

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA 6831
<b>2: Name of authorisation holder</b>	GENUS PHARMACEUTICALS LIMITED
<b>3: Address(es) of manufacturing site(s)</b>	GENUS PHARMACEUTICALS LIMITED, MANCHESTER ROAD, LINTHWAITE, HUDDERSFIELD, HD7 5QH, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	GENUS PHARMACEUTICALS LIMITED, MANCHESTER ROAD, LINTHWAITE, HUDDERSFIELD, HD7 5QH, 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>	10/10/2024
<b>9: Annexes attached</b>	Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

**GENUS PHARMACEUTICALS LIMITED, MANCHESTER ROAD, LINTHWAITE, HUDDERSFIELD, HD7 5QH, 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>[ 1.2 ] Non-sterile products</b> [ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms) [ 1.2.1.11 ] Semi-solids [ 1.2.2 ] Batch certification <b>[ 1.4 ] Other products or manufacturing activity</b> [ 1.4.1 ] Manufacture of: [ 1.4.1.3 ] Other Medical Devices <b>[ 1.5 ] Packaging</b> [ 1.5.2 ] Secondary packaging

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

[ 2.2.2 ] Non-sterile products

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