

# **NOBAMED Paul Danz AG**

# NOBADRAPE® Drape

### **REF 760045**

# **Product Description, Intended use, Application**

The 2-ply blue drape (45 cm x 75 cm) is individually packed in sterile packaging. It is used for sterile covering of patients and equipment in the OR. The top layer consists of an absorbent polypropylene nonwoven. The bottom layer consists of a moisture-impermeable polyethylene film providing a reliable barrier protection. Easily removable control stickers on the packaging facilitate surgical documentation.

### Composition

Polypropylene, polyethylene

### **Contraindications**

The product should not be used in the case of a known allergy against the material.

#### Incident reporting

According to MDR (EU) 2017/745, if serious incidents occur in relation to the device, they must be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

# Normative and Regulative Requirements, Common Standards

Medical Device according to MDD 93/42/EEC, MDR (EU) 2017/745.

The product complies with the requirements of the **DIN EN 13795-1**, "Surgical clothing and drapes – Requirements and test methods – Part 1: Surgical drapes and gowns (requirements of the performance level 'high' and 'standard').

Sterilization of the product complies with DIN EN 11135.

The product does not contain dangerous toxic substances according to REACH.

## **Packaging**

Primary: paper-film

packaging

Secondary: folding box

made of cellulose

Tertiary: carton made of

cellulose

### Symbols used in labelling

Explanation at <a href="https://www.nobamed.com">www.nobamed.com</a>



Marking on all packaging levels with CE and according to DIN EN ISO 15223-1- and DIN EN ISO 20417

#### **Storage and transport**

To be stored in a dry and dust-free environment, protect from sunlight.

### Sterile device

Before using a sterile product, visually inspect the packaging to ensure that it is intact.

### Single use device

Reusing a single use medical device can lead to microbiological danger. Reprocessing for reuse can decrease the product's performance significantly.

### Disposal

According to locally applicable legal regulations and standards of infection prophylaxis.

Date of information: March 21, 2023 [REV 7] Replaced: January 08, 2021 [Rev6]