# 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) 735 BOC LIMITED

BOC LIMITED, 28 DEER PARK ROAD, LONDON, SW19 3UF, UNITED KINGDOM

BOC LIMITED, FORGE, 43 CHURCH STREET WEST, WOKING, GU21 6HT, UNITED KINGDOM

BOC LIMITED, FORGE, 43 CHURCH STREET WEST, WOKING, GU21 6HT, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Part 6 of The Medicines for Human Use (Clinical Trials)

Regulations 2004 [SI 2004/1031]

Confidential

14/11/2022

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

#### Annex 2

Name and address of the site:

BOC LIMITED, 28 DEER PARK ROAD, LONDON, SW19 3UF, UNITED KINGDOM

**Human Investigational Medicinal Products** 

**Authorised Operations** 

MANUFACTURING OPERATIONS (according to part 1)

#### Part 1 - MANUFACTURING OPERATIONS

### [ 1.2 ] Non-sterile investigational medicinal products

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

[1.2.1.7] Medicinal gases

[ 1.2.2 ] Batch certification

### [ 1.5 ] Packaging

[1.5.1] Primary packaging

[1.5.1.7] Medicinal gases

[ 1.5.2 ] Secondary packaging

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# [ 1.6 ] Quality control testing

[1.6.3] Chemical/Physical

# SCOPE OF AUTHORISATION

#### Annex 2

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