

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA(IMP) 25076
<b>2: Name of authorisation holder</b>	MEDPHARM LIMITED
<b>3: Address(es) of manufacturing site(s)</b>	MEDPHARM LIMITED, UNIT 1 & UNIT 3 CHANCELLOR COURT, 50 OCCAM ROAD, SURREY RESEARCH PARK, GUILDFORD, GU2 7AB, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	MEDPHARM LIMITED, UNIT 1 & UNIT 3 CHANCELLOR COURT, 50 OCCAM ROAD, SURREY RESEARCH PARK, GUILDFORD, GU2 7AB, UNITED KINGDOM
<b>5: Scope of authorisation and dosage forms</b>	ANNEX 1 and/ or ANNEX 2
<b>6: Legal Basis of authorisation</b>	Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]
<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>	27/04/2026
<b>9: Annexes attached</b>	Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 2

Name and address of the site:

**MEDPHARM LIMITED**, UNIT 1 & UNIT 3 CHANCELLOR COURT, 50 OCCAM ROAD, SURREY RESEARCH PARK, GUILDFORD, GU2 7AB, UNITED KINGDOM

Human Investigational 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>[ 1.2 ] Non-sterile investigational medicinal products</b> [ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms) [ 1.2.1.4 ] Impregnated matrices [ 1.2.1.5 ] Liquids for external use [ 1.2.1.6 ] Liquids for internal use [ 1.2.1.8 ] Other solid dosage forms

[ 1.2.1.9 ] Pressurised preparations

[ 1.2.1.11 ] Semi-solids

[ 1.2.1.12 ] Suppositories

[ 1.2.1.14 ] Transdermal patches

[ 1.2.2 ] Batch certification

**[ 1.5 ] Packaging**

[ 1.5.1 ] Primary packaging

[ 1.5.1.4 ] Impregnated matrices

[ 1.5.1.5 ] Liquids for external use

[ 1.5.1.6 ] Liquids for internal use

[ 1.5.1.8 ] Other solid dosage forms

[ 1.5.1.9 ] Pressurised preparations

[ 1.5.1.11 ] Semi-solids

[ 1.5.1.12 ] Suppositories

[ 1.5.1.14 ] Transdermal patches

[ 1.5.2 ] Secondary packaging

**[ 1.6 ] Quality control testing**

[ 1.6.2 ] Microbiological: non-sterility

[ 1.6.3 ] Chemical/Physical

**Part 2 - IMPORTATION OF MEDICINAL PRODUCTS**

**[ 2.1 ] Quality control testing of imported medicinal products**

[ 2.1.2 ] Microbiological: non-sterility

[ 2.1.3 ] Chemical/Physical

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

[ 2.2.2 ] Non-sterile products