Ewa Koscielniak

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3122969/publications.pdf

Version: 2024-02-01

69 papers 2,149 citations

20 h-index 254184 43 g-index

70 all docs

70 docs citations

times ranked

70

2262 citing authors

#	Article	IF	CITATIONS
1	Results of Treatment for Soft Tissue Sarcoma in Childhood and Adolescence: A Final Report of the German Cooperative Soft Tissue Sarcoma Study CWS-86. Journal of Clinical Oncology, 1999, 17, 3706-3719.	1.6	250
2	Cooperative Trial CWS-91 for Localized Soft Tissue Sarcoma in Children, Adolescents, and Young Adults. Journal of Clinical Oncology, 2009, 27, 1446-1455.	1.6	163
3	Rituximab for High-Risk, Mature B-Cell Non-Hodgkin's Lymphoma in Children. New England Journal of Medicine, 2020, 382, 2207-2219.	27.0	157
4	Treatment of soft tissue sarcoma in childhood and adolescence: A report of the german cooperative soft tissue sarcoma study. Cancer, 1992, 70, 2557-2567.	4.1	132
5	Treatment of children with metastatic soft tissue sarcoma with oral maintenance compared to high dose chemotherapy: Report of the HD CWSâ€96 trial. Pediatric Blood and Cancer, 2008, 50, 739-745.	1.5	115
6	Initial Patient Characteristics Can Predict Pattern and Risk of Relapse in Localized Rhabdomyosarcoma. Journal of Clinical Oncology, 2008, 26, 406-413.	1.6	101
7	Soft Tissue Sarcoma in Children. Paediatric Drugs, 2002, 4, 21-28.	3.1	87
8	Recurrent intragenic rearrangements of EGFR and BRAF in soft tissue tumors of infants. Nature Communications, 2018, 9, 2378.	12.8	72
9	Prognostic value of PAX–FKHR fusion status in alveolar rhabdomyosarcoma: A report from the cooperative soft tissue sarcoma study group (CWS). Pediatric Blood and Cancer, 2011, 57, 406-414.	1.5	61
10	Treatment of children with relapsed soft tissue sarcoma: Report of the German CESS/CWS REZ 91 Trial. , 1998, 30, 269-275.		60
11	Kaposiform hemangioendothelioma in children: a benign vascular tumor with multiple treatment options. World Journal of Pediatrics, 2018, 14, 322-329.	1.8	53
12	Fifty years of rhabdomyosarcoma studies on both sides of the pond and lessons learned. Cancer Treatment Reviews, 2018, 68, 94-101.	7.7	48
13	Heme Oxygenase-1 Controls an HDAC4-miR-206 Pathway of Oxidative Stress in Rhabdomyosarcoma. Cancer Research, 2016, 76, 5707-5718.	0.9	46
14	Biallelic loss of <i>CDKN2A </i> is associated with poor response to treatment in pediatric acute lymphoblastic leukemia. Leukemia and Lymphoma, 2017, 58, 1162-1171.	1.3	43
15	Desmoplastic small round cell tumors: Multimodality treatment and new risk factors. Cancer Medicine, 2019, 8, 527-542.	2.8	39
16	Primary Metastatic Synovial Sarcoma: Experience of the CWS Study Group. Pediatric Blood and Cancer, 2016, 63, 1198-1206.	1.5	37
17	Does the timeâ€point of relapse influence outcome in pediatric rhabdomyosarcomas?. Pediatric Blood and Cancer, 2009, 52, 772-776.	1.5	35
18	Epithelioid hemangioendotheliomas of the liver and lung in children and adolescents. Pediatric Blood and Cancer, 2017, 64, e26675.	1.5	31

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19	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
20	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
21	Impact of Hemiscrotectomy on Outcome of Patients with Embryonal Paratesticular Rhabdomyosarcoma: Results from the Cooperative Soft Tissue Sarcoma Group Studies CWS-86, 91, 96 and 2002P. Journal of Urology, 2014, 192, 902-907.	0.4	23
22	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
23	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
24	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
25	Oral lowâ€dose chemotherapy: Successful treatment of an alveolar rhabdomyosarcoma during pregnancy. Pediatric Blood and Cancer, 2012, 58, 104-106.	1.5	19
26	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
27	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
28	COVID-19 in pediatric cancer patients is associated with treatment interruptions but not with short-term mortality: a Polish national study. Journal of Hematology and Oncology, 2021, 14, 163.	17.0	19
29	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
30	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
31	Dose-Adjusted Etoposide, Doxorubicin, and Cyclophosphamide With Vincristine and Prednisone Plus Rituximab Therapy in Children and Adolescents With Primary Mediastinal B-Cell Lymphoma: A Multicenter Phase II Trial. Journal of Clinical Oncology, 2021, 39, 3716-3724.	1.6	18
32	Pediatric Cancer Data Commons: Federating and Democratizing Data for Childhood Cancer Research. JCO Clinical Cancer Informatics, 2021, 5, 1034-1043.	2.1	18
33	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
34	Treatment and outcome of patients with localized intrathoracic and chest wall rhabdomyosarcoma: a report of the Cooperative Weichteilsarkom Studiengruppe (CWS). Journal of Cancer Research and Clinical Oncology, 2018, 144, 925-934.	2.5	16
35	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
36	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

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37	Pleuropulmonary blastoma in children and adolescents: The EXPERT/PARTNER diagnostic and therapeutic recommendations. Pediatric Blood and Cancer, 2021, 68, e29045.	1.5	15
38	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
39	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
40	Tumour expressions of hypoxic markers predict the response to neo-adjuvant chemotherapy in children with inoperable rhabdomyosarcoma. Biomarkers, 2019, 24, 538-548.	1.9	13
41	Outcome, Treatment, and Treatment Failures in Patients Suffering Localized Embryonal Paratesticular Rhabdomyosarcoma. Annals of Surgery, 2016, 264, 1148-1155.	4.2	12
42	Tumor expression of survivin, p53, cyclin D1, osteopontin and fibronectin in predicting the response to neo-adjuvant chemotherapy in children with advanced malignant peripheral nerve sheath tumor. Journal of Cancer Research and Clinical Oncology, 2018, 144, 519-529.	2.5	12
43	Analysis of the rRNA methylation complex components in pediatric B-cell precursor acute lymphoblastic leukemia: A pilot study. Advances in Clinical and Experimental Medicine, 2020, 29, 107-113.	1.4	12
44	Clinical features and treatment outcomes of peripheral T-cell lymphoma in children. A current data report from Polish Pediatric Leukemia/Lymphoma Study Group (PPLLSG). Advances in Medical Sciences, 2016, 61, 311-316.	2.1	11
45	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
46	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
47	Surface Expression of CRLF2 Protein Is Associated with Lower Minimal Residual Disease (MRD) Among Children with IKZF1-deleted Acute Lymphoblastic Leukemia (ALL). Blood, 2014, 124, 2400-2400.	1.4	10
48	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
49	<i>GATA3</i> germline variant is associated with <i>CRLF2</i> expression and predicts outcome in pediatric Bâ€cell precursor acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2019, 58, 619-626.	2.8	9
50	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
51	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 Bloo and Cancer, 2019, 66, e27884.	d1.5	8
52	The influence of different intensity of treatment on hormonal markers of gonadal function in acute lymphoblastic leukemia survivors. Hematological Oncology, 2019, 37, 609-616.	1.7	8
53	Multicolor flow cytometry immunophenotyping and characterization of aneuploidy in pediatric B-cell precursor acute lymphoblastic leukemia. Central-European Journal of Immunology, 2021, 46, 365-374.	1.2	8
54	Prognostic significance of <i>IKZF1</i> deletions and IKZF1 ^{plus} profile in children with Bâ€cell precursor acute lymphoblastic leukemia treated according to the ALLâ€IC BFM 2009 protocol. Hematological Oncology, 2022, 40, 430-441.	1.7	8

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55	Secondary malignant neoplasms after bone and soft tissue sarcomas in children, adolescents, and young adults. Cancer, 2022, 128, 1787-1800.	4.1	8
56	Congenital spindle cell rhabdomyosarcoma: An international cooperative analysis. European Journal of Cancer, 2022, 168, 56-64.	2.8	8
57	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
58	Nutritional status at the moment of diagnosis in childhood cancer patients. Pediatric Endocrinology, Diabetes and Metabolism, 2017, 23, 77-82.	0.7	6
59	Paratesticular alveolar rhabdomyosarcomas do not harbor typical translocations: a distinct entity with favorable prognosis?. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2018, 472, 441-449.	2.8	5
60	Heterozygous carriers of germline c.657_661del5 founder mutation in <i>NBN</i> gene are at risk of central nervous system relapse of B-cell precursor acute lymphoblastic leukemia. Haematologica, 2018, 103, e200-e203.	3.5	5
61	Treatment and outcome of patients with thoracic tumors of the Ewing sarcoma family: A report from the Cooperative Weichteilsarkom Studiengruppe CWSâ \in 81, â \in 86, â \in 91, â \in 96, and â \in 2002P trials. Pediatric Bloo and Cancer, 2019, 66, e27537.	d1.5	5
62	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
63	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
64	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
65	Chronic metastatic neuroblastoma. Pediatric Blood and Cancer, 2008, 50, 898-900.	1.5	4
66	Children with progressive and relapsed pleuropulmonary blastoma: A European collaborative analysis. Pediatric Blood and Cancer, 2021, 68, e29268.	1.5	4
67	Metabolic Disturbances in Children Treated for Solid Tumors. Nutrients, 2019, 11, 3062.	4.1	2
68	Analysis of incidence and risk factors of the multidrug resistant gastrointestinal tract infection in children and adolescents undergoing allogeneic and autologous hematopoietic cell transplantation: a nationwide study. Annals of Hematology, 2021, 101, 191.	1.8	1
69	HGG-11. Clinical characteristics and clinical evolution of a large cohort of pediatric patients with primary central nervous system (CNS) tumors and tropomyosin receptor kinase (TRK) fusion Neuro-Oncology, 2022, 24, i61-i62.	1.2	0