## 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(IMP) 8829 CHIESI LIMITED

CHIESI LIMITED, 333 STYAL ROAD, MANCHESTER, M22 5LG, UNITED KINGDOM

CHIESI LIMITED, 333 STYAL ROAD, MANCHESTER, M22 5LG, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

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

Confidential

09/05/2024 Annex 1 and/or Annex 2

Annex 2 Name and address of the site:

CHIESI LIMITED, 333 STYAL ROAD, MANCHESTER, M22 5LG, UNITED KINGDOM

Human Investigational Medicinal Products	
Authorised Operations	
IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)	
Part 2 - IMPORTATION OF MEDICINAL PRODUCTS	
[ 2.3 ] Other Importation Activities	
[ 2.3.4 ] Other	
Importation of QP certified IMPs from a country on the approved country for import list	
	NHR