

ORDER

X document registered by

for the amendment of Order No. RD-01-49 of 28.01.2022

Pursuant to Art. 61, para. 2, art. 63, para. 4, 5 and 11 and Art. 63c of the Health Act, Art. 73 of the Code of Administrative Procedure, and in connection with Regulation (EU) 2021/953 of the European Parliament and of the Council of 14 June 2021 on a framework for the issuance, verification and acceptance of interoperable COVID-19 vaccination, test and recovery certificates (EU Digital COVID Certificate) to facilitate free movement during the COVID-19 pandemic and Decision of the Council of Ministers No. 826 of 25 November 2021 extending the term announced by Decision of the Council of Ministers No. 325 of 14 May 2020 epidemic emergency situation, extended by Decision of the Council of Ministers No. 378 of 12 June 2020, Decision of the Council of Ministers No. 418 of 25 June 2020, Decision of the Council of Ministers No. 482 of 15 July 2020, Decision of the Council of Ministers No. 525 of 30 July 2020, Decision of the Council of Ministers No. 609 of 28 August 2020, Decision of the Council of Ministers No. 673 of 25 September 2020, Decision of the Council of Ministers No. 855 of 25 November 2020, Decision of the Council of Ministers No. 72 of 26 January 2021, Decision of the Council of Ministers No. 395 of 28 April 2021, Decision of the Council of Ministers No. 426 of 26 May 2021, Decision of the Council of Ministers No. 547 of 28 July 2021 and Decision of the Council of Ministers No. 629 of 26 August 2021, and a proposal by the Chief State Health Inspector,

I hereby

ORDER:

- I. In Order No. RD-01-49 of 28.01.2022 the following amendments shall be made:
- 1. In item I, 10, the word "states" shall be replaced by "states (territories)".
- 2. Appendix No. 3 to item I, 9.2.1. shall be amended as follows:

"Annex No. 3 to point I, 9.2.1.

Trade name of the vaccine	Name of the manufacturer /	Completed
according to the EU marketing	holder of the marketing	vaccination
authorization / WHO list	authorization	course
Comirnaty/ (/ BNT162b2 Pfizer-	BioNTech Manufacturing GmbH/	2 doses
BioNTech Covid-19 vaccine)	Pfizer-Biontech	
Vaxzevria/ AZD1222	AstraZeneca AB	2 doses
Spikevax/COVID-19 VACCINE	MODERNA BIOTECH	2 doses
Moderna / mRNA-1273		
Janssen / Ad26.COV2.S	Janssen-Cilag International NV	1 dose
- / Covishield (ChAdOx1_nCoV-	Serum Institute of India Pvt. Ltd	2 doses
19)		
- / SARS-CoV-2 Vaccine (Vero	Sinopharm / BIBP1	2 doses
Cell), Inactivated (InCoV)		
- / COVID-19 Vaccine (Vero Cell),	Sinovac Life Sciences Co., Ltd.	2 doses
Inactivated/Coronavac		
- / Sputnik V	Russian Direct Investment Fund	2 doses
Nuvaxovid /NVX-CoV2373	Novavax	2 doses
- /NVX-CoV2373/Covovax	Serum Institute of India Pvt. Ltd	2 doses

3. In the title of Appendix No. 4 to item I, 10, the word "states" shall be replaced by "states (territories)".

II. This Order shall come into force on February 3, 2022.

III. The order shall be published on the website of the Ministry of Health.

The order is subject to appeal within one month from the publication on the website of the Ministry of Health, before the relevant administrative court under the Administrative Procedure Code.

Х

PROF. ASENA SERBEZOVA, PhD, MSCPHARM, MPH

Minister of Health