



MD UreSil ® Universal Funnel Adapter - Luer Lock Medical device

INTENDED USE: For connexing a percutaneous drainage athlefer to a drainage bag or system. This product is intended for short-term use (less than 30 days). INTENDED PATTENT POPULATION: This product is intended to be a PATTENT POPULATION: This product is intended to be intended products. CUINCAL BEENETTES: Work as an accessory in conjunction with a percutaneous drainage cathlefer and a Careful product is intended only for use by full state drainage. CUINCAL BEENETTES: Work as an accessory in conjunction with a percutaneous full as using gravity or in conjunction with a percutaneous full as a product and the state of the system to remove full as using active scalar on or gravity. PERFORMANCE CHARACTERISTIC: UFALL is comprised of a tapend full met and a female leur connector and is used

with funneled catheters, for example, UreSil Ulti-flo Large Bore non-locking catheters (REF LBNL-series), to connect to a Gravity Drainage or Suction Drainage bag or system with standard luer. 8-20-218 rev. 04/10/2023 INTENDED FOR USE WITH: This accessory for a medical device is intended for use with funneled catheters for example. UreSil Ulti-flo Large Bore non-locking catheters (REF LBNL-series) to connect to a Gravity Drainage or Suction Drainage bag or system with standard luer. INSTRUCTIONS FOR USE: Used with funneled catheters, for example, UreSil Ulti-flo Large Bore non-locking catheters (REF LBNL-series), to connect to a Gravity Drainage or Suction Drainage bag or system with standard luer. CONTRAINDICATIONS: Currently no specific contraindications are known for this accessory. for a medical device. WARNING: The reuse of this single-use device can lead to patient infection and/or device malfunction. Sterile if package is unopened and undamaged. Do not use if the sterile package is damaged or is unintentionally opened before use. SAFE DISPOSAL: Dispose of used device in container marked for biohazard (i.e. contaminated with potentially infectious substances of human origin) REPORTING SERIOUS INCIDENTS: Any serious incident that has occurred in relation to the device should be reported to UreSil as the manufacturer and the Competent Authority of the Member State in which the user and/or natient is established

UreSil is a registered trademark of UreSil, LLC.



Importer in the European Community/ European Union: MedEnvoy, Prinses Margrietplantsoen 33, Suite 123, 2595 AM, The Hague, The Netherlands