

Matthew Rabinowitz

List of Publications by Year in descending order

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9
papers

1,191
citations

1040056

9
h-index

1474206

9
g-index

9
all docs

9
docs citations

9
times ranked

890
citing authors

#	ARTICLE	IF	CITATIONS
1	Cell-free DNA screening for prenatal detection of 22q11.2 deletion syndrome. American Journal of Obstetrics and Gynecology, 2022, 227, 79.e1-79.e11.	1.3	35
2	Cell-free DNA screening for trisomies 21, 18, and 13 in pregnancies at low and high risk for aneuploidy with genetic confirmation. American Journal of Obstetrics and Gynecology, 2022, 227, 259.e1-259.e14.	1.3	30
3	Validation of a Single-Nucleotide Polymorphism-Based Non-Invasive Prenatal Test in Twin Gestations: Determination of Zygosity, Individual Fetal Sex, and Fetal Aneuploidy. Journal of Clinical Medicine, 2019, 8, 937.	2.4	32
4	An Economic Analysis of Cell-Free DNA Non-Invasive Prenatal Testing in the US General Pregnancy Population. PLoS ONE, 2015, 10, e0132313.	2.5	44
5	Detection of triploid, molar, and vanishing twin pregnancies by a single-nucleotide polymorphism-based noninvasive prenatal test. American Journal of Obstetrics and Gynecology, 2015, 212, 79.e1-79.e9.	1.3	149
6	Single-Nucleotide Polymorphism-Based Noninvasive Prenatal Screening in a High-Risk and Low-Risk Cohort. Obstetrics and Gynecology, 2014, 124, 210-218.	2.4	254
7	Clinical experience and follow-up with large scale single-nucleotide polymorphism-based noninvasive prenatal aneuploidy testing. American Journal of Obstetrics and Gynecology, 2014, 211, 527.e1-527.e17.	1.3	242
8	SNP-based noninvasive prenatal testing detects sex chromosome aneuploidies with high accuracy. Prenatal Diagnosis, 2013, 33, 643-649.	2.3	121
9	Noninvasive prenatal aneuploidy testing of chromosomes 13, 18, 21, X, and Y, using targeted sequencing of polymorphic loci. Prenatal Diagnosis, 2012, 32, 1233-1241.	2.3	284