

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA(IMP) 59749
<b>2: Name of authorisation holder</b>	4BASEBIO UK LTD
<b>3: Address(es) of manufacturing site(s)</b>	4BASEBIO UK LTD, UNIT 12 & UNIT 18, MERIDIAN, BUCKINGWAY BUSINESS PARK, ANDERSON ROAD, SWAVESEY, CAMBRIDGE, CB24 4AE, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	4BASEBIO UK LTD, UNIT 25, NORMAN WAY INDUSTRIAL ESTATE, NORMAN WAY, OVER, CAMBRIDGE, CB24 5QE, UNITED KINGDOM
<b>5: Scope of authorisation and dosage forms</b>	ANNEX 1 and/ or ANNEX 2
<b>6: Legal Basis of authorisation</b>	Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]
<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>	31/03/2025
<b>9: Annexes attached</b>	Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 2

Name and address of the site:

**4BASEBIO UK LTD**, UNIT 12 & UNIT 18, MERIDIAN, BUCKINGWAY BUSINESS PARK, ANDERSON ROAD, SWAVESEY, CAMBRIDGE, CB24 4AE, UNITED KINGDOM

Human Investigational Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)
<b>Part 1 - MANUFACTURING OPERATIONS</b> <b>[ 1.3 ] Biological investigational medicinal products</b> [ 1.3.1 ] Biological medicinal products [ 1.3.1.8 ] Other biological medicinal products ATMP Manufacture- a) Critical Starting Material- CDMO Manufacturing of Synthetic DNA template for IMP drug substance manufacturing b) Drug Substance- CDMO Manufacturing of Synthetic DNA for IMP drug product