

Ledia Brunga

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7345277/publications.pdf>

Version: 2024-02-01

12
papers

703
citations

1163117

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h-index

1281871

11
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13
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docs citations

13
times ranked

2152
citing authors

#	ARTICLE	IF	CITATIONS
1	Perspectives and Experiences of Parents and Adolescents Who Participate in a Pediatric Precision Oncology Program: "When You Feel Helpless, This Kind of Thing Is Very Helpful". JCO Precision Oncology, 2022, 6, e2100444.	3.0	6
2	Abstract 1428: DNA methylation predicts early onset of primary tumor in patients with Li-Fraumeni syndrome. Cancer Research, 2022, 82, 1428-1428.	0.9	0
3	Resolving driver events in MLL-r negative high-risk infant ALL.. Journal of Clinical Oncology, 2021, 39, 10030-10030.	1.6	0
4	DNA Polymerase and Mismatch Repair Exert Distinct Microsatellite Instability Signatures in Normal and Malignant Human Cells. Cancer Discovery, 2021, 11, 1176-1191.	9.4	46
5	Performance of the McGill Interactive Pediatric OncoGenetic Guidelines for Identifying Cancer Predisposition Syndromes. JAMA Oncology, 2021, 7, 1806.	7.1	22
6	Mutant p53 induces Golgi tubulo-vesiculation driving a prometastatic secretome. Nature Communications, 2020, 11, 3945.	12.8	52
7	IQSEC2-related encephalopathy in males and females: a comparative study including 37 novel patients. Genetics in Medicine, 2019, 21, 837-849.	2.4	47
8	mTOR Inhibitors as a New Therapeutic Strategy in Treatment Resistant Epilepsy in Hemimegalencephaly: A Case Report. Journal of Child Neurology, 2019, 34, 132-138.	1.4	24
9	Prevalence of Genetic Disorders and GLUT1 Deficiency in a Ketogenic Diet Clinic. Canadian Journal of Neurological Sciences, 2018, 45, 93-96.	0.5	5
10	Rearrangement bursts generate canonical gene fusions in bone and soft tissue tumors. Science, 2018, 361, .	12.6	121
11	High Rate of Recurrent De Novo Mutations in Developmental and Epileptic Encephalopathies. American Journal of Human Genetics, 2017, 101, 664-685.	6.2	337
12	Germline and somatic mutations in <i>STXBP1</i> with diverse neurodevelopmental phenotypes. Neurology: Genetics, 2017, 3, e199.	1.9	41