

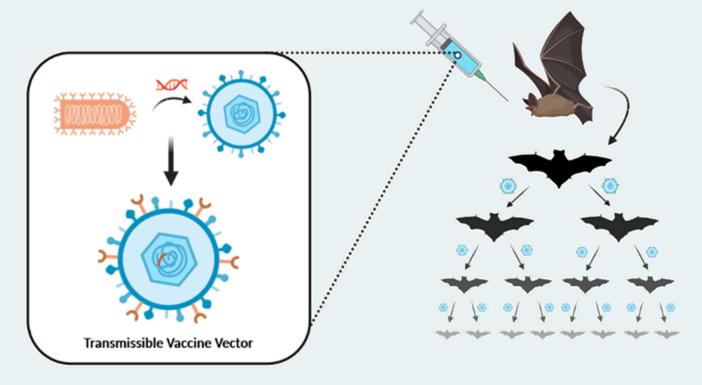




Investigating a **betaherpesvirus** of vampire bat *Desmodus* rotundus as a transmissible vaccine vector against rabies virus

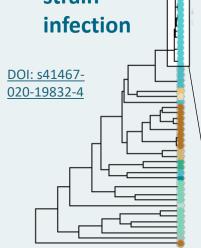
Aims:

- (1) Evaluate DrBHV for key characteristics of a transmissible vaccine.
 - (i) Host specificity
 - (ii) Transmissibility
 - (iii) Super-infection
- (2) Use deep sequencing and longitudinal samples to explore intra- and inter-host transmission dynamics.
- (3) Model DrBHV transmission and **simulate the effects of vaccination** on rabies virus
 outbreaks (ongoing).

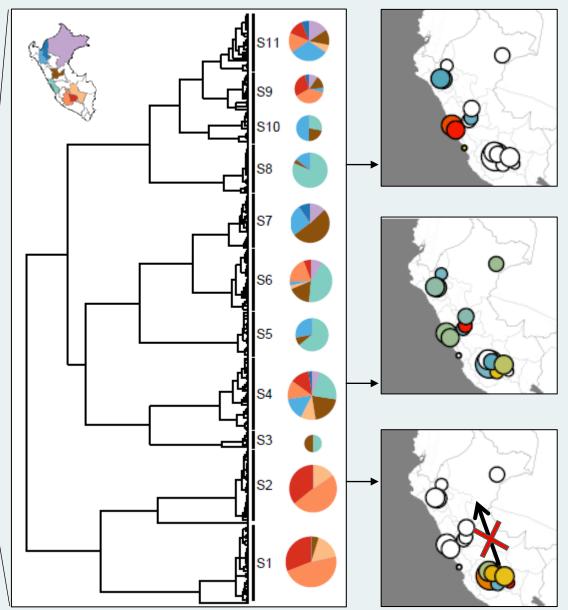


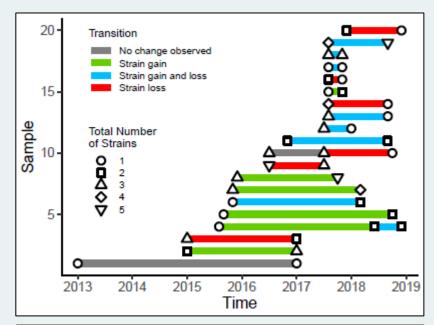
Desmodus rotundus betaherpesvirus: key characteristics

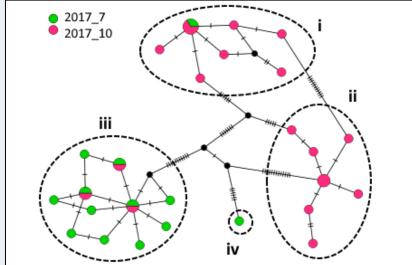
- Host specific at the bat family level
- 97% overall prevalence in all bats
- ~230kbp genome
- Multiple strain infection



Between host dynamics: 11 strains with unique geographic profiles. Unidirectional transmission in some areas. Strain prevalence influenced by strain age.







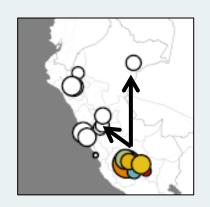
Within host dynamics: longitudinal sequencing shows evidence of strain gain, loss, and long term infection with latency and reactivation.

Conclusions:

✓ Host specific – unlikely to transmit outside of the target family



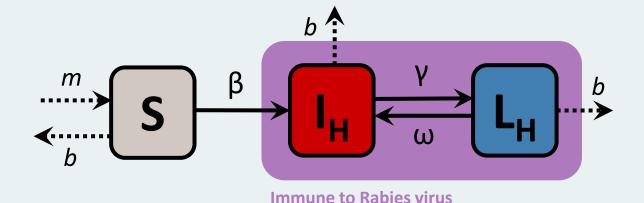
- ✓ **Transmissible** can reach even strain specific prevalences of up to 65%.
- ✓ No interference with wildtype can't show same strain superinfection, but can use strains from different areas



Future work:

Model the equilibrium levels of DrBHV, and the impact of vaccination on rabies outbreaks.

Identify the most important parameters that we don't currently have data for, e.g. how long vaccination lasts.



- Realistic strain-specific prevalence of **30-50%**.
- Active infection in 20-40% of samples.
- Latent period could be as short as 5 months.

