Jean Louis Mandel

List of Publications by Year
in descending order

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7 papers	863 citations	1307594 7 h-index	1720034 7 g-index
7 all docs	7 docs citations	7 times ranked	640 citing authors

#	Article	IF	Citations
1	The gene responsible for adrenoleukodystrophy encodes a peroxisomal membrane protein. Human Molecular Genetics, 1994, 3, 265-271.	2.9	263
2	Late onset neurological phenotype of the X-ALD gene inactivation in mice: a mouse model for adrenomyeloneuropathy. Human Molecular Genetics, 2002, 11, 499-505.	2.9	176
3	Functional overlap between ABCD1 (ALD) and ABCD2 (ALDR) transporters: a therapeutic target for X-adrenoleukodystrophy. Human Molecular Genetics, 2004, 13, 2997-3006.	2.9	170
4	Inactivation of the peroxisomal ABCD2 transporter in the mouse leads to late-onset ataxia involving mitochondria, Golgi and endoplasmic reticulum damage. Human Molecular Genetics, 2005, 14, 3565-3577.	2.9	90
5	Valproic acid induces antioxidant effects in X-linked adrenoleukodystrophy. Human Molecular Genetics, 2010, 19, 2005-2014.	2.9	90
6	Missense mutations are frequent in the gene for X-chromosomal adrenoleukodystrophy (ALD). Human Molecular Genetics, 1994, 3, 1903-1905.	2.9	57
7	Exon organisation of the mouse gene encoding the Adrenoleukodystrophy related protein (ALDRP). European Journal of Human Genetics, 1998, 6, 638-641.	2.8	17