



TauroLockTM U25000

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 prevent CRBSI and bio-film formation. TauroLock™-U25.000 maintains patency without undesired side-effects.

Ingredients / Specification:

TauroLock™-U25.000 contains substances to ensure patency and provide infection control in the device. Active ingredients in TauroLock™-U25.000 are (cyclo)-taurolidine, citrate (4%) and urokinase (25.000 IU). Other components include water for injection.

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 contents of the ampoule just before use. Each single-use ampoule contains 5 mL.

It is to be used with a port or a catheter-based vascular access device. TauroLock™-U25.000 is instilled into the device lumen between treatments in order to make the internal flow passages resistant to clot formation and hostile to bacterial and fungal growth.

Instillation:

TauroLock™-U25.000 is prepared by adding TauroLock™ into the vial to get a clear solution. TauroLock™-U25.000 acts prophylactically against occlusions within the central venous access device. The clear and colourless solution is to be instilled into the device lumen between two dialysis treatments. The solution is withdrawn prior to the next treatment.

Contraindications:

TauroLock™-U25.000 is contraindicated for patients with a known adverse effect to (cyclo)-taurolidine, citrate or urokinase or when a patient is currently taking medication with known adverse interaction to citrate, (cyclo)-taurolidine or urokinase.

Storage:

TauroLock™-U25.000 must be stored at 15–25 °C. 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 use)	5 x 5 ml

Manufacturer:



TauroPharm GmbH
August-Bebel-Str. 51
D-97297 Waldbüttelbrunn
Tel.: +49 (0) 931 304 299-0

ISO 13485

Contact Us:

JTS Medcare Solutions (Pty) Ltd

cell: +27 722 178 183

email: info@jtsmedsol.co.za

web: www.jtsmedsol.co.za

WWW.TAUROLOCK.COM

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