

# Medicines and Healthcare products Regulatory Agency

## MANUFACTURER'S AUTHORISATION

<b>1: Authorisation Number</b>	UK MIA 32496
<b>2: Name of authorisation holder</b>	AESICA QUEENBOROUGH LIMITED AESICA QUEENBOROUGH LIMITED, BUILDING 55, NORTH ROAD, QUEENBOROUGH, ME11 5EL, UNITED KINGDOM
<b>3: Address(es) of manufacturing site(s)</b>	AESICA QUEENBOROUGH LIMITED, NORTH ROAD, QUEENBOROUGH, ME11 5EL, UNITED KINGDOM
<b>4: Legally registered address of authorisation holder</b>	AESICA QUEENBOROUGH LIMITED, NORTH ROAD, QUEENBOROUGH, ME11 5EL, UNITED KINGDOM
<b>5: Scope of authorisation and dosage forms</b>	ANNEX 1 and/ or ANNEX 2
<b>6: Legal Basis of authorisation</b>	Regulation 17 of The Human Medicines Regulations 2012 (SI 2012/1916)
<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>	30/01/2023
<b>9: Annexes attached</b>	Annex 1 and/or Annex 2

### SCOPE OF AUTHORISATION

#### Annex 1

Name and address of the site:

**AESICA QUEENBOROUGH LIMITED, BUILDING 55, NORTH ROAD, QUEENBOROUGH, ME11 5EL, UNITED KINGDOM**

Human Medicinal Products
Authorised Operations
MANUFACTURING OPERATIONS (according to part 1)
<b>Part 1 - MANUFACTURING OPERATIONS</b> <b>[ 1.2 ] Non-sterile products</b> [ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms) [ 1.2.1.1 ] Capsules, hard shell <b>[ 1.5 ] Packaging</b> [ 1.5.1 ] Primary packaging [ 1.5.1.1 ] Capsules, hard shell [ 1.5.2 ] Secondary packaging

## SCOPE OF AUTHORISATION

### Annex 1

Name and address of the site:

**AESICA QUEENBOROUGH LIMITED**, NORTH ROAD, QUEENBOROUGH, ME11 5EL, UNITED KINGDOM

Human 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>[ 1.2 ] Non-sterile products</b> [ 1.2.1 ] Non-Sterile Products (processing operations for the following dosage forms) [ 1.2.1.6 ] Liquids for internal use [ 1.2.1.8 ] Other solid dosage forms [ 1.2.1.13 ] Tablets <b>Special Requirements</b> Hormonal tablets [ 1.2.1.17 ] Other non-sterile medicinal products Inhalation Anaesthetics [ 1.2.2 ] Batch certification <b>[ 1.5 ] Packaging</b> [ 1.5.1 ] Primary packaging [ 1.5.1.1 ] Capsules, hard shell [ 1.5.1.2 ] Capsules, soft shell [ 1.5.1.6 ] Liquids for internal use [ 1.5.1.8 ] Other solid dosage forms [ 1.5.1.13 ] Tablets <b>Special Requirements</b> Hormonal tablets [ 1.5.1.17 ] Other non-sterile medicinal products Inhalation Anaesthetics [ 1.5.2 ] Secondary packaging <b>[ 1.6 ] Quality control testing</b> [ 1.6.2 ] Microbiological: non-sterility [ 1.6.3 ] Chemical/Physical <b>Part 2 - IMPORTATION OF MEDICINAL PRODUCTS</b> <b>[ 2.1 ] Quality control testing of imported medicinal products</b> [ 2.1.2 ] Microbiological: non-sterility [ 2.1.3 ] Chemical/Physical <b>[ 2.2 ] Batch certification of imported medicinal products</b> [ 2.2.2 ] Non-sterile products

[ 2.3 ] Other Importation Activities

[ 2.3.2 ] Importation of Intermediate which undergoes further processing

MHRA-GMDP

MHRA

MHRA-GMDP

MHRA