

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA 16369
<b>2: Name of authorisation holder</b>	G-PHARMA LIMITED
<b>3: Address(es) of manufacturing site(s)</b>	G-PHARMA LIMITED, UNITS 1-4, DAKOTA AVENUE, DAKOTA PARK, SALFORD, M50 2PU, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	G-PHARMA LIMITED, RIVINGTON ROAD, WHITEHOUSE INDUSTRIAL ESTATE, PRESTON BROOK, RUNCORN, WA7 3DJ, 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>	08/11/2024
<b>9: Annexes attached</b>	Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

**G-PHARMA LIMITED**, UNITS 1-4, DAKOTA AVENUE, DAKOTA PARK, SALFORD, M50 2PU, UNITED KINGDOM

Human Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)
<b>Part 1 - MANUFACTURING OPERATIONS</b> <b>[ 1.5 ] Packaging</b> [ 1.5.2 ] Secondary packaging