

CITATION REPORT

List of articles citing

Health consumer groups in the UK: a new social movement?

DOI: 10.1111/j.0141-9889.2004.00416.x

Sociology of Health and Illness, 2004, 26, 737-56.

Source: <https://exaly.com/paper-pdf/36806518/citation-report.pdf>

Version: 2024-04-28

This report has been generated based on the citations recorded by exaly.com for the above article. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

#	Paper	IF	Citations
135	"SLANG"--Sensitive Language and the New Genetics--an exploratory study. 2005 , 14, 415-21		16
134	New social and health movements: Issues of representation and change. 2006 , 16, 219-231		38
133	Introduction: Patient organization movements and new metamorphoses in patienthood. 2006 , 62, 529-37		62
132	Regaining Trust in Medicine: Professional and State Strategies. 2006 , 54, 621-636		56
131	Parental food allergy information needs: a qualitative study. 2007 , 92, 771-5		50
130	Bringing 'the public' into health technology assessment and coverage policy decisions: from principles to practice. 2007 , 82, 37-50		171
129	'Trusting blindly can be the biggest risk of all': organised resistance to childhood vaccination in the UK. <i>Sociology of Health and Illness</i> , 2007 , 29, 198-215	3	142
128	'No one agrees except for those of us who have it': endometriosis patients as an epistemological community. <i>Sociology of Health and Illness</i> , 2007 , 29, 957-82	3	53
127	The Expert Patients Programme: a paradox of patient empowerment and medical dominance. <i>Health and Social Care in the Community</i> , 2007 , 15, 426-38	2.6	93
126	Mental health promotion and non-profit health organisations. <i>Health and Social Care in the Community</i> , 2007 , 15, 553-60	2.6	14
125	In whose interest? Relationships between health consumer groups and the pharmaceutical industry in the UK. <i>Sociology of Health and Illness</i> , 2008 , 30, 929-43	3	38
124	Health consumer and patients' organizations in Europe: towards a comparative analysis. 2008 , 11, 85-94		59
123	The patient movement as an emancipation movement. 2008 , 11, 102-12		24
122	Engaging the public in priority-setting for health technology assessment: findings from a citizens' jury. 2008 , 11, 282-93		86
121	Attributes and views of families with food allergic children recruited from allergy clinics and from a consumer organization. 2008 , 19, 264-9		11
120	Withering the Citizen, Managing the Consumer: Complaints in Healthcare Settings. 2008 , 7, 233-243		17
119	Organizing homeless people: Exploring the emergence of a user organization in Denmark. 2008 , 28, 27-50		10

118	An exploratory study of canadian aboriginal online health care forums. 2008 , 23, 270-81		23
117	Reconciling Two Conflicting Tales of the English Health Policy Process Since 1997. 2008 , 3, 183-203		16
116	Disability activism: social model stalwarts and biological citizens. 2009 , 24, 677-688		74
115	Lay Involvement and Legitimacy: The Construction of Expertise and Participation within HEART UK. 2009 , 38, 254-273		15
114	Health professionals' enactment of their accountability obligations: doing the best they can. 2009 , 69, 1063-71		24
113	Consumer health organisations for people with diabetes and arthritis: who contacts them and why?. <i>Health and Social Care in the Community</i> , 2009 , 17, 628-35	2.6	11
112	Why carers of frail older people are not using available respite services: an Australian study. 2010 , 19, 2057-64		29
111	Towards a history of choice in UK health policy. <i>Sociology of Health and Illness</i> , 2009 , 31, 309-24	3	45
110	Who rules rare disease associations? A framework to understand their action. <i>Sociology of Health and Illness</i> , 2009 , 31, 979-93	3	14
109	Habermas and Social Movement Theory. 2009 , 3, 381-393		8
108	Social Aspects of Genetic Testing Technologies. 2009 , 3, 972-985		5
107	Alliances in action: Opportunities and threats to solidarity between workers and service users in health and social care disputes. 2009 , 7, 148-169		5
106	Harmed patients gaining voice: challenging dominant perspectives in the construction of medical harm and patient safety reforms. 2010 , 71, 510-516		44
105	Genetic risks and healthy choices: creating citizen-consumers of genetic services through empowerment and facilitation. <i>Sociology of Health and Illness</i> , 2010 , 32, 365-81	3	20
104	Patient organizations in Finland: increasing numbers and great variation. 2010 , 13, 221-33		8
103	Patient participation in collective healthcare decision making: the Dutch model. 2010 , 13, 73-85		101
102	Investigating referral pathways from primary care to consumer health organisations. 2010 , 16, 260-7		8
101	If they want to risk the health and well-being of their child, that's up to them—Long-term breastfeeding, risk and maternal identity. 2010 , 12, 357-367		39

100 Health Activism in the Age of Governance.

99	Exploring genetic responsibility for the self, family and kin in the case of hereditary raised cholesterol. 2011 , 72, 1760-7	32
98	Medicalization and Biomedicalization Revisited: Technoscience and Transformations of Health, Illness and American Medicine. 2011 , 173-199	35
97	Institutional change at the frontlines. 2011 , 6, 26-45	9
96	Increasing access to consumer health organisations among patients with chronic disease - a randomised trial of a print-based intervention. 2011 , 12, 245-54	3
95	Health activism. 2012 , 27, 429-34	15
94	When Subjects Bite Back: The Bristol Cancer Help Centre Study and Increasing Consumer Involvement in UK Medical Research in the 1990s. 2012 , 25, 500-519	4
93	Women Reclaiming Sustainable Livelihoods. 2012 ,	9
92	Disease Politics and Medical Research Funding: Three Ways Advocacy Shapes Policy. 2012 , 77, 780-803	85
91	Greater Choice and Control? Health Policy in England and the Online Health Consumer. 2012 , 4, 1	13
90	Assessing stakeholder opinion on relations between cancer patient groups and pharmaceutical companies in Europe. 2012 , 5, 127-39	5
89	Finding the positive in loss: stillbirth and its potential for parental empowerment. 2012 , 31, 98-103	12
88	Increasing patient centredness in outpatient care through closer collaboration with patient groups?: an exploratory study on the views of health care professionals working in quality management for office-based physicians in Germany. 2012 , 107, 249-57	13
87	Reclaiming a moral identity: stillbirth, stigma and 'moral mothers'. 2012 , 28, 476-80	38
86	Les associations d'usagers dans le domaine de la périnatalité: un engagement fondé sur la connaissance ?. 2012 , 4, 198-205	
85	Information Flow and Health Policy Literacy: The Role of the Media. 2012 , 3, 391-402	6
84	Unpacking the social media phenomenon: towards a research agenda. 2012 , 12, 109-119	117
83	Epilepsy health consumer groups and charities; how representative of patients are they? The results of a pilot study. 2013 , 22, 472-5	

82	Treatment burden among people with chronic illness: what are consumer health organizations saying?. 2013 , 9, 220-32	13
81	Extended roles and the dietitian: community adult enteral tube care. 2013 , 26, 298-305	6
80	Clinicians, researchers and community activism: lessons for mental health services from another field of medicine [HIV/AIDS]. 2013 , 37, 81-84	1
79	Social Movements and Protest. 182-212	
78	Social Movements and Protest. xi-xiv	
77	Social Movements and Protest. 151-181	
76	Social Movements and Protest. 213-234	
75	Social Movements and Protest. 235-248	
74	Social Movements and Protest. 249-272	
73	Social Movements and Protest. 1-9	
72	Social Movements and Protest. 10-41	
71	Social Movements and Protest. 42-76	
70	Social Movements and Protest. 77-111	
69	Social Movements and Protest. 112-150	
68	Practising childbirth activism: A politics of evidence. 2014 , 9, 129-152	17
67	The voluntary sector and health policy: the role of national level health consumer and patients' organisations in the UK. 2014 , 123, 202-9	21
66	The emergent modes of dementia activism. 2014 , 34, 623-644	34
65	Consumer health organisations for chronic conditions: why do some people access them and others don't?. 2014 , 15, 418-29	2

64	Self-help friendliness: A German approach for strengthening the cooperation between self-help groups and health care professionals. 2014 , 123, 217-25	17
63	Evidence-based activism: Patients' users' and activists' groups in knowledge society. 2014 , 9, 111-128	112
62	Goals and organisational structure of the movement for global mental health. 2014 , 8, 31	3
61	The Bodies Politic: Chronic Health Conditions and Voter Turnout in the 2008 Election. 2015 , 40, 1115-55	33
60	What binds biosociality? The collective effervescence of the parent-led conference. 2015 , 126, 1-8	9
59	Protest, politics and produce: a resource account of anti-genetically modified organism activism. 2015 , 20, 34-49	3
58	Exploring Human Rights-Based Activism as a Social Determinant of Health: Insights from Brazil and South Africa. 2016 , 8, 198-218	2
57	The sociology of childbirth: an autobiographical journey through four decades of research. <i>Sociology of Health and Illness</i> , 2016 , 38, 689-705	3 15
56	Gender, Health and Healthcare. 2016 ,	1
55	The politics of health mobilization in the United States: The promise and pitfalls of "disease constituencies". 2016 , 165, 246-254	18
54	Publics and Their Health Systems. 2016 ,	24
53	An Ethnographic-Discursive Approach to Parental Self-Help Groups: The Case of ADHD. 2016 , 26, 935-50	19
52	Regulatory theory: commercially sustainable markets rely upon satisfying the public interest in obtaining credible goods. 2017 , 12, 471-493	1
51	Listen to me, I'm talking: involvement and recovery. 2017 , 22, 111-123	1
50	Resisting Throughput Pressures: Physicians' and Patients' Strategies to Manage Hospital Discharge. 2017 , 58, 116-130	5
49	Governing new global health-care markets: the case of stem cell treatments. 2017 , 22, 76-91	10
48	Queering activism through the lens of the sociology of everyday life. 2018 , 66, 1194-1208	5
47	Health democracy in Europe: Cancer patient organization participation in health policy. 2018 , 21, 474-484	8

46	Assessing Patient Organization Participation in Health Policy: A Comparative Study in France and Italy. <i>International Journal of Health Policy and Management</i> , 2018 , 7, 48-58	2.5	12
45	Cancer patients' organisation participation in health policy decision-making: a snapshot/cluster analysis of the EU-28 countries. 2018 , 8, e018896		3
44	Representing Whom? U.K. Health Consumer and Patients' Organizations in the Policy Process. 2018 , 15, 341-349		5
43	Share and protect our health data: an evidence based approach to rare disease patients' perspectives on data sharing and data protection - quantitative survey and recommendations. 2019 , 14, 175		34
42	The Cult of the Expert 2019 , 3-35		
41	Empowerment. 2019 , 107-138		
40	Index. 2019 , 261-270		
39	I'd failed to produce a baby and I'd failed to notice when the baby was in distress \square The social construction of bereaved motherhood. 2019 , 74, 35-41		1
38	Inclusion. 2019 , 139-164		
37	Frenetically Standing Still. 2019 , 192-192		
36	Hyper-active Governance and Social Acceleration. 2019 , 230-232		
35	Notes. 2019 , 236-237		
34	Managing the Expert \square Politics Nexus 2019 , 36-66		
33	Defence. 2019 , 69-106		
32	Defend, Empower and Include. 2019 , 165-190		
31	Experts, Politics and Co-Production. 2019 , 214-225		
30	Data Collection. 2019 , 226-229		
29	Crosstabs of Media Coverage of Herceptin. 2019 , 233-235		

28	Preface. 2019 , xi-xiii		
27	Transactional Experiences of Existential Anxiety as a Barrier to Effective Humanistic Intervention. <i>Journal of Humanistic Psychology</i> , 2019 , 59, 185-210	0.9	3
26	Expertise, advocacy and activism: A qualitative study on the activities of prostate cancer peer support workers. <i>Health (United Kingdom)</i> , 2020 , 24, 21-37	1.9	6
25	Knowledge, evidence, expertise? The epistemics of experience in contemporary healthcare. <i>Evidence and Policy</i> , 2020 , 16, 267-284	2.1	3
24	Citizens under the umbrella: citizenship projects and the development of genetic umbrella organizations in the USA and the UK. <i>New Genetics and Society</i> , 2020 , 39, 148-172	1.9	1
23	Patient involvement in agenda-setting processes in health research policy: A boundary work perspective. <i>Science and Public Policy</i> , 2020 , 47, 246-255	1.8	2
22	"It is a different world in here": collective identification and shared experiential knowledge between psychiatric inpatients. <i>Sociology of Health and Illness</i> , 2020 , 42, 724-738	3	1
21	Peer navigators' role in supporting people living with human immunodeficiency virus in Australia: Qualitative exploration of general practitioners' perspectives. <i>Health and Social Care in the Community</i> , 2021 ,	2.6	1
20	Structural injustice and dismantling racism in health and healthcare. <i>Nursing Inquiry</i> , 2021 , e12441	2.2	2
19	Hyper-active Governance: How Governments Manage the Politics of Expertise. 2019 ,		17
18	Social Movements and Protest. 2014 ,		31
17	The Biopolitics of Chronic Illness: Biology, Power and Personhood. 2010 , 205-224		2
16	Biocapital, Biopolitics and Biosocialities: Reframing Health, Livelihoods and Environments with New Genetics and Biotechnology. 2012 , 221-237		2
15	Safe Enough to Share. <i>Proceedings of the ACM on Human-Computer Interaction</i> , 2019 , 3, 1-23	3.4	5
14	Negotiating Leadership in Interdisciplinary Co-Productive Research: A Case Study of An International Community-Based Project Between Collaborators From South Africa and the United Kingdom. <i>SAGE Open</i> , 2020 , 10, 215824402097159	1.5	1
13	Pourquoi s'associer? Quatre motifs d'entr� dans un collectif dans les associations de maladies rares. <i>Revue Francaise De Sociologie</i> , 2011 , 52, 719	0.9	2
12	Quand la puissance publique fait surgir et �quipe une mobilisation protestataire. <i>Revue Fran�aise De Science Politique</i> , 2011 , 61, 183	1.4	7
11	Why do Patients Protest? Collective Action Processes in People with Chronic Illnesses. <i>Advances in Medical Diagnosis, Treatment, and Care</i> , 2016 , 128-150	0.2	3

10	Pressure Groups. 2014 , 161-163		
9	Self-Help Groups. 2014 , 172-174		
8	Anxious Subjectivities and Spaces of Care: Therapeutic Geographies of the UK National Phobics Society. 2017 , 95-110		1
7	The Rise of the Consucrat. <i>International Journal of Health Policy and Management</i> , 2021 , 10, 176-180	2.5	5
6	Why Do Patients Protest? Collective Action Processes in People With Chronic Illnesses. 1197-1219		1
5	Coalition Priorit[Cancer and the pharmaceutical industry in Quebec: conflicts of interest in the reimbursement of expensive cancer drugs?. <i>Healthcare Policy</i> , 2013 , 9, 52-64	1.1	8
4	Safe Enough to Share: Setting the Dementia Agenda Online. 2019 , 3,		0
3	Health, Wellbeing, and Democratic Citizenship: A Review and Research Agenda. <i>LSE Public Policy Review</i> , 2021 , 2,	1.5	
2	Social Movements and Stakeholder Engagement. 2022 , 67-74		
1	Negotiating mental illness across the lay-professional divide: Role play in peer work consultations.. <i>Sociology of Health and Illness</i> , 2022 ,	3	0