# 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 25

ORGANON PHARMA (UK) LIMITED

ORGANON PHARMA (UK) LIMITED, SHOTTON LANE, CRAMLINGTON, NE23 3JU, UNITED KINGDOM

ORGANON PHARMA (UK) LIMITED, SHOTTON LANE, CRAMLINGTON, NE23 3JU, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)

Confidential

12/02/2025

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

### Annex 1

Name and address of the site:

## ORGANON PHARMA (UK) LIMITED, SHOTTON LANE, CRAMLINGTON, NE23 3JU, UNITED KINGDOM

**Human Medicinal Products** 

**Authorised Operations** 

MANUFACTURING OPERATIONS (according to part 1)

#### Part 1 - MANUFACTURING OPERATIONS

#### [ 1.2 ] Non-sterile products

[ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms)

[ 1.2.1.8 ] Other solid dosage forms

### Special Requirements

Bulk Granules and bulk blend for oral solution.

[ 1.2.1.13 ] Tablets

### [ 1.5 ] Packaging

[1.5.1] Primary packaging

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[ 1.5.1.8 ] Other solid dosage forms

### **Special Requirements**

Bulk Granules and bulk blend for oral solution.

[ 1.5.1.13 ] Tablets

[ 1.5.2 ] Secondary packaging

# [ 1.6 ] Quality control testing

[ 1.6.2 ] Microbiological: non-sterility

[ 1.6.3 ] Chemical/Physical

