

Ove Bruland

List of Publications by Year in descending order

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Version: 2024-02-01

23
papers

601
citations

759233

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794594

19
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24
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24
times ranked

1011
citing authors

#	ARTICLE	IF	CITATIONS
1	Clinical features and molecular genetics of patients with ABCA4-related retinal dystrophies. <i>Acta Ophthalmologica</i> , 2021, 99, e733-e746.	1.1	6
2	Temperature-dependent autoactivation associated with clinical variability of <i>PDGFRB</i> Asn666 substitutions. <i>Human Molecular Genetics</i> , 2021, 30, 72-77.	2.9	6
3	The SH3PXD2A-HTRA1 fusion transcript is extremely rare in Norwegian sporadic vestibular schwannoma patients. <i>Journal of Neuro-Oncology</i> , 2021, 154, 35-40.	2.9	4
4	K ⁺ regulates relocation of Pellino2 to the site of NLRP3 inflammasome activation in macrophages. <i>FEBS Letters</i> , 2021, 595, 2437-2446.	2.8	6
5	Pellino2 in nonimmune cells: novel interaction partners and intracellular localization. <i>FEBS Letters</i> , 2021, 595, 2909-2921.	2.8	3
6	Genetic alterations associated with malignant transformation of sporadic vestibular schwannoma. <i>Acta Neurochirurgica</i> , 2021, , 1.	1.7	0
7	Gamma Knife Radiosurgery does not alter the copy number aberration profile in sporadic vestibular schwannoma. <i>Journal of Neuro-Oncology</i> , 2020, 149, 373-381.	2.9	1
8	Upregulated PDK4 expression is a sensitive marker of increased fatty acid oxidation. <i>Mitochondrion</i> , 2019, 49, 97-110.	3.4	75
9	B-Lymphocyte Depletion in Patients With Myalgic Encephalomyelitis/Chronic Fatigue Syndrome. <i>Annals of Internal Medicine</i> , 2019, 170, 585.	3.9	60
10	A tyrosine kinase-activating variant Asn666Ser in <i>PDGFRB</i> causes a progeria-like condition in the severe end of Penttinen syndrome. <i>European Journal of Human Genetics</i> , 2019, 27, 574-581.	2.8	20
11	Recurrent, Activating Variants in the Receptor Tyrosine Kinase <i>DDR2</i> Cause Warburg-Cinotti Syndrome. <i>American Journal of Human Genetics</i> , 2018, 103, 976-983.	6.2	17
12	Screening for viral nucleic acids in vestibular schwannoma. <i>Journal of NeuroVirology</i> , 2018, 24, 730-737.	2.1	4
13	<i>GBA2</i> Mutations Cause a Marinesco-Sjögren-Like Syndrome: Genetic and Biochemical Studies. <i>PLoS ONE</i> , 2017, 12, e0169309.	2.5	17
14	Adenoviral Mediated Expression of <i>BMP2</i> by Bone Marrow Stromal Cells Cultured in 3D Copolymer Scaffolds Enhances Bone Formation. <i>PLoS ONE</i> , 2016, 11, e0147507.	2.5	13
15	Metabolic profiling indicates impaired pyruvate dehydrogenase function in myalgic encephalopathy/chronic fatigue syndrome. <i>JCI Insight</i> , 2016, 1, e89376.	5.0	140
16	Serum BAFF and APRIL Levels, T-Lymphocyte Subsets, and Immunoglobulins after B-Cell Depletion Using the Monoclonal Anti-CD20 Antibody Rituximab in Myalgic Encephalopathy/Chronic Fatigue Syndrome. <i>PLoS ONE</i> , 2016, 11, e0161226.	2.5	18
17	<i>S100A16</i> promotes differentiation and contributes to a less aggressive tumor phenotype in oral squamous cell carcinoma. <i>BMC Cancer</i> , 2015, 15, 631.	2.6	43
18	Development of Congenital Stromal Corneal Dystrophy Is Dependent on Export and Extracellular Deposition of Truncated Decorin. , 2015, 56, 2909.		15

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19	B-Lymphocyte Depletion in Myalgic Encephalopathy/ Chronic Fatigue Syndrome. An Open-Label Phase II Study with Rituximab Maintenance Treatment. PLoS ONE, 2015, 10, e0129898.	2.5	103
20	Serum/Glucocorticoid-Regulated Kinase 1 (SGK1) Is a Prominent Target Gene of the Transcriptional Response to Cytokines In Multiple Myeloma and Supports the Growth of Myeloma Cells. Blood, 2010, 116, 1915-1915.	1.4	0
21	Inverse correlation between PDGFC expression and lymphocyte infiltration in human papillary thyroid carcinomas. BMC Cancer, 2009, 9, 425.	2.6	18
22	One-tube restriction enzyme digest and fluorescent labeling for restriction endonuclease fingerprinting single-strand conformational polymorphism. BioTechniques, 2004, 37, 906-912.	1.8	0
23	Accurate determination of the number of CAG repeats in the Huntington disease gene using a sequence-specific internal DNA standard. Clinical Genetics, 1999, 55, 198-202.	2.0	32