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**U.S. FOOD & DRUG
ADMINISTRATION**

DATE November 17, 2021

FROM (b) (6)

THROUGH (b) (6)

THROUGH (b) (6)

THROUGH (b) (6)

TO (b) (6), STN 125752/0
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SUBJECT (b) (6) Final Review Memo

SPONSOR ModernaTX, Inc.

PRODUCT mRNA-1273 COVID-19 vaccine (SPIKEVAX)

BLA STN 125752/0

FINAL SUMMARY STATEMENT

One (b) (6) inspection was conducted at a domestic Clinical Investigator (CI) site participating in the conduct of study protocol mRNA-1273-P301. The inspection did not reveal problems impacting the data submitted in support of this Biologics License Application (BLA).

Background

One CI study site was inspected in support of this BLA. The inspection was conducted in accordance with FDA's Compliance Program (CP) 7348.811, Inspection Program for Clinical Investigators. The inspection assignment was issued for the following study protocol, which included Parts A and B:

mRNA-1273-P301: A Phase 3, Randomized, Stratified, Observer-Blind, Placebo-Controlled Study to Evaluate the Efficacy, Safety, and Immunogenicity of mRNA-1273 SARS-CoV-2 Vaccine in Adults Aged 18 Years and Older

The site was selected based on previous inspectional history, geographic location, and the data submitted in this BLA. The inspection assignment included specific questions concerning the study protocol and requested the investigators to compare source documents at the site with information submitted in this BLA. The domestic CI site inspected in support of this BLA covered 3% of the subjects enrolled in the study inclusive

of Parts A and B. Study mRNA-1273-P301 was conducted at 99 study centers enrolling a total of 30,346 subjects aged 18 years and older in Part A; of these subjects that participated in Part A, a total of 28,964 subjects started Part B.

Inspection Outcome

Site ID	Number of subjects enrolled	Location	Form FDA 483 issued	Final Inspection Classification
319	928	Quality of Life Medical & Research Center, LLC Tucson, Arizona	No	No Action Indicated

The inspection verified the data reported in the BLA, including but not limited to subject's eligibility, protocol deviations, study drug administration, immunogenicity data, safety and reactogenicity events, concomitant medication administration, and adverse events for the randomly and equitably selected subjects enrolled at the inspected clinical site. The inspection further evaluated the adequacy of the study and site monitoring by the sponsor. No Form FDA 483 was issued at the conclusion of the inspection.

Noteworthy inspectional findings

None.

Sponsor Issues

No significant sponsor issues were noted.

Financial Disclosure

The Clinical Investigator Compliance Program directs the FDA investigators 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, spouses and dependent children, and if and when the information was updated. The information submitted to the BLA was verified at the inspected clinical site and found no deviations in the submitted data.

Administrative follow-up

No administrative follow-up is warranted at this time from (b) (6) for the inspected clinical investigator. Should you have any questions about the contents of this memo or any aspect of (b) (6) please contact me at (b) (6).

(b) (6)