

Anne F Townsend

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6353403/publications.pdf>

Version: 2024-02-01

24
papers

1,125
citations

430874

18
h-index

610901

24
g-index

24
all docs

24
docs citations

24
times ranked

1781
citing authors

#	ARTICLE	IF	CITATIONS
1	Self-managing and managing self: practical and moral dilemmas in accounts of living with chronic illness. <i>Chronic Illness</i> , 2006, 2, 185-194.	1.5	154
2	Managing multiple morbidity in mid-life: a qualitative study of attitudes to drug use. <i>BMJ: British Medical Journal</i> , 2003, 327, 837-0.	2.3	147
3	‘I want to know what's in Pandora's box’ Comparing stakeholder perspectives on incidental findings in clinical whole genomic sequencing. <i>American Journal of Medical Genetics, Part A</i> , 2012, 158A, 2519-2525.	1.2	135
4	Trust in Health Research Relationships: Accounts of Human Subjects. <i>Journal of Empirical Research on Human Research Ethics</i> , 2008, 3, 35-47.	1.3	86
5	eHealth, Participatory Medicine, and Ethical Care: A Focus Group Study of Patients’ and Health Care Providers’ Use of Health-Related Internet Information. <i>Journal of Medical Internet Research</i> , 2015, 17, e155.	4.3	85
6	Genetics professionals' perspectives on reporting incidental findings from clinical genome-wide sequencing. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 542-549.	1.2	55
7	Paternalism and the ACMG recommendations on genomic incidental findings: patients seen but not heard. <i>Genetics in Medicine</i> , 2013, 15, 751-752.	2.4	50
8	eHealth Technologies, Multimorbidity, and the Office Visit: Qualitative Interview Study on the Perspectives of Physicians and Nurses. <i>Journal of Medical Internet Research</i> , 2018, 20, e31.	4.3	45
9	Accessing health services through the back door: a qualitative interview study investigating reasons why people participate in health research in Canada. <i>BMC Medical Ethics</i> , 2013, 14, 40.	2.4	43
10	Putting context centre stage: evidence from a systems evaluation of an area based empowerment initiative in England. <i>Critical Public Health</i> , 2017, 27, 477-489.	2.4	39
11	Self-managing and managing self: practical and moral dilemmas in accounts of living with chronic illness. <i>Chronic Illness</i> , 2006, 2, 185-194.	1.5	35
12	Frequent consulting and multiple morbidity: a qualitative comparison of 'high' and 'low' consulters of GPs. <i>Family Practice</i> , 2008, 25, 168-175.	1.9	32
13	Applying Bourdieu’s theory to accounts of living with multimorbidity. <i>Chronic Illness</i> , 2012, 8, 89-101.	1.5	31
14	Exploring occupational disruption among women after onset of rheumatoid arthritis. <i>Arthritis Care and Research</i> , 2012, 64, 197-205.	3.4	31
15	A qualitative interview study: patient accounts of medication use in early rheumatoid arthritis from symptom onset to early postdiagnosis: Table 1. <i>BMJ Open</i> , 2013, 3, e002164.	1.9	29
16	Qualitative Research Ethics: Enhancing Evidence-Based Practice in Physical Therapy. <i>Physical Therapy</i> , 2010, 90, 615-628.	2.4	26
17	Reframing ‘participation’ and ‘inclusion’ in public health policy and practice to address health inequalities: Evidence from a major resident-led neighbourhood improvement initiative. <i>Health and Social Care in the Community</i> , 2019, 27, 199-206.	1.6	25
18	Exploring eHealth Ethics and Multi-Morbidity: Protocol for an Interview and Focus Group Study of Patient and Health Care Provider Views and Experiences of Using Digital Media for Health Purposes. <i>JMIR Research Protocols</i> , 2013, 2, e38.	1.0	25

#	ARTICLE	IF	CITATIONS
19	Women's accounts of help-seeking in early rheumatoid arthritis from symptom onset to diagnosis. <i>Chronic Illness</i> , 2014, 10, 259-272.	1.5	12
20	Using Physical Activity Trackers in Arthritis Self-Management: A Qualitative Study of Patient and Rehabilitation Professional Perspectives. <i>Arthritis Care and Research</i> , 2019, 71, 227-236.	3.4	12
21	Autonomy and the patient's right "not to know" in clinical whole-genomic sequencing. <i>European Journal of Human Genetics</i> , 2014, 22, 6-6.	2.8	9
22	Communications Between Volunteers and Health Researchers during Recruitment and Informed Consent: Qualitative Content Analysis of Email Interactions. <i>Journal of Medical Internet Research</i> , 2011, 13, e84.	4.3	8
23	Ethical issues experienced by persons with rheumatoid arthritis in a wearable-enabled physical activity intervention study. <i>Health Expectations</i> , 2022, 25, 1418-1431.	2.6	6
24	Evolving Patient-Researcher Collaboration: An Illustrative Case Study of a Patient-Led Knowledge Translation Event. <i>Journal of Participatory Medicine</i> , 2017, 9, e13.	1.3	5