

Gdansk, 1<sup>st</sup> February 2022

To whom it may concern,

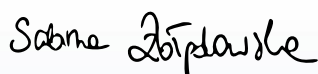
The GeneMe considering the latest information on the mutation of the SARS-CoV-2 genome and its influence on the results of RT-PCR assays. We hereby present our Research and Development report the objective of which was to determine if the recently observed mutations in the SARS-CoV-2 affect the loss of specificity of the SAVD+ RT-PCR test. This research and development report summarizes our findings regarding the influence of ORF1ab and N1 gene mutations in SARS-CoV-2 on SAVD+ primers hybridization. For this purpose, bioinformatic analysis of the mutated sequences of the coronavirus ORF1ab and the human N1 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 the innovative and patented *Taq* polymerase and specific primers and probes made it possible to create a highly specific and sensitive test. The individually designed primers are compatible with the SARS-CoV-2 genomic RNA sequence of the ORF1ab and N gene as deposited in the NCBI database and human RNase P gene. 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 Żółędowska,  
CQO



Dr Eng. Marta Skwarecka,  
Head of Research and Development Department



## Research and Development

**NAME:** *Marta Skwarecka, Head of RnD GeneMe*

**DATE:** *23.01.2022*

**PROPOSED PRODUCT:** *SAVD+*

**REVISION:** *1.0*

### 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

<b>Date of the test:</b>	--
<b>Place of the test:</b>	<i>GeneMe, ul. Kampinoska 25, 80-180 Gdansk, Poland</i>
<b>Test conditions (temperature, humidity):</b>	<i>Temp: 22°C Humidity: 36%</i>
<b>The person performing the tests:</b>	<i>Dr Eng. Marta Skwarecka</i>
<b>LOT of reagents analyzed:</b>	--
<b>LOT of reference reagents and trade name:</b>	--

### 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

#### 5. Results

Table 1 and 2 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+ test	Detection with the SAVD+ test
Reference Strain: Wuhan-Hu-1, nCoV	China	-	-	absence	Yes
B.1.617	India	synonymous mutation	C3037T	absence	Yes
		synonymous mutation	C3457T	absence	
		T1567I	C4965T	absence	
		synonymous mutation	G8491A	absence	
		T3646A	A11201G	absence	
		P4715L	C14408T	absence	

		<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>K6711R</i>	<i>A20396G</i>	<i>absence</i>	
		<i>S6713A</i>	<i>T20401G</i>	<i>absence</i>	
<i>B.1.1.28 P1, P2</i>	<i>Brazil</i>	<i>synonymous mutation</i>	<i>T733C</i>	<i>absence</i>	<i>Yes</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>B.1.351 (S.501Y.V2)</i>	<i>South Africa</i>	<i>Thr265I</i>	<i>C1059T</i>	<i>absence</i>	<i>Yes</i>
		<i>L1655Asn</i>	<i>G5230T</i>	<i>absence</i>	
		<i>L3353R</i>	<i>A10323AG</i>	<i>absence</i>	

<i>B.1.1.7</i>	<i>UK</i>	<i>T1001I</i>	<i>C3267T</i>	<i>absence</i>	<i>Yes</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>11288-11296 deletion</i>	<i>absence</i>	
<i>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>B.1.617.2+ (Delta+)</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>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>B.1.1.529 BA.1</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>	
		<i>P4715L</i>	<i>14407-14409</i>	<i>absence</i>	
		<i>I5967V</i>	<i>18163-18165</i>	<i>absence</i>	
<i>B.1.1.529 BA.2</i>	<i>South Africa</i>	-	-	<i>absence</i>	<i>Yes</i>
<i>B.1.525</i>	<i>Worldwide</i>	<i>L4715F</i>	<i>14407-14409</i>	<i>absence</i>	<i>Yes</i>
<i>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>	12052-12054	<i>absence</i>	
		<i>P4715L</i>	14407-14409	<i>absence</i>	
<i>B.1.526</i>	<i>USA</i>	<i>del3675-3677</i>	11287-11295	<i>absence</i>	<i>Yes</i>
<i>B.1.1.1.C37</i>	<i>Peru</i>	-	-	<i>absence</i>	<i>Yes</i>

Table 2. Mutations in the N1 gene of popular variants of the SARS-CoV-2 virus and their impact on the possibility of identification with the SAVD+ test.

<b>Virus variant</b>	<b>Country of origin (emergence)</b>	<b>Amino-acid mutation</b>	<b>Nucleotide mutation</b>	<b>Location of mismatch (5'-3') in the SAVD+ test</b>	<b>Detection with the SAVD+ test</b>
<i>Reference Strain: Wuhan-Hu-1, nCoV</i>	<i>China</i>	-	-	<i>absence</i>	<i>Yes</i>
<i>B.1.617</i>	<i>India</i>	-	-	<i>absence</i>	<i>Yes</i>
<i>B.1.1.28 P1, P2</i>	<i>Brazil</i>	<i>P80R</i>	28513-28515	<i>absence</i>	<i>Yes</i>
		<i>R203K</i>	28882-28884	<i>absence</i>	
		<i>G204R</i>	28885-28887	<i>absence</i>	
<i>B.1.351 (S.501Y.V2)</i>	<i>South Africa</i>	<i>T205I</i>	28978-28980	<i>absence</i>	<i>Yes</i>
<i>B.1.1.7</i>	<i>UK</i>	<i>D3L</i>	28282-28284	<i>absence</i>	<i>Yes</i>
		<i>S235F</i>	28978-28980	<i>absence</i>	
<i>B.1.617.2</i>	<i>India</i>	<i>D63G</i>	28462-28464	<i>absence</i>	<i>Yes</i>

		<i>R203M</i>	28882-28884	<i>absence</i>	
		<i>D377Y</i>	29404-29406	<i>absence</i>	
<i>B.1.617.2+ (Delta+)</i>	<i>Vietnamese</i>	<i>D63G</i>	28462-28464	<i>absence</i>	Yes
		<i>R203M</i>	28882-28884	<i>absence</i>	
		<i>D377Y</i>	29404-29406	<i>absence</i>	
<i>B.1.429, B.1.427</i>	<i>California (USA)</i>	<i>T205I</i>	28888-28890	<i>absence</i>	Yes
<i>B.1.1.529 BA.1</i>	<i>South Africa</i>	<i>31-33del</i>	28366-28374	<i>absence</i>	Yes
		<i>R203K</i>	28882-28884	<i>absence</i>	
		<i>G204R</i>	28885-28887	<i>absence</i>	
<i>B.1.1.529 BA.2</i>	<i>South Africa</i>	-	-	<i>absence</i>	Yes
<i>B.1.525</i>	<i>Worldwide</i>	<i>L4715F</i>	14407-14409	<i>absence</i>	Yes
<i>B.1.1.28.3, P3</i>	<i>Philippines</i>	<i>R203K</i>	28882-28884	<i>absence</i>	Yes
		<i>G204R</i>	28885-28887	<i>absence</i>	
<i>B.1.526</i>	<i>USA</i>	-	-	<i>absence</i>	Yes
<i>B.1.1.1.C37</i>	<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( B.1.1.7), Beta (B.1.351, S.501Y.V2), Gamma (B.1.1.28.1, P1), Delta (B.1.617.2), Delta+ (B.1.617.2+), Omicron BA.1 (B.1.1.529), Omicron BA.2 (BA.2), Epsilon (B.1.429, B.1.427), Zeta (B.1.1.28.2, P2), Eta (B.1.525), Theta (B.1.1.28.3, P3), Iota (B.1.526), Kappa (B.1.617.1), Lambda (B.1.1.1.C37) 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

Date of approval: 01.02.2022

Signature 