

Prenatal exome sequencing analysis in fetal structural anomalies and ultrasonography (PAGE): a cohort study

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Citation Report

#	ARTICLE	IF	CITATIONS
1	Ultrasound examination: The key to maximising the benefits of advances in molecular diagnostic technologies. <i>Prenatal Diagnosis</i> , 2019, 39, 663-665.	1.1	2
2	Prenatal chromosomal microarray testing of fetuses with ultrasound structural anomalies: A prospective cohort study of over 1000 consecutive cases. <i>Prenatal Diagnosis</i> , 2019, 39, 1064-1069.	1.1	16
4	Fetal arthrogryposis: Challenges and perspectives for prenatal detection and management. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2019, 181, 327-336.	0.7	29
5	Evaluation of Diagnostic Yield in Fetal Whole-Exome Sequencing: A Report on 45 Consecutive Families. <i>Frontiers in Genetics</i> , 2019, 10, 425.	1.1	33
6	Prenatally diagnosed developmental abnormalities of the central nervous system and genetic syndromes: A practical review. <i>Prenatal Diagnosis</i> , 2019, 39, 666-678.	1.1	27
7	Fetal phenotypes emerge as genetic technologies become robust. <i>Prenatal Diagnosis</i> , 2019, 39, 811-817.	1.1	37
8	Prenatal genetic considerations of congenital anomalies of the kidney and urinary tract (CAKUT). <i>Prenatal Diagnosis</i> , 2019, 39, 679-692.	1.1	39
9	Looking Back at Fetal Medicine in India in 2018, and Looking Forward to 2019. <i>Journal of Fetal Medicine</i> , 2019, 06, 47-50.	0.1	2
10	Is it feasible to select fetuses for prenatal WES based on the prenatal phenotype?. <i>Prenatal Diagnosis</i> , 2019, 39, 1039-1040.	1.1	14
11	Human Genetics and Fetal Disease: Assessment of the Fetal Genome. , 2019, , 36-47.		0
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13	Prenatal Diagnosis of Fetuses With Increased Nuchal Translucency by Genome Sequencing Analysis. <i>Frontiers in Genetics</i> , 2019, 10, 761.	1.1	52
14	Introduction of genomics into prenatal diagnostics. <i>Lancet, The</i> , 2019, 393, 719-721.	6.3	13
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16	A systematic-based approach to the genetic etiologies of non-immune hydrops fetalis. <i>Prenatal Diagnosis</i> , 2019, 39, 732-750.	1.1	34
17	From diagnostic yield to clinical impact: a pilot study on the implementation of prenatal exome sequencing in routine care. <i>Genetics in Medicine</i> , 2019, 21, 2303-2310.	1.1	41
18	Antenatal Fetal Assessment: 75 Years Later (1945-2019). <i>Journal of Obstetrics and Gynaecology Canada</i> , 2019, 41, S276-S280.	0.3	0
19	Relationships between the clinical phenotypes and genetic variants associated with the immunological mechanism in childhood idiopathic nephrotic syndrome: protocol for a prospective observational single-centre cohort study. <i>BMJ Open</i> , 2019, 9, e028717.	0.8	0

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21	Chromosomal copy number variations in products of conception from spontaneous abortion by next-generation sequencing technology. Medicine (United States), 2019, 98, e18041.	0.4	17
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40	Single nucleotide polymorphism array analysis of 102 patients with developmental delay and/or intellectual disability from Fujian, China. <i>Clinica Chimica Acta</i> , 2020, 510, 638-643.	0.5	2
41	Evidence for 28 genetic disorders discovered by combining healthcare and research data. <i>Nature</i> , 2020, 586, 757-762.	13.7	343
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54	Genetic diagnosis and clinical evaluation of severe fetal akinesia syndrome. <i>Prenatal Diagnosis</i> , 2020, 40, 1532-1539.	1.1	15
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#	ARTICLE	IF	CITATIONS
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#	ARTICLE	IF	CITATIONS
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287	Prenatal ultrasound finding of atypical genitalia: Counseling, genetic testing and outcomes. Prenatal Diagnosis, 0, , .	1.1	4
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#	ARTICLE	IF	CITATIONS
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