Christian Carpéné

List of Publications by Year in descending order

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77 papers

3,098 citations

29 h-index

172457

54 g-index

79 all docs

79 docs citations

times ranked

79

3524 citing authors

#	Article	IF	CITATIONS
1	Apelin, a Newly Identified Adipokine Up-Regulated by Insulin and Obesity. Endocrinology, 2005, 146, 1764-1771.	2.8	761
2	Resveratrol, Metabolic Syndrome, and Gut Microbiota. Nutrients, 2018, 10, 1651.	4.1	181
3	Role of Semicarbazide-sensitive Amine Oxidase on Glucose Transport and GLUT4 Recruitment to the Cell Surface in Adipose Cells. Journal of Biological Chemistry, 1998, 273, 8025-8032.	3.4	148
4	Regulation of glucose metabolism by bioactive phytochemicals for the management of type 2 diabetes mellitus. Critical Reviews in Food Science and Nutrition, 2019, 59, 830-847.	10.3	123
5	Adipose Tissue Proadipogenic Redox Changes in Obesity. Journal of Biological Chemistry, 2006, 281, 12682-12687.	3.4	93
6	Substrates of semicarbazide-sensitive amine oxidase co-operate with vanadate to stimulate tyrosine phosphorylation of insulin-receptor-substrate proteins, phosphoinositide 3-kinase activity and GLUT4 translocation in adipose cells. Biochemical Journal, 2000, 350, 171-180.	3.7	90
7	Resveratrol directly affects in vitro lipolysis and glucose transport in human fat cells. Journal of Physiology and Biochemistry, 2013, 69, 585-593.	3.0	68
8	Adipogenesis-related increase of semicarbazide-sensitive amine oxidase and monoamine oxidase in human adipocytes. Biochimie, 2007, 89, 916-925.	2.6	63
9	Semicarbazide-sensitive amine oxidase activity exerts insulin-like effects on glucose metabolism and insulin-signaling pathways in adipose cells. Biochimica Et Biophysica Acta - Proteins and Proteomics, 2003, 1647, 3-9.	2.3	62
10	Isopropylnorsynephrine is a stronger lipolytic agent in human adipocytes than synephrine and other amines present in Citrus aurantium. Journal of Physiology and Biochemistry, 2011, 67, 443-452.	3.0	62
11	High expression of monoamine oxidases in human white adipose tissue: evidence for their involvement in noradrenaline clearance. Biochemical Pharmacology, 1999, 58, 1735-1742.	4.4	61
12	Semicarbazide-Sensitive Amine Oxidase/Vascular Adhesion Protein-1 Activity Exerts an Antidiabetic Action in Goto-Kakizaki Rats. Diabetes, 2003, 52, 1004-1013.	0.6	60
13	Oral Insulin-Mimetic Compounds That Act Independently of Insulin. Diabetes, 2007, 56, 486-493.	0.6	60
14	Amine oxidase substrates mimic several of the insulin effects on adipocyte differentiation in 3T3 F442A cells. Biochemical Journal, 2001, 356, 769-777.	3.7	58
15	Tyramine Stimulates Glucose Uptake in Insulin-Sensitive Tissues in Vitro and in Vivo via Its Oxidation by Amine Oxidases. Journal of Pharmacology and Experimental Therapeutics, 2002, 303, 1238-1247.	2.5	56
16	Lipolytic Effects of beta1-, beta2-, and beta3-Adrenergic Agonists in White Adipose Tissue of Mammals. Annals of the New York Academy of Sciences, 1998, 839, 186-189.	3.8	46
17	Inhibition of rat fat cell lipolysis by monoamine oxidase and semicarbazide-sensitive amine oxidase substrates. European Journal of Pharmacology, 2003, 466, 235-243.	3.5	44
18	Amine oxidase substrates mimic several of the insulin effects on adipocyte differentiation in 3T3 F442A cells. Biochemical Journal, 2001, 356, 769.	3.7	44

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19	Adrenergic Receptors and Fat Cells: Differential Recruitment by Physiological Amines and Homologous Regulation. Obesity, 1995, 3, 507S-514S.	4.0	43
20	Chronic benzylamine administration in the drinking water improves glucose tolerance, reduces body weight gain and circulating cholesterol in high-fat diet-fed mice. Pharmacological Research, 2010, 61, 355-363.	7.1	42
21	Semicarbazide-Sensitive Amine Oxidase/Vascular Adhesion Protein-1 Deficiency Reduces Leukocyte Infiltration into Adipose Tissue and Favors Fat Deposition. American Journal of Pathology, 2009, 174, 1075-1083.	3.8	41
22	Resveratrol Anti-Obesity Effects: Rapid Inhibition of Adipocyte Glucose Utilization. Antioxidants, 2019, 8, 74.	5.1	40
23	Alteration of Amine Oxidase Activity in the Adipose Tissue of Obese Subjects. Obesity, 2004, 12, 547-555.	4.0	39
24	Advances in Integrating Traditional and Omic Biomarkers When Analyzing the Effects of the Mediterranean Diet Intervention in Cardiovascular Prevention. International Journal of Molecular Sciences, 2016, 17, 1469.	4.1	35
25	Short- and long-term insulin-like effects of monoamine oxidases and semicarbazide-sensitive amine oxidase substrates in cultured adipocytes. Metabolism: Clinical and Experimental, 2006, 55, 1397-1405.	3.4	34
26	Pomegranate juice and its main polyphenols exhibit direct effects on amine oxidases from human adipose tissue and inhibit lipid metabolism in adipocytes. Journal of Functional Foods, 2017, 33, 323-331.	3.4	33
27	Natriuretic peptides promote glucose uptake in a cGMP-dependent manner in human adipocytes. Scientific Reports, 2018, 8, 1097.	3.3	33
28	Anti-obesity effects of resveratrol: comparison between animal models and humans. Journal of Physiology and Biochemistry, 2016, 73, 417-429.	3.0	32
29	Piceatannol and resveratrol share inhibitory effects on hydrogen peroxide release, monoamine oxidase and lipogenic activities in adipose tissue, but differ in their antilipolytic properties. Chemico-Biological Interactions, 2016, 258, 115-125.	4.0	32
30	Substrates of semicarbazide-sensitive amine oxidase co-operate with vanadate to stimulate tyrosine phosphorylation of insulin-receptor-substrate proteins, phosphoinositide 3-kinase activity and GLUT4 translocation in adipose cells. Biochemical Journal, 2000, 350, 171.	3.7	30
31	Methylamine but not mafenide mimics insulin-like activity of the semicarbazide-sensitive amine oxidase-substrate benzylamine on glucose tolerance and on human adipocyte metabolism. Pharmacological Research, 2005, 52, 475-484.	7.1	28
32	Benzylamine Exhibits Insulin-Like Effects on Glucose Disposal, Glucose Transport, and Fat Cell Lipolysis in Rabbits and Diabetic Mice. Journal of Pharmacology and Experimental Therapeutics, 2004, 309, 1020-1028.	2.5	27
33	Exploring the Binding Mode of Semicarbazide-Sensitive Amine Oxidase/VAP-1:Â Identification of Novel Substrates with Insulin-like Activity. Journal of Medicinal Chemistry, 2004, 47, 4865-4874.	6.4	27
34	Glucose handling in streptozotocin-induced diabetic rats is improved by tyramine but not by the amine oxidase inhibitor semicarbazide. European Journal of Pharmacology, 2005, 522, 139-146.	3.5	27
35	Antidepressant Phenelzine Alters Differentiation of Cultured Human and Mouse Preadipocytes. Molecular Pharmacology, 2009, 75, 1052-1061.	2.3	26
36	The imidazoline I2-site ligands BU 224 and 2-BFI inhibit MAO-A and MAO-B activities, hydrogen peroxide production, and lipolysis in rodent and human adipocytes. European Journal of Pharmacology, 2006, 552, 20-30.	3.5	25

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37	Past, Present and Future Anti-Obesity Effects of Flavin-Containing and/or Copper-Containing Amine Oxidase Inhibitors. Medicines (Basel, Switzerland), 2019, 6, 9.	1.4	24
38	Oral Administration of Semicarbazide Limits Weight Gain together with Inhibition of Fat Deposition and of Primary Amine Oxidase Activity in Adipose Tissue. Journal of Obesity, 2011, 2011, 1-10.	2.7	23
39	Insulin-mimetic compound hexaquis (benzylammonium) decavanadate is antilipolytic in human fat cells. World Journal of Diabetes, 2017, 8, 143.	3.5	22
40	The Dietary Antioxidant Piceatannol Inhibits Adipogenesis of Human Adipose Mesenchymal Stem Cells and Limits Glucose Transport and Lipogenic Activities in Adipocytes. International Journal of Molecular Sciences, 2018, 19, 2081.	4.1	22
41	Increased primary amine oxidase expression and activity in white adipose tissue of obese and diabetic dbâ°'/â°' mice. Journal of Neural Transmission, 2011, 118, 1071-1077.	2.8	21
42	SSAO substrates exhibiting insulin-like effects in adipocytes as a promising treatment option for metabolic disorders. Future Medicinal Chemistry, 2010, 2, 1735-1749.	2.3	20
43	Oxidation of high doses of serotonin favors lipid accumulation in mouse and human fat cells. Molecular Nutrition and Food Research, 2013, 57, 1089-1099.	3.3	20
44	Pterostilbene Inhibits Lipogenic Activity similar to Resveratrol or Caffeine but Differently Modulates Lipolysis in Adipocytes. Phytotherapy Research, 2017, 31, 1273-1282.	5.8	20
45	Body fat reduction without cardiovascular changes in mice after oral treatment with the <scp>MAO</scp> inhibitor phenelzine. British Journal of Pharmacology, 2018, 175, 2428-2440.	5.4	18
46	The amine oxidase inhibitor phenelzine limits lipogenesis in adipocytes without inhibiting insulin action on glucose uptake. Journal of Neural Transmission, 2013, 120, 997-1003.	2.8	15
47	Mechanisms of the antilipolytic response of human adipocytes to tyramine, a trace amine present in food. Journal of Physiology and Biochemistry, 2018, 74, 623-633.	3.0	15
48	Potential renoprotective effects of piceatannol in ameliorating the early-stage nephropathy associated with obesity in obese Zucker rats. Journal of Physiology and Biochemistry, 2016, 72, 555-566.	3.0	14
49	Obesity of mice lacking VAP-1/SSAO by Aoc3 gene deletion is reproduced in mice expressing a mutated vascular adhesion protein-1 (VAP-1) devoid of amine oxidase activity. Journal of Physiology and Biochemistry, 2021, 77, 141-154.	3.0	14
50	Benzylamine antihyperglycemic effect is abolished by AOC3 gene invalidation in mice but not rescued by semicarbazide-sensitive amine oxidase expression under the control of aP2 promoter. Journal of Physiology and Biochemistry, 2012, 68, 651-662.	3.0	13
51	Dietary Phenolic Compounds Interfere with the Fate of Hydrogen Peroxide in Human Adipose Tissue but Do Not Directly Inhibit Primary Amine Oxidase Activity. Oxidative Medicine and Cellular Longevity, 2016, 2016, 1-15.	4.0	13
52	Lack of functional antilipolytic α-adrenoceptor in rat fat cell: Comparison with hamster adipocyte. Comparative Biochemistry and Physiology Part C: Comparative Pharmacology, 1983, 74, 41-45.	0.2	12
53	Anatomical distribution of primary amine oxidase activity in four adipose depots and plasma of severely obese women with or without a dysmetabolic profile. Journal of Physiology and Biochemistry, 2016, 73, 475-486.	3.0	12
54	Is there an optimal dose for dietary linoleic acid? Lessons from essential fatty acid deficiency supplementation and adipocyte functions in rats. Journal of Physiology and Biochemistry, 2014, 70, 615-627.	3.0	11

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55	Oral Phenelzine Treatment Mitigates Metabolic Disturbances in Mice Fed a High-Fat Diet. Journal of Pharmacology and Experimental Therapeutics, 2019, 371, 555-566.	2.5	11
56	Short-term and rapid effects of lysophosphatidic acid on human adipose cell lipolytic and glucose uptake activities. AIMS Molecular Science, 2016, 3, 222-237.	0.5	10
57	Methylamine Activates Glucose Uptake in Human Adipocytes Without Overpassing Action of Insulin or Stimulating its Secretion in Pancreatic Islets. Medicines (Basel, Switzerland), 2019, 6, 89.	1.4	9
58	Metabolic Effects of Oral Phenelzine Treatment on High-Sucrose-Drinking Mice. International Journal of Molecular Sciences, 2018, 19, 2904.	4.1	8
59	Vanadium-dependent activation of glucose transport in adipocytes by catecholamines is not mediated <i>via </i> adrenoceptor stimulation or monoamine oxidase activity. World Journal of Diabetes, 2020, 11, 622-643.	3.5	8
60	Randomized Clinical Trial: Effects of β-Hydroxy-β-Methylbutyrate (HMB)-Enriched vs. HMB-Free Oral Nutritional Supplementation in Malnourished Cirrhotic Patients. Nutrients, 2022, 14, 2344.	4.1	8
61	5-hydroxytryptamine actions in adipocytes: involvement of monoamine oxidase-dependent oxidation and subsequent PPAR \hat{I}^3 activation. Journal of Neural Transmission, 2013, 120, 919-926.	2.8	7
62	High intake of dietary tyramine does not deteriorate glucose handling and does not cause adverse cardiovascular effects in mice. Journal of Physiology and Biochemistry, 2016, 72, 539-553.	3.0	6
63	Effects of the amino acid derivatives, î²-hydroxy-l²-methylbutyrate, taurine, and N-methyltyramine, on triacylglycerol breakdown in fat cells. Journal of Physiology and Biochemistry, 2019, 75, 263-273.	3.0	6
64	Novel Facet of an Old Dietary Molecule? Direct Influence of Caffeine on Glucose and Biogenic Amine Handling by Human Adipocytes. Molecules, 2021, 26, 3831.	3.8	6
65	Methylxanthines Inhibit Primary Amine Oxidase and Monoamine Oxidase Activities of Human Adipose Tissue. Medicines (Basel, Switzerland), 2020, 7, 18.	1.4	5
66	High doses of tyramine stimulate glucose transport in human fat cells. Journal of Physiology and Biochemistry, 2022, 78, 543-556.	3.0	5
67	Glitazones inhibit human monoamine oxidase but their anti-inflammatory actions are not mediated by VAP-1/semicarbazide-sensitive amine oxidase inhibition. Journal of Physiology and Biochemistry, 2015, 71, 487-496.	3.0	4
68	Opipramol Inhibits Lipolysis in Human Adipocytes without Altering Glucose Uptake and Differently from Antipsychotic and Antidepressant Drugs with Adverse Effects on Body Weight Control. Pharmaceuticals, 2020, 13, 41.	3.8	4
69	High doses of catecholamines activate glucose transport in human adipocytes independently from adrenoceptor stimulation or vanadium addition. World Journal of Diabetes, 2022, 13, 37-53.	3.5	4
70	Comparative effects of idazoxan, efaroxan, and BU 224 on insulin secretion in the rabbit: Not only interaction with pancreatic imidazoline I2 binding sites. Health, 2010, 02, 112-123.	0.3	3
71	Short-term effects of obestatin on hexose uptake and triacylglycerol breakdown in human subcutaneous adipocytes. World Journal of Diabetes, 2018, 9, 25-32.	3.5	2
72	Hypercholesterolemia of obese mice with deletion of vascular adhesion protein-1 occurs without other atherosclerosis risk factor. , 0, , .		2

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73	Oral Supplementation with Benzylamine Delays the Onset of Diabetes in Obese and Diabetic db-/- Mice. Nutrients, 2021, 13, 2622.	4.1	2
74	Tyramine activates lipid accumulation in rat adipocytes: influences of in vitro and in vivo administration. AIMS Molecular Science, 2017, 4, 339-351.	0.5	2
75	Increased monoamine oxidase activity and imidazoline binding sites in insulin-resistant adipocytes from obese Zucker rats. World Journal of Biological Chemistry, 2022, 13, 15-34.	4.3	2
76	Engineering and Biomedical Effects of Commercial Juices of Berries, Cherries, and Pomegranates With High Polyphenol Content., 2019,, 259-283.		1
77	Editorial Special Issue: 2020 consortium for trans-pyrenean investigations on obesity and diabetes. Journal of Physiology and Biochemistry, 2022, , 1.	3.0	0