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	The fetus in the age of the genome. Human Genetics, 2022, 141, 1017-1026.	3.8	3
2	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
3	A classification of the aims of vaccination and its relevance to transgenerational justice. Journal of Global Health, 2020, 10, 010341.	2.7	4
4	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
5	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
6	New insights into the genetics of glioblastoma multiforme by familial exome sequencing. Oncotarget, 2015, 6, 5918-5931.	1.8	28
7	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
8	Establishment of a molecular cytogenetic analysis for native tumor tissue of meningiomas-suitable for clinical application. Molecular Cytogenetics, 2014, 7, 12.	0.9	12
9	Whole-Genome Sequencing in Diagnostic Medicine: Too Much Information for Doctors and Patients?. Transfusion Medicine and Hemotherapy, 2009, 36, 280-281.	1.6	2
10	Wie viele Verbote brauchen wir? Der Gesetzgeber und das Ärztliche Ethos. Ethik in Der Medizin, 2009, 21, 85-87.	1.0	0
11	Reply: Non-invasive prenatal diagnosis: an ethical imperative. Nature Reviews Genetics, 2009, 10, 733-733.	16.3	6
12	Monosomy 7p in meningiomas: a rare constituent of tumor progression. Cancer Genetics and Cytogenetics, 2003, 144, 65-68.	1.0	8
13	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
14	Malignant odontogenic myxoma of the maxilla: case with cytogenetic confirmation. Journal of Laryngology and Otology, 2000, 114, 533-535.	0.8	51
15	Fluorescence in-situ hybridization and immunophenotyping on smear preparations of collagenase-disaggregated fibrous solid tissues. Chromosome Research, 1999, 7, 663-665.	2.2	0
16	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
17	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
18	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

#	Article	IF	Citations
19	Chromosomal findings andp53-mutation analysis in chromophilic renal-cell carcinomas. , 1996, 68, 47-50.		19
20	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
21	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