

Zerrin Onadim

List of Publications by Year in descending order

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Version: 2024-02-01

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	The management of retinoblastoma. <i>Oncogene</i> , 2018, 37, 1551-1560.	5.9	164
2	Oncogenic point mutations in exon 20 of the RB1 gene in families showing incomplete penetrance and mild expression of the retinoblastoma phenotype.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1992, 89, 6177-6181.	7.1	122
3	A novel real-time PCR assay for quantitative analysis of methylated alleles (QAMA): analysis of the retinoblastoma locus. <i>Nucleic Acids Research</i> , 2004, 32, e125-e125.	14.5	120
4	Molecular mechanisms of oncogenic mutations in tumors from patients with bilateral and unilateral retinoblastoma.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1993, 90, 7351-7355.	7.1	79
5	Comparative genomic hybridization of 49 primary retinoblastoma tumors identifies chromosomal regions associated with histopathology, progression, and patient outcome. <i>Genes Chromosomes and Cancer</i> , 2003, 36, 121-128.	2.8	79
6	Spectrum of RB1 mutations identified in 403 retinoblastoma patients. <i>Journal of Medical Genetics</i> , 2014, 51, 208-214.	3.2	62
7	Application of intragenic DNA probes in prenatal screening for retinoblastoma gene carriers in the United Kingdom.. <i>Archives of Disease in Childhood</i> , 1990, 65, 651-656.	1.9	40
8	Genomic Alterations in Blastic Natural Killer/Extranodal Natural Killer-Like T Cell Lymphoma with Cutaneous Involvement. <i>Journal of Investigative Dermatology</i> , 2003, 121, 618-627.	0.7	40
9	Mechanisms of oncogenesis in patients with familial retinoblastoma. <i>British Journal of Cancer</i> , 1993, 68, 958-964.	6.4	37
10	Primary intravenous chemotherapy for group D retinoblastoma: a 13-year retrospective analysis. <i>British Journal of Ophthalmology</i> , 2017, 101, 82-88.	3.9	33
11	Follow-up of retinoblastoma patients having prenatal and perinatal predictions for mutant gene carrier status using intragenic polymorphic probes from the RB1 gene. <i>British Journal of Cancer</i> , 1992, 65, 711-716.	6.4	27
12	Genetic counselling in retinoblastoma: importance of ocular fundus examination of first degree relatives and linkage analysis.. <i>British Journal of Ophthalmology</i> , 1991, 75, 147-150.	3.9	23
13	Application of PCR amplification of DNA from paraffin embedded tissue sections to linkage analysis in familial retinoblastoma.. <i>Journal of Medical Genetics</i> , 1991, 28, 312-316.	3.2	23
14	Single cell detection of inherited retinoblastoma predisposition. <i>Prenatal Diagnosis</i> , 1999, 19, 1231-1236.	2.3	17
15	Extensive Variation in the Mutation Rate Between and Within Human Genes Associated with Mendelian Disease. <i>Human Mutation</i> , 2016, 37, 488-494.	2.5	17
16	Whole-Genome Sequencing of Retinoblastoma Reveals the Diversity of Rearrangements Disrupting RB1 and Uncovers a Treatment-Related Mutational Signature. <i>Cancers</i> , 2021, 13, 754.	3.7	16
17	Detection and reporting of RB1 promoter hypermethylation in diagnostic screening. <i>Ophthalmic Genetics</i> , 2018, 39, 526-531.	1.2	11
18	The Chemosensitivity Profile of Retinoblastoma. <i>Recent Results in Cancer Research</i> , 2003, 161, 73-80.	1.8	11

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19	The RB1 gene mutation in a child with ectopic intracranial retinoblastoma. British Journal of Cancer, 1997, 76, 1405-1409.	6.4	8
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	THE RECOGNITION OF CAVITARY RETINOBLASTOMA TUMORS. Retina, 2018, 38, 782-787.	1.7	7
22	Prognostic Information for Known Genetic Carriers of RB1 Pathogenic Variants (Germline and) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 622	2.4	7
23	Strabismus in retinoblastoma survivors with long-term follow-up. Journal of AAPOS, 2018, 22, 276.e1-276.e7.	0.3	5
24	<i>MYCN</i> amplification levels in primary retinoblastoma tumors analyzed by Multiple Ligation-dependent Probe Amplification. Ophthalmic Genetics, 2021, 42, 604-611.	1.2	5
25	Patient understanding of genetic information influences reproductive decision making in retinoblastoma. Clinical Genetics, 2017, 92, 587-593.	2.0	4
26	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
27	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
28	Carrier detection and prenatal screening of the retinoblastoma gene. Journal of Pathology, 1990, 161, 3-5.	4.5	3
29	The chemosensitivity profile of retinoblastoma. Anti-Cancer Drugs, 2001, 12, A6-A7.	1.4	0
30	The legacy of retinoblastoma: Three unusual tumors in a woman with a history of heritable retinoblastoma – a case report. , 2018, 37, 151-155.		0