

Lucie Sedlářková

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8402736/publications.pdf>

Version: 2024-02-01

13
papers

140
citations

1307594

7
h-index

1199594

12
g-index

13
all docs

13
docs citations

13
times ranked

279
citing authors

#	ARTICLE	IF	CITATIONS
1	Heat-shock protein expression in leukemia. <i>Tumor Biology</i> , 2011, 32, 33-44.	1.8	30
2	Cell surface and relative mRNA expression of heat shock protein 70 in human synovial cells. <i>Autoimmunity</i> , 2009, 42, 17-24.	2.6	23
3	UBTF Mutation Causes Complex Phenotype of Neurodegeneration and Severe Epilepsy in Childhood. <i>Neuropediatrics</i> , 2019, 50, 057-060.	0.6	21
4	Novel variant in the KCNK9 gene in a girl with Birk Barel syndrome. <i>European Journal of Medical Genetics</i> , 2020, 63, 103619.	1.3	16
5	Fetal cells of mesenchymal origin in cultures derived from synovial tissue and skin of patients with rheumatoid arthritis. <i>Joint Bone Spine</i> , 2008, 75, 563-566.	1.6	11
6	Expression of heat shock protein receptors on fibroblast-like synovial cells derived from rheumatoid arthritis-affected joints. <i>Rheumatology International</i> , 2008, 28, 837-844.	3.0	10
7	Analysis of cell surface and relative gene expression of heat shock protein 70 in human leukemia cell lines. <i>Leukemia and Lymphoma</i> , 2008, 49, 570-576.	1.3	9
8	Severe neurodevelopmental disorder with intractable seizures due to a novel SLC1A4 homozygous variant. <i>European Journal of Medical Genetics</i> , 2021, 64, 104263.	1.3	8
9	Peripheral blood mononuclear cell responses to heat shock proteins in patients undergoing stem cell transplantation. <i>Pediatric Transplantation</i> , 2006, 10, 178-186.	1.0	4
10	Cellules fœtales mésenchymateuses d'origine de tissu synovial et de peau de patients atteints de polyarthrite rhumatoïde. <i>Revue Du Rhumatisme (Edition Francaise)</i> , 2008, 75, 835-838.	0.0	3
11	Expression of heat shock protein 70 and NKG2D ligands in acute myeloid leukemia cell lines. <i>Journal of Receptor and Signal Transduction Research</i> , 2010, 30, 161-169.	2.5	3
12	A novel variant in <i>YWHAG</i> further supports phenotype of developmental and epileptic encephalopathy. <i>American Journal of Medical Genetics, Part A</i> , 2021, 185, 1363-1365.	1.2	2
13	Whole-Exome Sequencing in Czech Patients with Neurogenetic Diseases. <i>Genetic Testing and Molecular Biomarkers</i> , 2020, 24, 264-273.	0.7	0