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 18799

GOWRIE LAXMICO LIMITED

GOWRIE LAXMICO LIMITED, UNIT 4, BRADFIELD ROAD, RUISLIP, HA4 ONU, UNITED KINGDOM

GOWRIE LAXMICO LIMITED, UNIT 4, BRADFIELD ROAD, RUISLIP, HA4 0NU, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations

2012 (SI 2012/1916)

Confidential

31/03/2025

Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

GOWRIE LAXMICO LIMITED, UNIT 4, BRADFIELD ROAD, RUISLIP, HA4 0NU, UNITED KINGDOM

Human Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

[1.5] Packaging

[1.5.2] Secondary packaging

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