# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

| | |
|---|---|
| **1: Authorisation Number** | UK MIA 50674 |
| **2: Name of authorisation holder** | BIOCON PHARMA UK LIMITED |
| **3: Address(es) of manufacturing site(s)** | BIOCON PHARMA UK LIMITED, GROUND FLOOR, CAVENDISH HOUSE, 369 BURNT OAK BROADWAY, EDGWARE, HA8 5AW, UNITED KINGDOM |
| **4: Legally registered address of authorisation holder** | BIOCON PHARMA UK LIMITED, GROUND FLOOR, CAVENDISH HOUSE, 369 BURNT OAK BROADWAY, EDGWARE, HA8 5AW, UNITED KINGDOM |
| **5: Scope of authorisation and dosage forms** | ANNEX 1 and/ or ANNEX 2 |
| **6: Legal Basis of authorisation** | Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916) |
| **7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation** | Confidential |
| **8: Authorisation Date** | 13/03/2025 |
| **9: Annexes attached** | Annex 1 and/or Annex 2 |

**SCOPE OF AUTHORISATION**

**Annex 1**

Name and address of the site:

**BIOCON PHARMA UK LIMITED**, GROUND FLOOR, CAVENDISH HOUSE, 369 BURNT OAK BROADWAY, EDGWARE, HA8 5AW, UNITED KINGDOM

| |
|---|
| Human Medicinal Products |
| Authorised Operations |
| IMPORTATION OF MEDICINAL PRODUCTS (according to part 2) |
| Part 2 - IMPORTATION OF MEDICINAL PRODUCTS<br>**[ 2.2 ] Batch certification of imported medicinal products**<br>[ 2.2.1 ] Sterile Products<br>[ 2.2.1.1 ] Aseptically prepared<br>[ 2.2.1.2 ] Terminally sterilised<br>[ 2.2.2 ] Non-sterile products |