

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA 20912
<b>2: Name of authorisation holder</b>	DRUGSRUS LIMITED
<b>3: Address(es) of manufacturing site(s)</b>	DRUGSRUS LIMITED, 5 SANDRIDGE CLOSE, HARROW, HA1 1XD, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	DRUGSRUS LIMITED, 5 SANDRIDGE CLOSE, HARROW, HA1 1XD, 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>	28/12/2023
<b>9: Annexes attached</b>	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)
<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