# 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 20912

**DRUGSRUS LIMITED** 

DRUGSRUS LIMITED, 5 SANDRIDGE CLOSE, HARROW, HA1 1XD, UNITED KINGDOM

DRUGSRUS LIMITED, 5 SANDRIDGE CLOSE,

HARROW, HA1 1XD, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations

2012 (SI 2012/1916)

Confidential

28/12/2023

Annex 1 and/or Annex 2

## SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

DRUGSRUS LIMITED, 5 SANDRIDGE CLOSE, HARROW, HA1 1XD, 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

## [ 1.5 ] Packaging

[1.5.2] Secondary packaging

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

[ 2.2 ] Batch certification of imported medicinal products

[2.2.1] Sterile Products

[2.2.1.1] Aseptically prepared

[2.2.1.2] Terminally sterilised

[2.2.2] Non-sterile products

Issue Date: 28 Dec 2023