# 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 33155

RIVOPHARM (UK) LIMITED

RIVOPHARM (UK) LIMITED, 100 BISHOPSGATE,

LONDON, EC2N 4AG, UNITED KINGDOM

RIVOPHARM (UK) LIMITED, 100 BISHOPSGATE,

LONDON, EC2N 4AG, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations

2012 (SI 2012/1916)

Confidential

04/06/2025

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

RIVOPHARM (UK) LIMITED, 100 BISHOPSGATE, LONDON, EC2N 4AG, UNITED KINGDOM

**Human Medicinal Products** 

**Authorised Operations** 

MANUFACTURING OPERATIONS (according to part 1)

IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)

### Part 1 - MANUFACTURING OPERATIONS

#### [ 1.2 ] Non-sterile products

[ 1.2.2 ] Batch certification

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

[ 2.2 ] Batch certification of imported medicinal products

[2.2.2] Non-sterile products

Issue Date: 04 Jun 2025