Zhu Zhang

List of Publications by Year
in descending order

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

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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	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
2	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
3	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
4	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
5	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
6	Association of interleukin-23 receptor gene polymorphisms with risk of ovarian cancer. Cancer Genetics and Cytogenetics, 2010, 196, 146-152.	1.0	30
7	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
8	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
9	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
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	IL-27 suppresses SKOV3 cells proliferation by enhancing STAT3 and inhibiting the Akt signal pathway. Molecular Immunology, 2016, 78, 155-163.	2.2	13
12	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
13	Prenatal diagnosis of chromosomal aberrations by chromosomal microarray analysis in foetuses with ventriculomegaly. Scientific Reports, 2020, 10, 20765.	3.3	11
14	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
15	Pregnancy outcomes of fetuses with congenital heart disease after a prenatal diagnosis with chromosome microarray. Prenatal Diagnosis, 2022, 42, 79-86.	2.3	6
16	Genetic diagnoses in pediatric patients with epilepsy and comorbid intellectual disability. Epilepsy Research, 2021, 170, 106552.	1.6	5
17	Novel biallelic loss-of-function variants in CEP290 cause Joubert syndrome in two siblings. Human Genomics, 2020, 14, 26.	2.9	4
18	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

#	Article	IF	Citations
19	Mirror syndrome associated with Fetal Hemoglobin Bartâ€™s disease: a case report. Archives of Gynecology and Obstetrics, 2013, 288, 1183-1185.	1.7	3
20	Infertility in a man with oligoasthenozoospermia associated with mosaic chromosome 22q11 deletion. Molecular Genetics & Genomic Medicine, 2018, 6, 1249-1254.	1.2	1
21	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
22	Transcriptome changes induced by RUNX3 in cervical cancer cells inï¿½vitro. Oncology Letters, 2020, 19, 651-662.	1.8	0