

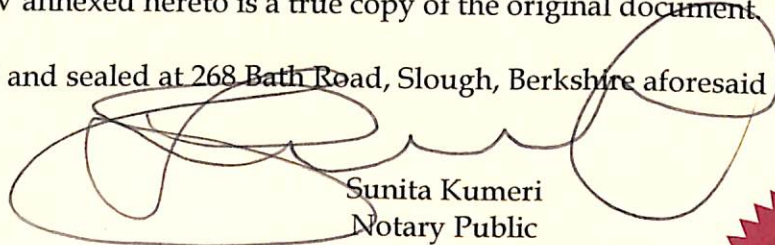


BE IT KNOWN that I, Sunita Kumeri of, 268 Bath Road, Slough, Berkshire United Kingdom a duly authorised Notary Public

CERTIFY that

1. The signature set and subscribed to the certificate at the foot of the first page of the copy document annexed hereto is genuine having been subscribed thereto by Zoe Bruce whose identity I the Notary attest and who is duly authorised by Pfizer Limited ("the Company") to represent them in this matter, and
2. Zoe Bruce has thereby certified on behalf of the company that the Certificate of GMP Compliance of a Manufacturer issued to Pfizer Manufacturing Belgium NV annexed hereto is a true copy of the original document.

SIGNED and sealed at 268 Bath Road, Slough, Berkshire aforesaid on 13th May 2020.


Sunita Kumeri
Notary Public
England and Wales



Protocol No. 13/20



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 ha sido firmado por	Sunita Kumeri
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 à / en	London
6. the le / el día	14 May 2020
7. by par / por	Her Majesty's Principal Secretary of State for Foreign and Commonwealth Affairs
8. Number sous no / bajo el numero	APO-1906024
9. Seal / stamp Sceau / timbre Sello / timbre 	10. Signature R. James Signature Firma 

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229378

Federal Agency for Medicines and Health Products

CERTIFICATE NUMBER: *BE/GMP/2020/009*

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER^{1, 2}

Part 1

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

The competent authority of Belgium confirms the following:

The manufacturer: *Pfizer Manufacturing Belgium NV*

Site address: *Rijksweg 12, Puurs, 2870, Belgium*

Has been inspected under the national inspection programme in connection with manufacturing authorisation no. *277 H* in accordance with Art. 40 of Directive 2001/83/EC transposed in the following national legislation:

Article 12 bis, § 1 of the Law of 25th March 1964 related to the Medicinal Products

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on *2020-01-29*, it is considered that it complies with :

- The principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC³

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 valid only 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.

¹ The certificate referred to in paragraph 111(5) of Directive 2001/83/EC and 80(5) of Directive 2001/82/EC, shall also be required for imports coming from third countries into a Member State.

² Guidance on the interpretation of this template can be found in the Help menu of EudraGMDP database.

³ These requirements fulfil the GMP recommendations of WHO.

I, Zoe Bruce, on behalf of Pfizer,
certify this document
to be a true copy
of the original

1 MANUFACTURING OPERATIONS

1.1	Sterile products
	<i>1.1.1 Aseptically prepared (processing operations for the following dosage forms)</i> 1.1.1.2 Lyophilisates Special Requirements 7 Other: highly potent products/certain hormones(en) 1.1.1.3 Semi-solids Special Requirements 7 Other: highly potent products(en) 1.1.1.4 Small volume liquids Special Requirements 7 Other: highly potent products/ certain hormones(en)
	<i>1.1.2 Terminally Sterilised (processing operations for the following dosage forms)</i> 1.1.2.3 Small volume liquids
	<i>1.1.3 Batch certification</i>
1.3	Biological medicinal products (list of product types)
	<i>1.3.1 Biological medicinal products (list of product types)</i> 1.3.1.2 Immunological products 1.3.1.5 Biotechnology products 1.3.1.6 Human or animal extracted products Special Requirements 7 Other: low molecular weight heparin(en)
	<i>1.3.2 Batch Certification (list of product types)</i> 1.3.2.2 Immunological products 1.3.2.5 Biotechnology products 1.3.2.6 Human or animal extracted products

Clarifying remarks (for public users)

Inspected area: 'Aseptic Area South' / 1.3.1.2: formulation and filling only 1.3.1.5: formulation and filling only 1.3.1.6: formulation and filling only

2020-04-22

Name and signature of the authorised person of the
Competent Authority of Belgium

in name of



Séverine Brasseur
(Signature)
2020.04.23 09:57:03
+02'00'

Mr. Xavier De Cuyper
Federal Agency for Medicines and Health Products
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Fax: +32 2 5284001