

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

|                                                                                                                        |                                                                                                                                                                                                                                                                                                              |
|------------------------------------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| 1: Authorisation Number                                                                                                | UK MIA(IMP) 1384                                                                                                                                                                                                                                                                                             |
| 2: Name of authorisation holder                                                                                        | UNIVERSITY OF EDINBURGH<br><br>UNIVERSITY OF EDINBURGH - INVESTIGATIONAL SUPPLIES GROUP, 1<br>GEORGE SQUARE, EDINBURGH, EH8 9JZ, UNITED KINGDOM<br><br>UNIVERSITY OF EDINBURGH - PET RADIOCHEMISTRY, QUEENS<br>MEDICAL RESEARCH INSTITUTE, 47 LITTLE FRANCE CRESCENT,<br>EDINBURGH, EH16 4TJ, UNITED KINGDOM |
| 3: Address(es) of manufacturing site(s)                                                                                | STERILE FILL UNIT - HEALTHCARE TECHNOLOGY ACCELERATOR<br>FACILITY, LEVEL 4, INSTITUTE FOR REGENERATION AND REPAIR, 4-5<br>LITTLE FRANCE DRIVE, EDINBURGH, EH16 4UU, UNITED KINGDOM                                                                                                                           |
| 4: Legally registered address of authorisation holder                                                                  | UNIVERSITY OF EDINBURGH, OLD COLLEGE, SOUTH BRIDGE,<br>EDINBURGH, EH8 9YL, 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<br>2004/1031]                                                                                                                                                                                                                   |
| 7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation | Confidential                                                                                                                                                                                                                                                                                                 |
| 8: Authorisation Date                                                                                                  | 25/11/2025                                                                                                                                                                                                                                                                                                   |
| 9: Annexes attached                                                                                                    | Annex 1 and/or Annex 2                                                                                                                                                                                                                                                                                       |

### SCOPE OF AUTHORISATION

#### Annex 2

Name and address of the site:

**UNIVERSITY OF EDINBURGH - INVESTIGATIONAL SUPPLIES GROUP, 1 GEORGE SQUARE, EDINBURGH, EH8 9JZ, UNITED KINGDOM**

|                                                                                                           |
|-----------------------------------------------------------------------------------------------------------|
| Human Investigational Medicinal Products                                                                  |
| Authorised Operations                                                                                     |
| MANUFACTURING OPERATIONS (according to part 1)<br>IMPORTATION OF MEDICINAL PRODUCTS (according to part 2) |
| <b>Part 1 - MANUFACTURING OPERATIONS</b>                                                                  |

**[ 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

**[ 1.5 ] Packaging**

[ 1.5.1 ] Primary packaging

[ 1.5.1.2 ] Capsules, soft shell

[ 1.5.1.12 ] Suppositories

[ 1.5.1.13 ] Tablets

[ 1.5.1.15 ] Other non-sterile medicinal products

Powders

[ 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

[ 2.2.3.4 ] Gene therapy products

[ 2.2.3.5 ] Biotechnology products

[ 2.2.3.6 ] Human or animal extracted products

**SCOPE OF AUTHORISATION**

**Annex 2**

Name and address of the site:

**UNIVERSITY OF EDINBURGH - PET RADIOCHEMISTRY**, QUEENS MEDICAL RESEARCH INSTITUTE, 47 LITTLE FRANCE CRESCENT, EDINBURGH, EH16 4TJ, UNITED KINGDOM

|                                                                                                           |
|-----------------------------------------------------------------------------------------------------------|
| Human Investigational Medicinal Products                                                                  |
| Authorised Operations                                                                                     |
| MANUFACTURING OPERATIONS (according to part 1)<br>IMPORTATION OF MEDICINAL PRODUCTS (according to part 2) |
| <b>Part 1 - MANUFACTURING OPERATIONS</b>                                                                  |
| <b>[ 1.1 ] Sterile Investigational Medicinal Products</b>                                                 |
| [ 1.1.1 ] Aseptically prepared (processing operations for the following dosage forms)                     |

[ 1.1.1.4 ] Small volume liquids

**Special Requirements**

Radiopharmaceuticals

[ 1.1.2 ] Terminally Sterilised (processing operations for the following dosage forms)

[ 1.1.2.3 ] Small volume liquids

[ 1.1.2.5 ] Other terminally sterilised prepared products

Radiopharmaceuticals

**[ 1.4 ] Other investigational medicinal products or manufacturing activity**

[ 1.4.2 ] Sterilisation of active substances/excipients/finished products:

[ 1.4.2.1 ] Filtration

[ 1.4.2.3 ] Moist heat

**[ 1.6 ] Quality control testing**

[ 1.6.2 ] Microbiological: non-sterility

[ 1.6.3 ] Chemical/Physical

**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

**SCOPE OF AUTHORISATION**

**Annex 2**

Name and address of the site:

**STERILE FILL UNIT - HEALTHCARE TECHNOLOGY ACCELERATOR FACILITY**, LEVEL 4, INSTITUTE FOR REGENERATION AND REPAIR, 4-5 LITTLE FRANCE DRIVE, EDINBURGH, EH16 4UU, UNITED KINGDOM

|                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           |
|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Human Investigational Medicinal Products                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |
| Authorised Operations                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     |
| MANUFACTURING OPERATIONS (according to part 1)<br>IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)                                                                                                                                                                                                                                                                                                                                                                                                                                 |
| <b>Part 1 - MANUFACTURING OPERATIONS</b><br><b>[ 1.1 ] Sterile Investigational Medicinal Products</b><br>[ 1.1.1 ] Aseptically prepared (processing operations for the following dosage forms)<br>[ 1.1.1.1 ] Large volume liquids<br>[ 1.1.1.4 ] Small volume liquids<br>[ 1.1.3 ] Batch certification<br><b>[ 1.2 ] Non-sterile investigational medicinal products</b><br>[ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms)<br>[ 1.2.1.1 ] Capsules, hard shell<br>[ 1.2.1.2 ] Capsules, soft shell |

- [ 1.2.1.5 ] Liquids for external use
- [ 1.2.1.6 ] Liquids for internal use
- [ 1.2.1.11 ] Semi-solids
- [ 1.2.1.12 ] Suppositories
- [ 1.2.1.13 ] Tablets

[ 1.2.2 ] Batch certification

### **[ 1.3 ] Biological investigational medicinal products**

- [ 1.3.1 ] Biological medicinal products
  - [ 1.3.1.2 ] Immunological products
- [ 1.3.2 ] Batch certification
  - [ 1.3.2.2 ] Immunological products

### **[ 1.4 ] Other investigational medicinal products or manufacturing activity**

- [ 1.4.2 ] Sterilisation of active substances/excipients/finished products:
  - [ 1.4.2.1 ] Filtration

### **[ 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
  - [ 1.5.1.8 ] Other solid dosage forms
  - [ 1.5.1.11 ] Semi-solids
  - [ 1.5.1.12 ] Suppositories
  - [ 1.5.1.13 ] Tablets
- [ 1.5.2 ] Secondary packaging

### **[ 1.6 ] Quality control testing**

- [ 1.6.3 ] Chemical/Physical

## **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
- [ 2.3.2 ] Importation of Intermediate which undergoes further processing
- [ 2.3.3 ] Biological Active Substance