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) 18351 ALCURA UK LIMITED

ALCURA UK LIMITED, ALCURA HOUSE, CASWELL ROAD, BRACKMILLS INDUSTRIAL ESTATE, NORTHAMPTON, NN4 7PU, UNITED KINGDOM

ALCURA UK LIMITED, ALCURA HOUSE, CASWELL ROAD, BRACKMILLS INDUSTRIAL ESTATE, NORTHAMPTON, NN4 7PU, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

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

Confidential

12/05/2025

Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 2 Name and address of the site:

ALCURA UK LIMITED, ALCURA HOUSE, CASWELL ROAD, BRACKMILLS INDUSTRIAL ESTATE, NORTHAMPTON, NN4 7PU, UNITED KINGDOM

Human Investigational Medicinal Products

Authorised Operations

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

[2.3] Other Importation Activities

[2.3.1] Site of Physical Importation

[2.3.4] Other

Importation of QP certified IMPs from a country on the 'approved country for import list'/Returns and destruction of IMPs