NDC 63323-349-25

Imipenem and Cilastatin

for Injection, USP (I.V.)

250 mg / 250 mg* per vial

Not to be divided

***Each vial contains:** Imipenem USP 250 mg (Anhydrous Equivalent) and Cilastatin Sodium USP equivalent to 250 mg Cilastatin.
Inactive ingredient: sodium bicarbonate 10 mg added to each vial as a buffer. The sodium content is approximately 18.8 mg (0.8 mEq).
After suspension, vial contents must be transferred to 100 mL of infusion solution prior to intravenous infusion.

CAUTION: SINGLE-DOSE VIAL / FOR I.V. USE ONLY / NOT FOR DIRECT INFUSION

Rx only

1 PACKAGE (25 Single-Dose Vials)

FRESENIUS KABI

964

AGA137102

GTIN:
S/N
LOT
EXP.

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

F30009A01146

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964

Store dry material at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature].

Each imipenem and Cilastatin for Injection, USP (I.V.) vial contains Imipenem USP 250 mg (Anhydrous Equivalent) and Cilastatin Sodium USP equivalent to 250 mg Cilastatin. In addition, each vial contains 10 mg of sodium bicarbonate.
The sodium content is approximately 18.8 mg (0.8 meq).

For the Preparation of Intravenous Solutions and USUAL ADULT DOSAGE: See accompanying package insert.

Color Changes in solution from colorless to yellow do not affect potency.

After suspension, vial contents must be transferred to 100 mL of infusion solution prior to intravenous infusion.

Discard unused portion of the infusion solution where applicable.

Mfd for:
FRESENIUS KABI
Lake Zurich, IL 60047
Made in Italy

www.fresenius-kabi.com/us

(01)20363233349251

621263F

NDC 63323-349-25

349025

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1 PACKAGE (25 Single-Dose Vials)

FRESENIUS KABI