



**NOBAMED Paul Danz AG**

## **NOBASORB<sup>®</sup>**

**REF 840015**

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

The non-sterile, absorbent wound pad with a size of **15cm x 25cm** consists of a soft nonwoven which wraps an absorbent core of soft, bleached fluff pulp and a cellulose layer for the distribution of secretions. On its back, the pad is provided with a nonwoven layer that is impenetrable to liquids. The absorbent wound pad is used for the treatment and cleansing of heavily exuding, infected or chronic wounds. It may also be used in the case of weeping wound healing disorders, for wound padding, and as secondary dressing for heavily exuding wounds. It is for single use.

### **Composition**

Polypropylene, cellulose

### **Contraindications**

All medium to lightly exuding wounds, in particular during granulation and epithelization stage, as well as deep and undermined wounds, should not be treated with absorbent wound pads.

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

### **Note**

The product must be sterilized in accordance with a validated sterilization method before use on open wounds. The product can be sterilized with moist heat at 121°C/ 134°C, 2 to 3 bar, according to DIN EN 17665 or ethylene oxide according to DIN EN ISO 11135.

### **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 *MDR (EU) 2017/745*.

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

### **Packaging**

Primary packaging: paper bag  
Secondary packaging: carton made of cellulose

### **Symbols used in labelling**

Explanation at [www.nobamed.com](http://www.nobamed.com)



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

### **Storage and Transport**

Dry and dustfree

### **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.