

Milena Paneque

List of Publications by Year in descending order

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49
papers

1,022
citations

516710

16
h-index

454955

30
g-index

53
all docs

53
docs citations

53
times ranked

1208
citing authors

#	ARTICLE	IF	CITATIONS
1	The need for recognition of core professional groups in genetics healthcare services in Europe. <i>European Journal of Human Genetics</i> , 2022, 30, 639-640.	2.8	2
2	Äœber die Notwendigkeit der Anerkennung von sog. Kernberufsgruppen innerhalb der genetischen Gesundheitsversorgung in Europa. <i>Medizinische Genetik</i> , 2022, 34, 81-83.	0.2	1
3	Erratum zu: Äœber die Notwendigkeit der Anerkennung von sog. Kernberufsgruppen innerhalb der genetischen Gesundheitsversorgung in Europa. <i>Medizinische Genetik</i> , 2022, 34, 189-191.	0.2	0
4	A new scale informed by the Reciprocal-Engagement Model for quality evaluation of genetic counselling by patients: Development and initial validation. <i>European Journal of Medical Genetics</i> , 2021, 64, 104375.	1.3	2
5	A experiÄªncia psicossocial de cuidadores informais de pessoas com paramiloidose: Um estudo qualitativo. <i>Psicologia</i> , 2021, 35, 45-62.	0.3	1
6	Developing a national certification pathway for genetic counselors in SwedenÄª”a short report. <i>Journal of Community Genetics</i> , 2020, 11, 113-117.	1.2	2
7	The Gen-Equip Project: evaluation and impact of genetics e-learning resources for primary care in six European languages. <i>Genetics in Medicine</i> , 2019, 21, 718-726.	2.4	17
8	Psychiatric genetic counseling: A mapping exercise. <i>American Journal of Medical Genetics Part B: Neuropsychiatric Genetics</i> , 2019, 180, 523-532.	1.7	20
9	Twenty Years of a Pre-Symptomatic Testing Protocol for Late-Onset Neurological Diseases in Portugal. <i>Acta Medica Portuguesa</i> , 2019, 32, 295.	0.4	14
10	Choosing not to know: accounts of non-engagement with pre-symptomatic testing for Machado-Joseph disease. <i>European Journal of Human Genetics</i> , 2019, 27, 353-359.	2.8	8
11	The Global State of the Genetic Counseling Profession. <i>European Journal of Human Genetics</i> , 2019, 27, 183-197.	2.8	215
12	How practical experiences, educational routes and multidisciplinary teams influence genetic counselors' clinical practice in Europe. <i>Clinical Genetics</i> , 2018, 93, 891-898.	2.0	11
13	Communication of Information about Genetic Risks: Putting Families at the Center. <i>Family Process</i> , 2018, 57, 836-846.	2.6	37
14	The recognition of the profession of Genetic Counsellors in Europe. <i>European Journal of Human Genetics</i> , 2018, 26, 1719-1720.	2.8	14
15	The Contribution of the ReciprocalÄªEngagement Model as a Theoretical Framework of a Portuguese Scale for Quality Assessment of Genetic Counseling. <i>Journal of Genetic Counseling</i> , 2018, 27, 1005-1007.	1.6	2
16	Complementarity between medical geneticists and genetic counsellors: its added value in genetic services in Europe. <i>European Journal of Human Genetics</i> , 2017, 25, 918-923.	2.8	29
17	The perceived impact of the European registration system for genetic counsellors and nurses. <i>European Journal of Human Genetics</i> , 2017, 25, 1075-1077.	2.8	8
18	Implementing genetic education in primary care: the Gen-Equip programme. <i>Journal of Community Genetics</i> , 2017, 8, 147-150.	1.2	16

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19	A systematic review of interventions to provide genetics education for primary care. BMC Family Practice, 2016, 17, 89.	2.9	52
20	Development of a registration system for genetic counsellors and nurses in health-care services in Europe. European Journal of Human Genetics, 2016, 24, 312-314.	2.8	27
21	Components of genetic counsellor education: A systematic review of the peer-reviewed literature. Journal of Community Genetics, 2016, 7, 107-118.	1.2	12
22	The need to develop an evidence base for genetic counselling in Europe. European Journal of Human Genetics, 2016, 24, 504-505.	2.8	17
23	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.	2.8	66
24	SCA2 predictive testing in Cuba: challenging concepts and protocol evolution. Journal of Community Genetics, 2015, 6, 265-273.	1.2	9
25	Quality issues concerning genetic counselling for presymptomatic testing: a European Delphi study. European Journal of Human Genetics, 2015, 23, 1468-1472.	2.8	9
26	Genetic Counseling in Portugal: Education, Practice and a Developing Profession. Journal of Genetic Counseling, 2015, 24, 548-552.	1.6	16
27	Genetics Health Professionals' Views on Quality of Genetic Counseling Service Provision for Presymptomatic Testing in Late-Onset Neurological Diseases in Portugal: Core Components, Specific Challenges and the Need for Assessment Tools. Journal of Genetic Counseling, 2015, 24, 616-625.	1.6	12
28	[NO TITLE AVAILABLE]. Genetics and Molecular Biology, 2014, 37, 263-270.	1.3	16
29	Large Normal and Intermediate Alleles in the Context of SCA2 Prenatal Diagnosis. Journal of Genetic Counseling, 2014, 23, 89-96.	1.6	7
30	Insufficient Referral for Genetic Counseling in the Management of Hereditary Haemochromatosis in Portugal: A Study of Perceptions of Health Professionals Requesting <i>HFE</i> Genotyping. Journal of Genetic Counseling, 2014, 23, 770-777.	1.6	4
31	The importance of the general practitioner as an information source for patients with hereditary haemochromatosis. Patient Education and Counseling, 2014, 96, 86-92.	2.2	11
32	What Counts as Effective Genetic Counselling for Presymptomatic Testing in Late-Onset Disorders? A Study of the Consultant's Perspective. Journal of Genetic Counseling, 2013, 22, 437-447.	1.6	28
33	Couples at risk for spinocerebellar ataxia type 2: the Cuban prenatal diagnosis experience. Journal of Community Genetics, 2013, 4, 451-460.	1.2	5
34	From Constraints to Opportunities? Provision of Psychosocial Support in Portuguese Oncogenetic Counseling Services. Journal of Genetic Counseling, 2013, 22, 771-783.	1.6	7
35	Non-syndromic Sensorineural Prelingual Deafness: The Importance of Genetic Counseling in Demystifying Parents' Beliefs About the Cause of Their Children's Deafness. Journal of Genetic Counseling, 2013, 22, 448-454.	1.6	6
36	The Cuban program for predictive testing of <i>SCA2</i> : 11 years and 768 individuals to learn from. Clinical Genetics, 2013, 83, 518-524.	2.0	19

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37	Predictive testing for two neurodegenerative disorders (FAP and HD): A psychological point of view. <i>Open Journal of Genetics</i> , 2013, 03, 270-279.	0.1	6
38	Cuban Adolescents Requesting Presymptomatic Testing for Spinocerebellar Ataxia Type 2. <i>ISRN Genetics</i> , 2013, 2013, 1-5.	0.2	1
39	Are family-oriented interventions in Portuguese genetics services a remote possibility? Professionals' views on a multifamily intervention for cancer susceptibility families. <i>Journal of Community Genetics</i> , 2012, 3, 311-318.	1.2	4
40	Quality Assessment of Genetic Counseling Process in the Context of Presymptomatic Testing for Late-Onset Disorders: A Thematic Analysis of Three Review Articles. <i>Genetic Testing and Molecular Biomarkers</i> , 2012, 16, 36-45.	0.7	13
41	The wide variation of definitions of genetic testing in international recommendations, guidelines and reports. <i>Journal of Community Genetics</i> , 2012, 3, 113-124.	1.2	16
42	Ethical Dilemmas in Genetic Testing: Examples from the Cuban Program for Predictive Diagnosis of Hereditary Ataxias. <i>Journal of Genetic Counseling</i> , 2011, 20, 241-248.	1.6	9
43	Estimation of the age at onset in spinocerebellar ataxia type 2 Cuban patients by survival analysis. <i>Clinical Genetics</i> , 2010, 78, 169-174.	2.0	44
44	Role of the Disease in the Psychological Impact of Pre-Symptomatic Testing for SCA2 and FAP ATTRV30M: Experience with the Disease, Kinship and Gender of the Transmitting Parent. <i>Journal of Genetic Counseling</i> , 2009, 18, 483-493.	1.6	24
45	Molecular epidemiology of spinocerebellar ataxias in Cuba: Insights into SCA2 founder effect in Holguin. <i>Neuroscience Letters</i> , 2009, 454, 157-160.	2.1	101
46	Age-dependent risks in genetic counseling for spinocerebellar ataxia type 2. <i>Clinical Genetics</i> , 2008, 74, 571-573.	2.0	3
47	Psychological Aspects of Presymptomatic Diagnosis of Spinocerebellar Ataxia Type 2 in Cuba. <i>Public Health Genomics</i> , 2007, 10, 132-139.	1.0	23
48	Psychological Follow-up of Presymptomatic Genetic Testing for Spinocerebellar Ataxia Type 2 (SCA2) in Cuba. <i>Journal of Genetic Counseling</i> , 2007, 16, 469-479.	1.6	13
49	Psychological aspects of pre-symptomatic testing for Machado-Joseph disease and familial amyloid polyneuropathy type I. <i>Clinical Genetics</i> , 2006, 69, 297-305.	2.0	33