

NEONATAL INFECTION SEPSIS

DEFINITION

NEONATAL SEPSIS IS DEFINED AS
A CLINICAL SYNDROME OF
BACTEREMIA

WITH
SYSTEMIC SIGNS AND SYMPTOMS
OF INFECTION

IN THE FIRST 4 WEEKS OF LIFE

CLASSIFICATION

ACCORDING TO THE ONSET OF SYMPTOMS

THE NEONATAL SEPSIS CAN BE CLASSIFIED

INTO

TWO SUB-TYPES

EARLY ONSET SEPSIS < 72 HOURS

LATE ONSET SEPSIS > 72 HOURS

ACCORDING TO THE ONSET OF SYMPTOMS

EARLY ONSET SEPSIS < 72 HOURS

85% OF EARLY-ONSET SEPSIS
APPEAR WITHIN 24 HOURS

5% PRESENT AT 24-48 HOURS

AND A SMALLER PERCENTAGE
PRESENT
WITHIN 48-72 HOURS

THE ONSET IS MOST RAPID IN
PREMATURE NEONATES

LATE ONSET SEPSIS > 72 HOURS

OCCURS

AT 4-90

DAYS OF LIFE

TRANSMISSION

EARLY-ONSET INFECTIONS

ARE ACQUIRED THROUGH THE
COLONIZED BIRTH CANAL

MATERNAL GENITAL TRACT

OR

IN THE DELIVERY AREA

TRANSPLACENTAL INFECTION

OR

ASCENDING INFECTION

EARLY-ONSET INFECTIONS

ARE ACQUIRED

FROM

THE CAREGIVING

ENVIRONMENT

ETIOLOGY

EARLY-ONSET SEPSIS

- **GROUP B STREPTOCOCCUS**
 - **GRAM-NEGATIVE**
 - **ORGANISMS**
 - **STAPH. AUREUS**
- **LISTERIA MONOCYTOGENES**

LATE-ONSET SEPSIS

TERM:

- **GROUP B STREPTOCOCCUS**
- **GRAM-NEGATIVE ORGANISMS**

PRETERM:

- **COAGULASE NEG. STAPH.**
- **GRAM-NEGATIVE ORGANISMS**
- **GROUP B STREPTOCOCCUS**
 - **STAPH. AUREUS**
 - **ENTEROCOCCUS**
 - **FUNGAL**

SINGS AND SYMPTOMS

RISK FACTORS

EARLY-ONSET SEPSIS

LOW BIRTH WEIGHT

*PROLONGED RUPTURE OF
MEMBRANES*

FOUL SMELLING LIQUOR

*MULTIPLE PER VAGINUM
EXAMINATIONS*

MATERNAL FEVER

*DIFFICULT OR PROLONGED
LABOUR*

ASPIRATION OF MECONIUM

LATE-ONSET SEPSIS

LOW BIRTH WEIGHT

LACK OF BREASTFEEDING

*SUPERFICIAL INFECTIONS
(PYODERMA, UMBILICAL SEPSIS)*

ASPIRATION OF FEEDS

*DISRUPTION OF SKIN INTEGRITY
WITH NEEDLE PRICKS AND USE
OF INTRAVENOUS FLUIDS*

CLINICAL MANIFESTATIONS OF NEONATAL SEPSIS

LETHARGY

REFUSAL TO SUCKLE

POOR CRY

NOT AROUSABLE, COMATOSED

ABDOMINAL DISTENSION

DIARRHEA

VOMITING

HYPOTHERMIA

POOR PERFUSION

SCLEREMA

POOR WEIGHT GAIN

SHOCK

BLEEDING

RENAL FAILURE

CYANOSIS*

TACHYPNEA*

CHEST RETRACTIONS*

GRUNT*

APNEA/GASPING*

FEVER⁺

SEIZURES⁺

BLANK LOOK⁺

HIGH PITCHED CRY⁺

EXCESSIVE CRYING/IRRITABILITY⁺

NECK RETRACTION⁺

BULGING FONTANEL⁺

*** PARTICULARLY SUGGESTIVE OF
PNEUMONIA**

**⁺PARTICULARLY SUGGESTIVE OF
MENINGITIS**

CLINICAL MANIFESTATIONS

EARLY ONSET SEPSIS
MANIFESTS FREQUENTLY AS
RESPIRATORY DISTRESS
(PNEUMONIA)

AND LESS COMMONLY
AS
SEPTICEMIA OR
MENINGITIS

LATE-ONSET SEPTICEMIA
IS CAUSED BY THE ORGANISMS
THRIVING IN THE EXTERNAL
ENVIRONMENT OF THE HOME OR THE
HOSPITAL

THE INFECTION IS OFTEN TRANSMITTED
THROUGH

THE HANDS OF THE CARE-PROVIDERS

THE PRESENTATION IS
THAT OF
SEPTICEMIA
PNEUMONIA
MENINGITIS

DANGER SIGNS AND SYMPTOMS OF INFECTIONS IN THE NEWBORN

DANGER SIGNS

- **UNABLE TO FEED**
- **CONVULSIONS**
- **DROWSINESS OR ALTERED CONSCIOUSNESS (EXCLUDE HYPOGLYCAEMIA !)**
- **RR < 20/MIN OR APNOEA (>15 SEC)**
 - **RR > 60/MIN**
 - **GRUNTING (PROLONGED > 4 HRS)**
- **SEVERE CHEST IN-DRAWING**
- **CENTRAL CYANOSIS**

OTHER SIGNS OF INFECTION

- **DEEP JAUNDICE**
- **SEVERE ABDOMINAL DISTENSION**
- **PERI-UMBILICAL REDNESS OR UMBILICUS DRAINING PUS**
- **BULGING FONTANELLE**
- **HYPOTHERMIA OR FEVER**
 - **IRRITABLE**
- **PAINFUL JOINTS, SWELLING OF THE JOINTS, REDUCED MOVEMENT AND IRRITABILITY IF HANDLING THESE JOINTS**
- **MANY OR SEVERE SKIN PUSTULES**

A large WHO study published in 2003 identified nine clinical features which predict severe bacterial illness in young infants

- 1. FEEDING ABILITY REDUCED**
- 2. NO SPONTANEOUS MOVEMENT**
- 3. TEMPERATURE >38 C**
- 4. PROLONGED CAPILLARY REFILL TIME**
- 5. LOWER CHEST WALL IN DRAWING**
- 6. RESP RATE > 60/MINUTE**
- 7. GRUNTING**
- 8. CYANOSIS**
- 9. CONVULSIONS**

DIAGNOSIS

DIRECT METHOD

➤ ISOLATION OF MICROORGANISMS
FROM BLOOD

➤ CSF

➤ URINE

➤ PLEURAL FLUID OR PUS

IS DIAGNOSTIC

INDIRECT METHOD I

THERE ARE A VARIETY OF TESTS WHICH ARE HELPFUL FOR SCREENING OF NEONATES WITH SEPSIS THE MOST USEFUL AND WIDELY USED IS

➤ ***THE WHITE BLOOD CELL COUNT AND DIFFERENTIAL COUNT***

LEUKOPENIA (TLC $< 5000/\text{cmm}$)

NEUTROPENIA (ABSOLUTE NEUTROPHIL COUNT ANC $< 1800/\text{cmm}$)

**IMMATURE NEUTROPHIL TO TOTAL NEUTROPHIL
I/T RATIO > 0.2**

➤ ***PLATELET COUNT OF LESS THAN 100,000 PER cmm***

INDIRECT METHOD II

➤ *C-REACTIVE PROTEIN (CRP) +*

➤ *ERYTHROCYTE SEDIMENTATION RATE*

ESR > 15mm 1ST HOUR

➤ *LUMBAR PUNCTURE*

**SHOULD BE DONE IN ALL CASES OF LATE ONSET
SEPSIS (>72 HOURS)**

BECAUSE

**10-15 % OF THEM MAY HAVE ASSOCIATED
MENINGITIS**

MANAGEMENT

IF DANGER SIGNS ALWAYS TREAT

EMERGENCY MANAGEMENT

- **O2 (NASAL PRONGS) IF RESPIRATORY DISTRESS OR CYANOSIS**
- **MASK VENTILATION IF RR < 20/MIN**
- **CHECK GLUCOSE IF < 45 MG/DL OR < 2.5 MMOL/L 2 ML/KG G 10% IV; IF YOU CANNOT CHECK GLUCOSE AND CHILD IS DROWSY OR CONVULSING TREAT FOR HYPOGLYCAEMIA (IF NO IV ACCESS GIVE G 10% OR BREAST-MILK VIA NG TUBE)**
- **IF CONVULSING: GIVE PHENOBARBITAL 20 MG/KG IV OR IM STAT**
- **IF SUSPECTED MENINGITIS PERFORM A LUMBAR PUNCTURE IF CLINICAL CONDITION ALLOWS AND BEGIN TREATMENT**
- **ANTIBIOTIC TREATMENT OF NEONATAL SEPSIS (AMPICILLIN + GENTAMYCIN OR AMPICILLIN + CEFOTAXIME (CEFTRIAZONE IF CEFOTAXIME NOT AVAILABLE))**
- **CHECK CHILD RECEIVED VITAMIN K**
- **ENSURE PROPER NUTRITION & FLUID MANAGEMENT. MONITOR THE CHILD CLOSELY.**

**NO DANGER SIGNS BUT *RISK FACTORS* OR OTHER
*SIGNS OF INFECTION***

**➤ CONSIDER TREATMENT
(ACCORDING TO CLINICAL JUDGEMENT)**

**IF UNSURE OR SEVERAL RISK OR OTHER FACTORS
EXIST**

➤ “PROPHYLACTIC TREATMENT” RECOMMENDED

DRUGS

➤ **ANTIBIOTIC TREATMENT IV
WITH AMPICILLIN + GENTAMYCIN**

OR

CEFOTAXIME + AMPICILLIN

(USE CEFTRIAZONE IF CEFOTAXIME NOT AVAILABLE)

REPLACE

**AMPICILLIN WITH CLOXACILLIN IF EXTENSIVE SKIN
PUSTULES OR ABSCESSES**

➤ **GIVE VITAMIN K IM IF NOT GIVEN BEFORE OR
UNCLEAR**

➤ **G 10% FOR TREATMENT OF HYPOGLYCEMIA**

DOSAGES OF ANTIBIOTICS

➤ AMPICILLIN

Age < 1 week 50 mg/kg x 2 IV

Age > 1 week 50 mg/kg x 3 IV

(meningitis: increase dose: < 1 week 150 mg/kg/day, > 1 week 200 mg/kg/day, see meningitis protocol)

➤ GENTAMYCIN

5 mg/kg OD IM or slowly IV (< 2.500 g 3 mg/kg OD IM or slowly IV), **Duration 3-5 days**

➤ CEFOTAXIME

Age < 1 week 50 mg/kg x 3 IV (premature infants x 2)

Age > 1 week 50 mg/kg x 4 IV

➤ CEFTRIAXONE

Age < 1 week 50 mg/kg OD IV

Age > 1 week 100 mg/kg x 1 IV or IM or 50 mg x 2 IV

(cefotaxime preferred for neonates)

➤ CLOXACILLIN

Age < 1 week: 50 mg/kg x 2 IV

Age > 1 week: 50 mg/kg x 3 IV

DURATION OF ANTIBIOTIC TREATMENT

IF TREATMENT WAS A PRECAUTION

BECAUSE OF RISK FACTORS (SEE ABOVE) AND NEWBORN NEVER SHOWED ANY SIGNS OF SICKNESS, IS ACTIVE AND DRINKS WELL

DISCONTINUE AFTER MINIMUM OF THREE FULL DAYS OF PARENTERAL TREATMENT

IF INFANT WAS SICK

**TREAT AT LEAST 7 DAYS, AIM AT 10 DAYS OF ANTIBIOTIC TREATMENT
AT LEAST 5 FULL DAYS PARENTERAL TREATMENT FOLLOWED BY ORAL TREATMENT IF BABY IS TAKING ORAL FLUIDS, AND HAS BEEN FREE OF FEVER FOR AT LEAST 24 HOURS AND THERE ARE NO NEUROLOGICAL SIGNS.**

IF THE CONDITION IS NOT IMPROVING WITHIN 48-72 HOURS

THE TREATMENT MAY NEED TO BE CHANGED. (CONSIDER OTHER OR ADDITIONAL CAUSES OF INFECTIONS INCLUDING MALARIA, FUNGAL INFECTION ETC

➤ **SUPPORTIVE CARE**

**THE PURPOSE OF SUPPORTIVE CARE IS
TO NORMALIZE THE TEMPERATURE
TO STABILIZE THE CARDIOPULMONARY
STATUS**

**TO CORRECT HYPOGLYCEMIA
TO PREVENT BLEEDING TENDENCY
THERMOREGULATION**

THANKS