Julia Schmidt

List of Publications by Year
in descending order

8 papers	117 citations	1684188 5 h-index	1720034 7 g-index
10 all docs	10 docs citations	10 times ranked	191 citing authors

#	Article	IF	Citations
1	A novel single-cell RNA-sequencing approach and its applicability connecting genotype to phenotype in ageing disease. Scientific Reports, 2022, 12, 4091.	3.3	12
2	Biallelic variants in YRDC cause a developmental disorder with progeroid features. Human Genetics, 2021, 140, 1679-1693.	3.8	3
3	Familial cleft tongue caused by a unique translation initiation codon variant in TP63. European Journal of Human Genetics, 2021, , .	2.8	7
4	Identification of the recently described plasminogen gene mutation p.Lys330Glu in a family from Northern Germany with hereditary angioedema. Clinical and Translational Allergy, 2019, 9, 9.	3.2	31
5	Hallermann-Streiff syndrome: A missing molecular link for a highly recognizable syndrome. , 2018, 178, 398-406.		18
6	An assessment of the experiences and needs of adolescents with chronic conditions in transitional care: a qualitative study to develop a patient education programme. Health and Social Care in the Community, 2017, 25, 652-666.	1.6	36
7	Molecular analysis of a novel intragenic deletion in GPC3 in three cousins with Simpsonâ€“Golabiâ€“Behmel syndrome. American Journal of Medical Genetics, Part A, 2017, 173, 1400-1405.	1.2	7
8	Progressive frontal intraosseous lipoma: Detection of the mosaic <scp> AKT1 </scp> variant discloses <scp>Proteus</scp> syndrome. Clinical Genetics, 0, , .	2.0	1