Alisa FÃ¶rster

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/6148623/publications.pdf

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9 papers	106 citations	1684188 5 h-index	1720034 7 g-index
9 all docs	9 docs citations	9 times ranked	265 citing authors

#	Article	IF	Citations
1	Validation and clinical application of transactivation assays for RUNX1 variant classification. Blood Advances, 2022, , .	5.2	5
2	A SUMO4 initiator codon variant in amyotrophic lateral sclerosis reduces SUMO4 expression and alters stress granule dynamics. Journal of Neurology, 2022, 269, 4863-4871.	3.6	3
3	Beyond Pathogenic RUNX1 Germline Variants: The Spectrum of Somatic Alterations in RUNX1-Familial Platelet Disorder with Predisposition to Hematologic Malignancies. Cancers, 2022, 14, 3431.	3.7	5
4	Rare germline variants in the E-cadherin gene CDH1 are associated with the risk of brain tumors of neuroepithelial and epithelial origin. Acta Neuropathologica, 2021, 142, 191-210.	7.7	6
5	FOCAD loss impacts microtubule assembly, G2/M progression and patient survival in astrocytic gliomas. Acta Neuropathologica, 2020, 139, 175-192.	7.7	15
6	SPG7 mutations in amyotrophic lateral sclerosis: a genetic link to hereditary spastic paraplegia. Journal of Neurology, 2020, 267, 2732-2743.	3.6	14
7	INNV-06. TREATMENT RESPONSE TO BEVACIZUMAB OVER TWO YEARS IN A PATIENT WITH GENETICALLY PROVEN SOMATIC NEUROFIBROMATOSIS TYPE 2 MOSAICISM. Neuro-Oncology, 2019, 21, vi131-vi131.	1.2	0
8	CBMT-12. FOCAD LOSS IMPACTS MICROTUBULE ASSEMBLY, G2/M PROGRESSION AND PATIENT SURVIVAL IN ASTROCYTIC GLIOMAS. Neuro-Oncology, 2019, 21, vi35-vi35.	1.2	0
9	Whole-Genome Sequencing of Cytogenetically Balanced Chromosome Translocations Identifies Potentially Pathological Gene Disruptions and Highlights the Importance of Microhomology in the Mechanism of Formation. Human Mutation, 2017, 38, 180-192.	2.5	58