Kathryn Pritchard-Jones

List of Publications by Year in descending order

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308 papers

17,783 citations

72 h-index 22488 117 g-index

323 all docs 323 docs citations

times ranked

323

15674 citing authors

#	Article	IF	Citations
1	International variation in childhood cancer mortality rates from 2001 to 2015: Comparison of trends in the International Cancer Benchmarking Partnership countries. International Journal of Cancer, 2022, 150, 28-37.	2.3	6
2	Surgical management, staging, and outcomes of Wilms tumours with intravascular extension: Results of the IMPORT study. Journal of Pediatric Surgery, 2022, 57, 572-578.	0.8	7
3	The epithelial splicing regulator <i>ESRP2</i> is epigenetically repressed by DNA hypermethylation in Wilms tumour and acts as a tumour suppressor. Molecular Oncology, 2022, 16, 630-647.	2.1	3
4	Long-term kidney function in children with Wilms tumour and constitutional WT1 pathogenic variant. Pediatric Nephrology, 2022, 37, 821-832.	0.9	5
5	Impact of the COVIDâ€19 pandemic on pediatric oncology providers globally: A mixedâ€methods study. Cancer, 2022, 128, 1493-1502.	2.0	17
6	Characteristics and outcomes of preoperatively treated patients with anaplastic Wilms tumors registered in the UK SIOPâ€WTâ€2001 and IMPORT study cohorts (2002â€2020). Cancer, 2022, 128, 1666-1675	2.0	6
7	Outcome of SIOP patients with low- or intermediate-risk Wilms tumour relapsing after initial vincristine and actinomycin-D therapy only Ⱐthe SIOP 93–01 and 2001 protocols. European Journal of Cancer, 2022, 163, 88-97.	1.3	8
8	SIOP Strategy 2021–2025: Cure for more, care for all. Pediatric Blood and Cancer, 2022, 69, e29577.	0.8	2
9	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.	1.3	0
10	A Summary of the Inaugural WHO Classification of Pediatric Tumors: Transitioning from the Optical into the Molecular Era. Cancer Discovery, 2022, 12, 331-355.	7.7	70
11	Loss associated with subtractive health service change: The case of specialist cancer centralization in England. Journal of Health Services Research and Policy, 2022, 27, 301-312.	0.8	6
12	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.	0.8	14
13	How we approach paediatric renal tumour core needle biopsy in the setting of preoperative chemotherapy: A Review from the SIOP Renal Tumour Study Group. Pediatric Blood and Cancer, 2022, 69, e29702.	0.8	9
14	The clinical impact of observer variability in lung nodule classification in children with Wilms tumour. Pediatric Blood and Cancer, 2022, 69, .	0.8	2
15	Implementing major system change in specialist cancer surgery: The role of provider networks. Journal of Health Services Research and Policy, 2021, 26, 4-11.	0.8	11
16	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.	2.0	30
17	Outcome of Stage IV Completely Necrotic Wilms Tumour and Local Stage III Treated According to the SIOP 2001 Protocol. Cancers, 2021, 13, 976.	1.7	6
18	Characteristics and outcome of pediatric renal cell carcinoma patients registered in the International Society of Pediatric Oncology (<scp>SIOP</scp>) 93â€01, 2001 and <scp>UKâ€IMPORT</scp> database: A report of the <scp>SIOPâ€Renal</scp> Tumor Study Group. International Journal of Cancer, 2021, 148, 2724-2735.	2.3	26

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19	The Global COVIDâ€19 Observatory and Resource Center for Childhood Cancer: A response for the pediatric oncology community by SIOP and St. Jude Global. Pediatric Blood and Cancer, 2021, 68, e28962.	0.8	8
20	The threat of the COVID-19 pandemic on reversing global life-saving gains in the survival of childhood cancer: a call for collaborative action from SIOP, IPSO, PROS, WCC, CCI, St Jude Global, UICC and WHPCA. Ecancermedicalscience, 2021, 15, 1187.	0.6	4
21	Predisposition to cancer in children and adolescents. The Lancet Child and Adolescent Health, 2021, 5, 142-154.	2.7	53
22	Global effect of the COVID-19 pandemic on paediatric cancer care: a cross-sectional study. The Lancet Child and Adolescent Health, 2021, 5, 332-340.	2.7	83
23	How to Cost the Implementation of Major System Change for Economic Evaluations: Case Study Using Reconfigurations of Specialist Cancer Surgery in Part of London, England. Applied Health Economics and Health Policy, 2021, 19, 797-810.	1.0	4
24	Comparative analysis of the clinical characteristics and outcomes of patients with Wilms tumor in the United Kingdom and Japan. Pediatric Blood and Cancer, 2021, 68, e29143.	0.8	7
25	Staging childhood cancers in Europe: Application of the Toronto stage principles for neuroblastoma and Wilms tumour. The JARC pilot study. Pediatric Blood and Cancer, 2021, 68, e29020.	0.8	7
26	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€CLG and GPOH studies. International Journal of Cancer, 2021, 149, 1332-1340.	2.3	16
27	Oral etoposide as a single agent in childhood and young adult cancer in England: Still a poorly evaluated palliative treatment. Pediatric Blood and Cancer, 2021, 68, e29204.	0.8	2
28	Wilms tumour surveillance in at-risk children: Literature review and recommendations from the SIOP-Europe Host Genome Working Group and SIOP Renal Tumour Study Group. European Journal of Cancer, 2021, 153, 51-63.	1.3	25
29	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.	0.6	14
30	Pediatric cancer care in Africa: SIOP Global MappingÂProgramÂreport on economic and population indicators. Pediatric Blood and Cancer, 2021, 68, e29345.	0.8	10
31	Global characteristics and outcomes of SARS-CoV-2 infection in children and adolescents with cancer (GRCCC): a cohort study. Lancet Oncology, The, 2021, 22, 1416-1426.	5.1	93
32	The multidisciplinary, theory-based co-design of a new digital health intervention supporting the care of oesophageal cancer patients. Digital Health, 2021, 7, 205520762110384.	0.9	7
33	Wilms tumour. Nature Reviews Disease Primers, 2021, 7, 75.	18.1	75
34	Is radiotherapy required in firstâ€line treatment of stage I diffuse anaplastic Wilms tumor? A report of SIOPâ€RTSG, AIEOP, JWiTS, and UKCCSG. Pediatric Blood and Cancer, 2020, 67, e28039.	0.8	14
35	Characteristics and Outcome of Children with Renal Cell Carcinoma: A Narrative Review. Cancers, 2020, 12, 1776.	1.7	29
36	Working Together to Build a Better Future for Children With Cancer in Africa. JCO Global Oncology, 2020, 6, 1076-1078.	0.8	13

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37	Pediatric and young adult renal cell carcinoma. Pediatric Blood and Cancer, 2020, 67, e28675.	0.8	14
38	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.	0.8	113
39	Incidence of childhood renal tumours: An international populationâ€based study. International Journal of Cancer, 2020, 147, 3313-3327.	2.3	73
40	Early advice on managing children with cancer during the COVIDâ€19 pandemic and a call for sharing experiences. Pediatric Blood and Cancer, 2020, 67, e28327.	0.8	93
41	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.	1.3	24
42	An organoid biobank for childhood kidney cancers that captures disease and tissue heterogeneity. Nature Communications, 2020, 11, 1310.	5.8	183
43	The role of imaging in the initial investigation of paediatric renal tumours. The Lancet Child and Adolescent Health, 2020, 4, 232-241.	2.7	25
44	Sustainable care for children with cancer: a Lancet Oncology Commission. Lancet Oncology, The, 2020, 21, e185-e224.	5.1	177
45	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.	1.1	33
46	The genetic changes of Wilms tumour. Nature Reviews Nephrology, 2019, 15, 240-251.	4.1	159
47	Highlights from the 13th African Continental Meeting of the International Society of Paediatric Oncology (SIOP), 6–9 March 2019, Cairo, Egypt. Ecancermedicalscience, 2019, 13, 932.	0.6	6
48	The European study on centralisation of childhood cancer treatment. European Journal of Cancer, 2019, 115, 120-127.	1.3	12
49	Evaluation of needle biopsy as a potential risk factor for local recurrence of Wilms tumour in the SIOP WT 2001 trial. European Journal of Cancer, 2019, 116, 13-20.	1.3	24
50	Reply to the Letter to the Editor: Renal tumors in children older than 10 yearsâ€"Should we be doing upfront nephrectomy?. Pediatric Blood and Cancer, 2019, 66, e27760.	0.8	0
51	Stage at diagnosis for childhood solid cancers in Australia: A population-based study. Cancer Epidemiology, 2019, 59, 208-214.	0.8	12
52	Science and health for all children with cancer. Science, 2019, 363, 1182-1186.	6.0	200
53	WARNING: Gâ€401 and SKâ€NEPâ€1 cell lines are not Wilms tumor cell lines. Pediatric Blood and Cancer, 2019, 66, e27741.	0.8	10
54	Comment on: "Indications and results of diagnostic biopsy in pediatric renal tumors: A retrospective analysis of 317 patients with critical review of SIOP guidelines― Pediatric Blood and Cancer, 2019, 66, e27746.	0.8	1

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55	The diagnostic accuracy and clinical utility of pediatric renal tumor biopsy: Report of the UK experience in the SIOP UK WT 2001 trial. Pediatric Blood and Cancer, 2019, 66, e27627.	0.8	41
56	Embryonal precursors of Wilms tumor. Science, 2019, 366, 1247-1251.	6.0	101
57	Pharmacotherapeutic Management of Wilms Tumor: An Update. Paediatric Drugs, 2019, 21, 1-13.	1.3	23
58	Outcomes of non-anaplastic stage III and †inoperable' Wilms tumour treated in the UKW3 trial. Radiotherapy and Oncology, 2019, 131, 1-7.	0.3	7
59	Evidence-based data and rare cancers: The need for a new methodological approach in research and investigation. European Journal of Surgical Oncology, 2019, 45, 22-30.	0.5	16
60	Rationale for the treatment of children with CCSK in the UMBRELLA SIOP–RTSG 2016 protocol. Nature Reviews Urology, 2018, 15, 309-319.	1.9	43
61	Childhood cancer incidence and survival in Japan and England: A populationâ€based study (1993â€⊋010). Cancer Science, 2018, 109, 422-434.	1.7	73
62	Evaluation of boost irradiation in patients with intermediateâ€risk stage III Wilms tumour with positive lymph nodes only: Results from the SIOPâ€WTâ€2001 Registry. Pediatric Blood and Cancer, 2018, 65, e27085.	0.8	10
63	Irinotecan for relapsed Wilms tumor in pediatric patients: SIOP experience and review of the literature—A report from the SIOP Renal Tumor Study Group. Pediatric Blood and Cancer, 2018, 65, e26849.	0.8	11
64	Follow-up surveillance of Wilm's tumour – Authors' reply. Lancet Oncology, The, 2018, 19, e503.	5.1	0
65	The UMBRELLA SIOP–RTSG 2016 Wilms tumour pathology and molecular biology protocol. Nature Reviews Urology, 2018, 15, 693-701.	1.9	152
66	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.	5.1	59
67	Congenital mesoblastic nephroma 50 years after its recognition: A narrative review. Pediatric Blood and Cancer, 2017, 64, e26437.	0.8	84
68	Review of phase I and II trials for Wilms' tumour – Can we optimise the search for novel agents?. European Journal of Cancer, 2017, 79, 205-213.	1.3	25
69	Nephrogenic rests in Wilms tumors treated with preoperative chemotherapy: The UK SIOP Wilms Tumor 2001 Trial experience. Pediatric Blood and Cancer, 2017, 64, e26547.	0.8	28
70	Role of CD56 in Normal Kidney Development and Wilms Tumorigenesis. Fetal and Pediatric Pathology, 2017, 36, 62-75.	0.4	18
71	Bilateral Wilms tumour: a review of clinical and molecular features. Expert Reviews in Molecular Medicine, 2017, 19, e8.	1.6	79
72	Integrating genomics to dig deeper into Wilms tumour biology. Nature Reviews Urology, 2017, 14, 703-704.	1.9	4

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73	Rationale for the treatment of Wilms tumour in the UMBRELLA SIOP–RTSG 2016 protocol. Nature Reviews Urology, 2017, 14, 743-752.	1.9	249
74	Wilms' Tumor., 2017,, 4850-4853.		O
75	The clinical phenotype of <scp><i>YWHAEâ€NUTM2B/E</i></scp> positive pediatric clear cell sarcoma of the kidney. Genes Chromosomes and Cancer, 2016, 55, 143-147.	1.5	14
76	Paediatric cancer stage in population-based cancer registries: the Toronto consensus principles and guidelines. Lancet Oncology, The, 2016, 17, e163-e172.	5.1	56
77	Risk of Adverse Health and Social Outcomes Up to 50 Years After Wilms Tumor: The British Childhood Cancer Survivor Study. Journal of Clinical Oncology, 2016, 34, 1772-1779.	0.8	59
78	European Survey on Standards of Care in paediatric oncology centres. European Journal of Cancer, 2016, 61, 11-19.	1.3	25
79	Ethical issues of clinical trials in paediatric oncology from 2003 to 2013: a systematic review. Lancet Oncology, The, 2016, 17, e187-e197.	5.1	14
80	Biology and treatment of Wilms' tumours in childhood. Revue D'Oncologie Hématologie Pédiatrique, 2016, 4, 170-181.	0.1	3
81	Intra-Tumor Genetic Heterogeneity in Wilms Tumor: Clonal Evolution and Clinical Implications. EBioMedicine, 2016, 9, 120-129.	2.7	61
82	Wilms tumor: "State-of-the-art―update, 2016. Seminars in Pediatric Surgery, 2016, 25, 250-256.	0.5	85
83	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.	0.8	105
84	Biology and treatment of renal tumours in childhood. European Journal of Cancer, 2016, 68, 179-195.	1.3	107
85	PWE-131â€The First UK Multidisciplinary Diagnostic Centre: A Novel Cancer Diagnostic Service. Gut, 2016, 65, A202.2-A203.	6.1	1
86	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.	1.0	35
87	Reorganising specialist cancer surgery for the twenty-first century: a mixed methods evaluation (RESPECT-21). Implementation Science, 2016, 11, 155.	2.5	18
88	The SIOPE strategic plan: A European cancer plan for children and adolescents. Journal of Cancer Policy, 2016, 8, 17-32.	0.6	57
89	WT1 Mutation in Childhood Cancer. Methods in Molecular Biology, 2016, 1467, 1-14.	0.4	17
90	256 Integrated analysis of DNA methylation, copy number and expression data in Wilms Tumour identifies subtype-specific molecular signatures. European Journal of Cancer, 2015, 51, S45.	1.3	0

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91	1407 The European Strategic Plan for children and adolescents with cancer. European Journal of Cancer, 2015, 51, S200.	1.3	О
92	Response to the letter to the editor: 1q gain is a frequent finding in preoperatively treated <scp>W</scp> ilms tumors, but of limited prognostic value for risk satisfaction in the <scp>SIOP</scp> 2009/ <scp>Gesellschaft für PÃdiatrische Onkologie und HÃmatologie (GPOH)</scp> trial. Genes Chromosomes and Cancer, 2015, 54, 397-399.	1.5	2
93	A multiâ€Gaussian model for apparent diffusion coefficient histogram analysis of Wilms' tumour subtype and response to chemotherapy. NMR in Biomedicine, 2015, 28, 948-957.	1.6	34
94	The yin and yang of kidney development and Wilms' tumors. Genes and Development, 2015, 29, 467-482.	2.7	129
95	Risk factors for local recurrence in Wilms tumour and the potential influence of biopsy – The United Kingdom experience. European Journal of Cancer, 2015, 51, 225-232.	1.3	47
96	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.	7.7	248
97	Biomarkers to detect Wilms tumors in pediatric patients: where are we now?. Future Oncology, 2015, 11, 2221-2234.	1.1	18
98	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.	1.3	67
99	Response to letter commenting on "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, 995-996.	1.3	0
100	Comparative methylome analysis identifies new tumour subtypes and biomarkers for transformation of nephrogenic rests into Wilms tumour. Genome Medicine, 2015, 7, 11.	3.6	39
101	Long-term effects of Wilms tumour therapy on renal function. Nature Reviews Urology, 2015, 12, 423-424.	1.9	2
102	Advances in Wilms Tumor Treatment and Biology: Progress Through International Collaboration. Journal of Clinical Oncology, 2015, 33, 2999-3007.	0.8	281
103	Omission of doxorubicin from the treatment of stage Il–III, intermediate-risk Wilms' tumour (SIOP WT) Tj ETQq	1 1 0.7843 6.3	314 rgBT / <mark>O</mark> v 165
104	Multiple mechanisms of MYCN dysregulation in Wilms tumour. Oncotarget, 2015, 6, 7232-7243.	0.8	85
105	Wilms' Tumor. , 2015, , 1-4.		0
106	Abstract A1-59: Multiple mechanisms of MYCN dysregulation in Wilms tumor., 2015,,.		1
107	Abstract A1-67: Prognostic significance of copy number aberrations in Wilms tumor., 2015,,.		0
108	TP53 Mutational Status Is a Potential Marker for Risk Stratification in Wilms Tumour with Diffuse Anaplasia. PLoS ONE, 2014, 9, e109924.	1.1	82

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109	Methylome analysis identifies a Wilms tumor epigenetic biomarker detectable in blood. Genome Biology, 2014, 15, 434.	3.8	33
110	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.	2.9	49
111	Why should survivors of childhood renal tumor and others with only one kidney be denied the chance to play contact sports?. Expert Review of Anticancer Therapy, 2014, 14, 363-366.	1.1	5
112	Drug discovery in paediatric oncology: roadblocks to progress. Nature Reviews Clinical Oncology, 2014, 11, 732-739.	12.5	28
113	Declining childhood and adolescent cancer mortality: Great progress but still much to be done. Cancer, 2014, 120, 2388-2391.	2.0	27
114	Toxicity and Outcome of Children and Adolescents Participating in Phase I/II Trials of Novel Anticancer Drugs. Journal of Pediatric Hematology/Oncology, 2014, 36, 218-223.	0.3	25
115	Towards reducing inequalities: European Standards of Care for Children with Cancer. European Journal of Cancer, 2014, 50, 481-485.	1.3	36
116	Germline mutations in the PAF1 complex gene CTR9 predispose to Wilms tumour. Nature Communications, 2014, 5, 4398.	5.8	85
117	Nephron sparing surgery (NSS) for unilateral wilms tumor (UWT): The SIOP 2001 experience. Pediatric Blood and Cancer, 2014, 61, 2175-2179.	0.8	85
118	Lin28 sustains early renal progenitors and induces Wilms tumor. Genes and Development, 2014, 28, 971-982.	2.7	149
119	Outcome of localized blastemal-type nephroblastoma patients treated according to intensified treatment in the SIOP 2001 protocol: A report of the SIOP-RTSG Journal of Clinical Oncology, 2014, 32, 10002-10002.	0.8	1
120	Learning what high quality compassionate care means for cancer patients and translating that into practice. Patient Experience Journal, 2014, 1, 124-131.	0.3	5
121	Wilms' tumor: biology, diagnosis and treatment. Translational Pediatrics, 2014, 3, 12-24.	0.5	100
122	Treatment and outcome of patients with relapsed clear cell sarcoma of the kidney (CCSK): A combined SIOP and AIEOP study Journal of Clinical Oncology, 2014, 32, 10041-10041.	0.8	1
123	Incidence and outcomes of patients with late recurrence of Wilms' tumor. Pediatric Blood and Cancer, 2013, 60, 1612-1615.	0.8	43
124	New policies to address the global burden of childhood cancers. Lancet Oncology, The, 2013, 14, e125-e135.	5.1	96
125	Surgical complications after immediate nephrectomy versus preoperative chemotherapy in non-metastatic Wilms' tumour: Findings from the 1991–2001 United Kingdom Children's Cancer Study Group UKW3 Trial. Journal of Pediatric Surgery, 2013, 48, 2181-2186.	0.8	31
126	Is Wilms Tumor a Candidate Neoplasia for Treatment with WNT∫β-Catenin Pathway Modulators?—A Report from the Renal Tumors Biology-Driven Drug Development Workshop. Molecular Cancer Therapeutics, 2013, 12, 2619-2627.	1.9	28

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127	Sustaining innovation and improvement in the treatment of childhood cancer: lessons from high-income countries. Lancet Oncology, The, 2013, 14, e95-e103.	5.1	175
128	Children with cancer: driving the global agenda. Lancet Oncology, The, 2013, 14, 189-191.	5.1	35
129	An international strategy to determine the role of high dose therapy in recurrent Wilms' tumour. European Journal of Cancer, 2013, 49, 194-210.	1.3	61
130	SIOP PODC: Clinical guidelines for the management of children with Wilms tumour in a low income setting. Pediatric Blood and Cancer, 2013, 60, 5-11.	0.8	81
131	Treatment of high risk Sertoli–Leydig cell tumors of the ovary using a gonadotropin releasing hormone (GnRH) analog. Pediatric Blood and Cancer, 2013, 60, E16-8.	0.8	9
132	Treatment of Wilms tumor in low-income countries: challenges and potential solutions. Future Oncology, 2013, 9, 1057-1059.	1.1	7
133	Weaver syndrome and <i>EZH2</i> mutations: Clarifying the clinical phenotype. American Journal of Medical Genetics, Part A, 2013, 161, 2972-2980.	0.7	119
134	Gain of 1q is a marker of poor prognosis in Wilms' tumors. Genes Chromosomes and Cancer, 2013, 52, 1065-1074.	1.5	54
135	Bilateral Wilms Tumor with <i>TP53</i> -Related Anaplasia. Pediatric and Developmental Pathology, 2013, 16, 217-223.	0.5	13
136	Challenges in incentivizing the pharmaceutical industry to supporting pediatric oncology clinical trials. Clinical Investigation, 2013, 3, 101-103.	0.0	2
137	miRNA Profiles as a Predictor of Chemoresponsiveness in Wilms' Tumor Blastema. PLoS ONE, 2013, 8, e53417.	1.1	71
138	Abstract 3829: TP53 mutation status defines two distinct classes of diffuse anaplastic Wilms tumor, 2013, , .		0
139	Abstract B36: DNA methylation profiling describes Wilms tumor evolution from its precursor lesion. , 2013, , .		0
140	Pragmatic approach to quality metrics development in cancer Journal of Clinical Oncology, 2013, 31, 60-60.	0.8	0
141	Treatment and outcome of Wilms' tumour patients: an analysis of all cases registered in the UKW3 trial. Annals of Oncology, 2012, 23, 2457-2463.	0.6	79
142	Population survival from childhood cancer in Britain during 1978–2005 by eras of entry to clinical trials. Annals of Oncology, 2012, 23, 2464-2469.	0.6	57
143	Consent to tissue banking for research: qualitative study and recommendations. Archives of Disease in Childhood, 2012, 97, 632-636.	1.0	8
144	A genome-wide association study identifies susceptibility loci for Wilms tumor. Nature Genetics, 2012, 44, 681-684.	9.4	72

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145	Reply to S. Stegmaier et al. Journal of Clinical Oncology, 2012, 30, 4040-4041.	0.8	4
146	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.	1.3	53
147	Stratification of Wilms tumor by genetic and epigenetic analysis. Oncotarget, 2012, 3, 327-335.	0.8	101
148	<i>PAX3/FOXO1</i> Fusion Gene Status Is the Key Prognostic Molecular Marker in Rhabdomyosarcoma and Significantly Improves Current Risk Stratification. Journal of Clinical Oncology, 2012, 30, 1670-1677.	0.8	297
149	ecancermedicalscience. Ecancermedicalscience, 2011, 5, 210.	0.6	16
150	4103 ORAL Doxorubicin Can Be Safely Omitted From the Treatment of Stage II/III, Intermediate Risk Histology Wilms Tumour – Results of the SIOP WT 2001 Randomised Trial, on Behalf of the SIOP Renal Tumours Study Group. European Journal of Cancer, 2011, 47, S284-S285.	1.3	1
151	4107 ORAL The State of Research Into Children With Cancer Across Europe: Results of Key Opinion Leaders Survey and Innovative Policy Strategies for the Next Decade. European Journal of Cancer, 2011, 47, S286.	1.3	0
152	171 INVITED The European Network for Cancer Research in Children and Adolescents(ENCCA) FP7 Project. European Journal of Cancer, 2011, 47, S41.	1.3	0
153	172 INVITED â€~Spread of Excellence' (Dissemination) Activities of the European Network of Excellence for Cancer Research in Children and Adolescents (ENCCA). European Journal of Cancer, 2011, 47, S41-S42.	1.3	0
154	Suggested change in definitional criteria for stage I Wilms tumour treated by surgery onlyâ€"Response. Pediatric Blood and Cancer, 2011, 56, 169-170.	0.8	0
155	Mesoblastic nephroma: A report of the United Kingdom children's cancer and leukaemia group (CCLG). Pediatric Blood and Cancer, 2011, 56, 744-748.	0.8	58
156	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.	0.8	125
157	Molecular profiling reveals frequent gain of <i>MYCN</i> and anaplasiaâ€specific loss of 4q and 14q in wilms tumor. Genes Chromosomes and Cancer, 2011, 50, 982-995.	1.5	54
158	Management of adults with Wilms' tumor: recommendations based on international consensus. Expert Review of Anticancer Therapy, 2011, 11, 1107-1115.	1.1	37
159	DICER1 syndrome: clarifying the diagnosis, clinical features and management implications of a pleiotropic tumour predisposition syndrome. Journal of Medical Genetics, 2011, 48, 273-278.	1.5	312
160	Long-term Risks of Subsequent Primary Neoplasms Among Survivors of Childhood Cancer. JAMA - Journal of the American Medical Association, 2011, 305, 2311.	3.8	289
161	Renal Tumors in Children Aged 10–16 Years: A Report from the United Kingdom Children's Cancer and Leukaemia Group. Pediatric and Developmental Pathology, 2011, 14, 189-193.	0.5	32
162	Wilms Tumor., 2011,, 3947-3950.		0

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163	Abstract 5343: Aberrant activation of hedgehog signaling confers a poor prognosis in embryonal and fusion gene negative alveolar rhabdomyosarcoma. , $2011, \dots$		O
164	Incidence and outcomes of patients with late recurrence of Wilms tumor (WT): The international experience Journal of Clinical Oncology, 2011, 29, 9544-9544.	0.8	O
165	Research and Drug Development in Paediatric Oncology. , 2010, , 155-159.		O
166	Verification of the susceptibility loci on 7p12.2, 10q21.2, and 14q11.2 in precursor B-cell acute lymphoblastic leukemia of childhood. Blood, 2010, 115, 1765-1767.	0.6	142
167	Hereditary leiomyomatosis and renal cell carcinoma: very early diagnosis of renal cancer in a paediatric patient. Familial Cancer, 2010, 9, 239-243.	0.9	56
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