SHORT REPORT

Two novel adenoviruses found in Cave Myotis bats (*Myotis velifer*) in Oklahoma

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Abstract

Bats are carriers of potentially zoonotic viruses, therefore it is crucial to identify viruses currently found in bats to better understand how they are maintained in bat populations and evaluate risks for transmission to other species. Adenoviruses have been previously detected in bats throughout the world, but sampling is still limited. In this study, 30 pooled-guano samples were collected from a cave roost of *Myotis velifer* in Oklahoma. A portion of the DNA polymerase gene from Adenoviridae was amplified successfully in 18 *M. velifer* samples; however, DNA sequence was obtained from only 6 of these *M. velifer* samples. One was collected in October 2016, one in March 2017, and 4 in July 2017. The October and March samples contained viral DNA that was 3.1% different from each other but 33% different than the novel viral sequence found in the July 2017 samples. Phylogenetic analysis of these fragments confirmed our isolates were from the genus *Mastadenovirus* and had genetic diversity ranging from 20 to 50% when compared to other bat adenoviruses.

Keywords

Adenovirus · Bat · Myotis · Mastadenovirus

Bats make up 20% of all mammals, and they are the second richest mammalian order in respect to number of species [1, 2]. In recent years, bats have emerged as a rich source of novel viruses [3, 4]. They have been found to host more zoonotic viruses per species than rodents [5], and even documented to harbor viruses from two different viral families simultaneously [6]. Viruses in bats can switch hosts to other bat species [4] and they are known to carry pathogenic viruses that can infect humans such as rabies, lyssaviruses, nipah and hendra viruses, ebola, and SARS coronavirus [7, 8]. However, in most cases bats serve as reservoirs for viruses with immunological tolerance and without transmission to other humans [7, 9]. Consequently, it is important to first identify viruses housed in bats in order to better understand the ecology of bat-borne viruses, how they are maintained in bat populations, and then evaluate risks for host transmission to other species.

Adenoviruses (AdVs) are double stranded DNA viruses found in vertebrate hosts of many different species [8, 10]. The family Adenoviridae consists of five genera [11] with members in the genus *Mastadenovirus* infecting mammals [12]. AdVs are widespread in the human population and cause a variety of usually minor symptoms, such as respiratory illnesses, conjunctivitis, and gastroenteritis [8]. Generally, these viruses are host-specific [13] and thought to have low zoonotic risk [14]; however, Chen et al. [15] discovered a novel adenovirus (TMAdV) with the ability to infect both monkeys and humans. Since bats are known reservoirs of numerous viruses and cross-species transmission has been documented for an AdV, it will be useful to know which AdVs bats carry. AdV strains have been found in more than 45 species of bats across their global distribution [6, 8, 12, 16–34] with seven species proposed by the International Committee on Taxonomy of Viruses [35]. These studies represent a start at investigating bat AdVs, but there is a need for additional studies considering there are over 1300 species of bats [2] and few North American bats have been investigated.

In this study, *Myotis velifer* guano samples were tested for the presence of AdVs. We expected to find AdVs in *M. velifer* because that genus had the most AdVs in a study on 19 bat species in China [8]. Guano was collected from *M.
velifer individuals in Washita Bat Cave (Washita Co, OK). Samples were either collected from a plastic tarp left laying overnight at the entrance of the cave (in March or July) or after bats were captured and placed in a sterile cup for 1 h (in October). Bats were handled following guidelines from Sikes et al. [36], and white nose syndrome decontamination protocols were followed [37]. Regardless of method of collection, four guano pellets were stored collectively in 500 µl of RNA Later®, and stored at −20 °C. We obtained a total of 120 guano pellets and this provided 30 pooled samples for analysis.

DNA extraction was carried out with QIAamp® DNA Mini Kit (Qiagen) following the manufacturer’s protocol with minor modifications. Nested PCR of the partial Adenoviridae DNA polymerase gene was carried out on each sample following Li et al. [8] using primers pol-F (5′ CAGCCCK-CKTTRTGYAGGGT 3′) and pol-R (5′ GCHACCATY AGCTCCAACTC 3′). Cycling profile consisted of 94 °C for 5 min, 30 cycles of 94 °C for 30 s, 60 °C for 30 s, 72 °C for 30 s, and then a final extension of 72 °C for 5 min. The second round of amplification used 1 µl of first round PCR product as template, primers pol-nf (5′ GGGCTCRRTGT CCAGCA 3′) and pol-nr (5′ TAYGACATCTGYYGGCATGTA 3′) and the same cycling steps. Positive (Human Adenovirus D DNA from American Type Culture Collection) and negative controls were used for each PCR. Positive PCR products were purified with Wizard® SV Gel and PCR Clean-up System (Promega). Species verification of bats with positive AdV samples was performed using nested PCR with primers SFF_145f (5′ GTHACHGCRYAGCHHTYGTAAAT 3′) and SFF_351r (CTCCWGCRTGDGCWAGRTTCCC 3′) from [38] and thermocycler steps consisting of 95 °C for 5 min, 38 cycles of 95 °C for 60 s, 60 °C for 30 s, 72 °C for 30 s, and final extension of 72 °C for 10 min to amplify a region of the cytochrome c oxidase gene that is highly diagnostic among bats.

Sanger sequencing of positive samples for bat and AdV identification was performed by Oklahoma Medical Research Foundation, and fragments were aligned and manually edited in Geneious v. 10.1.3 [39]. AdV sequences (131) isolated from other bat species, turkey, canine, bovine, and human AdVs A, B, C, and D on GenBank were added to the final alignment. Model of sequence evolution, maximum likelihood analysis, and uncorrected P nucleotide distance were performed in Mega v. 7.0.26 using all sites, including gaps, and 1000 bootstrap replicates [40].

There was at least one positive sample for each collection date, but only 6 of 18 positive samples had viral DNA quantities necessary for successful sequencing. The alignment of our DNA sequences with only the recognized viral species followed a Hasegawa–Kishino–Yano model of evolution with a gamma distribution of 0.4847. The maximum likelihood analysis indicated that the AdVs were of the Mastadenovirus genus and our proposed AdVs form separate clusters in distinct clades (Fig. 1). When all sequences from GenBank were included in the alignment the model of nucleotide evolution was the General Time Reversible model with a gamma distribution of 0.6376 and invariant sites. In this analysis (not shown), the AdV sequences did not form clusters according to their host family. This suggests transmission between host species is more common than coevolution with the host.

Myotis velifer samples from October 2016 (Guano 61) and March 2017 (Guano 2) were only 3.1% different from each other, while they were ~33% different from the 4 sequences extracted from July 2017 samples (Guano 21, 22, 24, 25). Guano 21 was identical to Guano 22 and 24, and it was only 1 nucleotide different than Guano 25. DNA sequences from Guano 2, 61, and 21 have been deposited in GenBank (Accession MN240005–MN240007). We do recognize the sequenced fragment is short (241 basepairs) and only provides preliminary viral classification. AdVs species are designated if amino acid sequence is >5% for the DNA polymerase gene. Based on this criterion, we suggest Guano 61 and Guano 2 are different strains of the same AdV species and are further referred to as Cave Myotis AdV1-1 and Cave Myotis AdV1-2. DNA sequence from Guano sample 21, 22, 24, and 25 represent a separate Adenovirus species and are further referred to as Cave Myotis AdV2. There were 2 non-synonymous mutations and 2 synonymous mutations between Cave Myotis AdV1-1 and AdV1-2. Cave Myotis AdV1-1 and AdV1-2 are most similar to GU226951 isolated from Myotis horsfieldii [8] with genetic differences of 21.7% and 23.3%, respectively, and most different from HQ529709 isolated from Rousettus leschenaultii [21] with genetic differences of 48.45 and 49.2%, respectively. Cave Myotis AdV2 is most similar to MF404977, 80–82, 84–87 isolated from Pipistrellus pygmaeus [32] with a genetic difference of 20.2%. Cave Myotis AdV2 is most different from KC692424, 28 isolated from Pteropus giganteus [24] and HQ529709 isolated from R. leschenaultii [21] by 44.5%. These two new AdVs are ≥27% different than any currently recognized bat mastadenovirus A–G (Table 1). When AdV sequences were compared to other AdVs from bats, genetic diversity ranged from 20 to 50%. This study demonstrates that there is great genetic diversity of DNA viruses within the same species of bats found in the same location, which is relatively uncommon for other vertebrate viruses [8].

We sampled caves during 3 seasons and found greater prevalence of viral DNA in M. velifer guano during summer (July; 14/16 samples = 87%) than spring (March; 1/4 samples = 25%) or autumn (October; 3/10 samples = 30%). This is the highest percentage of positive AdV samples detected in bats to date from a single sampling period. Drexler et al. [20] collected guano samples from M. myotis in Germany during May, June, and July for 3 years, and the highest percentage
of positive samples for 1 sample date (67.5%, 27/40 samples) was collected in May and the same frequency in July. It is likely our high percentage of positive samples from one sample date is because *M. velifer* give birth to their young in May–June [41]. In summer there are many young bats present with weaker immune systems and a greater risk of lactating females sharing viruses with their young. Drexler et al. [20] found a significant increase in prevalence of coronaviruses one month after parturition during summer months but AdV detection was not significantly higher in any particular month within the summer.

Little work has been done to investigate AdVs in North American species of bats; however, these studies [19, 29] and ours highlight the importance of identifying viruses housed in bats to better understand viral evolution, how viruses are maintained in bat colonies and evaluate risks for host transmission to other species. Li et al. [8] found their novel bat AdV (BtAdV-TJM) was capable of infecting several mammalian cells from different species, including humans, which indicates that bat AdVs possibly have a wide host range. They also suggest some bat AdVs have similar amino acid sequences for structural proteins to those in human AdVs and a high GC content, which suggest bat AdVs might be an ideal vector for gene therapy and vaccine delivery in humans [8]. Future studies should include sequencing the entire viral genomes and isolating the viruses

**Table 1** Percent nucleotide difference between strains of bat mastadenovirus isolated from *Myotis velifer* and recognized bat mastadenovirus species based on the 241-bp fragment of DNA polymerase gene

| Strain of bat mastadenovirus | Bat species                  | GenBank ID       | Cave Myotis AdV1-1 | Cave Myotis AdV1-2 | Cave Myotis AdV2 |
|-----------------------------|------------------------------|------------------|--------------------|--------------------|------------------|
| A                           | *Myotis ricketti*            | GU226970 [8]     | 32.6               | 33.3               | 27.1             |
| B                           | *Pipistrellus pipistrellus*  | JN252129 [17]    | 34.1               | 35.7               | 28.9             |
| C                           | *Rhinolophus sinicus*        | KT69853 [28]     | 35.7               | 35.7               | 28.6             |
| D                           | *Miniopterus schreibersii*   | KT698856 [12]    | 40.2               | 41.7               | 40.2             |
| E                           | *Miniopterus schreibersii*   | KT698852 [12]    | 43.3               | 43.3               | 44.1             |
| F                           | *Rousettus leschenaultia*    | KX961095 [12]    | 36.6               | 36.6               | 36.6             |
| G                           | *Corynorhinus rafinesquii*   | KX871230 [29]    | 32.6               | 32.6               | 28.7             |

Genetic difference values estimated in Mega7

**Fig. 1** Phylogenetic relationships of novel bat viruses and described bat mastadenovirus sequences available on GenBank. Maximum likelihood phylogeny was generated in MEGA7 using a 241-bp fragment of the DNA polymerase gene. Tree was inferred using the Hasegawa-Kishino-Yano G+I model and 1000 bootstrap replicates. Values > 50 indicated for nodal support. Sequences obtained in this study named with the proposed AdV strain.
to test possible transfections in other species to better characterize the viruses discovered here.

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Author contributions DNL and MA have contributed to data collection, analysis, and writing the manuscript.

Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

Ethical approval All applicable national guidelines for the care and use of animals were followed.

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