Clinical implementation of cellâ€free <scp>DNA</scp>â perspectives from a national audit

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Citation Report

#	Article	IF	Citations
1	Contingent non-invasive prenatal testing: an opportunity to improve non-genetic aspects of fetal aneuploidy screening. Prenatal Diagnosis, 2015, 35, 1347-1352.	1.1	14
2	National decline in invasive prenatal diagnostic procedures in association with uptake of combined first trimester and cellâ€free <scp>DNA</scp> aneuploidy screening. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2015, 55, 507-510.	0.4	38
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4	Nonâ€invasive prenatal testing: the new era in reproductive medicine. Medical Journal of Australia, 2015, 203, 57-58.	0.8	6
5	Knowledge of nonâ€invasive prenatal testing among pregnant women. Medical Journal of Australia, 2015, 203, 76-76.	0.8	3
6	Ushering in a new dawn in obstetrics and gynecology: the industry of cellâ€free <scp>DNA</scp> testing. Ultrasound in Obstetrics and Gynecology, 2015, 45, 1-3.	0.9	2
7	Cell-free DNA screening for fetal aneuploidy as a clinical service. Clinical Biochemistry, 2015, 48, 932-941.	0.8	75
8	Early clinical experience of cell-free DNA-based aneuploidy screening: A survey of obstetric sonologists in Australia and New Zealand. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2015, 55, 138-143.	0.4	12
9	Analysis of cellâ€free <scp>DNA</scp> in maternal blood in screening for fetal aneuploidies: updated metaâ€analysis. Ultrasound in Obstetrics and Gynecology, 2015, 45, 249-266.	0.9	547
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11	Clinical experience from Thailand: noninvasive prenatal testing as screening tests for trisomies 21, 18 and 13 in 4736 pregnancies. Prenatal Diagnosis, 2016, 36, 224-231.	1.1	18
12	Screening for Down syndrome in the second trimester of pregnancy. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2016, 56, 19-21.	0.4	2
13	â€œI think weâ€™ve got too many tests!â€: Prenatal providersâ€™ reflections on ethical and clinical challenges in the practice integration of cell-free DNA screening. Ethics, Medicine and Public Health, 2016, 2, 334-342.	0.5	38
14	Cell-free DNA testing for 22q11.2 deletion syndrome: appraising the viability, effectiveness and appropriateness of screening. Ultrasound in Obstetrics and Gynecology, 2016, 47, 137-141.	0.9	19
15	Noninvasive prenatal testing in routine clinical practice â€“ An audit of <scp>NIPT</scp> and combined firstâ€trimester screening in an unselected Australian population. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2016, 56, 22-28.	0.4	40
16	Population-based impact of noninvasive prenatal screening on screening and diagnostic testing for fetal aneuploidy. Genetics in Medicine, 2017, 19, 1338-1345.	1.1	59
17	Diagnostic performance and costs of contingent screening models for trisomy 21 incorporating nonâ€invasive prenatal testing. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2017, 57, 432-439.	0.4	11
18	Declining invasive prenatal diagnostic procedures: A comparison of tertiary hospital and national data from 2012 to 2015. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2017, 57, 152-156.	0.4	12

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19	Genomics-based non-invasive prenatal testing for detection of fetal chromosomal aneuploidy in pregnant women. The Cochrane Library, 2017, 11, CD011767.	1.5	67
20	Cell-free fetal DNA testing in singleton IVF conceptions. Human Reproduction, 2018, 33, 572-578.	0.4	47
21	Prenatal diagnosis and socioeconomic status in the nonâ€invasive prenatal testing era: A populationâ€based study. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2018, 58, 404-410.	0.4	34
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28	Fetal fraction and noninvasive prenatal testing: What clinicians need to know. Prenatal Diagnosis, 2020, 40, 155-163.	1.1	82
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38	Ethical issues associated with prenatal screening using nonâ€invasive prenatal testing for sex chromosome aneuploidy. Prenatal Diagnosis, 2023, 43, 226-234.	1.1	9
39	Positive predictive value of noninvasive prenatal testing for sex chromosome abnormalities. Molecular Biology Reports, 2022, 49, 9251-9256.	1.0	4
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