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# Cardiovascular Genetics ICD-10 Code Reference Sheet*

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|---|---|---|---|---|
| CARDIOMYOPATHY | I42.0 | Dilated cardiomyopathy | I42.1 | Hypertrophic obstructive cardiomyopathy |
| | I42.2 | Hypertrophic non-obstructive cardiomyopathy | I42.5 | Cardiomyopathy, other restrictive |
| | I42.8 | Other cardiomyopathies | I42.9 | Cardiomyopathy, unspecified |
| | I43 | Syndromic cardiomyopathy | I51.7 | Ventricular hypertrophy |
| ARRHYTHMIA | I42.8 | Arrhythmogenic right ventricular dysplasia (ARVD) | I44.2 | Atrioventricular block, complete |
| | I45.81 | Long QT syndrome | I45.89 | Other specified conduction disorders |
| | I46.2 | Cardiac arrest due to underlying cardiac condition | I47.20 | Ventricular tachycardia, unspecified |
| | I47.21 | Torsades de pointes | I47.29 | Other ventricular tachycardia |
| | I48.0 | Paroxysmal atrial fibrillation | I48.2 | Chronic atrial fibrillation |
| | I48.91 | Unspecified atrial fibrillation | I49.01 | Ventricular fibrillation |
| | I49.1 | Atrial premature depolarization (PACs) | I49.3 | Ventricular premature depolarization (PVCs) |
| | I49.5 | Sick sinus syndrome | I49.8 | Other specified cardiac arrhythmias |
| | Q23.81 | Bicuspid aortic valve | Q23.82 | Congenital mitral valve cleft leaflet |
| | Q23.88 | Other congenital malformations of aortic and mitral valves | R00.1 | Bradycardia, unspecified |
| | R00.2 | Palpitations | R94.31 | Abnormal electrocardiogram (ECG)(EKG) |
| AORTIC ANEURYSMS/MARFAN SYNDROME | H27.10 | Unspecified dislocation of the lens | H52.11 | Myopia, right eye |
| | H52.12 | Myopia, left eye | H52.13 | Myopia, bilateral eyes |
| | I34.1 | Mitral valve prolapse | I71.00 | Dissection of unspecified site of aorta |
| | I71.01 | Dissection of thoracic aorta | I71.02 | Dissection of abdominal aorta |
| | I71.10 | Thoracic aortic aneurysm, ruptured, unspecified | I71.11 | Aneurysm of the ascending aorta, ruptured |
| | I71.12 | Aneurysm of the aortic arch, ruptured | I71.13 | Aneurysm of the descending thoracic aorta, ruptured |
| | I71.20 | Thoracic aortic aneurysm, without rupture, unspecified | I71.21 | Aneurysm of the ascending aorta, without rupture |
| | I71.22 | Aneurysm of the aortic arch, without rupture | I71.23 | Aneurysm of the descending thoracic aorta, without rupture |
| | I71.8 | Aortic aneurysm of unspecified site, ruptured | I71.9 | Aortic aneurysm of unspecified site, without rupture |
| | I77.810 | Thoracic aortic ectasia | J93.0 | Spontaneous tension pneumothorax |
| | J93.11 | Primary spontaneous pneumothorax | J93.12 | Secondary spontaneous pneumothorax |
| | J93.81 | Chronic pneumothorax | J93.83 | Other pneumothorax |
| | L90.6 | Stria Atrophicae (stretch marks) | L98.8 | Other specified disorders of skin and subcutaneous tissue |
| | M35.7 | Hypermobility Syndrome | Q12.1 | Congenital displaced lens |
| | Q67.5 | Congenital deformity of the spine (scoliosis) | Q67.6 | Pectus excavatum |
| | Q67.7 | Pectus Carinatum | Q68.1 | Arachnodactyly ("Congenital deformity of finger(s) and hand") |
| | Q87.40 | Marfan syndrome | Q87.410 | Marfan syndrome with aortic dilation |
| | Q87.418 | Marfan syndrome with other cardiovascular manifestations | Q87.42 | Marfan syndrome with ocular manifestations |
| | Q87.43 | Marfan syndrome with skeletal manifestations | R23.3 | Spontaneous Ecchymoses (easy bruising) |

*This is not a comprehensive list of ICD-10 codes, but these codes are more commonly used for cardiovascular genetic testing. There are more specific codes available. Please visit cms.gov/medicare-coverage-database/staticpages/icd-10-code-lookup.aspx for a complete list of codes.

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| CHOLESTEROL/CORONARY ARTERY DISEASE | E75.5 | Other lipid storage disorders | E78.00 | Pure hypercholesterolemia, unspecified |
| | E78.010 | Homozygous familial hypercholesterolemia HoFH | E78.011 | Heterozygous familial hypercholesterolemia HeFH |
| | E78.019 | Familial hypercholesterolemia, unspecified | E78.1 | Pure hyperglyceridemia |
| | E78.2 | Mixed hyperlipidemia | E78.3 | Hyperchylomicronemia |
| | E78.41 | Elevated Lipoprotein(a) | E78.49 | Other hyperlipidemia |
| | E78.5 | Hyperlipidemia, unspecified | H18.411 | Arcus senilis, right eye |
| | H18.412 | Arcus senilis, left eye | H18.413 | Arcus senilis, bilateral eyes |
| | I20.0 | Unstable angina | I25.10 | Atherosclerotic heart disease of native coronary artery without angina pectoris |
| | I25.110 | Atherosclerotic heart disease of native coronary artery with unstable angina pectoris | I25.700 | Atherosclerosis of coronary artery bypass graft(s), unspecified, with unstable angina pectoris |
| | I25.710 | Atherosclerosis of autologous vein coronary artery bypass graft(s) with unstable angina pectoris | I25.720 | Atherosclerosis of autologous artery coronary artery bypass graft(s) with unstable angina pectoris |
| | I25.730 | Atherosclerosis of nonautogous biological coronary artery bypass graft(s) with unstable angina pectoris | I25.750 | Atherosclerosis of native coronary artery of transplanted heart with unstable angina pectoris |
| | I25.760 | Atherosclerosis of bypass graft of coronary artery of transplanted heart with unstable angina pectoris | I25.790 | Atherosclerosis of other coronary artery bypass graft(s) with unstable angina pectoris |
| | T46.6X5A | Adverse effect of antihyperlipidemic and antiarteriosclerotic drugs (statin intolerance) | | |
| OTHER | G71.00 | Muscular dystrophy, unspecified | G71.01 | Duchenne or Becker muscular dystrophy |
| | G71.02 | Facioscapulohumeral muscular dystrophy | I10 | Essential (primary) hypertension |
| | I34.0 | Mitral valve prolapse | I46.9 | Cardiac arrest, cause unspecified |
| | Q23.0 | Aortic stenosis, congenital | Q20.9 | Congenital heart disease (NOS) |
| | Q66.00 | Congenital talipes equinovarus, unspecified foot | Q66.01 | Congenital talipes equinovarus, right foot |
| | Q66.02 | Congenital talipes equinovarus, left foot | Q87.89 | Noonan syndrome |
| | R55 | Syncope and collapse | R06.02 | Shortness of breath |
| | Z01.810 | Encounter for preprocedural cardiovascular examination | Z13.6 | Encounter for screening for cardiovascular conditions |
| | Z79.01 | Long term (current) use of anticoagulants | Z86.74 | Personal history of sudden cardiac arrest |
| | Z95.0 | Presence of cardiac pacemaker | Z95.2 | Presence of prosthetic heart valve |
| FAMILY HISTORY | Z14.8 | Genetic carrier of other disease | Z15.89 | Genetic susceptibility to other disease |
| | Z82.41 | Family history of sudden cardiac death | Z82.49 | Family history of ischemic heart disease and other diseases of the circulatory system |
| | Z82.69 | Family history of other diseases of the musculoskeletal system and connective tissue | Z82.79 | Family history of other congenital malformations, deformations and chromosomal abnormalities |
| | Z83.42 | Family history of familial hypercholesterolemia | Z84.81 | Family history of carrier of genetic disease |
| | Z84.89 | Family history of other specified conditions | | |

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