

## 2020 ESC Guidelines for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation

**The Task Force for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation of the European Society of Cardiology (ESC)**

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These algorithms should always be integrated with a detailed clinical assessment and 12-lead ECG, and repeat blood sampling is mandatory in case of ongoing or recurrent chest pain.

The same concept applies to the 0 h/2 h algorithm. Cut-off levels are assay-specific and shown in Table 5. Cut-off levels for other hs-cTn assays are in development.

### 3.3.4 Observe

Patients who do not qualify for 'rule-out' or 'rule-in', are assigned to observe. They represent a heterogeneous group that usually requires a third measurement of cardiac troponin at 3 h and echocardiography as the next steps.<sup>85</sup> ICA should be considered in patients for whom there is a high degree of clinical suspicion of NSTEMI-ACS (e.g. relevant increase in cardiac troponin from presentation to 3 h), while in patients with low-to-intermediate likelihood for this condition according to clinical judgment, non-invasive imaging using CCTA or stress testing [stress echocardiography, positron emission tomography, single-photon-emission tomography (SPECT), or CMR for the detection of ACS features (oedema, late gadolinium enhancement, perfusion defect, etc.)] should be considered after discharge from the emergency department to the ward. No further diagnostic testing is indicated when alternative conditions, such as rapid ventricular rate response to

- ii. The ESC 0 h/1h and 0 h/2 h algorithms apply to all patients irrespective of chest pain onset. The safety (as quantified by the NPV) and sensitivity are very high (>99%), including in the subgroup of patients presenting very early (e.g. <2 h).<sup>69</sup> However, due to the time dependency of troponin release and the only moderate number of patients presenting <1 h after chest pain onset in previous studies, obtaining an additional cardiac troponin concentration at 3 h in patients presenting <1 h and triaged towards rule-out should be considered.
- iii. As late increases in cardiac troponin have been described in ~1% of patients, serial cardiac troponin testing should be pursued if the clinical suspicion remains high or whenever the patient develops recurrent chest pain.<sup>35,36,39,68,69,75,76,86</sup>

3.3.4.2 *Confounders of cardiac troponin concentration.* In patients presenting with suspected NSTEMI-ACS, beyond the presence or absence of MI, four clinical variables affect hs-cTn concentrations:<sup>35,36,39,69,79,87–93</sup>

- i. Age (to a large extent as a surrogate for pre-existing cardiac disease).
- ii. Renal dysfunction (to a large extent as a surrogate for pre-existing cardiac disease).
- iii. Time from chest pain onset.
- iv. Sex.

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