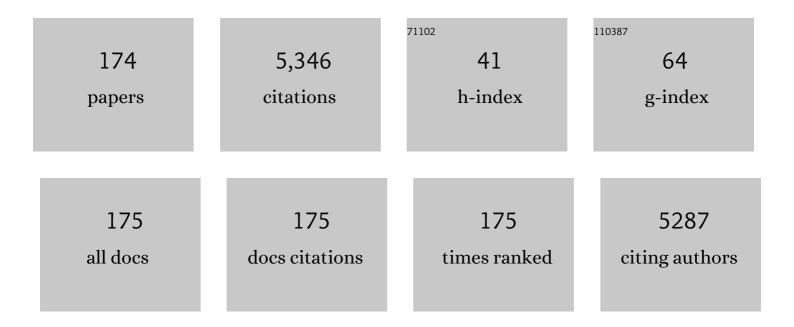
Filippo Spreafico

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Advances in Wilms Tumor Treatment and Biology: Progress Through International Collaboration. Journal of Clinical Oncology, 2015, 33, 2999-3007.	1.6	281
2	Rationale for the treatment of Wilms tumour in the UMBRELLA SIOP–RTSG 2016 protocol. Nature Reviews Urology, 2017, 14, 743-752.	3.8	249
3	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
4	High Response Rate to Cisplatin/Etoposide Regimen in Childhood Low-Grade Glioma. Journal of Clinical Oncology, 2002, 20, 4209-4216.	1.6	171
5	Hyperfractionated Accelerated Radiotherapy in the Milan Strategy for Metastatic Medulloblastoma. Journal of Clinical Oncology, 2009, 27, 566-571.	1.6	140
6	Adult-Type Soft Tissue Sarcomas in Pediatric-Age Patients: Experience at the Istituto Nazionale Tumori in Milan. Journal of Clinical Oncology, 2005, 23, 4021-4030.	1.6	130
7	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
8	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
9	Hyperfractionated radiotherapy and chemotherapy for childhood ependymoma: final results of the first prospective AIEOP (Associazione Italiana di Ematologia-Oncologia Pediatrica) study. International Journal of Radiation Oncology Biology Physics, 2004, 58, 1336-1345.	0.8	93
10	Distinct Methylation Changes at the IGF2-H19 Locus in Congenital Growth Disorders and Cancer. PLoS ONE, 2008, 3, e1849.	2.5	93
11	Brain tumors in children and adolescents: Cognitive and psychological disorders at different ages. Psycho-Oncology, 2005, 14, 386-395.	2.3	87
12	Children with cancer in the time of COVIDâ€19: An 8â€week report from the six pediatric oncoâ€hematology centers in Lombardia, Italy. Pediatric Blood and Cancer, 2020, 67, e28410.	1.5	82
13	How young patients with cancer perceive the COVIDâ€19 (coronavirus) epidemic in Milan, Italy: Is there room for other fears?. Pediatric Blood and Cancer, 2020, 67, e28318.	1.5	81
14	Adult Wilms' tumor: A monoinstitutional experience and a review of the literature. Cancer, 2004, 101, 289-293.	4.1	77
15	Treatment of relapsed Wilms tumors: lessons learned. Expert Review of Anticancer Therapy, 2009, 9, 1807-1815.	2.4	77
16	Wilms tumour. Nature Reviews Disease Primers, 2021, 7, 75.	30.5	75
17	A lower-dose, lower-toxicity cisplatin–etoposide regimen for childhood progressive low-grade glioma. Journal of Neuro-Oncology, 2010, 100, 65-71.	2.9	74
18	Vinorelbine in previously treated advanced childhood sarcomas. Cancer, 2002, 94, 3263-3268.	4.1	73

#	Article	IF	CITATIONS
19	Wilms' tumor: past, present and (possibly) future. Expert Review of Anticancer Therapy, 2006, 6, 249-258.	2.4	68
20	Functional inactivation of the WTX gene is not a frequent event in Wilms' tumors. Oncogene, 2008, 27, 4625-4632.	5.9	63
21	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
22	Results of nimotuzumab and vinorelbine, radiation and re-irradiation for diffuse pontine glioma in childhood. Journal of Neuro-Oncology, 2014, 118, 305-312.	2.9	61
23	Whole transcriptome sequencing identifies BCOR internal tandem duplication as a common feature of clear cell sarcoma of the kidney. Oncotarget, 2015, 6, 40934-40939.	1.8	61
24	Papillary thyroid carcinoma of childhood and adolescence: A 30-year experience at the istituto nazionale tumori in Milan. Pediatric Blood and Cancer, 2006, 46, 300-306.	1.5	60
25	Diffuse pontine gliomas in children: changing strategies, changing results? A mono-institutional 20-year experience. Journal of Neuro-Oncology, 2008, 87, 355-361.	2.9	59
26	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
27	Comparison of the Prognostic Value of Assessing Tumor Diameter Versus Tumor Volume at Diagnosis or in Response to Initial Chemotherapy in Rhabdomyosarcoma. Journal of Clinical Oncology, 2010, 28, 1322-1328.	1.6	58
28	The Youth Project at the Istituto Nazionale Tumori in Milan. Tumori, 2012, 98, 399-407.	1.1	58
29	Sequential chemotherapy, high-dose thiotepa, circulating progenitor cell rescue, and radiotherapy for childhood high-grade glioma. Neuro-Oncology, 2005, 7, 41-48.	1.2	56
30	Survival of adults treated for medulloblastoma using paediatric protocols. European Journal of Cancer, 2005, 41, 1304-1310.	2.8	56
31	Soft Tissue Sarcomas of Childhood and Adolescence: The Prognostic Role of Tumor Size in Relation to Patient Body Size. Journal of Clinical Oncology, 2009, 27, 371-376.	1.6	55
32	Psychological intervention in young brain tumor survivors: The efficacy of the cognitive behavioural approach. Disability and Rehabilitation, 2009, 31, 1066-1073.	1.8	50
33	Prognostic determinants in epithelioid sarcoma. European Journal of Cancer, 2011, 47, 287-295.	2.8	50
34	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
35	Intensive, Very Short-Term Chemotherapy for Advanced Burkitt's Lymphoma in Children. Journal of Clinical Oncology, 2002, 20, 2783-2788.	1.6	47
36	Supratentorial primitive neuroectodermal tumors (S-PNET) in children: A prospective experience with adjuvant intensive chemotherapy and hyperfractionated accelerated radiotherapy. International Journal of Radiation Oncology Biology Physics, 2006, 64, 1031-1037.	0.8	47

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37	Temozolomide is an active agent in children with recurrent medulloblastoma/primitive neuroectodermal tumor: an Italian multi-institutional phase II trial. Neuro-Oncology, 2014, 16, 748-753.	1.2	47
38	Clouds of Oxygen: Adolescents With Cancer Tell Their Story in Music. Journal of Clinical Oncology, 2015, 33, 218-221.	1.6	47
39	Mature and immature teratoma: A report from the second Italian pediatric study. Pediatric Blood and Cancer, 2015, 62, 1202-1208.	1.5	47
40	The Sooner the Better? How Symptom Interval Correlates With Outcome in Children and Adolescents With Solid Tumors: Regression Tree Analysis of the Findings of a Prospective Study. Pediatric Blood and Cancer, 2016, 63, 479-485.	1.5	45
41	No Salvage Using High-Dose Chemotherapy Plus/Minus Reirradiation for Relapsing Previously Irradiated Medulloblastoma. International Journal of Radiation Oncology Biology Physics, 2009, 73, 1358-1363.	0.8	44
42	Adult-type soft tissue sarcomas in paediatric age: A nomogram-based prognostic comparison with adult sarcoma. European Journal of Cancer, 2007, 43, 2691-2697.	2.8	43
43	Incidence and outcomes of patients with late recurrence of Wilms' tumor. Pediatric Blood and Cancer, 2013, 60, 1612-1615.	1.5	43
44	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
45	A collateral effect of the COVIDâ€19 pandemic: Delayed diagnosis in pediatric solid tumors. Pediatric Blood and Cancer, 2020, 67, e28640.	1.5	43
46	Conservative surgical approach for thyroid and lymph-node involvement in papillary thyroid carcinoma of childhood and adolescence. Pediatric Blood and Cancer, 2006, 46, 307-313.	1.5	40
47	Local lymph node involvement in pediatric renal cell carcinoma: A report from the Italian TREP project. Pediatric Blood and Cancer, 2008, 51, 475-478.	1.5	39
48	Germline mutations of thePOU6F2 gene in Wilms tumors with loss of heterozygosity on chromosome 7p14. Human Mutation, 2004, 24, 400-407.	2.5	38
49	Treatment of highâ€risk relapsed Wilms tumor with doseâ€intensive chemotherapy, marrowâ€ablative chemotherapy, and autologous hematopoietic stem cell support: Experience by the Italian association of pediatric hematology and oncology. Pediatric Blood and Cancer, 2008, 51, 23-28.	1.5	38
50	Brain Magnetic Resonance Imaging After High-Dose Chemotherapy and Radiotherapy for Childhood Brain Tumors. International Journal of Radiation Oncology Biology Physics, 2008, 70, 1011-1019.	0.8	38
51	Histological variants of medulloblastoma are the most powerful clinical prognostic indicators. Pediatric Blood and Cancer, 2013, 60, 210-216.	1.5	38
52	Management of adults with Wilms' tumor: recommendations based on international consensus. Expert Review of Anticancer Therapy, 2011, 11, 1107-1115.	2.4	37
53	Quantitative DNA methylation analysis improves epigenotype-phenotype correlations in Beckwith-Wiedemann syndrome. Epigenetics, 2013, 8, 1053-1060.	2.7	33
54	Primary metastatic osteosarcoma: results of a prospective study in children given chemotherapy and interleukin-2. Medical Oncology, 2017, 34, 191.	2.5	33

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55	Salvage treatment for childhood ependymoma after surgery only: Pitfalls of omitting "at once― adjuvant treatment. International Journal of Radiation Oncology Biology Physics, 2006, 65, 1440-1445.	0.8	31
56	Renal cell carcinoma in children and adolescents. Expert Review of Anticancer Therapy, 2010, 10, 1967-1978.	2.4	31
57	Paediatric renal tumours: perspectives from the SIOP–RTSG. Nature Reviews Urology, 2017, 14, 3-4.	3.8	31
58	Loss of Heterozygosity Analysis at Different Chromosome Regions in Wilms Tumor Confirms 1p Allelic Loss as a Marker of Worse Prognosis: A Study from the Italian Association of Pediatric Hematology and Oncology. Journal of Urology, 2013, 189, 260-267.	0.4	30
59	Immunomodulation in a Treatment Program Including Pre- and Post-Operative Interleukin-2 and Chemotherapy for Childhood Osteosarcoma. Tumori, 2003, 89, 263-268.	1.1	29
60	Characteristics and Outcome of Children with Renal Cell Carcinoma: A Narrative Review. Cancers, 2020, 12, 1776.	3.7	29
61	Genomic profiling by wholeâ€genome single nucleotide polymorphism arrays in Wilms tumor and association with relapse. Genes Chromosomes and Cancer, 2012, 51, 644-653.	2.8	28
62	Measuring the efficacy of a project for adolescents and young adults with cancer: A study from the Milan Youth Project. Pediatric Blood and Cancer, 2016, 63, 2197-2204.	1.5	28
63	Searching for Happiness. Journal of Clinical Oncology, 2017, 35, 2209-2212.	1.6	28
64	The Youth Project at the Istituto Nazionale Tumori in Milan. Tumori, 2012, 98, 399-407.	1.1	28
65	Prognostic Factors for Wilms Tumor Recurrence: A Review of the Literature. Cancers, 2021, 13, 3142.	3.7	27
66	Endodermal sinus tumor of the vagina. Pediatric Blood and Cancer, 2007, 48, 577-578.	1.5	26
67	Chromosomal anomalies at 1q, 3, 16q, and mutations of <i>SIX1</i> and <i>DROSHA</i> genes underlie Wilms tumor recurrences. Oncotarget, 2016, 7, 8908-8915.	1.8	26
68	CHILDHOOD LIPOSARCOMA: A Single-Institutional Twenty-Year Experience. Pediatric Hematology and Oncology, 1999, 16, 415-421.	0.8	25
69	End of life in children with cancer: Experience at the Pediatric Oncology Department of the Istituto Nazionale Tumori in Milan. Pediatric Blood and Cancer, 2010, 54, 88-91.	1.5	25
70	Teratoma with a malignant somatic component in pediatric patients: The Associazione Italiana Ematologia Oncologia Pediatrica (AIEOP) experience. Pediatric Blood and Cancer, 2010, 54, 532-537.	1.5	25
71	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.	2.8	25
72	Genetic and epigenetic analyses guided by high resolution whole-genome SNP array reveals a possible role of <i>CHEK2</i> in Wilms tumour susceptibility. Oncotarget, 2018, 9, 34079-34089.	1.8	25

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73	Proteomic analysis of cerebrospinal fluid from children with central nervous system tumors identifies candidate proteins relating to tumor metastatic spread. Oncotarget, 2017, 8, 46177-46190.	1.8	24
74	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.	2.8	24
75	Neuroblastoma in Patients over 12 Years Old: A 20-Year Experience at the Istituto Nazionale Tumori of Milan. Tumori, 2010, 96, 684-689.	1.1	23
76	Evolving of therapeutic strategies for CNS-PNET. Pediatric Blood and Cancer, 2013, 60, 2031-2035.	1.5	23
77	Synchronous bilateral Wilms tumor. Cancer, 2013, 119, 1586-1592.	4.1	22
78	Wilms tumor, medulloblastoma, and rhabdomyosarcoma in adult patients: lessons learned from the pediatric experience. Cancer and Metastasis Reviews, 2019, 38, 683-694.	5.9	22
79	Heterogeneity of Disease Classified as Stage III in Wilms Tumor: A Report From the Associazione Italiana Ematologia Oncologia Pediatrica (AIEOP). International Journal of Radiation Oncology Biology Physics, 2012, 82, 348-354.	0.8	21
80	Long-term results of combined preradiation chemotherapy and age-tailored radiotherapy doses for childhood medulloblastoma. Journal of Neuro-Oncology, 2012, 108, 163-171.	2.9	20
81	Rhabdomyosarcoma of the Head and Neck Region: Experience at the Pediatric Unit of the Istituto Nazionale Tumori, Milan. The Journal of Otolaryngology, 2006, 35, 53.	0.6	19
82	SARSâ€CoVâ€⊋ disease and children under treatment for cancer. Pediatric Blood and Cancer, 2020, 67, e28346.	1.5	19
83	Thyroid-Stimulating Hormone Suppression for Protection Against Hypothyroidism Due to Craniospinal Irradiation for Childhood Medulloblastoma/Primitive Neuroectodermal Tumor. International Journal of Radiation Oncology Biology Physics, 2007, 69, 404-410.	0.8	18
84	Radiation-induced thyroid changes: A retrospective and a prospective view. European Journal of Cancer, 2009, 45, 2546-2551.	2.8	18
85	Unmet needs for relapsed or refractory Wilms tumour: Mapping the molecular features, exploring organoids and designing early phase trials – A collaborative SIOP-RTSG, COGÂand ITCC session at the first SIOPE meeting. European Journal of Cancer, 2021, 144, 113-122.	2.8	18
86	Axial skeletal osteosarcoma: a 25-year monoinstitutional experience in children and adolescents. Medical Oncology, 2014, 31, 875.	2.5	17
87	Amelanotic melanoma in a child with oculocutaneous albinism. Medical and Pediatric Oncology, 2003, 41, 179-180.	1.0	16
88	Evolving treatment strategies for parameningeal rhabdomyosarcoma: The experience of the istituto nazionale tumori of Milan. Head and Neck, 2005, 27, 49-57.	2.0	16
89	SCT for Wilms' tumour. Bone Marrow Transplantation, 2008, 41, S128-S130.	2.4	16
90	Results of the Third AIEOP Cooperative Protocol on Wilms Tumor (TW2003) and Related Considerations. Journal of Urology, 2017, 198, 1138-1145.	0.4	16

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#	Article	IF	CITATIONS
91	Childhood Malignant Ovarian Germ Cell Tumors: A Monoinstitutional Experience. Gynecologic Oncology, 2001, 81, 436-440.	1.4	15
92	Telomere maintenance in wilms tumors: First evidence for the presence of alternative lengthening of telomeres mechanism. Genes Chromosomes and Cancer, 2011, 50, 823-829.	2.8	15
93	Clinical Stage I Nonseminomatous Germ Cell Tumors of the Testis in Childhood and Adolescence: An Analysis of 31 Cases. Journal of Pediatric Hematology/Oncology, 2002, 24, 454-458.	0.6	14
94	A Case of Relapsing Glioblastoma Multiforme Responding to Vinorelbine. Journal of Neuro-Oncology, 2006, 80, 195-201.	2.9	14
95	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.	2.8	14
96	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.	1.5	14
97	Prognostic significance of p80 and visceral involvement in childhood CD30 anaplastic large cell lymphoma (ALCL). Medical and Pediatric Oncology, 2001, 37, 97-102.	1.0	13
98	Malignant testicular germ cell tumors in children and adolescents: The AIEOP (Associazione Italiana) Tj ETQq0 0 Investigations, 2018, 36, 502.e7-502.e13.	0 rgBT /Ov 1.6	verlock 10 Tf 13
99	Metastatic Renal Cell Carcinoma in Children and Adolescents. Journal of Pediatric Hematology/Oncology, 2012, 34, e277-e281.	0.6	12
100	Occurrence of Breast Cancer After Chest Wall Irradiation for Pediatric Cancer, as Detected by a Multimodal Screening Program. International Journal of Radiation Oncology Biology Physics, 2013, 85, 35-39.	0.8	12
101	Treatment of relapsed Wilms tumour (WT) patients: Experience with topotecan. A report from the SIOP Renal Tumour Study Group (RTSG). Pediatric Blood and Cancer, 2015, 62, 598-602.	1.5	12
102	A case of relapsing spinal atypical teratoid/rhabdoid tumor (AT/RT) responding to vinorelbine, cyclophosphamide, and celecoxib. Child's Nervous System, 2015, 31, 1621-1623.	1.1	12
103	Sport activities and exercise as part of routine cancer care in children and adolescents. Pediatric Blood and Cancer, 2019, 66, e27826.	1.5	12
104	Undifferentiated nasopharyngeal carcinoma in children and adolescents: Comparison between staging systems. Annals of Oncology, 2001, 12, 1157-1162.	1.2	11
105	Stage 4 neuroblastoma: sequential hemi-body irradiation or high-dose chemotherapy plus autologous haemopoietic stem cell transplantation to consolidate primary treatment. British Journal of Cancer, 2005, 92, 1984-1988.	6.4	11
106	Should we encourage exercise and sports in children and adolescents with cancer?. Pediatric Blood and Cancer, 2014, 61, 2125-2125.	1.5	11
107	Thyroid carcinoma after treatment for malignancies in childhood and adolescence: from diagnosis through follow-up. Medical Oncology, 2014, 31, 121.	2.5	11
108	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.	1.5	11

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109	The Murine Pou6f2 Gene is Temporally and Spatially Regulated During Kidney Embryogenesis and its Human Homolog is Overexpressed in a Subset of Wilms Tumors. Journal of Pediatric Hematology/Oncology, 2006, 28, 791-797.	0.6	10
110	Constitutional ring chromosome 11 mosaicism in a Wilms tumor patient: Cytogenetic, molecular and clinicoâ€pathological studies. American Journal of Medical Genetics, Part A, 2010, 152A, 1756-1763.	1.2	10
111	A novel <i>WT1</i> mutation in familial wilms tumor. Pediatric Blood and Cancer, 2013, 60, 1388-1389.	1.5	10
112	FIVE QUESTIONS FOR ASSESSING PSYCHOLOGICAL PROBLEMS IN PEDIATRIC PATIENTS CURED OF NEOPLASTIC DISEASE. Pediatric Hematology and Oncology, 2004, 21, 481-487.	0.8	9
113	Analysis of the mutational status of SIX1/2 and microRNA processing genes in paired primary and relapsed Wilms tumors and association with relapse. Cancer Gene Therapy, 2021, 28, 1016-1024.	4.6	9
114	Positive Impact of Organized Physical Exercise on Quality of Life and Fatigue in Children and Adolescents With Cancer. Frontiers in Pediatrics, 2021, 9, 627876.	1.9	9
115	Clinical Experience with Psychological Aspects in Pediatric Patients Amputated for Malignancies. Tumori, 2004, 90, 399-404.	1.1	8
116	WT1 Gene Analysis in Sporadic Early-Onset and Bilateral Wilms Tumor Patients Without Associated Abnormalities. Journal of Pediatric Hematology/Oncology, 2005, 27, 197-201.	0.6	8
117	Value and difficulties of a common European strategy for recurrent Wilms' tumor. Expert Review of Anticancer Therapy, 2009, 9, 693-696.	2.4	8
118	Severe polyuria and polydipsia in hyponatremicâ€hypertensive syndrome associated with Wilms tumor. Pediatric Blood and Cancer, 2010, 55, 566-569.	1.5	8
119	Primary Renal Soft Tissue Sarcoma in Children. Urology, 2012, 80, 698-702.	1.0	8
120	Factors possibly affecting prognosis in children with Wilms' tumor diagnosed before 24 months of age: A report from the Associazione Italiana Ematologia Oncologia Pediatrica (AIEOP) Wilms Tumor Working Group. Pediatric Blood and Cancer, 2017, 64, e26644.	1.5	8
121	Reduced-dose craniospinal irradiation is feasible for standard-risk adult medulloblastoma patients. Journal of Neuro-Oncology, 2020, 148, 619-628.	2.9	8
122	Outcome of SIOP patients with low- or intermediate-risk Wilms tumour relapsing after initial vincristine and actinomycin-D therapy only â^' the SIOP 93–01 and 2001 protocols. European Journal of Cancer, 2022, 163, 88-97.	2.8	8
123	Unusual primary secreting germ cell tumor of the spine. Journal of Neurosurgery: Spine, 2006, 5, 65-67.	1.7	7
124	Molecular evidence of the independent origin of multiple Wilms tumors in a case of WAGR syndrome. Pediatric Blood and Cancer, 2008, 51, 344-348.	1.5	7
125	Is There a Role for FDG-PET for the Assessment of Treatment Efficacy in Wilms' Tumor? A Case Report and Literature Review. Pediatric Hematology and Oncology, 2013, 30, 633-639.	0.8	7
126	Longâ€ŧerm renal outcome in adolescent and young adult patients nephrectomized for unilateral Wilms tumor. Pediatric Blood and Cancer, 2014, 61, 1136-1137.	1.5	7

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127	Experiencing Social Isolation (Even in the Era of COVID-19 Pandemic Lockdown): Teachings Through Arts from Adolescents with Cancer. Journal of Adolescent and Young Adult Oncology, 2021, 10, 346-350.	1.3	7
128	Revised SIOP working classification of renal tumors of childhood. Medical and Pediatric Oncology, 2003, 41, 102-102.	1.0	6
129	ETOPOSIDE, CISPLATIN, EPIRUBICIN CHEMOTHERAPY IN THE TREATMENT OF PEDIATRIC LIVER TUMORS. Pediatric Hematology and Oncology, 2005, 22, 189-198.	0.8	6
130	Assistance to Parents who have Lost their Child with Cancer. Tumori, 2006, 92, 306-310.	1.1	6
131	Psychological Assessment of Women on an Early Breast Screening Program after Radiotherapy to the Chest Wall for Childhood Cancer. Tumori, 2008, 94, 568-573.	1.1	6
132	Mixed Epithelial and Stromal Tumor of Kidney: An Exceptional Renal Neoplasm in an 8-Year-Old Prepubertal Girl with Isolated Clitoral Hypertrophy. Pediatric Hematology and Oncology, 2012, 29, 89-91.	0.8	6
133	Oral Etoposide in Relapsed or Refractory Ewing Sarcoma: A Monoinstitutional Experience in Children and Adolescents. Tumori, 2016, 102, 84-88.	1.1	6
134	An Analysis of Treatment Failure in Wilms Tumor (WT): A Report from the Central American Association of Pediatric Hematology/Oncology (AHOPCA). Journal of Global Oncology, 2016, 2, 2s-2s.	0.5	6
135	Malignant sacrococcygeal germ cell tumors in childhood: The Associazione Italiana Ematologia Oncologia Pediatrica (AIEOP) experience. Pediatric Blood and Cancer, 2021, 68, e28812.	1.5	6
136	Adult-type non-rhabdomyosarcoma soft tissue sarcomas in pediatric age: Salvage rates and prognostic factors after relapse. European Journal of Cancer, 2022, 169, 179-187.	2.8	6
137	Bilateral preaxial polydactyly in a WAGR syndrome patient. American Journal of Medical Genetics, Part A, 2005, 134A, 426-429.	1.2	5
138	Nonâ€chromosome 11â€p syndromes in Wilms tumor patients: Clinical and cytogenetic report of two Down syndrome cases and one Turner syndrome case. American Journal of Medical Genetics, Part A, 2007, 143A, 85-88.	1.2	5
139	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.	2.4	5
140	Rehabilitation for children and young people surviving a brain tumor, and their transition to adult services: the main challenges. Expert Review of Quality of Life in Cancer Care, 2017, 2, 137-152.	0.6	5
141	Adolescents with cancer on privacy: Fact-finding survey on the need for confidentiality and space. Tumori, 2021, 107, 452-457.	1.1	5
142	Medulloblastoma and familial adenomatous polyposis: Good prognosis and good quality of life in the longâ€ŧerm?. Pediatric Blood and Cancer, 2021, 68, e28912.	1.5	5
143	Celiac Disease and Childhood Cancer. Journal of Pediatric Hematology/Oncology, 2006, 28, 346-349.	0.6	4
	Comment on: The LIK Experience of a Treatment Strategy for Pediatric Metastatic Medulloblastoma		

Comment on: The UK Experience of a Treatment Strategy for Pediatric Metastatic Medulloblastoma Comprising Intensive Induction Chemotherapy, Hyperfractionated Accelerated Radiotherapy, and Response Directed Highâ€Dose Myeloablative Chemotherapy or Maintenance Chemotherapy (Milan) Tj ETQq0 0 0 15BT /Overlock 10 Tf 144

#	Article	IF	CITATIONS
145	Salvage treatment for children with relapsed/refractory germ cell tumors: The Associazione Italiana Ematologia Oncologia Pediatrica (AIEOP) experience. Pediatric Blood and Cancer, 2020, 67, e28125.	1.5	4
146	Psychological support in children and adolescents with cancer when amputation is required. Medical and Pediatric Oncology, 2002, 38, 261-265.	1.0	3
147	Case Report: Pseudomonas aeruginosa-related Intervertebral Discitis in a Young Boy with Medulloblastoma. Journal of Neuro-Oncology, 2004, 68, 245-248.	2.9	3
148	Wilms Tumor in Monozygous Twins. Journal of Pediatric Hematology/Oncology, 2005, 27, 521-525.	0.6	3
149	A novel WT1 mutation in a 46,XY boy with congenital bilateral cryptorchidism, nystagmus and Wilms tumor. Pediatric Nephrology, 2009, 24, 1413-1417.	1.7	3
150	A female survivor of childhood medulloblastoma presenting with growth-hormone-induced edema and inflammatory lesions: a case report. Journal of Medical Case Reports, 2009, 3, 17.	0.8	3
151	Clinical and molecular description of a Wilms tumor in a patient with tuberous sclerosis complex. American Journal of Medical Genetics, Part A, 2011, 155, 1419-1424.	1.2	3
152	Winners' Cup: A National Football Tournament Brings Together Adolescent Patients with Cancer from all over Italy. Tumori, 2017, 103, e25-e29.	1.1	3
153	Cancer treatment in disabled children. European Journal of Pediatrics, 2020, 179, 1353-1360.	2.7	3
154	Long-term results of suppressing thyroid-stimulating hormone during radiotherapy to prevent primary hypothyroidism in medulloblastoma/PNET and Hodgkin lymphoma: a prospective cohort study. Frontiers of Medicine, 2021, 15, 101-107.	3.4	3
155	Mirror therapy for phantom limb pain in an adolescent cancer survivor. Tumori, 2012, 98, e27-30.	1.1	3
156	Extraosseous Ewing sarcoma in children and adolescents: A retrospective series from a referral pediatric oncology center. Pediatric Blood and Cancer, 2022, 69, e29512.	1.5	3
157	Transitory, spontaneously recovering, peripheral facial nerve palsy after vinorelbine administration. Neurological Sciences, 2006, 27, 110-113.	1.9	2
158	Cancer of the Kidney, Bladder, and Prostate. Pediatric Oncology, 2017, , 429-451.	0.5	2
159	Rectal Burkitt Lymphoma in Childhood. Journal of Pediatric Hematology/Oncology, 2008, 30, 176-178.	0.6	1
160	Bilateral testicular germ cell tumors. Journal of Pediatric Surgery, 2014, 49, 1341.	1.6	1
161	Towards evidence-based management of paediatric RCC. Nature Reviews Urology, 2015, 12, 426-428.	3.8	1
162	The Role of Alfa Fetoprotein in the Risk Management of Pediatric Germ Cell Tumors. Journal of Pediatric Biochemistry, 2016, 05, 157-160.	0.2	1

#	Article	IF	CITATIONS
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