

Risk Assessment, Genetic Counseling, and Genetic Testi

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Hereditary Cancer Evaluation in 2019—a Rapidly Evolving Landscape. <i>JAMA Oncology</i> , 2019, 5, 1539.	3.4	0
2	Maximizing the Outcomes of Breast Cancer Prevention. <i>JAMA Internal Medicine</i> , 2019, 179, 1621.	2.6	0
3	Blending Insights from Implementation Science and the Social Sciences to Mitigate Inequities in Screening for Hereditary Cancer Syndromes. <i>International Journal of Environmental Research and Public Health</i> , 2019, 16, 3899.	1.2	8
4	Regarding the Yin and Yang of Precision Cancer- Screening and Treatment: Are We Creating a Neglected Majority?. <i>International Journal of Environmental Research and Public Health</i> , 2019, 16, 4168.	1.2	5
6	Task force releases updated guidance on BRCA1/2 assessments. <i>Pharmacy Today</i> , 2019, 25, 4.	0.0	0
7	Hot Topic: Should all Women with Breast Cancer Undergo Genetic Testing?. <i>Current Breast Cancer Reports</i> , 2019, 11, 381-384.	0.5	1
8	Personalized breast cancer screening strategies: A systematic review and quality assessment. <i>PLoS ONE</i> , 2019, 14, e0226352.	1.1	38
9	Calling on Primary Care to Prevent BRCA-Related Cancers. <i>Journal of General Internal Medicine</i> , 2020, 35, 903-905.	1.3	2
10	<i>BRCA1/BRCA2</i> Pathogenic Variant Breast Cancer: Treatment and Prevention Strategies. <i>Annals of Laboratory Medicine</i> , 2020, 40, 114-121.	1.2	67
11	From Genetic Testing to Treatment and Prevention of BRCA-Related Breast Cancer. <i>Annals of Laboratory Medicine</i> , 2020, 40, 99-100.	1.2	0
12	Points to consider: is there evidence to support BRCA1/2 and other inherited breast cancer genetic testing for all breast cancer patients? A statement of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2020, 22, 681-685.	1.1	20
13	Genomic testing is best integrated into clinical practice when it is actionable. <i>Personalized Medicine</i> , 2020, 17, 5-8.	0.8	8
14	Implementing universal cancer screening programs can help sustain genomic medicine programs. <i>Personalized Medicine</i> , 2020, 17, 9-13.	0.8	1
15	DNA-Based Population Screening. <i>JAMA - Journal of the American Medical Association</i> , 2020, 323, 307.	3.8	31
16	Importance of family history and indications for genetic testing. <i>Breast Journal</i> , 2020, 26, 100-104.	0.4	13
17	A Pre-Test—Post-Test Trial of a Breast Cancer Risk Report for Women in Their 40s. <i>American Journal of Preventive Medicine</i> , 2020, 59, 343-354.	1.6	4
18	Frequency and spectrum of mutations across 94 cancer predisposition genes in African American women with invasive breast cancer. <i>Familial Cancer</i> , 2021, 20, 181-187.	0.9	2
19	Economic Evaluation of Population-Based BRCA1/BRCA2 Mutation Testing across Multiple Countries and Health Systems. <i>Cancers</i> , 2020, 12, 1929.	1.7	49

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20	Breast cancer risk assessment and management programs: A practical guide. <i>Breast Journal</i> , 2020, 26, 1556-1564.	0.4	7
21	Genetic Counseling, Testing, and Management of HBOC in India: An Expert Consensus Document from Indian Society of Medical and Pediatric Oncology. <i>JCO Global Oncology</i> , 2020, 6, 991-1008.	0.8	9
22	The contemporary landscape of genetic testing and breast cancer: Emerging issues. <i>Breast Journal</i> , 2020, 26, 1549-1555.	0.4	6
23	Patient With Questions About Cancer Risk. <i>Journal for Nurse Practitioners</i> , 2020, 16, e109-e112.	0.4	0
24	Pathogenic Variants in Breast Cancer Susceptibility Genes in Older Women. <i>JAMA - Journal of the American Medical Association</i> , 2020, 324, 397.	3.8	0
25	Cancer Prevention and Screening for Older Adults: Part 2. Interventions to Prevent and Screen for Breast, Prostate, Cervical, Ovarian, and Endometrial Cancer. <i>Journal of the American Geriatrics Society</i> , 2020, 68, 2684-2691.	1.3	8
26	Ovarian cancer predisposition beyond BRCA1 and BRCA2 genes. <i>International Journal of Gynecological Cancer</i> , 2020, 30, 1803-1810.	1.2	35
27	Breast cancer screening for women at high risk: review of current guidelines from leading specialty societies. <i>Breast Cancer</i> , 2021, 28, 1195-1211.	1.3	4
28	Integrating Personalized Medicine With Population Health Management. <i>JAMA - Journal of the American Medical Association</i> , 2020, 324, 631.	3.8	5
29	Parent of Origin Effects on Family Communication of Risk in BRCA+ Women: A Qualitative Investigation of Human Factors in Cascade Screening. <i>Cancers</i> , 2020, 12, 2316.	1.7	11
30	Strategies to enhance identification of hereditary breast cancer gene carriers. <i>Expert Review of Molecular Diagnostics</i> , 2020, 20, 861-865.	1.5	1
31	Implementing NGS-based BRCA tumour tissue testing in FFPE ovarian carcinoma specimens: hints from a real-life experience within the framework of expert recommendations. <i>Journal of Clinical Pathology</i> , 2021, 74, 596-603.	1.0	10
32	Molecular Features and Clinical Management of Hereditary Gynecological Cancers. <i>International Journal of Molecular Sciences</i> , 2020, 21, 9504.	1.8	13
33	New perspectives on the genetic causes of diminished ovarian reserve and opportunities for genetic screening: systematic review and meta-analysis. <i>F&S Reviews</i> , 2020, 1, 1-15.	0.7	4
34	Cost-effectiveness of Population-Wide Genomic Screening for Hereditary Breast and Ovarian Cancer in the United States. <i>JAMA Network Open</i> , 2020, 3, e2022874.	2.8	44
35	The Increasing Complexity of Preventive Services. <i>Journal of Women's Health</i> , 2020, 29, 748-749.	1.5	0
36	The Implementation Chasm Hindering Genome-informed Health Care. <i>Journal of Law, Medicine and Ethics</i> , 2020, 48, 119-125.	0.4	7
37	Identifying Ashkenazi Jewish BRCA1/2 founder variants in individuals who do not self-report Jewish ancestry. <i>Scientific Reports</i> , 2020, 10, 7669.	1.6	10

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38	Disparities in Genetic Testing and Care Among Black Women with Hereditary Breast Cancer. <i>Current Breast Cancer Reports</i> , 2020, 12, 125-131.	0.5	29
39	Association of Rare Pathogenic DNA Variants for Familial Hypercholesterolemia, Hereditary Breast and Ovarian Cancer Syndrome, and Lynch Syndrome With Disease Risk in Adults According to Family History. <i>JAMA Network Open</i> , 2020, 3, e203959.	2.8	75
40	Population-based Genetic Testing for Precision Prevention. <i>Cancer Prevention Research</i> , 2020, 13, 643-648.	0.7	20
41	Breast Cancer Risk Assessment. <i>Mayo Clinic Proceedings</i> , 2020, 95, 1268-1275.	1.4	12
42	Personalized early detection and prevention of breast cancer: ENVISION consensus statement. <i>Nature Reviews Clinical Oncology</i> , 2020, 17, 687-705.	12.5	178
43	Panel Testing for Hereditary Breast Cancer: More or Less?. <i>Current Breast Cancer Reports</i> , 2020, 12, 45-50.	0.5	0
44	Prevalence of Pathogenic Variants in Cancer Susceptibility Genes Among Women With Postmenopausal Breast Cancer. <i>JAMA - Journal of the American Medical Association</i> , 2020, 323, 995.	3.8	26
45	Peridiagnostic and cascade cancer genetic testing. <i>Nature Reviews Clinical Oncology</i> , 2020, 17, 277-278.	12.5	2
46	Use of an Online Breast Cancer Risk Assessment and Patient Decision Aid in Primary Care Practices. <i>Journal of Women's Health</i> , 2020, 29, 763-769.	1.5	15
47	Evaluation of Germline Genetic Testing Criteria in a Hospital-Based Series of Women With Breast Cancer. <i>Journal of Clinical Oncology</i> , 2020, 38, 1409-1418.	0.8	64
48	<p>Subsequent Development of Epithelial Ovarian Cancer After Ovarian Surgery for Benign Ovarian Tumor: A Population-Based Cohort Study</p>. <i>Clinical Epidemiology</i> , 2020, Volume 12, 637-649.	1.5	10
49	Implementation of interventions targeting the uptake of genetic testing services for breast cancer risk: protocol for a systematic review. <i>BMJ Open</i> , 2020, 10, e031727.	0.8	0
50	Updates in hereditary breast cancer genetic testing and practical high risk breast management in gene carriers. <i>Seminars in Oncology</i> , 2020, 47, 182-186.	0.8	5
51	Emerging Opportunity of Cascade Genetic Testing for Population-Wide Cancer Prevention and Control. <i>Journal of Clinical Oncology</i> , 2020, 38, 1371-1374.	0.8	18
52	Implications of the evidence for breast conservation therapy in BRCA-gene mutation carriers. <i>British Journal of Radiology</i> , 2020, 93, 20200038.	1.0	0
53	Disparities in BRCA counseling across providers in a diverse population of young breast cancer survivors. <i>Genetics in Medicine</i> , 2020, 22, 1088-1093.	1.1	10
54	Germline and Somatic Tumor Testing in Epithelial Ovarian Cancer: ASCO Guideline. <i>Journal of Clinical Oncology</i> , 2020, 38, 1222-1245.	0.8	202
55	Experts say more women should be tested for <i>BRCA1</i> and <i>BRCA2</i> genetic mutations. <i>Cancer</i> , 2020, 126, 693-693.	2.0	0

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56	Recommendations Related to Genetic Testing for Breast Cancer. JAMA - Journal of the American Medical Association, 2020, 323, 188.	3.8	0
57	Cascading After Peridiagnostic Cancer Genetic Testing: An Alternative to Population-Based Screening. Journal of Clinical Oncology, 2020, 38, 1398-1408.	0.8	60
58	Recommendations Related to Genetic Testing for Breast Cancer. JAMA - Journal of the American Medical Association, 2020, 323, 187.	3.8	0
59	Recommendations Related to Genetic Testing for Breast Cancerâ€”Reply. JAMA - Journal of the American Medical Association, 2020, 323, 188.	3.8	1
60	Accurate and Scalable Construction of Polygenic Scores in Large Biobank Data Sets. American Journal of Human Genetics, 2020, 106, 679-693.	2.6	80
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62	Genetic testing in male infertility â€” reassessing screening thresholds. Current Opinion in Urology, 2020, 30, 317-323.	0.9	7
63	Population Screening for Inherited Predisposition to Breast and Ovarian Cancer. Annual Review of Genomics and Human Genetics, 2020, 21, 373-412.	2.5	31
64	A Systematic Review on Cost-effectiveness Studies Evaluating Ovarian Cancer Early Detection and Prevention Strategies. Cancer Prevention Research, 2020, 13, 429-442.	0.7	10
65	Genetic counselorsâ€™ perspectives on populationâ€”based screening for <i>BRCA</i> -related hereditary breast and ovarian cancer and Lynch syndrome. Journal of Genetic Counseling, 2021, 30, 158-169.	0.9	8
66	Management of menopausal symptoms and ovarian function preservation in women with gynecological cancer. International Journal of Gynecological Cancer, 2021, 31, 352-359.	1.2	15
67	Characterizing germline APC and MUTYH variants in Ashkenazi Jews compared to other individuals. Familial Cancer, 2021, 20, 111-116.	0.9	5
68	Impact of Age, Race, and Socioeconomic Status on Women's Perceptions and Preferences Regarding Communication of Estimated Breast Cancer Risk. Academic Radiology, 2021, 28, 655-663.	1.3	4
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70	The role of genomics in global cancer prevention. Nature Reviews Clinical Oncology, 2021, 18, 116-128.	12.5	22
71	Rapid Genetic Testing for BRCA1 and BRCA2 Mutations at the Time of Breast Cancer Diagnosis: An Observational Study. Annals of Surgical Oncology, 2021, 28, 2219-2226.	0.7	7
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73	The current landscape of molecular profiling in the treatment of epithelial ovarian cancer. Gynecologic Oncology, 2021, 160, 333-345.	0.6	40

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74	Preimplantation genetic testing for carriers of BRCA1/2 pathogenic variants. <i>Critical Reviews in Oncology/Hematology</i> , 2021, 157, 103201.	2.0	26
75	A state-based approach to genomics for rare disease and population screening. <i>Genetics in Medicine</i> , 2021, 23, 777-781.	1.1	19
76	Training courses on hereditary breast and ovarian cancer to strengthen cross-sectoral care in underserved areas. <i>Patient Education and Counseling</i> , 2021, 104, 1431-1437.	1.0	2
77	Experiences and attitudes of hereditary cancer screening patients in a consumer directed testing model. <i>Patient Education and Counseling</i> , 2021, 104, 473-479.	1.0	2
78	Hereditary Gynecological Malignancy and Molecular Features. <i>Current Human Cell Research and Applications</i> , 2021, , 145-165.	0.1	0
79	Management of a Woman at Elevated Risk for Breast Cancer. , 2021, , 107-136.		0
80	Hereditary Cancer Counseling and Germline Genetic Testing. , 2021, , 305-317.		0
81	Timing and type of menopause and risk of cardiovascular disease. <i>Menopause</i> , 2021, 28, 477-479.	0.8	1
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83	A Population-Based Study of Genes Previously Implicated in Breast Cancer. <i>New England Journal of Medicine</i> , 2021, 384, 440-451.	13.9	414
84	Assessing Risk of Breast Cancer: A Review of Risk Prediction Models. <i>Journal of Breast Imaging</i> , 2021, 3, 144-155.	0.5	47
85	A prospective controlled study of sexual function and sexually related personal distress up to 12 months after premenopausal risk-reducing bilateral salpingo-oophorectomy. <i>Menopause</i> , 2021, 28, 748-755.	0.8	6
86	Clinical practice guidelines for BRCA1 and BRCA2 genetic testing. <i>European Journal of Cancer</i> , 2021, 146, 30-47.	1.3	81
87	Should the BCRA1/2-mutations healthy carriers be valid candidates for hematopoietic stem cell donation?. <i>Hereditary Cancer in Clinical Practice</i> , 2021, 19, 22.	0.6	0
88	Controversies in Hereditary Cancer Management. <i>Obstetrics and Gynecology</i> , 2021, 137, 941-955.	1.2	4
89	The Screen Project: Guided Direct-To-Consumer Genetic Testing for Breast Cancer Susceptibility in Canada. <i>Cancers</i> , 2021, 13, 1894.	1.7	8
91	Clinical management among individuals with variant of uncertain significance in hereditary cancer: A systematic review and meta-analysis. <i>Clinical Genetics</i> , 2021, 100, 119-131.	1.0	12
93	Outcomes of incidentally detected ovarian cancers diagnosed at time of risk-reducing salpingo-oophorectomy in BRCA mutation carriers. <i>Gynecologic Oncology</i> , 2021, 161, 521-526.	0.6	2

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94	Clinical validity and utility of preconception expanded carrier screening for the management of reproductive genetic risk in IVF and general population. <i>Human Reproduction</i> , 2021, 36, 2050-2061.	0.4	27
95	A Collaborative Model to Implement Flexible, Accessible and Efficient Oncogenetic Services for Hereditary Breast and Ovarian Cancer: The C-MOnGene Study. <i>Cancers</i> , 2021, 13, 2729.	1.7	8
96	Cancer Progress and Priorities: Breast Cancer. <i>Cancer Epidemiology Biomarkers and Prevention</i> , 2021, 30, 822-844.	1.1	47
97	Current status and future directions of U.S. genomic nursing health care policy. <i>Nursing Outlook</i> , 2021, 69, 471-488.	1.5	8
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103	Optimization of prediction methods for risk assessment of pathogenic germline variants in the Japanese population. <i>Cancer Science</i> , 2021, 112, 3338-3348.	1.7	3
104	Disparities in Breast Cancer Associated With African American Identity. <i>American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting</i> , 2021, 41, e29-e46.	1.8	27
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106	Association of Family History with the Development of Breast Cancer: A Cohort Study of 129,374 Women in KoGES Data. <i>International Journal of Environmental Research and Public Health</i> , 2021, 18, 6409.	1.2	5
107	A pitfall in targeted Sanger sequencing of BRCA splicing variants in at-risk individuals. <i>Pathology Research and Practice</i> , 2021, 222, 153456.	1.0	0
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110	Oncologic Anthropology: An Interdisciplinary Approach to Understanding the Association Between Genetically Defined African Ancestry and Susceptibility for Triple Negative Breast Cancer. <i>Current Breast Cancer Reports</i> , 2021, 13, 247-258.	0.5	7
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112	Nanotechnology in Tumor Biomarker Detection: The Potential of Liganded Nanoclusters as Nonlinear Optical Contrast Agents for Molecular Diagnostics of Cancer. <i>Cancers</i> , 2021, 13, 4206.	1.7	27
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119	Management of breast cancer patients with BRCA gene mutations in Lebanon of the Middle East: perspectives and challenges. <i>Hospital Practice (1995)</i> , 2021, , 1-5.	0.5	0
120	Clinical Implications of Combinatorial Pharmacogenomic Tests Based on Cytochrome P450 Variant Selection. <i>Frontiers in Genetics</i> , 2021, 12, 719671.	1.1	8
121	Design and user experience testing of a polygenic score report: a qualitative study of prospective users. <i>BMC Medical Genomics</i> , 2021, 14, 238.	0.7	29
122	The Need to Improve the Clinical Utility of Direct-to-Consumer Genetic Tests. <i>JAMA - Journal of the American Medical Association</i> , 2020, 323, 1443.	3.8	5
124	Assessment of and Interventions for Women at High Risk for Breast or Ovarian Cancer: A Survey of Primary Care Physicians. <i>Cancer Prevention Research</i> , 2021, 14, 205-214.	0.7	4
125	Mental Illness and BRCA1/2 Genetic Testing Intention Among Multiethnic Women Undergoing Screening Mammography. <i>Oncology Nursing Forum</i> , 2020, 47, E13-E24.	0.5	4
126	Effects of Marital Status on Prognosis in Women with Infiltrating Ductal Carcinoma of the Breast: A Real-World 1: 1 Propensity-Matched Study. <i>Medical Science Monitor</i> , 2020, 26, e923630.	0.5	6
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131	USPSTF Approach to Addressing Sex and Gender When Making Recommendations for Clinical Preventive Services. <i>JAMA - Journal of the American Medical Association</i> , 2021, 326, 1953.	3.8	27
132	Multiple imputation with missing data indicators. <i>Statistical Methods in Medical Research</i> , 2021, 30, 2685-2700.	0.7	30
133	Positive experiences of healthcare professionals with a mainstreaming approach of germline genetic testing for women with ovarian cancer. <i>Familial Cancer</i> , 2022, 21, 295-304.	0.9	15
134	Disparities in Genetic Testing for Heritable Solid-Tumor Malignancies. <i>Surgical Oncology Clinics of North America</i> , 2022, 31, 109-126.	0.6	9
135	Identification of women at risk of hereditary breast-ovarian cancer among participants in a population-based breast cancer screening. <i>Familial Cancer</i> , 2021, , 1.	0.9	1
136	Fertility Considerations for Reproductive-Aged Carriers of Deleterious BRCA Mutations: A Call for Early Intervention. <i>JCO Oncology Practice</i> , 2022, 18, 165-168.	1.4	4
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140	Hereditary Cancer Risk Using a Genetic Chatbot Before Routine Care Visits. <i>Obstetrics and Gynecology</i> , 2021, 138, 860-870.	1.2	37
141	Survey on Physicians' Knowledge and Training Needs in Genetic Counseling in Germany. <i>Breast Care</i> , 2021, 16, 389-395.	0.8	0
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144	Implication and Influence of Multigene Panel Testing with Genetic Counseling in Korean Patients with BRCA1/2 Mutation-Negative Breast Cancer. <i>Cancer Research and Treatment</i> , 2022, 54, 1099-1110.	1.3	1
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148	Survey on Physicians' Knowledge and Training Needs in Genetic Counseling in Germany. <i>Breast Care</i> , 2021, 16, 389-395.	0.8	3
149	Self-rated family health history knowledge among All of Us program participants. <i>Genetics in Medicine</i> , 2022, 24, 955-961.	1.1	8

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150	Automated Clinical Practice Guideline Recommendations for Hereditary Cancer Risk Using Chatbots and Ontologies: System Description. <i>JMIR Cancer</i> , 2022, 8, e29289.	0.9	3
151	Psychiatric symptoms in a Spanish sample with hereditary cancer risk. <i>Journal of Community Genetics</i> , 2022, , 1.	0.5	0
152	Retrospective assessment of barriers and access to genetic services for hereditary cancer syndromes in an integrated health care delivery system. <i>Hereditary Cancer in Clinical Practice</i> , 2022, 20, 7.	0.6	7
153	Hereditary Breast and Ovarian Cancer. <i>Obstetrics and Gynecology Clinics of North America</i> , 2022, 49, 117-147.	0.7	3
154	Comparison of a Cancer Family History Collection and Risk Assessment Tool “ ItRunsInMyFamily ” with Risk Assessment by Health-Care Professionals. <i>Public Health Genomics</i> , 2022, 25, 80-88.	0.6	2
155	Hereditary breast cancer “ what we have learned in the last decade. <i>Mastology</i> , 0, 31, .	0.1	1
156	A universal probe system for low-abundance point mutation detection based on endonuclease IV. <i>Analyst, The</i> , 2022, 147, 1534-1539.	1.7	1
157	Psychosocial outcome and health behaviour intent of breast cancer patients with BRCA1/2 and PALB2 pathogenic variants unselected by a priori risk. <i>PLoS ONE</i> , 2022, 17, e0263675.	1.1	0
158	Prevalence of Tumor Genomic Alterations in Homologous Recombination Repair Genes Among Taiwanese Breast Cancers. <i>Annals of Surgical Oncology</i> , 2022, 29, 3578-3590.	0.7	9
160	Factors Associated with Mammography Screening Choices by Women Aged 40“49 at Average Risk. <i>Journal of Women's Health</i> , 2022, 31, 1120-1126.	1.5	5
161	Hereditary gynecologic tumors and precision cancer medicine. <i>Journal of Obstetrics and Gynaecology Research</i> , 2022, 48, 1076-1090.	0.6	0
162	IPRS: Leveraging Gene-Environment Interaction to Reconstruct Polygenic Risk Score. <i>Frontiers in Genetics</i> , 2022, 13, 801397.	1.1	4
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