### FOOD AND DRUG ADMINISTRATION (FDA)

# Center for Biologics Evaluation and Research (CBER) 165<sup>th</sup> Meeting of the Vaccines and Related Biological Products Advisory Committee Silver Spring, MD March 5, 2021 AGENDA

**Topic:** The Committee will meet in open session to discuss and make recommendations on the selection of strains to be included in the influenza virus vaccines for the 2021 - 2022 influenza season.

Time	Presentation/Presenter
9:00am – 9:10am	Opening Remarks: Call to Order, Introduction of Committee
	Hana El Sahly, M.D., Chair, VRBPAC
	Administrative Announcements, Roll Call, Introduction of Committee,
9:10am – 9:30am	Conflict of Interest Statement
	Kathleen Hayes, M.P.H.
	Designated Federal Officer, VRBPAC
	CBER, FDA
9:30am – 9:45am	Influenza Virus Vaccine Strain Selection 2021-2022 Northern
	Hemisphere
	Jerry Weir, Ph.D. Director
	Division of Viral Products (DVP)
	Office of Vaccines Research and Review (OVRR)
	CBER, FDA
	Q &A: 5 min
9:45am – 10:10am	<u>U.S. Influenza Surveillance</u>
	Lisa Grohskopf, M.D., M.P.H.
	CAPT USPHS
	Associate Chief for Policy & Liaison
	Activities, Epidemiology & Prevention Branch, Influenza Division
	Centers for Disease Control and Prevention (CDC)
	Q &A: 5 min
10:10am – 11:20am	Global Influenza Virus Surveillance and Characterization
10.10am - 11.20am	David Wentworth, Ph.D.
	Branch Chief
	Influenza Division
	Virology, Surveillance, and Diagnostic Branch
	Centers for Disease Control and Prevention (CDC)
	Q &A: 10 min

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	BREAK
11:30am – 11:55am	<b>DoD Influenza Surveillance and Mid-Season Vaccine Effectiveness</b>
	LTC Kevin Taylor, M.D., MTM&H
	Focus Area Chief
	Global Emerging Infections Surveillance Branch
	Armed Forces Health Surveillance Division
	Public Health Division - DHA
	Kathleen Creppage, DrPh., M.P.H
	Senior Epidemiologist
	Armed Forces Health Surveillance Division
	Global Emerging Infections Surveillance Branch
	Q &A: 5 min
11:55am – 12:20pm	Candidate Vaccine Strains & Potency Reagents
	Manju Joshi, Ph.D.
	Lead Biologist
	Division of Biological Standards & Quality
	Office of Compliance and Biologics Quality
	CBER/FDA
	Q &A: 5 min
12:20pm – 12:45pm	Comments from Manufacturer Representative
	Lauren Parker, Ph.D.
	AstraZeneca
	O & A · 5 min
12:45pm – 1:30pm	Q &A: 5 min
12.45pm – 1.50pm	Lunch (45 min.)
1:30pm – 2:30pm	Open Public Hearing (60 min.)
2:30pm - 3:30pm	Committee Discussion, Recommendations, and Vote
	Committee of the Commit
3:30 pm	Adjourn Meeting
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