## Myra J Wick

## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9156985/publications.pdf

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		1163117	888059
17	366	8	17
papers	citations	h-index	g-index
19	19	19	870
all docs	docs citations	times ranked	citing authors

#	Article	IF	CITATIONS
1	Maternal obesity is associated with phenotypic alterations in fetal immune cells by singleâ€cell mass cytometry. American Journal of Reproductive Immunology, 2021, 85, e13358.	1.2	5
2	Management of individuals with germline variants in PALB2: a clinical practice resource of the American College of Medical Genetics and Genomics (ACMG). Genetics in Medicine, 2021, 23, 1416-1423.	2.4	34
3	Quantitative Alterations in Complement Alternative Pathway and Related Genetic Analysis in Severe Phenotype Preeclampsia. Kidney360, 2021, 2, 1463-1472.	2.1	2
4	Pregnancy and birth outcomes after SARS-CoV-2 vaccination in pregnancy. American Journal of Obstetrics & Synecology MFM, 2021, 3, 100467.	2.6	124
5	Points to consider: is there evidence to support BRCA1/2 and other inherited breast cancer genetic testing for all breast cancer patients? A statement of the American College of Medical Genetics and Genomics (ACMG). Genetics in Medicine, 2020, 22, 681-685.	2.4	20
6	Fetal glycosylation defect due to ALG3 and COG5 variants detected via amniocentesis: Complex glycosylation defect with embryonic lethal phenotype. Molecular Genetics and Metabolism, 2020, 131, 424-429.	1.1	6
7	<i>Int22h1/Int22h2</i> â€mediated Xq28 duplication syndrome: de novo duplications, prenatal diagnoses, and additional phenotypic features. Human Mutation, 2020, 41, 1238-1249.	2.5	9
8	Cell-Free DNA Screening During Pregnancy—Reply. JAMA - Journal of the American Medical Association, 2019, 321, 309.	7.4	1
9	Increasing genetic counseling referral rates through bundled interventions after ovarian cancer diagnosis. Gynecologic Oncology, 2018, 149, 121-126.	1.4	15
10	Implementing Group Prenatal Counseling for Expanded Noninvasive Screening Options. Journal of Genetic Counseling, 2018, 27, 894-901.	1.6	14
11	Use of Genetic Testing after Abnormal Screening Ultrasound: A Descriptive Cohort Study. Gynecologic and Obstetric Investigation, 2018, 83, 466-470.	1.6	6
12	Noninvasive Prenatal Genetic Screening Using Cell-free DNA. JAMA - Journal of the American Medical Association, 2018, 320, 591.	7.4	28
13	What do the new American College of Medical Genetics and Genomics (ACMG) guidelines mean for the provision of non-invasive prenatal genetic screening?. Journal of Obstetrics and Gynaecology, 2017, 37, 795-798.	0.9	2
14	Offering Prenatal Screening in the Age of Genomic Medicine: A Practical Guide. Journal of Women's Health, 2017, 26, 755-761.	3.3	9
15	False Negative Cell-Free DNA Screening Result in a Newborn with Trisomy 13. Case Reports in Genetics, 2016, 2016, 1-5.	0.2	2
16	Functional characterization of a <i><scp>GFAP</scp></i> variant of uncertain significance in an Alexander disease case within the setting of an individualized medicine clinic. Clinical Case Reports (discontinued), 2016, 4, 885-895.	0.5	3
17	Outcome of Whole Exome Sequencing for Diagnostic Odyssey Cases of an Individualized Medicine Clinic. Mayo Clinic Proceedings, 2016, 91, 297-307.	3.0	83