Erdal Tunç

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9557703/publications.pdf

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

| # | Article | IF | CITATIONS |
|----|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-----|-----------|
| 1 | Chromosomal analyses of 1510 couples who have experienced recurrent spontaneous abortions. Reproductive BioMedicine Online, 2016, 32, 414-419. | 2.4 | 43 |
| 2 | 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 |
| 3 | 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, | | 23 |
| 4 | Genetic alterations of chromosomes, p53 and p16 genes in low- and high-grade bladder cancer. Oncology Letters, 2014, 8, 25-32. | 1.8 | 18 |
| 5 | 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 |
| 6 | 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 |
| 7 | 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 |
| 8 | Frequency and Types of Chromosomal Abnormalities in Turkish Women with Amenorrhea. Journal of Pediatric and Adolescent Gynecology, 2014, 27, 274-277. | 0.7 | 5 |
| 9 | Microchimeric Cells, Sex Chromosome Aneuploidies and Cancer. Pathology and Oncology Research, 2015, 21, 1157-1165. | 1.9 | 4 |
| 10 | 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 |
| 11 | 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 |
| 12 | 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 |
| 13 | 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 |
| 14 | 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 |
| 15 | 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 |
| 16 | 16. kromozomun q kolunda gözlenen kalıtımsal mozaisizm ve fenotipik etkileri. Cukurova Medical Journal, 2018, 43, 1023-1027. | 0.2 | 1 |
| 17 | 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 | O |
| 18 | Kanserli hÃ⅓cre hatları, pasaj sayısı arttıkça genomik organizasyonunu ve karyotipini deÄŸiÅŸtirir: sitogenetik bir çalışma. Cukurova Medical Journal, 2018, 43, 1-1. | 0.2 | 0 |

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|----|--------------------------------------------------------------------------------------------------|-----|-----------|
| 19 | Are there fetal stem cells in the maternal brain?. Neural Regeneration Research, 2013, 8, 593-8. | 3.0 | 0 |