

Cinzia Colombo

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/784663/publications.pdf>

Version: 2024-02-01

40
papers

562
citations

759233

12
h-index

677142

22
g-index

45
all docs

45
docs citations

45
times ranked

877
citing authors

#	ARTICLE	IF	CITATIONS
1	Online health information seeking: how people with multiple sclerosis find, assess and integrate treatment information to manage their health. <i>Health Expectations</i> , 2016, 19, 727-737.	2.6	68
2	Web Search Behavior and Information Needs of People With Multiple Sclerosis: Focus Group Study and Analysis of Online Postings. <i>Interactive Journal of Medical Research</i> , 2014, 3, e12.	1.4	51
3	Industry funding of patient and health consumer organisations: systematic review with meta-analysis. <i>BMJ, The</i> , 2020, 368, l6925.	6.0	44
4	Critical observations on and suggested ways forward for healthcare communication during COVID-19: pEACH position paper. <i>Patient Education and Counseling</i> , 2021, 104, 217-222.	2.2	41
5	Menopause: Knowledge, attitude and practice among Italian women. <i>Maturitas</i> , 2009, 63, 246-252.	2.4	40
6	Patient Organizations' Funding from Pharmaceutical Companies: Is Disclosure Clear, Complete and Accessible to the Public? An Italian Survey. <i>PLoS ONE</i> , 2012, 7, e34974.	2.5	33
7	PartecipaSalute, an Italian project to involve lay people, patients' associations and scientific-medical representatives in the health debate. <i>Health Expectations</i> , 2007, 10, 194-204.	2.6	30
8	Patient empowerment as a component of health system reforms: rights, benefits and vested interests. <i>Internal and Emergency Medicine</i> , 2012, 7, 183-187.	2.0	23
9	A dynamic web-based decision aid to improve informed choice in organised breast cancer screening. A pragmatic randomised trial in Italy. <i>British Journal of Cancer</i> , 2020, 123, 714-721.	6.4	22
10	Presenting evidence-based health information for people with multiple sclerosis: the IN-DEEP project protocol. <i>BMC Medical Informatics and Decision Making</i> , 2012, 12, 20.	3.0	19
11	Informing women about hormone replacement therapy: the consensus conference statement. <i>BMC Women's Health</i> , 2009, 9, 14.	2.0	17
12	Development and assessment of a website presenting evidence-based information for people with multiple sclerosis: the IN-DEEP project. <i>BMC Neurology</i> , 2016, 16, 30.	1.8	15
13	Sharing individual participant data from clinical studies: a cross-sectional online survey among Italian patient and citizen groups. <i>BMJ Open</i> , 2019, 9, e024863.	1.9	13
14	International Health Consumers in The Cochrane Collaboration. <i>Journal of Ambulatory Care Management</i> , 2010, 33, 182-189.	1.1	12
15	Family Burden after Severe Brain Injury. <i>Patient</i> , 2011, 4, 55-65.	2.7	12
16	A new informed consent form model for cancer patients: Preliminary results of a prospective study by the Italian Association of Medical Oncology (AIOM). <i>Patient Education and Counseling</i> , 2012, 87, 243-249.	2.2	12
17	Informing Women on Menopause and Hormone Therapy: Know the Menopause a Multidisciplinary Project Involving Local Healthcare System. <i>PLoS ONE</i> , 2013, 8, e85121.	2.5	12
18	A European multi-language initiative to make the general population aware of independent clinical research: the European Communication on Research Awareness Need project. <i>Trials</i> , 2016, 17, 19.	1.6	11

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19	Personalised informed choice on evidence and controversy on mammography screening: study protocol for a randomized controlled trial. <i>BMC Cancer</i> , 2017, 17, 429.	2.6	10
20	Involving a Citizensâ€™ Jury in Decisions on Individual Screening for Prostate Cancer. <i>PLoS ONE</i> , 2016, 11, e0143176.	2.5	10
21	Press coverage of hormone replacement therapy and menopause. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , 2010, 153, 56-61.	1.1	8
22	Consumers and health providers working in partnership for the promotion of person-centred health services: a co-produced qualitative evidence synthesis. <i>The Cochrane Library</i> , 0, .	2.8	8
23	Fostering a Strategic Alliance Between Patients' Associations and Health Care Professionals. <i>Journal of Ambulatory Care Management</i> , 2010, 33, 223-230.	1.1	7
24	Producing an evidence-based treatment information website in partnership with people affected by multiple sclerosis. <i>Health Science Reports</i> , 2018, 1, e24.	1.5	7
25	Does a consumer training work? a follow-up survey of the PartecipaSalute training programs. <i>Health Research Policy and Systems</i> , 2012, 10, 27.	2.8	6
26	Deciding on cystic fibrosis carrier screening: three citizensâ€™ juries and an online survey. <i>European Journal of Public Health</i> , 2018, 28, 973-977.	0.3	6
27	Oncologists' opinions about research ethics committees in Italy: an update, 2004. <i>European Journal of Cancer Prevention</i> , 2006, 15, 91-94.	1.3	5
28	CIGNOweb.it. <i>Tumori</i> , 2011, 97, 133-135.	1.1	3
29	Identifying unanswered questions and setting the agenda for future systematic research in Multiple Sclerosis. A worldwide, multi-stakeholder Priority Setting project. <i>Multiple Sclerosis and Related Disorders</i> , 2022, 60, 103688.	2.0	3
30	Role of Hormone Therapy in the Management of Menopause. <i>Obstetrics and Gynecology</i> , 2010, 116, 442.	2.4	2
31	Choosing Wisely Italy: online survey on opinions and behaviors of 1006 people and 355 volunteers of healthcare advocacy associations. <i>European Journal of Public Health</i> , 2019, 29, 396-401.	0.3	2
32	The CORBEL matrix on informed consent in clinical studies: a multidisciplinary approach of Research Infrastructures Building Enduring Life-science Services. <i>BMC Medical Ethics</i> , 2021, 22, 95.	2.4	2
33	Citizens' juries in health care. <i>BMJ: British Medical Journal</i> , 2010, 341, c5141-c5141.	2.3	2
34	HIV priorities by Italian AIDS advocacy groups: information on prevention (still) comes first. An on-line survey. <i>Epidemiologia E Prevenzione</i> , 2019, 43, 270-274.	1.1	2
35	Informing Women About Hormone Replacement Therapy. <i>Cancer Journal (Sudbury, Mass)</i> , 2009, 15, 344.	2.0	1
36	Introduction. <i>Journal of Ambulatory Care Management</i> , 2010, 33, 178-181.	1.1	1

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37	Patient-reported outcomes: nothing without engagement. <i>Acta Oncologica</i> , 2016, 55, 1494-1495.	1.8	1
38	The IN-DEEP project – Integrating and Deriving Evidence, Experiences, Preferences – a web information model on magnetic resonance imaging for people with multiple sclerosis. <i>Journal of Neurology</i> , 2020, 267, 2421-2431.	3.6	1
39	Transparency of funding of patient groups is mandatory but is not enough. <i>BMJ, The</i> , 2014, 349, g6301-g6301.	6.0	0
40	Citizens’ juries could help to guide screening policy. <i>BMJ, The</i> , 2016, 353, i3520.	6.0	0