

Jerome Ausseil

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3860780/publications.pdf>

Version: 2024-02-01

55
papers

1,301
citations

430874

18
h-index

377865

34
g-index

64
all docs

64
docs citations

64
times ranked

1564
citing authors

#	ARTICLE	IF	CITATIONS
1	Extracellular Vesicles From LPS-Treated Macrophages Aggravate Smooth Muscle Cell Calcification by Propagating Inflammation and Oxidative Stress. <i>Frontiers in Cell and Developmental Biology</i> , 2022, 10, 823450.	3.7	10
2	Synthesis of new sulfated disaccharides for the modulation of TLR4-dependent inflammation. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 4346-4351.	2.8	0
3	Cell-Mediated Immunity to NAGLU Transgene Following Intracerebral Gene Therapy in Children With Mucopolysaccharidosis Type IIIB Syndrome. <i>Frontiers in Immunology</i> , 2021, 12, 655478.	4.8	16
4	Surrogate Cerebrospinal Fluid Biomarkers for Assessing the Efficacy of Gene Therapy in Hurler Syndrome. <i>Frontiers in Neurology</i> , 2021, 12, 640547.	2.4	0
5	Can antidepressants unlock prescription of rimonabant in the fight against COVID-19?. <i>Molecular Psychiatry</i> , 2021, 26, 7091-7092.	7.9	3
6	Intracerebral Gene Therapy in Four Children with Sanfilippo B Syndrome: 5.5-Year Follow-Up Results. <i>Human Gene Therapy</i> , 2021, 32, 1251-1259.	2.7	9
7	Human Cytomegalovirus Infection Changes the Pattern of Surface Markers of Small Extracellular Vesicles Isolated From First Trimester Placental Long-Term Histocultures. <i>Frontiers in Cell and Developmental Biology</i> , 2021, 9, 689122.	3.7	7
8	AAVrh10 Vector Corrects Disease Pathology in MPS IIIA Mice and Achieves Widespread Distribution of SGSH in Large Animal Brains. <i>Molecular Therapy - Methods and Clinical Development</i> , 2020, 17, 174-187.	4.1	21
9	GFOGER Peptide Modifies the Protein Content of Extracellular Vesicles and Inhibits Vascular Calcification. <i>Frontiers in Cell and Developmental Biology</i> , 2020, 8, 589761.	3.7	8
10	Glycerophosphodiesterase 3 (GDE3) is a lysophosphatidylinositol-specific ectophospholipase C acting as an endocannabinoid signaling switch. <i>Journal of Biological Chemistry</i> , 2020, 295, 15767-15781.	3.4	7
11	Possible Role of Adipose Tissue and the Endocannabinoid System in Coronavirus Disease 2019 Pathogenesis: Can Rimobant Return?. <i>Obesity</i> , 2020, 28, 1580-1581.	3.0	12
12	Effects of Chronic Kidney Disease and Uremic Toxins on Extracellular Vesicle Biology. <i>Toxins</i> , 2020, 12, 811.	3.4	11
13	Analysis of Mucopolysaccharidosis Type VI through Integrative Functional Metabolomics. <i>International Journal of Molecular Sciences</i> , 2019, 20, 446.	4.1	18
14	Predominant role of microglia in brain iron retention in Sanfilippo syndrome, a pediatric neurodegenerative disease. <i>Glia</i> , 2018, 66, 1709-1723.	4.9	21
15	Unveiling metabolic remodeling in mucopolysaccharidosis type III through integrative metabolomics and pathway analysis. <i>Journal of Translational Medicine</i> , 2018, 16, 248.	4.4	19
16	High urinary ferritin reflects myoglobin iron evacuation in DMD patients. <i>Neuromuscular Disorders</i> , 2018, 28, 564-571.	0.6	13
17	Oligogalacturonic Acid Inhibits Vascular Calcification by Two Mechanisms. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2017, 37, 1391-1401.	2.4	32
18	Intracerebral administration of rAAV2/5hNAGLU vector in children with MPS IIIB: results at 30 months of a phase I/II trial. <i>Molecular Genetics and Metabolism</i> , 2017, 120, S130.	1.1	2

#	ARTICLE	IF	CITATIONS
19	Urinary metabolic phenotyping of mucopolysaccharidosis type I combining untargeted and targeted strategies with data modeling. <i>Clinica Chimica Acta</i> , 2017, 475, 7-14.	1.1	19
20	Magnesium Sulfate Prevents Neurochemical and Long-Term Behavioral Consequences of Neonatal Excitotoxic Lesions: Comparison Between Male and Female Mice. <i>Journal of Neuropathology and Experimental Neurology</i> , 2017, 76, 883-897.	1.7	18
21	Evaluation of the Diagnostic and Prognostic Value of Procalcitonin in Acute Colitis. <i>Gastroenterology</i> , 2017, 152, S805.	1.3	0
22	Intracerebral gene therapy in children with mucopolysaccharidosis type IIIB syndrome: an uncontrolled phase 1/2 clinical trial. <i>Lancet Neurology</i> , The, 2017, 16, 712-720.	10.2	149
23	Efficient recovery of glycosaminoglycan oligosaccharides from polyacrylamide gel electrophoresis combined with mass spectrometry analysis. <i>Analytical and Bioanalytical Chemistry</i> , 2017, 409, 1257-1269.	3.7	11
24	Interactions between Flow Oscillations and Biochemical Parameters in the Cerebrospinal Fluid. <i>Frontiers in Aging Neuroscience</i> , 2016, 8, 154.	3.4	20
25	CSF protein variations correlates with CSF oscillations in hydrocephalus patients. <i>Fluids and Barriers of the CNS</i> , 2015, 12, O34.	5.0	0
26	Neuroinflammation, mitochondrial defects and neurodegeneration in mucopolysaccharidosis III type C mouse model. <i>Brain</i> , 2015, 138, 336-355.	7.6	113
27	Oxidative stress is independent of inflammation in the neurodegenerative sanfilippo syndrome type B. <i>Journal of Neuroscience Research</i> , 2015, 93, 424-432.	2.9	23
28	Brain disease in mucopolysaccharidosis III C mouse: Neuroinflammation, mitochondrial defects and neurodegeneration. <i>Molecular Genetics and Metabolism</i> , 2015, 114, S97.	1.1	0
29	Su1249 Evaluation of NT-proBNP in Inflammatory Bowel Disease. <i>Gastroenterology</i> , 2015, 148, S-451.	1.3	0
30	Heparan Sulfate Saccharides Modify Focal Adhesions: Implication in Mucopolysaccharidosis Neuropathophysiology. <i>Journal of Molecular Biology</i> , 2015, 427, 775-791.	4.2	31
31	Removal of albumin and immunoglobulins from canine cerebrospinal fluid using depletion kits: a feasibility study. <i>Fluids and Barriers of the CNS</i> , 2014, 11, 14.	5.0	6
32	Chemistry of free radicals produced by oxidation of endogenous α -aminoketones. A study of 5-aminolevulinic acid and α -aminoacetone by fast kinetics spectroscopy. <i>Biochimica Et Biophysica Acta - General Subjects</i> , 2014, 1840, 3190-3197.	2.4	0
33	GM130 gain-of-function induces cell pathology in a model of lysosomal storage disease. <i>Human Molecular Genetics</i> , 2012, 21, 1481-1495.	2.9	26
34	Oxidized low density lipoprotein induces cyclin a synthesis. Involvement of ERK, JNK and NFkappaB. <i>Atherosclerosis</i> , 2011, 218, 308-313.	0.8	15
35	Barhl2 limits growth of the diencephalic primordium through Caspase3 inhibition of β -catenin activation. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 2288-2293.	7.1	24
36	Safe, Efficient, and Reproducible Gene Therapy of the Brain in the Dog Models of Sanfilippo and Hurler Syndromes. <i>Molecular Therapy</i> , 2011, 19, 251-259.	8.2	129

#	ARTICLE	IF	CITATIONS
37	Methods for Noninvasive Monitoring of Muscle Fiber Survival with an AAV Vector Encoding the mSEAP Reporter Gene. <i>Methods in Molecular Biology</i> , 2011, 709, 63-74.	0.9	2
38	Storage problems in lysosomal diseases. <i>Biochemical Society Transactions</i> , 2010, 38, 1442-1447.	3.4	9
39	GAP43 overexpression and enhanced neurite outgrowth in mucopolysaccharidosis type IIIB cortical neuron cultures. <i>Journal of Neuroscience Research</i> , 2010, 88, 202-213.	2.9	23
40	Storage Vesicles in Neurons Are Related to Golgi Complex Alterations in Mucopolysaccharidosis IIIB. <i>American Journal of Pathology</i> , 2010, 177, 2984-2999.	3.8	39
41	Enhanced degradation of synaptophysin by the proteasome in mucopolysaccharidosis type IIIB. <i>Molecular and Cellular Neurosciences</i> , 2009, 41, 8-18.	2.2	37
42	O2-P001 Caspase3 and the homeodomain-containing gene Barhl2 act as brakes on neuroepithelial cell proliferation by inhibiting β -catenin activation. <i>Mechanisms of Development</i> , 2009, 126, S60.	1.7	0
43	Abnormal expression of truncated CRMP-1 protein in the brain cortex of MPSIIIB mice. <i>Molecular Genetics and Metabolism</i> , 2008, 94, 135-138.	1.1	1
44	Early Neurodegeneration Progresses Independently of Microglial Activation by Heparan Sulfate in the Brain of Mucopolysaccharidosis IIIB Mice. <i>PLoS ONE</i> , 2008, 3, e2296.	2.5	114
45	Mutations in TMEM76* Cause Mucopolysaccharidosis IIIC (Sanfilippo C Syndrome). <i>American Journal of Human Genetics</i> , 2006, 79, 807-819.	6.2	77
46	An acetylated 120-kDa lysosomal transmembrane protein is absent from mucopolysaccharidosis IIIC fibroblasts: A candidate molecule for MPS IIIC. <i>Molecular Genetics and Metabolism</i> , 2006, 87, 22-31.	1.1	17
47	β -L-Iduronidase transport in neurites. <i>Molecular Genetics and Metabolism</i> , 2006, 87, 349-358.	1.1	20
48	Gene therapy of the brain in the dog model of Hurler's syndrome. <i>Annals of Neurology</i> , 2006, 60, 204-213.	5.3	94
49	Localisation of a gene for mucopolysaccharidosis IIIC to the pericentromeric region of chromosome 8. <i>Journal of Medical Genetics</i> , 2004, 41, 941-945.	3.2	20
50	Dinoflagellate centrosome: Associated proteins old and new. <i>European Journal of Protistology</i> , 2000, 36, 1-19.	1.5	7
51	Cyclic Expression of A Nuclear Protein In A Dinoflagellate. <i>Journal of Eukaryotic Microbiology</i> , 1999, 46, 259-267.	1.7	17
52	Characterization of p80, a Novel Nuclear and Cytoplasmic Protein in Dinoflagellates. <i>Protist</i> , 1999, 150, 197-211.	1.5	13
53	Preservation of viable biological samples for experiments in space laboratories. <i>Journal of Biotechnology</i> , 1996, 47, 377-393.	3.8	8
54	Nuclear and cytoplasmic actin in dinoflagellates. <i>Biology of the Cell</i> , 1996, 87, 17-35.	2.0	8

#	ARTICLE	IF	CITATIONS
55	Proteins related to mitosis in unicellular dinoflagellates, a biochemical study. <i>Biology of the Cell</i> , 1995, 84, 103-103.	2.0	0