# Katrina A B Goddard

List of Publications by Year
in descending order



| | | | |
|---|---|---|---|
| 46 papers | 1,565 citations | 304743 22 h-index | 315739 38 g-index |
| 47 all docs | 47 docs citations | 47 times ranked | 2806 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Return of Genomic Results to Research Participants: The Floor, the Ceiling, and the Choices In Between. American Journal of Human Genetics, 2014, 94, 818-826. | 6.2 | 342 |
| 2 | The evolving landscape of expanded carrier screening: challenges and opportunities. Genetics in Medicine, 2019, 21, 790-797. | 2.4 | 90 |
| 3 | A standardized, evidence-based protocol to assess clinical actionability of genetic disorders associated with genomic variation. Genetics in Medicine, 2016, 18, 1258-1268. | 2.4 | 89 |
| 4 | Systematic review of the predictive effect of MSI status in colorectal cancer patients undergoing 5FU-based chemotherapy. BMC Cancer, 2015, 15, 156. | 2.6 | 81 |
| 5 | A survey of current practices for genomic sequencing test interpretation and reporting processes in US laboratories. Genetics in Medicine, 2017, 19, 575-582. | 2.4 | 68 |
| 6 | Awareness and use of direct-to-consumer nutrigenomic tests, United States, 2006. Genetics in Medicine, 2007, 9, 510-517. | 2.4 | 56 |
| 7 | Dengue hemorrhagic fever is associated with polymorphisms in JAK1. European Journal of Human Genetics, 2010, 18, 1221-1227. | 2.8 | 45 |
| 8 | Underutilization of Lynch syndrome screening in a multisite study of patients with colorectal cancer. Genetics in Medicine, 2013, 15, 933-940. | 2.4 | 45 |
| 9 | â€œIs It Worth Knowing?â€•Focus Group Participantsâ€™ Perceived Utility of Genomic Preconception Carrier Screening. Journal of Genetic Counseling, 2016, 25, 135-145. | 1.6 | 44 |
| 10 | Description and pilot results from a novel method for evaluating return of incidental findings from next-generation sequencing technologies. Genetics in Medicine, 2013, 15, 721-728. | 2.4 | 40 |
| 11 | Public Awareness and Use of Direct-to-Consumer Genetic Tests: Results From 3 State Population-Based Surveys, 2006. American Journal of Public Health, 2009, 99, 442-445. | 2.7 | 37 |
| 12 | Reasons for Declining Preconception Expanded Carrier Screening Using Genome Sequencing. Journal of Genetic Counseling, 2017, 26, 971-979. | 1.6 | 37 |
| 13 | Building the evidence base for decision making in cancer genomic medicine using comparative effectiveness research. Genetics in Medicine, 2012, 14, 633-642. | 2.4 | 36 |
| 14 | Preconception Carrier Screening by Genome Sequencing: Results from the Clinical Laboratory. American Journal of Human Genetics, 2018, 102, 1078-1089. | 6.2 | 35 |
| 15 | Evidenceâ€based assessments of clinical actionability in the context of secondary findings: Updates from ClinGen's Actionability Working Group. Human Mutation, 2018, 39, 1677-1685. | 2.5 | 34 |
| 16 | Comparative Effectiveness Research in Cancer Genomics and Precision Medicine: Current Landscape and Future Prospects. Journal of the National Cancer Institute, 2013, 105, 929-936. | 6.3 | 30 |
| 17 | Universal tumor screening for Lynch syndrome: Assessment of the perspectives of patients with colorectal cancer regarding benefits and barriers. Cancer, 2015, 121, 3281-3289. | 4.1 | 30 |
| 18 | Evidence of linkage and association on chromosome 20 for late-onset Alzheimer disease. Neurogenetics, 2004, 5, 121-128. | 1.4 | 27 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Design of a randomized controlled trial for genomic carrier screening in healthy patients seeking preconception genetic testing. Contemporary Clinical Trials, 2017, 53, 100-105. | 1.8 | 26 |
| 20 | Generating a taxonomy for genetic conditions relevant to reproductive planning. American Journal of Medical Genetics, Part A, 2016, 170, 565-573. | 1.2 | 25 |
| 21 | Stakeholder assessment of the evidence for cancer genomic tests: insights from three case studies. Genetics in Medicine, 2012, 14, 656-662. | 2.4 | 24 |
| 22 | Genome sequencing and carrier testing: decisions on categorization and whether to disclose results of carrier testing. Genetics in Medicine, 2017, 19, 803-808. | 2.4 | 24 |
| 23 | Time Costs for Genetic Counseling in Preconception Carrier Screening with Genome Sequencing. Journal of Genetic Counseling, 2018, 27, 823-833. | 1.6 | 22 |
| 24 | Patient perspectives on the use of categories of conditions for decision making about genomic carrier screening results. American Journal of Medical Genetics, Part A, 2018, 176, 376-385. | 1.2 | 21 |
| 25 | Patients' ratings of genetic conditions validate a taxonomy to simplify decisions about preconception carrier screening via genome sequencing. American Journal of Medical Genetics, Part A, 2016, 170, 574-582. | 1.2 | 20 |
| 26 | The NextGen Study: patient motivation for participation in genome sequencing for carrier status. Molecular Genetics & Genomic Medicine, 2017, 5, 508-515. | 1.2 | 20 |
| 27 | Validity of Eight Integrated Healthcare Delivery Organizations' Administrative Clinical Data to Capture Breast Cancer Chemotherapy Exposure. Cancer Epidemiology Biomarkers and Prevention, 2012, 21, 673-680. | 2.5 | 19 |
| 28 | It's complicated: criteria for policy decisions for the clinical integration of genomeâ€scale sequencing for reproductive decision making. Molecular Genetics & Genomic Medicine, 2015, 3, 239-242. | 1.2 | 19 |
| 29 | Assessment of willingness to pay for expanded carrier screening among women and couples undergoing preconception carrier screening. PLoS ONE, 2018, 13, e0200139. | 2.5 | 19 |
| 30 | Beta 2 adrenergic receptor polymorphisms in cystic fibrosis. Pediatric Pulmonology, 2005, 39, 544-550. | 2.0 | 16 |
| 31 | A Confidenceâ€Limitâ€Based Approach to the Assessment of Hardyâ€“Weinberg Equilibrium. Biometrical Journal, 2010, 52, 253-270. | 1.0 | 15 |
| 32 | Approaches to carrier testing and results disclosure in translational genomics research: The clinical sequencing exploratory research consortium experience. Molecular Genetics & Genomic Medicine, 2018, 6, 898-909. | 1.2 | 15 |
| 33 | Recommended care and care adherence following a diagnosis of Lynch syndrome: a mixed-methods study. Hereditary Cancer in Clinical Practice, 2019, 17, 31. | 1.5 | 14 |
| 34 | Universal screening for Lynch syndrome among patients with colorectal cancer: patient perspectives on screening and sharing results with at-risk relatives. Familial Cancer, 2017, 16, 377-387. | 1.9 | 11 |
| 35 | Anticipated responses of early adopter genetic specialists and nongenetic specialists to unsolicited genomic secondary findings. Genetics in Medicine, 2018, 20, 1186-1195. | 2.4 | 11 |
| 36 | Lessons Learned From A Study Of Genomics-Based Carrier Screening For Reproductive Decision Making. Health Affairs, 2018, 37, 809-816. | 5.2 | 10 |

| # | Article | IF | Citations |
|---|---|---|---|
| 37 | Utilization of HER2 genetic testing in a multi-institutional observational study. American Journal of Managed Care, 2012, 18, 704-12. | 1.1 | 10 |
| 38 | HLA-DQA Is Associated with Abdominal Aortic Aneurysms in the Belgian Population. Annals of the New York Academy of Sciences, 2006, 1085, 392-395. | 3.8 | 9 |
| 39 | Adapting the logical basis of tests for Hardyâ€Weinberg Equilibrium to the real needs of association studies in human and medical genetics. Genetic Epidemiology, 2009, 33, 569-580. | 1.3 | 9 |
| 40 | Association mapping by generalized linear regression with densityâ€based haplotype clustering. Genetic Epidemiology, 2009, 33, 16-26. | 1.3 | 6 |
| 41 | Implementation of a Systematic Tumor Screening Program for Lynch Syndrome in an Integrated Health Care Setting. Familial Cancer, 2019, 18, 317-325. | 1.9 | 6 |
| 42 | Density-based clustering in haplotype analysis for association mapping. BMC Proceedings, 2007, 1, S27. | 1.6 | 5 |
| 43 | Issues in association mapping with high-density SNP data and diverse family structures. Genetic Epidemiology, 2007, 31, S22-S33. | 1.3 | 4 |
| 44 | Expert and lay perspectives on burden, risk, tolerability, and acceptability of clinical interventions for genetic disorders. Genetics in Medicine, 2019, 21, 2561-2568. | 2.4 | 4 |
| 45 | ClinGenâ€™s Pediatric Actionability Working Group: Clinical actionability of secondary findings from genome-scale sequencing in children and adolescents. Genetics in Medicine, 2022, 24, 1328-1335. | 2.4 | 4 |
| 46 | Identifying patients with Lynch syndrome using a universal tumor screening program in an integrated healthcare system. Hereditary Cancer in Clinical Practice, 2022, 20, 17. | 1.5 | 4 |