## Chapter 26

- The Child with a Cardiovascular Disorder
- Objectives
- Distinguish the difference between the cardiovascular system of a child and an adult.
- List the general signs and symptoms of congenital heart disease.
- Differentiate among patent ductus arteriosus, coarctation of the aorta, atrial septal defect, ventricular septal defect, and tetralogy of Fallot.
- Discuss six nursing goals relevant to the child with heart disease.
- Objectives (cont.)
- List the symptoms of rheumatic fever.
- Discuss the prevention of rheumatic fever.
- Discuss hypertension in childhood.
- Differentiate between primary and secondary hypertension.
- Objectives (cont.)
- Identify factors that can prevent hypertension.
- Recognize the manifestation of Kawasaki disease and the related nursing care.
- Describe heart-healthy guidelines for children older than 2 years.
- Cardiovascular System
- Signs Related to Suspected Cardiac Pathology
- Failure to thrive and/or poor weight gain
- Cyanosis, pallor

•	Visually observed pulsations in the neck veins
•	Tachypnea, dyspnea
•	Irregular pulse rate
•	Clubbing of fingers
•	Fatigue during feeding or activity
•	Excessive perspiration, especially over forehead
•	Congenital Heart Disease
•	Occurs in approximately 8 out of 1,000 births
•	50% of these infants show signs/symptoms within the first year of life
•	Can be caused by genetic, maternal, or environmental factors
	Not a problem for the fetus because of the fetal-maternal circulation
	At birth, the infant's circulatory system must take over and provide the child's oxygen needs
•	Congenital Heart Disease (cont.)
•	Of the congenital anomalies, heart defects are the principal cause of death during the first year of life
•	Diagnostic studies vary from noninvasive, such as an electrocardiogram, to invasive, such as angiogram
•	Once diagnosis is confirmed, most cardiac defects require surgical intervention
•	Congenital Heart Disease (cont.)
•	Classification
•	Two categories
	— Cyanotic

	Acyanotic
•	Blood always flows from area of high pressure to an area of low pressure and it also takes the path of least resistance
•	Congenital Heart Disease (cont.)
•	Physiologically, defects can be organized into lesions that
	Increase pulmonary blood flow
	Obstruct blood flow
	Decrease pulmonary blood flow
•	A <i>shunt</i> refers to the flow of blood through an abnormal opening between two vessels of the heart
•	The Normal Heart and Various Congenital Heart Defects
•	Congenital Heart Disease (cont.)
•	Defects that increase pulmonary blood flow
	Blood returns to the right ventricle and recirculates through the lungs before exiting the left ventricle through the aorta
•	Some defects that increase pulmonary flow are
	Atrial septal defect
	Ventricular septal defect and patent ductus arteriosus
•	The oxygenated blood recirculates to the lungs, and cyanosis is rare
•	Nursing Tip
•	In congenital heart disease, cyanosis is <i>not</i> always a clinical sign
•	Restrictive Defects
•	Restriction usually from some form of stenosis of the vessel

 Coarctation	of the	aorta
Coarciation	or the	аопа

- Narrowing or constriction of the aortic arch or of the descending aorta
- Hemodynamically, increased pressure proximal to the defect and decreased pressure distally
- Coarctation of the Aorta
- Characteristic symptoms
- Marked difference in the blood pressure and pulses of the upper and lower extremities
- May not develop symptoms until late childhood
- Treatment is dependent upon type and severity of the defect
- Best time for surgical intervention is between 2 and 4 years of age
- Coarctation of the Aorta (cont.)
- If left untreated
  - Hypertension
  - Congestive heart failure
  - Infective endocarditis may occur
- After surgery, the nurse should observe for
  - Hypertension
  - Abdominal pain associated with nausea and vomiting
  - Leukocytosis
  - GI bleeding or obstruction
- Treatment includes
  - Antihypertensive drugs
  - Steroids
  - NG tube for decompression of the stomach
- Defects that Decrease Pulmonary Blood Flow

•	Occurs when a congenital heart anomaly allows blood that has not passed through the lungs (unoxygenated blood) to enter the aorta and general circulation
•	Cyanosis caused by the presence of unoxygenated blood in the circulation is a characteristic feature of this type of congenital heart anomaly
•	Tetralogy of Fallot
•	Four defects
	Stenosis or narrowing of the pulmonary artery
	Decreases blood flow to the lungs
	Hypertrophy of the right ventricle
	<ul> <li>Enlarges because it must work harder to pump blood through the narrow pulmonary artery</li> </ul>
	• Dextroposition of the aorta
	The aorta is displaced to the right and blood from both ventricles enters it
	Ventral septal defect (VSD)
•	Tetralogy of Fallot (cont.)
•	Cyanosis increases with age
•	Clubbing of fingers and toes
	— Due to chronic hypoxia
•	Child rests in a "squatting" position to breathe more easily by altering systemic venous return
•	Prevalent symptoms include
	Feeding problems
	Failure to thrive
	Frequent respiratory infections

	Severe dyspnea on exertion
	Polycythemia develops to compensate for the lack of oxygen
•	Paroxysmal Hypercyanotic Episodes
•	Known as Tet spells
•	Occur during the first 2 years of life
•	Spontaneous cyanosis, respiratory distress, weakness, and syncope occur
•	They can last up to a few hours and are followed by lethargy and sleep
•	Place child in knee-chest position when Tet spell occurs
•	Tet Position
•	Diagnosis
•	Diagnosis confirmed by chest X-ray that shows a typical boot-shaped heart
•	Additional tests include
	EKG
	3-D echocardiography
	Cardiac catheterization
•	Complications and Treatments
•	Complications
	Cerebral thrombosis caused by polycythemia, especially if dehydration occurs
	Iron-deficiency anemia due to decreased appetite and increased energy required to suck or eat
	Bacterial endocarditis can occur
•	Treatment

	Designed to increase pulmonary blood flow to relieve hypoxia	
•	Surgery	
	In some cases, IV prostaglandin E therapy can open a constricted ductus arteriosu and allow for oxygenation of the body until surgery is performed	s
•	Defects that Cause Mixed Pathology	
•	Hypoplastic left heart syndrome	
	Underdevelopment of the left side of the heart	
	Usually results in an absent or nonfunctional left ventricle and hypoplasia of the ascending aorta	
	Can be diagnosed before birth and infant is placed on a heart transplant list early	
•	Defects that Cause Mixed Pathology (cont.)	
•	Hypoplastic left heart syndrome (cont.)	
	Initial survival depends on a patent foramen ovale and ductus arteriosus to provide pathway for oxygenated blood to the general body system	а
	Symptoms include	
	A grayish-blue color of the skin and mucous membranes	
	Signs of CHF	
	<ul> <li>Dyspnea</li> </ul>	
	<ul> <li>Weak pulses</li> </ul>	
	Cardiac murmur	
•	General Treatment and Nursing Care	
•	Assorted medical and surgical treatments are currently available	
•	After the procedure, the nursing care involves	
	Monitoring vital signs	

Observing for thrombosis formation
Neurovascular checks of the limb
Emotional support to child and family
General Treatment and Nursing Care (cont.)
Instruct parents that children with congenital heart disease should avoid competitive sports because the pressure for a team win can interfere with the child's need to stop activity if specific symptoms arise
Nutritional guidance aimed at preventing anemia and promoting optimal growth and development
Vacations to high altitudes or very cold environments may cause adverse responses in a child who is already hypoxic or has cardiac problems
Acquired Heart Disease
Occurs after birth
May be a complication of a congenital heart disease or a response to respiratory infection, sepsis, hypertension, or severe anemia
Heart failure is a decrease in cardiac output necessary to meet the metabolic needs of the body
Congestive Heart Failure (CHF)
Manifestations depend on the side of the heart affected
Right side of the heart moves unoxygenated blood to the pulmonary circulation
A failure results in the backup of blood in the systemic venous system

Left side of heart moves oxygenated blood from the pulmonary circulation to the systemic circulation

Congestive Heart Failure (CHF) (cont.)

Failure results in backup into the lungs

When body tries to compensate

	Peripheral vasoconstriction occurs
	Results in cold and/or blue hands and feet
_	Tachycardia
_	Tachypnea
Safe	ty Alert
Early	y signs of CHF in infants that should be reported
_	Tachycardia at rest
_	Fatigue during feedings
_	Sweating around scalp and forehead
_	Dyspnea
_	Sudden weight gain
CHF	Goals of Treatment
Goa	ls
_	Reduce the work of the heart
	Improve respiration
_	Maintain proper nutrition
_	Prevent infection
_	Reduce the anxiety of the patient
_	Support and instruct the parents
CHF	and Nursing Care
Orga	anize care so that infant is not unnecessarily disturbed

Feed early if crying and late if asleep

•	Feedings are small and frequent
•	Oxygen is administered to relieve dyspnea
•	Medications are given as prescribed, after dosages are checked for safety
•	Accurate recording of intake and output
•	Rheumatic Fever (RF)
•	Systemic disease involving the joints, heart, central nervous system, skin, and subcutaneous tissues
	Belongs to a group of disorders known as collagen diseases
•	Common feature is destruction of connective tissue
	Scars mitral valve in the heart
•	Peak incidence is 5 to 15 years of age
	More prevalent in winter and spring
•	Autoimmune disease occurring as a complication of an untreated group A beta hemolytic streptococcus infection of the throat
•	Manifestations of RF
•	Modified Jones Criteria
•	Minor criteria
	— Fever
	— Arthralgia
	Previous history of rheumatic heart disease
	Elevated erythrocyte sedimentation rate
	Leukocytosis
	Altered PR interval on electrocardiogram
	Positive C-reactive protein

•	A positive diagnosis of RF cannot be made without the presence of two major criteria or one major and two minor criteria, <i>plus</i> a history of streptococcal infection
•	Modified Jones Criteria (cont.)
•	Major Criteria
	— Carditis
	Polyarthritis
	Erythema marginatum
	— Chorea
	Subcutaneous nodules
•	Treatment of RF
•	Antimicrobial therapy initially, then followed by chemoprophylaxis monthly for a minimum of 5 years
•	Rest
•	Relief of pain and fever
	Antiinflammatory agents
	Steroids
	— Aspirin
•	Management of cardiac failure, should it occur
•	Nursing Care of RF
•	Care should be organized to ensure as few interruptions as possible to prevent tiring the patient
•	Special attention should be given to skin and back care; good oral hygiene; and small, frequent feedings

•	If dental therapy is needed, prophylactic antimicrobial treatment is required before the procedure
•	Prevention of RF
•	Prevention of infection and prompt treatment of group A beta-hemolytic streptococcal infections
•	Nurse stresses importance of completing all antimicrobial therapy as prescribed
•	Systemic Hypertension
•	More prevalent during childhood and adolescence
•	Significant hypertension (HTN) is considered when measurements are persistently at or above the 95 <sup>th</sup> percentile for patient's age and sex
•	Primary, or essential, HTN implies that no known underlying disease is present
•	When the cause of hypertension can be explained by a disease process, it is known as secondary
	Renal, congenital, vascular, and endocrine
•	Systemic Hypertension (cont.)
•	Heredity, obesity, stress, and poor diet and exercise patterns are some of the contributing factors to the development of HTN
•	HTN more prevalent in children whose parents have high blood pressure
•	Systemic Hypertension (cont.)
•	Treatment and nursing care involve
	— Nutritional counseling
	Weight reduction
	Age-appropriate program of aerobic exercise
	Adolescents should be counseled concerning the adverse effects of drugs, alcohol, and tobacco on blood pressure

- Focus of treatment of secondary HTN is the underlying disease causing the elevated blood pressure
- Nonpharmacological Approach to HTN
- Aerobic exercise
- Reduce sedentary activities
- Weight reduction
- Dietary management
- Adequate intake of potassium and calcium
- Avoid smoking and those who smoke
- Hyperlipidemia
- Refers to excess lipids (fat and fatlike substances in the blood)
- Lipoproteins contain lipids and proteins and include
  - Low-density lipoproteins (LDL) contain low amounts of triglycerides, high levels of cholesterol, and some protein
    - Carries cholesterol to the cells, which aids in cellular metabolism and steroid production
  - High-density lipoproteins (HDL) contain low amounts of triglycerides, little cholesterol, and high levels of protein
    - Carries cholesterol to the liver for excretion
- Hyperlipidemia (cont.)
- Children with two consecutive blood cholesterol levels exceeding 170 mg/dL should be followed closely and offered nutritional guidance
  - Parental history of cholesterol levels exceeding 240 mg/dL or a family history of early cardiac death (under age 55 years) should have their cholesterol levels tested
- Dietary intake of no more than 300 mg of cholesterol per day and no more than 30% total dietary calories from fat are recommended

<ul> <li>Children younger than 2 years of age should not have a fat-restricted diet, because calories and fat are necessary for CNS growth and development</li> </ul>
Kawasaki Disease (KD)
Also known as mucocutaneous lymph node syndrome
Leading cause of acquired cardiovascular disease in the U.S.
Usually affects children younger than 5 years of age
May be a reaction to toxins produced by a previous infection with an organism such as Staphylococci
Not spread from person to person
Kawasaki Disease (KD) (cont.)
Diagnosis is made by clinical signs and symptoms, no specific lab studies
KD causes inflammation of the vessels in the cardiovascular system
Weakens the walls of the vessels
Often results in an aneurysm (an abnormal dilation of the wall of a blood vessel)
Aneurysms can cause thrombi (blood clots) to form, which can be life-threatening
Kawasaki Disease (KD) (cont.)
Manifestations
Onset is abrupt with a sustained fever
• As high as 104° F (40° C)
Does not respond to antipyretics or antimicrobials
• Fever lasts for more than 5 days
Conjunctivitis without discharge
— Fissured lips
A "strawberry tongue"

	Inflamed mouth and pharyngeal membranes
	Enlarged nontender lymph nodes
•	Kawasaki Disease (KD) (cont.)
•	Erythematous skin rash develops
•	Swollen hands and desquamation (peeling) of the palms and soles
•	Child is very irritable
•	May develop signs of cardiac problems
•	Kawasaki Disease (KD) (cont.)
•	Treatment
	IV gamma globulin, if given early, can prevent the development of coronary artery pathology
	Salicylate therapy for antithrombus properties
	Warfarin therapy may be prescribed if aneurysms are detected
•	Kawasaki Disease (KD) (cont.)
•	Nursing care
	Symptomatic and supportive
	Parent teaching should be reinforced concerning need to postpone active routine immunizations for several months after the administration of immune globulin, which is an immunosuppressant
	Long-term, low-dose aspirin therapy may be prescribed
	Compliance may be a problem for any long-term regimen in which medications must be taken when the child feels "well."
•	Question for Review
•	How does the squatting (Tet) position relieve dyspnea?

- Review
- Objectives
- Key Terms
- Key Points
- Online Resources
- Review Questions