The case of a 71-year-old male patient, with symptoms of dizziness and atypical chest pain and a positive isotopic exercise stress test, is reported. Coronary angiography demonstrated an anomalous origin of the left circumflex coronary artery from right coronary ostium but no obstructive atherosclerotic coronary lesions. The possible relation between the congenital coronary anomaly and the clinical manifestations of the patient is discussed.

Key words: Catheterization. Congenital heart defects. Ischemia. Scintigraphy.
MEHR without angor, electrical changes, and with a good pressor response. The study of myocardial perfusion after the ergometric test and at rest revealed small subsegmental perfusion defects in the apical, inferior, and lower interventricular septal area and positive redistribution to the lower septum and inferior face (Figure 1). These disturbances suggested ischemic changes induced by the test. In view of these results, cardiac catheterization was requested, which revealed right dominance and coronary arteries free of angiographic lesions. The circumflex coronary artery arose from the ostium of the right coronary and had a retroaortic path (Figures 2 and 3); LVF was normal.

At follow-up 3 months later the patient referred occasional dizziness.

DISCUSSION

The incidence of congenital anomalies of the coronary arteries in different series ranges from 0.3% to 8.3%. The origin of the circumflex artery (Cx) in the right coronary sinus (from an ostium common to the right coronary or an independent ostium) or right coronary artery (as a proximal branch of this artery) is the most common anomaly of the origin of the coronary arteries. Thus, Effler in 1970 recommended calling it a «normal variant» rather than an anomaly.

The disposition of the Cx from its anomalous origin is always the same. From its origin the Cx goes backward and to the left, circling the aorta from behind, then passing between the posterior aortic wall and first...
the anterior right atrial wall, then the left atrial wall, until it reaches its location in the left part of the atrioventricular sulcus, where it is covered by the left atrial appendage and has its usual disposition. This anomaly has been and continues to be considered benign. Nevertheless, cases of association with sudden death, AMI, and angina pectoris in the absence of atherosclerotic lesions have been reported. The factor responsible for this pathogenicity could be repeated compression of this vessel by dilation of the aortic root or angling as a result of its retroaortic position, which would compress the coronary ostium into a «slit» that obstructs blood flow. It seems reasonable to carry out tests to detect possible myocardial ischemia before considering a coronary anomaly as benign. One of the tests most often used is the exercise stress test with thallium. However, for Piovessana, et al. and Molajo, et al. this test is not sensitive enough, as they have demonstrated in published reports of patients with Cx anomalies and positive conventional stress tests with negative thallium stress tests. This finding was attributed to a lack of sensitivity of this method in characterizing myocardial perfusion defects in these patients. In fact, Dunn, et al. questioned the sensitivity of the thallium exercise stress test in demonstrating ischemia in the territory irrigated by the Cx.

Our patient did not have clear clinical manifestations of angina. He presented dizziness and chest discomfort that remitted with occasional use of sublingual nitrates. His life was practically normal and in the last month he had only one episode of dizziness without loss of consciousness. We must also consider that his dizziness could be due to cervicoarthrosis, or that the «chest discomfort», which could be caused by ischemia, could precipitate dizziness. We found no arrhythmic cause of dizziness in Holter studies and the tilt-test was negative. A finding suggesting a possible ischemic cause was the positivity of the radionuclide test, although it could be considered a false positive coinciding with the area of perfusion of an anomalous Cx.

In fact, an anomalous Cx artery, paradigm of the «benignity» of coronary anomalies, can sometimes be non-benign. Compression of the retroaortic segment of the Cx, or angling at its origin, could narrow the ostium to a slit and cause ischemia.

In this case we decided not to take an aggressive therapeutic approach for the moment. Taylor, et al has studied the main features of patients with coronary anomalies that cause sudden death. Age under 35 years and an interarterial path were the two factors most commonly related with this fatal outcome. The age of our patient and the fact that his clinical manifestations did not interfere with a normal life motivated us to use a conservative approach with anxiolytic drugs and sublingual nitrates.

REFERENCES