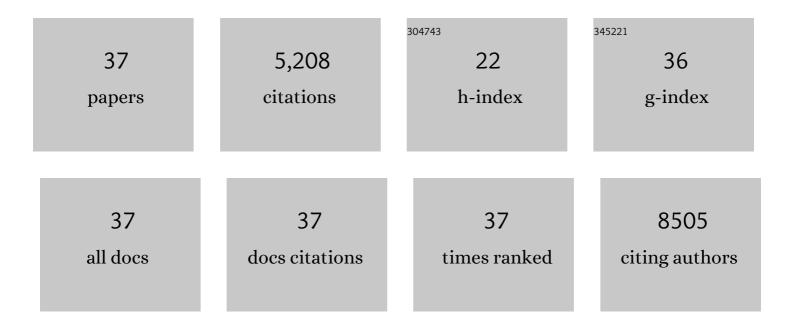
Gudrun Fleischhack

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	DNA methylation-based classification of central nervous system tumours. Nature, 2018, 555, 469-474.	27.8	1,872
2	The landscape of genomic alterations across childhood cancers. Nature, 2018, 555, 321-327.	27.8	1,068
3	Spectrum and prevalence of genetic predisposition in medulloblastoma: a retrospective genetic study and prospective validation in a clinical trial cohort. Lancet Oncology, The, 2018, 19, 785-798.	10.7	268
4	Divergent clonal selection dominates medulloblastoma at recurrence. Nature, 2016, 529, 351-357.	27.8	266
5	Next-generation personalised medicine for high-risk paediatric cancer patients – The INFORM pilot study. European Journal of Cancer, 2016, 65, 91-101.	2.8	262
6	Childhood cancer predisposition syndromes—A concise review and recommendations by the Cancer Predisposition Working Group of the Society for Pediatric Oncology and Hematology. American Journal of Medical Genetics, Part A, 2017, 173, 1017-1037.	1.2	200
7	Treatment of young children with localized medulloblastoma by chemotherapy alone: Results of the prospective, multicenter trial HIT 2000 confirming the prognostic impact of histology. Neuro-Oncology, 2011, 13, 669-679.	1.2	149
8	BET bromodomain protein inhibition is a therapeutic option for medulloblastoma. Oncotarget, 2013, 4, 2080-2095.	1.8	122
9	The Pediatric Precision Oncology INFORM Registry: Clinical Outcome and Benefit for Patients with Very High-Evidence Targets. Cancer Discovery, 2021, 11, 2764-2779.	9.4	110
10	Relapse patterns and outcome after relapse in standard risk medulloblastoma: a report from the HIT-SIOP-PNET4 study. Journal of Neuro-Oncology, 2016, 129, 515-524.	2.9	99
11	Pharmacokinetics Following Intraventricular Administration of Chemotherapy in Patients with Neoplastic Meningitis. Clinical Pharmacokinetics, 2005, 44, 1-31.	3.5	94
12	Evidence of H3 K27M mutations in posterior fossa ependymomas. Acta Neuropathologica, 2016, 132, 635-637.	7.7	73
13	High-dose chemotherapy (HDCT) with auto-SCT in children with atypical teratoid/rhabdoid tumors (AT/RT): a report from the European Rhabdoid Registry (EU-RHAB). Bone Marrow Transplantation, 2014, 49, 370-375.	2.4	58
14	Nonmetastatic Medulloblastoma of Early Childhood: Results From the Prospective Clinical Trial HIT-2000 and An Extended Validation Cohort. Journal of Clinical Oncology, 2020, 38, 2028-2040.	1.6	58
15	Nimotuzumab treatment of malignant gliomas. Expert Opinion on Biological Therapy, 2012, 12, 1649-1659.	3.1	53
16	Meropenem versus ceftazidime as empirical monotherapy in febrile neutropenia of paediatric patients with cancer. Journal of Antimicrobial Chemotherapy, 2001, 47, 841-853.	3.0	49
17	Feasibility of intraventricular administration of etoposide in patients with metastatic brain tumours. British Journal of Cancer, 2001, 84, 1453-1459.	6.4	48
18	Intraventricular methotrexate as part of primary therapy for children with infant and/or metastatic medulloblastoma: Feasibility, acute toxicity and evidence for efficacy. European Journal of Cancer, 2015, 51, 2634-2642.	2.8	44

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#	Article	IF	CITATIONS
19	Clinical Outcomes and Patient-Matched Molecular Composition of Relapsed Medulloblastoma. Journal of Clinical Oncology, 2021, 39, 807-821.	1.6	40
20	Cross-Species Genomics Reveals Oncogenic Dependencies in ZFTA/C11orf95 Fusion–Positive Supratentorial Ependymomas. Cancer Discovery, 2021, 11, 2230-2247.	9.4	39
21	Strategies to improve the quality of survival for childhood brain tumour survivors. European Journal of Paediatric Neurology, 2015, 19, 619-639.	1.6	36
22	Nimotuzumab and radiotherapy for treatment of newly diagnosed diffuse intrinsic pontine glioma (DIPG): a phase III clinical study. Journal of Neuro-Oncology, 2019, 143, 107-113.	2.9	25
23	Treatment of recurrent primitive neuroectodermal tumors (PNET) in children and adolescents with high-dose chemotherapy (HDC) and stem cell support: results of the HITREZ 97 multicentre trial. Journal of Neuro-Oncology, 2014, 120, 635-642.	2.9	21
24	Postponed Is Not Canceled: Role of Craniospinal Radiation Therapy in the Management of Recurrent Infant Medulloblastoma—An Experience From the HIT-REZ 1997 & 2005 Studies. International Journal of Radiation Oncology Biology Physics, 2014, 88, 1019-1024.	0.8	21
25	Local and systemic therapy of recurrent ependymoma in children and adolescents: short- and long-term results of the E-HIT-REZ 2005 study. Neuro-Oncology, 2021, 23, 1012-1023.	1.2	19
26	Intraventricular etoposide safety and toxicity profile in children and young adults with refractory or recurrent malignant brain tumors. Journal of Neuro-Oncology, 2016, 128, 463-471.	2.9	18
27	Functional loss of a noncanonical BCOR–PRC1.1 complex accelerates SHH-driven medulloblastoma formation. Genes and Development, 2020, 34, 1161-1176.	5.9	16
28	Treatment of children under 4 years of age with medulloblastoma and ependymoma in the HIT2000/HIT-REZ 2005 trials: Neuropsychological outcome 5 years after treatment. PLoS ONE, 2020, 15, e0227693.	2.5	14
29	Cyanotic Raynaud's phenomenon with conventional but not with liposomal amphotericin B: three case reports. Mycoses, 1997, 40, 359-361.	4.0	13
30	Hepatic sinusoidal obstruction syndrome and short-term application of 6-thioguanine in pediatric acute lymphoblastic leukemia. Leukemia, 2021, 35, 2650-2657.	7.2	13
31	Relapsed Medulloblastoma in Pre-Irradiated Patients: Current Practice for Diagnostics and Treatment. Cancers, 2022, 14, 126.	3.7	12
32	Local and Systemic Therapy of Recurrent Medulloblastomas in Children and Adolescents: Results of the P-HIT-REZ 2005 Study. Cancers, 2022, 14, 471.	3.7	9
33	Eye Tumors in Childhood as First Sign of Tumor Predisposition Syndromes: Insights from an Observational Study Conducted in Germany and Austria. Cancers, 2021, 13, 1876.	3.7	7
34	Systemic chemotherapy of pediatric recurrent ependymomas: results from the German HIT-REZ studies. Journal of Neuro-Oncology, 2021, 155, 193-202.	2.9	6
35	The Current Landscape of Targeted Clinical Trials in Non-WNT/Non-SHH Medulloblastoma. Cancers, 2022, 14, 679.	3.7	4
36	Procalcitonin in paediatric cancer patients: its diagnostic relevance is superior to that of Câ€reactive protein, interleukin 6, interleukin 8, soluble interleukin 2 receptor and soluble tumour necrosis factor receptor II. British Journal of Haematology, 2000, 111, 1093-1102.	2.5	2

#	Article	IF	CITATIONS
37	Treatment of relapsed AML in children. British Journal of Haematology, 1999, 104, 420-421.	2.5	0