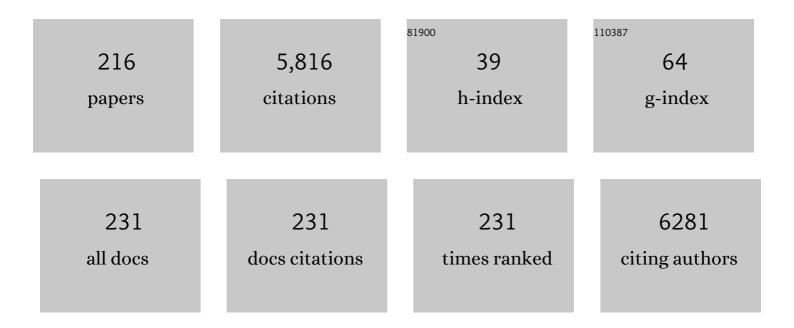
List of Publications by Year in descending order

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FAITH CIRSON

#	Article	IF	CITATIONS
1	Acceptability of a novel device to improve child patient experience during venepuncture for blood sampling: Intervention with â€~MyShield'. Journal of Child Health Care, 2024, 28, 53-68.	1.4	0
2	Culture, cognisance, capacity and capability: The interrelationship of individual and organisational factors in developing a research hospital. Journal of Clinical Nursing, 2022, 31, 362-377.	3.0	4
3	Increased admissions to paediatric wards with a primary mental health diagnosis: results of a survey of a network of eating disorder paediatricians in England. Archives of Disease in Childhood, 2022, 107, 309-310.	1.9	14
4	Nutritional support practices and opinions toward gastrostomy use in pediatric bone marrow transplant centers: A national survey. Nutrition, 2022, 95, 111556.	2.4	7
5	The experiences and perceptions of users of an electronic patient record system in a pediatric hospital setting: a systematic review. International Journal of Medical Informatics, 2022, 160, 104691.	3.3	13
6	Associations between diagnostic time intervals and health-related quality of life, clinical anxiety and depression in adolescents and young adults with cancer: cross-sectional analysis of the BRIGHTLIGHT cohort. British Journal of Cancer, 2022, 126, 1725-1734.	6.4	10
7	Developing, Nurturing, and Sustaining an Adolescent and Young Adult-Centered Culture of Care. Qualitative Health Research, 2022, 32, 956-969.	2.1	2
8	Nutritional assessment and dietetic resource for children and young people with cancer in the United Kingdom. Pediatric Blood and Cancer, 2022, 69, e29743.	1.5	7
9	QOL-34. The relationship between Psychological Flexibility, Quality-of-Life and psychological health in young people who have experienced a brain tumour. Neuro-Oncology, 2022, 24, i141-i141.	1.2	2
10	QOL-26. Exploring the experience of young people receiving remotely delivered Acceptance and Commitment Therapy following treatment for a brain tumour. Neuro-Oncology, 2022, 24, i139-i139.	1.2	0
11	Equal access to hospital care for children with learning disabilities and their families: a mixed-methods study. , 2022, 10, 1-168.		3
12	Care coordination, consistency and continuity: the case of the key worker role in children's cancer care. International Journal of Qualitative Studies on Health and Well-being, 2022, 17, .	1.6	1
13	Supporting families of children with an undiagnosed genetic condition: Using coâ€design to ensure the right person is in the right post doing the right job. Child: Care, Health and Development, 2021, 47, 300-310.	1.7	4
14	The BRIGHTLIGHT National Survey of the Impact of Specialist Teenage and Young Adult Cancer Care on Caregivers' Information and Support Needs. Cancer Nursing, 2021, 44, 235-243.	1.5	9
15	Mapping the current psychology provision for children and young people with juvenile dermatomyositis. Rheumatology Advances in Practice, 2021, 5, rkab062.	0.7	4
16	"Holistic Competenceâ€! How Is it Developed, Shared, and Shaped by Health Care Professionals Caring for Adolescents and Young Adults with Cancer?. Journal of Adolescent and Young Adult Oncology, 2021, 10, 503-511.	1.3	5
17	†We don't know what tomorrow will bring': Parents' experiences of caring for a child with an undiagnosed genetic condition. Child: Care, Health and Development, 2021, 47, 588-596.	1.7	6
18	Processes of care and survival associated with treatment in specialist teenage and young adult cancer centres: results from the BRIGHTLIGHT cohort study. BMJ Open, 2021, 11, e044854.	1.9	5

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19	Parent values and preferences underpinning treatment decision making in poor prognosis childhood cancer: a scoping review protocol. BMJ Open, 2021, 11, e046284.	1.9	2
20	Access to and experience of education for children and adolescents with cancer: a scoping review protocol. Systematic Reviews, 2021, 10, 167.	5.3	5
21	Support interventions provided during palliative care to families with dependent children when a parent has terminal illness: a scoping review protocol. JBI Evidence Synthesis, 2021, 19, 3163-3173.	1.3	0
22	Acceptance and commitment therapy for young brain tumour survivors: study protocol for an acceptability and feasibility trial. BMJ Open, 2021, 11, e051091.	1.9	2
23	Systematic review of gastrostomy complications and outcomes in pediatric cancer and bone marrow transplant. Nutrition in Clinical Practice, 2021, 36, 1185-1197.	2.4	4
24	Reporting the whole story: Analysis of the †outâ€ofâ€scope' questions from the James Lind Alliance Teenage and Young Adult Cancer Priority Setting Partnership Survey. Health Expectations, 2021, 24, 1593-1606.	2.6	8
25	Specialist cancer services for teenagers and young adults in England: BRIGHTLIGHT research programme. Programme Grants for Applied Research, 2021, 9, 1-82.	1.0	3
26	97â€The experiences and perceptions of users of an electronic patient record system in a paediatric hospital setting: a systematic review. , 2021, , .		0
27	96â€The going digital study: the views and perceptions of children and young people, parents, and staff before and after electronic patient record implementation – trust-wide survey findings. , 2021, , .		0
28	14â $\in$ Systematic review of gastrostomy complications and outcomes in paediatric cancer and bone marrow transplant. , 2021, , .		0
29	GPs and paediatric oncology palliative care: a Q methodological study. BMJ Supportive and Palliative Care, 2020, 10, e11-e11.	1.6	5
30	Relationships among resilience, self-esteem, and depressive symptoms in Chinese adolescents. Journal of Health Psychology, 2020, 25, 2396-2405.	2.3	21
31	Reducing unintentional injuries in under fives: Development and testing of a mobile phone app. Child: Care, Health and Development, 2020, 46, 203-212.	1.7	7
32	Renegotiation of identity in young adults with cancer: A longitudinal narrative study. International Journal of Nursing Studies, 2020, 102, 103465.	5.6	7
33	You give me a name that I can't say, but I have to explain what it is every day: the power of poetry to share stories from young people with a rare disease. Journal of Poetry Therapy, 2020, 33, 20-29.	0.7	3
34	Staying Connected: I Wonder, What Does Nurse Presence Look Like During This COVID-19 Pandemic?. Cancer Nursing, 2020, 43, 425-426.	1.5	2
35	BRIGHTLIGHT researchers as â€ <sup>~</sup> dramaturgs': creating There is a Light from complex research data. Research Involvement and Engagement, 2020, 6, 48.	2.9	3
36	Association of Self-reported Presenting Symptoms With Timeliness of Help-Seeking Among Adolescents and Young Adults With Cancer in the BRIGHTLIGHT Study. JAMA Network Open, 2020, 3, e2015437.	5.9	10

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37	Untellable tales and uncertain futures: the unfolding narratives of young adults with cancer. International Journal of Social Research Methodology: Theory and Practice, 2020, 23, 377-390.	4.4	6
38	Psychometric evaluation of the traditional Chinese version of the resilience Scale-14 and assessment of resilience in Hong Kong adolescents. Health and Quality of Life Outcomes, 2020, 18, 33.	2.4	9
39	Hospitalized children with intellectual disability: Parents as partners in their care. Journal of Applied Research in Intellectual Disabilities, 2020, 33, 917-926.	2.0	12
40	Longitudinal cohort study of the impact of specialist cancer services for teenagers and young adults on quality of life: outcomes from the BRIGHTLIGHT study. BMJ Open, 2020, 10, e038471.	1.9	17
41	Best Practice for Healthcare Transition: Development and Use of theÂBenchmarks for Transition. , 2020, , 127-146.		1
42	Bereavement support after the death of a child with cancer: implications for practice. Bereavement Care, 2020, 39, 12-20.	0.3	0
43	Gathering expert opinion to inform benchmarks to support transitional care. Journal of Child Health Care, 2019, 23, 131-146.	1.4	6
44	Health professional perceptions of communicating with adolescents and young adults about bone cancer clinical trial participation. Supportive Care in Cancer, 2019, 27, 467-475.	2.2	13
45	Children with Cancer: Communication, an Essential Component of Care. Principles of Specialty Nursing, 2019, , 329-341.	0.2	1
46	To flag or not to flag: Identification of children and young people with learning disabilities in English hospitals. Journal of Applied Research in Intellectual Disabilities, 2019, 32, 1176-1183.	2.0	22
47	Understanding care when cure is not likely for young adults who face cancer: a realist analysis of data from patients, families and healthcare professionals. BMJ Open, 2019, 9, e024397.	1.9	10
48	Being on the juvenile dermatomyositis rollercoaster: a qualitative study. Pediatric Rheumatology, 2019, 17, 30.	2.1	19
49	The culture of young people's cancer care: A narrative review and synthesis of the UK literature. European Journal of Cancer Care, 2019, 28, e13099.	1.5	6
50	Description of the BRIGHTLIGHT cohort: the evaluation of teenage and young adult cancer services in England. BMJ Open, 2019, 9, e027797.	1.9	17
51	Reducing pain in children with cancer: Methodology for the development of a clinical practice guideline. Pediatric Blood and Cancer, 2019, 66, e27698.	1.5	14
52	Measurement properties of instruments to assess pain in children and adolescents with cancer: a systematic review protocol. Systematic Reviews, 2019, 8, 33.	5.3	1
53	Research priorities for young people with cancer: a UK priority setting partnership with the James Lind Alliance. BMJ Open, 2019, 9, e028119.	1.9	42
54	How Far Do Research Priority Setting Exercises Influence What Research Is Undertaken: A Little, a Lot, or Not at All?. Cancer Nursing, 2019, 42, 89-90.	1.5	7

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55	Studying Children's Experiences in Interactions With Clinicians: Identifying Methods Fit for Purpose. Qualitative Health Research, 2019, 29, 393-403.	2.1	17
56	Consult, Negotiate, and Involve: Evaluation of an Advanced Communication Skills Program for Health Care Professionals. Journal of Pediatric Oncology Nursing, 2018, 35, 296-307.	1.5	10
57	Diagnostic timeliness in adolescents and young adults with cancer: a cross-sectional analysis of the BRIGHTLIGHT cohort. The Lancet Child and Adolescent Health, 2018, 2, 180-190.	5.6	42
58	Physical activity reduces fatigue in patients with cancer and hematopoietic stem cell transplant recipients: A systematic review and meta-analysis of randomized trials. Critical Reviews in Oncology/Hematology, 2018, 122, 52-59.	4.4	111
59	" <scp>LEARN</scp> â€ing what is important to children and young people with intellectual disabilities when they are in hospital. Journal of Applied Research in Intellectual Disabilities, 2018, 31, 792-803.	2.0	11
60	Management of fatigue in children and adolescents with cancer and in paediatric recipients of haemopoietic stem-cell transplants: a clinical practice guideline. The Lancet Child and Adolescent Health, 2018, 2, 371-378.	5.6	44
61	Protecting an adult identity: A grounded theory of supportive care for young adults recently diagnosed with cancer. International Journal of Nursing Studies, 2018, 81, 40-48.	5.6	27
62	The perceptions of teenagers, young adults and professionals in the participation of bone cancer clinical trials. European Journal of Cancer Care, 2018, 27, e12476.	1.5	24
63	Preventing work-related stress among staff working in children's cancer Principal Treatment Centres in the UK: a brief survey of staff support systems and practices. European Journal of Cancer Care, 2018, 27, e12535.	1.5	9
64	R11 Understanding juvenile dermatomyositis from a young person's perspective. Rheumatology, 2018, 57	' 1.9	0
65	Involving young people in BRIGHTLIGHT from study inception to secondary data analysis: insights from 10 years of user involvement. Research Involvement and Engagement, 2018, 4, 50.	2.9	17
66	How should health service researchers respect children's personal data under GDPR?. The Lancet Child and Adolescent Health, 2018, 2, 696-697.	5.6	2
67	Conceptualizing age-appropriate care for teenagers and young adults with cancer: a qualitative mixed-methods study. Adolescent Health, Medicine and Therapeutics, 2018, Volume 9, 149-166.	0.9	25
68	Pharmacologic Interventions for Fatigue in Cancer and Transplantation: A Meta-Analysis. Current Oncology, 2018, 25, 152-167.	2.2	50
69	Mapping staff perspectives towards the delivery of hospital care for children and young people with and without learning disabilities in England: a mixed methods national study. BMC Health Services Research, 2018, 18, 203.	2.2	22
70	Insights from parents of a child with leukaemia and healthcare professionals about sharing illness and treatment information: A qualitative research study. International Journal of Nursing Studies, 2018, 83, 91-102.	5.6	11
71	Parents' Experiences of Caring for Their Child at the Time of Discharge After Cardiac Surgery and During the Postdischarge Period: Qualitative Study Using an Online Forum. Journal of Medical Internet Research, 2018, 20, e155.	4.3	34
72	Transitional Care: Best Practices for Adolescent Cancer Care. , 2018, , 133-151.		1

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73	Experiences and Preferences for End-of-Life Care for Young Adults with Cancer and Their Informal Carers: A Narrative Synthesis. Journal of Adolescent and Young Adult Oncology, 2017, 6, 200-212.	1.3	28
74	Multi-Professional Perspectives on Adolescent and Young Adult Oncology Across Europe: An e-Delphi Survey. Journal of Adolescent and Young Adult Oncology, 2017, 6, 178-185.	1.3	6
75	Cancer-Related Fatigue in Adolescents and Young Adults After Cancer Treatment: Persistent and Poorly Managed. Journal of Adolescent and Young Adult Oncology, 2017, 6, 489-493.	1.3	38
76	Mind and body practices for fatigue reduction in patients with cancer and hematopoietic stem cell transplant recipients: A systematic review and meta-analysis. Critical Reviews in Oncology/Hematology, 2017, 120, 210-216.	4.4	39
77	Qualitative study to understand the barriers to recruiting young people with cancer to BRIGHTLIGHT: a national cohort study in England. BMJ Open, 2017, 7, e018291.	1.9	13
78	Identifying research priorities with nurses at a tertiary children's hospital in the United Kingdom. Child: Care, Health and Development, 2017, 43, 211-221.	1.7	7
79	105â€Blood quest – preparing children for blood tests: using arts-based techniques in creative research and app development. , 2017, , .		Ο
80	Improving the identification of cancer in young people: A scoping review. Expert Review of Quality of Life in Cancer Care, 2017, 2, 87-101.	0.6	11
81	Addressing the Ethical Challenges for Young Adults, from a Rights-Based Perspective. Pediatric Oncology, 2017, , 765-778.	0.5	1
82	"We Sometimes Hold on to Ours―– Professionals' Views on Factors that both Delay and Facilitate Transition to Adult Care. Frontiers in Pediatrics, 2016, 4, 125.	1.9	20
83	How young people describe the impact of living with and beyond a cancer diagnosis: feasibility of using social media as a research method. Psycho-Oncology, 2016, 25, 1317-1323.	2.3	24
84	Assent for children's participation in research: why it matters and making it meaningful. Child: Care, Health and Development, 2016, 42, 588-597.	1.7	30
85	Mapping Adolescent Cancer Services. Cancer Nursing, 2016, 39, 358-366.	1.5	23
86	Transition of care for adolescents from paediatric services to adult health services. The Cochrane Library, 2016, 2016, CD009794.	2.8	239
87	Interventions for promoting participation in shared decision-making for children with cancer. The Cochrane Library, 2016, 2016, CD008970.	2.8	38
88	Challenges encountered in end of life care research with young adults with cancer. European Journal of Surgical Oncology, 2016, 42, S251-S252.	1.0	0
89	Specialist nurse key worker in children's cancer care: Professionals' perspectives on the core characteristics of the role. European Journal of Oncology Nursing, 2016, 24, 70-78.	2.1	9
90	Psychosocial Interventions for Adolescents and Young Adults Diagnosed with Cancer During Adolescence: A Critical Review. Journal of Adolescent and Young Adult Oncology, 2016, 5, 310-321.	1.3	39

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91	Information-sharing between healthcare professionals, parents and children with cancer: more than a matter of information exchange. European Journal of Cancer Care, 2016, 25, 141-156.	1.5	80
92	G619â€The benefits and challenges of using creative methods to explore happiness with children with rare diseases. Archives of Disease in Childhood, 2016, 101, A367.1-A367.	1.9	0
93	Pay More Attention: a national mixed methods study to identify the barriers and facilitators to ensuring equal access to high-quality hospital care and services for children and young people with and without learning disabilities and their families. BMJ Open, 2016, 6, e012333.	1.9	14
94	Modified international e-Delphi survey to define healthcare professional competencies for working with teenagers and young adults with cancer. BMJ Open, 2016, 6, e011361.	1.9	57
95	Antiemetic medication for prevention and treatment of chemotherapy-induced nausea and vomiting in childhood. The Cochrane Library, 2016, 2016, CD007786.	2.8	46
96	From informed consent to dissemination: Using participatory visual methods with young people with long-term conditions at different stages of research. Global Public Health, 2016, 11, 636-650.	2.0	29
97	A prospective observational study of machine translation software to overcome the challenge of including ethnic diversity in healthcare research. Nursing Open, 2015, 2, 14-23.	2.4	22
98	Building the Evidence for Nursing Practice: Learning from a Structured Review of SIOP Abstracts, 2003-2012. Pediatric Blood and Cancer, 2015, 62, 2172-2176.	1.5	2
99	Pediatric Oncology Palliative Care: Experiences of General Practitioners and Bereaved Parents. Journal of Palliative Care & Medicine, 2015, 05, 214.	0.1	4
100	Be my guest! Challenges and practical solutions of undertaking interviews with children in the home setting. Journal of Child Health Care, 2015, 19, 432-443.	1.4	23
101	A Critical Review of the Use of Technology to Provide Psychosocial Support for Children and Young People with Long-Term Conditions. Journal of Pediatric Nursing, 2015, 30, 87-101.	1.5	20
102	Individualizing Hospital Care for Children and Young People With Learning Disabilities: It's the Little Things That Make the Difference. Journal of Pediatric Nursing, 2015, 30, 78-86.	1.5	25
103	Novel participatory methods of involving patients in research: naming and branding a longitudinal cohort study, BRIGHTLIGHT. BMC Medical Research Methodology, 2015, 15, 20.	3.1	32
104	Development and validation of the BRIGHTLIGHT Survey, a patient-reported experience measure for young people with cancer. Health and Quality of Life Outcomes, 2015, 13, 107.	2.4	50
105	Cancer-Related Pain and Pain Management. Journal of Pediatric Oncology Nursing, 2015, 32, 369-384.	1.5	60
106	Transition From Child to Adult Care – †It's Not a One-Off Event': Development of Benchmarks to Improve the Experience. Journal of Pediatric Nursing, 2015, 30, 638-647.	1.5	46
107	Selfâ€management and skills acquisition in boys with haemophilia. Health Expectations, 2015, 18, 1105-1113.	2.6	32
108	A relational understanding of sibling experiences of children with rare life-limiting conditions. Journal of Child Health Care. 2014. 18, 230-240.	1.4	27

FAITH GIBSON

#	Article	IF	CITATIONS
109	Guidance on clinical research involving infants, children and young people: an update for researchers and research ethics committees. Archives of Disease in Childhood, 2014, 99, 887-891.	1.9	58
110	What Is It Really Like to Be a Young Person, in Our Hospital, at This Moment?. Cancer Nursing, 2014, 37, 86-87.	1.5	2
111	Children's participation in shared decision-making: Children, adolescents, parents and healthcare professionals' perspectives and experiences. European Journal of Oncology Nursing, 2014, 18, 273-280.	2.1	217
112	A systematic review of models of care for the followâ€up of childhood cancer survivors. Pediatric Blood and Cancer, 2013, 60, 351-356.	1.5	38
113	An action research study to explore the nature of the nurse consultant role in the care of children and young people. Journal of Clinical Nursing, 2013, 22, 201-210.	3.0	11
114	Developing a conceptual model of teenage and young adult experiences of cancer through meta-synthesis. International Journal of Nursing Studies, 2013, 50, 832-846.	5.6	65
115	International collaboration: Working with and through professional groups to inform the science that underpins the care of children and young people. European Journal of Oncology Nursing, 2013, 17, 695-696.	2.1	0
116	Systematic review of basic oral care for the management of oral mucositis in cancer patients. Supportive Care in Cancer, 2013, 21, 3165-3177.	2.2	194
117	Collaboration between doctors and nurses in children's cancer care: Insights fromÂa European project. European Journal of Oncology Nursing, 2013, 17, 745-749.	2.1	8
118	Developing a national †low risk' febrile neutropenia framework for use in children and young people's cancer care. Supportive Care in Cancer, 2013, 21, 1241-1251.	2.2	16
119	Exploring the work of nurses who administer chemotherapy to children and young people. European Journal of Oncology Nursing, 2013, 17, 59-69.	2.1	19
120	Neutropenic sepsis: prevention and management of neutropenic sepsis in cancer patients (NICE Clinical) Tj ETQo	0 <u>8 9</u> rgB 0	T /Qyerlock 1
121	Why can't we improve the timeliness of cancer diagnosis in children, teenagers, and young adults?. BMJ, The, 2013, 347, f6493-f6493.	6.0	24
122	A reaudit of current febrile neutropenia practice in UK paediatric oncology centres prior to implementation of NICE guidance. Archives of Disease in Childhood, 2013, 98, 315-316.	1.9	15
123	Enhanced quality and productivity of long-term aftercare of cancer in young people. Archives of Disease in Childhood, 2013, 98, 818-824.	1.9	19
124	The Art of Age-Appropriate Care. Cancer Nursing, 2013, 36, E27-E38.	1.5	88
125	Pediatric Oncology Nursing Research Goes Global. Cancer Nursing, 2013, 36, 339.	1.5	4
126	Young people describe their prediagnosis cancer experience. Psycho-Oncology, 2013, 22, 2585-2592.	2.3	31

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127	G178 What Do GPs and Bereaved Parents Think About Paediatric Oncologists and Palliative Care?. Archives of Disease in Childhood, 2013, 98, A81-A81.	1.9	0
128	â€~I can always rely on them': the importance of social support for boys with haemophilia. The Journal of Haemophilia Practice, 2013, 1, 10-16.	0.4	6
129	Multimethodology research with boys with severe haemophilia. Nurse Researcher, 2013, 20, 40-44.	0.5	6
130	What's It Like When You Find Eating Difficult. Cancer Nursing, 2012, 35, 265-277.	1.5	25
131	Practical approaches to seeking assent from children. Nurse Researcher, 2012, 19, 23-27.	0.5	22
132	Guideline for the Management of Fever and Neutropenia in Children With Cancer and/or Undergoing Hematopoietic Stem-Cell Transplantation. Journal of Clinical Oncology, 2012, 30, 4427-4438.	1.6	311
133	The benefits of prophylaxis: views of adolescents with severe haemophilia. Haemophilia, 2012, 18, e286-9.	2.1	32
134	Challenging symptoms in children with rare lifeâ€limiting conditions: findings from a prospective diary and interview study with families. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, 985-992.	1.5	51
135	A scoping exercise of favourable characteristics of professionals working in teenage and young adult cancer care: †thinking outside of the box'. European Journal of Cancer Care, 2012, 21, 330-339.	1.5	29
136	Initial Evaluation of an Electronic Symptom Diary for Adolescents with Cancer. JMIR Research Protocols, 2012, 1, e23.	1.0	73
137	Children's understanding of cancer and views on health-related behaviour: a â€~draw and write' study. Child: Care, Health and Development, 2011, 37, 289-299.	1.7	37
138	â€Just an unfortunate coincidence': children's understanding of haemophilia genetics and inheritance. Haemophilia, 2011, 17, 470-475.	2.1	7
139	Parental assessment of adolescent quality of life: can it replace self-assessment?. Quality of Life Research, 2011, 20, 1715-1720.	3.1	15
140	Audit to monitor the uptake of national mouth care guidelines for children and young people being treated for cancer. Supportive Care in Cancer, 2011, 19, 1335-1341.	2.2	3
141	Working in collaboration with young people and health professionals. A staged approach to the implementation of a randomised controlled trial. Journal of Research in Nursing, 2011, 16, 561-576.	0.9	19
142	"Your Place or Mine?―Priorities for a Specialist Teenage and Young Adult (TYA) Cancer Unit: Disparity Between TYA and Professional Perceptions. Journal of Adolescent and Young Adult Oncology, 2011, 1, 145-151.	1.3	33
143	Challenging symptom profiles of life-limiting conditions in children: a survey of care professionals and families. Palliative Medicine, 2011, 25, 357-364.	3.1	38
144	Utilization of the Medical Research Council Evaluation Framework in the Development of Technology for Symptom Management. Cancer Nursing, 2010, 33, 343-352.	1.5	33

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145	Contemporary issues relating to assessment of pre-registration nursing students in practice. Nurse Education in Practice, 2010, 10, 158-163.	2.6	39
146	Children and young people's experiences of cancer care: A qualitative research study using participatory methods. International Journal of Nursing Studies, 2010, 47, 1397-1407.	5.6	149
147	Measuring quality of life in children and young people after transplantation: Methodological considerations. Pediatric Transplantation, 2010, 14, 445-458.	1.0	17
148	Antiemetic medication for prevention and treatment of chemotherapy induced nausea and vomiting in childhood. , 2010, , CD007786.		23
149	The Stories of Young People Living With a Liver Transplant. Qualitative Health Research, 2010, 20, 1076-1090.	2.1	56
150	A Systematic Review of Oral Assessment Instruments. Cancer Nursing, 2010, 33, E1-E19.	1.5	18
151	The development of evidence-based guidelines on mouth care for children, teenagers and young adults treated for cancer. European Journal of Cancer, 2010, 46, 1399-1412.	2.8	45
152	Refinement of the Children's International Mucositis Evaluation Scale (ChIMES): Child and parent perspectives on understandability, content validity and acceptability. European Journal of Oncology Nursing, 2010, 14, 29-41.	2.1	27
153	Assessment and management of nutritional challenges in children's cancer care: A survey of current practice in the United Kingdom. European Journal of Oncology Nursing, 2010, 14, 439-446.	2.1	48
154	Study of the Factors Affecting Health-Related Quality of Life in Adolescents After Liver Transplantation. American Journal of Transplantation, 2009, 9, 1179-1188.	4.7	73
155	Designing an oral mucositis assessment instrument for use in children: generating items using a nominal group technique. Supportive Care in Cancer, 2009, 17, 555-562.	2.2	22
156	Multiprofessional collaboration in children's cancer care: believed to be a good thing but how do we know when it works well?. European Journal of Cancer Care, 2009, 18, 327-329.	1.5	6
157	What is Important to Young Children Who Have Cancer While in Hospital?. Children and Society, 2009, 23, 85-98.	1.7	50
158	Involving health professionals in the development of an advanced symptom management system for young people: The ASyMS©-YG study. European Journal of Oncology Nursing, 2009, 13, 187-192.	2.1	32
159	What are the experiences of the child with a brain tumour and their parents?. European Journal of Oncology Nursing, 2009, 13, 255-261.	2.1	50
160	How do young people find out about their parent's cancer diagnosis: AÂphenomenologicalÂstudy. European Journal of Oncology Nursing, 2009, 13, 213-222.	2.1	46
161	Meeting the needs of young people with cancer: A lesson in change. European Journal of Oncology Nursing, 2009, 13, 147-148.	2.1	3
162	4170 The experiences of young people from first symptoms to the diagnosis of cancer: a narrative study. European Journal of Cancer, Supplement, 2009, 7, 234-235.	2.2	1

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163	Putting Both Patient Need and the Science of Nursing at the Heart of Our Research Agenda. Cancer Nursing, 2009, 32, 427-428.	1.5	0
164	Understandability, Content Validity, and Overall Acceptability of the Children's International Mucositis Evaluation Scale (ChIMES). Journal of Pediatric Hematology/Oncology, 2009, 31, 416-423.	0.6	20
165	A concept analysis of healthâ€related quality of life in young people with chronic illness. Journal of Clinical Nursing, 2008, 17, 1823-1833.	3.0	125
166	The experience of living with a chronic illness during adolescence: a critical review of the literature. Journal of Clinical Nursing, 2008, 17, 3083-3091.	3.0	147
167	Navigating the maze: parents' views and influence on a paediatric haematology and oncology day care service. Journal of Clinical Nursing, 2008, 17, 3159-3167.	3.0	7
168	Editorial: Getting it right for children and young people's health care services. Journal of Clinical Nursing, 2008, 17, 3081-3082.	3.0	4
169	The cultural context of communicating complex information to parents of children with cancer—not just a linguistic obstacle. European Journal of Oncology Nursing, 2008, 12, 2-3.	2.1	5
170	Challenges of mucositis assessment in children: Expert opinion. European Journal of Oncology Nursing, 2008, 12, 469-475.	2.1	22
171	Methodological Issues When Using the Draw and Write Technique With Children Aged 6 to 12 Years. Qualitative Health Research, 2008, 18, 1001-1011.	2.1	106
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