Jaime M Merino

List of Publications by Year in descending order

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LAIME M MEDINO

#	Article	IF	CITATIONS
1	Aryl Hydrocarbon Receptor: From Homeostasis to Tumor Progression. Frontiers in Cell and Developmental Biology, 2022, 10, 884004.	3.7	8
2	Aryl hydrocarbon receptor blocks aging-induced senescence in the liver and fibroblast cells. Aging, 2022, 14, 4281-4304.	3.1	10
3	The aryl hydrocarbon receptor promotes differentiation during mouse preimplantational embryo development. Stem Cell Reports, 2021, 16, 2351-2363.	4.8	9
4	Alu retrotransposons modulate Nanog expression through dynamic changes in regional chromatin conformation via aryl hydrocarbon receptor. Epigenetics and Chromatin, 2020, 13, 15.	3.9	12
5	The aryl hydrocarbon receptor in the crossroad of signalling networks with therapeutic value. , 2018, 185, 50-63.		72
6	Aryl Hydrocarbon Receptor Promotes Liver Polyploidization and Inhibits PI3K, ERK, and Wnt/β-Catenin Signaling. IScience, 2018, 4, 44-63.	4.1	26
7	Dioxin Receptor Adjusts Liver Regeneration After Acute Toxic Injury and Protects Against Liver Carcinogenesis. Scientific Reports, 2017, 7, 10420.	3.3	25
8	Lung regeneration after toxic injury is improved in absence of dioxin receptor. Stem Cell Research, 2017, 25, 61-71.	0.7	21
9	AhR-dependent 2,3,7,8-tetrachlorodibenzo- p -dioxin toxicity in human neuronal cell line SHSY5Y. NeuroToxicology, 2016, 56, 55-63.	3.0	12
10	<i>Alu</i> retrotransposons promote differentiation of human carcinoma cells through the aryl hydrocarbon receptor. Nucleic Acids Research, 2016, 44, 4665-4683.	14.5	45
11	Dioxin receptor regulates aldehyde dehydrogenase to block melanoma tumorigenesis and metastasis. Molecular Cancer, 2015, 14, 148.	19.2	31
12	2,3,7,8-Tetrachlorodibenzo-p-dioxin induces apoptosis by disruption of intracellular calcium homeostasis in human neuronal cell line SHSY5Y. Apoptosis: an International Journal on Programmed Cell Death, 2012, 17, 1170-1181.	4.9	36
13	Aryl hydrocarbon receptorâ€dependent induction of apoptosis by 2,3,7,8â€ŧetrachlorodibenzoâ€ <i>p</i> â€dioxin in cerebellar granule cells from mouse. Journal of Neurochemistry, 2011, 118, 153-162.	3.9	51
14	2,3,7,8-Tetrachlorodibenzo-p-dioxin induces apoptosis in neural growth factor (NGF)-differentiated pheochromocytoma PC12 cells. NeuroToxicology, 2010, 31, 267-276.	3.0	35
15	Nerve growth factor increases the sensitivity to zinc toxicity and induces cell cycle arrest in PC12 cells. Brain Research Bulletin, 2010, 81, 458-466.	3.0	19
16	The Dioxin Receptor Regulates the Constitutive Expression of the <i>Vav3</i> Proto-Oncogene and Modulates Cell Shape and Adhesion. Molecular Biology of the Cell, 2009, 20, 1715-1727.	2.1	72
17	NMDA-induced neuroprotection in hippocampal neurons is mediated through the protein kinase A and CREB (cAMP-response element-binding protein) pathway. Neurochemistry International, 2008, 53, 148-154.	3.8	42
18	Resveratrolâ€induced apoptosis in MCFâ€7 human breast cancer cells involves a caspaseâ€independent mechanism with downregulation of Bclâ€2 and NFâ€ĤB. International Journal of Cancer, 2005, 115, 74-84.	5.1	208

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19	Small peptides patterned after the Nâ€ŧerminus domain of SNAP25 inhibit SNARE complex assembly and regulated exocytosis. Journal of Neurochemistry, 2004, 88, 124-135.	3.9	39
20	Down-regulation of CYP1A2 induction during the maturation of mouse cerebellar granule cells in culture: role of nitric oxide accumulation. European Journal of Neuroscience, 2003, 18, 2265-2272.	2.6	13
21	Identification of SNARE complex modulators that inhibit exocytosis from an α-helix-constrained combinatorial library. Biochemical Journal, 2003, 375, 159-166.	3.7	23
22	A Novel N-Methyl-d-aspartate Receptor Open Channel Blocker with in Vivo Neuroprotectant Activity. Journal of Pharmacology and Experimental Therapeutics, 2002, 302, 163-173.	2.5	41
23	Neuroprotection Against Excitotoxicity by N-Alkylglycines in Rat Hippocampal Neurons. NeuroMolecular Medicine, 2002, 2, 271-280.	3.4	10
24	pH and ligand binding modulate the strength of protein–protein interactions in the Ca2+-ATPase from sarcoplasmic reticulum membranes. Biochimica Et Biophysica Acta - Biomembranes, 1999, 1420, 203-213.	2.6	3
25	Plausible Stoichiometry of the Interacting Nucleotide-Binding Sites in the Ca2+-ATPase from Sarcoplasmic Reticulum Membranes. Archives of Biochemistry and Biophysics, 1999, 368, 298-302.	3.0	4
26	Selected peptides targeted to the NMDA receptor channel protect neurons from excitotoxic death. Nature Biotechnology, 1998, 16, 286-291.	17.5	43
27	Structural determinants of the blocker binding site in glutamate and NMDA receptor channels. Neuropharmacology, 1998, 37, 139-147.	4.1	35
28	Structural Changes of the Sarcoplasmic Reticulum Ca(II)-ATPase Nucleotide Binding Domain by pH and La(III). Archives of Biochemistry and Biophysics, 1997, 348, 152-156.	3.0	3
29	Thermal unfolding of monomeric Ca(II),Mg(II)-ATPase from sarcoplasmic reticulum of rabbit skeletal muscle. FEBS Letters, 1994, 343, 155-159.	2.8	18
30	Fluorescence energy transfer as a tool to locate functional sites in membrane proteins. Biochemical Society Transactions, 1994, 22, 784-788.	3.4	14
31	Differential scanning calorimetry study of the thermal unfolding of sarcoplasmic reticulum Ca2+, Mg2+-ATPase from rabbit skeletal muscle. Biochemical Society Transactions, 1994, 22, 384S-384S.	3.4	2