Doc. No.:	RGI/IFU-09/405RK
Issue No.:	04
Rev. No.:	03
Date	0310.2022

UNIVERSAL BILE BAG

(Bile Collection Bag)

REF

UN405RK

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INSTRUCTIONS FOR USE

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

Device Description: Device Consist of a PVC Bag (Capacity 500 ml). Bile Bag has Graduation from 25 ml to 500 ml as per case. These graduations are only for indicative purpose. Bile Bag on its proximal end consists of an Inlet Tube. This tube is followed by a Non Return Valve (NRV) to disable accidental backflow of fluid attached to a double tapered step connector this connector get fixed to the distal end of the T-tube Bile Drainage adaptor .Distal end of the Bile Bag consist of a outlet tube attached to a one quarter turn open and close adaptor . Which helps in instant drainage of collected fluid in the bag .Bile Bag consists of fur eyelets at the top and two eyelets at the bottom to enable Hanging / Attachment of Bile Bag .

Population of Use: Adult & Paediatrics.

Intended User: To be administered by an expert qualified medical professional.

Intended Use: Intended for the collection of drained bile. Maximum use period not more than 7 days

Preferred Route of Administration: Attached to the T-Tube(Kehr's T-Tube)

Contents : Sterile, One Unit in a Peel Open Pack.

Size: Capacity-500ml.

Material of Construction: PVC, PP, PE, MB.

Storage: Store in a cool & dry place.

Indications : Bile Duct Drainage, percutaneous drainage, percutaneous transhepatic biliary drainage, transhepatic gallbladder drainage, subhepatic collection, etc.

Contraindications: Use in patient with a known allergic reaction to any of the product components, Massive ascites, multiple intrahepatic obstructions, bleeding diathesis, etc.

Potential Complications / Risks : Acute biliary sepsis, persistent hyperbilirubinemia, malignant biliary obstructions, acute hemobilia.

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 factors, 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 user. 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 Bile 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) Conduct procedure under strict surgical protocol and ensure complete asepsis.
- (iv) Destroy the device & its accessories after single use as bio-medical waste as per applicable laws.
- (v) Do not Re-sterilize. Do not Re-use. Single use only.
- (vi) Do not put the Device to Use after the Use by Date or Date of Expiry.

Adverse Events: Biliary leakage, Biliary obstructions, Bag dislodgement from connector, Bile Spillage & contamination / infection, etc.

General Instructions:

To be administered by an expert qualified medical professional. Use maximal sterile barrier precautions during administration Dispose the device after use as bio-medical waste as per applicable laws.

Use Instructions :

- (1) Check the packing carefully, if found damaged, torn or pierced discard the piece.
- (2) Wash up and scrub hands and preferably use pre sterile protective gloves.
- (3) Peel open the pack & draw out the device with care.
- (4) Close & Close the Bottom Outlet of the Bile Bag.
- (5) Connect the drainage tap to the Kehr's / Ryles Connector / Wide Bore double tapered connector.
- (6) To drain, Open the drainage tap & then close after use.

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(7) Discard the Bile Bag after Single Patient Use. Mfr.: Romsons International (Unit-II), 59J(C) & 9, Noida Special Economic Zone, Noida Dadri Road, Phase-II, Noida-201305 (INDIA) EC REP OBELIS S.A, Bd. Général Wahis 53, 1030 Brussels, Belgium, Email: mail@obelis.net **Basic UDI DI**: 8902120F005SK REF Catalogue **Medical Device Donot Re-use** i MD Consult Number **Instructions for Use** LOT **Batch Code** ΛΨ Country of Do Not Re-Keep away from Manufacture sterilize sunlight Date of **Sterilized Using** UDI Unique device STERILE EO **Keep Dry** Manufacture Identifier EO Use by date Non-pyrogenic Do Not Use if Caution, Pack is Damaged and consult **Instructions** for Use Non Return EC REP Single Sterile **Authorized Temperature Limit** Valve Representative barrier in the European system Union **C**€₂₄₆₀

Manufacture