TauroLock™ U 25000

CE 0123

ANTIMICROBIAL CATHETER LOCK SOLUTION

TauroLock™ U25.000

CE 0123

Antibiotic Free

Indication:
TauroLock™-U25.000 is approved to be instilled into catheter-based devices for central venous access in order to maintain patency of the device. Additionally, by creating a hostile environment to bacterial and fungal growth, the risk of catheter-related bloodstream infections (CRBSI) is reduced.

Ingredients / Specification:
TauroLock™-U25.000 contains substances to ensure patency and provide infection control in the device. Active ingredients in TauroLock™-U25.000 are taurolidine, citrate (4%) and urokinase (25.000 IU). The product is sterile filter processed and supplied as a clear, sterile, non-pyrogenic solution. Urokinase is delivered in a separate vial and is reconstituted with the content of the ampoule immediately before use. Each single-dose ampoule contains 5 mL.

Instillation:
TauroLock™-U25.000 is prepared by adding TauroLock™ into the vial containing urokinase to obtain a clear solution. After treatment (e.g. dialysis), the system is thoroughly flushed with saline (min. 10 mL). Then, the clear fluid is to be instilled into the device lumen by exactly respecting the filling volume. TauroLock™-U25.000 remains in the system until the next treatment. The solution must be aspirated before initiating the next treatment. Flush the device with 10 mL of saline.

Contraindications:
TauroLock™-U25.000 is contraindicated for patients with a known allergy to taurolidine, citrate or urokinase, or for patients currently taking medication with known adverse interaction with taurolidine, citrate or urokinase.

Storage:
TauroLock™-U25.000 must be stored at 15 – 25 °C. It must be neither transported nor stored under freezing conditions. The reconstituted solution must be used instantly.

Note:
This information does not replace the instruction for use.

article-no.	description	packaging
TP-05	TauroLock™-U25.000 catheter lock solution 5 mL vial (single-dose)	5 x 5 ml

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