# EU Quality Assurance Certificate

| Certificate no. | Final Assessment Report no. | Effective date | Expiry date |
|---|---|---|---|
| 0462GB449230328 | 0462AU28F | 2023-03-28 | 2027-10-19 |

This is to certify that the quality system of

## HANS HEPP GmbH & Co. KG Verbandstoff-Fabrik

Georgswerder Damm 16, 20539 Hamburg, Germany

SRN: DE-MF-000006024

For production, and final product inspection/testing of

**Medical devices/groups of medical devices at locations as listed on the following pages**

Has been assessed and found to comply with respect to

**The conformity assessment procedure described in Annex XI Part A of Regulation (EU) 2017/745 on Medical Devices**

Any applicable limitations for certain medical devices are included in the following list or recorded in the final assessment report. This certification is subject to surveillance by DNV MEDCERT.

Place and date
Hamburg, 2023-03-28

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-MDR-096

For the issuing office
**DNV MEDCERT GmbH – Notified Body 0482**
**Pilatuspool 2, 20355 Hamburg, Germany**

**Marcus Harder**
Director Certification

The certificate is only valid when provided entirely with all of its pages. To verify the validity of this certificate, contact info@medcert.de

Certificate no.: 0462GB449230328
Place and date: Hamburg, 2023-03-28

## Sites covered by this certificate

HANS HEPP GmbH & Co. KG Verbandstoff-Fabrik, Georgswerder Damm 16, 20539 Hamburg, Germany

Certificate no.: 0462GB449230328
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## Products covered by this certificate

### Class I medical devices

For class I medical devices placed on the market in sterile condition (class Is), the audit of the quality management system was limited to the aspects relating to establishing, securing, and maintaining sterile conditions.

| Category | Class | Medical devices/groups of medical devices |
|---|---|---|
| MDN 1204 | Is | Non-active non-implantable devices for wound and skin care |