

CADScor[®] System Billing and Coding Overview

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The following information is provided for physician coding and billing guidance for the CADScor[®] System which is an advanced acoustic-based diagnostic aid to rule out significant coronary artery disease. For more information, please visit Acarix.com

FDA Clearance

The CADScor[®] System is an FDA De Novo cleared device (DEN190047).

Intended Use

The intended use of the CADScor[®] System is to record heart sounds, i.e., murmurs and vibration, for calculation of a patient specific score, indicating the risk of presence of coronary stenosis, as an aid in cardiac analysis and diagnosis.¹

CPT[®] Code²

CPT[®] codes are reported by physicians for the services they perform. The American Medical Association has assigned Category III CPT code* 0716T for the CADScor[®] System.

CPT Code	Description
0716T	Cardiac acoustic waveform recording with automated analysis and generation of coronary artery disease risk score

*Category III codes are temporary codes for emerging technology, services and procedures that allow for specific data collection. There are no assigned Relative Value Unit's (RVU's) or established payment for these codes.



ICD-10 Diagnosis Code					
Primary Symptoms		Primary Risk Factors		Secondary Risk Factors*	
ICD-10	Description	ICD-10	Description	ICD-10	Description
I20.89	Other forms of angina pectoris	I10	Essential Hypertension	E66.3	Overweight
M79.601	Pain in right arm	E10.____	Type 1 Diabetes	E66.8	Other obesity
M79.602	Pain in left arm	E11.____	Type 2 Diabetes	E83.52	Hypercalcemia
R06.00	Dyspnea, unspecified	E78.0	Pure hypercholesterolemia, unspecified	Z72.0	Tobacco use
R06.02	Shortness of breath	E78.2	Mixed hyperlipidemia	Z82.49	Family history of ischemic heart disease
R06.89	Other abnormalities of breathing			Z82.40	Family History of coronary artery disease
R07.2	Precordial pain				
R07.82	Intercostal pain				
R07.89	Other chest pain				
R07.9	Chest pain, unspecified				
R55	Syncope and collapse				

*Must have primary symptom selected

1. User manual US-FDA v.12.Y

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