

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA(IMP) 42551
<b>2: Name of authorisation holder</b>	COPEA PHARMA LIMITED
<b>3: Address(es) of manufacturing site(s)</b>	COPEA PHARMA LIMITED, 25 COMPASS WEST, SPINDUS ROAD, LIVERPOOL, L24 1YA, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	COPEA PHARMA LIMITED, 25 COMPASS WEST, SPINDUS ROAD, LIVERPOOL, L24 1YA, UNITED KINGDOM
<b>5: Scope of authorisation and dosage forms</b>	ANNEX 1 and/ or ANNEX 2
<b>6: Legal Basis of authorisation</b>	Part 6 of The Medicines for Human Use (Clinical Trials) Regulations 2004 [SI 2004/1031]
<b>7: Name of responsible officer of the competent authority of the member state granting the manufacturing authorisation</b>	Confidential
<b>8: Authorisation Date</b>	29/07/2022
<b>9: Annexes attached</b>	Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 2

Name and address of the site:

**COPEA PHARMA LIMITED**, 25 COMPASS WEST, SPINDUS ROAD, LIVERPOOL, L24 1YA, UNITED KINGDOM

Human Investigational Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1) IMPORTATION OF MEDICINAL PRODUCTS (according to part 2)
<b>Part 1 - MANUFACTURING OPERATIONS</b> <b>[ 1.1 ] Sterile Investigational Medicinal Products</b> [ 1.1.3 ] Batch certification <b>[ 1.2 ] Non-sterile investigational medicinal products</b> [ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms) [ 1.2.1.5 ] Liquids for external use [ 1.2.1.6 ] Liquids for internal use [ 1.2.1.8 ] Other solid dosage forms [ 1.2.1.11 ] Semi-solids [ 1.2.2 ] Batch certification

**[ 1.3 ] Biological investigational medicinal products**

- [ 1.3.2 ] Batch certification
  - [ 1.3.2.2 ] Immunological products
  - [ 1.3.2.5 ] Biotechnology products

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

- [ 1.4.1 ] Manufacture of:
  - [ 1.4.1.1 ] Herbal products

**[ 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
  - [ 1.5.1.15 ] Other non-sterile medicinal products
    - Powders, Granules
- [ 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
  - [ 2.2.3.2 ] Immunological products
  - [ 2.2.3.5 ] Biotechnology products

**[ 2.3 ] Other Importation Activities**

- [ 2.3.1 ] Site of Physical Importation
- [ 2.3.2 ] Importation of Intermediate which undergoes further processing
- [ 2.3.4 ] Other
  - Importation of QP Certified IMPs from a country on the approved country for import list