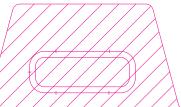


GTIN: 0036519120124

221012

200 mg per vial

Dalbavancin for Injection



Sample label. Please see package insert for complete prescribing information.

NDC 65219-120-12

Dalbavancin for Injection

500 mg per vial

For Intravenous Infusion Only Sterile Single-Dose Vial Discard Unused Portion

One Vial Rx only



Dalbavancin for Injection

500 mg per vial

For Intravenous Infusion Only Sterile Single-Dose Vial Discard Unused Portion

Store at 25°C (77°F); excursions permitted to 15°C to 30°C (59°F to 86°F) [see USP Controlled Room Temperature].

Dalbawancin for injection must be reconstituted under aseptic conditions, using 25 mL of either Sterile Water for Injection, USP, or 5% Dextrose Inlection, USP, for each 500 mm vial.

- or 5% Dextrose Injection, USP, for each 500 mg vial.

 The reconstituted solution contains dalbavancin at a concentration of 20 mg/mL.
- For infusion, dilute the reconstituted solution with 5% Dextrose Injection, USP, to a final dalbavancin concentration of 1 mg/mL to 5 mg/mL. The total time from reconstitution to infusion should not exceed 48 hours.
- DO NOT co-infuse with other medications or electrolytes.
- After reconstitution and dilution, Dalbavancin for Injection is to be administered via intravenous infusion, using a total infusion time of 30 minutes.

Dosage and Administration: See full prescribing information.

One Vial Rx only



NDC 65219-120-12

Dalbavancin for Injection

500 mg per vial

For Intravenous Infusion Only Sterile Single-Dose Vial Discard Unused Portion





www.fresenius-kabi.com/us

Dalbavancin for Injection

500 mg per vial

For Intravenous Infusion Only Sterile Single-Dose Vial Discard Unused Portion

Package contains one sterile vial of dalbavancin HCl powder equivalent to 500 mg of dalbavancin, plus lactose monohydrate (129 mg) and mannitol (129 mg) as excipients. Sodium hydroxide or hydrochloric acid may be added to adjust the pH at the time of manufacture.

One Vial Rx only



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