CLINICAL STUDY REPORT ERRATA

Protocol Number: C4591001

Study Report Title:	Interim Report – 6 Month Update: A Phase 1/2/3, Placebo-Controlled, Randomized, Observer-Blind, Dose- Finding Study to Evaluate the Safety, Tolerability, Immunogenicity, and Efficacy of SARS-COV-2 RNA Vaccine Candidates Against COVID-19 in Healthy Individuals

Minor errors were identified in the Clinical Study Report. Those errors are listed below.

Although these errors do not affect the conclusions of the study, they have been included in this Errata to ensure the accuracy of the report.

Section	Description of Error
Sections 10.1.2, 12.2.3, 12.2.4, and 14 Section 16.2.7 Narratives (contain corrected information) Appendix 16.2.7.4.1 Listing of Adverse Events – All Subjects ≥16 Years of Age Appendix 16.2.7.5 Listing of Serious Adverse Events – All Subjects ≥16 Years of Age[a] Appendix 16.2.7.6 Listing of Adverse Events Leading to Discontinuation – All Subjects ≥16 Years of Age	After the data cutoff date of 13 Mar 2021 (with database snapshot taken on 25 Mar 2021), it was determined that: • Participants C4591001 1044 10441163 and C4591001 1090 10901140 did not withdraw from the study because of the injection site pain and remain in the study • Participants C4591001 1090 10901492 and C4591001 1091 10911297 did not withdraw from the study because of the fatigue and remain in the study • Participant C4591001 1091 10911247 did not withdraw from the study because of the dermatitis and remains in the study • Participant C4591001 1092 10921021 did not withdraw from the study because of the pain and remains in the study • Participant C4591001 1254 12541006 did not withdraw from the study because of the chills and remains in the study • Participant C4591001 1254 12541189 did not withdraw from the study because of the headache and remains in the study • Participant C4591001 1096 10961036 did not discontinue from the study intervention because of the suicidal depression but discontinued from the study because she no longer met the eligibility criteria

	• Participant C4591001 1232 12321299 was not withdrawn from the study for safety reasons (hypertension) but was lost to follow-up • Participant C4591001 1037 10371141 was not pregnant and remains in the study

a. Participant C4591001 1096 10961036 only.