

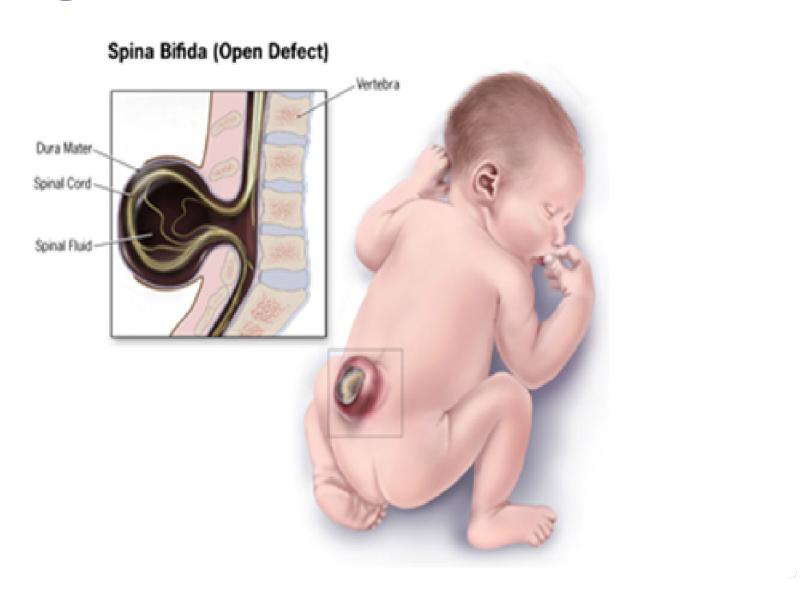
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#### FACULTY OF NURSING

## CHILD WITH CONGENITAL DISORDERS

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### SPINAL BIFIDA



#### **DEFINITION**

Spinal bifida is a neural tube defects where there is an incomplete closure of the vertebrae and neural tube.

#### OR

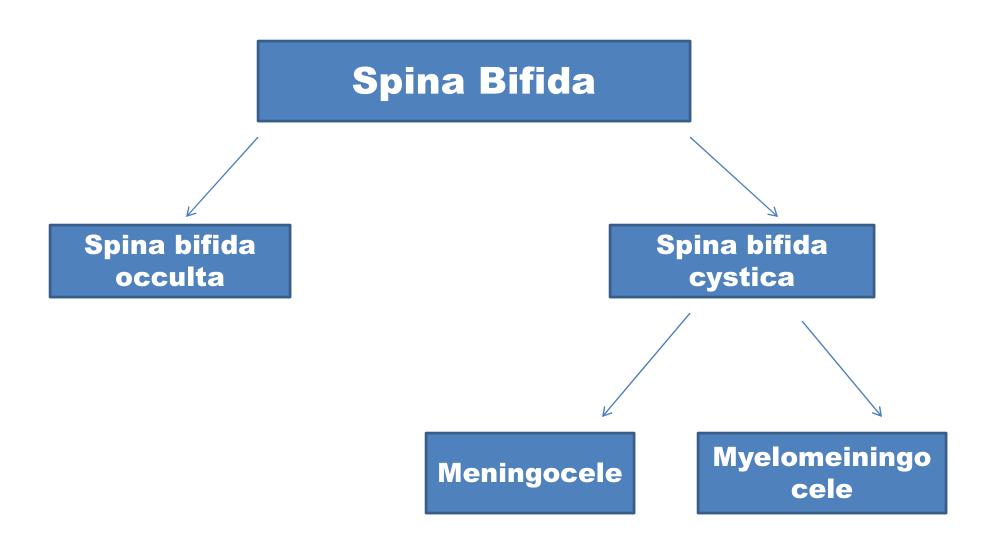
Spinal bifida is a malformation of spine in which posterior portion of lamina of vertebra fail to closure with or without defective development of spinal cord.

#### **ETIOLOGY**

- **≻** Genetic
- ➤ Maternal age (pregnant women >35 years and <20 years)
- ➤ Environmental (radiation defective neural tube dev
- ➤ Diet (folic acid deficiency
- ➤ Medication (anticonvulsa



#### **CLASSIFICATION/TYPES**



#### **TYPES OF SPINAL BIFIDA**

#### 1. Spinal Bifida Occulta

- An abnormality is confined to the vertebrae only and is due to an unclosed posterior vertebral arch.
- This has no visible defect to the external (no protrusion). It occurs most at the lumbasacral area.
- There is a dimple, hairy patch, dark spot or swelling over affected area
- Spinal cords and nerves usually normal

- It is the mildest type of spina bifida
- It has no symptoms
- There is gap in one or more of the vertebra of the spine
- No treatment needed

#### 2. Spinal bifida Cystica

- A more severe type of spinal bifida
- This refers to the visible defect with the external saclike protrusion
- It has two major forms;
- A- Meningocele
- **B- Myelomeningocele**

#### A. Meningocele

- Is a rarest form of which a cyst or fluid-filled sac pokes through an open part of the spine.
- The sac contains membranes that protect spinal cord but not spinal nerves i.e. there is no neural elements but only CSF and meninges.

#### B. Myelomeningocele

Most severe form of spinal bifida cystica

The cyst holds both membranes and nerve roots of

spinal cord and often the cord itself

Almost 96%

#### **Clinical Manifestations**

Signs and symptoms vary according to degree of spinal defect:

- Readily apparent on inspection!
- Loss of sensation below lesion
- Poor urinary and bladder control. There is an Overflow incontinence with constant dripling in urine due to nerve dysfunction that supplies the bladder.

- Joint deformities in lower extremities due to demolition to the muscles of the lower extremities.
- Orthopedic abnormalities (i.e. club foot, hip dislocation, scoliosis)
- Hydrocephalus
- Swelling
- Brain damage

#### **DIAGNOSTIC EVALUATION**

- ➤ Prenatal diagnosis-
  - -USG
  - -Fetal MRI
  - -Amniocentesis
- ➤ Diagnosis after birth-
  - Neonatal examination
  - CT scan and MRI
  - RFT
  - Urine
  - WBC count

➤ Neurological test (for motor response and sensory reaction, developmental assessment to detect any delay in milestone).

#### **MANAGEMENT**

- > Surgical management-
- a) Laminectomy (it is a surgical procedure that removes a portion of the vertebral bone called the lamina. Which is the root of spinal cord)
- b) T- closure of the second of

#### **Nursing Management**

#### Pre-Op

- Position the child in prone with legs abducted. This reduces tension and risk of sac trauma.
- Put the child in an incubator or warmer area without clothes. This maintains normal body temperature, and reduces trauma from the clothes.
- Apply dressing (moist, no adhesions), to avoid drying of the area due to heat in the incubator.
- Strictly use the sterile gauze so as to prevent reinfection.
- Change dressing two-four hourly, to avoid drying.

- Use normal saline or silver nitrate in dressing.
- Gentle handling of the child to avoid any risk of trauma.
- Change the child's position every two hours, to promote circulation and prevent development of decubitus sore.

- Check vital signs and signs of increased intracranial pressure.
- Assess for signs of hydrocephalus.
- Cover the sacrum with sterile surgical drape, but not latex tape.
- Measure the head circumference
- Prepare the mother psychologically.
- Apply gentle pressure to suprapubic area to facilitate urine emptying.
- Gently do a range of motion of the extremities to the child.

#### Post-Op

- Position the child in prone to avoid pressure on suture, or side lying position alternatively.
- Monitor the child's vital signs every 30 minutes until stable.
- Use all measures to avoid any infection e. g. hand washing.
- Monitor input and output.
- Encourage the mother to continue breastfeeding if the child is being breastfed.
- Resume the feeding after the effects of anaesthesia.
- Remove the dressings after 48hrs to check any signs of bleeding or bulging

#### **Complications**

- Meningitis due to infections
- Hydrocephalus due to Increased Intracranial Pressure
- Physical and neurological problems e.g. lack of normal bowel and bladder control, Paralysis of the legs
- Latex allergy

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