Irene C Huffnagel

List of Publications by Year
in descending order

18	572	840776 11	888059 17
papers	citations	h-index	g-index

19	19	19	701
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Magnetic resonance spectroscopy as marker for neurodegeneration in X-linked adrenoleukodystrophy. NeuroImage: Clinical, 2021, 32, 102793.	2.7	1
2	Optical coherence tomography to measure the progression of myelopathy in adrenoleukodystrophy. Annals of Clinical and Translational Neurology, 2021, 8, 1064-1072.	3.7	2
3	Imaging in X-Linked Adrenoleukodystrophy. Neuropediatrics, 2021, 52, 252-260.	0.6	8
4	Five men with arresting and relapsing cerebral adrenoleukodystrophy. Journal of Neurology, 2021, 268, 936-940.	3.6	2
5	Molecular Biomarkers for Adrenoleukodystrophy: An Unmet Need. Cells, 2021, 10, 3427.	4.1	14
6	Optical coherence tomography shows neuroretinal thinning in myelopathy of adrenoleukodystrophy. Journal of Neurology, 2020, 267, 679-687.	3.6	6
7	Plasma NfL and GFAP as biomarkers of spinal cord degeneration in adrenoleukodystrophy. Annals of Clinical and Translational Neurology, 2020, 7, 2127-2136.	3.7	19
8	Postural Body Sway as Surrogate Outcome for Myelopathy in Adrenoleukodystrophy. Frontiers in Physiology, 2020, 11, 786.	2.8	3
9	Spinal cord atrophy as a measure of severity of myelopathy in adrenoleukodystrophy. Journal of Inherited Metabolic Disease, 2020, 43, 852-860.	3.6	13
10	Overall intact cognitive function in male X-linked adrenoleukodystrophy adults with normal MRI. Orphanet Journal of Rare Diseases, 2019, 14, 217.	2.7	5
11	Disease progression in women with X-linked adrenoleukodystrophy is slow. Orphanet Journal of Rare Diseases, 2019, 14, 30.	2.7	58
12	Longitudinal diffusion MRI as surrogate outcome measure for myelopathy in adrenoleukodystrophy. Neurology, 2019, 93, e2133-e2143.	1.1	14
13	Progression of myelopathy in males with adrenoleukodystrophy: towards clinical trial readiness. Brain, 2019, 142, 334-343.	7.6	43
14	The Natural History of Adrenal Insufficiency in X-Linked Adrenoleukodystrophy: An International Collaboration. Journal of Clinical Endocrinology and Metabolism, 2019, 104, 118-126.	3.6	102
15	Comparison of C26:0-carnitine and C26:0-lysophosphatidylcholine as diagnostic markers in dried blood spots from newborns and patients with adrenoleukodystrophy. Molecular Genetics and Metabolism, 2017, 122, 209-215.	1.1	50
16	Mitochondrial Encephalopathy and Transient 3-Methylglutaconic Aciduria in ECHS1 Deficiency: Long-Term Follow-Up. JIMD Reports, 2017, 39, 83-87.	1.5	23
17	Adrenoleukodystrophy â€“ neuroendocrine pathogenesis and redefinition of natural history. Nature Reviews Endocrinology, 2016, 12, 606-615.	9.6	189
18	Rhizomelic chondrodysplasia punctata and cardiac pathology. Journal of Medical Genetics, 2013, 50, 419-424.	3.2	20