AlÄ^o Karaman

List of Publications by Year in descending order

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1040056 839539 28 350 9 18 citations h-index g-index papers 29 29 29 831 docs citations all docs times ranked citing authors

#	Article	IF	CITATIONS
1	Novel FBN1 mutation in a family with inherited Marfan Syndrome: p.Cys2672Arg. Congenital Anomalies (discontinued), 2018, 58, 41-43.	0.6	2
2	Evaluation of maternal serum folate, vitamin B12, and homocysteine levels andfactor V Leiden, factor II g.20210G>A, and MTHFR variations in prenatallydiagnosed neural tube defects. Turkish Journal of Medical Sciences, 2016, 46, 489-494.	0.9	9
3	Molecular etiology of arthrogryposis in multiple families of mostly Turkish origin. Journal of Clinical Investigation, 2016, 126, 762-778.	8.2	82
4	Evaluation of Factor V Leiden, Prothrombin G20210A, MTHFR C677T and MTHFR A1298C gene polymorphisms in retinopathy of prematurity in a Turkish cohort. Ophthalmic Genetics, 2016, 37, 415-418.	1.2	6
5	A novel missense mutation, p.(R102W) in WNT7A causes Al-Awadi Raas-Rothschild syndrome in a fetus. European Journal of Medical Genetics, 2016, 59, 604-606.	1.3	6
6	Exome sequencing identifies a homozygous <i>C5orf42</i> variant in a Turkish kindred with oralâ€facialâ€digital syndrome type VI. American Journal of Medical Genetics, Part A, 2015, 167, 2132-2137.	1.2	12
7	DNA damage is increased in lymphocytes of patients with metabolic syndrome. Mutation Research - Genetic Toxicology and Environmental Mutagenesis, 2015, 782, 30-35.	1.7	41
8	Whole-exome sequencing links TMCO1 defect syndrome with cerebro-facio-thoracic dysplasia. European Journal of Human Genetics, 2014, 22, 1145-1148.	2.8	19
9	Aplasia cutis congenita and limb anomaly: A case of non-scalp lesion. Goztepe Tip Dergisi, 2014, 28, 220-223.	0.0	1
10	Phototherapy causes a transient DNA damage in jaundiced newborns. Drug and Chemical Toxicology, 2013, 36, 88-92.	2.3	16
11	Jarcho-Levin syndrome (spondylocostal dysostosis) and hydrocephalia: case report. Medical Journal of Bakirkoy, 2013, , 183-185.	0.1	0
12	Duplication 10q syndrome: A new case. Goztepe Tip Dergisi, 2013, 28, 55-57.	0.0	0
13	Noonan syndrome: Case report. Goztepe Tip Dergisi, 2013, 28, 48-50.	0.0	0
14	Genetic alterations in benign, preneoplastic and malignant breast lesions. Indian Journal of Pathology and Microbiology, 2012, 55, 319.	0.2	2
15	Comet assay and analysis of micronucleus formation in patients with rheumatoid arthritis. Mutation Research - Genetic Toxicology and Environmental Mutagenesis, 2011, 721, 1-5.	1.7	24
16	Oro-fasio-dijital sendrom tip 1: Olgu sunumu / Oro-facio-digital syndrome type 1: case report. Medical Journal of Bakirkoy, 2011, , 120-121.	0.1	0
17	Genomic instability in patients with Barrett's esophagus. Cancer Genetics and Cytogenetics, 2010, 201, 88-93.	1.0	4
18	Exposure to bitumen fumes and genotoxic effects on Turkish asphalt workers. Clinical Toxicology, 2009, 47, 321-326.	1.9	25

#	Article	IF	CITATIONS
19	Sister chromatid exchange and micronucleus studies in patients with Behçet's disease. Journal of Cutaneous Pathology, 2009, 36, 831-837.	1.3	32
20	Sister chromatid exchange analysis in patients with psoriasis. Experimental Dermatology, 2008, 17, 524-529.	2.9	5
21	Double aneuploidy in a Turkish child: Down–Klinefelter syndrome. Congenital Anomalies (discontinued), 2008, 48, 45-47.	0.6	11
22	Micronucleus analysis in patients with colorectal adenocarcinoma and colorectal polyps. World Journal of Gastroenterology, 2008, 14, 6835.	3.3	30
23	Alteration of sister chromatid exchange frequencies in gastric cancer and chronic atrophic gastritis patients with and without H pylori infection. World Journal of Gastroenterology, 2008, 14, 2534.	3.3	9
24	Intraperitoneal dedifferentiated liposarcoma: A case report. World Journal of Gastroenterology, 2008, 14, 5927.	3.3	5
25	Bardet-Biedl syndrome: a case report. Dermatology Online Journal, 2008, 14, 9.	0.5	1
26	Frequency of sister chromatid exchanges in the lymphocytes of patients with atopic dermatitis. Journal of Dermatology, 2006, 33, 596-602.	1.2	7
27	Waardenburg syndrome type 1. Dermatology Online Journal, 2006, 12, 21.	0.5	1
28	A 45 Year Old Man With Cutaneous And Subcutaneous Neurofibromas Of Varying Size. Marmara Medical Journal, 0, , .	0.1	0