

Detection and characterization of the SARS-CoV-2 lineage B.1.526 in New York

Anthony P. West, Jr.^{1*}, Joel O. Wertheim², Jade C. Wang³, Tetyana I. Vasylyeva², Jennifer L. Havens⁴, Moinuddin A. Chowdhury³, Edimarlyn Gonzalez³, Courtney E. Fang³, Steve S. Di Lonardo³, Scott Hughes³, Jennifer L. Rakeman³, Henry H. Lee^{5,6} Christopher O. Barnes¹, Priyanthi N. P. Gnanapragasam¹, Zhi Yang¹, Christian Gaebler⁷, Marina Caskey⁷, Michel C. Nussenzweig^{7,8}, Jennifer R. Keeffe¹, Pamela J. Bjorkman¹

¹Division of Biology and Biological Engineering, California Institute of Technology, Pasadena, CA 91125, USA.

²Department of Medicine, University of California San Diego, La Jolla, CA 92093

³New York City Public Health Laboratory, New York City Department of Health and Mental Hygiene, New York, NY, 10016 USA

⁴Bioinformatics and Systems Biology Graduate Program, University of California San Diego, La Jolla, CA 92093

⁵Pandemic Response Laboratory, Long Island City, NY 11101

⁶Department of Genetics, Harvard Medical School, Boston, MA 02115

⁷Laboratory of Molecular Immunology, The Rockefeller University, New York, NY 10065, USA.

⁸Howard Hughes Medical Institute, The Rockefeller University, New York, NY, 10065 USA.

*Corresponding author: Anthony P. West, Jr., apwest@caltech.edu

1 Abstract

2 Wide-scale SARS-CoV-2 genome sequencing is critical to tracking viral evolution during the
3 ongoing pandemic. Variants first detected in the United Kingdom, South Africa, and Brazil have
4 spread to multiple countries. We developed the software tool, Variant Database (VDB), for
5 quickly examining the changing landscape of spike mutations. Using VDB, we detected an
6 emerging lineage of SARS-CoV-2 in the New York region that shares mutations with previously
7 reported variants. The most common sets of spike mutations in this lineage (now designated as
8 B.1.526) are L5F, T95I, D253G, E484K or S477N, D614G, and A701V. This lineage was first
9 sequenced in late November 2020 when it represented <1% of sequenced coronavirus genomes
10 that were collected in New York City (NYC). By February 2021, genomes from this lineage
11 accounted for ~32% of 3288 sequenced genomes from NYC specimens. Phylodynamic inference
12 confirmed the rapid growth of the B.1.526 lineage in NYC, notably the sub-clade defined by the
13 spike mutation E484K, which has outpaced the growth of other variants in NYC. Pseudovirus
14 neutralization experiments demonstrated that B.1.526 spike mutations adversely affect the
15 neutralization titer of convalescent and vaccinee plasma, indicating the public health
16 importance of this lineage.

17

18 Introduction

19 After the early months of the SARS-CoV-2 pandemic in 2020, the vast majority of sequenced
20 genomes contained the spike mutation D614G (along with 3 separate nucleotide changes)¹.
21 Following a period of gradual change, the fourth quarter of 2020 witnessed the emergence of
22 several variants containing multiple mutations, many within the spike gene²⁻⁵. Multiple lines of
23 evidence support escape from antibody selective pressure as a driving force for the
24 development of these variants⁶⁻⁹.

25
26 Genomic surveillance of SARS-CoV-2 is now focused on monitoring the emergence of these
27 variants and the functional impact that their mutations may have on the effectiveness of
28 passive antibody therapies and the efficacy of vaccines to prevent mild or moderate COVID-19.
29 While an increasing number of specimens are being sequenced, analysis of these genomes
30 remains a challenge¹⁰. Here, we developed a simple and fast utility that permits rapid
31 inspection of the mutational landscape revealed by genomic surveillance of SARS-CoV-2:
32 Variant Database (**vdb**). With this tool, we uncovered several groups of recently sequenced
33 genomes with mutations at critical antibody epitopes. Among this group is a new lineage
34 emerging in NYC that has increased in frequency to now account for ~32% of sequenced
35 genomes as of February 2021. We confirm the rapid spread of B.1.526 in NYC during early 2021
36 through phylodynamic inference. Furthermore, we evaluated the impact of the B.1.526 spike
37 mutations on the neutralization titer of convalescent and vaccinee plasma.

38 Results

39 vdb

40 Phylogenetic analysis is critical to understand the relationships of viral genomes. However,
41 other perspectives can be useful for detecting patterns in large numbers of sequences. We
42 developed **vdb** as a utility to query the sets of spike mutations observed during genomic
43 surveillance. Using the **vdb** tool to analyze SARS-CoV-2 sequences in the Global Initiative on
44 Sharing Avian Influenza Data (GISAID) dataset^{11,12}, we detected several clusters of sequences
45 distinct from variants B.1.1.7, B.1.351, B.1.1.248, and B.1.429²⁻⁵ with spike mutations at sites
46 known to be associated with resistance to antibodies against SARS-CoV-2^{8,13} (**Table 1**). The **vdb**
47 program can find clusters of virus sharing identical sets of spike mutations, and then these
48 patterns can be used to find potentially related sequences.

49 Defining mutations of B.1.526

50 One notable cluster of genome sequences was collected from the New York region and
51 represents a distinct lineage, now designated as B.1.526 (**Figure 1, Supplementary Figure 1**).
52 This variant is found within the 20.C clade and is distinguished by 3 defining spike mutations:
53 L5F, T95I, and D253G. Within B.1.526, the largest sub-clade is defined by E484K and two distinct
54 sub-clades are each defined by S477N; both of these mutations located within the receptor-
55 binding domain (RBD) of spike (**Figure 2 and Supplementary Table 1**). We note that the
56 evolutionary history at spike position 701 varies depending on whether the tree is rooted using
57 a molecular clock (**Figure 1**) versus its sister clade (characterized by an L452R mutation;
58 **Supplementary Figure 2**), the latter of which posits a substitution A701V followed by a
59 reversion V701A. Among the nucleotide mutations in lineage B.1.526, the most characteristic
60 include A16500C (NSP13 Q88H), A22320G (spike D253G), and T9867C (NSP4_L438P). Another

61 notable feature of the B.1.526 lineage is the deletion of nucleotides 11288-11296 (NSP6 106-
62 108), which also occurs in variants B.1.1.7, B.1.351, P.1, and B.1.525¹⁴.
63
64 Regarding four of the spike mutations prevalent in this lineage: (1) E484K is known to attenuate
65 neutralization of multiple anti-SARS-CoV-2 antibodies, particularly those found in class 2 anti-
66 RBD neutralizing antibodies^{13,15}, and is also present in variants B.1.351⁴ and P.1/B.1.1.248², (2)
67 D253G has been reported as an escape mutation from antibodies against the N-terminal
68 domain¹⁶, (3) S477N has been identified in several earlier lineages¹⁷, is near the epitopes of
69 multiple antibodies¹⁸, and has been implicated to increase viral infectivity through enhanced
70 interactions with ACE2^{19,20}, and (4) A701V sits adjacent to the S2' cleavage site of the
71 neighboring protomer and is shared with variant B.1.351⁴. The overall pattern of mutations in
72 lineage B.1.526 (**Figure 2**) suggests that it arose in part in response to selective pressure from
73 antibodies. Based on the dates of collection of these viruses, it appears that the frequency of
74 this lineage has increased rapidly in New York (**Table 2**).

75 [Trends in B.1.526 surveillance](#)

76 As part of public health surveillance conducted by the New York City Public Health Laboratory
77 (NYC PHL) and the Pandemic Response Lab (PRL) in New York, approximately 4.5 thousand
78 SARS-CoV-2 genomes have been sequenced by NYC PHL and PRL from December 1, 2020 to
79 February 28th, 2021. Of these genomes, approximately 25% are from lineage B.1.526. We
80 separately analyzed these genomes, because viral genomic surveillance by PHL and PRL
81 provides a less biased picture of viral diversity in NYC than genomes uploaded to GISAID. The
82 proportion of B.1.526 genomes in NYC has steadily increased since this variant was first
83 detected in NYC surveillance data in late 2020, and its weekly average exceeded 10% by 14

84 January 2021. From early January to early March, B.1.526 has been increasing by about 0.7%
85 per day (segmented linear regression) and was at 43% the week prior to 03 March 2021 (**Figure**
86 **3A**). Around 54% (n=678) of the B.1.526 genomes contain the E484K mutation, which has also
87 been rising in frequency since early 2021. The weekly average of B.1.526 genomes with E484K
88 has been above 10% since 01 February 2021 and has been increasing around 0.4% per day
89 (**Figure 3B**).

90
91 This increase in B.1.526 temporally coincides with the peak and subsequent decline of the
92 second epidemic wave in NYC (**Figure 3C**). If we separate the approximated number of B.1.526
93 cases from the rest of second wave SARS-CoV-2, the non-B.1.526 virus has steadily declined
94 since its peak in early January 2021. However, the increasing proportion of B.1.526 appears to
95 have slowed the rate of decline in total COVID-19 case counts in NYC.

96
97 [Geographic distribution of B.1.526 in NYC](#)
98 The New York City Public Health Laboratory and the PRL in New York have sequenced 4538
99 SARS-CoV-2 genomes from December 2020 thru February 2021 (**Figure 4A**). Geographic case
100 distribution of specimens received at PHL and PRL for SARS-CoV-2 diagnostic nucleic acid
101 amplification testing (NAAT) are representative of citywide testing efforts. Those SARS-CoV-2
102 positive specimens with NAAT cross-threshold values below 32 were selected at random to be
103 sequenced. On a month-to-month basis using data generated by NYC PHL and PRL, we have
104 observed an increasing number of B.1.526 genomes identified throughout NYC. The geographic
105 distribution of over 600 B.1.526 E484K cases is similar (**Figure 4B**). While the B.1.526 lineage is

106 not limited to NYC, almost 90% of genomes deposited to GISAID prior to March 2021, are from
107 the New York region.

108

109 [Phylodynamic analysis](#)

110 Other SARS-CoV-2 variants of concern or interest (B.1.1.7, B.1.427, and B.1.429) have also been
111 circulating in NYC contemporaneously with the rise of B.1.526 and have all risen in relative
112 frequency during the second wave of the NYC pandemic (**Figure 3D**). To compare the relative
113 growth rates of these variants during this time-period, we fitted an exponential population
114 growth model²¹ implemented in BEAST1.10²² to the sequences that correspond to these
115 lineages of interest. Specifically, we estimated the growth rate for the B.1.1.7, B.1.427, and
116 B.1.429 variants and for two subsets of the B.1.526 clade sequences (with and without the
117 E484K mutation).

118

119 The B.1.526 E484K clade experienced more rapid exponential growth compared with other
120 lineages: 23.2 (95% highest posterior density [HPD]: 19.6–27.1). B.1.526 with E484 and B.1.1.7
121 experienced similar growth rates: 14.3 (95% HPD: 11.7–16.9) and 14.5 (95% HPD 11.6 – 17.8),
122 respectively. The B.1.427 and B.1.429 lineages experienced lower growth rates that were
123 significantly greater than zero: 3.8 (95% HPD: 0.7–7.0) and 5.2 (95% HPD: 2.1–8.3), respectively.
124 We caution that these lineage growth rates do not distinguish between per-contact
125 transmissibility or per-virion infectiousness and speak only to the relative number of people
126 detected with these variants in NYC during late 2020 and early 2021.

127

128 As part of the phylodynamic analysis, we inferred the time of most recent common ancestor
129 (TMRCA) for the B.1.526 E484K clade to be 08 November 2020 (95% HPD: 22 October – 24
130 November). The TMRCA for the rest of the B.1.526 clade was estimated to be 15 September
131 2020 (95% HPD: 17 August – 08 October).

132

133 [Neutralization activity of convalescent and vaccinee plasma against B.1.526](#)

134 The identification of several mutations associated with resistance to anti-SARS-CoV-2
135 antibodies in B.1.526 sequences raises the question of the impact on SARS-CoV-2 immunity. We
136 generated HIV-based pseudoviruses expressing SARS-CoV-2 spike protein containing either the
137 most common B.1.526 mutation pattern (v.1: L5F, T95I, D253G, E484K, D614G, and A701V), the
138 2nd most common pattern (v.2: L5F, T95I, D253G, S477N, D614G, and Q957R), or only D614G.
139 Pseudovirus neutralization titers were determined for human plasma samples from vaccinees
140 [Moderna (mRNA-1273) or Pfizer-BioNTech(BNT162b2)]⁸ or convalescent plasma [at either
141 1.3¹⁵ or 6.2 months¹³ post-infection]. The E484K-containing B.1.526 pseudovirus had a
142 statistically significant reduced neutralization titer compared to the D614G control: for vaccinee
143 plasma, 4.5-fold reduced ($p = 0.00005$); for 1.3-month convalescent plasma, 6.0-fold reduced (p
144 = 0.03); and for 6.2-month convalescent plasma, 4.8-fold reduced ($p = 0.02$) (**Figure 5a and**
145 **Supplementary Table 2**). The smaller reduction of the titers in the 6.2-month convalescent
146 plasma samples compared to the 1.3-month samples is consistent with the greater resistance of
147 more matured anti-SARS-CoV-2 antibodies to viral escape mutations²³. The S477N/Q957R-
148 containing B.1.526 pseudovirus demonstrated a smaller effect on plasma neutralization (**Figure**
149 **5b**).

150

151 Discussion

152 Genomic surveillance is a critical tool to monitor the progression of the COVID-19 pandemic
153 and modelling suggests that sequencing at least 5% of specimens that test positive for SARS-
154 Cov-2 in a geographic region is necessary to reliably detect the emergence of novel variants at a
155 lower prevalence limit of between 0.1% to 1%²⁴. Through the combination of increased
156 sequencing efforts and the use of the software utility described here, we were able to identify
157 the B.1.526 lineage and to begin to characterize its phylogenetic and phylodynamic patterns in
158 NYC in early 2021. Based on sequences in GISAID as of March 2021, the majority of cases with
159 sequence data are in the NYC region, but it is expected that the prevalence B.1.526 variants will
160 continue to increase beyond the NYC region. The B.1.526 variant has also been described in
161 other recent studies^{25,26}.

162
163 Pseudovirus containing spike gene mutations associated with B.1.526 was significantly more
164 resistant to neutralization by either convalescent or vaccinee plasma. The presence of E484K
165 mutation likely plays a key role in facilitating increased viral transmission and reducing antibody
166 neutralizing titers, as previously shown in other studies^{7,27}. Continued monitoring for emerging
167 variants with mutations such as E484K is important to maximize the impact of public health
168 measures to mitigate the effects of the SARS-CoV-2 pandemic. For example, high frequencies of
169 SARS-CoV-2 variants has potential impacts on selection of appropriate antibody therapeutics
170 and vaccination strategies.

171 Methods

172 Variant Database Program

173 We developed a software tool named VDB (Variant Database). This tool consists of two Unix
174 command line utilities: (1) **vdb**, a program for examining spike mutation patterns in a collection
175 of sequenced viral genomes, and (2) **vdbCreate**, a program for generating a list of viral spike
176 mutations from a multiple sequence alignment for use by **vdb**. The design goal for the query
177 program **vdb** is to provide a fast, lightweight, and natural means to examine the landscape of
178 SARS-CoV-2 spike mutations. These programs are written in Swift and are available for MacOS
179 and Linux from the authors or from the Github repository: [https://github.com/variant-](https://github.com/variant-database/vdb)
180 [database/vdb](https://github.com/variant-database/vdb).

181 The **vdb** program implements a mutation pattern query language (see Supplemental Method)
182 as a command shell. The first-class objects in this environment are a collection of viruses (a
183 “cluster”) and a group of spike mutations (a “pattern”). These objects can be assigned to
184 variables and are the return types of various commands. Generally, clusters can be obtained
185 from searches for patterns, and patterns can be found by examining a given cluster. Clusters
186 can be filtered by geographical location, collection date, mutation count, or the presence or
187 absence of a mutation pattern. The geographic or temporal distribution of clusters can be
188 listed.

189 Results presented here are based on a multiple sequence alignment from GISAID^{11,12}
190 downloaded on February 10, 2021. Additional sequences downloaded from GISAID on February
191 22, 2021, were aligned with MAFFT v7.464²⁸.

192 Initial Phylogenetic Analysis

193 Multiple sequence alignments were performed with MAFFT v7.464²⁸. The phylogenetic tree
194 was calculated by IQ-TREE²⁹, and the tree diagram was generated using iTOL (Interactive Tree of
195 Life)³⁰. The Pango lineage nomenclature system³¹ provides systematic names for SARS-CoV-2
196 lineages. The Pango lineage designation for B.1.526 was supported by the phylogenetic tree
197 shown in **Supplementary Figure 1**.

198 Library preparation and sequencing

199 RNA was extracted from positive specimens collected at NYC PHL using the EZ1 (Qiagen, CA),
200 NUCLISENS® easyMAG® (bioMérieux Inc., Netherlands), or Kingfisher™ Flex Purification System
201 (Thermo Fisher Scientific, MA). RNA extracts were subjected to annealing reaction with random
202 hexamers and dNTPs (New England Biolabs Inc., NEB, MA), and reverse transcribed with
203 SuperScript IV Reverse Transcriptase at 42°C for 50 min. The resulting cDNA was amplified
204 using two separate multiplex PCRs with ARTIC V3 primer pools (Integrated DNA Technologies,
205 IA) per sample in the presence of Q5 2X Hot Start Master Mix (NEB) at 98°C for 30 secs,
206 followed by 35 cycles of 98°C for 15 secs and 65°C for 5 min^{32,33}. The resulting PCR products per
207 sample were combined and purified using Agencourt Ampure XP magnetic beads (Beckman
208 Coulter, IN), at a ratio of 1:1 sample to bead ratio and quantified using a Qubit 3.0 fluorometer
209 (Thermo Fisher Scientific, MA). The PCR products were normalized to 90 ng as input for the
210 NEBNext Ultra II Library Preparation Kit according to standard protocol (NEB): Briefly, the ARTIC
211 PCR products were subjected to simultaneous end-repair, 5'-phosphorylation, and dA-tailing
212 reaction at 20°C for 30 min, followed by heat inactivation at 65°C for 30 min. NEBNext Adaptor
213 was then ligated at 25°C for 30 min, and then cleaved by USER Enzyme at 37°C for 15 min. This
214 product was subjected to bead cleanup at a ratio of 0.6x sample to bed ratio. The eluted

215 product was amplified for 6 cycles using NEBNext Ultra II Q5 Master Mix in the presence of
216 NEBNext Multiplex Oligos for Illumina (NEB). The PCR product was purified with Ampure XP
217 beads at a 0.6x sample to bead ratio. The product was a barcoded library containing Illumina P5
218 and P7 adapters for sequencing on Illumina instruments. The individual libraries were
219 quantified, normalized and pooled at equimolar concentration and loaded onto the Illumina
220 MiSeq sequencing instrument using V3 600-cycle reagent kits and a V3 flow cell for 250-cycle
221 paired end sequencing (Illumina, CA).

222 [Genome Assembly](#)

223 All raw paired end sequence reads are trimmed using Trim Galore version 0.6.4_dev³⁴ removing
224 NEB adapters and quality score below 20 from ends of the reads. The trimmed reads were
225 assembled using the Burrows-Wheeler Aligner MEM algorithm (BWA-MEM) version 0.7.12³⁵
226 with SARS-CoV-2 Wuhan-Hu-1 (GenBank accession number MN908947.3) as the reference
227 sequence. Intrahost variant analysis of replicates (iVar)³⁶ tool was used to remove primer
228 sequences from the amplicon-based sequencing data. Finally, the mutation calls and consensus
229 genome were built using a combination of samtools mpileup³⁷ and iVar consensus, with a
230 minimum quality score of 20, frequency threshold of 0.6, and minimum depth of 15 to optimize
231 high quality variant calls. A sequence mapping quality control tool developed in-house was used
232 to assess depth of coverage across all sequences, percent of ambiguous bases in the consensus
233 genome and percent sequence mapped to the reference genome. Consensus genome with
234 more than 3% ambiguous bases or less than 95% reference mapped were excluded from any
235 further analyses.

236 Library preparation and sequencing (PRL)

237 Positive RNA specimens between cycle threshold of 15-30 were selected from all samples
238 tested at Pandemic Response Labs, NYC and cDNA for each specimen was generated using
239 LunaScript RT SuperMix (NEB, MA) according to manufacturer protocol. To target SARS-CoV-2
240 specifically, cDNA for each specimen was amplified in two separate pools, 28- and 30-plex
241 respectively, to generate 1200bp of overlapping amplicons³⁸ using Q5 2x Hot-Start Master Mix
242 (NEB, MA). The resulting pools are combined in equal volume and enriched for full length 1200
243 bp product using a SPRI-based magnetic bead cleanup. Enriched amplicons are tagmented
244 (Illumina, CA) and barcoded (IDT, IA) and paired-end sequenced on an Illumina MiSeq or
245 NextSeq 550.

246

247 Genome Assembly (PRL)

248 For each specimen, sequencing adapters are first trimmed using Trim Galore v0.6.6³⁴, then
249 aligned to the SARS-CoV-2 Wuhan-Hu-1 reference genome (NCBI Nucleotide NC_045512.2)
250 using BWA MEM 0.7.17-r1188³⁵. Reads that are unmapped or those that have secondary
251 alignments are discarded from the alignment. Consensus and mutations were called using
252 samtools³⁷ and Intrahost variant analysis of replicates (iVar)³⁶ with a minimum quality score of
253 20, frequency threshold of 0.6 and a minimum read depth of 10x coverage. A consensus
254 genome with $\geq 90\%$ breath-of-coverage with ≤ 3000 ambiguous bases is considered a successful
255 reconstruction (as per APHL recommendation).

256

257 [Genome alignment](#)

258 Complete genome sequences produced by the NYC PHL and the PRL with reported collection
259 dates on or before 04 March 2021 were analyzed. We restricted our analysis to genomes
260 produced by public health surveillance to NYC to reduce bias due to geography or preferential
261 sequencing of viral variants by academic institutions. Genomes were aligned to the Wuhan-Hu-
262 1 reference genome (GenBank Accession MN908947) using mafft v7.475 (mafft --6merpair --
263 keeplength --addfragments)²⁸. Pango lineage designations³¹ for variants were assigned using
264 Pangolin v2.3.2³⁹.

265

266 [Segmented regression analysis](#)

267 To estimate the timing and approximate linear slope of increase in B.1.526 and the E484K clade
268 prevalence, we employed a segmented regression analysis (segmented package in R).

269

270 [Maximum likelihood phylogenetic inference](#)

271 Maximum likelihood trees were inferred using IQTree2 for B.1.1.7, B.1.427, B.1.429, and
272 B.1.526 genomes using a GTR+F+ Γ_4 substitution model⁴⁰. Minimum branch length of 1e-9 was
273 enforced and an expanded NNI search (--allnni) was employed to improve topology search.
274 Preliminary molecular clock analyses were performed in TreeTime v0.8.1 using a fixed
275 substitution rate of 8×10^{-4} substitutions/site/year and a skyline coalescent model⁴¹. This
276 analysis identified 34 genomes whose root-to-tip genetic distance were flagged as problematic
277 and excluded from subsequent phylodynamic analyses. TreeTime was also used to root and
278 perform ancestral state reconstruction for a tree inferred from the 258 B.1.526 genomes
279 sampled by the NYC PHL used to display the history of spike mutations in B.1.526 (**Figure 1**).

280

281 [Bayesian phylodynamic inference](#)

282 We performed population growth rate inference in coalescence-based framework using an
283 exponential growth model in BEAST 1.10.4²². We used a strict molecular clock model with the
284 fixed substitution rate of 8×10^{-4} substitutions/site/year. We applied a GTR+F+ Γ_4 substitution
285 model and specified the following priors for the population growth model: OneOnX distribution
286 prior for the population size parameter and Laplace distribution prior (mean = 0.0, scale = 1.0)
287 for the growth rate prior. Markov chain Monte Carlo analyses were run for 100-300 million
288 generations; the first 10% of samples were discarded as burn-in. Separate inference was
289 performed for B.1.1.7 (n=354), B.1.427 (n=35), B.1.429 (n=69), B.1.526 E484 (n=569), and
290 B.1.526 E484K (n=678). For the B.1.526 phylodynamic inference, we did not include two
291 sequences most closely related to B.1.526 (hCoV-19/USA/NY-NYCPHL-001701/2020 and hCoV-
292 19/USA/NY-NYCPHL-002542/2021).

293

294 [Geocoding addresses](#)

295 To identify areas with the highest density of B.1.526 sequenced genomes in NYC from
296 December 2020 to March 2021, patient addresses were geocoded to be visualized on a map⁴².
297 Geocoding was performed using the NYC DOHMH's Geoportal application. Once geocoded, a
298 map representing the point locations of individuals with sequenced B.1.526 genomes was
299 created in ArcMap (v. 10.6.1) and exported as a point feature class.

300 [Point density method](#)

301 Point density maps of individuals with B.1.526 sequenced genomes were created by using the
302 point density tool in ArcMap. Point density calculates the density-per-unit area from point

303 features (individuals with a SARS-CoV-2 B.1.526 sequenced genome) that fall within a defined
304 neighborhood by totaling the number of points that fall within the neighborhood divided by the
305 neighborhood area. Density calculations result in the observed gradient patterns. The point
306 density map parameters were 4000 ft radius from the center of 250 square foot cells. The
307 symbology class for point density classification was set at equal intervals of 5.

308

309 [Human plasma samples](#)

310 Human plasma samples were among those collected in previously reported studies^{8,13,15}. The
311 study visits and blood draws were performed in compliance with all relevant ethical regulations
312 and the protocol for human participants was approved by the Institutional Review Board (IRB)
313 of the Rockefeller University (protocol #DRO-1006).

314

315 [Pseudovirus neutralization by human plasma samples](#)

316 Human plasma samples were assayed for neutralization activity against lentiviruses
317 pseudotyped with SARS-CoV-2 spike containing a 21-amino acid cytoplasmic tail deletion and
318 either D614G or mutations corresponding to lineage B.1.526 (L5F, T95I, D253G, E484K, D614G,
319 and A701V). Pseudotyped lentiviruses were generated and neutralizations assays were
320 conducted as previously described^{43,44}. Briefly, lentiviral particles were produced by co-
321 transfecting the gene encoding SARS-CoV-2 spike protein (D614G or B.1.526) and Env-deficient
322 HIV backbone expressing Luciferase-IRES-ZsGreen. Plasma samples were heat inactivated at
323 56°C for 1 hour, then 3-fold serial diluted and incubated with SARS-CoV-2 pseudotyped virus for
324 1 hour at 37°C. The virus/plasma mixture was added to 293T_{ACE2} target cells, which were
325 seeded the previous day on poly-L-lysine coated plates. After incubating for 48 hours at 37°C,

326 target cells were lysed with Britelite Plus (Perkin Elmer) and luciferase activity was measured as
327 relative luminescence units (RLUs) and normalized to values derived from cells infected with
328 pseudotyped virus in the absence of plasma. Data were fit to 2-parameter non-linear regression
329 in Antibody database⁴⁵.

330

331 [Data availability](#)

332 The data analyzed as part of this project were obtained from the GISAID database and through
333 a Data Use Agreement between NYC DOHMH and the University of California San Diego.
334 Sequences analyzed by using the **vdb** tool were downloaded from GISAID. No personally
335 identifying information were included as part of these analyses. SARS-CoV-2 genomes included
336 in these analyses have been deposited in GISAID. See **Supplementary Data 1** for a list of
337 genomes, including which genomes were excluded from the phylogenetic analysis.
338 Data for **Figure 5** are provided in **Supplementary Table 2**.

339

340 [Code availability](#)

341 The source code for the vdb program is available at the Github repository:

342 <https://github.com/variant-database/vdb>.

343

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353 [Author Contributions](#)

354 A.P.W., J.O.W., J.L.H., T.I.V., H.H.L., S.H. and J.C.W. analyzed data. J.C.W., M.A.C., E.G. and H.H.L.
355 performed genome sequencing and assembly. J.O.W. curated data. C.G., M. Caskey and M.C.N.
356 provided clinical samples. P.N.P.G. and J.R.K. carried out experiments. A.P.W., C.O.B., Z.Y., S.H.,
357 S.S.D., C.E.F. and J.O.W. prepared figures. A.P.W., J.O.W., T.I.V., C.O.B., J.C.W. and S.H. wrote the
358 manuscript with input from all co-authors. A.P.W., P.J.B., J.O.W., J.L.R. and S.H. supervised the
359 study.

360

361 [Competing Interests](#)

362 P.J.B. is a co-inventor on a provisional application from the California Institute of Technology for
363 the use of mosaic nanoparticles as coronavirus immunogens. M.C.N., P.J.B., and C.O.B. are co-
364 inventors on provisional applications for several anti-SARS-CoV-2 monoclonal antibodies. J.O.W.
365 has received funding from Gilead Sciences, LLC (completed) and the CDC (ongoing) via grants
366 and contracts to his institution unrelated to this research.

367

368 **Tables**

369

370 **Table 1**

371 Mutation patterns of viruses with mutations at select Spike positions, excluding viruses related

372 to variants B.1.1.7, B.1.351, B.1.1.248, and B.1.429. Mutations included in this analysis were

373 E484K, N501Y, K417T, K417N, L452R, and A701V. In this table viruses are only included if their

374 spike mutation pattern exactly matches the given pattern. Note about P681H/P681R: variant

375 B.1.1.7 has P681H. Note about W152L: variant B.1.429 has W152C

376

377

378	Pattern	Number of genomes	Top Locations	First collection date
379	LSF T95I D253G E484K D614G A701V	243	US(240; NY 235)	12/16/2020
380	E484K D614G V1176F	235	Brazil(132), US(40)	4/15/2020
381	W152L E484K D614G G769V	49	US(32)	11/1/2020
382	E484K D614G P681H	37	US(37; MD 27)	11/18/2020
383	R102I F157L V367F E484K Q613H P681R	36	England(35)	12/27/2020
384	Q52R A67V H69-V70- Y144- E484K D614G Q677H F888L	36	England(22)	12/15/2020

385

386 [Table 2](#)

387 Counts of virus genomes in lineage B.1.526 by month in New York State. The total number of
388 sequenced genomes examined from GISAID from New York during these time periods is also
389 listed. *Latest viral collection date was March 4, 2021. Note that geographic sampling may have
390 varied over time as genome sequencing increased.

391
392
393 Viruses containing spike mutations T95I and D253G (earliest collection date Nov. 23, 2020)

394

395 Month	count	total sequences	fraction
396 Nov. 2020	2	524	0.4%
397 Dec. 2020	46	2209	2.1%
398 Jan. 2021	201	3148	6.4%
399 Feb. 2021	1207	3868	31.2%
400 March 2021*	124	274	45.3%

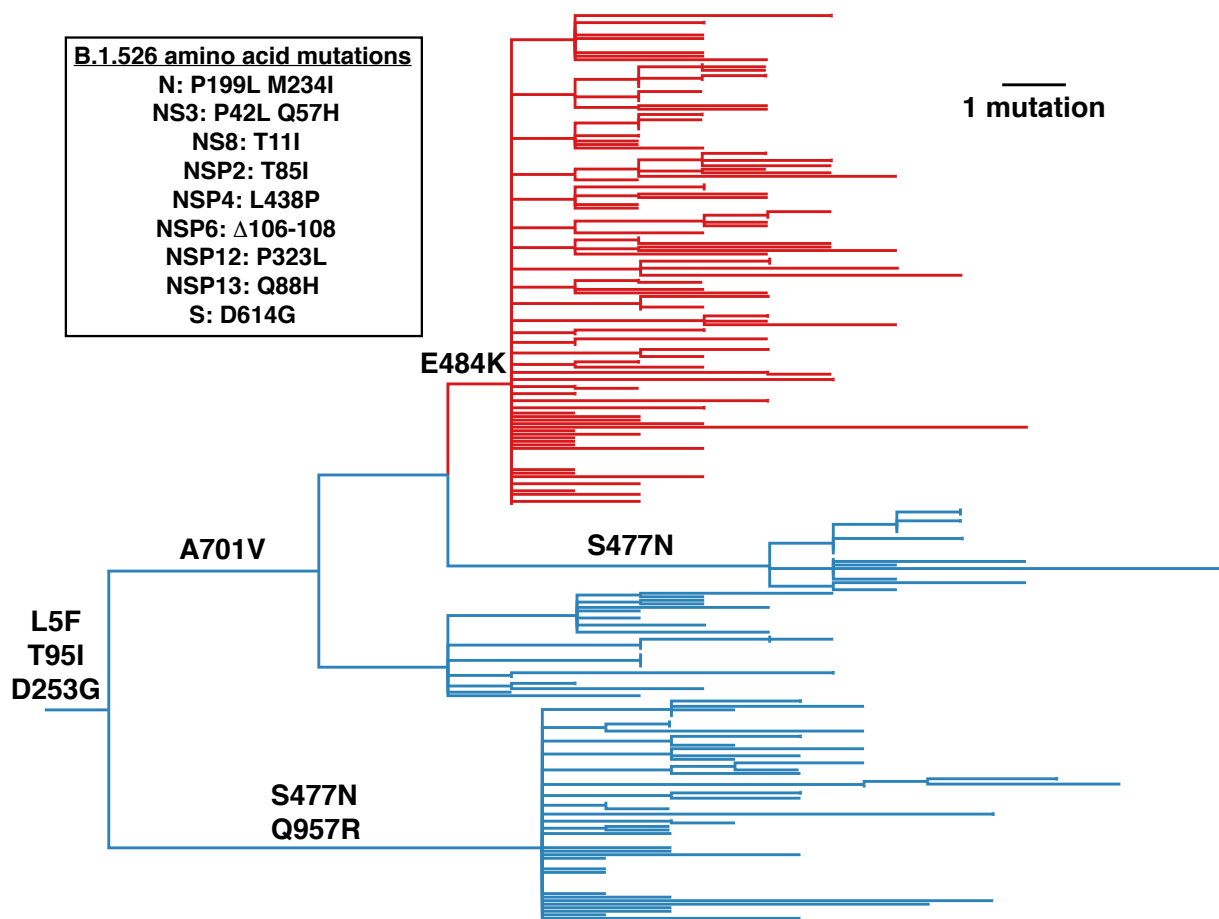
401
402
403
404 Viruses containing spike mutations L5F, T95I, D253G, E484K, D614G, and A701V (earliest
405 collection date Dec. 16, 2020)

406

407 Month	count	total sequences	fraction
408 Nov. 2020	0		
409 Dec. 2020	25	2209	1.1%
410 Jan. 2021	109	3148	3.5%
411 Feb. 2021	628	3868	16.2%
412 March 2021*	61	274	22.3%

413
414

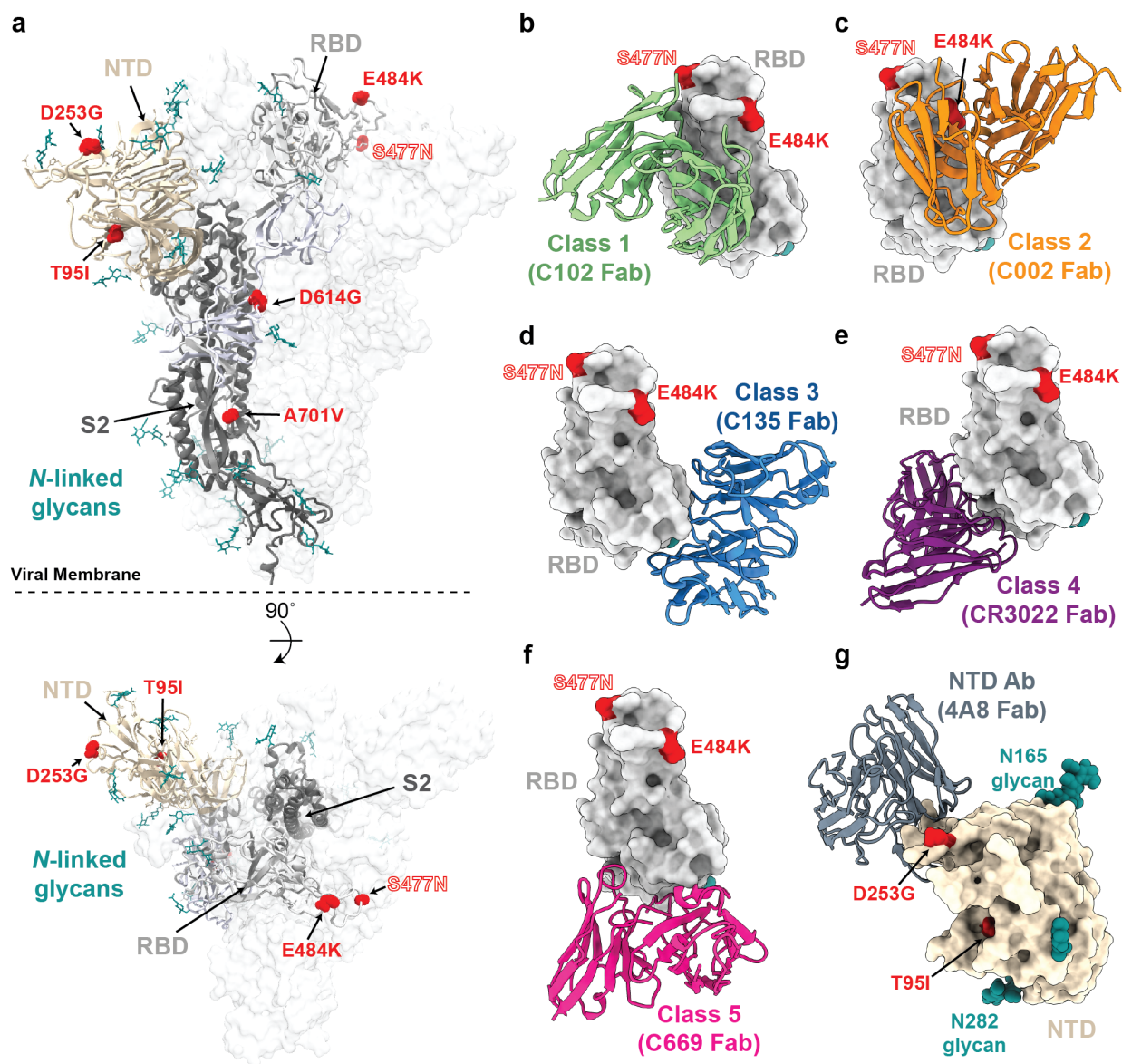
415 Figure 1.
416 Phylogenetic tree of lineage B.1.526 indicating spike mutations. Maximum likelihood phylogeny
417 of SARS-CoV-2 variant B.1.526 sampled by NYC PHL (n=258). Amino acid substitutions in the
418 spike protein occurring on internal branches are labeled, including the three spike mutations
419 characteristic of B.1.526. The B.1.526 clade defined by the E484K mutation is highlighted in red.
420 Inset highlights non-spike amino acid substitutions and deletions differentiating the B.1.526
421 clade from the Hu-1 reference genome. For display purposes, only NYC PHL genomes are
422 shown.
423



424

425

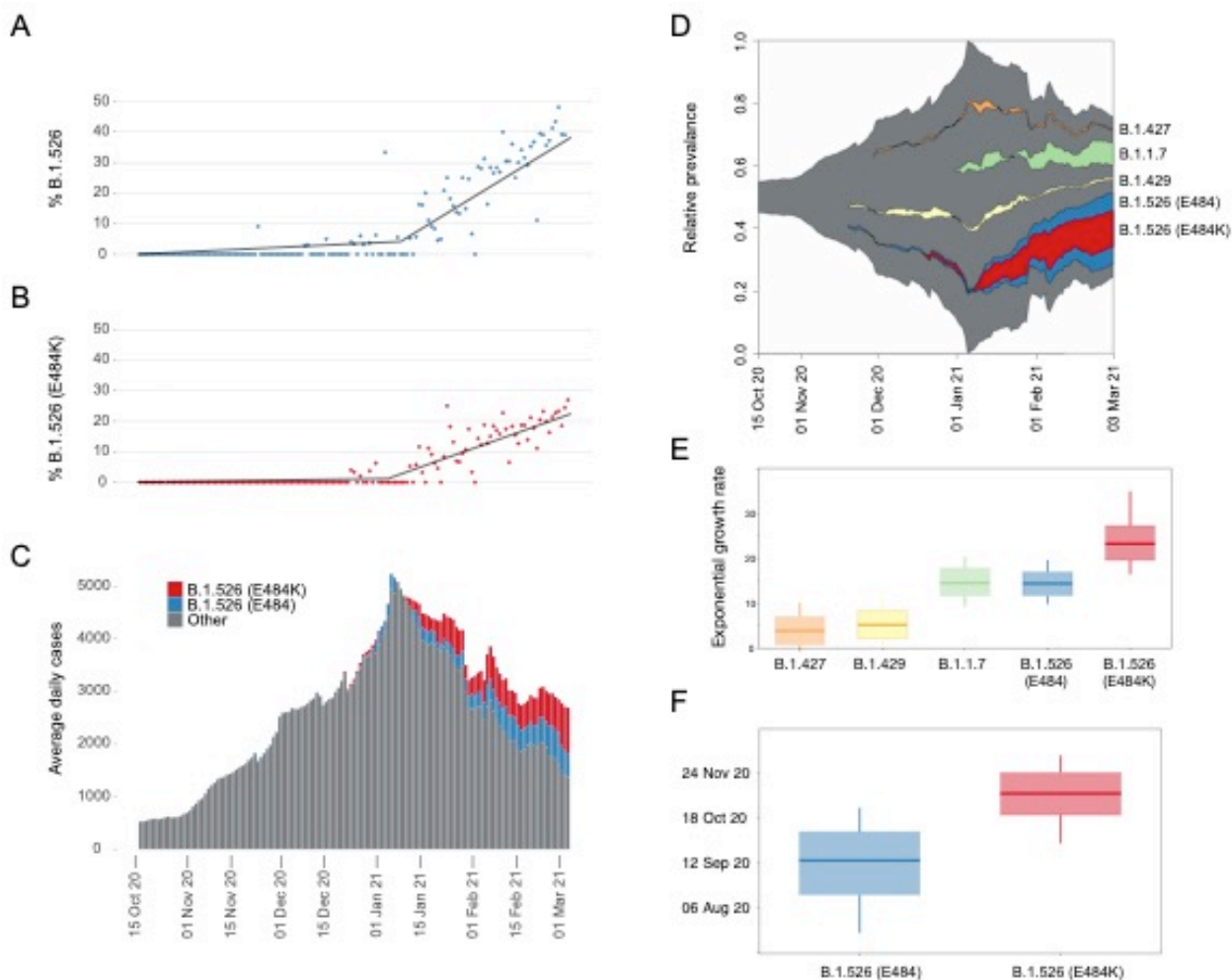
426 Figure 2.
427 Structural locations of the spike mutations of lineage B.1.526.



428
429 a, Side and top views of the SARS-CoV-2 spike trimer (PDB 7JJI) with mutations of lineage
430 B.1.526 shown as spheres. **b-g**, Models of representative neutralizing antibodies (cartoon,
431 VH-VL domain only) bound to RBD (**b-f**, gray surface) or NTD (**g**, wheat surface). Sites for
432 B.1.526 lineage mutations are shown as red spheres. The S477N site is also shown for the
433 branch containing this mutation instead of the E484K mutation (see **Figure 1**); **b**, Class 1 (PDB

434 7K8M); **c**, Class 2 (PDB 7K8S); **d**, Class 3 (PDB 7K8Z); **e**, Class 4 (PDB 6W41); **f**, Class 5⁸; **g**, NTD-
435 specific antibody 4A8 (PDB 7C2L).
436

437 Figure 3.



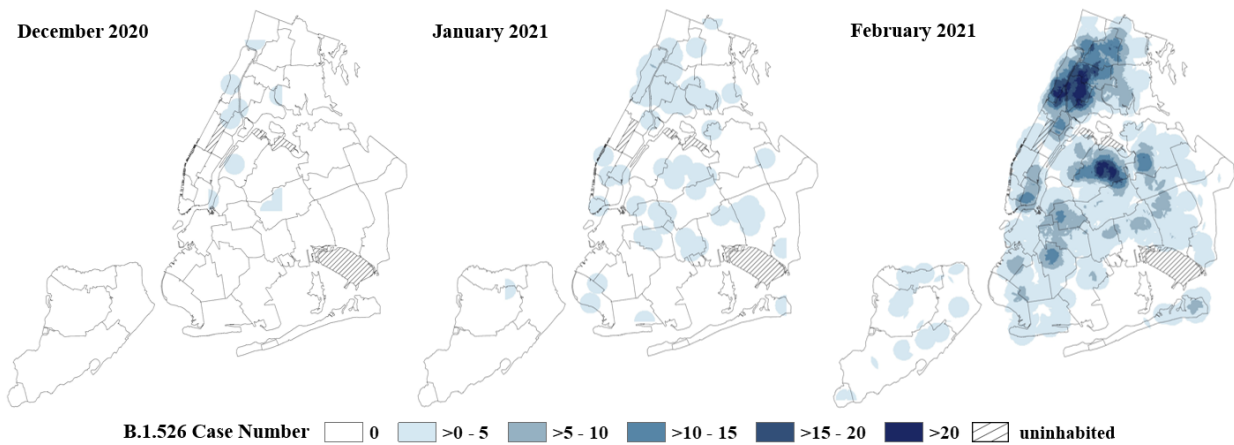
438

439 Rise of SARS-CoV-2 variants in New York City (NYC) in late-2020 and early 2021. (A) Relative
 440 frequency of B.1.526. Segmented linear regression is shown as a solid black line. (B) Relative
 441 frequency of B.1.526 with E484K mutation. Segmented linear regression is shown as a dashed
 442 gray line. (C) Rolling average number of total daily COVID-19 cases in NYC through time. Color
 443 indicates the estimated proportions of B.1.526 (blue) and B.1.526 E484K (red) extrapolated
 444 from a 7-day rolling average with an average of $n=236$ genomes sampled per week during this
 445 time period. (D) Muller plot depicting sampling, with pseudocounts, of SARS-CoV-2 variants
 446 scaled to the rolling average of total daily COVID-19 case counts. (E) Inferred exponential

447 growth rates for SARS-CoV-2 variants in NYC; the horizontal line indicates the median growth
448 rate estimate, the box outlines the interquartile range. **(F)** Inferred time of most recent
449 common ancestor (TMRCA) estimates for B.1.526 (E484) and B.1.526 (E484K).
450

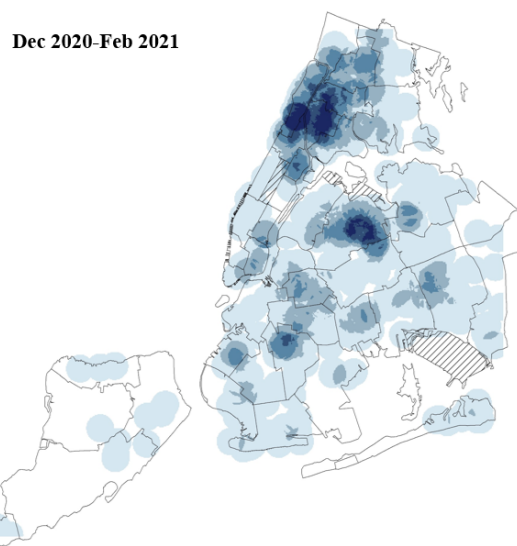
451 Figure 4.

452 **A**



453

454 **B**



455

B.1.526 E484K Case Number 0 >0 - 3 >3 - 6 >6 - 9 >9 - 12 >12 uninhabited

456

457 **(A)** Spatiotemporal increase of B.1.526 lineage in New York City (NYC). Point density of B.1.526

458 variants geo-located by case address overlaid on a map of NYC delineated by United Hospital

459 Fund areas. Data for each month is based on specimen collection date. The NYC PHL and the

460 PRL in New York have sequenced 4538 SARS-CoV-2 genomes from December 2020 thru

461 February 2021. Data represents 11 B.1.526 variants out of 515 sequenced genomes in
462 December 2020, 80 B.1.526 variants out of 735 sequenced genomes in January and 1063
463 B.1.526 variants identified out of a total of 3288 sequenced genomes in February 2021. **(B)**
464 Distribution of B.1.526 E484K cases in NYC. Point density map of 608 B.1.526 E484K variant
465 cases in NYC. Data is based on specimen collection period from December 1, 2020 through
466 February 28th, 2021.
467

468 **Figure 5.**

469 Plasma neutralizing activity against pseudoviruses with B.1.526 lineage spike mutations. SARS-

470 CoV-2 pseudovirus neutralization assays were used to determine neutralization titer (NT50) for

471 COVID-19 vaccinee (n=10) and convalescent plasma at 1.3 months (n=10) and 6.2 months (n=9)

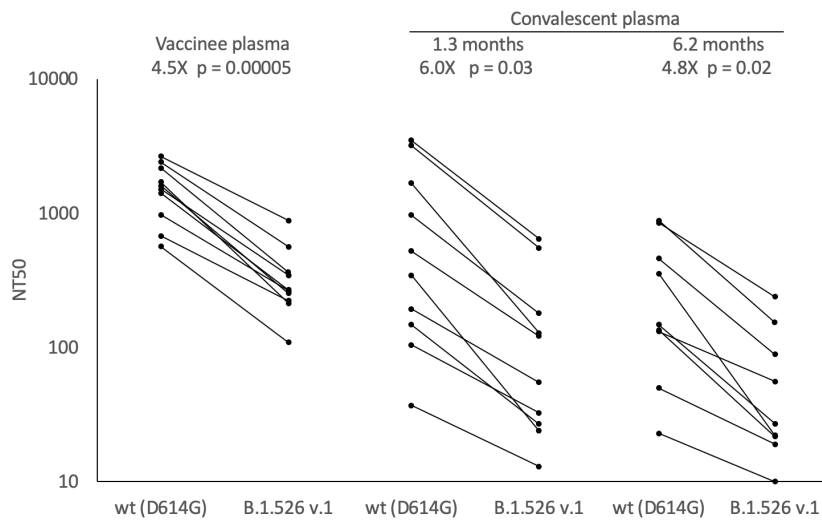
472 after infection. (A) Pseudovirus with spike mutations L5F, T95I, D253G, E484K, D614G

473 (B.1.526 v.1), and A701V, (B) Pseudovirus with spike mutations L5F, T95I, D253G, S477N,

474 D614G, and Q957R (B.1.526 v.2). Statistical significance was determined using paired two-tailed

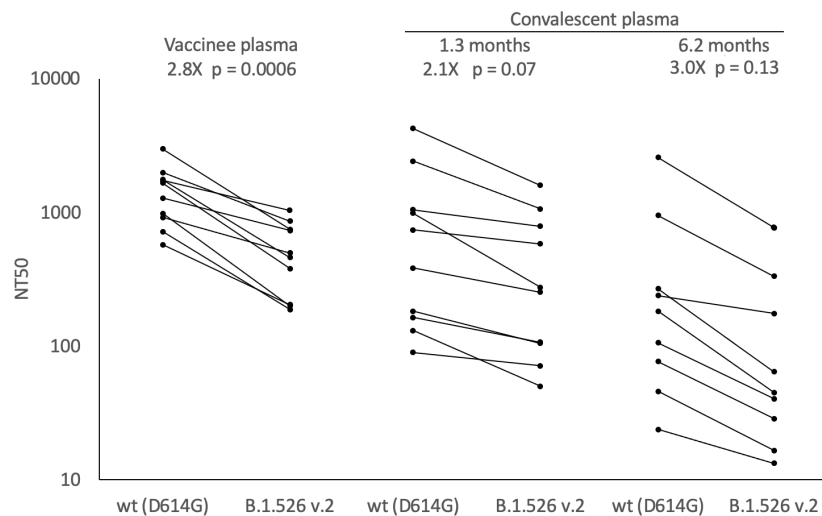
475 *t*-tests. Fold-differences of means are shown.

476 **A**



477

478 **B**



479

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489 [lineage-in-the-uk-defined-by-a-novel-set-of-spike-mutations/563](https://virological.org/t/preliminary-genomic-characterisation-of-an-emergent-sars-cov-2-lineage-in-the-uk-defined-by-a-novel-set-of-spike-mutations/563) (2020).
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- 599
600

601 Supplementary Material

602

603 Supplementary Methods.

604 Commands for the program **vdb**, implementing a mutation pattern query language:

605

606 Notation

607 cluster = group of viruses < > = user input n = an integer
608 pattern = group of mutations [] = optional () = explanation of command
609 "world" = all viruses in database -> result

610

611 To define a variable for a cluster or pattern: <name> = cluster or pattern

612 Set operations +, -, and * (intersection) can be applied to clusters or patterns

613 If no cluster is entered, all viruses will be used ("world")

614

615 Filter commands

616 <cluster> from <country or state> -> cluster
617 <cluster> containing [<n>] <pattern> -> cluster alias with (matches for >=n mutations)
618 <cluster> not containing <pattern> -> cluster alias without (considers whole pattern)
619 <cluster> before <date> -> cluster
620 <cluster> after <date> -> cluster
621 <cluster> > or <<n> -> cluster (filter by number of mutations)

622

623 Commands to find mutation patterns

624 consensus [for] <cluster or country or state> -> pattern
625 patterns [in] [<n>] <cluster> -> pattern (lists n patterns)

626

627 Listing commands

628 list [<n>] <cluster>
629 [list] countries [for] <cluster>
630 [list] states [for] <cluster>
631 [list] frequencies [for] <cluster> alias freq (frequency of individual mutations)
632 [list] monthly [for] <cluster> [<cluster2>] (number of viruses per month or week)
633 [list] weekly [for] <cluster> [<cluster2>] (as a fraction of number of viruses in cluster2)
634 [list] patterns (list built-in and user-defined patterns)
635 [list] clusters (list built-in and user-defined clusters)

636

637 Other commands

638 sort <cluster> (by date)

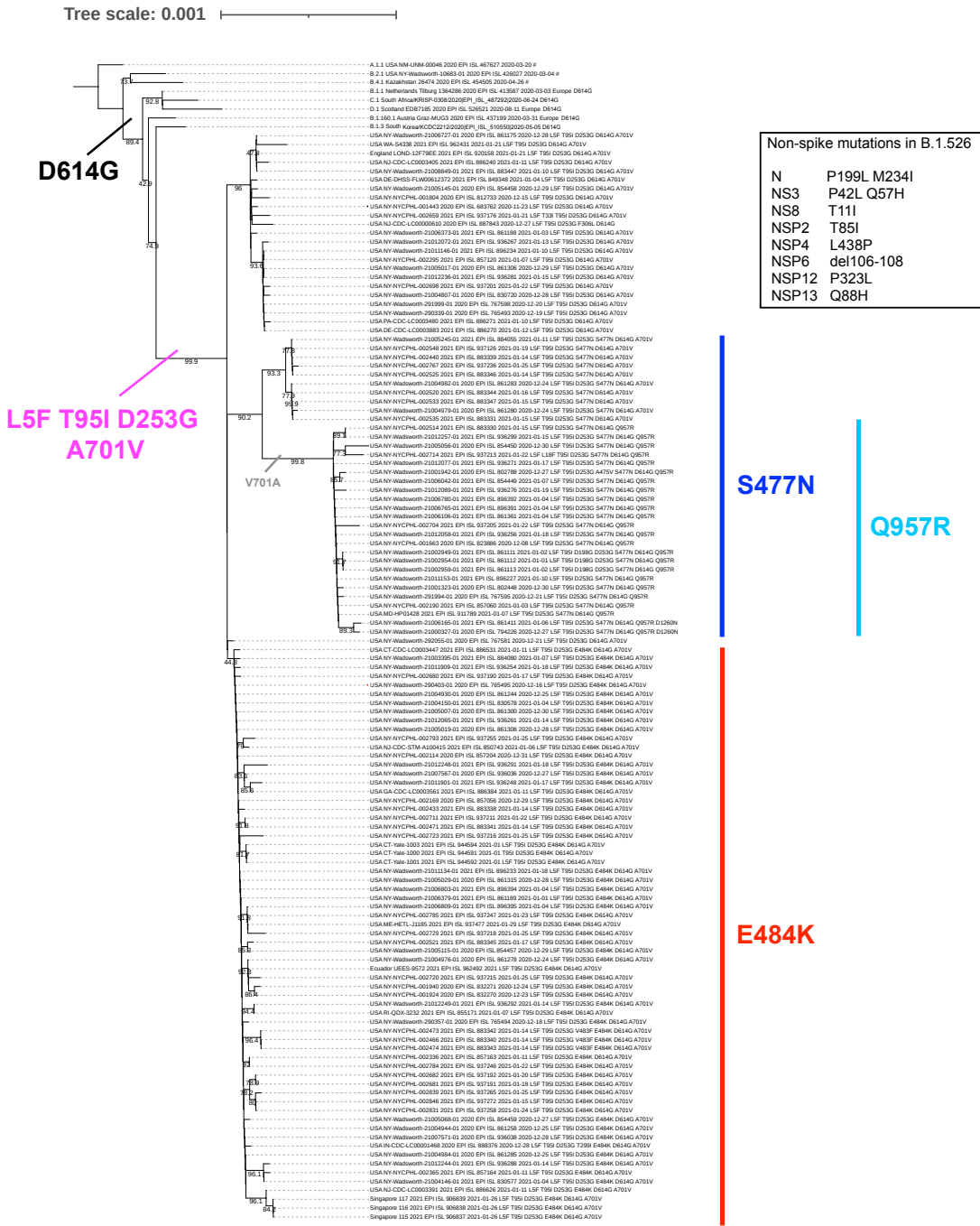
639 help

640 history

641 quit

642

643 **Supplementary Figure 1.**
 644 **Phylogenetic tree of lineage B.1.526 indicating spike mutations. The inset lists non-spike**
 645 **mutations common in this lineage.**



646

647 **Supplementary Figure 2.**

648 **Maximum likelihood phylogenetic tree of the B.1.526 lineage in relation to a sister clade**

649 defined by an L452R spike mutation and the 20C ancestral virus (both shown in gray). Tree was

650 rooted using the clade 20C ancestral viruses. Amino acid substitutions in the spike protein

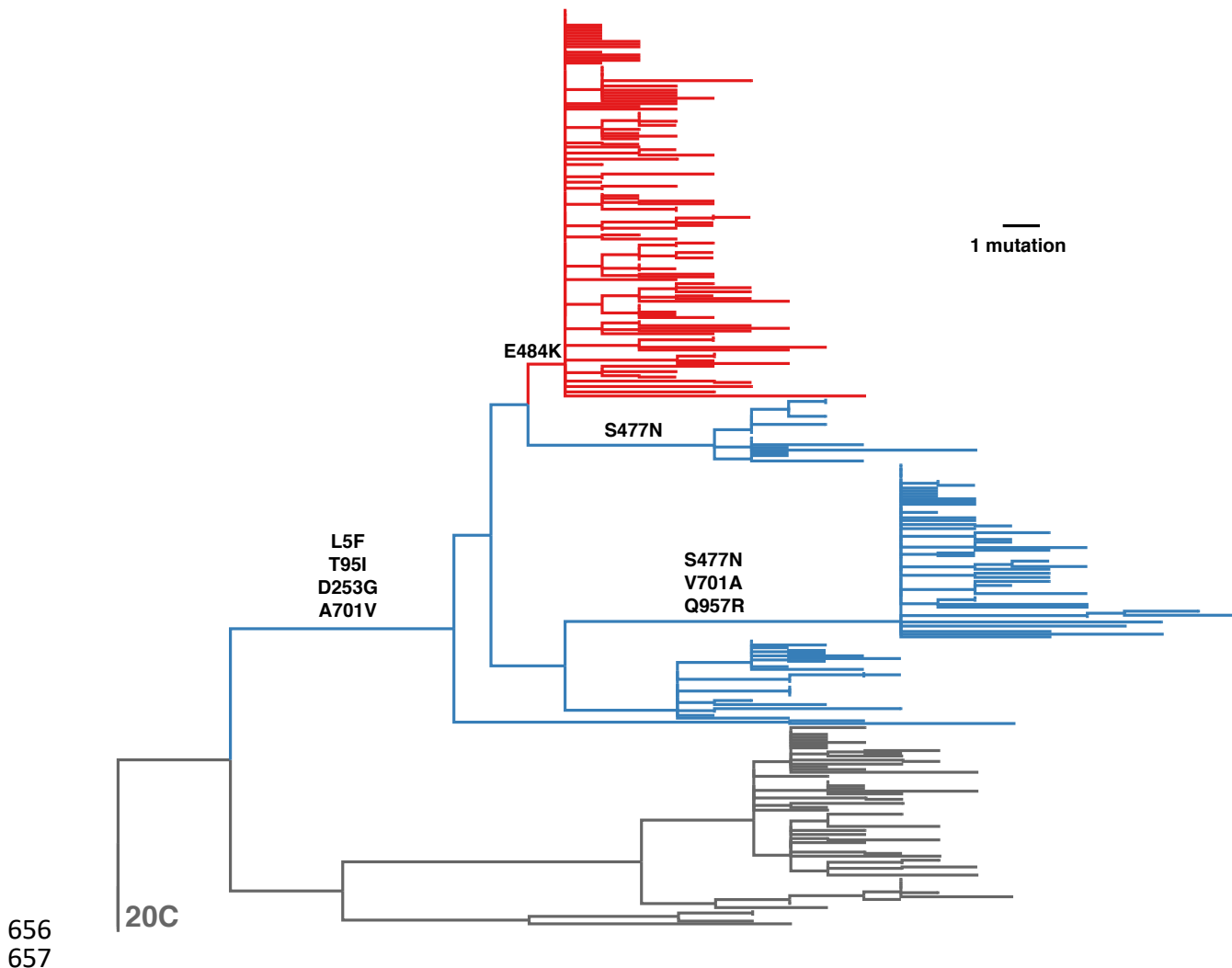
651 occurring on internal branches are labeled leading to and within B.1.526 are labeled. The

652 B.1.526 lineage is colored blue, except for the clade defined by the E484K mutation, which is

653 highlighted in red. The most common pattern of spike mutations in the sister clade is D80G,

654 Δ Y144, F157S, L452R, D614G, T859N, and D950H.

655



658 [Supplementary Table 1.](#)

659 List of 124 viral genomes (with their accession number, location, collection date, and spike

660 mutations) in lineage B.1.526. Mutations E484K, S477N, Q957R are highlighted in red, blue, and

661 cyan, respectively.

662

663 EPI_ISL_683762, USA/NY-NYCPHL-001443/2020-11-23 : L5F T95I D253G D614G A701V
664 EPI_ISL_823886, USA/NY-NYCPHL-001663/2020-12-08 : L5F T95I D253G **S477N** D614G **Q957R**
665 EPI_ISL_812733, USA/NY-NYCPHL-001804/2020-12-15 : L5F T95I D253G D614G A701V
666 EPI_ISL_765495, USA/NY-Wadsworth-290403-01/2020-12-16 : L5F T95I D253G **E484K** D614G A701V
667 EPI_ISL_765494, USA/NY-Wadsworth-290357-01/2020-12-18 : L5F T95I D253G **E484K** D614G A701V
668 EPI_ISL_765493, USA/NY-Wadsworth-290339-01/2020-12-19 : L5F T95I D253G D614G A701V
669 EPI_ISL_767598, USA/NY-Wadsworth-291999-01/2020-12-20 : L5F T95I D253G D614G A701V
670 EPI_ISL_767581, USA/NY-Wadsworth-292055-01/2020-12-21 : L5F T95I D253G D614G A701V
671 EPI_ISL_767595, USA/NY-Wadsworth-291994-01/2020-12-21 : L5F T95I D253G **S477N** D614G **Q957R**
672 EPI_ISL_832270, USA/NY-NYCPHL-001924/2020-12-23 : L5F T95I D253G **E484K** D614G A701V
673 EPI_ISL_832271, USA/NY-NYCPHL-001940/2020-12-24 : L5F T95I D253G **E484K** D614G A701V
674 EPI_ISL_861278, USA/NY-Wadsworth-21004976-01/2020-12-24 : L5F T95I D253G **E484K** D614G A701V
675 EPI_ISL_861280, USA/NY-Wadsworth-21004979-01/2020-12-24 : L5F T95I D253G **S477N** D614G A701V
676 EPI_ISL_861283, USA/NY-Wadsworth-21004982-01/2020-12-24 : L5F T95I D253G **S477N** D614G A701V
677 EPI_ISL_861258, USA/NY-Wadsworth-21004944-01/2020-12-25 : L5F T95I D253G **E484K** D614G A701V
678 EPI_ISL_861285, USA/NY-Wadsworth-21004984-01/2020-12-25 : L5F T95I D253G **E484K** D614G A701V
679 EPI_ISL_861244, USA/NY-Wadsworth-21004930-01/2020-12-25 : L5F T95I D253G **E484K** D614G A701V
680 EPI_ISL_802788, USA/NY-Wadsworth-21001942-01/2020-12-27 : L5F T95I D253G A475V **S477N** D614G **Q957R**
681 EPI_ISL_794226, USA/NY-Wadsworth-21000327-01/2020-12-27 : L5F T95I D253G **S477N** D614G **Q957R** D1260N
682 EPI_ISL_936036, USA/NY-Wadsworth-21007567-01/2020-12-27 : L5F T95I D253G **E484K** D614G A701V
683 EPI_ISL_854459, USA/NY-Wadsworth-21005068-01/2020-12-27 : L5F T95I D253G **E484K** D614G A701V
684 EPI_ISL_887843, USA/NJ-CDC-LC00000610/2020-12-27 : L5F T95I D253G F306L D614G
685 EPI_ISL_861315, USA/NY-Wadsworth-21005029-01/2020-12-28 : L5F T95I D253G **E484K** D614G A701V
686 EPI_ISL_888376, USA/IN-CDC-LC00001468/2020-12-28 : L5F T95I D253G T299I **E484K** D614G A701V
687 EPI_ISL_830720, USA/NY-Wadsworth-21004807-01/2020-12-28 : L5F T95I D253G D614G A701V
688 EPI_ISL_861308, USA/NY-Wadsworth-21005019-01/2020-12-28 : L5F T95I D253G **E484K** D614G A701V
689 EPI_ISL_861175, USA/NY-Wadsworth-21006727-01/2020-12-28 : L5F T95I D253G D614G A701V
690 EPI_ISL_936038, USA/NY-Wadsworth-21007571-01/2020-12-28 : L5F T95I D253G **E484K** D614G A701V
691 EPI_ISL_854458, USA/NY-Wadsworth-21005145-01/2020-12-29 : L5F T95I D253G D614G A701V
692 EPI_ISL_854457, USA/NY-Wadsworth-21005115-01/2020-12-29 : L5F T95I D253G **E484K** D614G A701V
693 EPI_ISL_857056, USA/NY-NYCPHL-002169/2020-12-29 : L5F T95I D253G **E484K** D614G A701V
694 EPI_ISL_861306, USA/NY-Wadsworth-21005017-01/2020-12-29 : L5F T95I D253G D614G A701V
695 EPI_ISL_802448, USA/NY-Wadsworth-21001323-01/2020-12-30 : L5F T95I D253G **S477N** D614G **Q957R**
696 EPI_ISL_861300, USA/NY-Wadsworth-21005007-01/2020-12-30 : L5F T95I D253G **E484K** D614G A701V
697 EPI_ISL_854450, USA/NY-Wadsworth-21005056-01/2020-12-30 : L5F T95I D253G **S477N** D614G **Q957R**
698 EPI_ISL_857204, USA/NY-NYCPHL-002114/2020-12-31 : L5F T95I D253G **E484K** D614G A701V
699 EPI_ISL_861112, USA/NY-Wadsworth-21002954-01/2021-01-01 : L5F T95I D198G D253G **S477N** D614G **Q957R**
700 EPI_ISL_861189, USA/NY-Wadsworth-21006379-01/2021-01-01 : L5F T95I D253G **E484K** D614G A701V
701 EPI_ISL_944591, USA/CT-Yale-1000/2021-01-01 : T95I D253G **E484K** D614G A701V
702 EPI_ISL_944592, USA/CT-Yale-1001/2021-01-01 : L5F T95I D253G **E484K** D614G A701V
703 EPI_ISL_944594, USA/CT-Yale-1003/2021-01-01 : L5F T95I D253G **E484K** D614G A701V
704 EPI_ISL_962492, Ecuador/UEES-9572/2021-01-01 : L5F T95I D253G **E484K** D614G A701V
705 EPI_ISL_962493, Ecuador/UEES-9602/2021-01-01 : L5F T95I D253G **E484K** D614G A701V

706 EPI_ISL_861113, USA/NY-Wadsworth-21002959-01/2021-01-02 : L5F T95I D198G D253G S477N D614G Q957R
707 EPI_ISL_861111, USA/NY-Wadsworth-21002949-01/2021-01-02 : L5F T95I D198G D253G S477N D614G Q957R
708 EPI_ISL_857060, USA/NY-NYCPHL-002190/2021-01-03 : L5F T95I D253G S477N D614G Q957R
709 EPI_ISL_861188, USA/NY-Wadsworth-21006373-01/2021-01-03 : L5F T95I D253G D614G A701V
710 EPI_ISL_896394, USA/NY-Wadsworth-21006803-01/2021-01-04 : L5F T95I D253G E484K D614G A701V
711 EPI_ISL_849348, USA/DE-DHSS-FLW00612372/2021-01-04 : L5F T95I D253G D614G A701V
712 EPI_ISL_896391, USA/NY-Wadsworth-21006765-01/2021-01-04 : L5F T95I D253G S477N D614G Q957R
713 EPI_ISL_896392, USA/NY-Wadsworth-21006780-01/2021-01-04 : L5F T95I D253G S477N D614G Q957R
714 EPI_ISL_830577, USA/NY-Wadsworth-21004146-01/2021-01-04 : L5F T95I D253G E484K D614G A701V
715 EPI_ISL_896395, USA/NY-Wadsworth-21006809-01/2021-01-04 : L5F T95I D253G E484K D614G A701V
716 EPI_ISL_861361, USA/NY-Wadsworth-21006106-01/2021-01-04 : L5F T95I D253G S477N D614G Q957R
717 EPI_ISL_830578, USA/NY-Wadsworth-21004150-01/2021-01-04 : L5F T95I D253G E484K D614G A701V
718 EPI_ISL_861411, USA/NY-Wadsworth-21006165-01/2021-01-06 : L5F T95I D253G S477N D614G Q957R D1260N
719 EPI_ISL_850743, USA/NJ-CDC-STM-A100415/2021-01-06 : L5F T95I D253G E484K D614G A701V
720 EPI_ISL_854449, USA/NY-Wadsworth-21006042-01/2021-01-07 : L5F T95I D253G S477N D614G Q957R
721 EPI_ISL_911789, USA/MD-HP01428/2021-01-07 : L5F T95I D253G S477N D614G Q957R
722 EPI_ISL_857120, USA/NY-NYCPHL-002295/2021-01-07 : L5F T95I D253G D614G A701V
723 EPI_ISL_855171, USA/RI-QDX-3232/2021-01-07 : L5F T95I D253G E484K D614G A701V
724 EPI_ISL_884080, USA/NY-Wadsworth-21003395-01/2021-01-07 : L5F T95I D253G E484K D614G A701V
725 EPI_ISL_896234, USA/NY-Wadsworth-21011146-01/2021-01-10 : L5F T95I D253G D614G A701V
726 EPI_ISL_883447, USA/NY-Wadsworth-21008849-01/2021-01-10 : L5F T95I D253G D614G A701V
727 EPI_ISL_896227, USA/NY-Wadsworth-21011153-01/2021-01-10 : L5F T95I D253G S477N D614G Q957R
728 EPI_ISL_886271, USA/PA-CDC-LC0003480/2021-01-10 : L5F T95I D253G D614G A701V
729 EPI_ISL_886626, USA/NJ-CDC-LC0003391/2021-01-11 : L5F T95I D253G E484K D614G A701V
730 EPI_ISL_857164, USA/NY-NYCPHL-002365/2021-01-11 : L5F T95I D253G E484K D614G A701V
731 EPI_ISL_886531, USA/CT-CDC-LC0003447/2021-01-11 : L5F T95I D253G E484K D614G A701V
732 EPI_ISL_857163, USA/NY-NYCPHL-002336/2021-01-11 : L5F T95I D253G E484K D614G A701V
733 EPI_ISL_886240, USA/NJ-CDC-LC0003405/2021-01-11 : L5F T95I D253G D614G A701V
734 EPI_ISL_884055, USA/NY-Wadsworth-21005245-01/2021-01-11 : L5F T95I D253G S477N D614G A701V
735 EPI_ISL_886384, USA/GA-CDC-LC0003561/2021-01-11 : L5F T95I D253G E484K D614G A701V
736 EPI_ISL_886270, USA/DE-CDC-LC0003883/2021-01-12 : L5F T95I D253G D614G A701V
737 EPI_ISL_936267, USA/NY-Wadsworth-21012072-01/2021-01-13 : L5F T95I D253G D614G A701V
738 EPI_ISL_936292, USA/NY-Wadsworth-21012249-01/2021-01-14 : L5F T95I D253G E484K D614G A701V
739 EPI_ISL_936261, USA/NY-Wadsworth-21012065-01/2021-01-14 : L5F T95I D253G E484K D614G A701V
740 EPI_ISL_883338, USA/NY-NYCPHL-002433/2021-01-14 : L5F T95I D253G E484K D614G A701V
741 EPI_ISL_883342, USA/NY-NYCPHL-002473/2021-01-14 : L5F T95I D253G V483F E484K D614G A701V
742 EPI_ISL_883339, USA/NY-NYCPHL-002440/2021-01-14 : L5F T95I D253G S477N D614G A701V
743 EPI_ISL_883340, USA/NY-NYCPHL-002466/2021-01-14 : L5F T95I D253G V483F E484K D614G A701V
744 EPI_ISL_936288, USA/NY-Wadsworth-21012244-01/2021-01-14 : L5F T95I D253G E484K D614G A701V
745 EPI_ISL_883343, USA/NY-NYCPHL-002474/2021-01-14 : L5F T95I D253G V483F E484K D614G A701V
746 EPI_ISL_883346, USA/NY-NYCPHL-002525/2021-01-14 : L5F T95I D253G S477N D614G A701V
747 EPI_ISL_883341, USA/NY-NYCPHL-002471/2021-01-14 : L5F T95I D253G E484K D614G A701V
748 EPI_ISL_883330, USA/NY-NYCPHL-002514/2021-01-15 : L5F T95I D253G S477N D614G Q957R
749 EPI_ISL_883347, USA/NY-NYCPHL-002533/2021-01-15 : L5F T95I D253G S477N D614G A701V
750 EPI_ISL_936299, USA/NY-Wadsworth-21012257-01/2021-01-15 : L5F T95I D253G S477N D614G Q957R
751 EPI_ISL_883331, USA/NY-NYCPHL-002535/2021-01-15 : L5F T95I D253G S477N D614G A701V
752 EPI_ISL_937272, USA/NY-NYCPHL-002846/2021-01-15 : L5F T95I D253G E484K D614G A701V
753 EPI_ISL_936281, USA/NY-Wadsworth-21012236-01/2021-01-15 : L5F T95I D253G D614G A701V
754 EPI_ISL_883344, USA/NY-NYCPHL-002520/2021-01-16 : L5F T95I D253G S477N D614G A701V
755 EPI_ISL_937190, USA/NY-NYCPHL-002680/2021-01-17 : L5F T95I D253G E484K D614G A701V
756 EPI_ISL_883345, USA/NY-NYCPHL-002521/2021-01-17 : L5F T95I D253G E484K D614G A701V
757 EPI_ISL_936248, USA/NY-Wadsworth-21011901-01/2021-01-17 : L5F T95I D253G E484K D614G A701V
758 EPI_ISL_936271, USA/NY-Wadsworth-21012077-01/2021-01-17 : L5F T95I D253G S477N D614G Q957R

759 EPI_ISL_936256, USA/NY-Wadsworth-21012058-01/2021-01-18 : L5F T95I D253G **S477N** D614G **Q957R**
760 EPI_ISL_936254, USA/NY-Wadsworth-21011909-01/2021-01-18 : L5F T95I D253G **E484K** D614G A701V
761 EPI_ISL_936291, USA/NY-Wadsworth-21012248-01/2021-01-18 : L5F T95I D253G **E484K** D614G A701V
762 EPI_ISL_896233, USA/NY-Wadsworth-21011134-01/2021-01-18 : L5F T95I D253G **E484K** D614G A701V
763 EPI_ISL_937191, USA/NY-NYCPHL-002681/2021-01-18 : L5F T95I D253G **E484K** D614G A701V
764 EPI_ISL_936276, USA/NY-Wadsworth-21012089-01/2021-01-19 : L5F T95I D253G **S477N** D614G **Q957R**
765 EPI_ISL_937126, USA/NY-NYCPHL-002548/2021-01-19 : L5F T95I D253G **S477N** D614G A701V
766 EPI_ISL_937192, USA/NY-NYCPHL-002682/2021-01-20 : L5F T95I D253G **E484K** D614G A701V
767 EPI_ISL_937176, USA/NY-NYCPHL-002659/2021-01-21 : L5F T33I T95I D253G D614G A701V
768 EPI_ISL_920158, England/LOND-12F79EE/2021-01-21 : L5F T95I D253G D614G A701V
769 EPI_ISL_962431, USA/WA-S4338/2021-01-21 : L5F T95I D253G D614G A701V
770 EPI_ISL_937205, USA/NY-NYCPHL-002704/2021-01-22 : L5F T95I D253G **S477N** D614G **Q957R**
771 EPI_ISL_937211, USA/NY-NYCPHL-002711/2021-01-22 : L5F T95I D253G **E484K** D614G A701V
772 EPI_ISL_937201, USA/NY-NYCPHL-002698/2021-01-22 : L5F T95I D253G D614G A701V
773 EPI_ISL_937213, USA/NY-NYCPHL-002714/2021-01-22 : L5F L18F T95I D253G **S477N** D614G **Q957R**
774 EPI_ISL_937246, USA/NY-NYCPHL-002784/2021-01-22 : L5F T95I D253G **E484K** D614G A701V
775 EPI_ISL_937247, USA/NY-NYCPHL-002785/2021-01-23 : L5F T95I D253G **E484K** D614G A701V
776 EPI_ISL_937258, USA/NY-NYCPHL-002831/2021-01-24 : L5F T95I D253G **E484K** D614G A701V
777 EPI_ISL_937236, USA/NY-NYCPHL-002767/2021-01-25 : L5F T95I D253G **S477N** D614G A701V
778 EPI_ISL_937218, USA/NY-NYCPHL-002729/2021-01-25 : L5F T95I D253G **E484K** D614G A701V
779 EPI_ISL_937216, USA/NY-NYCPHL-002723/2021-01-25 : L5F T95I D253G **E484K** D614G A701V
780 EPI_ISL_937265, USA/NY-NYCPHL-002839/2021-01-25 : L5F T95I D253G **E484K** D614G A701V
781 EPI_ISL_937215, USA/NY-NYCPHL-002720/2021-01-25 : L5F T95I D253G **E484K** D614G A701V
782 EPI_ISL_937255, USA/NY-NYCPHL-002793/2021-01-25 : L5F T95I D253G **E484K** D614G A701V
783 EPI_ISL_906839, Singapore/117/2021-01-26 : L5F T95I D253G **E484K** D614G A701V
784 EPI_ISL_906838, Singapore/116/2021-01-26 : L5F T95I D253G **E484K** D614G A701V
785 EPI_ISL_906837, Singapore/115/2021-01-26 : L5F T95I D253G **E484K** D614G A701V
786 EPI_ISL_937477, USA/ME-HETL-J1185/2021-01-29 : L5F T95I D253G **E484K** D614G A701V
787