Kanay Yararbas

List of Publications by Year
in descending order

21	82	1684188 5	1588992 8
papers	citations	h-index	g-index

21	21	21	138
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	DNA fragmentation index and human papilloma virus in males with previous assisted reproductive technology failures. Turkish Journal of Urology, 2019, 45, 12-16.	1.3	15
2	Sarcolemmal Alpha and Gamma Sarcoglycan Protein Deficiencies in Turkish Siblings With a Novel Missense Mutation in the Alpha Sarcoglycan Gene. Pediatric Neurology, 2014, 50, 640-647.	2.1	13
3	AUTS2 Syndrome in a 68â€yearâ€old female: Natural history and further delineation of the phenotype. American Journal of Medical Genetics, Part A, 2016, 170, 3231-3236.	1.2	8
4	Prenatal diagnosis of VACTERL syndrome and partial caudal regression syndrome: A previously unreported association. Journal of Clinical Ultrasound, 2009, 37, 464-466.	0.8	7
5	Unusual Chromosomal Rearrangement Resulted in Interstitial Monosomy 9p: Case Report. Cytogenetic and Genome Research, 2016, 148, 19-24.	1.1	7
6	Concomitant Alpha- and Gamma-Sarcoglycan Deficiencies in a Turkish Boy with a Novel Deletion in the Alpha-Sarcoglycan Gene. Case Reports in Genetics, 2014, 2014, 1-6.	0.2	5
7	Clinical research The significance of Y chromosome microdeletion analysis in subfertile men with clinical variocele. Archives of Medical Science, 2010, 3, 382-387.	0.9	4
8	PRENATAL DIAGNOSIS OF INTRACARDIAC HAMARTOMA AND TURNER SYNDROME. Fetal and Pediatric Pathology, 2010, 29, 330-337.	0.7	4
9	Sarcolemmal deficiency of sarcoglycan complex in an 18-month-old Turkish boy with a large deletion in the beta sarcoglycan gene. Balkan Journal of Medical Genetics, 2015, 18, 71-76.	0.5	4
10	First Observation of Hb South Florida [beta 1(NA1) Val>Met] in Turkey. Turkish Journal of Haematology, 2013, 30, 223-224.	0.2	3
11	A Large PROP1 Gene Deletion in a Turkish Pedigree. Case Reports in Endocrinology, 2018, 2018, 1-5.	0.4	3
12	The miRNA content of circulating exosomes in DLBCL patients and in vitro influence of DLBCL-derived exosomes on miRNA expression of healthy B-cells from peripheral blood. Cancer Biomarkers, 2021, 32, 519-529.	1.7	3
13	Trichorhinophalangeal syndrome type II presenting with short stature in a child. Archivos Argentinos De Pediatria, 2016, 114, e403-e407.	0.2	2
14	del5p/dup5q in a â€˜cri du chatâ€™ patient without parental chromosomal rearrangement. American Journal of Medical Genetics, Part A, 2006, 140A, 1016-1020.	1.2	1
15	First Observation of Hemoglobin Jabalpur [Beta 3 (NA3) Leu>Pro] in the Turkish Population. Turkish Journal of Haematology, 2014, 31, 319-320.	0.5	1
16	A novel TWIST1 gene mutation in a patient with Saethreâ€“Chotzen syndrome. Clinical Dysmorphology, 2017, 26, 175-178.	0.3	1
17	A case with isochromosome 18p and 2q13 deletion including the BUB1 gene. Clinical Dysmorphology, 2018, 27, 101-104.	0.3	1
18	Premarital Genetic Diagnosis Revealed Co-heredity Nature of Beta Globin Gene 25-26 del AA and 3â€™UTR+101 G-C Variants in Two Beta Thalassemia Heterozygotes. Turkish Journal of Haematology, 2017, 34, 105-106.	0.5	0

#	Article	IF	Citations
19	The factors affecting amniocentesis decision by pregnant women in the risk group and the influence of consultant. Perinatal Journal, 2019, 27, 6-13.	0.2	0
20	AÃ§Ä±klanamayan erkek infertilitesi: gerÃ§ekten idiopatik mi? Azoospermik erkeklerde karÅŸilaÅŸtirmali genomik hibridizasyon yÃ¶ntemi ile DNA kopya sayisi varyasyonlari ve aday kromozomal lokuslarin belirlenmesi. Pamukkale Medical Journal, 0, , .	0.2	0
21	Clonal evolution of monosomy 7 in acquired severe aplastic anemia: Two cases treated with allogeneic hematopoietic stem cell transplantation. Turkish Journal of Haematology, 2008, 25, 94-7.	0.5	0