Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number	UK MIA 13163
2: Name of authorisation holder	PACKPHARM LIMITED
3: Address(es) of manufacturing site(s)	PACKPHARM LIMITED, UNIT 1, 39 MAHONEY GREEN, RACKHEATH, NORWICH, NR13 6JY, UNITED KINGDOM
4: Legally registered address of authorisation holder	PACKPHARM LIMITED, UNIT 1, 39 MAHONEY GREEN, RACKHEATH, NORWICH, NR13 6JY, UNITED KINGDOM
5: Scope of authorisation and dosage forms	ANNEX 1 and/ or ANNEX 2
6: Legal Basis of authorisation	Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)
7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation	Confidential
8: Authorisation Date	18/10/2023
9: Annexes attached	Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

PACKPHARM LIMITED, UNIT 1, 39 MAHONEY GREEN, RACKHEATH, NORWICH, NR13 6JY, 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

[1.5] Packaging

[1.5.1] Primary packaging

[1.5.1.1] Capsules, hard shell

[1.5.1.2] Capsules, soft shell

[1.5.1.8] Other solid dosage forms

[1.5.1.13] Tablets

[1.5.1.14] Transdermal patches

[1.5.2] Secondary packaging

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

MHRA-GMDP MHRA

[2.2] Batch certification of imported medicinal products

[2.2.2] Non-sterile products

MHRA-GMDP

MHRA-GMDP