Exceptional meeting of the Steering Board on CureVac contract-18 November

An exceptional Steering Board meeting was held to inform the Members on the CureVac contract and on next steps.

The Members who had not yet done so were asked to confirm the decryption of the contract.

The **key elements of the contract** were presented again to the Members, following an extensive presentation at the previous meeting (Friday 13 November).

Members were informed that, as set out in the Agreement, Member States have the right to optout no later than 5 working days after the Commission communicated its intention to conclude the APA. In the case of this contract, therefore, the deadline to exercise the right to opt-out would expire on Tuesday 24 November, except for two countries that have national holidays during this period, for which the period would expire the following day.

The Members were reminded that the MSs needed to notify only if the intended to opt-out, otherwise they were considered part of the contract.

It was agreed among the MSs that that discussions on				should already start.	
Consequently,		would be circulated.			
At the request of the Members, a discussion followed on the state of play of t					
	on BioNTec	h/Pfizer, J&J with some c	arification from the	Commission on how	
to fill in the Orde	er Forms, na	mely regarding:			

- the number of doses;
- allocations;
- the date of the signature.

Minutes: Steering Board meeting, 20 November 2020

1. Update on the CureVac contract

Following the comprehensive presentations during the last two SB meetings, the Members were informed again on key elements of the CureVac contract and on the next steps.

A general discussion followed on the **process of authorisation**, during which it was outlined that:

- the safety, efficacy and quality of the vaccines would be heavily scrutinised, as the authorisation process normally requires;
- scientific communication (including from EMA) should be given more visibility
 and priority (especially on the new technologies), in order to inform and to
 reassure the public with scientific facts over the safety, effectiveness and quality
 of vaccines.

The Commission informed the Members on several aspects regarding the implementation

2. Short update on the AZ contract

Customer set up forms:

of the AZ contr	act, as outlined below.
Distribution H	ubs:
 Member authoris 	rs were reminded that the hubs needed to have a wholesale distribution sation;
	such an authorisation, AZ would not be able/authorised to ship the doses istribution hub;
•	

	AZ had asked the Member States to fill in the customer set up forms;				
	 without these forms, AZ cannot ship doses; Members were drawn the attention that only to AZ ighthat the state of the stat				
	 those Member States that did not send them, were encouraged to do so as soon as possible. 				
3.	Update on the				
	A discussion was held on the various and their state of play.				
	The Members were informed that: • a table was circulated for Johnson & Johnson, reflecting the arrangement amongst Member States to secure doses				
	 the for Pfizer/BioNTech was closed, following the feedback provided by all MSs; the was ongoing for the CureVac and Moderna contacts. The Members were encouraged to provide timely feedback. 				
4.	Update on contracts on discussions with other companies				
Moderna – intensive work on the contact was ongoing with the aim of reaching its conclusion the following days. The members were updated on discussions regarding key aspects, such as: liability, delivery schedule, termination rights, resale, volume, price etc.					
	Novavax - the Members were updated on discussions on liability and indemnification.				
- as discussions with the company were concluded, the steering should decide on launching the contract process.					
	Valneva - on the structure of the contract, on price, delivery and payment schedules and options. The steering should decide on launching the contract process.				
we	embers were informed about leaks in the press about elements from the contracts and are reminded about the very strict confidentiality rules that, if breached, could lead to rious repercussions				

5. Joint procurement for syringes and needles

The Commission informed that:

- the evaluation of the joint procurement procedure for vaccination supplies was ongoing;
- priority was given to evaluation of syringes and needles;
- depending on the quality of offers and reactivity of companies, the evaluation of offers for syringes and needles might be finalized by , followed by the contract signature ;

Offers received show that the needs of participating countries should be covered. However, final coverage will depend on the results of the evaluation.

Depending on the replies received, the evaluation could be finalised

6. COVAX

The Commission indicated that a working group would be created on market exchange at the in COVAX. This working group could serve to move forward on in-kind donations and humanitarian buffer.

The Commission also informed about the outcome of the EUCO held the day before and discussions on vaccine donation and COVAX.

Exceptional meeting of the Steering Board on Moderna contract 24 November

This session was dedicated to a comprehensive presentation of the Moderna APA.

Key elements of the contract were presented and explained in great detail to the members. Elements illustrated include:

- information about the characteristics of the vaccine;
- · volume (base and optional doses);
- · price, upfront payment and payment schedule/structure;
- delivery schedule;
- resale and donation clause;
- liability and indemnification provisions;
- exchange rate;
- termination clauses and refundability etc.

out no later than 5 working days after the Commission communicated its intention to conclude the APA. If the contract would be adopted on the 25 November, the deadline to exercise the right to opt-out would be set for Wednesday -2 December, except for countries that have national holidays during this period.
The Members welcomed the conclusion of the contract. Some strongly advocated for exercising immediately the options to commit already for doses
COVAX
The Commission informed about the meeting with the and the 27 EU Ambassadors in Geneva, where the Commission representative presented a state of play of the EU portfolio.
The Commission also informed about on-going work with and other MS to establish an EU-

wide approach on donations, which would be submitted to the SB Members soon.

Members were informed that, as set out in the Agreement, Member States had the right to opt-