Erdal TunÃ§

List of Publications by Year
in descending order

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19 papers	153 citations	1684188 5 h-index	1281871 11 g-index
19 all docs	19 docs citations	19 times ranked	273 citing authors

#	ARTICLE	IF	CITATIONS
1	Effects of high-molecular-weight polyvinyl chloride on Xenopus laevis adults and embryos: the mRNA expression profiles of Myf5, Esr1, Bmp4, Pax6, and Hsp70 genes during early embryonic development. Environmental Science and Pollution Research, 2022, 29, 14767-14779.	5.3	2
2	L-Glutamic acid monosodium salt reduces the harmful effect of lithium on the development of Xenopus laevis embryos. Environmental Science and Pollution Research, 2020, 27, 42124-42132.	5.3	3
3	16. kromozomun q kolunda gÃ¶zlenen kalÄ±tÄ±msal mozaisizm ve fenotipik etkileri. Cukurova Medical Journal, 2018, 43, 1023-1027.	0.2	1
4	Kanserli hÃ¼cre hatlarÄ±, pasaj sayÄ±sÄ± arttÄ±kÃ§a genomik organizasyonunu ve karyotipini deÄŸiÅŸtirir: sitogenetik bir Ã§alÄ±Åma. Cukurova Medical Journal, 2018, 43, 1-1.	0.2	0
5	Chromosomal findings and sequence analysis of target exons of calcium-sensingreceptor (CaSR) gene in patients with Sagliker syndrome. Turkish Journal of Medical Sciences, 2017, 47, 13-21.	0.9	7
6	Chromosomal analyses of 1510 couples who have experienced recurrent spontaneous abortions. Reproductive BioMedicine Online, 2016, 32, 414-419.	2.4	43
7	FP430WHOLE 13 EXONS OF GNAS1 GENE IN SAGLIKER SYNDROME(SS).COMBINATION-COMPULSION OF BONE DYSPLASIAS-HEREDITARY OSTEODISTROPHIES(BD),CHRONIC KIDNEY DISEASES (CKD) AND SECONDARY HYPERPARATHYROIDISM(SH). Nephrology Dialysis Transplantation, 2015, 30, iii214-iii214.	0.7	0
8	Microchimeric Cells, Sex Chromosome Aneuploidies and Cancer. Pathology and Oncology Research, 2015, 21, 1157-1165.	1.9	4
9	Frequency and Types of Chromosomal Abnormalities in Turkish Women with Amenorrhea. Journal of Pediatric and Adolescent Gynecology, 2014, 27, 274-277.	0.7	5
10	Genetic alterations of chromosomes, p53 and p16 genes in low- and high-grade bladder cancer. Oncology Letters, 2014, 8, 25-32.	1.8	18
11	Genetic polymorphisms of estrogen receptor alpha and catechol-O-methyltransferase genes in Turkish patients with familial prostate carcinoma. Indian Journal of Human Genetics, 2013, 19, 408.	0.7	5
12	Chromosome Imbalances and Alterations in the p53 Gene in Uterine Myomas from the Same Family Members: Familial Leiomyomatosis in Turkey. Asian Pacific Journal of Cancer Prevention, 2013, 14, 651-658.	1.2	1
13	Are there fetal stem cells in the maternal brain?. Neural Regeneration Research, 2013, 8, 593-8.	3.0	0
14	International Evaluation of Unrecognizably Uglifying Human Faces in Late and Severe Secondary Hyperparathyroidism in Chronic Kidney Disease. Sagliker Syndrome. A Unique Catastrophic Entity, Cytogenetic Studies for Chromosomal Abnormalities, Calcium-Sensing Receptor Gene and GNAS1 Mutations. Striking and Promising Missense Mutations on the GNAS1 Gene Exons 1, 4, 10, 4. , 2012, 22, 157-161.		23
15	Identification of chromosome abnormalities in screening of a family with manic depression and psoriasis: Predisposition to aneuploidy. Asian Journal of Psychiatry, 2012, 5, 169-174.	2.0	12
16	Diagnosis of chromosomal abnormalities in a case with thanatophoric dysplasia (TD) type I: the first report describing an important association between cytogenetic findings and TD.. American Journal of Case Reports, 2012, 13, 109-113.	0.8	1
17	The genotoxic effect of nicotine on chromosomes of human fetal cells: The first report described as an important study. Inhalation Toxicology, 2011, 23, 829-834.	1.6	23
18	The Reliability of Maternal Serum Triple Test in Prenatal Diagnosis of Fetal Chromosomal Abnormalities of Pregnant Turkish Women. Genetic Testing and Molecular Biomarkers, 2011, 15, 701-707.	0.7	3

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19	Inheritance of pericentric inversion in chromosome 7 through the three progenies and a newborn with congenital hydronephrosis diagnosed prenatally by fetal urine sampling. Fertility and Sterility, 2008, 89, 228.e1-228.e6.	1.0	2