



U.S. FOOD & DRUG
ADMINISTRATION

DATE: August 13, 2021

FROM: (b) (6)
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TO: (b) (6)
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SUBJECT: (b) (6) Discipline Review Memo
SPONSOR: BioNTech Manufacturing GmbH
PRODUCT: COVID-19 Vaccine, mRNA (COMIRNATY)
BLA: STN 125742/0

FINAL SUMMARY STATEMENT

(b) (6) inspection assignments were issued for a total of nine (9) clinical study sites that participated in the conduct of Study Protocol C4591001. Three (3) of these inspection assignments focused on clinical study sites that enrolled the pediatric population and six (6) of the study sites enrolled the adult population. The inspections did not reveal findings that impact the Biologics License Application (BLA).

BACKGROUND

On February 4, 2020, the Secretary of the Department of Health and Human Services (HHS) determined that there is a public health emergency (PHE) involving to a novel coronavirus named SARS-CoV-2 that causes Coronavirus Disease 2019 (COVID-19). On March 27, 2020, the Secretary of HHS issued a Notice of Emergency Use Authorization (EUA) Declaration pursuant to Section 564 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act or the Act).

In response to the PHE, (b) (6) reviewers proactively performed a review of the sponsor's investigational new drug application (IND 19736) and issued the necessary (b) (6) inspections to review the study conduct of Protocol C4591001, "A Phase 1/2/3 Study to Evaluate the Safety, Tolerability, Immunogenicity, and Efficacy of SARS-COV-2 RNA Vaccine Candidates Against COVID-19 in Healthy Individuals."

Protocol C4591001 was a multi-center study conducted at a total of 153 clinical sites: 131 study sites in the United States and 22 sites outside of the United States. Due to the COVID-19 pandemic travel restrictions, only the domestic sites were considered for an on-site (b) (6) inspection. Initially, six (6) study sites were inspected, before FDA issued the original Emergency Use Authorization for individuals 16 years of age and older. Subsequently, three (3) additional sites were inspected before FDA authorized use of the vaccine in those 12 and older. All of the study sites were selected based on subject enrollment, previous inspectional history, and other information submitted in IND 19736.

The inspections were conducted in accordance with FDA's Compliance Program 7348.811, Inspection Program for Clinical Investigators, focusing primarily on the study conduct, human subject protection and compliance with related FDA regulations. The data integrity and verification portion of the (b) (6) inspections were limited because the study was ongoing, and the data required for verification and comparison were not yet available to the IND. The table below summarizes the domestic study site information and the outcome of each (b) (6) inspection:

Site ID	Site Location	Form FDA 483 Issued	Final Classification
1007	Cincinnati Children's Hospital Medical Center Cincinnati Center for Clinical Research Cincinnati, OH	No	No Action Indicated (NAI)
1009	J. Lewis Research Inc./ Foothill Family Clinic South, Salt Lake City, UT	No	NAI
1044	Virginia Research Center, LLC. Midlothian, VA	No	NAI
1056	Indago Research and Health Center, Inc. Hialeah, FL	No	NAI
1109	Deland Clinical Research Unit DeLand, FL	No	NAI
1118	Meridian Clinical Research, LLC. Binghamton, NY	No	NAI
1125	Meridian Clinical Research, LLC Norfolk, NE	No	NAI
1133	Research Centers of America Hollywood, FL	No	NAI
1149	Collaborative Neuroscience Research, LLC at two locations: Long Beach & Garden Grove, CA	No	NAI

SIGNIFICANT INSPECTIONAL FINDINGS

No significant inspectional findings were noted.

SPONSOR/MONITORING ISSUES

No significant sponsor or monitoring issues were noted at the sites that were inspected.

FINANCIAL DISCLOSURE

The Clinical Investigator Compliance Program directs the FDA investigator to ask the clinical investigator if and when he/she disclosed information about his/her financial interests to the sponsor and/or interests of any sub-investigators, spouse(s) and dependent children, and if and when the information was updated. The information submitted to the BLA was verified for each of the inspected clinical sites.

ADMINISTRATIVE FOLLOW-UP

Should you have any questions or comments about the contents of this memo or any aspect of (b) (6) , please contact me at (b) (6) .

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