

## Quality of Vaccines and Blood Products

### 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.

<b>Trade name</b>	COMIRNATY
<b>International Non-proprietary Name / Ph.Eur. name / Common name</b>	<b>Pandemic COVID-19 Vaccine (mRNA) Comirnaty 30 µg/dose</b>
<b>Batch numbers appearing on the package and other identification numbers associated with this batch<sup>1</sup></b>	[REDACTED]
<b>Type of container</b>	Vial
<b>Total number of containers in this batch</b>	[REDACTED]
<b>Number of doses per container</b>	6 doses
<b>Date of start of period of validity</b>	03 December 2021
<b>Date of expiry</b>	31 May 2022
<b>Marketing Authorisation number (member state / EU) issued by</b>	EU/1/20/1528/002-003
<b>Name and address of manufacturer</b>	[REDACTED]
<b>Name and address of marketing authorisation holder if different</b>	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.**

<b>Signature</b>	Digitally signed by [REDACTED]
<b>Date of issue</b>	Date: 2022.01.20 11:12:49 +01'00'
<b>Name and function of signatory</b>	[REDACTED]

**Certificate number:** [REDACTED]

<sup>1</sup> Such as batch number of final bulk