

Suppl 4. List of excluded studies after full-text screening

| First Author & Year | Title | Reason for Exclusion |
|--------------------------------|---|-----------------------------|
| Bailly 2019 | Role of family history and clinical screening in the identification of families with idiopathic dilated cardiomyopathy in Johannesburg, South Africa | Family history |
| Gimeno 2009 | Penetrance and Risk Profile in Inherited Cardiac Diseases Studied in a Dedicated Screening Clinic | Family history |
| Goerss 1995 | Frequency of familial dilated cardiomyopathy | Family history |
| Huggins 2022 | Prevalence and Cumulative Risk of Familial Idiopathic Dilated Cardiomyopathy | Family history |
| Marume 2020 | Prognosis and clinical characteristics of dilated cardiomyopathy with family history via pedigree analysis | Family history |
| Michels 1991 | The frequency of familial dilated cardiomyopathy in a series of patients with idiopathic dilated cardiomyopathy | Family history |
| Monserrat 2002 | Familial dilated cardiomyopathy in patients transplanted for idiopathic dilated cardiomyopathy | Family history |
| Venugopalan 2001 | Low proportion of familial dilated cardiomyopathy in an Arab population with a high prevalence of consanguineous marriages | Family history |
| Akinrinade 2015 | Prevalence of titin truncating variants in general population | Outcomes |
| Andreasen 2013 | New population-based exome data are questioning the pathogenicity of previously cardiomyopathy-associated genetic variants | Outcomes |
| Bourfiss 2022 | Prevalence and disease expression of pathogenic and likely pathogenic variants associated with inherited cardiomyopathies in the general population | Outcomes |
| Cannata 2021 | Titin mutations and female sex characterize dilated cardiomyopathy in the elderly | Outcomes |
| Golbus 2012 | Population-based variation in cardiomyopathy genes | Outcomes |
| Mahmood 2022 | Disease Penetrance in Asymptomatic Carriers of Familial Cardiomyopathy Variants | Outcomes |
| Nallari 2009 | Epidemiology and genetics of dilated cardiomyopathy in the Indian context | Outcomes |
| Nouhravesh 2016 | Analyses of more than 60,000 exomes questions the role of numerous genes previously associated with dilated cardiomyopathy | Outcomes |
| Paldino 2021 | CLINICAL MANIFESTATION and PROGNOSIS of DIFFERENT CARDIOMYOPATHY TYPES on the BASIS of GENETIC BACKGROUND | Outcomes |
| Vaikhanskaya 2020 | Comparative genetic, clinical features, and 5-years outcomes for dilated vs noncompaction cardiomyopathy | Outcomes |
| Andrews 2008 | New-onset heart failure due to heart muscle disease in childhood: A prospective study in the United Kingdom and Ireland | Population |
| Bick 2012 | Burden of rare sarcomere gene variants in the Framingham and Jackson Heart study cohorts | Population |
| Keren 1988 | Features of mildly dilated congestive cardiomyopathy compared with idiopathic restrictive cardiomyopathy and typical dilated cardiomyopathy | Population |
| Pasotti 2008 | Long-Term Outcome and Risk Stratification in Dilated Cardiomyopathies | Population |
| Seidelmann 2016 | Familial dilated cardiomyopathy diagnosis is commonly overlooked at the time of transplant listing | Population |
| Seliem 2007 | Influence of consanguinity on the pattern of familial aggregation of congenital cardiovascular anomalies in an outpatient population: Studies from the Eastern Province of Saudi Arabia | Population |
| van Waning 2019 | Cardiac Phenotypes, Genetics, and Risks in Familial Noncompaction Cardiomyopathy | Population |
| Yeh 2019 | Targeted next generation sequencing for genetic mutations of dilated cardiomyopathy | Population |
| Yin 2015 | Association of the LMNA gene single nucleotide polymorphism rs4641 with dilated cardiomyopathy | Population |
| Zhao 2015 | Targeted Next-Generation Sequencing Reveals Hot Spots and Doubly Heterozygous Mutations in Chinese Patients with Familial Cardiomyopathy | Study design |
| Akinrinade 2016 | Relevance of truncating titin mutations in dilated cardiomyopathy | Other |
| Pena-Pena 2021 | Clinical utility of genetic testing in patients with dilated cardiomyopathy | Other |
| Ware 2016 | Shared genetic predisposition in Peripartum and dilated cardiomyopathies | Other |