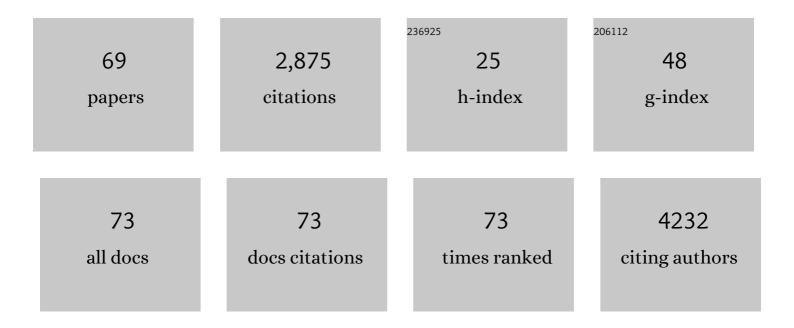
## Sophie Hill

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/11380638/publications.pdf Version: 2024-02-01



#	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	Ο
2	Interventions to improve safe and effective medicines use by consumers: an overview of systematic reviews. The Cochrane Library, 2022, 2022, CD007768.	2.8	140
3	Broadening the diversity of consumers engaged in guidelines: a scoping review. BMJ Open, 2022, 12, e058326.	1.9	4
4	Public perspectives on acquired brain injury rehabilitation and components of care: A Citizens' Jury. Health Expectations, 2021, 24, 352-362.	2.6	7
5	Assessing the information quality and usability of <i>My Health Record</i> within a health literacy framework: What's changed since 2016?. Health Information Management Journal, 2021, 50, 13-25.	1.2	21
6	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
7	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
8	The significance and expectations of HIV cure research among people living with HIV in Australia. PLoS ONE, 2020, 15, e0229733.	2.5	24
9	A conceptual framework for patient-directed knowledge tools to support patient-centred care: Results from an evidence-informed consensus meeting. Patient Education and Counseling, 2019, 102, 1898-1904.	2.2	14
10	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
11	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
12	OMERACT Development of a Core Domain Set of Outcomes for Shared Decision-making Interventions. Journal of Rheumatology, 2019, 46, 1409-1414.	2.0	14
13	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
14	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
15	Instruments that measure psychosocial factors related to vaccination: a scoping review protocol. BMJ Open, 2019, 9, e033938.	1.9	4
16	Reporting guideline for priority setting of health research (REPRISE). BMC Medical Research Methodology, 2019, 19, 243.	3.1	112
17	Research priorities for childhood chronic conditions: a workshop report. Archives of Disease in Childhood, 2019, 104, 237-245.	1.9	16
18	A content analysis of documentation on communication disability in hospital progress notes: diagnosis, function, and patient safety. Clinical Rehabilitation, 2019, 33, 943-956.	2.2	10

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19	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
20	Research priority setting in childhood chronic disease: a systematic review. Archives of Disease in Childhood, 2018, 103, 942-951.	1.9	41
21	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
22	A content analysis of the consumer-facing online information about My Health Record: Implications for increasing knowledge and awareness to facilitate uptake and use. Health Information Management Journal, 2018, 47, 106-115.	1.2	16
23	The health literacy demands of electronic personal health records (e-PHRs): An integrative review to inform future inclusive research. Patient Education and Counseling, 2018, 101, 2-15.	2.2	37
24	Identification of preliminary core outcome domains for communication about childhood vaccination: An online Delphi survey. Vaccine, 2018, 36, 6520-6528.	3.8	13
25	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
26	Stakeholder involvement in systematic reviews: a scoping review. Systematic Reviews, 2018, 7, 208.	5.3	81
27	Perceptions of HIV cure research among people living with HIV in Australia. PLoS ONE, 2018, 13, e0202647.	2.5	38
28	Qualitative focus groups with stakeholders identify new potential outcomes related to vaccination communication. PLoS ONE, 2018, 13, e0201145.	2.5	15
29	Face-to-face interventions for informing or educating parents about early childhood vaccination. The Cochrane Library, 2018, 2018, CD010038.	2.8	110
30	Research priority setting in organ transplantation: a systematic review. Transplant International, 2017, 30, 327-343.	1.6	30
31	Factors affecting the implementation of childhood vaccination communication strategies in Nigeria: a qualitative study. BMC Public Health, 2017, 17, 200.	2.9	75
32	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
33	Living systematic reviews: 4. Living guideline recommendations. Journal of Clinical Epidemiology, 2017, 91, 47-53.	5.0	184
34	Living systematic review: 1. Introduction—the why, what, when, and how. Journal of Clinical Epidemiology, 2017, 91, 23-30.	5.0	406
35	Living systematic reviews: 2. Combining human and machine effort. Journal of Clinical Epidemiology, 2017, 91, 31-37.	5.0	246
36	Living systematic reviews: 3. Statistical methods for updating meta-analyses. Journal of Clinical Epidemiology, 2017, 91, 38-46.	5.0	102

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37	Direct comparison of methionine restriction with leucine restriction on the metabolic health of C57BL/6J mice. Scientific Reports, 2017, 7, 9977.	3.3	54
38	Toward the Development of a Core Set of Outcome Domains to Assess Shared Decision-making Interventions in Rheumatology: Results from an OMERACT Delphi Survey and Consensus Meeting. Journal of Rheumatology, 2017, 44, 1544-1550.	2.0	21
39	Using the COMMVAC taxonomy to map vaccination communication interventions in Mozambique. Global Health Action, 2017, 10, 1321313.	1.9	3
40	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
41	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
42	HIV cure research: print and online media reporting in Australia. Journal of Virus Eradication, 2017, 3, 229-235.	0.5	4
43	Stakeholder perceptions of communication about vaccination in two regions of Cameroon: A qualitative case study. PLoS ONE, 2017, 12, e0183721.	2.5	17
44	Perceptions and experiences of childhood vaccination communication strategies among caregivers and health workers in Nigeria: A qualitative study. PLoS ONE, 2017, 12, e0186733.	2.5	25
45	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
46	Present and future of the undergraduate ophthalmology curriculum: a survey of UK medical schools. International Journal of Medical Education, 2017, 8, 389-395.	1.2	20
47	HIV cure research: print and online media reporting in Australia. Journal of Virus Eradication, 2017, 3, 229-235.	0.5	2
48	The E-health Literacy Demands of Australia's My Health Record: A Heuristic Evaluation of Usability. Perspectives in Health Information Management / AHIMA, American Health Information Management Association, 2017, 14, 1f.	0.0	16
49	"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
50	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
51	Dissemination of Clinical Practice Guidelines. Medical Decision Making, 2016, 36, 692-702.	2.4	23
52	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
53	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
54	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

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55	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
56	Use of the My Health Record by people with communication disability in Australia. Health Information Management Journal, 2016, 45, 107-115.	1.2	20
57	Communication strategies to promote the uptake of childhood vaccination in Nigeria: a systematic map. Global Health Action, 2016, 9, 30337.	1.9	32
58	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
59	Research Priorities in CKD: Report of a National Workshop Conducted in Australia. American Journal of Kidney Diseases, 2015, 66, 212-222.	1.9	73
60	Translating Falls Prevention Knowledge to Community-Dwelling Older PLWD: A Mixed-Method Systematic Review. Gerontologist, The, 2015, 55, 560-574.	3.9	20
61	Comparing Face-to-Face and Online Qualitative Research With People With Multiple Sclerosis. Qualitative Health Research, 2014, 24, 431-438.	2.1	52
62	"Communicate to vaccinate― the development of a taxonomy of communication interventions to improve routine childhood vaccination. BMC International Health and Human Rights, 2013, 13, 23.	2.5	41
63	Face to face interventions for informing or educating parents about early childhood vaccination. The Cochrane Library, 2013, , CD010038.	2.8	120
64	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
65	Consumer-oriented interventions for evidence-based prescribing and medicines use: an overview of systematic reviews. , 2011, , CD007768.		63
66	Development of a taxonomy of interventions to organise the evidence on consumers' medicines use. Patient Education and Counseling, 2011, 85, e101-e107.	2.2	26
67	'Communicate to vaccinate' (COMMVAC). building evidence for improving communication about childhood vaccinations in low- and middle-income countries: protocol for a programme of research. Implementation Science, 2011, 6, 125.	6.9	37
68	The Rx for Change database: a first-in-class tool for optimal prescribing and medicines use. Implementation Science, 2010, 5, 89.	6.9	25
69	Directions in health communication. Bulletin of the World Health Organization, 2009, 87, 648-648.	3.3	11