

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	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
2	The COMET Handbook: version 1.0. <i>Trials</i> , 2017, 18, 280.	1.6	1,128
3	How can systematic reviews incorporate qualitative research? A critical perspective. <i>Qualitative Research</i> , 2006, 6, 27-44.	3.5	603
4	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
5	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
6	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
7	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
8	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
9	Parenting in a crisis: conceptualising mothers of children with cancer. <i>Social Science and Medicine</i> , 2002, 55, 1835-1847.	3.8	222
10	Finding qualitative research: an evaluation of search strategies. <i>BMC Medical Research Methodology</i> , 2004, 4, 5.	3.1	212
11	A Systematic Review of Molecular and Biological Tumor Markers in Neuroblastoma. <i>Clinical Cancer Research</i> , 2004, 10, 4-12.	7.0	179
12	Development of the Verona coding definitions of emotional sequences to code health providers'™ responses (VR-CoDES-P) to patient cues and concerns. <i>Patient Education and Counseling</i> , 2011, 82, 149-155.	2.2	165
13	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
14	Creativity in clinical communication: from communication skills to skilled communication. <i>Medical Education</i> , 2011, 45, 217-226.	2.1	164
15	Parents' accounts of obtaining a diagnosis of childhood cancer. <i>Lancet, The</i> , 2001, 357, 670-674.	13.7	142
16	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
17	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
18	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

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19	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
20	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
21	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
22	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
23	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
24	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
25	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
26	Core assumptions and research opportunities in clinical communication. <i>Patient Education and Counseling</i> , 2005, 58, 225-234.	2.2	73
27	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
28	Partnerships with children. <i>BMJ: British Medical Journal</i> , 1999, 319, 778-780.	2.3	66
29	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
30	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
31	Doing challenging research studies in a patient-centred way: a qualitative study to inform a randomised controlled trial in the paediatric emergency care setting. <i>BMJ Open</i> , 2014, 4, e005045.	1.9	57
32	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
33	Parents' Agendas in Paediatric Clinical Trial Recruitment Are Different from Researchers' and Often Remain Unvoiced: A Qualitative Study. <i>PLoS ONE</i> , 2013, 8, e67352.	2.5	54
34	Teaching and learning about human sexuality in undergraduate medical education. <i>Medical Education</i> , 2002, 36, 432-440.	2.1	51
35	A qualitative study of the health-related quality of life of disabled children. <i>Developmental Medicine and Child Neurology</i> , 2007, 49, 660-665.	2.1	50
36	Standard 1: Consent and Recruitment. <i>Pediatrics</i> , 2012, 129, S118-S123.	2.1	49

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37	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
38	A new paradigm for clinical communication: critical review of literature in cancer care. <i>Medical Education</i> , 2017, 51, 258-268.	2.1	48
39	Faith and Protection: The Construction of Hope by Parents of Children with Leukemia and Their Oncologists. <i>Oncologist</i> , 2012, 17, 398-404.	3.7	45
40	How experience makes a difference: practitioners' views on the use of deferred consent in paediatric and neonatal emergency care trials. <i>BMC Medical Ethics</i> , 2013, 14, 45.	2.4	45
41	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
42	Integrating quantitative and qualitative data and findings when undertaking randomised controlled trials. <i>BMJ Open</i> , 2019, 9, e032081.	1.9	42
43	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
44	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
45	What Parents of Children Who Have Received Emergency Care Think about Deferring Consent in Randomised Trials of Emergency Treatments: Postal Survey. <i>PLoS ONE</i> , 2012, 7, e35982.	2.5	40
46	Is communication guidance mistaken? Qualitative study of parent- <i>oncologist communication in childhood cancer. British Journal of Cancer</i> , 2013, 109, 836-843.	6.4	39
47	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
48	Fifteen-minute consultation: an evidence-based approach to research without prior consent (deferred) <i>TJ ETQq0 0 0 rgBT /Overlock 10 T and Practice Edition</i> , 2016, 101, 49-53.	0.5	38
49	Identity and role in parenting a child with cancer. <i>Developmental Neurorehabilitation</i> , 2002, 5, 209-214.	1.1	37
50	The CONSENSUS study: protocol for a mixed methods study to establish which outcomes should be included in a core outcome set for oropharyngeal cancer. <i>Trials</i> , 2014, 15, 168.	1.6	36
51	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
52	Parents' Experiences of Their Children's Presence in Discussions With Physicians About Leukemia. <i>Pediatrics</i> , 2011, 127, e1230-e1238.	2.1	34
53	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
54	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

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55	Clinical Trial Decisions in Difficult Circumstances: Parental Consent Under Time Pressure. <i>Pediatrics</i> , 2015, 136, e983-e992.	2.1	32
56	Representing childhood cancer: accounts from newspapers and parents. <i>Sociology of Health and Illness</i> , 2003, 25, 143-164.	2.1	30
57	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
58	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
59	Primary Mental Health Workers within Youth Offending Teams: a new service model. <i>Journal of Adolescence</i> , 2003, 26, 185-199.	2.4	27
60	CONTRACT Study - CONservative TReatment of Appendicitis in Children (feasibility): study protocol for a randomised controlled Trial. <i>Trials</i> , 2018, 19, 153.	1.6	27
61	Tissue Samples as "Gifts" for Research: A Qualitative Study of Families and Professionals. <i>Medical Law International</i> , 2008, 9, 131-150.	1.1	26
62	The validity of education and guidance for clinical communication in cancer care: Evidence-based practice will depend on practice-based evidence. <i>Patient Education and Counseling</i> , 2013, 90, 193-199.	2.2	25
63	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
64	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
65	Examining the validity of the unitary theory of clinical relationships: Comparison of observed and experienced parent-doctor interaction. <i>Patient Education and Counseling</i> , 2011, 85, 60-67.	2.2	24
66	Telling "Everything" but not "Too Much": The Surgeon's Dilemma in Consultations about Breast Cancer. <i>World Journal of Surgery</i> , 2011, 35, 2187-2195.	1.6	24
67	Lay evaluation of services for childhood asthma. <i>Health and Social Care in the Community</i> , 2002, 10, 503-511.	1.6	23
68	Developing New Mental Health Services for Looked after Children: A Focus Group Study. <i>Adoption & Fostering</i> , 2003, 27, 51-63.	0.5	23
69	What can we learn from parents about enhancing participation in pharmacovigilance?. <i>British Journal of Clinical Pharmacology</i> , 2013, 75, 1109-1117.	2.4	23
70	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
71	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
72	Incorporating patients' 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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73	Comparing doctors' and nurses' accounts of how they provide emotional care for parents of children with acute lymphoblastic leukaemia. <i>Psycho-Oncology</i> , 2013, 22, 260-267.	2.3	20
74	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
75	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
76	What can we learn from trial decliners about improving recruitment? Qualitative study. <i>Trials</i> , 2016, 17, 494.	1.6	19
77	Qualitative methods can test and challenge what we think we know about clinical communication "if they are not too constrained by methodological "brands". <i>Patient Education and Counseling</i> , 2018, 101, 1515-1517.	2.2	19
78	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
79	Uncertainty and test anxiety: Psychometric properties of the Intolerance of Uncertainty Scale " 12 (IUS-12) among university students. <i>International Journal of Educational Research</i> , 2020, 104, 101672.	2.2	18
80	Patient participation in Delphi surveys to develop core outcome sets: systematic review. <i>BMJ Open</i> , 2021, 11, e051066.	1.9	18
81	Childhood cancer and users' views: a critical perspective. <i>European Journal of Cancer Care</i> , 2002, 11, 173-177.	1.5	17
82	Enhancing Communication about Paediatric Medicines: Lessons from a Qualitative Study of Parents' Experiences of Their Child's Suspected Adverse Drug Reaction. <i>PLoS ONE</i> , 2012, 7, e46022.	2.5	17
83	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
84	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
85	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
86	ADRIC: Adverse Drug Reactions In Children " a programme of research using mixed methods. <i>Programme Grants for Applied Research</i> , 2014, 2, 1-184.	1.0	17
87	In-depth qualitative interviews identified barriers and facilitators that influenced chief investigators' use of core outcome sets in randomised controlled trials. <i>Journal of Clinical Epidemiology</i> , 2022, 144, 111-120.	5.0	17
88	Why trials lose participants: A multitrail investigation of participants' perspectives using the theoretical domains framework. <i>Journal of Clinical Epidemiology</i> , 2021, 137, 1-13.	5.0	16
89	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
90	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

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91	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
92	Systematic Techniques to Enhance rEtention in Randomised controlled trials: the STEER study protocol. <i>Trials</i> , 2018, 19, 197.	1.6	14
93	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
94	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
95	Research with children and young people: not on them. <i>Archives of Disease in Childhood</i> , 2016, 101, 1086-1089.	1.9	13
96	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
97	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
98	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
99	Alleviating Emotional Distress in a Young Adult Survivor of Adolescent Cancer. <i>Clinical Case Studies</i> , 2013, 12, 22-38.	0.8	11
100	The psychological processes involved in patient empowerment. <i>Orphanet Journal of Rare Diseases</i> , 2012, 7, A31.	2.7	10
101	The "information spectrum"™: a qualitative study of how breast cancer surgeons give information and of how their patients experience it. <i>Psycho-Oncology</i> , 2013, 22, 2364-2371.	2.3	10
102	How do surgeons think they learn about communication? A qualitative study. <i>Medical Education</i> , 2015, 49, 408-416.	2.1	10
103	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
104	New European Union regulation of clinical trials is conflicting on deferred consent in emergency situations. <i>BMJ, The</i> , 2013, 346, f667-f667.	6.0	9
105	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
106	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
107	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
108	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

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109	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
110	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
111	Researching chronic childhood illness: the example of childhood cancer. <i>Chronic Illness</i> , 2006, 2, 165-177.	1.5	8
112	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
113	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
114	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
115	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
116	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
117	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
118	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
119	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
120	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
121	Patient and Public Participation in Health Care: Can We Do It Better?. <i>American Journal of Bioethics</i> , 2014, 14, 17-18.	0.9	4
122	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
123	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
124	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
125	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
126	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

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127	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
128	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
129	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
130	A mixture model for sucking patterns of breast-fed infants. , 1998, 17, 395-405.		2
131	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
132	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
133	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
134	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
135	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
136	63â€…Methods of mobilising collective intelligence through crowdsourcing in research:a scoping review. , 2018, , .		2
137	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
138	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
139	Recruitersâ€™ perspectives and experiences of trial recruitment processes: a qualitative evidence synthesis protocol. <i>BMJ Open</i> , 2021, 11, e045233.	1.9	1
140	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
141	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
142	Is clinical communication pedagogy really as good as itâ€™s going to get?. <i>Medical Education</i> , 2011, 45, 961-962.	2.1	0
143	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
144	P07â€fIdentifying 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

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145	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
146	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
147	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. Emotion, Space and Society, 2021, 41, 100837.	1.5	0
148	Incorporating Patients' Perspectives into the Initial Stages of Core Outcome Set Development: A Rapid Review of Qualitative Studies of Type 2 Diabetes. SSRN Electronic Journal, 0, , .	0.4	0
149	Patient participation in Delphi surveys to develop core outcome sets: systematic review. BMJ Open, 2021, 11, e051066.	1.9	0