

Anthony H Futerman

List of Publications by Year in descending order

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233
papers

16,026
citations

14614

66
h-index

19690

117
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240
all docs

240
docs citations

240
times ranked

14165
citing authors

#	ARTICLE	IF	CITATIONS
1	Silencing of ceramide synthase 2 in hepatocytes modulates plasma ceramide biomarkers predictive of cardiovascular death. <i>Molecular Therapy</i> , 2022, 30, 1661-1674.	3.7	9
2	GBA mutations, glucosylceramide and Parkinson's disease. <i>Current Opinion in Neurobiology</i> , 2022, 72, 148-154.	2.0	23
3	16pdel lipid changes in iPSC-derived neurons and function of FAM57B in lipid metabolism and synaptogenesis. <i>IScience</i> , 2022, 25, 103551.	1.9	8
4	Laurdan in live cell imaging: Effect of acquisition settings, cell culture conditions and data analysis on generalized polarization measurements. <i>Journal of Photochemistry and Photobiology B: Biology</i> , 2022, 228, 112404.	1.7	8
5	Dependence of ABCB1 transporter expression and function on distinct sphingolipids generated by ceramide synthases-2 and -6 in chemoresistant renal cancer. <i>Journal of Biological Chemistry</i> , 2022, 298, 101492.	1.6	6
6	A novel C-terminal DxRSDxE motif in ceramide synthases involved in dimer formation. <i>Journal of Biological Chemistry</i> , 2022, 298, 101517.	1.6	12
7	Fatty acid transport protein 2 interacts with ceramide synthase 2 to promote ceramide synthesis. <i>Journal of Biological Chemistry</i> , 2022, 298, 101735.	1.6	9
8	A Storm in a Primordial Teacup. <i>Inference</i> , 2022, 7, .	0.0	0
9	Generation of a ceramide synthase 6 mouse lacking the DDRSDIE C-terminal motif. <i>PLoS ONE</i> , 2022, 17, e0271675.	1.1	0
10	Brain pathology and cerebellar purkinje cell loss in a mouse model of chronic neuronopathic Gaucher disease. <i>Progress in Neurobiology</i> , 2021, 197, 101939.	2.8	6
11	Substrate reduction therapy using Genzâ€667161 reduces levels of pathogenic components in a mouse model of neuronopathic forms of Gaucher disease. <i>Journal of Neurochemistry</i> , 2021, 156, 692-701.	2.1	16
12	Ceramide Synthase 2 Null Mice Are Protected from Ovalbumin-Induced Asthma with Higher T Cell Receptor Signal Strength in CD4+ T Cells. <i>International Journal of Molecular Sciences</i> , 2021, 22, 2713.	1.8	11
13	The fineâ€tuning of cell membrane lipid bilayers accentuates their compositional complexity. <i>BioEssays</i> , 2021, 43, e2100021.	1.2	15
14	Ceramide synthases: Reflections on the impact of Dr. Lina M. Obeid. <i>Cellular Signalling</i> , 2021, 82, 109958.	1.7	19
15	Biophysical impact of sphingosine and other abnormal lipid accumulation in Niemann-Pick disease type C cell models. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2021, 1866, 158944.	1.2	1
16	The role of the â€˜sphingoid motifâ€™™ in shaping the molecular interactions of sphingolipids in biomembranes. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2021, 1863, 183701.	1.4	10
17	The Complex Tail of Circulating Sphingolipids in Atherosclerosis and Cardiovascular Disease. <i>Journal of Lipid and Atherosclerosis</i> , 2021, 10, 268.	1.1	10
18	Sphingolipids. , 2021, , 281-316.		5

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19	Ceramide synthase 2 deletion decreases the infectivity of HIV-1. <i>Journal of Biological Chemistry</i> , 2021, 296, 100340.	1.6	15
20	Proteomics analysis of a human brain sample from a mucopolipidosis type IV patient reveals pathophysiological pathways. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 39.	1.2	11
21	The role of ceramide in regulating endoplasmic reticulum function. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2020, 1865, 158489.	1.2	29
22	Lysosomal Storage Disorders Shed Light on Lysosomal Dysfunction in Parkinson's Disease. <i>International Journal of Molecular Sciences</i> , 2020, 21, 4966.	1.8	21
23	Different rates of flux through the biosynthetic pathway for long-chain versus very-long-chain sphingolipids. <i>Journal of Lipid Research</i> , 2020, 61, 1341-1346.	2.0	9
24	Mice defective in interferon signaling help distinguish between primary and secondary pathological pathways in a mouse model of neuronal forms of Gaucher disease. <i>Journal of Neuroinflammation</i> , 2020, 17, 265.	3.1	10
25	Innate immune response in neuronopathic forms of Gaucher disease confers resistance against viral-induced encephalitis. <i>Acta Neuropathologica Communications</i> , 2020, 8, 144.	2.4	8
26	Integrin Alpha E (CD103) Limits Virus-Induced IFN- β Production in Conventional Dendritic Cells. <i>Frontiers in Immunology</i> , 2020, 11, 607889.	2.2	1
27	The Lysosome and Nonmotor Symptoms: Linking Parkinson's Disease and Lysosomal Storage Disorders. <i>Movement Disorders</i> , 2020, 35, 2150-2155.	2.2	5
28	Absence of infiltrating peripheral myeloid cells in the brains of mouse models of lysosomal storage disorders. <i>Journal of Neurochemistry</i> , 2019, 148, 625-638.	2.1	20
29	Hepatic triglyceride accumulation via endoplasmic reticulum stress-induced SREBP-1 activation is regulated by ceramide synthases. <i>Experimental and Molecular Medicine</i> , 2019, 51, 1-16.	3.2	51
30	Yeast ceramide synthases, Lag1 and Lac1, have distinct substrate specificity. <i>Journal of Cell Science</i> , 2019, 132, .	1.2	26
31	TLR9-mediated dendritic cell activation uncovers mammalian ganglioside species with specific ceramide backbones that activate invariant natural killer T cells. <i>PLoS Biology</i> , 2019, 17, e3000169.	2.6	24
32	Ablation of the pro-inflammatory master regulator miR-155 does not mitigate neuroinflammation or neurodegeneration in a vertebrate model of Gaucher's disease. <i>Neurobiology of Disease</i> , 2019, 127, 563-569.	2.1	19
33	<i>In vivo</i> inactivation of glycosidases by conurititol B epoxide and cyclophellitol as revealed by activity-based protein profiling. <i>FEBS Journal</i> , 2019, 286, 584-600.	2.2	44
34	A Stroll Down the CerS Lane. <i>Advances in Experimental Medicine and Biology</i> , 2019, 1159, 49-63.	0.8	32
35	The Cell Biology of SARS-CoV-2. <i>Inference</i> , 2019, 5, .	0.0	4
36	Eleven residues determine the acyl chain specificity of ceramide synthases. <i>Journal of Biological Chemistry</i> , 2018, 293, 9912-9921.	1.6	50

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37	The brain lipidome in neurodegenerative lysosomal storage disorders. <i>Biochemical and Biophysical Research Communications</i> , 2018, 504, 623-628.	1.0	20
38	Sphingolipid regulation of lung epithelial cell mitophagy and necroptosis during cigarette smoke exposure. <i>FASEB Journal</i> , 2018, 32, 1880-1890.	0.2	59
39	Fingolimod phosphate inhibits astrocyte inflammatory activity in mucopolipidosis IV. <i>Human Molecular Genetics</i> , 2018, 27, 2725-2738.	1.4	22
40	Altered lysosome distribution is an early neuropathological event in neurological forms of Gaucher disease. <i>FEBS Letters</i> , 2017, 591, 774-783.	1.3	20
41	Jaspine B induces nonapoptotic cell death in gastric cancer cells independently of its inhibition of ceramide synthase. <i>Journal of Lipid Research</i> , 2017, 58, 1500-1513.	2.0	18
42	Signalome-wide RNAi screen identifies GBA1 as a positive mediator of autophagic cell death. <i>Cell Death and Differentiation</i> , 2017, 24, 1288-1302.	5.0	67
43	Regulation of very-long acyl chain ceramide synthesis by acyl-CoA-binding protein. <i>Journal of Biological Chemistry</i> , 2017, 292, 7588-7597.	1.6	35
44	Identification of a feedback loop involving β -glucosidase 2 and its product sphingosine sheds light on the molecular mechanisms in Gaucher disease. <i>Journal of Biological Chemistry</i> , 2017, 292, 6177-6189.	1.6	22
45	Oxidative stress elicited by modifying the ceramide acyl chain length reduces the rate of clathrin-mediated endocytosis. <i>Journal of Cell Science</i> , 2017, 130, 1486-1493.	1.2	15
46	Pathological levels of glucosylceramide change the biophysical properties of artificial and cell membranes. <i>Physical Chemistry Chemical Physics</i> , 2017, 19, 340-346.	1.3	28
47	The metabolism of glucocerebrosides " From 1965 to the present. <i>Molecular Genetics and Metabolism</i> , 2017, 120, 22-26.	0.5	28
48	Ablation of ceramide synthase 2 exacerbates dextran sodium sulphate-induced colitis in mice due to increased intestinal permeability. <i>Journal of Cellular and Molecular Medicine</i> , 2017, 21, 3565-3578.	1.6	22
49	Combining Deep Sequencing, Proteomics, Phosphoproteomics, and Functional Screens To Discover Novel Regulators of Sphingolipid Homeostasis. <i>Journal of Proteome Research</i> , 2017, 16, 571-582.	1.8	11
50	Sortilin Deficiency Reduces Ductular Reaction, Hepatocyte Apoptosis, and Liver Fibrosis in Cholestatic-Induced Liver Injury. <i>American Journal of Pathology</i> , 2017, 187, 122-133.	1.9	24
51	Clozapine Modulates Glucosylceramide, Clears Aggregated Proteins, and Enhances ATG8/LC3 in <i>Caenorhabditis elegans</i> . <i>Neuropsychopharmacology</i> , 2017, 42, 951-962.	2.8	9
52	Critical Role for Very-Long Chain Sphingolipids in Invariant Natural Killer T Cell Development and Homeostasis. <i>Frontiers in Immunology</i> , 2017, 8, 1386.	2.2	19
53	Induction of the type I interferon response in neurological forms of Gaucher disease. <i>Journal of Neuroinflammation</i> , 2016, 13, 104.	3.1	53
54	Delineating pathological pathways in a chemically induced mouse model of Gaucher disease. <i>Journal of Pathology</i> , 2016, 239, 496-509.	2.1	54

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55	Effect of the sphingosine kinase 1 selective inhibitor, PF-543 on arterial and cardiac remodelling in a hypoxic model of pulmonary arterial hypertension. <i>Cellular Signalling</i> , 2016, 28, 946-955.	1.7	37
56	Identification of Modifier Genes in a Mouse Model of Gaucher Disease. <i>Cell Reports</i> , 2016, 16, 2546-2553.	2.9	52
57	Making Sense of the Yeast Sphingolipid Pathway. <i>Journal of Molecular Biology</i> , 2016, 428, 4765-4775.	2.0	41
58	Perspective: Finding common ground. <i>Nature</i> , 2016, 537, S160-S161.	13.7	14
59	K _{Ca} 3.1 upregulation preserves endothelium-dependent vasorelaxation during aging and oxidative stress. <i>Aging Cell</i> , 2016, 15, 801-810.	3.0	15
60	Glucosylceramide Reorganizes Cholesterol-Containing Domains in a Fluid Phospholipid Membrane. <i>Biophysical Journal</i> , 2016, 110, 612-622.	0.2	24
61	Ceramide synthases in biomedical research. <i>Chemistry and Physics of Lipids</i> , 2016, 197, 25-32.	1.5	45
62	Sphingolipids. , 2016, , 297-326.		8
63	Altering sphingolipid composition with aging induces contractile dysfunction of gastric smooth muscle via $K_{Ca}1.1$ upregulation. <i>Aging Cell</i> , 2015, 14, 982-994.	3.0	22
64	Identification of a Biomarker in Cerebrospinal Fluid for Neuronopathic Forms of Gaucher Disease. <i>PLoS ONE</i> , 2015, 10, e0120194.	1.1	53
65	A rapid ceramide synthase activity using NBD-sphinganine and solid phase extraction. <i>Journal of Lipid Research</i> , 2015, 56, 193-199.	2.0	22
66	Development of pheochromocytoma in ceramide synthase 2 null mice. <i>Endocrine-Related Cancer</i> , 2015, 22, 623-632.	1.6	27
67	LPS-mediated septic shock is augmented in ceramide synthase 2 null mice due to elevated activity of TNF- α -converting enzyme. <i>FEBS Letters</i> , 2015, 589, 2213-2217.	1.3	27
68	Innate immune responses in the brain of sphingolipid lysosomal storage diseases. <i>Biological Chemistry</i> , 2015, 396, 659-667.	1.2	23
69	Preface to the Special Issue on brain lipids. <i>Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids</i> , 2015, 1851, 997-998.	1.2	1
70	Lack of ceramide synthase 2 suppresses the development of experimental autoimmune encephalomyelitis by impairing the migratory capacity of neutrophils. <i>Brain, Behavior, and Immunity</i> , 2015, 46, 280-292.	2.0	53
71	Sortilin deficiency improves the metabolic phenotype and reduces hepatic steatosis of mice subjected to diet-induced obesity. <i>Journal of Hepatology</i> , 2015, 62, 175-181.	1.8	59
72	Emerging therapeutic targets for Gaucher disease. <i>Expert Opinion on Therapeutic Targets</i> , 2015, 19, 321-334.	1.5	18

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73	Bcl2L13 is a ceramide synthase inhibitor in glioblastoma. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 5682-5687.	3.3	86
74	The HIV-1 Envelope Transmembrane Domain Binds TLR2 through a Distinct Dimerization Motif and Inhibits TLR2-Mediated Responses. PLoS Pathogens, 2014, 10, e1004248.	2.1	33
75	Sphingoid long chain bases prevent lung infection by <i>Pseudomonas aeruginosa</i> . EMBO Molecular Medicine, 2014, 6, 1205-1214.	3.3	109
76	Reduced ceramide synthase 2 activity causes progressive myoclonic epilepsy. Annals of Clinical and Translational Neurology, 2014, 1, 88-98.	1.7	50
77	Ceramide synthases as potential targets for therapeutic intervention in human diseases. Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids, 2014, 1841, 671-681.	1.2	183
78	Neuronal accumulation of glucosylceramide in a mouse model of neuronopathic Gaucher disease leads to neurodegeneration. Human Molecular Genetics, 2014, 23, 843-854.	1.4	123
79	From sheep to mice to cells: Tools for the study of the sphingolipidoses. Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids, 2014, 1841, 1189-1199.	1.2	16
80	Hepatic fatty acid uptake is regulated by the sphingolipid acyl chain length. Biochimica Et Biophysica Acta - Molecular and Cell Biology of Lipids, 2014, 1841, 1754-1766.	1.2	50
81	RIPK3 as a potential therapeutic target for Gaucher's disease. Nature Medicine, 2014, 20, 204-208.	15.2	147
82	CerS2 Haploinsufficiency Inhibits β -Oxidation and Confers Susceptibility to Diet-Induced Steatohepatitis and Insulin Resistance. Cell Metabolism, 2014, 20, 687-695.	7.2	379
83	Influence of Intracellular Membrane pH on Sphingolipid Organization and Membrane Biophysical Properties. Langmuir, 2014, 30, 4094-4104.	1.6	12
84	A Dynamic Interface between Vacuoles and Mitochondria in Yeast. Developmental Cell, 2014, 30, 95-102.	3.1	321
85	Changes in membrane biophysical properties induced by sphingomyelinase depend on the sphingolipid N-acyl chain. Journal of Lipid Research, 2014, 55, 53-61.	2.0	51
86	A combined fluorescence spectroscopy, confocal and 2-photon microscopy approach to re-evaluate the properties of sphingolipid domains. Biochimica Et Biophysica Acta - Biomembranes, 2013, 1828, 2099-2110.	1.4	38
87	Neuronal Forms of Gaucher Disease. Handbook of Experimental Pharmacology, 2013, , 405-419.	0.9	45
88	The complexity of sphingolipid biosynthesis in the endoplasmic reticulum. Biochimica Et Biophysica Acta - Molecular Cell Research, 2013, 1833, 2511-2518.	1.9	136
89	Effect of glucosylceramide on the biophysical properties of fluid membranes. Biochimica Et Biophysica Acta - Biomembranes, 2013, 1828, 1122-1130.	1.4	32
90	Identification of <i>N</i> -acetyl-fumonisin B ₁ as new cytotoxic metabolites of fumonisin mycotoxins. Molecular Nutrition and Food Research, 2013, 57, 516-522.	1.5	38

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91	Ablation of very long acyl chain sphingolipids causes hepatic insulin resistance in mice due to altered detergent-resistant membranes. <i>Hepatology</i> , 2013, 57, 525-532.	3.6	140
92	Accumulation of Ordered Ceramide-Cholesterol Domains in Farber Disease Fibroblasts. <i>JIMD Reports</i> , 2013, 12, 71-77.	0.7	14
93	The Yeast P5 Type ATPase, Spf1, Regulates Manganese Transport into the Endoplasmic Reticulum. <i>PLoS ONE</i> , 2013, 8, e85519.	1.1	62
94	Impaired Epidermal Ceramide Synthesis Causes Autosomal Recessive Congenital Ichthyosis and Reveals the Importance of Ceramide Acyl Chain Length. <i>Journal of Investigative Dermatology</i> , 2013, 133, 2202-2211.	0.3	138
95	Myristate-derived d16:0 Sphingolipids Constitute a Cardiac Sphingolipid Pool with Distinct Synthetic Routes and Functional Properties. <i>Journal of Biological Chemistry</i> , 2013, 288, 13397-13409.	1.6	63
96	Protection of a Ceramide Synthase 2 Null Mouse from Drug-induced Liver Injury. <i>Journal of Biological Chemistry</i> , 2013, 288, 30904-30916.	1.6	35
97	Ablation of Ceramide Synthase 2 Causes Chronic Oxidative Stress Due to Disruption of the Mitochondrial Respiratory Chain. <i>Journal of Biological Chemistry</i> , 2013, 288, 4947-4956.	1.6	165
98	Ceramide Synthases Expression and Role of Ceramide Synthase-2 in the Lung: Insight from Human Lung Cells and Mouse Models. <i>PLoS ONE</i> , 2013, 8, e62968.	1.1	69
99	Oxidized Phospholipids Induce Ceramide Accumulation in RAW 264.7 Macrophages: Role of Ceramide Synthases. <i>PLoS ONE</i> , 2013, 8, e70002.	1.1	36
100	Lysosomal storage disorders: old diseases, present and future challenges. <i>Pediatric Endocrinology Reviews</i> , 2013, 11 Suppl 1, 59-63.	1.2	12
101	Acyl Chain Specificity of Ceramide Synthases Is Determined within a Region of 150 Residues in the Tram-Lag-CLN8 (TLC) Domain. <i>Journal of Biological Chemistry</i> , 2012, 287, 3197-3206.	1.6	60
102	Contribution of brain inflammation to neuronal cell death in neuronopathic forms of Gaucher's disease. <i>Brain</i> , 2012, 135, 1724-1735.	3.7	132
103	Modulation of Ceramide Synthase Activity via Dimerization. <i>Journal of Biological Chemistry</i> , 2012, 287, 21025-21033.	1.6	98
104	Limonoid Compounds Inhibit Sphingomyelin Biosynthesis by Preventing CERT Protein-dependent Extraction of Ceramides from the Endoplasmic Reticulum. <i>Journal of Biological Chemistry</i> , 2012, 287, 24397-24411.	1.6	29
105	Ablation of ceramide synthase 2 strongly affects biophysical properties of membranes. <i>Journal of Lipid Research</i> , 2012, 53, 430-436.	2.0	62
106	Ceramide Kinase-Like (CERKL) Interacts with Neuronal Calcium Sensor Proteins in the Retina in a Cation-Dependent Manner. , 2012, 53, 4565.		15
107	Methylation of glycosylated sphingolipid modulates membrane lipid topography and pathogenicity of <i>Cryptococcus neoformans</i> . <i>Cellular Microbiology</i> , 2012, 14, 500-516.	1.1	67
108	Comparison of a molecular dynamics model with the X-ray structure of the N370S acid- β -glucosidase mutant that causes Gaucher disease. <i>Protein Engineering, Design and Selection</i> , 2011, 24, 773-775.	1.0	13

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109	Self-Segregation of Myelin Membrane Lipids in Model Membranes. <i>Biophysical Journal</i> , 2011, 101, 2713-2720.	0.2	38
110	Animal models for Gaucher disease research. <i>DMM Disease Models and Mechanisms</i> , 2011, 4, 746-752.	1.2	80
111	Effect of ceramide structure on membrane biophysical properties: The role of acyl chain length and unsaturation. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , 2011, 1808, 2753-2760.	1.4	172
112	Intracellular localization of organized lipid domains of C16-ceramide/cholesterol. <i>Journal of Structural Biology</i> , 2011, 175, 21-30.	1.3	17
113	Cyclodextrin-mediated crystallization of acid β -glucosidase in complex with amphiphilic bicyclic nojirimycin analogues. <i>Organic and Biomolecular Chemistry</i> , 2011, 9, 4160.	1.5	31
114	Lysosomal storage disorders and Parkinson's disease: Gaucher disease and beyond. <i>Movement Disorders</i> , 2011, 26, 1593-1604.	2.2	141
115	Spatial and temporal correlation between neuron loss and neuroinflammation in a mouse model of neuronopathic Gaucher disease. <i>Human Molecular Genetics</i> , 2011, 20, 1375-1386.	1.4	93
116	Encephalopathy Caused by Ablation of Very Long Acyl Chain Ceramide Synthesis May Be Largely Due to Reduced Galactosylceramide Levels. <i>Journal of Biological Chemistry</i> , 2011, 286, 30022-30033.	1.6	71
117	The Role of the Ceramide Acyl Chain Length in Neurodegeneration: Involvement of Ceramide Synthases. <i>NeuroMolecular Medicine</i> , 2010, 12, 341-350.	1.8	128
118	Stress-induced ER to Golgi translocation of ceramide synthase 1 is dependent on proteasomal processing. <i>Experimental Cell Research</i> , 2010, 316, 78-91.	1.2	39
119	Mammalian ceramide synthases. <i>IUBMB Life</i> , 2010, 62, 347-356.	1.5	377
120	The plasma membrane code. <i>Nature Chemical Biology</i> , 2010, 6, 487-488.	3.9	4
121	A Critical Role for Ceramide Synthase 2 in Liver Homeostasis. <i>Journal of Biological Chemistry</i> , 2010, 285, 10911-10923.	1.6	200
122	Altered expression and distribution of cathepsins in neuronopathic forms of Gaucher disease and in other sphingolipidoses. <i>Human Molecular Genetics</i> , 2010, 19, 3583-3590.	1.4	76
123	A Critical Role for Ceramide Synthase 2 in Liver Homeostasis. <i>Journal of Biological Chemistry</i> , 2010, 285, 10902-10910.	1.6	213
124	Cellular pathogenesis in sphingolipid storage disorders: the quest for new therapeutic approaches. <i>Clinical Lipidology</i> , 2010, 5, 255-265.	0.4	3
125	Molecular Basis of Reduced Glucosylceramidase Activity in the Most Common Gaucher Disease Mutant, N370S. <i>Journal of Biological Chemistry</i> , 2010, 285, 42105-42114.	1.6	31
126	Common and Uncommon Pathogenic Cascades in Lysosomal Storage Diseases. <i>Journal of Biological Chemistry</i> , 2010, 285, 20423-20427.	1.6	298

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127	Increased ceramide synthase 2 and 6 mRNA levels in breast cancer tissues and correlation with sphingosine kinase expression. <i>Biochemical and Biophysical Research Communications</i> , 2010, 391, 219-223.	1.0	70
128	Characterization of gene-activated human acid- β -glucosidase: Crystal structure, glycan composition, and internalization into macrophages. <i>Glycobiology</i> , 2010, 20, 24-32.	1.3	113
129	Ceramide Synthases: Roles in Cell Physiology and Signaling. <i>Advances in Experimental Medicine and Biology</i> , 2010, 688, 60-71.	0.8	142
130	Ceramide Synthesis Is Modulated by the Sphingosine Analog FTY720 via a Mixture of Uncompetitive and Noncompetitive Inhibition in an Acyl-CoA Chain Length-dependent Manner. <i>Journal of Biological Chemistry</i> , 2009, 284, 16090-16098.	1.6	108
131	No evidence for activation of the unfolded protein response in neuronopathic models of Gaucher disease. <i>Human Molecular Genetics</i> , 2009, 18, 1482-1488.	1.4	52
132	Ceramide synthase 1 is regulated by proteasomal mediated turnover. <i>Biochimica Et Biophysica Acta - Molecular Cell Research</i> , 2009, 1793, 1218-1227.	1.9	46
133	6- <i>N</i> -deoxy-5,6-dia- <i>N</i> -octyliminomethylidene)nojirimycin: Synthesis, Biological Evaluation, and Crystal Structure in Complex with Acid β -Glucosidase. <i>ChemBioChem</i> , 2009, 10, 1480-1485.	1.3	44
134	Impaired IL-10 transcription and release in animal models of Gaucher disease macrophages. <i>Blood Cells, Molecules, and Diseases</i> , 2009, 43, 134-137.	0.6	12
135	Lipid Raft Composition Modulates Sphingomyelinase Activity and Ceramide-Induced Membrane Physical Alterations. <i>Biophysical Journal</i> , 2009, 96, 3210-3222.	0.2	87
136	De Novo Ceramide Synthesis Is Required for N-Linked Glycosylation in Plasma Cells. <i>Journal of Immunology</i> , 2009, 182, 7038-7047.	0.4	20
137	An exposed carboxyl group on sialic acid is essential for gangliosides to inhibit calcium uptake via the sarco/endoplasmic reticulum Ca^{2+} -ATPase: relevance to gangliosidosis. <i>Journal of Neurochemistry</i> , 2008, 104, 140-146.	2.1	22
138	Control of the rate of evaporation in protein crystallization by the 'microbatch under oil' method. <i>Journal of Applied Crystallography</i> , 2008, 41, 969-971.	1.9	15
139	Acid β -glucosidase: insights from structural analysis and relevance to Gaucher disease therapy. <i>Biological Chemistry</i> , 2008, 389, 1361-1369.	1.2	43
140	Characterization of Ceramide Synthase 2. <i>Journal of Biological Chemistry</i> , 2008, 283, 5677-5684.	1.6	410
141	Regulation of (dihydro) ceramide synthase 1. <i>FASEB Journal</i> , 2008, 22, 299-299.	0.2	1
142	Crystal Structures of Complexes of N-Butyl- and N-Nonyl-Deoxynojirimycin Bound to Acid β -Glucosidase. <i>Journal of Biological Chemistry</i> , 2007, 282, 29052-29058.	1.6	109
143	A New Functional Motif in Hox Domain-containing Ceramide Synthases. <i>Journal of Biological Chemistry</i> , 2007, 282, 27366-27373.	1.6	58
144	(Dihydro)ceramide Synthase 1 Regulated Sensitivity to Cisplatin Is Associated with the Activation of p38 Mitogen-Activated Protein Kinase and Is Abrogated by Sphingosine Kinase 1. <i>Molecular Cancer Research</i> , 2007, 5, 801-812.	1.5	104

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145	Changes in macrophage morphology in a Gaucher disease model are dependent on CTP:phosphocholine cytidyltransferase β . <i>Blood Cells, Molecules, and Diseases</i> , 2007, 39, 124-129.	0.6	7
146	Kinetic characterization of mammalian ceramide synthases: Determination of K_m values towards sphinganine. <i>FEBS Letters</i> , 2007, 581, 5289-5294.	1.3	73
147	Antibody Labeling of Cholesterol/Ceramide Ordered Domains in Cell Membranes. <i>ChemBioChem</i> , 2007, 8, 2286-2294.	1.3	15
148	Taxi service for lipids. <i>Nature</i> , 2007, 449, 35-37.	13.7	6
149	Production of glucocerebrosidase with terminal mannose glycans for enzyme replacement therapy of Gaucher's disease using a plant cell system. <i>Plant Biotechnology Journal</i> , 2007, 5, 579-590.	4.1	371
150	The metabolism and function of sphingolipids and glycosphingolipids. <i>Cellular and Molecular Life Sciences</i> , 2007, 64, 2270-2284.	2.4	291
151	Genetic diseases of sphingolipid metabolism: Pathological mechanisms and therapeutic options. <i>FEBS Letters</i> , 2006, 580, 5510-5517.	1.3	38
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