Medicines and Healthcare products Regulatory Agency

WHOLESALE DISTRIBUTION AUTHORISATION

(MEDICINAL PRODUCTS FOR HUMAN USE)

1. Authorisation Number

2. Name of Authorisation Holder

3. Legally registered address of Authorisation Holder

4. Address(es) of Site(s)

5. Scope of authorisation (complete for each site under 4)

6. Legal basis of authorisation

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

8. Date

9. Annexes attached

UK WDA(H) 17901

ASTRAZENECA UK LIMITED

ASTRAZ<mark>ENECA UK LIMITED, 1 FRANCIS CRICK AVENUE, CAMBRIDGE BIOMEDICAL</mark> CAMPUS, CAMBRIDGE, CB2 0AA, UNITED KINGDOM

ASTRAZENECA UK LIMITED, UK OPERATIONS MACCLESFIELD, CHARTER WAY, SILK ROAD BUSINESS PARK, MACCLESFIELD, SK10 2NA, UNITED KINGDOM

ANNEX 1

Regulation 18 of the Human Medicines Regulations 2012

Confidential

15/05/2025

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:

ASTRAZENECA UK LIMITED, UK OPERATIONS MACCLESFIELD, CHARTER WAY, SILK ROAD BUSINESS PARK,

MACCLESFIELD, SK10 2NA, 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

- 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

3.1.2 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, 4.2 General Sales List, 4.4 Pharmacy

