

Mãrcia Em Oliveira

List of Publications by Year in descending order

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Version: 2024-02-01

19
papers

790
citations

623734

14
h-index

940533

16
g-index

19
all docs

19
docs citations

19
times ranked

607
citing authors

#	ARTICLE	IF	CITATIONS
1	Characterization of Peroxisomal Pex5p from Rat Liver. <i>Journal of Biological Chemistry</i> , 2000, 275, 32444-32451.	3.4	106
2	Characterization of the Peroxisomal Cycling Receptor, Pex5p, Using a Cell-free in Vitro Import System. <i>Journal of Biological Chemistry</i> , 2003, 278, 226-232.	3.4	92
3	Characterization of the Mammalian Peroxisomal Import Machinery. <i>Journal of Biological Chemistry</i> , 2001, 276, 29935-29942.	3.4	88
4	The Energetics of Pex5p-mediated Peroxisomal Protein Import. <i>Journal of Biological Chemistry</i> , 2003, 278, 39483-39488.	3.4	81
5	Insertion of Pex5p into the Peroxisomal Membrane Is Cargo Protein-dependent. <i>Journal of Biological Chemistry</i> , 2003, 278, 4389-4392.	3.4	79
6	<i>LAMA2</i> gene mutation update: Toward a more comprehensive picture of the laminin- α 2 variome and its related phenotypes. <i>Human Mutation</i> , 2018, 39, 1314-1337.	2.5	71
7	The Import Competence of a Peroxisomal Membrane Protein Is Determined by Pex19p before the Docking Step. <i>Journal of Biological Chemistry</i> , 2006, 281, 34492-34502.	3.4	53
8	Mammalian Pex14p: membrane topology and characterisation of the Pex14p-Pex14p interaction. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2002, 1567, 13-22.	2.6	45
9	Atypical phenotype in two patients with LAMA2 mutations. <i>Neuromuscular Disorders</i> , 2014, 24, 419-424.	0.6	30
10	Expanding the MTM1 mutational spectrum: novel variants including the first multi-exonic duplication and development of a locus-specific database. <i>European Journal of Human Genetics</i> , 2013, 21, 540-549.	2.8	29
11	Identification of a 24 kDa intrinsic membrane protein from mammalian peroxisomes. <i>Biochimica Et Biophysica Acta Gene Regulatory Mechanisms</i> , 1999, 1445, 337-341.	2.4	26
12	Protein Translocation Across the Peroxisomal Membrane. <i>Cell Biochemistry and Biophysics</i> , 2004, 41, 451-468.	1.8	25
13	Alkaline Density Gradient Floatation of Membranes: Polypeptide Composition of the Mammalian Peroxisomal Membrane. <i>Analytical Biochemistry</i> , 1999, 274, 270-277.	2.4	20
14	New massive parallel sequencing approach improves the genetic characterization of congenital myopathies. <i>Journal of Human Genetics</i> , 2016, 61, 497-505.	2.3	15
15	Reviewing Large LAMA2 Deletions and Duplications in Congenital Muscular Dystrophy Patients. <i>Journal of Neuromuscular Diseases</i> , 2014, 1, 169-179.	2.6	14
16	Characterization of the Peroxisomal Cycling Receptor Pex5p Import Pathway. <i>Advances in Experimental Medicine and Biology</i> , 2003, 544, 219-220.	1.6	13
17	Integrating Whole-Genome Sequencing in Clinical Genetics: A Novel Disruptive Structural Rearrangement Identified in the Dystrophin Gene (DMD). <i>International Journal of Molecular Sciences</i> , 2022, 23, 59.	4.1	3
18	Short-term complications after renal transplantation in AFibE526V (p.Glu545Val) amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2019, 26, 162-163.	3.0	0

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19	X-Linked Myotubular Myopathy: A Novel Mutation Expanding the Genotypic Spectrum of a Phenotypically Heterogeneous Myopathy. <i>Journal of Pediatric Genetics</i> , 0, , .	0.7	0