

Maria C Linder

List of Publications by Year in descending order

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91
papers

4,583
citations

117453

34
h-index

106150

65
g-index

92
all docs

92
docs citations

92
times ranked

4237
citing authors

#	ARTICLE	IF	CITATIONS
1	Secretion and uptake of copper via a small copper carrier in blood fluid. <i>Metallomics</i> , 2022, 14, .	1.0	10
2	Apoceruloplasmin: Abundance, Detection, Formation, and Metabolism. <i>Biomedicines</i> , 2021, 9, 233.	1.4	13
3	Intestinal uptake of Cu by processes that do not involve copper transporter 1 (CTR1), as determined with Caco2 cell monolayers. <i>FASEB Journal</i> , 2021, 35, .	0.2	0
4	A small copper carrier in blood plasma: purification, characterization, and metabolism. <i>FASEB Journal</i> , 2021, 35, .	0.2	0
5	Copper Homeostasis in Mammals, with Emphasis on Secretion and Excretion. A Review. <i>International Journal of Molecular Sciences</i> , 2020, 21, 4932.	1.8	53
6	Comparison of Alpha ₂ -Macroglobulins from Swine and Humans and their Copper Binding. <i>FASEB Journal</i> , 2019, 33, 825.3.	0.2	1
7	Mobilization of iron from ferritin: new steps and details. <i>Metallomics</i> , 2018, 10, 154-168.	1.0	40
8	Mechanism of Copper Uptake from Blood Plasma Ceruloplasmin by Mammalian Cells. <i>PLoS ONE</i> , 2016, 11, e0149516.	1.1	87
9	Ceruloplasmin and other copper binding components of blood plasma and their functions: an update. <i>Metallomics</i> , 2016, 8, 887-905.	1.0	194
10	Relation of Copper Toxicosis in Dogs and Wilson Disease to the Appearance of a Small Copper Carrier (SCC) in Blood Plasma and Urine. <i>FASEB Journal</i> , 2015, 29, 921.2.	0.2	1
11	Mechanism of Ceruloplasmin-Copper Uptake by, and Overexpression in, Mammalian Cells. <i>FASEB Journal</i> , 2015, 29, 1011.4.	0.2	0
12	Mechanism of Copper Absorption Investigated in Caco2 and HuTu80 Enterocyte Models. <i>FASEB Journal</i> , 2015, 29, 1011.3.	0.2	0
13	Mechanisms of Ferritin Iron Mobilization in Macrophages and Hepatocytes. <i>FASEB Journal</i> , 2015, 29, 1011.2.	0.2	0
14	Effects of ATP7A overexpression in mice on copper transport and metabolism in lactation and gestation. <i>Physiological Reports</i> , 2014, 2, e00195.	0.7	9
15	Mobilization of Stored Iron in Mammals: A Review. <i>Nutrients</i> , 2013, 5, 4022-4050.	1.7	104
16	Uptake of copper from plasma proteins in cells where expression of CTR1 has been modulated. <i>BioMetals</i> , 2012, 25, 697-709.	1.8	38
17	The relationship of copper to DNA damage and damage prevention in humans. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2012, 733, 83-91.	0.4	101
18	Lack of ceruloplasmin expression alters aspects of copper transport to the fetus and newborn, as determined in mice. <i>BioMetals</i> , 2012, 25, 373-382.	1.8	17

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19	Direct uptake of copper from ceruloplasmin by cells with and without CTR1, and potential involvement of a copper reductase. <i>FASEB Journal</i> , 2012, 26, 1112.1.	0.2	0
20	Release of iron from cellular iron stores in lysosomes: Potential involvement of divalent metal transporter 1 (DMT1) and common metabolites. <i>FASEB Journal</i> , 2012, 26, 1112.6.	0.2	1
21	Copper transporter 1 (CTR1) and the mechanisms of Cu uptake from blood plasma proteins by mammalian cells. <i>FASEB Journal</i> , 2012, 26, 641.19.	0.2	1
22	Trace metal dyshomeostasis is associated with loss of TRPML1 ion channel function. <i>FASEB Journal</i> , 2010, 24, 708.3.	0.2	0
23	The 10-fold increase in basal copper uptake by mammary gland in lactation is not induced by lactational hormones. <i>FASEB Journal</i> , 2010, 24, 719.3.	0.2	0
24	Common metabolites and reducing/chelating agents do not mobilize iron from intracellular ferritin but can (like lysosomal extracts) dissolve ferritin iron mineral. <i>FASEB Journal</i> , 2010, 24, 717.17.	0.2	0
25	DMT1 is not essential for Cu uptake from blood by internal organs. <i>FASEB Journal</i> , 2010, 24, 229.6.	0.2	0
26	Excess ATP7A reduces the copper content of the mammary gland in pregnancy and lactation but does not alter levels of plasma ceruloplasmin. <i>FASEB Journal</i> , 2010, 24, 719.4.	0.2	0
27	Copper proteins and ferroxidases in human plasma and that of wild-type and ceruloplasmin knockout mice. <i>Biochemical Journal</i> , 2009, 419, 237-245.	1.7	32
28	Forms of copper secreted by human mammary epithelial cells in response to lactational hormones. <i>FASEB Journal</i> , 2009, 23, 727.4.	0.2	0
29	Mechanisms of iron release from lysosomes. <i>FASEB Journal</i> , 2009, 23, 921.11.	0.2	0
30	Potential abnormalities in iron metabolism in hyperlipidemia patient fibroblasts. <i>FASEB Journal</i> , 2009, 23, 105.4.	0.2	0
31	Characterizing the chelatable/labile iron pool in mammalian cells. <i>FASEB Journal</i> , 2009, 23, 921.10.	0.2	0
32	Copper distribution and incorporation into ceruloplasmin and other plasma proteins in mice overexpressing ATP7A. <i>FASEB Journal</i> , 2009, 23, 727.5.	0.2	0
33	Copper binding components of blood plasma and organs, and their responses to influx of large doses of ⁶⁵ Cu, in the mouse. <i>BioMetals</i> , 2008, 21, 525-543.	1.8	47
34	Copper transport during lactation in transgenic mice expressing the human ATP7A protein. <i>Biochemical and Biophysical Research Communications</i> , 2008, 372, 613-617.	1.0	19
35	ATP7B Expression in Human Breast Epithelial Cells Is Mediated by Lactational Hormones. <i>Journal of Histochemistry and Cytochemistry</i> , 2008, 56, 389-399.	1.3	41
36	Copper is taken up efficiently from albumin and α_2 -macroglobulin by cultured human cells by more than one mechanism. <i>American Journal of Physiology - Cell Physiology</i> , 2008, 295, C708-C721.	2.1	86

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37	Copper is taken up efficiently from albumin and alpha α 2 α macroglobulin by cultured human cells by more than one mechanism. FASEB Journal, 2008, 22, 443.3.	0.2	0
38	Lactational hormones increase expression of milk proteins but do not enhance uptake of copper by mammary epithelial cells. FASEB Journal, 2008, 22, 1192.1.	0.2	0
39	Copper secretion from human breast epithelial cells is mediated by ATP7B and lactational hormones. FASEB Journal, 2008, 22, 443.6.	0.2	0
40	DMT1 is not involved in uptake of copper from the blood plasma by hepatic and mammary epithelial cells. FASEB Journal, 2008, 22, 692.9.	0.2	0
41	Quantitation and characterization of low molecular weight iron complexes in mammalian cells in conditions of iron overload and deficiency that may comprise components of the "labile iron pool" TM . FASEB Journal, 2008, 22, 1057.10.	0.2	0
42	Transcuprein is a macroglobulin regulated by copper and iron availability. Journal of Nutritional Biochemistry, 2007, 18, 597-608.	1.9	73
43	Unknown low molecular weight iron-peptide complexes and their possible contribution to the "labile iron pool" in mammalian cells. FASEB Journal, 2007, 21, A1117.	0.2	0
44	Copper is taken up efficiently from plasma albumin by polarized mammary epithelial cell monolayers. FASEB Journal, 2007, 21, A723.	0.2	0
45	How ferritin releases its iron through entry into lysosomes and proteolysis, and a potential role for DMT1. FASEB Journal, 2007, 21, A163.	0.2	0
46	Vesicular transport of Fe and interaction with other metal ions in polarized Caco2 Cell monolayers. Biological Research, 2006, 39, 143-56.	1.5	18
47	Release of iron from ferritin requires lysosomal activity. American Journal of Physiology - Cell Physiology, 2006, 291, C445-C455.	2.1	221
48	Vesicular transport and apotransferrin in intestinal iron absorption, as shown in the Caco-2 cell model. American Journal of Physiology - Renal Physiology, 2006, 290, G301-G309.	1.6	32
49	Transcupreins are serum copper-transporters of the macroglobulin family, and may be regulated by iron and copper. FASEB Journal, 2006, 20, A553.	0.2	1
50	Iron and copper homeostasis and intestinal absorption using the Caco2 cell model. BioMetals, 2003, 16, 145-160.	1.8	38
51	Regulation of copper absorption by copper availability in the Caco-2 cell intestinal model. American Journal of Physiology - Renal Physiology, 2003, 284, G739-G747.	1.6	18
52	Copper transport to mammary gland and milk during lactation in rats. American Journal of Physiology - Endocrinology and Metabolism, 2002, 283, E667-E675.	1.8	41
53	Dietary Iron Status Has Little Effect on Expression of Ceruloplasmin but Alters That of Ferritin in Rats. Journal of Nutrition, 2002, 132, 351-356.	1.3	13
54	Effects of copper and ceruloplasmin on iron transport in the Caco 2 cell intestinal model. Journal of Nutritional Biochemistry, 2002, 13, 138-148.	1.9	38

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55	Copper and genomic stability in mammals. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2001, 475, 141-152.	0.4	141
56	Iron Prevents Ferritin Turnover in Hepatic Cells. Journal of Biological Chemistry, 2001, 276, 48775-48780.	1.6	45
57	Milk Ceruloplasmin and Its Expression by Mammary Gland and Liver in Pigs. Archives of Biochemistry and Biophysics, 2000, 373, 451-461.	1.4	34
58	Copper Transport in Mammals. Advances in Experimental Medicine and Biology, 1999, 448, 1-16.	0.8	34
59	Measurement of acute phase proteins in the rat brain: contribution of vascular contents. Neurochemical Research, 1999, 24, 1313-1317.	1.6	8
60	Copper Transport and Ceruloplasmin During Lactation and Pregnancy. , 1999, , 117-129.		1
61	Ran-2, a glial lineage marker, is a GPI-anchored form of ceruloplasmin. Journal of Neuroscience Research, 1998, 54, 147-157.	1.3	63
62	Copper transport. American Journal of Clinical Nutrition, 1998, 67, 965S-971S.	2.2	251
63	Secretion of Ferritin by Rat Hepatoma Cells and Its Regulation by Inflammatory Cytokines and Iron. Blood, 1997, 90, 4979-4986.	0.6	208
64	Secretion of ferritin by rat hepatoma cells and its regulation by inflammatory cytokines and iron. Blood, 1997, 90, 4979-86.	0.6	78
65	Serum ferritin: Does it differ from tissue ferritin?. Journal of Gastroenterology and Hepatology (Australia), 1996, 11, 1033-1036.	1.4	42
66	Copper biochemistry and molecular biology. American Journal of Clinical Nutrition, 1996, 63, 797S-811S.	2.2	838
67	Copper transport in the Nagase analbuminemic rat. American Journal of Physiology - Renal Physiology, 1994, 267, G259-G269.	1.6	14
68	Synthesis and Turnover of Ceruloplasmin in Rats Treated with 17 β -Estradiol. Archives of Biochemistry and Biophysics, 1993, 302, 362-368.	1.4	28
69	Ceruloplasmin and Copper Transport During the Latter Part of Gestation in the Rat. Experimental Biology and Medicine, 1993, 203, 428-439.	1.1	59
70	Comparison of Copper Binding Components in Dog Serum with Those in Other Species. Experimental Biology and Medicine, 1992, 200, 321-329.	1.1	22
71	Ferritin synthesis on polyribosomes attached to the endoplasmic reticulum. Journal of Inorganic Biochemistry, 1992, 47, 229-240.	1.5	11
72	Biochemistry of Copper. , 1991, , .		303

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73	Dissociation of ferritins. Archives of Biochemistry and Biophysics, 1989, 269, 485-496.	1.4	24
74	Heart tissue contains small and large aggregates of ferritin subunits. Archives of Biochemistry and Biophysics, 1989, 273, 34-41.	1.4	13
75	Ferritin mRNA is found on bound as well as on free polyribosomes in rat heart. Archives of Biochemistry and Biophysics, 1989, 273, 89-98.	1.4	14
76	Binding and uptake of copper from ceruloplasmin. Biochemical and Biophysical Research Communications, 1986, 139, 822-829.	1.0	62
77	Turnover and excretion of copper in rats as measured with ⁶⁷ Cu. American Journal of Physiology - Endocrinology and Metabolism, 1986, 251, E551-E555.	1.8	7
78	Copper transport in rats involving a new plasma protein. American Journal of Physiology - Endocrinology and Metabolism, 1985, 249, E77-E88.	1.8	47
79	Distribution of copper among components of human serum. Journal of the National Cancer Institute, 1985, 75, 277-84.	3.0	68
80	Interactions of pH and Ascorbate in Intestinal Iron Absorption. Journal of Nutrition, 1983, 113, 2615-2622.	1.3	37
81	Concentration, Structure and Iron Saturation of Ferritins from Normal Human Lung and Lung Tumors with Graded Histopathology. Enzyme, 1982, 27, 189-198.	0.7	9
82	Circulating ceruloplasmin is an important source of copper for normal and malignant animal cells. Biochimica Et Biophysica Acta - General Subjects, 1981, 678, 27-38.	1.1	72
83	The size and shape of heart and muscle ferritins analyzed by sedimentation, gel filtration, and electrophoresis. Journal of Biological Chemistry, 1981, 256, 9104-10.	1.6	31
84	Copper Regulation of Ceruloplasmin in Copper-Deficient Rats. Enzyme, 1979, 24, 23-35.	0.7	72
85	Ferritin: structure, biosynthesis, and role in iron metabolism.. Physiological Reviews, 1978, 58, 317-396.	13.1	385
86	Sex difference in distribution and iron responsiveness of the two ferritins of rat cardiac and skeletal muscle. Biochimica Et Biophysica Acta - General Subjects, 1977, 497, 280-287.	1.1	13
87	Structural features of rat cardiac ferritins. Biochimica Et Biophysica Acta (BBA) - Protein Structure, 1977, 491, 67-75.	1.7	19
88	Structural differences in ferritins from normal and malignant rat tissues. Biochimica Et Biophysica Acta (BBA) - Protein Structure, 1975, 386, 409-421.	1.7	31
89	Size and charge heterogeneity of rat tissue ferritins. Biochimica Et Biophysica Acta (BBA) - Protein Structure, 1975, 412, 148-156.	1.7	26
90	Ferritin and intestinal iron absorption: pancreatic enzymes and free iron. American Journal of Physiology, 1975, 228, 196-204.	5.0	41

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91	Metabolic and chemical features of ferritins, a series of iron-inducible tissue proteins. American Journal of Pathology, 1973, 72, 263-82.	1.9	42