# 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 54373** 

PLUSULTRA PHARMA UK LTD

PLUSULTRA-PHARMA UK LTD, 20 WENLOCK ROAD, LONDON, N1 7GU, UNITED KINGDOM

PLUSULTRA PHARMA UK LTD, 149 WOODSTOCK ROAD, OXFORD, OX2 7NA, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)

Confidential

18/07/2025

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

# PLUSULTRA-PHARMA UK LTD, 20 WENLOCK ROAD, LONDON, N1 7GU, 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

## [ 1.1 ] Sterile Products

[ 1.1.3 ] Batch certification

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

[ 2.2 ] Batch certification of imported medicinal products

[2.2.2] Non-sterile products

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