

Calculation of B.1.1.7 (North American) substitutions

The constellation of substitutions carried by the B.1.1.7 lineage in comparison with Wuhan-Hu-1 were computed for isolates from North America using sequence records derived from Genbank so that the sequences can be released through ViPR and BV-BRC and so that they are relevant to circulation in the U.S, as follows:

1. Generate an S protein working set to use as a BLAST/Short Peptide Search database using query: SARS-related species, complete genome, S protein, sinceJUN2020, human, North America on 25JAN2021.
2. Use Short Peptide Search for GFQPTYGVGYQ to identify S proteins from North America since July 2020 carrying the N501Y substitution => 11 proteins
3. For quality control purposes, removed strains in which S protein sequences had any “X” ambiguous positions => 8 proteins from the following strains:
 - a. SARS-CoV-2/human/USA/SEARCH-5574-SAN/2020, SARS-CoV-2/human/USA/CO-CDPHE-2100156850/2020, SARS-CoV-2/human/USA/CA-CDC-STM-P017/2020, SARS-CoV-2/human/USA/FL-BPHL-2779/2021, SARS-CoV-2/human/USA/FL-CDC-STM-P012/2020, SARS-CoV-2/human/USA/NY-Wadsworth-291673-01/2020, SARS-CoV-2/human/USA/NMDOH-2021003333/2021, SARS-CoV-2/human/USA/FL-BPHL-2776/2021
4. Use Quick Search to pull out corresponding whole genome sequences.
5. Use SNP tool to calculate consensus genome sequence => “B.1.1.7_North America_genome_consensus.fasta”
6. Combine with “Wuhan-Hu-1_genome.fasta” with “B.1.1.7_North America_genome_consensus.fasta” => “Wuhan+B.1.1.7(QC)_North America_genome_consensus.fasta”
7. Run SNP on “Wuhan+B.1.1.7(QC)_North America_genome_consensus.fasta” to generate SNP report of all consensus substitutions in comparison with Wuhan-Hu-1 => 48 SNPs, including 19 deletions in “Wuhan-Hu-1 + B.1.1.7_North America_genome_consensus_SNP.xlsx”
8. Annotate “B.1.1.7_North America_genome_consensus.fasta” with VIGOR4 => “B.1.1.7_North America_genome_consensus_VIGOR4”
9. Align “Wuhan-Hu-1 S protein.fasta” with “B.1.1.7_North America_S protein_consensus.fasta” to determine S protein substitutions => Spike: H69-, V70-, N501Y, A570D, D614G, P681H, T716I, S982A, D1118H
10. Determine which SNPs were located within which ORF gene or mature peptide genomic region through Wuhan mapping table. Combine relevant CAL.20C_genome_consensus_VIGOR4 protein with corresponding Wuhan-Hu-1 protein. Perform multiple sequence alignment and determine amino acid substitutions for all genomic SNPs => nsp3: T183I, A890D, I1412T; nsp6: S106-, G107-, F108-; RNA-dependent RNA polymerase: P323L; helicase: K460R; ORF8: Q27stop
11. BLAST consensus against “B.1.1.7(QC)_North America_genome” working set => best representative (Identities = 29847/29861 (99%)) SARS-CoV-2/human/USA/FL-CDC-STM-P012/2020 - <https://www.viprbrc.org/brc/viprStrainDetails.spg?ncbiAccession=MW430974&decorator=corona&context=1611876836946>

Calculation of B.1.351 substitutions

1. Since no sequences carrying the PGQTGTIADYN (K417T) short peptide were available in ViPR, downloaded sequences from GISAID using search criteria: B.1.351 lineage; 501Y.V2 substitution; Africa; complete; low coverage excluded => "gisaid_B.1.351_501Y.V2_Africa_complete.fasta" with 485 records
2. In order to reduce the number of records for SNP analysis, selected the first 79 records => "gisaid_B.1.351_501Y.V2_Africa_complete_subset.fasta"
3. Ran SNP analysis to generate "gisaid_B.1.351_501Y.V2_Africa_complete_subset_SNP_consensus.fasta" a.k.a "B.1.351_genome_consensus.fasta"
4. Combine "Wuhan-Hu-1_genome.fasta" and "gisaid_B.1.351_501Y.V2_Africa_complete_subset_SNP_consensus.fasta" => "Wuhan-Hu-1+B.1.351_consensus.fasta"
5. Run SNP on "Wuhan-Hu-1+B.1.351_consensus.fasta" to generate SNP report of all consensus substitutions in comparison with Wuhan-Hu-1 => 37 SNPs including 18 deletions in "Wuhan-Hu-1+B.1.351_consensus_SNP.xlsx"
6. Capture specific substitutions from the SNP MSA in "Wuhan-Hu-1 + B.1.351_consensus_genome_SNP.xlsx"
7. Annotate "gisaid_B.1.351_501Y.V2_Africa_complete_subset_SNP_consensus.fasta" with VIGOR4 => "B.1.375_North America_genome_consensus_VIGOR4"
8. Align "Wuhan-Hu-1 S protein.fasta" with "B.1.351_S protein_consensus.fasta" to determine S protein substitutions => Spike: D80A, D215G, L241-, L242-, A243-, K417N, E484K, N501Y, D614G, A701V
9. Determine which SNPs were located within which ORF gene or mature peptide genomic region through Wuhan mapping table. Combine relevant "B.1.375_North America_genome_consensus_VIGOR4" protein with corresponding Wuhan-Hu-1 protein. Perform multiple sequence alignment and determine amino acid substitutions for all genomic SNPs => nsp2: T85I; nsp3: K837N; 3C-like proteinase: K90R; nsp6: S106-, G107-, F108-; RNA-dependent RNA polymerase: P323L; ORF3a: Q57H; S171L; envelope protein: P71L; nucleocapsid phosphoprotein: T205I

Calculation of P.1 (North American) substitutions

The constellation of substitutions carried by the B.1.1.7 lineage were computed for the first US isolate (SARS-CoV-2/human/USA/MN-MDH-2399/2021) in comparison with Wuhan-Hu-1 using the sequence record derived from Genbank so that the sequence can be released through ViPR and BV-BRC and so that they are relevant to circulation in the U.S, as follows, as follows:

1. Use Quick Search to pull out SARS-CoV-2/human/USA/MN-MDH-2399/2021 whole genome sequence
=> "P.1_US_genome_consensus.fasta"
2. Combine with "Wuhan-Hu-1_genome.fasta" with "P.1_US_genome_consensus.fasta" =>
"Wuhan+P.1_US_genome_consensus.fasta"
3. Run SNP on "Wuhan+P.1_US_genome_consensus.fasta" to generate SNP report of all consensus substitutions in comparison with Wuhan-Hu-1 => 49 SNPs, including 9 deletions and 5 insertions in
"Wuhan-Hu-1 + P.1_first US_genome_SNP.xlsx"
4. Annotate "P.1_US_genome_consensus.fasta" with VIGOR4 =>
"P.1_US_genome_consensus_VIGOR4"
5. Align "Wuhan-Hu-1 S protein.fasta" with "P.1_US_S protein_consensus.fasta" to determine S protein substitutions => Spike: L18F, T20N, P26S, D138Y, R190S, K417T, E484K, N501Y, D614G, H655Y, T1027I, V1176F
6. Determine which SNPs were located within which ORF gene or mature peptide genomic region through Wuhan mapping table. Combine relevant P.1_US_genome_consensus_VIGOR4 protein with corresponding Wuhan-Hu-1 protein. Perform multiple sequence alignment and determine amino acid substitutions for all genomic SNPs => nsp3: S370L, K977Q; nsp4: S184N; 3C-like proteinase: A260V; nsp6: S106-, G107-, F108-; RNA-dependent RNA polymerase: P323L; helicase: E341D; ORF3a protein: S253P; ORF8 protein: E92K; nucleocapsid phosphoprotein: P80R, R203K, G204R
7. Representative sequence: SARS-CoV-2/human/USA/MN-MDH-2399/2021 -
<https://www.viprbrc.org/brc/viprStrainDetails.spg?ncbiAccession=MW520923&decorator=corona&context=1611876882126>

Calculation of CAL.20C substitutions

1. Generate an S protein working set to use as a BLAST/Short Peptide Search database using query: SARS-related species, complete genome, S protein, sinceJUN2020, human, North America on 25JAN2021.
2. Use Short Peptide Search for NYNYRYRLFR to identify S proteins from North America since July 2020 carrying the L452R substitution => 55 proteins
3. For quality control purposes, removed strains in which S protein sequences had any “X” ambiguous positions => 17 proteins from the following strains:
 - a. SARS-CoV-2/human/USA/CA-CZB-12872/2020, SARS-CoV-2/human/USA/NMDOH-2021013232/2021, SARS-CoV-2/human/USA/CA-LACPHL-AF00003/2020, SARS-CoV-2/human/USA/CA-LACPHL-AF00014/2020, SARS-CoV-2/human/USA/UT-UPHL-2012360496/2020, SARS-CoV-2/human/USA/CA-LACPHL-AF00041/2020, SARS-CoV-2/human/USA/CA-LACPHL-AF00074/2020, SARS-CoV-2/human/USA/CA-LACPHL-AF00077/2020, SARS-CoV-2/human/USA/CA-LACPHL-AF00169/2021, SARS-CoV-2/human/USA/CA-LACPHL-AF00141/2021, SARS-CoV-2/human/USA/CA-LACPHL-AF00055/2020, SARS-CoV-2/human/USA/CA-LACPHL-AF00030/2020, SARS-CoV-2/human/USA/CA-LACPHL-AF00031/2020, SARS-CoV-2/human/USA/WA-S3353/2020, SARS-CoV-2/human/USA/WA-S3355/2020, SARS-CoV-2/human/USA/CA-CDC-STM-A100056/2021, SARS-CoV-2/human/USA/KY-CDC-STM-A100059/2021
4. Use Quick Search to pull out corresponding whole genome sequences.
5. Use SNP tool to calculate consensus genome sequence => “CAL.20C_genome_consensus.fasta”
6. Combine with Wuhan-Hu-1_genome.fasta with CAL.20C_genome_consensus.fasta => Wuhan+CAL.20C consensus_genome_25JAN2021.fasta
7. Run SNP on “Wuhan+CAL.20C consensus_genome_25JAN2021.fasta” to generate SNP report of all consensus substitutions in comparison with Wuhan-Hu-1 => 21 SNPs in “Wuhan-Hu-1 + CAL.20C consensus_genome_SNP.xlsx”
8. Annotate “CAL.20C_genome_consensus.fasta” with VIGOR4 => “CAL.20C_genome_consensus_VIGOR4”
9. Align “Wuhan-Hu-1 S protein.fasta” with “CAL.20C_S protein_consensus.fasta” to determine S protein substitutions => Spike: S13I, W152C, L452R, D614G
10. Determine which SNPs were located within which ORF gene or mature peptide genomic region through Wuhan mapping table. Combine relevant CAL.20C_genome_consensus_VIGOR4 protein with corresponding Wuhan-Hu-1 protein. Perform multiple sequence alignment and determine amino acid substitutions for all genomic SNPs.
11. BLAST consensus against “genome_North America_CAL.20C_QC” working set => best representative (Identities = 29893/29901 (99%)) “SARS-CoV-2/human/USA/CA-LACPHL-AF00141/2021” - <https://www.viprbrc.org/brc/viprStrainDetails.spg?ncbiAccession=MW485882&decorator=corona&context=1611876924806>

Calculation of B.1.375 substitutions

1. Generate an S protein working set to use as a BLAST/Short Peptide Search database using query: SARS-related species, complete genome, S protein, since JUN2020, human, North America on 25JAN2021.
2. Use Short Peptide Search for NVTWFHAI SGTNGTKRFD to identify S proteins from North America since July 2020 carrying the 69/70 deletion => 49 proteins
3. Use Short Peptide Search for YGFQPTNGVG to identify S proteins from North America since July 2020 that do not also carry the N501Y substitution of B.1.1.7 => 38 proteins from the following strains:
 - a. SARS-CoV-2/human/USA/FL-BPHL-2259/2020, SARS-CoV-2/human/USA/GA-CDC-STM-A100055/2021, SARS-CoV-2/human/USA/UT-UPHL-2012039803/2020, SARS-CoV-2/human/USA/UT-UPHL-2012566145/2020, SARS-CoV-2/human/USA/UT-UPHL-2012558654/2020, SARS-CoV-2/human/USA/UT-UPHL-2012218768/2020, SARS-CoV-2/human/USA/UT-UPHL-2012875547/2020, SARS-CoV-2/human/USA/UT-UPHL-2012522014/2020, SARS-CoV-2/human/USA/UT-UPHL-2012983373/2020, SARS-CoV-2/human/USA/UT-UPHL-2012045106/2020, SARS-CoV-2/human/USA/UT-UPHL-2012663924/2020, SARS-CoV-2/human/USA/WI-UW-2622/2020, SARS-CoV-2/human/USA/WI-UW-2590/2021, SARS-CoV-2/human/USA/WI-UW-2607/2021, SARS-CoV-2/human/USA/WI-UW-2634/2020, SARS-CoV-2/human/USA/WI-UW-2645/2020, SARS-CoV-2/human/USA/FL-CDC-STM-P004/2020, SARS-CoV-2/human/USA/FL-CDC-STM-P007/2020, SARS-CoV-2/human/USA/CA-CDC-STM-P002/2020, SARS-CoV-2/human/USA/CA-CDC-STM-P008/2020, SARS-CoV-2/human/USA/FL-CDC-STM-P005/2020, SARS-CoV-2/human/USA/FL-CDC-STM-P011/2020, SARS-CoV-2/human/USA/FL-CDC-STM-P013/2020, SARS-CoV-2/human/USA/FL-CDC-STM-P014/2020, SARS-CoV-2/human/USA/FL-CDC-STM-P015/2020, SARS-CoV-2/human/USA/FL-CDC-STM-P058/2020, SARS-CoV-2/human/USA/MA-CDC-STM-P006/2020, SARS-CoV-2/human/USA/MA-CDC-STM-P016/2020, SARS-CoV-2/human/USA/MA-CDC-STM-P018/2020, SARS-CoV-2/human/USA/PA-CDC-STM-P010/2020, SARS-CoV-2/human/USA/PA-CDC-STM-P020/2020, SARS-CoV-2/human/USA/CA-LACPHL-AF00166/2021, SARS-CoV-2/human/USA/WI-UW-2513/2020, SARS-CoV-2/human/USA/WI-UW-2527/2020, SARS-CoV-2/human/USA/MA-CDC-STM-A100010/2021, SARS-CoV-2/human/USA/FL-CDC-STM-A100001/2021, SARS-CoV-2/human/USA/FL-CDC-STM-A100090/2021, SARS-CoV-2/human/USA/UT-UPHL-2012257709/2020
4. Use Quick Search to pull out corresponding whole genome sequences.
5. Use SNP tool to calculate consensus genome sequence => "B.1.375_North America_genome_consensus.fasta"
6. Combine with Wuhan-Hu-1_genome.fasta with B.1.375_North America_genome_consensus.fasta=> "Wuhan+ B.1.375_North America_genome_consensus.fasta"
7. Run SNP on "Wuhan+ B.1.375_North America_genome_consensus.fasta" to generate SNP report of all consensus substitutions in comparison with Wuhan-Hu-1 => 38 SNPs, including 22 deletions in "Wuhan-Hu-1 + B.1.375 consensus_genome_SNP.xlsx"
8. Annotate "B.1.375_North America_genome_consensus.fasta" with VIGOR4 => "B.1.375_North America_genome_consensus_VIGOR4"

9. Align “Wuhan-Hu-1 S protein.fasta” with “B.1.375_North America_S protein_consensus.fasta” to determine S protein substitutions => Spike: H69-, V70-, D614G
10. Determine which SNPs were located within which ORF gene or mature peptide genomic region through Wuhan mapping table. Combine relevant B.1.375_North America_genome_consensus_VIGOR4 protein with corresponding Wuhan-Hu-1 protein. Perform multiple sequence alignment and determine amino acid substitutions for all genomic SNPs => nsp2: T85I; nsp3: T1010A; RNA-dependent RNA polymerase: P323L; helicase: E341D; 2'-O-ribose methyltransferase: A258V; ORF3a: Q57H; membrane glycoprotein: I48V; nucleocapsid phosphoprotein: T205I
11. BLAST consensus against “B.1.375_North America_genome” => best representative (Identities = 29869/29876 (99%) SARS-CoV-2/human/USA/FL-CDC-STM-P015/2020 - <https://www.viprbrc.org/brc/viprStrainDetails.spg?ncbiAccession=MW430977&decorator=corona>