Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number UK MIA 43086

2: Name of authorisation holder TARGET HEALTHCARE LIMITED

TARGET HEALTHCARE LIMITED, QUANTUM HOUSE, HOBSON INDUSTRIAL ESTATE, HOBSON, NEWCASTLE UPON TYNE,

NE16 6EA, UNITED KINGDOM

TARGET HEALTHCARE LIMITED, UNIT 19J, WHITE ROSE WAY, FOLLINGSBY PARK, GATESHEAD, NE10 8YX, UNITED KINGDOM

TARGET HEALTHCARE LIMITED, QUANTUM HOUSE, HOBSON

4: Legally registered address of authorisation holder INDUSTRIAL ESTATE, HOBSON, NEWCASTLE UPON TYNE,

NE16 6EA, UNITED KINGDOM

5: Scope of authorisation and dosage formsANNEX 1 and/ or ANNEX 2

6: Legal Basis of authorisation Regulation 17 of The Human Medicines Regulations 2012 (SI

2012/1916)

7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation

Confidential

8: Authorisation Date 12/06/2025

9: Annexes attached Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

3: Address(es) of manufacturing site(s)

Annex 1

Name and address of the site:

TARGET HEALTHCARE LIMITED, QUANTUM HOUSE, HOBSON INDUSTRIAL ESTATE, HOBSON, NEWCASTLE UPON TYNE, NE16 6EA, UNITED KINGDOM

Human Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

[1.2] Non-sterile products

[1.2.1] Non-Sterile Products (processing operations for the following dosage forms)

[1.2.1.6] Liquids for internal use

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[1.2.1.9] Pressurised preparations

Special Requirements

CANNABIDIOL/DRONABINOL

[1.2.2] Batch certification

[1.5] Packaging

[1.5.1] Primary packaging

[1.5.1.6] Liquids for internal use

[1.5.1.9] Pressurised preparations

[1.5.2] Secondary packaging

[1.6] Quality control testing

[1.6.3] Chemical/Physical

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

TARGET HEALTHCARE LIMITED, UNIT 19J, WHITE ROSE WAY, FOLLINGSBY PARK, GATESHEAD, NE10 8YX, UNITED KINGDOM

Human Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

Part 1 - MANUFACTURING OPERATIONS

[1.1] Sterile Products

[1.1.3] Batch certification

[1.2] Non-sterile products

[1.2.2] Batch certification

[1.5] Packaging

[1.5.2] Secondary packaging

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

[2.2] Batch certification of imported medicinal products

[2.2.2] Non-sterile products

[2.3] Other Importation Activities

[2.3.1] Site of Physical Importation

Issue Date: 12 Jun 2025