

Our Reference: EUA 27205 **EUA AMENDMENT – CONCURRENCE**

September 08, 2021

Janssen Biotech, Inc. Attention: Ms. Ruta Walawalkar 920 Route 202 Raritan, NJ 08869

Dear Ms. Walawalkar,

Please refer to your Emergency Use Authorization (EUA) for emergency use of Janssen COVID 19 Vaccine, re-issued on June 10, 2021, under section 564 of the Federal Food, Drug, and Cosmetic Act (FDCA) (21 U.S.C. 360bbb-3).

We also refer you to your EUA amendments:

submitted and received on March 5, 2021 submitted and received on March 29, 2021 submitted and received on April 12, 2021 submitted and received on August 13, 2021 submitted and received on August 25, 2021

Based on our review of the available data and information, we have determined that the Janssen's AD26.COV2.S DS Area 1 batch 21004323 (GMP 2) is suitable for use and meets the EUA standard, which is outlined in your Letter of Authorization. Thus, we concur with your request to add this batch to the EUA. Because the Bayview facility was not operating in compliance with Current Good Manufacturing Practice requirements at the time this batch was manufactured, through this concurrence letter, I am waiving Current Good Manufacturing Practice requirements for this batch, and only this batch, for the duration of this EUA. This concurrence does not add any other batches manufactured at this facility to the EUA at this time and does not add the facility itself to the EUA at this time. Nor does the EUA cover vaccine manufactured by combining this batch with different batches of drug substance that are not authorized explicitly under this EUA.

We remind you that any changes that you plan to implement to the description of the product, manufacturing process, facilities, or equipment will need to be submitted as an amendment to the EUA and not implemented without concurrence by the Agency.

Page 2 – EUA 27205 – Ruta Walawalkar

If you have any questions, please contact the Regulatory Project Manager, Sudhakar Agnihothram, PhD at 202-870-6949.

Sincerely,

--/S/--

Marion Gruber, PhD Director Office of Vaccines Research and Review Center for Biologics Evaluation and Research