

# Xin Yang

## List of Publications by Year in descending order

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Version: 2024-02-01

23  
papers

298  
citations

932766

10  
h-index

940134

16  
g-index

24  
all docs

24  
docs citations

24  
times ranked

470  
citing authors

#	ARTICLE	IF	CITATIONS
1	A novel splicing mutation of <i>ARHGAP29</i> is associated with nonsyndromic cleft lip with or without cleft palate. <i>Journal of Maternal-Fetal and Neonatal Medicine</i> , 2022, 35, 2499-2506.	0.7	7
2	The Application of Crown-Chin Length to Crown-Rump Length Ratio in Predicting Fetal Skeletal Dysplasia at First Trimester. <i>Journal of Ultrasound in Medicine</i> , 2022, , .	0.8	0
3	Prenatal exome sequencing in fetuses with callosal anomalies. <i>Prenatal Diagnosis</i> , 2022, 42, 744-752.	1.1	10
4	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.	1.4	2
5	Case Report: Two Novel L1CAM Mutations in Two Unrelated Chinese Families With X-Linked Hydrocephalus. <i>Frontiers in Genetics</i> , 2022, 13, 810853.	1.1	0
6	The Genetic and Clinical Outcomes in Fetuses With Isolated Fetal Growth Restriction: A Chinese Single-Center Retrospective Study. <i>Frontiers in Genetics</i> , 2022, 13, 856522.	1.1	4
7	Compound heterozygous mutation of the ASXL3 gene causes autosomal recessive congenital heart disease. <i>Human Genetics</i> , 2021, 140, 333-348.	1.8	17
8	Microarray analysis in fetuses with duodenal obstruction: It is not just trisomy 21. <i>Prenatal Diagnosis</i> , 2021, 41, 316-322.	1.1	10
9	Fetal Crown-Chin Length to Crown-Rump Length Ratio as a Prenatal Sonographic Marker for Triploidy at First Trimester. <i>Journal of Ultrasound in Medicine</i> , 2021, , .	0.8	1
10	Risk factors associated with fetal pleural effusion in prenatal diagnosis: a retrospective study in a single institute in Southern China. <i>Journal of Obstetrics and Gynaecology</i> , 2020, 40, 443-447.	0.4	5
11	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.	1.1	24
12	Genetic tests aid in counseling of fetuses with cerebellar vermis defects. <i>Prenatal Diagnosis</i> , 2020, 40, 1228-1238.	1.1	15
13	All-trans-retinoid acid induces the differentiation of P19 cells into neurons involved in the PI3K/Akt/GSK3 $\beta$ signaling pathway. <i>Journal of Cellular Biochemistry</i> , 2020, 121, 4386-4396.	1.2	12
14	Nonimmune hydrops fetalis: Genetic analysis and clinical outcome. <i>Prenatal Diagnosis</i> , 2020, 40, 803-812.	1.1	17
15	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.	0.3	21
16	Application of noninvasive prenatal testing in pregnancies with fetal double bubble sign: Is it feasible?. <i>Prenatal Diagnosis</i> , 2018, 38, 402-405.	1.1	2
17	How to make an accurate diagnosis of fetal pyriform sinus fistula in utero: experience at a single medical center in mainland China. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , 2018, 228, 76-81.	0.5	6
18	Outcome of isolated enlarged cisterna magna identified in utero: experience at a single medical center in mainland China. <i>Prenatal Diagnosis</i> , 2017, 37, 575-582.	1.1	9

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19	Chromosome microarray analysis in the investigation of children with congenital heart disease. <i>BMC Pediatrics</i> , 2017, 17, 117.	0.7	34
20	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.	0.4	54
21	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.	0.8	20
22	Non-invasive prenatal detection of haemoglobin Bart's disease by cardiothoracic ratio during the first trimester. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , 2015, 193, 92-95.	0.5	9
23	Prenatal diagnosis of foetuses with congenital abnormalities and duplication of the MECP2 region. <i>Gene</i> , 2014, 546, 222-225.	1.0	17