CONGENITAL HEART DISEASES



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FETAL CIRCULATION



CONGENITAL HEART DISEASES (CHD)

These are cardiac anomalies arising as a result of a defect in the structure or function of the heart and great vessels which is present at birth



These lesions either obstruct blood flow in the heart or vessels near it, or alter the pathway of blood circulating through the heart

CLASSIFICATION OF CHD



Atrial Septal Defect (ASD)

- An abnormal opening in the atrial septum which allows oxygenated blood from the left atrium to mix with deoxygenated blood in the right atrium at a minor pressure difference
- Right atrium recieves blood from SVC,IVC as well as from left atrium leading to volume overload and pulmonary congestion

- Occurs in about 4-10% of CHD
- More common in female child



I Types of ASD:

Ostium Secundum

most common- 50-70%,
In the middle of the septum in the region of the foramen ovale
Ostium primum
30% -Low position
Form of AV septal defect
Sinus venosus
Least common-10%
Site-at entry of superior venacava into right atrium



Clinical Presentation:

 Most infants and children are asymptomatic but over years to decades may experience the symptoms depending on type and severity of ASD

Infant gets tired during feeding
 Child gets tired with playing/eating
 Shortness of breath
 Fatigue
 Sweating
 Palpitations
 Stunted growth

Diagnosis:

On Auscultation-✓S1 :normal ✓S2: Widely split & fixed with P2 accentuated

Ejection systolic murmur is present



 Chest X-ray - Mild to moderate cardiomegaly with enlarged right atrium & right ventricle, prominent pulmonary artery segment, increased pulmonary vascular markings



ECG- Right Axis Deviation, Right ventricular strain pattern in lead V₁

Echocardiogram- position, size, signs of LDR shunt, flow



- Anagement :
- 20% of atrial septal defects will close spontaneously in the first year of life or as the child grows
- For defects of 3-8mm, or smaller, supportive medical management – Digoxin, diuretics and prophylactic antibiotics are sufficient up till spontaneous closure
- If defect is >8mm, surgical repair may be is required
- If spontaneous closure does not occur by schoolgoing age, surgical repair becomes essential to prevent lung problems that will develop from longtime exposure to extra blood flow
- Surgical repair- defect may be closed with stitches or a special patch. The material utilized for patch closure of ASD's may be the patient's own pericardium, commercially available bovine pericardium, or syn_{Dr}t_Nh_{id}e_{hi}t_Ai_hc_{ya(}m_{Ass}a_{tP}t_{eof}rial 13



Ventricular Septal Defect (VSD)

- An abnormal opening in the ventricular septum which allows oxygenated blood from the left ventricle to mix with deoxygenated blood in the right ventricle
- Right ventricle recieves blood from right atrium as well as from left ventricle leading to volume overload and pulmonary congestion
- VSDs are the most commonly occurring type of congenital heart defect, occurring in 14-17 % of babies born each year



I Types of VSD:

Supracristal VSD

occurs just beneath the aortic valve at the left ventricular outflow tract • Membranous VSD

- The most common type and originate inferior to the crista supraventricularis • Muscular VSD
- Occur in the mid to apical interventricular septum



Clinical Presentation: with the size of Signs and symptoms vary defect. the Clinical symptoms are usually not seen at birth continued high because of pulmonary vascular resistance in the newborn Infants with moderate to large defects will become symptomatic within the first few weeks of life. Shortness of breath while feeding Poor growth Failure to gain weight Pounding Heart Frequent respiratory tract infections ✓ If reversal of shuntoccurs- cynosis, clubbing, respiratory distress 18

Diagnosis:

On Auscultation Pansystolic murmur is present
 S1 is masked by the murmur
 S3 can be heard at the apex



Chest X-ray- Cardiomegaly and incresed pulmonary vascular markings



Management :

Medical management

digoxinDiuretics

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Adequate nutrition
 high-calorie formula or breast milk
 supplemental tube feedings

 Prophylactic antibiotics to prevent bacterial endocarditis



- Surgical repair closed stitches or special patch
- Interventional cardiac catheterization – Septal occluder
- Outcome of Surgery- 95% success rate, growth of child catches up in 1-2 years, size of the heart reduces, murmurs can be heard 2-3 months postoperative also but hold very little clinical importance

Patent Ductus Arteriosis (PDA)

- Failure of closure of ductus arteriosus
- Incidence: Mostly in premature infants or infants born to a mother who had rubella during the first trimester of pregnancy
- Through the PDA □ oxygenated blood passes from the aorta to the pulmonary artery & mixes with the deoxygenated blood which goes to the lungs □ □ blood volume to the lungs □ pulm_tonary hypertension & congestion



 As blood is pumped at high pressure through the PDA, the lining of the pulmonary artery will become irritated and inflamed. Bacteria in the bloodstream can easily infect this injured area bacterial endocarditis



Clinical Presentation:

- Shortness of breath
- Congested breathing
- Disinterest in feeding, or tiring while feeding
- Poor weight gain
- Sweating
- Tachypnea
- Bounding pulse

Diagnosis:

On Auscultation Continuous machinery murmur in the left infraclavicular region



Management: Medical Management

 Indomethacin IV (prostaglandin inhibitor) may help close a PDA. It works by stimulating the muscles inside the PDA to constrict, thereby closing the connection

- •Digoxin
- Diuretics
- Adequate nutrition
- •High-calorie formula or breast milk
- Special nutritional supplements may be added to formula or pumped breast milk that increase the number of calories in each ounce

Surgical Management

 Repair is usually indicated in infants younger than 6 months of age who have large defects that are causing symptoms, such as poor weight gain and rapid breathing

Transcatheter coil closure of the PDA

 PDA ligation-involves closing the open PDA with stitches or the vessel connecting the aorta and pulmonary artery may be cut and cauterized

CYNOTIC HEART DISEASE

- These type of defects lead to either increased or decreased pulmonary blood flow
- The primary pathology arises either due to an obstructive lesion; or due to abnormal anatomy or both
- The shunt present is predominantly from Right to Left leading to shunting of venous blood without passing through the lungs to be oxygenated

Unoxygenated blood circulates in arteries I cyanosis

Example: Tetralogy of Fallot, TGV

Tetralogy of Fallot (TOF)

- A complex condition of several congenital defects that occur due to abnormal devlopment of the fetal heart during the first 8 weeks of pregnancy
- 'Tetra' meaning 'four'
 Ventricular septal defect (VSD)
 Pulmonary valve stenosis
 Overriding aorta
 Right ventricular hypertrophy





- Due to pulmonary artery stenosis, RV has to work harder to push blood into the lungs, thereby increasing the RV pressure and size
- Presence of VSD facilitates blood to pass from the RV into the left ventricle, and mixing of blood takes place.
- Overriding of aorta- The aorta sits above both the left and right ventricles over the VSD, rather than just over the left ventricle. As a result, oxygen poor blood from the right ventricle can flow directly into the aorta instead of into the pulmonary artery to the lungs
- Decresed pulmonary blood flow and poorly oxygenated blood cirr ciulati(ng trhof)rough out the body leads to CYNOSIS

Clinical Presentation:

- Cyanosis (bluish color of the skin, lips, and nail beds) that occurs with such activity as crying or feeding
- Irritability
- Lethargic
- Reduced physical activity
- Fainting
- Clubbing of nails of fingers/toe
- Breathing difficulty

Children with Tetralogy of Fallot exhibit bluish skin during episodes of crying or feeding.



Diagnosis:

 On Auscultation- An ejection systolic murmur is present at the Left parasternal region 3rd ICS due to pulmonary stenosis.



- Management :
- Requires surgical repair usually undertaken at 6-18 months age
- It involves- closure of VSD with a tangential patch to correct the override and the pulmonary stenosis is relieved with a patch).



Transposition of Great Vessels (TOG)

- The aorta is connected to the right ventricle, and the pulmonary artery is connected to the left ventricle
- Oxygen-poor (blue) blood returns to the right atrium from the body
 passes through the right atrium and ventricle,
 into the misconnected aorta back to the body.
 - Oxygen-rich (red) blood returns to the left atrium from the lungs [] passes through the left atrium and ventricle, [] into the pulmonary artery and back to the lungs.



Clinical Presentation:

Cyanosis - (bluish color of the skin, lips, and nail beds) that occurs with such activity as crying or feeding
Rapid and laboured breathing
Cold and clammy skin
Failure to thrive

Management:

- Admitted to NICU
- On ventilator support
- Cardiac Catheterization
- Ballon Atrial Septostomy
- I.v. Prostaglandins administered
- By 2nd week of life, TGA repair
 is done
- 'Switch' operation



Summary

- Fetal Circulation Congenital Heart
- Diseases
- Classification of CHD
- Hemodynamics of Common CHD
- Assessment & Management



QUESTIONS

- 1. WRITE ABOUT FATEL CIRCULATIONS.
- 2. GIVE THE CLASSIFICATION OF CHD.

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3. WRITE THE ASSESMENT AND MANGEMENT OF THE CHD



Thank you...!!!