

## Cardiovascular Genetics ICD-10 Code Reference Sheet\*

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	I47.2	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
	Q23.8	Brugada syndrome	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	I71.01	Dissection of abdominal aorta
	I71.1	Thoracic aortic aneurysm, ruptured	I71.2	Thoracic aortic aneurysm, without rupture
	I71.9	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
	Q66.0	Congenital Talipes Equinovarus ("club foot")	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)			

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CHOLESTEROL/CORONARY ARTERY DISEASE	E75.5	Other lipid storage disorders	E78.0	Pure hypercholesterolemia
	E78.01	Familial hypercholesterolemia	E78.1	Pure hyperglyceridemia
	E78.2	Mixed hyperlipidemia	E78.3	Hyperchylomicronemia
	E78.4	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	G71.0	Hereditary progressive 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)
	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	Z82.69	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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