Zerrin Onadim

List of Publications by Year
in descending order

30 papers	975 citations	623734 14 h-index	501196 28 g-index
30 all docs	30 docs citations	30 times ranked	979 citing authors

#	Article	IF	Citations
1	Whole-Genome Sequencing of Retinoblastoma Reveals the Diversity of Rearrangements Disrupting RB1 and Uncovers a Treatment-Related Mutational Signature. Cancers, 2021, 13, 754.	3.7	16
2	Prognostic Information for Known Genetic Carriers of RB1 Pathogenic Variants (Germline and) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 702	2.4	7
3	MYCN amplification levels in primary retinoblastoma tumors analyzed by Multiple Ligation-dependent Probe Amplification. Ophthalmic Genetics, 2021, 42, 604-611.	1.2	5
4	Examinations under anaesthesia as a measure of disease burden in unilateral retinoblastoma: the London experience. British Journal of Ophthalmology, 2020, 104, 17-22.	3.9	4
5	Number, frequency and time interval of examinations under anesthesia in bilateral retinoblastoma. Graefe's Archive for Clinical and Experimental Ophthalmology, 2020, 258, 879-886.	1.9	4
6	The management of retinoblastoma. Oncogene, 2018, 37, 1551-1560.	5.9	164
7	THE RECOGNITION OF CAVITARY RETINOBLASTOMA TUMORS. Retina, 2018, 38, 782-787.	1.7	7
8	Detection and reporting of RB1 promoter hypermethylation in diagnostic screening. Ophthalmic Genetics, 2018, 39, 526-531.	1.2	11
9	Strabismus in retinoblastoma survivors with long-term follow-up. Journal of AAPOS, 2018, 22, 276.e1-276.e7.	0.3	5
10	The legacy of retinoblastoma: Three unusual tumors in a woman with a history of heritable retinoblastoma â€“ a case report. , 2018, 37, 151-155.		0
11	Primary intravenous chemotherapy for group D retinoblastoma: a 13-year retrospective analysis. British Journal of Ophthalmology, 2017, 101, 82-88.	3.9	33
12	Patient understanding of genetic information influences reproductive decision making in retinoblastoma. Clinical Genetics, 2017, 92, 587-593.	2.0	4
13	Extensive Variation in the Mutation Rate Between and Within Human Genes Associated with Mendelian Disease. Human Mutation, 2016, 37, 488-494.	2.5	17
14	Spectrum ofRB1mutations identified in 403 retinoblastoma patients. Journal of Medical Genetics, 2014, 51, 208-214.	3.2	62
15	A novel real-time PCR assay for quantitative analysis of methylated alleles (QAMA): analysis of the retinoblastoma locus. Nucleic Acids Research, 2004, 32, e125-e125.	14.5	120
16	Comparative genomic hybridization of 49 primary retinoblastoma tumors identifies chromosomal regions associated with histopathology, progression, and patient outcome. Genes Chromosomes and Cancer, 2003, 36, 121-128.	2.8	79
17	Genomic Alterations in Blastic Natural Killer/Extranodal Natural Killer-Like T Cell Lymphoma with Cutaneous Involvement. Journal of Investigative Dermatology, 2003, 121, 618-627.	0.7	40
18	The Chemosensitivity Profile of Retinoblastoma. Recent Results in Cancer Research, 2003, 161, 73-80.	1.8	11

#	Article	IF	Citations
19	The chemosensitivity profile of retinoblastoma. Anti-Cancer Drugs, 2001, 12, A6-A7.	1.4	0
20	A case of naevus sebaceous associated with familial retinoblastoma, multiple lipomata and meningioma. British Journal of Dermatology, 2000, 143, 211-214.	1.5	7
21	Single cell detection of inherited retinoblastoma predisposition. Prenatal Diagnosis, 1999, 19, 1231-1236.	2.3	17
22	The RB1 gene mutation in a child with ectopic intracranial retinoblastoma. British Journal of Cancer, 1997, 76, 1405-1409.	6.4	8
23	Mechanisms of oncogenesis in patients with familial retinoblastoma. British Journal of Cancer, 1993, 68, 958-964.	6.4	37
24	Molecular mechanisms of oncogenic mutations in tumors from patients with bilateral and unilateral retinoblastoma.. Proceedings of the National Academy of Sciences of the United States of America, 1993, 90, 7351-7355.	7.1	79
25	Oncogenic point mutations in exon 20 of the RB1 gene in families showing incomplete penetrance and mild expression of the retinoblastoma phenotype.. Proceedings of the National Academy of Sciences of the United States of America, 1992, 89, 6177-6181.	7.1	122
26	Follow-up of retinoblastoma patients having prenatal and perinatal predictions for mutant gene carrier status using intragenic polymorphic probes from the RB1 gene. British Journal of Cancer, 1992, 65, 711-716.	6.4	27
27	Genetic counselling in retinoblastoma: importance of ocular fundus examination of first degree relatives and linkage analysis.. British Journal of Ophthalmology, 1991, 75, 147-150.	3.9	23
28	Application of PCR amplification of DNA from paraffin embedded tissue sections to linkage analysis in familial retinoblastoma.. Journal of Medical Genetics, 1991, 28, 312-316.	3.2	23
29	Carrier detection and prenatal screening of the retinoblastoma gene. Journal of Pathology, 1990, 161, 3-5.	4.5	3
30	Application of intragenic DNA probes in prenatal screening for retinoblastoma gene carriers in the United Kingdom.. Archives of Disease in Childhood, 1990, 65, 651-656.	1.9	40