

Dominik T Schneider

List of Publications by Year in descending order

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Version: 2024-02-01

105
papers

4,609
citations

109321

35
h-index

102487

66
g-index

122
all docs

122
docs citations

122
times ranked

4951
citing authors

#	ARTICLE	IF	CITATIONS
1	MicroRNA-profiling of miR-371~373- and miR-302/367-clusters in serum and cerebrospinal fluid identify patients with intracranial germ cell tumors. <i>Journal of Cancer Research and Clinical Oncology</i> , 2023, 149, 791-802.	2.5	9
2	Rare Tumors in Children and Adolescents – the STEP Working Group’s Evolution to a Prospective Registry. <i>Klinische Padiatrie</i> , 2022, 234, 146-153.	0.6	1
3	Diagnostic and prognostic classification of atypical spitzoid tumours based on histology and genomic aberrations: A prospective cohort study with long-term follow-up. <i>European Journal of Cancer</i> , 2022, 163, 200-210.	2.8	1
4	Multimodal Treatment of Nasopharyngeal Carcinoma in Children, Adolescents and Young Adults-Extended Follow-Up of the NPC-2003-GPOH Study Cohort and Patients of the Interim Cohort. <i>Cancers</i> , 2022, 14, 1261.	3.7	9
5	Gonadal and Extragenital Germ Cell Tumors, Sex Cord Stromal and Rare Gonadal Tumors. <i>Pediatric Oncology</i> , 2022, , 301-389.	0.5	1
6	Facharzt-Training PÄdiatrie – Vorbereitungskurs zur Facharztprüfung Kinder- und Jugendmedizin. <i>Monatsschrift Fur Kinderheilkunde</i> , 2022, 170, 33-35.	0.1	0
7	Rare pediatric tumors in Germany – Not as rare as expected: a study based on data from the Bavarian Cancer Registry and the German Childhood Cancer Registry. <i>European Journal of Pediatrics</i> , 2022, , 1.	2.7	0
8	Incidences and characteristics of primary lung malignancies in childhood in Germany: An analysis of population-based data from German cancer registries. <i>Pediatric Blood and Cancer</i> , 2022, 69, e29744.	1.5	3
9	Seltene Tumoren. <i>Springer Reference Medizin</i> , 2021, , 1-8.	0.0	0
10	Consensus recommendations from the EXPeRT/PARTNER groups for the diagnosis and therapy of sex cord stromal tumors in children and adolescents. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29017.	1.5	13
11	Thymoma and thymic carcinoma in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29042.	1.5	5
12	Phasic and tonic alertness in preterm 5-year-old healthy children. <i>Child Neuropsychology</i> , 2021, 27, 1073-1087.	1.3	1
13	Nasopharyngeal carcinoma in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29018.	1.5	11
14	The European Paediatric Rare Tumours Network –European Registry (PARTNER) project for very rare tumors in children. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29072.	1.5	11
15	Pleuropulmonary blastoma in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29045.	1.5	15
16	Initial presenting manifestations in 16,486 patients with inborn errors of immunity include infections and noninfectious manifestations. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 148, 1332-1341.e5.	2.9	75
17	Salivary gland carcinoma in children and adolescents: The EXPeRT/PARTNER diagnosis and treatment recommendations. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29058.	1.5	7
18	Pancreatoblastoma in children: EXPeRT/PARTNER diagnostic and therapeutic recommendations. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29112.	1.5	9

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19	Cutaneous melanoma in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. <i>Pediatric Blood and Cancer</i> , 2021, 68, e28992.	1.5	9
20	Adrenocortical tumours in children and adolescents: The EXPeRT/PARTNER diagnostic and therapeutic recommendations. <i>Pediatric Blood and Cancer</i> , 2021, 68, e29025.	1.5	16
21	Facing the challenges of very rare tumors of pediatric age: The European Cooperative Study Group for Pediatric Rare Tumors (EXPeRT) background, goals, and achievements. <i>Pediatric Blood and Cancer</i> , 2021, 68, e28993.	1.5	10
22	The Pediatric Precision Oncology INFORM Registry: Clinical Outcome and Benefit for Patients with Very High-Evidence Targets. <i>Cancer Discovery</i> , 2021, 11, 2764-2779.	9.4	110
23	â€žFacharzt-Training PÄdiatrie â€“AVorbereitungskurs zur FacharztprÄ¼fung Kinder- und Jugendmedizinâ€œ. <i>Monatsschrift Fur Kinderheilkunde</i> , 2021, 169, 205-206.	0.1	0
24	Primary lung carcinoma in children and adolescents â€“ Clinical characteristics and outcome of 12 cases from the German registry for rare paediatric tumours (STEP). <i>Lung Cancer</i> , 2021, 160, 66-72.	2.0	5
25	Treating rare tumors with the assistance of the expert virtual consultation system: two cases of juvenile granulosa cell tumors. <i>Tumori</i> , 2021, 107, NP141-NP143.	1.1	1
26	Facharzt-Training PÄdiatrieâ€“ Vorbereitungskurs zur FacharztprÄ¼fung Kinder- und Jugendmedizin. <i>Monatsschrift Fur Kinderheilkunde</i> , 2021, 169, 305-306.	0.1	0
27	Genotyping circulating tumor DNA of pediatric Hodgkin lymphoma. <i>Leukemia</i> , 2020, 34, 151-166.	7.2	53
28	Mesothelioma in children and adolescents: the European Cooperative Study Group for Pediatric Rare Tumors (EXPeRT) contribution. <i>European Journal of Cancer</i> , 2020, 140, 63-70.	2.8	12
29	Age-Dependent Presentation and Clinical Course of 1465 Patients Aged 0 to Less than 18 Years with Ovarian or Testicular Germ Cell Tumors; Data of the MAKEI 96 Protocol Revisited in the Light of Prenatal Germ Cell Biology. <i>Cancers</i> , 2020, 12, 611.	3.7	23
30	ESGOâ€“SIOPE guidelines for the management of adolescents and young adults with non-epithelial ovarian cancers. <i>Lancet Oncology</i> , The, 2020, 21, e360-e368.	10.7	50
31	Pediatric Multisystemic Inflammatory Syndrome Associated With SARS-CoV-2 Infection. <i>Deutsches A&#x0308;rzteblatt International</i> , 2020, 117, 431.	0.9	2
32	The German National Registry of Primary Immunodeficiencies (2012â€“2017). <i>Frontiers in Immunology</i> , 2019, 10, 1272.	4.8	71
33	DICER1 and Associated Conditions: Identification of At-risk Individuals and Recommended Surveillance Strategiesâ€“Response. <i>Clinical Cancer Research</i> , 2019, 25, 1689-1690.	7.0	8
34	Defining and listing very rare cancers of paediatric age: consensus of the Joint Action on Rare Cancersâ€“in cooperation with the European Cooperative Study Group for Pediatric Rare Tumors. <i>European Journal of Cancer</i> , 2019, 110, 120-126.	2.8	61
35	Reduction of the event-related potential P3 in preterm born 5-year-old healthy children. <i>Clinical Neurophysiology</i> , 2019, 130, 675-682.	1.5	1
36	Specialized pediatric palliative care services for children dying from cancer: A repeated cohort study on the developments of symptom management and quality of care over a 10-year period. <i>Palliative Medicine</i> , 2019, 33, 381-391.	3.1	31

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37	Joining forces for pediatric very rare tumors. <i>Oncotarget</i> , 2019, 10, 3084-3085.	1.8	5
38	Keimzelltumoren und seltene gonadale Tumoren bei Kindern und Jugendlichen. <i>Springer Reference Medizin</i> , 2019, , 1-8.	0.0	0
39	<i>DICER1</i> and Associated Conditions: Identification of At-risk Individuals and Recommended Surveillance Strategies. <i>Clinical Cancer Research</i> , 2018, 24, 2251-2261.	7.0	260
40	Pediatric patients with cutaneous melanoma: A European study. <i>Pediatric Blood and Cancer</i> , 2018, 65, e26974.	1.5	26
41	Brentuximab vedotin exerts profound antiproliferative and proapoptotic efficacy in CD30-positive as well as cocultured CD30-negative germ cell tumour cell lines. <i>Journal of Cellular and Molecular Medicine</i> , 2018, 22, 568-575.	3.6	17
42	The challenge of very rare childhood cancers in developed and developing countries. <i>Expert Opinion on Orphan Drugs</i> , 2017, 5, 331-341.	0.8	12
43	Clinical characteristics and outcome of 60 pediatric patients with malignant melanoma registered with the German Pediatric Rare Tumor Registry (STEP). <i>Klinische Padiatrie</i> , 2017, 229, 322-328.	0.6	16
44	DICER1-related Sertoli-Leydig cell tumor and gynandroblastoma: Clinical and genetic findings from the International Ovarian and Testicular Stromal Tumor Registry. <i>Gynecologic Oncology</i> , 2017, 147, 521-527.	1.4	87
45	Ovarian Tumors During Childhood and Adolescence. , 2017, , 3328-3334.		0
46	Pediatric Colorectal Carcinoma is Associated With Excellent Outcome in the Context of Cancer Predisposition Syndromes. <i>Pediatric Blood and Cancer</i> , 2016, 63, 611-617.	1.5	22
47	Identification of a Cryptic Insertion ins(11;X)(q23;q28q12) Resulting in a <i>KMT2A</i> and <i>FLNA</i> Fusion in a 13-Month-Old Child with Acute Myelomonocytic Leukemia. <i>Cytogenetic and Genome Research</i> , 2016, 150, 281-286.	1.1	2
48	Ovarian Sex Cord-Stromal Tumors. <i>Journal of Oncology Practice</i> , 2016, 12, 940-946.	2.5	71
49	ATTENTION FUNCTIONING IN CHILDREN WITH PRENATAL DRUG EXPOSURE. <i>Infant Mental Health Journal</i> , 2015, 36, 522-530.	1.8	4
50	Sex Cord Stromal Tumors: It is Networking-or Not Working. <i>Pediatric Blood and Cancer</i> , 2015, 62, 2065-2066.	1.5	2
51	Ovarian Sertoli Leydig cell tumours in children and adolescents: An analysis of the European Cooperative Study Group on Pediatric Rare Tumors (EXPeRT). <i>European Journal of Cancer</i> , 2015, 51, 543-550.	2.8	62
52	443 paediatric cases of malignant melanoma registered with the German Central Malignant Melanoma Registry between 1983 and 2011. <i>European Journal of Cancer</i> , 2015, 51, 861-868.	2.8	45
53	Rare malignant pediatric tumors registered in the German Childhood Cancer Registry 2001-2010. <i>Pediatric Blood and Cancer</i> , 2014, 61, 1202-1209.	1.5	29
54	Ovarian and Testicular Sex Cord-Stromal Tumors. <i>Pediatric Oncology</i> , 2014, , 101-113.	0.5	2

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55	Brain metastases during follow-up of children and adolescents with extracranial malignant germ cell tumors: Risk adapted management decision tree analysis based on data of the MAHO/MAKEI registry. <i>Pediatric Blood and Cancer</i> , 2013, 60, 217-223.	1.5	7
56	Regional deep hyperthermia for salvage treatment of children and adolescents with refractory or recurrent non-testicular malignant germ-cell tumours: an open-label, non-randomised, single-institution, phase 2 study. <i>Lancet Oncology</i> , The, 2013, 14, 843-852.	10.7	72
57	The founding of the European Cooperative Study Group on Pediatric Rare Tumors – EXPeRT. <i>Expert Review of Anticancer Therapy</i> , 2013, 13, 1-3.	2.4	32
58	Salvage treatment of relapsed or refractory germ-cell tumours – Authors' reply. <i>Lancet Oncology</i> , The, 2013, 14, e486-e487.	10.7	0
59	EPCAM – A novel molecular target for the treatment of pediatric and adult germ cell tumors. <i>Genes Chromosomes and Cancer</i> , 2013, 52, 24-32.	2.8	10
60	Testicular sex cord stromal tumors: Analysis of patients from the MAKEI study. <i>Pediatric Blood and Cancer</i> , 2013, 60, 1651-1655.	1.5	32
61	Management of Ovarian and Testicular Sex Cord-stromal Tumors in Children and Adolescents. <i>Journal of Pediatric Hematology/Oncology</i> , 2012, 34, S55-S63.	0.6	34
62	Prevalence of c-KIT Mutations in Gonadoblastoma and Dysgerminomas of Patients with Disorders of Sex Development (DSD) and Ovarian Dysgerminomas. <i>PLoS ONE</i> , 2012, 7, e43952.	2.5	50
63	Rare Tumors: A Different Perspective on Oncology. <i>Pediatric Oncology</i> , 2012, , 3-13.	0.5	5
64	Gonadal and Extragenital Germ Cell Tumors, Sex Cord Stromal and Rare Gonadal Tumors. <i>Pediatric Oncology</i> , 2012, , 327-402.	0.5	18
65	Germ Cell Tumors of the Head and Neck. <i>Pediatric Oncology</i> , 2012, , 169-173.	0.5	0
66	Mediastinal Germ Cell Tumors. <i>Pediatric Oncology</i> , 2012, , 205-211.	0.5	0
67	Gastrointestinal stromal tumours in children and young adults: A clinicopathologic series with long-term follow-up from the database of the Cooperative Weichteilsarkom Studiengruppe (CWS). <i>European Journal of Cancer</i> , 2011, 47, 1692-1698.	2.8	26
68	Analysis of the adenomatous polyposis coli (APC) gene in childhood and adolescent germ cell tumors. <i>Pediatric Blood and Cancer</i> , 2011, 56, 384-391.	1.5	18
69	Ovarian Tumors during Childhood and Adolescence. , 2011, , 2722-2727.		0
70	Clinical spectrum of the pseudotumor cerebri complex in children. <i>Child's Nervous System</i> , 2010, 26, 313-321.	1.1	54
71	Care for Rare Cancers: Improved Care Requires Improved Communication. <i>Klinische Padiatrie</i> , 2010, 222, 124-126.	0.6	16
72	Cisplatin and Etoposide in Childhood Germ Cell Tumor: Brazilian Pediatric Oncology Society Protocol GCT-91. <i>Journal of Clinical Oncology</i> , 2009, 27, 1297-1303.	1.6	39

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73	Germ cell tumors of the head and neck: Report from the MAKEI Study Group. <i>Pediatric Blood and Cancer</i> , 2009, 52, 223-226.	1.5	27
74	Improvement in the outcome of children with germ cell tumors. <i>Pediatric Blood and Cancer</i> , 2008, 50, 250-253.	1.5	16
75	Mesenchymal chondrosarcoma of soft tissues and bone in children, adolescents, and young adults. <i>Cancer</i> , 2008, 112, 2424-2431.	4.1	133
76	Further characterization of the first seminoma cell line TCamâ€2. <i>Genes Chromosomes and Cancer</i> , 2008, 47, 185-196.	2.8	126
77	Microarray analysis of Ewingâ€™s sarcoma family of tumours reveals characteristic gene expression signatures associated with metastasis and resistance to chemotherapy. <i>European Journal of Cancer</i> , 2008, 44, 699-709.	2.8	87
78	Pediatric Malignant Germ Cell Tumors Show Characteristic Transcriptome Profiles. <i>Cancer Research</i> , 2008, 68, 4239-4247.	0.9	83
79	An Immunodeficiency Disease with <i>RAG</i> Mutations and Granulomas. <i>New England Journal of Medicine</i> , 2008, 358, 2030-2038.	27.0	219
80	Genotype and Protein Expression After Bone Marrow Transplantation for Adrenoleukodystrophy. <i>Archives of Neurology</i> , 2007, 64, 651.	4.5	18
81	JKTâ€1 is not a human seminoma cell line. <i>Journal of Developmental and Physical Disabilities</i> , 2007, 30, 350-365.	3.6	20
82	Fatal glioblastoma multiforme in a patient with neurofibromatosis type I: the dilemma of systematic medical follow-up. <i>Child's Nervous System</i> , 2007, 23, 343-347.	1.1	14
83	Constitutive Activation of Neuregulin/ERBB3 Signaling Pathway in Clear Cell Sarcoma of Soft Tissue. <i>Neoplasia</i> , 2006, 8, 613-622.	5.3	26
84	Molecular genetic analysis of central nervous system germ cell tumors with comparative genomic hybridization. <i>Modern Pathology</i> , 2006, 19, 864-873.	5.5	83
85	Imbalances of chromosome arm 1p in pediatric and adult germ cell tumors are caused by true allelic loss: A combined comparative genomic hybridization and microsatellite analysis. <i>Genes Chromosomes and Cancer</i> , 2006, 45, 995-1006.	2.8	28
86	Ovarian small cell carcinoma of the hypercalcemic type in children and adolescents. <i>Cancer</i> , 2006, 107, 2298-2306.	4.1	70
87	Activation of Wnt/ β -Catenin Signaling in Distinct Histologic Subtypes of Human Germ Cell Tumors. <i>Pediatric and Developmental Pathology</i> , 2006, 9, 115-131.	1.0	47
88	Genomic and Expression Profiling of Human Spermatocytic Seminomas: Primary Spermatocyte as Tumorigenic Precursor and DMRT1 as Candidate Chromosome 9 Gene. <i>Cancer Research</i> , 2006, 66, 290-302.	0.9	208
89	Keimzelltumoren. , 2006, , 922-938.		1
90	IGF2/H19 imprinting analysis of human germ cell tumors (GCTs) using the methylation-sensitive single-nucleotide primer extension method reflects the origin of GCTs in different stages of primordial germ cell development. <i>Genes Chromosomes and Cancer</i> , 2005, 44, 256-264.	2.8	85

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91	Ovarian sex cord-stromal tumors in children and adolescents. Journal of reproductive medicine, The, 2005, 50, 439-46.	0.2	30
92	Epidemiologic analysis of 1,442 children and adolescents registered in the German germ cell tumor protocols. Pediatric Blood and Cancer, 2004, 42, 169-175.	1.5	198
93	High Frequency of Human Papillomavirus 6/11, 16, and 18 Infections in Precancerous Lesions and Squamous Cell Carcinoma of the Conjunctiva in Subtropical Tanzania. American Journal of Clinical Pathology, 2004, 122, 938-943.	0.7	23
94	Ovarian sex cord?stromal tumors?a clinicopathological study of 72 cases from the Kiel Pediatric Tumor Registry. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2003, 443, 549-560.	2.8	81
95	Genetic and Genetic Expression Analyses of Clear Cell Sarcoma of the Kidney. Laboratory Investigation, 2003, 83, 1293-1299.	3.7	49
96	POU5F1 (OCT3/4) identifies cells with pluripotent potential in human germ cell tumors. Cancer Research, 2003, 63, 2244-50.	0.9	487
97	Renal medullary carcinoma: clinical, pathologic, immunohistochemical, and genetic analysis with pathogenetic implications. Urology, 2002, 60, 1083-1089.	1.0	224
98	Genetic analysis of mediastinal nonseminomatous germ cell tumors in children and adolescents. Genes Chromosomes and Cancer, 2002, 34, 115-125.	2.8	90
99	DIAGNOSTIC VALUE OF ALPHA₁-FETOPROTEIN AND BETA-HUMAN CHORIONIC GONADOTROPIN IN INFANCY AND CHILDHOOD. Pediatric Hematology and Oncology, 2001, 18, 11-26.	0.8	111
100	Efficacy and outcome of intensive care in pediatric oncologic patients. Critical Care Medicine, 2001, 29, 2276-2280.	0.9	87
101	Deletion mapping of 6q21-26 and frequency of 1p36 deletion in childhood endodermal sinus tumors by microsatellite analysis. Oncogene, 2001, 20, 8042-8044.	5.9	24
102	Primary Mediastinal Germ Cell Tumors in Children and Adolescents: Results of the German Cooperative Protocols MAKEI 83/86, 89, and 96. Journal of Clinical Oncology, 2000, 18, 832-832.	1.6	110
103	Acute Myelogenous Leukemia After Treatment for Malignant Germ Cell Tumors in Children. Journal of Clinical Oncology, 1999, 17, 3226-3233.	1.6	60
104	DIAGNOSTIC AND THERAPEUTIC PITFALLS IN INFANTS WITH LARGE SACROCOCCYGEAL TUMORS. Pediatric Hematology and Oncology, 1999, 16, 481-482.	0.8	0
105	Acute febrile neutrophilic dermatosis (Sweet syndrome) as initial presentation in a child with acute myelogenous leukemia. , 1998, 31, 178-181.		18