

Ting-Ying Lei

List of Publications by Year in descending order

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12
papers

212
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1163117

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docs citations

14
times ranked

375
citing authors

#	ARTICLE	IF	CITATIONS
1	Whole-exome sequencing for prenatal diagnosis of fetuses with congenital anomalies of the kidney and urinary tract. <i>Nephrology Dialysis Transplantation</i> , 2017, 32, 1665-1675.	0.7	54
2	Whole-exome sequencing in the evaluation of fetal congenital anomalies of the kidney and urinary tract detected by ultrasonography. <i>Prenatal Diagnosis</i> , 2020, 40, 1290-1299.	2.3	24
3	Application of chromosome microarray analysis in patients with unexplained developmental delay/intellectual disability in South China. <i>Pediatrics and Neonatology</i> , 2019, 60, 35-42.	0.9	21
4	Clinical application of SNP array analysis in fetuses with ventricular septal defects and normal karyotypes. <i>Archives of Gynecology and Obstetrics</i> , 2017, 296, 929-940.	1.7	20
5	Prenatal diagnosis of foetuses with congenital abnormalities and duplication of the MECP2 region. <i>Gene</i> , 2014, 546, 222-225.	2.2	17
6	Nonimmune hydrops fetalis: Genetic analysis and clinical outcome. <i>Prenatal Diagnosis</i> , 2020, 40, 803-812.	2.3	17
7	Compound heterozygous mutation of the ASXL3 gene causes autosomal recessive congenital heart disease. <i>Human Genetics</i> , 2021, 140, 333-348.	3.8	17
8	All-trans-retinoid acid induces the differentiation of P19 cells into neurons involved in the PI3K/Akt/GSK3 β signaling pathway. <i>Journal of Cellular Biochemistry</i> , 2020, 121, 4386-4396.	2.6	12
9	Microarray analysis in fetuses with duodenal obstruction: It is not just trisomy 21. <i>Prenatal Diagnosis</i> , 2021, 41, 316-322.	2.3	10
10	Prenatal exome sequencing in fetuses with callosal anomalies. <i>Prenatal Diagnosis</i> , 2022, 42, 744-752.	2.3	10
11	Prenatal diagnosis of Smith-Magenis syndrome in two fetuses with increased nuchal translucency, mild lateral ventriculomegaly, and congenital heart defects. <i>Taiwanese Journal of Obstetrics and Gynecology</i> , 2016, 55, 886-890.	1.3	7
12	Identification of differential microRNAs and messenger RNAs resulting from ASXL transcriptional regulator 3 knockdown during during heart development. <i>Bioengineered</i> , 2022, 13, 9948-9961.	3.2	2