

Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number	UK MIA(IMP) 5866
2: Name of authorisation holder	GLAXOSMITHKLINE RESEARCH & DEVELOPMENT LIMITED GLAXOSMITHKLINE RESEARCH & DEVELOPMENT, HARRIS'S LANE, WARE, SG12 0GX, UNITED KINGDOM
3: Address(es) of manufacturing site(s)	GLAXOSMITHKLINE RESEARCH & DEVELOPMENT LIMITED, ADDENBROOKE'S HOSPITAL BOX 128, CLINICAL RESEARCH UNIT, HILLS ROAD, CAMBRIDGE, CB2 0GG, UNITED KINGDOM GLAXOSMITHKLINE RESEARCH & DEVELOPMENT LIMITED, THIRD AVENUE, HARLOW, CM19 5AW, UNITED KINGDOM
4: Legally registered address of authorisation holder	GLAXOSMITHKLINE RESEARCH & DEVELOPMENT LIMITED, GSK HOUSE, 980 GREAT WEST ROAD, BRENTFORD, TW8 9GS, UNITED KINGDOM
5: Scope of authorisation and dosage forms	ANNEX 1 and/ or ANNEX 2
6: Legal Basis of authorisation	Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]
7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation	Confidential
8: Authorisation Date	10/10/2024
9: Annexes attached	Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 2

Name and address of the site:

GLAXOSMITHKLINE RESEARCH & DEVELOPMENT, HARRIS'S LANE, WARE, SG12 0GX, UNITED KINGDOM

Human Investigational Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)
Part 1 - MANUFACTURING OPERATIONS [1.2] Non-sterile investigational medicinal products [1.2.1] Non-Sterile Products (processing operations for the following dosage forms) [1.2.1.1] Capsules, hard shell [1.2.1.8] Other solid dosage forms

[1.2.1.13] Tablets

[1.2.1.15] Other non-sterile medicinal products

Solid non-sterile multi dose forms (including powders and granules), powder inhalation products and devices.

[1.2.2] Batch certification

[1.6] Quality control testing

[1.6.2] Microbiological: non-sterility

[1.6.3] Chemical/Physical

SCOPE OF AUTHORISATION

Annex 2

Name and address of the site:

GLAXOSMITHKLINE RESEARCH & DEVELOPMENT LIMITED, ADDENBROOKE'S HOSPITAL BOX 128, CLINICAL RESEARCH UNIT, HILLS ROAD, CAMBRIDGE, CB2 0GG, UNITED KINGDOM

Human Investigational Medicinal Products

Authorised Operations

MANUFACTURING OPERATIONS (according to part 1)

Part 1 - MANUFACTURING OPERATIONS

[1.2] Non-sterile investigational medicinal products

[1.2.1] Non-Sterile Products (processing operations for the following dosage forms)

[1.2.1.1] Capsules, hard shell

[1.2.1.5] Liquids for external use

[1.2.1.6] Liquids for internal use

[1.2.1.11] Semi-solids

[1.2.1.15] Other non-sterile medicinal products

Powder in bottle

[1.5] Packaging

[1.5.1] Primary packaging

[1.5.1.1] Capsules, hard shell

[1.5.1.2] Capsules, soft shell

[1.5.1.5] Liquids for external use

[1.5.1.6] Liquids for internal use

[1.5.1.11] Semi-solids

[1.5.1.13] Tablets

[1.5.2] Secondary packaging

SCOPE OF AUTHORISATION

Annex 2

Name and address of the site:

Human Investigational Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1) IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)
<p>Part 1 - MANUFACTURING OPERATIONS</p> <p>[1.1] Sterile Investigational Medicinal Products</p> <p>[1.1.3] Batch certification</p> <p>[1.2] Non-sterile investigational medicinal products</p> <p>[1.2.2] Batch certification</p> <p>[1.3] Biological investigational medicinal products</p> <p>[1.3.2] Batch certification</p> <p>[1.3.2.5] Biotechnology products</p> <p>[1.5] Packaging</p> <p>[1.5.1] Primary packaging</p> <p>[1.5.1.1] Capsules, hard shell</p> <p>[1.5.1.13] Tablets</p> <p>[1.5.2] Secondary packaging</p> <p>Part 2 - IMPORTATION OF MEDICINAL PRODUCTS</p> <p>[2.2] Batch certification of imported medicinal products</p> <p>[2.2.1] Sterile Products</p> <p>[2.2.1.1] Aseptically prepared</p> <p>[2.2.1.2] Terminally sterilised</p> <p>[2.2.2] Non-sterile products</p> <p>[2.2.3] Biological medicinal products</p> <p>[2.2.3.5] Biotechnology products</p> <p>[2.2.3.8] Other biological medicinal products</p> <p>Synthetic Proteins</p> <p>[2.3] Other Importation Activities</p> <p>[2.3.1] Site of Physical Importation</p> <p>[2.3.2] Importation of Intermediate which undergoes further processing</p> <p>[2.3.4] Other</p> <p>Authorised for Importation of QP certified IMPs from a country on the approved country for import list.</p>