

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA 8215
<b>2: Name of authorisation holder</b>	KENT PHARMACEUTICALS LIMITED
<b>3: Address(es) of manufacturing site(s)</b>	KENT PHARMACEUTICALS LIMITED, UNIT 200, WESTMINSTER 42, WESTMINSTER INDUSTRIAL ESTATE, REPTON ROAD, MEASHAM, SWADLINCOTE, DE12 7DT, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	KENT PHARMACEUTICALS LIMITED, UNIT 200, WESTMINSTER 42, WESTMINSTER INDUSTRIAL ESTATE, REPTON ROAD, MEASHAM, SWADLINCOTE, DE12 7DT, UNITED KINGDOM
<b>5: Scope of authorisation and dosage forms</b>	ANNEX 1 and/ or ANNEX 2
<b>6: Legal Basis of authorisation</b>	Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)
<b>7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation</b>	Confidential
<b>8: Authorisation Date</b>	31/03/2025
<b>9: Annexes attached</b>	Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

**KENT PHARMACEUTICALS LIMITED**, UNIT 200, WESTMINSTER 42, WESTMINSTER INDUSTRIAL ESTATE, REPTON ROAD, MEASHAM, SWADLINCOTE, DE12 7DT, UNITED KINGDOM

Human Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1) IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)
<b>Part 1 - MANUFACTURING OPERATIONS</b> <b>[ 1.5 ] Packaging</b> [ 1.5.2 ] Secondary packaging <b>Part 2 - IMPORTATION OF MEDICINAL PRODUCTS</b> <b>[ 2.2 ] Batch certification of imported medicinal products</b> [ 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**

[ 2.3.1 ] Site of Physical Importation

[ 2.3.2 ] Importation of Intermediate which undergoes further processing

MHRA-GMDP

MHRA

MHRA-GMDP

MHRA