Harm van Tinteren

List of Publications by Year in Descending Order

34 papers	1,691 citations	21 h-index	37 g-index
37 ext. papers	2,097 ext. citations	6.6 avg, IF	3.76 L-index

#	Paper	IF	Citations
34	Characteristics and outcome of children with renal tumors in the Netherlands: The first five-year experience of national centralization.. *PLoS ONE*, **2022**, 17, e0261729	3.7	2
33	Prognostic Factors for Wilms Tumor Recurrence: A Review of the Literature. *Cancers*, **2021**, 13,	6.6	2
32	Locoregional control using highly conformal flank target volumes and volumetric-modulated arc therapy in pediatric renal tumors: Results from the Dutch national cohort. *Radiotherapy and Oncology*, **2021**, 159, 249-254	5.3	3
31	Clinical characteristics and outcomes of children with WAGR syndrome and Wilms tumor and/or nephroblastomatosis: The 30-year SIOP-RTSG experience. *Cancer*, **2021**, 127, 628-638	6.4	9
30	Outcome of Stage IV Completely Necrotic Wilms Tumour and Local Stage III Treated According to the SIOP 2001 Protocol. *Cancers*, **2021**, 13,	6.6	2
29	Characteristics and outcome of pediatric renal cell carcinoma patients registered in the International Society of Pediatric Oncology (SIOP) 93-01, 2001 and UK-IMPORT database: A report of the SIOP-Renal Tumor Study Group. *International Journal of Cancer*, **2021**, 148, 2724-2735	7.5	8
28	Outcome of patients with stage IV high-risk Wilms tumour treated according to the SIOP2001 protocol: A report of the SIOP Renal Tumour Study Group. *European Journal of Cancer*, **2020**, 128, 38-46	7.5	12
27	Tumour-infiltrating lymphocytes (TILs) and BRCA-like status in stage III breast cancer patients randomised to adjuvant intensified platinum-based chemotherapy versus conventional chemotherapy. *European Journal of Cancer*, **2020**, 127, 240-250	7.5	9
26	Is radiotherapy required in first-line treatment of stage I diffuse anaplastic Wilms tumor? A report of SIOP-RTSG, AIEOP, JWiTS, and UKCCSG. *Pediatric Blood and Cancer*, **2020**, 67, e28039	3	9
25	Evaluation of needle biopsy as a potential risk factor for local recurrence of Wilms tumour in the SIOP WT 2001 trial. *European Journal of Cancer*, **2019**, 116, 13-20	7.5	12
24	Prognostic significance of age in 5631 patients with Wilms tumour prospectively registered in International Society of Paediatric Oncology (SIOP) 93-01 and 2001. *PLoS ONE*, **2019**, 14, e0221373	3.7	11
23	Evaluation of boost irradiation in patients with intermediate-risk stage III Wilms tumour with positive lymph nodes only: Results from the SIOP-WT-2001 Registry. *Pediatric Blood and Cancer*, **2018**, 65, e27085	3	6
22	Adjuvant dose-dense doxorubicin-cyclophosphamide versus docetaxel-doxorubicin-cyclophosphamide for high-risk breast cancer: First results of the randomised MATADOR trial (BOOG 2004-04). *European Journal of Cancer*, **2018**, 102, 40-48	7.5	6
21	Position paper: Rationale for the treatment of Wilms tumour in the UMBRELLA SIOP-RTSG 2016 protocol. *Nature Reviews Urology*, **2017**, 14, 743-752	5.5	150
20	Gain of 1q As a Prognostic Biomarker in Wilms Tumors (WTs) Treated With Preoperative Chemotherapy in the International Society of Paediatric Oncology (SIOP) WT 2001 Trial: A SIOP Renal Tumours Biology Consortium Study. *Journal of Clinical Oncology*, **2016**, 34, 3195-203	2.2	75
19	Outcome of localised blastemal-type Wilms tumour patients treated according to intensified treatment in the SIOP WT 2001 protocol, a report of the SIOP Renal Tumour Study Group (SIOP-RTSG). *European Journal of Cancer*, **2015**, 51, 498-506	7.5	47
18	Omission of doxorubicin from the treatment of stage II-III, intermediate-risk Wilms tumour (SIOP WT 2001): an open-label, non-inferiority, randomised controlled trial. *Lancet, The*, **2015**, 386, 1156-64	40	127

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17	Mutations in the SIX1/2 pathway and the DROSHA/DGCR8 miRNA microprocessor complex underlie high-risk blastemal type Wilms tumors. *Cancer Cell*, **2015**, 27, 298-311	24.3	183
16	Multiple mechanisms of MYCN dysregulation in Wilms tumour. *Oncotarget*, **2015**, 6, 7232-43	3.3	66
15	Minimally invasive nephrectomy for Wilms tumors in children - data from SIOP 2001. *Journal of Pediatric Surgery*, **2014**, 49, 1544-8	2.6	48
14	Nephron sparing surgery (NSS) for unilateral wilms tumor (UWT): the SIOP 2001 experience. *Pediatric Blood and Cancer*, **2014**, 61, 2175-9	3	62
13	Treatment and outcome of patients with relapsed clear cell sarcoma of the kidney: a combined SIOP and AIEOP study. *British Journal of Cancer*, **2014**, 111, 227-33	8.7	40
12	Incidence and outcomes of patients with late recurrence of WilmsTtumor. *Pediatric Blood and Cancer*, **2013**, 60, 1612-5	3	27
11	Clear cell sarcomas of the kidney registered on International Society of Pediatric Oncology (SIOP) 93-01 and SIOP 2001 protocols: a report of the SIOP Renal Tumour Study Group. *European Journal of Cancer*, **2013**, 49, 3497-506	7.5	80
10	Gain of 1q is a marker of poor prognosis in WilmsTtumors. *Genes Chromosomes and Cancer*, **2013**, 52, 1065-74	5	45
9	Treatment of pulmonary metastases in children with stage IV nephroblastoma with risk-based use of pulmonary radiotherapy. *Journal of Clinical Oncology*, **2012**, 30, 3533-9	2.2	76
8	The contribution of chest CT-scan at diagnosis in children with unilateral WilmsTtumour. Results of the SIOP 2001 study. *European Journal of Cancer*, **2012**, 48, 1060-5	7.5	46
7	Characteristics and outcome of stage II and III non-anaplastic WilmsTtumour treated according to the SIOP trial and study 93-01. *European Journal of Cancer*, **2012**, 48, 3240-8	7.5	64
6	Malignant rhabdoid tumours of the kidney (MRTKs), registered on recent SIOP protocols from 1993 to 2005: a report of the SIOP renal tumour study group. *Pediatric Blood and Cancer*, **2011**, 56, 733-7	3	93
5	Nephroblastoma: does the decrease in tumor volume under preoperative chemotherapy predict the lymph nodes status at surgery?. *Pediatric Blood and Cancer*, **2011**, 57, 1266-9	3	24
4	Stromal and epithelial predominant Wilms tumours have an excellent outcome: the SIOP 93 01 experience. *Pediatric Blood and Cancer*, **2010**, 55, 233-8	3	35
3	Characteristics and survival of 750 children diagnosed with a renal tumor in the first seven months of life: A collaborative study by the SIOP/GPOH/SFOP, NWTSG, and UKCCSG Wilms tumor study groups. *Pediatric Blood and Cancer*, **2008**, 50, 1130-4	3	112
2	Progression of localised WilmsTtumour during preoperative chemotherapy is an independent prognostic factor: a report from the SIOP 93-01 nephroblastoma trial and study. *European Journal of Cancer*, **2007**, 43, 131-6	7.5	26
1	Reduction of postoperative chemotherapy in children with stage I intermediate-risk and anaplastic WilmsTtumour (SIOP 93-01 trial): a randomised controlled trial. *Lancet, The*, **2004**, 364, 1229-35	40	172