

Chest Pain • Evaluation & Diagnosis

Acute & stable chest pain: triage, risk stratification & testing pathway · point-of-care reference for internal medicine

- Class I – recommended ● IIa – reasonable ● IIb – may consider ● III: Harm / No benefit – do not use

1 First: don't miss the life-threatening causes

△ Immediately life-threatening → act now

Five emergencies drive the initial workup: **STEMI** (ECG within 10 min of arrival **I**) · **NSTE-ACS** · **acute aortic syndrome** (dissection / intramural haematoma) · **pulmonary embolism** · **oesophageal rupture** / **tension pneumothorax**. Also consider tamponade, fulminant myopericarditis & sickle-cell chest crisis. Get a 12-lead ECG & **high-sensitivity troponin** early; transport by EMS, not private vehicle **I**.

△ Chest pain ≠ pain in the chest – read the pattern

Anginal equivalents include pressure / tightness / heaviness in chest, shoulders, arms, neck, back, upper abdomen or jaw, plus dyspnoea & fatigue. **Favours ischaemia**: central, exertional, retrosternal, pressure/squeezing. **Lowers probability**: pleuritic, positional, sharp/fleeting, reproduced by palpation. Drop the word **"atypical"** – describe pain as **cardiac, possibly cardiac, or noncardiac** **I**.

2 Initial evaluation – every patient

History + focused exam + 12-lead ECG + cTn **I**

Characterise nature, onset/duration, location/radiation, provoking & relieving factors, associated symptoms; assess CV risk factors. ECG within **10 min** of arrival **I**; serial ECGs if nondiagnostic & suspicion high **I**. Add leads V7–V9 to exclude posterior MI **IIa**. Chest radiograph for alternative causes **I**.

▼ is the ECG diagnostic?

STEMI / ST-depression / new LBBB

→ follow STEMI / NSTE-ACS guidelines **I**.

Nondiagnostic / normal ECG

→ serial cTn & risk-stratify (a normal ECG does not exclude ACS).

Biomarkers – high-sensitivity troponin preferred **I**

hs-cTn enables faster rule-in / rule-out & higher accuracy than conventional assays **I**. Repeat at **1–3 h** (hs-cTn) or **3–6 h** (conventional) after time-zero **I**. A single hs-cTn below the limit of detection can exclude injury if symptoms began ≥3 h before arrival **IIa**. Know your assay's 99th-percentile URL **I**. **CK-MB & myoglobin add nothing** once cTn is available **III: No benefit**.

Implement a **clinical decision pathway (CDP)** to stratify low / intermediate / high risk **I** – HEART, EDACS, ADAPT, NOTR, ESC 0/1-h.

3 Risk strata & ischaemic vs nonischaemic clues

FAVOURS ISCHAEMIA / HIGHER RISK

- Central, retrosternal, pressure, squeezing, heaviness
- Provoked by exertion / emotional stress; relieved by rest
- Builds gradually over minutes; radiates to arm, neck, jaw
- Dyspnoea, diaphoresis, nausea; diabetes, women, elderly
- Ischaemic ECG changes · troponin-confirmed injury · EF < 40%

FAVOURS NONCARDIAC / LOWER RISK

- Sharp, fleeting (seconds), pleuritic or positional
- Reproduced by palpation (costochondral tenderness)
- Localised to a small area; radiates below umbilicus/hip
- Ripping pain to back → suspect aortic syndrome instead
- Normal serial ECG + non-elevated hs-cTn → low risk

CDP risk tiers → disposition

Low (<1% 30-day death/MACE): discharge without admission or urgent testing **IIa**; outpatient CAC optional.

Intermediate: bedside TTE **I**; observation-unit care **IIa**; further testing per known/unknown CAD.

High (ischaemic ECG, +troponin, EF < 40%, moderate-severe ischaemia, instability, high CDP score): **ICA** **I**.

4 Additional testing – guided by the clue (after Hx / exam / ECG / cTn)

SCENARIO	FIRST-LINE	FURTHER / ADD-ON OPTIONS
Intermediate-risk, no known CAD	CCTA to exclude plaque / obstructive CAD I , or stress imaging (echo · PET/SPECT MPI · CMR) or exercise ECG I	FFR-CT for 40–90% proximal/mid stenosis IIa ; stress imaging if CCTA inconclusive IIa ; ICA if moderate-severe ischaemia I
Intermediate-risk, known CAD	Optimise GDMT before more testing I ; stress imaging for new/worse symptoms IIa	ICA for high-risk CAD / worsening symptoms I ; CCTA for nonobstructive progression IIa ; FFR-CT IIa
Suspected acute aortic syndrome	CT angiography of chest/abdomen/pelvis I	TEE or CMR if CT contraindicated/unavailable I ; TTE may show effusion / AR / flap
Suspected pulmonary embolism	CTA with PE protocol in stable patients I	Pretest probability + D-dimer guides imaging I ; V/Q as second-line · echo for RV strain
Suspected myopericarditis	CMR with gadolinium to distinguish from MINOCA I	TTE for wall-motion / effusion I ; cardiac CT for pericardial thickening IIb
Suspected valvular disease	TTE for presence, severity & cause of VHD I	TEE (3D) if TTE inadequate I ; CMR if echo nondiagnostic IIa
Stable, no known CAD	Pretest-probability model to find low-risk I ; CCTA for intermediate-high risk I	CAC or exercise ECG if low risk IIa ; stress imaging I ; FFR-CT for 40–90% stenosis IIa

Don't routinely test in low-risk patients: for acute or stable chest pain deemed low risk, urgent/early stress imaging or anatomic testing is *not* needed **III**. Honour warranty periods – normal CCTA (no plaque/stenosis) 2 y; normal stress test 1 y. CK-MB / myoglobin added to cTn **III**; delayed transfer from office for testing **III: Harm**.

5 Pathway-specific & special populations

High-risk acute chest pain – stepwise

- Designate high risk: new ischaemic ECG, +troponin, EF < 40%, moderate-severe ischaemia, instability, high CDP score **I**
- Invasive coronary angiography for the full extent & severity of obstructive CAD **I**
- If troponin+ but CCTA/ICA show no obstruction → CMR or echo for MINOCA / myocarditis **IIa**

Known CAD with stable chest pain

- Optimise GDMT first; defer testing where possible **I**
- Moderate-severe ischaemia despite GDMT → ICA to guide therapy **I**
- Stress PET/SPECT, CMR or echo for diagnosis & risk **I**; add MBFR on PET/CMR **I**

Special populations – one-liners

- Prior CABG, no ACS** → stress imaging or CCTA for graft patency **I**; ICA if nondiagnostic **I**
- On dialysis, unremitting pain** → EMS transfer to acute care **I**
- Cocaine / methamphetamine** → consider as cause; standard risk stratification **IIa**
- Sickle cell disease** → EMS transfer **I**; exclude ACS / acute chest syndrome **I**
- Recurrent, workup negative** → evaluate GI causes **IIa**; refer to CBT for anxiety/panic **IIa**
- INOCA** → invasive coronary function testing **IIa**; stress PET/CMR with MBFR **IIa**

6 Shared decisions & cost-value

CHOOSING THE RIGHT TEST (FIGURE 6)

CCTA favoured	age < 65 · rule out obstructive CAD
Stress imaging favoured	age ≥ 65 · suspect scar/ischaemia
PET over SPECT (if available)	↑ accuracy · MBFR
Lowest-cost test	exercise ECG / CAC
Pregnant / child-bearing age	minimise ionising radiation

SHARED DECISION-MAKING

Low-risk acute chest pain → patient decision aids (e.g. Chest Pain Choice) improve understanding & risk communication **I**. Intermediate-risk → shared decision-making on admit vs observe vs outpatient follow-up reduces low-value testing **I**.

KEY CAUTIONS

Relief with nitroglycerin is **not** diagnostic of ischaemia. Up to 6% of evolving ACS have a normal initial ECG. "Known CAD" now includes nonobstructive plaque – optimise prevention, don't overlook it. High-risk CAD = left main ≥50% or 3-vessel ≥70%.

Remember

The chest-pain trap is **over-testing the low-risk & under-recognising the dangerous**. Get an ECG within **10 minutes** and **hs-cTn** early on everyone; a normal ECG does not exclude ACS. Use a **structured CDP** to stratify risk – low-risk patients (<1% 30-day MACE) need **no urgent testing** and can go home. Reserve CCTA, stress imaging & ICA for intermediate-to-high pretest probability. Say **cardiac / possibly cardiac / noncardiac** – never "atypical".