Ting-Ying Lei

List of Publications by Year
in descending order

12	212	1163117 8	1125743 13
papers	citations	h-index	g-index

14	14	14	375
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Prenatal exome sequencing in fetuses with callosal anomalies. Prenatal Diagnosis, 2022, 42, 744-752.	2.3	10
2	Identification of differential microRNAs and messenger RNAs resulting from ASXL transcriptional regulator 3 knockdown during during heart development. Bioengineered, 2022, 13, 9948-9961.	3.2	2
3	Compound heterozygous mutation of the ASXL3 gene causes autosomal recessive congenital heart disease. Human Genetics, 2021, 140, 333-348.	3.8	17
4	Microarray analysis in fetuses with duodenal obstruction: It is not just trisomy 21. Prenatal Diagnosis, 2021, 41, 316-322.	2.3	10
5	Wholeâ€exome sequencing in the evaluation of fetal congenital anomalies of the kidney and urinary tract detected by ultrasonography. Prenatal Diagnosis, 2020, 40, 1290-1299.	2.3	24
6	Allâ€transâ€retinoid acid induces the differentiation of P19 cells into neurons involved in the PI3K/Akt/GSK3Î² signaling pathway. Journal of Cellular Biochemistry, 2020, 121, 4386-4396.	2.6	12
7	Nonimmune hydrops fetalis: Genetic analysis and clinical outcome. Prenatal Diagnosis, 2020, 40, 803-812.	2.3	17
8	Application of chromosome microarray analysis in patients with unexplained developmental delay/intellectual disability in South China. Pediatrics and Neonatology, 2019, 60, 35-42.	0.9	21
9	Whole-exome sequencing for prenatal diagnosis of fetuses with congenital anomalies of the kidney and urinary tract. Nephrology Dialysis Transplantation, 2017, 32, 1665-1675.	0.7	54
10	Clinical application of SNP array analysis in fetuses with ventricular septal defects and normal karyotypes. Archives of Gynecology and Obstetrics, 2017, 296, 929-940.	1.7	20
11	Prenatal diagnosis of Smithâ€“Magenis syndrome in two fetuses with increased nuchal translucency, mild lateral ventriculomegaly, and congenital heart defects. Taiwanese Journal of Obstetrics and Gynecology, 2016, 55, 886-890.	1.3	7
12	Prenatal diagnosis of foetuses with congenital abnormalities and duplication of the MECP2 region. Gene, 2014, 546, 222-225.	2.2	17