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

UK MIA 55644 RED KNIGHTS PHARMA LIMITED

RED KNIGHTS PHARMA LIMITED, 5300 LAKESIDE, CHEADLE ROYAL BUSINESS PARK, CHEADLE, SK8 3GP, UNITED KINGDOM

RED KNIGHTS PHARMA LIMITED, 5300 LAKESIDE, CHEADLE ROYAL BUSINESS PARK, CHEADLE, SK8 3GP, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

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

Confidential

09/09/2024

Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1 Name and address of the site:

RED KNIGHTS PHARMA LIMITED, 5300 LAKESIDE, CHEADLE ROYAL BUSINESS PARK, CHEADLE, SK8 3GP, UNITED KINGDOM

Human Medicinal Products	
Authorised Operations	
MANUFACTURING OPERATIONS (according to part 1)	
IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)	
Part 1 - MANUFACTURING OPERATIONS	
[1.2] Non-sterile products	
[1.2.2] Batch certification	
Part 2 - IMPORTATION OF MEDICINAL PRODUCTS	
[2.2] Batch certification of imported medicinal products	
[2.2.2] Non-sterile products	