Classification of *Bartonella* Strains Associated with Straw-Colored Fruit Bats (*Eidolon helvum*) across Africa Using a Multi-locus Sequence Typing Platform

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Abstract

*Bartonella*ae are facultative intracellular bacteria and are highly adapted to their mammalian host cell niches. Straw-colored fruit bats (*Eidolon helvum*) are commonly infected with several bartonella strains. To elucidate the genetic diversity of these bartonella strains, we analyzed 79 bartonella isolates from straw-colored fruit bats in seven countries across Africa (Cameroon, Annobon island of Equatorial Guinea, Ghana, Kenya, Nigeria, Tanzania, and Uganda) using a multi-locus sequencing typing (MLST) approach based on nucleotide sequences of eight loci (*ftsZ*, *gltA*, *nuoG*, *ribC*, *rpoB*, *ssrA*, ITS, and 16S rRNA). The analysis of each locus but *ribC* demonstrated clustering of the isolates into six genogroups (*E1*–*E5* and *Ew*), while *ribC* was absent in the isolates belonging to the genogroup *Ew*. In general, grouping of all isolates by each locus was mutually supportive; however, *nuoG*, *gltA*, and *rpoB* showed some incongruity with other loci in several strains, suggesting a possibility of recombination events, which were confirmed by network analyses and recombination/mutation rate ratio (*r/m*) estimations. The MLST scheme revealed 45 unique sequence types (ST1–45) among the analyzed bartonella isolates. Phylogenetic analysis of concatenated sequences supported the discrimination of six phylogenetic lineages (*E1*–*E5* and *Ew*) corresponding to separate and unique *Bartonella* species. One of the defined lineages, *Ew*, consisted of only two STs (ST1 and ST2), and comprised more than one-quarter of the analyzed isolates, while other lineages contained higher numbers of STs with a smaller number of isolates belonging to each lineage. The low number of allelic polymorphisms of isolates belonging to *Ew* suggests a more recent origin for this species. Our findings suggest that at least six *Bartonella* species are associated with straw-colored fruit bats, and that distinct STs can be found across the distribution of this bat species, including in populations of bats which are genetically distinct.
Author Summary

Bats, with over 1000 recognized species, represent about 20% of all classified mammalian species worldwide. These mammals have a wide range of ecologies and life-history traits, and are now widely recognized as important reservoirs of many pathogens. *Bartonella* species have been found distributed in a wide range of mammalian species, including bats. About half of recognized *Bartonella* species, including one bat-associated species, have been associated with human illness. Previous studies have shown that *Bartonella* species are extremely diverse, with or without evident specificity to their mammalian hosts. Possessing many unique aspects, bartonellae can serve as a useful biological marker to study how microorganisms have evolved and diversified along with their animal hosts in evolutionary history. In this study, we applied multi-locus sequence typing, or MLST, to study the genetic differences of straw-colored fruit bat (*Eidolon helvum*)-associated *Bartonella* species. Our studies suggest *Bartonella* species have both exchanged genetic materials among species through recombination events and lost genes that are perhaps superfluous to their life cycles, which includes an intracellular stage in mammals.

Introduction

Bartonellae are both Gram-negative alpha-proteobacteria and hemotropic bacteria highly adapted to facultative intracellular lifestyle in a wide variety of mammals, such as rodents, bats, insectivores, carnivores, ungulates, and other vertebrates. During the last two decades, progressively more bacterial species belonging to the genus *Bartonella* have been recognized with over 30 species described from different mammalian hosts. A number of *Bartonella* species were found to be associated with human illnesses and are associated with a growing spectrum of emerging diseases, including life-threatening endocarditis [1–9]. Animal reservoirs have been identified for some of the human pathogens, while remain unknown for others. Knowledge of the transmission of bartonella bacteria between mammalian hosts is incomplete. However, hematophagous arthropods such as fleas, flies, lice, mites, and ticks have been found naturally infected and are frequently implicated in transmitting *Bartonella* species [10–15].

Increasing recognition of bats as natural reservoirs of many emerging pathogens has drawn considerable attentions to study these mammals [16]. Multiple investigations of bartonella in bats have been conducted in different regions of the world [17–23]. These studies reported that bartonella infections are highly prevalent in many bat species and bartonella communities associated with bats are extremely diverse with co-circulation of numerous *Bartonella* species in the same bat populations. The data on relationships between *Bartonella* species and bats are quite contradictory depending on the investigated geographic region. Specifically, in Central and South America, multiple bat species may share the same *Bartonella* species without an evident host-specificity [19,20], while investigations from Asia and Africa demonstrated that a bat population typically harbors one or few *Bartonella* species specific for a particular bat species [17–18,21–22].

The straw-colored fruit bat (*Eidolon helvum*) is widely distributed in Africa. Colonies of these bats can be large and often found near human populations, with some roosts containing millions of individuals. Like other bats, straw-colored fruit bats have long life spans, with some individuals reaching at least 14 years of age [24]. Local human residents have a close contact with the bats, as they are frequently hunted for “bush meat”. A previous study in Kenya showed that straw-colored fruit bats are infected with bartonellae with high prevalence reaching 26% [18]. The same study demonstrated that *Bartonella* associated with the bats were genetically
distant and belonged to four distinct genogroups based on sequence variation in the citrate synthase gene (gltA) [18]. Similar observations were reported in a very recent study conducted in Nigeria [22]. Considering the broad distribution and other ecological characteristics of straw-colored fruit bats, it was expected that additional Bartonella genogroups associated with this bat species can be identified in Africa.

In the present study, we aim to expand our knowledge of bartonella infections in straw-colored fruit bats and to better understand how multiple Bartonella species can co-habit populations of one bat species. We compared genetic differences of bartonella isolates obtained from straw-colored fruit bats captured in seven countries across Africa using a multi-locus sequence typing (MLST) approach. Based on comparison of nucleotide sequences derived from multiple loci, MLST has been shown to provide high discriminatory power in epidemiological and genetic analysis of bacterial strain populations, while retaining signatures of longer-term evolutionary relationships or clonal stability [25–28]. In addition, sequencing of multiple loci can detect evidences of micro-evolutionary events, particularly homologous recombination, among identified sequence types [29,30]. The main objective of the current study was to analyze bartonella isolates obtained from naturally infected straw-colored fruit bats from different parts of Africa to determine whether well-defined phylogenetic lineages correspond to currently accepted criteria for discrimination of bacterial species [31]. This, in turn, can help to enhance our understanding of population structure in the bacteria and the relationships between allelic profiles and the animal host. To achieve this goal, we developed a MLST scheme that incorporates eight loci which have been previously proposed for characterization of Bartonella species. In this study, clusters defined by the phylogenetic analysis were called either a “genogroup” or a “lineage”. A genogroup is defined when a single locus sequence was applied for the phylogenetic analysis, while a lineage is defined from concatenated sequences of all loci.

**Materials and Methods**

**Bartonella isolates selection**

A total of 79 isolates (Table 1) originating from straw-colored fruit bats from seven countries in different regions across Africa were selected for the MLST analysis, including Annobón Island of Equatorial Guinea (n = 9), Cameroon (n = 3), Ghana (n = 20), Kenya (n = 16), Nigeria (n = 11), Tanzania (n = 15), and Uganda (n = 5). These isolates were obtained from previous studies completed by Bartonella Laboratory, CDC at Fort Collins, Colorado, and culturing procedures were published elsewhere [18,20].

**Multi-locus sequence typing (MLST)**

Eight loci (ftsZ, gltA, nuoG, ribC, rpoB, ssrA, ITS, and 16S rRNA) that have been previously used for bartonella description [27] were selected for MLST characterization of the bartonella isolates. Information on primers is provided in Table 2. A specific fragment of each locus was amplified by PCR for each of the 79 isolates. PCR products of each locus were purified with the QIAquick PCR Purification Kit (Qiagen, Germantown, MD) and sequenced in both directions using an Applied Biosystems Model 3130 Genetic Analyzer (Applied Biosystems, Foster City, CA). Using the Lasergene software package (DNASTAR, Madison, WI), obtained sequences were aligned by each locus and compared between the isolates and with other bat-originated Bartonella strains (S1 Table) and known Bartonella species (S2 Table). Based on the allelic profile (Table 1), each unique variant was designated as a sequence type (ST) and sequences for the eight loci were concatenated.
Table 1. Allelic profiles, sequence types (ST), and lineage identification for the 79 *Bartonella* isolates from different geographic locations.

| #  | Isolate  | Country/Region | ftsZ | gitA | nuoG | ribC | rpoB | ssrA | ITS 16S rRNA | Total length | ST  | Lineage |
|----|----------|----------------|------|------|------|------|------|------|-------------|--------------|-----|---------|
| 1  | B32133   | Nigeria        | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 2  | B37799   | Annobón       | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 3  | B37802   | Annobón       | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 4  | B37890   | Annobón       | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 5  | B37893   | Annobón       | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 6  | B37895   | Annobón       | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 7  | B37897   | Annobón       | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 8  | B39231   | Ghana         | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 9  | B39235   | Ghana         | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 10 | B39243   | Ghana         | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 11 | B39248   | Ghana         | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 12 | B39957   | Cameroon      | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 13 | B39958   | Cameroon      | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 14 | B40389   | Tanzania       | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 15 | B40404   | Tanzania       | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 16 | B40906   | Uganda        | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 17 | B40907   | Uganda        | 1    | 1    | 1    | neg  | 1    | 1    | 1            | 4622         | ST1 | Ew      |
| 18 | B32135   | Nigeria       | 1    | 1    | 2    | neg  | 1    | 1    | 1            | 4622         | ST2 | Ew      |
| 19 | B23978   | Kenya         | 1    | 1    | 2    | neg  | 1    | 1    | 1            | 4622         | ST2 | Ew      |
| 20 | B39238   | Ghana         | 1    | 1    | 2    | neg  | 1    | 1    | 1            | 4622         | ST2 | Ew      |
| 21 | B40386   | Tanzania       | 1    | 1    | 2    | neg  | 1    | 1    | 1            | 4622         | ST2 | Ew      |
| 22 | B23975   | Kenya         | 2    | 2    | 3    | 1*   | 2    | 4    | 2            | 5007         | ST3 | E5      |
| 23 | B39233   | Ghana         | 2    | 2    | 4    | 1*   | 2    | 5    | 4            | 5007         | ST4 | E5      |
| 24 | B39296   | Ghana         | 2    | 3    | 5    | 2*   | 3    | 3    | 5            | 5007         | ST5 | E5      |
| 25 | B40908   | Uganda        | 3    | 4    | 6    | 3*   | 4    | 2    | 5            | 5007         | ST6 | E5      |
| 26 | B39286   | Ghana         | 4    | 5    | 10   | 4*   | 10   | 3    | 2            | 5007         | ST7 | E5      |
| 27 | B39249   | Ghana         | 4    | 6    | 6    | 4*   | 4    | 2    | 2            | 5007         | ST8 | E5      |
| 28 | B40014   | Tanzania       | 5    | 7    | 7    | 4*   | 5    | 5    | 3            | 5007         | ST9 | E5      |
| 29 | B40406   | Tanzania       | 6    | 8    | 7    | 5*   | 6    | 4    | 2            | 5007         | ST10| E5      |
| 30 | B40408   | Tanzania       | 6    | 8    | 7    | 5*   | 6    | 4    | 2            | 5007         | ST10| E5      |
| 31 | B40391   | Tanzania       | 7    | 9    | 6    | 5*   | 4    | 6    | 2            | 5008         | ST11| E5      |
| 32 | B23976   | Kenya         | 8    | 10   | 8    | 6    | 7    | 7    | 6            | 5160         | ST12| E1      |
| 33 | B32112   | Nigeria       | 8    | 10   | 8    | 6    | 8    | 7    | 6            | 5160         | ST13| E1      |
| 34 | B23983   | Kenya         | 8    | 10   | 8    | 6    | 8    | 8    | 6            | 5160         | ST14| E1      |
| 35 | B24219   | Kenya         | 8    | 10   | 8    | 6    | 8    | 8    | 6            | 5160         | ST14| E1      |
| 36 | B39239   | Ghana         | 8    | 10   | 9    | 6    | 7    | 8    | 6            | 5160         | ST15| E1      |
| 37 | B23987   | Kenya         | 8    | 10   | 9    | 6    | 8    | 7    | 6            | 5160         | ST16| E1      |
| 38 | B24003   | Kenya         | 8    | 10   | 9    | 6    | 8    | 7    | 6            | 5160         | ST16| E1      |
| 39 | B23977   | Kenya         | 8    | 11   | 8    | 6    | 7    | 7    | 6            | 5160         | ST17| E1      |
| 40 | B32314   | Nigeria       | 8    | 11   | 8    | 6    | 7    | 8    | 6            | 5160         | ST18| E1      |
| 41 | B23973   | Kenya         | 8    | 12   | 8    | 6    | 7    | 7    | 6            | 5160         | ST19| E1      |
| 42 | B23982   | Kenya         | 8    | 12   | 8    | 6    | 7    | 7    | 6            | 5160         | ST19| E1      |
| 43 | B40402   | Tanzania       | 8    | 13   | 8    | 6    | 7    | 8    | 6            | 5160         | ST20| E1      |
| 44 | B40006   | Tanzania       | 8    | 14   | 8    | 6    | 7    | 7    | 6            | 5160         | ST21| E1      |
| 45 | B23979   | Kenya         | 9    | 15   | 11   | 7    | 9    | 9    | 7            | 5159         | ST22| E2      |
| 46 | B39302   | Ghana         | 9    | 15   | 11   | 7    | 9    | 9    | 7            | 5159         | ST22| E2      |

(Continued)
Analysis of nucleotide polymorphism and diversity

Sequence polymorphisms of the eight MLST loci were examined in MEGA v6.06 separately for each Bartonella lineage and among all 45 STs. For protein-coding loci (ftsZ, gltA, nuoG, ribC, and rpoB), the appropriate open reading frame that contained no stop codons was selected for each locus by checking all possible starting positions. To test for evidence of selection in these loci, $d_N/d_S$ ratios were calculated using the Nei-Gojobori method for each of the loci considering each lineage separately and for all 45 STs together. Nucleotide diversity ($\pi$), determined by the number of nucleotide differences per site, was calculated as p-distance for all eight loci individually and for concatenated sequences considering each lineage separately and for all 45 STs
together. For concatenated sequences, the total length included gap regions and missing genes, thus the pairwise deletion option was used.

Phylogenetic analysis

A neighbor-joining tree based on the concatenated MLST alleles alone was constructed using the Clustal W program within the Lasergene 11 package of DNASTAR (version 8). An additional phylogeny was constructed, in which known *Bartonella* species and *Bartonella* strains isolated from bats in Old World were included in addition to representative strains from straw-colored fruit bats (E1—E5, Ew), and *Brucella abortus* was included as an outgroup. This phylogeny was inferred using sequences of *ftsZ*, *gltA*, *nuoG*, *ribC*, *rpoB*, *ssrA*, and 16S rRNA; ITS sequences were not included due to the large number of gaps among the strains that could not be resolved. Sequences from each locus were aligned using Clustal X v2.1, trimmed to equal lengths, and concatenated. The best model of nucleotide substitution was determined using MEGA. Based on this model, a maximum-likelihood tree was generated in MEGA with 1000 bootstrap replicates. Due to missing genes among the strains as well as some alignment gaps, we used the pairwise deletion option when inferring the tree.

Recombination tests

To visualize potential recombination events among the sequence types, a phylogenetic network was inferred from concatenated sequences from the 45 STs using the Neighbor-Net algorithm in SplitsTree v4.13.1 with 1000 bootstrap replicates. Gaps in aligned sequences and from missing genes were included. The pairwise homoplasy index (PHI) was implemented in SplitsTree to test for significant recombination among the isolates.

ClonalFrame v1.1 was used to estimate the relative contribution of recombination and mutation in generating polymorphisms among the 45 *Bartonella* STs. Based on recommendations by Vos and Didelot [32], ITS and genes coding for RNA (*ssrA* and 16S rRNA) were not included because of potential confounding effects of selection on the detection of homologous recombination rates. Therefore, we used only the five protein-coding loci (*ftsZ*, *gltA*, *nuoG*, *ribC*, and *rpoB*) in the study. Two independent runs were performed in ClonalFrame using 200,000 MCMC iterations. The initial mutation rate (θ) was set using Watterson’s moment estimator while the remaining initial parameters used the default settings in ClonalFrame. We assessed convergence and mixing properties of the dataset through visual inspection of the traces for the likelihood and model parameters.
Nucleotide sequence accession numbers
Newly-identified alleles from the current study were submitted to GenBank with the following accession numbers: KJ999677 to KJ999694 (ftsZ), KM030503 to KM030526 (gltA), KM030527 to KM030544 (nuoG), KM215178 to KM215188 (ribC), KM215189 to KM215208 (rpoB), KM233456 to KM233470 (ssrA), KM233471 to KM233471 (ITS), and KJ944271 to KJ944279 (16S rRNA).

Results
Individual sequence analysis of gltA and other loci
All 79 isolates were first compared based on the 751bp gltA fragment for the initial identification. The gltA sequence alignment revealed 247 variable sites with 24 variants, delineating six unique genogroups with divergence of 7.3–23% among the genogroups and with similarities of ≥ 96.8% within a genogroup. Four genogroups were the same as previously identified in straw-colored fruit bats from Kenya, which were named E1, E2, E3, and Ew [18]; whereas the remaining two genogroups were distinct from these four and from any other previously reported Bartonella genotypes. Continuing the same naming system proposed for discrimination of Bartonella strains discovered in Kenya [18], these two new genogroups were designated as E4 and E5.

Following the initial identification, additional analyses were performed with the other seven loci. Each analyzed locus, except ribC, showed that all bartonella isolates obtained from straw-colored fruit bats also fell into the same six genogroups identified by gltA. Unexpectedly, analysis of the ribC locus revealed a large variation in length of the examined fragment among the isolates. Depending on the group identified by other markers, the examined ribC fragment was either fully presented (535bp), or reduced (382bp), or non-amplifiable (PCR negative). The phylogenetic analysis based on ribC sequences of the 79 isolates revealed five genogroups related to E1—E5 clusters that correspond to the grouping by other loci. Absence of the ribC locus was indicative for the genogroup Ew. All isolates with a reduced ribC fragment (partially missing) belonged to the genogroup E5 (Table 1).

Allelic profiles, sequence types (ST), and phylogenetic analysis
The size of sequenced fragments ranged between 282bp and 1172bp at different loci. The investigated loci showed different degrees of variation, with 38–249 variable sites and 9–24 alleles (Table 2). The length of concatenated sequences ranged from 4,622bp to 5,160bp as a result of the variation in the fragment length of ribC (from 0 to 535bp), ITS (from 315 to 352bp), and ssrA (from 282 to 289bp). Based on the allelic profiles, the MLST analysis distinguished 45 sequence types (ST) among the 79 isolates, showing high heterogeneity. Most STs were represented by a single isolate, while some were represented by 2–4 isolates, and ST1 was found in 17 isolates (Table 1). Phylogenetic analysis demonstrated that all of the 45 STs resolved into six lineages, namely, E1, E2, E3, E4, E5, and Ew, to match the names proposed for grouping the strains based on sequences of one locus (Fig. 1). The divergence was 5.1–14% among lineages and ≤ 4.2% within a lineage.

Lineage Ew contained 21 isolates of two similar sequence types (ST1 and ST2) with only one nucleotide difference between the two STs. The 17 isolates presenting ST1 were discovered in each of the seven studied countries. ST2 was identified in four isolates from Ghana, Kenya, Nigeria, and Tanzania (one isolate per country) (Table 1). None of the isolates belonging to this lineage possessed the fragment for ribC. The lineage Ew was very common among the six lineages identified and accounted for 26.6% (21/79) of all analyzed isolates.
Lineage E1 contained 13 isolates of 10 sequence types (ST12—ST21). The distance among the STs was less than 2.1%. Eight isolates were from Kenya, and the others were from Ghana, Nigeria, and Tanzania (Table 1).

Lineage E2 included seven isolates of six sequence types (ST22—ST27). The distance among the STs was less than 2.1%. Isolates belonging to this lineage were from Kenya, Ghana, Tanzania, and Uganda (Table 1).

Lineage E3 contained 24 isolates of 15 sequence types (ST28—ST42). The distance among the STs was less than 4.2%. ST30, ST36, ST37, and ST40 were represented in four, three, two, and three isolates, respectively. The remaining STs were each represented by a single strain. Lineage E3 was the most common among all bartonella lineages detected in straw-colored fruit bats, and accounted for 30.4% (26/79) of all isolates analyzed. Notably, mismatches in assignment of isolates to specific ST lineages were observed for a few loci on several occasions. Specifically, isolate B32119 (ST38) from a Kenyan bat and isolate B23797 (ST39) from a Nigerian bat were assigned to lineage E3 by ST classification, but separate analyses of the gltA and nuoG sequences of the two isolates indicated their closeness to genogroup Ew. Similarly, isolate B40005 (ST28) from a Cameroonian bat was identified as E3 by ST classification, but was closer to genogroup E1 by the gltA and nuoG sequences.

Lineage E4 contained four isolates of three sequence type (ST43—ST45). These STs are very similar among themselves with distance < 0.1%. The isolates were from Ghana, Tanzania, and Uganda (Table 1).

Lineage E5 contained 10 isolates of eight sequence types (ST3—ST11). Compared to other lineages, the STs within this lineage were more distant (0.2–3.5%). All isolates/STs in this lineage presented shorter concatenated sequences due to the partial fragment missing in ribC. One isolate (B39286) from this lineage also showed a mismatch in lineage assignment when analyzed by individual loci. The isolate was from a Ghanaian bat and belonged to lineage E5 by ST classification, but assigned to genogroup E2 by the rpoB and nuoG sequences.

Based on MEGA results, the best nucleotide substitution model for the concatenated sequences of gltA, ftsZ, nuoG, ribC, rpoB, ssrA, ITS, and 16S rRNA was determined to be GTR+G+I [33]. Comparison with known Bartonella species and other bat-associated Bartonella strains demonstrated that lineages E1, E2, E3, and E5 belong to a clade that includes other Bartonella strains found in Old World bats, while E4 and Ew appear to have arisen independently (Fig. 2).

Patterns of selection and diversity in nucleotide sequences

The $d_{st}/d_{s}$ ratios calculated for protein-coding sequences ranged from 0.022 for ftsZ to 0.242 for ribC when comparing all 45 STs. The values for each locus differed when each Bartonella lineage was analyzed separately. Nucleotide diversity among STs varied by locus, ranging from 1.02% in 16S rRNA to 14.38% in ribC. Among the Bartonella lineages, E2 and E5 had the highest nucleotide diversity when all loci were considered together, with 1.56% and 1.24%, respectively, while Ew had the lowest (0.02%).

Recombination test

A network phylogeny based on the concatenated sequences for the 45 STs in SplitsTree (v4.13.1) shows that the 45 STs fall into the same six identified lineages identified by single loci.
However, isolates B32119 from Nigeria, B23797 from Kenya, B40005 from Cameroon, and B39286 from Ghana are split from the network, reflecting their mixed ancestry (Fig. 3). The pairwise homoplasy index (PHI) \cite{34} found significant evidence of recombination in the network (mean = 0.2, variance = 4.5x10^{-6}, p < 0.01). Using ClonalFrame, the r/m value (the ratio of probabilities that a site is altered by recombination or mutation) was estimated to be 1.14 (95% CI 0.63, 1.82) and 1.06 (95% CI 0.53, 1.67) based on two independent runs (average = 1.1), while the ρ/θ value (the ratio of rates at which recombination and mutation occur) was estimated to be 0.049 (95% CI 0.023, 0.087) and 0.037 (95% CI 0.016, 0.065).

**Discussion**

Straw-colored fruit bats have been reported to be commonly infected with multiple *Bartonella* strains based on *gltA* sequences \cite{18,22}. In this study, we developed a MLST scheme using...
eight loci which have been commonly used for Bartonella species description to characterize 79 bartonella isolates obtained from straw-colored fruit bats that were captured in different regions across Africa to better elucidate the genetic diversity among these isolates.

The analyses of selection pattern showed none of the \( d_s/d_\beta \) ratios exceeded 1. This suggested that all of the protein-coding loci in the study are undergoing purifying selection. The analyzed bartonella isolates exhibited a high level of heterogeneity with 9–24 alleles identified by different loci. The MLST scheme resolved 45 STs among the 79 isolates and these isolates/STs clustered into six distinct lineages (\( E1–E5 \), and \( Ew \)) with genetic distances of 5.1–14% among the lineages. MLST has been regarded as a highly discriminatory tool by a number of previous studies. According to the widely accepted criteria for a new bacterial species based on nucleotide divergence greater than 5% in housekeeping genes [31], the MLST analyses in the present study demonstrated that each lineage identified in straw-colored fruit bats in this study likely represents a new Bartonella species. These Bartonella species have never been found in other bat species, suggesting their specific co-adaptation to Eidolon bats, though future studies are necessary to confirm this. The performed analyses allow us to conclude that there are at least six Bartonella species associated with straw-colored fruit bats. Phylogenetic analysis based on multiple loci demonstrates that four of these lineages, \( E1, E2, E3, \) and \( E5 \), belong to a phylogenetic lineage that includes other Bartonella strains isolated from Old World bats [18,21–23], while \( E4 \) and \( Ew \) appear to have evolved independently.

Lineages \( E3 \) and \( Ew \), containing 24 and 21 isolates, respectively, and accounting for 57% of all isolates, are likely to be the more common Bartonella species circulating among this bat species. However, because the original bat samples from which the analyzed bartonella isolates obtained were collected in different settings and during different seasons by multiple investigators, there could be some bias in temporal or seasonal variations. Furthermore, the culturing procedure that was necessary for performing such bacterial classification may also contribute to misrepresentation of comparative prevalence of Bartonella species as some of them could hardly be cultured [35]. Due to the limited number of isolates selected from each part of Africa, we did not attempt to investigate any patterns in the geographic distribution of Bartonella species in this study. The presented classification, however, can be used for future studies aimed at describing geographic patterns of bartonella distribution among Eidolon bats.

Most lineages consisted of multiple STs (from 4 to 14). Nevertheless, lineage \( Ew \) consisted of only two STs with a large number of isolates, and in fact there is only one nucleotide difference between the two STs. This indicates a lower diversity of allelic polymorphisms in the isolates from this lineage and allows a speculation that \( Ew \) has diverged more recently compared to the other Bartonella lineages. Further, our observations of partially missing or non-amplifiable sequences of ribC suggests gene loss, a common evolutionary process in bacteria [36], possibly has occurred and resulted in the appearance of two new Bartonella species (\( E5 \) and \( Ew \)). As a housekeeping gene, ribC participates in the regulation of riboflavin biosynthesis. The non-amplifiability of the gene might not be true but suggest the gene is in some undetected form. If this gene is truly missing, its loss may be compensated by the fact that bartonellae are intracellular parasites which can perhaps utilize host riboflavin or they have lost the need of it. If this were true, fitness costs may lead to selection against this superfluous genetic material [37]. Overall, little is known about the mechanisms of regulation of bacterial riboflavin genes [38]. Although we cannot explain biological consequences of the loss of ribC, the ‘presence/absence’ characteristics make ribC a marker for the differentiation and classification of Eidolon-associated Bartonella strains. In addition, the dramatic variation in ribC among the straw-colored fruit bat-associated Bartonella species presents an interesting model for understanding evolutionary trends in developing host-bacterial parasite relations [36].
Another intriguing observation in this study was the set of mismatches between MLST lineages and some genogroups comparing phylograms built based either on entire concatenated sequences or individual locus sequences for four isolates. Particularly, three isolates belonging to E3 based on the ST classification were either Ew (two isolates) or E1 (one isolate) by nuoG or gltA alone and one isolate of E5 by ST classification would be E2 by nuoG or rpoB alone. The loci rpoB and gltA have been shown to be the most potent for species discrimination in other studies [31]. According to our observations in the present study, an analysis limited to two loci may still lead to a misclassification. Thus characterization of multiple loci should be preferably used for Bartonella species identification. The mismatches between lineages and genogroups identified by different loci indicate that recombination and/or lateral gene transfer events between different Bartonella species are ongoing processes.

Our recombination test results indicate that while recombination happens infrequently, an individual recombination event introduces a large number of polymorphisms. Recombination in rpoB, gltA, and other housekeeping loci has been observed in other Bartonella species [29,30,39]. Estimated r/m values for B. grahamii alone (1.7 [38] and 6.81 [30]), B. taylorii alone (3.77 [27]), or B. grahamii and B. taylorii considered together (4.06 [30]) compare favorably with our estimate for recombination among six species of Bartonella in E. helvum (r/m = 1.1). These estimates contrast with results from B. henselae (r/m = 0.1) [40] and the perception that intracellular bacteria have low rates of recombination and horizontal gene transfer [32]. Intermediate to high rates of recombination in these studies suggest that multiple Bartonella species may coexist at some point in the infection cycle, potentially in their mammalian hosts and/or arthropod vectors, thus facilitating gene exchange. Moreover, gene exchange appears to play an important role in generating sequence diversity among co-circulating Bartonella species. Further studies that could measure co-infection of Bartonella species and resulting rates of recombination would help to explain these interesting dynamics.

Our study provides information on the genetic diversity of straw-colored fruit bat-associated Bartonella species that can be used for the discrimination of the lineages at the level corresponding to separate species. Additional information is needed to understand the interaction between these Bartonella species and their bat hosts under natural conditions, as well as their putative ectoparasite vectors [41]. A biological approach to define Bartonella species has been discussed [42] with emphasis on host-association as a phenomenon that promotes biological isolation of Bartonella species. However, such an approach has limitations for situations when several Bartonella species are associated with one animal host. A similar situation was also described among Bartonella strains in grasshopper mice [43] and cotton rats [44]. Nevertheless, among all mammalian species described as a host of one or multiple Bartonella species, the number of presumptive species associated with E. helvum bats is clearly the highest. Although there is no evidence suggesting that Bartonella species associated with African fruit bats may cause illnesses in humans or in bats themselves, a recent study reported the presence of Bartonella mayotimonensis, a reported etiologic agent of endocarditis in humans, in the Dauben-ton’s bat (Myotis daubentonii) and the Northern bat (Eptesicus nilssonii) in Europe [23], suggesting a potential role of bats as reservoirs for human bacterial pathogens. It is important to understand the role of straw-colored fruit bats in biotic communities and their importance as reservoir hosts contributing to the maintenance and transmission of bartonellae to other animals and humans.

The great diversity of Bartonella species associated with E. helvum, however, clearly presents interesting questions from the perspective of microbial evolution. What are the mechanisms that generate and maintain such diversity? How can these bacteria share the same ecological niche? Analyzing such relationships, Chan and Kosoy [45] hypothesized that co-circulation of independent Bartonella species in populations of one host species may represent an escape
mechanism to circumvent the host immune responses. We ultimately believe this system will be very informative for understanding the evolution of bacteria in a host-vector-pathogen system.

Supporting Information

S1 Table. GenBank accession numbers for ftsZ, gltA, nuoG, ribC, rpoB, ssrA, ITS, and 16S rRNA sequences for bat-associated Bartonella strains.

S2 Table. GenBank accession numbers for ftsZ, gltA, nuoG, ribC, rpoB, ssrA, ITS, and 16S rRNA sequences of reference Bartonella species.

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Author Contributions

Conceived and designed the experiments: YB MYK DTSH. Performed the experiments: YB. Analyzed the data: YB CDM. Contributed reagents/materials/analysis tools: MYK. Wrote the paper: YB CDM MYK DTSH.

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