Clinical Classification of Myocardial Infarction:

<table>
<thead>
<tr>
<th>Type</th>
<th>Description</th>
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<tr>
<td>Type 1</td>
<td>MI caused by acute atherothrombotic coronary artery disease and usually precipitated by atherosclerotic plaque disruption.</td>
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<td>Type 2</td>
<td>MI consequent to a mismatch between oxygen supply and demand.</td>
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<td>Type 3</td>
<td>Patients with the typical presentation of myocardial ischemia with unexpected death before samples for biomarkers could be drawn or before the appearance in the blood.</td>
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<td>Type 4a</td>
<td>MI associated with percutaneous coronary intervention.</td>
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<td>Type 4b</td>
<td>A subcategory of PCI-related MI due to stent thrombosis.</td>
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<tr>
<td>Type 5</td>
<td>Myocardial Infarction associated with cardiac surgery.</td>
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Presentation: The most common symptom of STEMI is chest discomfort. In fact, chest pain and diaphoresis have the highest LR and PPV for STEMI. Chest discomfort is typically retrosternal but may also present as left parasternal, left precordial or across the anterior chest. Patients describe it as heavy, pressing, crushing, tightening, aching, or burning. Sites of radiation can include the left arm, right arm, shoulder, neck, jaw, teeth, epigastrium, and interscapular areas. Although the classic presentation of chest pressure with radiation to the jaw, left arm, or both arms is 96% specific, it is only about 11% sensitive. Do not forget the atypical presentation of ACS. In nearly 20% of cases, MI’s are either painless (silent) or atypical. Elderly patients, women, and diabetics are particularly prone to atypical MI’s. Atypical MI presents with pain predominantly in the neck, jaw, ear, arm, or epigastrium. Other signs to look for STEMI: sudden dyspnea, weakness, diaphoresis, lightheadedness, hypotension, syncope, new arrhythmias, nausea, and vomiting.

In your initial evaluation, look for evidence of systemic hypoperfusion (hypotension, tachycardia, altered mental status, clammy/pale skin). This is concerning for cardiogenic shock associated with MI (5-10% of cases). For more details on cardiogenic shock and its management, see our other board bomb here.

Volume overload is not an expected finding as the onset is rapid. Rather, look for dyspnea, hypoxia, rales/wheezing. However, pulmonary congestion is an uncommon finding, only seen in <30% of patients.

TIMI score is a specific scoring system designed to determine risk of in-hospital mortality. But honestly, this is rarely used in the ED as the speed at which STEMI patients are addressed and given reperfusion therapy means they should (optimistically), be out of the ED no sooner than 45 minutes or less.

Diagnosis: Electrocardiogram and clinical symptoms are all that is needed to diagnose a STEMI. Biomarkers may be normal early on.

ECG Criteria: Remember the goal to get an ECG within 10 minutes of first encounter.

- ≥1mm ST-segment elevation in two contiguous chest or limb leads
Tombstone: STEMI & Acute Coronary Syndrome

- ≥2mm ST-segment elevation in men ≥40 years in leads V2-V3, ≥2.5mm in men <40 years in leads V2-V3, and ≥1.5mm ST-segment elevation in women in lead V2-V3

Localization of ischemia can be determined by what ECG leads show ST- elevation.

- Anterior wall ischemia: two or more of the precordial leads (V1-V6); LAD V1-V4
- Anteroseptal ischemia: Lead V1-V4; Proximal LAD
- Apical or lateral ischemia: Leads aVL and I, V4 to V6; Distal/Diagonal LAD or LCX
- Inferior wall ischemia: Lead II, II, and aVF; 90% RCA, 10% LCX

Don’t forget about STEMI mimickers such as pericarditis (look for diffuse ST elevation without reciprocal T wave changes), myocarditis, hyperkalemia, LBBB, Brugada, Osborne waves in hypothermia, and early repolarization (look for J point notching). When reading EKGs for STE and STD, look at the J point. The J point is used to determine the magnitude of the ST segment elevation, and is compared to the isoelectric part of the tracing. In addition, AV block has been found in about 7% of cases and can result in bradycardia and low cardiac output. Have your pacer pads ready!

Speaking of LBBB, remember that it can obscure the ST-segment analysis and complicate ECG interpretation. The presence of a new LBBB is not a STEMI equivalent, but when there is a new LBBB and other clinical and laboratory findings suggestive of an acute MI, it is associated with a high mortality. How can we tell? Thanks to the Sgarbossa criteria that was developed in the late 90s, a score of ≥3 has specificity of 98% but a sensitivity of 20%, increasing to a specificity of 100% and a sensitivity of 14% with score ≥5. There is now a modified Sgarbossa rule which highlights the criteria yielding only 2 points (seen in the middle of the photo on the right). Smith et al found an ST/S ratio of < -0.25 as a replacement for the absolute ST elevation of ≥5 mm. The most concerning finding is discordant STE. A simple way to apply the Sgarbossa is to look at each lead on the EKG for concordance. Each lead in a LBBB should be discordant or isoelectric. If there is concordance, stop and evaluate for Sgarbossa further.

Unique cases

STE in aVR is associated with >1 mm STD in other multiple leads may suggest LMCA stenosis or occlusion. However, more and more recently this has come under fire. It seems more commonly aVR elevation is due to subendocardial ischemia or evidence of triple vessel disease. In fact, tachycardia-related depression can even cause STE in lead aVR.

STE in aVR is not specific for LMCA, and the patient’s clinical condition should be taken into consideration.

Isolated Posterior MIs are rare: 3% of total MIs. More commonly, they are seen as an extension of an anterior infarction into much larger area involving the posterior. These will present as ST depression in leads V1-3 with a positive T wave. Get a posterior EKG (V7-9). This is easy. Move leads V4-6 to their mirror position on the back. ST elevation >0.5 mm in any posterior lead is diagnostic.

RV infarction is seen in up to 40% of inferior STEMIs. On a typical 12-lead, suspect when there is an inferior STEMI with ST elevation in V1 or V1 > V2, ST elevation in Lead III > II, or ST elevation in V1 with ST depression in V2. The latter is the most specific for RV infarct. Confirm RV infarction by performing a “Right sided EKG”. Place V1-V6 in a mirror image pattern on the R chest. These patients are especially preload sensitive! These are the people you are avoiding giving Nitrates. Treat hypotension with IV fluids.

We cannot emphasize serial EKGs enough. We recommend getting one every 15-30 minutes for the first 1-2 hours as indicated, or anytime the patient’s pain changes. One report showed 8% of all STEMI were identified on repeat EKG.

What about troponins?

Elevations in cardiac troponin is usually present in those presenting with a STEMI, however, if patients present soon enough (under an hour since symptom onset), the initial troponin can be negative. An elevation in cTnI above the 99th percentile of normal has is both sensitive and specific for myocardial damage. Troponins are detectable from 1 to 4 hours after the onset of an acute MI. They are maximally sensitive at 8 to 12 hours and peak at 10 to 24 hours.

Never wait for a positive laboratory test to proceed with a percutaneous coronary intervention (PCI) if ECG and clinical symptoms are highly suspicious for STEMI.

Imaging: In most places, a chest x-ray can be performed quickly not delaying transporting to the cath lab. If it delays transport, omit it. Yes, you should think about aortic dissection, but an abnormal CXR is only present in about 68% of cases so make sure you know that if you are highly suspicious, get a CTA chest. A bedside echo can help as studies in patients presenting to the ED with chest pain have found it to be very sensitive with regional wall motion abnormalities present in about 93% of cases with AMI, they are also found in 43% of those without AMI.

Check out our next handout on STEMI management as we wrap up this high yield topic here!

References: for a complete list of references please visit our website under this topic’s heading. We do not want to waste paper printing them! (We know your hospital is probably responsible for wasting an equivalent of an acre from the rain forest daily).