

Mar 24th, 11:00 AM - 2:30 PM

Characterization of a New mtDNA Minicircle in a Chewing Louse, *Geomydoecus aurei*

Ashley L. Campbell
University of Northern Iowa, campbaap@uni.edu

Dino Bolic
University of Northern Iowa, bolicd@uni.edu

See next page for additional authors

Let us know how access to this document benefits you

Copyright ©2020 Ashley Campbell, Dino Bolic, James W. Demastes, and Theresa A. Spradling
Follow this and additional works at: <https://scholarworks.uni.edu/rcapitol>

 Part of the [Biology Commons](#)

Recommended Citation

Campbell, Ashley L.; Bolic, Dino; Demastes, James W.; and Spradling, Theresa A., "Characterization of a New mtDNA Minicircle in a Chewing Louse, *Geomydoecus aurei*" (2020). *Research in the Capitol*. 3.
<https://scholarworks.uni.edu/rcapitol/2020/all/3>

This Open Access Poster Presentation is brought to you for free and open access by the Honors Program at UNI ScholarWorks. It has been accepted for inclusion in Research in the Capitol by an authorized administrator of UNI ScholarWorks. For more information, please contact scholarworks@uni.edu.

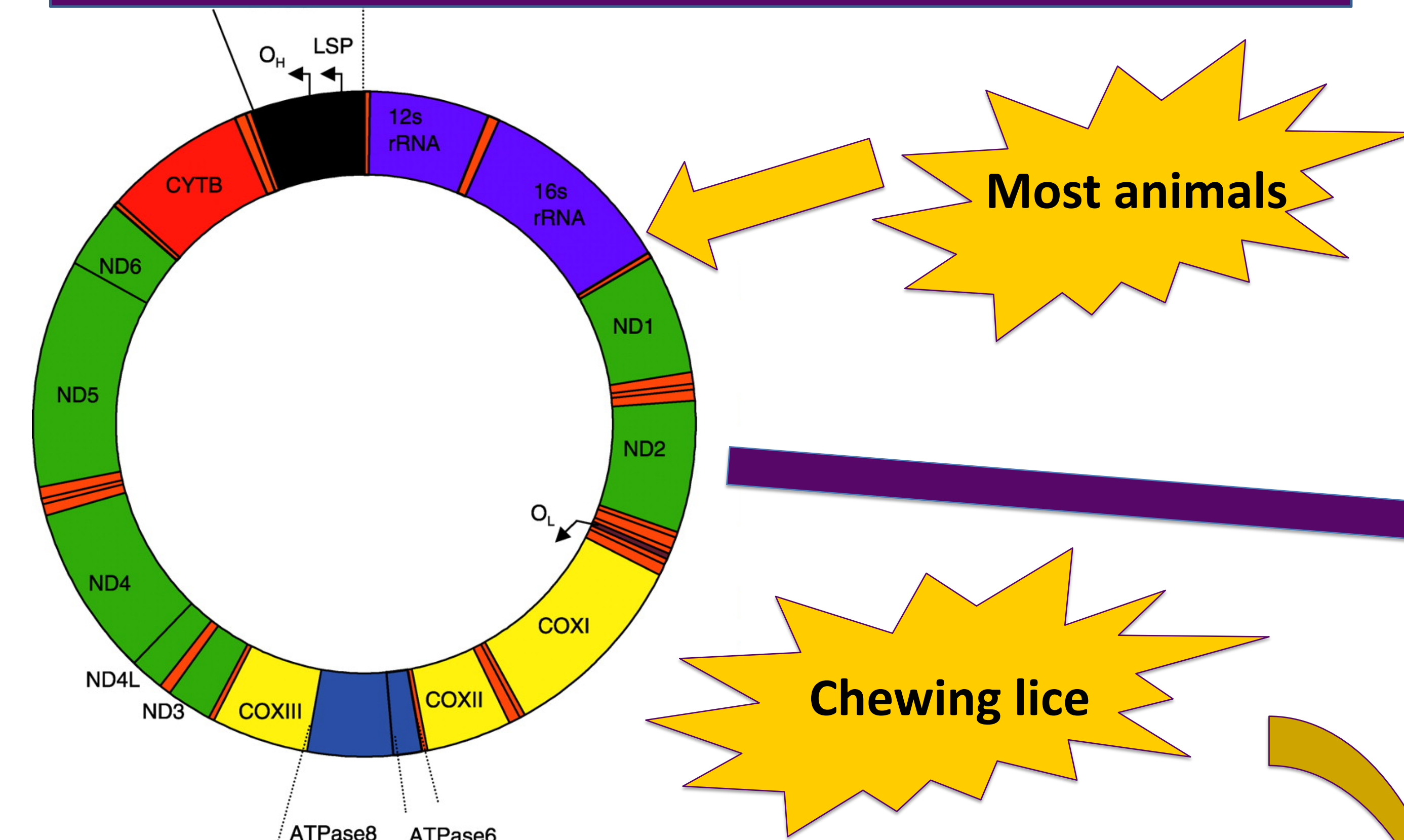
Author

Ashley L. Campbell, Dino Bolic, James W. Demastes, and Theresa A. Spradling



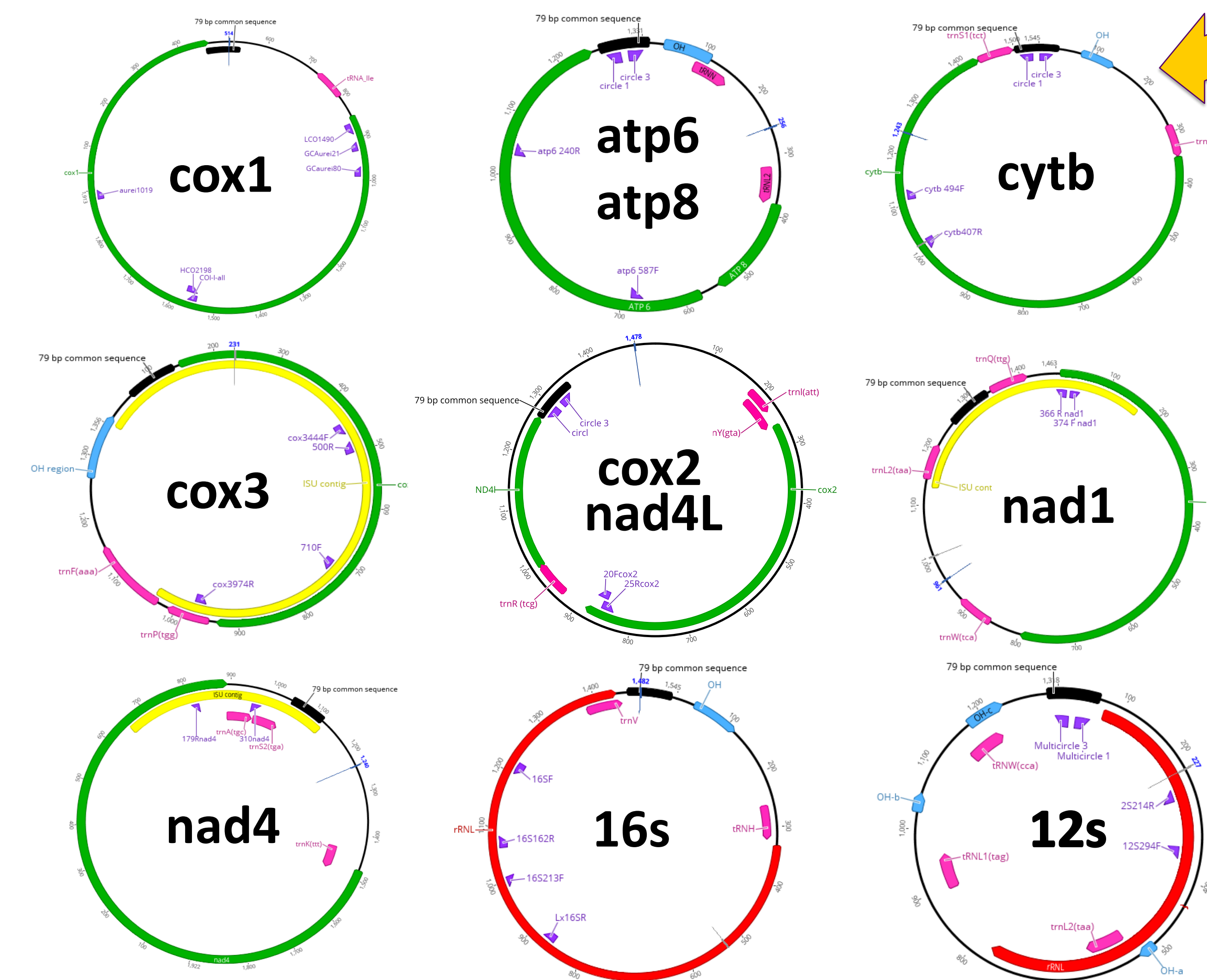
Background

Typical Animal mtDNA Genome

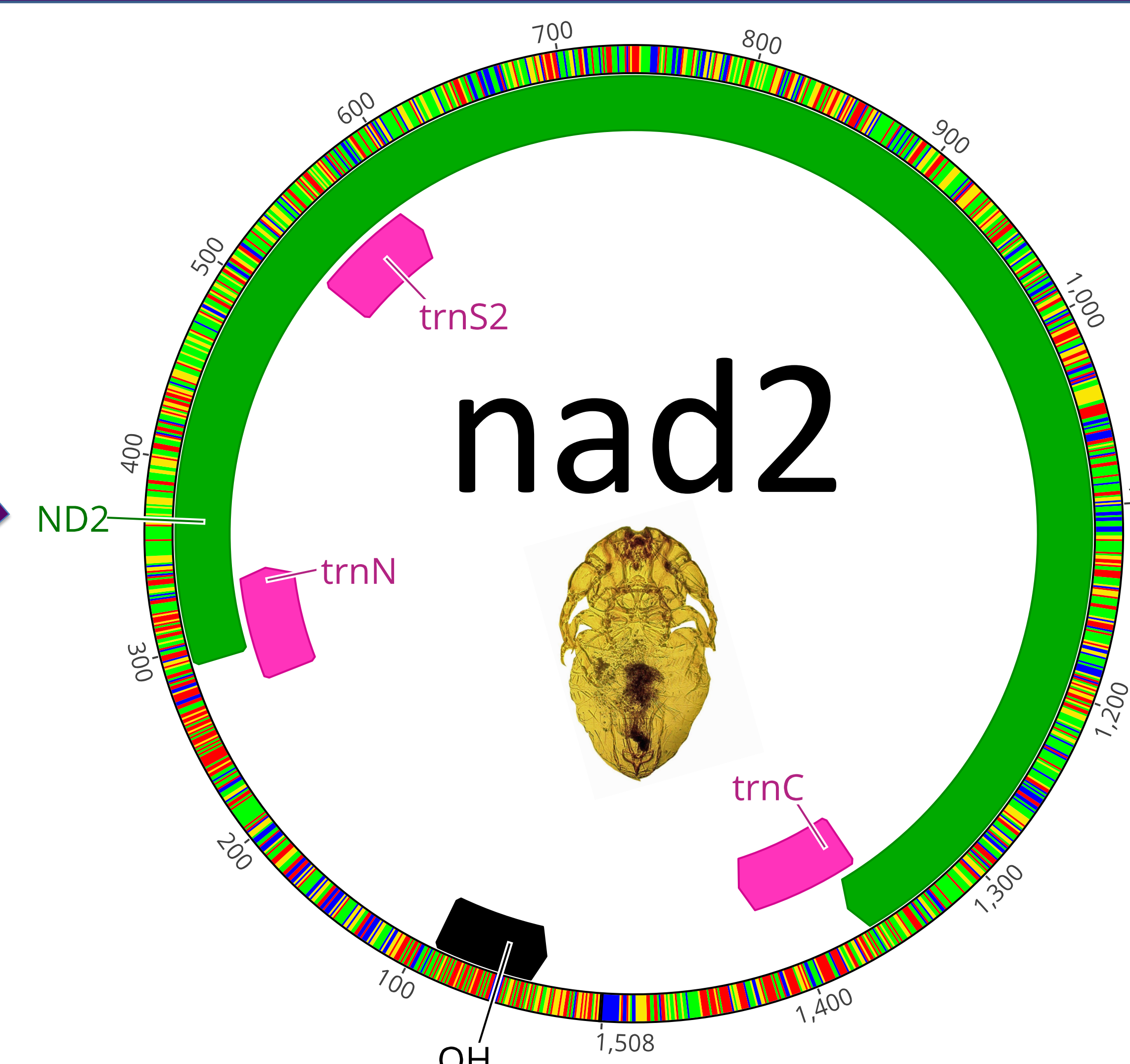


- In most animals, the mitochondrial genome is very conserved with 13 protein coding genes, 22 tRNAs, and two rRNAs in a single circular chromosome (Cameron et al., 2011).
- Many species of lice are unique in the sense that they contain multiple minicircles of fragmented mtDNA instead of a single circular chromosome.

Characterized Minicircles in *G. aurei*



Nad2 Minicircle of *Geomydoecus aurei*



- The newly characterized *nad2* minicircle contains the *nad2* gene and three putative tRNAs: *trnN*, *trnS2*, and *trnC*.
- The *nad2* gene is 1077 bp in length.
- The *nad2* minicircle contains the conserved 79-bp region, shown above in black, that has been identified in all *G. aurei* minicircles characterized to date.
- The *nad2* minicircle was identified and characterized using PCR, cloning reactions, and Sanger sequencing.

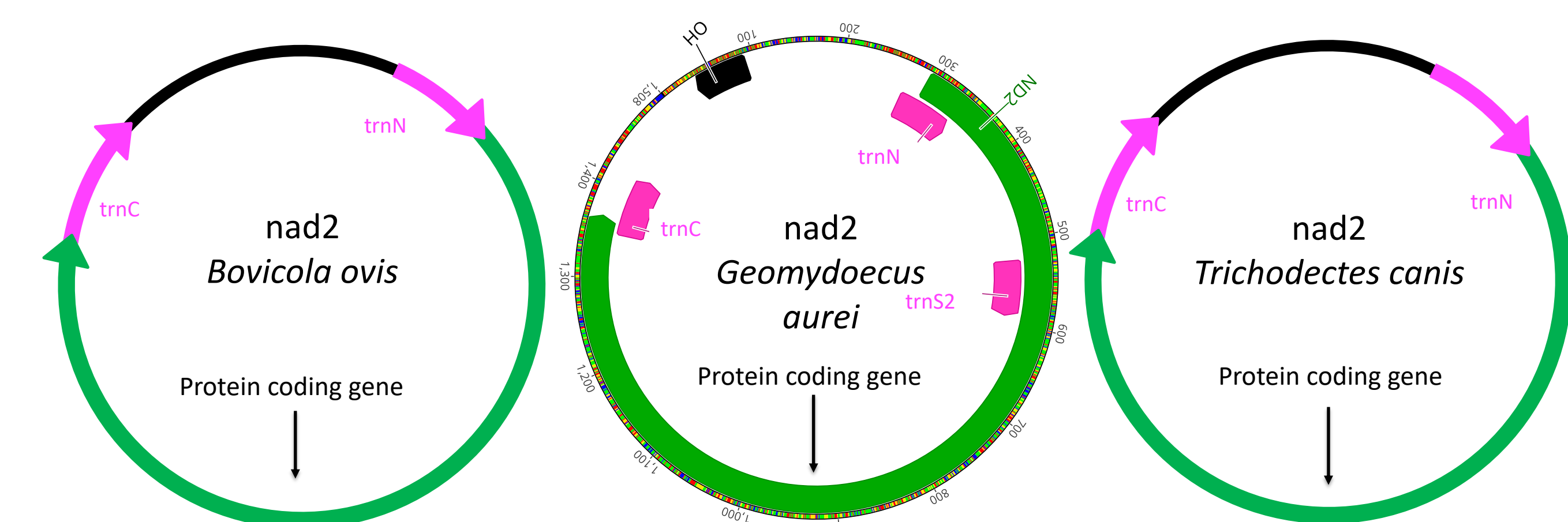
Literature

- Cameron, S. L., Yoshizawa, K., Mizukoshi, A., Whiting, M. F., & Johnson, K. P. (2011). Mitochondrial genome deletions and minicircles are common in lice (Insecta: Phthiraptera). *BMC genomics*, 12(1), 394.
- Herd, K., Barker, S. C., & Shao, R. (2012). Letter: High-level of Heteroplasmy in the Mitochondrial Cox1-Minicromosome of the Human Body Louse, *Pediculus humanus*, and the Human Head Louse, *Pediculus capitis*. *Open Genomics Journal*, 5, 14-17.
- Lightowlers, R. N., Chinnery, P. F., Turnbull, D. M., & Howell, N. (1997). Mammalian mitochondrial genetics: heredity, heteroplasmy and disease. *Trends in Genetics*, 13(11), 450-455.
- Shao, R., Zhu, X. Q., Barker, S. C., & Herd, K. (2012). Evolution of extensively fragmented mitochondrial genomes in the lice of humans. *Genome biology and evolution*, 4(11), 1088-1101.
- Song F, Li H, Liu GH, Wang W, James P, Colwell DD, Tran A, Gong S, Cai W, Shao R. 2018. Mitochondrial genome fragmentation unites the parasite lice of eutherian mammals. *Syst Biol*. 0(0):1-11.
- St John, J. C., Lloyd, R. E., Bowles, E. J., Thomas, E. C., & El Shourbagy, S. (2004). The consequences of nuclear transfer for mammalian foetal development and offspring survival. A mitochondrial DNA perspective. *Reproduction*, 127(6), 631-641.
- Xiong, H., Barker, S. C., Burger, T. D., Raoult, D., & Shao, R. (2013). Heteroplasmy in the mitochondrial genomes of human lice and ticks revealed by high throughput sequencing. *PLoS one*, 8(9), e73329.

Results

Conclusions

- The 79-bp DNA sequence found on this circle is identical to sequence found on nine other *G. aurei* minichromosomes, suggesting functional importance.
- The majority of the tRNAs identified in minicircles containing protein coding genes have been found nearly adjacent to the coding region or to very minimally overlap with the coding sequence.
- The gene order of the *G. aurei nad2* minichromosome (*trnN-nad2-trnC*) is seen in other mammalian louse species (Song et al, 2018), including *B. ovis*, and *T. canis* (shown below).
- The position of *trnS2* within the interior of the *nad2* gene is unusual and requires further investigation. Another DNA sequence that is a likely *tRNS2* gene is found on the *G. aurei nad4* minichromosome in a position also seen in other louse species. *tRNS2* has not been associated with *nad2* in other lice.



Next Steps

- Two protein coding genes, *nad3* and *nad5* have been identified and are in the process of being characterized and mapped.
- There are four tRNAs (*trnD*, *trnG*, *trnM*, and *trnT*) expected to be found upon the mapping of the remaining minicircles.
- Discovery of the remaining gene, *nad6*, has been problematic in this and other studies of many insects.
- Additional PCR, cloning reactions, and Sanger sequencing will be performed in an effort to identify *nad6* and map the mitochondrial genome of *G. aurei* to completion.

Acknowledgements

- Dr. Alan Orr Research Award
- NSF Award (DEB-1445708)
- UNI Intercollegiate Academics Travel Fund