

Stephanie Puget

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/12057339/publications.pdf>

Version: 2024-02-01

93
papers

7,321
citations

76326

40
h-index

58581

82
g-index

94
all docs

94
docs citations

94
times ranked

8935
citing authors

#	ARTICLE	IF	CITATIONS
1	New Brain Tumor Entities Emerge from Molecular Classification of CNS-PNETs. <i>Cell</i> , 2016, 164, 1060-1072.	28.9	702
2	Reduced H3K27me3 and DNA Hypomethylation Are Major Drivers of Gene Expression in K27M Mutant Pediatric High-Grade Gliomas. <i>Cancer Cell</i> , 2013, 24, 660-672.	16.8	633
3	Histone H3F3A and HIST1H3B K27M mutations define two subgroups of diffuse intrinsic pontine gliomas with different prognosis and phenotypes. <i>Acta Neuropathologica</i> , 2015, 130, 815-827.	7.7	482
4	Recurrent activating ACVR1 mutations in diffuse intrinsic pontine glioma. <i>Nature Genetics</i> , 2014, 46, 457-461.	21.4	423
5	Divergent clonal selection dominates medulloblastoma at recurrence. <i>Nature</i> , 2016, 529, 351-357.	27.8	266
6	Craniopharyngioma. <i>Nature Reviews Disease Primers</i> , 2019, 5, 75.	30.5	255
7	Clinical, Radiologic, Pathologic, and Molecular Characteristics of Long-Term Survivors of Diffuse Intrinsic Pontine Glioma (DIPG): A Collaborative Report From the International and European Society for Pediatric Oncology DIPG Registries. <i>Journal of Clinical Oncology</i> , 2018, 36, 1963-1972.	1.6	250
8	Pediatric craniopharyngiomas: classification and treatment according to the degree of hypothalamic involvement. <i>Journal of Neurosurgery: Pediatrics</i> , 2007, 106, 3-12.	1.3	225
9	Mesenchymal Transition and PDGFRA Amplification/Mutation Are Key Distinct Oncogenic Events in Pediatric Diffuse Intrinsic Pontine Gliomas. <i>PLoS ONE</i> , 2012, 7, e30313.	2.5	200
10	Frequent <i>hSNF5/INI1</i> Germline Mutations in Patients with Rhabdoid Tumor. <i>Clinical Cancer Research</i> , 2011, 17, 31-38.	7.0	191
11	Embryonal tumor with abundant neuropil and true rosettes (ETANTR), ependymoblastoma, and medulloepithelioma share molecular similarity and comprise a single clinicopathological entity. <i>Acta Neuropathologica</i> , 2014, 128, 279-289.	7.7	191
12	Locoregionally administered B7-H3-targeted CAR T cells for treatment of atypical teratoid/rhabdoid tumors. <i>Nature Medicine</i> , 2020, 26, 712-719.	30.7	172
13	Biopsy in a series of 130 pediatric diffuse intrinsic Pontine gliomas. <i>Child's Nervous System</i> , 2015, 31, 1773-1780.	1.1	145
14	Craniopharyngioma: the pendulum of surgical management. <i>Child's Nervous System</i> , 2005, 21, 691-695.	1.1	129
15	Pemetrexed and Gemcitabine as Combination Therapy for the Treatment of Group3 Medulloblastoma. <i>Cancer Cell</i> , 2014, 25, 516-529.	16.8	128
16	Stereotactic biopsy of diffuse pontine lesions in children. <i>Journal of Neurosurgery: Pediatrics</i> , 2007, 107, 1-4.	1.3	126
17	Craniopharyngioma. <i>Orphanet Journal of Rare Diseases</i> , 2007, 2, 18.	2.7	125
18	Injuries to inferior vermis and dentate nuclei predict poor neurological and neuropsychological outcome in children with malignant posterior fossa tumors. <i>Cancer</i> , 2009, 115, 1338-1347.	4.1	118

#	ARTICLE	IF	CITATIONS
19	<i>IDH1</i> and <i>IDH2</i> Mutations in Gliomas. <i>New England Journal of Medicine</i> , 2009, 360, 2248-2249.	27.0	112
20	Critical oncogenic mutations in newly diagnosed pediatric diffuse intrinsic pontine glioma. <i>Pediatric Blood and Cancer</i> , 2012, 58, 489-491.	1.5	111
21	New outlook on the diagnosis, treatment and follow-up of childhood-onset craniopharyngioma. <i>Nature Reviews Endocrinology</i> , 2017, 13, 299-312.	9.6	105
22	Aberrant ERBB4-SRC Signaling as a Hallmark of Group 4 Medulloblastoma Revealed by Integrative Phosphoproteomic Profiling. <i>Cancer Cell</i> , 2018, 34, 379-395.e7.	16.8	104
23	Histone H3 wild-type DIPG/DMG overexpressing EZHIP extend the spectrum diffuse midline gliomas with PRC2 inhibition beyond H3-K27M mutation. <i>Acta Neuropathologica</i> , 2020, 139, 1109-1113.	7.7	104
24	Radiotherapy with concurrent and adjuvant temozolomide in children with newly diagnosed diffuse intrinsic pontine glioma. <i>Journal of Neuro-Oncology</i> , 2012, 106, 399-407.	2.9	100
25	Clonally Expanded T Cells Reveal Immunogenicity of Rhabdoid Tumors. <i>Cancer Cell</i> , 2019, 36, 597-612.e8.	16.8	100
26	Germline Elongator mutations in Sonic Hedgehog medulloblastoma. <i>Nature</i> , 2020, 580, 396-401.	27.8	94
27	The occurrence of intracranial rhabdoid tumours in mice depends on temporal control of <i>Smarb1</i> inactivation. <i>Nature Communications</i> , 2016, 7, 10421.	12.8	92
28	Long-Term Outcome of 106 Consecutive Pediatric Ruptured Brain Arteriovenous Malformations After Combined Treatment. <i>Stroke</i> , 2014, 45, 1664-1671.	2.0	86
29	Transcriptomic and epigenetic profiling of diffuse midline gliomas, H3 K27M-mutant™ discriminate two subgroups based on the type of histone H3 mutated and not supratentorial or infratentorial location. <i>Acta Neuropathologica Communications</i> , 2018, 6, 117.	5.2	83
30	Co-occurrence of histone H3 K27M and BRAF V600E mutations in paediatric midline grade I ganglioglioma. <i>Brain Pathology</i> , 2018, 28, 103-111.	4.1	80
31	Thalamic tumors in children: a reappraisal. <i>Journal of Neurosurgery: Pediatrics</i> , 2007, 106, 354-362.	1.3	75
32	TP53 Pathway Alterations Drive Radioresistance in Diffuse Intrinsic Pontine Gliomas (DIPG). <i>Clinical Cancer Research</i> , 2019, 25, 6788-6800.	7.0	66
33	Clinical Relevance of Tumor Cells with Stem-Like Properties in Pediatric Brain Tumors. <i>PLoS ONE</i> , 2011, 6, e16375.	2.5	57
34	Preclinical evaluation of dasatinib alone and in combination with cabozantinib for the treatment of diffuse intrinsic pontine glioma. <i>Neuro-Oncology</i> , 2015, 17, 953-964.	1.2	56
35	Diagnostics of pediatric supratentorial RELA ependymomas: integration of information from histopathology, genetics, DNA methylation and imaging. <i>Brain Pathology</i> , 2019, 29, 325-335.	4.1	55
36	Neuronal differentiation distinguishes supratentorial and infratentorial childhood ependymomas. <i>Neuro-Oncology</i> , 2010, 12, 1126-1134.	1.2	54

#	ARTICLE	IF	CITATIONS
37	Clinical, Imaging, Histopathological and Molecular Characterization of Anaplastic Ganglioglioma. <i>Journal of Neuropathology and Experimental Neurology</i> , 2016, 75, 971-980.	1.7	54
38	A driver role for GABA metabolism in controlling stem and proliferative cell state through GHB production in glioma. <i>Acta Neuropathologica</i> , 2017, 133, 645-660.	7.7	53
39	High-grade gliomas in adolescents and young adults highlight histomolecular differences from their adult and pediatric counterparts. <i>Neuro-Oncology</i> , 2020, 22, 1190-1202.	1.2	50
40	Loss of SMARCE1 expression is a specific diagnostic marker of clear cell meningioma: a comprehensive immunophenotypical and molecular analysis. <i>Brain Pathology</i> , 2018, 28, 466-474.	4.1	46
41	Hypothalamic syndrome. <i>Nature Reviews Disease Primers</i> , 2022, 8, 24.	30.5	42
42	New <i>in vivo</i> avatars of diffuse intrinsic pontine gliomas (DIPG) from stereotactic biopsies performed at diagnosis. <i>Oncotarget</i> , 2017, 8, 52543-52559.	1.8	41
43	Treatment Strategies in Childhood Craniopharyngioma. <i>Frontiers in Endocrinology</i> , 2012, 3, 64.	3.5	40
44	Is Biopsy Safe in Children with Newly Diagnosed Diffuse Intrinsic Pontine Glioma?. <i>American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting</i> , 2012, , 629-633.	3.8	35
45	ERAP1 promotes Hedgehog-dependent tumorigenesis by controlling USP47-mediated degradation of β TrCP. <i>Nature Communications</i> , 2019, 10, 3304.	12.8	35
46	Cognitive and Academic Outcome After Benign or Malignant Cerebellar Tumor in Children. <i>Cognitive and Behavioral Neurology</i> , 2009, 22, 270-278.	0.9	32
47	Computation of reliable textural indices from multimodal brain MRI: suggestions based on a study of patients with diffuse intrinsic pontine glioma. <i>Physics in Medicine and Biology</i> , 2018, 63, 105003.	3.0	32
48	Predictors of Outcome in Patients with Pediatric Intracerebral Hemorrhage: Development and Validation of a Modified Score. <i>Radiology</i> , 2018, 286, 651-658.	7.3	31
49	Clear cell meningiomas are defined by a highly distinct DNA methylation profile and mutations in SMARCE1. <i>Acta Neuropathologica</i> , 2021, 141, 281-290.	7.7	31
50	Blood-brain barrier disruption with low-intensity pulsed ultrasound for the treatment of pediatric brain tumors: a review and perspectives. <i>Neurosurgical Focus</i> , 2020, 48, E10.	2.3	31
51	ATOH1 Promotes Leptomeningeal Dissemination and Metastasis of Sonic Hedgehog Subgroup Medulloblastomas. <i>Cancer Research</i> , 2017, 77, 3766-3777.	0.9	29
52	Habit learning dissociation in rats with lesions to the vermis and the interpositus of the cerebellum. <i>Neurobiology of Disease</i> , 2007, 27, 228-237.	4.4	27
53	Historadiological correlations in high-grade glioma with the histone 3.3 G34R mutation. <i>Journal of Neuroradiology</i> , 2018, 45, 316-322.	1.1	26
54	Isavuconazole Diffusion in Infected Human Brain. <i>Antimicrobial Agents and Chemotherapy</i> , 2019, 63, .	3.2	24

#	ARTICLE	IF	CITATIONS
55	Pediatric methylation class HGNET-MN1: unresolved issues with terminology and grading. <i>Acta Neuropathologica Communications</i> , 2019, 7, 176.	5.2	24
56	Intellectual, educational, and situation-based social outcome in adult survivors of childhood medulloblastoma. <i>Developmental Neurorehabilitation</i> , 2019, 22, 19-26.	1.1	22
57	Supratentorial non-RELA, ZFTA-fused ependymomas: a comprehensive phenotype genotype correlation highlighting the number of zinc fingers in ZFTA-NCOA1/2 fusions. <i>Acta Neuropathologica Communications</i> , 2021, 9, 135.	5.2	21
58	Brain abscess in children, a two-centre audit: outcomes and controversies. <i>Archives of Disease in Childhood</i> , 2020, 105, 288-291.	1.9	20
59	Multimodal Magnetic Resonance Imaging of Treatment-Induced Changes to Diffuse Infiltrating Pontine Gliomas in Children and Correlation to Patient Progression-Free Survival. <i>International Journal of Radiation Oncology Biology Physics</i> , 2017, 99, 476-485.	0.8	18
60	Pediatric Chordomas: Results of a Multicentric Study of 40 Children and Proposal for a Histopathological Prognostic Grading System and New Therapeutic Strategies. <i>Journal of Neuropathology and Experimental Neurology</i> , 2018, 77, 207-215.	1.7	18
61	Biological material collection to advance translational research and treatment of children with CNS tumours: position paper from the SIOPE Brain Tumour Group. <i>Lancet Oncology</i> , The, 2018, 19, e419-e428.	10.7	16
62	Developmental venous anomaly in adult patients with diffuse glioma. <i>Neurology</i> , 2019, 92, e55-e62.	1.1	15
63	An integrative histopathological and epigenetic characterization of primary intracranial mesenchymal tumors, FET:CREB fused broadening the spectrum of tumor entities in comparison with their soft tissue counterparts. <i>Brain Pathology</i> , 2022, 32, e13010.	4.1	15
64	The Management of Birth-Related Posterior Fossa Hematomas in Neonates. <i>Neurosurgery</i> , 2013, 72, 755-762.	1.1	14
65	Role of neoadjuvant chemotherapy in metastatic medulloblastoma: a comparative study in 92 children. <i>Neuro-Oncology</i> , 2020, 22, 1686-1695.	1.2	14
66	Reconstruction of a large calvarial traumatic defect using a custom-made porous hydroxyapatite implant covered by a free latissimus dorsi muscle flap in an 11-year-old patient. <i>Journal of Neurosurgery: Pediatrics</i> , 2017, 19, 51-55.	1.3	13
67	A kinome-wide shRNA screen uncovers vaccinia-related kinase 3 (VRK3) as an essential gene for diffuse intrinsic pontine glioma survival. <i>Oncogene</i> , 2019, 38, 6479-6490.	5.9	13
68	Child dermoid cyst mimicking a craniopharyngioma: the benefit of MRI T2-weighted diffusion sequence. <i>Child's Nervous System</i> , 2018, 34, 359-362.	1.1	12
69	Circular RNA profiling distinguishes medulloblastoma groups and shows aberrant RMST overexpression in WNT medulloblastoma. <i>Acta Neuropathologica</i> , 2021, 141, 975-978.	7.7	12
70	Intratumoral heterogeneity of MYC drives medulloblastoma metastasis and angiogenesis. <i>Neuro-Oncology</i> , 2022, 24, 1509-1523.	1.2	12
71	Pattern of loco-regional relapses and treatment in pediatric esthesioneuroblastoma: The French very rare tumors group (<i>Fracture</i>) contribution. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28154.	1.5	11
72	Radiogenomics of diffuse intrinsic pontine gliomas (DIPGs): correlation of histological and biological characteristics with multimodal MRI features. <i>European Radiology</i> , 2021, 31, 8913-8924.	4.5	11

#	ARTICLE	IF	CITATIONS
73	Management of advanced uniâ€•or bilateral retinoblastoma with macroscopic optic nerve invasion. <i>Pediatric Blood and Cancer</i> , 2020, 67, e27998.	1.5	10
74	A CBF decrease in the left supplementary motor areas: New insight into postoperative pediatric cerebellar mutism syndrome using arterial spin labeling perfusion MRI. <i>Journal of Cerebral Blood Flow and Metabolism</i> , 2021, 41, 3339-3349.	4.3	10
75	Prognostic Clinical and Biologic Features for Overall Survival after Relapse in Childhood Medulloblastoma. <i>Cancers</i> , 2021, 13, 53.	3.7	10
76	Tumor dissemination through surgical tracts in diffuse intrinsic pontine glioma. <i>Journal of Neurosurgery: Pediatrics</i> , 2018, 22, 678-683.	1.3	9
77	Deep intronic hotspot variant explaining rhabdoid tumor predisposition syndrome in two patients with atypical teratoid and rhabdoid tumor. <i>European Journal of Human Genetics</i> , 2017, 25, 1170-1172.	2.8	8
78	Diagnostic Accuracy of a Reduced Immunohistochemical Panel in Medulloblastoma Molecular Subtyping, Correlated to DNA-methylation Analysis. <i>American Journal of Surgical Pathology</i> , 2021, 45, 558-566.	3.7	7
79	Clinical and molecular analysis of smoothed inhibitors in Sonic Hedgehog medulloblastoma. <i>Neuro-Oncology Advances</i> , 2021, 3, vdab097.	0.7	5
80	DIPG-35. BIOLOGICAL MEDICINE FOR DIFFUSE INTRINSIC PONTINE GLIOMA (DIPG) ERADICATION: RESULTS OF THE THREE ARM BIOMARKER-DRIVEN RANDOMIZED BIOMEDE 1.0 TRIAL. <i>Neuro-Oncology</i> , 2020, 22, iii293-iii294.	1.2	5
81	Molecular screening for cancer treatment optimization (MOSCATO 01) in pediatric patients: First feasibility results of a prospective molecular stratification trial. <i>Journal of Clinical Oncology</i> , 2014, 32, 10050-10050.	1.6	5
82	Deciphering the genetic and epigenetic landscape of pediatric bithalamic tumors. <i>Brain Pathology</i> , 2022, 32, e13039.	4.1	5
83	A novel case of cribriform neuroepithelial tumor: A potential diagnostic pitfall in the ventricular system. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29037.	1.5	3
84	Posterior Fossa Arachnoid Cyst in a Pediatric Population is Associated with Social Perception and Rest Cerebral Blood Flow Abnormalities. <i>Cerebellum</i> , 2020, 19, 58-67.	2.5	2
85	Acute surgical management of children with ruptured brain arteriovenous malformation. <i>Journal of Neurosurgery: Pediatrics</i> , 2021, 27, 437-445.	1.3	2
86	Hydrocephalus in children with ruptured cerebral arteriovenous malformation. <i>Journal of Neurosurgery: Pediatrics</i> , 2020, 26, 283-287.	1.3	2
87	The dark matter of diffuse intrinsic pontine gliomas: an update. <i>Expert Opinion on Orphan Drugs</i> , 2019, 7, 11-20.	0.8	1
88	CNS tumors with YWHAE:NUTM2 and KDM2B-fusions present molecular similarities to extra-CNS tumors having BCOR internal tandem duplication or alternative fusions. <i>Acta Neuropathologica Communications</i> , 2021, 9, 176.	5.2	1
89	HG-46RECURRENT DIFFUSE INTRINSIC PONTINE GLIOMAS: CLINICAL, BIOLOGICAL, RADIOLOGICAL AND THERAPEUTIC FACTORS CORRELATING WITH THE SURVIVAL. <i>Neuro-Oncology</i> , 2016, 18, iii57.4-iii58.	1.2	0
90	HGG-42. GLIOMA ONCOGENESIS IN CONSTITUTIONNAL MISMATCH REPAIR DEFICIENCY (CMMRD) SYNDROME: A CLINICO-PATHOLOGICAL AND MOLECULAR STUDY IN 15 PATIENTS. <i>Neuro-Oncology</i> , 2018, 20, i97-i98.	1.2	0

#	ARTICLE	IF	CITATIONS
91	MBCL-21. GERMLINE ELONGATOR MUTATIONS IN SONIC HEDGEHOG MEDULLOBLASTOMA. Neuro-Oncology, 2020, 22, iii392-iii393.	1.2	0
92	DIPG-61. RESCUE REGIMENS AFTER BIOMEDE: POSSIBLE INFLUENCE ON OS ASSESSMENT. Neuro-Oncology, 2020, 22, iii299-iii299.	1.2	0
93	HGG-41. Glioma oncogenesis in the constitutional mismatch repair deficiency (CMMRD) syndrome. Neuro-Oncology, 2022, 24, i70-i70.	1.2	0