

# Nayuta Higa

## List of Publications by Year in descending order

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25  
papers

151  
citations

1683354

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1199166

12  
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docs citations

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times ranked

218  
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#	ARTICLE	IF	CITATIONS
1	An oncogenic splice variant of PDGFR $\beta$ in adult glioblastoma as a therapeutic target for selective CDK4/6 inhibitors. <i>Scientific Reports</i> , 2022, 12, 1275.	1.6	6
2	Prognostic impact of PDGFRA gain/amplification and MGMT promoter methylation status in patients with IDH wild-type glioblastoma. <i>Neuro-Oncology Advances</i> , 2022, 4, .	0.4	3
3	Fronto-Orbital Advancement and Posterior Cranial Vault Expansion Using Distraction Osteogenesis in Patients With Multiple Craniosynostosis. <i>Journal of Craniofacial Surgery</i> , 2021, 32, 1882-1885.	0.3	3
4	Roles for hENT1 and dCK in gemcitabine sensitivity and malignancy of meningioma. <i>Neuro-Oncology</i> , 2021, 23, 945-954.	0.6	11
5	The Incidence of Depressed Skull Fractures Due to the Use of Pin-Type Head Frame Systems in the Adult Population: 10-year Experience of a Single Neurosurgical Center. <i>World Neurosurgery</i> , 2021, 155, e395-e401.	0.7	2
6	Image-guided confirmation of a precision pull-through procedure during laparoscopically assisted anorectoplasty in an open MRI operating theater: first application in an infantile case with anorectal malformation. <i>Surgical Case Reports</i> , 2021, 7, 211.	0.2	1
7	Diffuse Large B-Cell Lymphoma of the Central Nervous System Manifesting with Intratumoral Hemorrhage: A Case Report and Literature Review. <i>World Neurosurgery</i> , 2020, 143, 490-494.	0.7	3
8	A tailored next-generation sequencing panel identified distinct subtypes of wildtype IDH and TERT promoter glioblastomas. <i>Cancer Science</i> , 2020, 111, 3902-3911.	1.7	34
9	A case of developing obstructive hydrocephalus following aqueductal stenosis caused by developmental venous anomalies. <i>Child's Nervous System</i> , 2020, 36, 1549-1555.	0.6	3
10	Histopathological variation in the demyelinating sentinel lesion of primary central nervous system lymphoma. , 2020, 11, 342.		5
11	A sellar neuroblastoma showing rapid growth and causing syndrome of inappropriate secretion of antidiuretic hormone: A case report. , 2020, 11, 165.		1
12	LGG-20. CLINICAL FEATURES AND TREATMENT RESULTS FOR PEDIATRIC OPTICO-HYPOTHALAMIC ASTROCYTOMA. <i>Neuro-Oncology</i> , 2020, 22, iii370-iii370.	0.6	0
13	RARE-04. INTELLECTUAL DEVELOPMENT IN CHILDREN WITH PEDIATRIC CRANIOPHARYNGIOMA AFTER TUMOR REMOVAL. <i>Neuro-Oncology</i> , 2020, 22, iii442-iii442.	0.6	0
14	LGG-32. CLINICAL OUTCOME OF PEDIATRIC GLIOMAS IN SINGLE INSTITUTION. <i>Neuro-Oncology</i> , 2020, 22, iii372-iii372.	0.6	0
15	High filamin-C expression predicts enhanced invasiveness and poor outcome in glioblastoma multiforme. <i>British Journal of Cancer</i> , 2019, 120, 819-826.	2.9	28
16	ANGI-06. FUNCTION OF FORMIN-LIKE 1 (FMNL1) IN GLIOBLASTOMA MULTIFORME. <i>Neuro-Oncology</i> , 2019, 21, vi31-vi31.	0.6	0
17	PATH-50. ROUTINE MOLECULAR DIAGNOSIS OF GLIOMA PATIENTS USING A GLIOMA-SPECIFIC NGS PANEL. <i>Neuro-Oncology</i> , 2019, 21, vi154-vi155.	0.6	0
18	COT-14 CLINICAL FEATURES OF PEDIATRIC CENTRAL NERVOUS SYSTEM TUMORS. <i>Neuro-Oncology Advances</i> , 2019, 1, ii43-ii43.	0.4	0

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19	MPC-15 FEASIBILITY OF GLIOMA SPECIFIC ONCOPANEL IN THE DIAGNOSIS OF GLIOMA. <i>Neuro-Oncology Advances</i> , 2019, 1, ii24-ii24.	0.4	0
20	Formin-like 1 (FMNL1) Is Associated with Glioblastoma Multiforme Mesenchymal Subtype and Independently Predicts Poor Prognosis. <i>International Journal of Molecular Sciences</i> , 2019, 20, 6355.	1.8	12
21	Calcifying pseudoneoplasm of the neuraxis in direct continuity with a low-grade glioma: A case report and review of the literature. <i>Neuropathology</i> , 2017, 37, 446-451.	0.7	21
22	ATP7B expression in human glioblastoma is related to temozolomide resistance. <i>Oncology Letters</i> , 2017, 14, 7777-7782.	0.8	1
23	Ganglioglioma in the Third Ventricle: A Case Report and Literature Review. <i>NMC Case Report Journal</i> , 2016, 3, 97-101.	0.2	5
24	Concurrent occurrence of an inherited 16p13.11 microduplication and a de novo 19p13.3 microdeletion involving MAP2K2 in a patient with developmental delay, distinctive facial features, and lambdoid synostosis. <i>European Journal of Medical Genetics</i> , 2016, 59, 559-563.	0.7	5
25	Surgical technique for preventing subcutaneous migration of distal lumboperitoneal shunt catheters. <i>Innovative Neurosurgery</i> , 2013, 1, .	0.1	7