# 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(IMP) 27436

PROPHARMA GROUP MIS LIMITED

PROPHARMA GROUP MIS LIMITED, OLLIVER, ASKE, RICHMOND, DL10 5HX, UNITED KINGDOM

PROPHARMA GROUP MIS LIMITED, OLLIVER, ASKE, RICHMOND, DL10 5HX, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

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

Confidential

25/09/2024

Annex 1 and/or Annex 2

#### SCOPE OF AUTHORISATION

## Annex 2

Name and address of the site:

## PROPHARMA GROUP MIS LIMITED, OLLIVER, ASKE, RICHMOND, DL10 5HX, 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

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