Medicines and Healthcare products Regulatory Agency

MANUFACTURER'S AUTHORISATION

1: Authorisation Number UK MIA(IMP) 45848

2: Name of authorisation holder QUOTIENT SCIENCES (ALNWICK) LIMITED

3: Address(es) of manufacturing site(s)

QUOTIENT SCIENCES (ALNWICK) LIMITED, TAYLOR DRIVE,

ALNWICK, NE66 2DH, UNITED KINGDOM

QUOTIENT SCIENCES (ALNWICK) LIMITED, TRENT HOUSE, MERE

WAY, RUDDINGTON FIELDS BUSINESS PARK, RUDDINGTON,

NOTTINGHAM, NG11 6JS, 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

4: Legally registered address of authorisation holder

authority of the member state granting the

manufacturing authorisation

Confidential

8: Authorisation Date 02/12/2024

9: Annexes attached Annex 1 and/or Annex 2

SCOPE OF AUTHORISATION

Annex 2

Name and address of the site:

QUOTIENT SCIENCES (ALNWICK) LIMITED, TAYLOR DRIVE, ALNWICK, NE66 2DH, UNITED KINGDOM

Human Investigational 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 Investigational Medicinal Products

[1.1.1] Aseptically prepared (processing operations for the following dosage forms)

[1.1.1.4] Small volume liquids

[1.1.1.6] Other aseptically prepared products

Radiolabelled Products & Cytotoxic Products

[1.1.3] Batch certification

[1.2] Non-sterile investigational medicinal products

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

[1.2.1.1] Capsules, hard shell

[1.2.1.2] Capsules, soft shell

[1.2.1.5] Liquids for external use

[1.2.1.6] Liquids for internal use

[1.2.1.8] Other solid dosage forms

[1.2.1.11] Semi-solids

[1.2.1.15] Other non-sterile medicinal products

Radiolabelled Products & Cytotoxic Products, Moulded Tablets, Drug in a Vial, Over encapsulated product.

[1.2.2] Batch certification

[1.4] Other investigational medicinal products or manufacturing activity

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

[1.4.2.1] Filtration

[1.6] Quality control testing

[1.6.1] Microbiological: sterility

[1.6.2] Microbiological: non-sterility

[1.6.3] Chemical/Physical

Part 2 - IMPORTATION OF MEDICINAL PRODUCTS

[2.1] Quality control testing of imported medicinal products

[2.1.1] Microbiological: sterility

[2.1.2] Microbiological: non-sterility

[2.1.3] Chemical/Physical

MHRA: GMDP MHRA

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