Paola Forabosco

List of Publications by Year
in descending order

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41	2,523	218381	288905
papers	citations	26 h-index	40 g-index

43	43	43	4854
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Identification of Candidate Parkinson Disease Genes by Integrating Genome-Wide Association Study, Expression, and Epigenetic Data Sets. JAMA Neurology, 2021, 78, 464.	4.5	95
2	Dystonia genes functionally converge in specific neurons and share neurobiology with psychiatric disorders. Brain, 2020, 143, 2771-2787.	3.7	50
3	A Sardinian founder mutation in glycoprotein Ib platelet subunit beta (GP1BB) that impacts thrombocytopenia. British Journal of Haematology, 2020, 191, e124-e128.	1.2	2
4	An additional k-means clustering step improves the biological features of WGCNA gene co-expression networks. BMC Systems Biology, 2017, 11, 47.	3.0	253
5	Gene co-expression networks shed light into diseases of brain iron accumulation. Neurobiology of Disease, 2016, 87, 59-68.	2.1	24
6	Frontotemporal dementia: insights into the biological underpinnings of disease through gene co-expression network analysis. Molecular Neurodegeneration, 2016, 11, 21.	4.4	39
7	A Missense Mutation in KCTD17 Causes Autosomal Dominant Myoclonus-Dystonia. American Journal of Human Genetics, 2015, 96, 938-947.	2.6	109
8	Insights From Cerebellar Transcriptomic Analysis Into the Pathogenesis of Ataxia. JAMA Neurology, 2014, 71, 831.	4.5	60
9	Rare coding variants in the phospholipase D3 gene confer risk for Alzheimerâ€™s disease. Nature, 2014, 505, 550-554.	13.7	425
10	Genetic evidence for a pathogenic role for the vitamin D3 metabolizing enzyme CYP24A1 in multiple sclerosis. Multiple Sclerosis and Related Disorders, 2014, 3, 211-219.	0.9	44
11	Insights into TREM2 biology by network analysis of human brain gene expression data. Neurobiology of Aging, 2013, 34, 2699-2714.	1.5	145
12	Predicting the Risk of Rheumatoid Arthritis and Its Age of Onset through Modelling Genetic Risk Variants with Smoking. PLoS Genetics, 2013, 9, e1003808.	1.5	55
13	Association of the IL-10 Gene Family Locus on Chromosome 1 with Juvenile Idiopathic Arthritis (JIA). PLoS ONE, 2012, 7, e47673.	1.1	26
14	Application of a New Method for GWAS in a Related Case/Control Sample with Known Pedigree Structure: Identification of New Loci for Nephrolithiasis. PLoS Genetics, 2011, 7, e1001281.	1.5	10
15	Meta-analysis of 20 genome-wide linkage studies evidenced new regions linked to asthma and atopy. European Journal of Human Genetics, 2010, 18, 700-706.	1.4	54
16	Meta-Analysis of Genome-Wide Linkage Studies in Celiac Disease. Human Heredity, 2009, 68, 223-230.	0.4	10
17	Meta-analysis of genome-wide linkage studies across autoimmune diseases. European Journal of Human Genetics, 2009, 17, 236-243.	1.4	39
18	Meta-analysis of linkage studies for Alzheimer's diseaseâ€”A web resource. Neurobiology of Aging, 2009, 30, 1037-1047.	1.5	58

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19	Metaâ€analysis of genomeâ€wide linkage scans of attention deficit hyperactivity disorder. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2008, 147B, 1392-1398.	1.1	160
20	Data Acquisition for Meta-Analysis of Genome-Wide Linkage Studies Using the Genome Search Meta-Analysis Method. Human Heredity, 2007, 64, 74-81.	0.4	11
21	Serotonin transporter gene (5-Htt): Association analysis with temporal lobe epilepsy. Neuroscience Letters, 2007, 421, 52-56.	1.0	32
22	Further evidence of genetic heterogeneity in families with autosomal dominant nocturnal frontal lobe epilepsy. Epilepsy Research, 2007, 74, 70-73.	0.8	29
23	Meta-analysis of genome-wide linkage studies for multiple sclerosis, using an extended GSMA method. European Journal of Human Genetics, 2007, 15, 703-710.	1.4	16
24	Genetic Heterogeneity in Italian Families with IgA Nephropathy: Suggestive Linkage for Two Novel IgA Nephropathy Loci. American Journal of Human Genetics, 2006, 79, 1130-1134.	2.6	111
25	Statistical tools for linkage analysis and genetic association studies. Expert Review of Molecular Diagnostics, 2005, 5, 781-796.	1.5	11
26	A Genomewide Search Using an Original Pairwise Sampling Approach for Large Genealogies Identifies a New Locus for Total and Low-Density Lipoprotein Cholesterol in Two Genetically Differentiated Isolates of Sardinia. American Journal of Human Genetics, 2004, 75, 1015-1031.	2.6	48
27	Genetic heterogeneity in inherited spastic paraplegia associated with epilepsy. , 2003, 117A, 116-121.		7
28	Identification of a Novel Gene and a Common Variant Associated with Uric Acid Nephrolithiasis in a Sardinian Genetic Isolate. American Journal of Human Genetics, 2003, 72, 1479-1491.	2.6	65
29	A New Essential Hypertension Susceptibility Locus on Chromosome 2p24-p25, Detected by Genomewide Search. American Journal of Human Genetics, 2002, 71, 893-905.	2.6	74
30	Not all isolates are equal: linkage disequilibrium analysis on Xq13.3 reveals different patterns in Sardinian sub-populations. Human Genetics, 2002, 111, 9-15.	1.8	39
31	Optic disc drusen, angioid streaks, and mottled fundus in various combinations in a Sicilian family. Graefe's Archive for Clinical and Experimental Ophthalmology, 2002, 240, 771-776.	1.0	6
32	Identification of a New Candidate Locus for Uric Acid Nephrolithiasis. American Journal of Human Genetics, 2001, 68, 1119-1129.	2.6	36
33	Archival, demographic and genetic studies define a Sardinian sub-isolate as a suitable model for mapping complex traits. Human Genetics, 2001, 109, 198-209.	1.8	63
34	Combined segregation and linkage analysis of inflammatory bowel disease in the IBD1 region using severity to characterise Crohn's disease and ulcerative colitis. European Journal of Human Genetics, 2000, 8, 846-852.	1.4	18
35	A refined physical and transcriptional map of the SPG9 locus on 10q23.3â€“q24.2. European Journal of Human Genetics, 2000, 8, 777-782.	1.4	17
36	Linkage study of early-onset obesity to leptin receptor gene in Italian children. Nutrition Research, 2000, 20, 1059-1063.	1.3	2

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37	Genetic analysis in Italian families with inflammatory bowel disease supports linkage to the IBD1 locus â€“ A GISC study. European Journal of Human Genetics, 1999, 7, 567-573.	1.4	81
38	Exclusion of the Sonic Hedgehog gene as responsible for Currarino syndrome and anorectal malformations with sacral hypodevelopment. Human Genetics, 1999, 104, 108-110.	1.8	23
39	Genetic Mapping to 10q23.3-q24.2, in a Large Italian Pedigree, of a New Syndrome Showing Bilateral Cataracts, Gastroesophageal Reflux, and Spastic Paraparesis with Amyotrophy. American Journal of Human Genetics, 1999, 64, 586-593.	2.6	108
40	Inheritance of Astigmatism: Evidence for a Major Autosomal Dominant Locus. American Journal of Human Genetics, 1998, 63, 825-830.	2.6	54
41	Inheritance of cleft palate in Italy. Evidence for a major autosomal recessive locus. Human Genetics, 1997, 100, 204-209.	1.8	13