

Regierungspraesidium Tuebingen Leitstelle
Arzneimittelueberwachung Baden-Wuerttemberg

UNION FORMAT FOR A WHOLESALE DISTRIBUTION AUTHORISATION

Human medicinal products

1. Authorisation Number : DE_BW_01_WDA_2024_0001_Docpharm GmbH
2. Name of Authorisation Holder : Docpharm GmbH
(ORG-100040197 / LOC-100065411)
3. Address(es) of Site(s) : Borsigstrasse 3, Muenklingen, Weil Der Stadt, Baden-Wuerttemberg, 71263, Germany
Rheinstrasse 219, Oos, Baden-Baden, Baden-Wuerttemberg, 76532, Germany
4. Legally registered address of Authorisation Holder : Borsigstrasse 3, Muenklingen, Weil Der Stadt, Baden-Wuerttemberg, 71263
5. Scope of authorisation (complete for each site under 4) : ANNEX 1
6. Legal basis of authorisation : Art. 77(1) of Directive 2001/83/EC
7. Name of responsible officer of the competent authority of the member state granting the wholesaling authorisation : Confidential, Confidential
8. Signature :
9. Date : 2024-07-03
10. Annexes attached : 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: Docpharm GmbH
(ORG-100040197 / LOC-100065411)
, Borsigstrasse 3, Muenklingen, Weil Der Stadt, Baden-Wuerttemberg, 71263, Germany

Human medicinal products

1. MEDICINAL PRODUCTS 1.1. with a Marketing Authorisation in EEA country(s)
2. AUTHORISED WHOLESALE DISTRIBUTION OPERATIONS 2.1. Procurement 2.3. Supply
3. MEDICINAL PRODUCTS WITH ADDITIONAL REQUIREMENTS 3.1.3. Immunological medicinal products 3.3. Cold chain products(requiring low temperature handling)

Name and address of the site: Docpharm GmbH
(ORG-100040197 / LOC-100091930)
, Rheinstrasse 219, Oos, Baden-Baden, Baden-Wuerttemberg, 76532, Germany

Human medicinal products

1. MEDICINAL PRODUCTS 1.1. with a Marketing Authorisation in EEA country(s)
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*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