# 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(IMP) 51381 ROYLANCE STABILITY STORAGE LIMITED

ROYLANCE STABILITY STORAGE LIMITED, BIOCITY SCOTLAND, BO'NESS ROAD, MOTHERWELL, ML1 5UH, UNITED KINGDOM

ROYLANCE STABILITY STORAGE LIMITED, BIOCITY SCOTLAND, BO'NESS ROAD, MOTHERWELL, ML1 5UH, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]

Confidential

31/01/2024

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

Annex 2 Name and address of the site:

ROYLANCE STABILITY STORAGE LIMITED, BIOCITY SCOTLAND, BO'NESS ROAD, MOTHERWELL, ML1 5UH, UNITED KINGDOM

Human Investigational Medicinal Products

Authorised Operations

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

#### Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

- [ 2.2 ] Batch certification of imported medicinal products
  - [2.2.1] Sterile Products
    - [2.2.1.1] Aseptically prepared
    - [2.2.1.2] Terminally sterilised

[2.2.2] Non-sterile products

[2.2.3] Biological medicinal products

[2.2.3.2] Immunological products

[2.2.3.3] Cell therapy products

- [ 2.2.3.4 ] Gene therapy products
- [ 2.2.3.5 ] Biotechnology products
- [2.2.3.6] Human or animal extracted products

### [ 2.3 ] Other Importation Activities

- [2.3.1] Site of Physical Importation
- [ 2.3.2 ] Importation of Intermediate which undergoes further processing

[ 2.3.4 ] Other

Importation of QP certified IMPs from a country on the approved country for import list