

The psychological impact of genetic information on chil

Genetics in Medicine

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Genetic testing for the risk of developing late effects among survivors of childhood cancer: Consumer understanding, acceptance, and willingness to pay. <i>Cancer</i> , 2016, 122, 2876-2885.	4.1	6
2	Predictive genetic testing of minors: evidence and experience with families. <i>Genetics in Medicine</i> , 2016, 18, 763-764.	2.4	9
3	Ethical issues in pediatric genetic testing and screening. <i>Current Opinion in Pediatrics</i> , 2016, 28, 700-704.	2.0	42
4	Psychosocial effects in parents and children 12 years after newborn genetic screening for type 1 diabetes. <i>European Journal of Human Genetics</i> , 2017, 25, 397-403.	2.8	6
5	Parents' attitudes toward genetic testing of children for health conditions: A systematic review. <i>Clinical Genetics</i> , 2017, 92, 569-578.	2.0	34
6	Cancer Screening Recommendations for Individuals with Li-Fraumeni Syndrome. <i>Clinical Cancer Research</i> , 2017, 23, e38-e45.	7.0	358
7	Psychosocial care and cardiac genetic counseling following sudden cardiac death in the young. <i>Progress in Pediatric Cardiology</i> , 2017, 45, 31-36.	0.4	15
8	The Precision Medicine Nation. <i>Hastings Center Report</i> , 2017, 47, 19-29.	1.0	49
9	Genetic Counselor Recommendations for Cancer Predisposition Evaluation and Surveillance in the Pediatric Oncology Patient. <i>Clinical Cancer Research</i> , 2017, 23, e91-e97.	7.0	92
10	Talking with Children About Adult-Onset Hereditary Cancer Risk: A Developmental Approach for Parents. <i>Journal of Genetic Counseling</i> , 2018, 27, 533-548.	1.6	24
11	Cancer-Related Genetic Testing and Personalized Medicine for Adolescents: A Narrative Review of Impact and Understanding. <i>Journal of Adolescent and Young Adult Oncology</i> , 2018, 7, 259-262.	1.3	20
12	Development of an Educational Program Integrating Concepts of Genetic Risk and Preventive Strategies for Children with a Family History of Melanoma. <i>Journal of Cancer Education</i> , 2018, 33, 774-781.	1.3	6
13	Reconceptualizing harms and benefits in the genomic age. <i>Personalized Medicine</i> , 2018, 15, 419-428.	1.5	13
14	Genetic Basis for Congenital Heart Disease: Revisited: A Scientific Statement From the American Heart Association. <i>Circulation</i> , 2018, 138, e653-e711.	1.6	387
15	When to Offer Predictive Genetic Testing to Children at Risk of an Inherited Arrhythmia or Cardiomyopathy. <i>Circulation Genomic and Precision Medicine</i> , 2018, 11, e002300.	3.6	5
16	Adolescent and Parental Attitudes About Return of Genomic Research Results: Focus Group Findings Regarding Decisional Preferences. <i>Journal of Empirical Research on Human Research Ethics</i> , 2018, 13, 371-382.	1.3	36
17	Parent-child communication surrounding genetic testing for Li-Fraumeni syndrome: Living under the cloud of cancer. <i>Pediatric Blood and Cancer</i> , 2018, 65, e27350.	1.5	15
18	Secondary findings from next-generation sequencing: what does actionable in childhood really mean?. <i>Genetics in Medicine</i> , 2019, 21, 124-132.	2.4	18

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19	Practice Variation among an International Group of Genetic Counselors on when to Offer Predictive Genetic Testing to Children at Risk of an Inherited Arrhythmia or Cardiomyopathy. <i>Journal of Genetic Counseling</i> , 2019, 28, 70-79.	1.6	4
20	Healthcare professionals' attitudes toward cancer precision medicine: A systematic review. <i>Seminars in Oncology</i> , 2019, 46, 291-303.	2.2	17
21	Qualitative Research on Expanded Prenatal and Newborn Screening: Robust but Marginalized. <i>Hastings Center Report</i> , 2019, 49, S72-S81.	1.0	5
22	Health professional and at-risk BRCA young adult perspectives about information needs: What does Gen Y need to know?. <i>Journal of Genetic Counseling</i> , 2019, 28, 1154-1165.	1.6	8
23	What Is the Psychosocial Impact of Providing Genetic and Genomic Health Information to Individuals? An Overview of Systematic Reviews. <i>Hastings Center Report</i> , 2019, 49, S88-S96.	1.0	33
24	Genome Diagnostics: Novel Strategies for Measuring Value. <i>Journal of Managed Care & Specialty Pharmacy</i> , 2019, 25, 1096-1101.	0.9	3
25	Psychological outcomes related to exome and genome sequencing result disclosure: a meta-analysis of seven Clinical Sequencing Exploratory Research (CSER) Consortium studies. <i>Genetics in Medicine</i> , 2019, 21, 2781-2790.	2.4	55
26	I can remember how I felt, but I don't remember the gene: Families' experiences of cancer-related genetic testing in childhood. <i>Pediatric Blood and Cancer</i> , 2019, 66, e27762.	1.5	26
27	Genetic Counseling and Genomic Sequencing. , 2019, , 125-142.		0
28	Rethinking the "open future" argument against predictive genetic testing of children. <i>Genetics in Medicine</i> , 2019, 21, 2190-2198.	2.4	43
29	Parents', Health Care Professionals', and Scientists' Experiences of a Precision Medicine Pilot Trial for Patients With High-Risk Childhood Cancer: A Qualitative Study. <i>JCO Precision Oncology</i> , 2019, 3, 1-11.	3.0	8
30	Families' and health care professionals' attitudes towards Li-Fraumeni syndrome testing in children: A systematic review. <i>Clinical Genetics</i> , 2019, 95, 140-150.	2.0	12
31	Genetics-related service and information needs of childhood cancer survivors and parents: a mixed-methods study. <i>European Journal of Human Genetics</i> , 2020, 28, 6-16.	2.8	7
32	Cancer-predisposing germline variants and childhood cancer. , 2020, , 221-232.		0
33	Identifying the challenges to successfully teaching about genetic diversity among Japanese junior high school students. <i>SAGE Open Medicine</i> , 2020, 8, 205031212096065.	1.8	4
34	"A sword of Damocles": patient and caregiver beliefs, attitudes and perspectives on presymptomatic testing for autosomal dominant polycystic kidney disease: a focus group study. <i>BMJ Open</i> , 2020, 10, e038005.	1.9	5
35	Human Genetics Society of Australasia Position Statement: Predictive and Presymptomatic Genetic Testing in Adults and Children. <i>Twin Research and Human Genetics</i> , 2020, 23, 184-189.	0.6	13
36	Discouraging Elective Genetic Testing of Minors: A Norm under Siege in a New Era of Genomic Medicine. <i>Cold Spring Harbor Perspectives in Medicine</i> , 2020, 10, a036657.	6.2	8

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37	Pediatric reporting of genomic results study (PROGRESS): a mixed-methods, longitudinal, observational cohort study protocol to explore disclosure of actionable adult- and pediatric-onset genomic variants to minors and their parents. BMC Pediatrics, 2020, 20, 222.	1.7	11
38	Current controversies in prenatal diagnosis 2: The 59 genes ACMG recommends reporting as secondary findings when sequencing postnatally should be reported when detected on fetal (and) Tj ETQq1 1 0.784314 rgBTt/Overlo	3.1	14
39	Parent and child perspectives on family interactions related to melanoma risk and prevention after CDKN2A/p16 testing of minor children. Journal of Community Genetics, 2020, 11, 321-329.	1.2	5
40	Factors Associated With Declining to Participate in a Pediatric Oncology Next-Generation Sequencing Study. JCO Precision Oncology, 2020, 4, 202-211.	3.0	15
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43	ACG Clinical Report and Recommendations on Transition of Care in Children and Adolescents With Hereditary Polyposis Syndromes. American Journal of Gastroenterology, 2021, 116, 638-646.	0.4	7
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48	Psychosocial interventions and needs among individuals and families with <scp>Liâ€Fraumeni</scp> syndrome: A scoping review. Clinical Genetics, 2022, 101, 161-182.	2.0	2
49	The Ethics of Predicting Autism Spectrum Disorder in Infancy. Journal of the American Academy of Child and Adolescent Psychiatry, 2021, 60, 942-945.	0.5	6
50	Polygenic risk scores in the clinic: new perspectives needed on familiar ethical issues. Genome Medicine, 2021, 13, 14.	8.2	79
51	"We Should View Him as an Individual": The Role of the Child's Future Autonomy in Shared Decision-Making About Unsolicited Findings in Pediatric Exome Sequencing. Health Care Analysis, 2021, 29, 249-261.	2.2	4
52	Psychosocial Aspects of Childhood Cancer Genetics. , 2021, , 445-471.		0
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55	A Person-Centered Approach to Cardiovascular Genetic Testing. Cold Spring Harbor Perspectives in Medicine, 2020, 10, a036624.	6.2	6
56	Predictive Genetic Testing for Hereditary Cancers in Children: An Argumentation. Journal of Comprehensive Pediatrics, 2019, 11, .	0.3	0
57	Genetic Testing in the Pediatric Nephrology Clinic: Understanding Families' Experiences. Journal of Pediatric Genetics, 2022, 11, 117-125.	0.7	7
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63	Fetal information as shared information: using NIPT to test for adult-onset conditions. Monash Bioethics Review, 2021, 39, 82-102.	0.8	2
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69	Parents' decision-making regarding whether to receive adult-onset only genetic findings for their children: Findings from the BabySeq Project. Genetics in Medicine, 2023, 25, 100002.	2.4	5
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75	A multidisciplinary approach to inherited retinal dystrophies from diagnosis to initial care: a narrative review with inputs from clinical practice. Orphanet Journal of Rare Diseases, 2023, 18, .	2.7	2
76	The Psychosocial Impact of Familial Endocrine Cancer Syndromes (FECS) on the Patient and Caregiver. , 2023, , 1-35.		0
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79	Psychiatric and educational aspects of familial adenomatous polyposis - a nationwide Danish cohort study with matched non-exposed individuals. American Journal of Gastroenterology, 2023, , .	0.4	0
80	Stakeholder Perception of the Implementation of Genetic Risk Testing for Twelve Multifactorial Diseases. Genes, 2024, 15, 49.	2.4	0
81	The psychological impact of genetic testing in childhood cancer: A systematic review. Psycho-Oncology, 2024, 33, .	2.3	0
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83	Human Genetics Society of Australasia Position Statement: Predictive and Presymptomatic Genetic Testing in Adults and Children. Twin Research and Human Genetics, 0, , 1-8.	0.6	0