Latvian State Agency of Medicines

UNION FORMAT FOR A WHOLESALE DISTRIBUTION AUTHORISATION

(MEDICINAL PRODUCTS FOR HUMAN USE)

1. Authorisation Number	:	L00120
2. Name of Authorisation Holder	:	SIA "GP Nord"
3. Legally registered address of Authorisation Holder	:	Kurzemes prospekts 23, Rīga, LV-1067, Latvia
4. Address(es) of Site(s)	:	Kurzemes prospekts 23, Rīga, LV-1067, Latvia
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	:	2019-04-04
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: SIA "GP Nord" zāļu lieltirgotava, Kurzemes prospekts 23, Rīga, LV-1067, Latvia

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
3.3 Cold chain products(requiring low temperature handling)
3.4 Other products: 3.4.1 veterinary medicinal products

Any restrictions or clarifying remarks (for all users): 3.1.1.1. psychotropic medicinal products; 3.1.1.3. narcotic and equivalent thereto psychotropic medicinal products
1)☐Annex 1 Paragraph 3.3. - it is permitted to distribute only medicines requiring cold temperature storage (from 2°C to 8°C)

*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