D-Z Li

List of Publications by Year
in descending order

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		471509	477307
222	1,452	17	29
papers	citations	h-index	g-index

222	222	222	1533
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Whole exome sequencing as a diagnostic adjunct to clinical testing in fetuses with structural abnormalities. Ultrasound in Obstetrics and Gynecology, 2018, 51, 493-502.	1.7	113
2	Noninvasive prenatal diagnosis of common aneuploidies by semiconductor sequencing. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 7415-7420.	7.1	110
3	Efficacy and safety of cordocentesis for prenatal diagnosis. International Journal of Gynecology and Obstetrics, 2006, 93, 13-17.	2.3	64
4	Detection of fetal copy number variants by nonâ€invasive prenatal testing for common aneuploidies. Ultrasound in Obstetrics and Gynecology, 2016, 47, 53-57.	1.7	61
5	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
6	Prenatal Diagnosis of Î²-Thalassemia by Reverse Dot-Blot Hybridization in Southern China. Hemoglobin, 2006, 30, 365-370.	0.8	42
7	Rapid prenatal diagnosis of skeletal dysplasia using medical trio exome sequencing: Benefit for prenatal counseling and pregnancy management. Prenatal Diagnosis, 2020, 40, 577-584.	2.3	42
8	Prenatal diagnosis of fetal multicystic dysplastic kidney via high-resolution whole-genome array. Nephrology Dialysis Transplantation, 2016, 31, 1693-1698.	0.7	31
9	Pre Gestational Thalassemia Screening in Mainland China: The First Two Years of a Preventive Program. Hemoglobin, 2017, 41, 248-253.	0.8	26
10	Pregnancy outcome of autosomal aneuploidies other than common trisomies detected by noninvasive prenatal testing in routine clinical practice. Prenatal Diagnosis, 2018, 38, 849-857.	2.3	24
11	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
12	Prenatal exome sequencing in fetuses with congenital heart defects. Clinical Genetics, 2020, 98, 215-230.	2.0	23
13	Prenatal diagnosis of 17q12 duplication and deletion syndrome in two fetuses with congenital anomalies. Taiwanese Journal of Obstetrics and Gynecology, 2014, 53, 579-582.	1.3	22
14	Prenatal diagnosis of 17q12 deletion syndrome: a retrospective case series. Journal of Obstetrics and Gynaecology, 2019, 39, 323-327.	0.9	22
15	Exome sequencing improves genetic diagnosis of fetal increased nuchal translucency. Prenatal Diagnosis, 2020, 40, 1426-1431.	2.3	22
16	Detection of alpha-thalassemia in beta-thalassemia carriers and prevention of Hb Bart's hydrops fetalis through prenatal screening. Haematologica, 2006, 91, 649-51.	3.5	22
17	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
18	Detection of rare thalassemia mutations using long-read single-molecule real-time sequencing. Gene, 2022, 825, 146438.	2.2	19

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19	Delineation of the molecular basis of borderline hemoglobin A2 in Chinese individuals. Blood Cells, Molecules, and Diseases, 2014, 53, 261-264.	1.4	18
20	Prenatal diagnosis of foetuses with congenital abnormalities and duplication of the MECP2 region. Gene, 2014, 546, 222-225.	2.2	17
21	A novel selective deletion of the major Î±â€globin regulatory element (MCSâ€R2) causing Î±â€thalassaemia. British Journal of Haematology, 2017, 176, 984-986.	2.5	17
22	Compound heterozygous mutation of the ASXL3 gene causes autosomal recessive congenital heart disease. Human Genetics, 2021, 140, 333-348.	3.8	17
23	Association of an Î±-Globin Gene Cluster Duplication and Heterozygous Î²-Thalassemia in a Patient with a Severe Thalassemia Syndrome. Hemoglobin, 2015, 39, 102-106.	0.8	15
24	Screening for common Î²â€globin gene cluster deletions in Chinese individuals with increased hemoglobin F. International Journal of Laboratory Hematology, 2015, 37, 752-757.	1.3	15
25	Rare autosomal trisomies on nonâ€invasive prenatal testing: not as adverse as expected. Ultrasound in Obstetrics and Gynecology, 2019, 54, 838-839.	1.7	15
26	Prenatal diagnosis of Î²-thalassemia in Southern China. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2006, 128, 81-85.	1.1	14
27	Thalassemia Intermedia Caused by 16p13.3 Sectional Duplication in a Î²-Thalassemia Heterozygous Child. Pediatric Hematology and Oncology, 2015, 32, 349-353.	0.8	14
28	Noninvasive prenatal testing: impact on invasive prenatal diagnosis at a mainland Chinese tertiary medical center. Journal of Maternal-Fetal and Neonatal Medicine, 2016, 29, 1-12.	1.5	12
29	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
30	A case of transfusionâ€dependent nondeletional Hb H disease undiagnosed during prenatal screening for thalassemia. Prenatal Diagnosis, 2008, 28, 165-166.	2.3	11
31	Maternal serum PlGF (placental growth factor) in Chinese women in the first trimester undergoing screening for Down syndrome. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2016, 201, 166-170.	1.1	11
32	Fetal Anemia and Hydrops Fetalis Associated with Homozygous Hb Constant Spring (HBA2: c.427Tâ€‰>â€‰C). Hemoglobin, 2016, 40, 97-101.	0.8	11
33	What would be missed in the first trimester if nuchal translucency measurement is replaced by cell free DNA foetal aneuploidy screening?. Journal of Obstetrics and Gynaecology, 2018, 38, 498-501.	0.9	11
34	Clinical outcome of pregnancies with the prenatal double bubble sign â€“ a five-year experience from one single centre in mainland China. Journal of Obstetrics and Gynaecology, 2018, 38, 206-209.	0.9	11
35	Recurrent hypoplasia of corpus callosum as a prenatal phenotype of Xia-Gibbs syndrome caused by maternal germline mosaicism of an AHDC1 variant. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2020, 244, 208-210.	1.1	11
36	Evidence of Selection for theÎ±-Globin Gene Deletions and Triplications in a Southern Chinese Population. Hemoglobin, 2015, 39, 442-444.	0.8	10

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37	Hematological Characteristics of Î²-Globin Gene Mutation â€“50 (G>A) (HBB: c.-100G>A) Carriers in Mainland China. Hemoglobin, 2020, 44, 240-243.	0.8	10
38	First-trimester cystic hygroma and neurodevelopmental disorders: The association to remember. Taiwanese Journal of Obstetrics and Gynecology, 2020, 59, 960-962.	1.3	10
39	Microarray analysis in fetuses with duodenal obstruction: It is not just trisomy 21. Prenatal Diagnosis, 2021, 41, 316-322.	2.3	10
40	Prenatal exome sequencing in fetuses with callosal anomalies. Prenatal Diagnosis, 2022, 42, 744-752.	2.3	10
41	Non-invasive prenatal detection of haemoglobin Bart's disease by cardiothoracic ratio during the first trimester. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2015, 193, 92-95.	1.1	9
42	Outcome of isolated enlarged cisterna magna identified in utero: experience at a single medical center in mainland China. Prenatal Diagnosis, 2017, 37, 575-582.	2.3	9
43	Prenatal diagnosis of Wolf-Hirschhorn syndrome: Ultrasonography and molecular karyotyping results. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2018, 225, 19-21.	1.1	8
44	A cost-effectiveness analysis comparing two different strategies in advanced maternal age: Combined first-trimester screening and maternal blood cell-free DNA testing. Taiwanese Journal of Obstetrics and Gynecology, 2018, 57, 536-540.	1.3	8
45	Identification of thalassemia gene cluster deletion by longâ€read wholeâ€genome sequencing (LRâ€WGS). International Journal of Laboratory Hematology, 2021, 43, 859-865.	1.3	8
46	Fetal micrognathia in the first trimester: An ominous finding even after a normal array. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2021, 263, 176-180.	1.1	8
47	The Codon 37 (TGGâ†'TAG)Î²0-Thalassemia Mutation Found in a Chinese Family. Hemoglobin, 2006, 30, 171-173.	0.8	7
48	Misdiagnosis of Hb Constant Spring (Î±142, Termâ†'Gln,TAAâ†'CAA in Î±2) in a Hb H (Î²4) Disease Child. Hemoglobin, 2007, 31, 105-108.	0.8	7
49	Four Cases of Hb Q-H Disease Found in Southern China. Hemoglobin, 2007, 31, 109-111.	0.8	7
50	Incidental Discovery of Nonpaternity during Prenatal Testing of Genetic Disease. Fetal Diagnosis and Therapy, 2008, 24, 39-41.	1.4	7
51	Congenital Nonspherocytic Hemolytic Anemia Caused by KrÃ¼ppel-Like Factor 1 Gene Variants: Another Case Report. Hemoglobin, 2019, 43, 292-295.	0.8	7
52	A novel splicing mutation of ARHGAP29 is associated with nonsyndromic cleft lip with or without cleft palate. Journal of Maternal-Fetal and Neonatal Medicine, 2022, 35, 2499-2506.	1.5	7
53	Prenatal diagnosis and molecular analysis of type 1 thanatophoric dysplasia. International Journal of Gynecology and Obstetrics, 2005, 91, 268-270.	2.3	6
54	Chorionic villus sampling for early prenatal diagnosis: Experience at a mainland Chinese hospital. Journal of Obstetrics and Gynaecology, 2014, 34, 669-672.	0.9	6

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55	Compound Heterozygosity for HKÎ±Î±and anin CisDeletion of DoubleÎ±Genes Presents asÎ±-Thalassemia Trait. Hemoglobin, 2015, 39, 256-259.	0.8	6
56	Identification of NondeletionalÎ±-Thalassemia in a Prenatal Screening Program by Reverse Dot-Blot in Southern China. Hemoglobin, 2015, 39, 42-45.	0.8	6
57	The Frequency of Î±-Globin Gene Triplication in a Southern Chinese Population. Indian Journal of Hematology and Blood Transfusion, 2016, 32, 320-322.	0.6	6
58	Increased first-trimester nuchal translucency associated with a dicentric chromosome and 9q34.3 microdeletion syndrome. Journal of Obstetrics and Gynaecology, 2017, 37, 327-329.	0.9	6
59	Results of Coexistence of Î²-Thalassemia Minor in Hb H Disease Patients. Hemoglobin, 2018, 42, 306-309.	0.8	6
60	Early prenatal diagnosis of lysosomal storage disorders by enzymatic and molecular analysis. Prenatal Diagnosis, 2018, 38, 779-787.	2.3	6
61	How to make an accurate diagnosis of fetal pyriform sinus fistula in utero: experience at a single medical center in mainland China. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2018, 228, 76-81.	1.1	6
62	The role of ultrasound in the choice between chorionic villus sampling and amniocentesis for patients with a positive NIPT result for trisomy 18/13. Prenatal Diagnosis, 2019, 39, 1155-1158.	2.3	6
63	The role of ultrasound in women with a positive NIPT result for trisomy 18 and 13. Taiwanese Journal of Obstetrics and Gynecology, 2019, 58, 798-800.	1.3	6
64	Detection of confined placental trisomy 16 using non-invasive prenatal testing in a pregnancy associated with intrauterine growth restriction and normal karyotype. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2019, 233, 81-83.	1.1	6
65	Hb Westmead (HBA2: c.369C>G): Hematological Characteristics in Heterozygotes with and without Î±0-Thalassemia. Hemoglobin, 2020, 44, 153-155.	0.8	6
66	Neonatal screening for Î±â€‰thalassemia by cord hemoglobin Barts: how effective is it?. International Journal of Laboratory Hematology, 2015, 37, 649-653.	1.3	5
67	Hb Alesha [Î²67(E11)Valâ†'Met (GTG>ATG);HBB: c.202Gâ€‰>â€‰A] Found in a Chinese Girl. Hemoglobin, 2016, 40, 420-421.	0.8	5
68	Novel FREM1 mutations are associated with severe hydrocephalus and shortened limbs in a prenatal case. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2017, 215, 262-264.	1.1	5
69	A de novo ankyrin mutation (ANK1 Q109X) causing severe hereditary spherocytosis from preterm neonatal period. Annals of Hematology, 2017, 96, 1067-1068.	1.8	5
70	Two Î±1-Globin Gene Point Mutations Causing Severe Hb H Disease. Hemoglobin, 2017, 41, 293-296.	0.8	5
71	KFL1 Gene Variants in Î±-Thalassemia Individuals with Increased Fetal Hemoglobin in a Chinese Population. Hemoglobin, 2018, 42, 161-165.	0.8	5
72	Risk factors associated with fetal pleural effusion in prenatal diagnosis: a retrospective study in a single institute in Southern China. Journal of Obstetrics and Gynaecology, 2020, 40, 443-447.	0.9	5

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73	Detection of an Î±-Globin Fusion Gene Using Real-Time Polymerase Chain Reaction-Based Multicolor Melting Curve. Hemoglobin, 2020, 44, 427-431.	0.8	5
74	Prospective ultrasonographic diagnosis of orofacial clefts during the first trimester. Ultrasound in Obstetrics and Gynecology, 2020, 58, 134-137.	1.7	5
75	Hematological Characteristics of Hb Constant Spring (HBA2: c.427T>C) Carriers in Mainland China. Hemoglobin, 2020, 44, 86-88.	0.8	5
76	Prenatal genetic diagnosis of cardiac rhabdomyoma: A single-center experience. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2020, 249, 7-10.	1.1	5
77	Fetal akinesia: The application of clinical exome sequencing in cases with decreased fetal movement. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2021, 260, 59-63.	1.1	5
78	GATA zinc finger domainâ€containing protein 2A (GATAD2A) deficiency reactivates fetal haemoglobin in patients with Î²â€thalassaemia through impaired formation of methylâ€binding domain protein 2 (MBD2)â€containing nucleosome remodelling and deacetylation (NuRD) complex. British Journal of Haematology, 2021, 193, 1220-1227.	2.5	5
79	Prenatal diagnosis of Miller-Dieker syndrome/PAFAH1B1-related lissencephaly: Ultrasonography and genetically investigative results. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2022, 274, 28-32.	1.1	5
80	Prenatal diagnosis of sex chromosome aneuploidies: experience at a mainland Chinese hospital. Journal of Obstetrics and Gynaecology, 2013, 33, 827-829.	0.9	4
81	First Report of a Chinese Family Carrying a Double Heterozygosity for Hb Q-Thailand and Hb J-Bangkok. Hemoglobin, 2016, 40, 425-427.	0.8	4
82	Prenatal DNA diagnosis of Noonan syndrome in a fetus with increased nuchal translucency using next-generation sequencing. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2016, 201, 229-230.	1.1	4
83	First Case of a Compound Heterozygosity for Two NondeletionalÎ±-Thalassemia mutations, Hb Constant Spring and Hb Quong Sze. Hemoglobin, 2016, 40, 210-212.	0.8	4
84	Prenatal diagnosis of Ectrodactylyâ€“Ectodermal dysplasiaâ€“Cleft (EEC) syndrome in a Chinese woman with a TP63 mutation. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2017, 213, 146-147.	1.1	4
85	Treatment of fetal congenital chylothorax: Report of eight cases at a mainland Chinese medical center. Taiwanese Journal of Obstetrics and Gynecology, 2017, 56, 867-869.	1.3	4
86	Hb A₂-Tianhe (HBD: c.323G>A): First Report in a Chinese Family with Normal Hb A₂-Î²-Thalassemia Trait. Hemoglobin, 2017, 41, 291-292.	0.8	4
87	Fetal-onset congenital dyserythropoietic anemia type 1 due to CDAN1 mutations presenting as hydrops fetalis. Pediatric Hematology and Oncology, 2018, 35, 447-450.	0.8	4
88	Regulatory Single Nucleotide Polymorphism rs368698783 (G>A): a Genetic Modifier of Hb F Production Only under Erythropoietic Stress Characteristic for Î²-Globin Chain Deficiency?. Hemoglobin, 2019, 43, 73-75.	0.8	4
89	A KrÃ¼ppel-Like Factor 1 Gene Mutation Ameliorates the Severity of Î²-Thalassemia: A Case Report. Hemoglobin, 2019, 43, 137-139.	0.8	4
90	Chromosomal microarray analysis detects trisomy 9 mosaicism in a prenatal case not revealed by conventional cytogenetic analysis of cord blood. Journal of Obstetrics and Gynaecology, 2019, 39, 123-125.	0.9	4

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91	Prenatal diagnosis and postnatal management of congenital mesoblastic nephroma: Experience at a single center in China. Prenatal Diagnosis, 2021, 41, 766-771.	2.3	4
92	Fetal akinesia: The need for clinical vigilance in first trimester with decreased fetal movements. Taiwanese Journal of Obstetrics and Gynecology, 2021, 60, 559-562.	1.3	4
93	Contribution of maternal mosaicism to false-positive chromosome X loss associated with noninvasive prenatal testing. Journal of Maternal-Fetal and Neonatal Medicine, 2022, 35, 9647-9653.	1.5	4
94	The Genetic and Clinical Outcomes in Fetuses With Isolated Fetal Growth Restriction: A Chinese Single-Center Retrospective Study. Frontiers in Genetics, 2022, 13, 856522.	2.3	4
95	Hydrops Fetalis Caused by Homozygous Alpha-Thalassemia and Kidd Antigen Alloimmunization in a Chinese Woman. Fetal Diagnosis and Therapy, 2008, 24, 331-333.	1.4	3
96	Monozygotic Twins Discordant for Dandy-Walker Malformation. Fetal Diagnosis and Therapy, 2009, 25, 141-143.	1.4	3
97	Consequences of Delayed Prenatal Diagnosis of Î²-Thalassemia in Mainland China. Hemoglobin, 2016, 40, 191-193.	0.8	3
98	Generation of Induced Pluripotent Stem Cells from Amniotic Fluid Cells of a Fetus with Hb Bartâ€™s Disease. Hemoglobin, 2017, 41, 198-202.	0.8	3
99	First Report of the Rare IVS-II-705 (T>G) Î²-Thalassemia Mutation in a Chinese Family. Hemoglobin, 2017, 41, 286-287.	0.8	3
100	Apparent germline mosaicism for a 15q11-q13 deletion causing recurrent Angelman syndrome in a Chinese family. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2019, 236, 255-257.	1.1	3
101	Foetal phenotype of ALG1-CDG caused by paternal uniparental disomy 16. Journal of Obstetrics and Gynaecology, 2020, 41, 1-3.	0.9	3
102	Can cell-free DNA testing be used in pregnancies with isolated fetal omphalocele? Preliminary evidence from cytogenetic results of prenatal cases. Journal of Maternal-Fetal and Neonatal Medicine, 2021, 34, 624-628.	1.5	3
103	A novel deletion of the major regulatory element flanking the Î±â€globin gene cluster as a cause of Î±⁰â€thalassemia. International Journal of Laboratory Hematology, 2021, 43, O190-O192.	1.3	3
104	Association of Hb Q-Thailand with Heterozygous Hb E in a Chinese Patient. Hemoglobin, 2008, 32, 319-321.	0.8	2
105	Homozygous Î±-Thalassemia-1 Presenting in a Fetus without Anemia. Acta Haematologica, 2010, 123, 207-209.	1.4	2
106	Association of Hb New York with Hb E and Î±⁰-Thalassemia in a Chinese Woman Identified by Sebia Capillarys2 System. Hemoglobin, 2012, 36, 157-160.	0.8	2
107	Î±0-Thalassemia Trait with Normal Red Cell Indices: A Report of Two Cases. Hemoglobin, 2012, 36, 589-591.	0.8	2
108	<scp>H</scp>b <scp>A</scp>₂ levels in patients with nondeletional <scp>H</scp>b <scp>H</scp> disease. International Journal of Laboratory Hematology, 2012, 34, 663-664.	1.3	2

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109	Heterozygous Î²-thalassemia with complete absence of hemoglobin A2in a Chinese adult. International Journal of Laboratory Hematology, 2015, 37, e147-e149.	1.3	2
110	Implementation of Newborn Screening for Hemoglobin H Disease in Mainland China. Indian Journal of Hematology and Blood Transfusion, 2015, 31, 242-246.	0.6	2
111	A Program on Noninvasive Prenatal Diagnosis of Î±-Thalassemia in Mainland China: A Costâ€“Benefit Analysis. Hemoglobin, 2016, 40, 247-249.	0.8	2
112	A Novel Frameshift Mutation at Codons 138/139 (HBB: c.417_418insT) on the Î²-Globin Gene Leads to Î²-Thalassemia. Hemoglobin, 2017, 41, 59-60.	0.8	2
113	Application of noninvasive prenatal testing in pregnancies with fetal double bubble sign: Is it feasible?. Prenatal Diagnosis, 2018, 38, 402-405.	2.3	2
114	Complex interactions between thalassemia defective alleles compromise screening and cause severe anemia in a Chinese family. International Journal of Laboratory Hematology, 2018, 40, e55-e58.	1.3	2
115	A <scp>KLF</scp>1 gene mutation causes Î²â€thalassemia minor in a Chinese family. International Journal of Laboratory Hematology, 2018, 40, e35-e37.	1.3	2
116	Î´-Thalassemia with Complete Absence of Hb A₂in a Chinese Family. Hemoglobin, 2018, 42, 135-137.	0.8	2
117	Early prenatal detection of Bardetâ€Biedl syndrome in a case with postaxial polydactyly and hyperechoic kidneys confirmed by next generation sequencing. Congenital Anomalies (discontinued), 2019, 59, 142-144.	0.6	2
118	Unstable Hemoglobin Variants: The Need for Clinical Vigilance in Infants with Congenital Jaundice. Hemoglobin, 2019, 43, 60-62.	0.8	2
119	Germline mosaicism in an Î±â€thalassemia family: Incidental identification by prenatal ultrasound. Prenatal Diagnosis, 2019, 39, 1166-1169.	2.3	2
120	A Î²-Thalassemia Trait with Two Mutations in Cis in a Chinese Family. Hemoglobin, 2019, 43, 289-291.	0.8	2
121	Diagnostic value of chromosomal microarray in fetuses with increased nuchal translucency. Ultrasound in Obstetrics and Gynecology, 2019, 53, 554-555.	1.7	2
122	Early prenatal detection of triploidy: a 9-year experience in mainland China. Journal of Maternal-Fetal and Neonatal Medicine, 2021, 34, 4072-4076.	1.5	2
123	Prenatal diagnosis of trisomy 22 at the first trimester of pregnancy. Journal of Obstetrics and Gynaecology, 2020, 40, 440-442.	0.9	2
124	Recurrent Wilms tumor in a Chinese family caused by a novel WT1 variant inherited from a mosaic parent. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2020, 254, 333-334.	1.1	2
125	Use of noninvasive prenatal screening with cell-free DNA in late pregnancy with sonographic soft markers. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2020, 252, 431-433.	1.1	2
126	First-trimester detection of micrognathia as a presentation of mandibulofacial dysostosis with microcephaly. Journal of Obstetrics and Gynaecology, 2021, 41, 821-823.	0.9	2

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127	The Trend in Timing of Prenatal Diagnosis for Thalassemia at a Chinese Tertiary Obstetric Center. Hemoglobin, 2020, 44, 325-328.	0.8	2
128	Chromosomal microarray analysis in pregnancies at risk for a molecular disorder. Journal of Maternal-Fetal and Neonatal Medicine, 2021, 34, 159-162.	1.5	2
129	Prenatal detection of 1p36 deletion syndrome: ultrasound findings and microarray testing results. Journal of Maternal-Fetal and Neonatal Medicine, 2021, 34, 2180-2184.	1.5	2
130	Unmasking a recessive allele by a deletion: Early prenatal diagnosis of <scp>Bardetâ€Biedl</scp> syndrome in a Chinese family. Congenital Anomalies (discontinued), 2021, 61, 138-139.	0.6	2
131	Pregnancies with trisomy 2 cells in chorionic villi: Ultrasound determines the outcome. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2021, 261, 247-248.	1.1	2
132	Is there a role for nuchal translucency in the detection of rare chromosomal abnormalities inÂtheÂera of noninvasive prenatal testing?. American Journal of Obstetrics and Gynecology, 2021, 225, 463-464.	1.3	2
133	A genetic approach to the etiologic investigation of isolated intrauterine growth restriction. American Journal of Obstetrics and Gynecology, 2021, 225, 695-696.	1.3	2
134	Impact of cell-free fetal DNA on early invasive prenatal diagnosis at a Chinese reference maternal medicine center. Journal of Maternal-Fetal and Neonatal Medicine, 2020, , 1-5.	1.5	2
135	Idiopathic polyhydramnios and foetal macrosomia in the absence of maternal diabetes: clinical vigilance for costello syndrome. Journal of Obstetrics and Gynaecology, 2022, 42, 704-706.	0.9	2
136	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
137	Congenital extrahepatic portosystemic shunt: An unusual feature in cardio-facio-cutaneous syndrome. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2022, 273, 107-108.	1.1	2
138	The Codon 35 (Aâ€‰>â€‰G) (HBB: c.107Aâ€‰>â€‰G) at theÎ±-Î²Chain Interface of theÎ²-Globin Gene: A Silent Mutation?. Hemoglobin, 2016, 40, 56-58.	0.8	1
139	Incidental prenatal detection of DMD gene deletion using array comparative genomic hybridization. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2016, 203, 338-340.	1.1	1
140	A New Î´-Globin Gene Variant: Hb A₂-Fengshun [Î´121(GH4)Gluâ†'Lys (HBD: c.364Gâ€‰>â€‰A)]. Hemoglobin, 2016, 40, 213-214.	0.8	1
141	Diagnostic Dilemma of Hb Perth [Î²32(B14)Leuâ†'Pro;HBB: c.98Tâ€‰>â€‰C] in Mainland China. Hemoglobin, 2016, 40, 202-205.	0.8	1
142	First Detection of the â^'27 (Aâ€‰>â€‰G) (HBB: c.-77Aâ€‰>â€‰G) Mutation of theÎ²-Globin Gene in a Chinese Family. Hemoglobin, 2016, 40, 59-60.	0.8	1
143	A prenatal case of osteogenesis imperfecta diagnosed with next-generation sequencing. Journal of Obstetrics and Gynaecology, 2017, 37, 809-810.	0.9	1
144	Characterization of a novel Î²â€globin gene cluster deletion causing (^AÎ³Î²)⁰â€thalassemia by nextâ€generation sequencing. International Journal of Laboratory Hematology, 2017, 39, e19-e22.	1.3	1

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145	Germline mosaicism in a DMD family: incidental identification in prenatal diagnosis. Journal of Obstetrics and Gynaecology, 2018, 38, 1166-1168.	0.9	1
146	First Report of a Case with Nondeletional Hb H Disease Caused by IVS-I-116 (A>G) of the Î±2-Globin Gene. Hemoglobin, 2018, 42, 344-346.	0.8	1
147	Early prenatal diagnosis of 49,XXXXY: two case reports. Journal of Obstetrics and Gynaecology, 2019, 39, 275-277.	0.9	1
148	Fetal phenotype of Galloway-Mowat syndrome 3 caused by a specific OSGEP variant. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2019, 242, 182-184.	1.1	1
149	Prenatal Ultrasound Presentations in Late Pregnancies Affected With Alpha Thalassemia Major. Pediatric and Developmental Pathology, 2019, 22, 603-604.	1.0	1
150	Clinical utility of noninvasive prenatal screening for pathogenic copy number variants. American Journal of Obstetrics and Gynecology, 2019, 221, 660.	1.3	1
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