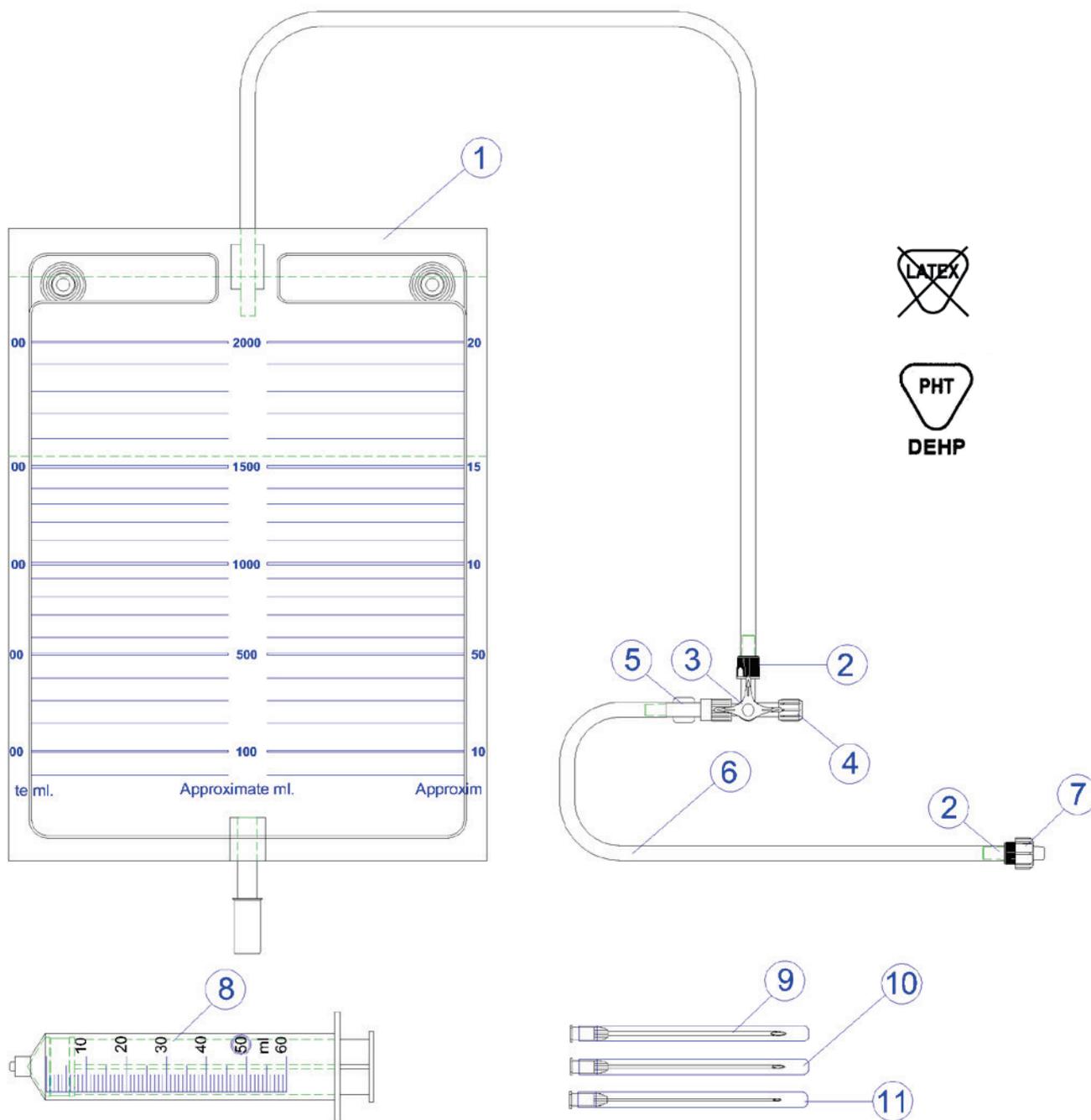


# PRODUCT TECHNICAL DATA SHEET



## DESCRIPTION

### TORASET – Thoracentesis drainage set with 14G-16G-19G needles



CODE	BAG VOLUME ml	NEEDLES	CARTON pcs.	GTIN-UDI SINGLE PRODUCT	GTIN-UDI CARTON
M032010S	2000	14G/16G/19G – 80 mm	30	18053676294524	38053676294528
M032000S	2000	14G/16G/19G – 55 mm	30	18053676294494	38053676294498

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## ● PRODUCT COMPOSITION:

The device consists of:

1. **2000 ml collection bag** in **PVC**, with non-return valve, "push-pull" drain tap, connection tube I.D. 5.2 mm x O.D. 6.5 mm and length cm 90;
2. **Male Luer Lock connector** in **ABS**;
3. **3-way stopcock** (360° rotation) made in **Polyethylene** and **Polycarbonate**, with fluid direction indicators to identify the different procedure phases (aspiration of pleural effusion from the patient; discharge into collection bag);
4. **Protective cap** in blue **ABS**;
5. **Female luer lock connector** in **ABS**;
6. **Tube** in **DEHP FREE PVC**, I.D. 4.8 mm x O.D. 6.8 mm and length cm 40;
7. **Vented cap** in **Polyethylene**;
8. **60 ml syringe** to be connected to the 3-way stopcock, made in **Polypropylene** and **Synthetic Isoprene rubber**;
9. **Stainless Steel 14G Needle** with a length of 55 or 80 mm (see table), packaged in microperforated bag;
10. **Stainless Steel 16G Needle** with a length of 55 or 80 mm (see table), packaged in microperforated bag;
11. **Stainless Steel 19G Needle** with a length of 55 or 80 mm (see table), packaged in microperforated bag.

## ● INTENDED USE:

Closed circuit system thoracentesis, particularly suitable for the aspiration of pleural effusion liquid.

Following an evaluation by the healthcare professional, and under his direct control, the device can be used for paracentesis procedures.

## ● PACKAGING:

Primary packaging: single device in micro-perforated **PE** inner pouch bag + blister in medical grade paper/**PP-PE** film.  
Secondary packaging: carton with 30 pieces.

## ● PRODUCTION PROCESS:

The device is manufactured in accordance with HMC Premedical S.p.A. Quality System and in comply with the requirements of the standard EN ISO 13485.

## ● CONTROL ON THE PRODUCT:

In all stages of processing, according to internal procedures and to sampling plans defined by norm ISO 2859-1.

## ● CLASSIFICATION:

**Class IIa** sterile.

## ● STERILIZATION:

Product sterilized by ethylene oxide according to a validated course of treatment in accordance with current regulations.

Shelf life: **5 years** from the date of sterilization.

For single use only and non re-sterilizable.

## ● STORAGE:

Standard storage procedures and conditions.

## ● DISPOSAL:

The user must follow the legal regulations regarding disposal of hospital waste.

## ● WARNINGS:

The device must be used exclusively by healthcare professionals.

## ● REGISTRATION TO ITALIAN MEDICAL DEVICES REPERTOIRE:

CND: **A060204**

RDM: **1575159**

GMDN: **36787**

## ● UNIT OF SALE:

Carton with **30 pieces**.

## ● GTIN-UDI:

See table.



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# PRODUCT TECHNICAL DATA SHEET



REV.	CHANGES	ISSUED	VERIFIED AND APPROVED	DATE
00	First issue – english language	E. Benassi	D. Bosetti	24/10/2017
01	Data sheet revision; drawing and product composition update	E. Benassi	S. Tralli	31/01/2020
02	Intended use updated	E. Benassi	S. Tralli	29/09/2020
03	Leggl manufacturer's address update	E. Benassi	S. Tralli	28/04/2021
08	Data sheet update; jump to rev. 08 to align with italian version	E. Benassi	S. Tralli	09/09/2021
09	Table update with GTIN-UDI	E. Benassi	S. Tralli	02/02/2022
10	Drawing update	E. Benassi	S. Tralli	11/02/2022