

Bin Yu

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

69

papers

669

citations

16

h-index

22

g-index

89

ext. papers

977

ext. citations

3

avg, IF

4.32

L-index

#	Paper	IF	Citations
69	Noninvasive prenatal screening for fetal common sex chromosome aneuploidies from maternal blood. <i>Journal of International Medical Research</i> , 2017 , 45, 621-630	1.4	52
68	Prenatal chromosomal microarray analysis in fetuses with congenital heart disease: a prospective cohort study. <i>American Journal of Obstetrics and Gynecology</i> , 2018 , 218, 244.e1-244.e17	6.4	37
67	Expression profile of circular RNAs in placentas of women with gestational diabetes mellitus. <i>Endocrine Journal</i> , 2019 , 66, 431-441	2.9	36
66	Newborn Screening and Molecular Profile of Congenital Hypothyroidism in a Chinese Population. <i>Frontiers in Genetics</i> , 2018 , 9, 509	4.5	30
65	LincRNA-p21 knockdown reversed tumor-associated macrophages function by promoting MDM2 to antagonize* p53 activation and alleviate breast cancer development. <i>Cancer Immunology, Immunotherapy</i> , 2020 , 69, 835-846	7.4	26
64	Overall evaluation of the clinical value of prenatal screening for fetal-free DNA in maternal blood. <i>Medicine (United States)</i> , 2017 , 96, e7114	1.8	26
63	Down-regulated circPAPPA suppresses the proliferation and invasion of trophoblast cells via the miR-384/STAT3 pathway. <i>Bioscience Reports</i> , 2019 , 39,	4.1	25
62	Targeted next-generation sequencing of thirteen causative genes in Chinese patients with congenital hypothyroidism. <i>Endocrine Journal</i> , 2018 , 65, 1019-1028	2.9	24
61	Establishment of self-sequential longitudinal reference intervals of maternal thyroid function during pregnancy. <i>Experimental Biology and Medicine</i> , 2010 , 235, 1212-5	3.7	24
60	The profile analysis of circular RNAs in human placenta of preeclampsia. <i>Experimental Biology and Medicine</i> , 2018 , 243, 1109-1117	3.7	22
59	Luteolin induces hippocampal neurogenesis in the Ts65Dn mouse model of Down syndrome. <i>Neural Regeneration Research</i> , 2019 , 14, 613-620	4.5	20
58	Higher fetal insulin resistance in Chinese pregnant women with gestational diabetes mellitus and correlation with maternal insulin resistance. <i>PLoS ONE</i> , 2013 , 8, e59845	3.7	19
57	Application of Next-Generation Sequencing Following Tandem Mass Spectrometry to Expand Newborn Screening for Inborn Errors of Metabolism: A Multicenter Study. <i>Frontiers in Genetics</i> , 2019 , 10, 86	4.5	18
56	Iron deficiency in late pregnancy and its associations with birth outcomes in Chinese pregnant women: a retrospective cohort study. <i>Nutrition and Metabolism</i> , 2019 , 16, 30	4.6	17
55	Sequencing shorter cfDNA fragments improves the fetal DNA fraction in noninvasive prenatal testing. <i>American Journal of Obstetrics and Gynecology</i> , 2019 , 221, 345.e1-345.e11	6.4	17
54	Non-invasive prenatal testing to detect chromosome aneuploidies in 57,204 pregnancies. <i>Molecular Cytogenetics</i> , 2019 , 12, 29	2	16
53	Assessment of thyroid function during pregnancy: the advantage of self-sequential longitudinal reference intervals. <i>Archives of Medical Science</i> , 2011 , 7, 679-84	2.9	16

52	Preliminary proteomic-based identification of a novel protein for Down's syndrome in maternal serum. <i>Experimental Biology and Medicine</i> , 2012 , 237, 530-9	3.7	14
51	Downregulation of hsa_circ_0005243 induces trophoblast cell dysfunction and inflammation via the Eatenin and NF- κ B pathways. <i>Reproductive Biology and Endocrinology</i> , 2020 , 18, 51	5	13
50	Increased secreted frizzled-related protein 4 and ficolin-3 levels in gestational diabetes mellitus women. <i>Endocrine Journal</i> , 2018 , 65, 499-508	2.9	13
49	Ficolin-3/adiponectin ratio for the prediction of gestational diabetes mellitus in pregnant women. <i>Journal of Diabetes Investigation</i> , 2018 , 9, 403-410	3.9	12
48	High false-positive non-invasive prenatal screening results for sex chromosome abnormalities: Are maternal factors the culprit?. <i>Prenatal Diagnosis</i> , 2020 , 40, 463-469	3.2	12
47	Associations of serum markers screening for Down's syndrome with pregnancy outcomes: A Chinese retrospective cohort study. <i>Clinica Chimica Acta</i> , 2019 , 489, 130-135	6.2	11
46	Clinical features and pregnancy outcomes of women with abnormal cell-free fetal DNA test results. <i>Annals of Translational Medicine</i> , 2019 , 7, 317	3.2	10
45	Epidemiology of birth defects based on a birth defect surveillance system in Southern Jiangsu, China, 2014-2018. <i>Journal of Maternal-Fetal and Neonatal Medicine</i> , 2020 , 1-7	2	9
44	Mechanisms Underlying Footshock and Psychological Stress-Induced Abrupt Awakening From Posttraumatic "Nightmares". <i>International Journal of Neuropsychopharmacology</i> , 2016 , 19,	5.8	9
43	Transcriptomic Profiling of Human Placenta in Gestational Diabetes Mellitus at the Single-Cell Level. <i>Frontiers in Endocrinology</i> , 2021 , 12, 679582	5.7	9
42	Complicated Relationship between Genetic Mutations and Phenotypic Characteristics in Transient and Permanent Congenital Hypothyroidism: Analysis of Pooled Literature Data. <i>International Journal of Endocrinology</i> , 2020 , 2020, 6808517	2.7	8
41	Tissue Infiltrating Immune Cells as Prognostic Biomarkers in Endometrial Cancer: A Meta-Analysis. <i>Disease Markers</i> , 2020 , 2020, 1805764	3.2	8
40	FTO, GCKR, CDKAL1 and CDKN2A/B gene polymorphisms and the risk of gestational diabetes mellitus: a meta-analysis. <i>Archives of Gynecology and Obstetrics</i> , 2018 , 298, 705-715	2.5	8
39	Not all chromosome aberrations can be detected by NIPT in women at advanced maternal age: A multicenter retrospective study. <i>Clinica Chimica Acta</i> , 2018 , 486, 232-236	6.2	8
38	The assessment of combined karyotype analysis and chromosomal microarray in pregnant women of advanced maternal age: a multicenter study. <i>Annals of Translational Medicine</i> , 2019 , 7, 318	3.2	8
37	Investigating the changes in amino acid values in premature infants: a pilot study. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2018 , 31, 435-441	1.6	7
36	Sequencing Shorter cfDNA Fragments Decreases the False Negative Rate of Non-invasive Prenatal Testing. <i>Frontiers in Genetics</i> , 2020 , 11, 280	4.5	7
35	Early second-trimester plasma cell free DNA levels with subsequent risk of pregnancy complications. <i>Clinical Biochemistry</i> , 2019 , 71, 46-51	3.5	6

34	Birth Defects Data From Population-Based Birth Defects Surveillance System in a District of Southern Jiangsu, China, 2014-2018. <i>Frontiers in Public Health</i> , 2020 , 8, 378	6	6
33	MiR-384 inhibits proliferation and migration of trophoblast cells via targeting PTBP3. <i>Pregnancy Hypertension</i> , 2020 , 21, 132-138	2.6	5
32	Clinical evaluation of NIPS for women at advanced maternal age: a multicenter retrospective study. <i>Journal of Maternal-Fetal and Neonatal Medicine</i> , 2019 , 32, 4080-4085	2	5
31	Analysis of Down syndrome failed to be diagnosed after prenatal screening: A multicenter study. <i>Medicine (United States)</i> , 2017 , 96, e7166	1.8	5
30	Association between low fetal fraction of cell free DNA at the early second-trimester and adverse pregnancy outcomes. <i>Pregnancy Hypertension</i> , 2020 , 22, 101-108	2.6	5
29	Lower detectability of non-invasive prenatal testing compared to prenatal diagnosis in high-risk pregnant women. <i>Annals of Translational Medicine</i> , 2019 , 7, 319	3.2	4
28	The Necessity of Prenatal Diagnosis by CMA for the Women with NIPS-Positive Results. <i>International Journal of Genomics</i> , 2020 , 2020, 2145701	2.5	4
27	Combined identification of lncRNA NONHSAG004550 and NONHSAT125420 as a potential diagnostic biomarker of perinatal depression. <i>Journal of Clinical Laboratory Analysis</i> , 2021 , 35, e23890	3	4
26	Bioinformatics characterization of differential proteins in serum of mothers carrying Down syndrome fetuses: combining bioinformatics and ELISA. <i>Archives of Medical Science</i> , 2012 , 8, 183-91	2.9	3
25	Association of maternal D-dimer level in late pregnancy with birth outcomes in a Chinese cohort. <i>Clinica Chimica Acta</i> , 2020 , 501, 258-263	6.2	3
24	Effect quantification and value prediction of factors in noninvasive detection for specific fetal copy number variants by semiconductor sequencing. <i>Molecular Genetics & Genomic Medicine</i> , 2019 , 7, e00718	2.3	2
23	Placental metabolic profiling in gestational diabetes mellitus: An important role of fatty acids. <i>Journal of Clinical Laboratory Analysis</i> , 2021 , 35, e24096	3	2
22	Fibrin/fibrinogen degradation products in late pregnancy promote macrosomia prediction in normal uncomplicated pregnancy. <i>Placenta</i> , 2020 , 96, 27-33	3.4	2
21	The Optimal Cutoff Value of Z-scores Enhances the Judgment Accuracy of Noninvasive Prenatal Screening. <i>Frontiers in Genetics</i> , 2021 , 12, 690063	4.5	2
20	3-Methylcrotonyl-CoA carboxylase deficiency newborn screening in a population of 536,008: is routine screening necessary?. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2019 , 32, 1321-1326	1.6	2
19	Second-trimester Maternal Serum Screening Biomarkers in the Risk Assessment for Preeclampsia. <i>Annals of Clinical and Laboratory Science</i> , 2018 , 48, 308-313	0.9	2
18	Two Infants With Beta-Ketothiolase Deficiency Identified by Newborn Screening in China. <i>Frontiers in Genetics</i> , 2019 , 10, 451	4.5	1
17	Effect of the inflammatory response on serum indices of iron status in late pregnancy. <i>Journal of Trace Elements in Medicine and Biology</i> , 2020 , 61, 126516	4.1	1

16	Assessing apoptosis gene expression profiling with a PCR array in the hippocampus of Ts65Dn mice. <i>BioMed Research International</i> , 2015 , 2015, 214618	3	1
15	NeoSeq: a new method of genomic sequencing for newborn screening. <i>Orphanet Journal of Rare Diseases</i> , 2021 , 16, 481	4.2	1
14	Estimating the frequency of causal genetic variants in foetuses with congenital heart defects: a Chinese cohort study.. <i>Orphanet Journal of Rare Diseases</i> , 2022 , 17, 2	4.2	1
13	Comprehensive Evaluation of Non-invasive Prenatal Screening to Detect Fetal Copy Number Variations. <i>Frontiers in Genetics</i> , 2021 , 12, 665589	4.5	1
12	Genetic and Phenotypic Characteristics of Congenital Hypothyroidism in a Chinese Cohort. <i>Frontiers in Endocrinology</i> , 2021 , 12, 705773	5.7	1
11	Prenatal Diagnostic Testing Following High-Risk Result from Serological Screening: Which Shall We Select?. <i>International Journal of Women's Health</i> , 2021 , 13, 879-888	2.8	1
10	Impact of maternal thyroid hormone in late pregnancy on adverse birth outcomes: A retrospective cohort study in China. <i>Endocrine Journal</i> , 2021 , 68, 317-328	2.9	1
9	Increased METTL3-mediated mA methylation inhibits embryo implantation by repressing HOXA10 expression in recurrent implantation failure.. <i>Reproductive Biology and Endocrinology</i> , 2021 , 19, 187	5	1
8	Investigation and Application of Risk Factors of Macrosomia Based on 10,396 Chinese Pregnant Women.. <i>Frontiers in Endocrinology</i> , 2022 , 13, 837816	5.7	0
7	Does ceruloplasmin differential express in the brain of Ts65Dn: a mouse mode of Down syndrome?. <i>Neurological Sciences</i> , 2014 , 35, 589-93	3.5	
6	Alterations in the metabolic status of amino acids in newborns of pre-eclampsia women.. <i>Pregnancy Hypertension</i> , 2022 , 27, 170-172	2.6	
5	Bioinformatic characterization of differential proteins in the hippocampus of Ts65Dn: A mouse model of down syndrome. <i>Archives of Biological Sciences</i> , 2014 , 66, 1157-1162	0.7	
4	Variant of TSHR is Not a Frequent Cause of Congenital Hypothyroidism in Chinese Han Patients. <i>International Journal of General Medicine</i> , 2021 , 14, 4135-4143	2.3	
3	More attention should be paid to pregnant women who fail non-invasive prenatal screening. <i>Clinical Biochemistry</i> , 2021 , 96, 33-37	3.5	
2	Association of Maternal Serum Uric Acid and Cystatin C Levels in Late Pregnancy with Adverse Birth Outcomes: An Observational Cohort Study in China.. <i>International Journal of Women's Health</i> , 2022 , 14, 213-223	2.8	
1	Molecular Genetic Screening of Neonatal Intensive Care Units: Hyperbilirubinemia as an Example. <i>The Application of Clinical Genetics</i> , Volume 15, 39-48	3.1	