## 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

UK MIA 15764 STRANDHAVEN LIMITED

STRANDHAVEN LIMITED, 641 HIGH ROAD, ILFORD, IG3 8RA, UNITED KINGDOM

STRANDHAVEN LIMITED, 600 HIGH ROAD, SEVEN KINGS, ILFORD, IG3 8BS, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

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

Confidential

14/03/2024 Annex 1 and/or Annex 2

Annex 1 Name and address of the site:

## STRANDHAVEN LIMITED, 641 HIGH ROAD, ILFORD, IG3 8RA, UNITED KINGDOM

Human Medicinal Products	
Authorised Operations	
IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)	
Part 2 - IMPORTATION OF MEDICINAL PRODUCTS	
[ 2.2 ] Batch certification of imported medicinal products	
[ 2.2.1 ] Sterile Products	
[ 2.2.1.1 ] Aseptically prepared	0
[ 2.2.1.2 ] Terminally sterilised	
[ 2.2.2 ] Non-sterile products	
	NHM