

Bridget Young

List of Publications by Year in descending order

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Version: 2024-02-01

149
papers

9,486
citations

66343

42
h-index

40979

93
g-index

156
all docs

156
docs citations

156
times ranked

12946
citing authors

#	ARTICLE	IF	CITATIONS
1	Engaging children, young people, parents and health professionals in interviews: Using an interactive ranking exercise within the co-design of multimedia websites. <i>Journal of Child Health Care</i> , 2024, 28, 181-195.	1.4	1
2	Assessing metacognitive beliefs in test anxiety: Psychometric properties of the metacognitions questionnaire, 30 (MCQ-30) among university students. <i>Current Psychology</i> , 2022, 41, 1425-1433.	2.8	7
3	Patientsâ€™ Perspectives on Transforming Clinical Trial Participation: Large Online Vignette-based Survey. <i>Journal of Medical Internet Research</i> , 2022, 24, e29691.	4.3	3
4	Testing times: the association of intolerance of uncertainty and metacognitive beliefs to test anxiety in college students. <i>BMC Psychology</i> , 2022, 10, 6.	2.1	7
5	In-depth qualitative interviews identified barriers and facilitators that influenced chief investigatorsâ€™ use of core outcome sets in randomised controlled trials. <i>Journal of Clinical Epidemiology</i> , 2022, 144, 111-120.	5.0	17
6	Assessing the effectiveness of social network interventions for adults with a diagnosis of mental health problems: a systematic review and narrative synthesis of impact. <i>Social Psychiatry and Psychiatric Epidemiology</i> , 2022, 57, 907-925.	3.1	9
7	Enhancing patient and public contribution in health outcome selection during clinical guideline development: an ethnographic study. <i>BMC Health Services Research</i> , 2022, 22, 361.	2.2	4
8	The development of theory-informed participant-centred interventions to maximise participant retention in randomised controlled trials. <i>Trials</i> , 2022, 23, 268.	1.6	3
9	Conservative treatment for uncomplicated appendicitis in children: the CONTRACT feasibility study, including feasibility RCT. <i>Health Technology Assessment</i> , 2021, 25, 1-192.	2.8	10
10	SAVER: sodium valproate for the epigenetic reprogramming of high-risk oral epithelial dysplasiaâ€™ a phase II randomised control trial study protocol. <i>Trials</i> , 2021, 22, 428.	1.6	4
11	Strategies to enhance recruitment and consent to intensive care studies: a qualitative study with researchers and patientâ€™public involvement contributors. <i>BMJ Open</i> , 2021, 11, e048193.	1.9	7
12	Why trials lose participants: A multitrial investigation of participantsâ€™ perspectives using the theoretical domains framework. <i>Journal of Clinical Epidemiology</i> , 2021, 137, 1-13.	5.0	16
13	Longitudinal Outcomes of Gender Identity in Children (LOGIC): protocol for a prospective longitudinal cohort study of children referred to the UK gender identity development service. <i>BMJ Open</i> , 2021, 11, e045628.	1.9	3
14	â€œIt shows we felt safe because a few of us had tearsâ€ Exploring how perceptions of the group space can influence experiences of parenting programmes. <i>Emotion, Space and Society</i> , 2021, 41, 100837.	1.5	0
15	CONservative TRreatment of Appendicitis in Children: a randomised controlled feasibility Trial (CONTRACT). <i>Archives of Disease in Childhood</i> , 2021, 106, 764-773.	1.9	15
16	â€œSometimes I think my frustration is the real issueâ€ A qualitative study of parentsâ€™ experiences of transformation after a parenting programme. <i>PLoS ONE</i> , 2021, 16, e0258528.	2.5	2
17	Inclusion of participants from low-income and middle-income countries in core outcome sets development: a systematic review. <i>BMJ Open</i> , 2021, 11, e049981.	1.9	13
18	Recruitersâ€™ perspectives and experiences of trial recruitment processes: a qualitative evidence synthesis protocol. <i>BMJ Open</i> , 2021, 11, e045233.	1.9	1

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19	Longitudinal outcomes of gender identity in children (LOGIC): a study protocol for a prospective longitudinal qualitative study of the experiences and well-being of families referred to the UK Gender Identity Development Service. <i>BMJ Open</i> , 2021, 11, e047875.	1.9	1
20	Patient participation in Delphi surveys to develop core outcome sets: systematic review. <i>BMJ Open</i> , 2021, 11, e051066.	1.9	0
21	Longitudinal Outcomes of Gender Identity in Children (LOGIC): study protocol for a retrospective analysis of the characteristics and outcomes of children referred to specialist gender services in the UK and the Netherlands. <i>BMJ Open</i> , 2021, 11, e054895.	1.9	2
22	Patient participation in Delphi surveys to develop core outcome sets: systematic review. <i>BMJ Open</i> , 2021, 11, e051066.	1.9	18
23	Heterogeneity in outcome assessment for inflammatory bowel disease in routine clinical practice: a mixed-methods study in a sample of English hospitals. <i>BMJ Open</i> , 2021, 11, e056413.	1.9	3
24	Informing the personalisation of interventions for parents of children with conduct problems: a qualitative study. <i>BMC Psychiatry</i> , 2020, 20, 513.	2.6	7
25	Controversy and Debate Series on Core Outcome Sets. Paper 4: Debate on Paper 1 from the perspective of COMET [Core Outcome Measures in Effectiveness Trials]. <i>Journal of Clinical Epidemiology</i> , 2020, 125, 222-224.	5.0	7
26	Uncertainty and test anxiety: Psychometric properties of the Intolerance of Uncertainty Scale â€“ 12 (IUS-12) among university students. <i>International Journal of Educational Research</i> , 2020, 104, 101672.	2.2	18
27	Ensuring young voices are heard in core outcome set development: international workshops with 70 children and young people. <i>Research Involvement and Engagement</i> , 2020, 6, 19.	2.9	12
28	Challenges Conveying Clinical Equipoise and Exploring Patient Treatment Preferences in an Oncology Trial Comparing Active Monitoring with Radiotherapy (ROAM/EORTC 1308). <i>Oncologist</i> , 2020, 25, e691-e700.	3.7	17
29	The research burden of randomized controlled trial participation: a systematic thematic synthesis of qualitative evidence. <i>BMC Medicine</i> , 2020, 18, 6.	5.5	67
30	Enhancing communication, informed consent and recruitment in a paediatric urgent care surgical trial: a qualitative study. <i>BMC Pediatrics</i> , 2020, 20, 140.	1.7	8
31	Ketogenic diets as an adjuvant therapy for glioblastoma (KEATING): a randomized, mixed methods, feasibility study. <i>Journal of Neuro-Oncology</i> , 2020, 147, 213-227.	2.9	42
32	Different corticosteroid induction regimens in children and young people with juvenile idiopathic arthritis: the SIRJIA mixed-methods feasibility study. <i>Health Technology Assessment</i> , 2020, 24, 1-152.	2.8	3
33	Trials need participants but not their feedback? A scoping review of published papers on the measurement of participant experience of taking part in clinical trials. <i>Trials</i> , 2019, 20, 381.	1.6	25
34	Identifying the primary outcome measure and protocol components for a prospective feasibility study of corticosteroid regimens for CYP with JIA using consensus methods with young people, families and professionals. <i>Rheumatology</i> , 2019, 58, .	1.9	0
35	How could we know if communication skills training needed no more evaluation? The case for rigour in research design. <i>Patient Education and Counseling</i> , 2019, 102, 1401-1403.	2.2	2
36	Incorporating patientsâ€™ perspectives into the initial stages of core outcome set development: a rapid review of qualitative studies of type 2 diabetes. <i>BMJ Open Diabetes Research and Care</i> , 2019, 7, e000615.	2.8	22

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37	A scoping review provided a framework for new ways of doing research through mobilizing collective intelligence. <i>Journal of Clinical Epidemiology</i> , 2019, 110, 1-11.	5.0	13
38	Supporting children and young people when making decisions about joining clinical trials: qualitative study to inform multimedia website development. <i>BMJ Open</i> , 2019, 9, e023984.	1.9	19
39	The efficacy of interventions for test-anxious university students: A meta-analysis of randomized controlled trials. <i>Journal of Anxiety Disorders</i> , 2019, 63, 36-50.	3.2	34
40	AB0936â€¦A NATIONAL SURVEY OF CLINICAL PRACTICE OF CORTICOSTEROID USE IN NEWLY DIAGNOSED OR FLARING CASES OF JUVENILE IDIOPATHIC ARTHRITIS ACROSS THE UK. , 2019, , .		0
41	AB0937â€¦IDENTIFYING THE PRIMARY OUTCOME MEASURE AND PROTOCOL COMPONENTS FOR A PROSPECTIVE FEASIBILITY STUDY OF CORTICOSTEROID REGIMENS FOR CHILDREN AND YOUNG PEOPLE WITH JUVENILE IDIOPATHIC ARTHRITIS USING CONSENSUS METHODS WITH YOUNG PEOPLE, FAMILIES AND PROFESSIONALS. , 2019, , .		0
42	Integrating quantitative and qualitative data and findings when undertaking randomised controlled trials. <i>BMJ Open</i> , 2019, 9, e032081.	1.9	42
43	Participating in core outcome set development via Delphi surveys: qualitative interviews provide pointers to inform guidance. <i>BMJ Open</i> , 2019, 9, e032338.	1.9	20
44	Overcoming Barriers to Mobilizing Collective Intelligence in Research: Qualitative Study of Researchers With Experience of Collective Intelligence. <i>Journal of Medical Internet Research</i> , 2019, 21, e13792.	4.3	4
45	The association of metacognitive beliefs with emotional distress and trauma symptoms in adolescent and young adult survivors of cancer. <i>Journal of Psychosocial Oncology</i> , 2018, 36, 545-556.	1.2	7
46	Training health professionals to recruit into challenging randomized controlled trials improved confidence: the development of the QuinteT randomized controlled trial recruitment training intervention. <i>Journal of Clinical Epidemiology</i> , 2018, 95, 34-44.	5.0	28
47	Systematic Techniques to Enhance rEtention in Randomised controlled trials: the STEER study protocol. <i>Trials</i> , 2018, 19, 197.	1.6	14
48	CONTRACT Study - CONservative TRreatment of Appendicitis in Children (feasibility): study protocol for a randomised controlled Trial. <i>Trials</i> , 2018, 19, 153.	1.6	27
49	Perspectives of patients with haematological cancer on how clinicians meet their information needs: â€œManagingâ€ information versus â€œgivingâ€ it. <i>Psycho-Oncology</i> , 2018, 27, 1719-1726.	2.3	8
50	MNGI-17. THE ROAM / EORTC 1308 INFORMATION STUDY RESULTS: HOW QUALITATIVE RESEARCH METHODS CAN OPTIMISE PATIENT RECRUITMENT FOR MENINGIOMA TRIALS. <i>Neuro-Oncology</i> , 2018, 20, vi152-vi152.	1.2	0
51	Protective parents and permissive children: what qualitative interviews with parents and children can tell us about the feasibility of juvenile idiopathic arthritis trials. <i>Pediatric Rheumatology</i> , 2018, 16, 76.	2.1	6
52	Qualitative methods can test and challenge what we think we know about clinical communication â€œ if they are not too constrained by methodological â€˜brandsâ€™. <i>Patient Education and Counseling</i> , 2018, 101, 1515-1517.	2.2	19
53	Survey indicated that core outcome set development is increasingly including patients, being conducted internationally and using Delphi surveys. <i>Trials</i> , 2018, 19, 113.	1.6	45
54	Letter to the Editor. Phase III randomized controlled trials are essential to properly evaluate the role of radiotherapy in WHO grade II meningioma. <i>Journal of Neurosurgery</i> , 2018, 129, 1104-1105.	1.6	2

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55	Children's views on research without prior consent in emergency situations: a UK qualitative study. <i>BMJ Open</i> , 2018, 8, e022894.	1.9	15
56	Pervasive, hard-wired and male: Qualitative study of how UK adolescents view alcohol-related aggression. <i>PLoS ONE</i> , 2018, 13, e0191269.	2.5	1
57	63...Methods of mobilising collective intelligence through crowdsourcing in research:a scoping review. , 2018, , .		2
58	When trust is threatened: Qualitative study of parents' perspectives on problematic clinical relationships in child cancer care. <i>Psycho-Oncology</i> , 2017, 26, 1301-1306.	2.3	12
59	Improving core outcome set development: qualitative interviews with developers provided pointers to inform guidance. <i>Journal of Clinical Epidemiology</i> , 2017, 86, 140-152.	5.0	41
60	Priorities for methodological research on patient and public involvement in clinical trials: A modified Delphi process. <i>Health Expectations</i> , 2017, 20, 1401-1410.	2.6	36
61	A new paradigm for clinical communication: critical review of literature in cancer care. <i>Medical Education</i> , 2017, 51, 258-268.	2.1	48
62	GPs' experiences of children with anxiety disorders in primary care: a qualitative study. <i>British Journal of General Practice</i> , 2017, 67, e888-e898.	1.4	17
63	The inseparability of emotional and instrumental care in cancer: Towards a more powerful science of clinical communication. <i>Patient Education and Counseling</i> , 2017, 100, 2138-2140.	2.2	8
64	Developing and evaluating multimedia information resources to improve engagement of children, adolescents, and their parents with trials (TRECA study): Study protocol for a series of linked randomised controlled trials. <i>Trials</i> , 2017, 18, 265.	1.6	23
65	The COMET Handbook: version 1.0. <i>Trials</i> , 2017, 18, 280.	1.6	1,128
66	Development of a core outcome set to determine the overall treatment success of acute uncomplicated appendicitis in children: a study protocol. <i>BMJ Paediatrics Open</i> , 2017, 1, e000151.	1.4	14
67	Is clinical communication the one area of clinical oncology that needs no new ideas?. <i>Medical Education</i> , 2017, 51, 1291-1293.	2.1	2
68	The efficacy of interventions for test anxiety in university students: A protocol for a systematic review and meta-analysis. <i>International Journal of Educational Research</i> , 2016, 77, 92-98.	2.2	9
69	Questioning assent: how are children's views included as families make decisions about clinical trials?. <i>Child: Care, Health and Development</i> , 2016, 42, 900-908.	1.7	17
70	Towards understanding problems in the parent-practitioner relationship when a child has cancer: meta-synthesis of the qualitative literature. <i>Psycho-Oncology</i> , 2016, 25, 1252-1260.	2.3	2
71	What can we learn from trial decliners about improving recruitment? Qualitative study. <i>Trials</i> , 2016, 17, 494.	1.6	19
72	The impact of advertising patient and public involvement on trial recruitment: embedded cluster randomised recruitment trial. <i>Trials</i> , 2016, 17, 586.	1.6	13

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73	Including patients in core outcome set development: issues to consider based on three workshops with around 100 international delegates. <i>Research Involvement and Engagement</i> , 2016, 2, 25.	2.9	84
74	A patient and public involvement (PPI) toolkit for meaningful and flexible involvement in clinical trials – a work in progress. <i>Research Involvement and Engagement</i> , 2016, 2, 15.	2.9	140
75	Fifteen-minute consultation: an evidence-based approach to research without prior consent (deferred) Tj ETQq1 1 0.784314 rgBT /Ov and Practice Edition, 2016, 101, 49-53.	0.5	38
76	Research with children and young people: not on them. <i>Archives of Disease in Childhood</i> , 2016, 101, 1086-1089.	1.9	13
77	The ROAM/EORTC-1308 trial: Radiation versus Observation following surgical resection of Atypical Meningioma: study protocol for a randomised controlled trial. <i>Trials</i> , 2015, 16, 519.	1.6	165
78	A little more conversation please? Qualitative study of researchers'™ and patients'™ interview accounts of training for patient and public involvement in clinical trials. <i>Trials</i> , 2015, 16, 190.	1.6	76
79	Alleviating Emotional Distress in Adolescent and Young Adult Cancer Survivors: An Open Trial of Metacognitive Therapy. <i>Journal of Adolescent and Young Adult Oncology</i> , 2015, 4, 64-69.	1.3	20
80	Using qualitative research to understand what outcomes matter to patients: direct and indirect approaches to outcome elicitation. <i>Trials</i> , 2015, 16, .	1.6	5
81	Factors affecting recruitment into depression trials: Systematic review, meta-synthesis and conceptual framework. <i>Journal of Affective Disorders</i> , 2015, 172, 274-290.	4.1	91
82	How do surgeons think they learn about communication? A qualitative study. <i>Medical Education</i> , 2015, 49, 408-416.	2.1	10
83	How parents and practitioners experience research without prior consent (deferred consent) for emergency research involving children with life threatening conditions: a mixed method study. <i>BMJ Open</i> , 2015, 5, e008522.	1.9	84
84	Clinical Trial Decisions in Difficult Circumstances: Parental Consent Under Time Pressure. <i>Pediatrics</i> , 2015, 136, e983-e992.	2.1	32
85	Maximising the impact of qualitative research in feasibility studies for randomised controlled trials: guidance for researchers. <i>Pilot and Feasibility Studies</i> , 2015, 1, 32.	1.2	240
86	What Difference Does Patient and Public Involvement Make and What Are Its Pathways to Impact? Qualitative Study of Patients and Researchers from a Cohort of Randomised Clinical Trials. <i>PLoS ONE</i> , 2015, 10, e0128817.	2.5	97
87	An evidence base to optimise methods for involving patient and public contributors in clinical trials: a mixed-methods study. <i>Health Services and Delivery Research</i> , 2015, 3, 1-142.	1.4	23
88	Interventions to improve recruitment and retention in clinical trials: a survey and workshop to assess current practice and future priorities. <i>Trials</i> , 2014, 15, 399.	1.6	279
89	Patient and public involvement in the early stages of clinical trial development: a systematic cohort investigation. <i>BMJ Open</i> , 2014, 4, e005234-e005234.	1.9	48
90	From plans to actions in patient and public involvement: qualitative study of documented plans and the accounts of researchers and patients sampled from a cohort of clinical trials. <i>BMJ Open</i> , 2014, 4, e006400.	1.9	83

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91	Patient and Public Participation in Health Care: Can We Do It Better?. American Journal of Bioethics, 2014, 14, 17-18.	0.9	4
92	The CONSENSUS study: protocol for a mixed methods study to establish which outcomes should be included in a core outcome set for oropharyngeal cancer. Trials, 2014, 15, 168.	1.6	36
93	Doing challenging research studies in a patient-centred way: a qualitative study to inform a randomised controlled trial in the paediatric emergency care setting. BMJ Open, 2014, 4, e005045.	1.9	57
94	ADRIC: Adverse Drug Reactions In Children – a programme of research using mixed methods. Programme Grants for Applied Research, 2014, 2, 1-184.	1.0	17
95	Comparing doctors' and nurses' accounts of how they provide emotional care for parents of children with acute lymphoblastic leukaemia. Psycho-Oncology, 2013, 22, 260-267.	2.3	20
96	The validity of education and guidance for clinical communication in cancer care: Evidence-based practice will depend on practice-based evidence. Patient Education and Counseling, 2013, 90, 193-199.	2.2	25
97	Alleviating Emotional Distress in a Young Adult Survivor of Adolescent Cancer. Clinical Case Studies, 2013, 12, 22-38.	0.8	11
98	What can we learn from parents about enhancing participation in pharmacovigilance?. British Journal of Clinical Pharmacology, 2013, 75, 1109-1117.	2.4	23
99	The –information spectrum–™: a qualitative study of how breast cancer surgeons give information and of how their patients experience it. Psycho-Oncology, 2013, 22, 2364-2371.	2.3	10
100	Is communication guidance mistaken? Qualitative study of parent–oncologist communication in childhood cancer. British Journal of Cancer, 2013, 109, 836-843.	6.4	39
101	How experience makes a difference: practitioners–™ views on the use of deferred consent in paediatric and neonatal emergency care trials. BMC Medical Ethics, 2013, 14, 45.	2.4	45
102	New European Union regulation of clinical trials is conflicting on deferred consent in emergency situations. BMJ, The, 2013, 346, f667-f667.	6.0	9
103	Parents–™ Agendas in Paediatric Clinical Trial Recruitment Are Different from Researchers–™ and Often Remain Unvoiced: A Qualitative Study. PLoS ONE, 2013, 8, e67352.	2.5	54
104	Faith and Protection: The Construction of Hope by Parents of Children with Leukemia and Their Oncologists. Oncologist, 2012, 17, 398-404.	3.7	45
105	Standard 1: Consent and Recruitment. Pediatrics, 2012, 129, S118-S123.	2.1	49
106	The psychological processes involved in patient empowerment. Orphanet Journal of Rare Diseases, 2012, 7, A31.	2.7	10
107	What Parents of Children Who Have Received Emergency Care Think about Deferring Consent in Randomised Trials of Emergency Treatments: Postal Survey. PLoS ONE, 2012, 7, e35982.	2.5	40
108	Enhancing Communication about Paediatric Medicines: Lessons from a Qualitative Study of Parents' Experiences of Their Child's Suspected Adverse Drug Reaction. PLoS ONE, 2012, 7, e46022.	2.5	17

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109	LUCAS: a theoretically informed instrument to assess clinical communication in objective structured clinical examinations. <i>Medical Education</i> , 2012, 46, 267-276.	2.1	30
110	Incorporating Children's and Their Parents' Perspectives into Condition-Specific Quality-of-Life Instruments for Children with Cerebral Palsy: A Qualitative Study. <i>Value in Health</i> , 2011, 14, 705-711.	0.3	8
111	Creativity in clinical communication: from communication skills to skilled communication. <i>Medical Education</i> , 2011, 45, 217-226.	2.1	164
112	Is clinical communication pedagogy really as good as itâ€™s going to get?. <i>Medical Education</i> , 2011, 45, 961-962.	2.1	0
113	Examining the validity of the unitary theory of clinical relationships: Comparison of observed and experienced parentâ€™doctor interaction. <i>Patient Education and Counseling</i> , 2011, 85, 60-67.	2.2	24
114	Telling â€œEverythingâ€ but not â€œToo Muchâ€: The Surgeonâ€™s Dilemma in Consultations about Breast Cancer. <i>World Journal of Surgery</i> , 2011, 35, 2187-2195.	1.6	24
115	Integrative qualitative communication analysis of consultation and patient and practitioner perspectives: Towards a theory of authentic caring in clinical relationships. <i>Patient Education and Counseling</i> , 2011, 82, 448-454.	2.2	66
116	What parents think about being approached about childrenâ€™s trials, how this differs from what practitioners expect, and what this tells us about enhancing recruitment. <i>Trials</i> , 2011, 12, .	1.6	4
117	Development of the Verona coding definitions of emotional sequences to code health providersâ€™ responses (VR-CoDES-P) to patient cues and concerns. <i>Patient Education and Counseling</i> , 2011, 82, 149-155.	2.2	165
118	Parents' Experiences of Their Children's Presence in Discussions With Physicians About Leukemia. <i>Pediatrics</i> , 2011, 127, e1230-e1238.	2.1	34
119	Communication about Children's Clinical Trials as Observed and Experienced: Qualitative Study of Parents and Practitioners. <i>PLoS ONE</i> , 2011, 6, e21604.	2.5	57
120	The ethics of responsibility and ownership in decision-making about treatment for breast cancer: Triangulation of consultation with patient and surgeon perspectives. <i>Social Science and Medicine</i> , 2010, 70, 1904-1911.	3.8	74
121	Balancing high accrual and ethical recruitment in paediatric oncology: a qualitative study of the 'look and feel' of clinical trial discussions. <i>BMC Medical Research Methodology</i> , 2010, 10, 101.	3.1	24
122	How do parents experience being asked to enter a child in a randomised controlled trial?. <i>BMC Medical Ethics</i> , 2009, 10, 1.	2.4	88
123	Dependence and caring in clinical communication: The relevance of attachment and other theories. <i>Patient Education and Counseling</i> , 2009, 74, 331-338.	2.2	93
124	Tissue Samples as â€˜Giftsâ€™ for Research: A Qualitative Study of Families and Professionals. <i>Medical Law International</i> , 2008, 9, 131-150.	1.1	26
125	Appraising qualitative research for inclusion in systematic reviews: a quantitative and qualitative comparison of three methods. <i>Journal of Health Services Research and Policy</i> , 2007, 12, 42-47.	1.7	410
126	A qualitative study of the healthâ€™related quality of life of disabled children. <i>Developmental Medicine and Child Neurology</i> , 2007, 49, 660-665.	2.1	50

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127	Decision-making in community-based paediatric physiotherapy: a qualitative study of children, parents and practitioners. <i>Health and Social Care in the Community</i> , 2006, 14, 116-124.	1.6	38
128	How can systematic reviews incorporate qualitative research? A critical perspective. <i>Qualitative Research</i> , 2006, 6, 27-44.	3.5	603
129	Researching chronic childhood illness: the example of childhood cancer. <i>Chronic Illness</i> , 2006, 2, 165-177.	1.5	8
130	Synthesising qualitative and quantitative evidence: A review of possible methods. <i>Journal of Health Services Research and Policy</i> , 2005, 10, 45-53.	1.7	1,270
131	Core assumptions and research opportunities in clinical communication. <i>Patient Education and Counseling</i> , 2005, 58, 225-234.	2.2	73
132	Synthesising qualitative and quantitative evidence: a review of possible methods. <i>Journal of Health Services Research and Policy</i> , 2005, 10, 45-53.	1.7	511
133	Finding qualitative research: an evaluation of search strategies. <i>BMC Medical Research Methodology</i> , 2004, 4, 5.	3.1	212
134	A Systematic Review of Molecular and Biological Tumor Markers in Neuroblastoma. <i>Clinical Cancer Research</i> , 2004, 10, 4-12.	7.0	179
135	Mental health support for youth offending teams: a qualitative study. <i>Health and Social Care in the Community</i> , 2003, 11, 55-63.	1.6	33
136	Representing childhood cancer: accounts from newspapers and parents. <i>Sociology of Health and Illness</i> , 2003, 25, 143-164.	2.1	30
137	Primary Mental Health Workers within Youth Offending Teams: a new service model. <i>Journal of Adolescence</i> , 2003, 26, 185-199.	2.4	27
138	Managing communication with young people who have a potentially life threatening chronic illness: qualitative study of patients and parents. <i>BMJ: British Medical Journal</i> , 2003, 326, 305-305.	2.3	249
139	Developing New Mental Health Services for Looked after Children: A Focus Group Study. <i>Adoption & Fostering</i> , 2003, 27, 51-63.	0.5	23
140	Identity and role in parenting a child with cancer. <i>Developmental Neurorehabilitation</i> , 2002, 5, 209-214.	1.1	37
141	Parenting in a crisis: conceptualising mothers of children with cancer. <i>Social Science and Medicine</i> , 2002, 55, 1835-1847.	3.8	222
142	Lay evaluation of services for childhood asthma. <i>Health and Social Care in the Community</i> , 2002, 10, 503-511.	1.6	23
143	Teaching and learning about human sexuality in undergraduate medical education. <i>Medical Education</i> , 2002, 36, 432-440.	2.1	51
144	Childhood cancer and users' views: a critical perspective. <i>European Journal of Cancer Care</i> , 2002, 11, 173-177.	1.5	17

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145	Parents' accounts of obtaining a diagnosis of childhood cancer. <i>Lancet, The</i> , 2001, 357, 670-674.	13.7	142
146	Choosing and using services for sexual health: a qualitative study of women's views. <i>Sexually Transmitted Infections</i> , 2001, 77, 335-339.	1.9	63
147	Partnerships with children. <i>BMJ: British Medical Journal</i> , 1999, 319, 778-780.	2.3	66
148	A mixture model for sucking patterns of breast-fed infants. , 1998, 17, 395-405.		2
149	Incorporating Patients' Perspectives into the Initial Stages of Core Outcome Set Development: A Rapid Review of Qualitative Studies of Type 2 Diabetes. <i>SSRN Electronic Journal</i> , 0, , .	0.4	0