

Bin Yu

List of Publications by Year in Descending Order

Source: <https://exaly.com/author-pdf/1339435/bin-yu-publications-by-year.pdf>

Version: 2024-04-27

This document has been generated based on the publications and citations recorded by exaly.com. For the latest version of this publication list, visit the link given above.

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	Alterations in the metabolic status of amino acids in newborns of pre-eclampsia women.. <i>Pregnancy Hypertension</i> , 2022 , 27, 170-172	2.6	
68	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
67	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	
66	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
65	NeoSeq: a new method of genomic sequencing for newborn screening. <i>Orphanet Journal of Rare Diseases</i> , 2021 , 16, 481	4.2	1
64	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
63	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
62	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
61	Comprehensive Evaluation of Non-invasive Prenatal Screening to Detect Fetal Copy Number Variations. <i>Frontiers in Genetics</i> , 2021 , 12, 665589	4.5	1
60	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
59	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	
58	Genetic and Phenotypic Characteristics of Congenital Hypothyroidism in a Chinese Cohort. <i>Frontiers in Endocrinology</i> , 2021 , 12, 705773	5.7	1
57	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
56	More attention should be paid to pregnant women who fail non-invasive prenatal screening. <i>Clinical Biochemistry</i> , 2021 , 96, 33-37	3.5	
55	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
54	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
53	MiR-384 inhibits proliferation and migration of trophoblast cells via targeting PTBP3. <i>Pregnancy Hypertension</i> , 2020 , 21, 132-138	2.6	5

52	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
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	Tissue Infiltrating Immune Cells as Prognostic Biomarkers in Endometrial Cancer: A Meta-Analysis. <i>Disease Markers</i> , 2020 , 2020, 1805764	3.2	8
49	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
48	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
47	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
46	Fibrin/fibrinogen degradation products in late pregnancy promote macrosomia prediction in normal uncomplicated pregnancy. <i>Placenta</i> , 2020 , 96, 27-33	3.4	2
45	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
44	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
43	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
42	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
41	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
40	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
39	Non-invasive prenatal testing to detect chromosome aneuploidies in 57,204 pregnancies. <i>Molecular Cytogenetics</i> , 2019 , 12, 29	2	16
38	Two Infants With Beta-Ketothiolase Deficiency Identified by Newborn Screening in China. <i>Frontiers in Genetics</i> , 2019 , 10, 451	4.5	1
37	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
36	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
35	Expression profile of circular RNAs in placentas of women with gestational diabetes mellitus. <i>Endocrine Journal</i> , 2019 , 66, 431-441	2.9	36

34	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
33	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
32	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
31	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
30	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
29	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
28	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
27	Luteolin induces hippocampal neurogenesis in the Ts65Dn mouse model of Down syndrome. <i>Neural Regeneration Research</i> , 2019 , 14, 613-620	4.5	20
26	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
25	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
24	Associations of serum markers screening for Down syndrome with pregnancy outcomes: A Chinese retrospective cohort study. <i>Clinica Chimica Acta</i> , 2019 , 489, 130-135	6.2	11
23	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
22	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
21	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
20	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
19	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
18	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
17	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

16	Newborn Screening and Molecular Profile of Congenital Hypothyroidism in a Chinese Population. <i>Frontiers in Genetics</i> , 2018 , 9, 509	4.5	30
15	The profile analysis of circular RNAs in human placenta of preeclampsia. <i>Experimental Biology and Medicine</i> , 2018 , 243, 1109-1117	3.7	22
14	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
13	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
12	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
11	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
10	Mechanisms Underlying Footshock and Psychological Stress-Induced Abrupt Awakening From Posttraumatic "Nightmares". <i>International Journal of Neuropsychopharmacology</i> , 2016 , 19,	5.8	9
9	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
8	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	
7	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	
6	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
5	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
4	Preliminary proteomic-based identification of a novel protein for Down syndrome in maternal serum. <i>Experimental Biology and Medicine</i> , 2012 , 237, 530-9	3.7	14
3	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
2	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
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	