Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number	UK MIA 16363
2: Name of authorisation holder	MILPHARM LIMITED
3: Address(es) of manufacturing site(s)	MILPHARM LIMITED, 1 ROUNDWOOD AVENUE, STOCKLEY PARK, UXBRIDGE, UB11 1AF, UNITED KINGDOM MILPHARM LIMITED, ARES BLOCK, ODYSSEY BUSINESS PARK, WEST END ROAD, RUISLIP, HA4 6QD, UNITED KINGDOM
4: Legally registered address of authorisation holder	MILPHARM LIMITED, 1 ROUNDWOOD AVENUE, STOCKLEY PARK, UXBRIDGE, UB11 1AF, 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	30/01/2026
9: Annexes attached	Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

MILPHARM LIMITED, 1 ROUNDWOOD AVENUE, STOCKLEY PARK, UXBRIDGE, UB11 1AF, 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

MHRA-GMDP MHRA

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