Doc. No.:	RGI/IFU-09/1020T			
Issue No.:	04			
Rev. No.:	03			
Date	03.10.2022			

<u>Romsons®</u>

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URINE COLLECTING BAG

(With 'T' Tap Valve - Bag Capacity 2000ml)

REF GS-1020T

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i INSTRUCTIONS FOR USE

The device is biocompatible, Non-Toxic, Non-Pyrogenic , Phthalate Free, Sterile, Single Use.

Caution : Carefully read all instructions prior to use. Observe all warnings & precautions noted throughout these instructions.

Failure to do so may result in complications. To be administered by an expert qualified medical professional. This device is Sterile & Ready for Use. Sterility is guaranteed - if pack is undamaged. The device is for Single Use Only.

Device Description : Closed urine drainage system for short and long term use.Manufactured from soft clinical grade PVC sheeting. Reinforced eyelets allow easy hanging and provide maximum protection against tearing. Consist a 'T 'tap valve -.Kink resistant 90cm tube provided with universal tapered connector. Easy to read scale for quick determination of urine volume.Capacity-2000ml

Population of Use: Adult.

Intended User: To be administered by an expert qualified medical professional, Urologist & Trained nurse.

Intended Use : Used for short term urine collection & drainage. Maximum use period not beyond 7 days.

Preferred Route of Administration: Attached to the In-dewelled urethral catheter.

Contents : Urine Collection Bag (2000ml) With 'T 'tap valve .

Size : 2000ml.

Material of Construction : Polyvinyl Chloride, PVC Sheet, LDPE, Master Batch.

Storage : Store in a cool & dry place.

Indications : Any medical condition requiring management of collection & drainage of urine.

Contra-indications : Hypersensitivity to device material , Any abnormality associated with using of a urine bag, lower abdominal wounds or scarring overlying cellulitis.

Potential Complications / Risks : Dislodgement of connection from the urethral catheter, Temporary blockage due to folding of tubing, contamination of the bottom outlet

Warnings : This device is intended for Single use only.

DO NOT resterilize and /or reuse the device, as this can compromise the device performance (functionality) and may cause inadequacy, deterioration of the device technical factor, rendering the device non-functional and unfit for intended use and also this may increase the risk of cross contamination due to several aspects including inappropriate reprocessing.

Do not put the Device to Use after the Use by Date or Date of Expiry.

Re-use of single use device creates a potential risk for patient or use. It may lead to contamination and/or impairment of functional capability. Contamination and / or limited functionality of the device may lead to injury, illness of the patient.

Precautions :

- ((i) Check the integrity and functionality of the urine collecting bag before use. Do not use if the Unit Pack is open or damaged.
- (ii) Determine patient's condition and vitals status During device application / Operation.
- (iii) Dispose the device & its accessories after single use as bio-medical waste as per applicable laws.
- (iv) Do not Resterilize. Do not Re-use. Discard after single use.

(v) Do not put the Device to Use after the Use by Date or Date of Expiry.

Adverse Events : CAUTI, Microbial colonization, etc.

General Instructions : To be administered by an expert qualified medical professional. Use maximal sterile barrier precautions during administration. (These are only the guidelines & hence, user's medical experience / expertise in important).

Use Instructions :

- Check the packing carefully, if found damaged, torn or pierced discard the piece. Do not use if pack is damaged. (1)
- Wash up and scrub hands and preferably use pre sterile protective gloves. (2)
- Tear open the pack & drawout the device with care. (3)
- Close the Bottom Outlet 'T Type' of the Urine Bag. Remove the cap from the Tapered Connector. (4)
- (5)
- Connect the Tapered Connector to the Uretheral Catheter. (6)
- (7)Discard the Urine Bag after Use.

Mfr.: Romsons International (Unit-II), 59J(C) & 9, Noida Special Economic Zone, Noida Dadri Road, Phase-II, Noida-201305 (INDIA) ECREP OBELIS S.A, Bd. Général Wahis 53, 1030 Brussels, Belgium, Email: mail@obelis.net Basic UDI DI : 8902120A012RD

REF	Catalogue Number	MD	Medical Device	(2)	Donot Re-use	i	Consult Instructions for Use
LOT	Batch Code		Country of Manufacture	STERRIZE	Do Not Re- sterilize	溇	Keep away from sunlight
\sim	Date of Manufacture	STERILE EO	Sterilized Using Ethyele Oxide	UDI	Unique device Identifier	-	Keep Dry
Σ	Use by date	EC REP	Authorized Representative in the European Union		Do Not Use if Pack is Damaged	Λ	Caution

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~~	Manufacturer	0	Single Sterile barrier system	10 °C (104 °F)	Temperature Limit		
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