blacks. *The American Society of Human Genetics 2012 Annual Meeting*, November 6-10, San Francisco.
226. Wang X, Reis SE, Barmada MM, Halder I, Demirci FY, **Kamboh MI**. Genetic association of CVD-related loci with lipid traits in the Heart Strategies Concentrating on Risk Evaluation (Heart SCORE) study. *The American Society of Human Genetics 2012 Annual Meeting*, November 6-10, San Francisco.
227. Jalil SF, Bhatti A, Demirci FY, Wang X, Ahmed I, Ahmed M, Barmada MM, Malik JM, John P, **Kamboh MI**. A replication study of genome-wide significant rheumatoid arthritis susceptibility loci in the Pakistani population. *The American Society of Human Genetics 2012 Annual Meeting*, November 6-10, San Francisco.
228. McClain L, Wang X, Barmada MM, Demirci FY, Lopez OL, **Kamboh MI**. Investigation of lipid pathway genes in late onset Alzheimer's Disease. *The American Society of Human Genetics 2012 Annual Meeting*, November 6-10, San Francisco.
229. Rosenthal SL, Wang X, Barmada MM, Demirci FY, Lopez OL, **Kamboh MI**. Beta-amyloid toxicity modifier genes and the risk of Alzheimer's disease. *The American Society of Human Genetics 2012 Annual Meeting*, November 6-10, San Francisco.
230. Reitz C, Jun G, Buros J, Vardarajan B, Wang L-S, Busbaum JD, Larson EB, Graff-Radford N, Evans D, Ertekin-Tanner, N, Logue M, Baldwin TC, Green RC, Barnes LL, Cantwell LB, Fallin MD, Manly J, Lunetta KL, **Kamboh MI**, Bennet DA, Hall K, Goate AM, Byrd GS, Kukul; WA, Foroud TM, Haines JL, Pericak-Vance MA, Farrer L, Schellenberg G, Mayeux, ADGC Consortium: Common variants in ABCA7 and GRIN3B, HMHA1 and SBNO2, are associated with late-onset Alzheimer's disease in African Americans. *The American Society of Human Genetics 2012 Annual Meeting*, November 6-10, San Francisco.
231. Rosenthal, SL, Barmada MM, Wang X, Demirci FY, Lopez OL, **Kamboh MI**. An examination of regulatory function for GWAS confirmed and suggestive loci of late-onset Alzheimer's disease. *Alzheimer's Association International Conference (AAIC)*, July 13-18, 2013, Boston, MA.
232. Wang X, Lopez, O, Sweet R, Barmada, MM, Demirci FY, Kamboh MI. Genetic Determinants of Progression of Alzheimer's Disease. *Alzheimer's Association International Conference (AAIC)*, July 13-18, 2013, Boston, MA.
233. Wang M-S, Fiocchi C, Zhu X, Ripke S, **Kamboh MI**, Robert N, Duerr RH, Achkar J-P. Novel gene-gene and gene-environment interactions in ulcerative colitis. *American College of Gastroenterology Annual Meeting*, October 2013, San Diego.
234. Pirim D, Demirci FY, Wang X, Hokanson JE, Hamman RF, Bunker CH, Barmada MM, **Kamboh MI**.

Identification of sequence variants of hepatic lipase (LIPC) gene in individuals with extreme HDL-C/TG levels. *The American Society of Human Genetics 2013 Annual Meeting*, October 22-26, Boston.

235. Wang X, Lopez OL, Sweet RA, Becker J, Barmada MM, Demirci FY, **Kamboh MI**. Genetic determinants of the natural history of Alzheimer's disease. *The American Society of Human Genetics 2013 Annual Meeting*, October 22-26, Boston.
236. Rosenthal SL, Barmada MM, Wang X, Demirci FY, Lopez OL, **Kamboh MI**. Potential regulatory functions of late-onset Alzheimer's disease associated variants. *The American Society of Human Genetics 2013 Annual Meeting*, October 22-26, Boston.
237. Radwan ZH, Wang X, Waqar F, Hokanson JE, Hamman RF, Barmada MM, Demirci FY, **Kamboh MI**. A comprehensive association study of *APOE-C1-C4-C2* gene cluster variation with plasma lipoprotein traits in U.S. Whites. *The American Society of Human Genetics 2013 Annual Meeting*, October 22-26, Boston.
238. Niemsiri V, Wang X, Barmada MM, Hokanson JE, Hamman RF, Demirci FY, Kamboh MI. Genetic Influence of Scavenger Receptor Class B Type 1 (SCARB1) on Plasma Lipid Traits in non-Hispanic White Americans. *The American Society of Human Genetics 2013 Annual Meeting*, October 22-26, Boston.
239. Chaudhry M, Hasnain S, Snitz B, Wang X, Winger D, Wang L, Rosenthal S, Demirci FY, **Kamboh MI**. Association of *APOE* Polymorphism and Stressful Life Events with Dementia in the Pakistani Population. *The American Society of Human Genetics 2013 Annual Meeting*, October 22-26, Boston.
240. Pirim D, Wang X, Hokanson JE, Hamman RF, Barmada MM, Demirci FY, Kamboh MI. *CETP* gene variation and its association with plasma lipoprotein-lipid traits in U.S. Whites. *The XIII National Congress of Medical Biology and Genetics*, Kusadasi, Turkey, October, 2013
241. Minhas DS, Gregg NM, Kim AE, Gurol ME, Lopez OL, Aizenstein HJ, Price JC, Mathis CA, James JA, Snitz BE, Cohen AD, **Kamboh MI**, Weissfeld LA, Tamburo EL, Klunk WE. Cerebral microbleeds are associated with cerebral blood flow and metabolism but not amyloid burden or brain atrophy in cognitively normal elderly. *Alzheimer's Association International Conference (AAIC)*, *Alzheimer's Imaging Consortium (AIC)*, July 12-17, 2014, Copenhagen, Denmark
242. French S, Taylor KE, Chung SA, Nititham J, Petri M, Gregersen PK, Orthmann W, Lee A, Behrens TW, Manxi S, Demirci FY, **Kamboh MI**, Graham R, Seldin MF. The impact of northern European ancestry and susceptibility loci on the risk of lupus nephritis. *The American College of Rheumatology (ACR) and the Association for Rheumatology Health Professionals (ARHP) meeting*, November 2014, Boston
243. Ombrello MJ, Remmers EF, Tachmazidou I, Grom A, Föll D, Martini A, Gattorno M, Ozen S, Prahalad S, Zeft A, Bohnsack J, Ilowite N, Park J, Mellins ED, Russo R, Len C, Oliveira S, Yeung R, Wedderburn LR, Anton J, Schwarz T, Han B, Duerr R, Achkar JP, **Kamboh MI**, Kaufman K, Kotyan K, Pinto D, Scherer S, Alarcon-Riquelme M, Martinez ED, Estivill X, Güll A, Satorius C, de Bakker P, Raychaudhuri S, Langefeld C, Thompson S, Zeggini E, Thomson W, Kastner D, Woo P, on behalf of the International Childhood Arthritis Genetics (INCHARGE) Consortium. *HLA-DRB1*1101*, regulatory variants of the MHC, and a regulatory region near an intergenic long noncoding RNA on chromosome 1 are risk factors for systemic juvenile idiopathic arthritis. *The American College of Rheumatology (ACR) and the Association for Rheumatology Health Professionals (ARHP) meeting*, November 2014, Boston
244. Demirci FY, Niemsiri V, Wang X, Barmada MM, Hokanson JE, Hamman RF, **Kamboh MI**.

- Contribution of genetic variation of ATP-binding cassette transporter A1 (*ABCA1*) to the regulation of plasma lipid/lipoprotein levels in US Non-Hispanic Whites. *American Society of Human Genetics 2014 Annual Meeting*, October 18-22, 2014. San Diego, CA.
245. Niemsiri V, Wang X, Barmada MM, Bunker CH, Demirci FY, **Kamboh MI**. Genetic variation of Scavenger Receptor Class B Type I (SCARB1) and Plasma Lipid Traits: An Association Study in a Nigerian Population. *American Society of Human Genetics 2014 Annual Meeting*, October 18-22, 2014. San Diego, CA.
246. Wang X, Lopez OL, Sweet RA, Becker JT, Barmada MM, Feingold E, Demirci FY, **Kamboh MI**. Genetic determinants of survival in patients with Alzheimer's disease. *American Society of Human Genetics 2014 Annual Meeting*, October 18-22, 2014. San Diego, CA.
247. Rosenthal SL, Bamne MN, Wang X, Berman S, Snitz BE, Klunk WE, Sweet RA, Demirci FY, Lopez OL, **Kamboh MI**. More evidence for association of a rare TREM2 variant (R47H) with Alzheimer's disease risk. *American Society of Human Genetics 2014 Annual Meeting*, October 18-22, 2014. San Diego, CA.
248. Cheema AN, Bhatti A, Turi J, Wang X, Demirci F, **Kamboh MI**. Association analysis of putative loci with coronary artery disease. *International Conference on Cardiology and Cardiac Surgery*, Dubai, UAE, May 2015.
249. Niemsiri V, Wang X, Radwan ZH, Pirim D, Hokanson JE, Hamman RF, Barmada MM, Demirci FY, **Kamboh MI**. The contribution of common and low-frequency/rare variants in ATP-binding cassette A1 (*ABCA1*) to lipoprotein-lipid traits. *American Society of Human Genetics 2015 Annual Meeting*, October 6-10, 2015. Baltimore, MD.
250. Wang X, Reis SE, Halder I, Barmada MM, Demirci FY, **Kamboh MI**. Comprehensive gene-centric association analysis for lipid traits in the Heart SCORE (Heart Strategies Concentrating on Risk Evaluation) study using the HumanCVD BeadChip. *American Society of Human Genetics 2015 Annual Meeting*, October 6-10, 2015. Baltimore, MD.
251. Bamne MN, McDade E, Wang X, Demirci FY, Feingold E, Klunk WE, **Kamboh MI**. Copy number variation in early-onset Alzheimer's disease. *American Society of Human Genetics 2015 Annual Meeting*, October 6-10, 2015. Baltimore, MD.
252. Lopez OL, Kofler J, Steinberg A, Becker JT, Sweet R, Berman S, McDade E, Rodriguez E, **Kamboh MI**, Klunk WE. Relationship between Hippocampal sclerosis and TDP-43, and duration of the symptoms of dementia in Alzheimer's disease patients. *American Academy of Neurology Annual Meeting*, Vancouver, BC, Canada, April 15-21, 2016.
253. **Kamboh MI**, Wang X, Feingold E, Barmada MM, Bamne MN, Demirci FY, Klunk WE, Ganguli M, Lopez OL. Gender and the genetics of Alzheimer's disease. *Alzheimer's Association International Conference (AAIC) Meeting*, Toronto, Canada, July 24-28, 2016.
254. Prim D, Radwan Z, Wang X, Waqar F, Niemsiri V, Hokanson JE, Hamman RF, Feingold E, Bunker CH, Barmada MM, Demirci FY, **Kamboh MI**. A Comprehensive association study of apolipoprotein E-C1-C4-C2 gene cluster variation with plasma lipoprotein traits. *European Atherosclerosis Society*, Parague, Czech Republic, April 23-26, 2017.
255. Rosenthal S, Demirci FY, Yan Q, Bamne M, Lopez OL, Feingold E, **Kamboh MI**. Targeted sequencing

of GWAS-implicated loci in Alzheimer's disease. *Alzheimer's Association International Conference*, July 16-20, 2017. London, England.

256. Demirci FY, Yan Q, Nho K, Del-Aguila J, Wang X, Risacher S, Snitz B, Aizenstein H, Mathis C, Lopez O, Feingold E, Klunk W, Saykin AJ, Cruchaga C, **Kamboh MI**. Genome-wide association study of brain amyloid deposition as measured by PiB-PET imaging and the assessment of genetic variance of amyloid deposition. *American Society of Human Genetics 2017 Annual Meeting*, October 17-21, 2017. Orlando, FL.
257. Niemsiri V, Wang X, Radwan ZH, Pirim D, Halder I, Reis SE, Hokanson JE, Hamman RF, Barmada MM, Feingold E, Demirci FY, **Kamboh MI**. Contribution of *ABCA1* Genetic Variation to Plasma Lipid Levels in Non-Hispanic White Americans. *American Heart Association Scientific Sessions 2017 Annual Meeting*, November 11-15, 2017. Anaheim, CA.
258. Cheema A, Rosenthal S, **Kamboh MI**. Proficiency of data interpretation: identification of signaling single nucleotides polymorphism for coronary artery disease. *12th Asia-Pacific Conference on Human Genetics*, November 8-10, 2017. Bangkok, Thailand. *Ann Transl Med* 2017;5(Suppl 2): AB069.
259. Cheema A, Pirim D, Wang X, Demirci FY, **Kamboh MI**. Association of genome-wide significant single-nucleotide polymorphisms with coronary artery disease in Pakistani population: a case-control study. *12th Asia-Pacific Conference on Human Genetics*, November 8-10, 2017. Bangkok, Thailand. *Ann Transl Med* 2017;5(Suppl 2): AB070.
260. Rajakumar K, Yan Q, Khalid A, Lewis D, Moore CG, Feingold E, Demirci FY, **Kamboh MI**. Gene expression profiles as related to body mass index and cardiometabolic phenotypes in vitamin D-deficient black children. *Pediatric Academic Societies (PAS) 2018 Meeting*, May 5-8, Toronto, Canada
261. **Kamboh MI**, Yan Q, Beer JC, Fan K, Wang X, Demirci FY, Snitz BE, Chang CH, Feingold E, Ganguli M. Population-based genome-wide association study of cognitive decline in older adults free of dementia. *American Society of Human Genetics 2018 Annual Meeting*, October 16-20, 2018. San Diego, CA.
262. Fan K-S, Rosenthal S, Feingold E, Demirci FY, Lopez O, **Kamboh MI**. Whole-exome sequencing (WES) analysis of Alzheimer's disease in non-*APOE*4* carriers. *American Society of Human Genetics 2018 Annual Meeting*, October 16-20, 2018. San Diego, CA.
263. Aslam MM, John P, Fan K, Bhatti A, Feingold E, Demirci FY, **Kamboh MI**. Investigating the GWAS-implicated loci for Rheumatoid Arthritis in Pakistani population. *American Society of Human Genetics 2018 Annual Meeting*, October 16-20, 2018. San Diego, CA.
264. Snitz BE, Teodorescu DL, Lopez OL, Kuller LH, Cohen AD, DeKosky ST, **Kamboh MI**, Lopresti B, Mathis C, Klunk WE. Predicting resiliency against amyloid-beta deposition, cognitive impairment, and their combination in the oldest-old. *Alzheimer's Association International Conference*, July 22-26, 2018. Chicago, IL.
265. Kofler J, Fan K, Yan Q, Sweet RA, Feingold E, Lopez O, **Kamboh MI**. Association of Alzheimer's disease genetic risk variants with pathology endophenotypes. *Alzheimer's Association International Conference*, July 22-26, 2018. Chicago, IL.
266. Aslam MM, John P, Fan, K-H, Bhatti A, Feingold E, Demirci FY, **Kamboh MI**. Exploration of shared genetic susceptibility loci between type 1 diabetes and rheumatoid arthritis in the Pakistani population.

American Society of Human Genetics 2019 Annual Meeting, October 15-19, 2019. Houston TX.

267. Aslam MM, Jalil F, John P, Fan KH, Bhatti A, Feingold E, Demirci FY, **Kamboh MI**. A sequencing study of CTLA4 in Pakistani rheumatoid arthritis cases. *Annual American Society of Human Genetics Meeting*, October 27-30, 2020, Virtual Meeting.
268. Acharya V, Fan KH, Feingold E, Kofler J, **Kamboh MI**. Somatic mutation analysis reveals higher mutational load in APOE and ABCA7 loci in Alzheimer's disease. *Annual American Society of Human Genetics Meeting*, October 27-30, 2020, Virtual Meeting.
269. Fan KH, Aslam MM, Feingold E, Rosenthal SL, Demirci FY, Gangui M, Lopez OL, **Kamboh MI**. Targeted sequencing of Alzheimer's disease-associated loci. *Annual American Society of Human Genetics Meeting*, October 27-30, 2020, Virtual Meeting.
270. Acharya V, Fan K-H, Snitz BE, Ganguli M, DeKosky ST, Lopez OL, Feingold E, **Kamboh MI**. Genome-wide meta-analysis of age-related cognitive decline in population based older individuals. *Alzheimer's Association International Conference (AAIC) Neuroscience Next 2021*, October 12-13, 2021. *Alzheimers Dement* 2021; 17:e058723.
271. Ngo M, Donnelly CJ, Fan K-H, **Kamboh MI**, Kofler J. Role of GWAS candidate gene, SCYL3, in TDP-43 expression and aggregation. Virtual meeting about age-related TDP-43 proteinopathy, February 11, 2022.
272. Fongang B, Weinstein G, Guðjónsson A, Mishra A, Bis JC, Yang Q, Winsvold B, Muralidharan S, Fan KH, **Kamboh MI**, Li G, Yang J, Hilal S, Satizabal CL, Jian X, Knol MJ, Concas MP, Girotto G, Riaz M, Lacaze P, Ruiz, Naj AC, Schellenberg GD, Kehoe PG, Lee SJ, Skrobot OA, Gudnason V, Lopez OL, Haan M, Bosnes I, Dufouil C, Ganguli M, Cheung CL, Bennett DA, Lambert JC, Chen C, Ikram MA, Debette S, Fornage M, Seshadri S, Cohorts for Heart and Aging Research in Genomic Epidemiology (CHARGE). A meta-analysis of genome-wide association studies identifies new genetic loci associated with all-cause and vascular dementia. *Alzheimers Dement* 2021;17:e056081.
273. Harper JD, Fan KH, Aslam MM, Snitz BE, DeKosky ST, Lopez OL, Feingold E, **Kamboh MI**. Genome-wide association study of incident dementia in a community-based sample of older subjects. *Alzheimer's Association International Conference (AAIC)*, July 31-August 4, 2022
274. Shi R, Fan KH, Lopez OL, Feingold E, **Kamboh MI** A whole-genome-sequencing study to identify genetic variants associated with Alzheimer's disease progression. *Alzheimer's Association International Conference (AAIC)*, July 31-August 4, 2022
275. Fan KH, Aslam MM, Francis L, Bedison A, Lawrence E, Acharya V, Snitz BE, Ganguli M, DeKosky ST, Lopez OL, Feingold E, **Kamboh MI**. Re-evaluation of the genetic role of a rare APOE variant (L28P; *APOE*4Pittsburgh*) in late-onset Alzheimer disease. *Alzheimer's Association International Conference (AAIC)*, July 31-August 4, 2022.
276. Acharya V, Fan KH, Snitz BE, Ganguli M, DeKosky ST, Lopez OL, Feingold E, **Kamboh MI**. Meta-analysis of age-related cognitive decline reveals a novel locus for the attention domain and implicates a COVID-19 related gene for global cognitive. *Alzheimer's Association International Conference (AAIC)*, July 31-August 4, 2022.
277. Aslam MM, Fan KH, Lawrence E, Bedison A, Snitz BE, DeKosky ST, Lopez OL, Feingold E, **Kamboh**

MI. Genome-wide association study of plasma apolipoprotein E levels and risk of dementia in older adults. *Alzheimer's Association International Conference (AAIC)*, July 31-August 4, 2022.

278. Han X, Moore MR, Chung J, Durape SS, Rosenthaler M, Uretsky M, Abdolmohammadi B, Farrell J, Zhang X, Jun G, Qiu W, Logue M, Stein T, Bennett DA, Paul K, Crane PK, **Kamboh MI**, Kukull WA, Larson EB, Au R, Haines JL, Pericak-Vance MA, Schellenberg GD, Mayeux R, Lunetta KL, Lindsay A, Farrer LA, Mez J. Genome-wide interaction study with Smoking identifies *FHIT* and *SLC22A23* associated with Alzheimer's disease. *Annual American Society of Human Genetics Meeting*, October 25-29, 2022.

279. Acharya V, Fan KH, Snitz BE, Ganguli M, DeKosky ST, Lopez OL, Feingold E, **Kamboh MI**. Sex-stratified meta-analysis of age-related cognitive decline across neurocognitive domains. *Annual American Society of Human Genetics Meeting*, October 25-29, 2022.

280. Aslam MM, Fan KH, Snitz BE, DeKosky ST, Lopez OL, Feingold E, **Kamboh MI**. Genome-wide association study of plasma amyloid β levels in older adults. *Annual American Society of Human Genetics Meeting*, October 25-29, 2022.

SERVICE

University and Professionally Related

a. Department, School or University Committees

Position	Committee	Specify Department, School or University	Year(s)
Member	Faculty advancement, Promotion and Tenure Committee(FAPTC)	Graduate School of Public Health, University of Pittsburgh	2015-2018
Juror	Dean's Day	Graduate School of Public Health, University of Pittsburgh	2015, 2016
Juror	Dean's Day	Graduate School of Public Health, University of Pittsburgh	2014
Member	Graduate School of Public Health Council	Graduate School of Public Health, University of Pittsburgh	2005- 2013
Member	The Jay L. Foster Memorial Lecture Series	Graduate School of Public Health, University of Pittsburgh	2002-2007, 2009-2013
Member	Executive Committee, Alzheimer's Disease Research Center	University of Pittsburgh	2000-Present
Member	Institutional Review Board (IRB)	University of Pittsburgh	1998-2001
Member	M.D./Ph.D. Selection Committee	School of Medicine, University of Pittsburgh	1997-2000

Member	Planning and Budgeting Committee	Graduate School of Public Health, University of Pittsburgh	1993-1995
Member	Faculty Advancement Committee	Graduate School of Public Health, University of Pittsburgh	1993-1998
Member	Advisory Committee on the GSPH Multidisciplinary M.P.H. degree	Graduate School of Public Health, University of Pittsburgh	1991-1996
Member	Safety Committee	Graduate School of Public Health, University of Pittsburgh	1992-1994
Member	Appointment/Promotion ad-hoc Committee	School of Medicine, University of Pittsburgh	1994-1999
Member	Appointment/Promotion ad-hoc Committee	Graduate School of Public Health, University of Pittsburgh	1994-1999
Member	Ad hoc Search and Appointment Committee	Department of Human Genetics, Graduate School of Public Health, University of Pittsburgh	1991-Present
Member	Curriculum Revision Committee	Department of Human Genetics, Graduate School of Public Health, University of Pittsburgh	1991
Member	Graduate Faculty	University of Pittsburgh	1987-Present

b. Editorial Boards

Type of Activity	Position	Title	Date
Editorial Board	Associate Editor	<i>Journal of Alzheimer's Disease</i>	2015-2017, 2021
Editorial Board	Founding Editor-in-Chief	<i>Current Genetic Medicine Reports</i>	2013-2018
Editorial Board	Editor-in-Chief	<i>The Open Genomics Journal</i>	2013
Editorial Board	Senior Member	<i>American Journal of Neurodegenerative Disease</i>	2012-Present
Editorial Board	Section Editor	<i>BioMed Research International</i> (formerly <i>Journal of Biomedicine and Biotechnology</i>)	2009-2015
Editorial Board	Member	<i>The Open Translational Medicine Journal</i>	2008-Present

Editorial Board	Member	<i>The Open Geriatric Medicine Journal</i>	2008-Present
Editorial Board	Member	<i>The Open Genomics Journal</i>	2007-2012
Editorial Board	Member	<i>The Open Rheumatology Journal</i>	2007-Present
International Advisory Board	Member	<i>Journal of the Dow University of Health Sciences</i>	2007-Present
Editorial Board	Associate Editor	<i>Annals of Human Genetics</i>	2003-2013
Editorial Board	Member	<i>Human Biology</i>	1994-2008
Editorial Board	Member	<i>Ethnicity and Disease</i>	1991-1993

c. Manuscript/Publication Reviewer

Title	Title
<i>Acta Neuropathologia</i>	<i>European Journal of Human Genetics</i>
<i>American Journal of Human Genetics</i>	<i>Experimental Gerontology</i>
<i>American Journal of Epidemiology</i>	<i>Genetic Epidemiology</i>
<i>American Journal of Medical Genetics</i>	<i>Human Biology</i>
<i>American Journal of Physical Anthropology</i>	<i>Human Genetics</i>
<i>Annals of Human Genetics</i>	<i>Human Heredity</i>
<i>Annals of Neurology</i>	<i>Human Immunology</i>
<i>Arteriosclerosis, Thrombosis and Vascular Biology</i>	<i>Human Mutation</i>
<i>Atherosclerosis</i>	<i>Experimental Biology & Medicine</i>
<i>Arthritis & Rheumatism</i>	<i>Journal of Alzheimer's Disease</i>
<i>Biochemical Genetics</i>	<i>Journal of Lipid Research</i>
<i>Biochimica Et Biophysica Acta</i>	<i>Journal of Medical Genetics</i>
<i>BMC Genetics</i>	<i>Journal of Neurology, Neurosurgery & Psychiatry</i>
<i>Biological Psychiatry</i>	
<i>Brain Research</i>	<i>Journal of Neurological Sciences</i>
<i>Circulation</i>	<i>Lancet</i>
<i>Clinical Chemistry</i>	<i>Lancet Neurology</i>
<i>Clinical Chemistry and Laboratory Medicine</i>	<i>Life Sciences</i>

<i>Clinical Genetics</i>	<i>Molecular Psychiatry</i>
<i>Clinical & Experimental Medicine</i>	<i>Mutation Research</i>
<i>Clinical Lipidology</i>	<i>Neurobiology of Aging</i>
<i>Diabetes</i>	<i>Neurogenetics</i>
<i>Diabetologia</i>	<i>Neurology</i>
<i>Disease Markers</i>	<i>Neuroscience Letters</i>
	<i>Neurotherapeutics</i>
<i>Electrophoresis</i>	<i>Pharmacogenetics</i>
<i>European Journal of Clinical Investigation</i>	<i>PLoS Genetics</i>
<i>Future Neurology</i>	<i>Psychiatry Research</i>
	<i>Scientific Reports</i>

d. Study Sections/Review Panels and Related Advisory Boards

Type of Service	Position	Agency	Year(s)
Research Grant Proposal	Committee Chair	NIH –Special Emphasis Panel –ZRG1 GGG-S(02) Genetics, Genomics, and Therapeutics of Diseases	2019
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH –Special Emphasis Panel – Interdisciplinary Research to Understand the Complex Biology of Resilience to Alzheimer’s Disease Risk	2018
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH –Special Emphasis Panel – Interdisciplinary Research to Understand the Complex Biology of Resilience to Alzheimer’s Disease Risk	2017
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH –Special Emphasis Panel –Major Opportunities for Research in Epidemiology of Alzheimer’s Disease and Cognitive Resilience	2016
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH – National Institute on Aging Special Emphasis Panel - Inflammation and Alzheimer’s Disease	2015

Research Grant Proposal	<i>Ad-hoc</i> Member	NIH – ZRG1 GGGE (02) Member Conflict: Genetic Variants in Disease	2015
Research Grant Proposal	Committee Chair	NIH – National Heart, Lung, and Blood Institute: Genetic Basis of Monogenic Blood Disorder ZHL1 CSR-C (F1)	2013
Research Grant Proposal	Committee Chair	NIH – ZRG1 PSE-P(02) Member Conflict: Chronic Disease, Aging and Genetics Review Committee	2013
7-Year Review of the Human and Molecular Genetics Ph.D. Program	External Reviewer	University of Texas Graduate School of Biomedical Sciences at Houston	2013
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH – ZRG1 PSE-D (03) NAME conflict Study Section	2012
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH – ZRG1 BBBP-E(53)/NIH Director's Early Independence Award Review	2012
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH – ZRG1 EMNR-Q 50/Specialized Centers of Research (SCOR) on Sex differences	2012
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH – ZRG1 GGG0M 50/Epigenomics of Human Health and Disease	2011
Medical Advisory Committee	Member	Brain Injury Research Institute (BIRI), West Virginia University, Morgantown, WV	2010
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH – Genetics of Health and Disease (GHD) Study Section	2009
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH Special Emphasis Panel ZRG1 HOP-D(05) M, /CASE Member SEP	2008
Research Grant Proposal	Chartered Member	National Institutes of Health (NIH)/Cardiovascular and Sleep Epidemiology Study Section (formerly ECD and EDC1 Study Sections)	2003-2007
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH Special Emphasis Panel ZRG1 HOP-S 02 M, KNOD/CASE Member SEP	2007

Research Grant Proposal	<i>Ad-hoc</i> Member	NIH Special Emphasis Panel ZRG1 HOP-U(40)---SCORS	2007
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH/Heart, Lung, and Blood Program Project (HLBP) Review Committee of NHLBI	2007
Research Grant Proposal	<i>Ad-hoc</i> Member	NIH/ZRG1 Clinical Cardiovascular Sciences (CVS) C 90S	2004
Research Grant Proposals	<i>Ad-hoc</i> Member	NIH/Social Sciences, Nursing, Epidemiology and Methods (SNEM5) (02) Special Emphasis Panel	2002-2003
Research Grant Proposals	<i>Ad-hoc</i> Member	NIH/Epidemiology and Disease Control –1 (EDC1) Study Section	2001-2003
Research Grant Proposals	Member	NIH/National Institute of Environmental Health Sciences (NIEHS), Special Emphasis Panel Review Group	1998
Abstract Selection Committee for the category “Lipid and Lipoprotein Metabolism: Basic & Cellular (ATVB)”	Member	American Heart Association Annual Scientific Meeting	2006
Abstract Selection Committee for the category “Genetics and Genomics of Cardiovascular Disease (FGTB)”	Member	American Heart Association Annual Scientific Meeting	2008
Scientific Advisory Board	Founding Member	Cure Alzheimer’s Fund	2005-2011
Research Grant Proposal	Reviewer	Biomedical Research Council (BMRC) of the Agency for Science, Technology and Research (A*STAR) , Singapore	2005
Research Grant Proposals	External Reviewer	Research Grants Council (RGC) of Hong Kong	2007
Research Grant Proposal	Reviewer	Medical Research Council, United Kingdom	2003, 2009
Research Grant Proposals	Member	Alzheimer’s Association, Initial Review Board of the Medical and Scientific Advisory Council.	1999-Present
Abstract Selection Committee	Member	American College of Rheumatology	2002

for the category
“Antiphospholipid Syndrome”

Annual Scientific Meeting

Research Grant Proposals	Reviewer	Swiss National Science Foundation	2002
Research Grant Proposal	Reviewer	Arthritis Research Campaign, United Kingdom	2001
Research Grant Proposals	Reviewer	The Wellcome Trust, United Kingdom	2000
Research Grant Proposals	Reviewer	The Wellcome Trust, United Kingdom	2005
Research Grant Proposals	Reviewer	The Wellcome Trust, United Kingdom	2007
Research Grant Proposals	Reviewer	The Wellcome Trust, United Kingdom	2009
Research Grant Proposals	Reviewer	University of Pittsburgh Alzheimer's Disease Research Center	2001- Present
Research Grant Proposals	Reviewer	National Science Foundation	1999
Research Grant Proposals	Reviewer	The Israel Science Foundation	1994
Research Grant Proposals	Chartered Member	National Dairy Promotion and Research Board	1993- 1996
Research Grant Proposals	Reviewer	Evaluation and Site Visit of a Program Project of the National Dairy Board, Dairy Research Institute for Genetics and Nutrition, Berkeley, CA	1991
Research Grant Proposal	Reviewer	Ministry of Public Health, Kuwait	1986

e. Patents

Title	US Patent No.	Issued Date	Inventors
Determination of Alzheimer's Disease Risk Using Apolipoprotein E and α -1 Antichymotrypsin Genotype Analysis	5,773,220	June 30, 1998	Steven, T. DeKosky, M. Ilyas Kamboh
Identification of Apolipoprotein H Mutations and Their Diagnostic Uses	6,203,980 B1	March 20 2001	M. Ilyas Kamboh, D. K. Sanghera, S. Manzi

f. Other

Position	Type of Service	Agency	Year(s)
Member	To select the best	Gabriel W. Lasker Award Committee	1998

	paper published in <i>Human Biology</i>		
External Examiner	Ph.D. Dissertation (Dr. Lin Yanaun)	National University of Singapore	1997
External Examiner	Ph.D. Dissertation (Dr. Tan Hui Hui Jenny)	National University of Singapore	2004
Member, Board of Examiners	Ph.D. Dissertation (S. Kanthimathi)	University of Madras, India	2006
External Examiner	Ph.D. Dissertation (I. Ahmad)	The Islamia University of Bahawalpur, Pakistan	2009
External Examiner	M. Phil Thesis (S.Mahmood)	University of Health Sciences, Lahore, Pakistan	2009
External Examiner	Ph.D. Dissertation (Ghulab Sher)	Quaid-i-Azam University, Islamabad, Pakistan	2014