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List of Publications by Year
in descending order

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46 papers	1,796 citations	471509 17 h-index	276875 41 g-index
49 all docs	49 docs citations	49 times ranked	2287 citing authors

#	Article	IF	Citations
1	Ewing Sarcoma: Current Management and Future Approaches Through Collaboration. Journal of Clinical Oncology, 2015, 33, 3036-3046.	1.6	516
2	Results of the EICESS-92 Study: Two Randomized Trials of Ewing's Sarcoma Treatmentâ€"Cyclophosphamide Compared With Ifosfamide in Standard-Risk Patients and Assessment of Benefit of Etoposide Added to Standard Treatment in High-Risk Patients. Journal of Clinical Oncology, 2008, 26, 4385-4393.	1.6	236
3	Cyclophosphamide Compared With Ifosfamide in Consolidation Treatment of Standard-Risk Ewing Sarcoma: Results of the Randomized Noninferiority Euro-EWING99-R1 Trial. Journal of Clinical Oncology, 2014, 32, 2440-2448.	1.6	136
4	High-Dose Chemotherapy and Blood Autologous Stem-Cell Rescue Compared With Standard Chemotherapy in Localized High-Risk Ewing Sarcoma: Results of Euro-E.W.I.N.G.99 and Ewing-2008. Journal of Clinical Oncology, 2018, 36, 3110-3119.	1.6	107
5	Initial Patient Characteristics Can Predict Pattern and Risk of Relapse in Localized Rhabdomyosarcoma. Journal of Clinical Oncology, 2008, 26, 406-413.	1.6	101
6	Recurrent intragenic rearrangements of EGFR and BRAF in soft tissue tumors of infants. Nature Communications, 2018, 9, 2378.	12.8	72
7	Kaposiform hemangioendothelioma in children: a benign vascular tumor with multiple treatment options. World Journal of Pediatrics, 2018, 14, 322-329.	1.8	53
8	Desmoplastic small round cell tumors: Multimodality treatment and new risk factors. Cancer Medicine, 2019, 8, 527-542.	2.8	39
9	Treatment for soft tissue sarcoma in childhood and adolescence. Wiener Klinische Wochenschrift, 2005, 117, 196-209.	1.9	38
10	Primary Metastatic Synovial Sarcoma: Experience of the CWS Study Group. Pediatric Blood and Cancer, 2016, 63, 1198-1206.	1.5	37
11	Epithelioid hemangioendotheliomas of the liver and lung in children and adolescents. Pediatric Blood and Cancer, 2017, 64, e26675.	1.5	31
12	Synovial Sarcoma Recurrence in Children and Young Adults. Annals of Surgical Oncology, 2016, 23, 618-626.	1.5	29
13	The prognostic impact of SYTâ€SSX fusion type and histological grade in pediatric patients with synovial sarcoma treated according to the CWS (Cooperative Weichteilsarkom Studie) trials. Pediatric Blood and Cancer, 2017, 64, 89-95.	1.5	29
14	Tumour volume reduction after neoadjuvant chemotherapy impacts outcome in localised embryonal rhabdomyosarcoma. Pediatric Blood and Cancer, 2015, 62, 16-23.	1.5	26
15	The impact of local control in the treatment of type II/III pleuropulmonary blastoma. Experience of the Cooperative Weichteilsarkom Studiengruppe (CWS). Journal of Surgical Oncology, 2017, 115, 164-172.	1.7	21
16	The prognostic value of early radiographic response in children and adolescents with embryonal rhabdomyosarcoma stage IV, metastases confined to the lungs: A report from the Cooperative Weichteilsarkom Studiengruppe (CWS). Pediatric Blood and Cancer, 2017, 64, e26510.	1.5	19
17	Importance of whole-body imaging with complete coverage of hands and feet in alveolar rhabdomyosarcoma staging. Pediatric Radiology, 2018, 48, 648-657.	2.0	19
18	Systemic therapy of aggressive fibromatosis in children and adolescents: Report of the Cooperative Weichteilsarkom Studiengruppe (CWS). Pediatric Blood and Cancer, 2018, 65, e26943.	1.5	19

#	Article	IF	Citations
19	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
20	Local treatment of rhabdomyosarcoma of the female genital tract: Expert consensus from the Children's Oncology Group, the European Softâ€Tissue Sarcoma Group, and the Cooperative Weichteilsarkom Studiengruppe. Pediatric Blood and Cancer, 2023, 70, e28601.	1.5	18
21	RhabdomyosarcomaÂdiagnosed in the first year of life: Localized, metastatic, and relapsed disease. Outcome data from five trials and one registry of the Cooperative Weichteilsarkom Studiengruppe (CWS). Pediatric Blood and Cancer, 2019, 66, e27652.	1.5	17
22	Alveolar softâ€part sarcoma: Primary metastatic disease and metastatic relapse occurring during longâ€term followâ€up. Pediatric Blood and Cancer, 2018, 65, e27405.	1.5	16
23	The impact of local control in the treatment of children with advanced infantile and adult-type fibrosarcoma: Experience of the cooperative weichteilsarkom studiengruppe (CWS). Journal of Pediatric Surgery, 2020, 55, 1740-1747.	1.6	16
24	Treatment and outcome of the patients with rhabdomyosarcoma of the biliary tree: Experience of the Cooperative Weichteilsarkom Studiengruppe (CWS). BMC Cancer, 2019, 19, 945.	2.6	13
25	Epithelioid sarcoma in children, adolescents, and young adults: Localized, primary metastatic and relapsed disease. Treatment results of five Cooperative Weichteilsarkom Studiengruppe (CWS) trials and one registry. Pediatric Blood and Cancer, 2019, 66, e27879.	1.5	13
26	Genetic testing and surveillance in infantile myofibromatosis: a report from the SIOPE Host Genome Working Group. Familial Cancer, 2021, 20, 327-336.	1.9	13
27	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
28	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
29	Extraskeletal Ewing sarcoma in children, adolescents, and young adults. An analysis of three prospective studies of the Cooperative Weichteilsarkomstudiengruppe (CWS). Pediatric Blood and Cancer, 2021, 68, e29145.	1.5	11
30	Longâ€term 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
31	Localized synovial sarcoma of the foot or ankle: A series of 32 Cooperative Weichteilsarkom Study Group patients. Journal of Surgical Oncology, 2019, 119, 109-119.	1.7	10
32	Synovial sarcoma disease characteristics and primary tumor sites differ between patient age groups: a report of the Cooperative Weichteilsarkom Studiengruppe (CWS). Journal of Cancer Research and Clinical Oncology, 2020, 146, 953-960.	2.5	10
33	Fusion transcripts as liquid biopsy markers in alveolar rhabdomyosarcoma and synovial sarcoma: A report of the Cooperative Weichteilsarkom Studiengruppe (CWS). Pediatric Blood and Cancer, 2022, , e29652.	1.5	10
34	Rhabdomyosarcoma of the female genitourinary tract: Primary and relapsed disease in infants and older children. Treatment results of five Cooperative Weichteilsarkom Studiengruppe (CWS) trials and one registry. Pediatric Blood and Cancer, 2021, 68, e28889.	1.5	9
35	Treatment and outcome of patients with thoracic tumors of the Ewing sarcoma family: A report from the Cooperative Weichteilsarkom Studiengruppe CWSâ€81, â€86, â€91, â€96, and â€2002P trials. Pediatric Blood and Cancer, 2019, 66, e27884.	1.5	8
36	Lowâ€grade fibromyxoid sarcoma: A report of the Cooperative Weichteilsarkom Studiengruppe (CWS). Pediatric Blood and Cancer, 2020, 67, e28009.	1.5	8

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37	Congenital spindle cell rhabdomyosarcoma: An international cooperative analysis. European Journal of Cancer, 2022, 168, 56-64.	2.8	8
38	Dermatofibrosarcoma protuberans in children and adolescents: Primary and Relapsed diseaseâ€"Experience of the Cooperative Weichteilsarkomstudiengruppe (CWS). Journal of Surgical Oncology, 2020, 122, 263-272.	1.7	6
39	Treatment and outcome of patients with thoracic tumors of the Ewing sarcoma family: A report from the Cooperative Weichteilsarkom Studiengruppe CWSâ€81, â€86, â€91, â€96, and â€2002P trials. Pediatric Blood and Cancer, 2019, 66, e27537.	1.5	5
40	Endothelial cell malignancies in infants, children and adolescents: Treatment results of three Cooperative Weichteilsarkom Studiengruppe (CWS) trials and one registry. Pediatric Blood and Cancer, 2020, 67, e28095.	1.5	5
41	Patterns of Prior and Subsequent Neoplasms in Children and Adolescents With Soft Tissue Sarcomas. Journal of Pediatric Hematology/Oncology, 2020, 42, e265-e270.	0.6	5
42	Infantile myofibromatosis: Excellent prognosis but also rare fatal progressive disease. Treatment results of five Cooperative Weichteilsarkom Studiengruppe (CWS) trials and one registry. Pediatric Blood and Cancer, 2021, , e29403.	1.5	5
43	Children with progressive and relapsed pleuropulmonary blastoma: A European collaborative analysis. Pediatric Blood and Cancer, 2021, 68, e29268.	1.5	4
44	Metronomic oral maintenance chemotherapy in patients with localized high-risk rhabdomyosarcoma (RMS) and RMS-like tumors: A report from a randomized, multicenter, phase III trial CWS-2007HR.. Journal of Clinical Oncology, 2022, 40, 10033-10033.	1.6	2
45	What is the best therapy for grossly resected synovial sarcoma? Experience of the CWS Study Group.. Journal of Clinical Oncology, 2019, 37, 10042-10042.	1.6	0
46	Pre-operative radiotherapy is associated with superior local relapse-free survival in advanced synovial sarcoma. Journal of Cancer Research and Clinical Oncology, 0, , .	2.5	0