

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

**1: Authorisation Number**

UK MIA 52811

**2: Name of authorisation holder**

KINDEVA DRUG DELIVERY LIMITED

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

KINDEVA DRUG DELIVERY LIMITED, DERBY ROAD,  
LOUGHBOROUGH, LE11 5SF, UNITED KINGDOM

**4: Legally registered address of authorisation holder**

KINDEVA DRUG DELIVERY LIMITED, DERBY ROAD,  
LOUGHBOROUGH, LE11 5SF, UNITED KINGDOM

**5: Scope of authorisation and dosage forms**

ANNEX 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**

15/01/2026

**9: Annexes attached**

Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

**KINDEVA DRUG DELIVERY LIMITED, DERBY ROAD, LOUGHBOROUGH, LE11 5SF, UNITED KINGDOM**

Human Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)
<b>Part 1 - MANUFACTURING OPERATIONS</b>
<b>[ 1.2 ] Non-sterile products</b>
[ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms)
[ 1.2.1.9 ] Pressurised preparations
<b>[ 1.4 ] Other products or manufacturing activity</b>
[ 1.4.1 ] Manufacture of:
[ 1.4.1.3 ] Other
Micronisation of active pharmaceutical ingredient for use in the manufacture of finished products
<b>[ 1.5 ] Packaging</b>
[ 1.5.1 ] Primary packaging
[ 1.5.1.9 ] Pressurised preparations

[ 1.5.2 ] Secondary packaging

**[ 1.6 ] Quality control testing**

[ 1.6.2 ] Microbiological: non-sterility

[ 1.6.3 ] Chemical/Physical

MHRA-GMDP

MHRA

MHRA-GMDP

MHRA