Kelly E Ormond

List of Publications by Year
in descending order

112 papers	9,523 citations	116194 36 h-index	45040 94 g-index
117 all docs	117 docs citations	117 times ranked	13218 citing authors

#	Article	IF	Citations
1	â€œI wish that there was more infoâ€: characterizing the uncertainty experienced by carriers of pathogenic ATM and/or CHEK2 variants. Familial Cancer, 2022, 21, 143-155.	0.9	7
2	Dealing with uncertainty in prenatal genomics. , 2022, , 69-81.		0
3	Treatment decision-making in sickle cell disease patients. Journal of Community Genetics, 2022, 13, 143-151.	0.5	3
4	Assessing womenâ€™s preferences towards tests that may reveal uncertain results from prenatal genomic testing: Development of attributes for a discrete choice experiment, using a mixed-methods design. PLoS ONE, 2022, 17, e0261898.	1.1	4
5	Factors that impact on women's decisionâ€making around prenatal genomic tests: An international discrete choice survey. Prenatal Diagnosis, 2022, 42, 934-946.	1.1	5
6	Parental experiences of uncertainty following an abnormal fetal anomaly scan: Insights using Hanâ€™s taxonomy of uncertainty. Journal of Genetic Counseling, 2021, 30, 198-210.	0.9	20
7	Dealing with uncertain results from chromosomal microarray and exome sequencing in the prenatal setting: An international crossâ€sectional study with healthcare professionals. Prenatal Diagnosis, 2021, 41, 720-732.	1.1	13
8	Improving reporting standards for polygenic scores in risk prediction studies. Nature, 2021, 591, 211-219.	13.7	265
9	Application of a framework to guide genetic testing communication across clinical indications. Genome Medicine, 2021, 13, 71.	3.6	14
10	Public willingness to participate in personalized health research and biobanking: A large-scale Swiss survey. PLoS ONE, 2021, 16, e0249141.	1.1	31
11	U.S. Genetic counselorsâ€™ perceptions of inpatient genetic counseling: A valuable model for medically complex patients. Journal of Genetic Counseling, 2021, 30, 1683-1694.	0.9	2
12	â€œDoctors can read about it, they can know about it, but they've never lived with itâ€: How parents use social media throughout the diagnostic odyssey. Journal of Genetic Counseling, 2021, 30, 1707-1718.	0.9	10
13	How to deal with uncertainty in prenatal genomics: A systematic review of guidelines and policies. Clinical Genetics, 2021, 100, 647-658.	1.0	15
14	Physician-directed genetic screening to evaluate personal risk for medically actionable disorders: a large multi-center cohort study. BMC Medicine, 2021, 19, 199.	2.3	17
15	Defining the Critical Components of Informed Consent for Genetic Testing. Journal of Personalized Medicine, 2021, 11, 1304.	1.1	6
16	Clinical Genetics Lacks Standard Definitions and Protocols for the Collection and Use of Diversity Measures. American Journal of Human Genetics, 2020, 107, 72-82.	2.6	52
17	Informed Consent in the Genomics Era. Cold Spring Harbor Perspectives in Medicine, 2020, 10, a036582.	2.9	16
18	â€œI donâ€™t want to be Henrietta Lacksâ€: diverse patient perspectives on donating biospecimens for precision medicine research. Genetics in Medicine, 2019, 21, 107-113.	1.1	67

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19	Developing a conceptual, reproducible, rubric-based approach to consent and result disclosure for genetic testing by clinicians with minimal genetics background. Genetics in Medicine, 2019, 21, 727-735.	1.1	40
20	Genetics and ethics. Journal of Community Genetics, 2019, 10, 1-2.	0.5	0
21	The clinical application of gene editing: ethical and social issues. Personalized Medicine, 2019, 16, 337-350.	0.8	25
22	Attitudes of Members of Genetics Professional Societies Toward Human Gene Editing. CRISPR Journal, 2019, 2, 331-339.	1.4	15
23	Attitudes Toward Hypothetical Uses of Gene-Editing Technologies in Parents of People with Autosomal Aneuploidies. CRISPR Journal, 2019, 2, 324-330.	1.4	10
24	Developing a genomics rotation: Practical training around variant interpretation for genetic counseling students. Journal of Genetic Counseling, 2019, 28, 466-476.	0.9	10
25	Attitudes of people with inherited retinal conditions toward gene editing technology. Molecular Genetics & Genomic Medicine, 2019, 7, e00803.	0.6	15
26	A genetic counseling needs assessment of Mexico. Molecular Genetics & Genomic Medicine, 2019, 7, e668.	0.6	12
27	Secondary findings: How did we get here, and where are we going?. Journal of Genetic Counseling, 2019, 28, 326-333.	0.9	20
28	Much ado about nothing: A qualitative study of the experiences of an averageâ€risk population receiving results of exome sequencing. Journal of Genetic Counseling, 2019, 28, 428-437.	0.9	15
29	What do we do now?: Responding to claims of germline gene editing in humans. Genetics in Medicine, 2019, 21, 2181-2183.	1.1	15
30	Assessing genetic counselorsâ€™ experiences with physician aidâ€inâ€dying and practice implications. Journal of Genetic Counseling, 2019, 28, 164-173.	0.9	4
31	The Responsibility to Recontact Research Participants after Reinterpretation of Genetic and Genomic Research Results. American Journal of Human Genetics, 2019, 104, 578-595.	2.6	91
32	Genetic counselors' perceptions of uncertainty in pretest counseling for genomic sequencing: A qualitative study. Journal of Genetic Counseling, 2019, 28, 292-303.	0.9	10
33	Challenges of infertility genetic counseling: Impact on counselors' personal and professional lives. Journal of Genetic Counseling, 2019, 28, 626-640.	0.9	1
34	Perspectives of Sickle Cell Disease Stakeholders on Heritable Genome Editing. CRISPR Journal, 2019, 2, 441-449.	1.4	9
35	Genetic Counselors' and Genetic Counseling Students' Implicit and Explicit Attitudes toward Homosexuality. Journal of Genetic Counseling, 2019, 28, 91-101.	0.9	12
36	The Global State of the Genetic Counseling Profession. European Journal of Human Genetics, 2019, 27, 183-197.	1.4	215

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37	Operationalizing the Reciprocal Engagement Model of Genetic Counseling Practice: a Framework for the Scalable Delivery of Genomic Counseling and Testing. Journal of Genetic Counseling, 2018, 27, 1111-1129.	0.9	25
38	Mindfulness Among Genetic Counselors Is Associated with Increased Empathy and Work Engagement and Decreased Burnout and Compassion Fatigue. Journal of Genetic Counseling, 2018, 27, 1175-1186.	0.9	40
39	Genetic counseling globally: Where are we now?. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2018, 178, 98-107.	0.7	109
40	Beyond Consent: Building Trusting Relationships With Diverse Populations in Precision Medicine Research. American Journal of Bioethics, 2018, 18, 3-20.	0.5	152
41	Exploring the Medical and Psychosocial Concerns of Adolescents and Young Adults With Craniofacial Microsomia. Cleft Palate-Craniofacial Journal, 2018, 55, 1430-1439.	0.5	21
42	Patient and provider perspectives on the development of personalized medicine: a mixed-methods approach. Journal of Community Genetics, 2018, 9, 283-291.	0.5	17
43	Metaphors matter: from biobank to a library of medical information. Genetics in Medicine, 2018, 20, 802-805.	1.1	6
44	Ethical considerations in prenatal testing: Genomic testing and medical uncertainty. Seminars in Fetal and Neonatal Medicine, 2018, 23, 1-6.	1.1	35
45	National Society of Genetic Counselors Code of Ethics: Explication of 2017 Revisions. Journal of Genetic Counseling, 2018, 27, 9-15.	0.9	7
46	Predictors of adverse psychological experiences surrounding genome-wide profiling for disease risk. Journal of Community Genetics, 2018, 9, 217-225.	0.5	15
47	Trustworthiness in Untrustworthy Times: Response to Open Peer Commentaries on Beyond Consent. American Journal of Bioethics, 2018, 18, W6-W8.	0.5	6
48	Human Germline Genome Editing. American Journal of Human Genetics, 2017, 101, 167-176.	2.6	168
49	Recommendations for reporting of secondary findings in clinical exome and genome sequencing, 2016 update (ACMG SF v2.0): a policy statement of the American College of Medical Genetics and Genomics. Genetics in Medicine, 2017, 19, 249-255.	1.1	1,398
50	Exercise restrictions trigger psychological difficulty in active and athletic adults with hypertrophic cardiomyopathy. Open Heart, 2016, 3, e000488.	0.9	29
51	2013 Review and Update of the Genetic Counseling Practice Based Competencies by a Task Force of the Accreditation Council for Genetic Counseling. Journal of Genetic Counseling, 2016, 25, 868-879.	0.9	44
52	Points to Consider: Ethical, Legal, and Psychosocial Implications of Genetic Testing in Children and Adolescents. American Journal of Human Genetics, 2015, 97, 6-21.	2.6	453
53	Non-invasive prenatal testing for aneuploidy and beyond: challenges of responsible innovation in prenatal screening. Summary and recommendations. European Journal of Human Genetics, 2015, , .	1.4	13
54	Non-invasive prenatal testing for aneuploidy and beyond: challenges of responsible innovation in prenatal screening. European Journal of Human Genetics, 2015, 23, 1438-1450.	1.4	260

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55	Standardizing Variant Interpretation in Genomic Sequencing: Implications for Genetic Counseling Practice. Current Genetic Medicine Reports, 2015, 3, 137-142.	1.9	1
56	Whole-Exome Sequencing of 10 Scientists: Evaluation of the Process and Outcomes. Mayo Clinic Proceedings, 2015, 90, 1327-1337.	1.4	10
57	Translating personalized medicine using new genetic technologies in clinical practice: the ethical issues. Personalized Medicine, 2014, 11, 211-222.	0.8	34
58	Views of Genetics Health Professionals on the Return of Genomic Results. Journal of Genetic Counseling, 2014, 23, 531-538.	0.9	43
59	Clinical Interpretation and Implications of Whole-Genome Sequencing. JAMA - Journal of the American Medical Association, 2014, 311, 1035.	3.8	398
60	Teaching Genomic Counseling: Preparing the Genetic Counseling Workforce for the Genomic Era. Journal of Genetic Counseling, 2014, 23, 445-451.	0.9	60
61	Attitudes of Mothers of Children with Down Syndrome Towards Noninvasive Prenatal Testing. Journal of Genetic Counseling, 2014, 23, 805-813.	0.9	63
62	Evaluating the utilization of educational materials in communicating about Lynch syndrome to at-risk relatives. Familial Cancer, 2014, 13, 381-389.	0.9	32
63	Genetic Testing of Children for Predisposition to Mood Disorders: Anticipating the Clinical Issues. Journal of Genetic Counseling, 2014, 23, 566-577.	0.9	10
64	PATH-SCAN: a reporting tool for identifying clinically actionable variants. Pacific Symposium on Biocomputing Pacific Symposium on Biocomputing, 2014, , 229-40.	0.7	10
65	The Views of Pakistani Doctors Regarding Genetic Counseling Services â€“ Is there a Future?. Journal of Genetic Counseling, 2013, 22, 721-732.	0.9	11
66	Personal genome testing in medical education: student experiences with genotyping in the classroom. Genome Medicine, 2013, 5, 24.	3.6	53
67	ACMG recommendations for reporting of incidental findings in clinical exome and genome sequencing. Genetics in Medicine, 2013, 15, 565-574.	1.1	2,186
68	Noninvasive Prenatal Testing/Noninvasive Prenatal Diagnosis: the Position of the National Society of Genetic Counselors. Journal of Genetic Counseling, 2013, 22, 291-295.	0.9	168
69	Best ethical practices for clinicians and laboratories in the provision of noninvasive prenatal testing. Prenatal Diagnosis, 2013, 33, 656-661.	1.1	47
70	PATH-SCAN: A REPORTING TOOL FOR IDENTIFYING CLINICALLY ACTIONABLE VARIANTS. , 2013, , .		9
71	From genetic counseling to â€œgenomic counselingâ€• Molecular Genetics & Genomic Medicine, 2013, 1, 189-193.	0.6	59
72	Attitudes towards Social Networking and Sharing Behaviors among Consumers of Direct-to-Consumer Personal Genomics. Journal of Personalized Medicine, 2013, 3, 275-287.	1.1	7

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73	Evidence That Personal Genome Testing Enhances Student Learning in a Course on Genomics and Personalized Medicine. PLoS ONE, 2013, 8, e68853.	1.1	84
74	Customers or research participants?: Guidance for research practices in commercialization of personal genomics. Genetics in Medicine, 2012, 14, 833-835.	1.1	12
75	Genetic Counseling for Prenatal Testing: Where is the Discussion About Disability?. Journal of Genetic Counseling, 2012, 21, 814-824.	0.9	39
76	Concurrent Use of Cultural Health Practices and Western Medicine During Pregnancy: Exploring the Mexican Experience in the United States. Journal of Genetic Counseling, 2011, 20, 609-624.	0.9	14
77	Practical considerations to guide development of access controls and decision support for genetic information in electronic medical records. BMC Health Services Research, 2011, 11, 294.	0.9	8
78	Noninvasive prenatal diagnosis: pregnant women's interest and expected uptake. Prenatal Diagnosis, 2011, 31, 1292-1299.	1.1	117
79	Medical and graduate students' attitudes toward personal genomics. Genetics in Medicine, 2011, 13, 400-408.	1.1	41
80	Phased Whole-Genome Genetic Risk in a Family Quartet Using a Major Allele Reference Sequence. PLoS Genetics, 2011, 7, e1002280.	1.5	137
81	Health insurance coverage of genetic services in Illinois. Genetics in Medicine, 2010, 12, 525-531.	1.1	14
82	Clinical assessment incorporating a personal genome. Lancet, The, 2010, 375, 1525-1535.	6.3	637
83	Challenges in the clinical application of whole-genome sequencing. Lancet, The, 2010, 375, 1749-1751.	6.3	207
84	Prenatal Screening and Diagnosis. Issues in Clinical Child Psychology, 2010, , 221-240.	0.2	5
85	What Is the Role of Nongeneticist Physicians, and Are They Prepared for It?. AMA Journal of Ethics, 2009, 11, 678-682.	0.4	0
86	699: First trimester screening versus diagnostic testing; how important is prenatal counseling in the decision-making process?. American Journal of Obstetrics and Gynecology, 2009, 201, S253.	0.7	0
87	Assessing the understanding of biobank participants. American Journal of Medical Genetics, Part A, 2009, 149A, 188-198.	0.7	118
88	Information preferences of high literacy pregnant women regarding informed consent models for genetic carrier screening. Patient Education and Counseling, 2009, 75, 244-250.	1.0	27
89	Letter to the Editor. Journal of Genetic Counseling, 2009, 18, 197-199.	0.9	3
90	Association of spinocerebellar ataxia type 3 and spinocerebellar ataxia type 8 microsatellite expansions: Genetic counseling implications. Movement Disorders, 2008, 23, 154-155.	2.2	11

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91	Medical Ethics for the Genome World. Journal of Molecular Diagnostics, 2008, 10, 377-382.	1.2	3
92	The next exclusion debate: Assessing technology, ethics, and intellectual disability after the human genome project. Mental Retardation and Developmental Disabilities Research Reviews, 2007, 13, 121-128.	3.5	20
93	Outcome of chromosomally abnormal pregnancies in Lebanon: obstetricians' roles during and after prenatal diagnosis. Prenatal Diagnosis, 2007, 27, 525-534.	1.1	18
94	What do Patients Prefer: Informed Consent Models for Genetic Carrier Testing. Journal of Genetic Counseling, 2007, 16, 539-550.	0.9	40
95	Genetic Risk Assessment and BRCA Mutation Testing. Annals of Internal Medicine, 2006, 144, 303.	2.0	0
96	Implementing prenatal screening for cystic fibrosis in routine obstetric practice. American Journal of Obstetrics and Gynecology, 2006, 194, 904.	0.7	2
97	Knowledge and Attitudes Toward a Free Education and Ashkenazi Jewish Carrier Testing Program. Journal of Genetic Counseling, 2006, 15, 61-70.	0.9	15
98	Attitudes of genetic counselors towards expanding newborn screening and offering predictive genetic testing to children. American Journal of Medical Genetics, Part A, 2006, 140A, 2312-2319.	0.7	24
99	Disclosing Genetic Research Results: Examples from Practice. American Journal of Bioethics, 2006, 6, 30-32.	0.5	23
100	Marshall-Smith syndrome: Natural history and evidence of an osteochondrodysplasia with connective tissue abnormalities. American Journal of Medical Genetics, Part A, 2005, 137A, 117-124.	0.7	30
101	NSGC Foundationsâ€"Then, Now, and Tomorrow. Journal of Genetic Counseling, 2005, 14, 85-88.	0.9	2
102	Adjunct Prenatal Testing: Patient Decisions Regarding Ethnic Carrier Screening and Fluorescence In Situ Hybridization. Journal of Genetic Counseling, 2004, 13, 45-63.	0.9	5
103	Outline of a medical genetics curriculum for internal medicine residency training programs. Genetics in Medicine, 2004, 6, 543-547.	1.1	40
104	Attitudes of Health Care Trainees About Genetics and Disability: Issues of Access, Health Care Communication, and Decision Making. Journal of Genetic Counseling, 2003, 12, 333-349.	0.9	28
105	Effect of family history on disclosure patterns of cystic fibrosis carrier status. American Journal of Medical Genetics Part A, 2003, 119C, 70-77.	2.4	42
106	The genetic family history as a risk assessment tool in internal medicine. Genetics in Medicine, 2003, 5, 84-91.	1.1	95
107	The Impact of Genetic Technologies on Perceptions of Disability. Quality Management in Health Care, 2000, 8, 19-26.	0.4	11
108	Recommendations for Telephone Counseling. Journal of Genetic Counseling, 2000, 9, 63-71.	0.9	32

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109	Pregnancy Outcome Following Maternal Use of the New Selective Serotonin Reuptake Inhibitors. JAMA - Journal of the American Medical Association, 1998, 279, 609.	3.8	371
110	Accidental electric shock in pregnancy: A prospective cohort study. American Journal of Obstetrics and Gynecology, 1997, 176, 678-681.	0.7	64
111	Update and Review: Maternal Serum Screening. Journal of Genetic Counseling, 1997, 6, 395-417.	0.9	11
112	Pre-screening education in multiple marker screening programs: The effect on patient anxiety and knowledge. Journal of Genetic Counseling, 1996, 5, 69-80.	0.9	17