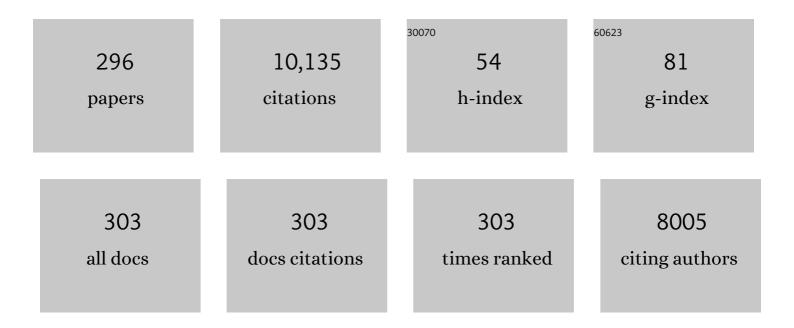
Gianni Bisogno

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Local treatment of rhabdomyosarcoma of the female genital tract: Expert consensus from the Children's Oncology Group, the European Softâ€īssue Sarcoma Group, and the Cooperative Weichteilsarkom Studiengruppe. Pediatric Blood and Cancer, 2023, 70, e28601.	1.5	18
2	Surgical management of extremity rhabdomyosarcoma: A consensus opinion from the Children's Oncology Group, the European Pediatric Softâ€Tissue Sarcoma Study Group, and the Cooperative Weichteilsarkom Studiengruppe. Pediatric Blood and Cancer, 2023, 70, e28608.	1.5	14
3	Introducing INSTRuCT: An international effort to promote cooperation and data sharing. Pediatric Blood and Cancer, 2023, 70, e28701.	1.5	20
4	Congenital rhabdomyosarcoma: A report from the European paediatric Soft tissue sarcoma Study Group. Pediatric Blood and Cancer, 2022, 69, e29376.	1.5	3
5	Localised rhabdomyosarcoma in infants (<12 months) and young children (12–36 months of age) treated on the EpSSG RMS 2005 study. European Journal of Cancer, 2022, 160, 206-214.	2.8	8
6	Factors associated with postoperative hypocalcemia following thyroidectomy in childhood. Pediatric Blood and Cancer, 2022, , e29576.	1.5	3
7	Reply to H. B et al. Journal of Clinical Oncology, 2022, , JCO2102612.	1.6	1
8	Phase 2 study for nonmetastatic extremity highâ€grade osteosarcoma in pediatric and adolescent and young adult patients with a riskâ€adapted strategy based on ABCB1/Pâ€glycoprotein expression: An Italian Sarcoma Group trial (ISC/OSâ€2). Cancer, 2022, 128, 1958-1966.	4.1	12
9	Clinical significance of circulating tumor cells and cellâ€free DNA in pediatric rhabdomyosarcoma. Molecular Oncology, 2022, 16, 2071-2085.	4.6	7
10	Radiomic features as biomarkers of soft tissue paediatric sarcomas: preliminary results of a PET/MR study. Radiology and Oncology, 2022, 56, 138-141.	1.7	4
11	In transit metastases in children, adolescents and young adults with localized rhabdomyosarcoma of the distal extremities: Analysis of the EpSSG RMS 2005 study. European Journal of Surgical Oncology, 2022, 48, 1536-1542.	1.0	6
12	Longâ€ŧerm results from the multicentric European randomized phase 3 trial CWS/RMSâ€96 for localized highâ€risk soft tissue sarcoma in children, adolescents, and young adults. Pediatric Blood and Cancer, 2022, 69, e29691.	1.5	11
13	Congenital spindle cell rhabdomyosarcoma: An international cooperative analysis. European Journal of Cancer, 2022, 168, 56-64.	2.8	8
14	Perianal/perineal rhabdomyosarcoma: Results of the SIOP MMT 95, Italian RMS 96, and EpSSG RMS 2005 studies. Pediatric Blood and Cancer, 2022, 69, e29739.	1.5	4
15	Metastatic Rhabdomyosarcoma: Results of the European <i>Paediatric</i> Soft Tissue Sarcoma Study Group MTS 2008 Study and Pooled Analysis With the Concurrent BERNIE Study. Journal of Clinical Oncology, 2022, 40, 3730-3740.	1.6	22
16	Immunoreactivity against fibroblast growth factor 8 in alveolar rhabdomyosarcoma patients and its involvement in tumor aggressiveness. Oncolmmunology, 2022, 11, .	4.6	2
17	Adolescents and young adults with rhabdomyosarcoma treated in the European paediatric Soft tissue sarcoma Study Croup (EpSSC) protocols: a cohort study. The Lancet Child and Adolescent Health, 2022, 6, 545-554.	5.6	11
18	Molecular testing of rhabdomyosarcoma in clinical trials to improve risk stratification and outcome: A consensus view from European paediatric Soft tissue sarcoma Study Group, Children's Oncology Group and Cooperative Weichteilsarkom-Studiengruppe. European Journal of Cancer, 2022, 172, 367-386.	2.8	19

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19	Therapy and prognostic significance of regional lymph node involvement in embryonal rhabdomyosarcoma: aÂreport from the European paediatric Soft tissue sarcoma Study Group. European Journal of Cancer, 2022, 172, 119-129.	2.8	4
20	Impact of Rhabdomyosarcoma Treatment Modalities by Age in a Population-Based Setting. Journal of Adolescent and Young Adult Oncology, 2021, 10, 309-315.	1.3	7
21	Alveolar rhabdomyosarcoma with regional nodal involvement: Results of a combined analysis from two cooperative groups. Pediatric Blood and Cancer, 2021, 68, e28832.	1.5	13
22	Pathology of childhood rhabdomyosarcoma: A consensus opinion document from the Children's Oncology Group, European Paediatric Soft Tissue Sarcoma Study Group, and the Cooperative Weichteilsarkom Studiengruppe. Pediatric Blood and Cancer, 2021, 68, e28798.	1.5	38
23	Surgical management of paratesticular rhabdomyosarcoma: A consensus opinion from the Children's Oncology Group, European paediatric Soft tissue sarcoma Study Group, and the Cooperative Weichteilsarkom Studiengruppe. Pediatric Blood and Cancer, 2021, 68, e28938.	1.5	20
24	Extraâ€appendicular neuroendocrine tumors: A report from the TREP project (2000â€⊋020). Pediatric Blood and Cancer, 2021, 68, e28880.	1.5	1
25	Integrating irinotecan in standard chemotherapy: A novel doseâ€density combination for highâ€risk pediatric sarcomas. Pediatric Blood and Cancer, 2021, 68, e28951.	1.5	10
26	Consensus recommendations from the EXPeRT/PARTNER groups for the diagnosis and therapy of sex cord stromal tumors in children and adolescents. Pediatric Blood and Cancer, 2021, 68, e29017.	1.5	13
27	Volumetric histograms-based analysis of apparent diffusion coefficients and standard uptake values for the assessment of pediatric sarcoma at staging: preliminary results of a PET/MRI study. Radiologia Medica, 2021, 126, 878-885.	7.7	18
28	Embryonal rhabdomyosarcoma completely resected at diagnosis: The European paediatric Soft tissue sarcoma Study Group RMS2005 experience. European Journal of Cancer, 2021, 146, 21-29.	2.8	20
29	Thymoma and thymic carcinoma in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. Pediatric Blood and Cancer, 2021, 68, e29042.	1.5	5
30	Case Report: Circulating Tumor Cells as a Response Biomarker in ALK-Positive Metastatic Inflammatory Myofibroblastic Tumor. Frontiers in Pediatrics, 2021, 9, 652583.	1.9	3
31	Nasopharyngeal carcinoma in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. Pediatric Blood and Cancer, 2021, 68, e29018.	1.5	11
32	The European Paediatric Rare Tumours Network ―European Registry (PARTNER) project for very rare tumors in children. Pediatric Blood and Cancer, 2021, 68, e29072.	1.5	11
33	Pleuropulmonary blastoma in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. Pediatric Blood and Cancer, 2021, 68, e29045.	1.5	15
34	Efficacy of dose intensification in induction therapy for localized Ewing sarcoma: Italian Sarcoma Group (ISG) and Associazione Italiana Ematologia ed Oncologia Pediatrica (AIEOP) ISG/AIEOP EW-1 study Journal of Clinical Oncology, 2021, 39, 11501-11501.	1.6	2
35	Salivary gland carcinoma in children and adolescents: The EXPeRT/PARTNER diagnosis and treatment recommendations. Pediatric Blood and Cancer, 2021, 68, e29058.	1.5	7
36	Pancreatoblastoma in children: EXPeRT/PARTNER diagnostic and therapeutic recommendations. Pediatric Blood and Cancer, 2021, 68, e29112.	1.5	9

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37	Cutaneous melanoma in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. Pediatric Blood and Cancer, 2021, 68, e28992.	1.5	9
38	Adrenocortical tumours in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. Pediatric Blood and Cancer, 2021, 68, e29025.	1.5	16
39	Whole Lung Irradiation after High-Dose Busulfan/Melphalan in Ewing Sarcoma with Lung Metastases: An Italian Sarcoma Group and Associazione Italiana Ematologia Oncologia Pediatrica Joint Study. Cancers, 2021, 13, 2789.	3.7	1
40	Facing the challenges of very rare tumors of pediatric age: The European Cooperative Study Group for Pediatric Rare Tumors (EXPeRT) background, goals, and achievements. Pediatric Blood and Cancer, 2021, 68, e28993.	1.5	10
41	The Impact of Radiation Therapy in Children and Adolescents With Metastatic Rhabdomyosarcoma. International Journal of Radiation Oncology Biology Physics, 2021, 111, 968-978.	0.8	15
42	Role of centers with different patient volumes in the management of rhabdomyosarcoma. An analysis by the Italian Pediatric Soft Tissue Sarcoma Committee. Pediatric Blood and Cancer, 2021, 68, e29234.	1.5	1
43	Non-parameningeal head and neck rhabdomyosarcoma in children, adolescents, and young adults: Experience of the European paediatric Soft tissue sarcoma Study Group (EpSSC) – RMS2005 study. European Journal of Cancer, 2021, 151, 84-93.	2.8	21
44	Paediatric non-rhabdomyosarcoma soft tissue sarcomas: the prospective NRSTS 2005 study by the European Pediatric Soft Tissue Sarcoma Study Group (EpSSG). The Lancet Child and Adolescent Health, 2021, 5, 546-558.	5.6	28
45	Randomized Phase II Trial of Vincristine-Irinotecan With or Without Temozolomide, in Children and Adults With Relapsed or Refractory Rhabdomyosarcoma: A European Paediatric Soft Tissue Sarcoma Study Group and Innovative Therapies for Children With Cancer Trial. Journal of Clinical Oncology, 2021, 39, 2979-2990.	1.6	38
46	Role of 18F-FDG-PET/CT in the staging of metastatic rhabdomyosarcoma: a report from the European paediatric Soft tissue sarcoma Study Group. European Journal of Cancer, 2021, 155, 155-162.	2.8	17
47	Children with progressive and relapsed pleuropulmonary blastoma: A European collaborative analysis. Pediatric Blood and Cancer, 2021, 68, e29268.	1.5	4
48	Melanoma in Adolescents and Young Adults: Evaluation of the Characteristics, Treatment Strategies, and Prognostic Factors in a Monocentric Retrospective Study. Frontiers in Oncology, 2021, 11, 725523.	2.8	2
49	Autoantibody profiling of alveolar rhabdomyosarcoma patients unveils tumor-associated antigens with diagnostic and prognostic significance. Oncolmmunology, 2021, 10, 1954765.	4.6	1
50	Pediatric Cancer Data Commons: Federating and Democratizing Data for Childhood Cancer Research. JCO Clinical Cancer Informatics, 2021, 5, 1034-1043.	2.1	18
51	Treatment Strategies for Children With Relapsed Pancreatoblastoma: A Literature Review. Journal of Pediatric Hematology/Oncology, 2021, 43, 288-293.	0.6	5
52	Treatment of Rhabdomyosarcoma. Pediatric Oncology, 2021, , 53-63.	0.5	0
53	Pediatric airway tumors: A report from the International Network of Pediatric Airway Teams (INPAT). Laryngoscope, 2020, 130, E243-E251.	2.0	17
54	High dose chemotherapy and autologous hematopoietic cell transplantation for Wilms tumor: a study of the European Society for Blood and Marrow Transplantation. Bone Marrow Transplantation, 2020, 55, 376-383.	2.4	7

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55	Where Are Adolescents with Soft Tissue Sarcomas Treated? An Italian Nationwide Study on Referrals Based on Hospital Discharge Records. Journal of Adolescent and Young Adult Oncology, 2020, 9, 190-195.	1.3	7
56	Pleuropulmonary blastoma: a report from the TREP (Tumori Rari in Età Pediatrica) Project. Tumori, 2020, 106, 126-132.	1.1	6
57	Inequalities in diagnosis and registration of pediatric very rare tumors: a European study on pleuropulmonary blastoma. European Journal of Pediatrics, 2020, 179, 749-756.	2.7	9
58	A new standard of care for patients with high-risk rhabdomyosarcoma? – Authors' reply. Lancet Oncology, The, 2020, 21, e3.	10.7	2
59	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
60	Is surveillance imaging in pediatric patients treated for localized rhabdomyosarcoma useful? The European experience. Cancer, 2020, 126, 823-831.	4.1	21
61	Mesothelioma in children and adolescents: the European Cooperative Study Group for Pediatric Rare Tumors (EXPeRT) contribution. European Journal of Cancer, 2020, 140, 63-70.	2.8	12
62	Pediatric sarcomas display a variable EpCAM expression in a histology-dependent manner. Translational Oncology, 2020, 13, 100846.	3.7	8
63	Clinical Characteristics and Outcome of Severe Acute Respiratory Syndrome Coronavirus 2 Infection in Italian Pediatric Oncology Patients: A Study From the Infectious Diseases Working Group of the Associazione Italiana di Oncologia e Ematologia Pediatrica. Journal of the Pediatric Infectious Diseases Society. 2020. 9. 530-534.	1.3	81
64	Spotlight on the treatment of infantile fibrosarcoma in the era of neurotrophic tropomyosin receptor kinase inhibitors: International consensus and remaining controversies. European Journal of Cancer, 2020, 137, 183-192.	2.8	28
65	Pediatric Soft Tissue Tumors With BCOR ITD Express EGFR but Not OLIG2. Pediatric and Developmental Pathology, 2020, 23, 424-430.	1.0	7
66	Liquid Biopsy in Pediatric Renal Cancer: Stage I and Stage IV Cases Compared. Diagnostics, 2020, 10, 810.	2.6	1
67	Defining the impact of prognostic factors at the time of relapse for nonmetastatic rhabdomyosarcoma. Pediatric Blood and Cancer, 2020, 67, e28674.	1.5	7
68	Circulating miR-26a as Potential Prognostic Biomarkers in Pediatric Rhabdomyosarcoma. Frontiers in Genetics, 2020, 11, 606274.	2.3	10
69	Pediatric Rhabdomyosarcomas: Three-Dimensional Radiological Assessments after Induction Chemotherapy Predict Survival Better than One-Dimensional and Two-Dimensional Measurements. Cancers, 2020, 12, 3808.	3.7	9
70	Undifferentiated round cell sarcoma with BCOR internal tandem duplications (ITD) or YWHAE fusions: a clinicopathologic and molecular study. Modern Pathology, 2020, 33, 1669-1677.	5.5	29
71	Phase II results from a phase I/II study to assess the safety and efficacy of weekly nab-paclitaxel in paediatric patients with recurrent or refractory solid tumours: A collaboration with the European Innovative Therapies for Children with Cancer Network. European Journal of Cancer, 2020, 135, 89-97.	2.8	13
72	Paratesticular rhabdomyosarcoma—Impact of locoregional approach on patient outcome: A report from the European paediatric Soft tissue sarcoma Study Group (EpSSG). Pediatric Blood and Cancer, 2020, 67, e28479.	1.5	13

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73	Pheochromocytomas and paragangliomas in children: Data from the Italian Cooperative Study (TREP). Pediatric Blood and Cancer, 2020, 67, e28332.	1.5	12
74	Secondary Osteosarcoma After Bone Marrow Transplant: An Aggressive Disease. Journal of Adolescent and Young Adult Oncology, 2020, 9, 672-675.	1.3	4
75	Plerixafor combined with standard regimens for hematopoietic stem cell mobilization in pediatric patients with solid tumors eligible for autologous transplants: two-arm phase I/II study (MOZAIC). Bone Marrow Transplantation, 2020, 55, 1744-1753.	2.4	12
76	Prognostic Value of Circulating IGFBP2 and Related Autoantibodies in Children with Metastatic Rhabdomyosarcomas. Diagnostics, 2020, 10, 115.	2.6	8
77	Genetic Polymorphisms Affecting Cardiac Biomarker Concentrations in Children with Cancer: an Analysis from the "European Paediatric Oncology Off-patents Medicines Consortium―(EPOC) Trial. European Journal of Drug Metabolism and Pharmacokinetics, 2020, 45, 413-422.	1.6	7
78	Inflammatory myofibroblastic tumor: The experience of the European pediatric Soft Tissue Sarcoma Study Group (EpSSG). European Journal of Cancer, 2020, 127, 123-129.	2.8	71
79	Outcomes of metastatic non-rhabdomyosarcoma soft tissue sarcomas (NRSTS) treated within the BERNIE study: a randomised, phase II study evaluating the addition of bevacizumab to chemotherapy. European Journal of Cancer, 2020, 130, 72-80.	2.8	18
80	Treatment of rhabdomyosarcoma in children and adolescent from four low health expenditures average rates countries. Radiology and Oncology, 2020, 54, 455-460.	1.7	2
81	An unresolved issue in rhabdomyosarcoma treatment: The duration of chemotherapy. Pediatric Blood and Cancer, 2020, 67, e28174.	1.5	8
82	Topotecan/carboplatin regimen for refractory/recurrent rhabdomyosarcoma in children: Report from the AIEOP Soft Tissue Sarcoma Committee. Tumori, 2019, 105, 138-143.	1.1	9
83	Prognostic role of pleural effusion or ascites in localized rhabdomyosarcoma. Pediatric Blood and Cancer, 2019, 66, e27932.	1.5	3
84	Outcome and prognostic factors in pediatric malignant peripheral nerve sheath tumors: An analysis of the European Pediatric Soft Tissue Sarcoma Group (EpSSG) NRSTSâ€2005 prospective study. Pediatric Blood and Cancer, 2019, 66, e27833.	1.5	40
85	Vinorelbine and continuous low-dose cyclophosphamide as maintenance chemotherapy in patients with high-risk rhabdomyosarcoma (RMS 2005): a multicentre, open-label, randomised, phase 3 trial. Lancet Oncology, The, 2019, 20, 1566-1575.	10.7	161
86	PAlâ€1 and protein C as early markers of venoâ€occlusive disease in patients treated for Wilms tumor. Pediatric Blood and Cancer, 2019, 66, e27695.	1.5	2
87	Indeterminate Pulmonary Nodules at Diagnosis in Rhabdomyosarcoma: Are They Clinically Significant? A Report From the European Paediatric Soft Tissue Sarcoma Study Group. Journal of Clinical Oncology, 2019, 37, 723-730.	1.6	24
88	Clinical features and outcomes of young patients with epithelioid sarcoma: an analysis from the Children's Oncology Group and the European paediatric soft tissue Sarcoma Study Group prospective clinical trials. European Journal of Cancer, 2019, 112, 98-106.	2.8	21
89	Outcome of localized liverâ€bile duct rhabdomyosarcoma according to local therapy: A report from the European Paediatric Softâ€īssue Sarcoma Study Group (EpSSG)â€RMS 2005 study. Pediatric Blood and Cancer, 2019, 66, e27725.	1.5	11
90	Defining and listing very rare cancers of paediatric age: consensus of the Joint Action on Rare CancersÂin cooperation with the European Cooperative Study Group for Pediatric Rare Tumors. European Journal of Cancer, 2019, 110, 120-126.	2.8	61

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91	Cell-Free DNA in Pediatric Rhabdomyosarcoma: Potential and Challenges. Methods in Molecular Biology, 2019, 1909, 165-175.	0.9	4
92	Outcome after surgery for solid pseudopapillary pancreatic tumors in children: Report from the TREP project—Italian Rare Tumors Study Group. Pediatric Blood and Cancer, 2019, 66, e27519.	1.5	16
93	Evidence of hydroxyurea activity in children with pretreated desmoidâ€ŧype fibromatosis: A new option in the armamentarium of systemic therapies. Pediatric Blood and Cancer, 2019, 66, e27472.	1.5	14
94	Joining forces for pediatric very rare tumors. Oncotarget, 2019, 10, 3084-3085.	1.8	5
95	Genomic complexity in pediatric synovial sarcomas (Synobio study): the European pediatric soft tissue sarcoma group (EpSSG) experience. Cancer Medicine, 2018, 7, 1384-1393.	2.8	22
96	Pediatric patients with cutaneous melanoma: A European study. Pediatric Blood and Cancer, 2018, 65, e26974.	1.5	26
97	Alveolar soft part sarcoma in children and adolescents: The European Paediatric Soft Tissue Sarcoma study group prospective trial (EpSSG NRSTS 2005). Pediatric Blood and Cancer, 2018, 65, e26942.	1.5	21
98	Appraisal of the role of radical prostatectomy for rhabdomyosarcoma in children: oncological and urological outcome. Therapeutic Advances in Urology, 2018, 10, 189-196.	2.0	7
99	Salvage rates and prognostic factors after relapse in children and adolescents with malignant peripheral nerve sheath tumors. Pediatric Blood and Cancer, 2018, 65, e26816.	1.5	14
100	Longâ€ŧerm results in children with head and neck rhabdomyosarcoma: A report from the Italian Soft Tissue Sarcoma Committee. Pediatric Blood and Cancer, 2018, 65, e26876.	1.5	13
101	Demographic and Treatment Variables Influencing Outcome for Localized Paratesticular Rhabdomyosarcoma: Results From a Pooled Analysis of North American and European Cooperative Groups. Journal of Clinical Oncology, 2018, 36, 3466-3476.	1.6	21
102	Malignant testicular germ cell tumors in children and adolescents: The AIEOP (Associazione Italiana) Tj ETQq0 0 Investigations, 2018, 36, 502.e7-502.e13.	D rgBT /Ov 1.6	verlock 10 Tf 5 13
103	Fusion status in patients with lymph nodeâ€positive (N1) alveolar rhabdomyosarcoma is a powerful predictor of prognosis: Experience of the European Paediatric Soft Tissue Sarcoma Study Group (EpSSG). Cancer, 2018, 124, 3201-3209.	4.1	51
104	Localized vaginal/uterine rhabdomyosarcoma—results of a pooled analysis from four international cooperative groups. Pediatric Blood and Cancer, 2018, 65, e27096.	1.5	40
105	Central nervous system involvement in children and adolescents with rhabdomyosarcoma. A report from the AIEOP Soft Tissue Sarcoma Committee. Neuro-Oncology Practice, 2018, 5, 241-245.	1.6	3
106	Paraneoplastic Cytopenia in Ovarian Germ Cell Tumor: A Novel Presentation. Journal of Pediatric Hematology/Oncology, 2018, 40, e432-e434.	0.6	1
107	Fifty years of rhabdomyosarcoma studies on both sides of the pond and lessons learned. Cancer Treatment Reviews, 2018, 68, 94-101.	7.7	48
108	Phase I results of a phase I/II study of weekly nab-paclitaxel in paediatric patients with recurrent/refractory solid tumours: A collaboration with innovative therapies for children with cancer. European Journal of Cancer, 2018, 100, 27-34.	2.8	22

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109	Addition of dose-intensified doxorubicin to standard chemotherapy for rhabdomyosarcoma (EpSSG) Tj ETQq1 1 (19, 1061-1071.	0.784314 r 10.7	gBT /Overlo 137
110	Biliary tract rhabdomyosarcoma: a report from the Soft Tissue Sarcoma Committee of the Associazione Italiana Ematologia Oncologia Pediatrica. Tumori, 2018, 104, 232-237.	1.1	12
111	Maintenance low-dose chemotherapy in patients with high-risk (HR) rhabdomyosarcoma (RMS): A report from the European Paediatric Soft Tissue Sarcoma Study Group (EpSSG) Journal of Clinical Oncology, 2018, 36, LBA2-LBA2.	1.6	23
112	Pediatric nonrhabdomyosarcoma soft tissue sarcomas arising at visceral sites. Pediatric Blood and Cancer, 2017, 64, e26490.	1.5	6
113	The challenge of very rare childhood cancers in developed and developing countries. Expert Opinion on Orphan Drugs, 2017, 5, 331-341.	0.8	12
114	Extraskeletal osteosarcoma: A European Musculoskeletal Oncology Society study on 266 patients. European Journal of Cancer, 2017, 74, 9-16.	2.8	67
115	Surgery alone is sufficient therapy for children and adolescents with low-risk synovial sarcoma: A joint analysis from the European paediatric soft tissue sarcoma Study Group and the Children's Oncology Group. European Journal of Cancer, 2017, 78, 1-6.	2.8	62
116	Malignant ovarian germ cell tumors in pediatric patients: The AIEOP (Associazione Italiana Ematologia) Tj ETQqO	0	Overlock 10
117	Alveolar Rhabdomyosarcoma Decellularization. Methods in Molecular Biology, 2017, 1577, 317-325.	0.9	4
118	Imatinib mesylate in desmoplastic small round cell tumors. Future Oncology, 2017, 13, 1233-1237.	2.4	13
119	Attachment orientations and psychological adjustment of parents of children with cancer: A matched-group comparison. Journal of Psychosocial Oncology, 2017, 35, 726-740.	1.2	6
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	Gastrointestinal tract carcinoma in pediatric and adolescent age: The Italian TREP project experience. Pediatric Blood and Cancer, 2017, 64, e26658.	1.5	22
122	Outcome and prognostic factors in highâ€risk childhood adrenocortical carcinomas: A report from the European Cooperative Study Group on Pediatric Rare Tumors (EXPeRT). Pediatric Blood and Cancer, 2017, 64, e26368.	1.5	47
123	Radiologic Response Assessment in Pediatric Soft Tissue Sarcoma: Computed-Assisted Volume Evaluation. Journal of Pediatrics, 2017, 182, 327-334.e2.	1.8	7
124	Open-label, multicentre, randomised, phase II study of the EpSSG and the ITCC evaluating the addition of bevacizumab to chemotherapy in childhood and adolescent patients with metastatic soft tissue sarcoma (the BERNIE study). European Journal of Cancer, 2017, 83, 177-184.	2.8	70
125	The EpSSG NRSTS 2005 treatment protocol for desmoid-type fibromatosis in children: an international prospective case series. The Lancet Child and Adolescent Health, 2017, 1, 284-292.	5.6	43
126	New insights into sinusoidal obstruction syndrome. Internal Medicine Journal, 2017, 47, 1173-1183.	0.8	24

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127	Osteopathia striata with cranial sclerosis and Wilms tumor: Coincidence or consequence?. Clinical Genetics, 2017, 92, 674-675.	2.0	9
128	Towards a Model-Based Dose Recommendation for Doxorubicin in Children. Clinical Pharmacokinetics, 2017, 56, 215-223.	3.5	5
129	Access to clinical trials for adolescents with soft tissue sarcomas: Enrollment in European pediatric Soft tissue sarcoma Study Group (EpSSG) protocols. Pediatric Blood and Cancer, 2017, 64, e26348.	1.5	32
130	Winners' Cup: A National Football Tournament Brings Together Adolescent Patients with Cancer from all over Italy. Tumori, 2017, 103, e25-e29.	1.1	3
131	NELL1, whose high expression correlates with negative outcomes, has different methylation patterns in alveolar and embryonal rhabdomyosarcoma. Oncotarget, 2017, 8, 33086-33099.	1.8	14
132	Outcome of extracranial malignant rhabdoid tumours in children registered in the European Paediatric Soft Tissue Sarcoma Study Group Non-Rhabdomyosarcoma Soft Tissue Sarcoma 2005 Study—EpSSG NRSTS 2005. European Journal of Cancer, 2016, 60, 69-82.	2.8	63
133	Prognostic factors in children undergoing salvage surgery for bladder/prostate rhabdomyosarcoma. Journal of Pediatric Urology, 2016, 12, 265.e1-265.e8.	1.1	4
134	Surgical management of papillary thyroid carcinoma in childhood and adolescence: an Italian multicenter study on 250 patients. Journal of Endocrinological Investigation, 2016, 39, 1055-1059.	3.3	38
135	Pharmacokinetic and pharmacodynamic study of doxorubicin in children with cancer: results of a "European Pediatric Oncology Off-patents Medicines Consortium―trial. Cancer Chemotherapy and Pharmacology, 2016, 78, 1175-1184.	2.3	25
136	A Molecular Study of Pediatric Spindle and Sclerosing Rhabdomyosarcoma. American Journal of Surgical Pathology, 2016, 40, 224-235.	3.7	208
137	Clobal DNA methylation profiling uncovers distinct methylation patterns of protocadherin alpha4 in metastatic and non-metastatic rhabdomyosarcoma. BMC Cancer, 2016, 16, 886.	2.6	13
138	Conservative strategy in infantile fibrosarcoma is possible: The European paediatric Soft tissue sarcoma Study GroupÂexperience. European Journal of Cancer, 2016, 57, 1-9.	2.8	94
139	Anaplastic lymphoma kinase aberrations correlate with metastatic features in pediatric rhabdomyosarcoma. Oncotarget, 2016, 7, 58903-58914.	1.8	15
140	Papillary Thyroid Carcinoma in Pediatric Age: An Example of a Rare Tumour Managed Within a Cooperative Comprehensive Project. Current Pediatric Reviews, 2016, 12, 265-271.	0.8	1
141	Prognostic Factors for Outcome in Localized Extremity Rhabdomyosarcoma. Pooled Analysis from Four International Cooperative Groups. Pediatric Blood and Cancer, 2015, 62, 2125-2131.	1.5	23
142	Molecular Cytogenetics Detect an Unbalanced t(2;13)(q36;q14) and <i>PAX3</i> -FOXO1 Fusion in Rhabdomyosarcoma With Mixed Embryonal/Alveolar Features. Pediatric Blood and Cancer, 2015, 62, 2238-2241.	1.5	5
143	MicroRNA-27a Contributes to Rhabdomyosarcoma Cell Proliferation by Suppressing RARA and RXRA. PLoS ONE, 2015, 10, e0125171.	2.5	26
144	DAX-1 Expression in Pediatric Rhabdomyosarcomas: Another Immunohistochemical Marker Useful in the Diagnosis of Translocation Positive Alveolar Rhabdomyosarcoma. PLoS ONE, 2015, 10, e0133019.	2.5	0

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145	Mature and immature teratoma: A report from the second Italian pediatric study. Pediatric Blood and Cancer, 2015, 62, 1202-1208.	1.5	47
146	Ovarian Sertoli Leydig cell tumours in children and adolescents: An analysis of the European Cooperative Study Group on Pediatric Rare Tumors (EXPeRT). European Journal of Cancer, 2015, 51, 543-550.	2.8	62
147	Management and followâ€up of urothelial neoplasms of the bladder in children: A report from the TREP project. Pediatric Blood and Cancer, 2015, 62, 1000-1003.	1.5	19
148	Epithelial Tumors of the Ovary in Children and Teenagers: A Prospective Study from the Italian TREP Project. Journal of Pediatric and Adolescent Gynecology, 2015, 28, 441-446.	0.7	17
149	The Role of Registries and Tumor Banking in Rare Pediatric Tumors. Current Pediatrics Reports, 2015, 3, 128-136.	4.0	1
150	Thymoma and thymic carcinoma in children and adolescents: A report from the European Cooperative Study Group for Pediatric Rare Tumors (EXPeRT). European Journal of Cancer, 2015, 51, 2444-2452.	2.8	52
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