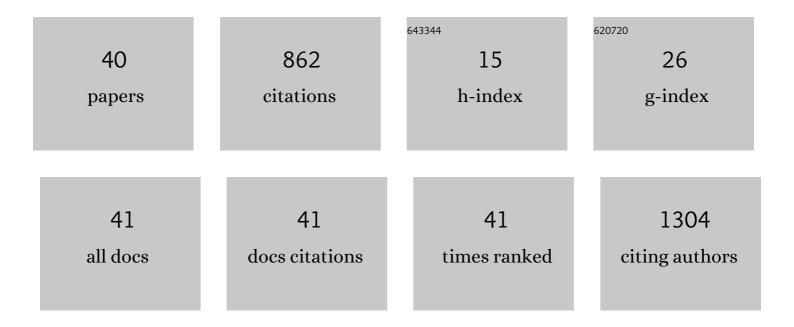
Ålvaro Mendes

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6590646/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Return of genomic results does not motivate intentÂtoÂparticipate in research for all: Perspectives across 22 countries. Genetics in Medicine, 2022, 24, 1120-1129.	1.1	8
2	The use of polygenic risk scores in pre-implantation genetic testing: an unproven, unethical practice. European Journal of Human Genetics, 2022, 30, 493-495.	1.4	38
3	From older to younger generations: Intergenerational transmission of health-related roles in families with Huntington's disease. Journal of Aging Studies, 2022, 61, 101027.	0.7	3
4	Opportunistic genomic screening. Recommendations of the European Society of Human Genetics. European Journal of Human Genetics, 2021, 29, 365-377.	1.4	76
5	Role of older generations in the family's adjustment to Huntington disease. Journal of Community Genetics, 2021, 12, 469-477.	0.5	3
6	Between responsibility and desire: Accounts of reproductive decisions from those at risk for or affected by lateâ€onset neurological diseases. Journal of Genetic Counseling, 2021, 30, 1480-1490.	0.9	4
7	Demonstrating trustworthiness when collecting and sharing genomic data: public views across 22 countries. Genome Medicine, 2021, 13, 92.	3.6	39
8	Impacto dos papéis dos mais velhos na promoção da saúde em famÃ l ias com paramiloidose. Psicologia, 2021, 35, 17-26.	0.1	0
9	ESHG PPPC Comments on postmortem use of genetic data for research purposes. European Journal of Human Genetics, 2020, 28, 144-146.	1.4	3
10	Global Public Perceptions of Genomic Data Sharing: What Shapes the Willingness to Donate DNA and Health Data?. American Journal of Human Genetics, 2020, 107, 743-752.	2.6	76
11	Web-based return of BRCA2 research results: one-year genetic counselling experience in Iceland. European Journal of Human Genetics, 2020, 28, 1656-1661.	1.4	12
12	Patients' perceptions of family engagement in health information practices: influences on the self-management of asthma. Journal of Communication in Healthcare, 2020, 13, 17-26.	0.8	1
13	Management of information within Portuguese families with Huntington disease: a transgenerational process for putting the puzzle together. European Journal of Human Genetics, 2020, 28, 1210-1217.	1.4	4
14	European recommendations integrating genetic testing into multidisciplinary management of sudden cardiac death. European Journal of Human Genetics, 2019, 27, 1763-1773.	1.4	78
15	Twenty Years of a Pre-Symptomatic Testing Protocol for Late-Onset Neurological Diseases in Portugal. Acta Medica Portuguesa, 2019, 32, 295.	0.2	14
16	Choosing not to know: accounts of non-engagement with pre-symptomatic testing for Machado-Joseph disease. European Journal of Human Genetics, 2019, 27, 353-359.	1.4	8
17	Recontacting patients in clinical genetics services: recommendations of the European Society of Human Genetics. European Journal of Human Genetics, 2019, 27, 169-182.	1.4	65
18	A diabetes não dói, não se sente, mas está cá dentro!. Psychologica, 2019, 62, 7-21.	0.2	0

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#	Article	IF	CITATIONS
19	Recontacting or not recontacting? A survey of current practices in clinical genetics centres in Europe. European Journal of Human Genetics, 2018, 26, 946-954.	1.4	33
20	Communication of Information about Genetic Risks: Putting Families at the Center. Family Process, 2018, 57, 836-846.	1.4	37
21	The challenges of the expanded availability of genomic information: an agenda-setting paper. Journal of Community Genetics, 2018, 9, 103-116.	0.5	45
22	Discredited legacy: Stigma and familial amyloid polyneuropathy in Northwestern Portugal. Social Science and Medicine, 2017, 182, 73-80.	1.8	12
23	From older to younger: intergenerational promotion of health behaviours in Portuguese families affected by familial amyloid polyneuropathy. European Journal of Human Genetics, 2017, 25, 687-693.	1.4	12
24	Direct-to-consumer genetic testing: where and how does genetic counseling fit?. Personalized Medicine, 2017, 14, 249-257.	0.8	44
25	"That Should Be Left to Doctors, That's What They are There For!â€â€"Exploring the Reflexivity and Trust of Young Adults When Seeking Health Information. Health Communication, 2017, 32, 1076-1081.	1.8	30
26	LIVING WITH DIABETES IS "CARRYING A CROSS― METAPHORS OF ELDERLY DIABETIC. Psicologia, Saúde & Doenças, 2017, 18, 747-759.	0.0	0
27	How communication of genetic information within the family is addressed in genetic counselling: a systematic review of research evidence. European Journal of Human Genetics, 2016, 24, 315-325.	1.4	66
28	Supporting families in genetic counselling services: a psychoeducational multifamily discussion group for atâ€risk colorectal cancer families. Journal of Family Therapy, 2015, 37, 343-360.	0.5	9
29	Genetic Counseling in Portugal: Education, Practice and a Developing Profession. Journal of Genetic Counseling, 2015, 24, 548-552.	0.9	16
30	Genetics Health Professionals' Views on Quality of Genetic Counseling Service Provision for Presymptomatic Testing in Lateâ€Onset Neurological Diseases in Portugal: Core Components, Specific Challenges and the Need for Assessment Tools. Journal of Genetic Counseling, 2015, 24, 616-625.	0.9	12
31	Material inheritances: an affective story in the history of elderly persons. International Journal of Ageing and Later Life, 2015, 9, 35-52.	0.4	7
32	From Constraints to Opportunities? Provision of Psychosocial Support in Portuguese Oncogenetic Counseling Services. Journal of Genetic Counseling, 2013, 22, 771-783.	0.9	7
33	A resiliência da saúde migrante: itinerários terapêuticos plurais e transnacionais. REMHU: Revista Interdisciplinar Da Mobilidade Humana, 2013, 21, 69-92.	0.0	3
34	Are family-oriented interventions in Portuguese genetics services a remote possibility? Professionals' views on a multifamily intervention for cancer susceptibility families. Journal of Community Genetics, 2012, 3, 311-318.	0.5	4
35	ProFamilies-dementia: A programme for elderly people with dementia and their families. Dementia, 2012, 11, 589-596.	1.0	7
36	Families' experience of oncogenetic counselling: accounts from a heterogeneous hereditary cancer risk population. Familial Cancer, 2012, 11, 291-306.	0.9	12

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#	Article	IF	CITATIONS
37	proFamilies: a psycho-educational multi-family group intervention for cancer patients and their families. European Journal of Cancer Care, 2011, 20, 337-344.	0.7	28
38	Experiencing genetic counselling for hereditary cancers: the client's perspective. European Journal of Cancer Care, 2011, 20, 204-211.	0.7	16
39	Family matters: examining a multi-family group intervention for women with BRCA mutations in the scope of genetic counselling. Journal of Community Genetics, 2010, 1, 161-168.	0.5	25
40	The Experience of Receiving and Transmitting a Genetic Disease. Journal of Constructivist Psychology, 0, , 1-16.	0.7	1