# Medicines and Healthcare products Regulatory Agency

# WHOLESALE DISTRIBUTION AUTHORISATION

# (MEDICINAL PRODUCTS FOR HUMAN USE)

1. Authorisation Number UK WDA(H) 49718

2. Name of Authorisation Holder NEURAXPHARM UK LIMITED

3. Legally registered address of NEURAXPHARM UK LIMITED, SUITE 2, ARLINGTON FLEX, THIRD FLOOR, BUILDING

Authorisation Holder 1420, ARLINGTON BUSINESS PARK, THEALE, READING, RG7 4SA, UNITED KINGDOM

NEURAXPHARM UK LTD, SUITE 2, ARLINGTON FLEX, THIRD FLOOR, BUILDING 1420, 4. Address(es) of Site(s)

ARLINGTON BUSINESS PARK, THEALE, READING, RG7 4SA, UNITED KINGDOM

5. Scope of authorisation (complete for

each site under 4)

ANNEX 1

6. Legal basis of authorisation Regulation 18 of the Human Medicines Regulations 2012

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

authorisation

9. Annexes attached

Confidential

10/09/2024 8. Date

> 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 Additional provisions

# ANNEX 1

#### SCOPE OF WHOLESALE DISTRIBUTION AUTHORISATION

Name and address of the site:

NEURAXPHARM UK LTD, SUITE 2, ARLINGTON FLEX, THIRD FLOOR, BUILDING 1420, ARLINGTON BUSINESS PARK, THEALE, READING, RG7 4SA, UNITED KINGDOM

#### 1. MEDICINAL PRODUCTS

- 1.1 With "an authorisation" (a UK, Great Britain or Northern Ireland Marketing Authorisation, an Article 126a authorisation, a certificate of registration or traditional herbal registration)
- 1.2 Without "an authorisation" (a UK, Great Britain or Northern Ireland Marketing Authorisation, an Article 126a authorisation, a certificate of registration or traditional herbal registration) in GB or EEA and intended for the UK market

#### 2. AUTHORISED WHOLESALE DISTRIBUTION OPERATIONS

2.1 Procurement

Issue Date: 10 Sep 2024

- 2.3 Supply
- 2.6 Products imported from countries on a list
- 2.6a Products certified under Article 51 of Directive 2001/83/EC
- 2.6b Products not certified under Article 51 of Directive 2001/83/EC

# 3. MEDICINAL PRODUCTS WITH ADDITIONAL REQUIREMENTS

- 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)

# Any restrictions or clarifying remarks (for all users)

4 Categories of Products Handled at this Site: 4.1 Prescription Only Medicines

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