

# Myra J Wick

## List of Publications by Year in descending order

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

Version: 2024-02-01

17  
papers

366  
citations

1163117

8  
h-index

888059

17  
g-index

19  
all docs

19  
docs citations

19  
times ranked

870  
citing authors

#	ARTICLE	IF	CITATIONS
1	Pregnancy and birth outcomes after SARS-CoV-2 vaccination in pregnancy. American Journal of Obstetrics & Gynecology MFM, 2021, 3, 100467.	2.6	124
2	Outcome of Whole Exome Sequencing for Diagnostic Odyssey Cases of an Individualized Medicine Clinic. Mayo Clinic Proceedings, 2016, 91, 297-307.	3.0	83
3	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
4	Noninvasive Prenatal Genetic Screening Using Cell-free DNA. JAMA - Journal of the American Medical Association, 2018, 320, 591.	7.4	28
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	Increasing genetic counseling referral rates through bundled interventions after ovarian cancer diagnosis. Gynecologic Oncology, 2018, 149, 121-126.	1.4	15
7	Implementing Group Prenatal Counseling for Expanded Noninvasive Screening Options. Journal of Genetic Counseling, 2018, 27, 894-901.	1.6	14
8	Offering Prenatal Screening in the Age of Genomic Medicine: A Practical Guide. Journal of Women's Health, 2017, 26, 755-761.	3.3	9
9	CR-mediated Xq28 duplication syndrome: de novo duplications, prenatal diagnoses, and additional phenotypic features. Human Mutation, 2020, 41, 1238-1249.	2.5	9
10	Use of Genetic Testing after Abnormal Screening Ultrasound: A Descriptive Cohort Study. Gynecologic and Obstetric Investigation, 2018, 83, 466-470.	1.6	6
11	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
12	Maternal obesity is associated with phenotypic alterations in fetal immune cells by single-cell mass cytometry. American Journal of Reproductive Immunology, 2021, 85, e13358.	1.2	5
13	Functional characterization of a GFAP 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
14	False Negative Cell-Free DNA Screening Result in a Newborn with Trisomy 13. Case Reports in Genetics, 2016, 2016, 1-5.	0.2	2
15	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
16	Quantitative Alterations in Complement Alternative Pathway and Related Genetic Analysis in Severe Phenotype Preeclampsia. Kidney360, 2021, 2, 1463-1472.	2.1	2
17	Cell-Free DNA Screening During Pregnancy—Reply. JAMA - Journal of the American Medical Association, 2019, 321, 309.	7.4	1