CASE REPORT

Swallow syncope associated with complete atrioventricular block and vasovagal syncope

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Abstract
A 21 year old man was referred to our hospital for evaluation of multiple, recurrent episodes of syncope associated with swallowing. He noted an augmentation and subsequent attenuation of symptoms during and after swallowing, which resolved as he stopped swallowing. Multiple episodes of symptomatic and asymptomatic paroxysmal atrioventricular (AV) block without escape beats were recorded repeatedly during meals or while drinking water on Holter ECG. He achieved a Bruce stage 5 during a treadmill exercise test and his heart rate reached 178 beats per minute without prolongation of the PQ interval. Sinus bradycardia was induced by carotid sinus massage, Valsalva manoeuvre, and gag reflex, but AV block was not induced by any of these.

Figure 1 shows complete AV block, of which the maximum RR interval was three seconds, and was induced by swallowing large amounts of carbonated beverages. Intravenous atropine at a dose of 2.5 mg completely abolished the AV block, but the effect of isoprenaline was incomplete. On the other hand, 0.2 mg of propranolol worsened the AV block. Oral atropine was discontinued due to the patient’s complaint of thirstiness.

Figure 2 shows the results of the head up tilt test. Tilting to 80° caused an initial increase in heart rate without a significant drop in blood pressure. Then, systolic blood pressure decreased slowly with rhythmic fluctuations, finally leading to syncope. Shortly before the onset of syncope, the heart rate gradually decreased. The spectral indices of heart rate variability (HRV) using fast Fourier analysis revealed that the ratio of the low frequency spectrum (LF) to the high frequency spectrum (HF) increased immediately after the head up tilt while the HF decreased. Finally, the HF increased and the LF/HF decreased starting from just one minute before the onset of syncope. Oral disopyramide and intravenous atropine were not effective in inhibiting the syncope induced by the head up tilt test.

Keywords: swallow syncope; AV block; electrophysiological study; head up tilt test

Figure 1 ECG during swallowing carbonated beverage. (A) Swallowing is followed by three seconds of complete AV block. (B) After the intravenous administration of 2 mg of atropine, the AV block is completely blocked.
An electrophysiological study demonstrated normal AH and HV intervals (75 and 50 ms, respectively). Incremental atrial pacing revealed the AV nodal decremental conduction property at a pacing rate of 150 beats per minute. Atrial extrastimulus testing revealed normal AV nodal function and effective refractory periods. Swallowing of carbonated beverages and inflation of a rubber balloon situated in the lower oesophagus to a varying degree of inflation volumes produced paroxysmal AV block (fig 3), which was shown to be an AH block by the His bundle ECG study. The cardiac catheterisation study, left ventriculography, and coronary angiography were normal.

**Discussion**

The neurally mediated syncopal syndromes encompass a number of apparently related disturbances of reflex cardiovascular control characterised by transient inappropriate bradycardia and/or vasodilatation of the vascular bed. Although vasovagal syncope occurs very frequently, swallow or deglutition syncope is an unusual condition whereby temporary loss of consciousness occurs with swallowing.

We present a case of swallow syncope associated with complete AV block and vasovagal syncope. Although about 30 patients with swallow syncope have been reported in the literature, this is the first case that presented with both swallow syncope and vasovagal syncope.

It has been shown that most patients with swallow syncope have a variety of disorders, such as myocardial infarction, rheumatic carditis, digoxin toxicity, and gastro-oesophageal disorders (for example, hiatus hernia, gastric and oesophageal diverticulae, achalasia, spasm, and carcinoma of the oesophagus). A minority of patients are found to have no cause. Recent studies suggest that cardioinhibitory control neurons lie in an area between the nucleus ambiguus and the ventromedial aspect of the trigeminal spinal nucleus. Sensory fibres lie in fascicles in close proximity to each other as they pass the jugular ganglion and then disperse evenly within the nerve after passing the nodose ganglion. Vagal fibres innervate primarily the SA node, atrial musculature, and AV node, inhibiting AV conduction without any direct effect on the ventricular conduction system or myocardial contraction. Physiologically, swallow syncope is produced by a vagovagal reflex. With balloon inflation at the lower oesophagus, the afferent barrage is from distension of the oesophagus. Further from the results of the atropine administration, at least some of the necessary synapses in the arc were central and could be interrupted by atropine.

Head up tilt test suggests that this patient has a disturbance of the system described above. The normal compensatory response to head up posture is reflex tachycardia, more forceful contraction of the left ventricle, and vasoconstriction. However, in individuals with neurally mediated syncope, this forceful ventricular contraction in the setting of a relatively empty cavity may activate the cardiac mechanoreceptors. An afferent pathway consisting of unmyelinated left ventricular vagal C fibres transmits signals to specific central nervous system sites. This reduces the efferent sympathetic tone and results in reflex hypotension and/or bradycardia. Hayano et al reported that the HF amplitude of the heart rate rapidly decayed with head up tilt and increased quickly with tilt back to the supine position. It has been shown that the LF amplitude of the blood pressure and heart rate shows rhythmic fluctuations at an interval of 48 to 100 seconds during head up tilt. This patient showed a rhythmic fluctuation in the LF amplitude of the heart rate. Furthermore, the LF/HF ratio was greater and the HF amplitude was smaller than those in normal subjects immediately after head up tilt. These findings mean that the first amplitude of the continual fluctuations in response to a variety of stimuli and derangement from both intrinsic and extrinsic environments in this
patient is greater than that in normal subjects. Thus, a disturbance in the autonomic feedback loop of the cardiovascular reflex led to the observed neurally mediated syncope and swallow syncope in this patient.

Swallow syncope has been successfully treated with anticholinergic agents, surgical denervation of portions of the oesophagus, and demand mode transvenous pacemaker placement. Planned treatment for our patient was permanent pacemaker implantation, with β-blockers.


IMAGES IN CARDIOLOGY

Giant coronary sinus and absent right superior vena cava

This 66 year old woman underwent investigation for atypical chest pain. Chest radiography showed cardiomegaly, and echocardiography showed a giant coronary sinus 4 cm in diameter and larger than the aortic root and the left atrium, as shown on the parasternal long axis view (top). Intravenous injection of agitated saline from either the left or right brachial vein showed contrast enhancement of the giant coronary sinus. Venous angiography with contrast material injected simultaneously from both the right and left brachial veins showed that the right superior vena cava was absent and the contrast material entered the right atrium via a persistent left superior vena cava and the giant coronary sinus (bottom).

A persistent left superior vena cava is not an uncommon anomaly and usually drains into the right atrium through the coronary sinus which is dilated, measuring about 1 to 2 cm in diameter. In the rare setting of an absent right superior vena cava, this anomaly produces a truly giant coronary sinus because of the severely increased flow into the coronary sinus. Although this malformation can be associated with other cardiac anomalies, it may be an isolated finding as in this case and not associated with significant haemodynamic consequences. Recognition of this anomaly is important to avoid problems in venous catheterisation, pacemaker insertion, and cardiopulmonary bypass. (Ao, aorta; CS, coronary sinus; LA, left atrium; LV, left ventricle; RA, right atrium.)

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