## Landesamt für soziale Dienste Schleswig-Holstein

# UNION FORMAT FOR A WHOLESALE DISTRIBUTION AUTHORISATION (MEDICINAL PRODUCTS FOR HUMAN USE)

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т.	Aut	horisation	munn	ושט

2. Name of Authorisation Holder

3. Legally registered address of Authorisation Holder

4. Address(es) of Site(s)

5. Scope of authorisation (complete for each site under 4)

6. Legal basis of authorisation

7. Name of responsible officer of the competent authority of the member state granting the wholesaling authorisation

8. Signature

9. Date

10. Annexes attached

: DE\_SH\_01\_WDA\_2015\_0017

: Care Competence Pharmazeutische Großhandels- und Beratungs-GmbH

: Henstedter Straße 1, Wakendorf II, 24558, Germany

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: ANNEX 1

: Art.77(1) of Directive 2001/83/EC

: Confidential, Confidential

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: Annex 1 Scope of wholesale distribution authorisation

Annex 2 (Optional) Address(es) of contract wholesale distribution sites and their authorisation number

Annex 3 (Optional) Name(s) of responsible person(s)

Annex 4 (Optional) Date of Inspection on which authorisation was granted

Annex 5 (Optional) Additional provisions based on national requirements

#### **ANNEX 1**

#### SCOPE OF WHOLESALE DISTRIBUTION AUTHORISATION

**Name and address of the site:** Care Competence Pharmazeutische Großhandels- und Beratungs-GmbH, Henstedter Straße 1, Wakendorf II, 24558, Germany

### 1. MEDICINAL PRODUCTS

- 1.1 with a Marketing Authorisation in EEA country(s)
- 1.2 without a Marketing Authorisation in the EEA and intended for EEA market\*
- 1.3 without a Marketing Authorisation in the EEA and intended for exportation

## 2. AUTHORISED WHOLESALE DISTRIBUTION OPERATIONS

- 2.1 Procurement
- 2.2 Holding
- 2.3 Supply
- 2.4 Export

## 3. MEDICINAL PRODUCTS WITH ADDITIONAL REQUIREMENTS

- 3.1 Products according to Art. 83 of 2001/83/EC \*\*
  - 3.1.2 Medicinal products derived from blood
  - 3.1.3 Immunological medicinal products
- 3.3 Cold chain products(requiring low temperature handling)

Any restrictions or clarifying remarks (for all users): National legal basis of authorisation: § 52 a Section 1 of German Drug Law (Arzneimittelgesetz)

\*Art 5 of Directive 2001/83/EC or Art 83 of Regulation EC/726/2004

\*\*Without prejudice to further authorisations as may be required according to national legislation



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