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 1 Name and address of the site:

SMIRO QUALITAS LIMITED, 4 FLATFORD ROAD, HAVERHILL, CB9 7WW, 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	
[2.2.1.2] Terminally sterilised	
[2.2.2] Non-sterile products	
[2.2.3] Biological medicinal products	
[2.2.3.2] Immunological products	
[2.3] Other Importation Activities	

UK MIA 44646 SMIRO QUALITAS LIMITED

SMIRO QUALITAS LIMITED, 4 FLATFORD ROAD, HAVERHILL, CB9 7WW, UNITED KINGDOM

SMIRO QUALITAS LIMITED, 4 FLATFORD ROAD, HAVERHILL, CB9 7WW, UNITED KINGDOM ANNEX 1 and/ or ANNEX 2

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

Confidential

16/12/2024 Annex 1 and/or Annex 2 [2.3.4] Products authorised under regulation 174 (supply in response to spread of pathogenic agents etc) GMP Certification and Batch release of imported biological imported Product- Monoclonal antibodies, Vaccines, oral vaccines, recombinant proteins, MVAs and adjuvants