

Thorsten Simon

List of Publications by Year in descending order

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93
papers

6,726
citations

101543

36
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62596

80
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98
all docs

98
docs citations

98
times ranked

6047
citing authors

#	ARTICLE	IF	CITATIONS
1	Cardiovascular Health Status And Genetic Risk In Survivors of Childhood Neuroblastoma and Nephroblastoma Treated With Doxorubicin: Protocol of the Pharmacogenetic Part of the LESS-Anthra Cross-Sectional Cohort Study. <i>JMIR Research Protocols</i> , 2022, 11, e27898.	1.0	3
2	Circulating Cell-Free DNA Assessment in Biofluids from Children with Neuroblastoma Demonstrates Feasibility and Potential for Minimally Invasive Molecular Diagnostics. <i>Cancers</i> , 2022, 14, 2080.	3.7	6
3	Infants and Newborns with Atypical Teratoid Rhabdoid Tumors (ATRT) and Extracranial Malignant Rhabdoid Tumors (eMRT) in the EU-RHAB Registry: A Unique and Challenging Population. <i>Cancers</i> , 2022, 14, 2185.	3.7	9
4	ETMR-04. Embryonal tumor with multi-layered rosettes (ETMR) located in the brainstem: a case report on clinical decision-making and a multimodal, interdisciplinary treatment approach including interstitial brachytherapy. <i>Neuro-Oncology</i> , 2022, 24, i49-i50.	1.2	0
5	NFB-13. Rhabdoid Tumor Predisposition Syndrome (RTPS) – Finding Evidence by systematic Analyses. <i>Neuro-Oncology</i> , 2022, 24, i130-i131.	1.2	0
6	ATRT-05. Infants and newborns with atypical teratoid/rhabdoid tumors (ATRT) and extracranial malignant rhabdoid tumors: a unique and challenging population. <i>Neuro-Oncology</i> , 2022, 24, i2-i3.	1.2	0
7	Clinical and genetic risk factors define two risk groups of extracranial malignant rhabdoid tumours (eMRT/RTK). <i>European Journal of Cancer</i> , 2021, 142, 112-122.	2.8	15
8	Long-term follow-up of children with neuroblastoma receiving radiotherapy to metastatic lesions within the German Neuroblastoma Trials NB97 and NB2004. <i>Strahlentherapie Und Onkologie</i> , 2021, 197, 683-689.	2.0	6
9	Neuroblastom. <i>Springer Reference Medizin</i> , 2021, , 1-14.	0.0	0
10	Clinical and molecular characterization of patients with stage 4(M) neuroblastoma aged less than 18 months without MYCN amplification. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29038.	1.5	4
11	Genetic Alterations and Resectability Predict Outcome in Patients with Neuroblastoma Assigned to High-Risk Solely by MYCN Amplification. <i>Cancers</i> , 2021, 13, 4360.	3.7	1
12	Hypercalcemia is a frequent side effect of 13-cis-retinoic acid treatment in patients with high-risk neuroblastoma. <i>Pediatric Blood and Cancer</i> , 2021, , e29374.	1.5	1
13	Genotyping circulating tumor DNA of pediatric Hodgkin lymphoma. <i>Leukemia</i> , 2020, 34, 151-166.	7.2	53
14	Extended induction chemotherapy does not improve the outcome for high-risk neuroblastoma patients: results of the randomized open-label GPOH trial NB2004-HR. <i>Annals of Oncology</i> , 2020, 31, 422-429.	1.2	36
15	Synovial sarcoma disease characteristics and primary tumor sites differ between patient age groups: a report of the Cooperative Weichteilsarkom Studiengruppe (CWS). <i>Journal of Cancer Research and Clinical Oncology</i> , 2020, 146, 953-960.	2.5	10
16	Proton Beam Therapy for Children With Neuroblastoma: Experiences From the Prospective KiProReg Registry. <i>Frontiers in Oncology</i> , 2020, 10, 617506.	2.8	8
17	Mesenchymal Neuroblastoma Cells Are Undetected by Current mRNA Marker Panels: The Development of a Specific Neuroblastoma Mesenchymal Minimal Residual Disease Panel. <i>JCO Precision Oncology</i> , 2019, 3, 1-11.	3.0	17
18	The long noncoding RNA lncNB1 promotes tumorigenesis by interacting with ribosomal protein RPL35. <i>Nature Communications</i> , 2019, 10, 5026.	12.8	67

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19	A new risk score for patients after first recurrence of stage 4 neuroblastoma aged $\geq 18\text{ months}$ at first diagnosis. <i>Cancer Medicine</i> , 2019, 8, 7236-7243.	2.8	12
20	Brainstem biopsy in pediatric diffuse intrinsic pontine glioma in the era of precision medicine: the INFORM study experience. <i>European Journal of Cancer</i> , 2019, 114, 27-35.	2.8	51
21	Phase I/II intra-patient dose escalation study of vorinostat in children with relapsed solid tumor, lymphoma, or leukemia. <i>Clinical Epigenetics</i> , 2019, 11, 188.	4.1	27
22	Telomerase Is a Prognostic Marker of Poor Outcome and a Therapeutic Target in Neuroblastoma. <i>JCO Precision Oncology</i> , 2019, 3, 1-20.	3.0	29
23	Recurrence of Ewing sarcoma: Is detection by imaging follow-up protocol associated with survival advantage?. <i>Pediatric Blood and Cancer</i> , 2018, 65, e27011.	1.5	22
24	Surgical Implications for Diagnosis and Treatment of Intestinal Aspergillosis in Pediatric Patients with ALL. <i>European Journal of Pediatric Surgery</i> , 2018, 28, 477-483.	1.3	4
25	Retrospective analysis of relapsed abdominal high-risk neuroblastoma. <i>Journal of Pediatric Surgery</i> , 2018, 53, 558-566.	1.6	3
26	A mechanistic classification of clinical phenotypes in neuroblastoma. <i>Science</i> , 2018, 362, 1165-1170.	12.6	213
27	Dominant SCN2A Mutation Causes Familial Episodic Ataxia and Impairment of Speech Development. <i>Neuropediatrics</i> , 2018, 49, 379-384.	0.6	12
28	ATRT-16. CONGENITAL RHABDOID TUMORS AS A MAJOR CLINICAL CHALLENGE - A COLLABORATIVE EUROPEAN EFFORT. <i>Neuro-Oncology</i> , 2018, 20, i30-i31.	1.2	0
29	Long-term outcomes of the GPOH NB97 trial for children with high-risk neuroblastoma comparing high-dose chemotherapy with autologous stem cell transplantation and oral chemotherapy as consolidation. <i>British Journal of Cancer</i> , 2018, 119, 282-290.	6.4	30
30	Childhood cancer predisposition syndromes—A concise review and recommendations by the Cancer Predisposition Working Group of the Society for Pediatric Oncology and Hematology. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 1017-1037.	1.2	200
31	Update on the diagnostic value and safety of stereotactic biopsy for pediatric brainstem tumors: a systematic review and meta-analysis of 735 cases. <i>Journal of Neurosurgery: Pediatrics</i> , 2017, 20, 261-268.	1.3	90
32	Feasibility, Risk Profile and Diagnostic Yield of Stereotactic Biopsy in Children and Young Adults with Brain Lesions. <i>Klinische Padiatrie</i> , 2017, 229, 133-141.	0.6	14
33	2017 GPOH Guidelines for Diagnosis and Treatment of Patients with Neuroblastic Tumors. <i>Klinische Padiatrie</i> , 2017, 229, 147-167.	0.6	76
34	Molecular Classification Substitutes for the Prognostic Variables Stage, Age, and MYCN Status in Neuroblastoma Risk Assessment. <i>Neoplasia</i> , 2017, 19, 982-990.	5.3	26
35	Spinal Canal Involvement in Neuroblastoma. <i>Journal of Pediatrics</i> , 2017, 188, 294-298.	1.8	14
36	Lack of immunocytological GD2 expression on neuroblastoma cells in bone marrow at diagnosis, during treatment, and at recurrence*. <i>Pediatric Blood and Cancer</i> , 2017, 64, 46-56.	1.5	44

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37	Computer-Based Exercise Program: Effects of a 12-Week Intervention on Mood and Fatigue in Pediatric Patients With Cancer. <i>Clinical Journal of Oncology Nursing</i> , 2017, 21, E280-E286.	0.6	7
38	Complete surgical resection improves outcome in INRG high-risk patients with localized neuroblastoma older than 18 months. <i>BMC Cancer</i> , 2017, 17, 520.	2.6	63
39	Treatment and outcome of Ganglioneuroma and Ganglioneuroblastoma intermixed. <i>BMC Cancer</i> , 2016, 16, 542.	2.6	110
40	Prognostic significance of pattern and burden of metastatic disease in patients with stage 4 neuroblastoma: A study from the International Neuroblastoma Risk Group database. <i>European Journal of Cancer</i> , 2016, 65, 1-10.	2.8	56
41	Neuroblastoma messenger RNA is frequently detected in bone marrow at diagnosis of localised neuroblastoma patients. <i>European Journal of Cancer</i> , 2016, 54, 149-158.	2.8	10
42	I-131-mIBG therapy in neuroblastoma: established role and prospective applications. <i>Clinical and Translational Imaging</i> , 2016, 4, 87-101.	2.1	11
43	Assessment of Primary Site Response in Children With High-Risk Neuroblastoma: An International Multicenter Study. <i>Journal of Clinical Oncology</i> , 2016, 34, 740-746.	1.6	37
44	MYCN and HDAC5 transcriptionally repress <i>CD9</i> to trigger invasion and metastasis in neuroblastoma. <i>Oncotarget</i> , 2016, 7, 66344-66359.	1.8	30
45	Minimal residual disease detection in autologous stem cell grafts from patients with high risk neuroblastoma. <i>Pediatric Blood and Cancer</i> , 2015, 62, 1368-1373.	1.5	22
46	Osteosarcoma in Patients with Rothmund-Thomson Syndrome. <i>Pediatric Hematology and Oncology</i> , 2015, 32, 32-40.	0.8	16
47	Revised Risk Estimation and Treatment Stratification of Low- and Intermediate-Risk Neuroblastoma Patients by Integrating Clinical and Molecular Prognostic Markers. <i>Clinical Cancer Research</i> , 2015, 21, 1904-1915.	7.0	80
48	Chromosome 17/17q gain and unaltered profiles in high resolution array-CGH are prognostically informative in neuroblastoma. <i>Genes Chromosomes and Cancer</i> , 2014, 53, 639-649.	2.8	34
49	Significance of clinical and biologic features in Stage 3 neuroblastoma: A report from the International Neuroblastoma Risk Group project. <i>Pediatric Blood and Cancer</i> , 2014, 61, 1932-1939.	1.5	32
50	Role of Surgery in the Treatment of Patients With Stage 4 Neuroblastoma Age 18 Months or Older at Diagnosis. <i>Journal of Clinical Oncology</i> , 2013, 31, 752-758.	1.6	115
51	Iodine-123 Metaiodobenzylguanidine Scintigraphy Scoring Allows Prediction of Outcome in Patients With Stage 4 Neuroblastoma: Results of the Cologne Interscore Comparison Study. <i>Journal of Clinical Oncology</i> , 2013, 31, 944-951.	1.6	80
52	Diagnostic Value and Safety of Stereotactic Biopsy for Brainstem Tumors. <i>Neurosurgery</i> , 2013, 72, 873-882.	1.1	83
53	The RIST design: A molecularly targeted multimodal approach for the treatment of patients with relapsed and refractory neuroblastoma. <i>Journal of Clinical Oncology</i> , 2013, 31, 10017-10017.	1.6	14
54	Stereotactic iodine-125 brachytherapy for treatment of inoperable focal brainstem gliomas of WHO grades I and II: feasibility and long-term outcome. <i>Journal of Neuro-Oncology</i> , 2012, 109, 273-283.	2.9	29

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55	Short- and long-term outcome of patients with symptoms of spinal cord compression by neuroblastoma. <i>Developmental Medicine and Child Neurology</i> , 2012, 54, 347-352.	2.1	46
56	Chemotherapy-Associated Angiogenesis in Neuroblastoma Tumors. <i>American Journal of Pathology</i> , 2012, 180, 1370-1377.	3.8	13
57	Stereotactic Brachytherapy With Iodine-125 Seeds for the Treatment of Inoperable Low-Grade Gliomas in Children: Long-Term Outcome. <i>Journal of Clinical Oncology</i> , 2011, 29, 4151-4159.	1.6	66
58	Long term outcome of high-risk neuroblastoma patients after immunotherapy with antibody ch14.18 or oral metronomic chemotherapy. <i>BMC Cancer</i> , 2011, 11, 21.	2.6	113
59	Treatment and outcomes of patients with relapsed, high-risk neuroblastoma: Results of German trials. <i>Pediatric Blood and Cancer</i> , 2011, 56, 578-583.	1.5	110
60	Analysis of Ribosomal Protein Genes Associated with Diamond Blackfan Anemia (DBA) In German DBA Patients and Their Relatives. <i>Blood</i> , 2011, 118, 729-729.	1.4	2
61	Dosimetry for ¹³¹ I-MIBG therapies in metastatic neuroblastoma, pheochromocytoma and paraganglioma. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2010, 37, 1279-1290.	6.4	27
62	<i>HAX1</i> mutations causing severe congenital neuropenia and neurological disease lead to cerebral microstructural abnormalities documented by quantitative MRI. <i>American Journal of Medical Genetics, Part A</i> , 2010, 152A, 3157-3163.	1.2	23
63	Prognostic Impact of Gene Expression-Based Classification for Neuroblastoma. <i>Journal of Clinical Oncology</i> , 2010, 28, 3506-3515.	1.6	129
64	RESIDUAL LYMPH NODE METASTASIS IN STAGE 4 NEUROBLASTOMA—ADVANTAGE OF RADIO-GUIDED SURGERY?. <i>Pediatric Hematology and Oncology</i> , 2010, 27, 471-475.	0.8	2
65	Focal nodular hyperplasia of the liver in longterm survivors of neuroblastoma. <i>European Journal of Radiology</i> , 2010, 74, e1-e5.	2.6	35
66	The International Neuroblastoma Risk Group (INRG) Staging System: An INRG Task Force Report. <i>Journal of Clinical Oncology</i> , 2009, 27, 298-303.	1.6	869
67	Heterogeneity of the MYCN Oncogene in Neuroblastoma. <i>Clinical Cancer Research</i> , 2009, 15, 2085-2090.	7.0	52
68	Co-regulated expression of HAND2 and DEIN by a bidirectional promoter with asymmetrical activity in neuroblastoma. <i>BMC Molecular Biology</i> , 2009, 10, 28.	3.0	36
69	The International Neuroblastoma Risk Group (INRG) Classification System: An INRG Task Force Report. <i>Journal of Clinical Oncology</i> , 2009, 27, 289-297.	1.6	1,540
70	Review of image defined risk factors in localized neuroblastoma patients: Results of the GPOH NB97 trial. <i>Pediatric Blood and Cancer</i> , 2008, 50, 965-969.	1.5	79
71	The prognostic impact of functional imaging with ¹²³ I-mIBG in patients with stage 4 neuroblastoma >1 year of age on a high-risk treatment protocol: Results of the German Neuroblastoma Trial NB97. <i>European Journal of Cancer</i> , 2008, 44, 1552-1558.	2.8	88
72	Localized Infant Neuroblastomas Often Show Spontaneous Regression: Results of the Prospective Trials NB95-S and NB97. <i>Journal of Clinical Oncology</i> , 2008, 26, 1504-1510.	1.6	263

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73	Identification of DEIN, a Novel Gene with High Expression Levels in Stage IVS Neuroblastoma. <i>Molecular Cancer Research</i> , 2007, 5, 1276-1284.	3.4	22
74	Topotecan and Etoposide in the Treatment of Relapsed High-risk Neuroblastoma. <i>Journal of Pediatric Hematology/Oncology</i> , 2007, 29, 101-106.	0.6	35
75	Activation of Akt Predicts Poor Outcome in Neuroblastoma. <i>Cancer Research</i> , 2007, 67, 735-745.	0.9	218
76	Topotecan, cyclophosphamide, and etoposide (TCE) in the treatment of high-risk neuroblastoma. Results of a phase-II trial. <i>Journal of Cancer Research and Clinical Oncology</i> , 2007, 133, 653-661.	2.5	60
77	Solide Tumoren. , 2007, , 805-870.		0
78	Risk estimation in localized unresectable single copy MYCN neuroblastoma by the status of chromosomes 1p and 11q. <i>Cancer Letters</i> , 2006, 237, 215-222.	7.2	21
79	Intensified External-Beam Radiation Therapy Improves the Outcome of Stage 4 Neuroblastoma in Children > 1 Year with Residual Local Disease. <i>Strahlentherapie Und Onkologie</i> , 2006, 182, 389-394.	2.0	76
80	Oligonucleotide array-based comparative genomic hybridization (aCGH) of 90 neuroblastomas reveals aberration patterns closely associated with relapse pattern and outcome. <i>Genes Chromosomes and Cancer</i> , 2006, 45, 1130-1142.	2.8	72
81	Loss in Chromosome 11q Identifies Tumors with Increased Risk for Metastatic Relapses in Localized and 4S Neuroblastoma. <i>Clinical Cancer Research</i> , 2006, 12, 3368-3373.	7.0	92
82	Clinical Presentation. , 2005, , 63-85.		9
83	Myeloablative megatherapy with autologous stem-cell rescue versus oral maintenance chemotherapy as consolidation treatment in patients with high-risk neuroblastoma: a randomised controlled trial. <i>Lancet Oncology</i> , The, 2005, 6, 649-658.	10.7	350
84	The role of age in neuroblastoma risk stratification: the German, Italian, and children's oncology group perspectives. <i>Cancer Letters</i> , 2005, 228, 257-266.	7.2	48
85	Epidural compression in neuroblastoma: Diagnostic and therapeutic aspects. <i>Cancer Letters</i> , 2005, 228, 283-299.	7.2	53
86	Consolidation Treatment With Chimeric Anti-GD2-Antibody ch14.18 in Children Older Than 1 Year With Metastatic Neuroblastoma. <i>Journal of Clinical Oncology</i> , 2004, 22, 3549-3557.	1.6	140
87	New definition of low-risk neuroblastoma using stage, age, and 1p and MYCN status. <i>Journal of Pediatric Hematology/Oncology</i> , 2004, 26, 791-6.	0.6	23
88	Asymmetric salivary gland ¹²³ I-meta-iodobenzylguanidine uptake in a patient with cervical neuroblastoma and horner syndrome. <i>Medical and Pediatric Oncology</i> , 2001, 36, 489-490.	1.0	4
89	Metastatic neuroblastoma in infancy: What does the pattern of metastases contribute to prognosis?. <i>Medical and Pediatric Oncology</i> , 2000, 35, 683-687.	1.0	35
90	Testicular and paratesticular involvement by metastatic neuroblastoma. <i>Cancer</i> , 2000, 88, 2636-2641.	4.1	33

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91	Metastatic neuroblastoma in infancy: What does the pattern of metastases contribute to prognosis?. , 2000, 35, 683.		2
92	Clinical Features of Progressive Neuroblastoma. Pediatric and Adolescent Medicine, 0, , 1-9.	0.4	0
93	Adrenocortical Tumors and Pheochromocytoma/Paraganglioma Initially Mistaken as Neuroblastomaâ€”Experiences From the GPOH-MET Registry. Frontiers in Endocrinology, 0, 13, .	3.5	4