

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA 20848
<b>2: Name of authorisation holder</b>	THORPE LABORATORIES LIMITED
<b>3: Address(es) of manufacturing site(s)</b>	THORPE LABORATORIES LIMITED, GOLF ROAD INDUSTRIAL ESTATE, MABLETHORPE, LN12 1NB, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	THORPE LABORATORIES LIMITED, GOLF ROAD INDUSTRIAL ESTATE, MABLETHORPE, LN12 1NB, 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>	13/03/2024
<b>9: Annexes attached</b>	Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

**THORPE LABORATORIES LIMITED**, GOLF ROAD INDUSTRIAL ESTATE, MABLETHORPE, LN12 1NB, UNITED KINGDOM

Human Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)
<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) [ 1.2.1.5 ] Liquids for external use [ 1.2.1.6 ] Liquids for internal use [ 1.2.1.11 ] Semi-solids <b>[ 1.4 ] Other products or manufacturing activity</b> [ 1.4.1 ] Manufacture of: [ 1.4.1.1 ] Herbal products <b>[ 1.5 ] Packaging</b> [ 1.5.1 ] Primary packaging

[ 1.5.1.2 ] Capsules, soft shell

[ 1.5.1.5 ] Liquids for external use

[ 1.5.1.6 ] Liquids for internal use

[ 1.5.1.8 ] Other solid dosage forms

[ 1.5.1.11 ] Semi-solids

[ 1.5.2 ] Secondary packaging

**[ 1.6 ] Quality control testing**

[ 1.6.3 ] Chemical/Physical