# Zhu Zhang

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5979472/publications.pdf

Version: 2024-02-01

| 22 | 386 | 840776 | 794594 |
|---|---|---|---|
| papers | citations | 11 | 19 |
| | | h-index | g-index |

| 30 | 30 | 30 | 552 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Pregnancy outcomes of fetuses with congenital heart disease after a prenatal diagnosis with chromosome microarray. Prenatal Diagnosis, 2022, 42, 79-86. | 2.3 | 6 |
| 2 | Prenatal chromosomal microarray analysis inÂ2466Âfetuses with ultrasonographic soft markers: aÂprospective cohort study. American Journal of Obstetrics and Gynecology, 2021, 224, 516.e1-516.e16. | 1.3 | 35 |
| 3 | Genetic diagnoses in pediatric patients with epilepsy and comorbid intellectual disability. Epilepsy Research, 2021, 170, 106552. | 1.6 | 5 |
| 4 | Prenatal diagnosis of genetic aberrations in fetuses with short femur detected by ultrasound: A prospective cohort study. Prenatal Diagnosis, 2021, 41, 1153-1163. | 2.3 | 6 |
| 5 | New mechanism of partial duplication and deletion of chromosome 8: A case report. World Journal of Clinical Cases, 2021, 9, 7139-7145. | 0.8 | 0 |
| 6 | Prenatal diagnosis of chromosomal aberrations by chromosomal microarray analysis in foetuses with ventriculomegaly. Scientific Reports, 2020, 10, 20765. | 3.3 | 11 |
| 7 | Novel biallelic loss-of-function variants in CEP290 cause Joubert syndrome in two siblings. Human Genomics, 2020, 14, 26. | 2.9 | 4 |
| 8 | Comparison of spontaneous fetal loss rates between women with singleton and twin pregnancies after midâ€trimester amniocentesis â€•A historical cohort study. Prenatal Diagnosis, 2020, 40, 1315-1320. | 2.3 | 4 |
| 9 | Transcriptome changes induced by RUNX3 in cervical cancer cells inï¿½vitro. Oncology Letters, 2020, 19, 651-662. | 1.8 | 0 |
| 10 | Prenatal diagnosis of chromosomal aberrations by chromosomal microarray analysis in fetuses with ultrasound anomalies in the urinary system. Prenatal Diagnosis, 2019, 39, 1096-1106. | 2.3 | 16 |
| 11 | Prenatal Diagnostic Value of Chromosomal Microarray in Fetuses with Nuchal Translucency Greater than 2.5â€‰mm. BioMed Research International, 2019, 2019, 1-9. | 1.9 | 33 |
| 12 | Chromosomal Aberrations in Pediatric Patients with Developmental Delay/Intellectual Disability: A Single-Center Clinical Investigation. BioMed Research International, 2019, 2019, 1-16. | 1.9 | 18 |
| 13 | Infertility in a man with oligoasthenozoospermia associated with mosaic chromosome 22q11 deletion. Molecular Genetics & Genomic Medicine, 2018, 6, 1249-1254. | 1.2 | 1 |
| 14 | Prospective chromosome analysis of 3429 amniocentesis samples in China using copy number variation sequencing. American Journal of Obstetrics and Gynecology, 2018, 219, 287.e1-287.e18. | 1.3 | 71 |
| 15 | Clinical Significance of Serum Interleukin-31 and Interleukin-33 Levels in Patients of Endometrial Cancer: A Case Control Study. Disease Markers, 2016, 2016, 1-7. | 1.3 | 39 |
| 16 | Insertion/deletion polymorphism in IL1A 3â€²â€‰UTR is associated with susceptibility to endometrial cancer in Chinese Han women. Journal of Obstetrics and Gynaecology Research, 2016, 42, 983-989. | 1.3 | 11 |
| 17 | IL-27 suppresses SKOV3 cells proliferation by enhancing STAT3 and inhibiting the Akt signal pathway. Molecular Immunology, 2016, 78, 155-163. | 2.2 | 13 |
| 18 | A polymorphism at miRNA-122-binding site in the IL-1Î± 3â€²UTR is associated with risk of epithelial ovarian cancer. Familial Cancer, 2014, 13, 595-601. | 1.9 | 18 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Prognostic value of IL-27 polymorphisms and the susceptibility to epithelial ovarian cancer in a Chinese population. Immunogenetics, 2014, 66, 85-92. | 2.4 | 35 |
| 20 | Association of an insertion/deletion polymorphism in IL1A 3â€²-UTR with risk for cervical carcinoma in Chinese Han Women. Human Immunology, 2014, 75, 740-744. | 2.4 | 24 |
| 21 | Mirror syndrome associated with Fetal Hemoglobin Bartâ€™s disease: a case report. Archives of Gynecology and Obstetrics, 2013, 288, 1183-1185. | 1.7 | 3 |
| 22 | Association of interleukin-23 receptor gene polymorphisms with risk of ovarian cancer. Cancer Genetics and Cytogenetics, 2010, 196, 146-152. | 1.0 | 30 |