

Statement of Procedure Pack Producer

(as stipulated by Regulation (EU) 2017/745 (MDR))

We herewith declare, that for the products listed in Table 1:

- We have verified the mutual compatibility of the devices and, if applicable other products, in accordance with their manufacturers' instructions and have carried out our activities in accordance with those instructions;
- We package the procedure pack and supply relevant information to users incorporating the information supplied by the manufacturers of the devices or other products that are put together;
- The of combining devices and, if applicable, other products, is subject to appropriate methods of internal monitoring, verification and validation.
- Sterilization, if applicable, is carried out in accordance with the manufacturer's instructions. The sterilization process is under the supervision of the Notified Body: TÜV NORD CERT GmbH, Identification No. 0044

Table 1: Product List

REF	Description
160030	Verband-Set standard, steril

Date: 2024-07-08

Promedical AG



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