Fetal Heterotaxy with Tricuspid Atresia, Pulmonary Atresia, and Isomerism of the Right Atrial Appendages at 22 Weeks

Julia E. Solomon, MD1 John H. Stock, MD2 Randy R. Richardson, MD3 Norman H. Silverman, MD4

1 Division of Maternal Fetal Medicine, Fetal Diagnostic Center, St. Joseph’s Hospital and Medical Center–Dignity Health, Phoenix, Arizona
2 Arizona Pediatric Cardiology Consultants, Pediatrix Medical Group, Phoenix, Arizona
3 Department of Radiology, St. Joseph’s Hospital and Medical Center–Dignity Health, Phoenix, Arizona
4 Division of Pediatric Cardiology, University of California at San Francisco, San Francisco, California


Case Report

A 31-year-old gravida 1 was seen for routine evaluation of fetal anatomy at 22 weeks. A complex cardiac abnormality was diagnosed. Findings at the time of the detailed exam included levocardia with a normal left-sided stomach bubble, but a midline-appearing liver. An absent right atrioventricular valve connection typical of tricuspid atresia was identified with a patent and normal-sized left atrioventricular (mitral) valve (► Fig. 1A). A large unrestricted ventricular septal defect was noted, communicating to a relatively hypoplastic right ventricle. The atrial septum demonstrated normal bowing of the atrial septum from the right to the left atrium with evidence of flow on color flow Doppler, with a moderate-sized ostium primum atrial septal defect. Atrial morphology was suggestive of bilateral right atrial appendages but inconclusive (► Fig. 1B). There was D-transposition with the aorta arising from the anterior, morphologically right ventricle (► Fig. 1C). The aortic arch was right-sided with mirror image branching. A discrete main pulmonary artery was not identified in the usual mediastinal position; however, retrograde flow was noted in the ductus arteriosus both in two-dimensional imaging and echocardiography.

We report the accurate prenatal diagnosis at 22 weeks gestation of right atrial isomerism in association with tricuspid atresia. Several distinctive sonographic features of isomerism of the right atrial appendages were present in this fetus: complex cardiac abnormality, ventriculoarterial discordance, juxtaposition of the aorta and the inferior vena cava to the right side, pulmonary atresia, and anomalous pulmonary venous return to the morphological right atrium. Tricuspid atresia, which is an extremely rare lesion within heterotaxy spectrum disorders, was present. Postnatal investigations confirmed all prenatally diagnosed abnormalities, with additional findings of pulmonary atresia with discontinuous pulmonary arteries and bilateral arterial ducts, asplenia, and bilateral eparterial bronchi. To our knowledge, tricuspid atresia in the setting of isomerism of the right atrial appendages has not previously been diagnosed or reported prenatally. Because of the complexity of cardiac lesions that may be present in cases of atrial isomerism, these disorders should be considered even if sonographic findings are uncommon or atypical.

Address for correspondence J. E. Solomon, MD, Fetal Diagnostic Center, 1727 W Frye Road, Suite 210, Chandler, AZ 85224 (e-mail: julia.solomon@dignityhealth.org).

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Abstract

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four-dimensional spatiotemporal image correlation imaging, both with superimposed color flow Doppler, consistent with presumptive pulmonary atresia. Bilateral superior venae cavae were also identified, without evidence of a bridging innominate vein. All four pulmonary veins were seen to drain into the right atrium without obstruction (►Fig. 1D). Finally, assessment of the abdominal vessel configuration demonstrated both an aorta and inferior vena cava that were right-sided. Based on the presence of a complex cardiac malformation and the latter two findings of total anomalous pulmonary venous return and juxtaposition of both the aorta and inferior vena cava to the right side, our prenatal diagnosis was that of isomerism of the right atrial appendages.

Formal karyotype testing with microarray was offered to the parents and declined. The pregnancy progressed otherwise uneventfully with normal interval growth and reassuring parameters of fetal well-being. The patient labored spontaneously at 39\(\frac{1}{7}\) weeks; the fetus was delivered vaginally with a birth weight of 3,430 g and Apgar scores of 8 at 1 minute and 9 at 5 minutes. Postnatal prostaglandin infusion was initiated and an echocardiogram was performed, which confirmed of all of the prenatal ultrasound findings (►Fig. 2). The patent atrioventricular valve was confirmed to be a mitral valve morphologically, without evidence of a septal papillary muscle. Ventricular walls were predominately smooth, again consistent with left ventricular morphology. Cardiac catheterizations confirmed a right-sided aorta (►Fig. 3) and
discontinuous pulmonary arteries arising from bilateral arterial ducts.

Both abdominal ultrasonography and computed tomography scan failed to identify a spleen in the usual or any other location. Barium studies did not identify any malrotation abnormalities of the gut. Cardiac computerized tomographic angiography with three-dimensional reconstruction also confirmed the echocardiographic findings (►Fig. 4A). Origin of the pulmonary arteries/ducts was the brachiocephalic trunk for the left side and the inferior aspect of the aortic arch for the right side (►Fig. 4B). Imaging of the bronchial tree was conclusive for bilateral eparterial (right-sided) bronchi (►Fig. 4C), all supporting the prenatal diagnosis of isomerism of the right atrial appendages.

The neonate underwent bilateral ductal-pulmonary artery stenting at 8 days of life. Bilateral bidirectional cavopulmonary shunts were performed at 4 months of age. The child is currently 11 months of age and thriving. There has been no postnatal evidence of associated dysmorphology or other unrelated abnormalities, and no further genetic testing was performed.

Discussion

Visceroatrial heterotaxy syndrome is a complex developmental condition that is frequently associated with serious congenital cardiac disease. This set of disorders is typically characterized by abnormal arrangement of cardiac and visceral organs across a left–right axis, different from situs solitus or complete situs inversus. Historically, heterotaxy has been divided into the subcategories of right or left isomerism based on the appearance of the atrial appendages; isomerism refers to the symmetric development of structures that are normally asymmetric. More recently, the extent of pectinate muscle proximity to the atrioventricular valves has been found to be more accurate in assessing right- or left-sidedness than atrial appendage morphology, but this cannot be defined by pre- or postnatal echocardiography. Isomerism of the right atrial appendages has classically been associated with asplenia and isomerism of the left atrial appendages with polysplenia. However, spleen status may be discordant in either right or left isomerism, and splenic anatomy is less well correlated with atrial appendage arrangement than bronchopulmonary anatomy, which is highly consistent.

Both right and left atrial isomerism are associated with complex congenital cardiac defects in addition to situs abnormalities of thoracic and abdominal organs. With the exception of stomach bubble and gall bladder laterality, abdominal visceral situs can be difficult to assess in a fetus.
In a small case series, color flow Doppler identification of a splenic artery has been successful in documenting a normally located spleen but can represent a false-negative in cases of multiple small spleens. In the absence of pleural effusions, lung lobation can be extremely hard to assess. Because the bronchial tree is fluid-filled in the fetus, bronchopulmonary situs may potentially be determined by ultrasound, but this is time-consuming and subject to limitations of fetal positioning and is therefore infrequently possible. Intestinal malrotation is not amenable to prenatal ultrasound diagnosis.

Because of the frequent segregation of often–typical cardiac abnormalities in either form of isomerism of the atrial appendages, the correct identification of these findings affords the prenatal sonologist the best opportunity at an antenatal diagnosis. Ultrasound assessment of fetal atrial appendage shape has been performed correctly in a small series, but remains difficult and not completely reliable. Because of the differential flow and pressure characteristics noted during fetal life, isomeric appendages may also be dissimilar in size and shape, making the diagnosis difficult. The pectinate muscles cannot be evaluated at all. Aside from analysis of segmental anatomy, knowledge of the associations of intracardiac morphology and extracardiac findings affords the best opportunity to arrive at a correct diagnosis in these syndromes.

In cases of isomerism of the right atrial appendages, there is substantial literature devoted to the documentation of the specific malformations. These include abnormalities of cardiac position, systemic and pulmonary venous return, forms of atrioventricular canal defects, abnormalities of ventriculoarterial connections, and arterial atresias or stenoses. Some of these malformations are extremely prevalent in the syndrome of right atrial isomerism, many of which are noted in this fetus. These include abnormal ventriculoarterial connections, pulmonary artery outflow obstruction, bilateral superior venae cavae, juxtaposition of the aorta and inferior vena cava, anomalous pulmonary venous return, and an absent coronary sinus. The presence of numerous abnormalities typical for the disorder allowed us to counsel the parents accurately.

There were several uncommon features. First, this fetus demonstrated anomalous pulmonary venous return directly to the right atrium. In most cases of isomerism of the right atrial appendages, anomalous pulmonary venous return is supra- or infracardiac. The combination of bilateral systemic venous return and anomalous pulmonary venous return to the right-sided atrium was seen in only 10% of cases in a large series of right atrial isomerism.

Second, the atrioventricular connection is extremely atypical. In up to 90% of cases of right atrial isomerism, this connection is in the form of a complete atrioventricular canal defect. In our patient, tricuspid atresia with a patent mitral valve is noted. To our knowledge, there has only been one prior reported case of tricuspid atresia associated with fetal right atrial isomerism, and this specific abnormality was only diagnosed postnatally. In fact, the two largest fetal series of tricuspid atresia (totaling 142 patients) did not identify a single case that was associated with a heterotaxy abnormality, even when postnatal investigations were considered. Our case represents the first accurate prenatal diagnosis of tricuspid atresia on a background of right atrial isomerism. Because of this rare association, the larger diagnosis of isomerism of the right atrial appendages could be potentially dismissed, leading to suboptimal counseling regarding outcome and prognosis.

The implications of isomerism of the right atrial appendages extend beyond the complexity of the cardiac abnormality itself. Most children are functionally asplenic and have a lifelong susceptibility to encapsulated bacteria. Despite advances in surgical intervention, the mortality rate for patients requiring surgical intervention in the neonatal period is 75%, with those requiring pulmonary vein repair having a surgical mortality of 95%.

Our diagnosis confirms that recognition of this constellation of abnormalities permits accurate prenatal diagnosis and counseling even when atypical findings are present.

References