# 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 46113** 

**AUTOLUS LIMITED** 

AUTOLUS LIMITED THE NUCLEUS, THE NUCLEUS,

MARSHGATE, STEVENAGE, SG1 1FR, UNITED KINGDOM

AUTOLUS LIMITED, THE MEDIAWORKS, 191 WOOD LANE,

LONDON, W12 7FP, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Regulation 17 of The Human Medicines Regulations 2012 (SI

2012/1916)

Confidential

22/03/2024

Annex 1 and/or Annex 2

## SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

AUTOLUS LIMITED THE NUCLEUS, THE NUCLEUS, MARSHGATE, STEVENAGE, SG1 1FR, UNITED KINGDOM

**Human Medicinal Products** 

**Authorised Operations** 

MANUFACTURING OPERATIONS (according to part 1)

### Part 1 - MANUFACTURING OPERATIONS

## [ 1.1 ] Sterile Products

[ 1.1.1 ] Aseptically prepared (processing operations for the following dosage forms)

[1.1.1.1] Large volume liquids

**Special Requirements** 

Live Cells

[1.1.1.4] Small volume liquids

**Special Requirements** 

Live Cells

Issue Date: 22 Mar 2024

[ 1.1.1.6 ] Other aseptically prepared products Gene therapy **Special Requirements** Live Cells [ 1.1.3 ] Batch certification [ 1.3 ] Biological medicinal products [ 1.3.1 ] Biological medicinal products [1.3.1.4] Gene therapy products **Special Requirements** Live Cells [ 1.3.2 ] Batch certification [1.3.2.4] Gene therapy products **Special Requirements** Live Cells [ 1.5 ] Packaging [ 1.5.2 ] Secondary packaging [ 1.6 ] Quality control testing [ 1.6.1 ] Microbiological: sterility [ 1.6.2 ] Microbiological: non-sterility [ 1.6.4 ] Biological

