Claire E Wakefield

List of Publications by Year in descending order

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CLAIDE E MAKEELELD

#	Article	IF	CITATIONS
1	Barriers and facilitators of exercise experienced by cancer survivors: a mixed methods systematic review. Supportive Care in Cancer, 2018, 26, 685-700.	2.2	172
2	Parental adjustment to the completion of their child's cancer treatment. Pediatric Blood and Cancer, 2011, 56, 524-531.	1.5	145
3	A systematic review of psychological interventions for adolescents and young adults living with chronic illness Health Psychology, 2012, 31, 380-393.	1.6	130
4	Understanding the school experiences of children and adolescents with serious chronic illness: a systematic metaâ€review. Child: Care, Health and Development, 2017, 43, 645-662.	1.7	125
5	The Psychosocial Impact of Completing Childhood Cancer Treatment: A Systematic Review of the Literature. Journal of Pediatric Psychology, 2010, 35, 262-274.	2.1	121
6	Primary Care Physicians' Perspectives of Their Role in Cancer Care: A Systematic Review. Journal of General Internal Medicine, 2016, 31, 1222-1236.	2.6	115
7	Attitudes toward Organ Donation and Donor Behavior: A Review of the International Literature. Progress in Transplantation, 2010, 20, 380-391.	0.7	114
8	Health Literacy in Adolescents and Young Adults: An Updated Review. Journal of Adolescent and Young Adult Oncology, 2016, 5, 106-118.	1.3	113
9	Hospital-based bereavement services following the death of a child: A mixed study review. Palliative Medicine, 2015, 29, 193-210.	3.1	103
10	Survivorship Care Plans in Cancer: A Meta-Analysis and Systematic Review of Care Plan Outcomes. Oncologist, 2020, 25, e351-e372.	3.7	101
11	Supporting children facing a parent's cancer diagnosis: a systematic review of children's psychosocial needs and existing interventions. European Journal of Cancer Care, 2017, 26, e12432.	1.5	89
12	Attitudes toward organ donation and donor behavior: a review of the international literature. Progress in Transplantation, 2010, 20, 380-391.	0.7	85
13	The psychological impact of genetic information on children: a systematic review. Genetics in Medicine, 2016, 18, 755-762.	2.4	83
14	Family information needs at childhood cancer treatment completion. Pediatric Blood and Cancer, 2012, 58, 621-626.	1.5	81
15	Educational and vocational goal disruption in adolescent and young adult cancer survivors. Psycho-Oncology, 2018, 27, 532-538.	2.3	74
16	Online group-based cognitive-behavioural therapy for adolescents and young adults after cancer treatment: A multicenter randomised controlled trial of Recapture Life-AYA. BMC Cancer, 2012, 12, 339.	2.6	71
17	Assessment of Psychosocial Outcomes in Genetic Counseling Research: An Overview of Available Measurement Scales. Journal of Genetic Counseling, 2007, 16, 693-712.	1.6	70
18	A randomized controlled trial of a decision aid for women considering genetic testing for breast and ovarian cancer risk. Breast Cancer Research and Treatment, 2008, 107, 289-301.	2.5	70

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19	Fertility concerns among child and adolescent cancer survivors and their parents: A qualitative analysis. Journal of Psychosocial Oncology, 2016, 34, 347-362.	1.2	70
20	"Forewarned and forearmedâ€: Long-term childhood cancer survivors' and parents' information needs and implications for survivorship models of care. Patient Education and Counseling, 2017, 100, 355-363.	2.2	69
21	The impact of long-term follow-up care for childhood cancer survivors: A systematic review. Critical Reviews in Oncology/Hematology, 2017, 114, 131-138.	4.4	68
22	Returning to School After Adolescent Cancer: A Qualitative Examination of Australian Survivors' and Their Families' Perspectives. Journal of Adolescent and Young Adult Oncology, 2011, 1, 87-94.	1.3	66
23	Dietary intake after treatment in child cancer survivors. Pediatric Blood and Cancer, 2012, 58, 752-757.	1.5	66
24	Distress and adjustment among adolescents and young adults with cancer: an empirical and conceptual review. Translational Pediatrics, 2013, 2, 167-97.	1.2	66
25	Participation in psychosocial oncology and quality-of-life research: a systematic review. Lancet Oncology, The, 2017, 18, e153-e165.	10.7	64
26	Healthâ€related quality of life of children on treatment for acute lymphoblastic leukemia: A systematic review. Pediatric Blood and Cancer, 2017, 64, e26489.	1.5	62
27	Health-related quality of life of survivors of childhood acute lymphoblastic leukemia: a systematic review. Quality of Life Research, 2018, 27, 1431-1443.	3.1	60
28	It's More than Dollars and Cents: The Impact of Childhood Cancer on Parents' Occupational and Financial Health. Journal of Psychosocial Oncology, 2014, 32, 602-621.	1.2	59
29	Acceptability and feasibility of an e-mental health intervention for parents of childhood cancer survivors: "Cascade― Supportive Care in Cancer, 2016, 24, 2685-2694.	2.2	55
30	School Experiences of Siblings of Children with Chronic Illness: A Systematic Literature Review. Journal of Pediatric Nursing, 2017, 33, 23-32.	1.5	53
31	Cancer care disruption and reorganisation during the COVID-19 pandemic in Australia: A patient, carer and healthcare worker perspective. PLoS ONE, 2021, 16, e0257420.	2.5	52
32	Randomized trial of a decision aid for individuals considering genetic testing for hereditary nonpolyposis colorectal cancer risk. Cancer, 2008, 113, 956-965.	4.1	51
33	Decision-making in childhood cancer: parents' and adolescents' views and perceptions. Supportive Care in Cancer, 2019, 27, 4331-4340.	2.2	49
34	Childhood cancer survivors' school (re)entry: Australian parents' perceptions. European Journal of Cancer Care, 2013, 22, 484-492.	1.5	48
35	Support after the completion of cancer treatment: perspectives of Australian adolescents and their families. European Journal of Cancer Care, 2013, 22, 530-539.	1.5	47
36	Narrative Review of the Educational, Vocational, and Financial Needs of Adolescents and Young Adults with Cancer: Recommendations for Support and Research. Journal of Adolescent and Young Adult Oncology, 2018, 7, 143-147.	1.3	47

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37	Development and Pilot Testing of Two Decision Aids for Individuals Considering Genetic Testing for Cancer Risk. Journal of Genetic Counseling, 2007, 16, 325-339.	1.6	46
38	A randomized trial of a breast/ovarian cancer genetic testing decision aid used as a communication aid during genetic counseling. Psycho-Oncology, 2008, 17, 844-854.	2.3	46
39	Patient-reported depression measures in cancer: a meta-review. Lancet Psychiatry,the, 2015, 2, 635-647.	7.4	46
40	Feasibility, acceptability, and safety of the Recapture Life videoconferencing intervention for adolescent and young adult cancer survivors. Psycho-Oncology, 2019, 28, 284-292.	2.3	46
41	Psychological Outcomes, Health-Related Quality of Life, and Neurocognitive Functioning in Survivors of Childhood Cancer and Their Parents. Pediatric Clinics of North America, 2020, 67, 1103-1134.	1.8	45
42	Parent feeding interactions and practices during childhood cancer treatment. A qualitative investigation. Appetite, 2015, 89, 219-225.	3.7	43
43	Development and Pilot Testing of a Fertility Decision Aid for Young Women Diagnosed with Early Breast Cancer. Breast Journal, 2011, 17, 112-114.	1.0	42
44	Religious and ethnic influences on willingness to donate organs and donor behavior: an Australian perspective. Progress in Transplantation, 2011, 21, 161-168.	0.7	42
45	Cognitive Compensations for Blindness in Children: An Investigation Using Odour Naming. Perception, 2004, 33, 429-442.	1.2	41
46	Quantifying family dissemination and identifying barriers to communication of risk information in Australian BRCA families. Genetics in Medicine, 2017, 19, 1323-1331.	2.4	41
47	A Narrative Review of Models of Care for Adolescents and Young Adults with Cancer: Barriers and Recommendations. Journal of Adolescent and Young Adult Oncology, 2018, 7, 148-152.	1.3	41
48	Financial toxicity of childhood cancer and changes to parents' employment after treatment completion. Pediatric Blood and Cancer, 2020, 67, e28345.	1.5	41
49	Models of childhood cancer survivorship care in Australia and New Zealand: Strengths and challenges. Asia-Pacific Journal of Clinical Oncology, 2017, 13, 407-415.	1.1	40
50	Results of a Nurse-Led Intervention. Journal of Pediatric Oncology Nursing, 2013, 30, 333-341.	1.5	39
51	Distance-delivered physical activity interventions for childhood cancer survivors: A systematic review and meta-analysis. Critical Reviews in Oncology/Hematology, 2017, 118, 27-41.	4.4	39
52	Communication and Consent: Discussion and Organ Donation Decisions for Self and Family. Transplantation Proceedings, 2013, 45, 10-12.	0.6	37
53	Perceived cancer-related pain and fatigue, information needs, and fear of cancer recurrence among adult survivors of childhood cancer. Patient Education and Counseling, 2019, 102, 2270-2278.	2.2	37
54	Sexual and Romantic Relationships: Experiences of Adolescent and Young Adult Cancer Survivors. Journal of Adolescent and Young Adult Oncology, 2016, 5, 286-291.	1.3	36

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55	School students with chronic illness have unmet academic, social, and emotional school needs School Psychology, 2019, 34, 627-636.	2.4	35
56	Neuropsychology reports for childhood brain tumor survivors: Implementation of recommendations at home and school. Pediatric Blood and Cancer, 2014, 61, 1080-1087.	1.5	34
57	Parents' attitudes toward genetic testing of children for health conditions: A systematic review. Clinical Genetics, 2017, 92, 569-578.	2.0	34
58	The Role of Primary Care Physicians in Childhood Cancer Survivorship Care: Multiperspective Interviews. Oncologist, 2019, 24, 710-719.	3.7	34
59	Ethical and Clinical Challenges Delivering Groupâ€based Cognitiveâ€Behavioural Therapy to Adolescents and Young Adults with Cancer Using Videoconferencing Technology. Australian Psychologist, 2015, 50, 271-278.	1.6	33
60	Nutritional interventions for survivors of childhood cancer. The Cochrane Library, 2016, 2016, CD009678.	2.8	33
61	Family Communication, Risk Perception and Cancer Knowledge of Young Adults from <i>BRCA1/2</i> Families: a Systematic Review. Journal of Genetic Counseling, 2017, 26, 1179-1196.	1.6	33
62	Parent-targeted home-based interventions for increasing fruit and vegetable intake in children: a systematic review and meta-analysis. Nutrition Reviews, 2018, 76, 154-173.	5.8	33
63	Consensus Among International Ethical Guidelines for the Provision of Videoconferencing-Based Mental Health Treatments. JMIR Mental Health, 2016, 3, e17.	3.3	31
64	An exploration of the communication preferences regarding genetic testing in individuals from families with identified breast/ovarian cancer mutations. Familial Cancer, 2011, 10, 97-105.	1.9	30
65	Parental sleep experiences on the pediatric oncology ward. Supportive Care in Cancer, 2013, 21, 557-564.	2.2	30
66	Grandparents of children with cancer: a controlled study of distress, support, and barriers to care. Psycho-Oncology, 2014, 23, 855-861.	2.3	30
67	Online parent-targeted cognitive-behavioural therapy intervention to improve quality of life in families of young cancer survivors: study protocol for a randomised controlled trial. Trials, 2015, 16, 153.	1.6	30
68	Changes in body mass index in long-term survivors of childhood acute lymphoblastic leukemia treated without cranial radiation and with reduced glucocorticoid therapy. Pediatric Blood and Cancer, 2017, 64, e26344.	1.5	29
69	Attitudes and experiences of childhood cancer survivors transitioning from pediatric care to adult care. Supportive Care in Cancer, 2018, 26, 2743-2750.	2.2	29
70	Psychometric evaluation of the Sibling Cancer Needs Instrument (SCNI): an instrument to assess the psychosocial unmet needs of young people who are siblings of cancer patients. Supportive Care in Cancer, 2014, 22, 653-665.	2.2	27
71	What About School? Educational Challenges for Children and Adolescents With Cancer. Educational and Developmental Psychologist, 2015, 32, 23-40.	0.7	27
72	Cancer survivorship services for indigenous peoples: where we stand, where to improve? A systematic review. Journal of Cancer Survivorship, 2016, 10, 330-341.	2.9	27

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73	Strategies to facilitate shared decision-making about pediatric oncology clinical trial enrollment: A systematic review. Patient Education and Counseling, 2018, 101, 1157-1174.	2.2	27
74	Children and young people's understanding of inherited conditions and their attitudes towards genetic testing: A systematic review. Clinical Genetics, 2019, 95, 10-22.	2.0	27
75	Parent, patient and health professional perspectives regarding enteral nutrition in paediatric oncology. Nutrition and Dietetics, 2017, 74, 476-487.	1.8	26
76	"l remember how I felt, but I don't remember the gene― Families' experiences of cancerâ€related genetic testing in childhood. Pediatric Blood and Cancer, 2019, 66, e27762.	1.5	26
77	Development and pilot testing of an online screening decision aid for men with a family history of prostate cancer. Patient Education and Counseling, 2011, 83, 64-72.	2.2	25
78	A Study Protocol for the Australasian Oncofertility Registry: Monitoring Referral Patterns and the Uptake, Quality, and Complications of Fertility Preservation Strategies in Australia and New Zealand. Journal of Adolescent and Young Adult Oncology, 2016, 5, 215-225.	1.3	25
79	Therapeutic Alliance and Group Cohesion in an Online Support Program for Adolescent and Young Adult Cancer Survivors: Lessons from "Recapture Life― Journal of Adolescent and Young Adult Oncology, 2017, 6, 568-572.	1.3	25
80	Talking across generations: Family communication about <i>BRCA1</i> and <i>BRCA2</i> genetic cancer risk. Journal of Genetic Counseling, 2019, 28, 516-532.	1.6	25
81	Interactive video games to reduce paediatric procedural pain and anxiety: a systematic review and meta-analysis. British Journal of Anaesthesia, 2021, 127, 608-619.	3.4	25
82	Recommendations for the surveillance of mental health problems in childhood, adolescent, and young adult cancer survivors: a report from the International Late Effects of Childhood Cancer Guideline Harmonization Group. Lancet Oncology, The, 2022, 23, e184-e196.	10.7	25
83	Adolescent and young adult cancer survivors' memory and future thinking processes place them at risk for poor mental health. Psycho-Oncology, 2018, 27, 2709-2716.	2.3	24
84	Barriers and enablers to physical activity and aerobic fitness deficits among childhood cancer survivors. Pediatric Blood and Cancer, 2020, 67, e28339.	1.5	24
85	Long-term care for people treated for cancer during childhood and adolescence. Lancet, The, 2022, 399, 1561-1572.	13.7	24
86	Women's Knowledge, Beliefs, and Information Needs in Relation to the Risks and Benefits Associated with Use of the Oral Contraceptive Pill. Journal of Women's Health, 2011, 20, 635-642.	3.3	23
87	Impact of physical appearance changes reported by adolescent and young adult cancer survivors: A qualitative analysis. European Journal of Cancer Care, 2019, 28, e13052.	1.5	23
88	A Systematic Review Summarizing the State of Evidence on Bullying in Childhood Cancer Patients/Survivors. Journal of Pediatric Oncology Nursing, 2019, 36, 55-68.	1.5	23
89	The $6\hat{a}\in$ minute walk test is a good predictor of cardiorespiratory fitness in childhood cancer survivors when access to comprehensive testing is limited. International Journal of Cancer, 2020, 147, 847-855.	5.1	23
90	Online prostate cancer screening decision aid for at-risk men: A randomized trial Health Psychology, 2014, 33, 986-997.	1.6	22

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91	Challenges and strategies proposed by genetic health professionals to assist with family communication. European Journal of Human Genetics, 2019, 27, 1630-1638.	2.8	22
92	Supportive oncofertility care, psychological health and reproductive concerns: a qualitative study. Supportive Care in Cancer, 2020, 28, 809-817.	2.2	22
93	The information needs of parents of children with early-onset epilepsy: A systematic review. Epilepsy and Behavior, 2020, 112, 107382.	1.7	22
94	"We needed thisâ€: perspectives of parents and healthcare professionals involved in a pilot newborn screening program for spinal muscular atrophy. EClinicalMedicine, 2021, 33, 100742.	7.1	22
95	Time Perspective in Hereditary Cancer: Psychometric Properties of a Short Form of the Zimbardo Time Perspective Inventory in a Community and Clinical Sample. Genetic Testing and Molecular Biomarkers, 2010, 14, 617-627.	0.7	21
96	"For All My Family's Sake, I Should Go and Find Out― An Australian Report on Genetic Counseling and Testing Uptake in Individuals at High Risk of Breast and/or Ovarian Cancer. Genetic Testing and Molecular Biomarkers, 2011, 15, 379-385.	0.7	21
97	Strategies to improve adherence to treatment in adolescents and young adults with cancer: a systematic review. Clinical Oncology in Adolescents and Young Adults, 0, , 35.	0.8	21
98	Online, Group-Based Psychological Support for Adolescent and Young Adult Cancer Survivors: Results from the Recapture Life Randomized Trial. Cancers, 2021, 13, 2460.	3.7	21
99	Usefulness of the Threatening Medical Situations Inventory in individuals considering genetic testing for cancer risk. Patient Education and Counseling, 2007, 69, 29-38.	2.2	20
100	Physical activity and screenâ€ŧime of childhood haematopoietic stem cell transplant survivors. Acta Paediatrica, International Journal of Paediatrics, 2015, 104, e455-9.	1.5	20
101	Health Literacy in Adolescents and Young Adults: Perspectives from Australian Cancer Survivors. Journal of Adolescent and Young Adult Oncology, 2017, 6, 150-158.	1.3	20
102	Decision Aid for Nutrition Support in Pediatric Oncology: A Pilot Study. Journal of Parenteral and Enteral Nutrition, 2017, 41, 1336-1347.	2.6	20
103	Cancer-Related Genetic Testing and Personalized Medicine for Adolescents: A Narrative Review of Impact and Understanding. Journal of Adolescent and Young Adult Oncology, 2018, 7, 259-262.	1.3	20
104	Parents' experiences of requests for organ and tissue donation: the value of asking. Archives of Disease in Childhood, 2019, 104, 837-843.	1.9	20
105	"Balancing Expectations with Actual Realities― Conversations with Clinicians and Scientists in the First Year of a High-Risk Childhood Cancer Precision Medicine Trial. Journal of Personalized Medicine, 2020, 10, 9.	2.5	20
106	â€~What they're not telling you': A new scale to measure grandparents' information needs when their grandchild has cancer. Patient Education and Counseling, 2014, 94, 351-355.	2.2	19
107	Exploring the views of parents regarding dietary habits of their young cancer-surviving children. Supportive Care in Cancer, 2015, 23, 463-471.	2.2	19
108	Longâ€ŧerm healthâ€ŧelated quality of life in young childhood cancer survivors and their parents. Pediatric Blood and Cancer, 2021, 68, e29398.	1.5	19

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109	Utilizing the Experience of Consumers in Consultation to Develop the Australasian Oncofertility Consortium Charter. Journal of Adolescent and Young Adult Oncology, 2016, 5, 232-239.	1.3	18
110	Grandparents of children with cancer: a controlled comparison of perceived family functioning. Supportive Care in Cancer, 2019, 27, 2087-2094.	2.2	18
111	Understanding parents' communication experiences in childhood cancer: a qualitative exploration and model for future research. Supportive Care in Cancer, 2020, 28, 4467-4476.	2.2	18
112	Psychosocial impact of genetic testing on parents of children with developmental and epileptic encephalopathy. Developmental Medicine and Child Neurology, 2022, 64, 95-104.	2.1	18
113	Healthcare professionals' attitudes toward cancer precision medicine: A systematic review. Seminars in Oncology, 2019, 46, 291-303.	2.2	17
114	The â€~value' of values clarification in cancer-related decision aids. Patient Education and Counseling, 2013, 90, 281-283.	2.2	16
115	How Parents of Childhood Cancer Survivors Perceive Support From Their Extended Families. Journal of Child and Family Studies, 2019, 28, 1537-1547.	1.3	16
116	Childhood cancer survivorship: barriers and preferences. BMJ Supportive and Palliative Care, 2019, , bmjspcare-2019-002001.	1.6	16
117	Siblings of young people with chronic illness: Caring responsibilities and psychosocial functioning. Journal of Child Health Care, 2022, 26, 581-596.	1.4	16
118	Facilitating engagement with school in students with chronic illness through positive education: A mixed-methods comparison study School Psychology, 2019, 34, 677-686.	2.4	16
119	Parents' Perspectives on Their Child's Social Experience in the Context of Childhood Chronic Illness: A Qualitative Study. Journal of Pediatric Nursing, 2018, 42, e10-e18.	1.5	15
120	Research Report: Early Blindness May be Associated with Changes in Performance on Verbal Fluency Tasks. Journal of Visual Impairment and Blindness, 2006, 100, 306-310.	0.7	14
121	The association of psychosocial screening and service provision in pediatric oncology: the Psychosocial Assessment Tool (PAT2.0) into clinical practice. Supportive Care in Cancer, 2016, 24, 2945-52.	2.2	14
122	â€~Re-engage' pilot study protocol: a nurse-led eHealth intervention to re-engage, educate and empower childhood cancer survivors. BMJ Open, 2018, 8, e022269.	1.9	14
123	Piloting a parent and patient decision aid to support clinical trial decision making in childhood cancer. Psycho-Oncology, 2019, 28, 1520-1529.	2.3	14
124	Reproductive Care of Childhood and Adolescent Cancer Survivors: A 12-Year Evaluation. Journal of Adolescent and Young Adult Oncology, 2021, 10, 131-141.	1.3	14
125	A sociology of precisionâ€inâ€practice: The affective and temporal complexities of everyday clinical care. Sociology of Health and Illness, 2021, 43, 2178-2195.	2.1	14
126	Improving mutation notification when new genetic information is identified in research: a trial of two strategies in familial breast cancer. Genetics in Medicine, 2013, 15, 187-194.	2.4	13

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127	A study protocol for Truce: a pragmatic controlled trial of a seven-week acceptance and commitment therapy program for young people who have a parent with cancer. BMC Psychology, 2015, 3, 31.	2.1	13
128	Grandparents of children with cancer: Quality of life, medication and hospitalizations. Pediatric Blood and Cancer, 2017, 64, 163-171.	1.5	13
129	How physically active do Australian and New Zealander childhood cancer survivors perceive themselves? A report from the ANZCHOG survivorship study. Complementary Therapies in Medicine, 2019, 44, 196-203.	2.7	13
130	The Development of Delta: Using Agile to Develop a Decision Aid for Pediatric Oncology Clinical Trial Enrollment. JMIR Research Protocols, 2018, 7, e119.	1.0	13
131	Re-Engage: A Novel Nurse-Led Program for Survivors of Childhood Cancer Who Are Disengaged From Cancer-Related Care. Journal of the National Comprehensive Cancer Network: JNCCN, 2020, 18, 1067-1074.	4.9	13
132	Accommodation in pediatric oncology: parental experiences, preferences and unmet needs. Rural and Remote Health, 2013, 13, 2005.	0.5	13
133	Development and Pilot Testing of a Decision Aid for Men Considering Genetic Testing for Breast and/or Ovarian Cancer–Related Mutations (BRCA1/2). Genetic Testing and Molecular Biomarkers, 2008, 12, 523-532.	1.7	12
134	Management of fever and neutropenia in children with cancer. Supportive Care in Cancer, 2015, 23, 2079-2087.	2.2	12
135	Establishing Australian school reâ€entry service guidelines for children diagnosed with cancer. Journal of Paediatrics and Child Health, 2017, 53, 529-533.	0.8	12
136	The Use of Activity Trackers in Interventions for Childhood Cancer Patients and Survivors: A Systematic Review. Journal of Adolescent and Young Adult Oncology, 2021, 10, 1-14.	1.3	12
137	Systematic Review of Educational Supports of Pediatric Cancer Survivors: Current Approaches and Future Directions. Journal of Clinical Oncology, 2021, 39, 1813-1823.	1.6	12
138	Issues Faced by Unaffected Men With a Family History of Prostate Cancer: A Multidisciplinary Overview. Journal of Urology, 2008, 180, 38-46.	0.4	11
139	Paediatric oncology patient preference for oral nutritional supplements in a clinical setting. Supportive Care in Cancer, 2011, 19, 1289-1296.	2.2	11
140	Assessment of Fundamental Movement Skills in Childhood Cancer Patients. Pediatric Blood and Cancer, 2015, 62, 2211-2215.	1.5	11
141	Development of Breast Cancer Choices: a decision support tool for young women with breast cancer deciding whether to have genetic testing for BRCA1/2 mutations. Supportive Care in Cancer, 2019, 27, 297-309.	2.2	11
142	Brief report: Bereaved parents informing research design: The place of a pilot study. Death Studies, 2019, 43, 62-69.	2.7	11
143	The COVIDâ€19 pandemic: Distanceâ€delivered care for childhood cancer survivors. Pediatric Blood and Cancer, 2020, 67, e28715.	1.5	11
144	Aligning intuition and theory: enhancing the replicability of behaviour change interventions in cancer genetics. Implementation Science Communications, 2020, 1, 90.	2.2	11

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145	Attitudes Toward Genetic Testing for Cancer Risk after Genetic Counseling and Decision Support: A Qualitative Comparison between Hereditary Cancer Types. Genetic Testing and Molecular Biomarkers, 2007, 11, 401-412.	1.7	10
146	Development and Evaluation of an Information Booklet for Grandparents of Children With Cancer. Journal of Pediatric Oncology Nursing, 2016, 33, 361-369.	1.5	10
147	Variables associated with grief and personal growth following the death of a child from cancer: A mixed method analysis. Death Studies, 2021, 45, 702-713.	2.7	10
148	Paediatric surgery for childhood cancer: Lasting experiences and needs of children and parents. European Journal of Cancer Care, 2019, 28, e13116.	1.5	10
149	The Cost of Cure: Chronic Conditions in Survivors of Child, Adolescent, and Young Adult Cancers. , 2016, , 371-420.		10
150	Patterns and drivers of health care use in long-term childhood cancer survivors: A systematic review. Critical Reviews in Oncology/Hematology, 2017, 120, 60-76.	4.4	9
151	Something's got to give: time–cost tradeâ€offs in siteâ€specific research approval can negatively impact patient recruitment in multiâ€institutional studies. Internal Medicine Journal, 2017, 47, 1088-1089.	0.8	9
152	Family historyâ€ŧaking practices and genetic confidence in primary and tertiary care providers for childhood cancer survivors. Pediatric Blood and Cancer, 2018, 65, e26923.	1.5	9
153	Childhood cancer survivors report preferring lifestyle interventions delivered in person rather than online: An adolescent and parent perspective. Pediatric Blood and Cancer, 2019, 66, e27922.	1.5	9
154	Dietary intake and diet quality in children receiving treatment for cancer. Nutrition Reviews, 2019, 77, 267-277.	5.8	9
155	Familial diagnostic experiences in paediatric oncology. British Journal of Cancer, 2015, 112, 20-23.	6.4	8
156	Transition of childhood cancer survivors to adult care: The survivor perspective. Pediatric Blood and Cancer, 2017, 64, e26354.	1.5	8
157	Recruiting primary care physicians to qualitative research: Experiences and recommendations from a childhood cancer survivorship study. Pediatric Blood and Cancer, 2018, 65, e26762.	1.5	8
158	Adolescent and Young Adult Cancer Survivors' Experiences of the Healthcare System: A Qualitative Study. Journal of Adolescent and Young Adult Oncology, 2018, 7, 88-96.	1.3	8
159	Health professional and atâ€risk BRCA young adult perspectives about information needs: What does Gen Y need to know?. Journal of Genetic Counseling, 2019, 28, 1154-1165.	1.6	8
160	Enrolment in paediatric oncology earlyâ€phase clinical trials: The healthâ€care professionals' perspective. Journal of Paediatrics and Child Health, 2019, 55, 561-566.	0.8	8
161	Perceptions of future health and cancer risk in adult survivors of childhood cancer: Implications for engagement in followâ€up care. Cancer, 2019, 125, 1008-1009.	4.1	8
162	â€~Why us?' Causal attributions of childhood cancer survivors, survivors' parents and community comparisons - a mixed methods analysis. Acta Oncológica, 2019, 58, 209-217.	1.8	8

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163	When to break the news and whose responsibility is it? A cross-sectional qualitative study of health professionals' views regarding disclosure of BRCA genetic cancer risk. BMJ Open, 2020, 10, e033127.	1.9	8
164	School and educational support programmes for paediatric oncology patients and survivors: A systematic review of evidence and recommendations for future research and practice. Psycho-Oncology, 2021, 30, 431-443.	2.3	8
165	Providing Psychological Support to Parents of Childhood Cancer Survivors: â€~Cascade' Intervention Trial Results and Lessons for the Future. Cancers, 2021, 13, 5597.	3.7	8
166	A systematic review of social support for siblings of children with neurodevelopmental disorders. Research in Developmental Disabilities, 2022, 126, 104234.	2.2	8
167	Importance of updating family cancer history in childhood cancer survivors. Familial Cancer, 2017, 16, 605-610.	1.9	7
168	Genetics-related service and information needs of childhood cancer survivors and parents: a mixed-methods study. European Journal of Human Genetics, 2020, 28, 6-16.	2.8	7
169	Meal planning values impacted by the cancer experience in families with school-aged survivors—a qualitative exploration and recommendations for intervention development. Supportive Care in Cancer, 2020, 28, 1305-1313.	2.2	7
170	Avatar acceptability: views from the Australian Cystic Fibrosis community on the use of personalised organoid technology to guide treatment decisions. ERJ Open Research, 2021, 7, 00448-2020.	2.6	7
171	A costâ€effective approach to increasing participation in patientâ€reported outcomes research in cancer: A randomized trial of video invitations. International Journal of Cancer, 2021, 148, 971-980.	5.1	7
172	Social Anxiety Symptoms in Survivors of Childhood and Adolescent Cancer. Journal of Adolescent and Young Adult Oncology, 2022, 11, 129-137.	1.3	7
173	Design and Rationale for a Parent-Led Intervention to Increase Fruit and Vegetable Intake in Young Childhood Cancer Survivors (Reboot): Protocol for a Pilot Study. JMIR Research Protocols, 2018, 7, e129.	1.0	7
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