Thanh T Le

List of Publications by Year
in descending order

9 papers	1,874 citations	1040056 9 h-index	1372567 10 g-index
10 all docs	10 docs citations	10 times ranked	1636 citing authors

#	Article	IF	Citations
1	SMNÎ”7, the major product of the centromeric survival motor neuron (SMN2) gene, extends survival in mice with spinal muscular atrophy and associates with full-length SMN. Human Molecular Genetics, 2005, 14, 845-857.	2.9	550
2	The Relationship between SMN, the Spinal Muscular Atrophy Protein, and Nuclear Coiled Bodies in Differentiated Tissues and Cultured Cells. Experimental Cell Research, 2000, 256, 365-374.	2.6	183
3	A transgene carrying an A2G missense mutation in the SMN gene modulates phenotypic severity in mice with severe (type I) spinal muscular atrophy. Journal of Cell Biology, 2003, 160, 41-52.	5.2	140
4	A role for complexes of survival of motor neurons (SMN) protein with gemins and profilin in neurite-like cytoplasmic extensions of cultured nerve cells. Experimental Cell Research, 2005, 309, 185-197.	2.6	118
5	Temporal requirement for high SMN expression in SMA mice. Human Molecular Genetics, 2011, 20, 3578-3591.	2.9	118
6	Absence of gemin5 from SMN complexes in nuclear Cajal bodies. BMC Cell Biology, 2007, 8, 28.	3.0	44
7	Plastin 3 Expression Does Not Modify Spinal Muscular Atrophy Severity in the âˆ†7 SMA Mouse. PLoS ONE, 2015, 10, e0132364.	2.5	41
8	Mild SMN missense alleles are only functional in the presence of SMN2 in mammals. Human Molecular Genetics, 2018, 27, 3404-3416.	2.9	15
9	Conditional deletion of SMN in cell culture identifies functional SMN alleles. Human Molecular Genetics, 2021, 29, 3477-3492.	2.9	9