## Sophie J Hill

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3216687/publications.pdf

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56	2,740	17 h-index	50
papers	citations		g-index
57	57	57	4901 citing authors
all docs	docs citations	times ranked	

#	Article	IF	CITATIONS
1	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	O
2	Public perspectives on acquired brain injury rehabilitation and components of care: A Citizens' Jury. Health Expectations, 2021, 24, 352-362.	2.6	7
3	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
4	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
5	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
6	Medications and doctor–patient communication. Australian Journal of General Practice, 2021, 50, 709-714.	0.8	7
7	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
8	Falls prevention for people with dementia: a knowledge translation intervention. Dementia, 2020, 19, 2267-2293.	2.0	7
9	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
10	The significance and expectations of HIV cure research among people living with HIV in Australia. PLoS ONE, 2020, 15, e0229733.	2.5	24
11	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
12	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
13	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
14	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
15	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
16	Instruments that measure psychosocial factors related to vaccination: a scoping review protocol. BMJ Open, 2019, 9, e033938.	1.9	4
17	Reporting guideline for priority setting of health research (REPRISE). BMC Medical Research Methodology, 2019, 19, 243.	3.1	112
18	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

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19	Producing an evidenceâ€based treatment information website in partnership with people affected by multiple sclerosis. Health Science Reports, 2018, 1, e24.	1.5	7
20	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
21	How could health information be improved? Recommended actions from the Victorian Consultation on Health Literacy. Australian Health Review, 2018, 42, 134.	1.1	14
22	Identification of preliminary core outcome domains for communication about childhood vaccination: An online Delphi survey. Vaccine, 2018, 36, 6520-6528.	3.8	13
23	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
24	Stakeholder involvement in systematic reviews: a scoping review. Systematic Reviews, 2018, 7, 208.	5.3	81
25	Perceptions of HIV cure research among people living with HIV in Australia. PLoS ONE, 2018, 13, e0202647.	2.5	38
26	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
27	Qualitative focus groups with stakeholders identify new potential outcomes related to vaccination communication. PLoS ONE, 2018, 13, e0201145.	2.5	15
28	Research priorities in health communication and participation: international survey of consumers and other stakeholders. BMJ Open, 2018, 8, e019481.	1.9	58
29	Factors affecting the implementation of childhood vaccination communication strategies in Nigeria: a qualitative study. BMC Public Health, 2017, 17, 200.	2.9	75
30	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
31	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
32	Using the COMMVAC taxonomy to map vaccination communication interventions in Mozambique. Global Health Action, 2017, 10, 1321313.	1.9	3
33	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
34	The comprehensive †Communicate to Vaccinate†to taxonomy of communication interventions for childhood vaccination in routine and campaign contexts. BMC Public Health, 2017, 17, 423.	2.9	32
35	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
36	Stakeholder perceptions of communication about vaccination in two regions of Cameroon: A qualitative case study. PLoS ONE, 2017, 12, e0183721.	2.5	17

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37	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
38	HIV cure research: print and online media reporting in Australia. Journal of Virus Eradication, 2017, 3, 229-235.	0.5	2
39	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
40	"The Right Way at the Right Time†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
41	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
42	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
43	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
44	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
45	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
46	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
47	Changing definitions altered multimorbidity prevalence, but not burden associations, in a musculoskeletal population. Journal of Clinical Epidemiology, 2016, 78, 116-126.	5.0	10
48	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
49	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
50	Communication Vulnerabilities in Working-age Australians with Musculoskeletal Conditions: A Cross-sectional Analysis. Journal of Rheumatology, 2015, 42, 1484-1493.	2.0	1
51	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
52	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
53	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
54	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

## SOPHIE J HILL

#	Article	IF	CITATIONS
55	Knowledge translation of research findings. Implementation Science, 2012, 7, 50.	6.9	1,586
56	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