# Medicines and Healthcare products Regulatory Agency

### WHOLESALE DISTRIBUTION AUTHORISATION

## (MEDICINAL PRODUCTS FOR HUMAN USE)

1. Authorisation Number UK WDA(H) 4425

2. Name of Authorisation Holder AVENTIS PHARMA LIMITED

3. Legally registered address of

AVENTIS PHARMA LIMITED, 410 THAMES VALLEY PARK DRIVE, READING, RG6 1PT, UNITED KINGDOM

Authorisation Holder

5....<u>-</u>2.......

4. Address(es) of Site(s)

AVENTIS PHARMA LIMITED T\A SANOFI, 410 THAMES VALLEY PARK DRIVE, READING,

RG6 1PT, 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 Confidential

granting the wholesaling authorisation

8. Date 10/11/2025

Annex 1 Scope of wholesale distribution authorisation Annex 2 (Optional) Address(es) of contract wholesale distribution sites and their authorisation number Annex 3 (Optional)

9. Annexes attached

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:

AVENTIS PHARMA LIMITED T\A SANOFI, 410 THAMES VALLEY PARK DRIVE, READING, RG6 1PT, 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
- 2.3 Supply
- 2.4 Export

Issue Date: 10 Nov 2025

- 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
- 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

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