A Review of Findings in Fetal Cardiac Section Drawings
Part 3: The 3-Vessel-Trachea View and Variants

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Objective. The goal of this presentation is to review some of the common and rare fetal heart abnormalities and to provide an easy approach to these findings with the schematic drawings. Methods. Over the past 10 years, we collected cases in which the common views of the heart were abnormal and the differential diagnoses that existed for each. This presentation shows the normal sonographic sections and then variations of these sections and the associated anomalies. We used illustrative drawings to present these findings, enabling us to point out the main sonographic features of abnormalities of the heart. Results. This work reviews 21 fetal heart abnormalities in schematic drawings. Conclusions. This short review highlights several of the anomalies that can be recognized on the common sonographic views. The drawings tend to simplify the findings but should serve as a basis for those doing fetal echocardiography when they encounter an unusual finding. Key words: drawings; fetal echocardiography; prenatal sonography.

In this third part of our 3-part series, we review the prenatal sonographic findings of some cardiac anomalies at the level of the 3-vessel-trachea view. Many of the cardiac anomalies result in modifications of the anatomy, position, or direction of flow at the level of the structures in the upper thoracic region. The purpose of this presentation is to provide schematic drawings of these findings to serve as an easy reference for those anomalies.

Materials and Methods

We have summarized the typical findings of some cardiac anomalies collected over the past 10 years.

The colors used in the drawings are conventional medical artist colors, with arteries in red and veins in purple or blue. The colors do not represent flow directions, as in color Doppler sonography, for instance, and they do not correlate with fetal blood oxygenation. A few drawings (cases 10–17) use colors that represent color Doppler images, and those drawings are clearly identified. The drawings are presented with the fetus presenting in the cephalic dorsal position (Figure 1). Although this does not represent the conventional position used in computed tomography and magnetic resonance imaging, it will be more familiar to those who do prenatal sonographic examinations.
In the following descriptions and drawings, we review some anomalies at the level of the 3-vessel-trachea view. Figure 2 represents a normal 3-vessel-trachea view to serve as a reference image for comparison with the subsequent drawings representing anomalies. The drawing shows the pulmonary artery, the aorta, and the superior vena cava. Some of the following drawings also show the trachea and esophagus, which we have represented here in Figure 3.

In this case, the 3-vessel view shows a thin pulmonary artery and a large aorta (Figure 4). This indicates an increase in flow in the left side with a concomitant decrease in the right. This could be present in tetralogy of Fallot, pulmonary atresia, pulmonary atresia with a ventricular septal defect (VSD), a double-outlet right ventricle, and a truncus arteriosus.\textsuperscript{1,2}

In this case, the 3-vessel view shows the reverse of case 1: a thin aorta and a large pulmonary artery (Figure 5). These findings can represent aortic atresia/stenosis, coarctation of the aorta, and a double-outlet right ventricle. The double-outlet right ventricle is in the previous category too: it depends on whether the aorta or the pulmonary artery shows an obstruction.\textsuperscript{1} Another rare condition with a huge dilated pulmonary artery is absent pulmonary valve syndrome\textsuperscript{3,4} (see case 1).

This case is a tricky one (Figure 6). If both great vessels are enlarged, it cannot be that one is enlarged at the expense of the other. Therefore, a different explanation is required. The only way to get increased flow in both vessels is to have incompetent valves in both vessels. Thus, the increase in flow is due to regurgitation of the incompetent aortic and pulmonary valves.
Occasionally, with atroventricular blocks, the vessels may also be dilated, as in this case.

**Case 4**
In this case, the little red “buglike” structure shown on the drawing is a double aortic arch (Figure 7). The appearance is due to its division into right and left aortic arches, each with its own common carotid and subclavian arteries.\(^5\)\(^-\)\(^7\) This diagnosis is almost impossible in sagittal and parasagittal views because they would appear as normal arches (though with only 2 brachiocephalic vessels). Note that the trachea and esophagus are located in the division of the 2 branches. This would be responsible for dysphagia and stridor after birth.

**Case 5**
In this case, the abnormal finding is a vein on the left side of the pulmonary artery (Figure 8). Although the finding is easy to recognize on this view, it is commonly missed because it is not sought. This is the classic finding of a persistent left superior vena cava.\(^7\) Lower at the level of the 4-chamber view, we could see a vessel in the atrioventricular groove. The persistent left superior vena cava will drain either in the back of the left atrium or in the coronary sinus.\(^8\)\^-\(^13\) The distension of the coronary sinus will appear as a transverse line (roughly parallel to the atrioventricular valve) in the lower aspect of the left atrium.
Case 6
This case represents an enlarged superior vena cava (Figure 9). This is the final level at which to recognize an inferior vena cava interruption with azygos or hemizygos continuation. Another possibility is an anomalous pulmonary venous return with a supracardiac connection or increased circulation of the brain, as found in a vein of Galen aneurysm.

Case 7
In this case, the great vessels arise in parallel, thus representing transposition of the great arteries (Figure 10). The drawing is a little simplified because the vessels are rarely at the same level but tend to curl under each other, usually with the transposed aorta more cephalad then the pulmonary artery.

Convergence of the Great Vessels
The next 2 images are at a slightly different level then the regular 3-vessel view. This is the level at which the confluence of the aorta and pulmonary artery can be observed. A normal section at this level is depicted in Figure 11.

Case 8
In this case, the “V-shaped” confluence of the aorta and ductus points to the right of the trachea (Figure 12). This is a sign of a right-sided aortic arch. Right-sided refers to the position of the arch compared with the trachea, not to the position of the descending aorta.

Figure 8. Case 5. Three-vessel-trachea view showing a supernumerary vascular structure (purple) on the left of the pulmonary artery (blue), which represents a persistent left superior vena cava.

Figure 9. Case 6. Three-vessel-trachea view showing an enlarged superior vena cava (purple).

Figure 10. Case 7. Three-vessel-trachea view showing both great vessels in parallel (aorta in red, pulmonary artery in blue, and superior vena cava in purple). In real life, they are more or less parallel but rarely on the same plane.

Figure 11. Normal confluence of the ductal and aortic arches. The descending aorta, which is on a lower plane, is “shown by transparency.” It is represented to indicate the position of the descending aorta in lower images. Again, note the positions of the trachea and esophagus (using the same symbols as in Figure 3).
Case 9
In this case, the “U-shaped” confluence of the aorta, pulmonary artery, and ductus encircles the trachea (Figure 13). This too is a sign of a right-sided aortic arch. This is one of the “rings and slings” that will cause dysphagia and stridor in the young child.14

Color Doppler Imaging
In the next few drawings, which are in a section similar to that in Figure 11, we represent the color Doppler information so that we can show findings in which a color Doppler signal is useful for the diagnosis. In these cases, the conventional assignment of color (blue going away from the transducer and red going toward the transducer) has been respected. The great vessels are in their expected positions.

Figure 12. Case 8. Upper transverse thoracic view showing a V-shaped confluence of the aorta (red) and ductus (blue). The trachea and esophagus are located to the left of the ductus.

Figure 13. Case 9. Upper transverse thoracic view showing a U-shaped confluence of the aorta (red) and pulmonary artery (blue). The trachea and esophagus are located between the aorta and the ductus.

Figure 14. Normal 3-vessel view. The colors given to the vascular structures in the following drawings correspond to color Doppler representations. Flow in the pulmonary artery (left) and aorta (right) is directed away from the transducer toward the spine. (We presume that the transducer is placed on the top of the image.)

Case 10
In this case, both vessels have antegrade flow, and the pulmonary artery is thinner than the aorta (Figure 15). This would thus suggest a right-sided restriction such as pulmonary stenosis, an Ebstein anomaly, tetralogy of Fallot, or a double-outlet right ventricle. This is similar to case 1.

Figure 15. Case 10. Upper transverse thoracic view. Both great vessels have antegrade flow, but the pulmonary artery (left) is thinner than the aorta (right).
Case 11
In this case, color Doppler sonography shows antegrade flow in the aorta and retrograde flow in the pulmonary artery (Figure 16). This would suggest pulmonary atresia with an intact septum or with a ventricular septal defect.15

Case 12
The difference between this case and case 11 is that only minimal retrograde flow exists in the pulmonary artery (Figure 17). This would suggest pulmonary atresia with a ventricular septal defect, and the flow is blood going to the lungs only. The main pulmonary artery is missing and not visible.15

Case 13
In this case, the thin pulmonary artery arises from the aorta (Figure 18), the hallmark of a truncus arteriosus.16

Case 14
In this case, both vessels have antegrade flow, but now the pulmonary artery is larger than the aorta (Figure 19). This finding can be seen in coarctation of the aorta.16–18

Case 15
With antegrade flow in the pulmonary artery but retrograde flow in the aorta (Figure 20), we are probably dealing with hypoplastic left heart syndrome with reversed flow to the coronary arteries.18–20

Figure 16. Case 11. Upper transverse thoracic view. The flow within the aorta has a normal antegrade direction (blue), but the flow within the pulmonary artery has a retrograde direction (red).

Figure 17. Case 12. Upper transverse thoracic view. The flow within the aorta has a normal antegrade direction (blue), but the minimal flow within the pulmonary artery has a retrograde direction (red).

Figure 18. Case 13. Upper transverse thoracic view showing a thin pulmonary artery arising from the aorta, the hallmark of a truncus arteriosus.

Figure 19. Case 14. Upper transverse thoracic view. Both great vessels have normal antegrade flow, but the pulmonary artery (left) is larger than the aorta (right).
Case 16
A partially visible aortic arch (Figure 21) should suggest an interrupted aortic arch. Because the pulmonary artery is arising from the arch, this is called a hemitruncus.\textsuperscript{21,22}

Case 17
Although this image (Figure 22) is quite similar to the previous one (Figure 21, case 16), the aorta in this case is independent from the pulmonary artery. This finding suggests a simple interruption of the aortic arch.

Differential Diagnoses of Overriding Aorta
In the next set of drawings, we review the approach to cases of an overriding aorta and concentrate on how to use the 3-vessel view to approach the differential diagnosis. Figure 23 depicts the overriding aorta in a 4-chamber view of the heart.

Case 18
When the pulmonary artery appears normal in size (Figure 24), we are probably dealing with a malalignment ventricular septal defect\textsuperscript{23} in a mild form of tetralogy of Fallot.
Case 19
When the pulmonary artery is thin (Figure 25), then classic tetralogy of Fallot is the most likely diagnosis.

Case 20
When the pulmonary artery and ductus are not only thin but also tortuous (Figure 26), and the flow is reversed (ductal dependent), then the likely diagnosis is pulmonary atresia with a ventricular septal defect.24

Case 21
This is another unusual case but so characteristic that it is well worth mentioning. Massive dilatation of the pulmonary artery (Figure 27) is due to “absent” pulmonary valve syndrome.4 (Although the valve is rarely totally absent, it is usually very incompetent.)

Conclusions
This short review highlights several of the anomalies that can be recognized on 3-vessel sonographic views and with color Doppler imaging. The drawings tend to simplify the findings, but our goal was to create simple and easily rememberable patterns of some anomalies that can serve as a basis for those doing fetal echocardiography when they encounter an unusual finding.
References


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