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

SCOPE OF AUTHORISATION

Annex 2 Name and address of the site:

EUROAPI UK LIMITED, 37 HOLLANDS ROAD, HAVERHILL, CB9 8PU, 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)	
[1.2.1.15] Other non-sterile medicinal products	
Partial manufacture only: formulation & spray-drying only	
[1.6] Quality control testing	
[1.6.2] Microbiological: non-sterility	
[1.6.3] Chemical/Physical	

UK MIA(IMP) 8596 EUROAPI UK LIMITED

EUROAPI UK LIMITED, 37 HOLLANDS ROAD, HAVERHILL, CB9 8PU, UNITED KINGDOM

EUROAPI UK LIMITED, 37 HOLLANDS ROAD, HAVERHILL, CB9 8PU, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]

Confidential

16/12/2024

Annex 1 and/or Annex 2