Zanamavir Intravenous Infusion for Adults



Who can administer

May be administered by registered competent doctor or nurse/midwife

Important information

- Zanamivir is indicated for the treatment of complicated and potentially life-threatening influenza A or B virus infection when:
 - \circ The patient's influenza virus is known or suspected to be resistant to anti-influenza medicinal products other than zanamivir $^{(ref \, 1)}$ OR
 - Other anti-viral medicinal products for treatment of influenza, are not suitable for the individual patient. (ref 1,3) AND the patient has multi-organ involvement, or requires intensive care (ref 2)
 - Use should be supervised by a consultant clinical microbiologist/virologist or infectious diseases consultant and a consultant in intensive care medicine (ref 3)Â
- Treatment should commence as soon as possible and usually within 6 days of the onset of symptoms (ref 2)
- See under 'Dose' for adjustments required in **renal impairment**

Available preparations

Dectova 200mg per 20ml (10mg/mL) vial

Reconstitution

Already in solution

Infusion fluids

Sodium Chloride 0.9% ONLY

Methods of intravenous administration

Intermittent intravenous infusion (using an electronically controlled infusion device)

- Add the required dose to infusion fluid to make a final volume of 100ml or 250ml
 - Withdraw and discard a volume (equal to the volume of drug solution) from the infusion bag
- The infusion bag should be gently manipulated by hand to ensure it is mixed thoroughly
- Administer over 30 minutes (ref 1,3)
- See also under Further information- re administration undiluted

Dose in adults

Usual dose

Give 600mg every 12 hours for 5 to 10 days (ref 2)

Renal impairment

See Table 1 below

Table 1: Doses in renal impairment			
CrCl (mL/min)	Initial dose	Maintenance dose	Maintenance dose schedule
50 to < 80	600mg	400mg every 12 hours	Begin Maintenance dose 12 hours AFTER initial dose
30 to < 50	600mg	250mg every 12 hours	
15 to < 30	600mg	150mg every 12 hours	Begin Maintenance dose 24 hours AFTER initial dose
< 15	600mg	60mg (SIXTY) every 12 hours	Begin Maintenance dose 48 hours AFTER initial dose

Hepatic impairment

• No dose modification is required

Elderly

· No dose modification is required

Monitoring

- Monitor temperature, blood pressure, heart rate, respiratory rate, O² saturation, behavioural changes, and injection site reaction (ref 3)
- Can increase ALT and AST and cause hepatocellular injury (1%)
- Common side effects are rash, ALT/AST increase and diarrhoea (ref 1,3)

Further information

- Can give undiluted over 30 minutes via electronically controlled device
- No dose adjustment is required for elderly patients

Storage

Store below 25°C

References

SPC - Dectova Â 01/10/2019

- 1. Zanamivir BNF, accessed online via medicines complete Â 17/01/2024
- 2. Guidance on the use of antiviral agents for the treatment and prophylaxis of Influenza, HSE, December 2023
- 3. Injectable medicines guide NHS Medusa Injectable Medicines Guide, downloaded 17/01/2024

Therapeutic classification

Direct Acting Antivirals - Neuraminidase inhibitors - J05AH