Minutes: Steering Board Meeting, 21 August 2020

1. Update on AstraZeneca contract

opt-out expired on Frie	sion (EC) reminded the Meml day-21 August 2020, except August- due to a National hol	for		y deadline to ose deadline
The European Commiss	sion stressed that no opt-out re	equests have been	received so fa	r.
decide to opt-out, the d States, which will not b	e, the European Commission oses becoming available work required to take more than the the possibility/option to take	ald not be impose their current pro r	ed on to the of ata. Instead, t	her Member he other MS
Commission drew the a order to decrease the nu	vishes to take the doses or attention to the fact that the mber of doses to which the E her conditions, including	contract might n EU may commit. I	eed to be reno	egotiated, in
a portfolio, reminding the SANOFI, J&J, CureVarepresents an insurance successful vaccine, on i	sion underlined, also as a prime intensive ongoing parallel ac, Moderna and BioNTec strategy as a) the successful ts own, can provide sufficien of solidarity, offering thus the successful that the successfu	processes with size th- hopefully next vaccine is not known t quantities (espec	x companies (Axt week). The cown in advance cially at the be	AstraZeneca, nis portfolio ce and b) no eginning). Its
Some Member States	explicitly supported this ap	pproach.		

Q&A on AZ contract

The objective of the Q&A session was to give MS an opportunity to ask any remaining questions on the AZ contract. Clarifications provided at the meeting are noted below.

Communication channels on an eventual opt-out

In case of an opt-out, Member States (a person that has the authority to bind the Government) should send a registered letter to the Commission and a scan of this letter to the EC vaccines mailbox.

Pro rata allocation

The pro rata allocation is the default distribution key for the doses. Member States will not be required to take more doses than their pro rata allocation. It was also clarified that the "opt in/out" was a matter or agreeing or not to take part in the Agreement with the company and was not about the number of doses.

Member States will need to			the national hub and		
delivery within their territory to vaccination	P a see	Member	States will		
from the hub to the final of	estination.				
The European Commission encouraged Member States to cooperate regarding the practicalities					
in distribution and transport.					
Cross references					
Optional/additional doses					
Member States will have to express interest in the optional / additional doses and in case of interest exceeding the available supply of doses, a pro-rata allocation will apply.					
As the term suggests it, this is an option and	was added to the c	ontract in	order to		
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<u>Order form – deadline for submission</u>					
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The Order Forms have to be entered into by each of the Participating Member States
So the timing for submission of the order form is .
will be slightly different from one contract to the other,
Effective date
The date of the signature of the Contract.
Other issues
Member States asked:
 if it would be possible to designate e different hub for each delivery. The EC is not in a position to reply, as this would depend on the company. if the number of doses could be disclosed publicly. The Commission confirmed this.
2. Update on the contracts in the SANOFI
The Member States were informed that the offer from Sanofi was received, a first meeting with
of took place and the work on the draft APA was in progress.
Johnson & Johnson
The deadline for submitting expires on 23 August.
was submitted from J&J by the time of the meeting.
CureVac
The European Commission sent on Thursday 20 August an invitation to CureVac to submit a by 30 th August.
BioNTech/Pfizer
The European Commission recalled the

The Commission and Member States	BioNTech could also be in
process soon.	
3. Update on discussions with Modern	1a
	were constructive and informed that the key points of discussions. This allowed the . The plan was to send the invitation on
The Commission would also communicate or	n the progress in the exploratory talks publicly.
The discussions have been focused on	with the Member States to
	resented, especially since the development of the als, making it one of the front-runners for the time
AOB	
1. COVAX	
The second secon	itment to the success of the COVAX facility, as firm toach to vaccines and indicated EU should have a OVAX governance structure.
The Commission stands ready to lend the fin development of the COVAX.	ancial support needed as well as the expertise in the
The state of the s	X, as the deadline for the Members States to react to the 31 of August. The COVAX secretariat informed and of the week.
Participants agreed that it was important	to keep a good flow of communication with MS

vaccines strategy.

The Member States called for an EU consolidated approach and not 27 individual commitments.

The Commission fully supported this and also argued in favour of a coordinated EU (EC + Member State) approach towards participation in and support to COVAX.

2. An update on the joint procurement of vaccination supplies was provided in writing following questions raised during the meeting.