

Acyclovir Sodium

Brand names	Generic
Medication error potential	ISMP reports that Zovirax has been confused with Doribax, Zyvox, and Zostrix. ⁽¹⁾
Contraindications and warnings	Contraindications: Hypersensitivity to acyclovir or valacyclovir ⁽²⁾ Warnings: Renal failure and thrombotic thrombocytopenic purpura (TTP)/hemolytic uremic syndrome (HUS) have both been reported with acyclovir. Deaths have occurred due to renal failure and TTP/HUS. Caution should be used in patients with underlying neurological abnormalities, as well as those with serious renal, hepatic, or electrolyte abnormalities or significant hypoxia. ⁽²⁾
Infusion-related cautions	To decrease the risk of nephrotoxicity, the patient should be adequately hydrated before and during the infusion. ^(2,3) Extravasation may cause inflammation and phlebitis at the injection site particularly at higher concentrations. ^(2,4) (See Appendix E for additional information regarding extravasation treatment.)
Dosage	Doses generally range from 15–60 mg/kg/day or 1500 mg/m ² /day divided q 8 hr depending on the infection. ^(2,5-12) Obese patients should be dosed using ideal body weight. ⁽²⁾ Herpes simplex virus (HSV) Neonatal HSV: Although product information continues to recommend 30 mg/kg/day divided q 8 hr for 10 days, 60 mg/kg/day divided q 8 hr for 14–21 days is currently recommended by most experts with duration based on severity of illness. ^(2,11,12,26) HSV encephalitis ≥3 months–12 years: 60 mg/kg/day divided q 8 hr for 10–21 days ^(2,24) ; some experts recommend 30–45 mg/kg/day divided q 8 hr for 14–21 days due to concern for nephrotoxicity with larger doses. ⁽¹¹⁾ ≥12 years: 30 mg/kg/day divided q 8 hr for 10–21 days ^(2,11) Genital HSV (first episode), ≥12 years: 15 mg/kg/day divided q 8 hr for 5–7 days ^(2,11) HSV in immunocompromised host (localized, progressive, or disseminated): 30 mg/kg/day divided q 8 hr for 7–14 days ⁽¹¹⁾ Prophylaxis in HSV-seropositive immunocompromised hosts: 15 mg/kg/day divided q 8 hr during risk period ⁽¹¹⁾ Varicella Immunocompetent hosts >2 years: 30 mg/kg/day divided q 8 hr or 1500 mg/m ² /day divided q 8 hr for 7–10 days ⁽¹¹⁾ Immunocompromised hosts <12 years: 60 mg/kg/day divided q 8 hr for 7 days ⁽²⁾ >12 years: 30 mg/kg/day divided q 8 hr for 7 days ⁽²⁾ or <1 year: 30 mg/kg/day divided q 8 hr for 7–10 days ⁽¹¹⁾ ≥1 year: 1500 mg/m ² /day (or 30 mg/kg/day) divided q 8 hr for 7–10 days ^(8,10,11) Initiate therapy as soon as possible after rash appears. Early initiation therapy is associated with more rapid improvement and less dissemination than late initiation therapy (>3–5 days). ⁽⁹⁾



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Dosage (cont.)

IV therapy is recommended for the duration of treatment in immunocompromised hosts.⁽¹¹⁾

Sixty-six pediatric renal transplant recipients with VZV were given 1500 mg/m²/day divided q 8 hr. Azathioprine was temporarily discontinued until the lesions crusted over and no new lesions appeared; at that time, azathioprine was restarted at the usual dose. Three recipients experienced acute rejection that responded to prednisone. One patient died.⁽¹³⁾

Zoster in immunocompetent or compromised host, all ages: 30 mg/kg/day divided q 8 hr for 7–10 days⁽¹¹⁾

Prevention or suppression of cytomegalovirus infection in allogeneic bone marrow transplant recipients (this use has largely been replaced by ganciclovir): 1500 mg/m²/day divided q 8 hr beginning 5 days before transplantation and continuing 30 days after^(7,14)

Aplastic anemia (use not established): 15 mg/kg/day (schedule not provided) for 10 days has been given to two patients (one, a 13-year-old girl) who had severe, possibly viral related, aplastic anemia refractory to standard treatment.⁽⁵⁾

Dosage adjustment in organ dysfunction

Adjust dosage in patients with renal dysfunction.^(2,11,15,16) If CrCl is 25–50 mL/min, give normal dose q 12 hr; if CrCl is 10–25 mL/min, give normal dose q 24 hr, and if CrCl is <10 mL/min, give 50% of the dose q 24 hr.⁽²⁾

Another source recommends the following dose adjustment based on GFR⁽¹⁶⁾:

GFR (mL/min/1.73 m ²)	Dose
30–50	10 mg/kg q 12 hr
10–29	10 mg/kg q 24 hr
<10	5 mg/kg q 24 hr

Neonates with hepatic or renal dysfunction and young premature infants may also require dose adjustment.⁽¹⁵⁾

Maximum dosage

Do not exceed 20 mg/kg q 8 hr.⁽²⁾

An 11-day-old neonate has received 258 mg/kg of IV acyclovir in a 24-hour period that was treated with NS hydration; the patient experienced only a transient increase in SCr.⁽¹⁷⁾

Additives

Aqueous solution contains 5.1 mg/mL sodium.⁽³⁾ Lyophilized powder contains 49 mg sodium/500 mg acyclovir.⁽²⁴⁾

Suitable diluents

D5W, NS, D5NS, LR⁽¹⁸⁾

Maximum concentration

≤7 mg/mL.^(2,18) Infusion of a solution ≥10 mg/mL increases the risk of phlebitis and extravasation.^(2,19)

Preparation and delivery

Stability: Vials should be reconstituted with SW for injection and should be used within 12 hours.⁽²⁴⁾ Once diluted for administration, doses should be used within 24 hours.^(2,18) Refrigeration may cause precipitation; however, the precipitate redissolves at room temperature and potency does not appear to be affected.^(18,24) BW for injection containing parabens or benzyl alcohol should not be used to dilute acyclovir sodium powder because precipitation could occur.^(18,24)

Compatibility: For PN compatibility information, see Appendix D.

