Norbert Graf

List of Publications by Year in descending order

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426 papers 17,813 citations

69 h-index 21540 114 g-index

525 all docs 525 docs citations

525 times ranked

15873 citing authors

#	Article	IF	CITATIONS
1	Dissecting the genomic complexity underlying medulloblastoma. Nature, 2012, 488, 100-105.	27.8	765
2	Treatment of Early Childhood Medulloblastoma by Postoperative Chemotherapy Alone. New England Journal of Medicine, 2005, 352, 978-986.	27.0	682
3	Neoadjuvant chemotherapy of osteosarcoma: results of a randomized cooperative trial (COSS-82) with salvage chemotherapy based on histological tumor response Journal of Clinical Oncology, 1988, 6, 329-337.	1.6	434
4	Toward the blood-borne miRNome of human diseases. Nature Methods, 2011, 8, 841-843.	19.0	339
5	Long-term results of the co-operative German–Austrian–Swiss osteosarcoma study group's protocol COSS-86 of intensive multidrug chemotherapy and surgery for osteosarcoma of the limbs. Annals of Oncology, 1998, 9, 893-899.	1.2	304
6	The impact of the methotrexate administration schedule and dose in the treatment of children and adolescents with B-cell neoplasms: a report of the BFM Group Study NHL-BFM95. Blood, 2004, 105, 948-958.	1.4	304
7	Improved treatment results in childhood B-cell neoplasms with tailored intensification of therapy: A report of the Berlin-Frankfurt-Münster Group Trial NHL-BFM 90. Blood, 1999, 94, 3294-306.	1.4	303
8	Advances in Wilms Tumor Treatment and Biology: Progress Through International Collaboration. Journal of Clinical Oncology, 2015, 33, 2999-3007.	1.6	281
9	Rationale for the treatment of Wilms tumour in the UMBRELLA SIOP–RTSG 2016 protocol. Nature Reviews Urology, 2017, 14, 743-752.	3.8	249
10	Mutations in the SIX1/2 Pathway and the DROSHA/DGCR8 miRNA Microprocessor Complex Underlie High-Risk Blastemal Type Wilms Tumors. Cancer Cell, 2015, 27, 298-311.	16.8	248
11	Obesity after childhood craniopharyngioma - German multicenter study on pre-operative risk factors and quality of life. Klinische Padiatrie, 2001, 213, 244-249.	0.6	242
12	Childhood cancer predisposition syndromes—A concise review and recommendations by the Cancer Predisposition Working Group of the Society for Pediatric Oncology and Hematology. American Journal of Medical Genetics, Part A, 2017, 173, 1017-1037.	1.2	200
13	Malignant renal tumours incidence and survival in European children (1978–1997): Report from the Automated Childhood Cancer Information System project. European Journal of Cancer, 2006, 42, 2103-2114.	2.8	197
14	Reduction of postoperative chemotherapy in children with stage I intermediate-risk and anaplastic Wilms' tumour (SIOP 93-01 trial): a randomised controlled trial. Lancet, The, 2004, 364, 1229-1235.	13.7	191
15	Making sense of big data in health research: Towards an EU action plan. Genome Medicine, 2016, 8, 71.	8.2	190
16	THE ROLE OF PREOPERATIVE CHEMOTHERAPY IN THE MANAGEMENT OF WILMS' TUMOR. Urologic Clinics of North America, 2000, 27, 443-454.	1.8	184
17	Less Toxicity by Optimizing Chemotherapy, but Not by Addition of Granulocyte Colony-Stimulating Factor in Children and Adolescents With Acute Myeloid Leukemia: Results of AML-BFM 98. Journal of Clinical Oncology, 2006, 24, 4499-4506.	1.6	173
18	Methotrexate pharmacokinetics and prognosis in osteosarcoma Journal of Clinical Oncology, 1994, 12, 1443-1451.	1.6	166

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19	Omission of doxorubicin from the treatment of stage II–III, intermediate-risk Wilms' tumour (SIOP WT) Tj ETQq1	. 1.0.7843 13.7	14 rgBT /0
20	Newcastle disease virotherapy induces longâ€term survival and tumorâ€specific immune memory in orthotopic glioma through the induction of immunogenic cell death. International Journal of Cancer, 2015, 136, E313-25.	5.1	165
21	Effect of intraarterial versus intravenous cisplatin in addition to systemic doxorubicin, high-dose methotrexate, and ifosfamide on histologic tumor response in osteosarcoma (study COSS-86). Cancer, 1990, 66, 1703-1710.	4.1	155
22	Constitutional 11p15 abnormalities, including heritable imprinting center mutations, cause nonsyndromic Wilms tumor. Nature Genetics, 2008, 40, 1329-1334.	21.4	154
23	The UMBRELLA SIOP–RTSG 2016 Wilms tumour pathology and molecular biology protocol. Nature Reviews Urology, 2018, 15, 693-701.	3.8	152
24	Characteristics and survival of 750 children diagnosed with a renal tumor in the first seven months of life: A collaborative study by the SIOP/GPOH/SFOP, NWTSG, and UKCCSG Wilms tumor study groups. Pediatric Blood and Cancer, 2008, 50, 1130-1134.	1.5	151
25	Randomized trial comparing liposomal daunorubicin with idarubicin as induction for pediatric acute myeloid leukemia: results from Study AML-BFM 2004. Blood, 2013, 122, 37-43.	1.4	151
26	Clinical impact of histologic subtypes in localized non-anaplastic nephroblastoma treated according to the trial and study SIOP-9/GPOH. Annals of Oncology, 2001, 12, 311-319.	1.2	144
27	Multi-omics enrichment analysis using the GeneTrail2 web service. Bioinformatics, 2016, 32, 1502-1508.	4.1	144
28	Wilms tumour: prognostic factors, staging, therapy and late effects. Pediatric Radiology, 2008, 38, 2-17.	2.0	140
29	Factor VII deficiency: clinical manifestation of 717 subjects from Europe and Latin America with mutations in the factor 7 gene. Haemophilia, 2009, 15, 267-280.	2.1	132
30	US, CT and MR imaging characteristics of nephroblastomatosis. Pediatric Radiology, 1998, 28, 435-443.	2.0	127
31	Wilms' Tumor in Adults: Results of the Society of Pediatric Oncology (SIOP) 93-01/Society for Pediatric Oncology and Hematology (GPOH) Study. Journal of Clinical Oncology, 2004, 22, 4500-4506.	1.6	126
32	Treatment of early childhood medulloblastoma by postoperative chemotherapy and deferred radiotherapy. Neuro-Oncology, 2009, 11, 201-210.	1.2	125
33	Malignant rhabdoid tumours of the kidney (MRTKs), registered on recent SIOP protocols from 1993 to 2005: A report of the SIOP renal tumour study group. Pediatric Blood and Cancer, 2011, 56, 733-737.	1.5	125
34	Characterization of the chromosomal translocation $t(10;17)(q22;p13)$ in clear cell sarcoma of kidney. Journal of Pathology, 2012, 227, 72-80.	4.5	125
35	Risk Stratification for Wilms Tumor: Current Approach and Future Directions. American Society of Clinical Oncology Educational Book / ASCO American Society of Clinical Oncology Meeting, 2014, , 215-223.	3.8	124
36	Results of the SIOP 93-01/GPOH Trial and Study for the Treatment of Patients with Unilateral Nonmetastatic Wilms Tumor. Klinische Padiatrie, 2004, 216, 132-140.	0.6	123

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37	Definition of a standard-risk group in children with AML. British Journal of Haematology, 1999, 104, 630-639.	2.5	122
38	Clear cell sarcoma of the kidney: A review. European Journal of Cancer, 2012, 48, 2219-2226.	2.8	118
39	Population-based study of renal cell carcinoma in children in Germany, 1980–2005. Cancer, 2006, 107, 2906-2914.	4.1	117
40	The COVIDâ€19 pandemic: A rapid global response for children with cancer from SIOP, COG, SIOPâ€E, SIOPâ€PODC, IPSO, PROS, CCI, and St Jude Global. Pediatric Blood and Cancer, 2020, 67, e28409.	1.5	113
41	Survival in nephroblastoma treated according to the trial and study SIOP-9/GPOH with respect to relapse and morbidity. Annals of Oncology, 2004, 15, 808-820.	1.2	112
42	Mesoblastic nephroma—A report from the Gesellschaft fur PÃ d iatrische Onkologie und HÃ m atologie (GPOH). Cancer, 2006, 106, 2275-2283.	4.1	111
43	Pathogen response-like recruitment and activation of neutrophils by sterile immunogenic dying cells drives neutrophil-mediated residual cell killing. Cell Death and Differentiation, 2017, 24, 832-843.	11.2	111
44	The Pediatric Precision Oncology INFORM Registry: Clinical Outcome and Benefit for Patients with Very High-Evidence Targets. Cancer Discovery, 2021, 11, 2764-2779.	9.4	110
45	Down's syndrome in childhood acute lymphoblastic leukemia: clinical characteristics and treatment outcome in four consecutive BFM trials. Leukemia, 1998, 12, 645-651.	7.2	106
46	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
47	Gain of 1q As a Prognostic Biomarker in Wilms Tumors (WTs) Treated With Preoperative Chemotherapy in the International Society of Paediatric Oncology (SIOP) WT 2001 Trial: A SIOP Renal Tumours Biology Consortium Study. Journal of Clinical Oncology, 2016, 34, 3195-3203.	1.6	105
48	Complementary and alternative treatment methods in children with cancer: A population-based retrospective survey on the prevalence of use in Germany. European Journal of Cancer, 2008, 44, 2233-2240.	2.8	100
49	<i>TP53</i> Mutation Is Frequently Associated With <i>CTNNB1</i> Mutation or <i>MYCN</i> Amplification and Is Compatible With Long-Term Survival in Medulloblastoma. Journal of Clinical Oncology, 2010, 28, 5188-5196.	1.6	100
50	Laser acupuncture in children with headache: A double-blind, randomized, bicenter, placebo-controlled trial. Pain, 2008, 137, 405-412.	4.2	97
51	Clinical and molecular features in patients with atypical teratoid rhabdoid tumor or malignant rhabdoid tumor. Genes Chromosomes and Cancer, 2010, 49, 176-181.	2.8	96
52	Treatment of Pulmonary Metastases in Children With Stage IV Nephroblastoma With Risk-Based Use of Pulmonary Radiotherapy. Journal of Clinical Oncology, 2012, 30, 3533-3539.	1.6	95
53	Preradiation chemotherapy for pediatric patients with high-grade glioma. Cancer, 2002, 94, 264-271.	4.1	91
54	Preradiation chemotherapy of children and young adults with malignant brain tumors: Results of the german pilot trial HIT'88/89 Klinische Padiatrie, 1998, 210, 227-233.	0.6	90

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55	Incidence of tuberous sclerosis and age at first diagnosis: new data and emerging trends from a national, prospective surveillance study. Orphanet Journal of Rare Diseases, 2018, 13, 117.	2.7	86
56	Expression profiling of Wilms tumors reveals new candidate genes for different clinical parameters. International Journal of Cancer, 2006, 118, 1954-1962.	5.1	85
57	Nephron sparing surgery (NSS) for unilateral wilms tumor (UWT): The SIOP 2001 experience. Pediatric Blood and Cancer, 2014, 61, 2175-2179.	1.5	85
58	Multiple mechanisms of MYCN dysregulation in Wilms tumour. Oncotarget, 2015, 6, 7232-7243.	1.8	85
59	Congenital mesoblastic nephroma 50 years after its recognition: A narrative review. Pediatric Blood and Cancer, 2017, 64, e26437.	1.5	84
60	Partial Nephrectomy for Unilateral Wilms Tumor: Results of Study SIOP 93–01/GPOH. Journal of Urology, 2003, 170, 939-944.	0.4	82
61	Target genes of the WNT/l²-catenin pathway in Wilms tumors. Genes Chromosomes and Cancer, 2006, 45, 565-574.	2.8	82
62	Acupuncture to Alleviate Chemotherapy-induced Nausea and Vomiting in Pediatric Oncology – A Randomized Multicenter Crossover Pilot Trial. Klinische Padiatrie, 2008, 220, 365-370.	0.6	81
63	Characteristics and outcome of stage II and III non-anaplastic Wilms' tumour treated according to the SIOP trial and study 93-01. European Journal of Cancer, 2012, 48, 3240-3248.	2.8	81
64	Treatment options in childhood pontine gliomas. Journal of Neuro-Oncology, 2006, 79, 281-287.	2.9	78
65	IDA-FLAG (idarubicin, fludarabine, cytarabine, G-CSF), an effective remission-induction therapy for poor-prognosis AML of childhood prior to allogeneic or autologous bone marrow transplantation: experiences of a phase II trial. British Journal of Haematology, 1998, 102, 647-655.	2.5	77
66	Treatment of relapsed Wilms tumors: lessons learned. Expert Review of Anticancer Therapy, 2009, 9, 1807-1815.	2.4	77
67	Initial presenting manifestations in 16,486 patients with inborn errors of immunity include infections and noninfectious manifestations. Journal of Allergy and Clinical Immunology, 2021, 148, 1332-1341.e5.	2.9	75
68	Allele loss in Wilms tumors of chromosome arms 11q, 16q, and 22q correlates with clinicopathological parameters. , 1998, 22, 287-294.		74
69	High-dose chemotherapy with autologous stem cell rescue in children with nephroblastoma. Bone Marrow Transplantation, 2002, 30, 893-898.	2.4	73
70	Use of complementary and alternative medicine in healthy children and children with chronic medical conditions in Germany. Complementary Therapies in Medicine, 2013, 21, S61-S69.	2.7	73
71	Improved 6â€year overall survival in <scp>AT</scp> / <scp>RT</scp> â€" results of the registry study Rhabdoid 2007. Cancer Medicine, 2016, 5, 1765-1775.	2.8	73
72	Recurrent intragenic rearrangements of EGFR and BRAF in soft tissue tumors of infants. Nature Communications, 2018, 9, 2378.	12.8	72

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73	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
74	Sorafenib and cisplatin/doxorubicin (PLADO) in pediatric hepatocellular carcinoma. Pediatric Blood and Cancer, 2012, 58, 539-544.	1.5	71
75	The German National Registry of Primary Immunodeficiencies (2012–2017). Frontiers in Immunology, 2019, 10, 1272.	4.8	71
76	miRNA Profiles as a Predictor of Chemoresponsiveness in Wilms' Tumor Blastema. PLoS ONE, 2013, 8, e53417.	2.5	71
77	Effective childhood cancer treatment: The impact of large scale clinical trials in Germany and Austria. Pediatric Blood and Cancer, 2013, 60, 1574-1581.	1.5	70
78	Subtype-Specific <i>FBXW7</i> Mutation and <i>MYCN</i> Copy Number Gain in Wilms' Tumor. Clinical Cancer Research, 2010, 16, 2036-2045.	7.0	69
79	Outcome of localised blastemal-type Wilms tumour patients treated according to intensified treatment in the SIOP WT 2001 protocol, a report of the SIOP Renal Tumour Study Group (SIOP-RTSG). European Journal of Cancer, 2015, 51, 498-506.	2.8	67
80	Rhabdoid tumors in children: prognostic factors in 70 patients diagnosed in Germany. Oncology Reports, 2008, 19, 819-23.	2.6	65
81	Surgical Aspects in the Treatment of Patients With Unilateral Wilms Tumor. Annals of Surgery, 2009, 249, 666-671.	4.2	63
82	Surgery of cavoatrial tumor thrombus in nephroblastoma: A report of the SIOP/GPOH study. Pediatric Blood and Cancer, 2004, 43, 40-45.	1.5	62
83	DNA Repair Alterations in Children With Pediatric Malignancies: Novel Opportunities to Identify Patients at Risk for High-Grade Toxicities. International Journal of Radiation Oncology Biology Physics, 2010, 78, 359-369.	0.8	61
84	An international strategy to determine the role of high dose therapy in recurrent Wilms' tumour. European Journal of Cancer, 2013, 49, 194-210.	2.8	61
85	Minimally invasive nephrectomy for Wilms tumors in children – data from SIOP 2001. Journal of Pediatric Surgery, 2014, 49, 1544-1548.	1.6	61
86	Combining miRNA and mRNA Expression Profiles in Wilms Tumor Subtypes. International Journal of Molecular Sciences, 2016, 17, 475.	4.1	61
87	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
88	Relapse of Wilms' tumour and detection methods: a retrospective analysis of the 2001 Renal Tumour Study Group–International Society of Paediatric Oncology Wilms' tumour protocol database. Lancet Oncology, The, 2018, 19, 1072-1081.	10.7	59
89	Circulating serum miRNAs as potential biomarkers for nephroblastoma. Pediatric Blood and Cancer, 2015, 62, 1360-1367.	1.5	56
90	Mutually exclusive <i>BCOR</i> internal tandem duplications and <i>YWHAEâ€NUTM2</i> fusions in clear cell sarcoma of kidney: not the full story. Journal of Pathology, 2016, 238, 617-620.	4.5	56

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91	<i>WTX</i> inactivation is a frequent, but late event in Wilms tumors without apparent clinical impact. Genes Chromosomes and Cancer, 2009, 48, 1102-1111.	2.8	54
92	mHealth and telemedicine apps: in search of a common regulation. Ecancermedicalscience, 2018, 12, 853.	1.1	54
93	The contribution of chest CT-scan at diagnosis in children with unilateral Wilms' tumour. Results of the SIOP 2001 study. European Journal of Cancer, 2012, 48, 1060-1065.	2.8	53
94	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
95	Genotyping circulating tumor DNA of pediatric Hodgkin lymphoma. Leukemia, 2020, 34, 151-166.	7.2	53
96	Propofol Versus Midazolam/Ketamine for Procedural Sedation in Pediatric Oncology. Journal of Pediatric Hematology/Oncology, 2005, 27, 471-476.	0.6	52
97	Clinical and Molecular Characterization of Patients with Heterozygous Mutations in Wilms Tumor Suppressor Gene 1. Clinical Journal of the American Society of Nephrology: CJASN, 2015, 10, 825-831.	4.5	52
98	Loss of 11q and 16q in Wilms tumors is associated with anaplasia, tumor recurrence, and poor prognosis. Genes Chromosomes and Cancer, 2007, 46, 163-170.	2.8	51
99	Acute pancreatitis induced by short-term propofol administration. Paediatric Anaesthesia, 2005, 15, 1006-1008.	1.1	50
100	Clinical relevance of mutations in the Wilms tumor suppressor 1 gene <i>WT1</i> and the cadherinâ€associated protein β1 gene <i>CTNNB1</i> for patients with Wilms tumors. Cancer, 2008, 113, 1080-1089.	4.1	50
101	Current Concepts in Surgery for Wilms Tumor—The Risk and Function-Adapted Strategy. European Journal of Pediatric Surgery, 2014, 24, 457-460.	1.3	50
102	Incidence, Trends, and Survival of Children With Embryonal Tumors. Pediatrics, 2015, 136, e623-e632.	2.1	50
103	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
104	Acute leukemia with chromosome translocation (4; 11): 7 new patients and analysis of 71 cases. Blut, 1987, 54, 325-335.	1.2	48
105	Two infants with life-threatening diffuse neonatal hemangiomatosis treated with cyclophosphamide. Pediatric Blood and Cancer, 2006, 46, 239-242.	1.5	48
106	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
107	The ACGT Master Ontology and its applications – Towards an ontology-driven cancer research and management system. Journal of Biomedical Informatics, 2011, 44, 8-25.	4.3	47
108	Impact of chemotherapy on disseminated lowâ€grade glioma in children and adolescents: Report from the HITâ€LGG 1996 trial. Pediatric Blood and Cancer, 2011, 56, 1046-1054.	1.5	47

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109	Hepatotoxicity in patients treated according to the Nephroblastoma Trial and Study SIOP-9/GPOH., 1999, 33, 462-469.		46
110	Tumor Biology Influences the Prognosis of Nephroblastoma Patients With Primary Pulmonary Metastases. Annals of Surgery, 2011, 254, 155-162.	4.2	46
111	iManageCancer: Developing a Platform for Empowering Patients and Strengthening Self-Management in Cancer Diseases. , 2017, , .		45
112	<i>TRIM28</i> haploinsufficiency predisposes to Wilms tumor. International Journal of Cancer, 2019, 145, 941-951.	5.1	45
113	All-trans retinoic acid treatment of Wilms tumor cells reverses expression of genes associated with high risk and relapse in vivo. Oncogene, 2005, 24, 5246-5251.	5.9	44
114	Assessing the risk of mortality in paediatric cancer patients admitted to the paediatric intensive care unit: a novel risk score?. European Journal of Pediatrics, 2005, 164, 563-567.	2.7	44
115	Improving data and knowledge management to better integrate health care and research. Journal of Internal Medicine, 2013, 274, 321-328.	6.0	44
116	Incidence and outcomes of patients with late recurrence of Wilms' tumor. Pediatric Blood and Cancer, 2013, 60, 1612-1615.	1.5	43
117	Rationale for the treatment of children with CCSK in the UMBRELLA SIOP–RTSG 2016 protocol. Nature Reviews Urology, 2018, 15, 309-319.	3.8	43
118	Networking for Children and Adolescents with Very Rare Tumors: Foundation of the GPOH Pediatric Rare Tumor Group. Klinische Padiatrie, 2009, 221, 181-185.	0.6	42
119	<i>ETV6</i> – <i>NTRK3</i> in congenital mesoblastic nephroma: A report of the SIOP/GPOH nephroblastoma study. Pediatric Blood and Cancer, 2018, 65, e26925.	1.5	41
120	Expression of serologically identified tumor antigens in acute leukemias. Leukemia Research, 2003, 27, 655-660.	0.8	37
121	Management of adults with Wilmsâ \in ^{\mathbb{M}} tumor: recommendations based on international consensus. Expert Review of Anticancer Therapy, 2011, 11, 1107-1115.	2.4	37
122	Multicentre prospective observational study on professional wound care using honey (Medihoneyâ,,¢). International Wound Journal, 2013, 10, 252-259.	2.9	37
123	Treatment of Cystic Nephroma and Cystic Partially Differentiated Nephroblastoma—A Report From the SIOP/GPOH Study Group. Journal of Urology, 2007, 177, 294-296.	0.4	36
124	Role of MRI in the management of patients with nephroblastoma. European Radiology, 2008, 18, 683-691.	4.5	36
125	New approaches to risk stratification for Wilms tumor. Current Opinion in Pediatrics, 2021, 33, 40-48.	2.0	36
126	Evidence for a delay in diagnosis of Wilms' tumour in the UK compared with Germany: implications for primary care for children. Archives of Disease in Childhood, 2016, 101, 417-420.	1.9	35

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127	Highâ€dose treatment for malignant rhabdoid tumor of the kidney: No evidence for improved survivalâ€"The Gesellschaft fÃ⅓r PÃ d iatrische Onkologie und HÃ m atologie (GPOH) experience. Pediatric Blood and Cancer, 2018, 65, e26746.	1.5	35
128	Patient empowerment for cancer patients through a novel ICT infrastructure. Journal of Biomedical Informatics, 2020, 101, 103342.	4.3	35
129	Effects of short-term propofol administration on pancreatic enzymes and triglyceride levels in children. Anaesthesia, 2005, 60, 660-663.	3.8	34
130	Feasibility of Intensive Multimodal Therapy in Infants Affected by Rhabdoid Tumors – Experience of the EU-RHAB registry. Klinische Padiatrie, 2014, 226, 143-148.	0.6	33
131	Prognostic significance of age in 5631 patients with Wilms tumour prospectively registered in International Society of Paediatric Oncology (SIOP) 93-01 and 2001. PLoS ONE, 2019, 14, e0221373.	2.5	33
132	The BFM-protocol for HIV-negative Burkitt's lymphomas and L3 ALL in adult patients: a high chance for cure. Annals of Hematology, 1992, 65, 201-205.	1.8	32
133	First experience of the AMLâ€Berlinâ€Frankfurtâ€MÃ⅓nster group in pediatric patients with standardâ€risk acute promyelocytic leukemia treated with arsenic trioxide and allâ€ <i>trans</i> retinoid acid. Pediatric Blood and Cancer, 2017, 64, e26461.	1.5	32
134	Outcome of relapses of nephroblastoma in patients registered in the SIOP/GPOH trials and studies. Oncology Reports, 2008, 20, 463-7.	2.6	32
135	Pulmonary Dysfunction in Pediatric Oncology Patients. Pediatric Hematology and Oncology, 2004, 21, 175-195.	0.8	31
136	Clinically driven design of multi-scale cancer models: the ContraCancrum project paradigm. Interface Focus, 2011, 1, 450-461.	3.0	31
137	Nephroblastoma: does the decrease in tumor volume under preoperative chemotherapy predict the lymph nodes status at surgery?. Pediatric Blood and Cancer, 2011, 57, 1266-1269.	1.5	31
138	Paediatric renal tumours: perspectives from the SIOP–RTSG. Nature Reviews Urology, 2017, 14, 3-4.	3.8	31
139	The Technologically Integrated Oncosimulator: Combining Multiscale Cancer Modeling With Information Technology in the In Silico Oncology Context. IEEE Journal of Biomedical and Health Informatics, 2014, 18, 840-854.	6.3	30
140	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
141	Clinical characteristics and outcomes of children with WAGR syndrome and Wilms tumor and/or nephroblastomatosis: The 30â€year SIOPâ€RTSG experience. Cancer, 2021, 127, 628-638.	4.1	30
142	Treatment-independent miRNA signature in blood of wilms tumor patients. BMC Genomics, 2012, 13, 379.	2.8	29
143	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
144	Rare malignant pediatric tumors registered in the German Childhood Cancer Registry 2001–2010. Pediatric Blood and Cancer, 2014, 61, 1202-1209.	1.5	29

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145	Characteristics and Outcome of Children with Renal Cell Carcinoma: A Narrative Review. Cancers, 2020, 12, 1776.	3.7	29
146	Arterial embolization of a secondary aneurysmatic bone cyst of the thoracic spine prior to surgical excision in a 15-year-old girl. European Journal of Radiology, 2002, 43, 79-81.	2.6	28
147	Ocular symptoms in children treated with human-mouse chimeric anti-GD2 mAb ch14.18 for neuroblastoma. Cancer Immunology, Immunotherapy, 2002, 51, 107-110.	4.2	28
148	The "Oncosimulator": a multilevel, clinically oriented simulation system of tumor growth and organism response to therapeutic schemes. Towards the clinical evaluation of in silico oncology. Annual International Conference of the IEEE Engineering in Medicine and Biology Society, 2007, 2007, 6629-32.	0.5	28
149	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
150	New insights into the genetics of glioblastoma multiforme by familial exome sequencing. Oncotarget, 2015, 6, 5918-5931.	1.8	28
151	Rhabdoid tumors in children: Prognostic factors in 70 patients diagnosed in Germany. Oncology Reports, 0, , .	2.6	27
152	Amplicons on chromosome 12q13â€21 in glioblastoma recurrences. International Journal of Cancer, 2010, 126, 2594-2602.	5.1	27
153	Knowledge engineering for health: A new discipline required to bridge the "ICT gap―between research and healthcare. Human Mutation, 2012, 33, 797-802.	2.5	27
154	DNA-Damage Foci to Detect and Characterize DNA Repair Alterations in Children Treated for Pediatric Malignancies. PLoS ONE, 2014, 9, e91319.	2.5	27
155	Semantic biomedical resource discovery: a Natural Language Processing framework. BMC Medical Informatics and Decision Making, 2015, 15, 77.	3.0	27
156	Integrated meta-omic analyses of the gastrointestinal tract microbiome in patients undergoing allogeneic hematopoietic stem cell transplantation. Translational Research, 2017, 186, 79-94.e1.	5.0	27
157	Prognostic Factors for Wilms Tumor Recurrence: A Review of the Literature. Cancers, 2021, 13, 3142.	3.7	27
158	Preoperative chemotherapy and local stage III in nephroblastoma. Translational Pediatrics, 2014, 3, 4-11.	1.2	27
159	Cellular drug resistance in acute myeloid leukemia: literature review and preliminary analysis of an ongoing collaborative study. Klinische Padiatrie, 1999, 211, 239-244.	0.6	26
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