Ashley Cannon

List of Publications by Year
in descending order

7 papers	191 citations	1478505 6 h-index	1588992 8 g-index
8 all docs	8 docs citations	8 times ranked	350 citing authors

#	Article	IF	Citations
1	Expanding the clinical phenotype of individuals with a 3-bp in-frame deletion of the NF1 gene (c.2970_2972del): an update of genotypeâ€“phenotype correlation. Genetics in Medicine, 2019, 21, 867-876.	2.4	62
2	Cutaneous neurofibromas in Neurofibromatosis type I: a quantitative natural history study. Orphanet Journal of Rare Diseases, 2018, 13, 31.	2.7	50
3	Recurrent arginine substitutions in the ACTG2 gene are the primary driver of disease burden and severity in visceral myopathy. Human Mutation, 2020, 41, 641-654.	2.5	27
4	Genome sequencing as a first-line diagnostic test for hospitalized infants. Genetics in Medicine, 2022, 24, 851-861.	2.4	22
5	Genetic Counseling for Neurofibromatosis 1, Neurofibromatosis 2, and Schwannomatosisâ€”Practice Resource of the National Society of Genetic Counselors. Journal of Genetic Counseling, 2020, 29, 692-714.	1.6	13
6	Clinical trial design for cutaneous neurofibromas. Neurology, 2018, 91, S31-S37.	1.1	11
7	Perspective of Adults With Neurofibromatosis 1 and Cutaneous Neurofibromas. Neurology, 2021, 97, S15-S24.	1.1	5