AlÄº Karaman

List of Publications by Year
in descending order

28	350	1040056 9	839539 18
papers	citations	h-index	g-index

29	29	29	831
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Molecular etiology of arthrogryposis in multiple families of mostly Turkish origin. Journal of Clinical Investigation, 2016, 126, 762-778.	8.2	82
2	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
3	Sister chromatid exchange and micronucleus studies in patients with BehÃ§etâ€™s disease. Journal of Cutaneous Pathology, 2009, 36, 831-837.	1.3	32
4	Micronucleus analysis in patients with colorectal adenocarcinoma and colorectal polyps. World Journal of Gastroenterology, 2008, 14, 6835.	3.3	30
5	Exposure to bitumen fumes and genotoxic effects on Turkish asphalt workers. Clinical Toxicology, 2009, 47, 321-326.	1.9	25
6	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
7	Whole-exome sequencing links TMCO1 defect syndrome with cerebro-facio-thoracic dysplasia. European Journal of Human Genetics, 2014, 22, 1145-1148.	2.8	19
8	Phototherapy causes a transient DNA damage in jaundiced newborns. Drug and Chemical Toxicology, 2013, 36, 88-92.	2.3	16
9	Exome sequencing identifies a homozygous C5orf42 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
10	Double aneuploidy in a Turkish child: Downâ€“Klinefelter syndrome. Congenital Anomalies (discontinued), 2008, 48, 45-47.	0.6	11
11	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
12	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
13	Frequency of sister chromatid exchanges in the lymphocytes of patients with atopic dermatitis. Journal of Dermatology, 2006, 33, 596-602.	1.2	7
14	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
15	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
16	Sister chromatid exchange analysis in patients with psoriasis. Experimental Dermatology, 2008, 17, 524-529.	2.9	5
17	Intraperitoneal dedifferentiated liposarcoma: A case report. World Journal of Gastroenterology, 2008, 14, 5927.	3.3	5
18	Genomic instability in patients with Barrett's esophagus. Cancer Genetics and Cytogenetics, 2010, 201, 88-93.	1.0	4

#	Article	IF	Citations
19	Novel FBN1 mutation in a family with inherited Marfan Syndrome: p.Cys2672Arg. Congenital Anomalies (discontinued), 2018, 58, 41-43.	0.6	2
20	Genetic alterations in benign, preneoplastic and malignant breast lesions. Indian Journal of Pathology and Microbiology, 2012, 55, 319.	0.2	2
21	Aplasia cutis congenita and limb anomaly: A case of non-scalp lesion. Goztepe Tip Dergisi, 2014, 28, 220-223.	0.0	1
22	Waardenburg syndrome type 1. Dermatology Online Journal, 2006, 12, 21.	0.5	1
23	Bardet-Biedl syndrome: a case report. Dermatology Online Journal, 2008, 14, 9.	0.5	1
24	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
25	Jarcho-Levin syndrome (spondylocostal dysostosis) and hydrocephalia: case report. Medical Journal of Bakirkoy, 2013, , 183-185.	0.1	0
26	A 45 Year Old Man With Cutaneous And Subcutaneous Neurofibromas Of Varying Size. Marmara Medical Journal, 0, , .	0.1	0
27	Duplication 10q syndrome: A new case. Goztepe Tip Dergisi, 2013, 28, 55-57.	0.0	0
28	Noonan syndrome: Case report. Goztepe Tip Dergisi, 2013, 28, 48-50.	0.0	0