# Medicines and Healthcare products Regulatory Agency

## WHOLESALE DISTRIBUTION AUTHORISATION

## (MEDICINAL PRODUCTS FOR HUMAN USE)

1. Authorisation Number UK WDA(H) 52206

2. Name of Authorisation Holder ZEYMOS PHARMA LTD

3. Legally registered address of ZEYMOS PHARMA LTD, UNIT 19, PARK ROYAL METRO CENTRE, BRITANNIA WAY,

Authorisation Holder LONDON, NW10 7PA, UNITED KINGDOM

4. Address(es) of Site(s) ZEYMOS PHARMA LTD, UNIT 19, PARK ROYAL METRO CENTRE, BRITANNIA WAY,

LONDON, NW10 7PA, UNITED KINGDOM

5. Scope of authorisation (complete for

each site under 4)

9. Annexes attached

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 Confidential

granting the wholesaling authorisation

8. Date 07/07/2025

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:

ZEYMOS PHARMA LTD, UNIT 19, PARK ROYAL METRO CENTRE, BRITANNIA WAY, LONDON, NW10 7PA, 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)
- 1.3 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

- 2.1 Procurement
- 2.2 Holding
- 2.3 Supply

Issue Date: 07 Jul 2025

- 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

Issue Date: 07 Jul 2025