Emilia Vitale

List of Publications by Year
in descending order

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27 papers	1,411 citations	858243 12 h-index	591227 27 g-index
27 all docs	27 docs citations	27 times ranked	1383 citing authors

#	Article	IF	Citations
1	CD33 rs2455069 SNP: Correlation with Alzheimerâ€™s Disease and Hypothesis of Functional Role. International Journal of Molecular Sciences, 2022, 23, 3629.	1.8	10
2	CD33 and SIGLECL1 Immunoglobulin Superfamily Involved in Dementia. Journal of Neuropathology and Experimental Neurology, 2020, 79, 891-901.	0.9	6
3	LINC00473 as an Immediate Early Gene under the Control of the EGR1 Transcription Factor. Non-coding RNA, 2020, 6, 46.	1.3	2
4	C9orf72, age at onset, and ancestry help discriminate behavioral from language variants in FTLD cohorts. Neurology, 2020, 95, e3288-e3302.	1.5	7
5	Clinical and Molecular Characterization of a Novel Progranulin Deletion Associated with Different Phenotypes. Journal of Alzheimer's Disease, 2020, 76, 341-347.	1.2	5
6	Identification, Characterization, and Regulatory Mechanisms of a Novel EGR1 Splicing Isoform. International Journal of Molecular Sciences, 2019, 20, 1548.	1.8	9
7	Circulating levels of IL-1 family cytokines and receptors in Alzheimerâ€™s disease: new markers of disease progression?. Journal of Neuroinflammation, 2018, 15, 342.	3.1	91
8	GRN deletion in familial frontotemporal dementia showing association with clinical variability in 3 familial cases. Neurobiology of Aging, 2017, 53, 193.e9-193.e16.	1.5	8
9	Metabolic response of SH-SY5Y cells to gold nanoparticles by NMR-based metabolomics analyses. Biomedical Physics and Engineering Express, 2016, 2, 045003.	0.6	4
10	Imidazole-stabilized gold nanoparticles induce neuronal apoptosis: Anin vitroandin vivostudy. Journal of Biomedical Materials Research - Part A, 2015, 103, 1436-1446.	2.1	13
11	Molecular responses of cells to 2-mercapto-1-methylimidazole gold nanoparticles (AuNPs)-mmi: investigations of histone methylation changes. Journal of Nanoparticle Research, 2014, 16, 1.	0.8	6
12	Donor spliceâ€site mutation in CUL4B is likely cause of Xâ€linked intellectual disability. American Journal of Medical Genetics, Part A, 2014, 164, 2294-2299.	0.7	13
13	Cortical Metabolic Deficits in a Rat Model of Cholinergic Basal Forebrain Degeneration. Neurochemical Research, 2013, 38, 2114-2123.	1.6	8
14	Structural Chromosomal Variations in Neurological Diseases. Neurologist, 2009, 15, 245-253.	0.4	10
15	Variants of ST8SIA1 Are Associated with Risk of Developing Multiple Sclerosis. PLoS ONE, 2008, 3, e2653.	1.1	10
16	Highly variable penetrance in subjects affected with cavernous cerebral angiomas (CCM) carrying novel CCM1 and CCM2 mutations. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2007, 144B, 691-695.	1.1	32
17	Arvanil and anandamide up-regulate CD36 expression in human peripheral blood mononuclear cells. Immunology Letters, 2007, 109, 145-154.	1.1	7
18	Second family with hearing impairment linked to 19q13 and refined DFNA4 localisation. European Journal of Human Genetics, 2002, 10, 95-99.	1.4	12

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19	Linkage analysis conditional on HLA status in a large North American pedigree supports the presence of a multiple sclerosis susceptibility locus on chromosome 12p12. Human Molecular Genetics, 2002, 11, 295-300.	1.4	33
20	Novel X-linked mental retardation syndrome with short stature maps to Xq24. American Journal of Medical Genetics Part A, 2001, 103, 1-8.	2.4	14
21	A second Middle Eastern kindred with autosomal recessive non-syndromic hearing loss segregates DFNB9. European Journal of Human Genetics, 1998, 6, 341-344.	1.4	14
22	Genomic Cloning and Characterization of the Human Thrombin Receptor Gene. Journal of Biological Chemistry, 1996, 271, 9307-9312.	1.6	45
23	Genetic Linkage of the Marfan Syndrome, Ectopia Lentis, and Congenital Contractural Arachnodactyly to the Fibrillin Genes on Chromosomes 15 and 5. New England Journal of Medicine, 1992, 326, 905-909.	13.9	257
24	Repating developmental expression of G-Hox 7, a novel homeobox-containing gene in the chicken. Developmental Biology, 1991, 148, 375-388.	0.9	98
25	Linkage of Marfan syndrome and a phenotypically related disorder to two different fibrillin genes. Nature, 1991, 352, 330-334.	13.7	676
26	Assignment of human aldolase C gene to chromosome 17, region cen?q21.1. Human Genetics, 1989, 82, 279-282.	1.8	12
27	Evidences that hemoglobin switch in the chick embryo depends on erythroid cell line substitution. Cell Differentiation, 1987, 20, 55-63.	1.3	9