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BELGIUM

Quality of Vaccines and Blood Products

BELAC
081-TEST

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EU OFFICIAL CONTROL AUTHORITY BATCH RELEASE CERTIFICATE FOR IMMUNOLOGICAL PRODUCTS

EU/EEA OFFICIAL CONTROL AUTHORITY BATCH RELEASE CERTIFICATE – Finished Product

Examined under Article 114 of Directive 2001/83/EC as amended by Directive 2004/27/EC (Immunological Medicinal Products) and in accordance with the Administrative Procedure for Official Control Authority Batch Release.

Trade name	**COMIRNATY**
International Non-proprietary Name / Ph.Eur. name / Common name	**Pandemic COVID-19 Vaccine (mRNA)**
Batch numbers appearing on the package and other identification numbers associated with this batch[1]	**FG7911**
Type of container	**Vial**
Total number of containers in this batch	[redacted]
Number of doses per container	**6 doses**
Date of start of period of validity	**22 July 2021**
Date of expiry	**31 December 2021**
Marketing Authorisation number (member state / EU) issued by	**EU/1/20/1528**
Name and address of manufacturer	**Pfizer Manufacturing Belgium NV / 2870 Puurs, Belgium**
Name and address of marketing authorisation holder if different	**BioNTech Manufacturing GmbH An der Goldgrube 12 55131 Mainz, Germany**

This batch has been examined using documented procedures which form part of a quality system which is in accordance with the ISO/IEC 17025 standard.
This examination is based on the relevant EU OCABR guideline for this product.

This batch is in compliance with the approved specifications laid down in the relevant European Pharmacopoeia monographs and the above marketing authorisation and is released.

Signature **Date of issue**	Digitally signed by [redacted] Date: 2021.08.23 16:21:55 +02'00'
Name and function of signatory	**p.o. [redacted], M.Sc., Head, Quality of Vaccines and Blood Products Service**

Certificate number: BE/21/1580

[1] Such as batch number of final bulk

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