

Gdansk, 27th January 2022

To whom it may concern,

The GeneMe considering the latest information on the mutation of the SARS-CoV-2 genome and its influence on the results of RT-PCR assays. We hereby present our Research and Development report the objective of which was to determine if the recently observed mutations in the SARS-CoV-2 affect the loss of specificity of the SAVD RT-PCR test. This research and development report summarizes our findings regarding the influence of ORF1ab-gene mutations in SARS-CoV-2 on SAVD primers hybridization. For this purpose, bioinformatic analysis of the mutated sequences of the coronavirus ORF1ab-gene was performed to assess if the point mutation or deletions lay in the hybridization region of SAVD RT-PCR primers.

The SAVD by GeneMe SARS-CoV-2 Direct Rapid Detection Kit is designed for the *in vitro* identification of the new coronavirus SARS-CoV-2, in a single reaction. The presence of an innovative and patented *Pwo* polymerase and specific primers in the kit has enabled the creation of a highly specific and sensitive SARS-CoV-2 rapid detection kit. The specifically designed primers are 100% compatible with the SARS-CoV-2 genomic RNA sequence of gene ORF1ab recommended by WHO and deposited in the NCBI database. Amplification of the targeted nucleic acids is observed by an increase of fluorescence signal during the reaction.

We, signed below, can definitively state that SAVD RT-PCR assay's ability to detect SARS-CoV-2 remains at the highest level regardless of these new mutations.

The GeneMe constantly cooperates with diagnostic laboratories in Poland and abroad (UK, Norway, Uganda, Mexico), regularly validating the test on clinical trials - swabs and saliva samples to make sure that the SAVD test maintains its sensitivity and specificity to the current SARS-Co-2 virus strains.

Yours sincerely,

Dr Sabina Żołędowska,
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Research and Development

NAME: *Marta Skwarecka, Head of RD GeneMe*

DATE: *27.01.2022*

PROPOSED PRODUCT: *SAVD*

REVISION: *2.0*

1. Title

Detection of different variants of SARS-CoV-2 virus by SAVD test.

2. Purpose and scope

The aim of the study is to check the universality of the SAVD test for the identification of known variants of the SARS-CoV-2 virus. The most popular variants of the mutant SARS-CoV-2 virus from Alfa to Omicron were analyzed in silico. The most popular of them are: B.1.1.7 United Kingdom, B.1.351 South Africa (also known as S.501Y.V2), B.1.1.28 Brazil P1, P2, B .1.617 India, B.1.429/ B.1.427 California variant (also known as epsilon variant), Vietnamese variant (Delta variant B.1.617.2 with additional mutations) and Omicron variant B.1.1.529.

3. Method

| | | | |
|--|---|-------------------|--------------------|
| Date of the test: | <i>27.01.2022</i> | | |
| Place of the test: | <i>GeneMe, ul. Kampinoska 25, 80-180 Gdansk, Poland</i> | | |
| Test conditions (temperature, humidity): | <i>Temp: -- Humidity: --</i> | | |
| The person performing the tests: | <i>Marta Skwarecka</i> | | |
| LOT of reagents analyzed: | LOT number | Trade name | Expiry date |
| | - | | |
| | - | | |
| | - | | |
| LOT of reference reagents and trade name: | LOT number | Trade name | Expiry date |
| | - | | |
| | - | | |
| | - | | |

Description of the tested method:

The study consisted of:

- 1. Finding in the analyzed variants of the SARS-CoV-2 virus the resulting mutations in the ORF1ab gene relative to the native strain and locating them in the genomic RNA of the virus.*
- 2. Assigning individual mutations to appropriate nucleotides.*

3. Comparison of the location of the mutated nucleotides with the location of the ORF1ab gene fragment, which is the target of the SAVD test.
4. Confirmation or exclusion of the effect of the mutation on the SAVD test identification capabilities.

4. Tested samples

| Sample number | Name | Supplier | Producer (as commercial material) | Concentration (as commercial material) |
|---------------|------|----------|-----------------------------------|--|
| 1. | n/a | n/a | n/a | n/a |
| 2. | n/a | n/a | n/a | n/a |

5. Results

Table 1 shows the popular variants of the SARS-CoV-2 virus along with the changed nucleotides and compared with the target sequence of the SAVD test.

Table 1. Mutations in the ORF1ab gene of popular variants of the SARS-CoV-2 virus and their impact on the possibility of identification with the SAVD test.

| Virus variant | Country of origin (emergence) | Amino-acid mutation | Nucleotide mutation | Location of mismatch (5'-3') in the SAVD Plus test | Detection with the SAVD test |
|------------------------------------|-------------------------------|------------------------|----------------------|--|------------------------------|
| Reference Strain: Wuhan-Hu-1, nCoV | China | - | - | absence | Yes |
| Alpha (B.1.1.7) | UK | T1001I | C3267T | absence | Yes |
| | | A1708D | C5388A | absence | |
| | | I2230T | T6954C | absence | |
| | | SGF 3675-3677 deletion | 11288-11296 deletion | absence | |
| Beta (B.1.351,S.501Y.V2) | South Africa | Thr265I | C1059T | absence | Yes |
| | | L1655Asn | G5230T | absence | |
| | | L3353R | A10323AG | absence | |
| Gamma | Brazil | synonymous mutation | T733C | absence | Yes |

| | | | | | |
|--------------------------------|-------------------|--------------------------------|-------------------------|----------------|------------|
| <i>(B.1.1.28.1, P1)</i> | | <i>synonymous mutation</i> | <i>C2749T</i> | <i>absence</i> | |
| | | <i>S1188L</i> | <i>C3828T</i> | <i>absence</i> | |
| | | <i>L1795Q</i> | <i>A5648C</i> | <i>absence</i> | |
| | | <i>synonymous mutation</i> | <i>A6319G</i> | <i>absence</i> | |
| | | <i>synonymous mutation</i> | <i>A6613G</i> | <i>absence</i> | |
| | | <i>synonymous mutation</i> | <i>C12778T</i> | <i>absence</i> | |
| | | <i>synonymous mutation</i> | <i>C13860T</i> | <i>absence</i> | |
| | | <i>E1264N</i> | <i>G17259T</i> | <i>absence</i> | |
| | | <i>synonymous mutation</i> | <i>C100T</i> | <i>absence</i> | |
| | | <i>L3468V</i> | <i>T10667G</i> | <i>absence</i> | |
| | | <i>synonymous mutation</i> | <i>C11824T</i> | <i>absence</i> | |
| | <i>L3930F</i> | <i>C12053T</i> | <i>absence</i> | | |
| <i>Delta (B.1.617.2)</i> | <i>India</i> | <i>P4715L</i> | <i>14408- 14410</i> | <i>absence</i> | <i>Yes</i> |
| | | <i>P5401L</i> | <i>16466- 16468</i> | <i>absence</i> | |
| | | <i>G5063S</i> | <i>20515- 20517</i> | <i>absence</i> | |
| <i>Delta+ (B.1.617.2+)</i> | <i>Vietnamese</i> | <i>synonymous mutation</i> | <i>C3037T</i> | <i>absence</i> | <i>Yes</i> |
| | | <i>synonymous mutation</i> | <i>C3457T</i> | <i>absence</i> | |
| | | <i>T1567I</i> | <i>C4965T</i> | <i>absence</i> | |
| | | <i>synonymous mutation</i> | <i>G8491A</i> | <i>absence</i> | |
| | | <i>T3646A</i> | <i>A11201G</i> | <i>absence</i> | |

| | | | | | |
|---------------------------------|---------------------|-------------------------------|-----------------------------|----------------|------------|
| | | <i>P4715L</i> | <i>C14408T</i> | <i>absence</i> | |
| | | <i>synonymous mutation</i> | <i>G14772A</i> | <i>absence</i> | |
| | | <i>synonymous mutation</i> | <i>C16134T</i> | <i>absence</i> | |
| | | <i>G5530C</i> | <i>G16852T</i> | <i>absence</i> | |
| | | <i>M5753I</i> | <i>G17523T</i> | <i>absence</i> | |
| | | <i>L6711R</i> | <i>A20396G</i> | <i>absence</i> | |
| | | <i>S6713A</i> | <i>T20401G</i> | <i>absence</i> | |
| | | <i>T1001I</i> | <i>C3267T</i> | <i>absence</i> | |
| | | <i>A1708D</i> | <i>C5388A</i> | <i>absence</i> | |
| | | <i>I2230T</i> | <i>T6954C</i> | <i>absence</i> | |
| | | <i>SGF 3675-3677 deletion</i> | <i>11287-11295 deletion</i> | <i>absence</i> | |
| | | <i>P4715L</i> | <i>14407-14409</i> | <i>absence</i> | |
| | | <i>P5401L</i> | <i>16465-16467</i> | <i>absence</i> | |
| | | <i>G5063S</i> | <i>15451-15453</i> | <i>absence</i> | |
| <i>Omicron BA.1 (B.1.1.529)</i> | <i>South Africa</i> | <i>K856R</i> | <i>2830-2832</i> | <i>absence</i> | <i>Yes</i> |
| | | <i>SL2083-2084I</i> | <i>6511-6516</i> | <i>absence</i> | |
| | | <i>A2710T</i> | <i>8392-8394</i> | <i>absence</i> | |
| | | <i>T3255I</i> | <i>10027-10029</i> | <i>absence</i> | |
| | | <i>P3395H</i> | <i>10447-10449</i> | <i>absence</i> | |
| | | <i>3674-3676 deletion</i> | <i>33056-11292</i> | <i>absence</i> | |
| | | <i>I3758V</i> | <i>11536-11538</i> | <i>absence</i> | |

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|-----------------------------------|-------------------------|----------------------------|--------------------|----------------|------------|
| | | <i>P4715L</i> | <i>14407-14409</i> | <i>absence</i> | |
| | | <i>I5967V</i> | <i>18163-18165</i> | <i>absence</i> | |
| <i>Omicron BA.2 (BA.2)</i> | <i>South Africa</i> | - | - | <i>absence</i> | <i>Yes</i> |
| <i>Epsilon (B.1.429, B.1.427)</i> | <i>California (USA)</i> | <i>T265I</i> | <i>1057-1059</i> | <i>absence</i> | <i>Yes</i> |
| | | <i>S3158T</i> | <i>9736-9738</i> | <i>absence</i> | |
| | | <i>I4205V</i> | <i>12877-1279</i> | <i>absence</i> | |
| | | <i>P314L</i> | <i>1204-1206</i> | <i>absence</i> | |
| | | <i>P976L</i> | <i>3190-3192</i> | <i>absence</i> | |
| | | <i>D1183T</i> | <i>3811-3813</i> | <i>absence</i> | |
| <i>Zeta (B.1.1.28.2, P2)</i> | <i>Brazil</i> | - | - | <i>absence</i> | <i>Yes</i> |
| <i>Eta (B.1.525)</i> | <i>Worldwide</i> | <i>L4715F</i> | <i>14407-14409</i> | <i>absence</i> | <i>Yes</i> |
| <i>Theta (B.1.1.28.3, P3)</i> | <i>Philippines</i> | <i>L3201P</i> | <i>9865-9867</i> | <i>absence</i> | <i>Yes</i> |
| | | <i>D3681E</i> | <i>11305-11307</i> | <i>absence</i> | |
| | | <i>L3930F</i> | <i>12052-12054</i> | <i>absence</i> | |
| | | <i>P4715L</i> | <i>14407-14409</i> | <i>absence</i> | |
| <i>Iota (B.1.526)</i> | <i>USA</i> | <i>del3675-3677</i> | <i>11287-11295</i> | <i>absence</i> | <i>Yes</i> |
| <i>Kappa (B.1.617.1)</i> | <i>India</i> | <i>synonymous mutation</i> | <i>C3037T</i> | <i>absence</i> | <i>Yes</i> |
| | | <i>synonymous mutation</i> | <i>C3457T</i> | <i>absence</i> | |
| | | <i>T1567I</i> | <i>C4965T</i> | <i>absence</i> | |

| | | | | | |
|--------------------------------|-------------|----------------------------|---------|----------------|-----|
| | | <i>synonymous mutation</i> | G8491A | <i>absence</i> | |
| | | T3646A | A11201G | <i>absence</i> | |
| | | P4715L | C14408T | <i>absence</i> | |
| | | <i>synonymous mutation</i> | G14772A | <i>absence</i> | |
| | | <i>synonymous mutation</i> | C16134T | <i>absence</i> | |
| | | G5530C | G16852T | <i>absence</i> | |
| | | M5753I | G17523T | <i>absence</i> | |
| | | K6711R | A20396G | <i>absence</i> | |
| | | S6713A | T20401G | <i>absence</i> | |
| <i>Lambda</i> (B.1.1.1.C37) | <i>Peru</i> | - | - | <i>absence</i> | Yes |

Link to the data repository kept in the cloud: --

6. Conclusions

The presented analysis shows that none of the mutations occurring in the variants of the SARS-CoV-2 virus, i.e., Alpha, Beta, Gamma, Delta, Omicron, Epsilon, Zeta, Theta, Iota, Kappa and Lambda did affect the effectiveness of the virus detection with the SAVD test. All analyzed variants are fully identifiable with the SAVD test.

7. References

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Approved for external release by Sabina Żołędowska, CQO

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Signature: 