

Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number	UK MIA 19819
2: Name of authorisation holder	BIOTEC SERVICES INTERNATIONAL LIMITED
3: Address(es) of manufacturing site(s)	BIOTEC SERVICES INTERNATIONAL LIMITED, BIOTEC HOUSE, CENTRAL PARK, WESTERN AVENUE, BRIDGEND INDUSTRIAL ESTATE, BRIDGEND, CF31 3RT, UNITED KINGDOM
4: Legally registered address of authorisation holder	BIOTEC SERVICES INTERNATIONAL LIMITED, UNITS 2100, 2110, 2120, 2130, 2010, 2430 AND 2500, PHASE 18, CENTRAL PARK, BRIDGEND INDUSTRIAL ESTATE, BRIDGEND, CF31 3TY, UNITED KINGDOM
5: Scope of authorisation and dosage forms	BIOTEC SERVICES INTERNATIONAL LIMITED, BIOTEC HOUSE, CENTRAL PARK, WESTERN AVENUE, BRIDGEND INDUSTRIAL ESTATE, BRIDGEND, CF31 3RT, UNITED KINGDOM
6: Legal Basis of authorisation	ANNEX 1 and/ or ANNEX 2
7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation	Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)
8: Authorisation Date	Confidential
9: Annexes attached	18/06/2025
	Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

BIOTEC SERVICES INTERNATIONAL LIMITED, BIOTEC HOUSE, CENTRAL PARK, WESTERN AVENUE, BRIDGEND INDUSTRIAL ESTATE, BRIDGEND, CF31 3RT, 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 [1.1] Sterile Products [1.1.3] Batch certification [1.2] Non-sterile products

- [1.2.2] Batch certification
- [1.3] Biological medicinal products**
 - [1.3.2] Batch certification
 - [1.3.2.1] Blood products
 - [1.3.2.2] Immunological products
 - [1.3.2.3] Cell therapy products
 - [1.3.2.4] Gene therapy products
 - [1.3.2.5] Biotechnology products
 - [1.3.2.6] Human or animal extracted products
 - [1.3.2.7] Tissue Engineered Products

Special Requirements

- Tissue engineered product
- [1.3.2.8] Other biological medicinal products
 - Tissue engineered products

[1.5] Packaging

- [1.5.2] Secondary packaging

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
- [2.2.3] Biological medicinal products
 - [2.2.3.1] Blood products
 - [2.2.3.2] Immunological products
 - [2.2.3.3] Cell therapy products
 - [2.2.3.4] Gene therapy products
 - [2.2.3.5] Biotechnology products
 - [2.2.3.6] Human or animal extracted products
 - [2.2.3.7] Tissue Engineered Products

[2.3] Other Importation Activities

- [2.3.1] Site of Physical Importation

SCOPE OF AUTHORISATION

Annex 1

Name and address of the site:

BIOTEC SERVICES INTERNATIONAL LIMITED, UNITS 2100, 2110, 2120, 2130, 2010, 2430 AND 2500, PHASE 18, CENTRAL PARK, BRIDGEND INDUSTRIAL ESTATE, BRIDGEND, CF31 3TY, UNITED KINGDOM

Human Medicinal Products
Authorised Operations

Part 1 - MANUFACTURING OPERATIONS

[1.1] Sterile Products

[1.1.3] Batch certification

[1.2] Non-sterile products

[1.2.2] Batch certification

[1.3] Biological medicinal products

[1.3.1] Biological medicinal products

[1.3.1.1] Blood products

[1.3.1.2] Immunological products

[1.3.1.3] Cell therapy products

[1.3.1.4] Gene therapy products

[1.3.1.5] Biotechnology products

[1.3.1.6] Human or animal extracted products

[1.3.2] Batch certification

[1.3.2.1] Blood products

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[1.3.2.7] Tissue Engineered Products

[1.5] Packaging

[1.5.2] Secondary packaging

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

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[2.2.3.5] Biotechnology products

[2.2.3.6] Human or animal extracted products

[2.2.3.7] Tissue Engineered Products

[2.2.3.8] Other biological medicinal products

Tissue engineered products

[2.3] Other Importation Activities

[2.3.1] Site of Physical Importation