## 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 45396

UNI HEALTH DISTRIBUTION LIMITED

PHARMPACT LIMITED, UNIT G2, G3 & G4, RIVERSIDE INDUSTRIAL ESTATE, RIVERSIDE WAY, DARTFORD, DA1 5BS, UNITED KINGDOM

UNI HEALTH DISTRIBUTION LIMITED, UNIT G4, RIVERSIDE INDUSTRIAL ESTATE, RIVERSIDE WAY, DARTFORD, DA1 5BS, UNITED KINGDOM

ANNEX 1 and/ or ANNEX 2

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

Confidential

07/05/2021

Annex 1 and/or Annex 2

## SCOPE OF AUTHORISATION

Annex 1 Name and address of the site:

PHARMPACT LIMITED, UNIT G2, G3 & G4, RIVERSIDE INDUSTRIAL ESTATE, RIVERSIDE WAY, DARTFORD, DA1 5BS, UNITED KINGDOM

Human Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

## Part 1 - MANUFACTURING OPERATIONS

[ 1.5 ] Packaging

[ 1.5.2 ] Secondary packaging