# Medicines and Healthcare products Regulatory Agency

### MANUFACTURER'S AUTHORISATION

1: Authorisation Number

2: Name of authorisation holder

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

4: Legally registered address of authorisation holder

5: Scope of authorisation and dosage forms

6: Legal Basis of authorisation

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

8: Authorisation Date

9: Annexes attached

UK MIA 11311

TILLOMED LABORATORIES LIMITED

TILLOMED LABORATORIES LIMITED, 220 BUTTERFIELD, GREAT MARLINGS, LUTON, LU2 8DL, UNITED KINGDOM

TILLOMED LABORATORIES LIMITED, 220 BUTTERFIELD, GREAT MARLINGS, LUTON, LU2 8DL, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations 2012 (SI

2012/1916)

Confidential

17/06/2025

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

#### TILLOMED LABORATORIES LIMITED, 220 BUTTERFIELD, GREAT MARLINGS, LUTON, LU2 8DL, UNITED KINGDOM

**Human Medicinal Products** 

**Authorised Operations** 

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

## Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

#### [ 2.2 ] Batch certification of imported medicinal products

[2.2.1] Sterile Products

[2.2.1.1] Aseptically prepared

[2.2.1.2] Terminally sterilised

[ 2.2.2 ] Non-sterile products

#### [ 2.3 ] Other Importation Activities

[ 2.3.2 ] Importation of Intermediate which undergoes further processing

[ 2.3.4 ] Products authorised under regulation 174 (supply in response to spread of pathogenic agents etc)
Semi finish / Bulk

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