# Mary M Jenkins

List of Publications by Year
in descending order



| 18 | 413 | 933447 10 | 888059 17 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 18 | 18 | 18 | 550 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Exome sequencing identifies variants in infants with sacral agenesis. Birth Defects Research, 2022, 114, 215-227. | 1.5 | 2 |
| 2 | A genomeâ€wide association study of obstructive heart defects among participants in the National Birth Defects Prevention Study. American Journal of Medical Genetics, Part A, 2022, 188, 2303-2314. | 1.2 | 3 |
| 3 | Exome sequencing identifies genetic variants in anophthalmia and microphthalmia. American Journal of Medical Genetics, Part A, 2022, 188, 2376-2388. | 1.2 | 2 |
| 4 | Paternal genetic variants and risk of obstructive heart defects: A parent-of-origin approach. PLoS Genetics, 2021, 17, e1009413. | 3.5 | 2 |
| 5 | Exome sequencing of childâ€“parent trios with bladder exstrophy: Findings in 26 children. American Journal of Medical Genetics, Part A, 2021, 185, 3028-3041. | 1.2 | 4 |
| 6 | A genome-wide association study implicates the BMP7 locus as a risk factor for nonsyndromic metopic craniosynostosis. Human Genetics, 2020, 139, 1077-1090. | 3.8 | 24 |
| 7 | Exome sequencing of family trios from the National Birth Defects Prevention Study: Tapping into a rich resource of genetic and environmental data. Birth Defects Research, 2019, 111, 1618-1632. | 1.5 | 9 |
| 8 | Genomeâ€wide association studies of structural birth defects: A review and commentary. Birth Defects Research, 2019, 111, 1329-1342. | 1.5 | 34 |
| 9 | Impact of sample collection participation on the validity of estimated measures of association in the National Birth Defects Prevention Study when assessing geneâ€environment interactions. Genetic Epidemiology, 2017, 41, 834-843. | 1.3 | 1 |
| 10 | The national birth defects prevention study: A review of the methods. Birth Defects Research Part A: Clinical and Molecular Teratology, 2015, 103, 656-669. | 1.6 | 188 |
| 11 | Challenges in Studying Modifiable Risk Factors for Birth Defects. Current Epidemiology Reports, 2015, 2, 23-30. | 2.4 | 31 |
| 12 | Maternal smoking, xenobiotic metabolizing enzyme gene variants, and gastroschisis risk. American Journal of Medical Genetics, Part A, 2014, 164, 1454-1463. | 1.2 | 23 |
| 13 | Qualitative assessment of study materials and communication strategies used in studies that include DNA collection. American Journal of Medical Genetics, Part A, 2011, 155, 2721-2731. | 1.2 | 10 |
| 14 | Evaluation of mailed pediatric buccal cytobrushes for use in a case-control study of birth defects. Birth Defects Research Part A: Clinical and Molecular Teratology, 2011, 91, 642-648. | 1.6 | 14 |
| 15 | Maternal attitudes toward DNA collection for geneâ€“environment studies: A qualitative research study. American Journal of Medical Genetics, Part A, 2009, 149A, 2378-2386. | 1.2 | 21 |
| 16 | Ethical issues raised by incorporation of genetics into the National Birth Defects Prevention Study. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2008, 148C, 40-46. | 1.6 | 14 |
| 17 | A novel stop codon mutation (X417L) of the ferrochelatase gene in bovine protoporphyria, a natural animal model of the human disease. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 1998, 1408, 18-24. | 3.8 | 15 |
| 18 | A high-frequency polymorphism of NADH-cytochrome b5 reductase in African-Americans. Human Genetics, 1997, 99, 248-250. | 3.8 | 16 |