



Prenatal diagnosis of congenital heart defects: echocardiography

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Abstract: Congenital heart defects (CHD) are the most common congenital anomaly, and the majority can be diagnosed during prenatal life. Prenatal detection rates remain highly variable, as most CHD occur in low risk pregnancies and therefore depend on the maternal obstetric provider to recognize fetal cardiac abnormality on obstetric screening anatomic ultrasound. Fetuses with abnormal findings on obstetric screening anatomic ultrasound and/or risk factors for cardiac disease should be referred for evaluation with fetal echocardiography. Fetal echocardiography should be performed by specialized sonographers and interpreted by physicians with knowledge of evolving fetal cardiac anatomy and physiology throughout gestation. A fetal echocardiography examination, which can be done from the late first trimester onward, utilizes a standardized and systemic approach to diagnose fetuses with CHD or other forms of primary or secondary cardiac disease. The field of fetal cardiology has advanced past the accurate prenatal diagnosis of simple and complex CHD, as fetal echocardiography enables understanding of dynamic fetal cardiac physiology and consideration of potential fetal/neonatal treatment. The greatest impact of fetal echocardiography remains identification of critical CHD before birth to allow immediate cardiac management after delivery to decrease neonatal morbidity and mortality. Analyzing the severity of abnormal cardiac physiology in various forms of CHD before birth allows the fetal cardiologist to prognosticate effects on the developing fetus, predict risk of postnatal hemodynamic instability, guide delivery planning through multidisciplinary collaboration, and anticipate how the disease will impact the neonate after delivery.

Keywords: Congenital heart defects (CHD); fetal echocardiography; fetal ultrasonography; fetal heart; prenatal diagnosis

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Introduction

Congenital heart defects (CHD) are the most common congenital anomaly, accounting for approximately 1% of live births (1). The majority of CHD can be diagnosed during prenatal life by fetal echocardiography, which has been utilized to detect CHD *in utero* since 1980 (2,3). With advances in ultrasound technology, details of fetal cardiac anatomy and physiology are able to be discerned leading to improved prenatal and perinatal management and prognostication of both simple and complex CHD (4). Prompt recognition of CHD in the fetus allows the family time to evaluate for potential associated extracardiac anomalies and/or genetic syndromes and make decisions regarding pregnancy continuation. Approximately 25%

of infants with CHD have critical CHD requiring timely procedural cardiac intervention after birth to prevent significant morbidity or mortality (5). Prenatal diagnosis of critical CHD leads to appropriate prenatal multidisciplinary maternal and fetal care and delivery planning in an effort to decrease the risk of neonatal hemodynamic instability while transitioning to postnatal management in a pediatric cardiac center (4).

Prenatal diagnosis of congenital heart defects

Screening for cardiac anomalies should occur in all pregnancies during routine obstetric anatomic ultrasound examination during the second trimester (6). Some pregnancies are affected by maternal and/or fetal factors

Table 1 Indications for fetal echocardiography

Maternal factors	Fetal factors
<ul style="list-style-type: none"> • Diabetes mellitus or gestational diabetes • Lupus or Sjogren's syndrome with anti-SSA/anti-SSB antibodies • Use of assisted reproduction technology • Maternal infection during pregnancy (i.e., Rubella, Coxsackie, Parvovirus) • Teratogen exposure • Phenylketonuria, uncontrolled • Family history of congenital heart disease in 1st degree relative (mother/father/sibling of fetus) 	<ul style="list-style-type: none"> • Suspected cardiac anomaly on obstetric ultrasound • Fetal arrhythmia • Extracardiac anomaly • Chromosomal abnormality • Increased NT (≥ 3 mm) • Fetal hydrops or pericardial effusion • Monochorionic twinning • Abnormal umbilical or placental venous anatomy

that increase the risk of fetal cardiac disease (7) and should therefore undergo fetal echocardiography regardless of obstetric anatomic ultrasound findings (*Table 1*). However, most CHD occurs in low risk pregnancies without maternal or fetal risk factors (8,9), and therefore most referrals for fetal echocardiography and subsequent prenatal diagnosis of CHD rely solely on detection of a cardiac anomaly on screening obstetric ultrasound (7,8).

Historically, obstetric anatomic ultrasound guidelines for fetal cardiac evaluation mandated only a four-chamber view, with “views of the outflow tracts attempted as part of the cardiac screening” (10). In the four-chamber view, anomalous pulmonary venous connections, atrioventricular canal defects, atrioventricular valve stenosis or atresia, Ebstein's anomaly, hypoplastic left heart syndrome (HLHS), and/or septal defects are examples of CHD that can be identified. However, many types of CHD can be missed by imaging only the four-chamber view. Outflow tract and conotruncal anomalies such as truncus arteriosus, double outlet right ventricle, D-transposition of the great arteries (d-TGA), tetralogy of Fallot (TOF), pulmonary stenosis, pulmonary atresia with intact ventricular septum, and aortic stenosis may only be detected on outflow tract imaging. In 2013, the American Institute of Ultrasound in Medicine officially recognized the importance of imaging ventricular outflow tracts by updating its guidelines on obstetric anatomic ultrasound to include these critical outflow tract views into its basic cardiac examination (6). These guidelines (developed in conjunction with the American College of Radiology, American College of Obstetricians and Gynecologists, and the Society of Radiologists in Ultrasound) are the standard of care reference for obstetric practices in the United States; similar guidelines have been published by The International Society of Ultrasound in

Obstetrics and Gynecology (11).

Despite advances in fetal cardiac imaging, prenatal detection rates of CHD remain highly variable (*Table 2*) (12-21). In particular, detection rates of CHD affecting the outflow tracts are particularly low (22,23) as compared to other types of critical CHD which affect the four-chamber view (such as single ventricle lesions) (12,24). Contemporary studies have shown a wide range of prenatal diagnosis rates for conotruncal anomalies, ranging from 25% to 75% (16-18,21,25-27). Prenatal detection rates of cardiac outflow tract anomalies have not increased significantly and remain suboptimal (28) despite the updated anatomic ultrasound guidelines in 2013.

To optimize prenatal detection of CHD, sonographers and practitioners who perform fetal anatomic screening ultrasounds must be, at a minimum, comfortable with assessing the four-chamber and ventricular outflow tract views (*Figure 1*) in accordance with the recommended guidelines. Studies have shown that less experienced obstetric sonographers have lower detection rates of cardiac abnormalities on screening ultrasound (29). Any concerns for a fetal cardiac abnormality raised by screening ultrasound should be directed to more comprehensive fetal echocardiography. Fetal echocardiography by fetal/pediatric cardiac sonographers, with interpretation by fetal/pediatric cardiologists, provides more accurate prenatal cardiac diagnosis (30,31) as compared to obstetric ultrasound and allows for specialized prenatal parental counseling as well as delivery management in an appropriate setting (32).

Current guidelines for fetal echocardiography

The fetal heart should be assessed in all pregnancies during the standard obstetric anatomic ultrasound, typically

Table 2 Contemporary rates of prenatal detection of select congenital heart defects

Congenital heart defect	Population location	Author	Years	Prenatal detection rate (%)
TAPVC	United States	Quartermain	2006–2012	9
Atrioventricular canal defect	United States	Quartermain	2006–2012	40
HLHS	Czech Republic	Marek	2000–2006	50
	Sweden	Ohman	2001–2010	63
	Finland	Markkanen	2003–2010	38
	United States	Quartermain	2006–2012	67
Critical Pulmonary Stenosis	Boston	Ronai	2000–2014	37
PA/IVS	Italy	Tuo	1993–2009	60
	Czech Republic	Marek	2000–2006	68
	Boston	Ronai	2000–2014	60
Aortic Stenosis	United States	Quartermain	2006–2012	25
Truncus Arteriosus	Netherlands	van Velzen	2002–2012	62
	United States	Quartermain	2006–2012	41
TOF	Netherlands	van Velzen	2002–2012	30
	United States	Quartermain	2006–2012	26
	Belgium	De Groote	2006–2014	23
D-TGA	Alberta	Ravi	2003–2015	37
	United States	Quartermain	2006–2012	31
	Netherlands	van Velzen	2007–2012	41
	Belgium	De Groote	2006–2014	24
cc-TGA	Toronto	Wan	1999–2006	29
	United States	Quartermain	2006–2012	45

TAPVC, total anomalous pulmonary venous connection; HLHS, hypoplastic left heart syndrome; PA/IVS, pulmonary atresia/intact ventricular septum; TOF, tetralogy of Fallot; D-TGA, d-transposition of the great arteries; cc-TGA, congenitally corrected transposition of the great arteries.

performed between 18–22 weeks gestation, with adequate visualization of the four-chamber and ventricular outflow tract views (6). Any fetus with abnormal cardiac views or increased maternal or fetal risk of CHD (*Table 1*) should undergo detailed fetal echocardiography (7). With normal cardiac views on screening anatomic ultrasound, fetal echocardiography can still increase detection of CHD (usually non-critical) in pregnancies at higher risk than the general population (33).

Performance of fetal echocardiography

Fetal echocardiography should be performed by trained and skilled sonographers and physicians (34,35) who have a detailed understanding of normal and abnormal fetal cardiac anatomy and physiology. The physician interpreting the fetal echocardiography examination should have comprehensive and contemporary knowledge of management options and outcomes of all forms of CHD, as counseling the family is an important component of

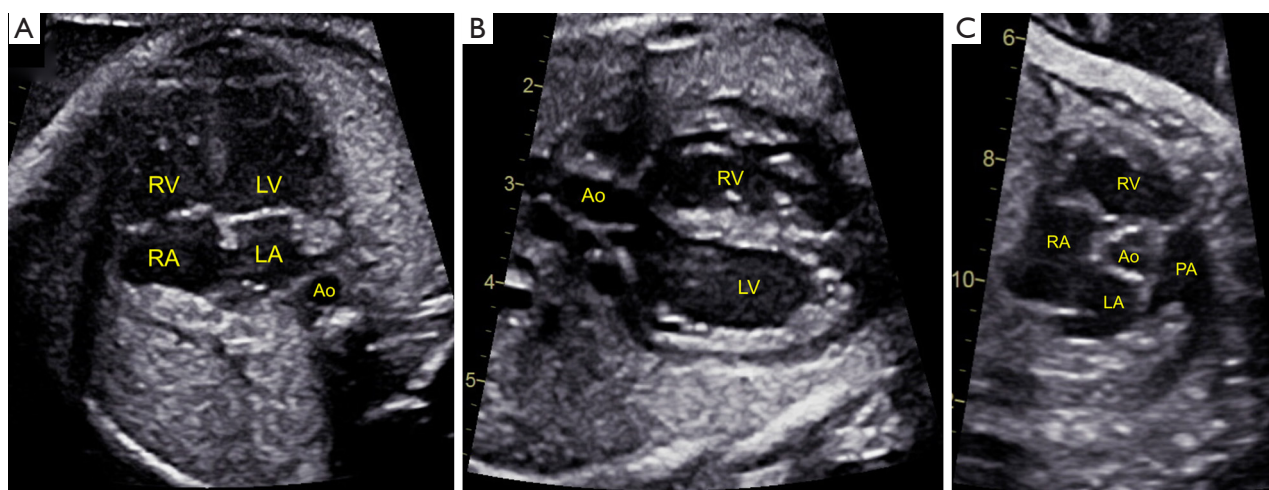


Figure 1 Cardiac screening views in routine obstetric anatomic ultrasound include the (A) four-chamber, (B) left ventricle (LV) outflow tract, and (C) right ventricle (RV) outflow tract views. RA, right atrium; LA, left atrium; PA, pulmonary artery; Ao, aorta.

caring for the fetus with CHD. The interpreting physician should also be integrated in a multidisciplinary team of fetal and pediatric care providers, such as perinatologists, neonatologists, geneticists, pediatric cardiologists, and pediatric cardiovascular surgeons for optimal care coordination of both fetus and mother.

The optimal timing for screening fetal echocardiography is typically between 18–22 weeks gestation (36). Early fetal echocardiography (performed before 16 weeks gestation) is typically reserved for those with fetuses at highest risk of having CHD, such as increased nuchal translucency ≥ 3.5 mm (37), abnormal ductus venosus (38), extracardiac or genetic abnormality, prior child with significant CHD, or monozygotic twinning (39). In this subset of pregnancies at highest risk for fetal CHD, early fetal echocardiography can be performed transvaginally (40) (as early as 11 weeks gestation) or transabdominally (41) (as early as 13–14 weeks gestation) to identify CHD. While early fetal echocardiography can accurately detect major CHD (Figure 2), some defects may not be apparent or manifest until later gestation (42,43). Therefore, early fetal echocardiography should supplement, but not replace, standard mid-gestation transabdominal fetal echocardiography (44). Fetal echocardiography should be performed as soon as possible following abnormal findings on obstetric ultrasound. It is important to note the inherent limitations of fetal cardiac imaging throughout pregnancy, as some defects may not be detected *in utero*, such as small atrial or ventricular septal defects, minor valvar abnormalities, partial anomalous

pulmonary venous return, coronary artery anomalies, or aortic coarctation. In addition, some forms of CHD may not be detected by mid-gestation fetal echocardiography due to manifestations in later gestation, such as isolated valvar pulmonary or aortic stenosis (45), cardiomyopathy, or cardiac tumors (46).

The ultrasound system used for fetal echocardiography should be capable of two-dimensional, color and pulse-wave Doppler, and M-mode imaging, with transducer probes with frequencies between 4–12 MHz. The highest frequency transducer should be utilized to maximize frame rate (with a preferred minimum of 50 Hz and optimum of 80–100 Hz) due to the fast fetal heart rate (typically between 120–160 bpm). The fetal sonographer should be cognizant to optimize acoustic windows, which can be affected by maternal body habitus, fetal lie, and fetal movement.

A comprehensive fetal echocardiography examination utilizes a systematic approach maximizing standard views of the fetal heart to evaluate each cardiac structure from multiple planes (11,35,47). The cardiac structures should be critically evaluated utilizing a combination of two-dimensional or gray scale imaging with color and pulse-wave Doppler interrogation. Due to the dynamic nature of cardiac structures and function, the views should be recorded with moving cine clips. Color Doppler should be used to evaluate for flow disturbances through each of the cardiac valves as well as systemic and pulmonary venous flow into and arterial flow out of the heart (48). Pulse wave spectral Doppler assessment is necessary to quantify flow

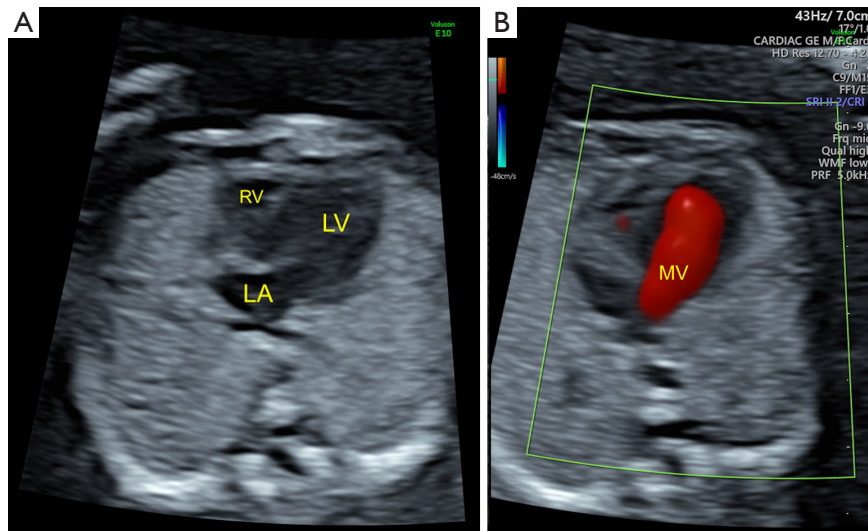


Figure 2 Early fetal echocardiography in a 15-week fetus with tricuspid atresia. (A) Four-chamber view showing hypoplastic right ventricle (RV) with normal left atrium (LA) and left ventricle (LV). (B) Absent tricuspid valve inflow and normal mitral valve (MV) inflow are demonstrated with color flow imaging.

through each valve as well as any abnormal flows detected by color Doppler. Ductus venosus and umbilical artery and vein flow are also typically assessed to monitor potential effects of altered fetal hemodynamics (49,50). The fetal heart rate and rhythm should be determined by pulse wave Doppler and/or M-mode imaging. Measurements of cardiac chambers, valve annuli, and vessel dimensions in two-dimensional imaging can be referenced to normative datasets based on fetal gestational age or biometry (51,52). Fetal cardiac function can be assessed qualitatively by two-dimensional imaging and/or quantitatively by M-mode or advanced techniques (such as strain or tissue Doppler imaging). Biometric measurements typically include at a minimum cardiothoracic ratio and femur length.

Fetal echocardiography: standard views

The methodical use of standard two-dimensional views provides a comprehensive evaluation of mid-gestation cardiac anatomy (53). The minimum standard views include definition of visceral and cardiac situs, four-chamber, right and left ventricular outflow tracts, three-vessel, three-vessel trachea, bicaval, ductal arch, aortic arch, and short-axis of the ventricles and great arteries. Additional non-traditional planes may be necessary to define abnormal cardiac anatomic connections in fetuses with complex CHD.

Each fetal echocardiography examination should begin

by establishing fetal lie followed by visceral and cardiac situs. The heart should sit in the left chest with a leftward apex, and a transverse sweep down to the abdomen should demonstrate a left sided stomach and right sided liver. In the transverse plane, the heart should comprise one third of the thoracic cavity area.

In the four-chamber view (*Figure 1A*), the position and size of atria and ventricles as well as the integrity of the interatrial and interventricular septum are evaluated. The foramen ovale flap is seen directing flow from the umbilical vein and should bow into the left atrium. At least one pulmonary vein from the right and one from the left lung should be seen entering the posterior wall of the left atrium. At the crux of the heart, the tricuspid valve attachment should be more apically displaced than the mitral valve attachment. The diameters of the tricuspid and mitral valve annuli (in diastole) and the right (RV) and left ventricles (LV) (at their maximal dimensions at end diastole) can be measured. The right and left side should be fairly symmetric in size. Color and pulse wave Doppler of the tricuspid and mitral inflows should show laminar flow with biphasic E and A waves. The moderator band should be seen in the anterior and rightward ventricle, identifying it as the true morphologic RV. In contrast, the posterior and leftward ventricle should be more smooth-walled, identifying it as the morphologic LV.

Evaluation of the outflow tracts is critical to detecting

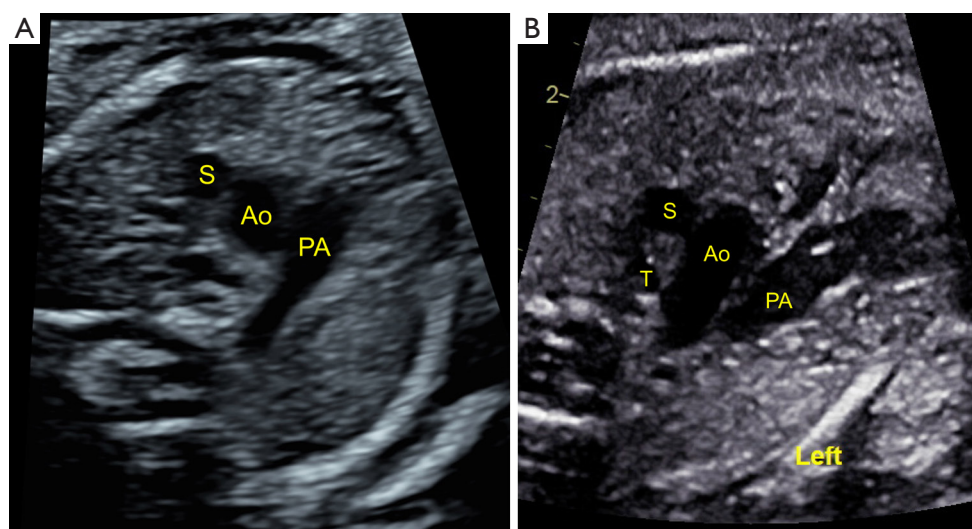


Figure 3 Normal (A) three-vessel and (B) three-vessel trachea view demonstrating a normal left-sided aortic arch. S, SVC; Ao, aorta; PA, pulmonary artery; T, trachea.

CHD, many of which can have a normal appearing four-chamber view, such as d-TGA, TOF, and truncus arteriosus amongst many others. A sweep from the four-chamber view superiorly should demonstrate the LV outflow tract crossing perpendicular to the more anterior RV outflow tract. The ventricular septum should be intact. The LV outflow tract view demonstrates the continuity of the ventricular septum with the great artery arising from it, which should not bifurcate (*Figure 1B*). In the RV outflow tract view, the main pulmonary artery should be seen bifurcating into the right and left branch pulmonary arteries (*Figure 1C*). The aortic and pulmonary valves should be assessed in a moving clip to demonstrate symmetric opening as well as with color flow and pulse wave Doppler to demonstrate normal laminar flow (with a velocity <100 cm/sec).

The three-vessel view is taken cranial to the four-chamber view, with the cross-sections of the superior vena cava, aorta, and pulmonary artery forming a linear configuration (*Figure 3A*). The superior vena cava has the smallest caliber and is more rightward and posterior than the aorta; the aorta is smaller than the pulmonary artery, which is largest and most leftward and anterior. In the three-vessel trachea view, the aortic and ductal arches should form a “V” shape alignment leftward of the midline echogenic trachea (*Figure 3B*) with flow from anterior to the posterior descending aorta.

A long-axis sagittal plane towards the right demonstrates the bicaval view, with the superior vena cava and inferior

vena cava joining into the right atrium. The foramen ovale flap can be seen deviating into the left atrium with flow from the umbilical vein and ductus venosus directed from the right towards the left atrium. More leftward, the ductal arch is seen arising from the short-axis of the right ventricle and directs flow from the pulmonary artery into the descending aorta. The aortic arch is more U-shaped and confirmed by demonstrating its head and neck vessel branches; the caliber of the ascending, transverse, and descending aorta should be relatively equal throughout. The short-axis view of the great arteries should show the tricuspid valve, RV outflow tract, and pulmonary artery wrapping around the short-axis of the aortic valve (*Figure 1C*). The short-axis view of the ventricles should be swept from apex to base with both two-dimensional and color flow imaging to demonstrate integrity of the ventricular septum.

Fetal echocardiography: advanced

Advanced fetal echocardiography techniques can supplement and aid the prenatal diagnosis of CHD. A thorough discussion of the available imaging modalities is beyond the scope of this review article. Strain or myocardial deformation imaging tracks movement of the myocardium throughout the cardiac cycle and analyzes global or regional abnormalities in myocardial function in various fetal cardiac diseases (54,55). Spectral or color tissue Doppler imaging quantifies the velocity of interrogated myocardial tissue

throughout the cardiac cycle, also evaluating for changes in fetal cardiac function in research and clinical practice (56). Current technologies in 3D/4D fetal echocardiography enable rapid acquisition of volumetric data of the heart that can be manipulated post-acquisition for improved diagnosis and understanding of spatial relationships in CHD (57,58).

Fetal cardiology

Most forms of CHD can be detected on fetal echocardiography, with some limitations as previously discussed. Several questions should be addressed upon diagnosis of a fetus with CHD: What is the risk of associated extracardiac anomalies or an underlying genetic syndrome? Are fetal cardiac interventions or *in utero* medical treatment options available? Is the fetus stable before birth, or is there risk of progressive heart failure and fetal demise? Will the family pursue pregnancy termination, cardiac intervention, or palliative care? After delivery, will the neonate be at high risk of hemodynamic decompensation upon separation from the maternal-placental circulation? What specialists need to be involved with the delivery and immediate postnatal care of the neonate? What postnatal cardiac interventions are anticipated? Addressing each type of CHD and its prenatal implications is outside the scope of this review, but select examples and the impact of prenatal diagnosis on perinatal and neonatal management will be highlighted here.

The prenatal diagnosis of CHD should prompt a thorough evaluation for extracardiac anomalies or genetic syndrome in the fetus. Fetuses can have associated malformations of or secondary developmental effects on the brain, lungs, kidneys, gastrointestinal and/or genitourinary systems which can impact perinatal morbidity as well as affect postnatal management. Genetic abnormalities have been found in up to 50% of fetuses with CHD (59), with certain types associated with higher risk of genetic abnormality (i.e., Atrioventricular canal defect with Trisomy 21 or truncus arteriosus, TOF, interrupted aortic arch, or right aortic arch with 22q11 deletion). Due to the association with genetic syndromes, all mothers of fetuses prenatally diagnosed with CHD should undergo consideration of genetic counseling and additional testing (i.e., amniocentesis with karyotyping or microarray).

Fetal cardiac intervention in a small subset of fetuses with CHD may be considered in an effort to alter natural *in utero* course, increase postnatal treatment options, and improve long-term outcomes (60-62). For example, a subset of mid-gestation fetuses with critical aortic stenosis may

evolve to develop HLHS by the end of gestation (63). Fetal balloon aortic valvuloplasty can relieve the aortic valve stenosis, promote flow through and potential growth of the left heart structures, and enable possibility of a biventricular circulation after birth (64) (Figure 4). Similarly, fetal balloon pulmonary valvuloplasty may be considered in a subset of mid-gestation fetuses with critical pulmonary stenosis or pulmonary atresia who may develop significant right heart hypoplasia without intervention (65,66). Families should be counseled that although fetal cardiac intervention may be technically successful (67), controversy exists weighing the risks versus benefits to the mother and fetus (68), who may or may not ultimately be a candidate for biventricular palliation after birth. Follow-up fetal echocardiography after intervention evaluates the impact on cardiac structural growth and physiology and provides up-to-date counseling regarding potential postnatal management and outcomes to the family.

Serial fetal echocardiography throughout gestation monitors for progressive fetal cardiac disease and the development of fetal heart failure. CHD at risk for fetal heart failure includes abnormal valves at risk for increasing valvar regurgitation or valvar stenosis, progressive obstructive lesions with risk for myocardial dysfunction, and structural abnormalities with risk for developing arrhythmias or heart block. Fetal heart failure can lead to valvar regurgitation, cardiomegaly, increased central venous pressure, and ultimately impede umbilical venous flow causing fetal hydrops with high risk of intrauterine demise (69). Assessment of the cardiovascular profile score by fetal echocardiography can be used to monitor fetuses at risk (49), with a low score predicting perinatal mortality. Options for transplacental medical therapy are limited in fetuses with structural or myocardial cardiac disease. Transplacental digoxin has been shown to improve heart failure, demonstrated by improved cardiovascular profile scores, in a small number of heterogeneous cases (70). In fetuses with Ebstein's anomaly or tricuspid valve dysplasia with circular shunt physiology due to significant tricuspid and pulmonary regurgitation, transplacental indomethacin and ibuprofen can cause ductal constriction, to diminish pulmonary regurgitation and systemic steal, prolong gestation, and increase perinatal survival (71,72). More successful treatment options exist for fetuses with arrhythmias, who may be given transplacental or more direct injections of antiarrhythmics (73). The decision regarding whether the presence of hydrops should prompt premature delivery should be weighed carefully against the

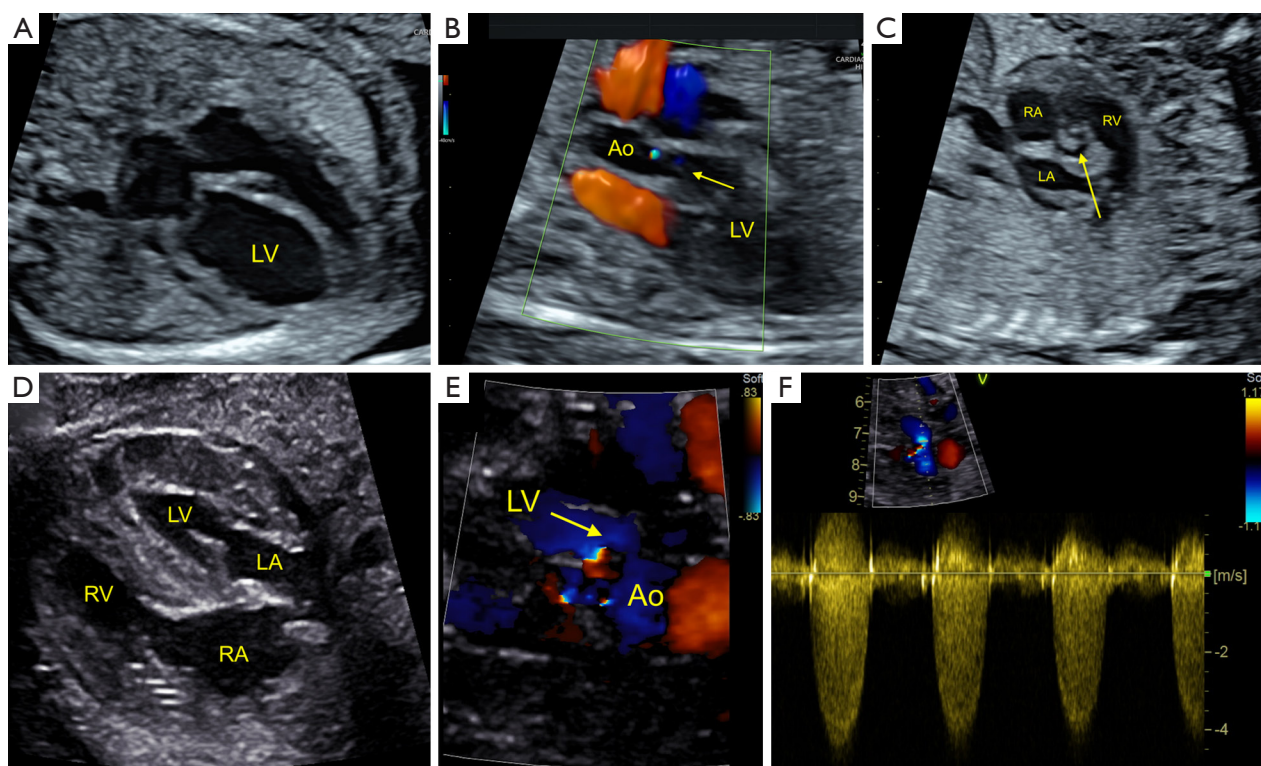


Figure 4 Fetus with critical aortic stenosis and evolving HLHS physiology at (A) 19 weeks gestation, with a dilated left ventricle (LV) with endocardial fibroelastosis and (B) minimal antegrade flow (arrow) across a (C) severely dysplastic and stenotic aortic valve. (D) After fetal balloon aortic valvuloplasty, the LV is hypertrophied with normalized function and (E) improved flow (arrow) across the (F) stenotic aortic valve at 34 weeks gestation. This infant subsequently required balloon aortic valvuloplasty in the neonatal period. Ao, aorta; RA, right atrium; LA, left atrium; RV, right ventricle.

known risks of prematurity associated with poor outcomes in major CHD (74).

Fetal echocardiography can guide delivery planning, as infants with critical CHD can have significant cyanosis, hypoxemia, and/or cardiogenic shock leading to morbidity/mortality after birth if not expeditiously treated. Fetuses can be stratified by risk of post-delivery hemodynamic compromise and need for urgent intervention upon separation from the maternal-placental circulation (32). For example, infants with total anomalous pulmonary venous connection (TAPVC) can manifest clinically with significant cyanosis and hypoxemia after delivery (75). While it can occur in isolation, TAPVC associated with other forms of critical CHD (such as heterotaxy) (Figure 5A) is more likely to be diagnosed prenatally (75). The low rate of prenatal diagnosis in isolated TAPVC is likely due to its more subtle appearance on screening cardiac views. Fetuses with obstructed TAPVC identified by fetal echocardiography (Figure 5B) are at high risk of significant hypoxemia

after birth without rapid cardiac intervention and should therefore undergo coordinated (with possible C-section) delivery to ensure prompt access to pediatric cardiology and cardiovascular surgery (76).

Prenatal detection improves pre-operative morbidity, mortality, and outcomes in certain types of CHD, such as d-TGA and HLHS (77-79). The majority of fetuses with d-TGA (Figure 6) remain stable *in utero*, supported by the placental circulation. After delivery, newborns with d-TGA can quickly become cyanotic and may require urgent balloon atrial septostomy to ensure adequate mixing between the parallel deoxygenated and oxygenated circulations. All fetuses with d-TGA are considered to be at high risk of hemodynamic decompensation after delivery and should prompt discussions to consider coordination of delivery close to a pediatric cardiac care center. Certain echocardiographic findings of abnormal septum primum, highly restrictive/intact atrial septum, and/or abnormal ductus arteriosus flow (80,81) in fetuses with d-TGA

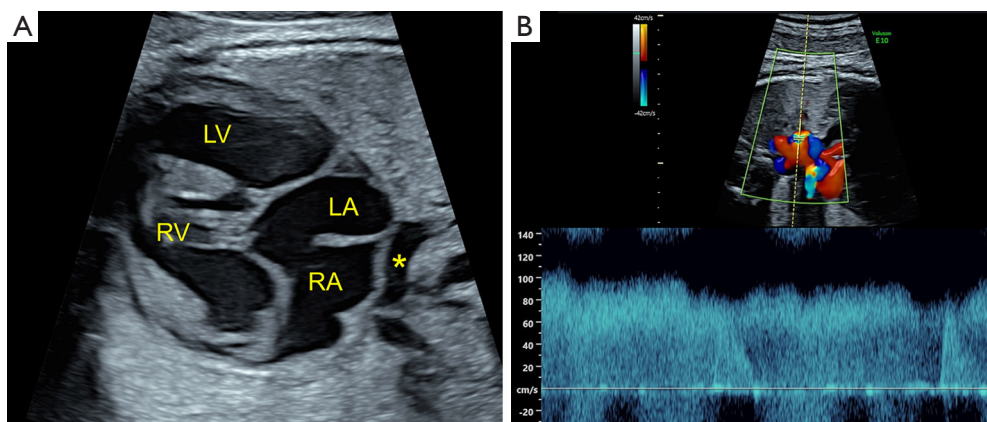


Figure 5 Total anomalous pulmonary venous connection in a fetus with heterotaxy, atrioventricular canal defect, and pulmonary stenosis. (A) The pulmonary venous confluence (*) is seen behind the right (RA) and left atrium (LA). (B) Pulse wave Doppler of the vertical vein showing non-phasic high velocity flow, consistent with obstruction, as it enters the innominate vein. RV, right ventricle; LV, left ventricle.

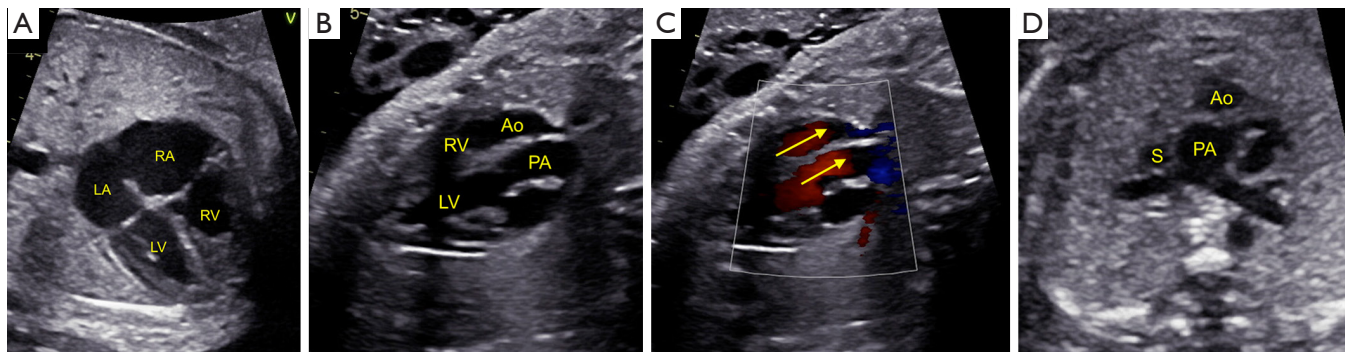


Figure 6 (A) Normal appearing four-chamber view in a fetus with D-transposition of the great arteries. (B) Abnormal parallel outflow tracts (arrows) are shown in two dimensional and (C) color flow imaging, with the right ventricle (RV) connected to the aorta (Ao) and the left ventricle (LV) connected to the pulmonary artery (PA). (D) Abnormal three-vessel view with the aorta anterior to the pulmonary artery. RA, right atrium; LA, left atrium; S, superior vena cava.

predict need for urgent balloon atrial septostomy. Similarly, in fetuses with HLHS, some develop restrictive or intact atrial septum impeding the return of oxygenated pulmonary venous blood flow causing severe hypoxemia after birth (82). Foramen ovale flow (Figure 7A) and pulmonary venous Doppler assessment (Figure 7B) can risk stratify these fetuses, with significant pulmonary venous flow reversal predicting need for urgent atrial septostomy after birth (83,84). HLHS fetuses with atrial restriction and significant pulmonary venous obstruction have worse outcomes (85,86) and survival (87) due to secondary effects on developing pulmonary vasculature (88). These findings *in utero* should guide medical team decision making regarding delivery (89) and parental desire for surgical versus palliative care

postnatally.

Some fetuses with CHD remain stable *in utero* with lower risk for hemodynamic compromise if ductal patency is maintained after birth (with the use of prostaglandin infusion) to preserve adequate pulmonary or systemic blood flow. Fetuses with right heart obstructive lesions, such as critical pulmonary stenosis, pulmonary atresia, severe TOF, or tricuspid stenosis/atresia with restrictive pulmonary flow, require a patent ductus arteriosus for pulmonary blood flow to prevent severe cyanosis after birth. The presence of fetal left to right (aorta to pulmonary) flow in the ductus arteriosus (instead of its normal flow from the pulmonary artery to descending aorta) predicts postnatal ductal dependency in right heart lesions (90) (Figure 8).

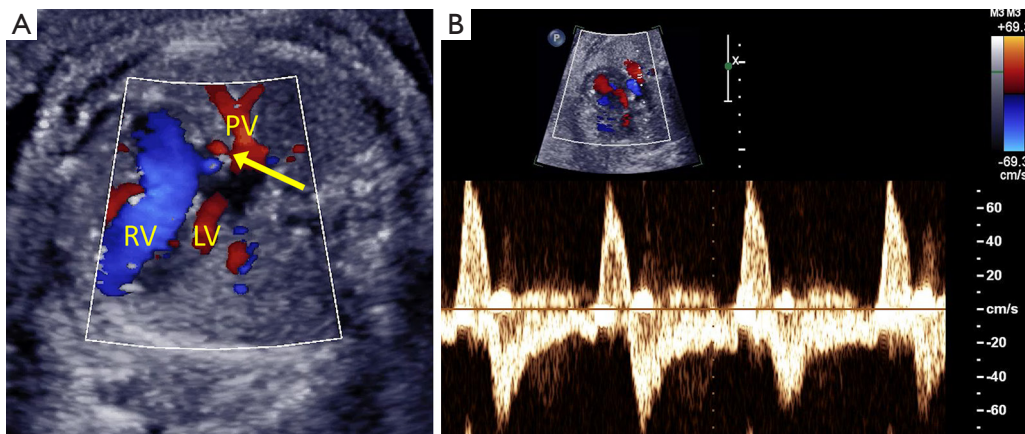


Figure 7 Fetus with hypoplastic left heart syndrome and (A) restrictive atrial septum (arrow), causing pulmonary vein (PV) flow reversal. (B) PV pulse wave Doppler shows significant A wave reversal (above the baseline). RV, right ventricle; LV, left ventricle.

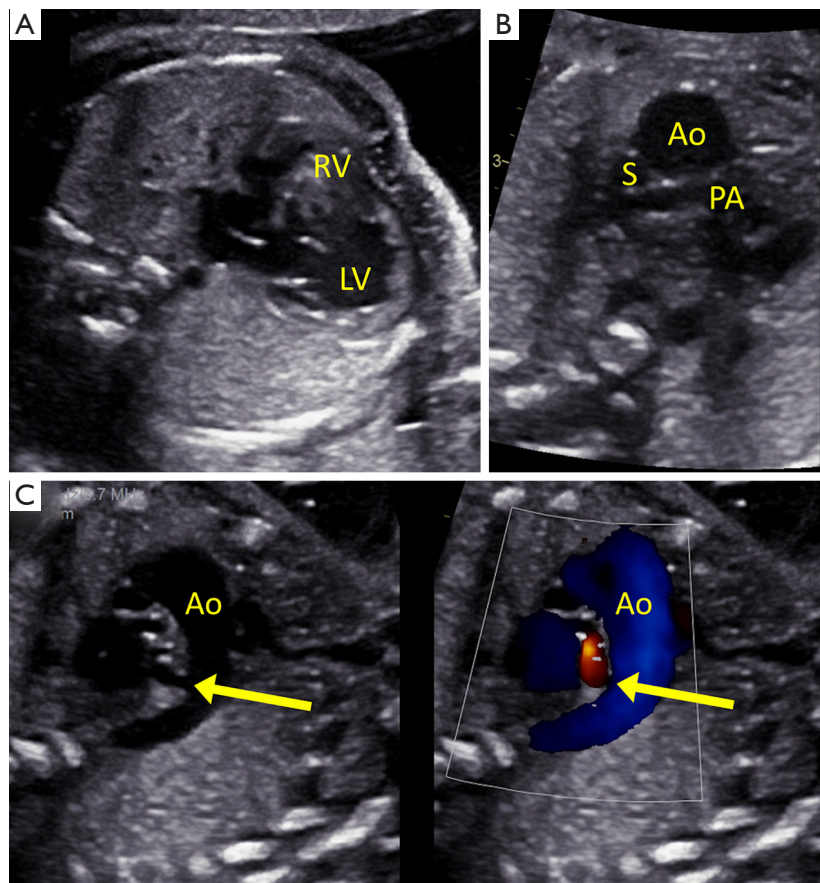


Figure 8 Fetus with pulmonary atresia and intact ventricular septum. (A) The right ventricle (RV) is severely hypoplastic and hypertrophied compared to the dilated left ventricle (LV). (B) Abnormal three-vessel view demonstrating a dilated aorta (Ao) and small pulmonary artery (PA). (C) Reverse oriented (arrow) ductus arteriosus with aorta to pulmonary artery flow predicting neonatal ductal dependence for pulmonary blood flow. S, superior vena cava.

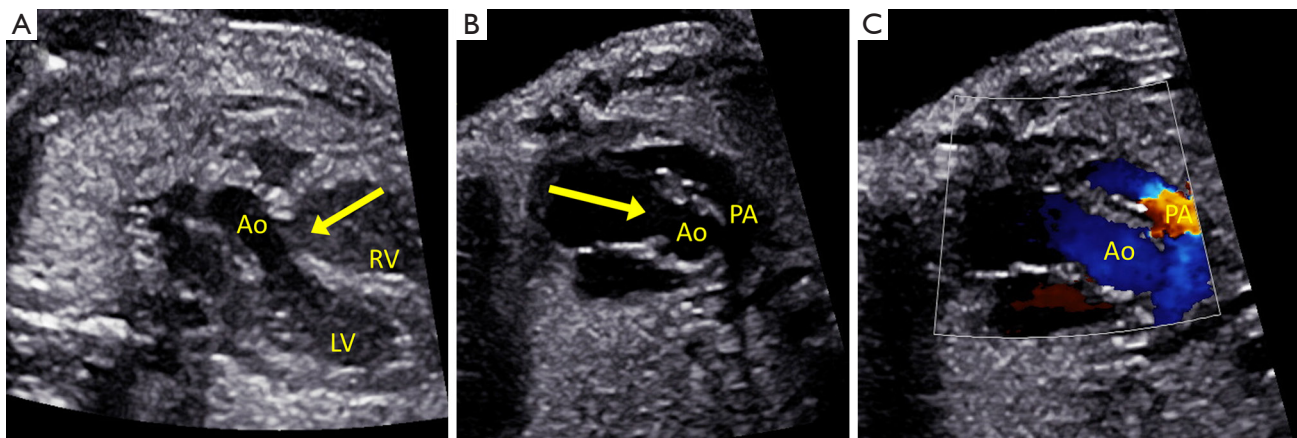


Figure 9 Fetus with non-ductal dependent tetralogy of Fallot. (A) Outflow tract imaging showing the overriding aorta (Ao) and ventricular septal defect (arrow). (B) Short axis view of the aorta and anterior malalignment ventricular septal defect (arrow), with the pulmonary artery (PA) smaller than the aorta. (C) Color flow shows mild flow acceleration across the mildly hypoplastic pulmonary valve.

Fetuses with left heart obstructive lesions, such as HLHS, critical aortic stenosis, aortic coarctation, or interrupted aortic arch, require a patent ductus arteriosus to maintain systemic perfusion after birth to prevent cardiogenic shock. Abnormal left to right flow across the foramen ovale predicts ductal dependent left sided CHD (91,92). Fetuses with aortic arch obstruction may have normal right to left flow across the foramen ovale, and determining true ductal dependence in fetuses with suspected aortic coarctation can be challenging by fetal echocardiography (93). Delivery planning for infants with ductal dependent CHD should consider their proximity to and need for admission to pediatric cardiac care centers with cardiac catheterization and surgical services.

Fetuses with minimal risk for ductal dependency or neonatal intervention can be safely delivered without alterations to planned location. Newborns with shunt lesions (i.e., isolated ventricular septal defects, balanced atrioventricular canal defects) or mild forms of TOF (Figure 9) typically remain hemodynamically stable *in utero* and can be evaluated with non-urgent/outpatient pediatric cardiology follow-up after delivery. These families benefit from prenatal counseling and anticipation of postnatal cardiac course, but can be reassured that the likelihood of neonatal cardiac intervention is low.

Conclusions

Fetal echocardiography plays an integral role in the accurate prenatal diagnosis of CHD, and the demonstration

of cardiac physiology *in utero* guides collaborative maternal and fetal care. The biggest limitation to the prenatal diagnosis of CHD remains its reliance on obstetrical providers to identify affected or at-risk pregnancies and therefore continued efforts in education should focus on these primary care sonographers and providers. When prenatal diagnosis of CHD is achieved, continuity of care for these families is established before birth, with the fetal cardiologist laying the foundation to guide the family and to optimize short and long-term outcomes.

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