

JesÃ³s Maria MartÃ­n-Campos

List of Publications by Year in descending order

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33
papers

1,067
citations

471477

17
h-index

395678

33
g-index

35
all docs

35
docs citations

35
times ranked

1572
citing authors

#	ARTICLE	IF	CITATIONS
1	Role of apoA-II in lipid metabolism and atherosclerosis: advances in the study of an enigmatic protein. <i>Journal of Lipid Research</i> , 2001, 42, 1727-1739.	4.2	118
2	Sitosterolemia: Diagnosis, Investigation, and Management. <i>Current Atherosclerosis Reports</i> , 2014, 16, 424.	4.8	92
3	A Genomewide Exploration Suggests a New Candidate Gene at Chromosome 11q23 as the Major Determinant of Plasma Homocysteine Levels: Results from the GAIT Project. <i>American Journal of Human Genetics</i> , 2005, 76, 925-933.	6.2	90
4	Molecular profiling related to poor prognosis in thyroid carcinoma. Combining gene expression data and biological information. <i>Oncogene</i> , 2008, 27, 1554-1561.	5.9	86
5	Identification of a novel mutation in the ANGPTL3 gene in two families diagnosed of familial hypobetalipoproteinemia without APOB mutation. <i>Clinica Chimica Acta</i> , 2012, 413, 552-555.	1.1	63
6	Overexpression of Human Apolipoprotein A-II in Transgenic Mice Does Not Impair Macrophage-Specific Reverse Cholesterol Transport In Vivo. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2005, 25, e128-32.	2.4	61
7	Chylomicrons: Advances in biology, pathology, laboratory testing, and therapeutics. <i>Clinica Chimica Acta</i> , 2016, 455, 134-148.	1.1	59
8	The Role of Nitric Oxide in the Local Antiallodynic and Antihyperalgesic Effects and Expression of μ -Opioid and Cannabinoid-2 Receptors during Neuropathic Pain in Mice. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2010, 334, 887-896.	2.5	58
9	Apolipoprotein A-II, genetic variation on chromosome 1q21-q24, and disease susceptibility. <i>Current Opinion in Lipidology</i> , 2004, 15, 247-253.	2.7	45
10	Peripheral Effects of Morphine and Expression of μ -Opioid Receptors in the Dorsal Root Ganglia during Neuropathic Pain: Nitric Oxide Signaling. <i>Molecular Pain</i> , 2011, 7, 1744-8069-7-25.	2.1	44
11	The Spinal Cord Expression of Neuronal and Inducible Nitric Oxide Synthases and Their Contribution in the Maintenance of Neuropathic Pain in Mice. <i>PLoS ONE</i> , 2010, 5, e14321.	2.5	40
12	Structural and functional analysis of APOA5 mutations identified in patients with severe hypertriglyceridemia. <i>Journal of Lipid Research</i> , 2013, 54, 649-661.	4.2	34
13	Molecular analysis of chylomicronemia in a clinical laboratory setting: Diagnosis of 13 cases of lipoprotein lipase deficiency. <i>Clinica Chimica Acta</i> , 2014, 429, 61-68.	1.1	34
14	ApoA-IMALLORCA impairs LCAT activation and induces dominant familial hypoalphalipoproteinemia. <i>Journal of Lipid Research</i> , 2002, 43, 115-123.	4.2	24
15	The Antinociceptive Effects of JWH-015 in Chronic Inflammatory Pain Are Produced by Nitric Oxide-cGMP-PKG-KATP Pathway Activation Mediated by Opioids. <i>PLoS ONE</i> , 2011, 6, e26688.	2.5	24
16	A rare STAP1 mutation incompletely associated with familial hypercholesterolemia. <i>Clinica Chimica Acta</i> , 2018, 487, 270-274.	1.1	19
17	Methionine-induced hyperhomocysteinemia impairs the antioxidant ability of high-density lipoproteins without reducing in vivo macrophage-specific reverse cholesterol transport. <i>Molecular Nutrition and Food Research</i> , 2013, 57, 1814-1824.	3.3	18
18	ApoA-I(MALLORCA) impairs LCAT activation and induces dominant familial hypoalphalipoproteinemia. <i>Journal of Lipid Research</i> , 2002, 43, 115-23.	4.2	18

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19	Molecular analysis of APOB, SAR1B, ANGPTL3, and MTTP in patients with primary hypocholesterolemia in a clinical laboratory setting: Evidence supporting polygenicity in mutation-negative patients. <i>Atherosclerosis</i> , 2019, 283, 52-60.	0.8	15
20	Apolipoprotein A5 S19W May Play a Role in Dysbetalipoproteinemia in Patients with the Apo E2/E2 Genotype. <i>Clinical Chemistry</i> , 2006, 52, 1974-1975.	3.2	14
21	Autosomal dominant hypercholesterolemia in Catalonia: Correspondence between clinical-biochemical and genetic diagnostics in 967 patients studied in a multicenter clinical setting. <i>Journal of Clinical Lipidology</i> , 2018, 12, 1452-1462.	1.5	14
22	Administration of CORM-2 inhibits diabetic neuropathy but does not reduce dyslipidemia in diabetic mice. <i>PLoS ONE</i> , 2018, 13, e0204841.	2.5	12
23	A novel germline mutation in exon 5 of the multiple endocrine neoplasia type 1 gene. <i>Journal of Molecular Medicine</i> , 1998, 76, 837-839.	3.9	11
24	Biological Age Acceleration Is Lower in Women With Ischemic Stroke Compared to Men. <i>Stroke</i> , 2022, 53, 2320-2330.	2.0	11
25	ICA1L Is Associated with Small Vessel Disease: A Proteome-Wide Association Study in Small Vessel Stroke and Intracerebral Haemorrhage. <i>International Journal of Molecular Sciences</i> , 2022, 23, 3161.	4.1	11
26	Molecular Pathology of Multiple Endocrine Neoplasia Type I. <i>Diagnostic Molecular Pathology</i> , 1999, 8, 195-204.	2.1	10
27	Phenytoin treatment reduces atherosclerosis in mice through mechanisms independent of plasma HDL-cholesterol concentration. <i>Atherosclerosis</i> , 2004, 174, 275-285.	0.8	9
28	Patients with MEN-1 are more insulin-resistant than their non-affected relatives. <i>European Journal of Internal Medicine</i> , 2005, 16, 507-509.	2.2	7
29	Human ApoA-I Overexpression Enhances Macrophage-Specific Reverse Cholesterol Transport but Fails to Prevent Inherited Diabetes in Mice. <i>International Journal of Molecular Sciences</i> , 2019, 20, 655.	4.1	6
30	Polygenic Markers in Patients Diagnosed of Autosomal Dominant Hypercholesterolemia in Catalonia: Distribution of Weighted LDL-c-Raising SNP Scores and Refinement of Variant Selection. <i>Biomedicines</i> , 2020, 8, 353.	3.2	6
31	Molecular Diagnosis of Lecithin: Cholesterol Acyltransferase Deficiency in a Presymptomatic Proband. <i>Clinical Chemistry and Laboratory Medicine</i> , 1998, 36, 443-8.	2.3	5
32	Genome-Wide Studies in Ischaemic Stroke: Are Genetics Only Useful for Finding Genes?. <i>International Journal of Molecular Sciences</i> , 2022, 23, 6840.	4.1	3
33	Genetic Determinants of Plasma Low-Density Lipoprotein Cholesterol Levels: Monogenicity, Polygenicity, and Missing Heritability. <i>Biomedicines</i> , 2021, 9, 1728.	3.2	0