Rhoikos FurtwÃ¤ngler

List of Publications by Year
in descending order

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91 papers	2,478 citations	201674 27 h-index	223800 46 g-index
111 all docs	111 docs citations	111 times ranked	2455 citing authors

#	Article	IF	Citations
1	Current and Emerging Therapeutic Approaches for Extracranial Malignant Rhabdoid Tumors. Cancer Management and Research, 2022, Volume 14, 479-498.	1.9	11
2	Treatment of patients with stage I focal anaplastic and diffuse anaplastic Wilms tumour: A report from the SIOP-WT-2001 GPOH and UK-CCLG studies. European Journal of Cancer, 2022, 166, 1-7.	2.8	0
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. Cancers, 2022, 14, 2185.	3.7	9
4	International Comparisons of Clinical Demographics and Outcomes in the International Society of Pediatric Oncology Wilms Tumor 2001 Trial and Study. JCO Global Oncology, 2022, 8, e2100425.	1.8	14
5	NFB-13. Rhabdoid Tumor Predisposition Syndrome (RTPS) â€“ Finding Evidence by systematic Analyses. Neuro-Oncology, 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. Neuro-Oncology, 2022, 24, i2-i3.	1.2	0
7	Persistent aseptic meningitis in aÂchildâ€”think patch. Wiener Medizinische Wochenschrift, 2021, 171, 38-40.	1.1	0
8	Clinical and genetic risk factors define two risk groups of extracranial malignant rhabdoid tumours (eMRT/RTK). European Journal of Cancer, 2021, 142, 112-122.	2.8	15
9	Clear cell sarcoma of the kidney in Austrian children: Longâ€term survival after relapse. Pediatric Blood and Cancer, 2021, 68, e28860.	1.5	7
10	Outcome of Stage IV Completely Necrotic Wilms Tumour and Local Stage III Treated According to the SIOP 2001 Protocol. Cancers, 2021, 13, 976.	3.7	6
11	Prognostic significance of histopathological response to preoperative chemotherapy in unilateral Wilms' tumor: An analysis of 899 patients treated on the SIOP WT 2001 protocol in the UKâ€CCLG and GPOH studies. International Journal of Cancer, 2021, 149, 1332-1340.	5.1	16
12	Fifty years of clinical and research studies for childhood renal tumors within the International Society of Pediatric Oncology (SIOP). Annals of Oncology, 2021, 32, 1327-1331.	1.2	14
13	Characteristics of Nephroblastoma/Nephroblastomatosis in Children with a Clinically Reported Underlying Malformation or Cancer Predisposition Syndrome. Cancers, 2021, 13, 5016.	3.7	6
14	MYCN and MAX alterations in Wilms tumor and identification of novel N-MYC interaction partners as biomarker candidates. Cancer Cell International, 2021, 21, 555.	4.1	10
15	High-risk blastemal Wilms tumor can be modeled by 3D spheroid cultures in vitro. Oncogene, 2020, 39, 849-861.	5.9	17
16	Age and DNA methylation subgroup as potential independent risk factors for treatment stratification in children with atypical teratoid/rhabdoid tumors. Neuro-Oncology, 2020, 22, 1006-1017.	1.2	72
17	Outcome of patients with stage IV high-risk Wilms tumour treated according to the SIOP2001 protocol: A report of the SIOP Renal Tumour Study Group. European Journal of Cancer, 2020, 128, 38-46.	2.8	24
18	Local Stage Dependent Necessity of Radiation Therapy in Rhabdoid Tumors of the Kidney (RTK). International Journal of Radiation Oncology Biology Physics, 2020, 108, 667-675.	0.8	6

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19	Nephroblastom. Springer Reference Medizin, 2019, , 1-12.	0.0	0
20	Solide Tumoren bei Kindern und Jugendlichen: Prinzipien der onkologischen Therapie. Springer Reference Medizin, 2019, , 701-715.	0.0	0
21	Highâ€dose treatment for malignant rhabdoid tumor of the kidney: No evidence for improved survivalâ€"The Gesellschaft fÃ¼r PÃdiatrische Onkologie und HÃmatologie (GPOH) experience. Pediatric Blood and Cancer, 2018, 65, e26746.	1.5	35
22	Rationale for the treatment of children with CCSK in the UMBRELLA SIOPâ€"RTSG 2016 protocol. Nature Reviews Urology, 2018, 15, 309-319.	3.8	43
23	ETV6â€"NTRK3 in congenital mesoblastic nephroma: A report of the SIOP/GPOH nephroblastoma study. Pediatric Blood and Cancer, 2018, 65, e26925.	1.5	41
24	The extraordinary challenge of treating patients with congenital rhabdoid tumorsâ€"a collaborative European effort. Pediatric Blood and Cancer, 2018, 65, e26999.	1.5	15
25	Outcome of two patients with bilateral nephroblastomatosis/Wilms tumour treated with an add-on 13-cis retinoic acid therapy â€" Case report. Pediatric Hematology and Oncology, 2018, 35, 218-224.	0.8	11
26	ATRT-17. ASSISTED REPRODUCTIVE TECHNOLOGIES AND THE DEVELOPMENT OF MALIGNANT RHABDOID TUMOURS. Neuro-Oncology, 2018, 20, i31-i31.	1.2	0
27	ATRT-16. CONGENITAL RHABDOID TUMORS AS A MAJOR CLINICAL CHALLENGE - A COLLABORATIVE EUROPEAN EFFORT. Neuro-Oncology, 2018, 20, i30-i31.	1.2	0
28	Relapse of Wilms' tumour and detection methods: a retrospective analysis of the 2001 Renal Tumour Study Groupâ€"International Society of Paediatric Oncology Wilms' tumour protocol database. Lancet Oncology, The, 2018, 19, 1072-1081.	10.7	59
29	Recurrent intragenic rearrangements of EGFR and BRAF in soft tissue tumors of infants. Nature Communications, 2018, 9, 2378.	12.8	72
30	Nierentumoren. , 2018, , 441-464.		2
31	Nosocomial legionellosis and invasive aspergillosis in a child with T-lymphoblastic leukemia. International Journal of Hygiene and Environmental Health, 2017, 220, 900-905.	4.3	6
32	TP53 alterations in Wilms tumour represent progression events with strong intratumour heterogeneity that are closely linked but not limited to anaplasia. Journal of Pathology: Clinical Research, 2017, 3, 234-248.	3.0	53
33	Rationale for the treatment of Wilms tumour in the UMBRELLA SIOPâ€"RTSG 2016 protocol. Nature Reviews Urology, 2017, 14, 743-752.	3.8	249
34	Improving outcomes of short peripheral vascular access in oncology and chemotherapy administration. Journal of Vascular Access, 2017, 18, 89-96.	0.9	24
35	WAGR(O) Syndrom â€" besondere Herausforderungen bei der Betreuung von Aniridie-Patienten. , 2017, 234, .		0
36	Association of FOXM1 expression with tumor histology and prognosis in Wilms tumor: Potential for a new prognostic marker. Oncology Letters, 2016, 12, 2854-2859.	1.8	3

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37	Genetic and epigenetic evaluation of potentially important subtypes of clear cell sarcoma of kidney (CCSK). European Journal of Cancer, 2016, 61, S25.	2.8	0
38	Mutually exclusive BCOR internal tandem duplications and YWHAEâ€NUTM2 fusions in clear cell sarcoma of kidney: not the full story. Journal of Pathology, 2016, 238, 617-620.	4.5	56
39	Nierentumoren beim Kind. , 2016, , 2025-2038.		2
40	Surveillance of bloodstream infections in pediatric cancer centers - what have we learned and how do we move on?. GMS Hygiene and Infection Control, 2016, 11, Doc11.	0.3	10
41	Tumoren der Blase und der Prostata beim Kind. , 2016, , 2043-2045.		0
42	Hodentumoren beim Kind. , 2016, , 2047-2051.		0
43	Neuroblastome in der pÃdiatrischen Urologie. , 2016, , 2039-2041.		0
44	Bloodstream infection in paediatric cancer centresâ€"leukaemia and relapsed malignancies are independent risk factors. European Journal of Pediatrics, 2015, 174, 675-686.	2.7	60
45	Development of Hypertension is Less Frequent after Bilateral Nephron Sparing Surgery for Bilateral Wilms Tumor in a Long-Term Survey. Journal of Urology, 2015, 193, 262-267.	0.4	30
46	Trichophagia: Rapunzel Syndrome in a 7-Year-Old Girl. Journal of Pediatrics, 2015, 166, 497.	1.8	7
47	New insights into the genetics of glioblastoma multiforme by familial exome sequencing. Oncotarget, 2015, 6, 5918-5931.	1.8	28
48	Impact of a modified Broviac maintenance care bundle on bloodstream infections in paediatric cancer patients. GMS Hygiene and Infection Control, 2015, 10, Doc15.	0.3	11
49	Mesoblastic Nephroma. , 2015, , 2745-2747.		0
50	DNA-Damage Foci to Detect and Characterize DNA Repair Alterations in Children Treated for Pediatric Malignancies. PLoS ONE, 2014, 9, e91319.	2.5	27
51	Treatment and outcome of patients with relapsed clear cell sarcoma of the kidney: a combined SIOP and AIEOP study. British Journal of Cancer, 2014, 111, 227-233.	6.4	49
52	Children with Relapsed or Refractory Nephroblastoma: Favorable Long-term Survival after High-dose Chemotherapy and Autologous Stem Cell Transplantation. Klinische Padiatrie, 2014, 226, 351-356.	0.6	7
53	Pretreatment for Bilateral Nephroblastomatosis is an Independent Risk Factor for Progressive Disease in Patients with Stage V Nephroblastoma. Klinische Padiatrie, 2014, 226, 175-181.	0.6	29
54	1q gain is a frequent finding in preoperatively treated Wilms tumors, but of limited prognostic value for risk stratification in the SIOP2001/GPOH trial. Genes Chromosomes and Cancer, 2014, 53, 960-962.	2.8	4

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55	Malignant Rhabdoid Tumor of the Kidney (MRTK) â€“ Data of 52 Patients Treated According to Protocols of the GPOH (German Society of Paediatric Oncology and Haematology). Cancer Genetics, 2014, 207, 454.	0.4	1
56	Malignant rhabdoid tumor of the kidney: significantly improved response to pre-operative treatment intensified with doxorubicin. Cancer Genetics, 2014, 207, 434-436.	0.4	14
57	Signals of neutropenia in human breath?. International Journal for Ion Mobility Spectrometry, 2014, 17, 19-23.	1.4	5
58	Preoperative chemotherapy and local stage III in nephroblastoma. Translational Pediatrics, 2014, 3, 4-11.	1.2	27
59	Hodentumoren beim Kind. , 2014, , 1-8.		0
60	Tumoren des Urogenitalsystems. , 2014, , 537-545.		0
61	Nierentumoren beim Kind. , 2014, , 1-19.		0
62	Clear Cell Sarcomas of the Kidney registered on International Society of Pediatric Oncology (SIOP) 93-01 and SIOP 2001 protocols: A report of the SIOP Renal Tumour Study Group. European Journal of Cancer, 2013, 49, 3497-3506.	2.8	105
63	An international strategy to determine the role of high dose therapy in recurrent Wilmsâ€™ tumour. European Journal of Cancer, 2013, 49, 194-210.	2.8	61
64	Results of a Multicentre Survey Evaluating Clinical Practice of Port and Broviac Management in Paediatric Oncology. Klinische Padiatrie, 2013, 225, 145-151.	0.6	19
65	Clear cell sarcoma of the kidney: A review. European Journal of Cancer, 2012, 48, 2219-2226.	2.8	118
66	Characteristics and outcome of stage II and III non-anaplastic Wilmsâ€™ tumour treated according to the SIOP trial and study 93-01. European Journal of Cancer, 2012, 48, 3240-3248.	2.8	81
67	Multicenter study identified molecular bloodâ€born protein signatures for Wilms Tumor. International Journal of Cancer, 2012, 131, 673-682.	5.1	4
68	Tumor Biology Influences the Prognosis of Nephroblastoma Patients With Primary Pulmonary Metastases. Annals of Surgery, 2011, 254, 155-162.	4.2	46
69	Update on Relapses in Unilateral Nephroblastoma Registered in 3 Consecutive SIOP/GPOH Studies - A Report from the GPOH-Nephroblastoma Study Group. Klinische Padiatrie, 2011, 223, 113-119.	0.6	28
70	Bloodstream Infections in Paediatric Cancer Patients. Prospective Comparative Study in 2 University Hospitals. Klinische Padiatrie, 2011, 223, 335-340.	0.6	15
71	Mesoblastic Nephroma. , 2011, , 2239-2240.		0
72	A floppy neonate with respiratory failure and burst suppression EEG (Case Presentation). Acta Paediatrica, International Journal of Paediatrics, 2010, 99, 326-327.	1.5	1

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73	Secondary Neoplasms After Wilmsâ€™ Tumor in Germany. Strahlentherapie Und Onkologie, 2009, 185, 11-12.	2.0	5
74	Surgical Aspects in the Treatment of Patients With Unilateral Wilms Tumor. Annals of Surgery, 2009, 249, 666-671.	4.2	63
75	KIT, PDGFRÎ± and EGFR analysis in nephroblastoma. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2008, 452, 637-650.	2.8	6
76	Mesenchymal chondrosarcoma of soft tissues and bone in children, adolescents, and young adults. Cancer, 2008, 112, 2424-2431.	4.1	133
77	New prognostic markers revealed by evaluation of genes correlated with clinical parameters in Wilms tumors. Genes Chromosomes and Cancer, 2008, 47, 386-395.	2.8	48
78	Human cytomegalovirus protein pp65: an efficient protein carrier system into human dendritic cells. Gene Therapy, 2008, 15, 318-325.	4.5	7
79	Surgical implications for liver metastases in nephroblastomaâ€”Data from the SIOP/GPOH study. Surgical Oncology, 2008, 17, 33-40.	1.6	17
80	A Clinical Perspective of Bilateral Renal Tumors - Data from Three Consecutive German Cooperative Trials. Journal of Pediatric Urology, 2008, 4, S64.	1.1	1
81	Consultation within the Nephroblastoma Trial SIOP 2001/GPOH as Part of the Workload in the Trial Office. Klinische Padiatrie, 2008, 220, 183-188.	0.6	6
82	Rhabdoid tumors in children: prognostic factors in 70 patients diagnosed in Germany. Oncology Reports, 2008, 19, 819-23.	2.6	65
83	Outcome of relapses of nephroblastoma in patients registered in the SIOP/GPOH trials and studies. Oncology Reports, 2008, 20, 463-7.	2.6	32
84	Treatment of Cystic Nephroma and Cystic Partially Differentiated Nephroblastomaâ€”A Report From the SIOP/GPOH Study Group. Journal of Urology, 2007, 177, 294-296.	0.4	36
85	Primary hepatic metastases in nephroblastomaâ€”a report of the SIOP/GPOH Study. Journal of Pediatric Surgery, 2006, 41, 168-172.	1.6	22
86	Mesoblastic nephromaâ€”A report from the Gesellschaft fur PÃdiatrische Onkologie und HÃmatologie (GPOH). Cancer, 2006, 106, 2275-2283.	4.1	111
87	Population-based study of renal cell carcinoma in children in Germany, 1980â€“2005. Cancer, 2006, 107, 2906-2914.	4.1	117
88	INTRAINDIVIDUAL PROPOFOL DOSAGE VARIABILITY IN CHILDREN UNDERGOING REPETITIVE PROCEDURAL SEDATIONS. Pediatric Hematology and Oncology, 2006, 23, 571-578.	0.8	8
89	Outcome of relapses of nephroblastoma in patients registered in the SIOP/GPOH trials and studies. Oncology Reports, 1994, 20, 463.	2.6	11
90	Rhabdoid tumors in children: Prognostic factors in 70 patients diagnosed in Germany. Oncology Reports, 0, , .	2.6	27

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91	Mesoblastic Nephroma. , 0, , 1834-1835.		0