# Annabelle Enriquez

List of Publications by Year
in descending order



| 12 | 523 | 1163117 8 | 1281871 11 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 12 | 12 | 12 | 1142 |
|---|---|---|---|
| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | NAD Deficiency, Congenital Malformations, and Niacin Supplementation. New England Journal of Medicine, 2017, 377, 544-552. | 27.0 | 177 |
| 2 | Altered Expression of HepaticCYP2E1andCYP4Ain Obese, Diabeticob/obMice, andfa/faZucker Rats. Biochemical and Biophysical Research Communications, 1999, 255, 300-306. | 2.1 | 105 |
| 3 | Constitutive and Inducible Expression of Hepatic CYP2E1 in Leptin-deficient ob/ob Mice. Biochemical and Biophysical Research Communications, 2000, 268, 337-344. | 2.1 | 66 |
| 4 | Identification of clinically actionable variants from genome sequencing of families with congenital heart disease. Genetics in Medicine, 2019, 21, 1111-1120. | 2.4 | 54 |
| 5 | Functional genomics and gene-environment interaction highlight the complexity of congenital heart disease caused by Notch pathway variants. Human Molecular Genetics, 2020, 29, 566-579. | 2.9 | 32 |
| 6 | Revealing hidden genetic diagnoses in the ocular anterior segment disorders. Genetics in Medicine, 2020, 22, 1623-1632. | 2.4 | 31 |
| 7 | KBG syndrome: An Australian experience. American Journal of Medical Genetics, Part A, 2017, 173, 1866-1877. | 1.2 | 25 |
| 8 | Heterozygous loss of WBP11 function causes multiple congenital defects in humans and mice. Human Molecular Genetics, 2021, 29, 3662-3678. | 2.9 | 14 |
| 9 | Recurrence of split hand/foot malformation, cleft lip/palate, and severe urogenital abnormalities due to germline mosaicism for TP63 mutation. American Journal of Medical Genetics, Part A, 2016, 170, 2372-2376. | 1.2 | 8 |
| 10 | A new era of genetic testing in congenital heart disease: A review. Trends in Cardiovascular Medicine, 2022, 32, 311-319. | 4.9 | 7 |
| 11 | CHDgene: A Curated Database for Congenital Heart Disease Genes. Circulation Genomic and Precision Medicine, 2022, 15, 101161CIRCGEN121003539. | 3.6 | 4 |
| 12 | Hemophagocytic Lymphohistiocytosis in Loeys-Dietz Syndrome. Journal of Clinical Immunology, 2018, 38, 234-236. | 3.8 | 0 |