Reem Al-Saadi

List of Publications by Year
in descending order

19	775	840776	794594
papers	citations	11 h-index	19 g-index

19	19	19	1487
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	An organoid biobank for childhood kidney cancers that captures disease and tissue heterogeneity. Nature Communications, 2020, 11, 1310.	12.8	183
2	Whole-exome sequencing reveals the mutational spectrum of testicular germ cell tumours. Nature Communications, 2015, 6, 5973.	12.8	161
3	Stratification of Wilms tumor by genetic and epigenetic analysis. Oncotarget, 2012, 3, 327-335.	1.8	101
4	Subtype-Specific FBXW7 Mutation and MYCN Copy Number Gain in Wilms' Tumor. Clinical Cancer Research, 2010, 16, 2036-2045.	7.0	69
5	Molecular profiling reveals frequent gain of MYCN and anaplasiaâ€specific loss of 4q and 14q in wilms tumor. Genes Chromosomes and Cancer, 2011, 50, 982-995.	2.8	54
6	Allele loss at 16q defines poorer prognosis Wilms tumour irrespective of treatment approach in the UKW1â€“3 clinical trials: A Childrenâ€™s Cancer and Leukaemia Group (CCLG) study. European Journal of Cancer, 2009, 45, 819-826.	2.8	50
7	Characteristics and outcome of pediatric renal cell carcinoma patients registered in the International Society of Pediatric Oncology (<scp>SIOP</scp>) 93â€01, 2001 and <scp>UKâ€IMPORT</scp> database: A report of the <scp>SIOPâ€Renal</scp> Tumor Study Group. International Journal of Cancer, 2021, 148, 2724-2735.	5.1	26
8	Defining a New Prognostic Index for Stage I Nonseminomatous Germ Cell Tumors Using CXCL12 Expression and Proportion of Embryonal Carcinoma. Clinical Cancer Research, 2016, 22, 1265-1273.	7.0	23
9	Impact of fusion gene status versus histology on risk-stratification for rhabdomyosarcoma: Retrospective analyses of patients on UK trials. Pediatric Blood and Cancer, 2017, 64, e26386.	1.5	21
10	Somatic TP53 Mutations Are Detectable in Circulating Tumor DNA from Children with Anaplastic Wilms Tumors. Translational Oncology, 2018, 11, 1301-1306.	3.7	21
11	Prognostic significance of histopathological response to preoperative chemotherapy in unilateral Wilms' tumor: An analysis of 899 patients treated on the SIOP WT 2001 protocol in the UKâ€CCLG and GPOH studies. International Journal of Cancer, 2021, 149, 1332-1340.	5.1	16
12	Outcome of SIOP patients with low- or intermediate-risk Wilms tumour relapsing after initial vincristine and actinomycin-D therapy only â˜ the SIOP 93â€“01 and 2001 protocols. European Journal of Cancer, 2022, 163, 88-97.	2.8	8
13	Comparative analysis of the clinical characteristics and outcomes of patients with Wilms tumor in the United Kingdom and Japan. Pediatric Blood and Cancer, 2021, 68, e29143.	1.5	7
14	Setting international standards for patient and parent involvement and engagement in childhood, adolescent and young adult cancer research: A report from a European Collaborative Workshop. Cancer Reports, 2021, , e1523.	1.4	7
15	Surgical management, staging, and outcomes of Wilms tumours with intravascular extension: Results of the IMPORT study. Journal of Pediatric Surgery, 2022, 57, 572-578.	1.6	7
16	Outcome of Stage IV Completely Necrotic Wilms Tumour and Local Stage III Treated According to the SIOP 2001 Protocol. Cancers, 2021, 13, 976.	3.7	6
17	Characteristics and outcomes of preoperatively treated patients with anaplastic Wilms tumors registered in the UK SIOPâ€WTâ€2001 and IMPORT study cohorts (2002â€2020). Cancer, 2022, 128, 1666-1675.	4.1	6
18	Long-term kidney function in children with Wilms tumour and constitutional WT1 pathogenic variant. Pediatric Nephrology, 2022, 37, 821-832.	1.7	5

#	Article	IF	Citations
19	Prediction of relapse in stage I nonseminomatous germ cell tumors (NSGCT) by CXCL12: Results from the MRC TE08 and TE22 clinical trials.. Journal of Clinical Oncology, 2013, 31, 319-319.	1.6	4