Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number	UK MIA 48428
2: Name of authorisation holder	TECHDOW PHARMA ENGLAND LIMITED
3: Address(es) of manufacturing site(s)	TECHDOW PHARMA ENGLAND LIMITED, SURREY TECHNOLOGY CENTRE, 40 OCCAM ROAD, SURREY RESEARCH PARK, GUILDFORD, GU2 7YG, UNITED KINGDOM
4: Legally registered address of authorisation holder	TECHDOW PHARMA ENGLAND LIMITED, SURREY TECHNOLOGY CENTRE, 40 OCCAM ROAD, SURREY RESEARCH PARK, GUILDFORD, GU2 7YG, 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	19/08/2026
9: Annexes attached	Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

TECHDOW PHARMA ENGLAND LIMITED, SURREY TECHNOLOGY CENTRE, 40 OCCAM ROAD, SURREY RESEARCH PARK, GUILDFORD, GU2 7YG, 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

[2.2.3] Biological medicinal products

[2.2.3.6] Human or animal extracted products