

Sandrine Silvente-Poirot

List of Publications by Year in descending order

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68
papers

3,830
citations

109321

35
h-index

123424

61
g-index

68
all docs

68
docs citations

68
times ranked

7066
citing authors

#	ARTICLE	IF	CITATIONS
1	Oxysterols are potential physiological regulators of ageing. <i>Ageing Research Reviews</i> , 2022, 77, 101615.	10.9	21
2	Targeting the liver X receptor with dendrogenin A differentiates tumour cells to secrete immunogenic exosome-enriched vesicles. <i>Journal of Extracellular Vesicles</i> , 2022, 11, e12211.	12.2	8
3	The 5,6-epoxycholesterol metabolic pathway in breast cancer: Emergence of new pharmacological targets. <i>British Journal of Pharmacology</i> , 2021, 178, 3248-3260.	5.4	27
4	Neutral Sphingomyelinase 2 Heightens Anti-Melanoma Immune Responses and Anti-PD-1 Therapy Efficacy. <i>Cancer Immunology Research</i> , 2021, 9, 568-582.	3.4	30
5	Dendrogenin A Enhances Anti-Leukemic Effect of Anthracycline in Acute Myeloid Leukemia. <i>Cancers</i> , 2020, 12, 2933.	3.7	7
6	Dendrogenin A Synergizes with Cytarabine to Kill Acute Myeloid Leukemia Cells In Vitro and In Vivo. <i>Cancers</i> , 2020, 12, 1725.	3.7	13
7	A fast UPLC-HILIC method for an accurate quantification of dendrogenin A in human tissues. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019, 194, 105447.	2.5	7
8	Oxysterols: An expanding family of structurally diversified bioactive steroids. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019, 194, 105443.	2.5	9
9	The cholesterol-derived metabolite dendrogenin A functionally reprograms breast adenocarcinoma and undifferentiated thyroid cancer cells. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019, 192, 105390.	2.5	22
10	Natural and semisynthetic oxyprenylated aromatic compounds as stimulators or inhibitors of melanogenesis. <i>Bioorganic Chemistry</i> , 2019, 87, 181-190.	4.1	9
11	Flavonoids differentially modulate liver X receptors activity-Structure-function relationship analysis. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019, 190, 173-182.	2.5	22
12	Chemistry, biochemistry, metabolic fate and mechanism of action of 6-oxo-cholestan-3 β ,5 α -diol (OCDO), a tumor promoter and cholesterol metabolite. <i>Biochimie</i> , 2018, 153, 139-149.	2.6	21
13	The tumor-suppressor cholesterol metabolite, dendrogenin A, is a new class of LXR modulator activating lethal autophagy in cancers. <i>Biochemical Pharmacology</i> , 2018, 153, 75-81.	4.4	48
14	Ligand-dependent transcriptional induction of lethal autophagy: A new perspective for cancer treatment. <i>Autophagy</i> , 2018, 14, 555-557.	9.1	25
15	Extracellular vesicles: lipids as key components of their biogenesis and functions. <i>Journal of Lipid Research</i> , 2018, 59, 1316-1324.	4.2	208
16	The Effects of Cholesterol-Derived Oncometabolites on Nuclear Receptor Function in Cancer. <i>Cancer Research</i> , 2018, 78, 4803-4808.	0.9	45
17	Circulating oxysterol metabolites as potential new surrogate markers in patients with hormone receptor-positive breast cancer: Results of the OXYTAM study. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2017, 169, 210-218.	2.5	48
18	Improvement of 5,6 α -epoxycholesterol, 5,6 β -epoxycholesterol, cholestane-3 β ,5 α ,6 β -triol and 6-oxo-cholestan-3 β ,5 α -diol recovery for quantification by GC/MS. <i>Chemistry and Physics of Lipids</i> , 2017, 207, 92-98.	3.2	7

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19	Identification of a tumor-promoter cholesterol metabolite in human breast cancers acting through the glucocorticoid receptor. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E9346-E9355.	7.1	96
20	Dendrogenin A drives LXR to trigger lethal autophagy in cancers. Nature Communications, 2017, 8, 1903.	12.8	84
21	Improving the efficacy of hormone therapy in breast cancer: The role of cholesterol metabolism in SERM-mediated autophagy, cell differentiation and death. Biochemical Pharmacology, 2017, 144, 18-28.	4.4	43
22	Quantitative analysis of the tumor suppressor dendrogenin A using liquid chromatography tandem mass spectrometry. Chemistry and Physics of Lipids, 2017, 207, 81-86.	3.2	8
23	When cholesterol meets histamine, it gives rise to dendrogenin A: a tumour suppressor metabolite ¹ . Biochemical Society Transactions, 2016, 44, 631-637.	3.4	17
24	From tamoxifen to dendrogenin A: The discovery of a mammalian tumor suppressor and cholesterol metabolite. Biochimie, 2016, 130, 109-114.	2.6	21
25	Dendrogenin A and B two new steroidal alkaloids increasing neural responsiveness in the deafened guinea pig. Frontiers in Aging Neuroscience, 2015, 7, 145.	3.4	11
26	Dendrogenin A: A Mammalian Metabolite of Cholesterol with Tumor Suppressor and Neurostimulating Properties. Current Medicinal Chemistry, 2015, 22, 3533-3549.	2.4	24
27	Cholesterol and Cancer, in the Balance. Science, 2014, 343, 1445-1446.	12.6	182
28	Emerging concepts on the role of exosomes in lipid metabolic diseases. Biochimie, 2014, 96, 67-74.	2.6	62
29	One step synthesis of 6-oxo-cholestan-3 β ,5 α -diol. Biochemical and Biophysical Research Communications, 2014, 446, 782-785.	2.1	11
30	The novel steroidal alkaloids dendrogenin A and B promote proliferation of adult neural stem cells. Biochemical and Biophysical Research Communications, 2014, 446, 681-686.	2.1	21
31	Dendrogenin_A : A Natural Liver X Receptor Modulator for the Treatment of Acute Myeloid Leukemia. Blood, 2014, 124, 3767-3767.	1.4	0
32	5,6-Epoxy-cholesterols contribute to the anticancer pharmacology of Tamoxifen in breast cancer cells. Biochemical Pharmacology, 2013, 86, 175-189.	4.4	56
33	Cholesterol-5,6-epoxides: Chemistry, biochemistry, metabolic fate and cancer. Biochimie, 2013, 95, 622-631.	2.6	69
34	Technical note: Hapten synthesis, antibody production and development of an enzyme-linked immunosorbent assay for detection of the natural steroidal alkaloid Dendrogenin A. Biochimie, 2013, 95, 482-488.	2.6	1
35	Dendrogenin A arises from cholesterol and histamine metabolism and shows cell differentiation and anti-tumour properties. Nature Communications, 2013, 4, 1840.	12.8	101
36	Antiestrogen-binding site ligands induce autophagy in myeloma cells that proceeds through alteration of cholesterol metabolism. Oncotarget, 2013, 4, 911-922.	1.8	27

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37	Surprising unreactivity of cholesterol-5,6-epoxides towards nucleophiles. <i>Journal of Lipid Research</i> , 2012, 53, 718-725.	4.2	36
38	Cholesterol epoxide hydrolase and cancer. <i>Current Opinion in Pharmacology</i> , 2012, 12, 696-703.	3.5	71
39	Cholesterol metabolism and resistance to tamoxifen. <i>Current Opinion in Pharmacology</i> , 2012, 12, 683-689.	3.5	49
40	[¹⁸ F]Siâ€RiboRGD: From Design and Synthesis to the Imaging of $\int_{\pm} v < /sub > \hat{I}^2 < /sub > 3 < /sub > \hat{a} \dots$ Integrins in Melanoma Tumors. <i>ChemPlusChem</i> , 2012, 77, 345-349.	2.8	11
41	Exosomes as intercellular signalosomes and pharmacological effectors. <i>Biochemical Pharmacology</i> , 2011, 81, 1171-1182.	4.4	471
42	Importance of cholesterol and oxysterols metabolism in the pharmacology of tamoxifen and other AEBS ligands. <i>Chemistry and Physics of Lipids</i> , 2011, 164, 432-437.	3.2	51
43	Development of a new radioligand for cholecystokinin receptor subtype 2 scintigraphy: From molecular modeling to in vivo evaluation. <i>Bioorganic and Medicinal Chemistry</i> , 2010, 18, 5400-5412.	3.0	12
44	Synthesis, characterization and in vitro evaluation of new oxorhenium- and oxotechnetium-CCK4 derivatives as molecular imaging agents for CCK2-receptor targeting. <i>European Journal of Medicinal Chemistry</i> , 2010, 45, 423-429.	5.5	8
45	Exosomes account for vesicle-mediated transcellular transport of activatable phospholipases and prostaglandins. <i>Journal of Lipid Research</i> , 2010, 51, 2105-2120.	4.2	528
46	Identification and pharmacological characterization of cholesterol-5,6-epoxide hydrolase as a target for tamoxifen and AEBS ligands. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 13520-13525.	7.1	109
47	Auraptene Is an Inhibitor of Cholesterol Esterification and a Modulator of Estrogen Receptors. <i>Molecular Pharmacology</i> , 2010, 78, 827-836.	2.3	50
48	Signaling through cholesterol esterification: a new pathway for the cholecystokinin 2 receptor involved in cell growth and invasion. <i>Journal of Lipid Research</i> , 2009, 50, 2203-2211.	4.2	64
49	Tamoxifen and AEBS ligands induced apoptosis and autophagy in breast cancer cells through the stimulation of sterol accumulation. <i>Autophagy</i> , 2009, 5, 1066-1067.	9.1	86
50	Synthesis of New Alkylaminooxysterols with Potent Cell Differentiating Activities: Identification of Leads for the Treatment of Cancer and Neurodegenerative Diseases. <i>Journal of Medicinal Chemistry</i> , 2009, 52, 7765-7777.	6.4	55
51	Microsomal antiestrogen-binding site ligands induce growth control and differentiation of human breast cancer cells through the modulation of cholesterol metabolism. <i>Molecular Cancer Therapeutics</i> , 2008, 7, 3707-3718.	4.1	56
52	Insights into the Cholecystokinin 2 Receptor Binding Site and Processes of Activation. <i>Molecular Pharmacology</i> , 2006, 70, 1935-1945.	2.3	8
53	The Prototypical Inhibitor of Cholesterol Esterification, Sah 58-035 [3-[Decyldimethylsilyl]-N-[2-(4-methylphenyl)-1-phenylethyl]propanamide], Is an Agonist of Estrogen Receptors. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2006, 319, 139-149.	2.5	20
54	Molecular Characterization of the Microsomal Tamoxifen Binding Site. <i>Journal of Biological Chemistry</i> , 2004, 279, 34048-34061.	3.4	84

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55	Tamoxifen Is a Potent Inhibitor of Cholesterol Esterification and Prevents the Formation of Foam Cells. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2004, 308, 1165-1173.	2.5	71
56	High tumorigenic potential of a constitutively active mutant of the cholecystokinin 2 receptor. <i>Oncogene</i> , 2003, 22, 6081-6089.	5.9	28
57	Identification of Tyrosine 189 and Asparagine 358 of the Cholecystokinin 2 Receptor in Direct Interaction with the Crucial C-Terminal Amide of Cholecystokinin by Molecular Modeling, Site-Directed Mutagenesis, and Structure/Affinity Studies. <i>Molecular Pharmacology</i> , 2003, 63, 973-982.	2.3	25
58	The Biologically Crucial C Terminus of Cholecystokinin and the Non-peptide Agonist SR-146,131 Share a Common Binding Site in the Human CCK1 Receptor. <i>Journal of Biological Chemistry</i> , 2002, 277, 7546-7555.	3.4	63
59	Structure of Cholecystokinin Receptor Binding Sites and Mechanism of Activation/Inactivation by Agonists/Antagonists. <i>Basic and Clinical Pharmacology and Toxicology</i> , 2002, 91, 313-320.	0.0	35
60	Mutation of Asn-391 within the Conserved NPXXY Motif of the Cholecystokinin B Receptor Abolishes Gq Protein Activation without Affecting Its Association with the Receptor. <i>Journal of Biological Chemistry</i> , 2000, 275, 17321-17327.	3.4	52
61	The Third Intracellular Loop of the Rat and Mouse Cholecystokinin-A Receptors Is Responsible for Different Patterns of Gene Activation. <i>Molecular Pharmacology</i> , 2000, 58, 1381-1388.	2.3	10
62	Evidence for a Direct Interaction between the Penultimate Aspartic Acid of Cholecystokinin and Histidine 207, Located in the Second Extracellular Loop of the Cholecystokinin B Receptor. <i>Journal of Biological Chemistry</i> , 1999, 274, 23191-23197.	3.4	42
63	Arginine 197 of the cholecystokinin receptor binding site interacts with the sulfate of the peptide agonist cholecystokinin. <i>Protein Science</i> , 1999, 8, 2347-2354.	7.6	50
64	Met-195 of the Cholecystokinin-A Receptor Interacts with the Sulfated Tyrosine of Cholecystokinin and Is Crucial for Receptor Transition to High Affinity State. <i>Journal of Biological Chemistry</i> , 1998, 273, 14380-14386.	3.4	71
65	Role of the Extracellular Domains of the Cholecystokinin Receptor in Agonist Binding. <i>Molecular Pharmacology</i> , 1998, 54, 364-371.	2.3	65
66	Ligand-induced Internalization of Cholecystokinin Receptors. <i>Journal of Biological Chemistry</i> , 1997, 272, 18179-18184.	3.4	38
67	A Segment of Five Amino Acids in the Second Extracellular Loop of the Cholecystokinin-B Receptor Is Essential for Selectivity of the Peptide Agonist Gastrin. <i>Journal of Biological Chemistry</i> , 1996, 271, 14698-14706.	3.4	58
68	A new probe for affinity labelling pancreatic cholecystokinin receptor with minor modification of its structure. <i>FEBS Journal</i> , 1989, 185, 397-403.	0.2	42