

Pablo Fuentes-Prior

List of Publications by Year in descending order

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54
papers

3,794
citations

136950

32
h-index

175258

52
g-index

57
all docs

57
docs citations

57
times ranked

5176
citing authors

#	ARTICLE	IF	CITATIONS
1	Structural basis for the anticoagulant activity of the thrombin-thrombomodulin complex. <i>Nature</i> , 2000, 404, 518-525.	27.8	304
2	Correlation between SMA type and SMN2 copy number revisited: An analysis of 625 unrelated Spanish patients and a compilation of 2834 reported cases. <i>Neuromuscular Disorders</i> , 2018, 28, 208-215.	0.6	273
3	Crystal structures of the membrane-binding C2 domain of human coagulation factor V. <i>Nature</i> , 1999, 402, 434-439.	27.8	258
4	Staphylocoagulase is a prototype for the mechanism of cofactor-induced zymogen activation. <i>Nature</i> , 2003, 425, 535-539.	27.8	234
5	Potent Phagocytic Activity with Impaired Antigen Presentation Identifying Lipopolysaccharide-Tolerant Human Monocytes: Demonstration in Isolated Monocytes from Cystic Fibrosis Patients. <i>Journal of Immunology</i> , 2009, 182, 6494-6507.	0.8	193
6	Ionomycin-activated Calpain Triggers Apoptosis. <i>Journal of Biological Chemistry</i> , 2002, 277, 27217-27226.	3.4	183
7	Metalloproteinases Shed TREM-1 Ectodomain from Lipopolysaccharide-Stimulated Human Monocytes. <i>Journal of Immunology</i> , 2007, 179, 4065-4073.	0.8	176
8	In vivo detection of <i>Staphylococcus aureus</i> endocarditis by targeting pathogen-specific prothrombin activation. <i>Nature Medicine</i> , 2011, 17, 1142-1146.	30.7	144
9	Mutation update of spinal muscular atrophy in Spain: molecular characterization of 745 unrelated patients and identification of four novel mutations in the SMN1 gene. <i>Human Genetics</i> , 2009, 125, 29-39.	3.8	139
10	Structure of the homodimeric androgen receptor ligand-binding domain. <i>Nature Communications</i> , 2017, 8, 14388.	12.8	131
11	Tumor Cells Deactivate Human Monocytes by Up-Regulating IL-1 Receptor Associated Kinase-M Expression via CD44 and TLR4. <i>Journal of Immunology</i> , 2005, 174, 3032-3040.	0.8	125
12	Catalytic Domain Structures of MT-SP1/Matriptase, a Matrix-degrading Transmembrane Serine Proteinase. <i>Journal of Biological Chemistry</i> , 2002, 277, 2160-2168.	3.4	105
13	Isolation, Cloning and Structural Characterisation of Boophilin, a Multifunctional Kunitz-Type Proteinase Inhibitor from the Cattle Tick. <i>PLoS ONE</i> , 2008, 3, e1624.	2.5	103
14	Tick-derived Kunitz-type inhibitors as antihemostatic factors. <i>Insect Biochemistry and Molecular Biology</i> , 2009, 39, 579-595.	2.7	86
15	Monocytes from Cystic Fibrosis Patients Are Locked in an LPS Tolerance State: Down-Regulation of TREM-1 as Putative Underlying Mechanism. <i>PLoS ONE</i> , 2008, 3, e2667.	2.5	76
16	New Insights into Binding Interfaces of Coagulation Factors V and VIII and their Homologues - Lessons from High Resolution Crystal Structures. <i>Current Protein and Peptide Science</i> , 2002, 3, 313-339.	1.4	66
17	Priming of SARS-CoV-2 S protein by several membrane-bound serine proteinases could explain enhanced viral infectivity and systemic COVID-19 infection. <i>Journal of Biological Chemistry</i> , 2021, 296, 100135.	3.4	63
18	Exploring the Collagen-binding Site of the DDR1 Tyrosine Kinase Receptor. <i>Journal of Biological Chemistry</i> , 2004, 279, 31462-31470.	3.4	61

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19	Subcellular Localization and in Vivo Subunit Interactions of Ubiquitous β -Calpain. <i>Journal of Biological Chemistry</i> , 2003, 278, 16336-16346.	3.4	60
20	Fibrinogen Substrate Recognition by Staphylocoagulase-(Pro)thrombin Complexes. <i>Journal of Biological Chemistry</i> , 2006, 281, 1179-1187.	3.4	56
21	The 2.2-Å... Crystal Structure of Human Pro-granzyme K Reveals a Rigid Zymogen with Unusual Features. <i>Journal of Biological Chemistry</i> , 2002, 277, 50923-50933.	3.4	55
22	Inflammatory responses associated with acute coronary syndrome up-regulate IRAK-M and induce endotoxin tolerance in circulating monocytes. <i>Journal of Endotoxin Research</i> , 2007, 13, 39-52.	2.5	55
23	Metazoan evolution of glutamate receptors reveals unreported phylogenetic groups and divergent lineage-specific events. <i>ELife</i> , 2018, 7, .	6.0	53
24	The 1.4 Å... Crystal Structure of Kumamolysin. <i>Structure</i> , 2002, 10, 865-876.	3.3	51
25	Leech-Derived Thrombin Inhibitors: From Structures to Mechanisms to Clinical Applications. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 3847-3861.	6.4	51
26	Unique thrombin inhibition mechanism by anophelin, an anticoagulant from the malaria vector. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, E3649-58.	7.1	49
27	K Domain CR9 of Low Density Lipoprotein (LDL) Receptor-related Protein 1 (LRP1) Is Critical for Aggregated LDL-induced Foam Cell Formation from Human Vascular Smooth Muscle Cells. <i>Journal of Biological Chemistry</i> , 2015, 290, 14852-14865.	3.4	48
28	Molecular basis of protein S deficiency. <i>Thrombosis and Haemostasis</i> , 2007, 98, 543-56.	3.4	48
29	The factor V C1 domain is involved in membrane binding: identification of functionally important amino acid residues within the C1 domain of factor V using alanine scanning mutagenesis. <i>Thrombosis and Haemostasis</i> , 2004, 91, 16-27.	3.4	43
30	C-Reactive Protein as a Therapeutic Target in Age-Related Macular Degeneration. <i>Frontiers in Immunology</i> , 2018, 9, 808.	4.8	42
31	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
32	Novel Fluorescent Prothrombin Analogs as Probes of Staphylocoagulase-Prothrombin Interactions. <i>Journal of Biological Chemistry</i> , 2006, 281, 1169-1178.	3.4	33
33	Practical guidelines to manage discordant situations of <i>SMN2</i> copy number in patients with spinal muscular atrophy. <i>Neurology: Genetics</i> , 2020, 6, e530.	1.9	32
34	Oleic acid is an endogenous ligand of TLX/NR2E1 that triggers hippocampal neurogenesis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022, 119, e2023784119.	7.1	30
35	Utility of two <i>SMN1</i> variants to improve spinal muscular atrophy carrier diagnosis and genetic counselling. <i>European Journal of Human Genetics</i> , 2018, 26, 1554-1557.	2.8	28
36	Plasminogen Substrate Recognition by the Streptokinase-Plasminogen Catalytic Complex Is Facilitated by Arg253, Lys256, and Lys257 in the Streptokinase β -Domain and Kringle 5 of the Substrate. <i>Journal of Biological Chemistry</i> , 2009, 284, 19511-19521.	3.4	27

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37	Pathophysiology of interleukin-1 receptor-associated kinase-M: implications in refractory state. <i>Current Opinion in Infectious Diseases</i> , 2006, 19, 237-244.	3.1	26
38	Rare Variants in Calcium Homeostasis Modulator 1 (CALHM1) Found in Early Onset Alzheimer's Disease Patients Alter Calcium Homeostasis. <i>PLoS ONE</i> , 2013, 8, e74203.	2.5	26
39	Beyond copy number: A new, rapid, and versatile method for sequencing the entire <i>SMN2</i> gene in SMA patients. <i>Human Mutation</i> , 2021, 42, 787-795.	2.5	23
40	6-Methylprednisolone down-regulates IRAK-M in human and murine osteoclasts and boosts bone-resorbing activity: a putative mechanism for corticoid-induced osteoporosis. <i>Journal of Leukocyte Biology</i> , 2007, 82, 700-709.	3.3	22
41	A novel gain-of-function STAT1 mutation resulting in basal phosphorylation of STAT1 and increased distal IFN- γ -mediated responses in chronic mucocutaneous candidiasis. <i>Molecular Immunology</i> , 2015, 68, 597-605.	2.2	21
42	Structural Basis for Reduced Staphylocoagulase-mediated Bovine Prothrombin Activation. <i>Journal of Biological Chemistry</i> , 2006, 281, 1188-1195.	3.4	19
43	A rare STAP1 mutation incompletely associated with familial hypercholesterolemia. <i>Clinica Chimica Acta</i> , 2018, 487, 270-274.	1.1	19
44	Non-canonical dimerization of the androgen receptor and other nuclear receptors: implications for human disease. <i>Endocrine-Related Cancer</i> , 2019, 26, R479-R497.	3.1	19
45	Contribution of globular death domains and unstructured linkers to MyD88-IRAK-4 heterodimer formation: An explanation for the antagonistic activity of MyD88s. <i>Biochemical and Biophysical Research Communications</i> , 2009, 380, 183-187.	2.1	18
46	Identification of 31 novel mutations in the F8 gene in Spanish hemophilia A patients: structural analysis of 20 missense mutations suggests new intermolecular binding sites. <i>Blood</i> , 2008, 111, 3468-3478.	1.4	16
47	Clinical and genetic findings in five female patients with haemophilia A: Identification of a novel missense mutation, p.Phe2127Ser. <i>Thrombosis and Haemostasis</i> , 2010, 104, 718-723.	3.4	15
48	Structural basis of thrombin-mediated factor V activation: the Glu666-Glu672 sequence is critical for processing at the heavy chain-B domain junction. <i>Blood</i> , 2011, 117, 7164-7173.	1.4	14
49	Diversity of Quaternary Structures Regulates Nuclear Receptor Activities. <i>Trends in Biochemical Sciences</i> , 2019, 44, 2-6.	7.5	13
50	TET2 missense variants in human neoplasia. A proposal of structural and functional classification. <i>Molecular Genetics & Genomic Medicine</i> , 2019, 7, e00772.	1.2	9
51	Position-dependent expression of GADD45 β in rat brain tumours. <i>Medical Oncology</i> , 2007, 24, 436-444.	2.5	3
52	Producción heteróloga y caracterización bioquímica del procoagulante humano Factor VIII para ensayos de cristalización de macromoléculas proteicas. <i>Tecnología En Marcha</i> , 2016, 29, 78.	0.1	0
53	Mutagenesis dirigida del conector interdominio Ácido FVIIIa3 del factor VIII de la coagulación como estrategia para favorecer la cristalización de sus complejos con la trombina. <i>Tecnología En Marcha</i> , 0, .	0.1	0
54	Confirmation of inherited protein S deficiency by PROS1 mutational screening: Identification of three novel PROS1 mutations and haplotype analysis of p.Q279X recurrence. <i>Thrombosis and Haemostasis</i> , 2008, 100, 721-4.	3.4	0