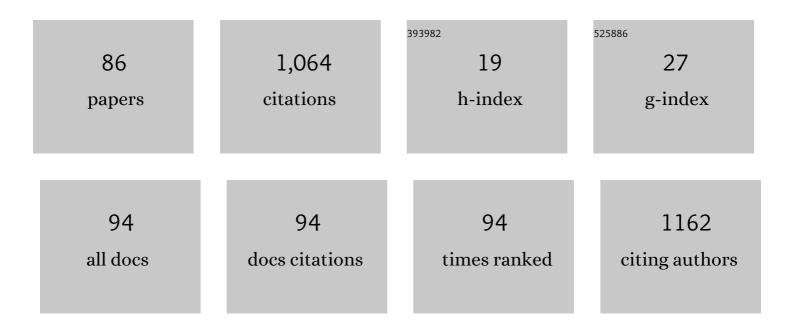


List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Parenting very preterm infants and stress in Neonatal Intensive Care Units. Early Human Development, 2016, 101, 3-9.	0.8	87
2	Distributed health literacy among people living with type 2 diabetes in Portugal: Defining levels of awareness and support. Health and Social Care in the Community, 2018, 26, 90-101.	0.7	60
3	Parents' views on factors that help or hinder breast milk supply in neonatal care units: systematic review. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2013, 98, F511-F517.	1.4	50
4	Factors Influencing Parent Reports of Facilitators and Barriers to Human Milk Supply in Neonatal Intensive Care Units. Journal of Human Lactation, 2016, 32, 695-703.	0.8	38
5	Factors associated with the donation and non-donation of embryos for research: a systematic review. Human Reproduction Update, 2014, 20, 641-655.	5.2	37
6	Socioeconomic Inequalities in the Prevalence of Nine Established Cardiovascular Risk Factors in a Southern European Population. PLoS ONE, 2012, 7, e37158.	1.1	35
7	Forensic DNA databases in European countries: is size linked to performance?. Life Sciences, Society and Policy, 2013, 9, .	3.1	35
8	Limited Health Literacy in Portugal Assessed with the Newest Vital Sign. Acta Medica Portuguesa, 2017, 30, 861-869.	0.2	33
9	Quality of life among parents of preterm infants: a scoping review. Quality of Life Research, 2018, 27, 1119-1131.	1.5	33
10	Public participation in genetic databases: crossing the boundaries between biobanks and forensic DNA databases through the principle of solidarity. Journal of Medical Ethics, 2015, 41, 820-824.	1.0	29
11	"Would you accept having your DNA profile inserted in the National Forensic DNA database? Why?― Results of a questionnaire applied in Portugal. Forensic Science International: Genetics, 2014, 8, 132-136.	1.6	28
12	What influences public views on forensic DNA testing in the criminal field? A scoping review of quantitative evidence. Human Genomics, 2019, 13, 23.	1.4	27
13	Awareness and motives for use and non-use of preimplantation genetic diagnosis in familial amyloid polyneuropathy mutation carriers. Prenatal Diagnosis, 2014, 34, 886-892.	1.1	25
14	Cross-cultural adaptation and validation of the health literacy assessment tool METER in the Portuguese adult population. Patient Education and Counseling, 2014, 97, 269-275.	1.0	24
15	Gold nanorods induce early embryonic developmental delay and lethality in zebrafish (<i>Danio) Tj ETQq1 1 0.7</i>	84314 rgE 1.1	BT /Qyerlock I
16	Socioeconomic position and quality of life among older people: The mediating role of social support. Preventive Medicine, 2020, 135, 106073.	1.6	24
17	Cancer screening in Portugal: sex differences in prevalence, awareness of organized programmes and perception of benefits and adverse effects. Health Expectations, 2017, 20, 211-220.	1.1	22
18	Needs of parents of very preterm infants in Neonatal Intensive Care Units: A mixed methods study. Intensive and Critical Care Nursing, 2019, 54, 88-95.	1.4	22

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19	Association between neighborhood deprivation and fruits and vegetables consumption and leisure-time physical activity: a cross-sectional multilevel analysis. BMC Public Health, 2013, 13, 1103.	1.2	21
20	Public and patient involvement in needs assessment and social innovation: a people-centred approach to care and research for congenital disorders of glycosylation. BMC Health Services Research, 2017, 17, 682.	0.9	20
21	Quality of life of parents of very preterm infants 4Âmonths after birth: a mixed methods study. Health and Quality of Life Outcomes, 2018, 16, 178.	1.0	19
22	Prisoners' expectations of the national forensic DNA database: Surveillance and reconfiguration of individual rights. Forensic Science International, 2011, 210, 139-143.	1.3	17
23	Patientâ€centered communication in type 2 diabetes: The facilitating and constraining factors in clinical encounters. Health Services Research, 2019, 54, 623-635.	1.0	17
24	Voluntary Participation in Forensic DNA Databases. Science Technology and Human Values, 2016, 41, 322-343.	1.7	16
25	Uncertainty, risks and ethics in unsuccessful in vitro fertilisation treatment cycles. Health, Risk and Society, 2010, 12, 531-545.	0.9	15
26	Informed Consent in Forensic DNA Databases: Volunteering, Constructions of Risk and Identity Categorization. BioSocieties, 2009, 4, 335-348.	0.8	14
27	Parenting roles and knowledge in neonatal intensive care units: protocol of a mixed methods study. BMJ Open, 2014, 4, e005941-e005941.	0.8	14
28	Assessment of the contents related to screening on Portuguese language websites providing information on breast and prostate cancer. Cadernos De Saude Publica, 2013, 29, 2163-2176.	0.4	13
29	Antenatal paternal adjustment and paternal attitudes after infertility treatment. Human Reproduction, 2018, 33, 109-115.	0.4	13
30	Public and patient involvement in health data governance (DATAGov): protocol of a people-centred, mixed-methods study on data use and sharing for rare diseases care and research. BMJ Open, 2021, 11, e044289.	0.8	13
31	Perspectives on access to in vitro fertilization in Portugal. Revista De Saude Publica, 2012, 46, 344-350.	0.7	12
32	Medicine and user involvement within European healthcare: a typology for European comparative research. International Journal of Clinical Practice, 2011, 65, 1218-1220.	0.8	11
33	Trust, morality and altruism in the donation of biological material: the case of Portugal. New Genetics and Society, 2009, 28, 103-118.	0.7	10
34	The construction of meaning by experts and wouldâ€be parents in assisted reproductive technology. Sociology of Health and Illness, 2011, 33, 853-868.	1.1	10
35	A short form of the neonatal intensive care unit family needs inventory. Jornal De Pediatria, 2016, 92, 73-80.	0.9	10
36	Factors associated with willingness to donate embryos for research among couples undergoing IVF. Reproductive BioMedicine Online, 2016, 32, 247-256.	1.1	10

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37	Public Perspectives on Risks and Benefits of Forensic DNA Databases. Bulletin of Science, Technology and Society, 2015, 35, 16-24.	1.1	9
38	The Role of Distributed Health Literacy in Asthma Integrated Care: A Public Medical Context from Portugal. International Journal of Integrated Care, 2018, 18, 18.	0.1	9
39	Patients' views on the embryo storage time limits. Reproductive BioMedicine Online, 2015, 31, 232-238.	1.1	8
40	Couples' willingness to donate embryos for research: a longitudinal study. Acta Obstetricia Et Gynecologica Scandinavica, 2016, 95, 912-919.	1.3	8
41	Dual consent? Donors' and recipients' views about involvement in decision-making on the use of embryos created by gamete donation in research. BMC Medical Ethics, 2019, 20, 90.	1.0	8
42	Childhood socioeconomic conditions are associated with increased chronic low-grade inflammation over adolescence: findings from the EPITeen cohort study. Archives of Disease in Childhood, 2020, 105, 677-683.	1.0	8
43	Social Trajectory and Body Image Satisfaction in Childbearing Women. Maternal and Child Health Journal, 2015, 19, 1237-1244.	0.7	7
44	In vitro genotoxicity assessment of an oxidized dextrinâ€based hydrogel for biomedical applications. Journal of Applied Toxicology, 2019, 39, 639-649.	1.4	7
45	Health-related knowledge of primary prevention of cancer in Portugal. European Journal of Cancer Prevention, 2016, 25, 50-53.	0.6	6
46	Heterosexual couples' uses and meanings of ovarian stimulation: Relatedness, embodiment and emotions. Health (United Kingdom), 2011, 15, 620-632.	0.9	5
47	Family structure and use of prenatal care. Cadernos De Saude Publica, 2015, 31, 1298-1304.	0.4	5
48	Concerns with educating the public about donating and receiving gametes. European Journal of Contraception and Reproductive Health Care, 2019, 24, 420-421.	0.6	5
49	Payment to gamete donors: equality, gender equity, or solidarity?. Journal of Assisted Reproduction and Genetics, 2020, 37, 133-140.	1.2	5
50	A compreensão jurÃdica, médica e "leiga" do embrião em Portugal: um alinhamento com a biologia?. Interface: Communication, Health, Education, 2009, 13, 31-43.	0.4	5
51	A patient-centred approach to embryo donation for research. Israel Journal of Health Policy Research, 2016, 5, 44.	1.4	4
52	Portuguese forensic DNA database: political enthusiasm, public trust and probable issues in future practice. , 0, , 218-239.		3
53	Qualitative Research in Epidemiology. , 0, , .		3
54	Socioeconomic Position Early in Adolescence and Mode of Delivery Later in Life: Findings from a Portuguese Birth Cohort. PLoS ONE, 2015, 10, e0119517.	1.1	3

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55	Validation of the Short Assessment of Health Literacy in Portuguese-speaking Adults in Portugal. Gaceta Sanitaria, 2020, 34, 435-441.	0.6	3
56	Obesity-related knowledge and body mass index: a national survey in Portugal. Eating and Weight Disorders, 2020, 25, 1437-1446.	1.2	3
57	Prematurityâ€related knowledge among mothers and fathers of very preterm infants. Journal of Clinical Nursing, 2020, 29, 2886-2896.	1.4	3
58	Public Views About Involvement in Decision-Making on Health Data Sharing, Access, Use and Reuse: The Importance of Trust in Science and Other Institutions. Frontiers in Public Health, 2022, 10, .	1.3	3
59	Reflexoes sobre a investigacao em embrioes de origem humana: o debate nas organizacoes de etica portuguesas. Historia, Ciencias, Saude - Manguinhos, 2013, 20, 1137-1151.	0.1	2
60	User involvement in assisted reproductive technologies: England and Portugal. Journal of Health Organization and Management, 2015, 29, 582-594.	0.6	2
61	Self-reported psychosocial factors among in vitro fertilization patients interviewed alone or with the partner. Psychology, Health and Medicine, 2016, 21, 431-438.	1.3	2
62	Resposta à Carta ao Editor: "Prevalência de Literacia em Saúde Inadequada em Portugal Medida com o Newest Vital Sign― Acta Medica Portuguesa, 2018, 31, 183.	0.2	2
63	Priority of Access to Fertility Treatments Based on Sexual Orientation and Marital Status: the Views of Gamete Donors and Recipients. Sexuality Research and Social Policy, 2021, 18, 368-376.	1.4	2
64	Medical ethics when moving towards non-anonymous gamete donation: the views of donors and recipients. Journal of Medical Ethics, 2021, , medethics-2020-106947.	1.0	2
65	The Influence of Question Wording on Interpersonal Trust. Methodology, 2019, 15, 56-66.	0.5	2
66	Single motherhood and prenatal care utilisation. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2011, 96, Fa46-Fa46.	1.4	1
67	Criminal Genomic Pragmatism: Prisoners' Representations of DNA Technology and Biosecurity. Journal of Biomedicine and Biotechnology, 2012, 2012, 1-5.	3.0	1
68	Parental involvement in infant caregiving in a Neonatal Intensive Care Unit: an ethnographic approach. European Journal of Public Health, 2013, 23, .	0.1	1
69	Regulação da Licença de Parentalidade em Pais de Crianças Muito Pré-termo. Acta Medica Portuguesa, 2015, 28, 281.	0.2	1
70	A short form of the neonatal intensive care unit family needs inventory. Jornal De Pediatria (Versão) Tj ETQqO O	0 rgBT /O	verlock 10 Tf
71	Consentir na criopreservação de embriões: perceção de casais usuários de medicina da reprodução. Interface: Communication, Health, Education, 2017, 21, 435-448.	0.4	1

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73	Participation in data generation and information sharing: are patients and carers willing to engage?. European Journal of Public Health, 2020, 30, .	0.1	1
74	Concerns and coping mechanisms during the first national COVID-19 lockdown: an online prospective study in Portugal. Public Health, 2022, 207, 24-27.	1.4	1
75	Biogenética e género na construção da intencionalidade da paternidade: o teste de DNA nas investigações judiciais de paternidade. Revista Estudos Feministas, 2011, 19, 823-848.	0.0	0
76	Public participation in genetic databases. European Journal of Public Health, 2014, 24, .	0.1	0
77	Trajectories of health-related behaviours among parents of very preterm infants: from preconceptional to postnatal period. European Journal of Public Health, 2014, 24, .	0.1	0
78	PB.12â€A Short Form of the Neonatal Intensive Care Unit Family Needs Inventory. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2014, 99, A36.1-A36.	1.4	0
79	PB.02â€Needs and stress in parents of very preterm infants hospitalised in Neonatal Intensive Care Units: the effect of the data collection period. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2014, 99, A33.2-A33.	1.4	0
80	Citizenship and gamete donation: socioethical issues in policy, care and research. Annals of Medicine, 2024, 51, 31-31.	1.5	0
81	Reproductive trajectories and social-biological dimensions in parenthood: Portuguese version of the Meaning of Parenthood scale. Porto Biomedical Journal, 2020, 5, e078.	0.4	0
82	Donor-centred care: the facilitating and constraining factors experienced by gamete donors in a public bank. Human Fertility, 2023, 26, 115-126.	0.7	0
83	Issues on Methods of Identification in Court Ordered Paternity Testing ?The Case of Portugal. Journal of Forensics Research, 0, s2, .	0.1	0
84	Scientific issues on embryo research in Portugal. Configurações, 2011, , 57-72.	0.0	0
85	Quality of websites for gamete donation. Revista Electronica De Comunicacao, Informacao & Inovacao Em Saude: RECIIS, 2019, 13, .	0.2	0
86	Support for decision-making on sharing health data for research: are data counsellors relevant?. European Journal of Public Health, 2021, 31, .	0.1	0