

Stephanie LaHaye

List of Publications by Year in descending order

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Version: 2024-02-01

23
papers

584
citations

840776

11
h-index

677142

22
g-index

25
all docs

25
docs citations

25
times ranked

1045
citing authors

#	ARTICLE	IF	CITATIONS
1	<sc><i>EGFR</i></sc> internal tandem duplications in fusion-negative congenital and neonatal spindle cell tumors. <i>Genes Chromosomes and Cancer</i> , 2023, 62, 17-26.	2.8	3
2	High early death rates, treatment resistance, and short survival of Black adolescents and young adults with AML. <i>Blood Advances</i> , 2022, 6, 5570-5581.	5.2	8
3	Novel morphologic findings in <sc>PLAG1-rearranged</sc> soft tissue tumors. <i>Genes Chromosomes and Cancer</i> , 2021, 60, 577-585.	2.8	9
4	Expanding the phenotypic spectrum of internal tandem duplications in somatic disease. <i>Molecular Genetics and Metabolism</i> , 2021, 132, S44.	1.1	0
5	KPT-330 Prevents Aortic Valve Calcification via a Novel C/EBP β Signaling Pathway. <i>Circulation Research</i> , 2021, 128, 1300-1316.	4.5	10
6	Molecular classification of a complex structural rearrangement of the RB1 locus in an infant with sporadic, isolated, intracranial, sellar region retinoblastoma. <i>Acta Neuropathologica Communications</i> , 2021, 9, 61.	5.2	5
7	Gastroblastoma with a novel <sc><i>EWSR1-CTBP1</i></sc> fusion presenting in adolescence. <i>Genes Chromosomes and Cancer</i> , 2021, 60, 640-646.	2.8	12
8	Endogenous retrovirus envelope as a tumor-associated immunotherapeutic target in murine osteosarcoma. <i>Science</i> , 2021, 24, 102759.	4.1	1
9	Evidence of pioneer factor activity of an oncogenic fusion transcription factor. <i>Science</i> , 2021, 24, 102867.	4.1	22
10	YAP1-FAM118B Fusion Defines a Rare Subset of Childhood and Young Adulthood Meningiomas. <i>American Journal of Surgical Pathology</i> , 2021, 45, 329-340.	3.7	14
11	Tgf β 1-Cthrc1 Signaling Plays an Important Role in the Short-Term Reparative Response to Heart Valve Endothelial Injury. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2021, 41, 2923-2942.	2.4	4
12	High Early Death Rates, Treatment Resistance and Short Survival of Black Adolescent and Young Adults (AYAs) with Acute Myeloid Leukemia (AML) (Alliance). <i>Blood</i> , 2021, 138, 221-221.	1.4	2
13	Discovery of clinically relevant fusions in pediatric cancer. <i>BMC Genomics</i> , 2021, 22, 872.	2.8	13
14	Clinically aggressive pediatric spinal ependymoma with novel MYC amplification demonstrates molecular and histopathologic similarity to newly described MYCN-amplified spinal ependymomas. <i>Acta Neuropathologica Communications</i> , 2021, 9, 192.	5.2	5
15	An evaluation of MGMT promoter methylation within the methylation subclasses of glioblastoma. <i>Neuro-Oncology Advances</i> , 2020, 2, vdaa117.	0.7	1
16	Developmental origins for semilunar valve stenosis identified in mice harboring congenital heart disease-associated <i>GATA4</i> mutation. <i>DMM Disease Models and Mechanisms</i> , 2019, 12, .	2.4	17
17	Postnatal and Adult Aortic Heart Valves Have Distinctive Transcriptional Profiles Associated With Valve Tissue Growth and Maintenance Respectively. <i>Frontiers in Cardiovascular Medicine</i> , 2018, 5, 30.	2.4	11
18	Epigenetic mechanisms underlying maternal diabetes-associated risk of congenital heart disease. <i>JCI Insight</i> , 2017, 2, .	5.0	59

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19	Notch1 haploinsufficiency causes ascending aortic aneurysms in mice. JCI Insight, 2017, 2, .	5.0	44
20	Utilization of Whole Exome Sequencing to Identify Causative Mutations in Familial Congenital Heart Disease. Circulation: Cardiovascular Genetics, 2016, 9, 320-329.	5.1	71
21	Rare GATA5 sequence variants identified in individuals with bicuspid aortic valve. Pediatric Research, 2014, 76, 211-216.	2.3	74
22	Genetics of Valvular Heart Disease. Current Cardiology Reports, 2014, 16, 487.	2.9	57
23	Endothelial nitric oxide signaling regulates Notch1 in aortic valve disease. Journal of Molecular and Cellular Cardiology, 2013, 60, 27-35.	1.9	142