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INTRODUCTION:

- Bats are a very diverse group of mammals known to host many pathogens of increasing domestic animal and human health concern.
- It is crucial to identify factors that may explain this variation in pathogen diversity.
- Known traits associated with viral diversity include:
- IUCN status
- population structure
- longer lifespan
- larger group size
- geographic distribution in the eastern hemisphere
- Bat roosting ecology, especially the ability to roost in anthropogenic structures, is often not included in past models owing to a sparsity of standardized data. Omitting this information could impact the prediction of spillover risk.

OBJECTIVE:

Using a novel roosting ecology dataset, this study is a species-level analysis that uses a machine learning approach to investigate if anthropogenic roosting:

- 1. is important in predicting viral richness
- 2. improves identification of undetected but likely bat reservoir species

METHODS: **Classifying bat roosting status**

Anthropogenic roosting bat species are defined as having record of roosting in any human-made structure (houses, bridges, attics, mines, etc.).



• Categories

 \circ This explanatory variable of interest is a binary variable of 0 and 1 (0 = natural roost, 1 = roosts in anthropogenic structures)

Sources

- IUCN
- Walker's Bats of the World 1994 edition
- Google Scholar
- Animal Diversity Web

• Incomplete or missing roost information

- Will be considered as NA if:
 - IUCN states natural history or roosting status unknown/data deficient Cannot find evidence of known roosting sites within the first 10 pages of GS or having no relevant search results

Dataset Building:



Resulting in a dataset with 1287 bat species and 56 explanatory variables

Analysis:

- For preliminary analysis, both overall virus and zoonotic virus response variables were turned into binary variables (0 for non-host and 1 for host).
- Boosted regression tree (BRT) models were run for both response variables, using a Bernoulli distribution for 50 partitions each of a 70:30 training/test split.
- Resulting variable importance and test AUC were averaged to view mean variable importance and standard error across models.

Anthropogenic Roosting and Virus Richness in Bats

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Scan to view/download data sources and additional info



Scan to view Briana's Website or go to babetke.github.io





PRELIMINARY RESULTS:

- identified and 470 non-NA values.
- anthropogenic roosting structures.
- overhangs of buildings.
- Vespertillionidae at 42%.



- human-made structures.
- are distinct across bat families.
- natural-roosting counterparts.



DISCUSSION:

This work emphasizes the need to characterize and synthesize species traits that relate directly to the wildlife-human interface, such as the use of anthropogenic structures as roosts, where spillover occurs. Further study consists of:

- roosting status to those that do.



• Current coverage of the roosting status variable is 40% with 544 bats

• Of the 470 bat species with a value (either 0 or 1), 57.7% were found to roost exclusively in natural structures and 42.3% were found to roost in

• These anthropogenic structures included bridges, attics, mines, and

• Preliminary roosting status data represent 11 bat families and show the highest percentage of anthropogenic roosting species to be Molossidae (69.7% roosting in anthropogenic structures). Second highest is

Natural (Grev)
Anthropogenic (Red)

• Preliminary analysis of known viral associations, 45.5% bats were found to not roost in human-made structures whereas 54.4% were found to roost in

• Boxplots suggest negligible differences in viral richness and proportion of zoonotic viruses, between natural and anthropogenic roosting bats.

• Ridgeplots reveal relationships between viral outcomes and roosting status

• Members of the Phyllostomidae and Hipposideridae do not show differences whereas anthropogenic roosting species in the Vespertilionidae,

Pteropodidae, and Molossidae have greater virus richness than their

• Improving coverage of roosting structure

• Change the response variables to the number of viruses and proportion of zoonotic virus rather than binary variables.

• Comparing performance/prediction between models that don't contain