

Methods and Applications

Co-Localization of Sampling and Sequencing for Zoonotic Pathogen Identification in the Field Monitoring Using Mobile Laboratories

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ABSTRACT

Introduction: Accurate etiological detection is needed to evaluate the risk of zoonotic diseases. Metagenomic next-generation sequencing (mNGS) can be used to monitor pathogens in animal species and identify potential zoonotic threats. The current sampling model for zoonotic pathogen monitoring in wild animals requires samples to be transferred from the field to a laboratory for further detection.

Methods: We constructed a zoonotic pathogen survey model using a set of mobile laboratories.

Results: The monitoring in this study was preplanned to detect *Yersinia pestis*, but the mNGS unexpectedly identified *Bartonella* spp. in the rodent samples, thus exposing the threat of bartonellosis to humans in this region. The co-localization of sampling and sequencing (CLOSS) model we tested required no long-distance transferring of samples and expands the regional coverage of zoonotic surveys by using a mobile laboratory.

Discussion: Using this mNGS technique will enable detection of more zoonotic pathogens beyond the preplanned monitoring targets. This may increase the surveillance efficiency compared with that of the previous workflow and expand the application of the mobile laboratories for infectious diseases identification and surveillance in the field.

surveillance system with strong laboratory investigation abilities is urgently needed to eliminate zoonoses and minimize the risk of entry, establishment, and spread of emerging zoonoses (2).

Metagenomic next-generation sequencing (mNGS) can detect various pathogenic microorganisms without bias. mNGS is increasingly used to diagnose infectious diseases, particularly when conventional diagnostic approaches are limited. One example involved the use of mNGS for a chronic neuroleptospirosis case: traditional methods failed to identify the pathogen; however, mNGS identified the sequence as *Leptospira santarosai* (3). mNGS has been applied for clinical diagnosis and pathogen discovery in wildlife, especially for rare, novel, and difficult-to-detect pathogens. Rapid pathogen screening facilitates identification to initiate timely interventions.

The importance of zoonoses and their public health effects are well recognized (2). Knowledge of zoonotic pathogens in wildlife hosts or vectors in frequent contact with humans and domestic animals is important for understanding pathogenic diversity in nature (4). Metagenomic sequencing has unprecedented value in these investigations because it can describe microbial communities in a relatively unbiased manner, as opposed to screening for specific taxa (5). In wildlife investigations for zoonotic diseases, the targeted pathogens are typically predetermined, and samples are collected in the field which are transferred to the laboratory. Some issues with these surveys include that they only test for predetermined pathogens, the survey samples must be preserved and transferred under biosafety protocol, and the procedure is time-consuming. Long field deployments in geographically isolated places where entire laboratory teams must be sent each sentinel site are not only costly, but also increase stress on laboratory employees.

During the coronavirus disease 2019 (COVID-19) pandemic, the mobile polymerase chain reactions

INTRODUCTION

Zoonotic diseases are increasing globally and pose a great threat to both livestock and human health. Zoonoses are estimated to cause over one billion cases of infection and millions of deaths annually worldwide. Over 30 new human pathogens have been detected in the last 3 decades with over 75% of these originating from animals (1). Establishment of a robust, integrated

(PCR) laboratories provided a rapid, accurate, and clinically effective option for hospital diagnostic capabilities. In an East African Community (including Kenya, Rwanda, Burundi, Uganda and South Sudan), the mobile laboratory network for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) diagnostics ensured trade was able to continue during border closures (6). Here, during a mobile laboratory survey for rodents carrying *Yersinia pestis* in the field, we performed mNGS onsite to detect possible pathogens in the rodent samples. We had planned to use the rodent samples for *Y. pestis*-associated plague monitoring, but *Bartonella* sequences were unexpectedly identified. With the sampling and sequencing performed onsite, mNGS provided a rapid and broad-spectrum survey model for detecting zoonotic pathogens in rodents.

METHODS

Sampling and Tissue Collection at Surveillance Sites

In September 2021, we conducted a field investigation for rodents carrying *Y. pestis* in the Inner Mongolia Autonomous Region. The rodents were sampled using rat clips. The vehicle containing the mobile laboratory arrived at the surveillance sentinel so laboratory testing could be carried out, such as

performing mNGS using a sequencer in the mobile laboratory (Figure 1). Heart, liver, lung, and serum samples were taken from the rodents.

Bacterial Cultures and Identification

The diagnostic methods included identifying *Y. pestis* from isolates with a culture by a special phage lysis assay. Immunological methods for detecting the F1 antigen, i.e., the reverse indirect hemagglutination assay (RIHA) and/or colloidal gold-immunochromatography assay, were performed on liver and lung samples. Serum samples collected from the rodents were tested for antibodies against the F1 antigen via an indirect hemagglutination assay (IHA).

Genomic DNA Extraction and Real-Time PCR

Total genomic DNA was extracted from the rodent hearts, livers and lungs using the QIAamp DNA Mini Kit (Qiagen, Hilden, Germany). All genomic DNA samples were tested for *caf1* and *Chr392* via real-time PCR to detect *Y. pestis*. The *ssrA* gene was tested using real-time PCR to detect *Bartonella* spp. (7).

Metagenomic Sequencing

DNA was quantified using Qubit v3. The sequencing libraries were constructed with the MGIEasy Fast PCR-FREE FS Library Prep Set (MGI,

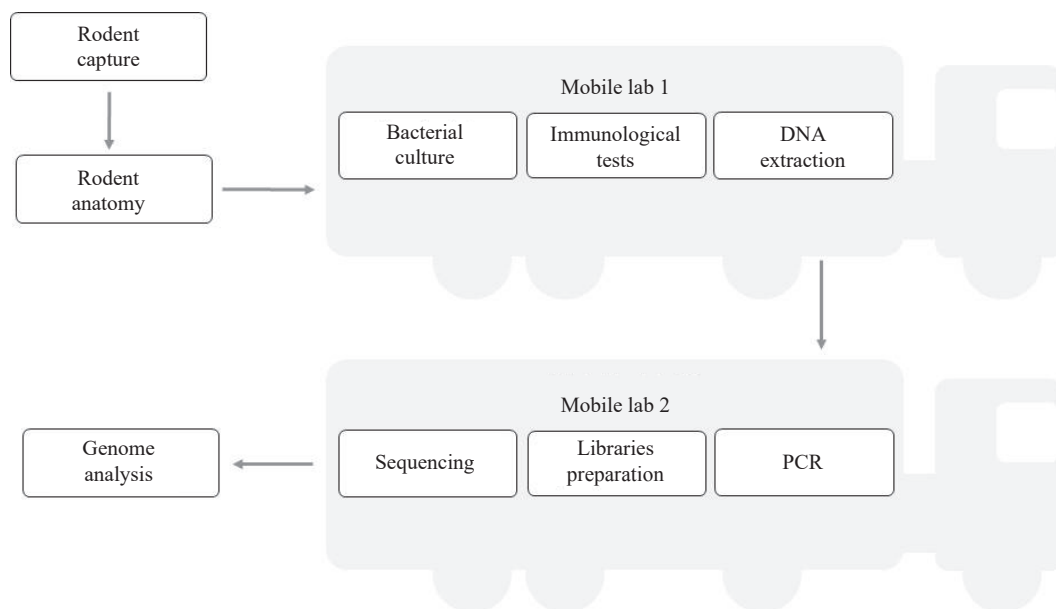


FIGURE 1. Laboratory procedure from rodent capture to pathogen genome identification in the field monitoring based on the mobile laboratories.

Abbreviations: DNA=deoxyribonucleic acid; PCR=polymerase chain reactions.

Wuhan, China), DNBSEQ OneStep DNB Make Reagent Kit V2.0 (OS-DB; MGI), Nextera XT DNA Sample Prep Kit (Illumina, California, USA) and Nextera XT Index Kit (Illumina). These libraries were sequenced on the portable DNBSEQ sequencing platforms (MGI) using a portable DNBSEQ Sequencing Set (SE100; MGI) and on the iSeq100 sequencing platforms (Illumina) using iSeq 100i1 Reagent (PE150; Illumina).

Bioinformatics

The mNGS data were processed using Kraken (8), the MGI Pathogen Fast Identification pipeline, and the MicroFuture multiple pathogen analysis for metagenomic identification of pathogens. Bowtie2 and samtools were used to extract the pathogen-related reads.

To determine the *Bartonella* species from the pathogen-related reads, the extracted reads were blasted against the NCBI refseq database, the NR database, and the *Bartonella* genus genome sequences. A *Bartonella* genus genome reference database was constructed by retrieving all the *Bartonella* genera with genome sequences in the NCBI Assemble database. The sequences were evaluated using the average nucleotide identity (ANI) to filter obviously abnormal genomes. Finally, 144 whole-genome sequences belonging to 39 species were detected and included in the reference database. The extracted reads were then blasted against the reference database. The BLASTN cut-offs for the E-value was 0.00001 and the identity was 80%.

RESULTS

The mobile laboratories were comprised of one BSL-3 laboratory for sample preparation and pathogen culture along with one BSL-1 laboratory for PCR and sequencing. When CLOSS was performed in the field, the power supply vehicle and camping vehicle will be included. Three to four technicians working in the laboratories and two technicians for operation and maintenance of the mobile laboratories and power supply vehicle were needed.

The mobile laboratories arrived at the field site where rodent samples were collected. The electrical supply allowed us to use a refrigerator to store reagents. In the BSL-3 mobile laboratory, tissue was taken and DNA was extracted. In the BSL-1 mobile laboratory, the library was prepared and sequencing was initiated.

The field-generated sequence data were analyzed using a variety of commercial and custom-developed bioinformatic workflows. The whole turnaround time of our protocol was less than 24 hours.

Real-Time PCR and Serological Detection of *Y. pestis*

Total DNA was extracted from two rodent samples and tested for *Y. pestis* using real-time PCR, and all samples were negative. Additionally, all rodent samples used to detect the F1 antigen via colloidal gold immunochromatography assay and RIHA and the F1 antibody using IHA were negative.

Metagenomic Sequencing

Metagenomic sequencing was performed on samples 11 and 12, which were identified as *Meriones unguiculatus* (gerbil) and *Spermophilus dauricus* (citellus). Sample 11 was the liver of a *Meriones unguiculatus*, which yielded about 1.4 Gb of sequencing data (5,017,499 reads, PE150) after 18 hours of sequencing. After filtering the quality reads and host reads, we obtained a dataset of 471,040 clean reads. As per the refseq database, 4,351 reads (including 253 unique reads) corresponded to *Bartonella grahamii* (relative abundance: 49.26%) and 2,722 reads (including 70 unique reads) corresponded to *B. tribocorum* (relative abundance: 30.82%). The reads were then mapped to the reference genomes of *B. grahamii* (NC_012846.1) and *B. tribocorum* (NC_010161.1), yielding approximately 22.4% and 7.8% coverages, respectively (Figure 2).

Virulence-associated genes such as *omp89* and *omp43* (coverage >70%) were identified as major putative Fn-binding proteins in the *B. henselae* outer membrane proteins (9). Compared to the NR database, all 323 unique reads extracted from the refseq database corresponded to *Bartonella* spp. The *Bartonella* genus database constructed comprised 144 whole-genome sequences belonging to 39 species. ANI analysis showed intraspecies nucleotide similarity rates of >97% (different strains within the same species; Supplementary Figure S1, available in <http://weekly.chinacdc.cn>). Finally, from the 323 unique reads, 22 were species-specific, including 21 corresponding to *B. grahamii* and one corresponding to *B. tribocorum*.

For sample 12, extracted from the liver of the *Spermophilus dauricus*, we obtained ~0.7 Gb of sequencing data (6,597,208 reads, SE100) after 9 hours of sequencing. After filtering the quality reads

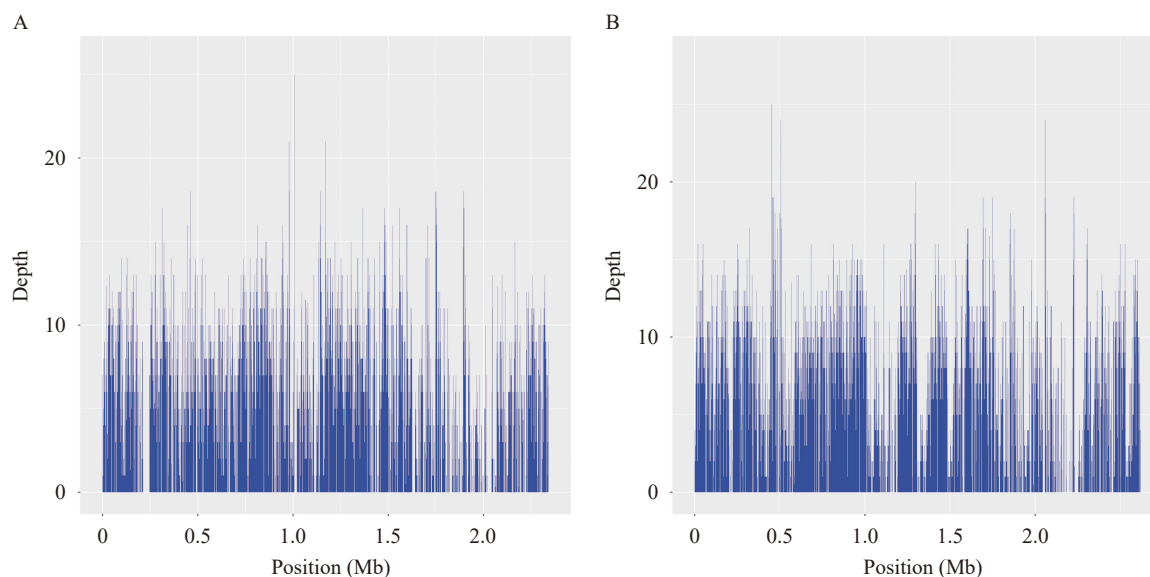


FIGURE 2. Mapping of genome assemblies for *B. grahamii* (A) and *B. tribocorum* (B) generated from metagenomic next-generation sequencing (mNGS) reads to the respective reference genomes.

Note: the reference genomes: *B. grahamii* NC_012846.1 and *B. tribocorum* NC_010161.1.

and the host reads, metagenomic sequencing yielded a dataset of 6,526,977 clean reads, of which only 29 corresponded to *Bartonella* spp. (relative abundance: 1.1%). We also obtained 2,673 reads related to *Aspergillus parasiticus* (relative abundance near 99%) in the sequencing data from this sample.

PCR Detection for *Bartonella*

We further detected the *Bartonella* from samples 11 and 12 using real-time PCR. Both samples were positive for *ssrA*, and the CT values were 27.24 for sample 11 and 32.88 for sample 12, indicating the correspondence of the relative abundances of the sequenced *Bartonella* reads from both samples.

DISCUSSION

Animal-based surveillance can identify zoonotic pathogens and provide public health authorities with sufficient warning to implement appropriate prevention measures before human cases occur (10). mNGS is an unbiased, timely, and accurate approach to pathogen detection (11–12). Cultivating pathogens can take several days; mNGS can reduce the time needed to obtain results to less than 24 h (13). In this study, we identified *Bartonella* spp. from rodent samples. *Bartonella* spp. are highly adaptable and can infect various hosts and vectors, with at least 13 species known to be pathogenic to humans (14). Many

Bartonella-associated illnesses are linked with arthropod or animal transmission, and most patients reported having animal contact prior to disease onset. In Inner Mongolia, 4 *Bartonella* species were detected from 188 small mammals sampled from 2012–2013, and the positive infection rates ranged from 38.37%–56.41% (15). Here, we identified *Bartonella* infections from rodent tissue samples. We detected only two samples via mNGS, and both were positive. In this study, species-specific reads corresponding to two *Bartonella* species were found in *Meriones unguiculatus*, suggesting possible complex *Bartonella* spp. infections in this rodent. Whether all *Bartonella* spp. were etiologic agents of human illness was unclear owing to the lack of accurate diagnostic methods in clinics, which were important for correct diagnoses and estimating disease burdens (16). *Bartonella* prevalence and transmission remain unknown in China. Previous data and our findings demonstrate the necessity of surveillance of wild animals and people who are exposed to wild animals and their surroundings that have tested positive for *Bartonella*.

In the present zoonosis surveillance system, transferring animal carcasses from sampling locations in the field to the laboratory is time-consuming and requires biosafety protection. Zoonotic pathogen surveillance, such as plague monitoring in grasslands and plateaus, covers vast areas; thus, transferring samples may lower working efficiency. Mobile

laboratory-based surveys save time and expand regional monitoring more effectively than previous strategies, which may include field sampling. Since mobile laboratory technologies are frequently designed to be durable, affordable, and mostly independent of energy, Internet access, or cold chain availability, they provide an excellent opportunity for local capacity building.

Additionally, mNGS can expand the monitored pathogen spectrum and is more advantageous for discovering new pathogens, especially for accurate laboratory diagnoses in regions without access to sophisticated laboratory techniques. We have named this zoonotic pathogen survey model “co-localization of sampling and sequencing (CLOSS),” which means performing animal-borne pathogen surveys in the field via metagenomic sequencing techniques in mobile laboratories. For monitoring zoonotic pathogen, CLOSS should be integrated into the Chinese pathogen identification net (China PIN). Technical laboratory staff should be trained in sequencing by China PIN. The CLOSS procedure will be further improved and practiced in order to evaluate and optimize this monitoring model. Due to COVID-19 outbreaks, many CDCs in China are now equipped with mobile laboratories. Our CLOSS model can expand the applications of these mobile laboratories to surveillance of other infectious diseases following the COVID-19 pandemic.

Conflicts of interest: No conflicts of interest.

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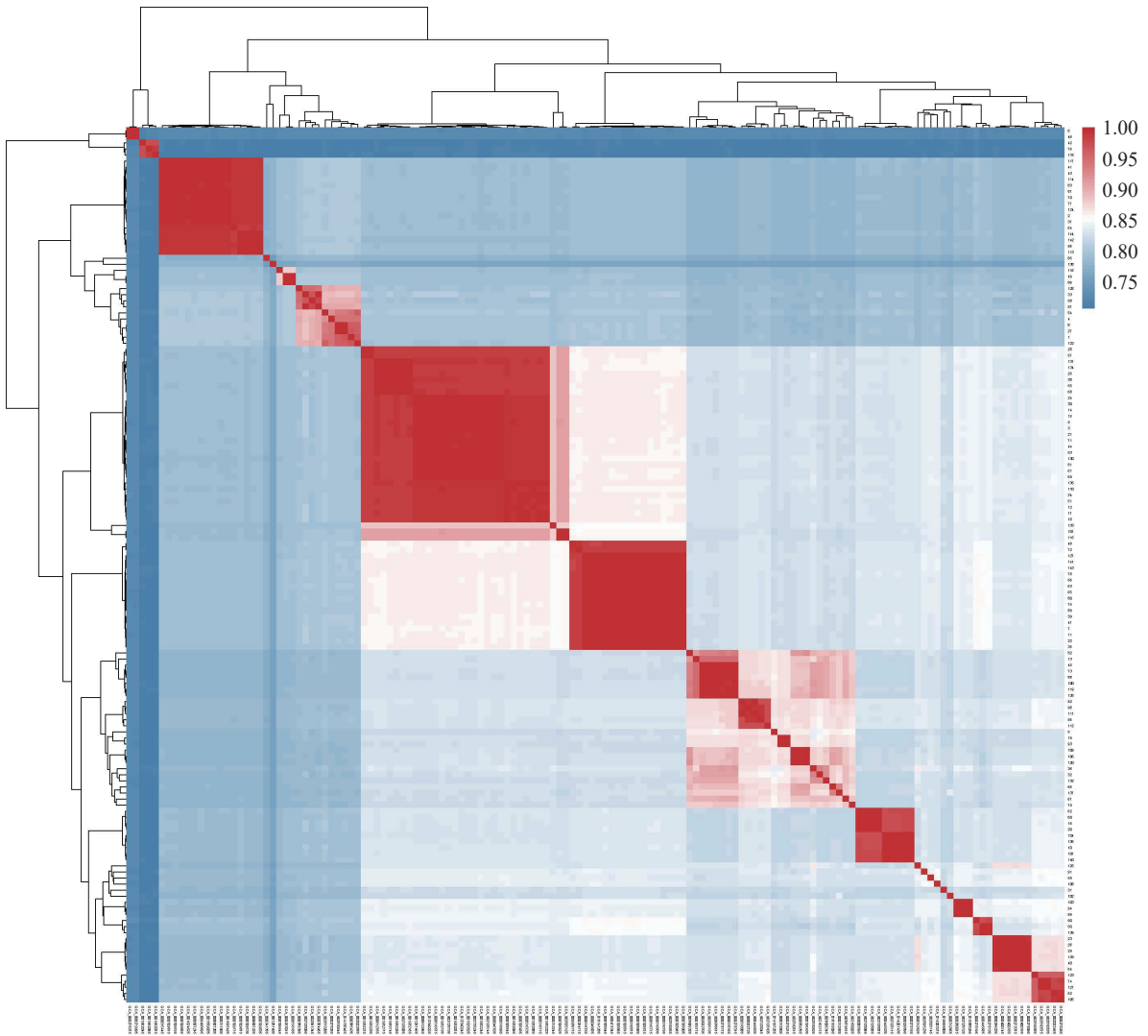
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SUPPLEMENTARY FIGURE S1. ANI analysis of the 144 whole-genome *Bartonella* spp. sequences from GenBank. Notes: The values generated by the Gegenees software Version 2 shown in the heat map indicate the percentage similarity between the analyzed genomes. The colors vary from blue (low similarity) to red (high similarity).