# 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 8596** 

**EUROAPI UK LIMITED** 

EUROAPI UK LIMITED, 37 HOLLANDS ROAD, HAVERHILL, CB9 8PU, UNITED KINGDOM

EUROAPI UK LIMITED, 37 HOLLANDS ROAD, HAVERHILL, CB9 8PU, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations

2012 (SI 2012/1916)

Confidential

10/04/2025

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

### Annex 1

Name and address of the site:

EUROAPI UK LIMITED, 37 HOLLANDS ROAD, HAVERHILL, CB9 8PU, UNITED KINGDOM

**Human Medicinal Products** 

**Authorised Operations** 

MANUFACTURING OPERATIONS (according to part 1)

#### Part 1 - MANUFACTURING OPERATIONS

[ 1.1 ] Sterile Products

[1.1.3] Batch certification

[ 1.2 ] Non-sterile products

[ 1.2.2 ] Batch certification

[ 1.3 ] Biological medicinal products

[ 1.3.2 ] Batch certification

[1.3.2.2] Immunological products

[ 1.3.2.3 ] Cell therapy products

[ 1.3.2.5 ] Biotechnology products

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[ 1.3.2.6 ] Human or animal extracted products [ 1.5 ] Packaging [1.5.1] Primary packaging

[ 1.5.1.6 ] Liquids for internal use

[1.5.2] Secondary packaging

[ 1.6 ] Quality control testing

[ 1.6.2 ] Microbiological: non-sterility

[ 1.6.3 ] Chemical/Physical

[ 1.6.4 ] Biological

