

BE IT KNOWN that I, Sunita Kumeri of, 18-24 Stoke Road, Slough, Berkshire United Kingdom a duly authorised Notary Public

CERTIFY ONLY that Brian Michael Howes who is well known to me and who is duly authorised by GlaxoSmithKline ("The Company") to represent them in this matter has today caused the annexed Certificate of GMP Compliance of a Manufacturer issued to Glaxo Operations UK Ltd Trading as Glaxo Wellcome Operations to be produced to me and that he has represented to me on behalf of the Company that the said document is a true copy of the original document.

SIGNED and sealed at 18-24 Stoke Road, Slough, Berkshire aforesaid on 15th May

2019.

Sunita Kumeri Notary Public

England and Wales

Protocol No. 3/19

	APOSTILLE (Convention de La Haye du 5 octobre 1961)			
1.	Country: Pays / Pais: United Kingdom of Great Britain and Northern Ireland			
	This public document Le présent acte public / El presente documento público			
2.	Has been signed by a été signé par Sunita Kumeri ha sido firmado por			
3.	Acting in the capacity of agissant en qualité de quien actúa en calidad de	Notary Public		
4.	Bears the seal / stamp of est revêtu du sceau / timbre de y está revestido del sello / timbre de	The Said Notary Public		
	Certified Attesté / Certificado			
5.	at London á / en	6. the 16 May 2019		
7.	by Her Majesty's Principal Secretary of State for Foreign and Commonwealth Affairs			
8.	Number sous no / bajo el numero	APO-1458798		
9.	Seal / stamp Sceau / timbre Sello / timbre	10. Signature A. Khan Signature Firma A. Khan A. Khan A. Khan		

This Apostille is not to be used in the UK and only confirms the authenticity of the signature, seal or stamp on the attached UK public document. It does not confirm the authenticity of the underlying document. Apostilles attached to documents that have been photocopied and certified in the UK confirm the signature of the UK official who conducted the certification only. It does not authenticate either the signature on the original document or the contents of the original document in any way.





Certificate No: UK MIA 4 Insp GMP/IMP 4/15159-0027

Medicines and Healthcare products Regulatory Agency

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER

Part 1

Issued following an inspection in accordance with Art. 111(5) of Directive 2001/83/EC.

The competent authority of the United Kingdom confirms the following:

The manufacturer

GLAXO OPERATIONS UK LTD TRADING AS GLAXO WELLCOME OPERATIONS

Site address

PRIORY STREET

WARE SG12 0DJ

UNITED KINGDOM

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. MIA 4 in accordance with Art. 40 of Directive 2001/83/EC transposed in the following national legislation: The Human Medicines Regulations 2012 (SI 2012/1916).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 21/01/2019, it is considered that it complies with the principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC.

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection. However, this period of validity may be reduced or extended using regulatory risk management principles by an entry in the Restrictions or Clarifying remarks field.

This certificate is only valid when presented with all pages and both parts 1 and 2.

The authenticity of this certificate may be verified in EudraGMDP. If it does not appear please contact the issuing authority.



Medicines & Healthcare products Reg<mark>u</mark>latory Agency



Certificate No: UK MIA 4 Insp GMP/IMP 4/15159-0027

Part 2

Human Medicinal Products

- 1. MANUFACTURING OPERATIONS
- 1.1 Sterile products

Not Authorised

- 1.2 Non-sterile products
- 1.2.1 Non-sterile products (processing operations for the following dosage forms)
 - 1.2.1.1 Capsules, hard shell
 - 1.2.1.8 Other solid dosage forms
 - 1.2.1.13 Tablets
 - 1.2.1.17 Other non-sterile medicinal products Licensable medical devices
- 1.3 Biological medicinal products
- 1.3.1 Biological medicinal products
 - 1.3.1.8 Other biological medicinal products Steroids
- 1.4 Other products or manufacturing activity
- 1.4.1 Manufacture of
 - 1.4.1.3 Other
 Micronisation of active substances
- 1.5 Packaging
- 1.5.2 Secondary packaging
- 1.6 Quality control testing
- 1.6.2 Microbiological: non-sterility
- 1.6.3 Chemical/physical
- 2. IMPORTATION OF MEDICINAL PRODUCTS
- 2.1 Quality control testing of imported medicinal products
- 2.1.2 Microbiological: non-sterility
- 2.1.3 Chemical/physical
- 2.2 Batch certification of imported medicinal products







- 2.2.2 Non-sterile products
- 2.3 Other importation activities

 Not Authorised





Medicines & Healthcare products Regulatory Agency



Certificate No: UK MIA 4 Insp GMP/IMP 4/15159-0027

3. MANUFACTURING OPERATIONS

- 3.1 Manufacture of Active Substance by Chemical Synthesis
 Not Authorised
- 3.2 Processing Activities of Active Substance from Natural Sources
 Not Authorised
- 3.3 Manufacture of Active Substance using Biological Processes
 Not Authorised
- 3.4 Manufacture of sterile active substance Not Authorised
- 3.5 General Finishing Steps Not Authorised
- 3.6 Quality Control Testing Not Authorised
- 4 Other Activities
 Not Authorised



Medicines & Healthcare products Regulatory Agency



Certificate No: UK MIA 4 Insp GMP/IMP 4/15159-0027

Any restrictions or clarifying remarks related to the scope of this certificate:

Scope of IMP packaging - For product protection purposes only; excludes clinical trial labelling activities

1. Building(s)/Area(s)

N/A

2. Room(s)

N/A

3. Line(s) Equipment(s)

N/A

4. QC testing

N/A

5. Medicinal Product(s)/IMP(s)

N/A

Name of the authorised person of the Competent Authority of the United Kingdom

Dr A J Gray Head of Inspectorate inspectionplanning@mhra.gov.uk

Date: 04/03/2019

