REPUBLIC OF BULGARIA
Ministry of Health
Minister of Health

O R D E R

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document.
registered by:

for amendment of Order № RD-01-733 dated 27.08.2021, amended by Order № RD-01-767 dated 10.09.2021, Order № RD-01-794 dated 24.09.2021, Order № RD -01-820 dated 08.10.2021 and Order № RD-01-874 dated 26.10.2021

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 relation to 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 vaccination certificates against, COVID testing and recovery -19 (EU Digital COVID Certificate) in order to facilitate free movement during the pandemic of COVID-19 and Council of Ministers Decision № 629 dated 26 August 2021 extending the term announced by Council of Ministers Decision № 325 dated 14 May 2020 extraordinary epidemic situation, extended by Decision № 378 of the Council of Ministers dated 12 June 2020, Decision № 418 of the Council of Ministers dated 25 June 2020, Decision № 482 of the Council of Ministers dated 15 July 2020, Decision № 525 of the Council of Ministers dated 30 July 2020, Decision № 609 of the Council of Ministers dated 23 August 2020, Decision № 673 of the Council of Ministers dated 25 September 2020, Decision № 855 of the Council of Ministers dated 25 November 2020., Decision № 72 of the Council of Ministers dated 26 January 2021, Decision № 395 of the Council of Ministers dated 28 April 2021, Decision № 426 of the Council of Ministers dated 26 May 2021 and Decision № 547 of the Council of Ministers dated 28 July 2021 and a proposal from the Chief State Health Inspector

O R D E R:

I. In Order № RD-01-733 dated 27.08.2021, amended by Order № RD-01-767 dated 10.09.2021, Order № RD-01-794 dated 24.09.2021, Order № RD -01-820 dated 08.10.2021 and Order № RD-01-874 dated 26.10.2021, the following amendments and additions are made:

1. In item **I**:

(a) point 4.1.2. is amended, as follows:

„4.1.2. the quarantined person may carry out a polymerase chain reaction test for the detection of COVID-19 or a rapid antigen test as specified in Annex № 2 by the end of the day

following his arrival in the country. In case of a negative result from the test, the quarantine of the person shall be considered terminated from the day following the registration of the result in the National Information System for Combating COVID-19. ";

(b) point 4.2.3. is amended, as follows:

„4.2.3. The quarantined person under item 4.2.2. may carry out a polymerase chain reaction test for the detection of COVID-19 or a rapid antigen test as specified in Annex 2 by the end of the day following his arrival in the country. In case of a negative result from the test, the quarantine of the person shall be considered terminated from the day following the registration of the result in the National Information System for Combating COVID-19. ";

c) in point 4.7. the second and third sentences are amended, as follows:

"In these cases, the quarantined person may carry out a polymerase chain reaction test to prove COVID-19 by the end of the day following his arrival in the country. In case of a negative result from the test, the quarantine of the person shall be considered terminated from the day following the registration of the result in the National Information System for Combating COVID-19. ";

d) Point 4.8.1. is amended, as follows:

„4.8.1. The quarantined person under item 4.8., who is a Bulgarian citizen or a person with the status of permanent, long-term or continuous residence on the territory of the Republic of Bulgaria, may within the end of the day following his arrival in the country make a test by the method of polymerase chain reaction for the detection of COVID-19. In case of a negative result from the conducted test, the quarantine of the person shall be considered terminated from the day following the registration of the result in the National Information System for Combating COVID-19. ".

2. Annex № 1 to points I, 3 is amended as follows:

„Annex № 1 to item I, 3

List of countries by color zones, countries for which there is information about a significant negative change in the epidemic situation and countries with a reciprocal basis

Green zone:

Vatican City State, Luxembourg, Italy, Kingdom of Spain, Republic of Malta, Kingdom of Sweden.

Orange zone:

All countries outside the green and red zone.

Red zone:

People's Republic of Bangladesh, Myanmar, Kingdom of Bhutan, Democratic Socialist Republic of Sri Lanka, Republic of Maldives, Republic of South Africa, Republic of Botswana, United Republic of Tanzania, Republic of Seychelles, Republic of Namibia, Republic of Zambia, Sultanate of Oman, Turkmenistan, Republic of Tajikistan, Afghanistan, Kyrgyz Republic, Mongolia, Democratic People's Republic of Korea, Republic of Chile, Eastern Republic of Uruguay, Federative Republic of Brazil, Republic of Paraguay, Multinational State of Bolivia, Republic of Suriname, Republic of Panama, Republic of Costa Rica, Republic of Guatemala, Belize, Republic of El Salvador, Republic of Cuba, Dominican Republic, Fiji, Saint Lucia, Federation of Saint Kitts and Nevis, Grenada, Barbados, Singapore, Brunei, United Kingdom of Great Britain and Northern Ireland, Georgia, Armenia, Montenegro, Republic of Serbia, Republic of Lithuania, Republic of Latvia, Republic of Estonia, Romania, Slovenia, Croatia, Slovak Republic, Ukraine, Republic of Austria, Kingdom of the Netherlands, Ireland, the Czech Republic, the Kingdom of Belgium, Hungary, the Kingdom of Denmark, Iceland, Greece, the Federal Republic of Germany, the Principality of Liechtenstein, the Principality of Andorra, the Republic of Poland and the Swiss Confederation. ".

3. Annex № 3 to point I, 10.2.1 is amended as follows:

"Annex № 3 to item I, 10.2.1

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Trade name of the vaccine according to the EU marketing authorization / WHO list	**Name of the manufacturer / marketing authorization holder**	**Completed vaccination course**
Comirnaty/ (/ BNT162b2 Pfizer-BioNTech Covid-19 vaccine)	BioNTech Manufacturing GmbH/ Pfizer-Biontech	2 doses
Vaxzevria/ AZD1222	AstraZeneca WCH/ AstraZeneca Canada Inc.	2 doses
- / AZD1222	SK Bioscience Co Ltd	2 doses
Spikevax/COVID-19 VACCINE Moderna / mRNA-1273	MODERNA BIOTECH	2 doses
Janssen / Ad26.COV2.S	Janssen-Cilag International NV	1 dose
- / Covishield (ChAdOx1_nCoV-19)	Serum Institute of India	2 doses
- / SARS-CoV-2 Vaccine (Vero Cell), Inactivated (lnCoV)	Sinopharm / BIBP1	2 doses
- / COVID-19 Vaccine (Vero Cell), Inactivated/Coronavac	Sinovac	2 doses
- / Sputnik V(Gam-COVID-VacComponent I Gam-COVID-Vac Component II)	The Gamaleya National Center of Epidemiology and Microbiology	2 doses
SARS-CoV-2 Vaccine, Inactivated (Vero Cell)/ COVAXIN	Bharat Biotech, India	2 doses

4. Annex № 4 to points I, 11 is amended as follows:

"Annex № 4 to item I, 11

List of countries whose COVID-19 vaccination, testing and getting over disease certificates are considered equivalent to the EU digital COVID certificate.

Republic of Northern Macedonia, Republic of San Marino, Swiss Confederation, Republic of Turkey, Ukraine, Vatican City State (only in respect of vaccination certificates issued), Principality of Andorra, Republic of Albania, Faroe Islands, Principality of Monaco, Republic of Panama, Kingdom of Morocco, Israel, Armenia and the United Kingdom of Great Britain and Northern Ireland."

II. The order enters into force on 23.11.2021.

III. The order ought to 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, in front of the relevant administrative court under the Administrative Procedure Code.

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DR. STOYCHO KATSAROV
Minister of Health