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# Investigation of COVID-19 Outbreak among Wildland Firefighters during Wildfire Response, Colorado, USA, 2020

Amanda Reiff Metz, Matthew Bauer, Chelsey Epperly, Ginger Stringer, Kristen E. Marshall, Lindsey Martin Webb, Molly Hetherington-Rauth, Shannon R. Matzinger, Sarah Elizabeth Totten, Emily A. Travanty, Kristen M. Good, Alexis Burakoff

A COVID-19 outbreak occurred among Cameron Peak Fire responders in Colorado, USA, during August 2020–January 2021. The Cameron Peak Fire was the largest recorded wildfire in Colorado history, lasting August–December 2020. At least 6,123 responders were involved, including 1,260 firefighters in 63 crews who mobilized to the fire camps. A total of 79 COVID-19 cases were identified among responders, and 273 close contacts were quarantined. State and local public health investigated the outbreak and coordinated with wildfire management teams to prevent disease spread. We performed whole-genome sequencing and applied social network analysis to visualize clusters and transmission dynamics. Phylogenetic analysis identified 8 lineages among sequenced specimens, implying multiple introductions. Social network analysis identified spread between and within crews. Strategies such as implementing symptom screening and testing of arriving responders, educating responders about overlapping symptoms of smoke inhalation and COVID-19, improving physical distancing of crews, and encouraging vaccinations are recommended.

The Cameron Peak Fire in Colorado, USA, began on August 13, 2020. Because of the magnitude of this wildfire, the response was coordinated by various Incident Management Teams (IMT); wildfire

responders included Colorado wildland firefighter crews as well as crews from around the country deployed to Colorado for the response. On August 25, 2020, the Larimer County Department of Health and Environment (LCDHE) and the Colorado Department of Public Health and Environment (CDPHE) received notification of a wildland firefighter responding to the Cameron Peak Fire who tested positive for SARS-CoV-2, the virus that causes COVID-19. This firefighter initially reported difficulty breathing and was transported to the local emergency department, then released. The next day, he was admitted to the hospital for continuing symptoms and tested positive for SARS-CoV-2 by reverse transcription PCR (RT-PCR). LCDHE, in partnership with the IMT, began contact tracing on the basis of the Centers for Disease Control and Prevention (CDC) definition of someone who was within 6 feet of an infected person for a cumulative 15 minutes or more over a 24-hour period (1). Two persons working on the same crew and 5 additional responders at the camp were identified as close contacts and quarantined. During contact interviews, it was reported that 2 crew members of the index case-patient were experiencing cough and headaches; both subsequently tested positive for SARS-CoV-2. An outbreak was declared and reported on September 2, 2020.

Wildfire response personnel operating across the state were in contact with CDPHE throughout the wildfire season regarding COVID-19 prevention and response plans. In July 2020, before the Cameron Peak Fire, CDPHE released public guidance documents addressing best practices for mitigating COVID-19 risks at wildfire camps (2). This document supplemented best practice guidance available from other sources

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Author affiliations: Colorado Department of Public Health and Environment, Denver, Colorado, USA (A.R. Metz, C. Epperly, G. Stringer, K.E. Marshall, L.M. Webb, M. Hetherington-Rauth, S.R. Matzinger, S.E. Totten, E.A. Travanty, K.M. Good, A. Burakoff); Larimer County Department of Health and Environment, Fort Collins, Colorado, USA (M. Bauer); Centers for Disease Control and Prevention, Atlanta, Georgia, USA (K.E. Marshall)

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such as CDC (3), United States Forest Service (USFS) (4), the Fire Management Board (5), and United States Department of the Interior (6). At the time that the Cameron Peak Fire started, a CDPHE occupational health epidemiologist regularly attended a morning safety briefing call organized by the USFS, in which incident management team representatives from all active fires in Colorado called in with updates on safety concerns including COVID-19.

**Methods**

**Case Investigations**

LCDHE and the Cameron Peak IMT collaborated to conduct case investigations and contact tracing activities. An outbreak case was defined as confirmed or probable COVID-19 (determined using the Council of State and Territorial Epidemiologists’ 2020 Interim COVID-19 Case Definition) (7) in a responder who was onsite at the Cameron Peak Fire within 14 days of symptom onset or positive test. Close contacts were identified on the basis of the CDC definition and quarantined. CDPHE and local hospital laboratories conducted SARS-CoV-2 RT-PCR testing using various platforms.

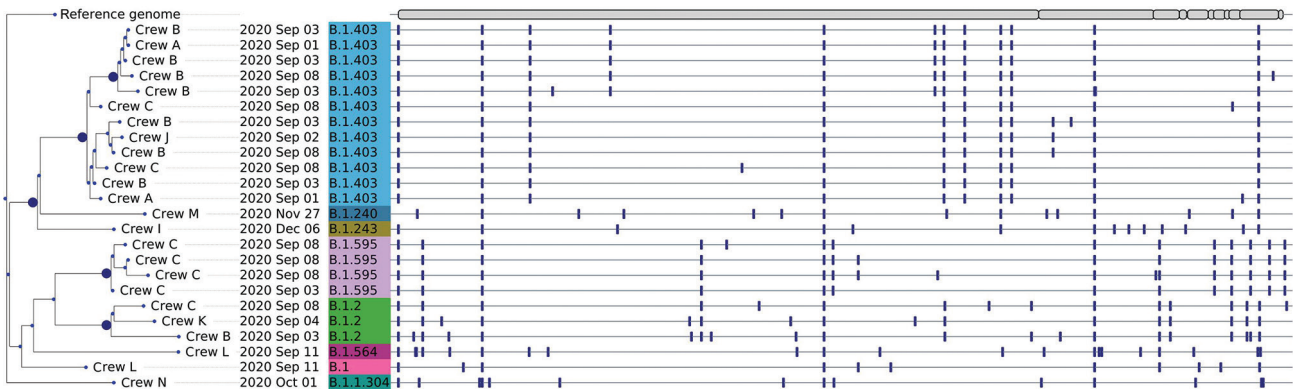
Outbreak response consultation calls among CDPHE, LCDHE, and IMT were held to provide recommendations for isolation of cases, quarantine of close contacts, and prevention practices such as improving physical distancing. CDPHE’s Rapid

Response Team hosted a testing event for Cameron Peak Fire responders before the first positive case was identified; surveillance and outbreak screening testing was offered to all Cameron Peak Fire responders starting August 24. Once the outbreak was identified, widespread testing was conducted 11 more times during August 26–October 25, 2020. After the fire, the USFS conducted a Facilitated Learning Analysis to identify lessons learned from the outbreak response (8).

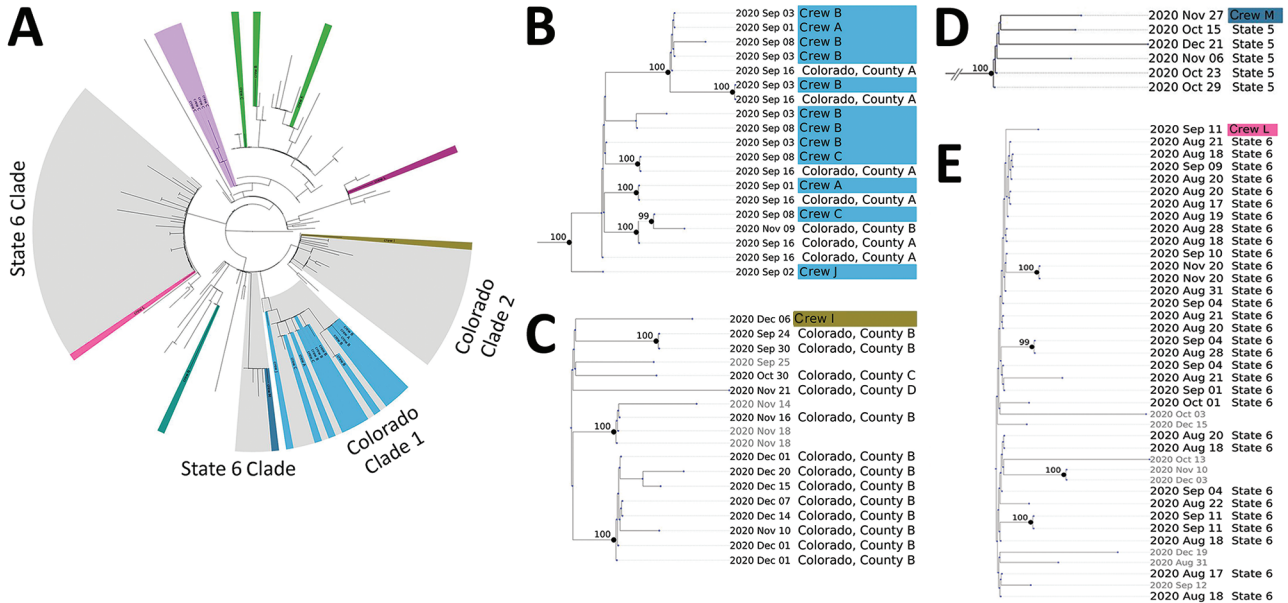
**Whole-Genome Sequencing**

CDPHE performed tiled amplicon whole-genome sequencing (WGS) on 40 (51%) available specimens from wildfire responders (Appendix, <https://wwwnc.cdc.gov/EID/article/28/8/22-0310-App1.pdf>); the remainder of the specimens were unavailable for sequencing because they were not sent to the CDPHE laboratory. We assembled sequencing data by using the Monroe workflow and CDPHE’s publicly available Nanopore data workflow (<https://github.com/CDPHE>).

Of the specimens available for WGS, 24 resulted in sequence determination; we used those sequences to construct a focal phylogenetic tree of the Cameron Peak Fire outbreak (Figure 1). In addition, to investigate the potential for multistate lineage introduction or community transmission, we constructed a contextual phylogenetic tree by using the 24 whole-genome sequences of the Cameron Peak Fire specimens and additional



**Figure 1.** Phylogenetic tree of SARS-CoV-2 consensus whole-genome sequences from 24 of 42 positive specimens from Cameron Peak firefighters available at the Colorado State Public Health Lab with ≥89% genome coverage. Nodes with at least 95% ultrafast bootstrap support are labeled. Firefighter crew, sample collection date, and lineage are displayed at the tips. A visualization of the reference genome is depicted at the top of the phylogeny. Vertical bars shown across each consensus sequence indicate positions of nucleotide changes relative to the reference genome. High-quality consensus sequences were defined as sequences with ≥89% genome coverage (10× sequence coverage depth for Illumina [<https://www.illumina.com>] and 20× for Oxford Nanopore [<https://nanoporetech.com>]) and minimum base quality of 20. Prior to phylogenetic inference, consensus sequences were aligned to the reference genome (Genbank accession no. NC\_045512.2), and insertions were removed so that all sequences were 29,903 nt in length. Phylogenetic inference of the consensus sequences was performed using IQTree version 2.0.3 (<http://www.iqtree.org>) with 1,000 ultrafast bootstrap replicates and phylogenetic tree visualization was performed using the python module ete3 version 3.1.2 (<https://pypi.org/project/ete3>). Pangolin v.2.4.2<sup>5</sup> (9) and Nextstrain’s Nextclade tools (10) were used to assign lineage and clade designations to each assembled genome.



**Figure 2.** Contextual phylogenetic tree and enlarged clades showing genetic relatedness of the Cameron Peak firefighter sequences to sequences of SARS-CoV-2 collected within the United States during September–December 2020. A) Full contextual tree constructed using 754 contextual sequences subsampled from GISAID (<https://www.gisaid.org>) plus 24 Cameron Peak firefighter consensus sequences. The phylogeny has been pruned to display 164 contextual sequences and Cameron Peak firefighter sequences. Cameron Peak sequences are highlighted in color according to their lineage assignment. Clades highlighted in gray represent potential community and interstate transmission events. Cameron Peak sequences assigned to lineage B.1.2 (green) do not cluster together on the contextual phylogeny to form a monophyletic group, suggesting that they are genetically divergent from one another and likely do not represent a single transmission event, despite belonging to the same lineage. Mutation differences among these sequences are shown in detail in Figure 1. B) Colorado clade 1. Twelve Cameron Peak firefighters formed a monophyletic group with sequences from 2 Colorado counties. C) Colorado clade 2. A single Cameron Peak firefighter sequence formed a clade with sequences collected from 3 Colorado counties and additional sequences collected from outside of Colorado (not labeled). Low support values for this clade may be expected because of low sequence diversity. D) State 5 clade. The Cameron Peak firefighter sequence formed a monophyletic clade with sequences collected from his or her state of deployment (State 5). E) State 6 clade. The Cameron Peak firefighter sequence formed a clade with sequences collected from his or her state of deployment (state 6) and additional sequences collected from outside of Colorado and not from his or her state of deployment (not labeled). Low support values for this clade may be caused by low sequence diversity. For panels B–E, all sequences within a clade are assigned the same lineage. Collection dates are labeled for all tips. Cameron Peak firefighter sequences are highlighted according to their lineage and labeled with crew. Nodes with at least 95% ultrafast bootstrap support values are labeled. Additional information is available in the Appendix (<https://wwwnc.cdc.gov/EID/article/28/8/22-0310-App1.pdf>).

whole-genome sequences that were either publicly available or additionally sequenced at the CDPHE State Public Health Laboratory (Figure 2; Appendix).

**Social Network Analysis**

We conducted social network analysis of all SARS-CoV-2–positive responders by using R Studio version 1.2.5033 (<https://www.rstudio.com>) and Gephi Graph Visualization and Manipulation software version 0.9.2 (<https://gephi.org>). We applied this analysis to WGS results to visualize clusters and transmission dynamics among Cameron Peak Fire crews (11). We assumed epidemiologic links of exposure between responders belonging to the same crew for network construction. Data showing potential exposure outside of crew assignments (i.e., socializing with members of other crews) were not available. This

activity was reviewed by CDC and was conducted consistent with applicable federal law and CDC policy (45 C.F.R. part 46, 21 C.F.R. part 56; 42 U.S.C. Sect. 241(d); 5 U.S.C. Sect. 552a; 44 U.S.C. Sect. 3501 et seq).

**Results**

The outbreak among wildfire responders occurred during August 25, 2020–January 8, 2021. (In Colorado, an outbreak is considered resolved 28 days after symptom onset of the last case.) A total of 6,123 responders were involved in the response. We identified 79 cases (78 confirmed and 1 probable); 73 of these were confirmed to be firefighters from 1 of the 63 crews, for an attack rate of 5.8% among 1,260 firefighters who were deployed full-time to the incident (Figure 3). The remainder of responder case-patients were persons from IMT, equipment operators, and





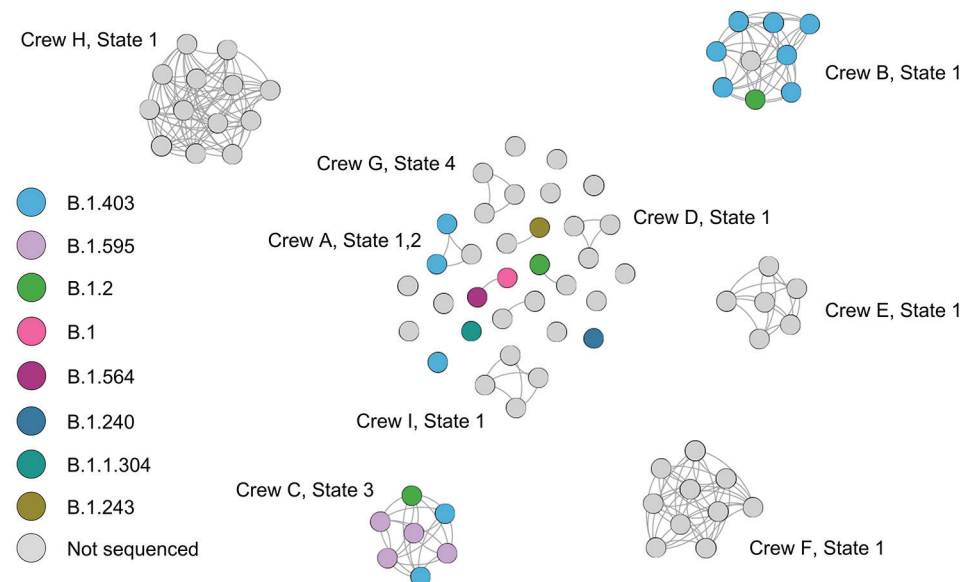
(Figure 2, panel A). The analysis revealed 4 clades that provided evidence of possible intrastate and interstate transmission (Figure 2, panels B–E). Twelve Cameron Peak Fire sequences formed a monophyletic clade with sequences collected from 2 Colorado counties with high support values (ultrafast bootstrap support >95% for nodes; Figure 2, panel B). Another sequence from a Cameron Peak Fire responder formed a clade with sequences collected from 3 Colorado counties and additional sequences collected from outside of Colorado but with low support values (ultrafast bootstrap support <95% for nodes; Figure 2, panel C). In addition, in 2 cases, sequences from 2 different responders formed a clade with contextual sequences collected from their state of deployment; 1 clade was supported with high support values but the other was not (Figure 2, panels D and E). Although not all clades were supported with high bootstrap values, low support values might be expected if sequence diversity is insufficient, which could result from either low diversity of SARS-CoV-2 circulating in the United States at the time, or low diversity among samples that were able to be sequenced and deposited in public repositories. Short branch lengths as observed on the tree are indicative of low divergence among sequences (12).

Social network analysis showed the 79 responders with COVID-19 clustered into 26 crews deploying from 17 states (Figure 4). Nine crews with responders from 10 states experienced  $\geq 3$  cases. We observed multiple lineages within single crews, suggesting multiple points of introduction, probable crew intermingling, and possible lapses in prevention measures such as social distancing.

**Discussion**

The Cameron Peak Fire was the largest recorded wildfire in Colorado’s history, burning 208,913 acres. A total of 79 cases of COVID-19 were identified among Cameron Peak Fire responders deployed from 17 states. Multiple points of SARS-CoV-2 introduction were likely because of frequent crew turnover as the wildfire grew, as suggested by WGS and social network analysis results.

Balancing management of a large-scale wildfire and control of COVID-19 among responders created several challenges for disease prevention and mitigation. Frequent responder turnover because of 2- to 3-week deployments, combined with the length of the fire, resulted in continuous opportunities for introduction of COVID-19 into wildfire camps (13). COVID-19 testing was available for incoming responders, but no testing or quarantine was required upon arrival, and no surveillance testing was required during the deployment period. In addition, turnover of responders resulted in several instances in which case-patients in isolation or contacts in quarantine were demobilized back to their home states or deployed to other wildfire responses before case investigation and contact tracing could be completed. In these situations, CDPHE notified the states to which responders were demobilized, and LCDHE coordinated with Cameron Peak IMT to ensure these responders were immediately notified and given instructions to proceed home immediately, avoiding contact with others and stops in indoor public settings during their travel. However, the potential for multistate spread was a major concern when responders were demobilized and sent home or to other responses.



**Figure 4.** Social network analysis of Cameron Peak firefighter crews with COVID-19, Colorado, USA, August–December 2020. All responders testing positive for SARS-CoV-2 (nodes) are included in this figure to show contact within crews (edges). Crews with  $\geq 3$  firefighters positive with SARS-CoV-2 are labeled.

The difficulty of screening responders for COVID-19 symptoms was compounded by challenges differentiating the effects of smoke and high altitude from symptoms of COVID-19. Smoke inhalation can cause several respiratory symptoms that are similar to COVID-19, including coughing, shortness of breath, sore throat, and chest pain (14). Altitude sickness symptoms also overlap with COVID-19 symptoms and can include headaches, fatigue, nausea, and vomiting, as well as, in more severe cases, shortness of breath, weakness, and cough (15). Symptoms of acute and chronic smoke exposure overlap with and can worsen COVID-19 symptoms, complicating symptom-based identification of COVID-19 (13,16). Elevations in the fire-affected area ranged from  $\approx$ 5,200 feet to >10,000 feet, resulting in the potential for altitude sickness for crews, particularly those coming to Colorado from states at lower elevations.

Often, responders continued to work while they were symptomatic and infectious and did not report symptoms until their illness became severe or they experienced a distinguishing symptom, such as loss of taste or smell. COVID-19 mitigation was further challenged by how fire camps were set up, potentially increasing exposure opportunities. Crews often camped together or worked geographically closely before implementation of mitigation and quarantine measures, potentially increasing exposure opportunities. Furthermore, because these camps were often located in areas with limited cell service, Wi-Fi hotspots provided relatively small areas where responders could access Wi-Fi, creating additional opportunities for exposure when responders gathered closely together in areas where Wi-Fi was available (9). Other barriers to the public health response included some responders' distrust of their positive SARS-CoV-2 test results because of lack of symptoms or overlap with smoke inhalation symptoms. Further, many responders were employed as contractors and were not provided paid sick leave to cover quarantine or isolation. Fire response coordinators and commanders indicated that some crew members might have been hesitant to report symptoms or get tested because of concerns over having to quarantine or isolate without pay. Challenges in gathering complete symptom information could be caused by responders' reluctance to be pulled from their crew, which could further strain resources during the response. Contact tracing was challenging early in the investigation because case-patients were unable to identify their close contacts or unwilling to provide names of close contacts to avoid quarantine. Further, responders and response commanders were resistant to implementing full

quarantines because staffing needs were strained by the severity of the Cameron Peak Fire and other wildfires happening concurrently in the region. Critical infrastructure-modified quarantine and testing-based strategies were used when full quarantines were not feasible, including release from quarantine after a negative RT-PCR result from a specimen collected 7 days after exposure (which was not a recommended practice under standard quarantine guidance at that time) or monitoring responders for symptoms while allowing them to continue working during quarantine (17).

The results of WGS and social network analysis suggest multiple SARS-CoV-2 introduction events throughout the wildfire response, as well as spread both between and within crews. The presence of sequences from a single lineage in >1 crew combined with near-identical nucleotide changes observed among these sequences suggest intercrew transmission or transmission between fire crews and nearby communities (Figure 2, panel B). Contextual analysis suggests possible transmission events linked to Cameron Peak Fire responders from both outside and within the state of Colorado; in a few instances, analysis suggested transmission from the state from which an individual was deployed and in other instances from surrounding counties within the state of Colorado. One state deployment introduction (state 5) and 1 Colorado county introduction (Colorado county A) are well supported by bootstrapping, but in the other 2 instances, support was weak. This result of low sequence diversity across many states present in sequences available in public repositories from this time period.

The first limitation of our study is that COVID-19 cases were likely underreported because of insufficient testing and lack of reporting of symptoms by responders. Surveillance testing was optional and the overlap between COVID-19 symptoms and symptoms associated with smoke inhalation and altitude sickness might have led some persons not to get tested when symptomatic. Second, only 51% of outbreak-related specimens were available for WGS because not all specimens were sent to the CDPHE laboratory, including those collected through the local hospital; therefore, results might not be complete. Finally, social network analysis epidemiologic links were assumed for responders on the same crew but lacked more robust data showing intercrew mingling during and outside of response activities.

Many lessons were learned in this COVID-19 outbreak during a wildfire response. Open communication between fire response agencies and public health agencies enabled enhanced prevention strategies. Fire

response agencies should consider symptom screening and testing of all arriving responders to limit introduction of SARS-CoV-2 into fire camps; educating responders about potentially overlapping symptoms of smoke inhalation, COVID-19, and altitude (when relevant); and improving physical distancing of crews onsite. Surveillance testing offers the ability to detect cases early and to prevent transmission before an outbreak occurs (18). Rapid testing options, such as the use of rapid antigen tests, can provide many benefits in wildfire response and other emergency management settings, including quick turnaround of results, which can minimize the need to quarantine critical responders while awaiting results; encouraging action in response to mild symptoms that might otherwise be dismissed as the result of smoke or altitude, because it is a quick and easy option to differentiate symptoms; and ease of implementation in remote and nonmedical settings, not requiring transport of persons off-site or coordination with nearby medical facilities. Response agencies should work with jurisdictional public health agencies at the beginning of each response to determine what testing options are currently available and how best to implement testing of responders. Rapidly identifying cases would lead to timely case investigations and contact tracing activities that could help mitigate spread of disease by enabling timely isolation of case-patients and quarantine of close contacts. Policies to compensate responders for time spent in isolation or quarantine could improve compliance with testing and screening procedures. During the response, fire response agencies recommended mask use, especially when other social distancing measures were difficult to maintain. Continuing the use of masks in indoor settings or close interactions with others could be considered in areas of high transmission even in the absence of local public health requirements. In current and future fire seasons, we encourage COVID-19 vaccination and surveillance testing, particularly given the challenges of implementing other mitigation techniques in resource-constrained fire responses. Response agencies should consider collaborating with public health agencies to ensure that appropriate disease control measures are put in place when COVID-19 has been identified among responders, including encouraging cooperation of persons who are identified as case-patients or close contacts to prevent the spread of disease. The lessons learned during this outbreak can contribute to developing best practices for managing wildfire response and outbreaks of COVID-19 and other communicable diseases among responders to large-scale emergency events.

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## About the Author

Ms. Metz works at the Colorado Department of Public Health and Environment as a COVID-19 Epi Response Team Unit Manager for metropolitan area counties in Colorado. Her research interests include infectious disease epidemiology and outbreak investigations.

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Address for correspondence: Amanda Metz, Colorado Department of Public Health and Environment, 4300 Cherry Creek Dr S, Denver, CO 80246, USA; email: [amanda.metz@state.co.us](mailto:amanda.metz@state.co.us)



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## Appendix

### Sequencing and Sequence Assembly

RNA was isolated from nasopharyngeal swab specimens using the MagMAX Viral/Pathogen Nucleic Acid Isolation Kit (Thermo Scientific, Waltham, MA, USA) with the 200  $\mu$ L sample input volume automated method using the KingFisher Flex Magnetic Particle Processor (Thermo Scientific, Waltham, MA, USA) in accordance with the TaqPath COVID-19 Combo Kit (Thermo Scientific, Waltham, MA, USA) (1). Samples were then screened with PCR for sequencing viability using the TaqPath COVID-19 Combo Kit PCR assay (Thermo Scientific, Waltham, MA, USA). Specimens were then sequenced on either Illumina MiSeq (Illumina, San Diego, California, USA) or Oxford Nanopore Technology (ONT) GridION (Oxford Nanopore Technologies, Oxford, UK) sequencing platforms with slight modifications following the ARTIC V3 tiled PCR amplicon sequencing protocol for Illumina (2) and ONT (3) respectively for cDNA preparation and tiled PCR amplification using the ARTIC V3 primer set. For specimens sequenced on Illumina MiSeq, we used the Illumina DNA Prep Kit (Illumina, San Diego, California, USA) for library preparation. For specimens sequenced on ONT GridION, we followed the ARTIC V3 protocol native barcoding method for library preparation. For Illumina MiSeq we used 150 bp paired-end reads and for ONT we used 400–700 bp reads. Each sequencing plate contained a non-template negative control.

We performed reference based viral genome assembly for both Illumina data and ONT gridiron data. For Illumina data we performed a reference based viral genome assembly in two parts. First we used the Monroe pipeline for the alignment of sequencing reads to the reference genome ([https://staphb.org/staphb\\_toolkit/workflow\\_docs/monroe/](https://staphb.org/staphb_toolkit/workflow_docs/monroe/); [https://github.com/StaPH-B/staphb\\_toolkit](https://github.com/StaPH-B/staphb_toolkit)). We then generated the consensus genome using the *consensus* function from

the package iVar (<https://github.com/andersen-lab/ivar>) (4), using a minimum allele frequency of 0.6, a minimum sequencing depth of 10x and a minimum base quality of 30. For ONT GridION data, we performed reference based viral genome assembly using a custom workflow based on the ARTIC bioinformatics nanopore protocol (<https://artic.network/ncov-2019/ncov2019-bioinformatics-sop.html>) and implemented on the Terra.bio cloud computer platform. Briefly, we filtered reads by length and removed low quality reads using guppy plex. Primer sequences were removed from reads, reads were mapped to the reference genome, and consensus genome sequences were generated using medaka as implemented in the ARTIC bioinformatics protocol.

For both Illumina and ONT data, we assessed the average sequencing depth and percent genome coverage of each consensus genome using custom python scripts ([https://github.com/CDPHE/python\\_scripts\\_for\\_terra\\_workflows](https://github.com/CDPHE/python_scripts_for_terra_workflows)). We removed insertions not previously documented in the genome as these were likely due to sequencing and/or assembly error using custom python scripts ([https://github.com/CDPHE/sars-cov-2\\_indel\\_finder](https://github.com/CDPHE/sars-cov-2_indel_finder)). We defined high quality assembled genomes as those with at least 90 percent coverage across the reference genome. In cases where we could not obtain high quality sequences from the specimen, it could have been due to low viral titer at the time of specimen collection resulting in low RT-PCR Ct values and/or specimen degradation.

All high quality assembled consensus genomes are published to the GISAID and NCBI Genbank repositories and sequencing read data has been published to the NCBI SRA repository (Appendix Tables 1, 2).

## **Lineage Characterization and Phylogenetic Analyses**

To determine relatedness among the Cameron Peak Fire consensus genome sequences and to infer possible transmission events, we first determined the lineage of each sequence using the software PANGOLIN v3.1.1 (<https://github.com/cov-lineages/pangolin>). Next we constructed a focal phylogenetic tree using the Cameron Peak Fire consensus genome sequences. We aligned consensus genomes using MAFFT v7.471 (5) and constructed a maximum likelihood tree using IQ-Tree v.1.6.1 (<http://www.iqtree.org>) using 1000 ultrafast bootstrap replicates. We visualized the phylogeny using the Python library ete3 v3.1.2 (<http://etetoolkit.org/>) and custom python scripts.

To investigate the potential for multistate lineage introduction and/or community transmission, we constructed a contextual phylogenetic tree. We first downloaded sequences from GISAID meeting the following criteria: 1) samples collected from human hosts, 2) sequences with complete coverage, 3) samples collected within the United States with locality information to at least the level of state, and 4) samples collected between August 15, 2020 and December 21, 2020, corresponding to 2 weeks before and 2 weeks following the Cameron Peak Fire outbreak (data downloaded July 7, 2021). This resulted in 44,814 sequences. To these sequences we added an additional 872 sequences from samples collected in Colorado between August 15, 2020 and December 21, 2020 that were sequenced at the CDPHE State Public Health Laboratory and for which we had county level collection localities. This resulted in a total of 45,686 contextual sequences. Next to determine which sequences among our contextual sequences were most genetically similar to our focal Cameron Peak Fire sequences, we used the *priorities.py* python script (6) from the NEXTSTRAIN workflow for SARS-CoV-2 (<https://github.com/nextstrain/ncov>). This script ranks sequences in order of their similarity to the set of focal sequences and is used as input for the AUGUR *filter* subcommand (<https://github.com/nextstrain/augur>). Next we filtered the contextual sequences using the *filter* subcommand from AUGUR v.12.0.0. We set the subsampling parameters to group samples by collection date and subsample a max of 1,000 sequences and set the minimum sequence length to 27,000 bp. This subsampling resulted in a total of 778 sequences (717 sequences from GISAID, 37 additional Colorado sequences sequenced at CDPHE State Public Health Laboratory, and 24 focal Cameron Peak Fire sequences) used to build our contextual phylogenetic tree. We aligned the sequences using MAFFT v7.471 and constructed a maximum likelihood tree using IQ-Tree v.1.6.1 using 1000 ultrafast bootstrap replicates. We visualized the phylogeny using the Python library ete3 v3.1.2 and custom python scripts. To help with visualization we pruned the final contextual phylogenetic tree to 200 sequences, while maintaining sequences forming monophyletic groups with the 24 Cameron Peak Fire sequences.

**Additional Information about Figure 3 (<https://wwwnc.cdc.gov/EID/article/28/8/22-0310-F3.htm>)**

The 24 Cameron Peak firefighter sequences served as focal sequences and a total of 45,686 sequences served as contextual sequences: 44,814 sequences downloaded from the



GISAID repository (7) collected from human specimens testing positive for SARS-CoV-2 within the United States during August 15–December 21, 2020 (representing 1 week before and 1 week after the first and last reported positive cases, respectively) and 872 sequences from human specimens collected from Colorado during the same period and sequenced internally at the Colorado Department of Public Health and Environment state laboratory. Next, all contextual sequences were ranked based on their sequence similarity to the 24 Cameron Peak sequences using Nextstrain’s *ncov priorities* python script (8). Contextual sequences were then filtered using Augur’s (Version 10.2.0) *filter* subcommand with parameters specified to group sequences by collection date and subsample a maximum of 1,000 sequences. The final sequence dataset for phylogenetic inference consisted of 754 contextual sequences and the 24 Cameron Peak firefighter focal sequences. Phylogenetic inference was performed using IQTree Version 2.0.3 (<http://www.iqtree.org>). To access branch support, 5,000 ultrafast bootstrap replicates were used. Phylogenetic tree visualization was performed using the python module ete3 Version 3.1.2 (<https://pypi.org/project/ete3>).

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8. Nextstrain. Ncov: GitHub [cited 2021 Aug 19]. <https://github.com/nextstrain/ncov>

**Appendix Table 1.** Accession IDs of 24 high-quality Cameron Peak Fire sequences, Colorado, USA\*

Sequence Name	NCBI BioProject Accession	NCBI BioSample Accession	NCBI SRA Accession	NCBI GenBank Accession	GISAIID Accession
Crew A sequences					
hCoV-19/USA/CO-CDPHE-2009021317/2020	PRJNA686984	SAMN17130086	SRR13374028	MW645995	EPI_ISL_677638
hCoV-19/USA/CO-CDPHE-2009021339/2020	PRJNA686984	SAMN17130087	SRR13374027	MW645996	EPI_ISL_677639
Crew B sequences					
hCoV-19/USA/CO-CDPHE-2009042396/2020	PRJNA686984	SAMN17130092	SRR13374021	MW645824	EPI_ISL_710222
hCoV-19/USA/CO-CDPHE-2009042404/2020	PRJNA686984	SAMN17130093	SRR13374020	MW645920	EPI_ISL_710331
hCoV-19/USA/CO-CDPHE-2009042416/2020	PRJNA686984	SAMN17130094	SRR13374019	MW645921	EPI_ISL_710283
hCoV-19/USA/CO-CDPHE-2009042507/2020	PRJNA686984	SAMN17130095	SRR13374018	MW645998	EPI_ISL_677641
hCoV-19/USA/CO-CDPHE-2009042555/2020	PRJNA686984	SAMN17130096	SRR13374017	MW645999	EPI_ISL_677642
hCoV-19/USA/CO-CDPHE-2009042743/2020	PRJNA686984	SAMN17130097	SRR13374016	MW645723	EPI_ISL_677313
hCoV-19/USA/CO-CDPHE-2009090328/2020	PRJNA686984	SAMN17130123	SRR13373986	MW646003	EPI_ISL_677646
hCoV-19/USA/CO-CDPHE-2009090330/2020	PRJNA686984	SAMN17130124	SRR13373985	MW646004	EPI_ISL_677647
Crew C sequences					
hCoV-19/USA/CO-CDPHE-2009042326/2020	PRJNA686984	SAMN17130091	SRR13374022	MW645722	EPI_ISL_677312
hCoV-19/USA/CO-CDPHE-2009090308/2020	PRJNA686984	SAMN17130117	SRR13373993	MW645733	EPI_ISL_677270
hCoV-19/USA/CO-CDPHE-2009090312/2020	PRJNA686984	SAMN17130118	SRR13373992	MW645734	EPI_ISL_677281
hCoV-19/USA/CO-CDPHE-2009090314/2020	PRJNA686984	SAMN17130119	SRR13373991	MW646000	EPI_ISL_677643
hCoV-19/USA/CO-CDPHE-2009090318/2020	PRJNA686984	SAMN17130120	SRR13373990	MW645735	EPI_ISL_677314
hCoV-19/USA/CO-CDPHE-2009090320/2020	PRJNA686984	SAMN17130121	SRR13373988	MW646001	EPI_ISL_677644
hCoV-19/USA/CO-CDPHE-2009090338/2020	PRJNA686984	SAMN17130125	SRR13373984	MW645736	EPI_ISL_677283
Crew I sequences					
hCoV-19/USA/CO-CDPHE-2100078312/2020	PRJNA686984	SAMN23283207	SRR17018872	OL678784	EPI_ISL_6581850
Crew J sequences					
hCoV-19/USA/CO-CDPHE-2009031031/2020	PRJNA686984	SAMN17130088	SRR13374026	MW645997	EPI_ISL_677640
Crew K sequences					
hCoV-19/USA/CO-CDPHE-2009044496/2020	PRJNA686984	SAMN17130098	SRR13374015	MW645724	EPI_ISL_677271
Crew L sequences					
hCoV-19/USA/CO-CDPHE-2009121107/2020	PRJNA686984	SAMN17130126	SRR13373983	MW645737	EPI_ISL_677262
hCoV-19/USA/CO-CDPHE-2009121110/2020	PRJNA686984	SAMN17130127	SRR13373982	MW645738	EPI_ISL_677279
Crew M sequences					
hCoV-19/USA/CO-CDPHE-2100042143/2020	PRJNA686984	SAMN19411787	SRR17435600	OM249427	EPI_ISL_2309210
Crew N sequences					
hCoV-19/USA/CO-CDPHE-2010022040/2020	PRJNA686984	SAMN19411784	SRR17435859	OM249281	EPI_ISL_2309207

\*Sequences listed by crew as referred to in Figure 2 (<https://wwwnc.cdc.gov/EID/article/28/8/22-0310-F2.htm>).



**Appendix Table 2.** Accession IDs of additional Colorado sequences sequenced at CDPHE State Public Health Laboratory\*

Sequence Name	NCBI BioProject Accession	NCBI BioSample Accession	NCBI SRA Accession	NCBI GenBank Accession	GI SAID Accession
<b>County A sequences</b>					
hCoV-19/USA/CO-CDPHE-2009170340/2020	PRJNA686984	SAMN17130186	SRR13373917	MW645951	EPI_ISL_710345
hCoV-19/USA/CO-CDPHE-2009170292/2020	PRJNA686984	SAMN17130181	SRR13373922	MW645946	EPI_ISL_710342
hCoV-19/USA/CO-CDPHE-2009163553/2020	PRJNA686984	SAMN17130143	SRR13373964	MW646006	EPI_ISL_677649
hCoV-19/USA/CO-CDPHE-2009164054/2020	PRJNA686984	SAMN17130161	SRR13373944	MW645928	EPI_ISL_710333
hCoV-19/USA/CO-CDPHE-2008161331/2020	PRJNA686984	SAMN17130051	SRR13374066	MW645817	EPI_ISL_710217
hCoV-19/USA/CO-CDPHE-2009170150/2020	PRJNA686984	SAMN17130176	SRR13373928	MW645943	EPI_ISL_710341
hCoV-19/USA/CO-CDPHE-2009170084/2020	PRJNA686984	SAMN17130172	SRR13373932	MW645939	EPI_ISL_710339
hCoV-19/USA/CO-CDPHE-2100008733/2020	PRJNA686984	SAMN17250951	SRR13404609	MW645561	EPI_ISL_771183
<b>County B Sequences</b>					
hCoV-19/USA/CO-CDPHE-2008170075/2020	PRJNA686984	SAMN17130056	SRR13374061	MW645710	EPI_ISL_677304
hCoV-19/USA/CO-CDPHE-2100060009/2020	PRJNA686984	SAMN17903010	SRR13703951	MW629399	EPI_ISL_983861
hCoV-19/USA/CO-CDPHE-2011221706/2020	PRJNA686984	SAMN18173874	SRR13867969	MW715242	EPI_ISL_1169713
hCoV-19/USA/CO-CDPHE-2100060046/2020	PRJNA686984	SAMN17903011	SRR13703950	MW629400	EPI_ISL_983862
hCoV-19/USA/CO-CDPHE-2011134677/2020	PRJNA686984	SAMN19413147	SRR17436226	OM171064	EPI_ISL_2310376
hCoV-19/USA/CO-CDPHE-2011134720/2020	PRJNA686984	SAMN19413148	SRR17436234	OM170758	EPI_ISL_2310377
hCoV-19/USA/CO-CDPHE-2100117646/2020	PRJNA686984	SAMN19413606	SRR17489341	OM211796	EPI_ISL_2310835
hCoV-19/USA/CO-CDPHE-2100106281/2020	PRJNA686984	SAMN19413558	SRR17489350	OM211622	EPI_ISL_2310787
hCoV-19/USA/CO-CDPHE-2100060065/2020	PRJNA686984	SAMN19413181	SRR17436998	OM170407	EPI_ISL_2310410
hCoV-19/USA/CO-CDPHE-2009262446/2020	PRJNA686984	SAMN19407744	SRR17350928	OM067237	EPI_ISL_1540131
hCoV-19/USA/CO-CDPHE-2100145835/2020	PRJNA686984	SAMN19413673	SRR17489016	OM211591	EPI_ISL_2310902
hCoV-19/USA/CO-CDPHE-2010022230/2020	PRJNA686984	SAMN19407745	SRR17350927	OM067433	EPI_ISL_1540132
hCoV-19/USA/CO-CDPHE-2100085221/2020	PRJNA686984	SAMN19413110	SRR17436315	OM170772	EPI_ISL_2310339
<b>County C Sequences</b>					
hCoV-19/USA/CO-CDPHE-2100013339/2020	PRJNA686984	SAMN17250967	SRR13404560	MW645576	EPI_ISL_771299
hCoV-19/USA/CO-CDPHE-2011044565/2020	PRJNA686984	SAMN19411925	SRR17435786	OM248836	EPI_ISL_2309348
<b>County D Sequences</b>					
hCoV-19/USA/CO-CDPHE-2100015283/2020	PRJNA686984	SAMN17250997	SRR13404525	MW645604	EPI_ISL_771179
<b>All Other Colorado Sequences</b>					
hCoV-19/USA/CO-CDPHE-2100116162/2020	PRJNA686984	SAMN20509402	SRR15488467	MZ830216	EPI_ISL_3160167
hCoV-19/USA/CO-CDPHE-2100036174/2020	PRJNA686984	SAMN17251037	SRR13404481	MW645994	EPI_ISL_771175
hCoV-19/USA/CO-CDPHE-2100001047/2020	PRJNA686984	SAMN17250939	SRR13404622	MW645549	EPI_ISL_771283
hCoV-19/USA/CO-CDPHE-2100001027/2020	PRJNA686984	SAMN17250938	SRR13404624	MW645548	EPI_ISL_771282
hCoV-19/USA/CO-CDPHE-2100014442/2020	PRJNA686984	SAMN17250979	SRR13404544	MW645587	EPI_ISL_771305
hCoV-19/USA/CO-CDPHE-2008170097/2020	PRJNA686984	SAMN17130057	SRR13374060	MW645711	EPI_ISL_677305
hCoV-19/USA/CO-CDPHE-2010191275/2020	PRJNA686984	SAMN17130202	SRR13373898	MW645964	EPI_ISL_710354
hCoV-19/USA/CO-CDPHE-2100038259/2020	PRJNA686984	SAMN17251055	SRR13404461	MW645657	EPI_ISL_771350
hCoV-19/USA/CO-CDPHE-2010294547/2020	PRJNA686984	SAMN20794120	SRR15460147	MZ832789	EPI_ISL_3403786
hCoV-19/USA/CO-CDPHE-2100036488/2020	PRJNA686984	SAMN17251043	SRR13404474	MW645645	EPI_ISL_771342
hCoV-19/USA/CO-CDPHE-2011200501/2020	PRJNA686984	SAMN17250909	SRR13404591	MW645519	EPI_ISL_771258
hCoV-19/USA/CO-CDPHE-2011193762/2020	PRJNA686984	SAMN17250899	SRR13404435	MW645509	EPI_ISL_771161
hCoV-19/USA/CO-CDPHE-2100280404/2020	PRJNA686984	SAMN23283277	SRR17019083	OL679077	EPI_ISL_6581924

\*Sequences are listed by county as referred to in Figure 3 (<https://wwwnc.cdc.gov/EID/article/28/8/22-0310-F3.htm>). CDPHE, Colorado Department of Public Health and Environment.