

Clinical Policy: Cardiac Biomarker Testing

Reference Number: LA.CP.MP.156

Date of Last Revision: 12/24

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Description

The release of cardiac biomarkers is among the cascade of events that occur during acute coronary syndromes and cardiac ischemia.¹ This policy discusses the medical necessity requirements for testing of these cardiac biomarkers.

Policy/Criteria

- I. It is the policy of Louisiana Healthcare Connections that troponin I or T testing is **medically necessary** and the appropriate cardiac biomarker for evaluating for suspected acute myocardial infarctions (AMI) or myocardial injury due to other mechanisms.

- II. It is the policy of Louisiana Healthcare Connections that creatine kinase myocardial isoenzyme (CK-MB) and myoglobin testing are **not medically necessary** in the evaluation for suspected AMI because troponin is the recommended biomarker due to its superior sensitivity and accuracy.

Background

Detection of specific cardiac biomarkers in blood serum is a useful clinical indication of acute myocardial infarctions (AMI), myocarditis, or heart failure.² Cardiac troponins I and T have become the preferred biomarkers used for diagnoses of acute coronary syndromes due to their high specificity and sensitivity and because these subunits are expressed in the myocardium.^{1,2,3,4,5} Furthermore, troponin levels are also elevated for acute and chronic decompensated heart failure in instances of myocyte injury and/or necrosis.⁶

Other cardiac peptides that were previously assessed for AMI include creatine kinase myocardial isoenzyme (CK-MB) and myoglobin.¹ However, recent evidence suggests that the sensitivity and specificity of these biomarkers are inferior compared to the troponins, suggesting that troponins are a more accurate biomarker of myocardial injury.^{1,2} According to the 2014 American College of Cardiologists/American Heart Association (ACC/AHA) clinical practice guidelines, CK-MB and myoglobin are no longer necessary for acute coronary syndrome diagnosis as a result of the advent of troponin assays.² CK-MB detection is comparatively less sensitive and less specific.^{1,2,3,4,5} A 2010 retrospective cohort study was performed in an emergency department over a 12 month period examining patients who had troponin testing.⁷ The study included 11,092 visits where at least one troponin test was ordered, and 97.9% of these patients also had a CK-MB ordered.⁷ The authors concluded that CK-MB testing can be omitted during the initial screening of AMIs since the study showed a 0% rate of positive CK-MB index with negative troponin.⁷ Eggers et al. evaluated the role of myoglobin with troponin I to detect AMI in a sample of 197 patients and determined that neither myoglobin nor CK-MB added clinical diagnostic value.⁸ Of note, Singh et al. measured CK-MB testing from 2007 to 2013 and found a dramatic decrease from 12,057 tests in 2007 to 36 tests in 2013.⁹

Coding Implications

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Table 1: CPT codes not medically necessary when billed with CPT 84484 Troponin

CPT Codes	Description
82553	Creatine kinase (CK), (CPK); MB fraction only
83874	Myoglobin

Reviews, Revisions, and Approvals	Revision Date	Approval Date	Effective Date
Converted corporate to local policy.	08/15/20		
Added “or myocardial injury due to other mechanisms” in addition to acute myocardial infarction for approval in criteria I. References reviewed and updated. Coding reviewed. Changed “review date” in the header to “date of last revision” and “date” in the revision log header to “revision date.” Added “and may not support medical necessity” to coding implications. Reviewed by specialist.	2/22	4/14/22	
Annual review. Background updated with no impact on criteria. References reviewed and updated.	10/22	1/14/23	
Annual review. Background updated with no impact on criteria. Coding reviewed. References reviewed and updated. Reviewed by external specialist.	10/23	1/5/23	
Annual review. References reviewed and updated.	12/24	1/27/25	2/27/25

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Important Reminder

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. LHCC makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved.

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CLINICAL POLICY

Cardiac Biomarker Testing for Acute Myocardial Infarction



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