Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number	UK MIA 48152				
2: Name of authorisation holder	CELL AND GENE THERAPY CATAPULT				
3: Address(es) of manufacturing site(s)	CELL AND GENE THERAPY CATAPULT MANUFACTURING CENTRE, GUNNELS WOOD ROAD, STEVENAGE, SG1 2FX, UNITED KINGDOM				
4: Legally registered address of authorisation holder	CELL AND GENE THERAPY CATAPULT, 12TH FLOOR TOWER WING, GUY'S HOSPITAL, GREAT MAZE POND, LONDON, SE1 9RT, UNITED KINGDOM				
5: Scope of authorisation and dosage forms	ANNEX 1 and/ or ANNEX 2				
6: Legal Basis of authorisation	Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)				
7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation	Confidential				
8: Authorisation Date	24/05/2019				
9: Annexes attached	Annex 1 and/or Annex 2				

SCOPE OF AUTHORISATION

Annex 1 Name and address of the site:

CELL AND GENE THERAPY CATAPULT MANUFACTURING CEN	TRE,	GUNNEL	s wo	OD ROAD,	STEVENAGE,	SG1 2FX,	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

Advanced Therapy Medicinal Products

[1.1.1.4] Small volume liquids
Special Requirements
Live Cells
Advanced Therapy Medicinal Products
[1.1.1.6] Other aseptically prepared products
Cell based therapies and viral vectors
Special Requirements
Live Cells
Advanced Therapy Medicinal Products
[1.1.3] Batch certification
[1.3] Biological medicinal products
[1.3.1] Biological medicinal products
[1.3.1.3] Cell therapy products
Special Requirements
Live Cells
Advanced Therapy Medicinal Products
[1.3.1.4] Gene therapy products
Special Requirements
Live Cells
Advanced Therapy Medicinal Products
[1.3.1.5] Biotechnology products
Special Requirements
Live Cells
Advanced Therapy Medicinal Products
[1.3.1.6] Human or animal extracted products
Special Requirements
Live Cells
Advanced Therapy Medicinal Products
[1.3.1.8] Other biological medicinal products
Viral vectors
Special Requirements
Live Cells
Advanced Therapy Medicinal Products
[1.3.2] Batch certification
[1.3.2.3] Cell therapy products
Special Requirements
Live Cells
Advanced Therapy Medicinal Products
[1.3.2.4] Gene therapy products
Special Requirements
Live Cells
Advanced Therapy Medicinal Products
[1.3.2.5] Biotechnology products
Special Requirements
Advanced Therapy Medicinal Products

	[1.3.2.6] Human or animal extracted products	
	Special Requirements	
	Live Cells	
	Advanced Therapy Medicinal Products	
	[1.3.2.8] Other biological medicinal products	
	Viral vectors	
	Special Requirements	
	Live Cells	
	Advanced Therapy Medicinal Products	
	[1.4] Other products or manufacturing activity	
	[1.4.2] Sterilisation of active substances/excipients/finished products:	
	[1.4.2.1] Filtration	
	[1.5] Packaging	
	[1.5.1] Primary packaging	
	[1.5.1.6] Liquids for internal use	
	[1.5.2] Secondary packaging	
	[1.6] Quality control testing	
	[1.6.1] Microbiological: sterility	
	[1.6.2] Microbiological: non-sterility	
	[1.6.3] Chemical/Physical	
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[1.6.4] Biological