Famotidine Intravenous for Adults



Who can administer

May be administered by registered competent doctor or nurse/midwife

Important information

• Unlicensed preparation

Available preparations

Famotidine 20mg per 2ml vial

Reconstitution

Already in solution

Dilute further prior to administration

Infusion fluids

Sodium chloride 0.9% or Glucose 5%

Methods of intravenous administration

Intermittent intravenous infusion (preferred route)

Dilute 20mg with 100ml infusion fluid, and administer over 15 to 30 minutess

Slow intravenous injection

• Dilute each 20mg to a volume of 5 to 10ml with infusion fluid and administer over at least two minutes

Dose in adults

Usual dose

• Give 20mg every 12 hours until oral therapy can be introduced

Premedication of infusion reactions (ref 1)

- Give 20mg thirty to sixty minutes prior to infusion
- Usually given in conjunction with an H1 antihistamine and glucocorticoid

Prophylaxis of acid aspiration (Mendelson's) syndrome (ref 1)

Give 20mg forty to ninety minutes before induction of general anaesthesia

Prophylaxis of upper gastrointestinal haemorrhage from stress ulceration in seriously ill patients $^{(\text{ref 1})}$

• Give 20mg twice daily

Renal impairment

- **Complete atrioventricular block** has been reported in association with administration of famotidine injection ^(ref 2)
- Prolonged QT interval has been reported in patients with moderate to severe renal impairment (ref 1)
- The manufacturers advise that CNS adverse effects have been reported in patients with moderate and severe renal insufficiency
- For patients with CrCl <50 mL/minute: advise increase the dosing interval to 36-48 hours (ref 1)

Storage

Storage in a **refrigerator** at 2-8°C

References

SPC for Famotidine IV (Mylan) revised April 2022

- 1: UpToDate, accessed 07/05/2025
- 2: Complete Atrioventricular Block and Cardiac Arrest following Intravenous Famotidine Administration Anesthesiology 2 1999, Vol.90, 623-626.Â

Therapeutic classification

H2-receptor antagonists