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Out with the Old and in with the New: A Comparison Between Molecular and Traditional Techniques to Identify Parasitized Birds

Christian Guerrero Department of Biological Sciences, Boise State University

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Introduction

Traditionally, the identification of blood parasites has been based on the visual examination of blood smears. This approach depends on individual expertise in making blood smears and identifying parasites which can vary widely from person to person. Recent work shows that reading blood smears is significantly less sensitive than using molecular studies in identification. Thus, the accuracy of data can fluctuate greatly. This project compares the ability of investigators to identify infected birds using both blood smears and microscopes with their ability to identify infected birds through molecular analysis of blood from the same sample.

Methods

During the Fall 2011 migration (September – October), raptors were captured at the Idaho Bird Observatory in Boise, ID.

- Blood was drawn from either the jugular or wing vein
- Part of the blood was used to make blood smears and remainder was stored for further molecular studies
- Blood smears were later stained using *Giemsa-Write* Hematological Stain
- Slides were examined for parasites
- DNA was extracted from the stored blood using an EZ BioResearch mini-kit
- DNA was screened for Avian Malaria using specific Plasmodium, Haemoproteus and Leucocytozoon primers: HaemNFI, R3, F, R2, FL, and R2L



Cooper's Hawk (Accipiter cooperi)



Northern Goshawk (Accipiter gentilis)



Sharp-Shinned Hawk (Accipiter striatus)

Traditional Approach



Red-Tailed Hawk (Buteo jamaicensis)

- Blood smears were analyzed through the use of a light microscope at 1,000 times magnification Morphological characteristics were used to verify
- parasitic organisms





Presence of Parasites in Blood Samples		
ID	Microscopy	PCR
02062	\checkmark	\checkmark
07766		\checkmark
07805		\checkmark
09147		\checkmark
09194		
15930		
16648		\checkmark
16682	\checkmark	\checkmark
16695		\checkmark
28688	\checkmark	\checkmark

Results

- Species within the *Plasmodium*, *Haemoproteus*, and Leucocytozoon genera were found to be parasitic Through Microscopy, three of ten samples were
- deemed parasitized
- Through PCR analysis, eight of ten samples were deemed parasitized
- The traditional method incorrectly passed five of eight parasitized samples as being free from parasites Only 37.5% of parasitized samples were read correctly



American Kestrel (Falco sparverius)

Fallon, Sylvia M., and Robert E. Ricklefs. 2008. "Parasitemia in PCRdetected Plasmodium and Haemoproteus infections in birds". Journal of Avian Biology. 39 (5): 514-522. Hellgren O, J Waldenström, and S Bensch. 2004. Figure 1 "A new PCR assay for simultaneous studies of Leucocytozoon, Plasmodium, and Haemoproteus from avian blood". The Journal of Parasitology. 90 (4): 797-802.

References

Molecular Approach

- Reaction (PCR)



HaemFi 5'-ATGGTGCTTTCGATATATGCATG-3 $HaemNE1^{+} \sim$

This comparative study reinforces the notion that visually inspecting blood smears is less accurate than confirming the presence of Avian Malaria through DNA analysis. This is due to the traditional method's limitations which "fail to register many malaria parasite infections that are picked up by PCR Screening" (Fallon & Ricklefs). By placing a preference towards the more accurate molecular approach, the misdiagnosis and improper treatment of a parasitized bird will be kept at a minimum. Further work can improve the management and conservation efforts of wild raptors and their vectors.



- The Idaho STEP Program



Extracted DNA was analyzed using Polymerase Chain

Extracted DNA with PCR reagents were put through a thermal cycler to amplify Avian Malaria DNA sequence Amplified DNA was analyzed through electrophoresis and bands were examined under a UV light The presence of bands in the examination of the gel determines if parasitic DNA was extracted

> Primers used for the Screening of DNA Haem NR3 -<u>GCATTATCTGGATGTGATAATGGT</u> Haem R2² 5'-CATTATCTGGATG<u>A</u>GATAATGG<u>IGC</u>-3' Haem R2L³

Discussion

Acknowledgements

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