Isoform-specific NF1 mRNA levels correlate with diseas
1

Orphanet Journal of Rare Diseases

14, 261

DOI: 10.1186/s13023-019-1223-1

Citation Report

#	Article	IF	Citations
1	Retrospective Multicentric Study on Non-Optic CNS Tumors in Children and Adolescents with Neurofibromatosis Type 1. Cancers, 2020, 12, 1426.	1.7	8
2	Hybrid Minigene Assay: An Efficient Tool to Characterize mRNA Splicing Profiles of NF1 Variants. Cancers, 2021, 13, 999.	1.7	7
3	The role of mRNA in the development, diagnosis, treatment and prognosis of neural tumors. Molecular Cancer, 2021, 20, 49.	7.9	21
4	Spectrum of splicing variants in disease genes and the ability of RNA analysis to reduce uncertainty in clinical interpretation. American Journal of Human Genetics, 2021, 108, 696-708.	2.6	43
5	Genotype-Phenotype Correlations in Neurofibromatosis Type 1: A Single-Center Cohort Study. Cancers, 2021, 13, 1879.	1.7	21
6	Pathogenic noncoding variants in the neurofibromatosis and schwannomatosis predisposition genes. Human Mutation, 2021, 42, 1187-1207.	1.1	5
7	Clinical variability of neurofibromatosis 1: A modifying role of cooccurring <scp>PTPN11</scp> variants and atypical brain <scp>MRI</scp> findings. Clinical Genetics, 2021, 100, 563-572.	1.0	6
8	Impacts of NF1 Gene Mutations and Genetic Modifiers in Neurofibromatosis Type 1. Frontiers in Neurology, 2021, 12, 704639.	1.1	14
9	Assessment of Mosaicism and Detection of Cryptic Alleles in CRISPR/Cas9-Engineered Neurofibromatosis Type 1 and TP53 Mutant Porcine Models Reveals Overlooked Challenges in Precision Modeling of Human Diseases. Frontiers in Genetics, 2021, 12, 721045.	1.1	5
10	Neurofibromin and suppression of tumorigenesis: beyond the GAP. Oncogene, 2022, 41, 1235-1251.	2.6	13
11	BoneÂmetabolism in patients with type 1 neurofibromatosis: key role of sun exposure and physical activity. Scientific Reports, 2022, 12, 4368.	1.6	6
12	Retinal ciliopathies through the lens of Bardet-Biedl Syndrome: Past, present and future. Progress in Retinal and Eye Research, 2022, 89, 101035.	7.3	17
13	Assessment of Rare Genetic Variants to Identify Candidate Modifier Genes Underlying Neurological Manifestations in Neurofibromatosis 1 Patients. Genes, 2022, 13, 2218.	1.0	0