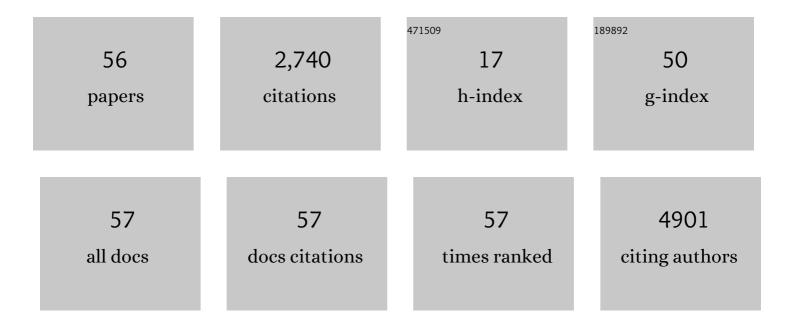
Sophie J Hill

List of Publications by Year in descending order

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SODHIE I HILL

#	Article	IF	CITATIONS
1	Knowledge translation of research findings. Implementation Science, 2012, 7, 50.	6.9	1,586
2	Reporting guideline for priority setting of health research (REPRISE). BMC Medical Research Methodology, 2019, 19, 243.	3.1	112
3	Development of the ACTIVE framework to describe stakeholder involvement in systematic reviews. Journal of Health Services Research and Policy, 2019, 24, 245-255.	1.7	84
4	Stakeholder involvement in systematic reviews: a scoping review. Systematic Reviews, 2018, 7, 208.	5.3	81
5	Factors affecting the implementation of childhood vaccination communication strategies in Nigeria: a qualitative study. BMC Public Health, 2017, 17, 200.	2.9	75
6	Online health information seeking: how people with multiple sclerosis find, assess and integrate treatment information to manage their health. Health Expectations, 2016, 19, 727-737.	2.6	68
7	Research priorities in health communication and participation: international survey of consumers and other stakeholders. BMJ Open, 2018, 8, e019481.	1.9	58
8	Shared Decision Making Interventions: Theoretical and Empirical Evidence with Implications for Health Literacy. Studies in Health Technology and Informatics, 2017, 240, 263-283.	0.3	58
9	Web Search Behavior and Information Needs of People With Multiple Sclerosis: Focus Group Study and Analysis of Online Postings. Interactive Journal of Medical Research, 2014, 3, e12.	1.4	51
10	Perceptions of HIV cure research among people living with HIV in Australia. PLoS ONE, 2018, 13, e0202647.	2.5	38
11	An integrative review of stakeholder views on Advance Care Directives (ACD): Barriers and facilitators to initiation, documentation, storage, and implementation. Patient Education and Counseling, 2019, 102, 1067-1079.	2.2	35
12	Stakeholder involvement in systematic reviews: a protocol for a systematic review of methods, outcomes and effects. Research Involvement and Engagement, 2017, 3, 9.	2.9	34
13	The comprehensive †Communicate to Vaccinate' taxonomy of communication interventions for childhood vaccination in routine and campaign contexts. BMC Public Health, 2017, 17, 423.	2.9	32
14	Selecting, refining and identifying priority Cochrane Reviews in health communication and participation in partnership with consumers and other stakeholders. Health Research Policy and Systems, 2019, 17, 45.	2.8	26
15	The significance and expectations of HIV cure research among people living with HIV in Australia. PLoS ONE, 2020, 15, e0229733.	2.5	24
16	An integrative review of patient safety in studies on the care and safety of patients with communication disabilities in hospital. Patient Education and Counseling, 2016, 99, 501-511.	2.2	22
17	Presenting evidence-based health information for people with multiple sclerosis: the IN-DEEP project protocol. BMC Medical Informatics and Decision Making, 2012, 12, 20.	3.0	19
18	A systematic review of falls in hospital for patients with communication disability: Highlighting an invisible population. Journal of Safety Research, 2019, 68, 89-105.	3.6	19

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19	Supporting implementation of Cochrane methods in complex communication reviews: resources developed and lessons learned for editorial practice and policy. Health Research Policy and Systems, 2019, 17, 32.	2.8	18
20	Mapping how information about childhood vaccination is communicated in two regions of Cameroon: What is done and where are the gaps?. BMC Public Health, 2015, 15, 1264.	2.9	17
21	Harnessing and supporting consumer involvement in the development and implementation of Models of Care for musculoskeletal health. Best Practice and Research in Clinical Rheumatology, 2016, 30, 420-444.	3.3	17
22	Stakeholder perceptions of communication about vaccination in two regions of Cameroon: A qualitative case study. PLoS ONE, 2017, 12, e0183721.	2.5	17
23	Development and assessment of a website presenting evidence-based information for people with multiple sclerosis: the IN-DEEP project. BMC Neurology, 2016, 16, 30.	1.8	15
24	Qualitative focus groups with stakeholders identify new potential outcomes related to vaccination communication. PLoS ONE, 2018, 13, e0201145.	2.5	15
25	Childhood vaccination communication outcomes unpacked and organized in a taxonomy to facilitate core outcome establishment. Journal of Clinical Epidemiology, 2017, 84, 173-184.	5.0	14
26	How could health information be improved? Recommended actions from the Victorian Consultation on Health Literacy. Australian Health Review, 2018, 42, 134.	1.1	14
27	Outcomes mapping study for childhood vaccination communication: too few concepts were measured in too many ways. Journal of Clinical Epidemiology, 2016, 72, 33-44.	5.0	13
28	Cross-sectional examination of musculoskeletal conditions and multimorbidity: influence of different thresholds and definitions on prevalence and association estimates. BMC Research Notes, 2017, 10, 51.	1.4	13
29	Identification of preliminary core outcome domains for communication about childhood vaccination: An online Delphi survey. Vaccine, 2018, 36, 6520-6528.	3.8	13
30	Consumer engagement critical to success in an Australian research project: reflections from those involved. Australian Journal of Primary Health, 2018, 24, 197.	0.9	13
31	Endorsement of the OMERACT core domain set for shared decision making interventions in rheumatology trials: Results from a multi-stepped consensus-building approach. Seminars in Arthritis and Rheumatism, 2021, 51, 593-600.	3.4	13
32	Effects of consumers and health providers working in partnership on health services planning, delivery and evaluation. The Cochrane Library, 2021, 2021, CD013373.	2.8	12
33	A call for dysphagia-related safety incident research in people with developmental disabilities. Journal of Intellectual and Developmental Disability, 2015, 40, 99-103.	1.6	11
34	Associations between multimorbidity and additional burden for working-age adults with specific forms of musculoskeletal conditions: a cross-sectional study. BMC Musculoskeletal Disorders, 2017, 18, 135.	1.9	11
35	Changing definitions altered multimorbidity prevalence, but not burden associations, in a musculoskeletal population. Journal of Clinical Epidemiology, 2016, 78, 116-126.	5.0	10
36	Legal, ethical, and rights issues in the adoption and use of the "My Health Record―by people with communication disability in Australia. Journal of Intellectual and Developmental Disability, 2018, 43, 506-514.	1.6	10

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37	"The Right Way at the Right Timeâ€: Insights on the Uptake of Falls Prevention Strategies from People with Dementia and Their Caregivers. Frontiers in Public Health, 2016, 4, 244.	2.7	9
38	Making rational choices about how best to support consumers' use of medicines: a perspective review. Therapeutic Advances in Drug Safety, 2016, 7, 159-164.	2.4	7
39	Producing an evidenceâ€based treatment information website in partnership with people affected by multiple sclerosis. Health Science Reports, 2018, 1, e24.	1.5	7
40	Falls prevention for people with dementia: a knowledge translation intervention. Dementia, 2020, 19, 2267-2293.	2.0	7
41	Public perspectives on acquired brain injury rehabilitation and components of care: A Citizens' Jury. Health Expectations, 2021, 24, 352-362.	2.6	7
42	Medications and doctor–patient communication. Australian Journal of General Practice, 2021, 50, 709-714.	0.8	7
43	Stakeholder Involvement in Systematic Reviews: Lessons From Cochrane's Public Health and Health Systems Network. American Journal of Public Health, 2021, 111, 1210-1215.	2.7	5
44	Instruments that measure psychosocial factors related to vaccination: a scoping review protocol. BMJ Open, 2019, 9, e033938.	1.9	4
45	Using the COMMVAC taxonomy to map vaccination communication interventions in Mozambique. Global Health Action, 2017, 10, 1321313.	1.9	3
46	Evaluation of the Cochrane Consumers and Communication Group's systematic review priority-setting project. Health Research Policy and Systems, 2020, 18, 98.	2.8	3
47	Sharing knowledge of falls prevention for people with dementia: insights for community care practice. Australian Journal of Primary Health, 2017, 23, 464.	0.9	2
48	Health Information and the Quality and Safety of Care for People With Disability. Journal of Patient Safety, 2018, Publish Ahead of Print, e1559-e1575.	1.7	2
49	Health Information Infrastructure for People with Intellectual and Developmental Disabilities (I/DD) Living in Supported Accommodation: Communication, Co-Ordination and Integration of Health Information. Health Communication, 2019, 34, 91-99.	3.1	2
50	HIV cure research: print and online media reporting in Australia. Journal of Virus Eradication, 2017, 3, 229-235.	0.5	2
51	Cochrane review ―interventions to improve safe and effective medicines use by consumers: an overview of systematic review. Journal of Evidence-Based Medicine, 2014, 7, 231-231.	2.4	1
52	Communication Vulnerabilities in Working-age Australians with Musculoskeletal Conditions: A Cross-sectional Analysis. Journal of Rheumatology, 2015, 42, 1484-1493.	2.0	1
53	Evaluation of a training program for medicines-oriented policymakers to use a database of systematic reviews. Health Research Policy and Systems, 2016, 14, 70.	2.8	1
54	Social Media as a Tool for Consumer Engagement in Hospital Quality Improvement and Service Design: Barriers and Enablers for Implementation. International Journal of Health Policy and Management, 2021, , .	0.9	1

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55	The Personally Controlled Electronic Health Record (PCEHR) for Adults with Severe Communication Impairments: Findings of Pilot Research. Studies in Health Technology and Informatics, 2015, 214, 100-6.	0.3	1
56	Mixed methods study to understand the experiences of adults with acquired brain injury and their family members who receive specialised rehabilitation. Brain Impairment, 2023, 24, 39-53.	0.7	0