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PERSONAL INFORMATION

Title: Professor
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HIGHER EDUCATION

1993-99 University of Michigan, Ph.D. in Clinical Psychology

1993-96 University of Michigan, M.A. in Clinical Psychology

1988-1992 Duke University, B.A. in English with Honors

1990 Duke in Oxford Program, Oxford University, England

POSTDOCTORAL TRAINING

2000-2001 Clinical Fellow in Geropsychology in the Department of Psychiatry, Harvard Medical School, Brockton (MA) VA Medical Center (Michele Karel, PhD, and Jennifer Moye, PhD, primary supervisors)

1999-2000 Research Fellow, Serious Mental Illness Treatment Research & Evaluation Center, Ann Arbor VA Health Services Research & Development Center of Excellence (Frederic Blow, PhD, advisor)

EMPLOYMENT & POSITIONS

2018-present Professor, with tenure, Department of Health Behavior & Health Education, University of Michigan School of Public Health (UM SPH)

2013-present Director, Outreach, Recruitment & Education Core, University of Michigan Alzheimer's Disease Center

2011-present Director, Public Health Genetics Certificate Program, UM SPH

2008-present Co-Director, Dual Degree Program in Public Health and Genetic Counseling, University of Michigan Schools of Public Health and Medicine

- 2007-present Core Faculty, University of Michigan Center for Bioethics and Social Sciences in Medicine (Director, Genomics, Health and Society Program)
- 2011-18 Associate Professor, with tenure, Department of Health Behavior & Health Education, University of Michigan School of Public Health (UM SPH)
- 2013 Visiting Scholar, University of California-Berkeley
- 2006-2011 Assistant Professor, Department of Health Behavior & Health Education, University of Michigan School of Public Health
- 2005-06 Co-Director, Education Core, Boston University Alzheimer's Disease Center
- 2003-06 Assistant Professor, Department of Neurology, Boston University School of Medicine
- 2005 Lecturer, Department of Gerontology, University of Massachusetts-Boston
- 2001-2003 Instructor, Department of Neurology, Boston University School of Medicine
- 1999-2000 Postdoctoral Psychology Fellow, Ann Arbor (MI) VA Medical Center
- 1998-99 Limited License Psychologist, St. Joseph's Mercy Hospital, Ann Arbor, MI
- 1996-97 Psychology Intern, Durham (NC) VA Medical Center

HONORS AND AWARDS

- 2004 Fellow, Hartford Geriatric Health Care Research Career Development Institute
- 2002 Fellow, Summer Research Institute, National Institute on Aging
- 2000 Fellow, Summer Research Institute, American Association for Geriatric Psychiatry/National Institute of Mental Health
- 1998 University of Michigan Rackham Dissertation Fellowship
- 1997-98 Predoctoral Fellowship, Ann Arbor VA Medical Center Department of Health Services Research and Development
- 1988-92 Dean's List, Duke University

PUBLICATIONS

Peer-Reviewed Journal Articles

Note: Underlined authors are students/trainees; * indicates senior or co-senior authorship

1. Mattos MK, Sereika SM, Beach SR, Klunk WE, Knox M, Nadkarni N, Parker LS, **Roberts JS**, Schulz R, Tamres L, Lingler JH. (2019, July 17). Research use of ecological momentary assessment for adverse event monitoring following amyloid- β imaging. *Journal of Alzheimer's Disease*, E-pub ahead of print. [PMID 31322563]
2. Leggett A, Connell CM, Dubin L, Dunkle R, Langa KM, Maust DT, **Roberts JS**, Spencer B, Kales HC. (2019, May 27). Dementia care across a tertiary-care health system: What

exists now and what needs to change. *Journal of the American Medical Directors Association*. E-pub ahead of print. [PMID 31147289]

3. Spector-Bagdady K, Fakih A, Krenz C, Marsh E, **Roberts JS***. (2019, June 17). Academic publications with privately owned or generated genetic data. *Genetics in Medicine*, E-pub ahead of print. [PMID 31204388]
4. **Roberts JS**. (2019, July 3). Assessing the psychological impact of genetic susceptibility testing: Where have we been, where do we go from here? *Hastings Center Report*, E-pub ahead of print.
5. **Roberts JS**, Gornick M, Le LQ, Bartnik N, Zikmund-Fisher B, Chinnaiyan A. (2019). Next-generation sequencing in precision oncology: Patient understanding and expectations. *Cancer Medicine*, 8(1), 227-37. [PMID 30600607]
6. Langbaum JB, Karlawish J, **Roberts JS**, et al. (2019). GeneMatch: A novel recruitment registry to conduct at-home *APOE* genotyping to enhance referrals to Alzheimer's prevention studies. *Alzheimer's & Dementia*, 15(4), 515-24. [PMID 30772251]
7. Gornick M, Ryan KA, Scherer AM, **Roberts JS**, De Vries RG, Uhlmann WR. (2019). Interpretations of the term "actionable" when discussing genetic test results: What you mean is not what I heard. *Journal of Genetic Counseling*, 28, 334-42. [PMID 30194681]
8. Grob R, **Roberts JS**, Timmermans S. (2018). Families' experiences with newborn screening: A critical source of evidence. *Hastings Center Report*, 48(S2), S29-S31. [PMID 30133722]
9. Guan Y, Roter DL, Erby LH, Wolff JL, Gitlin LN, **Roberts JS**, Green RC, Christensen KD. (2018). Communication predictors of patient and companion satisfaction with Alzheimer's disease genetic risk disclosure. *Journal of Health Communication*, 23(8), 807-14. [PMID 30325721]
10. Sturm AC, Schmidlen T, Scheinfeldt L, Hovick S, McElroy JP, Toland AE, **Roberts JS**, Sweet K. (2018). Early outcome data assessing utility of a post-test genomic counseling framework for the scalable delivery of precision health. *Journal of Personalized Medicine*, Jul 25; 8(3). [PMID 30046027]
11. Lingler JH, **Roberts JS**, Kim H, Morris J, Hu L, Mattos M, McDade E, Lopez O. (2018). Amyloid PET candidates may focus more on benefits than risks of results disclosure. *Alzheimer's & Dementia: Diagnosis, Assessment, and Disease Monitoring*, 10, 413-20. [PMID 30094328]
12. **Roberts JS**, Robinson JO, Diamond PM, Bharadwaj A, Christensen KD, Lee KB, Green RC, McGuire AL. (2018). Patient understanding of, satisfaction with, and perceived utility of whole genome sequencing: Findings from the MedSeq Project. *Genetics in Medicine*, 20(9), 1069-76. [PMID: 29300387]
13. Gornick M, Cobain E, Le LQ, Bartnik N, Stoffel E, Schuetze S, Talpaz M, Chinnaiyan A, **Roberts JS***. (2018, February 21). Oncologists' use of clinical sequencing data to inform clinical management. *JCO Precision Oncology*. DOI: 10.1200/PO.17.00122
14. Sturm AC, Hovick S, Schmidlen T, Scheinfeldt L, **Roberts JS**, et al. (2018). Operationalizing the reciprocal engagement model of genetic counseling practice: A framework for the scalable delivery of genomic counseling and testing. *Journal of Genetic Counseling*, 27(5), 1111-29. [PMID: 29460110]

15. Guan Y, Roter DL, Wolff JL, Gitlin LN, Christensen KD, **Roberts JS**, Green RC, Erby LH. (2018). The impact of genetic counselors' use of facilitative strategies on cognitive and emotional processing of genetic risk disclosure for Alzheimer's disease. *Patient Education & Counseling*, 101(5), 817-23. [PMID: 29203084]
16. Christensen KD, Uhlmann WR, **Roberts JS**, et al. (2018). A randomized controlled trial of disclosing genetic risk information for Alzheimer's disease via telephone. *Genetics in Medicine*, 20(1), 132-41. [PMID: 28726810]
17. Weipert C, Ryan K, Everett J, Yashar B, Chinnaiyan A, **Roberts JS**, De Vries R, Zikmund-Fisher B, Raymond V. (2018). Physician experiences and understanding of genomic sequencing in oncology. *Journal of Genetic Counseling*, 27(1), 187-96. [PMID: 28840409]
18. Landry L, Nielsen DE, Carere DA, **Roberts JS**, Green RC. (2017). Racial minority group interest in direct-to-consumer genetic testing: Findings from the PGen Study. *Journal of Community Genetics*, 8(4), 293-301. [PMID: 28868574]
19. Ostergren JE, Heeringa SG, Mendes de Leon CF, Connell CM, **Roberts JS***. (2017). The influence of psychosocial and cognitive factors on perceived threat of Alzheimer's disease. *American Journal of Alzheimer's Disease & Other Dementias*, 32(5), 289-99. [PMID: 28605999]
20. Robinson DR, Wu YM, Lonigro RJ, Cieslik M, Cobain E, Everett J, Cao X, Vats P, Rabban E, Kumar C, Raymond V, Schuetze S, Alva A, Siddiqui J, Chugh R, Worden F, Zalupski MM, Innis J, Mody RJ, Tomlins SA, Lucas D, Romnath N, Schott AF, Hayes DF, Stoffel EM, **Roberts JS**, Smith DC, Kunju LP, Talpaz M, Chinnaiyan AM. (2017). Integrative clinical genomics of metastatic cancer. *Nature*, 548(7667):297-303. [PMID: 28783718]
21. Gollust SE, Gray SW, Carere DA, Koenig BA, Lehmann LS, McGuire A, Sharp R, Spector-Bagdady K, Wang N, Green RC, **Roberts JS***. (2017). Consumer perspectives on access to direct-to-consumer genetic testing: Role of demographic factors and the testing experience. *Milbank Quarterly*, 95(2), 291-318. [PMID: 28589610] [PMCID: PMC5461393]
22. Koeller DR, Uhlmann WR, Carere DA, Green RC, **Roberts JS***. (2017). Utilization of genetic counseling after direct-to-consumer genetic testing: Findings from the Impact of Personal Genomics (PGen) Study. *Journal of Genetic Counseling*, 26(6), 1270-79. [PMID: 28512697] [PMCID: PMC5673568 Available on 2018-12-01]
23. Nielsen DE, Carere DA, Wang C, **Roberts JS***, Green RC. (2017). Diet and exercise changes following direct-to-consumer genetic testing. *BMC Medical Genomics*, 10(1):24. doi: 10.1186/s12920-017-0258-1. [PMID: 28464943] [PMCID: PMC5412058]
24. Grill JD, Apostolova LG, Bullain S, ...**Roberts JS**, et al. (2017). Communicating Mild Cognitive Impairment diagnosis with and without amyloid imaging. *Alzheimer's Research & Therapy*, 9:35 DOI 10.1186/s13195-017-0261-y [PMID: 28472970] [PMCID: PMC5418690]
25. Ryan K, De Vries R, Uhlmann WR, **Roberts JS***, Gornick MC. (2017). Public's views toward return of secondary results in genomic sequencing: It's (almost) all about the choice. *Journal of Genetic Counseling*, 26(6), 1197-1212. [PMID 28357777] [PMCID: PMC5620108]
26. Sweet K, Sturm AC, Schmidlen T, ... **Roberts JS**, et al. (2017). Outcomes of a randomized controlled trial of genomic counseling for patients receiving personalized and actionable complex disease reports. *Journal of Genetic Counseling*, 26(5), 980-98. [PMID: 28345121] [PMCID: PMC5617760 Available on 2018-10-01]
27. Qian J, Wolters F, Beiser A, Haan M, Ikram A, Karlawish J, Langbaum JB, Neuhaus JM, Reiman EM, **Roberts JS**, Seshadri S, Tariot PN, Betensky RA, Blacker D. (2017, March

- 21). APOE-related risk of mild cognitive impairment and dementia for prevention trials: An analysis of four cohorts. *PLoS Medicine*, e-pub ahead of print. [PMID: 28323826] [PMCID: PMC5360223]
28. Poey JL, Burr JA, **Roberts JS***. (2017). Social connectedness, perceived isolation, and dementia: Does the social environment moderate the relationship between genetic risk and cognitive well-being? *The Gerontologist*, 57(6), 1031-40. [PMID: 28329797]
29. **Roberts JS**, Gornick MC, Carere DA, Uhlmann WR, Ruffin MT, Green RC. (2017). Direct-to-consumer genetic testing: User motivations, decision making, and perceived utility of results. *Public Health Genomics*, 20(1), 36-45. [PMID: 28068660]
30. Gornick MC, Scherer AM, Sutton EJ, Ryan KA, Exe NL, Li M, Uhlmann WR, Kim SYH, **Roberts JS***, De Vries RG. (2017). Effect of public deliberation on attitudes toward return of secondary results in genomic sequencing. *Journal of Genetic Counseling*, 26(1), 122-32. [PMID: 27307100] [PMCID: PMC5161735]
31. Gray SW, Gollust SE, Carere DA, Chen CA, Cronin A, Kalia SS, Rana HQ, Ruffin MT, Wang C, **Roberts JS***, Green RC. (2017). Personal genomic testing for cancer risk: Results from the Impact of Personal Genomics (PGen) Study. *Journal of Clinical Oncology*, 35(6), 636-44. [PMID: 27937091] [PMCID: PMC5455805]
32. Katapodi MC, Duquette D, Yang JJ, Mendelsohn-Victor K, Anderson B, Nikolaidis C, Mancewicz E, Northouse LL, Duffy S, Ronis D, Milliron KJ, Meraver SD, Janz NK, Copeland G, Probst-Hensch N, **Roberts JS***. (2017). Recruiting families at risk for hereditary breast and ovarian cancer from a statewide cancer registry. *Cancer Causes and Control*, 28(3), 191-201. [PMID: 28197806]
33. Sweet K, Sturm AC, Schmidlen T, ... **Roberts JS**, Christman M. (2017). EMR documentation of physician-patient communication following genomic counseling for actionable complex disease and pharmacogenomic results. *Clinical Genetics*, 91(4), 545-56. [PMID: 27322592] [PMCID: PMC5173448 Available on 2018-04-01]
34. Guan Y, Roter DL, Erby LH, Wolff JL, Gitlin LN, **Roberts JS**, Christensen KD, Green RC. Disclosing genetic risk of Alzheimer's disease to cognitively impaired patients and visit companions: Findings from the REVEAL Study. (2017). *Patient Education and Counseling*, 100(5), 927-935. [PMID: 28012682] [PMCID: PMC5400683 Available on 2018-05-01]
35. Carere DA, VanderWeele T, Vassy JL, van der Wouden C, **Roberts JS**, Kraft P, Green RC. (2017). Prescription medication changes following direct-to-consumer personal genomic testing: Findings from the PGen Study. *Genetics in Medicine*, 19, 537-45. [PMID: 27657683] [PMCID: PMC5362351]
36. Sweet K, Hovick S, Sturm AC, Schmidlen T, Gordon ES, Bernhardt BA, Wawak L, Wernke K, McElroy J, Scheinfeldt L, Toland AE, **Roberts JS**, Christman M. (2017). Counselors' perspectives of genomic counseling following online receipt of multiple actionable complex disease and pharmacogenomic results: A qualitative research study. *Journal of Genetic Counseling*, 26(4), 738-51. [PMID: 27921197] [PMCID: PMC5459668 Available on 2018-08-01]
37. Krieger JL, Murray F, **Roberts JS***, Green RC. (2016). The impact of personal genomics on risk perceptions and medical decision-making. *Nature Biotechnology*, 34(9), 912-18. [PMID: 27606453]
38. Olfson E, Hartz S, Carere D, Green RC, **Roberts JS**, Bierut L. (2016). Implications of personal genomic testing for health behaviors: the case of smoking. *Nicotine & Tobacco Research*, 18(12), 2273-77. [PMID: 27613923] [PMCID: PMC5103936]

39. Christensen KD, **Roberts JS**, Whitehouse PJ, et al. (2016). Disclosing pleiotropic effects during genetic risk assessment for Alzheimer's disease: A randomized, controlled trial. *Annals of Internal Medicine*, 164, 155-63. [PMID: 26810768] [PMCID: PMC4979546]
40. Jones T, Lochhart JS, Mendelsohn-Victor KE,...**Roberts JS**, Katapodi MC. (2016). Use of cancer genetics services in African American young breast cancer survivors. *American Journal of Preventive Medicine*, 51(4), 427-36. [PMID: 27117712]
41. Green RC, Goddard KAB, Amendola LM,... **Roberts JS**, et al. (2016). The Clinical Sequencing Exploratory Research Consortium: Accelerating the evidence-based practice of genomic medicine. *American Journal of Human Genetics*, 98(6), 1051-66. [PMID: 27181682] [PMCID: PMC4908179]
42. Lingler JH, Butters MA, Hu L, Hunsaker AE, Klunk WE, Mattos MK, Parker LA, **Roberts JS**, Schulz R. (2016). Development of a standardized approach to disclosing amyloid imaging research results in mild cognitive impairment. *Journal of Alzheimer's Disease*, 52(1), 17-24. [PMID: 27060950] [PMCID: PMC4860948]
43. Baptista NM, Christensen KD, Carere DA, Broadley SA, **Roberts JS***, Green RC. (2016). Adopting genetics: Motivations and outcomes of personal genomic testing in adult adoptees. *Genetics in Medicine*, 18(9), 924-32. [PMID: 26820063] [PMCID: PMC4965328]
44. van der Wouden CH, Carere DA, van der Zee AHM, Ruffin MT, **Roberts JS***, Green RC. (2016). Consumer perceptions of primary care provider interactions following direct-to-consumer genetic testing. *Annals of Internal Medicine*, 164, 513-22. [PMID: 26928821]
45. Carere DA, VanderWeele T, Moreno TA, Mountain JL, **Roberts JS**, Kraft P, Green RC. (2015). The impact of direct-to-consumer personal genomic testing on perceived risk of breast, prostate, colorectal, and lung cancer: Findings from the PGen Study. *BMC Medical Genomics*, 8(1), 63. [PMID: 26468061] [PMCID: PMC4606558]
46. Mody RJ, Wu YM, Lonigro RJ,...**Roberts JS**,...Chinnaiyan AM. (2015). Integrative clinical sequencing in the management of refractory or relapsed cancer in youth. *JAMA*, 314(9), 913-25. [PMID: 26325560] [PMCID: PMC4758114]
47. Ostergen JE, Gornick MC, Carere DA, Kalia SS, Uhlmann WR, Ruffin MT, Mountain JL, Green RC, **Roberts JS*** (2015). How well do customers of direct-to-consumer personal genomic testing services comprehend genetic test results? Findings from the Impact of Personal Genomics (PGen) Study. *Public Health Genomics*, 18(4), 216-24. [PMID: 26087778] [PMCID: PMC4926310]
48. Meisel SM, Carere DA, Wardle J, Kalia SS, Moreno TA, Mountain JL, **Roberts JS***, Green RC (2015). Explaining, not just predicting, drives interest in personal genomics. *Genome Medicine*, 7(1):74. [PMCID: PMC4533947]
49. Christensen KD, **Roberts JS**, Zikmund-Fisher BJ, Kardia SLR, McBride CM, Linnenbringer E, Green RC (2015). Associations between self-referral and health behavior responses to genetic risk information. *Genome Medicine*, 7(1):10. [PMID: 25642295] [PMCID: PMC4311425]
50. Witte MM, Foster NL, Fleisher AS...**Roberts JS**, et al. Clinical use of amyloid PET neuroimaging: Practical and bioethical considerations. (2015). *Alzheimer's & Dementia: Diagnosis, Assessment and Disease Monitoring*, 1(3), 358-67. [PMID: 27239516] [PMCID: PMC4878065]
51. Carere DA, Kraft P, Kaphingst KA, **Roberts JS***, Green RC. (2016). Consumers report lower confidence in their genetics knowledge following direct-to-consumer genetic testing. *Genetics in Medicine*, 18(1), 65-72. [PMID: 25812042] [PMCID: PMC4583799]

52. Shiloh S, deHeer HD, Peleg S, Hensley Alford S, Skapinsky K, **Roberts JS**, Hadley DW. (2015). The impact of multiplex genetic testing on disease risk perceptions. *Clinical Genetics*, 87(2), 117-23. [PMID: 24720448]
53. **Besser AG**, Sanderson SC, **Roberts JS**, Chen CA, Christensen KD, Lautenbach DM, Cupples LA, Green RC. (2015). Factors affecting recall of different types of personal genetic information about Alzheimer's disease risk: The REVEAL Study. *Public Health Genomics*, 18(2), 78-86. [PMID: 25634646] [PMCID: PMC4470386]
54. Green RC, Christensen KD, Cupples LA, Relkin NR, Whitehouse PJ, Royal CDM, . . . **Roberts JS*** (2015). A randomized non-inferiority trial of condensed protocols for genetic risk disclosure of Alzheimer's disease. *Alzheimer's and Dementia*, 11(10), 1222-30. [PMID: 25499536] [PMCID: PMC4461546]
55. Khan C, Rini C, Bernhardt BA, **Roberts JS**, Christensen KD, Evans JP, Brothers KB, Roche MI, Berg JS, Henderson GE. (2015). How can psychological science inform research and clinical applications of next generation genomic sequencing? *Journal of Genetic Counseling*, 24(2), 193-204. [PMID: 25488723] [PMCID: PMC4777349]
56. **Carere DA**, Couper MP, Crawford SD, Kalia SS, Duggan JR, Moreno TA, Mountain JL, **Roberts JS***, Green RC. (2014). Design, methods and participant characteristics of the Impact of Personal Genomics (PGen) Study, a prospective cohort study of personal genomic testing customers. *Genome Medicine*, 6(12):96. [PMID 25484922] [PMCID: PMC4256737]
57. **Roberts JS**, Dolinoy DC, & Tarini BA. (2014). Emerging issues in public health genomics. *Annual Review of Genomics and Human Genetics*, 15, 461-80. [PMID: 25184533] [PMCID: PMC4229014]
58. **Roberts JS**, Connell CM, & **McLaughlin S**. (2014). Public knowledge and beliefs about risk and protective factors for Alzheimer's disease: Findings from the Health and Retirement Study. *Alzheimer's and Dementia*, 10(5 Suppl.), S381-9. [PMID: 24630852] [PMCID: PMC4163539]
59. Vassy JL, Lautenbach DM, . . . **Roberts JS**, et al. (2014). The MedSeq Project: A randomized trial of integrating whole genome sequencing into clinical medicine. *Trials*. March 20;15(1):85. [PMID: 24645908] [PMCID: PMC4113228]
60. **Lerner B**, **Roberts JS**, Shwartz M, Roter DL, Green RC, & Clark JA. (2014). Distinct communication patterns during genetic counseling for late-onset Alzheimer's risk assessment. *Patient Education and Counseling*, 94(2), 170-79. [PMID: 24316056] [PMCID: PMC3927403]
61. Robinson DR, Wu YM, Vars P, . . . **Roberts JS**, et al. (2013). Activating ESR1 mutations in hormone-resistant metastatic breast cancer. *Nature Genetics*, 45, 1446-51. [PMID: 24185510] [PMCID: PMC4009946]
62. **Roberts JS** & **Ostergren, J**. (2013). Direct-to-consumer genetic testing and personal genomics services: A review of recent empirical studies. *Current Genetic Medicine Reports*, 1(3), 182-200. [PMID: 24058877] [PMCID: PMC3777821]
63. **Roberts JS**, Dunn L, & Rabinovici G. (2013). Amyloid imaging, risk disclosure, and Alzheimer's disease: Ethical and practical issues. *Neurodegenerative Disease Management*, 3(3), 219-29. [PMID: 26167204] [PMCID: PMC4498486]
64. Molnar LJ, Eby DW, Langford J, Charlton JL, St. Louis RM, **Roberts JS**. (2013). Tactical, strategic, and life-goal self-regulation of driving by older adults: Development and testing of a questionnaire. *Journal of Safety Research*, 46, 107-117. [PMID: 23932692]

65. **Roberts JS** & Uhlmann WR. (2013). Genetic susceptibility testing for neurodegenerative diseases: Ethical and practice issues. *Progress in Neurobiology*, 110, 89-101. [PMID: 23583530] [PMCID: PMC3772971]
66. Shiloh S, Wade CH, **Roberts JS**, Alford SH, & Biesecker BB. (2013). On averages and peaks: how do people integrate information about multiple diseases to reach a decision about multiplex genetic testing? *Medical Decision Making*, 33, 71-77. [PMID: 23128581] [PMCID: PMC3799841]
67. Shiloh S, Wade C, **Roberts JS**, Alford SH, & Biesecker BB. (2013). Associations between risk perceptions and worry about common diseases: A between- and within-subjects examination. *Psychology and Health*, 28, 434-49. [PMID: 23121110] [PMCID: PMC3566271]
68. Wade C, Shiloh S, **Roberts JS**, Alford SH, Marteau T, & Biesecker BB. (2012). Preferences among diseases on a genetic susceptibility test for common health conditions: An ancillary study of the Multiplex Initiative. *Public Health Genomics*, 15(6), 322-26. [PMID: 22688356] [PMCID: PMC3514560]
69. **Roberts JS**, Chen C, Uhlmann WR, & Green RC. (2012). Effectiveness of a condensed protocol for disclosing *APOE* genotype and providing risk education for Alzheimer's disease: The REVEAL Study. *Genetics in Medicine*, 14, 742-48. [PMID: 22498844] [PMCID: PMC3718049]
70. **Roberts JS**. (2012). Genetic testing for Alzheimer's risk: Benefit or burden? *Neurodegenerative Disease Management*, 2(2), 141-44. PMCID: Not required
71. Lehmann L, Kaufman D, Sharp R, Moreno T, Mountain J, **Roberts JS***, & Green RC. (2012). Navigating a research partnership between academia and industry to assess the impact of personalized genetic testing. *Genetics in Medicine*, 14, 268-73. [PMID: 22241103] [PMCID: PMC3763722]
72. Langford A, Resnicow K, **Roberts JS**, & Zikmund-Fisher B. (2012). Racial and ethnic differences in genetic test Awareness in HINTS 2007: Sociodemographic and numeracy correlates. *Journal of Genetic Counseling*, 21, 440-47. [PMID: 22271378]
73. Roychowdhury S, Iyer MK, Robinson DR,...**Roberts JS**, et al. (2011). Personalized oncology through integrative high-throughput sequencing: A pilot study. *Science Translational Medicine*, 3, 111ra121. DOI: 10.1126/scitranslmed.3003161 [PMID: 22133722] [PMCID: PMC3476478]
74. **Roberts JS**, Christensen KD, & Green RC. (2011). Using Alzheimer's disease as a model for genetic risk disclosure: Implications for personal genomics. *Clinical Genetics*, 80, 407-14. [PMID: 21696382] [PMCID: PMC3191239]
75. Goldman J, Catania JW, LaRusse-Eckert S, Butson MB, Rumbaugh M, Strecker MN, Hahn SE, **Roberts JS**, Burke W, Mayeux R, & Bird TD. (2011). Genetic testing and genetic counseling for Alzheimer's disease: Joint Practice Guidelines of the American College of Medical Genetics and the National Society of Genetic Counselors. *Genetics in Medicine*, 6, 597-605. [PMID: 21577118] [PMCID: PMC3326653]
76. Kopits I, Chen CA, **Roberts JS**, Uhlmann WR, & Green RC. (2011). Willingness to pay for genetic testing for Alzheimer's disease: A measure of personal utility. *Genetic Testing and Molecular Biomarkers*, 15, 871-75. [PMID: 21749214] [PMCID: PMC3241735]

77. Wade C, Shiloh S, Woolford SW, **Roberts JS**, Alford SH, Marteau TM, & Biesecker BB. (2011). Modeling decisions to undergo genetic testing for susceptibility to common health conditions: An ancillary study of the Multiplex Initiative. *Psychology and Health, 27*(4), 430-44. [PMID: 21660870] [PMCID: PMC3175306]
78. Akinleye I, **Roberts JS**, Royal CDM, Linnenbringer E, Obisesan TO, Fasaye GA, & Green RC. (2011). Differences between African American and White research volunteers in their attitudes, beliefs and knowledge regarding genetic testing for Alzheimer's disease. *Journal of Genetic Counseling, 20*, 650-59. [PMID: 21656311] [PMCID: PMC3223287]
79. Christensen KD, **Roberts JS**, Shalowitz DI, Kim S, Raskin L, & Gruber SB. (2011). Disclosing CDKN2A genotypes from research to melanoma survivors: Interest, impact and demands on researchers. *Cancer Epidemiology, Biomarkers and Prevention, 20*, 522-29. [PMID: 21307304] [PMCID: PMC3833711]
80. Christensen KD, **Roberts JS**, Uhlmann WR, & Green RC. (2011). Changes in perceptions about the benefits, risks and limitations of genetic susceptibility testing for Alzheimer's disease risk. *Genetics in Medicine, 13*, 409-14. [PMID: 21270636] [PMCID: PMC3170997]
81. Hock KT, Christensen KD, Yashar BM, **Roberts JS**, Gollust SE, & Uhlmann WR. (2011). Direct-to-consumer genetic testing: An assessment of genetic counselors' knowledge and attitudes. *Genetics in Medicine, 13*, 325-32. [PMID: 21233722] [PMCID: PMC3804135]
82. **Roberts JS**, Shalowitz DI, Christensen KD, Everett J, Kim S, Raskin L, & Gruber SB. (2010). Returning individual research results: Development of a cancer genetics education and risk communication protocol. *Journal of Empirical Research on Human Research Ethics, 3*, 17-30. [PMID: 20831418] [PMCID: PMC3159194]
83. **Roberts JS**, Karlawish J, Uhlmann WR, Petersen R, & Green RC. (2010). Mild Cognitive Impairment in clinical care: A survey of neurologists' attitudes and practices. *Neurology, 75*, 425-31. [PMID: 20679636] [PMCID: PMC2918467]
84. **Roberts JS** & Tersegno S. (2010). Estimating and disclosing the risk of developing Alzheimer's disease: Challenges, controversies and future directions. *Future Neurology, 5*, 501-17. [PMID: 20856693] [PMCID: PMC2941213]
85. Ashida S, Koehly LM, **Roberts JS**, Chen CA, Hiraki S, & Green RC. (2010). The role of disease perceptions and results sharing on psychological adaptation after genetic susceptibility testing: The REVEAL Study. *European Journal of Human Genetics, 12*, 1296-1301. [PMID: 20664629] [PMCID: PMC2988099]
86. Vernarelli J, **Roberts JS**, Hiraki S, Chen CA, Cupples LA, & Green RC. (2010). Impact of Alzheimer's disease genetic risk disclosure on dietary supplement use. *American Journal of Clinical Nutrition, 91*, 1402-07. [PMID: 20219963] [PMCID: PMC2854909]
87. Christensen KD, Jayaratne TE, **Roberts JS**, Kardia SLR, & Petty EM. (2010). Understandings of basic genetics in the United States: Results from a national survey of Black and White men and women. *Public Health Genomics, 3*, 467-76. [PMID: 20203477] [PMCID: PMC3025896]
88. Linnenbringer E, **Roberts JS**, Hiraki S, Cupples LA, & Green RC. (2010). "I know what you told me, but this is what I think": Perceived risk of Alzheimer's disease among individuals who accurately recall their genetics-based risk estimate. *Genetics in Medicine, 12*, 219-27. [PMID: 20139767] [PMCID: PMC2921681]

89. McLaughlin S, Connell CM, Heeringa SG, Li LW, & **Roberts JS**. (2010). Successful aging in the United States: Prevalence estimates from a national sample of older adults. *Journals of Gerontology Series B: Psychological Sciences and Social Sciences*, 65, 216-26. [PMID: 20008481] [PMCID: PMC2981444]
90. Taylor DH, Cook-Deegan, R. M., Hiraki, S., **Roberts, J. S.**, Blazer, D. G., & Green, R. C. (2010). Genetic testing for Alzheimer's and long-term care insurance. *Health Affairs*, 29, 102-108. [PMID: 20048367] [PMCID: PMC2931337]
91. Green RC, **Roberts JS**, Cupples LA, et al. (2009). A randomized trial of APOE disclosure for risk of Alzheimer's disease: The REVEAL Study. *New England Journal of Medicine*, 361, 245-54. [PMID: 19605829] [PMCID: PMC2778270]
92. **Roberts JS** & Silverio EA. (2009). Evaluation of an education and support program for early-stage Alzheimer's disease. *Journal of Applied Gerontology*, 28, 419-35. DOI: 10.1177/0733464809333883.
93. Khoury MJ, McBride C, Schully SD, Ioannidis JPA, Feero WG, Janssens AC, **Roberts JS**, et al. (2009). The scientific foundation for personal genomics: Recommendations from an NIH-CDC multidisciplinary workshop. *Genetics in Medicine*, 11, 559-67. [PMID: 19617843] [PMCID: PMC2936269]
94. Connell CM, **Roberts JS**, McLaughlin S, & Carpenter BD. (2009). Black and white adult family members' attitudes toward a dementia diagnosis. *Journal of the American Geriatrics Society*, 57, 1562-68. [PMID: 19682136]
95. Connell CM, **Roberts JS**, McLaughlin SJ, & Akinleye D. (2009). Racial differences in knowledge and beliefs about Alzheimer disease. *Alzheimer Disease & Associated Disorders*, 23, 110-16. [PMID: 19474569]
96. Wain K, Uhlmann WR, Heidebrink J, & **Roberts JS**.* (2009). Living at risk: The sibling's perspective of early-onset Alzheimer's disease. *Journal of Genetic Counseling*, 18, 239-51. [PMID: 19132519]
97. Ashida S, Koehly LM, **Roberts JS**, Hiraki S, & Green RC. (2009). Disclosing the disclosure: Factors associated with communicating the results of genetic susceptibility testing for Alzheimer's disease. *Journal of Health Communications*, 14, 768-84. [PMID: 20029710] [PMCID: PMC2801901]
98. Hiraki S, Chen CA, **Roberts JS**, Cupples LA, & Green RC. (2009). Perceptions of familial risk in those seeking a genetic risk assessment for Alzheimer's disease. *Journal of Genetic Counseling*, 18, 130-36. [PMID: 18949541] [PMCID: PMC2919070]
99. Chung W, Chen CA, Cupples LA, **Roberts JS**, Hiraki S, Nair A, Green RC, & Stern RA. (2009). A new scale measuring psychological impact of genetic susceptibility testing for Alzheimer disease. *Alzheimer Disease & Associated Disorders*, 23, 50-56. [PMID: 19266699] [PMCID: PMC2743905]
100. Cassidy M, **Roberts JS**, Bird TD, Steinbart E, Cupples LA, Chen CA, Linnenbringer E, & Green RC. (2008). Comparing test-specific distress of susceptibility versus deterministic genetic testing for Alzheimer's disease. *Alzheimer's and Dementia*, 4, 406-13. [PMID: 19012865] [PMCID: PMC2610442]
101. Fanshawe T, Prevost AT, **Roberts JS**, Green RC, Armstrong D, & Marteau TM. (2008).

- Explaining behaviour change after genetic testing: Design solutions to the problem of collinearity between test and risk estimates. *Genetic Testing*, 12, 381-86. [PMID: 18666860] [PMCID: PMC2925186]
102. Chao S, **Roberts JS**, Marteau TM, Silliman R, Cupples LA, & Green RC. (2008). Health behavior changes after genetic risk assessment for Alzheimer's disease: The REVEAL Study. *Alzheimer Disease & Associated Disorders*, 22, 94-97. [PMID: 18317253] [PMCID: PMC2483341]
 103. Christensen KD, **Roberts JS**, Royal CDM, Fasaye GA, Obisesan T, Cupples LA, Whitehouse PJ, Butson MB, Linnenbringer E, Relkin NR, Farrer L, Cook-Deegan R, & Green RC. (2008). Incorporating ethnicity into genetic risk assessment for Alzheimer's disease: The REVEAL Study experience. *Genetics in Medicine*, 10, 207-14. [PMID: 18344711] [PMCID: PMC2483343]
 104. Jarvik L, LaRue A, Blacker D,.... **Roberts JS**, et al. (2008). Children of persons with Alzheimer's disease: What does the future hold? *Alzheimer Disease & Associated Disorders*, 22, 6-20. [PMID: 18317242] [PMCID: PMC3377487]
 105. Connell CM, **Roberts JS**, & McLaughlin SJ. (2007). Public opinion about Alzheimer disease among Blacks, Hispanics, and Whites: Results from a national survey. *Alzheimer Disease & Associated Disorders*, 21, 232-40. [PMID: 17804956]
 106. Eckert SL, Katzen H, **Roberts JS**, Barber M, Ravdin LD, Whitehouse PJ, Relkin NR, & Green RC. (2006). Recall of disclosed Apolipoprotein E genotype and lifetime risk estimate for Alzheimer's disease: The REVEAL Study. *Genetics in Medicine*, 8, 746-51. [PMID: 17172937]
 107. Gooding HC, Linnenbringer EP, Burack J, **Roberts JS**, Green RC, & Biesecker BB. (2006). Genetic susceptibility testing for Alzheimer disease: Motivation to obtain information and control as precursors to coping with increased risk. *Patient Education and Counseling*, 64, 259-67. [PMID: 16860524]
 108. Binetti G, Benussi L, **Roberts JS**, et al. (2006). Areas of intervention for genetic counselling of dementia: Cross-cultural comparison between Italians and Americans. *Patient Education and Counseling*, 64, 285-93. [PMID: 16850519]
 109. **Roberts JS**, Cupples LA, Relkin NR, Whitehouse PJ, & Green RC. (2005). Genetic risk assessment for adult children of people with Alzheimer's disease. *Journal of Geriatric Psychiatry and Neurology*, 18, 250-55. [PMID: 16306249]
 110. Hurley A, Harvey R, **Roberts JS**, Wilson-Chase C, Lloyd S, Prest J, Lock M, Horvath KJ, & Green RC. (2005). Genetic susceptibility for Alzheimer's disease: Why did adult offspring seek testing? *American Journal of Alzheimer's Disease and Other Dementias*, 20(6), 374-81. [PMID: 16396443]
 111. Marteau TM, **Roberts JS**, LaRusse S, & Green RC. (2005). Predicting risk of Alzheimer's disease: Impact upon perceived risk of including DNA testing. *Risk Analysis*, 25(2), 397-404. [PMID: 15876213]
 112. LaRusse S, **Roberts JS**, Marteau TM, Katzen H, Linnenbringer E, Barber M, Whitehouse PJ, Quaid K, Brown T, Green RC, & Relkin N. (2005). Genetic susceptibility testing versus family history-based risk assessment: Impact on perceived risk of Alzheimer's disease. *Genetics in Medicine*, 7(1), 48-53. [PMID: 15654228]

113. Zick C, Mathews C, **Roberts JS**, Cook-Deegan R, Pokorski R, & Green RC. (2005). Genetic susceptibility testing for Alzheimer's disease and its impact on insurance behavior. *Health Affairs*, 24(2), 483-90. [PMID: 15757934] [PMCID: PMC1761120]
114. **Roberts JS**, Barber M, Brown T, Cupples LA, Farrer LA, LaRusse S, Post SG, Quaid K, Ravdin L, Relkin N, Sadovnick D, Whitehouse PJ, Woodard J, & Green RC. (2004). Who seeks genetic susceptibility testing for Alzheimer's disease? Findings from a multisite, randomized clinical trial. *Genetics in Medicine*, 6, 197-203. [PMID: 15266207]
115. **Roberts JS**, La Russe SA, Katzen H, Whitehouse PJ, Barber M, Post SG, Relkin N, Quaid K, Piertzak RH, Cupples LA, Farrer LA, Brown T, & Green RC. (2003). Reasons for seeking genetic susceptibility testing among first-degree relatives of people with Alzheimer's disease. *Alzheimer Disease and Associated Disorders*, 17, 86-93. [PMID: 12794385]
116. Hipps Y, **Roberts JS**, Farrer LA, & Green RC. (2003). Differences between African Americans and Whites in their attitudes toward genetic testing for Alzheimer's disease. *Genetic Testing*, 7, 39-44. [PMID: 12820701]
117. **Roberts JS**, Connell CM, Cisewski D, Hipps YG, Demissie S, & Green RC. (2003). Differences between African Americans and Whites in their perceptions of Alzheimer disease. *Alzheimer Disease and Associated Disorders*, 17, 19-26. [PMID: 12621316]
118. **Roberts JS**. (2000). Anticipating response to predictive genetic testing for Alzheimer's disease: A survey of first-degree relatives. *The Gerontologist*, 40, 43-52. [PMID: 10750312]
119. **Roberts JS** & Connell CM. (2000). Illness representations among first-degree relatives of people with Alzheimer disease. *Alzheimer Disease and Associated Disorders*, 14, 129-136. [PMID: 10994653]
120. **Roberts JS**, Blow FC, Copeland LC, Barry KL, & Van Stone W. (2000). Age-group differences in schizophrenia treatment outcomes for male veterans: A three-year longitudinal study. *Journal of Geriatric Psychiatry and Neurology*, 13, 78-86. [PMID: 10912729]
121. Kales HC, Blow FC, Bingham CR, **Roberts JS**, Copeland LC, & Mellow AM. (2000). Race, psychiatric diagnosis and mental health care utilization in older patients. *American Journal of Geriatric Psychiatry*, 8, 301-309. [PMID: 11069270]

Book Chapters

1. **Roberts JS**, Gornick MR. (In press). Ethical, legal, and social implications of precision cancer medicine. In S Roychowdhury, E Van Allen (Eds.), *Precision Cancer Medicine: Challenges and Opportunities*. Springer Publishing.
2. Uhlmann WR, **Roberts JS**. (2018). Ethical issues in neurogenetics. In D Geschwind, C Klein, H Paulson (Eds.), *Handbook of Clinical Neurology*, 147, 23-36. Elsevier. [PMID 29325614]
3. **Roberts JS**. (2008). Psychological issues in genetic testing. In C Read, M Smyer, & RC Green (Eds.), *Aging, Biotechnology, & the Future*. Baltimore: Johns Hopkins University Press (pp. 173-87).
4. Hurley A, Harvey R, **Roberts JS**, & Horvath KJ. Alzheimer's disease genetic susceptibility in an aging society. (2008). In C Read, M Smyer, & RC Green (Eds.), *Aging, Biotechnology, & the Future*. Baltimore: Johns Hopkins University Press (pp. 157-72).

5. Read CY, **Roberts JS**, Linnenbringer E, & Green RC. (2008). Genetic testing for Alzheimer's disease: The REVEAL Study. In C Read, M Smyer, & RC Green (Eds.), *Aging, Biotechnology, & the Future*. Baltimore: Johns Hopkins University Press (pp. 127-44).
6. **Roberts JS**. (2001). Genetic testing. In MD Mezey (Ed.), *Encyclopedia of Care of the Elderly* (pp. 298-300). New York: Springer Publishing.
7. **Roberts JS** & Rosenwald GC. (2001). Ever upward and no turning back: Social mobility and identity formation among first-generation college students. In DP McAdams, R Josselson, & A Lieblich (Eds.), *Turns in the Road: Narrative Studies of Lives in Transition* (pp. 91-120). Washington, DC: American Psychological Association.

Manuscripts in Submission

Christensen KD, Schonman EF, Robinson JO, **Roberts JS**, Diamond PM, Lee KB, Green RC, McGuire AL. Genome sequencing does not cause distress and motivates health behavior improvements in patients with increased polygenic risk for cardiometabolic traits. Submitted to *Genetics in Medicine*.

Marzulla T, **Roberts JS**, De Vries R, et al. Be prepared: Understanding the motivations, expectations, and experiences of direct-to-consumer clients who seek genetic counseling. Submitted to *Journal of Genetic Counseling*.

Christensen KD, Karlawish J, **Roberts JS**, et al. Disclosure of genetic risk for Alzheimer's disease to patients with mild memory problems: A randomized controlled trial. Submitted to *Annals of Internal Medicine*.

Love-Nichols J, Uhlmann WR, Arscott P, Hornsby W, Willer C, **Roberts JS***. A prospective survey of aortic disease biorepository participants' preferences for return of research genetic results. Submitted to *Circulation: Genomic & Precision Medicine*.

FUNDED PROJECTS (Note: all funds listed are total costs of the overall project)

Currently funded

- | | |
|---------|---|
| 2019-24 | National Human Genome Research Institute career development award (K01 HG010496)
Title: <i>Genetic Data Partnerships: Enabling Equitable Access within Academic/Private Data Sharing Agreements</i>
Role: Primary Research Mentor (K. Spector-Bagdady, PI)
Project Funding: \$174,971 for FY 2019 |
| 2018-23 | National Human Genome Research Institute training grant (T32 HG010030)
Title: <i>University of Michigan ELSI Research Training Program</i>
Role: Program Director (PI equivalent for training grants)
Project Funding: \$114,579 for FY 2018 (with additional funding from the University of Michigan) |
| 2019-21 | National Institute of Aging R03 research grant (R03 AG063222)
Title: <i>Development of Culturally-Sensitive and Patient-Centered Feedback for Alzheimer's Dementia Risk Disclosure</i>
Role: Co-Investigator (A. Rahman, PI)
Project Funding: \$156,000 for FY 2019 |
| 2018-23 | National Institute on Aging U01 grant (AG057195) |

Title: *Early Onset Alzheimer's Disease Consortium*
Role: Advisory Board member (L. Apostolova, PI)
Project Funding: \$11,820,016 for FY 2019

- 2018-23 National Institute on Aging R01 research grant (R01 AG 058468)
Title: *Aducanumab Alzheimer's Prevention Trial*
Role: Co-Investigator (E. Reiman, PI)
Project Funding: \$5,670,134 for FY 2018
- 2018-23 National Institutes of Health R01 research grant (R01 HD095068)
Title: *Unresolved Issues in Newborn Screening*
Role: Advisory Board member (B. Tarini, PI)
Project Funding: \$795,449 for FY 2018
- 2016-21 National Institute on Aging P30 Center grant (AG053760)
Title: *Michigan Alzheimer's Disease Core Center*
Role: Core Director (H. Paulson, PI)
Project Funding: \$9 million / 5 years
- 2015-19 National Institute on Aging RF1 grant (AG047866-01A1)
Title: *Impact of Disclosing Amyloid Imaging Results to Cognitively Normal Individuals*
Role: Site PI (J. Karlawish & R. C. Green, Multiple PIs)
Project Funding: \$4,646,052 / 4 years
Note: This project has two funded supplements
- 2014-19 National Institute on Aging T32 training grant (AG027708-06A1)
Title: *Interdisciplinary Research Training on Health and Aging*
Roles: Research Mentor; Executive Committee member (C. M. de Leon, PI)
Project Funding: \$700K / 5 years

Completed

- 2014-19 National Institute on Aging R01 grant (AG046906-01)
Title: *Return of Amyloid Imaging Research Results in MCI*
Role: Co-Investigator (J. Lingler, PI)
Project Funding: \$1.6 million / 5 years
- 2016-19 National Human Genome Research Institute R03 grant (HG008809)
Title: *Developing a Genomics Literacy Measure*
Role: Consultant (M. Linderman, PI)
Project Funding: \$69,916 / 2 years
- 2013-18 National Institute on Aging UF1 award (AG046150)
Title: *Alzheimer's Prevention Initiative APOE4 Trial*
Role: Co-Investigator (E. Reiman, PI)
Project Funding: \$33.3 million / 5 years
- 2013-17 National Human Genome Research Institute (NIH) UM1 (HG006508-01A1)
Clinical Sequencing Exploratory Research (CSER) Consortium
Title: *Exploring Precision Cancer Medicine for Sarcoma and Rare Cancers*
Role: Multiple Principal Investigator (with A. Chinnaiyan)
Project Funding: \$7,844,077 / 4 years

- 2013-17 National Human Genome Research Institute (NIH) UM1 (HG006508-01A1)
Clinical Sequencing Exploratory Research (CSER) Consortium
Title: *Psychosocial and Ethical Issues in Precision Oncology*
Role: Principal Investigator
Note: *This project was part of the multi-component UM1 award listed above*
- 2012-17 National Human Genome Research Institute (NIH)
U01 Award (U01 HG006500), Clinical Sequencing Exploratory Research
Title: *Integration of Whole Genome Sequencing into Clinical Medicine*
Role: Consultant (R. C. Green, PI)
Project Funding: \$11,539,189 / 4 years
- 2016-17 University of Michigan MCubed Interdisciplinary Research Program
Title: Michigan Lean Thinking for Dementia Care
Role: Multiple Principal Investigator (H. Kales & R. Dunkle, M-PIs)
Project Funding: \$60,000 / 1 year
- 2012-15 Stand Up to Cancer / Prostate Cancer Foundation
Translational Cancer Research Grant
Title: *Precision Therapy of Advanced Prostate Cancer*
Role: Co-Investigator (A. Chinnaiyan, PI)
Project Funding: \$10,000,000 / 3 years
- 2013-15 National Human Genome Research Institute (NIH) R21 (HG006575-01A1)
Title: *Assessment and Development of a Genomic Counseling Service Delivery Model for Common Disease*
Role: Co-Investigator (K. Sweet, PI)
Project Funding: \$179,000 / 2 years
- 2000-12 National Human Genome Research Institute (NIH)
(4 awards)
R01 Award (R01 HG02213), Ethical, Legal, & Social Issues Grants Program
Title: *Risk Evaluation and Education for Alzheimer's Disease*
Roles: Co-Principal Investigator, 2002-06;
Site PI, 2007-12 (R. C. Green, PI)
Project Funding: \$8,199,934 / 12 years
- 2011-14 Centers for Disease Control U48 award
Title: Michigan Genomics Academic-Practice Partnership
Role: Advisory Board member (M. Katapodi, PI)
- 2010-13 National Human Genome Research Institute (NIH)
R01 Award (R01 HG05092), Ethical, Legal, & Social Issues Grants Program
Title: *Impact of Direct-to-Consumer Genetic Testing*
Project Funding: \$1,108,876 / 3 years
Role: Multiple Principal Investigator (with R. C. Green)
- 2012-13 National Institute for Deafness and Communication Disorders
R01 Administrative Supplement (R01 DC-005053-08-S1)
Title: *Collaborative Applied Research Supplement*
Project Funding: \$40,000 / 1 year
Role: Co-Investigator (S. Camper, PI)

- 2010-13 National Institute of Diabetes and Digestive and Kidney Diseases (NIH)
R01 Award (*R01 DK083347*)
Title: *Lay Public's Genetic Explanations for Type 2 Diabetes*
Project Funding: \$1,851,032 / 3 years
Role: Co-Investigator (T. Jayaratne, PI)
- 2009-12 United States Department of Veterans Affairs
Health Services Research & Development (*IBD-09-105*)
Title: *Preparing to Discuss Genetic Test Results for Colorectal Cancer Risk*
Project Funding: \$831,000 / 3 years
Role: Co-Investigator (A. Fagerlin, PI)
- 2011-12 University of Michigan Comprehensive Cancer Center Research Grants Program
Title: *Preparing Research on Ethical and Psychosocial Issues in Cancer Genome Sequencing*
Project Funding: \$40,000 / 1 year
Role: Principal Investigator
- 2011-12 University of Pittsburgh Aging Institute Research Grants Program
Title: *Disclosing Amyloid Imaging Results in Mild Cognitive Impairment*
Role: Consultant (J. Lingler, PI)
- 2009-11 National Institute of Diabetes and Digestive and Kidney Diseases (NIH)
Planning Grant for Educational, Behavioral, or Social Studies for Translation of Genetic Factors in Common Diseases (*U34 DK084542*)
Title: *Impact of Family History & Genomics-Based Risk Profiling on Primary Care*
Project Funding: \$856,051 / 2 years
Role: Principal Investigator
- 2008-11 National Alzheimer's Association
Investigator-Initiated Research Grants Program (*IIRG-07-58189*)
Title: *Communicating Diagnostic and Risk Information in MCI*
Project Funding: \$240,000 / 3 years
Role: Principal Investigator
- 2009-11 National Institute on Aging (NIH) R03 Award (*R03 AG033860*)
Title: *Disclosing a Dementia Diagnosis*
Project Funding: \$144,236 / 2 years
Role: Consultant (B. Carpenter, PI)
- 2009-10 University of Michigan Comprehensive Cancer Center
Research Grants Program
Title: *Development and evaluation of an interactive website for women considering genetic testing for breast and ovarian cancer*
Project Funding: \$40,000 / 1 year
Role: Principal Investigator
- 2008-09 University of Michigan Ethics in Public Life Initiative
University of Michigan Comprehensive Cancer Center (co-sponsors)
Research Grants Program
Title: *Returning Individual Genetic Test Results to Research Participants in the Genetics, Environment and Melanoma Study*
Project Funding: \$27,724 / 1 year

Role: Principal Investigator

- 2008-09 University of Michigan Transportation Research Institute
Title: *New approach to assessing self-regulation by older drivers*
Project Funding: \$100,000 / 1 year
Role: Co-Investigator (D. Eby, PI)
- 2008 United States Veterans Administration
Title: *Health Services Research in Genomics*
Project Funding: \$144,670 / 1 year
Role: Co-Investigator (B. Zikmund-Fisher, PI)
- 2007-08 University of Michigan Center for Health Communications Research
Developmental Project Award
Title: *Development of an Interactive Website to Provide Tailored Education and Risk Communication to Women at High Risk of Breast Cancer*
Project Funding: \$50,000 / 1 year
Role: Principal Investigator
- 2006-08 National Institute on Aging (NIH) R03 Award (R03 AG025914)
Title: *Disclosure of Genetic Risk for Alzheimer's Disease*
Project Funding: \$149,075 / 2 years
Role: Principal Investigator
- 2005-06 National Institute on Aging (NIH)
Boston University Alzheimer's Disease Core Center (P50 AG13846)
Title: *Education and Information Transfer Core*
Role: Co-Director, Education & Information Transfer Core (N. Kowall, PI)
- 2003-05 Charles Farnsworth Trust, Boston Medical Foundation
Elder Health Care Research Award
Title: *Evaluation of an Education and Support Program for Early-Stage Patients with Alzheimer's Disease and Their Care-Partners*
Project Funding: \$80,000 / 2 years
Role: Principal Investigator
- 2003-05 National Human Genome Research Institute (NIH)
Research Supplement for Underrepresented Minorities (D. Akinleye, fellow)
Title: *Risk Evaluation and Education for Alzheimer's Disease*
Role: Research Mentor
- 2002-04 Alzheimer's Association New Investigator Research Grant (NIRG-02-3712)
Title: *Differences in Illness Perceptions among African American and White Caregivers and Relatives of People with Alzheimer's Disease*
Project Funding: \$100,000 / 2 years
Role: Principal Investigator
- 2001-02 Boston University Alzheimer's Disease Center Pilot Grant Award
Title: *Perceptions of Alzheimer's Disease and Emerging Care Options*
Project Funding: \$30,000 / 1 year
Role: Principal Investigator
- 1998 Michigan Center for Applied Cognitive Research on Aging Seed Grant

Title: *Attitudes toward Predictive Genetic Testing for Alzheimer's Disease*
Role: Principal Investigator

Grants in submission

- 2019-24 National Institutes of Health U54 Rare Diseases Clinical Research consortium
Title: *Nephrotic Syndrome Rare Disease Clinical Research Network III*
Role: Co-Investigator (M. Kretzler, PI)
- 2019-24 National Human Genome Research Institute U24 center grant
Title: *CERA: Preserving the Legacy and Promoting the Future of ELSI Scholarship*
Role: Co-Investigator (B. Tarini, PI)
- 2019-24 National Cancer Institute U01 "Cancer Moonshot" research grant
Title: *Innovative Approaches to Expand Cancer Genetic Screening and Testing for Patients & Families in a Statewide Oncology Network through Community, State, & Payer Partnerships*
Role: Co-Investigator (E. Stoffel, J. Griggs, K. Resnicow, M-PIs)
- 2019-24 National Institute on Aging R01 research grant
Title: *Impacts of Receiving Alzheimer's Disease Genetic Risk Information among Latinos in Northern Manhattan*
Role: Site PI / Co-investigator (R. Ottman, PI)
- 2019-21 National Institute on Aging R21 research grant
Title: *A Scalable Model for Promoting Functioning and Well-Being among Older Adults with Mild Cognitive Impairment via Meaningful Social Interactions*
Role: Co-Investigator (J. Piette, PI)

PRESENTATIONS

Selected Conference Presentations

1. **Roberts, J. S.**, Welsh-Bohmer, K., Green, R. C., & Karlawish, J. (2017, July). Assessing the impact of disclosing amyloid imaging results to cognitively normal older adults: The REVEAL-SCAN study. Poster presentation, annual Alzheimer's Association International Conference, London.
2. **Roberts, J. S.** (2017, June). Disclosing genetic and neuroimaging risk information in the new era of Alzheimer's prevention trials: Practical and ethical challenges and opportunities. Oral presentation, 4th ELSI Congress, Jackson Laboratories, Farmington, Connecticut.
3. **Roberts, J. S.** (2016, May). Consumer perspectives on receiving personal genomic services. Oral presentation, annual meeting of the European Society of Human Genetics, Barcelona, Spain.
4. **Roberts, J.S.**, Bharadwaj, A., Diamond, P., et al. (2016, March). Patient understanding and satisfaction regarding clinical use of whole-genome sequencing. Poster presentation, annual meeting of the American College of Medical Genetics, Tampa, FL.
5. **Roberts, J. S.** (2015, July). Applying lessons learned from the REVEAL Study to genetic testing and counseling in Alzheimer's prevention research. Oral presentation, annual Alzheimer's Association International Conference, Washington DC.

6. **Roberts, J. S.** (2015, April). Giving patients incidental information from genomic sequencing. Oral presentation, annual meeting of the Society of Behavioral Medicine, San Antonio, Texas.
7. **Roberts, J. S.** (2014, July). Direct-to-consumer genetic testing for risk of Alzheimer's disease. Oral presentation at Alzheimer's Association International Conference, Copenhagen, Denmark.
8. **Roberts, J. S.** (2013, November). What findings should be offered to research participants whose data and specimens are archived in biobanks? Brocher Foundation workshop on "Returning Genetic Results in Biobanks: Opening an International Dialogue," Hermance, Switzerland.
9. Green, R.C., **Roberts, J. S.**, Hartz, S., & Kohane, I. (2013, October). Return of results in genetics: Integration with clinical care. Panel presentation, World Congress of Psychiatric Genetics, Boston, MA.
10. Bloss, C., Kaufman, D., Lee, S., & **Roberts, J. S.** (chair). (2012, November). Surveying customer responses to personal genetic (including DTC) services. Panel presentation, annual meeting of the American Society of Human Genetics, San Francisco, CA.
11. Green, R. C., Lingler, J., Karlawish, J., & **Roberts, J. S.** (chair). (2012, July). Disclosing risk information to individuals at imminent risk of Alzheimer's disease. Alzheimer's Association International Conference, Vancouver, Canada.
12. **Roberts, J. S.**, Gordon, E., & Green, R. C. (2011, April). Empirical research to inform practice and policy in personal genomics. Panel presentation, International ELSI Research Congress, Chapel Hill, North Carolina.
13. **Roberts, J. S.** (2010, November). Ten Years of the REVEAL Study: Informing the Bigger Picture of Genetic Susceptibility Risk Disclosure, Personal Genomics, and Ethical, Legal, and Social Issues in Genetics and Genomics. Panel presentation, annual meeting of the American Society of Human Genetics, Washington DC.
14. **Roberts, J. S.** (2010, October). Is the hope of benefit from personal genomic information materializing? Empirical results from patients, research participants, and the wider public. Panel presentation, annual meeting of the American Society for Bioethics and the Humanities, San Diego, California.
15. **Roberts, J. S.**, Milliron, K., Hoskins, K., & Merajver, S. (2010, April). Development and evaluation of a personalized, web-based decision aid for women considering genetic testing for risk of breast cancer. Meeting of the Centers of Excellence in Cancer Communications Research, Philadelphia, Pennsylvania.
16. **Roberts, J. S.**, & Christensen, K. D. (2009, November). Genetic susceptibility testing for individuals at risk for Alzheimer's disease: Findings from the REVEAL Study. Annual meeting of the Gerontological Society of America, Atlanta, Georgia.
17. **Roberts, J. S.** (2009, September). Genetic susceptibility testing for common diseases: Findings from the REVEAL (Risk Evaluation & Education for Alzheimer's Disease) Study. 5th International DNA Sampling Conference: The Age of Personalized Genomics, Banff, Alberta, Canada.

18. **Roberts, J. S.** (2009, July). The ethical implications of genetic susceptibility testing for individuals at risk for Alzheimer's disease. 13th Genetics & Ethics in the 21st Century Conference, Breckenridge, Colorado.
19. **Roberts, J. S.,** Uhlmann, W., Petersen, R., Karlawish, J., & Green, R. C. (2009, July). Clinical practices Regarding Mild Cognitive Impairment (MCI) among neurology service providers. 12th International Conference on Alzheimer's Disease (ICAD), Vienna, Austria.
20. **Roberts, J. S.,** et al. (2008, July). Psychological and behavioral impact of a condensed protocol for disclosing APOE genotype and risk information for Alzheimer's disease: The REVEAL Study. 11th International Conference on Alzheimer's Disease and related Disorders, Chicago, Illinois.
21. **Roberts, J. S.** (2008, May). Methodological challenges in assessing the process and impact of genetic susceptibility testing. International Conference on Ethical, Legal, and Social Implications of Genomics, Cleveland, Ohio.
22. **Roberts, J. S.,** et al. (2007, November). Age group differences in response to genetic risk assessment for Alzheimer's disease. Annual meeting of the Gerontological Society of America, San Francisco, California.
23. **Roberts, J. S.,** & Silverio, E. (2006, September). An education and support program for early stage Alzheimer's disease. 14th annual Alzheimer's Association Dementia Care Conference, Atlanta, Georgia.
24. **Roberts, J. S.** et al. (2006, July). Age group differences in response to genetic risk assessment for Alzheimer's disease. 10th biannual International Conference on Alzheimer's Disease and Related Disorders, Madrid, Spain
25. **Roberts, J. S.,** & Green, R. C. (2005, November). Genetic research in dementia: Risk evaluation and education for Alzheimer's disease. Annual meeting of the Gerontological Society of America, Orlando, Florida.
26. **Roberts, J. S.,** & Silverio, E. (2005, July). Education and support in Alzheimer's early stages. 13th annual Alzheimer's Association Dementia Care Conference, Chicago, Illinois.
27. **Roberts, J. S.,** Akinleye, D., Hipps, Y. G., & Green, R. C. (2005, June). Beliefs about prevention and treatment of Alzheimer's disease among African Americans. Alzheimer's Association International Conference on Prevention of Dementia, Washington DC.
28. **Roberts, J. S.** (2005, March). Genetic risk assessment for adult children of people with Alzheimer's disease. "Children of Alzheimer's Parents" workshop, annual meeting of the American Association for Geriatric Psychiatry, San Diego, California.
29. **Roberts, J. S.,** Lock, M., Prest, J., Barber, M., Whitehouse, P., LaRusse, S., Relkin, N., Brown, T., & Green, R. C. (2004, July). How does genetic testing affect anxiety about developing Alzheimer's disease? 9th International Conference on Alzheimer's Disease and Related Disorders, Philadelphia, Pennsylvania.
30. **Roberts, J.S.,** Akinleye, D., & Hipps, Y. G. (2004, July). Cultural differences in perceptions of Alzheimer's disease. Alzheimer's Association 12th National Alzheimer's Disease Education Conference, Philadelphia, Pennsylvania.

31. **Roberts, J.S.**, Silverio, E., & Kurra, C. (2004, July). Evaluation of an education and support program for early-stage patients with Alzheimer's disease and their care-partners. Alzheimer's Association 12th National Alzheimer's Disease Education Conference, Philadelphia, Pennsylvania.
32. **Roberts, J. S.**, Zick, C. Mathews, C., & Green, R. C. (2004, April). Genetic testing for Alzheimer's disease and its impact on insurance behavior. Annual meeting of the American Academy of Neurology, San Francisco, California.
33. **Roberts, J.S.**, Green, R. C., Relkin, N., Whitehouse, P., Brown, T., Barber, M., LaRusse, S., Katzen, H., Post, S., Sadovnick, D., Quaid, K. (2003, April). How do participants rate the impact of genetic susceptibility testing for Alzheimer's disease?. Annual meeting of the American Academy of Neurology, Honolulu, Hawaii.
34. **Roberts, J.S.**, Green, R. C., Relkin, N., Whitehouse, P., Brown, T., Barber, M., LaRusse, S., Post, S., Sadovnick, D., & Ravdin, L. (2002, July). Genetic susceptibility testing for adult children of patients with Alzheimer's disease: Participants' self-reports on the impact of risk assessment. 8th International Conference on Alzheimer's Disease and Related Disorders, Stockholm, Sweden.
35. **Roberts, J.S.**, Relkin, N., Whitehouse, P. J., Brown, T. C., Barber, M., LaRusse, S., & Green, R. C. (2002, April). Reasons for pursuing genetic susceptibility testing for Alzheimer's disease: Preliminary findings from the REVEAL Study. Annual meeting of the American Academy of Neurology, Denver, Colorado.
36. **Roberts, J.S.**, Kales, H.C., Blow, F.C., Connell, C.M., Visnic, S., & Mellow, A.M. (2001, February). Differences in psychological symptoms and caregiver burden among caregivers of older patients with depression, dementia, or coexisting dementia and depression. Annual meeting of the American Association for Geriatric Psychiatry, San Francisco, California.
37. **Roberts, J.S.**, Booth, B. M., Blow, F. C., & Bingham, R. C. (2000, November). Age group and psychiatric comorbidity as predictors of health-related quality of life: A 12-month study of medically hospitalized male veterans. Annual meeting of the Gerontological Society of America, Washington, DC.
38. **Roberts, J.S.** (1999, November). Anticipating response to predictive genetic testing for Alzheimer's disease. Annual meeting of the Gerontological Society of America, San Francisco, California.
39. **Roberts, J.S.**, Welsh-Bohmer, K. A., & Simons, M. I. (1998, August). Psychological issues in predictive testing for Alzheimer's disease. Annual meeting of the American Psychological Association, San Francisco, California.

Invited Oral Presentations

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| 2019 | <u>What's new in Alzheimer's research?</u> Presented as part of the Alzheimer's Association's (Greater Michigan chapter) "Research Night" series, Petoskey, MI. |
| 2019 | <u>Negotiation and faculty positions</u> . Panel discussion at National Human Genome Research Institute Annual Research Training and Career Development Meeting, St, Louis, MO |

- 2019 Disclosing genetic and biomarker information to individuals at risk for Alzheimer's disease. Presented at Michigan Center for Contextual Factors in Alzheimer's Disease.
- 2018 What's new in Alzheimer's research? Presented at Alzheimer's Caregiver Expo, Port Huron, MI
- 2018 Research Education Component of the Michigan Alzheimer's Disease Center. Biannual meeting of the NIA-funded national Alzheimer's Disease Centers program, Atlanta, GA.
- 2018 The promise and perils of precision health for cancer. Discussion panel as part of the 8th annual symposium of the Michigan Medicine-Peking University Health Science Center Joint Institute for Translational and Clinical Research, Ann Arbor MI
- 2018 Genetics and genomics research ethics. Guest lecture, "Legal Rules and Ethical Issues for Clinical Research" course (HMP 540), University of Michigan School of Public Health.
- 2018 Disclosure of APOE genotype and risk of Alzheimer's disease. 11th biennial Barcelona-Pittsburgh Conference on Alzheimer's Disease, Barcelona, Spain
- 2018 Alzheimer's Disease: Research Update. 22nd Annual Alzheimer's Association (Northwest Ohio Chapter) Professional Education Conference, Toledo, OH
- 2018 Assessing the impact of genetic susceptibility testing: Where do we go from here? Invited presentation, "Looking for the Psychosocial Impacts of Genomic Information," Columbia University, New York, NY
- 2018 Communicating genetic risk information. Invited presentation, Michigan Medicine Health Literacy and Patient Education Forum, Ann Arbor, MI
- 2018 What's new in Alzheimer's research? Public talk sponsored by the Alzheimer's Association (Greater Michigan Chapter), Southfield, MI
- 2018 Genetic testing for Alzheimer's disease. Podcast recorded for Michigan Alzheimer's Disease Center, Research Education Core series for postdoctoral trainees
- 2013-18 Challenges and controversies in direct-to-consumer genetic testing. Guest lecture, "Ethical Dilemmas in Health for Social Workers and Other Health Professionals" course (SW 705), University of Michigan School of Social Work.
- 2016-18 Genetic susceptibility testing for Alzheimer's disease. UM School of Public Health T32 Program in Public Health and Aging seminar series.
- 2017 Alzheimer's disease: An Overview. Seminar given to Corner Health Youth Leadership Council, Ypsilanti, MI

- 2017 What to do when participants bring their 23andMe results to their Alzheimer's Disease Center visit. Oral presentation, biannual national meeting of NIH-funded Alzheimer's Disease Centers, San Diego, CA.
- 2017 Finding common ground: A conversation on genetics and religion. Panel discussion, U-M Genetics & Faith public dialogue, Ann Arbor Public Library.
- 2017 Alive Inside film screening and dementia expert Q & A session. Panel discussion, Oakland University, Rochester MI.
- 2017 Disclosing biomarker information to individuals at elevated risk for Alzheimer's disease. Invited lecture, Michigan Alzheimer's Disease Center annual research symposium.
- 2017 Risk and protective factors for Alzheimer's disease. Invited lecture, Michigan Alzheimer's Disease Center annual Health & Aging Fair.
- 2017 Opportunities and challenges in precision oncology. Webinar for health professionals, hosted by the Michigan Department of Health & Human Services and the Michigan Cancer Genetics Alliance.
- 2014-17 Alzheimer's disease from a public health perspective. Guest lecture, "Memory, Aging and Expressive Arts" course (ARTDES 501 / SW 513), University of Michigan School of Art & Design.
- 2017 Patient and provider perspectives on precision oncology. Biannual meeting of the NIH Clinical Sequencing Exploratory Research (CSER) Consortium, Bethesda, MD.
- 2016 Ethical considerations in the implementation of precision medicine. National Cancer Institute / American Society for Radiation Oncology workshop on "Precision Medicine in Radiation Oncology," NIH Campus, Bethesda, MD.
- 2016 Emerging approaches in treatment and prevention of Alzheimer's disease: Ethical and practical challenges. Wayne State University Institute of Gerontology's 29th annual "Issues in Aging" conference.
- 2016 The impact of disclosing personal genomic information. University of Iowa Genetics PhD program speaker series.
- 2016 Research findings on the impact of returning genetic test results. Food & Drug Administration workshop on "Patient and Medical Professional Perspectives on the Return of Genetic Test Results," Bethesda MD
- 2016 Ethics of health behavior interventions. Guest lecture, doctoral seminar on health behavior interventions (HBHE 886), University of Michigan School of Public Health
- 2016 Mild Cognitive Impairment (MCI) in clinical practice. Meeting on "Developing Recommendations for Communicating Amyloid Imaging Results in MCI," University of California-Irvine.

- 2016 Ethical issues in research with children. Guest lecture, University of Michigan T32 Fellows' Seminar (Developmental Science to Improve Child Health).
- 2015 Bring your genes to Cal: How an educational program was drawn into the public debate over genetic testing. Panelist (with President Mark Schlissel and Professor David Ginsburg), University of Michigan Department of Human Genetics seminar series.
- 2015 Genetic susceptibility testing for Alzheimer's disease. Guest lecture, Alpha Epsilon Delta chapter, University of Michigan.
- 2015 How might genomic sequencing impact parents and children? UCSF-sponsored meeting on "Sequencing Newborns: Ethics & Policy," Hastings Center, New York.
- 2015 Ethical considerations in using biomarkers as an enrichment strategy. Session of the Clinical Trials Advancement Professional Interest Area, annual meeting of the Alzheimer's Association International Conference, Washington DC.
- 2015 Overview of Michigan Alzheimer's Disease Center. Meeting of the Mission Commission on Services to the Aging, Michigan Department of Health & Human Services.
- 2015 A conversation on the genomic revolution / evolution. Annual meeting, Institute for Functional Medicine, Austin TX.
- 2015 Genetic testing: Implications for health, illness and society. Guest lecture. "Health, Illness & Society" course (PSYCH 211), University of Michigan Department of Psychology.
- 2015 What makes clinical and public health research ethical? Guest lecture. "Advanced Research Methods" course (HBHE 621), University of Michigan School of Public Health.
- 2015 Understanding prevention trials. Webinar for Certificate Program in Advanced Clinical Dementia Practice, University of Michigan School of Social Work.
- 2015 Genetics, biomarkers and predictive testing. Webinar for Certificate Program in Advanced Clinical Dementia Practice, University of Michigan School of Social Work.
- 2014 Current research in Alzheimer's disease. Annual conference for dementia caregivers, Monroe Center for Healthy Aging, Monroe MI.
- 2014 MCI and risk / protective factors for dementia. Guest speaker, Living Well with Memory Loss program lecture series, Michigan Alzheimer's Disease Center.
- 2014 What's new in Alzheimer's disease? Guest speaker, Science Café community event sponsored by University of Michigan Museum of Natural History.
- 2013 Integrating whole-genome sequencing into clinical practice: Ethical and practical issues. Michigan State Medical Society 17th Annual Conference on Bioethics, Ann Arbor, Michigan.

- 2013 Communicating genetic testing and amyloid imaging results to individuals at risk for Alzheimer's disease: Ethical and practical challenges. Memory & Aging Center seminar series, University of California-San Francisco.
- 2013 Using biomarkers to disclose risk information for Alzheimer's disease: Ethical and psychosocial implications. Invited lecture, Vanderbilt Memory & Alzheimer's Center, Vanderbilt University Medical Center.
- 2013 Challenges and controversies in direct-to-consumer genetic testing. Guest lecture, "Public Health Ethics" course, University of California-Berkeley School of Public Health.
- 2013 Challenges and controversies in direct-to-consumer genetic testing. Guest lecture, Medical Genetics course, University of California-San Francisco School of Nursing.
- 2012 Ethical issues in genetics and genomics research. Guest lecture, "Legal Rules and Ethical Issues for Clinical Research" course (HMP 540), University of Michigan School of Public Health.
- 2012 Genetics, ethics and biobanking. Panel presentation on "Biobanking: Uses and Dilemmas," University of Michigan, Flint campus.
- 2012 Genetics in the media and popular culture. Michigan Association of Genetic Counselors annual meeting, Lansing, Michigan.
- 2012 Using biomarkers to disclose risk information for Alzheimer's disease: Ethical and psychosocial implications. University of Pittsburgh Alzheimer's Disease Research Center seminar series, Pittsburgh, Pennsylvania.
- 2012 Ethical issues in public health practice and research. Future Public Health Leaders summer program, University of Michigan School of Public Health.
- 2012 More than senior moments: Understanding early signs of memory changes. Ann Arbor District Library, Ann Arbor, Michigan.
- 2012 Current topics in Alzheimer's disease. Ann Arbor Rotary Club, Ann Arbor, MI.
- 2012 Genetic risk assessment for Alzheimer's disease. Baycrest Annual Conference on Cognitive Neuroscience, Toronto, Ontario, Canada.
- 2012 Ethics of personalized medicine. Mega Grand Rounds: Personalized Medicine. University of Michigan School of Dentistry.
- 2012 Disclosing diagnostic and risk information: How, when and to whom? 10th Annual Mild Cognitive Impairment Symposium and 1st Early Alzheimer's Diagnostic Workshop. Mount Sinai Medical Center of Miami Beach.
- 2011 Estimating and communicating risk of developing Alzheimer's disease. University of Michigan School of Public Health T32 Program in Public Health and Aging seminar series.

- 2011 Mild cognitive impairment. University of Michigan Alzheimer's Disease Center Community Appreciation Luncheon.
- 2011 Health education and genetic counseling for individuals with mild cognitive impairment. University of Michigan School of Public Health T32 Program in Public Health and Aging seminar series.
- 2011 Examining the benefits and harms of direct-to-consumer genetic testing. University of Michigan Center for Bioethics & Social Sciences in Medicine seminar series.
- 2011 Using CTools to Enhance Interactive Teaching. University of Michigan Teaching and Technology Collaborative, "Enriching Scholarship" conference.
- 2011 Examining the impact of direct-to-consumer personal genomics services. University of Michigan "Genomics and ELSI" lecture series.
- 2011 The impact of disclosing genetic risk information for Alzheimer's disease: Findings from the REVEAL Study. University of Michigan School of Public Health T32 Program in Public Health and Aging seminar series.
- 2010 Emerging health policy issues raised by genetic testing. University of Michigan School of Public Health, Executive Master's program, "Introduction to Health Policy" course.
- 2010 When do investigators need to share individual research results? Ethical and practical issues in disclosing genetic risk information to participants. University of Michigan Department of Psychology, Developmental Psychology Lecture Series.
- 2010 Disclosure of genetic research results: Practical and ethical issues. University of Michigan Medical School, Bioethics Grand Rounds.
- 2010 Psychological and behavioral responses to genetic risk assessment for common, complex diseases. University of Florida College of Health and Human Performance, Health Education and Behavior Seminar Series.
- 2010 Implications of personal genomics for cardiovascular medicine. University of Michigan Medical School, Cardiovascular Medicine Grand Rounds in the Division of Cardiovascular Medicine.
- 2010 Disclosing diagnostic and risk information about Alzheimer's disease. 9th Schreiber Memorial Conference, Alzheimer's Association Michigan Chapter, Kalamazoo, Michigan.
- 2009 Disclosing diagnostic and risk information for Alzheimer's disease: Perspectives of family members and providers. Michigan Alzheimer's Disease Research Center Community Conference on Advances in Prevention, Diagnosis and Treatment of AD. Ypsilanti, Michigan.
- 2009 Development of a tailored, web-based decision aid for women considering genetic testing for risk of breast cancer. Wayne State University, Karmanos Cancer Institute Clinical Cancer Genetics Monthly Conference.

- 2009 Ethical and practical issues in returning genetic research results. Guest lecture, University of Michigan Medical School, Department of Internal Medicine, "Clinical Research involving Special Populations" seminar.
- 2009 Understanding the process and impact of genetic susceptibility testing for Alzheimer's disease. University of Michigan Medical School, Genetic Counseling Program research seminar.
- 2009 Ethical and practical issues in returning genetic research results to participants. University of Michigan School of Medicine Bioethics Program lecture series.
- 2008 Ethical and practice issues raised by emerging cancer genetics technologies. Michigan Cancer Consortium annual meeting.
- 2007-08 Ethics and academic integrity within the School of Public Health. University of Michigan School of Public Health, Fall schoolwide orientation.
- 2008 Development of an interactive website to provide tailored education & risk communication to women at high risk of breast cancer. University of Michigan Center for Health Communication Research lecture series.
- 2008 Developing and evaluating genetic risk assessment programs. University of Michigan Medical School, Robert Wood Johnson Clinical Scholars Program.
- 2008 Advances in Alzheimer's research. 8th Schreiber Memorial Conference, Alzheimer's Association Michigan Chapter, Kalamazoo, Michigan.
- 2008 Understanding the process and impact of genetic susceptibility testing for common, complex conditions. Michigan Cancer Genetic Alliance meeting.
- 2008 Understanding the process and impact of genetic susceptibility testing for Alzheimer's disease. University of Michigan Medical School, Genetic Counseling Program research seminar.
- 2007 Understanding the process and impact of genetic susceptibility testing for Alzheimer's disease. University of Michigan Medical School, Geriatric Medicine Clinical Conference.
- 2007 Understanding the psychological and behavioral impact of genetic susceptibility testing for Alzheimer's disease. University of Michigan Medical School, Department of Human Genetics seminar series.
- 2006-07 Mental health and aging. Guest lecture, "Mental Health Policy in the United States" graduate level course, University of Michigan School of Public Health.
- 2007 Examining the process and impact of genetic susceptibility testing. University of Michigan Comprehensive Cancer Center Quality of Life Workgroup lecture series.
- 2007 Genetic testing for people at-risk for Alzheimer's disease: Findings from the REVEAL Study. Michigan Dementia Coalition, Howell, Michigan.
- 2007 Genetic susceptibility testing for adult-onset disorders. University of Michigan Center for Health Communication Research lecture series.

- 2006 Ethical issues in genetic testing. University of Michigan School of Medicine Bioethics Program lecture series.
- 2006 Genetic risk assessment for Alzheimer's disease. University of Michigan Health & Retirement Study lecture series.
- 2006 Genetic research in Alzheimer's disease: Implications for clinical practice. West Suburban Area Boston Alzheimer's Partnership lecture series.
- 2005 Genetic susceptibility testing for Alzheimer's disease. Guest lecture, "Genetics, Law, and Public Health" graduate level course, Boston University School of Public Health.
- 2005 Cultural differences in perceptions of dementia. Boston University Alzheimer's Disease Center Research Seminar Series.
- 2005 Psychological issues in genetic testing. NIA-funded "Aging, Genetic Technology, and the Future" conference, Boston College.
- 2005 Racial and ethnic differences in perceptions of dementia. Psychiatry Grand Rounds lecture series, Bedford (MA) VA Medical Center.
- 2004 Genetic risk assessment for individuals at risk for Alzheimer's disease. Psychology Lecture Series, Brockton VA Medical Center, Brockton, Massachusetts.
- 2004 Genetic risk assessment for Alzheimer's disease. University of Massachusetts-Boston Gerontology Institute speaker series.
- 2003 The REVEAL Study: Genetic testing for sporadic Alzheimer's disease. 3rd Genetics Update Workshop, Genetics Program for Nursing Faculty, Cincinnati Children's Hospital Medical Center.
- 2003 Genetic risk assessment for Alzheimer's disease. Alzheimer's Association Western Massachusetts Chapter annual Regional Professional Conference.
- 2001 Research update: 8th International Conference on Alzheimer's Disease and Related Disorders. Boston University Alzheimer's Disease Center Research Seminar series.
- 2002 Psychological research and Alzheimer's disease: An overview and recent trends. Geropsychology Lecture Series, Brockton VA Medical Center, Brockton, Massachusetts.
- 2001 Early detection and understanding of Alzheimer's disease. Boston University Gerontology Center's Annual Summer Institute.
- 2002 Working with patients with personality disorders in the nursing home. Brockton VA Medical Center Nursing Home Care Unit, Brockton, Massachusetts.
- 1999 Caring for a loved one with memory problems. Senior Interest Lecture Series, Brockton VA Medical Center, Brockton, Massachusetts.

- 2000 First-degree relatives' attitudes and beliefs about Alzheimer disease and genetic testing. Boston University Alzheimer's Disease Center Research Seminar series.
- 1999 Addiction, identity, and the life story. Guest lecture, University of Michigan undergraduate psychology seminar on substance abuse research, Ann Arbor, Michigan.
- 1997 Predictive testing for Alzheimer's disease: A psychological study of decision-making for at-risk individuals. University of Michigan Alzheimer's Disease Research Center Clinical Core lecture series.

THESES

- 2001 Medical decision-making in individuals at-risk for Alzheimer's disease: The case of predictive testing. Unpublished doctoral dissertation, University of Michigan. Dissertation Chair: Christopher Peterson, Ph.D.
- 1996 On the psychology of upward social mobility: A multiple-case study of first-generation college students. Unpublished master's thesis, University of Michigan. Thesis Advisor: George Rosenwald, Ph.D.

TEACHING & MENTORING

Courses Taught

- 2018-present Ethical, legal & social issues in genomics and health. Instructor for graduate level course (HBHE 715). University of Michigan School of Public Health, Department of Health Behavior & Health Education. This is a yearlong seminar.
- 2007-present Issues in public health ethics. Instructor for graduate level course (HBHE 610), University of Michigan School of Public Health, Department of Health Behavior & Health Education. This course is offered annually.
- 2006-present Genetics, health behavior, and health education. Instructor for graduate level course (HBHE 669), University of Michigan School of Public Health, Department of Health Behavior & Health Education. This course is offered annually.
- 2015 Mental health interventions. Instructor for graduate level course (HBHE 710), University of Michigan School of Public Health, Department of Health Behavior & Health Education.
- 2014-15 Issues in public health ethics (online version). Instructor for graduate level course (HBHE 610), University of Michigan School of Public Health, Department of Health Behavior & Health Education. Course offered as part of Executive Master's Program in Health Management & Policy.
- 2012-13 Readings in genetics, health behavior and health education. Instructor for graduate level seminar (HBHE 644), University of Michigan School of Public Health, Department of Health Behavior & Health Education.
- 2005 Mental health and aging. Instructor for graduate seminar in Gerontology doctoral program, University of Massachusetts-Boston.
- 1993-95 Introduction to psychopathology. Department of Psychology, University of Michigan. Graduate Student Instructor for five semesters.

Student Advising & Mentoring

- 2018-present Research advisor (Sara Feldman, PhD student), University of Michigan School of Public Health doctoral program in Health Behavior & Health Education
- 2018-19 Thesis advisor, University of Michigan Genetic Counseling Program (Adelyn Beil, master's student)
- 2016-18 Thesis advisor, University of Michigan Genetic Counseling Program (Jamie Love-Nichols, master's student)
- 2014-18 Dissertation committee member, University of Michigan School of Nursing (Cheryl Lee, doctoral student)
- 2016-17 Thesis committee member, University of Michigan Genetic Counseling Program (Tessa Marzulla, master's student)
- 2013-16 Dissertation committee chair (Jenny Ostergren, PhD candidate), University of Michigan School of Public Health doctoral program in Health Behavior & Health Education
- 2014-16 Thesis advisor, University of Michigan Genetic Counseling Program (Diane Koeller, master's student)
- 2014-16 Research mentor (Kayte Spector-Bagdady, JD, MBE), University of Michigan Center for Bioethics and Social Sciences in Medicine postdoctoral fellowship program
- 2013-15 Research mentor (Michele Gornick, PhD), University of Michigan Center for Bioethics and Social Sciences in Medicine postdoctoral fellowship program
- 2013-15 Research mentor (Janet Childerhose, PhD), University of Michigan Center for Bioethics and Social Sciences in Medicine postdoctoral fellowship program
- 2013-14 Dissertation committee member, University of Massachusetts-Boston Department of Gerontology (Judith Poey, doctoral candidate)
- 2010-12 Dissertation committee chair, University of Michigan School of Public Health doctoral program in Health Behavior & Health Education (Kurt Christensen, doctoral candidate)
- 2008- Faculty Advisor, University of Michigan dual degree program in Public Health and Genetic Counseling (11 MS/MPH students; 4 current)
- 2006- Faculty Advisor, University of Michigan School of Public Health doctoral program in Health Behavior & Health Education (4 PhD students)
- 2006- Faculty Advisor, University of Michigan School of Public Health master's program in Health Behavior & Health Education (39 MPH students; 10 current)
- 2007- Summer Field Placement Preceptor, University of Michigan School of Public Health master's program in Health Behavior & Health Education (7 MPH students)

- 2006- Independent Study Advisor, University of Michigan School of Public Health master's program in Health Behavior & Health Education (40 MPH students, 1 undergraduate)
- 2008-11 Dissertation committee member, Boston University School of Public Health (Barbara Lerner, doctoral candidate)
- 2010-11 Thesis advisor, University of Michigan Genetic Counseling Program (Barbara Hamlington, master's student)
- 2010-11 Thesis committee member, University of Michigan Genetic Counseling Program (Rebecca Frysinger, master's student)
- 2009-10 Thesis advisor, University of Michigan Genetic Counseling Program (Maureen Mork, master's student)
- 2008-09 Dissertation committee member, Boston University School of Public Health (Jacqueline Vernarelli, doctoral student)
- 2008-09 Thesis advisor, University of Michigan Genetic Counseling Program (Jessica Szymaniak, master's student)
- 2007-08 Thesis committee member, University of Michigan Genetic Counseling Program (Katie Hock, master's student)
- 2006-07 Thesis advisor, University of Michigan Genetic Counseling Program (Karen Wain, master's student)
- 2006-07 Dissertation committee member, University of Michigan School of Public Health doctoral program in Health Behavior & Health Education (Sara McLaughlin, doctoral student)
- 2002-06 Mentor, Boston University Alzheimer's Disease Center summer student internship program (3 medical students, 2 undergraduates, 1 master's in public policy student)
- 2003-05 Primary Mentor, NIH Research Supplement for Underrepresented Minorities (Dapo Akinleye, Boston University School of Public Health master's student)
- 2003-04 Master's thesis committee member, Boston University School of Public Health (Jaza Marina, MD; master's student)
- 2003-04 Master's thesis committee member, Boston University School of Public Health (Serena Chao, MD; master's student)

CLINICAL EXPERIENCE

- 2006-08 Licensed as clinical psychologist in state of Michigan
- 2005-2006 Student Support Counselor, Boston University School of Medicine Graduate Medical Sciences Program
- 2004-2006 Support group co-leader, Early Stage Program, Alzheimer's Association Massachusetts chapter

- 2002-2006 Licensed as clinical psychologist and health service provider in the states of Massachusetts and Rhode Island
- 2000-2001 Geropsychology Postdoctoral Fellowship, Brockton (MA) VA Medical Center. Provided psychological services to older veterans seen in inpatient psychiatry, nursing home, and specialty outpatient geriatric mental health settings. Supervisors: Michele Karel, Ph.D.; Jennifer Moye, Ph.D.
- 1999-2000 Postdoctoral Training, Ann Arbor (MI) VA Medical Center (Outpatient Psychiatry and PTSD clinic). Supervisor: Richard Amdur, Ph.D.
- 1997-98 Advanced Practicum, St. Joseph's Mercy Hospital (Outpatient Mental Health), Ann Arbor, MI. Supervisor: Daniel Greenberg, Ph.D.
- 1996-97 Internship, Durham (NC) VA Medical Center. Full-time, APA-approved internship with rotations in geropsychology, inpatient psychiatry, outpatient mental health, and medical psychology. Director of Training: Jill Hazlett, Psy.D.
- 1995-96 Clinical Practicum, Mood and Anxiety Disorders Clinic, University of Michigan Medical Center. Supervisor: James Coyne, Ph.D.
- 1995 Clinical Practicum, University of Michigan Psychological Clinic. Supervisor: James Hansell, Ph.D.
- 1994 Neuropsychology Practicum, Ann Arbor (MI) VA Medical Center. Supervisor: Linas Bieliauskas, Ph.D.

OTHER RESEARCH EXPERIENCE

- 2019-present External Advisory board member, national Alzheimer's Clinical Trials Consortium (Paul Aisen, Ron Petersen, Reisa Sperling, M-PIs)
- 2017-present Disclosure Monitoring Advisory Group member, Banner Alzheimer's Institute / Novartis Alzheimer's disease prevention trial
- 2015-19 Genetic Disclosure Monitoring Committee member, Banner Alzheimer's Institute Generation Study clinical trial (Eric Reiman & Pierre Tariot, M-PIs)
- 2017-18 Co-Investigator, Planning Committee, Imaging Dementia—Evidence for Amyloid Scanning (NEW IDEAS) Study (Gil Rabinovici, PI)
- 2015-18 Cardiovascular Health Improvement Project Biobank Medical Findings Committee member (Cristen Willer, PI)
- 2014-17 Data Safety & Monitoring Board member, R01 project *RCT to Improve Oncology Nurses' Protective Equipment Use* (Christopher Friese, PI)
- 2015-17 Hastings Center / UCSF Newborn Screening Ethics & Policy Board member (Barbara Koenig, PI)
- 2013-15 Ethics Advisory Board member, *NEPTUNE* project (Mathias Krenzler, PI)
- 2012-13 Expert Advisory Panel member, *Amyloid Imaging Disclosure* project, University of Pennsylvania (Jason Karlawish, MD, PI)

- 2009-11 Expert Advisory Panel member, *Presenting Diagnostic Results from Large-Scale Clinical Mutation Testing* study, Cleveland Clinic (Richard Sharp, Ph.D., PI)
- 2007 Expert Consultant, *Ethics of Surrogate Consent for Dementia Research* project, University of Michigan Medical School (Scott Kim, M.D., Ph.D., PI)
- 2006 Expert Panel member, *Fitness to Drive in Early Stage Dementia* study, University of Michigan Transportation Research Institute (David Eby, Ph.D., PI)
- 2000-2001 Research psychologist, dementia treatment study, Wellesley College Center for Research on Women (Nancy Emerson Lombardo, Ph.D., PI)
- 1995-96 Research Assistant, Department of Psychiatry, University of Michigan Medical Center (James Coyne, Ph.D., PI)
- 1994 Research Assistant, Duke University Bryan Alzheimer's Disease Research Center (Kathleen Welsh-Bohmer, Ph.D., PI)
- 1994-95 Research Assistant, *Adolescent Life Transitions* Study, Institute for Social Research, University of Michigan (Jacquelyne Eccles, Ph.D., PI)
- 1992-93 Research Assistant, Cancer Pain Management Program, University of North Carolina School of Nursing (Jo Ann Dalton, Ed.D., PI)
- 1991-93 Research Assistant, Community Guidance Clinic, Department of Psychiatry, Duke University Medical Center (John Lochman, Ph.D., PI)

PROFESSIONAL MEMBERSHIPS

- American Public Health Association
- American Society for Bioethics & Humanities
- International Society to Advance Alzheimer's Research & Treatment

PROFESSIONAL SERVICE ACTIVITIES

Scientific Review

Ad-hoc Member, Social & Ethical Issues in Research Study Section, National Institutes of Health

Special society journal reviewer, *Alzheimer's and Dementia*

Reviewer, *Alzheimer Disease and Associated Disorders*

Reviewer, *Alzheimer's Research & Therapy*

Reviewer, *American Journal of Geriatric Psychiatry*

Reviewer, *American Journal of Public Health*

Reviewer, *Archives of General Psychiatry*

Reviewer, *Community Genetics*

Reviewer, *Cross-Cultural Gerontology*

Reviewer, *Genetics in Medicine*

Reviewer, *Health Education & Behavior*

Reviewer, *IRB: Ethics and Human Research*

Reviewer, *International Journal of Geriatric Psychiatry*

Reviewer, *JAMA Neurology*

Reviewer, *Journal of Alzheimer's Disease*

Reviewer, *Journal of the American Geriatrics Society*

Reviewer, *Journal of Applied Gerontology*

Reviewer, *Journal of Genetic Counseling*

Reviewer, *Journal of Geriatric Psychiatry & Neurology*

Reviewer, *Journal of Gerontology: Psychological Sciences*

Reviewer, *Journal of the American Medical Association*

Reviewer, *Journal of Mental Health and Aging*

Reviewer, *Journal of Public Health Management and Practice*

Reviewer, *Neurology*

Reviewer, *Psychology and Aging*

Reviewer, *Research on Aging*

Reviewer, Alzheimer's Association scientific grants program

Reviewer, American Society for Bioethics and Humanities annual meeting

Reviewer, Cengage Learning, Career Education and Training Services

Reviewer, German Research Foundation research grants program

Reviewer, Pennsylvania Department of Health research grants program

Reviewer, United Kingdom National Institute for Health Research grants program

Reviewer, Wellcome Trust Postdoctoral Research Fellowship program in Biomedical Ethics

Reviewer, Wiley/Jossey-Bass, Public Health and Health Services

Department, School and University Level Service

2018-present Chair, Doctoral Admissions Committee, Department of Health Behavior & Health Education, University of Michigan School of Public Health

2018-present	Member, Precision Health Initiative Education Work Group, University of Michigan
2016-present	Member, Advisory Board, University of Michigan Genetic Counseling Program
2010-present	Director, Program in Genomics, Health and Society, University of Michigan Center for Bioethics and Social Sciences in Medicine
2008-present	Co-Director, Dual Degree (M.P.H./M.S.) Program in Public Health and Genetic Counseling, University of Michigan Schools of Public Health and Medicine
2011-19	Director, Public Health Genetics Certificate Program, University of Michigan School of Public Health
2017-18	Member, Doctoral Admissions Committee, Department of Health Behavior & Health Education, University of Michigan School of Public Health
2016-17	Co-Chair, Planning Committee, 75 th Anniversary of the University of Michigan School of Public Health
2016-17	Chair, Faculty Search Committee (2 open positions), Department of Health Behavior and Health Education, University of Michigan School of Public Health
2013-17	Member, University of Michigan Rackham Graduate School Integrity Board
2015-16	Member, Committee on the Undergraduate Program, University of Michigan School of Public Health
2006-16	Member (with service as Chair from 2011-15), Curriculum Committee, Department of Health Behavior & Health Education, University of Michigan School of Public Health
2010-15	Member, Advisory Committee for Academic Programs (ACAP), University of Michigan School of Public Health
2010-11	Chair, Faculty Search Committee, Department of Health Behavior and Health Education, University of Michigan School of Public Health
2006-11	Executive Committee Member, Public Health Genetics Certificate Program, University of Michigan School of Public Health
2009-10	Chair, Planning Committee, University of Michigan Distinguished Faculty & Graduate Student Seminars program: <i>The Ethical, Legal, and Social Implications of Personal Genomics</i>
2007-08	Steering Committee Member, University of Michigan Ethics in Public Life Initiative
2009	Chair, Ad-Hoc Ethics Subcommittee on Student Misconduct, University of Michigan School of Public Health

- 2009 Panel Member, Rackham-CRLT “Preparing Future Faculty” Seminar, University of Michigan
- 2009 Panel Member, Ann Arbor Community Forum on Biobanking, University of Michigan Life Sciences & Society Program
- 2005-06 Founding Webmaster, Boston University Alzheimer’s Disease Center website (www.bu.edu/alzresearch)
- 2002-06 Founding Editor, Boston University Alzheimer’s Disease Center newsletter, *BU ADC Bulletin*

State and National Level Service

- 2019-present External Advisory Board member, national Alzheimer’s Clinical Trial Consortium (ACTC)
- 2018-present Co-Chair, statewide Michigan Dementia Coalition
- 2018-present Member, Outreach, Recruitment, & Education Core Steering Committee, NIH Alzheimer’s Disease Centers program
- 2018-present Member, Genetic & Biomarker Disclosure Workgroup, NIH Alzheimer’s Disease Centers program
- 2018-present Member, Washtenaw County Health Department Ethics Committee
- 2015-18 Grand Champions level fundraiser, national Alzheimer’s Association Walk to End Alzheimer’s
- 2012-15 Empirical Research Review Committee, American Society for Bioethics and the Humanities Annual Meeting
- 2009-10 Chair, Public Health Ethics Review Committee, American Society for Bioethics and the Humanities Annual Meeting
- 2009 Advisory Committee Member, “Genetics for Social and Behavioral Scientists” Project, NCHPEG/NIH Office of Behavioral and Social Science Research
- 2008 Planning Committee, American Society for Bioethics and the Humanities Annual Meeting
- 2005-06 Planning Committee (Early Stage Track Co-Chair), Alzheimer’s Association National Dementia Care Conference