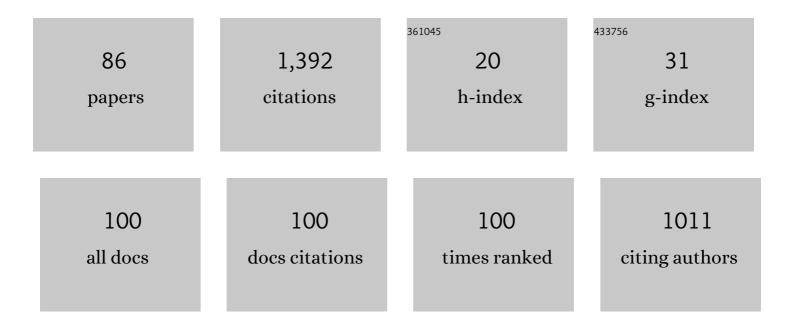
List of Publications by Year in descending order

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Δυίλο Ε Ρλζ

#	Article	IF	CITATIONS
1	Vibrational communication in subterranean mole rats (Spalax ehrenbergi). Behavioral Ecology and Sociobiology, 1987, 21, 31-33.	0.6	99
2	"Important to test, important to support― attitudes toward disability rights and prenatal diagnosis among leaders of support groups for genetic disorders in Israel. Social Science and Medicine, 2004, 59, 1857-1866.	1.8	87
3	Familial communication and cascade testing among relatives of BRCA population screening participants. Genetics in Medicine, 2018, 20, 1446-1454.	1.1	67
4	Carrier matching and collective socialization in community genetics: Dor Yeshorim and the reinforcement of stigma. Social Science and Medicine, 2008, 67, 1361-1369.	1.8	62
5	Population screening for BRCA1/BRCA2 founder mutations in Ashkenazi Jews: proactive recruitment compared with self-referral. Genetics in Medicine, 2017, 19, 754-762.	1.1	44
6	Representing autism: Challenges of collective representation in German and Israeli associations for and of autistic people. Social Science and Medicine, 2018, 200, 65-72.	1.8	42
7	Managerial Culture, Workplace Culture and Situated Curricula in Organizational Learning. Organization Studies, 2006, 27, 165-182.	3.8	40
8	Beyond cultural stereotyping: views on end-of-life decision making among religious and secular persons in the USA, Germany, and Israel. BMC Medical Ethics, 2017, 18, 13.	1.0	39
9	The mask of autism: Social camouflaging and impression management as coping/normalization from the perspectives of autistic adults. Social Science and Medicine, 2020, 248, 112826.	1.8	39
10	Diversity and uniformity in genetic responsibility: moral attitudes of patients, relatives and lay people in Germany and Israel. Medicine, Health Care and Philosophy, 2009, 12, 433-442.	0.9	36
11	The Mask of Dementia: Images of â€~Demented Residents' in a Nursing Ward. Ageing and Society, 1996, 16, 269-285.	1.2	33
12	Lay perceptions of genetic testing in Germany and Israel: the interplay of national culture and individual experience. New Genetics and Society, 2009, 28, 401-414.	0.7	31
13	Cousin marriage and premarital carrier matching in a Bedouin community in Israel: attitudes, service development and educational intervention. Journal of Family Planning and Reproductive Health Care, 2004, 30, 49-51.	0.9	29
14	Eugenic utopias/dystopias, reprogenetics, and community genetics. Sociology of Health and Illness, 2009, 31, 602-616.	1.1	29
15	Between Acculturation and Ambivalence: Knowledge of Genetics and Attitudes towards Genetic Testing in a Consanguineous Bedouin Community. Public Health Genomics, 2003, 6, 88-95.	0.6	26
16	The cultural context of patient's autonomy and doctor's duty: passive euthanasia and advance directives in Germany and Israel. Medicine, Health Care and Philosophy, 2010, 13, 363-369.	0.9	26
17	Population screening for BRCA1/BRCA2 mutations: lessons from qualitative analysis of the screening experience. Genetics in Medicine, 2017, 19, 628-634.	1.1	25
18	Pandora's pregnancy: NIPT, CMA, and genome sequencing—A new era for prenatal genetic testing. Prenatal Diagnosis, 2019, 39, 859-865.	1.1	25

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19	Doing gender in segregated and assimilative organizations: Ultraâ€Orthodox Jewish women in the Israeli highâ€tech labour market. Gender, Work and Organization, 2018, 25, 361-378.	3.1	24
20	One size does not fit all: Lessons from Israel's Covid-19 vaccination drive and hesitancy. Vaccine, 2021, 39, 4027-4028.	1.7	21
21	Abortions for fetuses with mild abnormalities. Israel Medical Association Journal, 2010, 12, 5-9.	0.1	21
22	Disability Rights, Prenatal Diagnosis and Eugenics: A Cross-Cultural View. Journal of Genetic Counseling, 2005, 14, 183-187.	0.9	20
23	Nondirectiveness and Its Lay Interpretations: The Effect of Counseling Style, Ethnicity and Culture on Attitudes Towards Genetic Counseling Among Jewish and Bedouin Respondents in Israel. Journal of Genetic Counseling, 2003, 12, 313-332.	0.9	19
24	"The Most Important Test You'll Ever Take"?: Attitudes toward confidential carrier matching and open individual testing among modern-religious Jews in Israel. Social Science and Medicine, 2011, 73, 1741-1747.	1.8	19
25	Between social hypocrisy and social responsibility: professional views of eugenics, disability and repro-genetics in Germany and Israel. New Genetics and Society, 2010, 29, 87-102.	0.7	18
26	Comparative Empirical Bioethics: Dilemmas of Genetic Testing and Euthanasia in Israel and Germany. SpringerBriefs in Ethics, 2016, , .	0.6	18
27	What a difference a role makes. Employee Relations, 2017, 39, 1131-1147.	1.5	18
28	Ambiguous professionalism: managing efficiency and service quality in an Israeli call centre. New Technology, Work and Employment, 2007, 22, 83-96.	2.6	16
29	The Cultural Context of End-of-Life Ethics: A Comparison of Germany and Israel. Cambridge Quarterly of Healthcare Ethics, 2010, 19, 381-394.	0.5	16
30	Domesticating Disney: Onstage Strategies of Adaptation in Tokyo Disneyland. Journal of Popular Culture, 2000, 33, 77-99.	0.0	15
31	Communities of practice or communities of coping?. Learning Organization, 2007, 14, 375-387.	0.7	15
32	Patient Representation and Advocacy for Alzheimer Disease in Germany and Israel. Journal of Bioethical Inquiry, 2018, 15, 369-380.	0.9	13
33	The hybridization of organizational culture in tokyo disneyland. Culture and Organization, 1999, 5, 235-264.	0.1	12
34	Exploring the Positions of German and Israeli Patient Organizations in the Bioethical Context of End-of-Life Policies. Health Care Analysis, 2014, 22, 143-159.	1.4	12
35	"What the patient wants…― Lay attitudes towards end-of-life decisions in Germany and Israel. Medicine, Health Care and Philosophy, 2015, 18, 329-340.	0.9	12
36	Donation of surplus frozen pre-embryos to research in Israel: underlying motivations. Israel Journal of Health Policy Research, 2016, 5, 25.	1.4	11

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37	When does a fetus become a person? An Israeli viewpoint. Journal of Family Planning and Reproductive Health Care, 2011, 37, 216-224.	0.9	10
38	Who Takes Part in the Political Game? The Sex Work Governance Debate in Israel. Sexuality Research and Social Policy, 2021, 18, 516-526.	1.4	10
39	Transparency, consent and trust in the use of customers' data by an online genetic testing company: an Exploratory survey among 23andMe users. New Genetics and Society, 2020, 39, 459-482.	0.7	10
40	Rituals of Exchange in the Social World of Israeli Beggars: An Exploratory Study. Symbolic Interaction, 1995, 18, 99-119.	0.7	9
41	The authorized self: How middle age defines old age in the postmodern. Semiotica, 1997, 113, .	0.2	9
42	Perceptions of Cousin Marriage Among Young Bedouin Adults in Israel. Marriage and Family Review, 2005, 37, 27-46.	0.7	9
43	Transplanting Management. Journal of Applied Behavioral Science, The, 2009, 45, 280-304.	2.0	9
44	"WE CAME TO TALK WITH THE PEOPLE BEHIND THE DISEASE:―COMMUNICATION AND CONTROL IN MEDIC EDUCATION. Culture, Medicine and Psychiatry, 2006, 30, 55-75.	CAL.7	8
45	Can Populationâ€Based Carrier Screening Be Left to the Community?. Journal of Genetic Counseling, 2009, 18, 114-118.	0.9	8
46	Attitudes of Israeli parents of children with Down syndrome toward nonâ€invasive prenatal screening and the scope of prenatal testing. Journal of Genetic Counseling, 2019, 28, 1119-1129.	0.9	8
47	`America' Meets `Japan'. Theory, Culture and Society, 1996, 13, 153-178.	1.3	7
48	Saving or Subordinating Life? Popular Views in Israel and Germany of Donor Siblings Created through PGD. Journal of Medical Humanities, 2017, 38, 191-207.	0.3	7
49	One For All, All For One? Collective Representation in Healthcare Policy. Journal of Bioethical Inquiry, 2018, 15, 337-340.	0.9	7
50	Coming to terms with the imperfectly normal child: attitudes of Israeli parents of screen-positive infants regarding subsequent prenatal diagnosis. Journal of Community Genetics, 2019, 10, 41-50.	0.5	6
51	Riding the Black Ship: Japan and Tokyo Disneyland. Monumenta Nipponica, 2000, 55, 315.	0.0	5
52	Chapter 8 Emotion Management in Cross-Cultural Perspective: "Smile Training―in Japanese and North American Service Organizations. Research on Emotion in Organizations, 0, , 199-220.	0.1	5
53	Commentary: a sociologist's view on community genetics. Journal of Community Genetics, 2010, 1, 3-10.	0.5	5
54	Parents like me: biosociality and lay expertise in self-help groups of parents of screen-positive newborns. New Genetics and Society, 2018, 37, 97-116.	0.7	5

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55	Reckless or Pioneering? Public Health Genetics Services in Israel. , 0, , 223-239.		5
56	Divergent evolution of newborn screening: Israel and the US as gene worlds. BioSocieties, 2018, 13, 580-600.	0.8	4
57	Epigenetic metaphors: an interdisciplinary translation of encoding and decoding. New Genetics and Society, 2019, 38, 264-288.	0.7	4
58	STATUS DISCLOSURE: GENETIC COUNSELING AS AN ARENA FOR NEGOTIATION. Studies in Symbolic Interaction, 2003, , 147-167.	0.3	4
59	Views of Israeli healthcare professionals regarding communication of genetic variants of uncertain significance to patients. Journal of Genetic Counseling, 2022, 31, 912-921.	0.9	4
60	Challenges for precision public health communication in the era of genomic medicine. Genetics in Medicine, 2022, 24, 1814-1820.	1.1	4
61	A Symbolic Interactionist User's Guide to the Answering Machine: 22 Reflections on Vocal Encounters in an Emerging Social World. Symbolic Interaction, 1994, 17, 411-429.	0.7	3
62	A NOTE ON INTER-VIEWING: USING SYMBOLIC INTERACTIONISM FOR INTERVIEW ANALYSIS. Studies in Symbolic Interaction, 0, , 323-339.	0.3	3
63	Frozen: social and bioethical aspects of cryopreservation. New Genetics and Society, 2020, 39, 243-249.	0.7	3
64	Unmet communication needs and moral work in the disposition decision concerning surplus frozen embryos: The perspectives of IVF users. Social Science and Medicine, 2021, 274, 113804.	1.8	3
65	Comparing Germany and Israel regarding debates on policy-making at the beginning of life: PGD, NIPT and their paths of routinization. Ethik in Der Medizin, 0, , 1.	1.0	3
66	Fragmented responsibility: views of Israeli HCPs regarding patient recontact following variant reclassification. Journal of Community Genetics, 2022, 13, 13-18.	0.5	3
67	"Donating with eyes shut― attitudes regarding DNA donation to a large-scale biobank in Israel. New Genetics and Society, 2022, 41, 47-65.	0.7	3
68	Poetry of unadulterated imagination: The late style of Akira Kurosawa Psychology of Aesthetics, Creativity, and the Arts, 2008, 2, 34-41.	1.0	2
69	Through the looking glass: engaging in a socio-ethical, cross-cultural dialogue. New Genetics and Society, 2010, 29, 55-59.	0.7	2
70	Life Stories, Status, and Symbolic Typing: Exploring Beggars As an Interpretive Community. Narrative Inquiry, 1996, 6, 123-143.	0.1	2
71	Views on disability and prenatal testing among families with Down syndrome and disability activists: A comparative analysis of interviews from Germany and Israel. Social Science and Medicine, 2022, 303, 115021.	1.8	2
72	Cancer patients' understandings of genetic variants of uncertain significance in clinical care. Journal of Community Genetics, 0, , .	0.5	2

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73	Emotions at Work: Normative Control, Organizations, and Culture in Japan and America. Contemporary Sociology, 2003, 32, 198.	0.0	1
74	Responsibility Revisited. Medicine Studies: an International Journal for History, Philosophy, and Ethics of Medicine and Allied Sciences, 2012, 3, 129-130.	0.1	1
75	Uncertain prophecies: How practitioners negotiate the prognostic ambivalence of 'disability' in prenatal diagnosis consultations. Social Science and Medicine, 2015, 146, 225-227.	1.8	1
76	Applying the Theoretical Tools: Being Affected, Responsibility, and Risk. SpringerBriefs in Ethics, 2016, , 21-43.	0.6	1
77	The Orient Strikes Back: A Global View of Cultural Display. American Ethnologist, 2001, 28, 947-948.	1.0	0
78	Testing Fate: Tay-Sachs and the Right to be Responsible by Shelley Z. Reuter. Shofar, 2017, 35, 135-137.	0.0	0
79	Patient Advocacy in Dementia: The Culture and Ethics of Policy-Making and Representation. Advances in Neuroethics, 2021, , 223-236.	0.1	0
80	Introduction: Engaging in Comparative Bioethics. SpringerBriefs in Ethics, 2016, , 1-11.	0.6	0
81	Risks and Responsibilities: Making Plans for Life and Death. SpringerBriefs in Ethics, 2016, , 95-105.	0.6	0
82	Lay Attitudes Towards End-of-Life Decision-Making in Germany and Israel. SpringerBriefs in Ethics, 2016, , 81-94.	0.6	0
83	Planning One's End of Life in an Expert Biomedical Culture. SpringerBriefs in Ethics, 2016, , 67-80.	0.6	0
84	Collective representation and the founders' culture in non-profit organisations: the case of Israel's national association for autism. Voluntary Sector Review, 2021, 12, 313-331.	0.2	0
85	Direct to consumer personal genomic testing and trust. , 2020, , 101-129.		0
86	A test of faith? Attitudes of ultraorthodox Jewish parents of children with down syndrome toward prenatal testing. Disability and Society, 2024, 39, 192-212.	1.4	0