Ove Bruland

List of Publications by Year
in descending order

23	601	759233 12	794594 19
papers	citations	h-index	g-index

24	24	24	1011
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Metabolic profiling indicates impaired pyruvate dehydrogenase function in myalgic encephalopathy/chronic fatigue syndrome. JCI Insight, 2016, 1, e89376.	5.0	140
2	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
3	Upregulated PDK4 expression is a sensitive marker of increased fatty acid oxidation. Mitochondrion, 2019, 49, 97-110.	3.4	75
4	B-Lymphocyte Depletion in Patients With Myalgic Encephalomyelitis/Chronic Fatigue Syndrome. Annals of Internal Medicine, 2019, 170, 585.	3.9	60
5	S100A16 promotes differentiation and contributes to a less aggressive tumor phenotype in oral squamous cell carcinoma. BMC Cancer, 2015, 15, 631.	2.6	43
6	Accurate determination of the number of CAG repeats in the Huntington disease gene using a sequenceâ€specific internal DNA standard. Clinical Genetics, 1999, 55, 198-202.	2.0	32
7	A tyrosine kinase-activating variant Asn666Ser in PDGFRB causes a progeria-like condition in the severe end of Penttinen syndrome. European Journal of Human Genetics, 2019, 27, 574-581.	2.8	20
8	Inverse correlation between PDGFC expression and lymphocyte infiltration in human papillary thyroid carcinomas. BMC Cancer, 2009, 9, 425.	2.6	18
9	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. PLoS ONE, 2016, 11, e0161226.	2.5	18
10	Recurrent, Activating Variants in the Receptor Tyrosine Kinase DDR2 Cause Warburg-Cinotti Syndrome. American Journal of Human Genetics, 2018, 103, 976-983.	6.2	17
11	GBA2 Mutations Cause a Marinesco-SjÃ¶gren-Like Syndrome: Genetic and Biochemical Studies. PLoS ONE, 2017, 12, e0169309.	2.5	17
12	Development of Congenital Stromal Corneal Dystrophy Is Dependent on Export and Extracellular Deposition of Truncated Decorin. , 2015, 56, 2909.		15
13	Adenoviral Mediated Expression of BMP2 by Bone Marrow Stromal Cells Cultured in 3D Copolymer Scaffolds Enhances Bone Formation. PLoS ONE, 2016, 11, e0147507.	2.5	13
14	Clinical features and molecular genetics of patients with ABCA4â€retinal dystrophies. Acta Ophthalmologica, 2021, 99, e733-e746.	1.1	6
15	Temperature-dependent autoactivation associated with clinical variability of PDGFRB Asn666 substitutions. Human Molecular Genetics, 2021, 30, 72-77.	2.9	6
16	K⁺ regulates relocation of Pellinoâ€2 to the site of NLRP3 inflammasome activation in macrophages. FEBS Letters, 2021, 595, 2437-2446.	2.8	6
17	Screening for viral nucleic acids in vestibular schwannoma. Journal of NeuroVirology, 2018, 24, 730-737.	2.1	4
18	The SH3PXD2A-HTRA1 fusion transcript is extremely rare in Norwegian sporadic vestibular schwannoma patients. Journal of Neuro-Oncology, 2021, 154, 35-40.	2.9	4

#	Article	IF	Citations
19	Pellinoâ€2 in nonimmune cells: novel interaction partners and intracellular localization. FEBS Letters, 2021, 595, 2909-2921.	2.8	3
20	Gamma Knife Radiosurgery does not alter the copy number aberration profile in sporadic vestibular schwannoma. Journal of Neuro-Oncology, 2020, 149, 373-381.	2.9	1
21	One-tube restriction enzyme digest and fluorescent labeling for restriction endonuclease fingerprinting single-strand conformational polymorphism. BioTechniques, 2004, 37, 906-912.	1.8	0
22	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
23	Genetic alterations associated with malignant transformation of sporadic vestibular schwannoma. Acta Neurochirurgica, 2021, , 1.	1.7	0