Μονάδα Κληρονομικών και Σπανίων Καρδιαγγειακών Παθήσεων



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| --- |
| Cardiomyopathies  |
|   | ORPHAcode  | ICD-10  |
| Hypertrophic cardiomyopathy  | 217569  | I42.1/I42.2  |
| Dilated cardiomyopathy  | 217604  | I42.0  |
| Arrhythmogenic right ventricular cardiomyopathy  | 247  | I42.8  |
| Left ventricular non-compaction  | 54260  | I42.8  |
| Restrictive cardiomyopathy  | 217635  | I42.5  |
| Peripartum cardiomyopathy  | 563  | O90.3  |
| Unclassified cardiomyopathy  | 217678  | -  |
| Myocarditis  | -  | I41.2  |
| Noonan syndrome  | 638/648  | Q87.1  |
| Costello syndrome  | 3071  | Q87.8  |
| Leopard syndrome  | 500  | Q87.1  |
| Neurofibromatosis  | 636  | Q85.0  |
| Cardiofaciocutaneous syndrome  | 1340  | Q87.8  |
| Danon disease  | 34587  | E74.0  |
| Pompe disease  |   | E74.0  |
| MPS/PRKAG2 without orpha codes  | --  |   |
| Fabry disease  | 324  | E75.2  |
| Hereditary familial Amyloidosis  | 271861  | --  |
| Barth syndrome  | 111  | E71.1  |
| MELAS syndrome  | 550  | E71.3  |
| Friedriech  | 95  | G11.1  |
| Becker syndrome  | 262  | G71.0  |
| Duchenne  | 262  | G71.0  |
| Myotonic dystrophy  | 206647  | G71.1  |
| Emery Dreifuss  | 261  | G71.0  |
| Down  | 870  | Q90.0 /Q90.1 Q90.2 / Q90.9  |
| Channelopathies  |
| Familial long QT syndrome  | 768  | I45.8  |
| Brugada syndrome  | 130  | I49.8  |
| Catecholaminergic polymorphic ventricular tachycardia  | 3286  | I47.2  |
| Familial short QT syndrome  | 51083  | I49.8  |
| Idiopathic ventricular fibrillation - not Brugada type  | 228140  | I49.0  |
| Aortopathies  |  |  |
| Marfan syndrome  | 558  | Q87.4  |
| Loeys-Dietz syndrome  | 60030  | Q87.4  |
| Ehlers-Danlos syndrome  | 98249  | Q79.6  |
| Rare disease with thoracic aortic aneurysm and aortic dissection  | 285014  | I71  |

Καρδιολόγος

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