

Medicines and Healthcare products Regulatory Agency

WHOLESALE DISTRIBUTION AUTHORISATION

(MEDICINAL PRODUCTS FOR HUMAN USE)

1. Authorisation Number UK WDA(H) 15694
2. Name of Authorisation Holder ALLOGA UK LIMITED
3. Legally registered address of Authorisation Holder
ALLOGA UK LIMITED, AMBER PARK 1, 2, AND 3, BERRISTOW LANE, SOUTH NORMANTON, ALFRETON, DE55 2FH, UNITED KINGDOM
ALLOGA UK LTD, AMBER PARK 9, UNIT 1, FARMWELL LANE, SUTTON-IN-ASHFIELD, NG17 1BX, UNITED KINGDOM
ALLOGA UK LIMITED, AMBER PARK 1, 2, AND 3, BERRISTOW LANE, SOUTH NORMANTON, ALFRETON, DE55 2FH, UNITED KINGDOM
ALLOGA UK LIMITED, AMBER PARK 7A, UNIT CW72, FARMWELL LANE, SUTTON-IN-ASHFIELD, NG17 1BX, UNITED KINGDOM
4. Address(es) of Site(s)
ALLOGA UK LIMITED, AMBER PARK 5 AND 6, UNIT C2, FARMWELL LANE, SOUTH NORMANTON, ALFRETON, DE55 2JX, UNITED KINGDOM
ALLOGA UK LIMITED, UNIT 5, FARMWELL LANE, SUTTON-IN-ASHFIELD, NG17 1BX, UNITED KINGDOM
ALLOGA UK LIMITED, AMBER PARK 8, DERBY 370, DERBY COMMERCIAL PARK, RAYNESWAY, DERBY, DE21 7HW, 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 Confidential
8. Date 09/07/2024
9. Annexes attached
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:

ALLOGA UK LTD, AMBER PARK 9, UNIT 1, FARMWELL LANE, SUTTON-IN-ASHFIELD, NG17 1BX, 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

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

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

3.1.4 Radiopharmaceutical (including radionuclide kits)

3.1.5 Products authorised under regulation 174 (supply in response to spread of pathogenic agents etc)

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, 4.5 Traditional Herbal Medicinal products

Name and address of the site:

ALLOGA UK LIMITED, AMBER PARK 1, 2, AND 3, BERRISTOW LANE, SOUTH NORMANTON, ALFRETON, DE55 2FH, UNITED KINGDOM

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3.1.2 Medicinal products derived from blood

3.1.3 Immunological medicinal products

3.1.4 Radiopharmaceutical (including radionuclide kits)

3.1.5 Products authorised under regulation 174 (supply in response to spread of pathogenic agents etc)

3.3 Cold chain products (requiring low temperature handling)

Any restrictions or clarifying remarks (for all users)

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Name and address of the site:

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2.4 Export

2.5 Other Activities

Thaw labelling of Pfizer and Moderna Covid-19 vaccine

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- 3.1.1 Narcotic or psychotropic products
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- 3.1.3 Immunological medicinal products
- 3.1.4 Radiopharmaceutical (including radionuclide kits)
- 3.1.5 Products authorised under regulation 174 (supply in response to spread of pathogenic agents etc)
- 3.3 Cold chain products (requiring low temperature handling)

Any restrictions or clarifying remarks (for all users)

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