Karen M Facey

List of Publications by Year in descending order

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623734 552781 29 859 14 26 citations g-index h-index papers 30 30 30 930 times ranked docs citations citing authors all docs

#	Article	IF	CITATIONS
1	Patients' perspectives in health technology assessment: A route to robust evidence and fair deliberation. International Journal of Technology Assessment in Health Care, 2010, 26, 334-340.	0.5	205
2	What principles should govern the use of managed entry agreements?. International Journal of Technology Assessment in Health Care, 2011, 27, 77-83.	0.5	123
3	Recommendations from the European Working Group for Value Assessment and Funding Processes in Rare Diseases (ORPH-VAL). Orphanet Journal of Rare Diseases, 2017, 12, 50.	2.7	72
4	EUPATI Guidance for Patient Involvement in Medicines Research and Development: Health Technology Assessment. Frontiers in Medicine, 2018, 5, 231.	2.6	47
5	HTA programme response to the challenges of dealing with orphan medicinal products: Process evaluation in selected European countries. Health Policy, 2019, 123, 140-151.	3.0	43
6	Real-world evidence to support Payer/HTA decisions about highly innovative technologies in the EU—actions for stakeholders. International Journal of Technology Assessment in Health Care, 2020, 36, 459-468.	0.5	35
7	EVALUATION OF PATIENT AND PUBLIC INVOLVEMENT INITIATIVES IN HEALTH TECHNOLOGY ASSESSMENT: A SURVEY OF INTERNATIONAL AGENCIES. International Journal of Technology Assessment in Health Care, 2017, 33, 715-723.	0.5	31
8	GENERATING HEALTH TECHNOLOGY ASSESSMENT EVIDENCE FOR RARE DISEASES. International Journal of Technology Assessment in Health Care, 2014, 30, 416-422.	0.5	30
9	Virtual community consultation? Using the literature and weblogs to link community perspectives and health technology assessment. Health Expectations, 2008, 11, 189-200.	2.6	27
10	IMPROVING THE EFFECTIVENESS AND EFFICIENCY OF EVIDENCE PRODUCTION FOR HEALTH TECHNOLOGY ASSESSMENT. International Journal of Technology Assessment in Health Care, 2015, 31, 201-206.	0.5	27
11	Are supplemental appraisal/reimbursement processes needed for rare disease treatments? An international comparison of country approaches. Orphanet Journal of Rare Diseases, 2020, 15, 189.	2.7	26
12	Implementing Outcomes-Based Managed Entry Agreements for Rare Disease Treatments: Nusinersen and Tisagenlecleucel. Pharmacoeconomics, 2021, 39, 1021-1044.	3.3	23
13	The Evolution of AIFA Registries to Support Managed Entry Agreements for Orphan Medicinal Products in Italy. Frontiers in Pharmacology, 2021, 12, 699466.	3.5	22
14	Putting Patients at the Centre of Healthcare: Progress and Challenges for Health Technology Assessments. Patient, 2018, 11, 581-589.	2.7	20
15	The use of nonrandomized evidence to estimate treatment effects in health technology assessment. Journal of Comparative Effectiveness Research, 2021, 10, 1035-1043.	1.4	18
16	Stories of Patient Involvement Impact in Health Technology Assessments: A Discussion Paper. International Journal of Technology Assessment in Health Care, 2019, 35, 266-272.	0.5	15
17	Patient-focused HTAs. International Journal of Technology Assessment in Health Care, 2011, 27, 273-274.	0.5	14
18	Designing and Implementing Deliberative Processes for Health Technology Assessment: A Good Practices Report of a Joint HTAi/ISPOR Task Force. Value in Health, 2022, 25, 869-886.	0.3	11

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19	Designing and Implementing Deliberative Processes for Health Technology Assessment: A Good Practices Report of a Joint HTAi/ISPOR Task Force. International Journal of Technology Assessment in Health Care, 2022, 38, .	0.5	8
20	As health technology assessment evolves so must its approach to patient involvement. Journal of Comparative Effectiveness Research, 2019, 8, 549-554.	1.4	7
21	Examining the impact of different country processes for appraising rare disease treatments: a case study analysis. International Journal of Technology Assessment in Health Care, 2021, 37, e65.	0.5	7
22	Developing the Mosaic of Patient Participation in HTA. , 2017, , 51-66.		7
23	Consideration of quality of life in the health technology assessments of rare disease treatments. European Journal of Health Economics, 2022, 23, 645-669.	2.8	7
24	The Imperative for Patient-Centred Research to Develop Better Quality Services in Rare Diseases. Patient, $2015, 8, 1-3$.	2.7	5
25	Patient Input to HTA., 2017, , 67-79.		5
26	Measuring What Matters: Little Evidence Supporting the Content Validity of EQ-5D in People with Duchenne Muscular Dystrophy and Their Caregivers. Medical Decision Making, 2022, 42, 139-140.	2.4	5
27	A framework for action to improve patient and public involvement in health technology assessment. International Journal of Technology Assessment in Health Care, 2022, 38, .	0.5	5
28	Factors Affecting Citizen Trust and Public Engagement Relating to the Generation and Use of Real-World Evidence in Healthcare. International Journal of Environmental Research and Public Health, 2022, 19, 1674.	2.6	4
29	Reflections for Future Development. , 2017, , 419-427.		3