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Bringing the public into health technology assessment and coverage policy decisions: from principles to practice

DOI: 10.1016/j.healthpol.2006.07.009
Health Policy, 2007, 82, 37-50.

Source: <https://exaly.com/paper-pdf/42043004/citation-report.pdf>

Version: 2024-04-20

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#	Paper	IF	Citations
186	The role of economic evidence in Canadian oncology reimbursement decision-making: to lambda and beyond. <i>Value in Health</i> , 2008 , 11, 771-83	3.3	80
185	Engaging the public in priority-setting for health technology assessment: findings from a citizens' jury. 2008 , 11, 282-93		86
184	Representation and legitimacy in health policy formulation at a national level: perspectives from a study of health technology eligibility procedures in the United Kingdom. <i>Health Policy</i> , 2008 , 85, 356-62	3.2	17
183	Public engagement in setting priorities in health care. <i>Cmaj</i> , 2008 , 179, 15-8	3.5	81
182	Guidance for considering ethical, legal, and social issues in health technology assessment: application to genetic screening. <i>International Journal of Technology Assessment in Health Care</i> , 2008 , 24, 412-22	1.8	22
181	Introducing patient perspective in health technology assessment at the local level. 2009 , 9, 54		28
180	New quality and quantity indices in science (NewQIS): the study protocol of an international project. 2009 , 4, 16		71
179	A knowledge synthesis of patient and public involvement in clinical practice guidelines: study protocol. 2009 , 4, 30		12
178	Health technology assessment in Canada: 20 years strong?. <i>Value in Health</i> , 2009 , 12 Suppl 2, S14-9	3.3	42
177	Public involvement in setting a national research agenda: a mixed methods evaluation. <i>Patient</i> , 2009 , 2, 179-90	3.7	32
176	Why consider patients' preferences? A discourse analysis of clinical practice guideline developers. 2009 , 47, 908-15		64
175	"It all depends": conceptualizing public involvement in the context of health technology assessment agencies. 2010 , 70, 1518-26		85
174	The Canadian biotechnology regulatory regime: The role of participation. 2010 , 32, 280-287		2
173	User's perspectives of barriers and facilitators to implementing quality colonoscopy services in Canada: a study protocol. 2010 , 5, 85		2
172	A Cost Decision Analysis for Diagnosing and Staging. 2010 , 101-110		
171	From actors to authors: a first account about the involvement of patients in the informed consent governance of a major Italian translational research hospital. 2010 , 33, 231-40		3
170	Patient and public involvement in clinical practice guidelines: a knowledge synthesis of existing programs. 2011 , 31, E45-74		107

169	Development of a framework for effective community engagement in Ontario, Canada. <i>Health Policy</i> , 2011 , 101, 59-69	3.2	15
168	Priority setting for systematic review of health care interventions in Nigeria. <i>Health Policy</i> , 2011 , 99, 244-9	3.2	9
167	Health technology assessment in Brazil: what do healthcare system players think about it?. 2011 , 129, 198-205		12
166	The integration of citizens into a science/policy network in genetics: governance arrangements and asymmetry in expertise. 2011 , 14, 261-71		23
165	Eliciting ethical and social values in health technology assessment: A participatory approach. 2011 , 73, 135-44		60
164	Introducing patients' and the public's perspectives to health technology assessment: A systematic review of international experiences. <i>International Journal of Technology Assessment in Health Care</i> , 2011 , 27, 31-42	1.8	106
163	Moving cautiously: Public involvement and the health technology assessment community. <i>International Journal of Technology Assessment in Health Care</i> , 2011 , 27, 43-9	1.8	28
162	Role of patient and public participation in health technology assessment and coverage decisions. 2011 , 11, 75-89		67
161	Stakeholder engagement in comparative effectiveness research: how will we measure success?. 2012 , 1, 397-407		29
160	Stakeholder participation in comparative effectiveness research: defining a framework for effective engagement. 2012 , 1, 181-194		175
159	The changing role of economic evaluation in valuing medical technologies. 2012 , 12, 711-23		14
158	An equity framework for health technology assessments. 2012 , 32, 428-41		29
157	Comparative effectiveness research: the experience of the National Institute for Health and Clinical Excellence. 2012 , 30, 4267-74		8
156	[Stakeholder participation in priority setting - a consideration of the normative status of quantitative and qualitative methods]. 2012 , 106, 412-7		
155	Stakeholders involvement by HTA Organisations: why is so different?. <i>Health Policy</i> , 2012 , 105, 236-45	3.2	19
154	Decision-making in healthcare: a practical application of partial least square path modelling to coverage of newborn screening programmes. <i>BMC Medical Informatics and Decision Making</i> , 2012 , 12, 83	3.6	17
153	Knowledge mobilization in the context of health technology assessment: an exploratory case study. 2012 , 10, 10		5
152	Patient involvement in a scientific advisory process: setting the research agenda for medical products. <i>Health Policy</i> , 2012 , 107, 231-42	3.2	31

151	A systematic review of coverage decision-making on health technologies-evidence from the real world. <i>Health Policy</i> , 2012 , 107, 218-30	3.2	42
150	The politics of health technology assessment in Poland. <i>Health Policy</i> , 2012 , 108, 178-93	3.2	20
149	What counts and how to count it: physicians' constructions of evidence in a disinvestment context. 2012 , 75, 2191-9		16
148	A pilot study to identify areas for further improvements in patient and public involvement in health technology assessments for medicines. <i>Patient</i> , 2012 , 5, 199-211	3.7	41
147	Involving patients in HTA activities at local level: a study protocol based on the collaboration between researchers and knowledge users. 2012 , 12, 14		15
146	Reflections on the evolution of health technology assessment in Europe. <i>Health Economics, Policy and Law</i> , 2012 , 7, 25-45	2.3	30
145	An international survey of the public engagement practices of health technology assessment organizations. <i>Value in Health</i> , 2013 , 16, 155-63	3.3	45
144	How are European birth-cohort studies engaging and consulting with young cohort members?. 2013 , 13, 56		12
143	Assessing the added value of health technologies: reconciling different perspectives. <i>Value in Health</i> , 2013 , 16, S7-13	3.3	50
142	Alternatives to seclusion and restraint in psychiatry and in long-term care facilities for the elderly: perspectives of service users and family members. <i>Patient</i> , 2013 , 6, 269-80	3.7	8
141	FDA Decisions and Public Deliberation:Challenges and Opportunities. 2013 , 73, S115-S126		15
140	Transparency vs. closed-door policy: do process characteristics have an impact on the outcomes of coverage decisions? A statistical analysis. <i>Health Policy</i> , 2013 , 112, 187-96	3.2	16
139	Ethics in Health Technology Assessment: Understanding Health Technologies as Policies. 2013 , 26, 72-76		5
138	Citizens' perspectives on personalized medicine: a qualitative public deliberation study. 2013 , 21, 1197-201		28
137	Public engagement in health priority setting in low- and middle-income countries: current trends and considerations for policy. 2013 , 10, e1001495		25
136	Link between process and appraisal in coverage decisions: an analysis with structural equation modeling. 2013 , 33, 1009-25		8
135	Enhancing citizen engagement in cancer screening through deliberative democracy. 2013 , 105, 380-6		46
134	Public engagement in health technology assessment and coverage decisions: a study of experiences in France, Germany, and the United Kingdom. <i>Journal of Health Politics, Policy and Law</i> , 2013 , 38, 89-122	2.6	29

133	Assessing the impacts of citizen deliberations on the health technology process. <i>International Journal of Technology Assessment in Health Care</i> , 2013 , 29, 282-9	1.8	28
132	Promotion of health sector reforms for health systems strengthening in Nigeria: perceptions of policy makers versus the general public on the Nigeria health systems performance. 2013 , 28, 541-53		2
131	Technological innovation and its effect on public health in the United States. 2013 , 6, 31-40		6
130	Emerging Therapeutic Enhancement Enabling Health Technologies and Their Discourses: What Is Discussed within the Health Domain?. 2013 , 1, 20-52		8
129	Towards a consumer-informed research agenda for aphasia: preliminary work. 2014 , 36, 1042-50		19
128	The use of research evidence on patient preferences in health care decision-making: issues, controversies and moving forward. 2014 , 14, 785-94		22
127	Australian Public Preferences for the Funding of New Health Technologies: A Comparison of Discrete Choice and Profile Case Best-Worst Scaling Methods. 2014 , 34, 638-54		35
126	Introducing the patient's perspective in hospital health technology assessment (HTA): the views of HTA producers, hospital managers and patients. 2014 , 17, 888-900		23
125	Mapping the impact of patient and public involvement on health and social care research: a systematic review. 2014 , 17, 637-50		650
124	The use of citizens' juries in health policy decision-making: a systematic review. 2014 , 109, 1-9		133
123	Involving patient in the early stages of health technology assessment (HTA): a study protocol. 2014 , 14, 273		12
122	'Practical' resources to support patient and family engagement in healthcare decisions: a scoping review. 2014 , 14, 175		59
121	New avenues within community engagement: addressing the ingenuity gap in our approach to health research and future provision of health care. 2014 , 1, 321-328		9
120	Barriers and facilitators influencing ethical evaluation in health technology assessment. <i>International Journal of Technology Assessment in Health Care</i> , 2015 , 31, 113-23	1.8	13
119	The ideal healthcare: priorities of people with chronic conditions and their carers. 2015 , 15, 551		16
118	Conceptualizing the use of public involvement in health policy decision-making. 2015 , 138, 14-21		33
117	Preferences for end-of-life care among community-dwelling older adults and patients with advanced cancer: A discrete choice experiment. <i>Health Policy</i> , 2015 , 119, 1482-9	3.2	35
116	Health Care Coverage Decision Making in Low- and Middle-Income Countries: Experiences from 25 Coverage Schemes. 2015 , 18, 265-71		7

115	Challenges in measuring the societal value of orphan drugs: insights from a canadian stated preference survey. <i>Patient</i> , 2015 , 8, 93-101	3.7	14
114	Which public and why deliberate?--A scoping review of public deliberation in public health and health policy research. 2015 , 131, 114-21		113
113	Why orphan drug coverage reimbursement decision-making needs patient and public involvement. <i>Health Policy</i> , 2015 , 119, 588-96	3.2	25
112	Participatory health councils and good governance: healthy democracy in Brazil?. <i>International Journal for Equity in Health</i> , 2015 , 14, 21	4.6	11
111	Participatory health system priority setting: Evidence from a budget experiment. 2015 , 146, 182-90		5
110	Prioritising patients for bariatric surgery: building public preferences from a discrete choice experiment into public policy. 2015 , 5, e008919		10
109	Expectations and values about expanded newborn screening: a public engagement study. 2015 , 18, 419-29		18
108	Deliberative Processes in Practice. <i>SSRN Electronic Journal</i> , 2016 ,	1	
107	Addressing Patient Needs and Public Commitments in Health Technology Innovation: A Review of Patient and Public Engagement in Technology Development and Assessment. <i>SSRN Electronic Journal</i> , 2016 ,	1	
106	The Future of Health Economics. 2016 ,		1
105	Towards a bioethics of innovation. 2016 , 42, 445-9		8
104	Developing a decision support system to link health technology assessment (HTA) reports to the health system policies in Iran. <i>Health Policy and Planning</i> , 2017 , 32, 504-515	3.4	3
103	PUBLIC AND PATIENT INVOLVEMENT IN HEALTH TECHNOLOGY ASSESSMENT: A FRAMEWORK FOR ACTION. <i>International Journal of Technology Assessment in Health Care</i> , 2016 , 32, 256-264	1.8	67
102	Attitudes to incorporating genomic risk assessments into population screening programs: the importance of purpose, context and deliberation. 2016 , 9, 25		7
101	Public participation in decision-making on the coverage of new antivirals for hepatitis C. <i>Journal of Health Organization and Management</i> , 2016 , 30, 769-85	1.9	15
100	Introduction: priority setting, equitable access and public involvement in health care. <i>Journal of Health Organization and Management</i> , 2016 , 30, 736-50	1.9	25
99	Deliberative Processes in Practice. <i>The International Library of Ethics, Law and Technology</i> , 2016 , 59-70	0.5	1
98	Big Picture Bioethics: Developing Democratic Policy in Contested Domains. <i>The International Library of Ethics, Law and Technology</i> , 2016 ,	0.5	1

97	Steps toward improving ethical evaluation in health technology assessment: a proposed framework. <i>BMC Medical Ethics</i> , 2016 , 17, 34	2.9	13
96	If you build it, they will come: unintended future uses of organised health data collections. <i>BMC Medical Ethics</i> , 2016 , 17, 54	2.9	33
95	Cognitive Enhancement: Social and Public Policy Issues. 2016 ,		5
94	Comparative Effectiveness Research in Health Technology Assessment. 2016 , 57-93		1
93	[Involving patients, the insured and the general public in healthcare decision making]. 2016 , 110-111, 36-44		4
92	Prioritization in Medicine. 2016 ,		3
91	Public and patient participation in health policy, care and research. 2017 , 2, 31-32		16
90	Identifying health system value dimensions: more than health gain?. <i>Health Economics, Policy and Law</i> , 2017 , 12, 387-400	2.3	3
89	Public consultation changes guidance on the use of health-care interventions. An observational study. 2017 , 20, 361-368		2
88	Value Assessment Frameworks for HTA Agencies: The Organization of Evidence-Informed Deliberative Processes. <i>Value in Health</i> , 2017 , 20, 256-260	3.3	50
87	STAKEHOLDER INVOLVEMENT THROUGHOUT HEALTH TECHNOLOGY ASSESSMENT: AN EXAMPLE FROM PALLIATIVE CARE. <i>International Journal of Technology Assessment in Health Care</i> , 2017 , 33, 552-561 ^{1,8}		5
86	Public drug policy for children in Canada. <i>Cmaj</i> , 2017 , 189, E990-E994	3.5	4
85	Criteria for the prioritization of public health interventions for climate-sensitive vector-borne diseases in Quebec. <i>PLoS ONE</i> , 2017 , 12, e0190049	3.7	5
84	OP35 Involving Members Of The Public In A National Screening Programme Health Technology Assessment. <i>International Journal of Technology Assessment in Health Care</i> , 2017 , 33, 16-16	1.8	
83	Engagement of Canadian Patients with Rare Diseases and Their Families in the Lifecycle of Therapy: A Qualitative Study. <i>Patient</i> , 2018 , 11, 353-359	3.7	4
82	Comparing Use of Health Technology Assessment in Pharmaceutical Policy among Earlier and More Recent Adopters in the European Union. <i>Value in Health Regional Issues</i> , 2018 , 16, 81-91	1.6	10
81	Health Technology Assessment. 2018 , 11-22		1
80	IMPACT OF HEALTH TECHNOLOGY ASSESSMENT REPORTS ON HOSPITAL DECISION MAKERS - 10-YEAR INSIGHT FROM A HOSPITAL UNIT IN SHERBROOKE, CANADA: IMPACT OF HEALTH TECHNOLOGY ASSESSMENT ON HOSPITAL DECISIONS. <i>International Journal of Technology Assessment in Health Care</i> , 2018 , 34, 383-389	1.8	6

79	The clinical application of gene editing: ethical and social issues. <i>Personalized Medicine</i> , 2019 , 16, 337-350.	2.2	15
78	Justice and public participation in universal health coverage: when is tiered coverage unfair and who should decide?. <i>Asian Bioethics Review</i> , 2019 , 11, 5-19	3.4	1
77	Multicriteria Decision Analysis to Support Health Technology Assessment Agencies: Benefits, Limitations, and the Way Forward. <i>Value in Health</i> , 2019 , 22, 1283-1288	3.3	47
76	Towards inclusive priority-setting for global health research projects: recommendations for sharing power with communities. <i>Health Policy and Planning</i> , 2019 , 34, 346-357	3.4	15
75	Understanding and using patient experiences as evidence in healthcare priority setting. <i>Cost Effectiveness and Resource Allocation</i> , 2019 , 17, 20	2.4	11
74	Involving patients and the public in medical and health care research studies: An exploratory survey on participant recruiting and representativeness from the perspective of study authors. <i>PLoS ONE</i> , 2019 , 14, e0204187	3.7	14
73	Citizens' juries can bring public voices on overdiagnosis into policy making. <i>BMJ, The</i> , 2019 , 364, l351	5.9	2
72	Methods Assessing Sociocultural Aspects of Health Technologies: Results of a Literature Review. <i>International Journal of Technology Assessment in Health Care</i> , 2019 , 35, 99-105	1.8	1
71	Inclusion of Marginalized Groups and Communities in Global Health Research Priority-Setting. <i>Journal of Empirical Research on Human Research Ethics</i> , 2019 , 14, 169-181	1.6	12
70	Creating sustainable health care systems. <i>Journal of Health Organization and Management</i> , 2019 , 33, 18-34	1.9	13
69	Identifying the Need for Good Practices in Health Technology Assessment: Summary of the ISPOR HTA Council Working Group Report on Good Practices in HTA. <i>Value in Health</i> , 2019 , 22, 13-20	3.3	45
68	Financial interests of patient organisations contributing to technology assessment at England's National Institute for Health and Care Excellence: policy review. <i>BMJ, The</i> , 2019 , 364, k5300	5.9	25
67	The Emerging Social Science Literature on Health Technology Assessment: A Narrative Review. <i>Value in Health</i> , 2020 , 23, 3-9	3.3	6
66	Preferences for Primary Healthcare Services Among Older Adults with Chronic Disease: A Discrete Choice Experiment. <i>Patient Preference and Adherence</i> , 2020 , 14, 1625-1637	2.4	1
65	[Beyond benefit evaluation: Considering the unintended consequences of telehealth]. <i>Ethics, Medicine and Public Health</i> , 2020 , 15, 100596	0.7	0
64	Developing a toolkit for engagement practice: sharing power with communities in priority-setting for global health research projects. <i>BMC Medical Ethics</i> , 2020 , 21, 21	2.9	7
63	Defining the role of the public in Health Technology Assessment (HTA) and HTA-informed decision-making processes. <i>International Journal of Technology Assessment in Health Care</i> , 2020 , 36, 87-95	1.8	8
62	Rethinking the electronic health record through the quadruple aim: time to align its value with the health system. <i>BMC Medical Informatics and Decision Making</i> , 2020 , 20, 32	3.6	10

61	(Re)defining legitimacy in Canadian drug assessment policy? Comparing ideas over time. <i>Health Economics, Policy and Law</i> , 2021 , 16, 424-439	2.3	3
60	What are Important Ways of Sharing Power in Health Research Priority Setting? Perspectives From People With Lived Experience and Members of the Public. <i>Journal of Empirical Research on Human Research Ethics</i> , 2021 , 16, 200-211	1.6	
59	Sharing power in global health research: an ethical toolkit for designing priority-setting processes that meaningfully include communities. <i>International Journal for Equity in Health</i> , 2021 , 20, 127	4.6	2
58	Health Technology Assessment Development in Vietnam: A Qualitative Study of Current Progress, Barriers, Facilitators, and Future Strategies. <i>International Journal of Environmental Research and Public Health</i> , 2021 , 18,	4.6	0
57	Implications of the Health Equity Perspective for the Right to Health. 2021 , 337-363		
56	What the UN Human Rights Treaty Bodies Tell Us about Economic Inequalities and Human Rights: An Empirical Analysis of Twenty Years of Practice. 2021 , 85-114		
55	Index. 2021 , 390-418		
54	Distributive Justice, and Economic and Social Rights. 2021 , 247-270		
53	Economic Inequality and the Right to Social Security: Contested Meanings and Potential Roles. 2021 , 295-315		
52	Human Rights and Economic Inequalities. 2021 ,		0
51	A Framework for Fiscal Justice: How Human Rights Can Change Public Finance. 2021 , 143-167		
50	How Can Economists Help Human Rights Practitioners to Measure Changes in Economic Inequalities?. 2021 , 115-140		
49	Constraints on Economic Inequality: Comparing Canada and the United States. 2021 , 63-84		
48	Global Tax Justice and Human Rights. 2021 , 168-192		
47	Education, Income Inequality and the Right to Participate in Cultural Life. 2021 , 316-336		
46	Fair Wages and a Decent Living: Paths to Greater Vertical Equality. 2021 , 271-294		
45	Growing Inequality and Risks to Social Rights in Our New Data Economy. 2021 , 193-216		
44	Caste, Economic Inequality, and Climate Justice in India. 2021 , 217-244		

43	Introduction. 2021 , 1-30		
42	Emerging Human Rights Norms and Standards on Vertical Inequalities. 2021 , 33-62		
41	The Potential Impact of the Right to Housing to Address Vertical Inequalities. 2021 , 364-389		
40	Achieving inclusive research priority-setting: what do people with lived experience and the public think is essential?. <i>BMC Medical Ethics</i> , 2021 , 22, 117	2.9	3
39	Methodology for constructing scenarios for health policy research: The case of coverage decision-making for drugs for rare diseases in Canada. <i>Technological Forecasting and Social Change</i> , 2021 , 171, 120960	9.5	0
38	Patient and public involvement in health technology assessment: update of a systematic review of international experiences. <i>International Journal of Technology Assessment in Health Care</i> , 2021 , 37, e36	1.8	3
37	Comparative Effectiveness Research in Health Technology Assessment. 2015 , 1-38		2
36	Involving Patients in Hospital-Based HTA: Experiences, Approaches, and Future Directions. 2016 , 345-359		1
35	Evaluation of Patient Involvement in HTA. 2017 , 201-213		6
34	Discussion: Research to Promote Patient-Based HTA. 2017 , 225-233		1
33	Reflections on Terms, Goals and Organisation. 2017 , 31-42		1
32	Just Access? Questions of Equity in Access and Funding for Assistive Technology. <i>Ethics and Behavior</i> , 2019 , 29, 172-191	1.4	16
31	"Getting to the Table": Changing Ideas about Public and Patient Involvement in Canadian Drug Assessment. <i>Journal of Health Politics, Policy and Law</i> , 2019 , 44, 631-663	2.6	9
30	Harnessing the potential to quantify public preferences for healthcare priorities through citizens' juries. <i>International Journal of Health Policy and Management</i> , 2014 , 3, 57-62	2.5	19
29	Collaboration Between Researchers and Knowledge Users in Health Technology Assessment: A Qualitative Exploratory Study. <i>International Journal of Health Policy and Management</i> , 2017 , 6, 437-446	2.5	4
28	Stakeholder Participation for Legitimate Priority Setting: A Checklist. <i>International Journal of Health Policy and Management</i> , 2018 , 7, 973-976	2.5	11
27	A Pilot Study to Identify Areas for Further Improvements in Patient and Public Involvement in Health Technology Assessments for Medicines. <i>Patient</i> , 2012 , 5, 199-211	3.7	13
26	Involving Citizen-Patients in the Development of Telehealth Services: Qualitative Study of Experts' and Citizen-Patients' Perspectives. <i>Journal of Participatory Medicine</i> , 2018 , 10, e10665	1.4	7

25	[Organizational and systemic conditions of citizen-patient involvement in the development of telehealth in Quebec]. <i>Sante Publique</i> , 2019 , Vol. 31, 125-135		0
24	Means of Knowledge Dissemination: Are the <i>Café Scientifique</i> and the Artistic Performance Equally Effective?. <i>Sociology Mind</i> , 2012 , 02, 191-199	0.1	8
23	Legal Governance in HTA: Environment, Health and Safety Issues / Ethical, Legal and Social Issues (EHSL/ELSI), the Ongoing Debate. <i>Canadian Journal of Bioethics</i> , 2020 , 3, 83	0.5	2
22	Pitfalls in reimbursement decisions for oncology drugs in South Korea: need for addressing the ethical dimensions in technology assessment. <i>Asian Pacific Journal of Cancer Prevention</i> , 2013 , 14, 3785-92	1.7	3
21	Where Public Health Meets Ethics. Conceptual Foundations and Practical Challenges of Public Health. <i>Public Health Ethics Analysis</i> , 2013 , 5-23	0.2	
20	Impacts of Public Participation on Public Budgeting Process of Kurdistan. <i>SSRN Electronic Journal</i> ,	1	0
19	Fair Innings as a Basis for Prioritization: An Empirical Perspective. 2016 , 179-196		
18	Seeking Community Views on Allocation of Scarce Resources in a Pandemic in Australia: Two Methods, Two Answers. <i>The International Library of Ethics, Law and Technology</i> , 2016 , 245-261	0.5	
17	Involving patients and publics in medical and health care research studies: an exploratory survey on participant recruiting and representativeness from the perspective of study authors.		1
16	Troutville: Where People Discuss Fairness Issues. <i>Canadian Journal of Bioethics</i> , 2020 , 3, 70	0.5	0
15	How good is good enough? Standards in policy decisions to cover new health technologies. <i>Healthcare Policy</i> , 2007 , 3, 91-101	1.1	14
14	Patient advocacy group involvement in health technology assessments: an observational study. <i>Research Involvement and Engagement</i> , 2021 , 7, 83	4.4	0
13	Solving the Evidence Interpretability Crisis in Health Technology Assessment: A Role for Mechanistic Models?. <i>Frontiers in Medical Technology</i> , 2022 , 4, 810315	1.9	0
12	Overcoming structural barriers to sharing power with communities in global health research priority-setting: Lessons from the Participation for Local Action project in Karnataka, India.. <i>Global Public Health</i> , 2022 , 1-19	3.5	1
11	Pharmacogenomics Implementation and Hurdles to Overcome; In the Context of a Developing Country.. <i>Iranian Journal of Pharmaceutical Research</i> , 2021 , 20, 92-106	1.1	
10	Mechanisms to Bridge the Gap Between Science and Politics in Evidence-Informed Policymaking: Mapping the Landscape. 2022 , 293-328		1
9	Potential Barriers of Patient Involvement in Health Technology Assessment in Central and Eastern European Countries. <i>Frontiers in Public Health</i> , 10,	6	
8	Public preferences regarding the priority setting criteria of health interventions for budget allocation: results of a survey of Iranian adults. 2022 , 22,		0

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- 6 Access to novel drugs and therapeutics for children and youth: Eliciting citizens' values to inform public funding decisions. ○
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- 3 Participation in health in the Americas: Bibliometric mapping of production, impact, visibility and collaboration. **2023**, 28, 487-500 ○
- 2 Towards conceptualizing patients as partners in health systems: a systematic review and descriptive synthesis. **2023**, 21, ○
- 1 Governance and Public Participation. **2023**, 29-45 ○