

**Government of Upper Bavaria - Central Authority for Supervision of  
Medicinal Products in Bavaria (GMP/GCP)**

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

1. Authorisation Number : DE\_BY\_04\_WDA\_2018/ROB-55Ph-2678.Ph\_3-122-17-2
2. Name of Authorisation Holder : ilapo Internationale Ludwigs-Arzneimittel GmbH & Co. KG
3. Legally registered address of Authorisation Holder : Friedenheimer Brücke 21, München, 80639, Germany
4. Address(es) of Site(s) : Friedenheimer Brücke 21, München, 80639, Germany
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 : 2018-03-14
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

EudraGMDP

## **ANNEX 1**

### **SCOPE OF WHOLESALE DISTRIBUTION AUTHORISATION**

**Name and address of the site:** ilapo Internationale Ludwigs-Arzneimittel GmbH & Co. KG,  
Friedenheimer Brücke 21, München, 80639, 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 \*\*
  - 3.1.1 Narcotic or psychotropic products
  - 3.1.2 Medicinal products derived from blood
  - 3.1.3 Immunological medicinal products
- 3.3 Cold chain products(requiring low temperature handling)

\*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