Bayram Toraman

List of Publications by Year
in descending order

10	400	1478505 6	1281871 11
papers	citations	h-index	g-index

11	11	11	1047
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	CEP152 is a genome maintenance protein disrupted in Seckel syndrome. Nature Genetics, 2011, 43, 23-26.	21.4	201
2	Mutations in RIPK4 Cause the Autosomal-Recessive Form of Popliteal Pterygium Syndrome. American Journal of Human Genetics, 2012, 90, 76-85.	6.2	80
3	Involvement ofDFNB59 mutations in autosomal recessive nonsyndromic hearing impairment. Human Mutation, 2007, 28, 718-723.	2.5	58
4	Analysis of centrosome and DNA damage response in PLK4 associated Seckel syndrome. European Journal of Human Genetics, 2017, 25, 1118-1125.	2.8	21
5	Novel splice-site and missense mutations in theALDH1A3gene underlying autosomal recessive anophthalmia/microphthalmia. British Journal of Ophthalmology, 2014, 98, 832-840.	3.9	18
6	Investigation of CYP21A2 mutations in Turkish patients with 21-hydroxylase deficiency and a novel founder mutation. Gene, 2013, 513, 202-208.	2.2	8
7	RIPK4 suppresses the TGFâ€Î²1 signaling pathway in HaCaT cells. Cell Biology International, 2020, 44, 848-860.	3.0	7
8	A novel homozygous RIPK4 variant in a family with severe Bartsocasâ€Papas syndrome. American Journal of Medical Genetics, Part A, 2021, 185, 1691-1699.	1.2	2
9	Finding underlying genetic mechanisms of two patients with autism spectrum disorder carrying familial apparently balanced chromosomal translocations. Journal of Gene Medicine, 2021, 23, e3322.	2.8	2
10	HLA-E*0101/0103X is Associated with Susceptibility to Pemphigus Vulgaris: A Case-control Study. Acta Dermatovenerologica Croatica, 2017, 25, 189-194.	0.1	2