Medicines and Healthcare products Regulatory Agency

WHOLESALE DISTRIBUTION AUTHORISATION

(MEDICINAL PRODUCTS FOR HUMAN USE)

TYPHARM LIMITED 2. Name of Authorisation Holder 3. Legally registered address of TYPHARM LIMITED, UNIT 39, MAHONEY GREEN, GREEN LANE WEST, RACKHEATH, Authorisation Holder NORWICH, NR13 6JY, UNITED KINGDOM TYPHARM LIMITED, UNIT 39, MAHONEY GREEN, GREEN LANE WEST, RACKHEATH 4. Address(es) of Site(s) NORWICH, NR13 6JY, UNITED KINGDOM 5. Scope of authorisation (complete for ANNEX 1 each site under 4) 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 Confidential state granting the wholesaling authorisation 8. Date 11/10/2024

UK WDA(H) 551

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:

TYPHARM LIMITED, UNIT 39, MAHONEY GREEN, GREEN LANE WEST, RACKHEATH, NORWICH, NR13 6JY, 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

1.3 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 the UK and not intended for the UK market

2. AUTHORISED WHOLESALE DISTRIBUTION OPERATIONS

Authorisation Number
Name of Authorisation Holds

9. Annexes attached

- 2.1 Procurement
- 2.2 Holding
- 2.3 Supply
- 2.4 Export
- 2.6 Products imported from countries on a list
- 2.6a Products certified under Article 51 of Directive 2001/83/EC

3. MEDICINAL PRODUCTS WITH ADDITIONAL REQUIREMENTS

3.1.1 Narcotic or psychotropic products

Any restrictions or clarifying remarks (for all users)

4 Categories of Products Handled at this Site: 4.1 Prescription Only Medicines, 4.2 General Sales List, 4.4 Pharmacy