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
4: Legally registered address of authorisation holder	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 authority of the member state granting the manufacturing authorisation	Confidential
8: Authorisation Date	23/02/2026
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

MHRA-GMDP MHRA-GMDP MHRA

[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 activitiy

[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