Daniel Orbach

List of Publications by Year
in descending order

231 papers	7,808 citations	50276 46 h-index	79698 73 g-index
304 all docs	304 docs citations	304 times ranked	7678 citing authors

#	Article	IF	Citations
1	Testicular Sertoli cell tumour and potentially testicular Leydig cell tumour are features of DICER1 syndrome. Journal of Medical Genetics, 2022, 59, 346-350.	3.2	4
2	Robotic Surgery in Pediatric Oncology: Lessons Learned from the First 100 Tumorsâ€"A Nationwide Experience. Annals of Surgical Oncology, 2022, 29, 1315-1326.	1.5	17
3	Lessons from a large nationwide cohort of 350 children with ovarian mature teratoma: A study in favor of ovarianâ€sparing surgery. Pediatric Blood and Cancer, 2022, 69, e29421.	1.5	2
4	Localised rhabdomyosarcoma in infants (<12 months) and young children (12â€"36 months of age) treated on the EpSSG RMS 2005 study. European Journal of Cancer, 2022, 160, 206-214.	2.8	8
5	PET metabolic tumor volume as a new prognostic factor in childhood rhabdomyosarcoma. PLoS ONE, 2022, 17, e0261565.	2.5	3
6	Extracranial rhabdoid tumours: Results of a SFCE series of patients treated with a dose compression strategy according to European Paediatric Soft tissue sarcoma Study Group recommendations. European Journal of Cancer, 2022, 161, 64-78.	2.8	7
7	Varicella post-exposure management for pediatric oncology patients. Bulletin Du Cancer, 2022, 109, 287-295.	1.6	4
8	Intraâ€•and extraâ€cranial <scp>BCORâ€ITD</scp> tumours are separate entities within the <scp>BCOR</scp>â€rearranged family. Journal of Pathology: Clinical Research, 2022, 8, 217-232.	3.0	10
9	Brachytherapy for Pediatric Patients at Gustave Roussy Cancer Campus: A Model of International Cooperation for Highly Specialized Treatments. International Journal of Radiation Oncology Biology Physics, 2022, 113, 602-613.	0.8	11
10	Lack of Prognostic Value of CTNNB1 Mutation Profile in Desmoid-Type Fibromatosis. Clinical Cancer Research, 2022, 28, 4105-4111.	7.0	11
11	Controversies and challenges in the management of paediatric non-rhabdomyosarcoma soft tissue sarcomas. The Lancet Child and Adolescent Health, 2022, 6, 221-223.	5.6	10
12	High Prevalence of Early Endocrine Disorders After Childhood Brain Tumors in a Large Cohort. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e2156-e2166.	3.6	6
13	Clinical, pathologic, and molecular features of inflammatory myofibroblastic tumors in children and adolescents. Pediatric Blood and Cancer, 2022, 69, e29460.	1.5	13
14	NUT carcinoma in children, adolescents and young adults. Bulletin Du Cancer, 2022, 109, 491-504.	1.6	2
15	Desmoid tumors located in the abdomen or associated with adenomatous polyposis: French intergroup clinical practice guidelines for diagnosis, treatment, and follow-up (SNFGE, FFCD, GERCOR,) Tj ETQq1 1 0.784314 gBT /Ov	0.7	3
16	The treatment approach to pediatric non-rhabdomyosarcoma soft tissue sarcomas: a critical review from the INternational Soft Tissue SaRcoma ConsorTium. European Journal of Cancer, 2022, 169, 10-19.	2.8	13
17	No Geographical Inequalities in Survival for Sarcoma Patients in France: A Reference Networksâ€™ Outcome?. Cancers, 2022, 14, 2620.	3.7	4
18	Metastatic Rhabdomyosarcoma: Results of the European Paediatric 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

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19	Clinical characteristics and outcome of a large cohort of patients with primary central nervous system (CNS) tumors and tropomyosin receptor kinase (TRK) fusion.. Journal of Clinical Oncology, 2022, 40, 2052-2052.	1.6	0
20	Dermatofibrosarcoma protuberans, fibrosarcomatous variant: A rare tumor in children. Pediatric Dermatology, 2021, 38, 217-222.	0.9	9
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	Major response to imatinib and chemotherapy in a newborn patient prenatally diagnosed with generalized infantile myofibromatosis. Pediatric Blood and Cancer, 2021, 68, e28576.	1.5	8
23	Secondâ€line therapy in young patients with relapsed or refractory orbital rhabdomyosarcoma. Acta Ophthalmologica, 2021, 99, 334-341.	1.1	7
24	Management of sarcomas in children, adolescents and adults: Interactions in two different age groups under the umbrellas of GSF-GETO and SFCE, with the support of the NETSARC+ network. Bulletin Du Cancer, 2021, 108, 163-176.	1.6	7
25	Rationale for the use of tyrosine kinase inhibitors in the treatment of paediatric desmoid-type fibromatosis. British Journal of Cancer, 2021, 124, 1637-1646.	6.4	12
26	Head and neck tumors in children and adolescents: Impact of a multidisciplinary tumor board. Oral Oncology, 2021, 114, 105145.	1.5	11
27	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
28	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
29	Infantile Rhabdomyosarcomas With VGLL2 Rearrangement Are Not Always an Indolent Disease. American Journal of Surgical Pathology, 2021, 45, 854-867.	3.7	12
30	Pattern of relapse in pediatric localized extremity rhabdomyosarcomas correlated with locoregional therapies administered. Strahlentherapie Und Onkologie, 2021, 197, 690-699.	2.0	1
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	Controversies on the possible role of immune checkpoint inhibitors in pediatric cancers: balancing irAEs and efficacy. Tumori, 2021, 107, 276-281.	1.1	6
33	The European Paediatric Rare Tumours Network â€•European Registry (PARTNER) project for very rare tumors in children. Pediatric Blood and Cancer, 2021, 68, e29072.	1.5	11
34	Pleuropulmonary blastoma in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. Pediatric Blood and Cancer, 2021, 68, e29045.	1.5	15
35	End-of-life care in children and adolescents with cancer: perspectives from a French pediatric oncology care network. Tumori, 2021, , 030089162110133.	1.1	5
36	Subcutaneous implantable pleural port catheter in the management of malignant pleural effusions in young patients with solid tumors: A new option in the armamentarium of symptomatic treatment. Pediatric Blood and Cancer, 2021, 68, e29109.	1.5	1

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37	Prognosis value of S45F mutation of CTNNB1 in desmoid-type fibromatosis (DF): Prospective analysis of 500 consecutive patients (pts) from ALTITUDES trial.. Journal of Clinical Oncology, 2021, 39, 11510-11510.	1.6	1
38	<scp>SMARCA4</scp>â€deficient rhabdoid tumours show intermediate molecular features between <scp>SMARCB1</scp>â€deficient rhabdoid tumours and small cell carcinomas of the ovary, hypercalcaemic type. Journal of Pathology, 2021, 255, 1-15.	4.5	14
39	Salivary gland carcinoma in children and adolescents: The EXPeRT/PARTNER diagnosis and treatment recommendations. Pediatric Blood and Cancer, 2021, 68, e29058.	1.5	7
40	Pancreatoblastoma in children: EXPeRT/PARTNER diagnostic and therapeutic recommendations. Pediatric Blood and Cancer, 2021, 68, e29112.	1.5	9
41	Cutaneous melanoma in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. Pediatric Blood and Cancer, 2021, 68, e28992.	1.5	9
42	Adrenocortical tumours in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. Pediatric Blood and Cancer, 2021, 68, e29025.	1.5	16
43	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
44	Desmoplastic Small Round Cell Tumors With EWS-WT1 Transcript Expression. Journal of Pediatric Hematology/Oncology, 2021, Publish Ahead of Print, .	0.6	4
45	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
46	Non-parameningeal head and neck rhabdomyosarcoma in children, adolescents, and young adults: Experience of the European paediatric Soft tissue sarcoma Study Group (EpSSG) â€“ RMS2005 study. European Journal of Cancer, 2021, 151, 84-93.	2.8	21
47	PD-0923 Increasing access to highly specialized radiation treatments: the example of pediatric brachytherapy. Radiotherapy and Oncology, 2021, 161, S763-S764.	0.6	0
48	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
49	Locally aggressive rarely metastazing tumors and low-grade sarcoma in children, adolescents and young adults: The benefits of a national network. European Journal of Surgical Oncology, 2021, , .	1.0	2
50	ASO Visual Abstract: Robotic Surgery in Pediatric Oncologyâ€"Lessons Learned from the First 100 Tumors: A Nationwide Experience. Annals of Surgical Oncology, 2021, 28, 730-731.	1.5	0
51	Children with progressive and relapsed pleuropulmonary blastoma: A European collaborative analysis. Pediatric Blood and Cancer, 2021, 68, e29268.	1.5	4
52	1522MO Hormonal contraception and pregnancy and risk of progression or relapse in desmoid-type fibromatosis (DF). Annals of Oncology, 2021, 32, S1112.	1.2	0
53	Childhood head and neck cancer in France: Incidence, survival and trends from 2000 to 2015. International Journal of Pediatric Otorhinolaryngology, 2021, 150, 110858.	1.0	7
54	Current Approaches to Therapy: Soft Tissue Sarcomas Other than Rhabdomyosarcoma in Children and Adolescents. Pediatric Oncology, 2021, , 65-85.	0.5	0

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55	Avoidance or adaptation of radiotherapy in patients with cancer with Li-Fraumeni and heritable TP53-related cancer syndromes. Lancet Oncology, The, 2021, 22, e562-e574.	10.7	22
56	Locoregional Control and Survival in Children, Adolescents, and Young Adults With Localized Head and Neck Alveolar Rhabdomyosarcomaâ€"The French Experience. Frontiers in Pediatrics, 2021, 9, 783754.	1.9	2
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	Exploring heterogeneity of adrenal cortical tumors in children: The French pediatric rare tumor group (Fracture) experience. Pediatric Blood and Cancer, 2020, 67, e28086.	1.5	12
59	Soft tissue sarcoma in children, adolescents and young adults: Outcomes according to compliance with international initial care guidelines. European Journal of Surgical Oncology, 2020, 46, 1277-1286.	1.0	19
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	Esthesioneuroblastoma in children, adolescents and young adults. Bulletin Du Cancer, 2020, 107, 934-945.	1.6	6
63	Rhabdomyosarcoma associated with germline TP53 alteration in children and adolescents: The French experience. Pediatric Blood and Cancer, 2020, 67, e28486.	1.5	19
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	Local staging and treatment in extremity rhabdomyosarcoma. A report from the EpSSGâ€RMS2005 study. Cancer Medicine, 2020, 9, 7580-7589.	2.8	16
66	1650P Desmoid type fibromatosis in patients. Annals of Oncology, 2020, 31, S985.	1.2	0
67	Dermatofibrosarcoma protuberans in children and adolescents: The European Paediatric Soft Tissue Sarcoma Study Group prospective trial (EpSSG NRSTS 2005). Pediatric Blood and Cancer, 2020, 67, e28351.	1.5	11
68	Functional analysis of young patients with desmoid-type fibromatosis: Initial surveillance does not jeopardize long term quality of life. European Journal of Surgical Oncology, 2020, 46, 1294-1300.	1.0	15
69	ESGOâ€"SIOPE guidelines for the management of adolescents and young adults with non-epithelial ovarian cancers. Lancet Oncology, The, 2020, 21, e360-e368.	10.7	50
70	Metronomic Maintenance for High-Risk Pediatric Malignancies: One Size Will Not Fit All. Trends in Cancer, 2020, 6, 819-828.	7.4	20
71	Integrative clinical and biopathology analyses to understand the clinical heterogeneity of infantile rhabdomyosarcoma: A report from the French MMT committee. Cancer Medicine, 2020, 9, 2698-2709.	2.8	28
72	Pure pediatric ovarian immature teratomas: The French experience. Pediatric Blood and Cancer, 2020, 67, e28186.	1.5	7

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73	Pattern of locoâ€regional relapses and treatment in pediatric esthesioneuroblastoma: The French very rare tumors group (Fracture) contribution. Pediatric Blood and Cancer, 2020, 67, e28154.	1.5	11
74	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
75	Disorder of sex development with germ cell tumors: Which is uncovered first?. Pediatric Blood and Cancer, 2020, 67, e28169.	1.5	6
76	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
77	Multicystic and diffuse malignant peritoneal mesothelioma in children. Pediatric Blood and Cancer, 2020, 67, e28286.	1.5	11
78	French Sarcoma Group proposals for management of sarcoma patients during the COVID-19 outbreak. Annals of Oncology, 2020, 31, 965-966.	1.2	24
79	Pheochromocytoma and Paraganglioma in Children and Adolescents: Experience of the French Society of Pediatric Oncology (SFCE). Journal of the Endocrine Society, 2020, 4, bvaa039.	0.2	21
80	New Born and Infant Soft Tissue Sarcomas. , 2020, , 145-164.		0
81	Update in pediatric nasopharyngeal undifferentiated carcinoma. British Journal of Radiology, 2019, 92, 20190107.	2.2	17
82	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
83	Fine-Needle Aspiration Features of BCOR-CCNB3 Sarcoma. American Journal of Clinical Pathology, 2019, 153, 315-321.	0.7	4
84	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
85	Reply to â€œPathological prognostication of pediatric adrenocortical tumors: Is a gold standard emerging?â€. Pediatric Blood and Cancer, 2019, 66, e27710.	1.5	5
86	Roboticâ€assisted laparoscopic management of renal tumors in children: Preliminary results. Pediatric Blood and Cancer, 2019, 66, e27867.	1.5	36
87	Desmoid-type fibromatosis of the head and neck in children: A changing situation. International Journal of Pediatric Otorhinolaryngology, 2019, 123, 33-37.	1.0	10
88	Can we cure patients with abdominal Desmoplastic Small Round Cell Tumor? Results of a retrospective multicentric study on 100 patients. Surgical Oncology, 2019, 29, 107-112.	1.6	46
89	Standard treatment and emerging drugs for managing synovial sarcoma: adultâ€™s and pediatric oncologist perspective. Expert Opinion on Emerging Drugs, 2019, 24, 43-53.	2.4	9
90	DICER1 and Associated Conditions: Identification of At-risk Individuals and Recommended Surveillance Strategiesâ€"Response. Clinical Cancer Research, 2019, 25, 1689-1690.	7.0	8

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91	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
92	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
93	Adapted strategy to tumor response in childhood nasopharyngeal carcinoma: the French experience. Strahlentherapie Und Onkologie, 2019, 195, 504-516.	2.0	20
94	Outcome of localized liverâ€bile duct rhabdomyosarcoma according to local therapy: A report from the European Paediatric Softâ€Tissue Sarcoma Study Group (EpSSG)â€RMS 2005 study. Pediatric Blood and Cancer, 2019, 66, e27725.	1.5	11
95	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
96	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, e27641.	1.5	31
97	Novel KHDRBS1-NTRK3 rearrangement in a congenital pediatric CD34-positive skin tumor: a case report. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2019, 474, 111-115.	2.8	15
98	Revisiting the role of the pathological grading in pediatric adrenal cortical tumors: results from a national cohort study with pathological review. Modern Pathology, 2019, 32, 546-559.	5.5	38
99	Evidence of hydroxyurea activity in children with pretreated desmoidâ€type fibromatosis: A new option in the armamentarium of systemic therapies. Pediatric Blood and Cancer, 2019, 66, e27472.	1.5	14
100	High Rates of Prescribing Antimicrobials for Prophylaxis in Children and Neonates: Results From the Antibiotic Resistance and Prescribing in European Children Point Prevalence Survey. Journal of the Pediatric Infectious Diseases Society, 2019, 8, 143-151.	1.3	33
101	Larotrectinib efficacy and safety in pediatric TRK fusion cancer patients.. Journal of Clinical Oncology, 2019, 37, 10010-10010.	1.6	14
102	Is alpha-fetoprotein decline a prognostic factor of childhood non-seminomatous germ cell tumours? Results of the French TGM95 study. European Journal of Cancer, 2018, 95, 11-19.	2.8	10
103	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
104	Enrollment in earlyâ€phase clinical trials in pediatric oncology: The experience at Institut Curie. Pediatric Blood and Cancer, 2018, 65, e26916.	1.5	6
105	DICER1 and Associated Conditions: Identification of At-risk Individuals and Recommended Surveillance Strategies. Clinical Cancer Research, 2018, 24, 2251-2261.	7.0	260
106	Pediatric patients with cutaneous melanoma: A European study. Pediatric Blood and Cancer, 2018, 65, e26974.	1.5	26
107	Cytoreductive surgery and hyperthermic intraperitoneal perfusion with chemotherapy in children with peritoneal tumor spread: A French nationwide study over 14 years. Pediatric Blood and Cancer, 2018, 65, e26934.	1.5	14
108	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

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109	Childhood Nasopharyngeal Carcinoma: State-of-the-Art, and Questions for the Future. Journal of Pediatric Hematology/Oncology, 2018, 40, 85-92.	0.6	30
110	Melanotic neuroectodermal tumor of infancy (MNTI) of the head and neck: A French multicenter study. Journal of Cranio-Maxillo-Facial Surgery, 2018, 46, 201-206.	1.7	36
111	Prognostic relevance of early radiologic response to induction chemotherapy in pediatric rhabdomyosarcoma: A report from the International Society of Pediatric Oncology Malignant Mesenchymal Tumor 95 study. Cancer, 2018, 124, 1016-1024.	4.1	25
112	Rhabdomyosarcoma and rhabdomyoma associated with nevoid basal cell carcinoma syndrome: Local treatment strategy. Pediatric Dermatology, 2018, 35, e245-e247.	0.9	5
113	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
114	Addition of dose-intensified doxorubicin to standard chemotherapy for rhabdomyosarcoma (EpSSG) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 5 19, 1061-1071.	10.7	137
115	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
116	The challenge of very rare childhood cancers in developed and developing countries. Expert Opinion on Orphan Drugs, 2017, 5, 331-341.	0.8	12
117	Soft tissue sarcomas in adolescents and young adults: a comparison with their paediatric and adult counterparts. Lancet Oncology, The, 2017, 18, e166-e175.	10.7	100
118	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
119	Pediatric Patient With Renal Cell Carcinoma Treated by Successive Antiangiogenics Drugs: A Case Report and Review of the Literature. Journal of Pediatric Hematology/Oncology, 2017, 39, e279-e284.	0.6	3
120	Primary mediastinal and retroperitoneal malignant germ cell tumors in children and adolescents: Results of the TGM95 trial, a study of the French Society of Pediatric Oncology (SociÃ©tÃ© FranÃ§aise des) Tj ETQq0 0 0 rgBT /Overloc	2.3	33
121	Feasibility and clinical integration of molecular profiling for target identification in pediatric solid tumors. Pediatric Blood and Cancer, 2017, 64, e26365.	1.5	56
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	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
124	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
125	Quiz a tous les Ã©tages. Quel est votre diagnosticÂ?. Revue D'Oncologie HÃ©matologie PÃ©diatrique, 2017, 5, 136-138.	0.1	0
126	Quiz Ã tous les Ã©tages. Quel est votre diagnosticÂ?. Revue D'Oncologie HÃ©matologie PÃ©diatrique, 2017, 5, 139-142.	0.1	0

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127	Congenital Infantile Fibrosarcoma Associated With a Lipofibromatosis-Like Component: One Train May Be Hiding Another. American Journal of Dermatopathology, 2017, 39, 463-467.	0.6	5
128	NUT carcinoma in children and adults: A multicenter retrospective study. Pediatric Blood and Cancer, 2017, 64, e26693.	1.5	65
129	Care management for foreign children, adolescents, young adults with cancer, and their families. Pediatric Blood and Cancer, 2017, 64, e26336.	1.5	2
130	Nonparameningeal head and neck rhabdomyosarcoma in children and adolescents: Lessons from the consecutive International Society of Pediatric Oncology Malignant Mesenchymal Tumor studies. Head and Neck, 2017, 39, 24-31.	2.0	20
131	Pediatric salivary gland carcinomas: Diagnostic and therapeutic management. Laryngoscope, 2017, 127, 140-147.	2.0	36
132	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
133	Synovial sarcoma presenting as colonic intussusception in a child. Pediatric Blood and Cancer, 2017, 64, 207-208.	1.5	9
134	Abdominal desmoplastic small round cell tumor without extraperitoneal metastases: Is there a benefit for HIPEC after macroscopically complete cytoreductive surgery?. PLoS ONE, 2017, 12, e0171639.	2.5	45
135	Two Tumors in 1: What Should be the Therapeutic Target? Pediatric Germ Cell Tumor With Somatic Malignant Transformation. Journal of Pediatric Hematology/Oncology, 2017, 39, 388-394.	0.6	13
136	Complete and Repeated Response of a Metastatic ALK-rearranged Inflammatory Myofibroblastic Tumor to Crizotinib in a Teenage Girl. Journal of Pediatric Hematology/Oncology, 2016, 38, 308-311.	0.6	32
137	Pleuropneumoblastome, tumeur de Sertoli-Leydig et autres maladies associÃ©es au spectre des mutations de DICER1. Revue D'Oncologie HÃ©matologie PÃ©diatrique, 2016, 4, 226-236.	0.1	0
138	Outcome of extracranial malignant rhabdoid tumours in children registered in the European Paediatric Soft Tissue Sarcoma Study Group Non-Rhabdomyosarcoma Soft Tissue Sarcoma 2005 Studyâ€"EpSSG NRSTS 2005. European Journal of Cancer, 2016, 60, 69-82.	2.8	63
139	Combined in situ hypothermic liver preservation and cardioplegia for resection of hepatoblastoma with intra-atrial extension in a 3 year old child. Journal of Pediatric Surgery Case Reports, 2016, 12, 44-49.	0.2	2
140	Cutaneous malignant melanoma in children and adolescents treated in pediatric oncology units. Pediatric Blood and Cancer, 2016, 63, 1922-1927.	1.5	21
141	PET/CT management in a pediatric oncology center. Medecine Nucleaire, 2016, 40, 341-348.	0.2	0
142	A Dilated Cardiomyopathy Revealing a Neuroblastoma: Which Link?. Journal of Pediatric Hematology/Oncology, 2016, 38, e251-e253.	0.6	5
143	Le rÃ©seau dâ€™ÃŽle-de-France dâ€™hÃ©matologie et dâ€™oncologie pÃ©diatrique (RIFHOP)Â: une structure au service des enfants et des professionnels de santÃ©. Revue D'Oncologie HÃ©matologie PÃ©diatrique, 2016, 4, 54-64.	0.1	4
144	Role of Adjuvant Radiation Therapy After Surgery for Abdominal Desmoplastic Small Round Cell Tumors. International Journal of Radiation Oncology Biology Physics, 2016, 95, 1244-1253.	0.8	24

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145	Surgical Management of Neuroendocrine Tumors of the Appendix in Children and Adolescents: A Retrospective French Multicenter Study of 114 Cases. Pediatric Blood and Cancer, 2016, 63, 598-603.	1.5	47
146	Recurrence of Solid Pseudopapillary Neoplasms of the Pancreas: Results of a Nationwide Study of Risk Factors and Treatment Modalities. Pediatric Blood and Cancer, 2016, 63, 1515-1521.	1.5	44
147	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
148	International randomized phase 2 study on the addition of docetaxel to the combination of cisplatin and 5-fluorouracil in the induction treatment for nasopharyngeal carcinoma in children and adolescents. Cancer Chemotherapy and Pharmacology, 2016, 77, 289-298.	2.3	57
149	Sex-Cord Stromal Tumors in Children and Teenagers: Results of the TGM-95 Study. Pediatric Blood and Cancer, 2015, 62, 2114-2119.	1.5	38
150	Rhabdomyosarcomas in children with neurofibromatosis type I: A national historical cohort. Pediatric Blood and Cancer, 2015, 62, 1733-1738.	1.5	55
151	Relapse after localized rhabdomyosarcoma: Evaluation of the efficacy of secondâ€line chemotherapy. Pediatric Blood and Cancer, 2015, 62, 1935-1941.	1.5	37
152	Considering chemotherapy in synovial sarcoma. Expert Opinion on Orphan Drugs, 2015, 3, 1111-1124.	0.8	4
153	Does aggressive local treatment have an impact on survival in children with metastatic rhabdomyosarcoma?. European Journal of Cancer, 2015, 51, 193-201.	2.8	27
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