

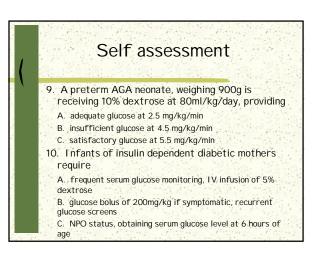
Self assessment

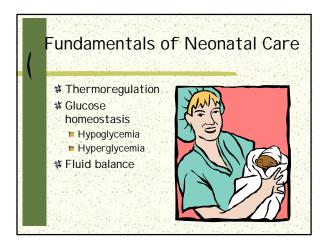
- 5. Insulated hats, often used on neonates at delivery, decrease heat loss since
 - A. a large percentage of heat production is via the brain
 - B. hats provide a conductive heat source for the neonateC. radiant warming beds will augment the insulated fabric
- A neonate with sepsis presents with hypothermia
- due to
- A. decreased ability to mount an immune response
- B. peripheral vasoconstriction to increase core temperature
- C. shock, vasodilation, loss of thermoregulatory response

Self assessment 7. Neonatal fluid overload is associated with A. chronic lung disease B. increased insensible water loss C. syndrome of inappropriate antidiuretic hormone 8. Neonatal insensible water loss can be decreased for preterm neonates by use of A. daily fluid maintenance at 180ml/kg B. incubators compared to radiant warming beds

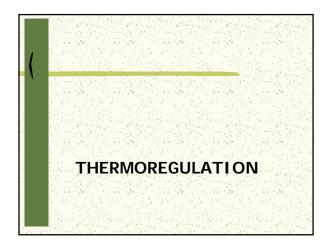
C. temperatures greater than the neutral thermal environment

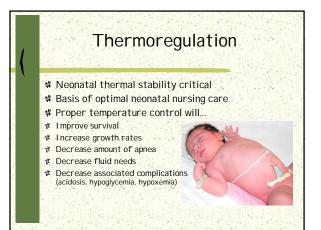
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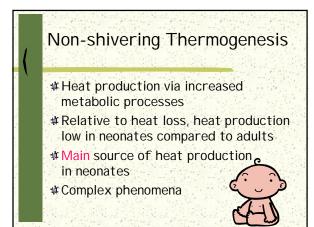




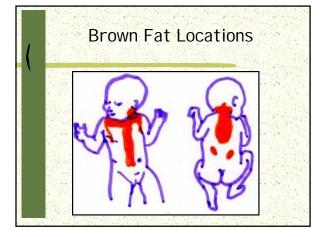






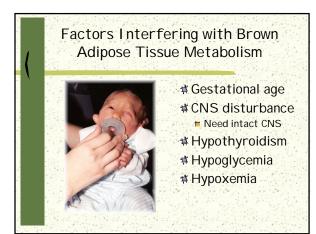


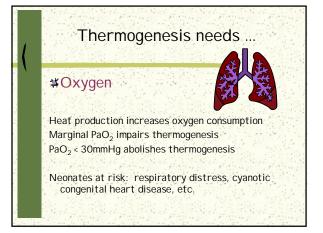
Human Temperature Regulation * Sympathetic nervous system * Preoptic area of anterior hypothalamus temperature "control" center * Heat gaining center vasoconstriction, increased metabolism, shivering, decreased sweating, increased thyroid hormone production * Heat losing center - vasodilatation, sweating, decreased muscle tone

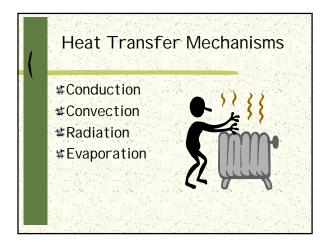


Brown Adipose Tissue

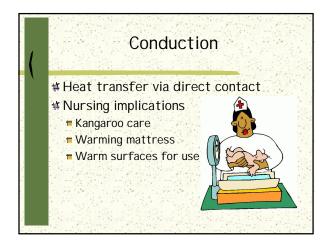
- Production begins 26-28 wks gestationStores increase until 3-5 wks after birth
- unless depleted by cold stress
- # Cannot be replenished once used
- # Metabolism initiated by skin thermal
- receptors (facial trigeminal area prominent) # Regulation by protein - thermogenin (located in mitochondria)

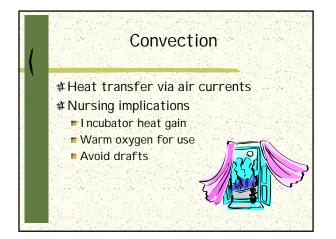




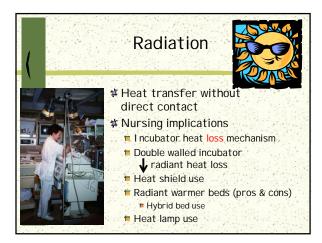


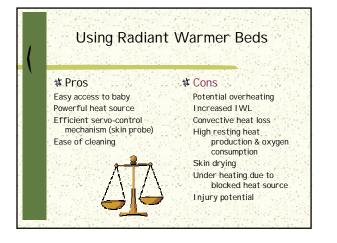


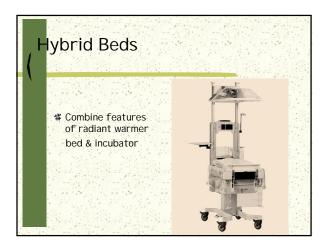




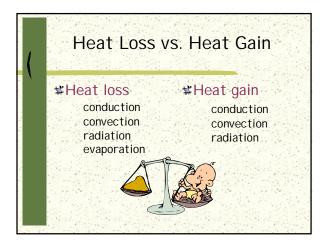


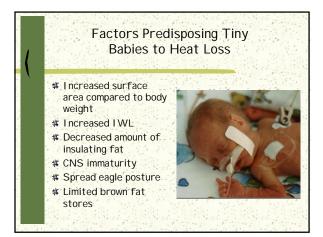


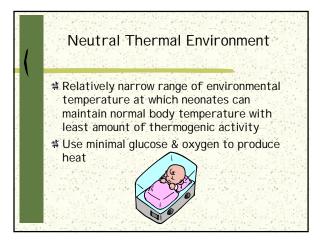


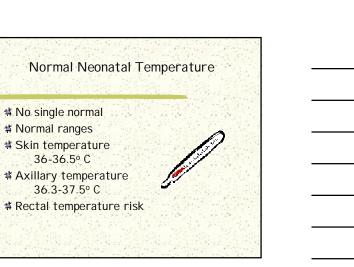


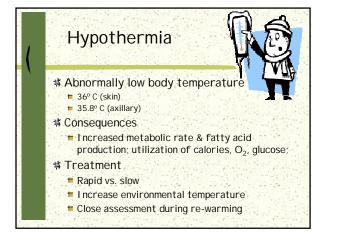


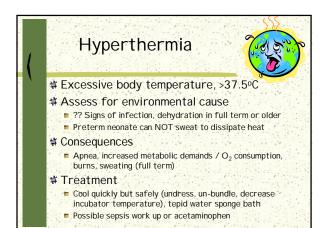


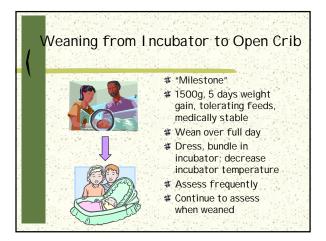


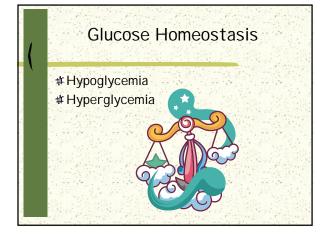


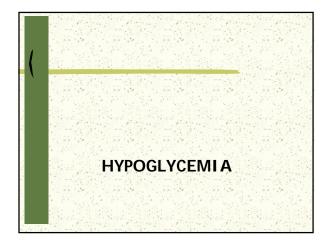














Fetal Glucose Metabolism



Glucose readily crosses placenta

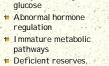
- # Fetal levels 70-80% of maternal
- # Glucose major fetal energy source
- # Glucose stored as glycogen
- # I nsulin does NOT cross placenta
- # Fetal insulin production begins 11-12 weeks

Pathogenesis



Excess utilization

- Hyperinsulinism
- Caloric demand
 - Thermoregulation Muscle activity
- Shift from aerobic to
- anaerobic
- Glucose demand by
 - tissues
- I nborn errors Acute brain injury



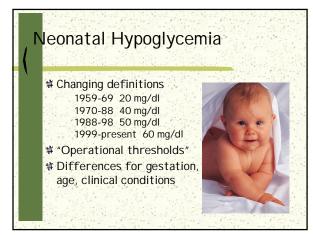
production, substrate

Delayed feeding, I V

- precursors, brain glucose transporters
- Suppressed

Inadequate

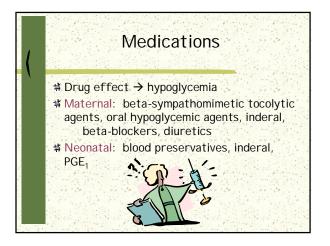
gluconeogenesis



Guidelines for Intervention

- # Asymptomatic: < 30-35 mg/dl, AGA
- \$ Symptomatic: < 45 mg/dl, signs abate with Rx
- # Sick neonates: < 45-60 mg/dl
- $\ensuremath{\sharp}$ > 24 hrs old: threshold increases to 40-50 mg/dl
- # At any age: < 20 mg/dl, need prompt IV glucose

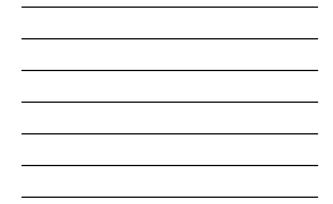
Neonates at Risk # Perinatal stress # Preterm # Fetal growth disorders # Sepsis (SGA, LBW, LGA) # Hypoxemia # Smaller of twins # Shock, hypoperfusion # Hypothermia # Severe anemia, # Maternal glucose polycythemia disorders / diabetes # Congenital anomalies # Maternal massive # Inborn errors obesity # I solated hepatomegaly # Excessive maternal # Hyperinsulinism intrapartum glucose # Some medications







Class	Age at Onset	Duration	Vascular disease	Insulir
Α	Any	Pregnancy	No	No
В	> 20 yrs	< 10 yrs	No	Yes
C	10-19 yrs	10-19 yrs	No	Yes
D	< 10 yrs	> 20 yrs	Benign retinopathy	Yes
F	Any	Any	Nephropathy	Yes
R	Any	Any	Proliferate retinopathy	Yes
Н	Any	Any	CV disease	Yes

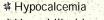


Etiology – Hypoglycemia in IDM

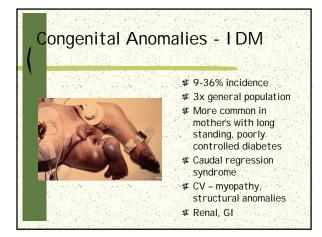
- Maternal hyperglycemia
- Fetal hyperglycemia (fetal pancreatic islet cell hypertrophy)
- Fetal hyperinsulinism (increase in hepatic glucose uptake, lipogenesis, glycogen & protein synthesis)
- # Delivery occurs
- # Neonatal hypoglycemia (glucose source ends)
- i I nadequate neonatal compensation
- * Continued hypoglycemia

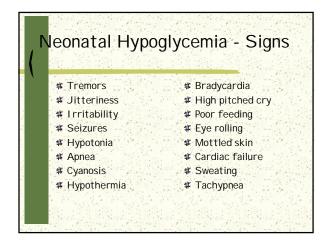
Common Problems of IDM

- Hypoglycemia
- # Altered growth
- LGA, SGA

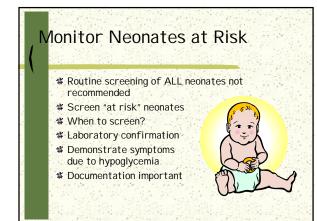


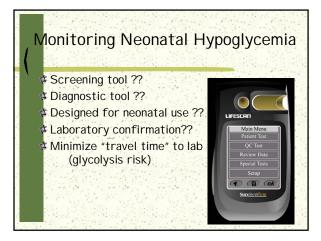
- # Hyperbilirubinemia
- # Surfactant deficiency
- # Polycythemia
- # Renal vein thrombosis
- # Congenital malformations











Asymptomatic Hypoglycemia

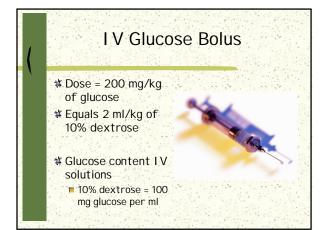
- Frequency < symptomatic hypoglycemia</p>
- # Risk groups: IDM, glycogen storage disease
- Lack of symptoms due to use of substrates supports brain metabolism & prevents clinical manifestations
- # Management: verify screen, begin feedings, recheck glucose, IV glucose if still hypoglycemic

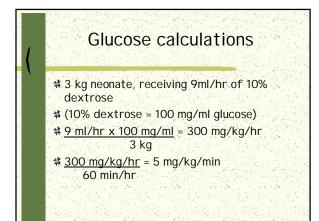
Symptomatic Hypoglycemia -Management

- * Verify laboratory analysis
- # Administer glucose bolus
- # Begin I V glucose (10 % dextrose, 6-8 mg/kg/min)

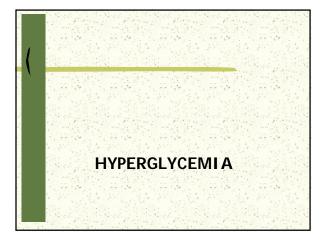
Recheck laboratory glucose (? etiology)
Symptoms clear → transient neonatal

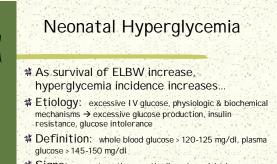
- hypoglycemia
- Symptoms clear but recur → increase IV glucose rate
 - etiology pitultary abnormality, hyperinsulinism, inborn error
- Symptoms persist with normoglycemia
 - ? etiology CNS, CV disease, sepsis, drug effect





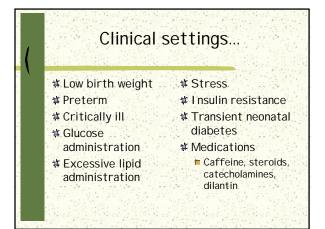






Signs: asymptomatic, osmotic diuresis, weight loss, failure to thrive, fever, glycosuria, ketosis, metabolic acidosis

Pathogenesis * Disordered glucose homeostasis with decreased maturity * I nadequate insulin release in response to hyperglycemia * Post-insulin receptor problems * Delays in insulin sensitivity due to hormone function * Excessive glucose production • Stress, common in ELBW



Complications * Hyperosmolarity * Osmotic diuresis * Polyuria, glucosuria * Dehydration * Cerebral hemorrhage (I VH) * I ncreased CO₂ * Electrolyte imbalance

Treatment



- # Confirm bedside screen
- # Note IV dextrose concentration, meds
- # Assess: urine output, SG, urine dipstick, serum electrolytes, weight, total fluid administration
- # Decrease exogenous glucose administration
- # Feed as appropriate
- # Exogenous insulin administration

Neonatal Insulin Use

- # I nitiation guidelines controversial
- # Most begin use: glucose > 300-400 mg/dl
- # Continuous infusion (0.02-0.05 units/kg/hr)
- SQ administration possible
- # Risk of hypoglycemia, hypokalemia
- # Variable I V delivery due to insulin
- adsorption to plastic tubing

