

# Zehra Ordulu

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

25  
papers

691  
citations

11  
h-index

26  
g-index

30  
ext. papers

866  
ext. citations

11.4  
avg, IF

3.02  
L-index

#	Paper	IF	Citations
25	Sertoli-Leydig Cell Tumors of the Ovary With Follicular Differentiation Often Resembling Juvenile Granulosa Cell Tumor: A Report of 38 Cases Including Comments on Sex Cord-Stromal Tumors of Mixed Forms (So-called Gynandroblastoma). <i>American Journal of Surgical Pathology</i> , <b>2021</b> , 45, 59-67	6.7	0
24	Case 9-2021: A 16-Year-Old Boy with Headache, Abdominal Pain, and Hypertension. <i>New England Journal of Medicine</i> , <b>2021</b> , 384, 1145-1155	59.2	1
23	Embryonal rhabdomyosarcoma of the uterine corpus: a clinicopathological and molecular analysis of 21 cases highlighting a frequent association with DICER1 mutations. <i>Modern Pathology</i> , <b>2021</b> , 34, 1758-1762 <sup>2</sup>	8.8	2
22	Hyperprogression of a mismatch repair-deficient colon cancer in a humanized mouse model following administration of immune checkpoint inhibitor pembrolizumab. <i>Oncotarget</i> , <b>2021</b> , 12, 2131-2148	3.3	1
21	Reproducibility of scoring criteria for HER2 immunohistochemistry in endometrial serous carcinoma: a multi-institutional interobserver agreement study. <i>Modern Pathology</i> , <b>2021</b> , 34, 1194-1202	9.8	5
20	Locally Recurrent Secretory Carcinoma of the Breast with NTRK3 Gene Fusion. <i>Oncologist</i> , <b>2021</b> , 26, 818-824	3.74	3
19	Molecular Pathology of Ovarian Epithelial Neoplasms: Predictive, Prognostic, and Emerging Biomarkers. <i>Surgical Pathology Clinics</i> , <b>2021</b> , 14, 415-428	3.9	0
18	Molecular and clinicopathologic characterization of intravenous leiomyomatosis. <i>Modern Pathology</i> , <b>2020</b> , 33, 1844-1860	9.8	6
17	Pediatric liver transplant following near catastrophic head bleed: Lessons learned. <i>Pediatric Transplantation</i> , <b>2020</b> , 24, e13646	1.8	
16	Phenotypic interpretation of complex chromosomal rearrangements informed by nucleotide-level resolution and structural organization of chromatin. <i>European Journal of Human Genetics</i> , <b>2018</b> , 26, 374-381	5.3	8
15	Clinical, pathologic, cytogenetic, and molecular profiling in self-identified black women with uterine leiomyomata. <i>Cancer Genetics</i> , <b>2018</b> , 222-223, 1-8	2.3	8
14	Computational Prediction of Position Effects of Apparently Balanced Human Chromosomal Rearrangements. <i>American Journal of Human Genetics</i> , <b>2017</b> , 101, 206-217	11	38
13	The genomic landscape of balanced cytogenetic abnormalities associated with human congenital anomalies. <i>Nature Genetics</i> , <b>2017</b> , 49, 36-45	36.3	172
12	Implication of LRRC4C and DPP6 in neurodevelopmental disorders. <i>American Journal of Medical Genetics, Part A</i> , <b>2017</b> , 173, 395-406	2.5	22
11	Structural Chromosomal Rearrangements Require Nucleotide-Level Resolution: Lessons from Next-Generation Sequencing in Prenatal Diagnosis. <i>American Journal of Human Genetics</i> , <b>2016</b> , 99, 1015-1033	11	43
10	Intravenous leiomyomatosis: an unusual intermediate between benign and malignant uterine smooth muscle tumors. <i>Modern Pathology</i> , <b>2016</b> , 29, 500-10	9.8	42
9	Fibroids: Genotype and Phenotype. <i>Clinical Obstetrics and Gynecology</i> , <b>2016</b> , 59, 25-9	1.7	6

8	Prenatal diagnosis of chromothripsis, with nine breaks characterized by karyotyping, FISH, microarray and whole-genome sequencing. <i>Prenatal Diagnosis</i> , <b>2015</b> , 35, 299-301	3.2	26
7	From GWAS to Therapy: Fatty Acid Synthase in Uterine Leiomyomata. <i>FASEB Journal</i> , <b>2015</b> , 29, 147.5	0.9	1
6	Describing sequencing results of structural chromosome rearrangements with a suggested next-generation cytogenetic nomenclature. <i>American Journal of Human Genetics</i> , <b>2014</b> , 94, 695-709	11	40
5	Approach to Helicobacter pylori infection in geriatric population. <i>World Journal of Gastrointestinal Pharmacology and Therapeutics</i> , <b>2014</b> , 5, 139-47	3	13
4	Haploinsufficiency of KDM6A is associated with severe psychomotor retardation, global growth restriction, seizures and cleft palate. <i>Human Genetics</i> , <b>2013</b> , 132, 537-52	6.3	48
3	Clinical diagnosis by whole-genome sequencing of a prenatal sample. <i>New England Journal of Medicine</i> , <b>2012</b> , 367, 2226-32	59.2	144
2	The selective expression of ret finger protein in endometrial cancer: can RFP be a marker of serous carcinomas?. <i>Turk Patoloji Dergisi</i> , <b>2012</b> , 28, 213-9	0.6	3
1	Disseminated peritoneal leiomyomatosis after laparoscopic supracervical hysterectomy with characteristic molecular cytogenetic findings of uterine leiomyoma. <i>Genes Chromosomes and Cancer</i> , <b>2010</b> , 49, 1152-60	5	57