# 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 27776

HELIOS HOMOEOPATHY LIMITED

HELIOS HOMOEOPATHY LIMITED, 89-97 CAMDEN ROAD, TUNBRIDGE WELLS, TN1 2QR, UNITED KINGDOM

HELIOS HOMOEOPATHY LIMITED, 89-97 CAMDEN ROAD,

TUNBRIDGE WELLS, TN1 2QR, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations 2012 (SI

2012/1916)

Confidential

30/01/2024

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

#### HELIOS HOMOEOPATHY LIMITED, 89-97 CAMDEN ROAD, TUNBRIDGE WELLS, TN1 2QR, UNITED KINGDOM

**Human Medicinal Products** 

**Authorised Operations** 

MANUFACTURING OPERATIONS (according to part 1)

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

#### Part 1 - MANUFACTURING OPERATIONS

#### [ 1.2 ] Non-sterile products

[ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms)

[ 1.2.1.6 ] Liquids for internal use

[ 1.2.1.8 ] Other solid dosage forms

[ 1.2.1.13 ] Tablets

#### [ 1.4 ] Other products or manufacturing activity

[1.4.1] Manufacture of:

[1.4.1.2] Homoeopathic products

[ 1.5 ] Packaging

Issue Date: 30 Jan 2024

[1.5.1] Primary packaging

[ 1.5.1.17 ] Other non-sterile medicinal products Pillules

[1.5.2] Secondary packaging

# [ 1.6 ] Quality control testing

[ 1.6.3 ] Chemical/Physical

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

# [ 2.2 ] Batch certification of imported medicinal products

[2.2.2] Non-sterile products

