Aortic intramural haematoma (IMH) originates from spontaneous rupture of the aorta vasavasorum or from a penetrating atherosclerotic ulcer. Recent advances in imaging techniques have significantly aided its diagnosis, placing the incidence between 10–30% of acute aortic syndromes. Clinical presentation may be identical to that of classical dissection. IMH affects patients with more atherosclerotic risk factors and older age than aortic dissection, and it is located in the descending aorta in 60–70% of cases.

**IMAGING TECHNIQUE INFORMATION**

Although the diagnostic sensitivity seems to be similar among computed tomography (CT), magnetic resonance imaging (MRI), and transoesophageal echocardiography (TOE), the diagnosis of IMH is more complex than that of classical dissection, with at least two diagnostic techniques being performed in most cases. In the International Registry of Aortic Dissection, the CT–TOE combination was the most used. Nevertheless, MRI has the advantage of permitting detection of new bleeds in the aorta wall, which may be important in subacute and chronic phases.

Imaging techniques provide significant prognostic information. Concurring with data reported by others,1–3 maximum aorta diameter in our series of 64 patients was an important predictor of adverse clinical events including death.4 Fluid extravasation, pericardial and pleural effusion, and mediastinal haemorrhage, more frequent in IMH than in classical dissection, worsen the prognosis. However, progression with small entry tears, are frequently detected with ulcer-like images. These images, secondaries to focal dissection, can be very useful for differentiating penetrating aortic ulcer from IMH. Careful follow up imaging study during acute and subacute phases is absolutely necessary. Transcathectomy echocardiography was performed daily and TOE within three days after the diagnosis and once a week until the third week. CT was repeated once a week. In this series, 43% of type A IMH required surgery. With this strategy, it is essential that diagnostic imaging modalities and urgent surgical intervention are readily available. The subgroup of patients at low risk for complications should be defined before treatment, other than surgery, is indicated, particularly in younger patients with low comorbidity.

Type B IMH should be treated with β blockers, closely followed up with imaging techniques, until complication-free absorption of the IMH is observed. The subgroup with progressive aortic dilatation or with images of localised (ulcer-like) dissection should be closely followed and treated more aggressively. Endovascular placement of stent grafts to cover some extent of IMH appears to be a promising treatment for this type of complication; however, it remains to be confirmed by results of large clinical series.

**REFERENCES**


**Abbreviations:** CT, computed tomography; IMH, intramural haematoma; MRI, magnetic resonance imaging; TOE, transoesophageal echocardiography
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Hypertrophic cardiomyopathy with persistent ST segment elevation simulating acute myocardial infarction

On May 2000, a 67 year old woman was admitted to the hospital because of right upper eyelid ptosis. She was normotensive and had never experienced chest pain, dyspnoea or other symptoms suggestive of heart disease. A routine 12 lead ECG (upper panel, A) revealed an ST segment elevation with associated negative T waves in several leads, suggesting a possible acute myocardial infarction. Physical examination did not reveal any abnormal heart finding, and cardiac enzymes were negative. Echocardiography (lower panels A and B: LA, left atrium; LV, left ventricle; RA, right atrium; RV, right ventricle) revealed a pronounced and generalised left ventricular hypertrophy with normal wall motion and absence of gradient across the left ventricular outflow tract. Several ECGs obtained in the following days were unchanged. The coronary angiogram showed normal coronary arteries. Accordingly, the apparent acute myocardial infarction pattern was likely to be dependent on hypertrophic cardiomyopathy. The patient was followed up and no changes in the ECG pattern were observed in the ensuing three years (upper panel, B).

This case shows that hypertrophic cardiomyopathy can be associated with an ECG falsely suggesting an acute myocardial infarction.

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