16.1.5.1 SPONSOR

CLINICAL STUDY REPORT APPROVAL FORM

| PROTOCOL NUMBER: | C4591001 |
|-------------------------|--|
| REPORT TITLE: | Final Analysis Interim Report: 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 |
| AUTHOR: | Dzung Nguyen |

I have reviewed this clinical study report, which is from its definitive source and final in content. I confirm that it is a complete and accurate representation of the relevant data and statistical analyses for this report.

Known deviations in the study did not impact overall compliance with Good Clinical Practice and local regulations.

Organization/Institution: BioNTech

Name/Position: Ugur Sahin, MD, Chief Executive Officer

Signature:

Date:_ 7. 11. 2020

16.1.5.1 SPONSOR AGENT

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STATISTICS LINE HEAD: Kenneth Koury, PhD

CLINICAL LEAD: John L. Perez, MD, MBA, MA

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