Anatomy of a Dextrocardia Case with Situs Solitus

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Introduction

Dextrocardia is an embryologic malformation characterized by the displacement of the largest axis (base to apex) of the heart to the right side of the chest, with reversion of the apical inclination. This malformation is caused by an anomalous rotation of the primitive heart tube to the left, in which the bulboventricular loop bends to the left, in a specular image of the normal condition, which occurs around the 8th week of embryonic life.¹,²

The incidence of dextrocardia associated with situs inversus in the general population is usually 1:10,000, whereas the one associated with situs solitus is 1:30,000 live births and only 1:900,000 in the adult population.³,⁴ This high incidence difference found in individuals with dextrocardia associated with situs solitus is due to the higher occurrence of associated cardiac and/or extracardiac malformations, such as tracheoesophageal fistula, pulmonary hypoplasia, imperforate anus, spina bifida and Kartagener syndrome.³,⁴

There is great discordance among authors regarding the classification of the several types of dextrocardia.⁵,⁶,⁷ Among the several studies, the one by Van Praagh et al⁶ is worth mentioning, as it describes and classifies in details the different types of dextrocardia. Dextrocardia must be differentiated from the dextroversion, as the latter is not a congenital condition, in which the heart can change sides due to a disease that pushes or pulls the organ into the right antimere, although the heart apex remains toward the left side.⁵,⁷,⁹,¹⁰

Case Report

Anatomical description

Dextrocardia was identified in the corpse of a female child, probably one year of age, which belonged to the Laboratory of Anatomy of the Campus of São José dos Campos - UNESP.

The thorax of the anatomical piece was partially dissected, without the presence of the pericardium and the lungs. The sternum was in its normal position. The inspection of the heart and the vessels of the base of the heart did not disclose possible position alterations caused by the removal of the lungs. The entire heart and vessels of the base were intact and had not been dissected. The heart and its apex were completely turned to the right.

After the removal of the sternum and the dissection of the vessels of the base, it was observed that due to the rotation of the heart, its entire right portion was further displaced to the posterior part and this fact was so accentuated in the right auricle that, with the heart in the observed position, it was not possible to visualize it (Figure 1). This anomalous rotation of the heart made the pulmonary trunk emerge from the right ventricle abnormally, appearing much more superficial and extensive (Figure 1).

During the inspection of the pulmonary trunk, a large ductus arteriosus (Botallo’s ductus) was found (Figure 2). The heart apex was positioned along the heart axis and was turned to the right side, thus demonstrating the dextrocardia.

With the help of a stereoscopic magnifying glass, all afferent and efferent branches of the vessels of the base of the heart were dissected in order to look for anomalies. With the exception of the brachiocephalic trunk, which presented a more accentuated angulation than the normal one, no other variations regarding position or trajectory were observed (Figure 1).

The diaphragm muscle was removed and subsequently, the liver was dissected, which presented normal aspect and morphology. The abdominal cavity was open and the position of the viscera was normal, without any type of alteration, which characterized situs solitus.

After this phase, the heart was then carefully removed from its place and again inspected with a magnifying glass. The auricles and atriums were opened and analyzed and disclosed normal anatomical aspects. The atrioventricular concordance was anatomically normal. An incision was made in each ventricle with the objective of verifying its internal morphology and the emergence of its vessels. The two ventricles presented...
normal internal morphology, atrioventricular valves and vessel emergence. The aorta and the pulmonary trunk presented normal semilunar valves.

**Discussion**

The literature presents a great diversity of types and classifications for dextrocardia cases. This diversity seems to be caused by the difficulty to establish criteria for a perfect classification.

Through simple anatomical observation, it can be determined that there was no change in the heart position after the corpse underwent fixation.

This case is characterized as a dextrocardia case, as the heart apex was completely turned to the right antimere. It cannot be classified as a dextroversion case, as in the latter, the heart is located in the right hemithorax with the apex turned to the left antimere, different from what was observed in the present case (Figure 1).

The study by Van Praagh et al. is the only one that describes two cases that were identical to the case reported here, with more anatomic details. Both cases, as in the present one, constituted necropsies of children younger than one year with a type of dextrocardia classified by the authors as dextrocardia with normally related great vessels. In these cases, as well as in the present case, situs solitus was found, the great vessels were normally related (with the bulboventricular loop to the right) and the absence of ventricular inversion was observed. The two cases described by these authors presented severe pulmonary alterations; as the lungs had been previously removed in the present case, it was not possible to study them.

Calceterra et al., studying several cases of dextrocardia, described one case with the same anatomical variations found in the present case; the authors, however, did not search for extracardiac alterations in the necropsies, thus not allowing a more detailed comparison with the present case.

Normally, dextrocardia is more frequent when associated to situs inversus, with the presence or absence of associated cardiac and/or extracardiac malformations. The case presented here is more uncommon as it was associated to situs solitus and for the fact that it did not present these types of malformation, which allows us to state that this variant of heart position is not necessarily associated to anomalies in atrioventricular communications, in the relations between the heart and its afferent and efferent vessels or even in the heart function.

The diagnosis of dextrocardia can be attained during the physical examination, through an electrocardiogram or imaging examination. Thus, the knowledge of this anomaly and its variants is of crucial importance for physicians due to the risk of atypical angina presentations, its frequent association with other cardiac and/or extracardiac diseases and also, for constituting a malformation of which the abnormalities,
despite their complexity, are currently possible of undergoing surgical correction\textsuperscript{8,9}.

Potential Conflict of Interest

No potential conflict of interest relevant to this article was reported.

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Study Association

This study is not associated with any graduation program.

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