

Prospecting for Zoonotic Pathogens by Using Targeted DNA Enrichment

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More than 60 zoonoses are linked to small mammals, including some of the most devastating pathogens in human history. Millions of museum-archived tissues are available to understand natural history of those pathogens. Our goal was to maximize the value of museum collections for pathogen-based research by using targeted sequence capture. We generated a probe panel that includes 39,916 80-bp RNA probes targeting 32 pathogen groups, including bacteria, helminths, fungi, and protozoans. Laboratory-generated, mock-control samples showed that we are capable of enriching targeted loci from pathogen DNA 2,882–6,746-fold. We identified bacterial species in museum-archived samples, including *Bartonella*, a known human zoonosis. These results showed that probe-based enrichment of pathogens is a highly customizable and efficient method for identifying pathogens from museum-archived tissues.

Many serious human pathogens result from zoonotic transmission, including 61% of known human pathogens and 75% of emerging human pathogens (1). For example, rabies virus is transmitted by saliva of infected animals (2). The plague bacteria (*Yersinia pestis*), the causative agent of the largest documented pandemic in human history that reduced the population of Europe by 30%–50%, was transmitted from rats to humans by fleas (3). Other zoonoses include Ebola virus (4), tularemia (*Francisella tularensis*) (5), and tuberculosis (6). The SARS-CoV-2 pandemic, thought to have a bat reservoir, has stimulated renewed emphasis on zoonotic pathogen surveillance (7,8).

Natural history museums are repositories of biologic information in the form of voucher specimens

that represent a major, underused resource for studying zoonotic pathogens (9–13). Originally, specimens were archived as dried skin and skeletal vouchers or preserved in fluids (ethanol) after fixation with formalin or formaldehyde. Now, best practices include preserving specimens and associated soft tissues in liquid nitrogen (–190°C) or mechanical freezers (–80°C) from the time they are collected (14). Those advances in preservation make it possible to extract high-quality DNA and RNA that can be used for pathogen surveillance. For example, retroactive sampling of archived tissues from the US Southwest found that Sin Nombre virus, a New World hantavirus, was circulating in wild rodent populations almost 20 years before the first human cases were reported (15).

It is critical to develop a range of tools for extracting pathogen information from museum-archived samples. Targeted sequencing using probe enrichment has become the tool of choice for medical genomics (16), population genetics (17), phylogenetics (18), and ancient DNA (19,20). Those methods are designed to enrich small amounts of DNA target from a background of contaminating DNA. Probe-based, targeted sequencing has been used to enrich pathogens from complex host–pathogen DNA mixtures (21). For example, Keller et al. used probes to capture and sequence complete *Y. pestis* genomes from burial sites >1,500 years old (22). Enrichment is frequently achieved by designing a panel of probes to specifically target a handful of pathogens of interest (23,24). Similarly, commercial probe sets are available for many types of viruses and human pathogens (23–25). However, many of these probe sets are limited to specific pathogens that might not infect other host species.

Our goal was to develop a panel of biotinylated baits, or probes, to identify the eukaryotic and bacterial pathogens responsible for 32 major zoonoses (Table 1). We aimed to capture both known and related pathogens, using the fact that probes can capture sequences that are ≤10% divergent. To perform this

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capture, we used a modified version of the ultraconserved element (UCE) targeted sequencing technique (26,27) to specifically enrich pathogen DNA. Biotinylated baits are designed to target conserved genomic regions among diverse groups of pathogens (Figure 1). The baits are hybridized to a library potentially containing pathogen DNA. Bait-bound DNA fragments are enriched during a magnetic bead purification step before sequencing (Figure 2). The final library contains hundreds or thousands of orthologous loci with single-nucleotide variants or indels from the targeted pathogen groups that can then be used for population or phylogenetic analyses.

Methods

We have compiled a detailed description of the methods used (Appendix 1, <https://wwwnc.cdc.gov/EID/article/29/8/22-1818-App1.pdf>; <https://doi.org/10.17504/protocols.io.5jyl8jnzrg2w/v1>). Code is available on GitHub (https://www.github.com/nealplatt/pathogen_probes; <https://doi.org/10.5281/zenodo.7319915>). Raw sequence data are available from the National Center for Biotechnology Information (BioProject PRJNA901509; Appendix 2, <https://wwwnc.cdc.gov/EID/article/29/8/22-1818-App2.xlsx>). A summary of our methods follows.

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Panel Development

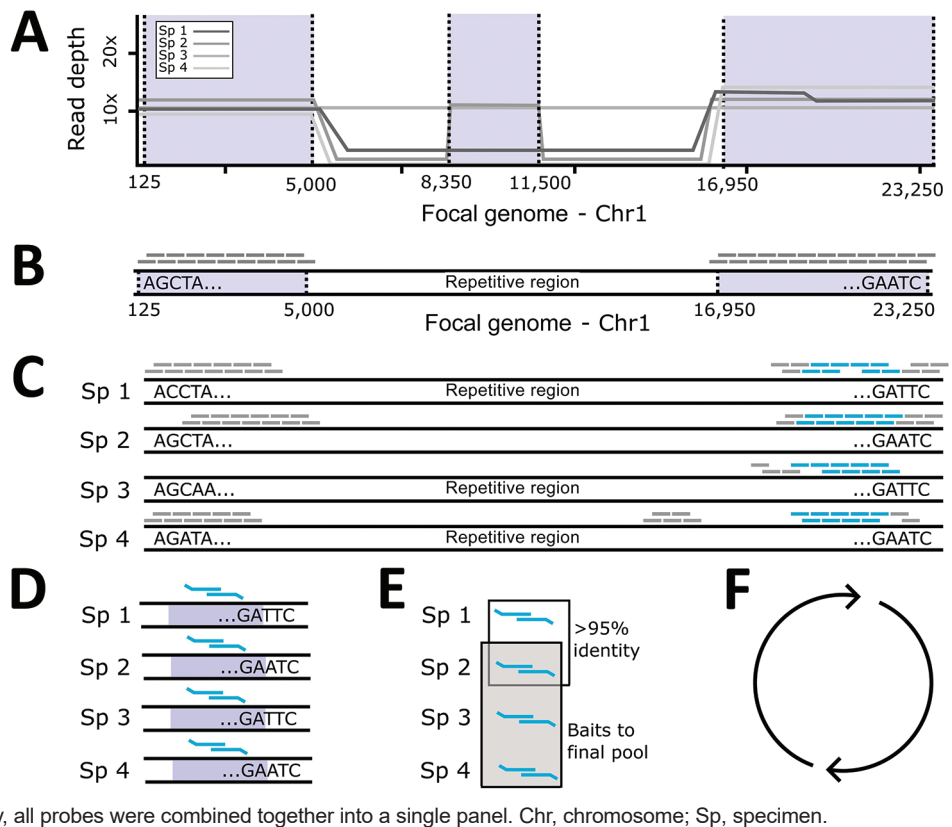
We developed a panel of baits for targeted sequencing of 32 zoonotic pathogens. To develop this panel, we used the Phyluce version 1.7.1 (26,27) protocol to design baits for conserved loci within each pathogen group. First, we simulated and mapped reads from each species within a pathogen group to a focal genome assembly (Table 1; Figure 1, panel A). We used the mapped reads to identify putative orthologous loci that were >80% similar across the group and generated in silico baits from the focal genome (Figure 1, panel B). These baits were mapped back to each member (Figure 1, panel C) to identify single-copy orthologs within the group. Next, we designed 2 overlapping 80-bp baits from loci in each member of the group (Figure 1, panel D) and removed baits with >95% sequence similarity (Figure 1, panel E). We repeated those steps for each pathogen group (Figure 1, panel F). We compared the remaining baits with mammalian genomes and replaced them to minimize

Table 1. Zoonotic pathogens targeted for DNA enrichment in study of prospecting for zoonotic pathogens by using targeted DNA enrichment

| Pathogen group | Taxonomic level | Focal pathogen | Zoonoses |
|-------------------------------|-----------------|-------------------------------------|--------------------|
| <i>Anaplasma</i> | Genus | <i>Anaplasma phagocytophilum</i> | Anaplasmosis |
| Apicomplexa | Phylum | <i>Plasmodium falciparum</i> | Malaria |
| <i>Bacillus cereus</i> group* | Species group | <i>Bacillus anthracis</i> | Anthrax |
| <i>Bartonella</i> | Genus | <i>Bartonella bacilliformis</i> | Cat-scratch fever |
| <i>Borrelia</i> | Genus | <i>Borrelia burgdorferi</i> | Lyme disease |
| <i>Burkholderia</i> | Genus | <i>Burkholderia mallei</i> | Glanders |
| <i>Campylobacter</i> | Genus | <i>Campylobacter jejuni</i> | Campylobacteriosis |
| Cestoda | Class | <i>Taenia multiceps</i> | Taeniasis |
| <i>Chlamydia</i> | Genus | <i>Chlamydia trachomatis</i> | Chlamydia |
| <i>Coxiella</i> | Genus | <i>Coxiella burnetii</i> | Q fever |
| <i>Ehrlichia</i> | Genus | <i>Ehrlichia canis</i> | Ehrlichiosis |
| Eurotiales | Order | <i>Talaromyces marneffeii</i> | Talaromycosis |
| <i>Francisella</i> | Genus | <i>Francisella tularensis</i> | Tularemia |
| Hexamitidae | Family | <i>Giardia intestinalis</i> | Giardiasis |
| Kinetoplastea | Class | <i>Leishmania major</i> | Leishmaniasis |
| <i>Leptospira</i> | Genus | <i>Leptospira interrogans</i> | Leptospirosis |
| <i>Listeria</i> | Genus | <i>Listeria monocytogenes</i> | Listeriaosis |
| <i>Mycobacterium</i> | Genus | <i>Mycobacterium tuberculosis</i> | Tuberculosis |
| Nematodes (clade I) | Phylum (clade) | <i>Trichinella spiralis</i> | Trichinosis |
| Nematodes (clade III) | Phylum (clade) | <i>Brugia malayi</i> | Filariasis |
| Nematodes (clade IVa) | Phylum (clade) | <i>Strongyloides stercoralis</i> | Strongyloidiasis |
| Nematodes (clade IVb) | Phylum (clade) | <i>Steinernema carpocapsae</i> | None |
| Nematodes (clade V) | Phylum (clade) | <i>Haemonchus contortus</i> | None |
| Onygenales | Order | <i>Histoplasma capsulatum</i> | Histoplasmosis |
| <i>Pasteurella</i> | Genus | <i>Pasteurella multocida</i> | Pasteurellosis |
| <i>Rickettsia</i> | Genus | <i>Rickettsia rickettsii</i> | Typhus |
| <i>Salmonella</i> | Genus | <i>Salmonella enterica</i> | Salmonellosis |
| <i>Streptobacillus</i> | Genus | <i>Streptobacillus moniliformis</i> | Rat-bite fever |
| Trematoda | Class | <i>Schistosoma mansoni</i> | Schistosomiasis |
| Tremellales | Order | <i>Cryptococcus neoformans</i> | Cryptococcosis |
| <i>Trypanosoma</i> * | Genus | <i>Trypanosoma cruzi</i> | Sleeping sickness |
| <i>Yersinia</i> | Genus | <i>Yersinia pestis</i> | Plague |

*Supplemented with additional probes/baits.

Figure 1. Probe panel design for study of prospecting for zoonotic pathogens by using targeted DNA enrichment. A) Simulated reads from each pathogen within a group were mapped back to a single focal genome. B) We identified regions with consistent coverage from each member of the pathogen group to identify putative, orthologous loci and generated a set of in silico probes from the focal genome. C) Those in silico probes were then mapped back to the genomes of each member in the pathogen group to find single copy, orthologous regions, present in most members. D, E) We designed 2 overlapping 80-bp baits to target the loci in each member of the pathogen group (D) and compared them with each another to remove highly similar probes (E). One probe was retained from each group of probes with high sequence similarity (>95%). F) We identified the probes necessary to capture 49 loci in that pathogen group. This process was repeated for the next pathogen group. Finally, all probes were combined together into a single panel. Chr, chromosome; Sp, specimen.

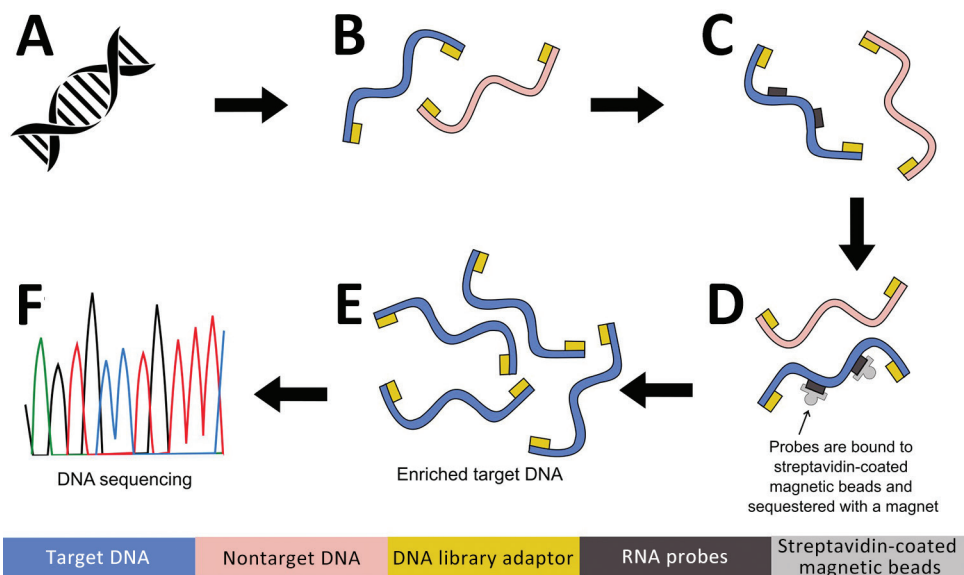


cross-reactivity with the host. Finally, we combined baits to capture 49 loci from each pathogen group into a panel that was synthesized by Daicel Arbor Biosciences (<https://arborbiosci.com>).

Museum-Archived and Control Samples

We extracted DNA from 38 museum samples by using the DNeasy Kit (QIAGEN, <https://www.qiagen.com>) (Table 2). We generated control samples

Figure 2. Targeted DNA enrichment workflow for study of prospecting for zoonotic pathogens by using targeted DNA enrichment. A) Genomic DNA extracted using the DNeasy Kit (QIAGEN, <https://www.qiagen.com>). B) Next-generation sequencing libraries prepared using KAPA Hyperplus Kit (<https://www.biocompare.com>) and barcoding each library with IDT xGen Stubby Adaptor-UDI Primers (<https://www.idtdna.com>). C) RNA probes hybridization using the high sensitivity protocol of myBaits version 5. (<https://arborbiosci.com>). D) Probes bound to streptavidin-coated magnetic beads and sequestered with a magnet (E) 15 cycles PCR amplification of enriched libraries. F) Libraries sequenced on an Illumina Hi-Seq 2500 platform (<https://www.illumina.com>).



by spiking naive mouse DNA with 1% microorganism DNA from *Mycobacterium bovis*, *M. tuberculosis*, *Plasmodium vivax*, *P. falciparum*, and *Schistosoma mansoni*. We then further diluted an aliquot of this 1% pathogen mixture into mouse DNA to create a 0.001% host-pathogen mixture. This range was designed to test the lower limits of detection but also represent a reasonable host-pathogen proportion.

For example, *Theileria parva*, a tick-transmitted apicomplexan, is present in samples from 0.9% through 3% (28), and 1.5% of DNA sequence reads in clinical blood samples is from *P. vivax* (29).

Library Preparation

We generated standard DNA sequencing libraries from 500 ng of DNA per sample. We combined

Table 2. Specimens examined using targeted sequencing in study of prospecting for zoonotic pathogens by using targeted DNA enrichment*

| Museum accession no. | Source species (common name) | Locality, country: state, county | Date | SRA ID |
|----------------------|--|---|-------------|--------------|
| TK48533 | <i>Myotis volans</i> (long-legged myotis) | Mexico: Durango, Arroyo El Triguero | 1995 May 18 | SAMN31718202 |
| TK49668 | <i>Didelphis virginiana</i> (Virginia opossum) | United States: Texas, Kerr | 1996 May 14 | SAMN31718203 |
| TK49674 | <i>Peromyscus attwateri</i> (Texas mouse) | United States: Texas, Kerr | 1996 May 14 | SAMN31718204 |
| TK49686 | <i>Peromyscus laceianus</i> (deer mouse) | United States: Texas, Kerr | 1996 May 14 | SAMN31718205 |
| TK49712 | <i>Dasyurus novemcinctus</i> (nine-banded armadillo) | United States: Texas, Kerr | 1996 May 16 | SAMN31718206 |
| TK49732 | <i>Lasiurus borealis</i> (eastern red bat) | United States: Texas, Kerr | 1996 May 17 | SAMN31718207 |
| TK49733 | <i>Myotis velifer</i> (vesper bat) | United States: Texas, Kerr | 1996 May 16 | SAMN31718208 |
| TK57832 | <i>P. attwateri</i> | United States: Texas, Kerr | 1997 May 14 | SAMN31718209 |
| TK70836 | <i>Desmodus rotundus</i> (common vampire bat) | Mexico: Durango, San Juan de Camarones | 1997 Jun 27 | SAMN31718210 |
| TK90542 | <i>Sigmodon hirsutus</i> (southern cotton rat) | Mexico: Chiapas, Comitán | 1999 Jul 9 | SAMN31718211 |
| TK93223 | <i>Peromyscus melanophrys</i> (plateau mouse) | Mexico: Oaxaca, Las Minas | 2000 Jul 13 | SAMN31718212 |
| TK93289 | <i>Carollia subrufa</i> (gray short-tailed bat) | Mexico: Chiapas, Ocozocoautla | 2000 Jul 16 | SAMN31718213 |
| TK93402 | <i>Chaetodipus eremicus</i> (Chihuahan pocket mouse) | Mexico: Coahuila | 2000 Jul 22 | SAMN31718214 |
| TK101275 | <i>Glossophaga commissarisi</i> (Commissaris' long-tongued bat) | Honduras: Comayagua, Playitas | 2001 Jul 10 | SAMN31718215 |
| TK136205 | <i>Heteromys desmarestianus</i> (Desmarest's spiny pocket mouse) | Honduras: Atlantida, Jardin Botanico Lancetilla | 2004 Jul 16 | SAMN31718216 |
| TK136222 | <i>Peromyscus mexicanus</i> (Mexican deer mouse) | Honduras: Colon, Trujillo | 2004 Jul 17 | SAMN31718217 |
| TK136228 | <i>H. desmarestianus</i> | Honduras: Colon, Trujillo | 2004 Jul 17 | SAMN31718218 |
| TK136240 | <i>Glossophaga soricine</i> (Pallas's long-tongued bat) | Honduras: Colon, Trujillo | 2004 Jul 16 | SAMN31718219 |
| TK136756 | <i>Eptesicus furinalis</i> (Argentine brown bat) | Honduras: Colon, Trujillo | 2004 Jul 17 | SAMN31718220 |
| TK136783 | <i>Glossophaga leachii</i> (gray long-tongued bat) | Honduras: Colon, Trujillo | 2004 Jul 17 | SAMN31718221 |
| TK148935 | <i>Rhogeessa tumida</i> (back-winged little yellow bat) | Mexico: Tamaulipas, Soto la Marina | 2008 Jul 27 | SAMN31718222 |
| TK148943 | <i>M. velifer</i> | Mexico: Tamaulipas, Soto la Marina | 2008 Jul 27 | SAMN31718223 |
| TK150290 | <i>Balantiopteryx plicata</i> (gray sac-winged bat) | Mexico: Michoacan, El Marqués | 2006 Jul 22 | SAMN31718224 |
| TK154677 | <i>Gerbilliscus leucogaster</i> (bushveld gerbil) | Botswana: Ngamiland, Koanaka Hills | 2008 Jun 29 | SAMN31718225 |
| TK154685 | <i>G. leucogaster</i> | Botswana: Ngamiland, Koanaka Hills | 2008 Jun 29 | SAMN31718226 |
| TK154687 | <i>G. leucogaster</i> | Botswana: Ngamiland, Koanaka Hills | 2008 Jun 29 | SAMN31718227 |
| TK164683 | <i>Mastomys natalensis</i> (Natal multimammate mouse) | Botswana: Ngamiland, Koanaka Hills | 2009 Jul 18 | SAMN31718228 |
| TK164686 | <i>M. natalensis</i> | Botswana: Ngamiland, Koanaka Hills | 2009 Jul 18 | SAMN31718229 |
| TK164689 | <i>M. natalensis</i> | Botswana: Ngamiland, Koanaka Hills | 2009 Jul 18 | SAMN31718230 |
| TK164690 | <i>M. natalensis</i> | Botswana: Ngamiland, Koanaka Hills | 2009 Jul 18 | SAMN31718231 |
| TK164702 | <i>M. natalensis</i> | Botswana: Ngamiland, Koanaka Hills | 2009 Jul 19 | SAMN31718232 |
| TK164714 | <i>M. natalensis</i> | Botswana: Ngamiland, Koanaka Hills | 2009 Jul 19 | SAMN31718233 |
| TK164728 | <i>M. natalensis</i> | Botswana: Ngamiland, Koanaka Hills | 2009 Jul 19 | SAMN31718234 |
| TK166246 | <i>P. attwateri</i> | United States: Texas, Kerr | 2010 May 17 | SAMN31718235 |
| TK179690 | <i>P. attwateri</i> | United States: Texas, Kerr | 2013 May 20 | SAMN31718236 |
| TK185677 | <i>P. attwateri</i> | United States: Texas, Kerr | 2018 May 21 | SAMN31718237 |
| TK197046 | <i>P. attwateri</i> | United States: Texas, Kerr | 2016 May 26 | SAMN31718238 |
| TK199855 | <i>P. attwateri</i> | United States: Texas, Kerr | 2019 May 21 | SAMN31718239 |

*ID, identification; SRA, National Center for Biotechnology Information Sequence Read Archive.

individual libraries with similar DNA concentrations into pools of 4 samples and used the myBaits version 5 (Daicel Arbor Biosciences) high sensitivity protocol to enrich target loci. We used 2 rounds of enrichment (24 h at 65°C), washed away unbound DNA, and amplified the remainder for 15 cycles before pooling for sequencing.

Classifying Reads

First, we generated a dataset of target loci by mapping the probes to representative and reference genomes in RefSeq v212 with BMap v38.96 (30). For each probe, we kept the 10 best sites that mapped with $\geq 85\%$ sequence identity along with 1,000 bp upstream and downstream. These sequences were combined into a database to classify reads by using Kraken2 version 2.1.1 (31) (Figure 3, panel A). Next, we extracted pathogen reads with KrakenTools version 1.2 (<https://github.com/jenniferlu717/KrakenTools>). We assembled those reads (Figure 3, panel B) with the SPAdes genome assembler version 3.14.1 (32) and filtered them to remove low quality contigs (<100 bp and <10 \times median coverage). We removed samples that had <2 contigs from downstream analyses. During this time, we extracted target loci in available reference genomes (Figure 3, panel C). Next, we identified (Figure 3, panel D), aligned and trimmed (Figure 3, panel E) orthologs before concatenating them into a single alignment (Figure 3, panel F). Finally, we

generated and bootstrapped a phylogenetic tree (Figure 3, panel G) by using RaxML-NG version 1.0.1 (33). We repeated those steps for each pathogen group (Figure 3, panel H).

Host Identification

There were sufficient mtDNA sequences from most samples to verify museum identifications by comparing reads to a Kraken2 version 2.1.2 (31) database of mammalian mitochondrial genomes. We filtered the classifications by removing samples with <50 classified reads and single-read, generic classifications.

Results

Panel Development

We used the ultraconserved element protocol developed by Faircloth et al. (26,27) to develop a set of 39,893 biotinylated baits that target 32 pathogen groups responsible for 32 zoonoses. Each pathogen group is targeted at 49 loci with a few diverse taxa, *Bacillus cereus* and *Trypanosoma* species, targeted at 98 loci. We compiled information on pathogen groups, focal taxa, genome accessions, and number of baits (Table 3).

Control Samples

We tested the efficacy of our bait set on laboratory-made host–pathogen mixtures containing DNA from

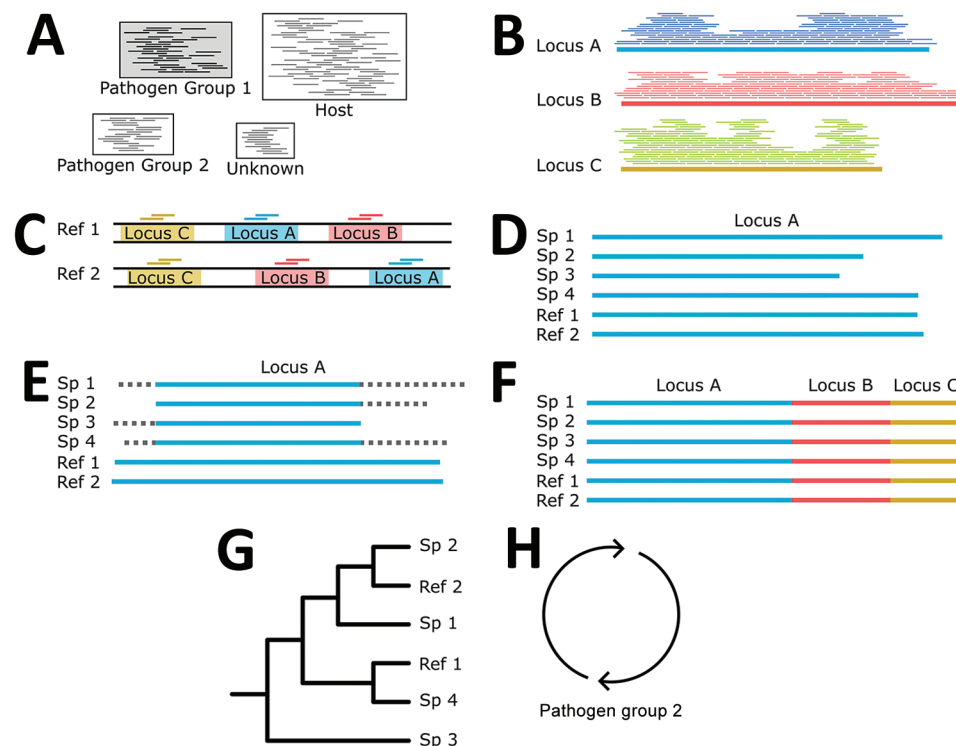


Figure 3. Building phylogenies from parasite reads for study of prospecting for zoonotic pathogens by using targeted DNA enrichment. A) After read classification, we extracted all the reads associated with a pathogen group. B) Those reads were assembled into contigs with a genome assembler. C) Simultaneously, we identified and extracted the target loci from all members of the pathogen group with available reference genomes to ensure that our final phylogeny has representatives from as many members of the pathogen group as possible. D, E) For each targeted locus, we combined the assembled contigs (D) and genome extracted loci for (E) multiple sequence alignment and trimming. F, G) Each aligned and trimmed locus is concatenated together (F) for phylogenetic analyses (G). H) If necessary, those steps are repeated for reads classified in other pathogen groups. Ref, reference; Sp, specimen.

Table 3. Summary of probes developed for targeted capture of pathogen DNA in study of prospecting for zoonotic pathogens by using targeted DNA enrichment

| Pathogen group | Type | Probe count | Locus count | RefSeq genome count | Focal pathogen | GenBank accession no. |
|-------------------------------|-----------|-------------|-------------|---------------------|-------------------------------------|-----------------------|
| <i>Anaplasma</i> | Bacteria | 368 | 49 | 57 | <i>Anaplasma phagocytophilum</i> | GCF000013125 |
| Apicomplexa | Eukaryote | 3,219 | 49 | 64 | <i>Plasmodium falciparum</i> | GCA000002765 |
| <i>Bacillus cereus</i> group* | Bacteria | 833 | 98 | 134 | <i>Bacillus anthracis</i> | GCF000008165 |
| <i>Bartonella</i> | Bacteria | 1,812 | 49 | 31 | <i>Bartonella bacilliformis</i> | GCF000015445 |
| <i>Borrelia</i> | Bacteria | 688 | 49 | 16 | <i>Borrelia burgdorferi</i> | GCF000502155 |
| <i>Burkholderia</i> | Bacteria | 683 | 49 | 39 | <i>Burkholderia mallei</i> | GCF000011705 |
| <i>Campylobacter</i> | Bacteria | 2,194 | 49 | 33 | <i>Campylobacter jejuni</i> | GCF000009085 |
| Cestoda | Eukaryote | 907 | 49 | 18 | <i>Taenia multiceps</i> | GCA001923025 |
| <i>Chlamydia</i> | Bacteria | 830 | 49 | 15 | <i>Chlamydia trachomatis</i> | GCF000008725 |
| <i>Coxiella</i> | Bacteria | 144 | 49 | 70 | <i>Coxiella burnetii</i> | GCF000007765 |
| <i>Ehrlichia</i> | Bacteria | 235 | 49 | 7 | <i>Ehrlichia canis</i> | GCF000012565 |
| Eurotiales | Eukaryote | 4,097 | 49 | 158 | <i>Talaromyces marneffeii</i> | GCF000001985 |
| <i>Francisella</i> | Bacteria | 470 | 49 | 14 | <i>Francisella tularensis</i> | GCF000008985 |
| Hexamitidae | Eukaryote | 782 | 49 | 19 | <i>Giardia intestinalis</i> | GCA000002435 |
| Kinetoplastea | Eukaryote | 2,917 | 49 | 49 | <i>Leishmania major</i> | GCF000002725 |
| <i>Leptospira</i> | Bacteria | 2,517 | 49 | 69 | <i>Leptospira interrogans</i> | GCF000092565 |
| <i>Listeria</i> | Bacteria | 765 | 49 | 23 | <i>Listeria monocytogenes</i> | GCF000196035 |
| <i>Mycobacterium</i> | Bacteria | 2,463 | 49 | 86 | <i>Mycobacterium tuberculosis</i> | GCF000195955 |
| Nematodes, clade I | Eukaryote | 357 | 49 | 13 | <i>Trichinella spiralis</i> | GCA000181795 |
| Nematodes, clade III | Eukaryote | 1,494 | 49 | 25 | <i>Brugia malayi</i> | GCA000002995 |
| Nematodes, clade IVa | Eukaryote | 252 | 49 | 7 | <i>Strongyloides stercoralis</i> | GCA000947215 |
| Nematodes, clade IVb | Eukaryote | 1,487 | 43 | 34 | <i>Steinernema carpocapsae</i> | GCA0000757645 |
| Nematodes, clade V | Eukaryote | 3,242 | 48 | 47 | <i>Haemonchus contortus</i> | GCA007637855 |
| Onygenales | Eukaryote | 1,973 | 49 | 38 | <i>Histoplasma capsulatum</i> | GCF000149585 |
| <i>Pasteurella</i> | Bacteria | 615 | 49 | 11 | <i>Pasteurella multocida</i> | GCF000754275 |
| <i>Rickettsia</i> | Bacteria | 394 | 49 | 37 | <i>Rickettsia rickettsii</i> | GCF001951015 |
| <i>Salmonella</i> | Bacteria | 145 | 49 | 35 | <i>Salmonella enterica</i> | GCF001159405 |
| <i>Streptobacillus</i> | Bacteria | 245 | 49 | 7 | <i>Streptobacillus moniliformis</i> | GCF000024565 |
| Trematoda | Eukaryote | 924 | 49 | 18 | <i>Schistosoma mansoni</i> | GCA000237925 |
| Tremellales | Eukaryote | 1,999 | 49 | 26 | <i>Cryptococcus neoformans</i> | GCF000091045 |
| Trypanosoma* | Eukaryote | 617 | 97 | 10 | <i>Trypanosoma cruzi</i> | GCF000209065 |
| <i>Yersinia</i> | Bacteria | 225 | 49 | 22 | <i>Yersinia pestis</i> | GCF000009065 |

*Supplemented.

Mus musculus, *Mycobacterium tuberculosis*, *Plasmodium falciparum*, *P. vivax*, and *Schistosoma mansoni*. We generated 4 control samples containing either 1% or 0.001% pathogen DNA that was enriched or not enriched. We classified reads against the database of target loci and found that 42.7% of all reads (*Mycobacterium* = 13.1%, *Plasmodium* = 28.1%, *Schistosoma* = 1.5%) were from control pathogens in the 1% enriched control sample. However, only 0.03% of the corresponding 1% unenriched control was from target loci. Aside from the raw percentages, we compared the coverage of each probed region in the 1% enriched and unenriched control samples (Figure 4, panels B–D) to understand how enrichment effected coverage at each locus. Mean coverage per *Mycobacterium* locus increased from 0.14× to 944.5× (6,746-fold enrichment), 0.53× to 1,527.4× for *Plasmodium* loci (2,882-fold enrichment), and 0.02× to 117.9× (5,895-fold enrichment) for schistosome loci. Because the sequencing library from the 0.001% unenriched sample did not work during the sequencing reaction, we do not have a baseline to examine enrichment in the 0.001% samples.

We extracted reads assigned to each pathogen group and assembled and aligned them with target loci extracted from reference genomes of closely related species by using tools from PhyLuce version 1.7.1 (26,27). We were able to assemble 0–23 target loci per pathogen group in the control samples (Table 4). Assembled loci varied in size from 109 to 1,991 bp (median 636.5 bp). For each sample/group with >2 loci captured, we generated a phylogenetic tree along with other members of the taxonomic group (Figure 5). In each case, pathogen loci from the control samples were sister groups to the appropriate reference genome with strong bootstrap support. For example, the *Schistosoma* loci assembled from the 1% enriched control sample were sister to the *S. mansoni* genome (GCA000237925) in 100% of bootstrap replicates.

Museum Samples

Next, we tested our bait set on museum-archived tissues. We generated 649.3 million reads across all 38 samples (mean 17.1 million reads/sample). An initial classification showed that, on average, 4.3% of reads

were assignable to loci in the database. Those reads were designated to 93 genera. However, 78 of those genera were at low frequency ($\leq 1,000$ reads/sample) (Figure 4). Many of the low frequency hits are

likely the result of bioinformatic noise. *Bartonella* and *Plasmodium* species were the most common genera; each was present in 36 of 38 museum samples. The distribution of *Bartonella* reads was strongly bimodal

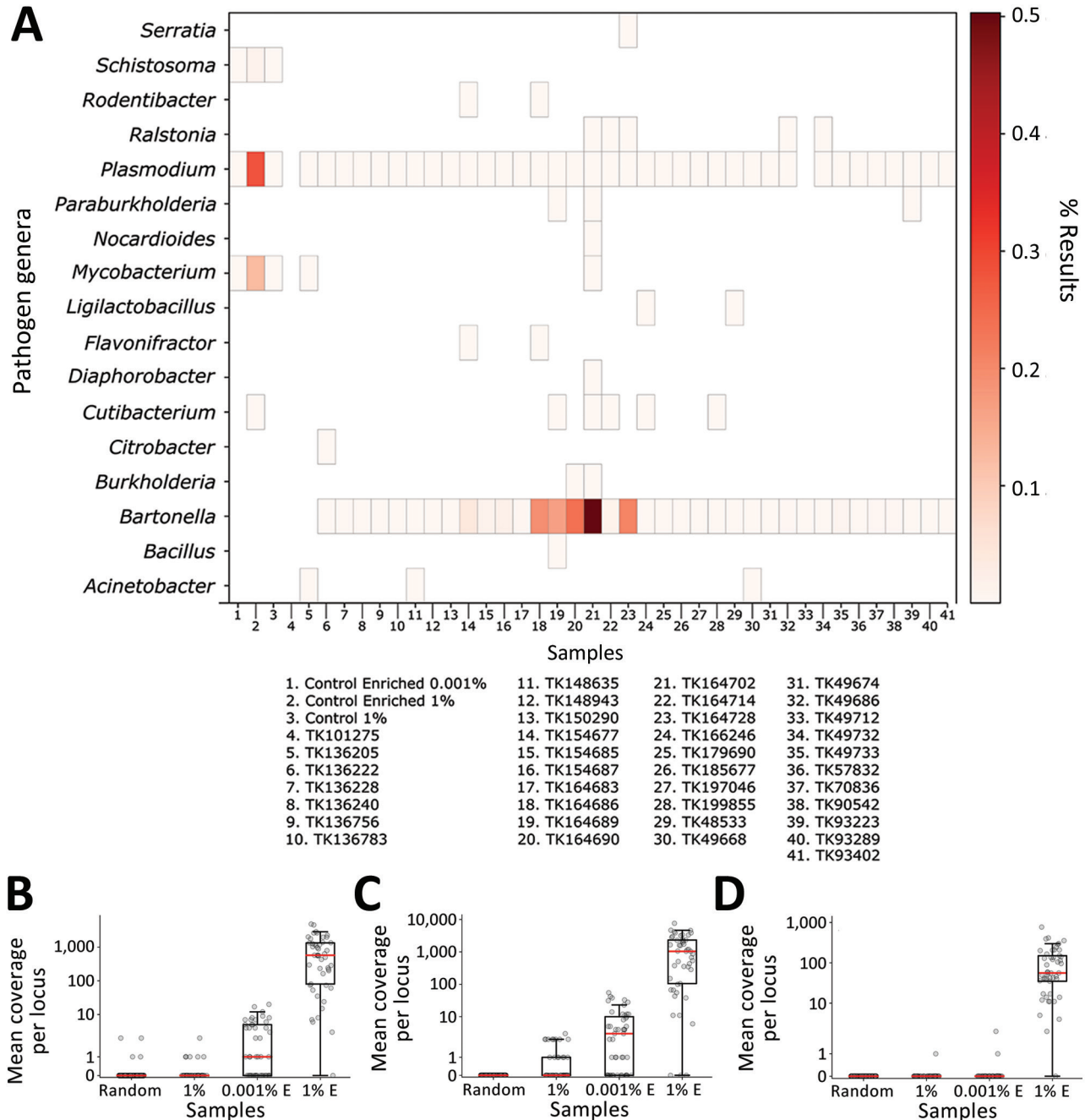


Figure 4. Identifying pathogen reads from controls and museum-archived tissue samples for study of prospecting for zoonotic pathogens by using targeted DNA enrichment. Control reads are indicated by the percentage of pathogen DNA 1% or 0.001%. A) Reads were compared with a database of target loci and assigned a taxonomic classification based on these results. Reads were assigned to 93 genera; of those, 17 (shown) were present in >1 sample, including controls, with $\geq 1,000$ reads. A heatmap of those results shows the relative proportion of reads assigned to each genus. Details of samples are provided in Table 2. B–D) Coverage at each probed locus is shown across all control samples for *Mycobacterium* (B), *Plasmodium* (C), and *Schistosoma* (D). Each point in the chart is coverage calculated at a single target locus. Horizontal lines within boxes indicate medians, box tops and bottoms indicate lower and upper quartiles, and whiskers represent minimum and maximum values, excluding outliers. Each sample is indicated with a circle. E, enriched.

Table 4. Parasite reads identified in and loci assembled from control samples

| Enriched | Pathogen concentration, % | Total reads | <i>Schistosoma</i> | | <i>Plasmodium</i> | | <i>Mycobacterium</i> | |
|----------|---------------------------|-------------|--------------------|------|-------------------|------|----------------------|------|
| | | | Reads | Loci | Reads | Loci | Reads | Loci |
| True | 0.001 | 509,672 | 3 | 0 | 168 | 7 | 556 | 0 |
| True | 1 | 398,469 | 5,879 | 23 | 52,274 | 8 | 112,141 | 23 |
| False | 1 | 375,786 | 15 | 0 | 17 | 0 | 83 | 0 |

such that 18 samples had <12 reads and 18 samples had >1,000 reads (median 552 reads/sample). In 5 samples, the percentage of *Bartonella* reads was exceedingly high (>10%). In comparison, the median number of *Plasmodium* reads never exceeded 0.04% of reads from a single museum sample (mean 158.5 reads/sample).

We used phylogenetic analyses and rules of monophyly to identify putative pathogens to species or strain for each of the 15 genera with $\geq 1,000$ reads (Figure 4, panel A). We were unable to assemble >1 target locus for any specimen in 13 genera. We were able to assemble 3–20 loci (mean 8 loci/sample) from 16 samples containing *Bartonella* (Figure 6), 3 loci from a sample containing *Paraburkholderia* reads (Figure 7), and 8 loci from a sample containing *Ralstonia* reads (Figure 8).

Host Identification

We compared reads from each sample to a database of mitochondrial genomes to identify the host. In gen-

eral, reads from the mitochondria comprised a small proportion ($\leq 1\%$, mean 0.04%) of each sample (Figure 9). Despite the low number of mitochondrial reads, generic classifications from the mitochondrial database coincided with the museum identifications after filtering samples with ≤ 50 mitochondrial reads. For the remaining samples, the correct genus was identified by >85% (mean 98%) of reads from that sample. Classifying reads less than the generic level is limited by mitochondrial genome availability, but where possible, we were able to confirm museum identifications at the species level.

Discussion

We developed a set of 39,893 biotinylated baits for targeted sequencing of >32 zoonotic pathogens, and their relatives, from host DNA samples. To test the efficacy of the bait panel, we used 4 control samples that contained either 1% or 0.001% pathogen DNA and further subdivided into pools that were enriched and unenriched. Our results (Figure 4) showed a

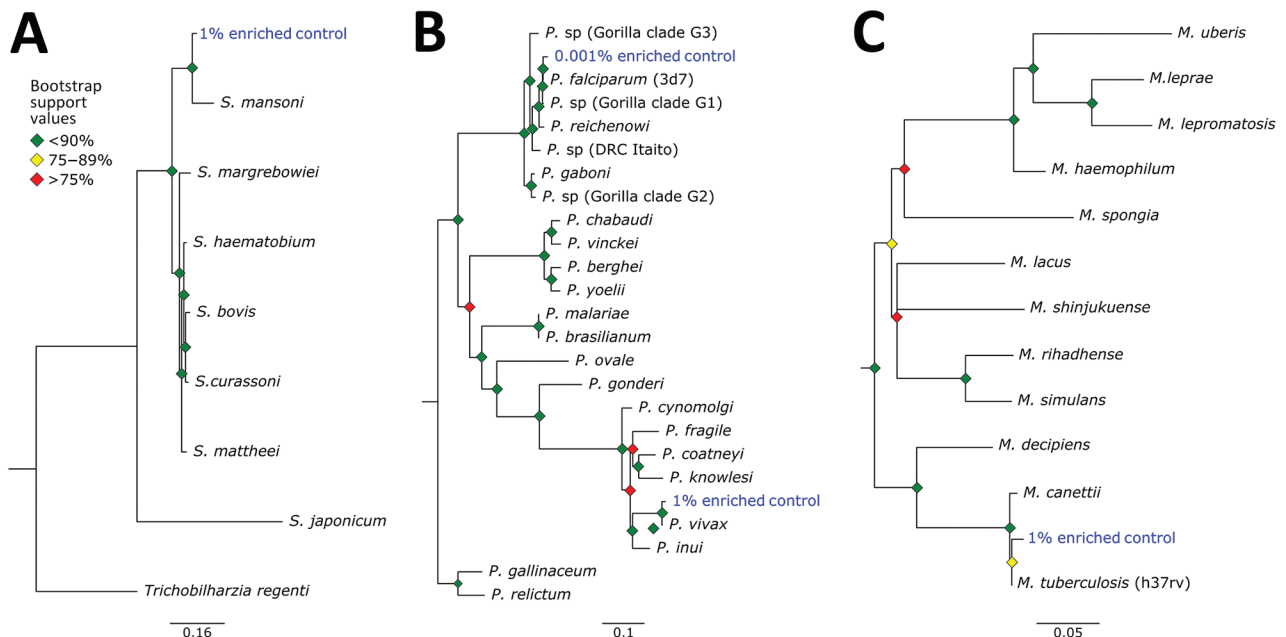


Figure 5. Phylogenetic analysis of pathogens used in control samples for study of prospecting for zoonotic pathogens by using targeted DNA enrichment. A) *Schistosoma*; B) *Plasmodium*; C) *Mycobacterium*. Reads from each control pathogen (*M. tuberculosis*, *P. falciparum*, *P. vivax*, and *S. mansoni*) were extracted, assembled, aligned, and trimmed for maximum-likelihood phylogenetic analyses. The phylogenies were used to identify the species or strain of pathogen used in the controls. Blue indicates control samples. Bootstrap support values are indicated by colored diamonds at each available node. Branches with <50% bootstrap support were collapsed. Nodal support is indicated by color coded diamonds. Scale bars indicate nucleotide substitutions per site. Assembly accession numbers (e.g., GCA902374465) and tree files are available from <https://doi.org/10.5281/zenodo.8014941>.

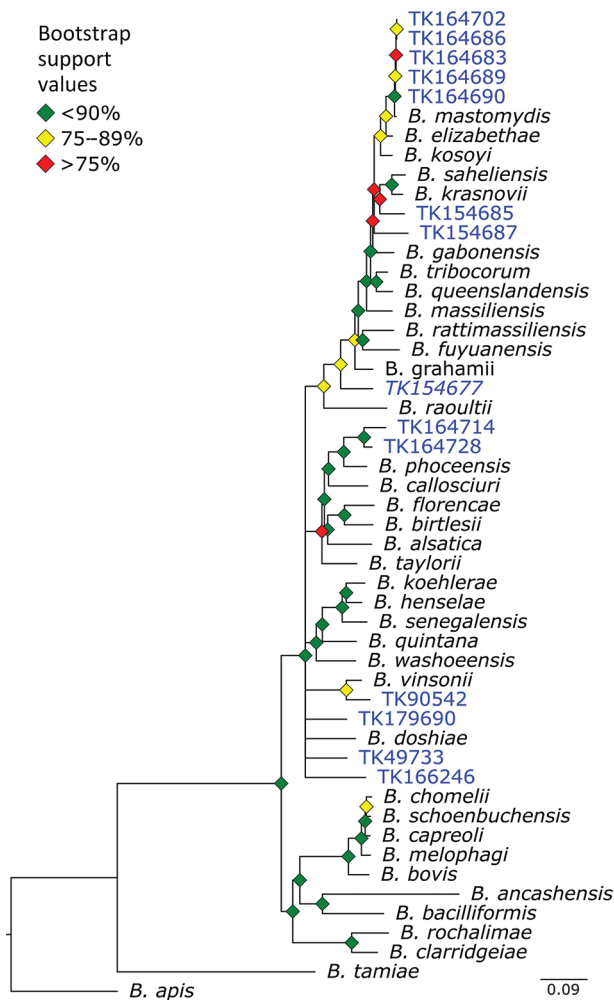


Figure 6. Phylogenetic analysis of *Bartonella* using museum archived samples in study of prospecting for zoonotic pathogens by using targeted DNA enrichment. Blue indicates museum archived samples; museum accession numbers are given (Table 1). Branches with <50% bootstrap support were collapsed. Nodal support is indicated by color coded diamonds. Scale bar indicates nucleotide substitutions per site. Assembly accession numbers (e.g., CA902374465) and tree files are available from <https://doi.org/10.5281/zenodo.8014941>.

large increase of pathogen DNA in the 1% enriched sample when compared with its unenriched counterpart. Specifically, enrichment increased the amount of pathogen DNA from 0.03% to 42.1%.

We were able to generate phylogenetically informative loci from *Plasmodium*, *Mycobacterium*, and *Schistosoma* species in the 1% enriched control sample. On the basis of genome size, we estimate genome copies as 91,611 for *Plasmodium*, 261,030 for *Mycobacterium*, and 3,159 for *Schistosoma* in the control sample. This finding indicates that the probe set is able to detect these pathogens from even a few thousand

genome copies per sample (*Schistosoma* species). In contrast, we were only able to generate phylogenetically informative loci from *P. falciparum* in 0.001% enriched sample, which would hypothetically contain ≈ 39 genome copies. This finding implies that the bait set might be capable of identifying pathogens present in samples with only a few hundred genome copies. However, there are limitations to *Plasmodium* detection that should be considered.

In each sample, reads were detected from only a few loci rather than from the entire genome. For example, in the 1% enriched sample, 5,879 of the 398,469 reads came from 32 loci totaling 19.6 kb. Had the unenriched sample contained the same number of reads, randomly distributed across the genome, it would have amounted to 1 read every 62 kb. We found that enrichment increased coverage at probed loci from 0.23 \times to 863.3 \times , a 3,732.3-fold increase when averaged across all pathogens/loci (Figure 4). Those results show that although large amounts of host DNA might remain in a sample, the targeted loci are greatly enriched.

We tested the panel of baits on 38, museum-archived, small mammal samples without previous knowledge of infection history. Reads from these samples were initially designated to 93 different genera, but most of these genera contained a limited number of reads. For example, almost half of the 93 genera ($n = 43$) were identified on the basis of a single read across all 38 samples, most likely a bioinformatic artifact. We identified 15 genera in which 1 sample had $\geq 1,000$ reads. For each of these 15 genera, we extracted any reads classified within the same family (e.g., genus *Bartonella*, family Bartonellaceae) and assembled, aligned, and trimmed them for phylogenetic analyses. In most cases, the reads failed the assembly step ($n = 6$), were filtered on the basis of locus size or coverage ($n = 5$), or assembled into multiple loci that were not targeted by our bait set ($n = 2$); we did not pursue those reads any further. However, we were able to generate phylogenies for specimens positive for *Bartonella*, *Ralstonia*, and *Paraburkholderia* species.

Bartonella is a bacterial genus responsible for cat-scratch disease, Carrión's disease, and trench fever (34). Transmission often occurs between humans and their pets or from infected fleas ticks, or other arthropod vectors (35). We were able to recover target loci for 14 of 36 specimens. A phylogeny of *Bartonella* species placed the museum samples in multiple clades (Figure 6). For example, 5 specimens formed a monophyletic clade sister to *B. mastomydis*. *B. mastomydis* recently was described from *Mastomys erythroleucus* mice collected in Senegal (36). Appropriately, the

samples we tested were collected from *M. natalensis* mice from Botswana (Table 2). Another clade contained *B. vinsonii* and a *Sigmodon* rat (TK90542) collected in Mexico. Zoonotic transmission of *B. vinsonii* has been implicated in neurologic disorders (37). Other museum samples probably contain novel *Bartonella* species/strains or at least represent species/strains without genomic references.

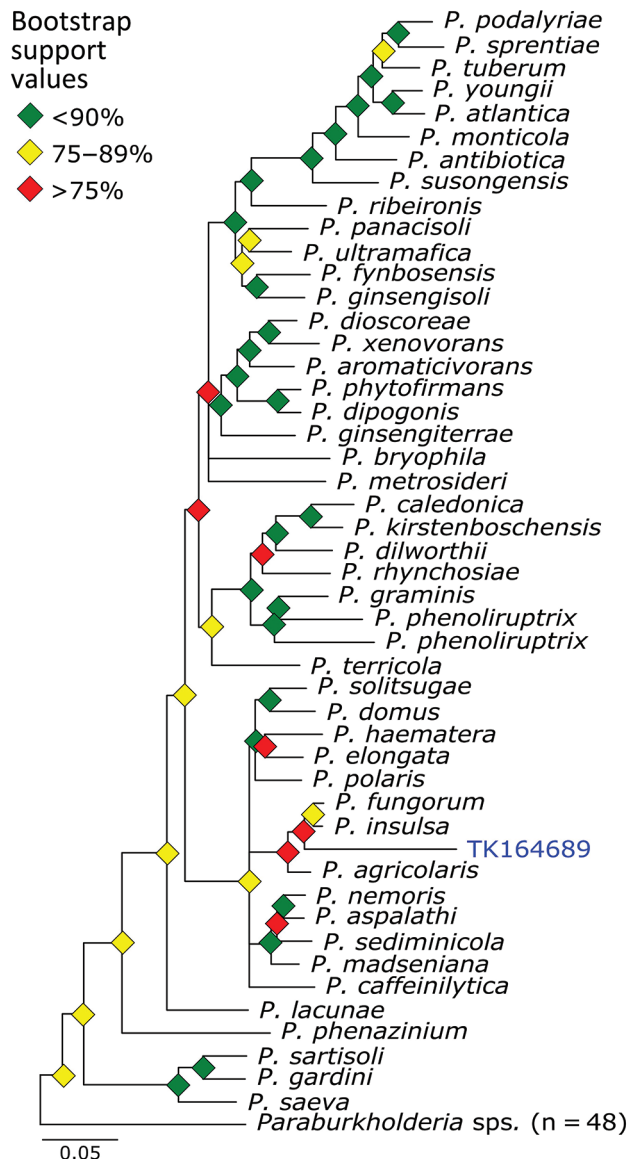


Figure 7. Phylogenetic analysis of *Paraburkholderia* using museum archived samples in study of prospecting for zoonotic pathogens by using targeted DNA enrichment. Blue indicates museum archived samples; museum accession numbers are given (Table 1). Branches with <50% bootstrap support were collapsed. Nodal support is indicated by color coded diamonds. Scale bar indicates nucleotide substitutions per site. Assembly accession numbers (e.g., GCA90237446) and tree files are available from <https://doi.org/10.5281/zenodo.8014941>.

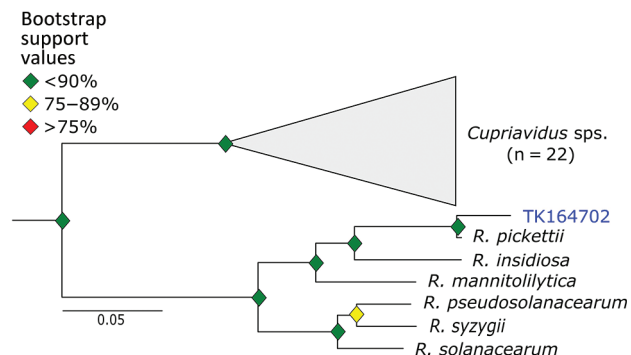


Figure 8. Phylogenetic analysis of *Ralstonia* using museum archived samples in study of prospecting for zoonotic pathogens by using targeted DNA enrichment. Blue indicates museum archived samples; museum accession numbers are given (Table 1). Branches with <50% bootstrap support were collapsed. Nodal support is indicated by color coded diamonds. Scale bar indicates nucleotide substitutions per site. Assembly accession numbers (e.g., GCA90237446) and tree files are available from <https://doi.org/10.5281/zenodo.8014941>.

Paraburkholderia is a genus of bacteria commonly associated with soil microbiomes and plant tissues. We identified *Paraburkholderia* reads in 3 specimens and were able to place 1 of those in a phylogeny sister to a clade containing *P. fungorum* and *P. insulsa*. Because bootstrap values across the phylogeny were moderate in general, and weak in this particular region (Figure 7), placement of this sample is tenuous. *P. fungorum* is the sole member of *Paraburkholderia* believed to be capable of infecting humans, but it is only a rare, opportunistic, human pathogen (38–40).

Ralstonia is a bacteria genus closely related to the genus *Pseudomonas*. We identified *Ralstonia* reads in 5 samples and were able to place a specimen on a phylogeny. This sample is closely affiliated with *R. pickettii* (Figure 8). We are unaware of any examples of zoonotic transmission of *R. pickettii*. Rather, *R. pickettii* has been identified as a common contaminant in laboratory reagents (41), and outbreaks have been caused by contaminated medical supplies (42). We failed to identify nucleic acids in any of our negative controls during library preparation. Furthermore, if there were systemic contamination, we would expect to find *Ralstonia* species in all of our samples, rather than the 5 of 36 observed. Thus, because we cannot rule out reagent contamination, the presence of *Ralstonia* species in the museum samples should be interpreted with caution.

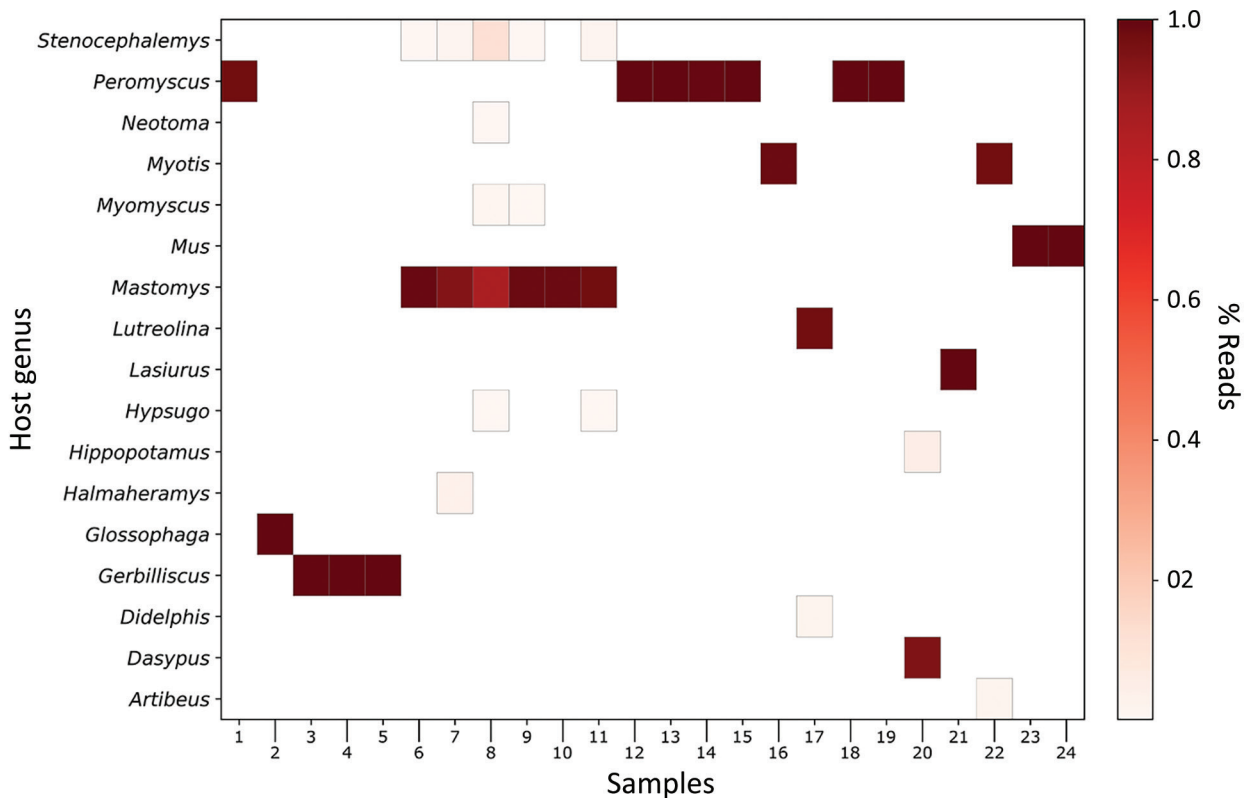
We were able to capture, sequence, and assemble loci from taxa that were not represented in the databases used to design the bait panel. This ability was possible for 2 reasons. First, the bait panel is highly redundant. The baits are sticky and able to capture

nucleic acid fragments that are $\leq 10\%$ – 12% divergent (43). We designed the panel with $\leq 5\%$ sequence divergence between any pair of baits at a particular locus (Figure 10). Second, sampled loci within each pathogen group spanned a range of divergences. Conserved loci were more likely to catch more divergent species that might not have been present in our initial dataset. For example, we recovered multiple species of *Bartonella* that were not present in our probe set, for which related genomes were available. However, for *Ralstonia* and *Paraburkholderia* species, we identified these samples from reads targeted by probes for the genus *Burkholderia*, a pathogenic taxon in the same family (Burkholderaceae). The ability to identify taxa at these distances is because of the more conserved loci targeted by the bait panel.

During the initial read classification stage, we identified low levels of *Plasmodium* species in all but 2 museum samples, which was unexpected. Museum

samples contained $\leq 3,221$ *Plasmodium* reads/sample (mean 428.3 reads/sample), but we were unable to assemble them into loci for phylogenetic analyses. This limitation effectively removed those samples from downstream analyses. The *P. falciparum* genome is extremely AT rich (82%, 44), which might result in bioinformatic false-positive results. We suspect that AT-rich, low-complexity regions of the host genome are misclassified as parasite reads. To test this hypothesis, we used fqtrim 0.9.7 (<https://ccb.jhu.edu/software/fqtrim>) to identify and remove low-complexity sequences within those reads. This filter by itself reduced the number of *Plasmodium* reads in the museum samples by 75.5% (maximum 298 reads, mean 57.2 reads). In comparison, only 8.2% of reads from 0.001% enriched control samples and 0.2% of reads from 1% enriched control samples were removed.

Several technical issues still need to be addressed. First, enrichment increases the targeted



- 1. TK136222 6. TK164683 11. TK164728 16. TK48533 21. TK49732
- 2. TK136240 7. TK164686 12. TK179690 17. TK49668 22. TK49733
- 3. TK154677 8. TK164689 13. TK185677 18. TK49674 23. Control enriched 0.001%
- 4. TK154685 9. TK164702 14. TK197046 19. TK49686 24. Control 1%
- 5. TK154687 10. TK164714 15. TK199855 20. TK49712

Figure 9. Genetic identification of mammal host from unenriched, mitochondrial reads in study of prospecting for zoonotic pathogens by using targeted DNA enrichment. Reads were compared with a database of mammalian mitochondria and assigned a taxonomic classification based on these results. A heatmap of the results shows the relative proportion of classified reads assigned to mammalian genera. Samples with <50 mitochondrial reads and single-read genera are not shown.

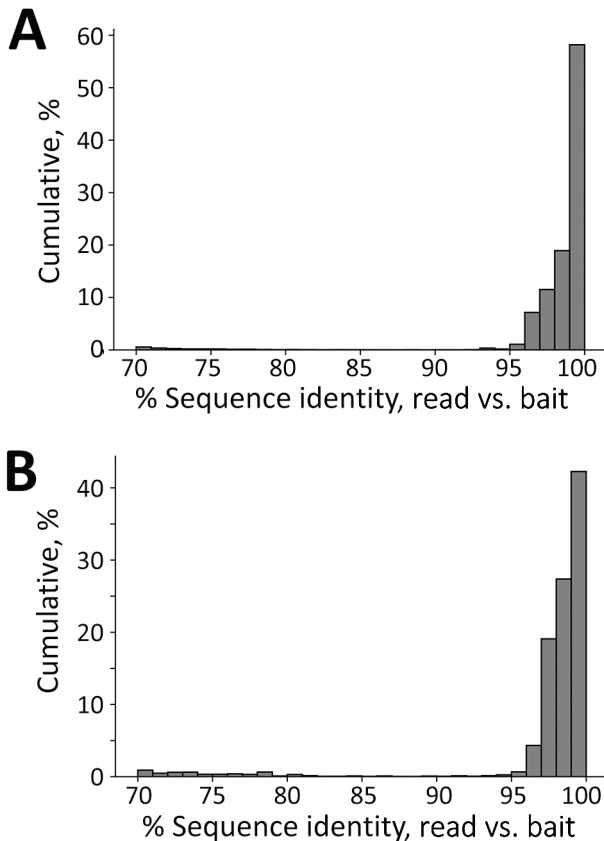


Figure 10. Sequence identity between enriched reads and baits in the probe panel used for targeting zoonotic pathogens in study of prospecting for zoonotic pathogens by using targeted DNA enrichment. Reads from each sample were classified against a database of target loci. Sequence identity between pathogen-derived reads and the most similar bait in the bait panel for all pathogens excluding *Bartonella* species (A) and for only *Bartonella* species (B). *Bartonella* was the most common pathogen in our samples, and the number of reads was biased toward a few individuals.

loci coverage by 3 orders of magnitude. However, the amount of host DNA remaining in each sample is still high. Ideally, host DNA would be rare or absent. Second, the bait panel requires relatively large up-front costs. Third, although the bait panel is developed to target a wide range of taxa, it is not possible to know which species are missed. The best way to circumvent that issue is to use controls spiked with various pathogens of interest, similar to how mock communities are used in other metagenomic studies (45). Those mock controls are commercially available for bacterial communities (e.g., ZymoBIOMICS Microbial Community Standards; Zymo Research, <http://www.zymoresearch.com>), but we have been unable to find similar products that contain eukaryotic pathogens. Solutions to those problems will make

targeted sequencing with bait panels a viable tool for pathogen surveillance. Fourth, the sensitivity of the probes will depend on the sequence divergence between the probes and pathogen DNA. The more diverged the 2 are, the less efficient the capture will be. This limitation indicates that pathogen groups that have biased or limited genomic data will be less likely to capture off-target species once divergence increases by >5%–10%. Finally, the current probe panel is capable of capturing and identifying pathogens if there are $\geq 3,000$ genome copies in the sample. Sensitivity needs to be improved in future iterations of the panel. One method could be to target pathogen-specific, repetitive sequences (46). Because those sequences are already present in the genome hundreds to thousands of times, it should be possible to greatly increase the sensitivity of the probe panel.

Although further effort is required to resolve these issues, we believe that enrichment of pathogen DNA from museum tissue samples is a viable tool worth further development. In its current form, enrichment represents a coarse tool that can be used to scan for various pathogens from archived tissues. More refined tests, such as quantitative PCR and targeted sequencing, can be used to answer taxon-specific questions. Target enrichment will be necessary for maximizing the pathogen data that are available from the hundreds of thousands of museum-archived tissues and will play a critical role in understanding our susceptibility to future zoonotic outbreaks.

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Prospecting for Zoonotic Pathogens by Using Targeted DNA Enrichment

Appendix 1

Materials and Methods

Host-Pathogen Control Samples

We isolated DNA using the QIAGEN blood and tissue kit following manufacturer's protocol and quantified DNA using Qubit. We prepared a cocktail of pathogen DNA mixtures comprising 200 ng DNA of each pathogen (*Mycobacterium bovis*, *M. tuberculosis*, *Plasmodium vivax*, *P. falciparum*, *Schistosoma mansoni*, and *S. bovis*). A mammalian-pathogen DNA mixture was prepared by mixing pathogen DNA in DNA from uninfected liver tissues of laboratory mouse (*Mus musculus*) to make 1% and 0.001% pathogen mixtures. The negative control was prepared from the same uninfected liver tissues of laboratory mouse (*Mus musculus*) without spiking with pathogens.

Museum-Archived Samples and Controls

We extracted DNA from 42 museum samples comprising of mammalian liver tissues (in lysed buffer or frozen in liquid nitrogen) collected between 1995 and 2018 in Africa, Southern America, and the United States. Control samples were as previously described (1% and 0.001% pathogens DNA in mammalian DNA). Information for each specimen are provided in Table 2.

Computing Environment and Reproducibility

All analyses were performed on a single compute node with 48 processors and limited to 100 Gb of RAM. Bioinformatic steps were documented in a series of BASH shell scripts or Jupyter v4.9.2 notebooks. These files along with conda v4.11.0 environments are available (github.com/nealplatt/pathogen_probes) and are archived: DOI:10.5281/zenodo.7319915.

Panel Development

We developed a set of biotinylated probes for UCE-based, targeted sequencing of 32 pathogen groups (Table 1). For example, given the large evolutionary distances covered by various pathogens, we generated sets of probes that target more discrete taxonomic groups (e.g., Nematoda, *Yersinia*). For bacterial pathogens, probes were designed to capture all species within the genus or species group. For eukaryotic pathogens, probes were designed to be effective at taxonomic ranks that ranged from species group to class. The taxonomic rank varied in eukaryotic pathogens based on the following criteria: 1) the number of available genomes, 2) sequence diversity - because this impacted the number of probes needed. Table 1 provides information on the pathogen group, targeted zoonotic agent and zoonoses.

For each group we used the Phyluce package v1.7.0 (1,2); we generated probes to target ≈ 49 loci using the methods described below. First, we identified orthologous loci between a focal pathogen and the remaining species in the pathogen group. Focal taxa were chosen based on their assembly contiguity or prominence as a zoonotic agent. To do this we downloaded a genome for each species in the pathogen group. Accession numbers for these assemblies are provided in Table 2. Next, we simulated 25x read coverage for each genome using the ART v2016.06.05 (3); read simulator with the following options: `art_illumina-paired-len 100-fcov 25-mflen 200-sdev 150 -ir 0.0 -ir2 0.0 -dr 0.0 -dr2 0.0 -qs 100 -qs2 100 -na`. Simulated reads from all query taxa were mapped back to a focal taxon with `bbmap v38.93` (4); enabling up to 10% sequence divergence (`minid = 0.9`). Unmapped, or multimapping reads were removed using `Bedtools v2.9.2` (5) and `phyluce_probe_strip_masked_loci_from_set (filter_mask 25%)`. The remaining reads were merged to generate a BED file containing orthologous regions between the query and focal taxa.

Then, we identified orthologous loci among all taxa within the pathogen group using `phyluce_probe_query_multi_merge_table`. Next, we filtered each set of loci to retain only those shared among 33% of taxa in the pathogen group using `phyluce_probe_query_multi_merge_table`. We extracted 160 bp from each locus and generated an initial set of in silico probes directly from the focal genome using `phyluce_probe_get_genome_sequences_from_bed` and `phyluce_probe_get_tiled_probes`. Additional options for probe design included generating two probes per locus (`-two_probes`) that overlapped in the middle (`-overlap-middle`). Focal probes with repetitive regions or skewed GC

content (<30% or >70%) content were removed. Next, the probes from the focal taxa were mapped back to each genome in the pathogen group with `phyluce_probe_run_multiple_lastzs_sqlite`. We used the `-identity` option to limit searches with a maximum divergence of 30%. Using these results, we extracted 120-bp loci from the probed regions in each representative genome extracted using `phyluce_probe_slice_sequence_from_genomes`. Theoretically, this dataset should contain orthologous 120-bp sequences from most taxa in each pathogen group. We verified this with `phyluce_probe_get_multi_fasta_table`, which provides a table showing the number of taxa identified at each locus. We used this information to identify the 100 loci capable of capturing most taxa from the pathogen group. Next, we generated two 80-bp probes from each of the 100-bp and 120-bp loci. We used `phyluce_probe_easy_lastz` to compare the probes to themselves and remove any that were possible duplicates. Then we reduced the probe set even further by clustering probes based on sequence identity with `cd-hit-est v4.8.1` (6). We identified sequence clusters with >95% similarity and retained only 1 probe per group. Finally, we recalculated the number of probes needed to capture each locus.

The proceeding steps were repeated for each pathogen group shown in Table 1. To generate a final panel, we selected 49 loci per pathogen group in a way that minimized the number of probes needed. In some cases, we needed to generate 2 sets of probes to adequately represent target pathogens. For example, Kinetoplastea contains 2 pathogens of interest, *Trypanosoma* and *Leishmania*. The baits designed for *Leishmania* were able to target all 49 loci in the most of the Kinetoplastea but only 23 loci in *Trypanosoma*. We then generated a second set of 617 *Trypanosoma*-specific baits to augment the kinetoplastid baits and ensure that *Trypanosoma* parasites were represented adequately in the final panel. Likewise, we doubled the number of baits used to capture loci from the *Bacillus cereus* group to effectively capture *B. cereus* and *B. anthracis*. The probe set was quality checked by Arbor Biosciences. This included comparing the probe set to mammal genomes with `blastn v2.12.0` (7) and checking for low-complexity sequences. Any probes that failed quality control were replaced before synthesis.

Library Preparation

Standard DNA sequencing libraries were generated from 500 ng of DNA per sample. We used the KAPA Hyperplus kit protocol with the following modifications: 1) enzymatic fragmentation at 37°C for 10 minutes, 2) adaptor ligation at 20°C for an hour, and 3) four cycles

of library PCR amplification. To minimize adaptor switching we used unique dual indexed (UDI) adaptors (IDT xGen Stubby Adaptor-UDI Primers). Each library was eluted in 20 μ L of sterile water and the base pairs sizes and concentration estimated by Agilent 4200 TapeStation (Figure 2).

Individual samples with similar DNA concentrations were combined together into pools of 4–16 samples and the total volume was reduced to 7 μ L with a speedvac vacuum concentrator. Next, we used the high sensitivity protocol of myBaits v.5 (Daicel Arbor Biosciences) to enrich target pathogen loci from the host/pathogen control and museum archived samples. We used 2 rounds of enrichment for each pool of samples. Probe concentration was 100 ng/ μ L. Each round was 24 hours at 65°C. After washing of unbound DNA, each library was amplified with a 15-cycle PCR amplification step and quantified using qPCR. Finally, the pools of 4–16 were combined into an equimolar pool for sequencing. All sequencing reactions were on single lanes of Illumina Hi-Seq 2500.

Bioinformatic Analyses

All analyses were performed on a single compute node with 48 processors and limited to 100 Gb of RAM. Bioinformatic steps were documented in a series of BASH shell scripts or Jupyter notebooks. These files along with conda environments are available at github.com/nealplatt/pathogen_probes and archived. The basic structure of the bioinformatic analyses are shown in Figure 3. In general, we used the Kraken2 v2.1.2 (8) to assign a taxonomic id to each read, the Phyluce v1.7.1 (1,2) pipeline to identify, assemble, and align loci, and RaxML-NG v1.0.1 to generate phylogenies from each pathogen group of interest.

First, we used Trimmomatic v0.39 (9) to trim and quality filter low-quality bases and Illumina adaptors. Then, we used Kraken2 v2.1.1 (8) to compare each read from our samples to a reduced dataset of target loci using a `-conf` cutoff of 0.2. We decided to compare our reads to a reduced dataset of target loci to minimize the computational expense of these comparison. To generate the reduced database of bait-targeted loci, we downloaded one representative or reference genome from all species in RefSeq v212 (10) with `genome_updater.sh` v0.5.1 (https://github.com/pirovc/genome_updater). Then we used BMAP v38.96 (4) to map all the baits to each genome and kept the 10 best sites that mapped with $\geq 85\%$ sequence identity.

Next, we extracted these hits along with 1,000 bp up and downstream. These sequences were combined into a single fasta file that should contain the major mapping locations for our baits.

Once reads were classified we identified genera that were known pathogens or were present in at least one sample with more than 1,000 reads. Next, we extracted reads from the relevant family with KrakenTools v1.2 (<https://github.com/jenniferlu717/KrakenTools/>). These reads were then assembled (Figure 3, panel B) with the SPAdes genome assembler v3.14.1 (11) using the `phyluce_assembly_assemblo_spades` wrapper script. We filtered out low quality contigs based on size (<100 bp) and median coverage (<10×) as calculated by the SPAdes genome assembler. Next, we filtered individuals even further by removing individuals with fewer <2 contigs.

While we were assembling and filtering contigs from each isolated target loci from species with available genome assemblies, we used `genome_updater.sh` v0.5.1 (https://github.com/pirovc/genome_updater) to download one (-A 1) reference or representative (-c reference genome, representative genome) genome from either refseq or Genbank (-d refseq.genbank) for the pathogen group. We also included at least 1 individual from an outlier genus to root downstream analyses. These genomes were converted to twobit format with `faToTwoBit`. Next, we used `phyluce_probe_run_multiple_lastzs_sqlite` to compare probes from the pathogen group to the genome assemblies with an identity cut off of 85% (-identity 0.85). These loci plus 1 kb of flanking sequence (-flank 1000) were extracted from the genome using `phyluce_probe_slice_sequence_from_genomes`. After extraction, the sliced loci were identified and counted using `phyluce_assembly_match_contigs_to_probes` (-min-identity 90) and `phyluce_assembly_get_match_counts`. Next, we combined the loci generated from our samples with those from representative and reference genomes and aligned them with `phyluce_align_seqcap_align`. The resulting alignments were trimmed with `gblocks` v0.91b (12) and `phyluce_align_get_gblocks_trimmed_alignments_from_untrimmed`. We then counted the number of taxa per locus alignment (`phyluce_align_get_taxon_locus_counts_in_alignments`) and removed taxa with fewer than 2 loci (`phyluce_align_extract_taxa_from_alignments`). Then we removed any loci that contain fewer than half of the expected number of taxa with `phyluce_align_get_only_loci_with_min_taxa` and concatenated the remaining loci into a single phylip alignment (`phyluce_align_concatenate_alignments`).

We used RaxML-NG v1.0.1 (13) to generate a maximum-likelihood phylogenetic tree from the concatenated alignment. We ran 100 parsimony tree searches and then another 1,000 replicates using the GTR + G substitution model. Branches with less than 50% support were collapsed with the Newick Utilities v1.6 (14), Newick editor (nw_ed <input_tree_file>'i and b< = 50'). These steps were then repeated with other pathogen groups identified in the samples.

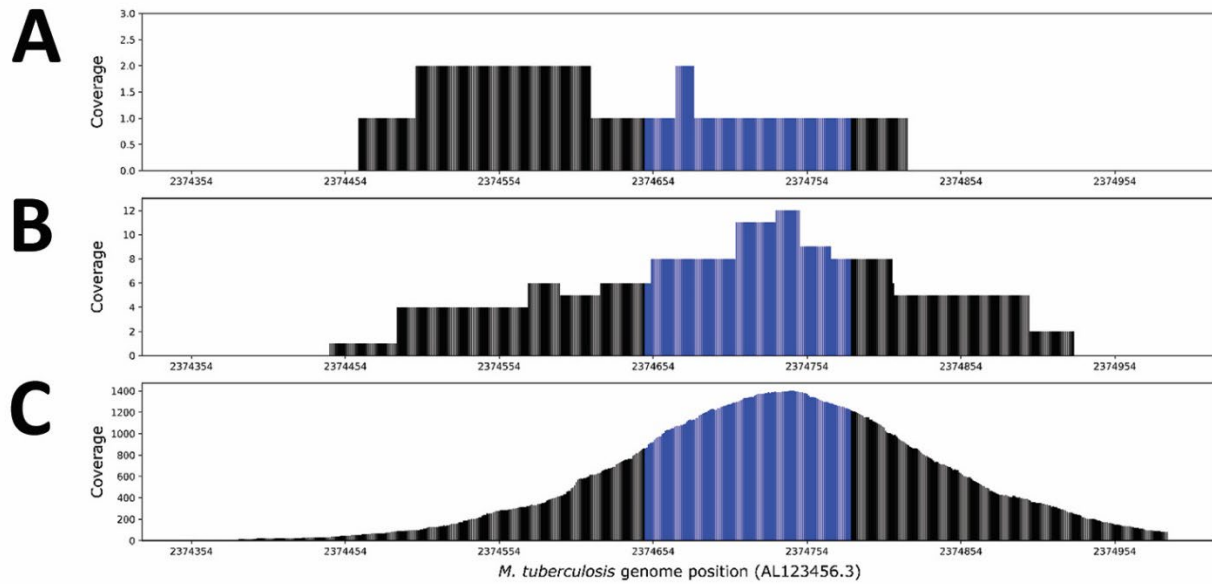
Host Identification

We verified museum identifications by comparing reads to a second Kraken2 v2.1.2 (8) database containing mammalian mitochondrial genomes. To do this, we downloaded all available mammalian mitochondrial genomes (n = 1,651) from <https://www.ncbi.nlm.nih.gov/genome/organelle/> (last accessed 3 November 2022). We then created a custom database and compared each of our samples using Kraken2 and no confidence cutoffs. The Kraken2 classifications were filtered by removing any samples with fewer than 50 classified reads and any single-read, generic classifications.

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Appendix Figure. Read depth at a targeted region in *Mycobacterium tuberculosis* in the A) 1%, unenriched, B) 0.001% enriched, and C) 1% enriched control, samples. This particular probe was designed for (AL123456.3:2,374,648–2,374,781; shown in blue). Median coverage at this locus increased from 1x in the 1% unenriched sample (A) to 8x in the 0.001% enriched sample (B) and 1,278x in the 1% enriched control sample (C).