



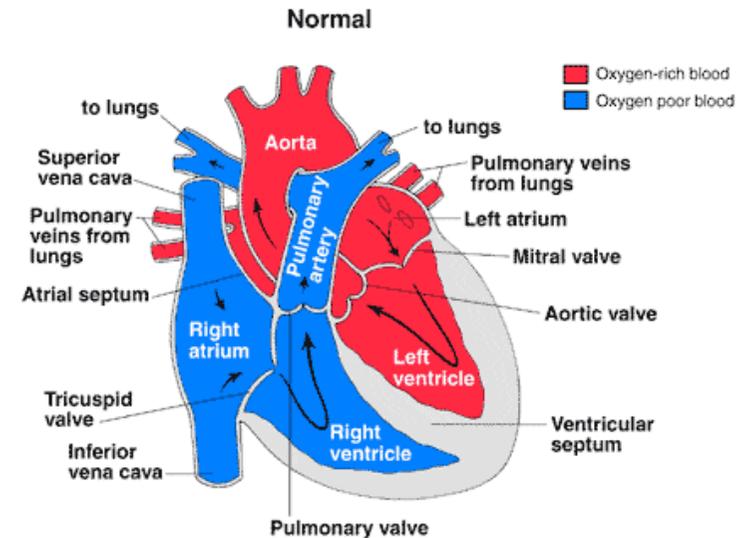
CARDIOVASCULAR ANOMALIES IN THE FETUS

Structural and Rhythm

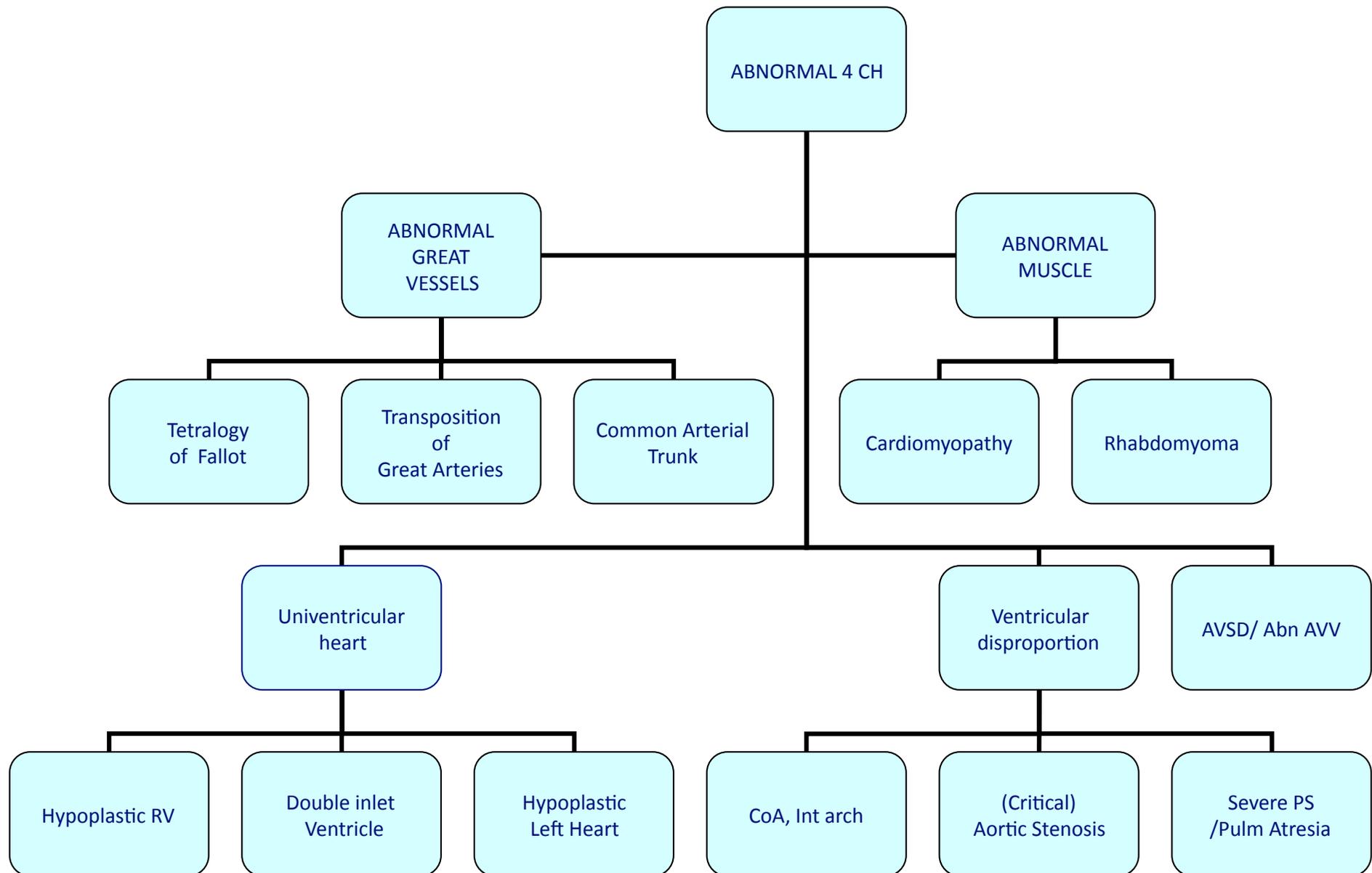
Identification of CHD

Sequential Segmental Analysis

- Pulmonary & Systemic veins
- Interatrial septum
- Atrio-ventricular connections
- Interventricular septum
- Ventriculo-Arterial connection
- Great Artery arrangement



Incredibly Simple Classification of CHD



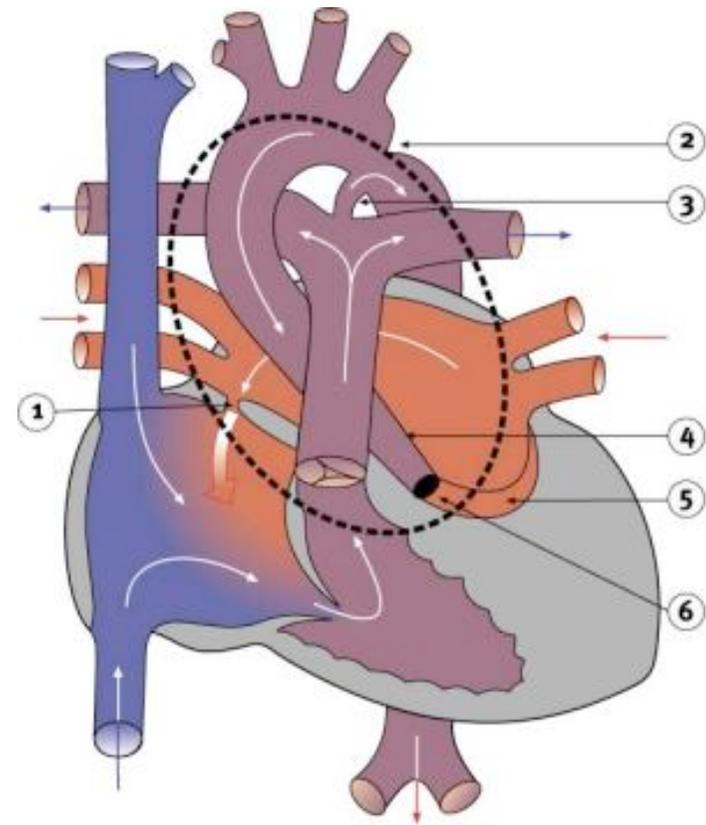
NB Classification modified by Dr S Barwick

Hypoplastic Left Heart Syndrome

- 5-10% of major CHD in infants
- Overrepresented in Fetal Series due to TOP, IUD, NND
- Readily detectable in 4 chamber view
- Diagnosis at $^{16-18}/_{40}$ or earlier
- Can be spectrum/progressive

Features of HLHS

- Disproportion
 - LA \ll RA; LV \ll RV; Ao \ll PA
- No forward flow through MV
- No forward flow through AV
- Reversal of flow in Ao Arch
- Reversal of flow at FO
 - ie. L to R

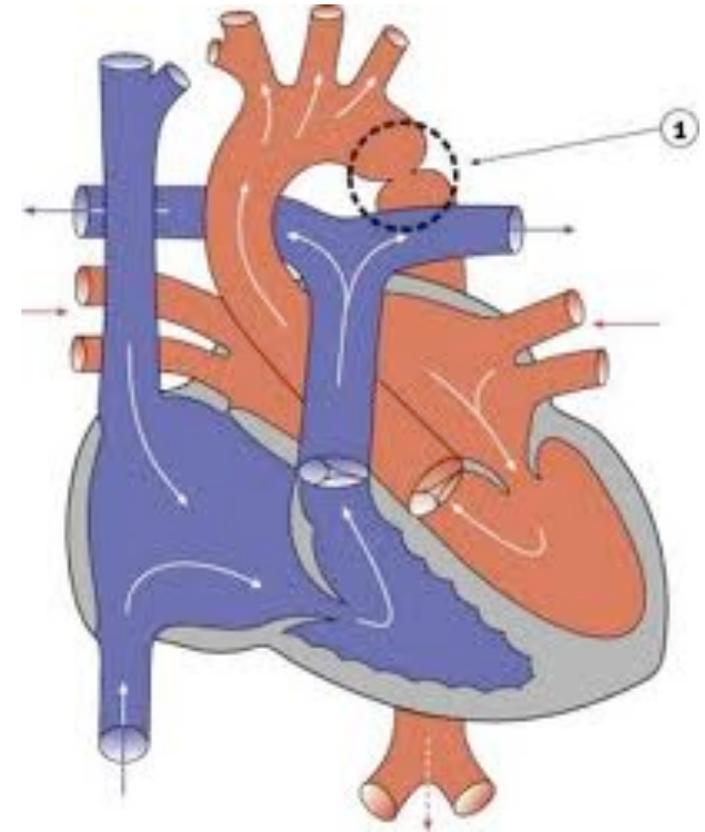


Abnormalities of the Aorta

- Coarctation of Aorta
- Interruption of Aortic arch
- Right Aortic arch
- AP window
 - ± Ventricular Septal defect
 - ± Ventricular Disproportion
 - Great vessel disproportion

Coarctation of the Aorta in the Fetus

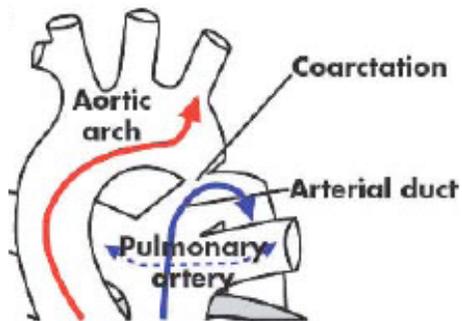
- Ventricular Disproportion
 - ie $LV < RV$
- Great vessel disproportion
 - ie $Ao < PA$



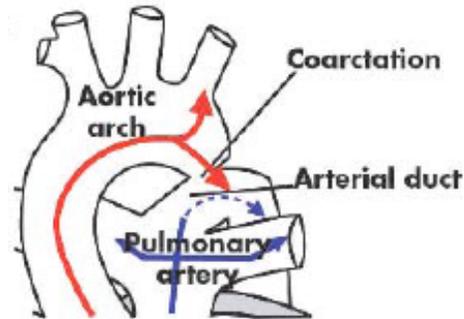
- “Isthmus of arch” $<$ Arterial Duct

Coarctation of the Aorta

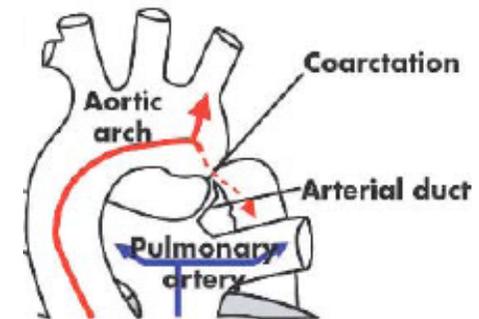
- Before birth fetal circulation able to cope with coarctation/ interruption



Coarctation in utero does not affect the fetal blood flow pattern



Fall in PVR with increased PBF and forward flow from aortic arch to dAo



Duct constricts Ao narrowing \uparrow
Increasing obstruction leads to gradient

Interruption of the Aorta in the Fetus

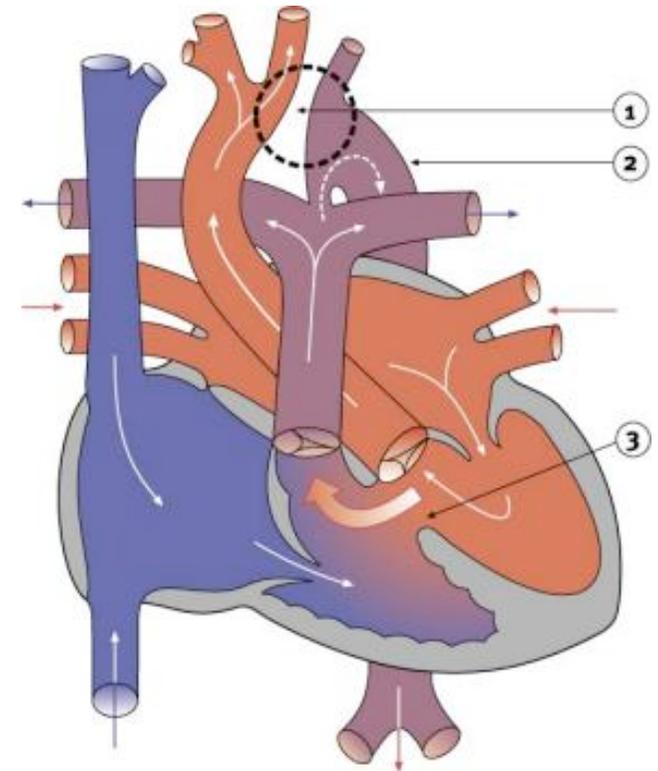
- No Ventricular Disproportion

– ie $LV = RV$

- Great vessel disproportion

– ie $Asc\ Ao \ll PA$

- Have to have ventricular septal defect



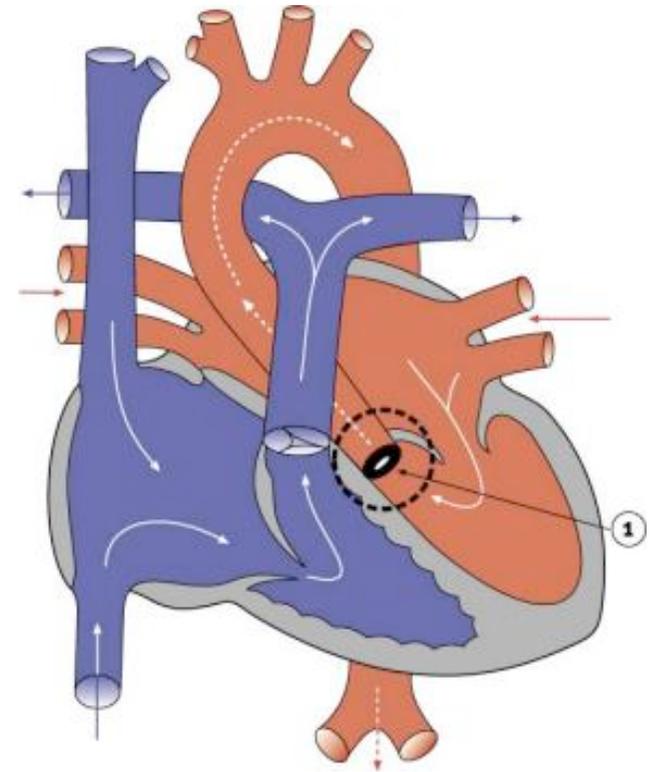
Aortic Valve abnormalities

Common in children

Less common in fetus

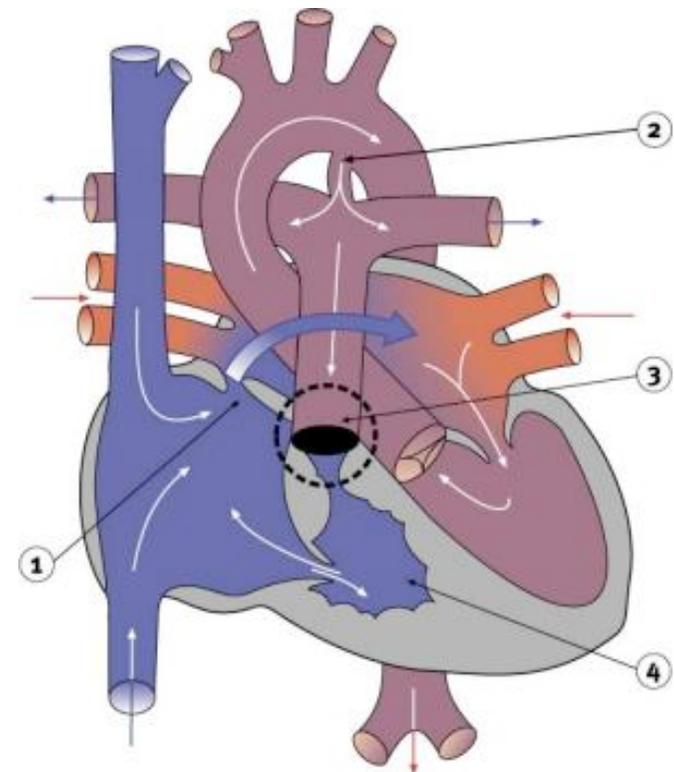
Aortic atresia/stenosis most commonly encountered aortic valve abnormality in fetal life

Assessed by Doppler (normal 30-100 cm/sec)



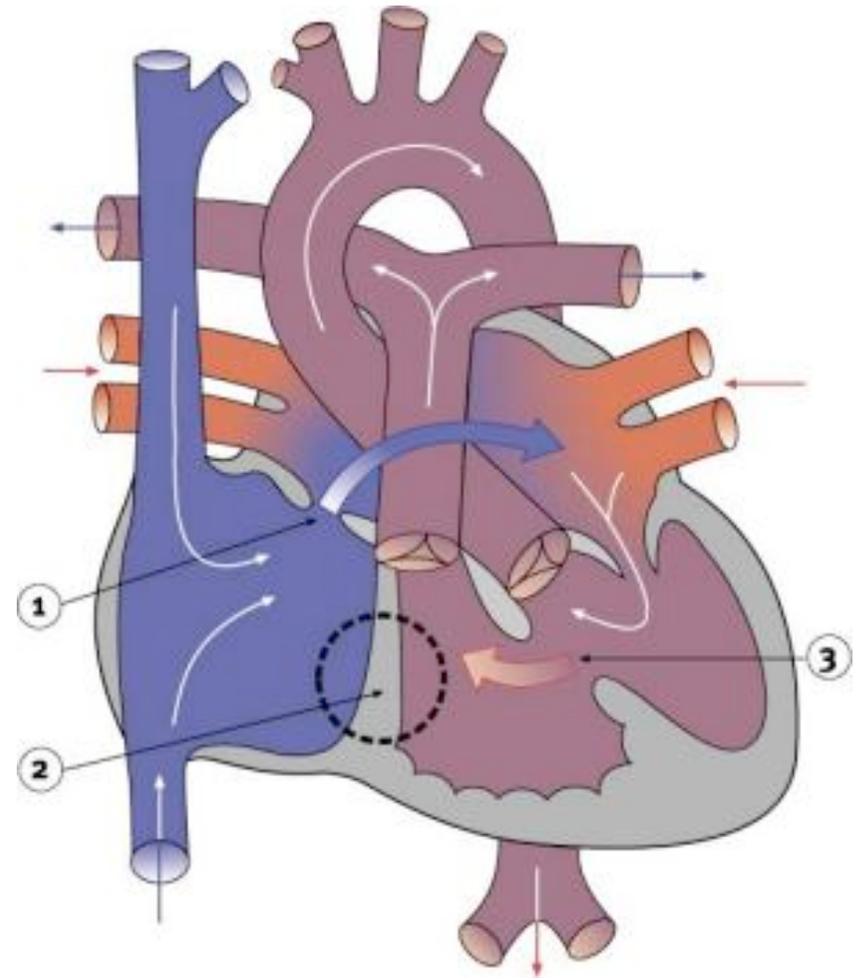
Pulmonary atresia

- Wide spectrum of disorders
- ± VSD
- ± Abnormal four chamber
- No forward flow out of RV



Tricuspid atresia

- No forward flow from RA to RV
- Always has VSD
- Abnormal four chamber
- Great arteries normally related OR transposed



Atrioventricular Septal Defect

- 50% of Trisomy 21 have congenital heart disease-
50% of these have AVSD
- ** Look for other features of Chr abnormality**

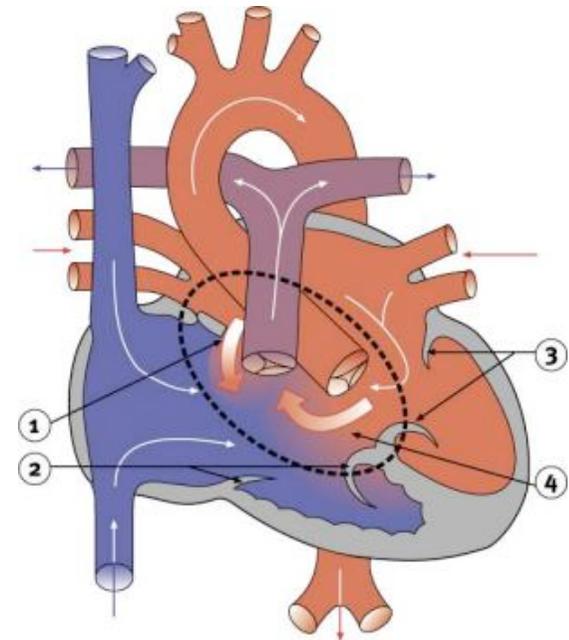
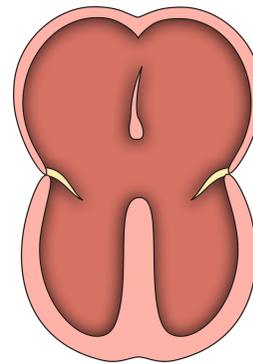
Cardinal Features

Common AV valve

Loss of offset of AV valves

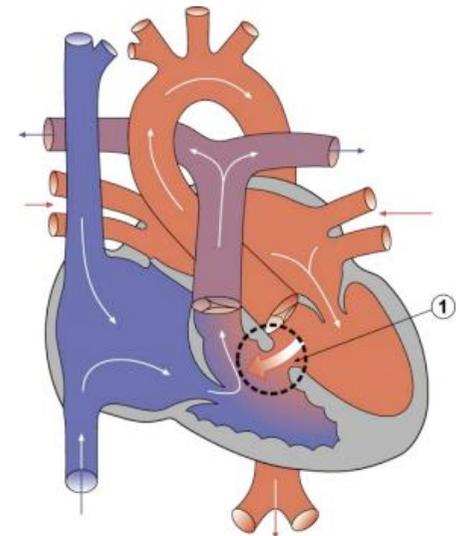
Absence of “primum septum”

Can be assoc with “laterality defects” so
check abdominal situs



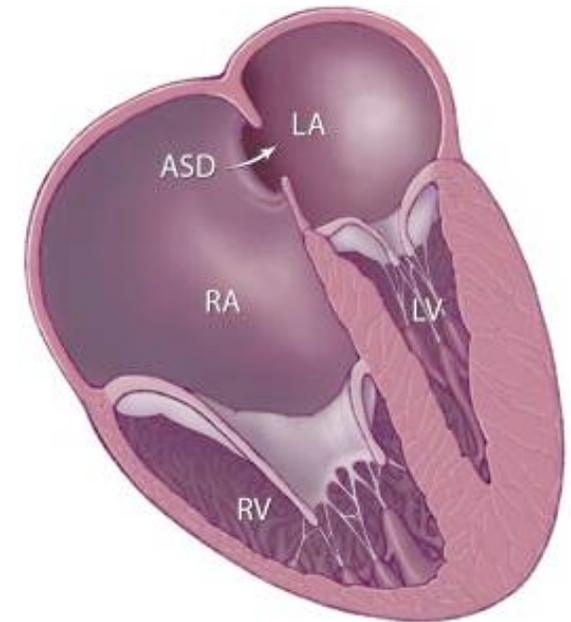
Ventricular Septal Defects

- Commonest form of CHD in infancy- 20% of all CHD
- Usually on only moderate sized/large seen in fetus
- Isolated/Associated with complex CHD
- Highly associated with extracardiac / chromosomal abnormalities
 - Perimembraneous (inlet/outlet)
 - Muscular
 - (Juxta-arterial)



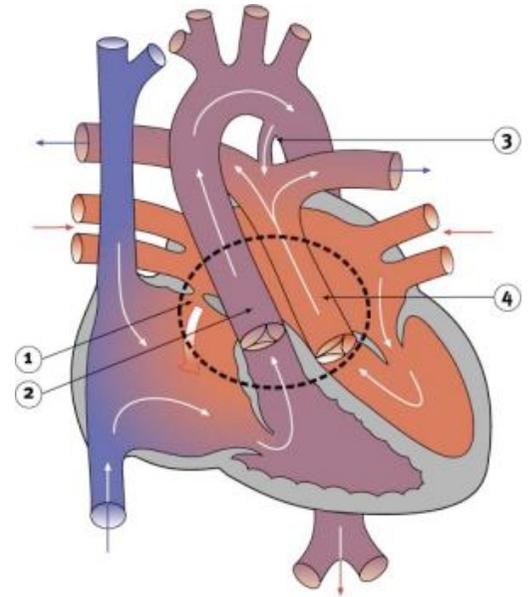
Ebstein's Anomaly/Dysplastic TV

- Displacement of Tricuspid valve leaflets of towards apex
- Worst end of the spectrum detected in the fetus,
- Significant Tricuspid regurgitation
 - RV volume load
 - Cardiomegaly- “Wall to wall heart”
 - Lung hypoplasia
 - Reduced PA flow
 - Atrial stretch- arrhythmia



Transposition of Great Arteries

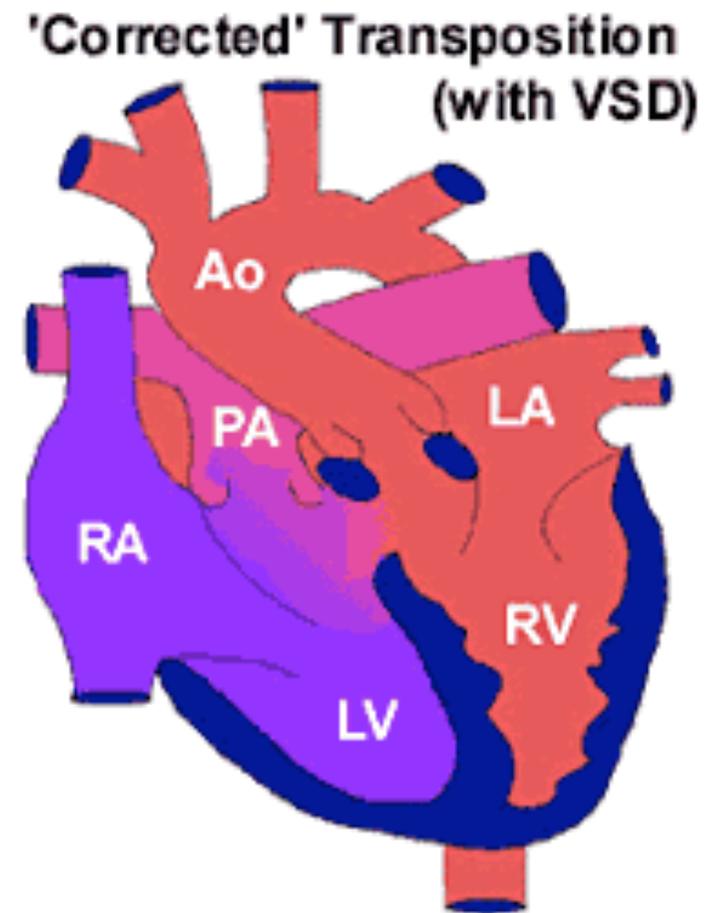
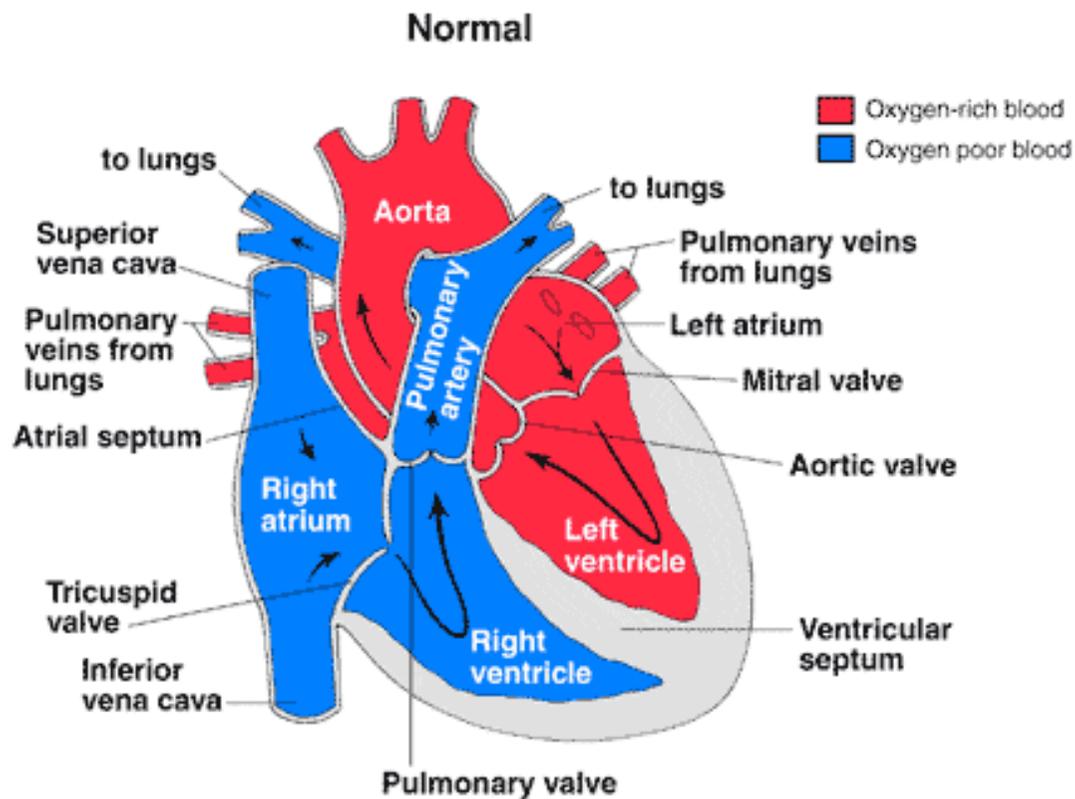
- SVC/IVC → RA → RV → Aorta
- Pulm veins → LA → LV → Pulm artery
- Great arteries arise in parallel arrangement
- Deoxygenated blood around body
- Up to 20% need urgent intervention
- Strong indication for delivery in Centre



Congenitally corrected Transposition of Great Arteries

- Rare
- “Double discordance” / Ventricular Inversion
- SVC/IVC → RA → LV → Pulm Artery
- Pulm veins → LA → RV → Aorta
 - Often have VSD
 - Association with Complete Heart Block
 - Assoc with valve abnormalities- pulmonary stenosis/
atresia, Ebstein’s anomaly

Congenitally corrected Transposition of Great Arteries



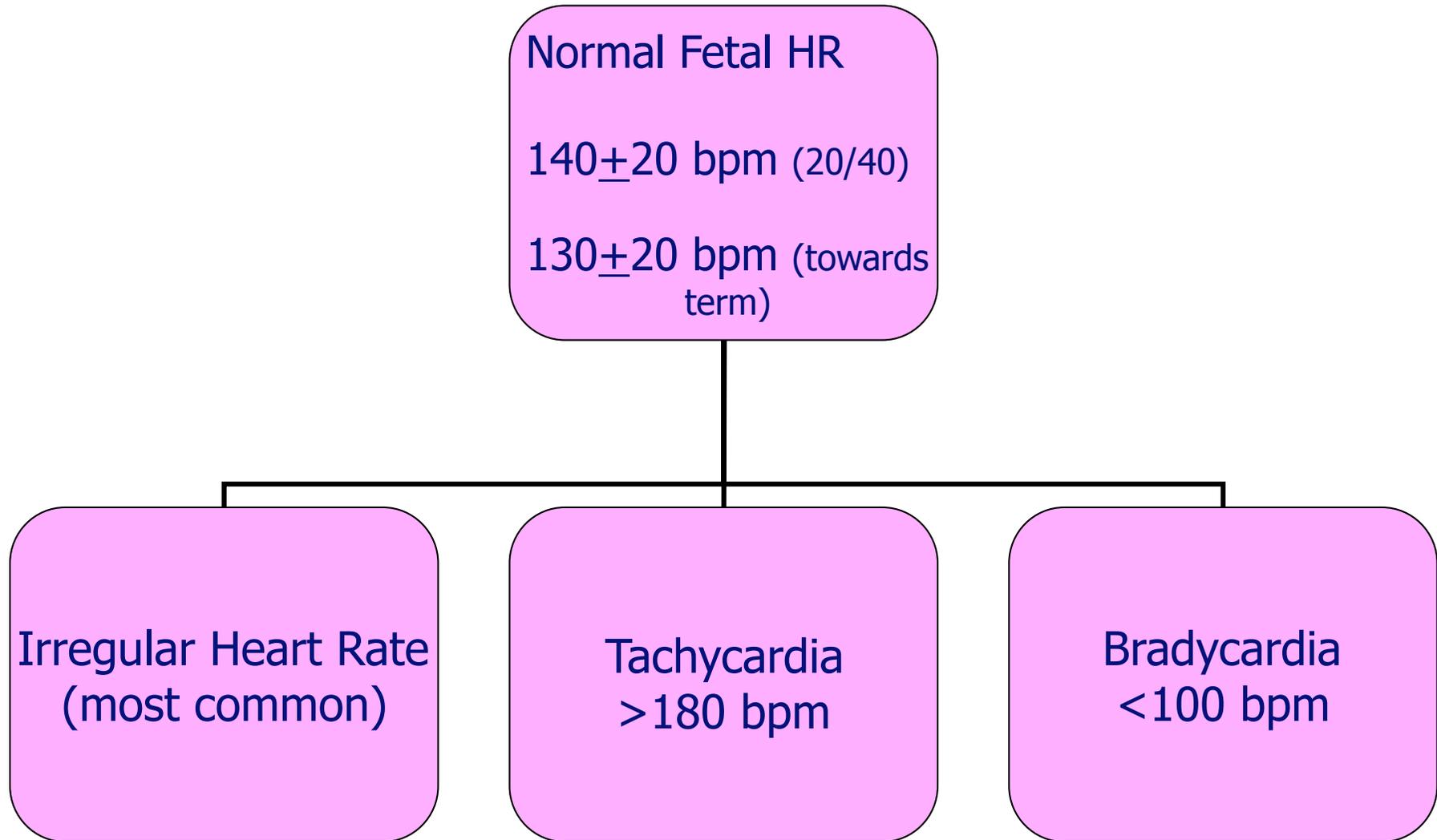
Tetralogy of Fallot ,Common Arterial Trunk, Double outlet RV

- “Conotruncal abnormalities” (also Pulm atresia)
- 30-40% association with 22q11 micordeletion
“DiGeorge Syndrome”
 - Learning difficulties, skeletal abn, Cleft lip/palate,
absent thymus
- Ventricular septal defect, Deviation of outlet
septum, Aortic override

Echogenic Foci

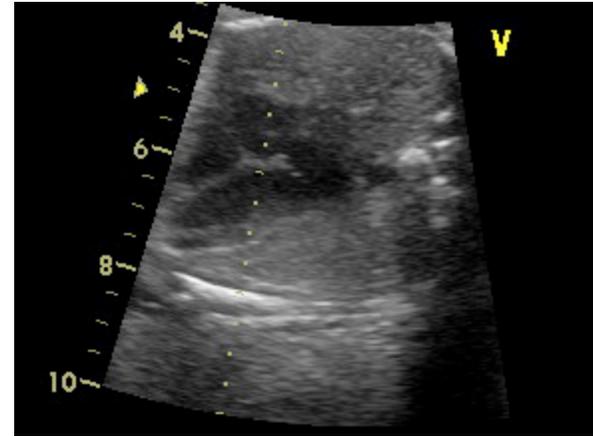
- Also know as “golf balls”, “peas”
- First described in mid 80’s
- Echogenicity similar to bone
- Incidence 3-8%
- Single, multiple, usually single focus LV
- Assoc with chromosomal abn-

Fetal Arrhythmias

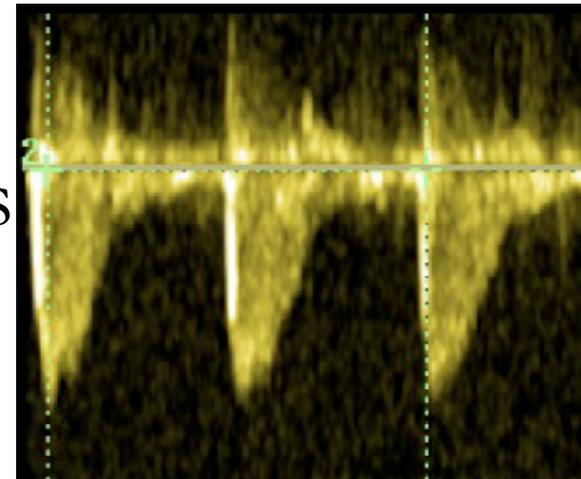


Evaluation of Arrhythmias

- M-mode Echocardiography
 - Atrial Wall contraction
 - Ventricular Systole



- Doppler interrogation of valves



Irregular Heart Rhythm/ Ectopics beats

- Most common “arrhythmia” esp in late pregnancy
- 1.7% of fetuses 36-41 weeks
- Atrial and Ventricular Extrasystoles
- Not associated with fetal hypoxia, distress or adverse perinatal outcome

Irregular Heart Rhythm -2

- Infrequent ectopic beats benign
 - 2% progress to tachy, 13% if blocked atrial ectopics
 - > 1 in 10 ectopics higher risk of arrhythmias
 - DON'T NEED REFERRAL TO CARDIOLOGY
- Monitor FHR 2 weekly by auscultation

Fetal Tachycardia

- Most commonly Supraventricular Tachycardia
- Also Atrial Flutter
- Ventricular Tachycardia incredibly rare
- Important cause of fetal morbidity and mortality
 - hydrops, cardiac failure, neurological, IUD
- Management
 - Transplacental (ie.maternal oral therapy)
 - Direct Fetal injection
 - Early delivery?

Fetal Bradycardia

- Most commonly due to CHB
Maternal connective tissue disease- SLE
- Prolonged sinus brady –may be sign of fetal distress
- Structural heart disease
- Long QT Syndrome

In Summary...

- Identify High risk groups
- 4 Chamber + Outlet views- 80% abnormalities
- Don't have to make diagnosis of complex CHD
- “Say what you see”
- Don't be afraid to ask opinion
- ** Echogenic Foci and Atrial Ectopics BENIGN