

Neonatal *Herpes Simplex* Virus Infection

HIGH-YIELD • NGN

SYSTEM — MATERNAL/NEWBORN

PATHOGEN — HSV-1 / HSV-2

ONSET — FIRST 4-6 WKS OF LIFE

TX — IV ACYCLOVIR

A vertically-transmitted viral infection that disguises itself as routine neonatal lethargy — until it doesn't.

01 Definition & Overview

Neonatal HSV is a potentially devastating viral infection of the newborn, most often acquired during passage through an infected maternal birth canal.

Caused by **HSV-2** (most common) or **HSV-1**, with three clinical presentations: skin/eye/mouth (SEM), CNS disease, and disseminated multi-organ infection.

Mortality and long-term neurologic injury remain high despite treatment — making **early empiric IV acyclovir** the single most important nursing/medical action.

~85%

INTRAPARTUM (BIRTH CANAL)

~10%

POSTNATAL (COLD SORE CONTACT)

~5%

INTRAUTERINE (CONGENITAL)

25–50%

TRANSMISSION IF PRIMARY MATERNAL HSV AT DELIVERY

02 Pathophysiology *simplified*

STEP 01

Maternal HSV

Primary or recurrent genital infection → viral shedding (often asymptomatic)

STEP 02

Exposure at Birth

Neonate contacts infected secretions during vaginal delivery

STEP 03

Viral Entry

Mucous membranes, skin breaks, conjunctiva, scalp lesions from monitors

STEP 04

Local Replication

Skin, eye, oral mucosa — vesicles appear 1–2 wks

STEP 05

Spread & Disease

Viremia → CNS or multi-organ dissemination if untreated

→ THREE CLINICAL PRESENTATIONS

~45%

SEM Disease

Skin · Eye · Mouth. Vesicles, conjunctivitis, oral ulcers. Best prognosis if treated — but untreated, 70% progress to CNS or disseminated.

~30%

CNS Disease

Encephalitis. Seizures, lethargy, bulging fontanelle, temp instability. High risk of permanent neurologic injury.

~25%

Disseminated

Multi-organ. Liver, lungs, adrenals, CNS, DIC, shock. Highest mortality; presents earliest (5–7 days).

Typical Onset Timeline



03 Signs & Symptoms

Early & Subtle

DAYS 1-14

- Lethargy, poor feeding, irritability
- Temperature instability — fever OR hypothermia
- **Vesicular skin lesions** — clustered on erythematous base (often on scalp/presenting part)
- Conjunctivitis, watery eye discharge, keratitis
- Oral ulcers, mouth vesicles
- Subtle apnea or tachypnea

Late • Red Flags

🚨 ACT NOW

- ➤ Seizures • focal or generalized
- ➤ Bulging fontanelle, ↑ ICP, coma
- ➤ Apnea, respiratory failure
- ➤ Jaundice, hepatomegaly, ↑ LFTs
- ➤ Thrombocytopenia, bleeding, DIC
- ➤ Septic-shock picture (often culture-negative)

“ A vesicular rash on any newborn is HSV until proven otherwise — and 40% of disseminated/CNS cases never develop skin lesions at all.

04 Priority Nursing Interventions

PRIORITY 01 • AIRWAY/BREATHING

Monitor Respiratory Status

Assess for apnea, tachypnea, retractions. Pulse ox continuous. Prepare for intubation if disseminated disease/shock develops.

PRIORITY 02 • CIRCULATION

Hemodynamic Support

Vital signs q1h initially. IV access × 2. Fluid resuscitation per protocol. Monitor for shock, DIC, bleeding.

PRIORITY 03 • TREAT EMPIRICALLY

Start IV Acyclovir NOW

Do **not** wait for culture/PCR results. Empiric IV acyclovir 20 mg/kg/dose q8h is the single most important intervention.

PRIORITY 04 • OBTAIN CULTURES

HSV PCR + Surface Cultures

Swab: nasopharynx, mouth, conjunctivae, rectum, vesicle fluid. Send blood & CSF HSV PCR; LFTs, CBC, coags.

PRIORITY 05 • NEURO & SEIZURE

Continuous Neuro Monitoring

Assess fontanelle, LOC, tone, pupillary response. Seizure precautions. EEG if available. Padded side rails.

PRIORITY 06 • INFECTION CONTROL

Contact Precautions

Gown + gloves. Cover skin lesions. Single-room isolation. Hand hygiene before/after every contact.

PRIORITY 07 • OCULAR PROTECTION

Ophthalmology Consult

Topical ophthalmic antiviral (trifluridine or ganciclovir gel) for eye involvement. Prevents corneal scarring & blindness.

PRIORITY 08 • NUTRITION & FAMILY

Feeding & Bonding

Strict I&O, daily weights. Support breastfeeding (safe unless breast lesions). Emotional support; involve parents in care.

05 Pharmacology — *the only drug that matters*

Acyclovir (IV)

ANTIVIRAL • NUCLEOSIDE ANALOG

MECHANISM

Phosphorylated by viral thymidine kinase → inhibits viral DNA polymerase → halts HSV replication. Selective for infected cells.

SIDE EFFECTS

Nephrotoxicity (crystalluria), neutropenia, thrombocytopenia, phlebitis & tissue necrosis at IV site, transient ↑ LFTs.

NURSING

Ensure adequate hydration. Monitor BUN/Cr & ANC. Infuse **over ≥1 hour**. Verify patent line (vesicant). Adjust dose for renal function.

KEY TEACHING

Course continues with **oral acyclovir suppression × 6 months** after IV completion to reduce skin recurrences and improve neurodev outcomes.

SEM: 20 mg/kg/dose IV q8h × 14 days **CNS / Disseminated:** 20 mg/kg/dose IV q8h × 21 days

Then: PO acyclovir 300 mg/m² TID × 6 months

↑ MATERNAL SUPPRESSION

Oral acyclovir or valacyclovir from 36 wks gestation for any HSV-positive mother. Reduces shedding, lesions, and C-section rates. *Not* used in neonate.

OCULAR ANTIVIRALS

Topical **trifluridine 1%** or **ganciclovir gel 0.15%** for HSV keratoconjunctivitis — given alongside IV acyclovir, never alone.

06 NCLEX Test Traps ⚠

~~"Mom has no symptoms, so the baby is safe."~~

TRAP 01

Truth: Most mothers transmitting HSV are **asymptomatic at delivery** with no known history. Absence of symptoms does *not* rule out HSV.

~~"Wait for the PCR to confirm before starting acyclovir."~~

TRAP 02

Truth: Empiric IV acyclovir is started **immediately on suspicion**. Delay = death or permanent neurologic injury.

~~"C-section eliminates the risk."~~

TRAP 03

Truth: Cesarean **reduces** but does *not* eliminate risk — especially if membranes have been ruptured >4–6 hours.

~~"Use a fetal scalp electrode to monitor — she has HSV but no active lesions."~~

TRAP 04

Truth: **Avoid scalp electrodes, vacuum, and forceps** with any HSV history. Any skin break is a portal of entry.

~~"Mom can't breastfeed because she has HSV."~~

TRAP 05

Truth: Breastfeeding is **allowed** unless there are HSV lesions on the breast. Strict handwashing required; cover oral cold sores.

~~"No vesicles — no HSV."~~

TRAP 06

Truth: Up to **40% of CNS & disseminated cases never show skin lesions**. Sepsis-like neonate <6 wks = include HSV in workup.

For the Pregnant Patient with HSV

- Disclose HSV history to **every** prenatal provider — even if dormant for years
- Take suppressive antiviral therapy from **36 weeks gestation** as prescribed
- At onset of labor: report any **prodromal symptoms** (tingling, burning, lesions) — may need C-section
- Avoid sexual contact in 3rd trimester if partner has HSV and you do not

For Parents of an HSV-Exposed Newborn

- Watch for: skin blisters, fever, hypothermia, poor feeding, lethargy, seizures, breathing changes
- Hand hygiene **every time** before holding baby
- Cover any cold sores with mask; **do not kiss** the baby on the face/mouth
- Family/visitors with active cold sores: no contact with baby
- Complete the full **6 months of oral acyclovir** — even when baby looks well
- Attend all neurodevelopmental, ophthalmology, and audiology follow-ups

Memory Boosters

THREE CLINICAL FORMS

S • C • D

- S** SEM — Skin · Eye · Mouth (~45%)
- C** CNS — Encephalitis · seizures (~30%)
- D** Disseminated — multi-organ + shock (~25%)

WHEN TO SUSPECT HSV

VESICLE

- V** Vesicles (clustered, on red base)
- E** Eye drainage / conjunctivitis
- S** Seizures & sepsis-like picture
- I** Irritability + Instability (temp)
- C** CSF pleocytosis (CNS form)
- L** Liver enzymes up · DIC
- E** Empiric Acyclovir — *don't wait*

ACYCLOVIR NURSING PRIORITIES

A • C • E

- A** Adequate hydration (prevent crystalluria)
- C** Check renal & CBC (BUN/Cr, ANC)
- E** Extended infusion ≥ 1 hr (vesicant; avoid extravasation)

MATERNAL DELIVERY RULE

"NO SCALP"

- Active lesions or prodrome → **C-section**
- × **No** scalp electrodes
- × **No** vacuum extraction
- × **No** forceps
- × **No** artificial ROM if avoidable
- ! Any skin break = portal of entry