Wolfram Henn

List of Publications by Year
in descending order

21	420	840776 11	940533 16
papers	citations	h-index	g-index

22	22	22	611
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Evidence of Focal Genetic Microheterogeneity in Glioblastoma Multiforme by Area-Specific CGH on Microdissected Tumor Cells. Journal of Neuropathology and Experimental Neurology, 1999, 58, 993-999.	1.7	89
2	High mutation detection rates in cerebral cavernous malformation upon stringent inclusion criteria: oneâ€‘third of probands are minors. Molecular Genetics & Genomic Medicine, 2014, 2, 176-185.	1.2	53
3	Malignant odontogenic myxoma of the maxilla: case with cytogenetic confirmation. Journal of Laryngology and Otology, 2000, 114, 533-535.	0.8	51
4	Atrichia with papular lesions resulting from compound heterozygous mutations in the hairless gene: A lesson for differential diagnosis of alopecia universalis. Journal of the American Academy of Dermatology, 2002, 47, 519-523.	1.2	42
5	New insights into the genetics of glioblastoma multiforme by familial exome sequencing. Oncotarget, 2015, 6, 5918-5931.	1.8	28
6	Clonal chromosome aberrations in cell cultures of synovial tissue from patients with rheumatoid arthritis. Genes Chromosomes and Cancer, 1993, 6, 232-234.	2.8	27
7	Chromosomal findings andp53-mutation analysis in chromophilic renal-cell carcinomas. , 1996, 68, 47-50.		19
8	Unbalanced translocation t(1;3)(p12-13;q11) in meningiomas as the unique feature of chordoid differentiation. Genes Chromosomes and Cancer, 1999, 26, 270-272.	2.8	17
9	Allocation criteria for an initial shortage of a future SARS-CoV-2 vaccine and necessary measures for global immunity. Vaccine, 2020, 38, 5396-5397.	3.8	16
10	Left Ventricular Systolic Dysfunction in Asymptomatic Marfan Syndrome Patients Is Related to the Severity of Gene Mutation: Insights from the Novel Three Dimensional Speckle Tracking Echocardiography. PLoS ONE, 2015, 10, e0124112.	2.5	16
11	Early proliferation enhancement by monosomy 10 and intratumor heterogeneity in malignant human gliomas as revealed by smear preparations from biopsies. , 1996, 16, 180-184.		14
12	A novel site of DNA amplification on chromosome 1p32-33 in a rhabdomyosarcoma revealed by comparative genomic hybridization. Human Genetics, 1996, 97, 87-90.	3.8	12
13	Establishment of a molecular cytogenetic analysis for native tumor tissue of meningiomas-suitable for clinical application. Molecular Cytogenetics, 2014, 7, 12.	0.9	12
14	Monosomy 7p in meningiomas: a rare constituent of tumor progression. Cancer Genetics and Cytogenetics, 2003, 144, 65-68.	1.0	8
15	Reply: Non-invasive prenatal diagnosis: an ethical imperative. Nature Reviews Genetics, 2009, 10, 733-733.	16.3	6
16	A classification of the aims of vaccination and its relevance to transgenerational justice. Journal of Global Health, 2020, 10, 010341.	2.7	4
17	The fetus in the age of the genome. Human Genetics, 2022, 141, 1017-1026.	3.8	3
18	Whole-Genome Sequencing in Diagnostic Medicine: Too Much Information for Doctors and Patients?. Transfusion Medicine and Hemotherapy, 2009, 36, 280-281.	1.6	2

#	Article	IF	Citations
19	General Cognitive Abilities and Psychosocial Development in Children and Adolescents Having a Co-Twin with Down Syndrome. Journal of Pediatrics, 2021, 232, 214-219.	1.8	1
20	Fluorescence in-situ hybridization and immunophenotyping on smear preparations of collagenase-disaggregated fibrous solid tissues. Chromosome Research, 1999, 7, 663-665.	2.2	0
21	Wie viele Verbote brauchen wir? Der Gesetzgeber und das Ãrztliche Ethos. Ethik in Der Medizin, 2009, 21, 85-87.	1.0	0