SHOCK IN THE NEONATES

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DEFINITION

• For cells to survive and function, they need oxygen.
• Inadequate tissue perfusion and oxygen delivery to vital organs → Shock
• Shock presents before hypotension
• Hypotension represents uncompensated shock
Acess Circulation and Cardiovascular status

- **Cardiac Output (CO)**
  \[ CO = HR \times SV \]

- **Oxygen Content in the Arterial Blood (CaO2)**
  \[ CaO2 = 1.36 \times Hgb \times SaO2 + (0.0031 \times PaO2) \]

- **Oxygen Delivery (DO2)**
  \[ DO2 = CO \times CaO2 \]
Systemic vascular resistance (SVR)
is the resistance that the left ventricle must
overcome to pump blood through the systemic
circulation

Blood pressure (BP)

\[ BP = CO \times SVR \]
# Stages of Shock

<table>
<thead>
<tr>
<th>Compensated shock</th>
<th>Decompensated shock</th>
<th>Irreversible shock</th>
</tr>
</thead>
<tbody>
<tr>
<td>- Normal blood pressure</td>
<td>- Anaerobic energy production of tissues</td>
<td>- Multiorgan dysfunction</td>
</tr>
<tr>
<td>- Tachycardia</td>
<td>- Worsening metabolic acidosis</td>
<td>- Poor response to inotropes</td>
</tr>
<tr>
<td>- Redistribution of blood to pivotal organs; brain, heart, adrenal glands</td>
<td>- Renal failure</td>
<td>- Intractable metabolic acidosis</td>
</tr>
<tr>
<td>- Delayed peripheral blood perfusion</td>
<td>- Decreased response to inotropes</td>
<td>- Death</td>
</tr>
<tr>
<td>- Decreased urine output</td>
<td>- Hypotension occurs</td>
<td></td>
</tr>
<tr>
<td>- May deteriorate if not treated</td>
<td>- Rapidly deteriorate to irreversible shock</td>
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</table>
Shock, a state of cellular and tissue hypoxia, is due to reduced oxygen delivery, increased oxygen consumption, and/or inadequate oxygen utilization.

Cellular hypoxia results in a switch to anaerobic metabolism and accumulation of lactic acid. Increasing levels of lactic acid causes metabolic acidosis, which interferes with cell and organ function and, if not addressed, cell death.
Blood Pressure
- 3 Main Causes of Shock

• Hypovolemia  --> **Hypovolemic Shock**

• Heart Failure  --> **Cardiogenic Shock**

• Infection  --> **Septic Shock**
PATHOPHYSIOLOGY OF SHOCK

• Hypovolemic shock
  – Feto-maternal or feto-fetal hemorrhage
  – Placental accidents
  – Birth injury – subgaleal hematoma

• Distributive shock
  – Septic shock

• Cardiogenic shock
  – Severe birth asphyxia with myocardial ischemia
  – Arrhythmia, congenital heart block
Hypovolemic

- Hypovolemic – Due to insufficient circulating blood volume, resulting in a reduction in cardiac output (CO) (reduced preload) or reduced oxygen delivery
# Etiology of Hypovolemic Shock

<table>
<thead>
<tr>
<th>Antenatal/Perinatal Hemorrhage</th>
<th>Postnatal Causes Hemorrhage</th>
<th>Water Loss</th>
</tr>
</thead>
<tbody>
<tr>
<td>- Placental abruptions</td>
<td>- Vitamin K deficiency</td>
<td>- Trans epidermal water loss</td>
</tr>
<tr>
<td>- TTTS</td>
<td>- DIC and coagulopathy</td>
<td>- Dehydration</td>
</tr>
<tr>
<td>- Fetomaternal hemorrhage</td>
<td>- IVH</td>
<td>- Diarrhea</td>
</tr>
<tr>
<td>- Cord accidents</td>
<td>- Surgical bleeding</td>
<td>- Polyuria</td>
</tr>
<tr>
<td>- Birth injuries</td>
<td></td>
<td>- Third space loss; NEC, septicemia</td>
</tr>
<tr>
<td>- Subgaleal hemorrhage</td>
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</tr>
</tbody>
</table>
Distributive – Severely decreased systemic vascular resistance (SVR) due to impairment of vascular tone, which results in maldistribution of blood within the microcirculation and regional and global hypoperfusion.
ETIOLOGY DISTRIBUTIVE SHOCK

- Sepsis
- Adrenal insufficiency
- include hydrops fetalis and toxic shock syndrome
CARDIOGENIC

Cardiac Dysfunction (Pump Failure) or Arrhythmia, Causing a decrease in CO primarily due to deceased stroke volume (SV) (eg, impaired contractility). In cases of complete heart block, the significant reduction in heart rate (HR) leads to inadequate CO.
ETIOLOGY CARDIOGENIC SHOCK

- Cardiogenic shock
- Myocardial ischemia/hypoxemia
- Congenital heart disease (CHD)
- Hypoplastic left heart syndrome
- Critical aortic valve stenosis
- Critical coarctation of the aorta (COA)
- Interrupted aortic arch
Etiology Cardiogenic Shock (Cont. . .)

- Cardiac arrhythmias
- Complete congenital heart block
- Supraventricular tachycardia.
- Ventricular tachycardia
- Myocarditis usually due to viral infection, most commonly coxsackievirus.
- Congenital cardiomyopathy that typically presents with hydrops fetalis.
Obstructive shock

- Occurs when extracardiac diseases lead to impaired CO. (eg, pulmonary embolus, severe pulmonary hypertension) or mechanical (eg, tension pneumothorax, pericardial tamponade, constrictive pericarditis).
- These are rare causes of neonatal shock.
Multifactorial shock

It is important to identify the multiple etiologies in order to guide treatment decision.
Vital Signs

- Abnormal heart rate
- Hypotension
- Abnormal body temperature
- Decreased peripheral perfusion
- Cool extremities, acrocyanosis, and pallor
- Delayed capillary refill >4 seconds
- Neurologic changes vary from lethargy (including poor feeding) to irritability
| CVS       | • Tachycardia, delayed CRT  
|          | • Decreased perfusion index, hypotension |
| CNS       | • Irritability, poor feeding or sucking  
|          | • Stupors or coma |
| KUB       | • Oliguria or anuria  
|          | • Oliguria within first 24 hour may be normal |
| RS        | • Unspecific - tachypnea or apnea  
|          | • Cyanosis |
| GI        | • Increased gastric residual  
|          | • Bowel ileus |
Basic Laboratory Studies

- Arterial blood gas
- Lactate
- Blood cultures
- Additional studies
- Echocardiography
- Electrocardiogram (ECG)
- Chest radiography
- Abdominal radiography
- Complete blood count (CBC)
Laboratory Findings

• Anemia due to blood loss
• Prolonged PT/INR and PTT tests
• Glucose levels may be elevated or decreased during neonatal shock.
• Hyperkalemia due to tissue injury and cell death
• Serum bilirubin levels and liver enzymes may be elevated due to hepatic injury and dysfunction
• Serum creatinine and BUN may be elevated due to renal injury and dysfunction.
Management

1. Airway/breathing
2. Vascular access
3. Fluid resuscitation
THERAPEUTIC INTERVENTIONS

• **Fluid resuscitation** — Fluid resuscitation aims to improve preload, thereby increasing stroke volume (SV) and cardiac output (CO).

• **Type of solution** — Several studies suggest that isotonic crystalloid solutions (eg, normal saline or Ringer's lactate) are preferable to colloid solutions (eg, albumin). Normal saline is the most commonly administered isotonic fluid in neonates and is the solution of choice.
• **Volume and Rate** — Infants with hypovolemic shock, as well as many infants with distributive shock, require aggressive fluid resuscitation. In these patients, we administer 10 mL/kg per bolus of normal saline infused over 10 to 15 minutes.

• **Therapy is Repeated**, as needed, up to four times if there is no clinical evidence of improvement and there are no signs of fluid overload (rales or hepatomegaly).

• Additional therapies, such as transfusion of blood products in neonates with shock due to hemorrhage
THERAPEUTIC INTERVENTIONS (Cont…)

- Fluid Resuscitation may be harmful for neonates with cardiogenic shock or those who have compensated shock with certain comorbidities (eg, prematurity, acute renal failure, intrinsic respiratory diseases).
Patients with obstructive shock may also require volume expanders to improve cardiac preload, but therapy should focus on emergent correction of the underlying cause (eg, needle or chest tube thoracostomy for tension pneumothorax or pericardiocentesis for cardiac tamponade).
Transfusion

- Transfusion — For neonates with shock due to acute blood loss, red blood cell (RBC) transfusion can be lifesaving
Antibiotic Therapy

• The empiric antibiotic regimen should include agents active against group B streptococcal and other organisms that cause neonatal sepsis (eg, Escherichia coli).

• The combination of ampicillin and gentamicin or ampicillin and a third generation cephalosporin (eg, cefotaxime, if available) are potential regimens that provide empiric coverage for these organisms until culture results are available.
Vasoactive agents

- **Vasopressor and inotropic** agents are used to support neonates with distributive shock who have not improved with initial fluid resuscitation and those with cardiogenic shock.
- Commonly used vasoactive agents include dopamine, epinephrine, dobutamine, and milrinone.
• **Dopamine** is the most commonly used agent in neonates, based on longer clinical experience and familiarity with its use.

• **Epinephrine** is often beneficial for the management of shock in older patients, but its pharmacologic properties are less well understood in neonates.

• **Dobutamine** is typically the first inotrope administered for cardiogenic neonatal shock.
Dopamine

- For infants with distributive shock who do not respond adequately to fluid resuscitation, dopamine is infused beginning at a rate of $5 \text{ mcg/kg/min}$ with titration up to a maximum of $15 \text{ mcg/kg/min}$ based on the infant's clinical response.

- **Dopamine** can also be used for the management of cardiogenic shock.
Dopamine

• The hemodynamic effects of dopamine are dose-dependent:
  • Low dosage: 1 to 5 mcg/kg/minute, increased renal blood flow and urine output
  • Intermediate dosage: 5 to 15 mcg/kg/minute, increased renal blood flow, heart rate, cardiac contractility, cardiac output, and blood pressure
  • High dosage: >15 mcg/kg/minute, alpha-adrenergic effects begin to predominate, vasoconstriction, increased blood pressure
Dobutamine

- Dobutamine increases CO via improved myocardial contractility and increased HR. Since it can also decrease SVR and cause hypotension, it is useful for patients with decreased myocardial function due to cardiogenic shock who are normotensiv
Dobutamine

- Dobutamine infusion begins at a rate of 5 mcg/kg per min with titration up to a maximum of 20 mcg/kg per min based on the infant's clinical response.
Epinephrine

- Epinephrine increases myocardial contractility and is a potent vasoconstrictor. It is sometimes used for distributive shock that is refractory to dopamine or as a first-line vasoactive therapy for cardiogenic shock.
Epinephrine

- Epinephrine is typically started at a rate of 0.1 mcg/kg per min and titrated up to a maximum of 1 mcg/kg per min based on the infant's clinical response.
Thank Youk