

Patrizia Cavadini

List of Publications by Year in descending order

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Version: 2024-02-01

32
papers

3,381
citations

279798

23
h-index

454955

30
g-index

32
all docs

32
docs citations

32
times ranked

3968
citing authors

#	ARTICLE	IF	CITATIONS
1	Ferritins: A family of molecules for iron storage, antioxidation and more. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2009, 1790, 589-599.	2.4	718
2	Mitochondrial processing peptidases. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2002, 1592, 63-77.	4.1	353
3	Identification of a common mutation in the carnitine palmitoyltransferase II gene in familial recurrent myoglobinuria patients. <i>Nature Genetics</i> , 1993, 4, 314-320.	21.4	214
4	Assembly and iron-binding properties of human frataxin, the protein deficient in Friedreich ataxia. <i>Human Molecular Genetics</i> , 2002, 11, 217-227.	2.9	180
5	Emergence of a new lagovirus related to rabbit haemorrhagic disease virus. <i>Veterinary Research</i> , 2013, 44, 81.	3.0	180
6	Altered leukocyte response to CXCL12 in patients with warts hypogammaglobulinemia, infections, myelokathexis (WHIM) syndrome. <i>Blood</i> , 2004, 104, 444-452.	1.4	172
7	RNA silencing of the mitochondrial ABCB7 transporter in HeLa cells causes an iron-deficient phenotype with mitochondrial iron overload. <i>Blood</i> , 2007, 109, 3552-3559.	1.4	156
8	Molecular characterization of inherited carnitine palmitoyltransferase II deficiency.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1992, 89, 8429-8433.	7.1	151
9	Proposal for a unified classification system and nomenclature of lagoviruses. <i>Journal of General Virology</i> , 2017, 98, 1658-1666.	2.9	148
10	AIRE deficiency in thymus of 2 patients with Omenn syndrome. <i>Journal of Clinical Investigation</i> , 2005, 115, 728-732.	8.2	146
11	Innate immunity defects in Hermansky-Pudlak type 2 syndrome. <i>Blood</i> , 2006, 107, 4857-4864.	1.4	136
12	The new French 2010 Rabbit Hemorrhagic Disease Virus causes an RHD-like disease in the Sardinian Cape hare (<i>Lepus capensis mediterraneus</i>). <i>Veterinary Research</i> , 2013, 44, 96.	3.0	113
13	Two-step Processing of Human Frataxin by Mitochondrial Processing Peptidase. <i>Journal of Biological Chemistry</i> , 2000, 275, 41469-41475.	3.4	101
14	Yeast and Human Frataxin Are Processed to Mature Form in Two Sequential Steps by the Mitochondrial Processing Peptidase. <i>Journal of Biological Chemistry</i> , 1999, 274, 22763-22769.	3.4	99
15	Carnitine palmitoyltransferase II deficiency: structure of the gene and characterization of two novel disease-causing mutations. <i>Human Molecular Genetics</i> , 1995, 4, 19-29.	2.9	89
16	Assignment of the Human Carnitine Palmitoyltransferase II Gene (CPT1) to Chromosome 1p32. <i>Genomics</i> , 1994, 24, 195-197.	2.9	65
17	TwoCPT2 mutations in three Japanese patients with carnitine palmitoyltransferase II deficiency: Functional analysis and association with polymorphic haplotypes and two clinical phenotypes. <i>Human Mutation</i> , 1998, 11, 377-386.	2.5	56
18	The effects of frataxin silencing in HeLa cells are rescued by the expression of human mitochondrial ferritin. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2008, 1782, 90-98.	3.8	56

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19	Clinical and molecular heterogeneity in very long-chain acyl-coenzyme a dehydrogenase deficiency. <i>Pediatric Neurology</i> , 2000, 22, 98-105.	2.1	44
20	Pantothenate kinase-2 (Pank2) silencing causes cell growth reduction, cell-specific ferroportin upregulation and iron deregulation. <i>Neurobiology of Disease</i> , 2010, 39, 204-210.	4.4	42
21	Emergence of Pathogenicity in Lagoviruses: Evolution from Pre-existing Nonpathogenic Strains or through a Species Jump?. <i>PLoS Pathogens</i> , 2015, 11, e1005087.	4.7	31
22	Recombinant human hepcidin expressed in <i>Escherichia coli</i> isolates as an iron containing protein. <i>Blood Cells, Molecules, and Diseases</i> , 2005, 35, 177-181.	1.4	29
23	Field and experimental data indicate that the eastern cottontail (<i>Sylvilagus floridanus</i>) is susceptible to infection with European brown hare syndrome (EBHS) virus and not with rabbit haemorrhagic disease (RHD) virus. <i>Veterinary Research</i> , 2015, 46, 13.	3.0	27
24	Molecular characterization of SG33 and Borghi vaccines used against myxomatosis. <i>Vaccine</i> , 2010, 28, 5414-5420.	3.8	18
25	Protein import and processing reconstituted with isolated rat liver mitochondria and recombinant mitochondrial processing peptidase. <i>Methods</i> , 2002, 26, 298-306.	3.8	16
26	Two novel sequence polymorphisms of the human carnitine palmitoyltransferase II (CPT1) gene. <i>Human Molecular Genetics</i> , 1993, 2, 334-334.	2.9	13
27	Red foxes (<i>Vulpes vulpes</i>) feeding brown hares (<i>Lepus europaeus</i>) infected by European brown hare syndrome virus (EBHSV) might be involved in the spread of the virus. <i>European Journal of Wildlife Research</i> , 2016, 62, 761-765.	1.4	10
28	Widespread occurrence of the non-pathogenic hare calicivirus (HaCV Lagovirus GII.2) in captive and free-living wild hares in Europe. <i>Transboundary and Emerging Diseases</i> , 2021, 68, 509-518.	3.0	8
29	Potential role of wolf (<i>Canis lupus</i>) as passive carrier of European brown hare syndrome virus (EBHSV). <i>Research in Veterinary Science</i> , 2018, 117, 81-84.	1.9	4
30	Evaluation of Three Rapid Diagnostic Tests Used in Bovine Spongiform Encephalopathy Monitoring in Italy. <i>Journal of Veterinary Diagnostic Investigation</i> , 2009, 21, 830-836.	1.1	3
31	Rabbit Hemorrhagic Disease Virus and European Brown Hare Syndrome Virus (Caliciviridae). , 2021, , 724-729.		3
32	Viral haemorrhagic disease: RHDV type 2 ten years later. <i>World Rabbit Science</i> , 2022, 30, 1-11.	0.6	0