

Ángel-Luis García-Otáñez

List of Publications by Year in descending order

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39
papers

934
citations

430442

18
h-index

454577

30
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42
all docs

42
docs citations

42
times ranked

1444
citing authors

#	ARTICLE	IF	CITATIONS
1	Human Apolipoprotein A-IV Reduces Secretion of Proinflammatory Cytokines and Atherosclerotic Effects of a Chronic Infection Mimicked by Lipopolysaccharide. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2004, 24, 756-761.	1.1	95
2	Mammalian genome targeting using site-specific recombinases. <i>Frontiers in Bioscience - Landmark</i> , 2006, 11, 1108.	3.0	86
3	Frequency of Low-Density Lipoprotein Receptor Gene Mutations in Patients With a Clinical Diagnosis of Familial Combined Hyperlipidemia in a Clinical Setting. <i>Journal of the American College of Cardiology</i> , 2008, 52, 1546-1553.	1.2	73
4	Individual Variation of Scavenger Receptor Expression in Human Macrophages with Oxidized Low-Density Lipoprotein Is Associated with a Differential Inflammatory Response. <i>Journal of Immunology</i> , 2007, 179, 3242-3248.	0.4	64
5	Myelination and motor coordination are increased in transferrin transgenic mice. <i>Journal of Neuroscience Research</i> , 2003, 72, 587-594.	1.3	57
6	Apo E variants in patients with type III hyperlipoproteinemia. <i>Atherosclerosis</i> , 1996, 127, 273-282.	0.4	46
7	Pharmacological activation of TRPV4 produces immediate cell damage and induction of apoptosis in human melanoma cells and HaCaT keratinocytes. <i>PLoS ONE</i> , 2018, 13, e0190307.	1.1	39
8	Oligodendrocyte differentiation is increased in transferrin transgenic mice. <i>Journal of Neuroscience Research</i> , 2006, 83, 403-414.	1.3	33
9	Increased Intestinal Cholesterol Absorption in Autosomal Dominant Hypercholesterolemia and No Mutations in the Low-Density Lipoprotein Receptor or Apolipoprotein B Genes. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2007, 92, 3667-3673.	1.8	32
10	Role of naturally-occurring plant sterols on intestinal cholesterol absorption and plasmatic levels. <i>Journal of Physiology and Biochemistry</i> , 2009, 65, 87-98.	1.3	27
11	Comparison of the hypolipidemic effect of gemfibrozil versus simvastatin in patients with type III hyperlipoproteinemia. <i>American Heart Journal</i> , 1999, 138, 156-162.	1.2	26
12	FABP4 plasma levels are increased in familial combined hyperlipidemia. <i>Journal of Lipid Research</i> , 2010, 51, 1173-1178.	2.0	26
13	A presumptive new locus for autosomal dominant hypercholesterolemia mapping to 8q24.22. <i>Clinical Genetics</i> , 2011, 79, 475-481.	1.0	25
14	Naturally-occurring phytosterols in the usual diet influence cholesterol metabolism in healthy subjects. <i>Nutrition, Metabolism and Cardiovascular Diseases</i> , 2012, 22, 849-855.	1.1	25
15	Novel Phenolic Inhibitors of Small/Intermediate-Conductance Ca ²⁺ -Activated K ⁺ Channels, KCa3.1 and KCa2.3. <i>PLoS ONE</i> , 2013, 8, e58614.	1.1	25
16	Allelic polymorphism ϵ 491A/T in apo E gene modulates the lipid-lowering response in combined hyperlipidemia treatment. <i>European Journal of Clinical Investigation</i> , 2002, 32, 421-428.	1.7	24
17	Association of plasma markers of cholesterol homeostasis with metabolic syndrome components. A cross-sectional study. <i>Nutrition, Metabolism and Cardiovascular Diseases</i> , 2011, 21, 651-657.	1.1	24
18	Plasma lipoprotein responses to enzyme-replacement in Gaucher's disease. <i>Lancet</i> , The, 1999, 353, 642-643.	6.3	22

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19	Analysis of apolipoprotein A-I, lecithin:cholesterol acyltransferase and glucocerebrosidase genes in hypoalphalipoproteinemia. <i>Atherosclerosis</i> , 2002, 163, 49-58.	0.4	19
20	An NPC1L1 gene promoter variant is associated with autosomal dominant hypercholesterolemia. <i>Nutrition, Metabolism and Cardiovascular Diseases</i> , 2010, 20, 236-242.	1.1	18
21	Proteomic study of macrophages exposed to oxLDL identifies a CAPG polymorphism associated with carotid atherosclerosis. <i>Atherosclerosis</i> , 2009, 207, 32-37.	0.4	14
22	ImageJ-based semiautomatic method to analyze senescence in cell culture. <i>Analytical Biochemistry</i> , 2018, 543, 30-32.	1.1	14
23	Hyperlipoproteinaemia(a) is a common cause of autosomal dominant hypercholesterolaemia. <i>Journal of Inherited Metabolic Disease</i> , 2007, 30, 970-977.	1.7	12
24	Novel antiangiogenic therapies against advanced hepatocellular carcinoma (HCC). <i>Clinical and Translational Oncology</i> , 2012, 14, 564-574.	1.2	12
25	A novel DNA polymorphism (4886C>T) in the human LCAT gene. <i>Human Mutation</i> , 2000, 15, 298-298.	1.1	11
26	Inhibition of Intermediate-Conductance Calcium-Activated K Channel (KCa3.1) and Fibroblast Mitogenesis by α -Linolenic Acid and Alterations of Channel Expression in the Lysosomal Storage Disorders, Fabry Disease, and Niemann Pick C. <i>Frontiers in Physiology</i> , 2017, 8, 39.	1.3	11
27	Atorvastatin Decreases Stearoyl-CoA Desaturase Gene Expression in THP-1 Macrophages Incubated with Oxidized LDL. <i>Lipids</i> , 2009, 44, 115-123.	0.7	10
28	A moderate intake of phytosterols from habitual diet affects cholesterol metabolism. <i>Journal of Physiology and Biochemistry</i> , 2009, 65, 397-404.	1.3	10
29	Overexpression of the CXCL3 gene in response to oxidized low-density lipoprotein is associated with the presence of tendon xanthomas in familial hypercholesterolemia. <i>Biochemistry and Cell Biology</i> , 2009, 87, 493-498.	0.9	10
30	Association and Linkage Disequilibrium Analyses of APOE Polymorphisms in Atherosclerosis. <i>Disease Markers</i> , 2008, 24, 65-72.	0.6	8
31	Haplotype analyses, mechanism and evolution of common double mutants in the human LDL receptor gene. <i>Molecular Genetics and Genomics</i> , 2010, 283, 565-574.	1.0	7
32	Vascular Reactivity Profile of Novel $Ca_v3.1$ -Selective Positive-Gating Modulators in the Coronary Vascular Bed. <i>Basic and Clinical Pharmacology and Toxicology</i> , 2016, 119, 184-192.	1.2	6
33	KCa3.1 Transgene Induction in Murine Intestinal Epithelium Causes Duodenal Chyme Accumulation and Impairs Duodenal Contractility. <i>International Journal of Molecular Sciences</i> , 2019, 20, 1193.	1.8	6
34	New contributions to the study of common double mutants in the human LDL receptor gene. <i>Die Naturwissenschaften</i> , 2011, 98, 943-949.	0.6	5
35	Conditional KCa3.1-transgene induction in murine skin produces pruritic eczematous dermatitis with severe epidermal hyperplasia and hyperkeratosis. <i>PLoS ONE</i> , 2020, 15, e0222619.	1.1	3
36	Síntesis y purificación de apolipoproteína apo A-I Zaragoza (L144R) recombinante. <i>Clínica e Investigación En Arteriosclerosis</i> , 2010, 22, 146-153.	0.4	1

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37	Expression and purification of recombinant apolipoprotein A-I Zaragoza (L144R) and formation of reconstituted HDL particles. <i>Protein Expression and Purification</i> , 2011, 80, 110-116.	0.6	1
38	Genetics and molecular biology. <i>Current Opinion in Lipidology</i> , 2003, 14, 531-535.	1.2	0
39	Estudio genético de la implicación del gen USF1 en el desarrollo del síndrome metabólico. <i>Clínica e Investigación en Arteriosclerosis</i> , 2011, 23, 78-87.	0.4	0