Medicines and Healthcare products Regulatory Agency

WHOLESALE DISTRIBUTION AUTHORISATION

(MEDICINAL PRODUCTS FOR HUMAN USE)

1. Authorisation Number UK WDA(H) 57452

2. Name of Authorisation Holder MHM PHARMACEUTICALS LIMITED

3. Legally registered address of MHM PHARMACEUTICALS LIMITED, UNIT 6D, LONGDEN ROAD, LEVENSHULME,

Authorisation Holder MANCHESTER, M12 5SJ, UNITED KINGDOM

MHM PHARMACEUTICALS LTD, UNIT 6D, LONGDEN ROAD, LEVENSHULME,

4. Address(es) of Site(s)

MANCHESTER, M12 5SJ, 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 granting the wholesaling authorisation

Confidential

8. Date 08/01/2024

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:

MHM PHARMACEUTICALS LTD, UNIT 6D, LONGDEN ROAD, LEVENSHULME, MANCHESTER, M12 5SJ, 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
- 1.4 With a Marketing Authorisation in EEA member state(s) and intended for the GB parallel import market

Issue Date: 08 Jan 2024

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
- 3.3 Cold chain products (requiring low temperature handling)

Any restrictions or clarifying remarks (for all users)

4 Categories of Products Handled at this Site: 4.3 General Sales List Only

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