HOW TO

PREPARE BIRD SPECIMENS

Part 9a – Washing bird skins for arthropod ectosymbionts

Part 9b - Drying a washed bird skin











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The Migratory Bird Conventions Act regulates the take and possession of birds in Canada. The Migratory Bird Treaty Act regulates the take and possession of birds in the United States. In addition, the provinces (in Canada) and the states (in the United States) also require permits. For some species SARA, ESA, or CITES permits may be required.

Always check the laws of your country and obtain the proper permits; failure to do so may result in civil and/or criminal penalties

When handling dead birds, it is probably impossible to tell if a bird is infected with a pathogen that may cause human illness even if you know the cause of death to be a wound or an injury. Take reasonable precautions to protect yourself. The Ornithological Council offers a peer-reviewed fact sheet on avian zoonotic disease and safety precautions for those who handle birds in the field and in the lab.

http://www.nmnh.si.edu/BIRDNET/docume nts/WNV&H5N1-FactSheet.pdf





Taxonomic relationships of arthropod ectosymbionts and their avian hosts are used to study speciation and to determine evolutionary trees of both arthropods and birds.

Record all ectosymbionts found on the avian data sheet. Heavy infestation may indicate other health issues.



The four major groups of arthropod ectosymbionts are:

- Lice (Mallophaga)
- Fleas (Siphonaptera)
- Mites including ticks (Acarina)
- Flat flies (Hippoboscidae)

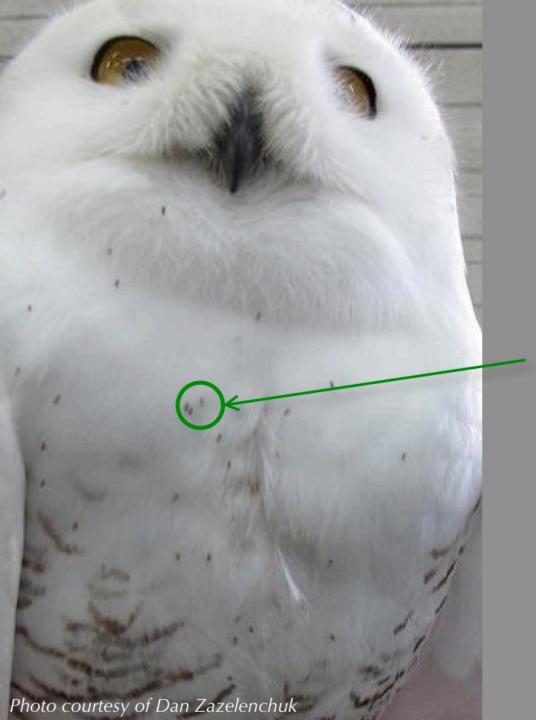


Lice (Mallophaga):

- · Live between feather shafts
- Feed on feathers or blood
- Females attach eggs to feathers

The Curlew Quill Louse (Actornithophilus patellatus) spends part of its life cycle in the quills of its host; this is the only known louse to live inside quills and feed on the pith (pulp) inside feather shafts.





Due to the white plumage, the heavy louse infestation is easy to see on this Snowy Owl.







Mites (Acarina) are divided into 4 groups:

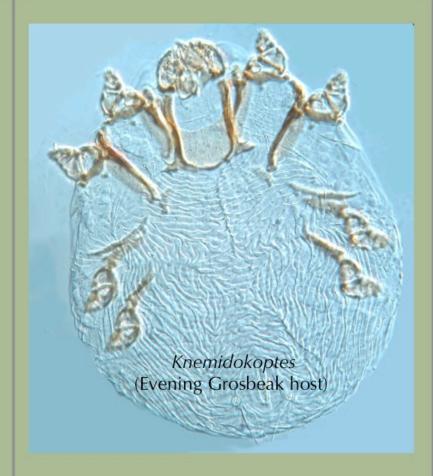
- 1. Feather mites (Astigmata: Analgoidea and Pterolichoidea):
- Most species live on feather surfaces and feed on uropygial (preening) oil
- A few species live on the skin's surface and feed on skin flakes
- A few species live inside quills and feed on feather pith





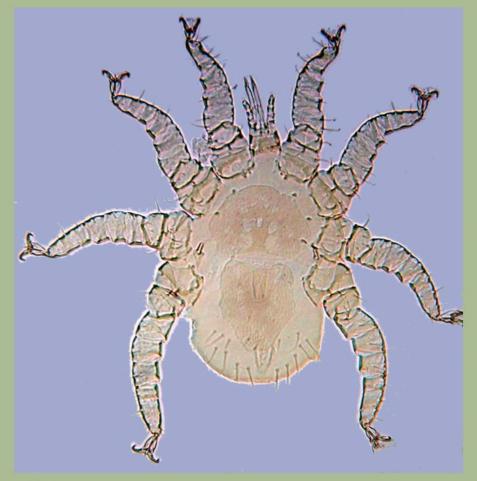
- 2. Quill Mites (Prostigmata: Syringophilidae):
- Live inside quills
- Feed on host body fluids by piercing quill wall into adjacent tissues with their long chelicerae





- 3. Scaly-leg mites
 (Astigmata: Epidermoptidae
 Knemidokoptinae):
- Feed on keratinous tissues of the skin, particularly legs and beaks

- 4. Nasal mites (Mesostigmata: Rhinonyssidae):
- Live inside the nasal passages and upper respiratory track
- Feed on host blood and tissues



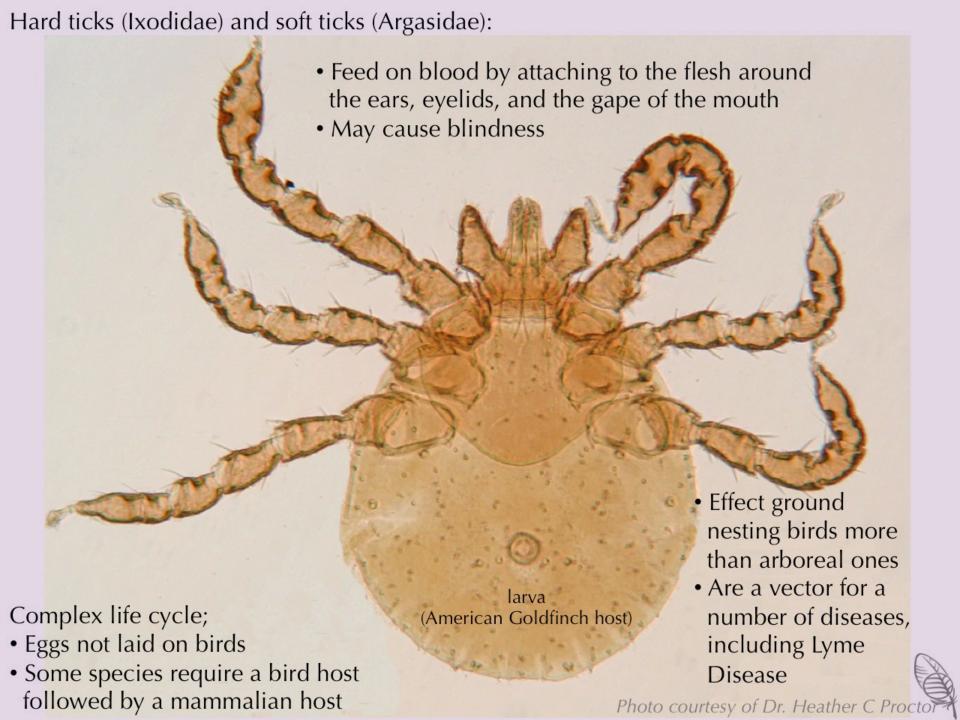
♀ *Tinaminyssus melloi* (Rock Pigeon host)

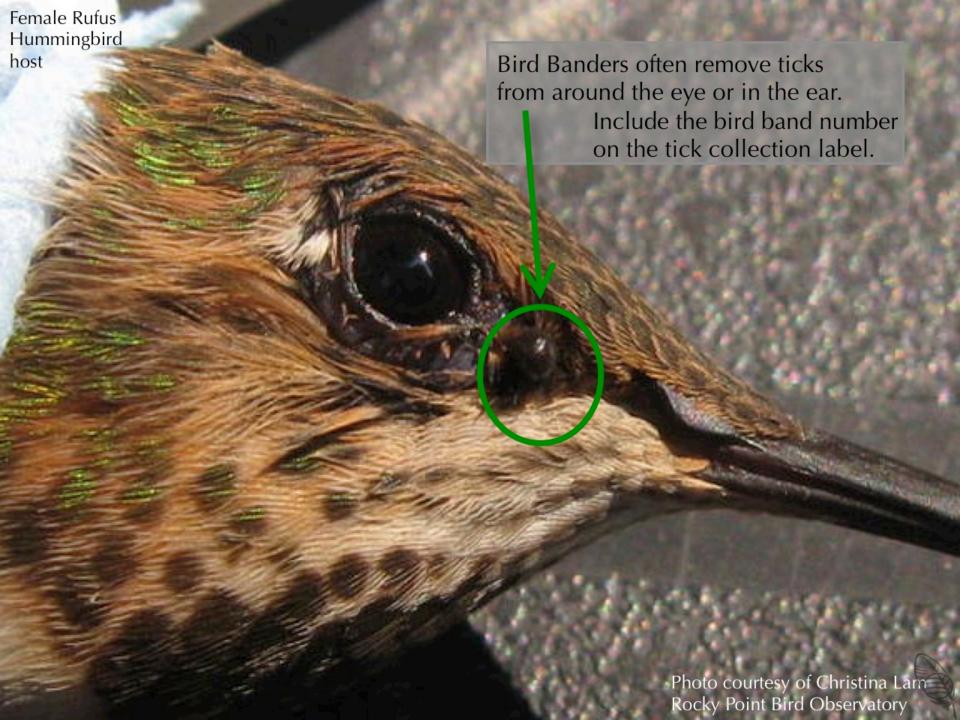
Dermanyssus quintus (Hairy Woodpecker host)

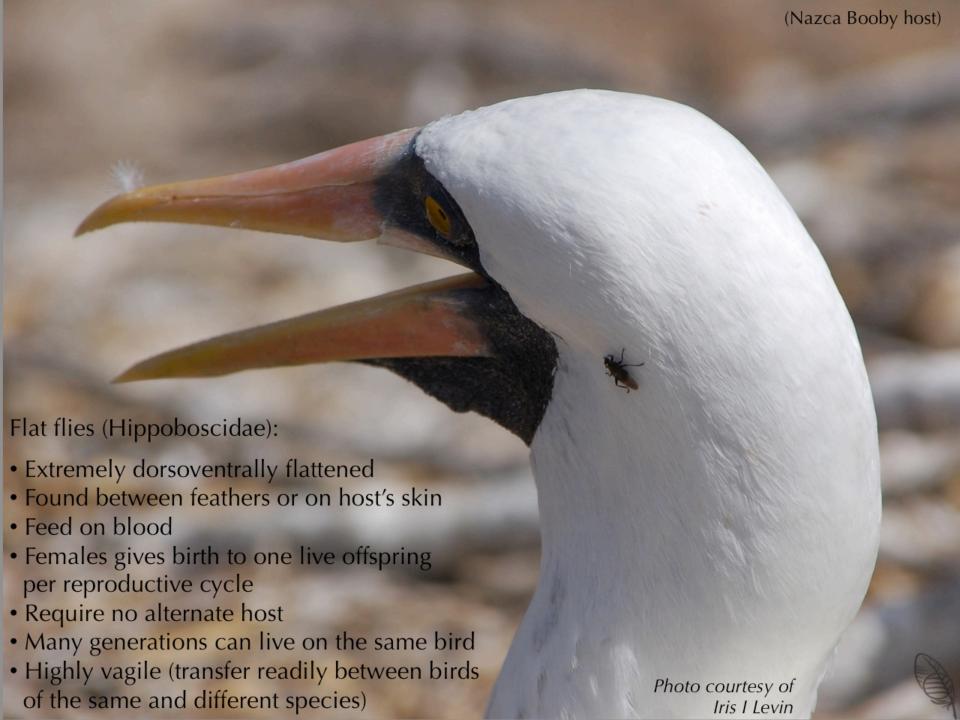


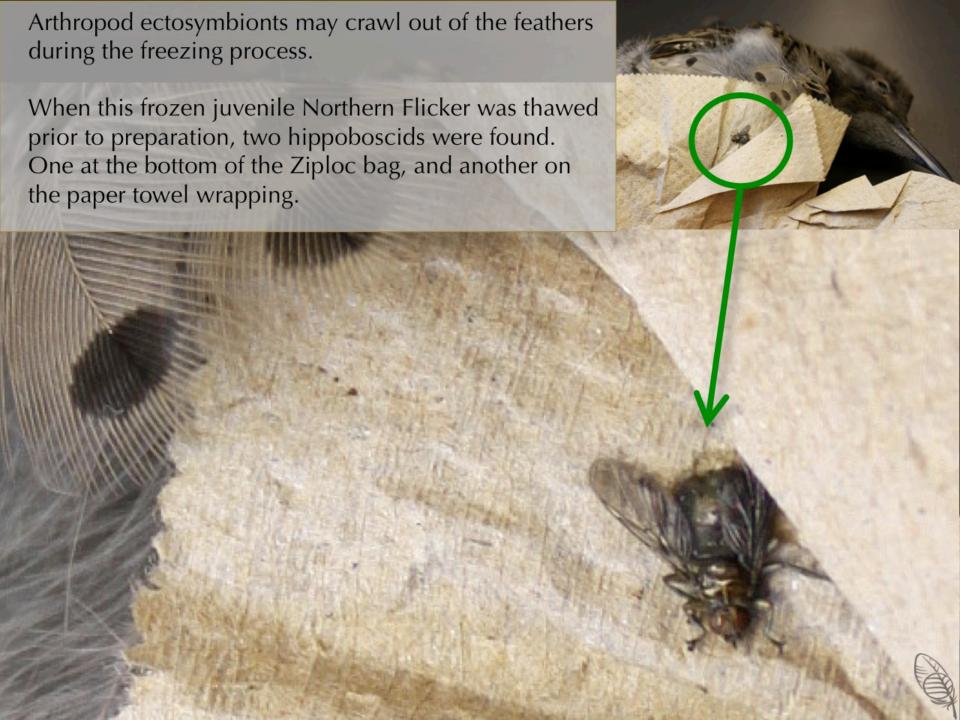
- 5. Fowl mites (Mesostigmata: Dermanyssdae and Macronyssidae):
- Feed on blood
- Depending on the species, live either on the bird or in the nest and surrounding area
- Heavy infestations kill nestlings

Photo courtesy of Dr. Heather C Proctor

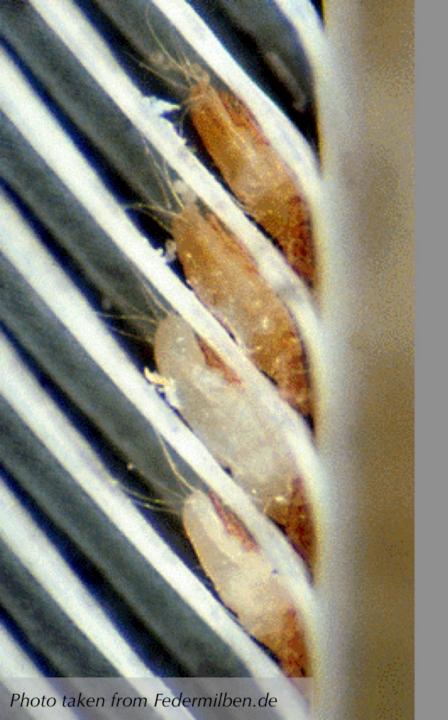












For additional information:

Dr. Heather C. Proctor web publication: List of Feather Mites and Their Hosts

http://www.biology.ualberta.ca/faculty/heath
er_proctor/?Page=5626

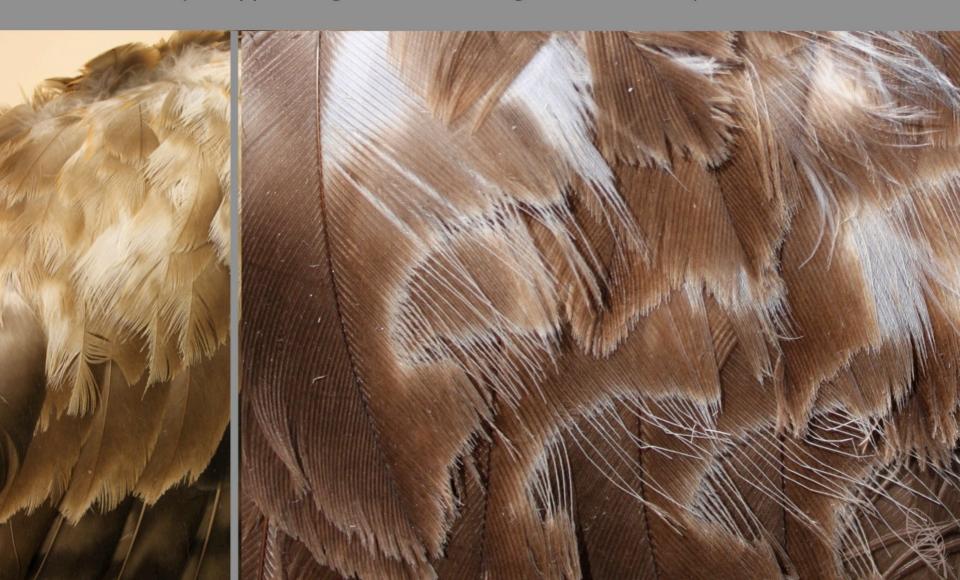
The following website does not appear to have been updates since 2007. Click on the ♀ and ♂ to view pictures of arthropod ectosymbiont species.

http://www.federmilben.de/en/gallery.html



Washing birds in poor heath or with one of the following two conditions will increase your changes of finding arthropod ectosymbionts:

(1) Look for frayed upper wing covers indicating a medium-heavy feather lice infestation



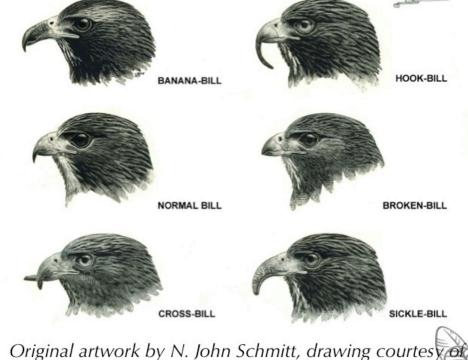
(2) Avian Keratin Disorder (a.k.a. Long-bill Syndrome)

Deformed beaks may decrease grooming efficiency resulting in higher than normal arthropod ectosymbionts loads.

This syndrome occurs in hummingbirds, woodpeckers, raptors, passerines, and probably other orders of birds.







Falcon Research Group (frg.org)

Gather the following:

- 1. Washing Tools:
- Sink (not shown)
- Wash basin
- Soap
- Tooth brush



- 2. Drying Tools:
- Cloth or paper towels
- Tooth brush

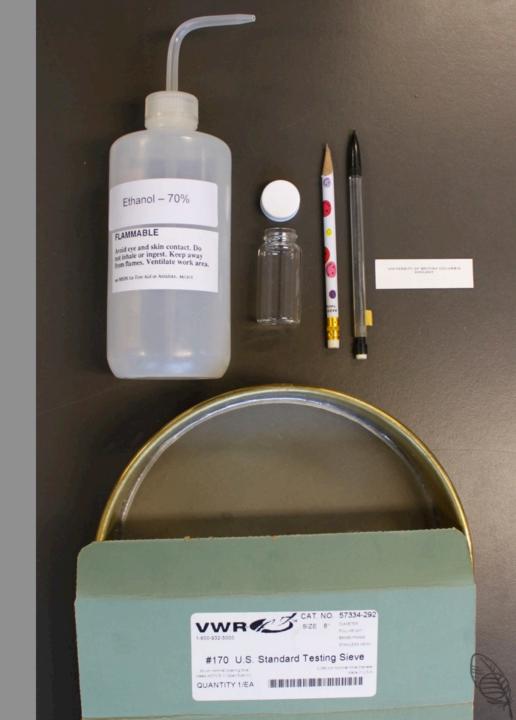
Plus one or more of the following:

- Hair dryer & chemistry stand
- Compressed air
- Mechanical or manual tumbler with an absorbent (corn meal, corn cob, or fine sawdust)



For arthropod symbionts sampling, these additional tools are required:

- Wash bottle filled with 70 or 95% ethanol
- Wash bottle filled with tap water
- Large container with a tight fitting cover
- Fine mesh sieve
- Vial (s)
- Internal label that is alcohol resistant
- Pencil or alcohol proof pen





OPENING MICROMETER

90





MADE IN U.S.A.

A 90 micrometer mesh sieve is commonly used. If available, use a 45 micrometer sieve. This is especially important when sampling for mites.

This PowerPoint can only give a quick overview.

Use this link to view a detailed 30 minute step-by-step video titled:

Washing Birds for Arthropod Ectosymbionts

http://youtu.be/iJZushKix2g

Watch Dr. Heather C. Proctor demonstrate how to wash a bird for arthropod ectosymbionts, decant and store the avian washings, examine and extract the arthropods, plus learn how to make and label arthropod microscope slides.

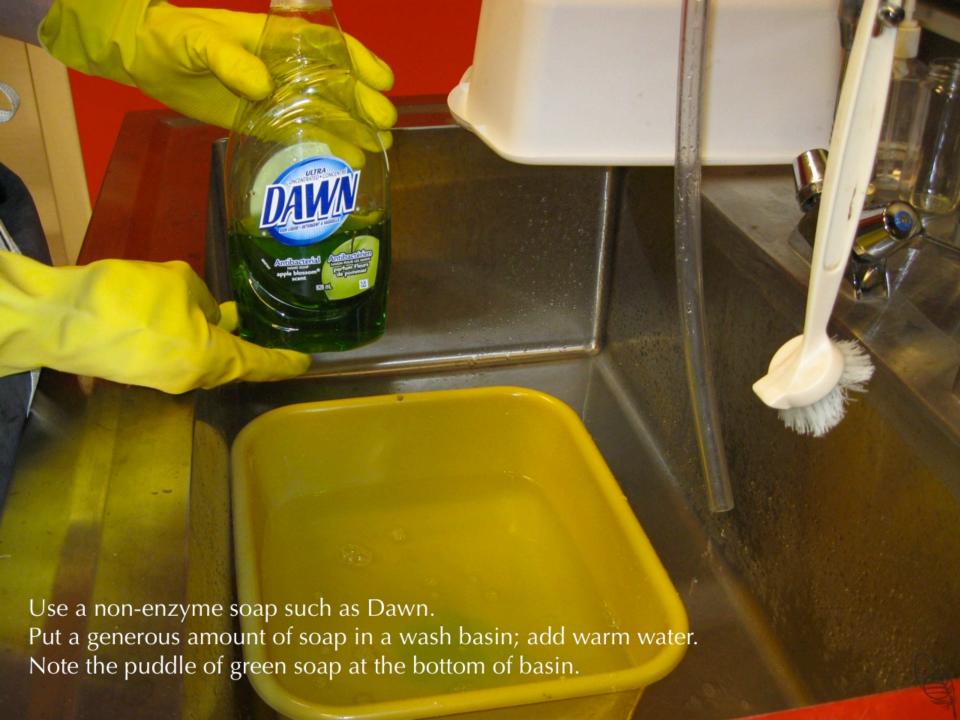


If possible, start with an ethanol pre-wash to help loosen arthropods:

- Using a wash bottle, liberally dose the bird with 70% or 95% ethanol
- Squirt Dawn soap directly on the bird
- Put the bird in a wash basin or bucket, turn the tap water on so the water runs through the feathers
- Soak the bird for 5 minutes
- Open the bird's beak and irrigate the mouth and nasal passages to sample for nasal mites
- Swirl the bird in the basin or, if you are using a container with a lid, shake for 5 minutes

Regardless of if you include an ethanol pre-wash or not, follow the same steps to rinse, sieve, decant the residue, and dry the bird.

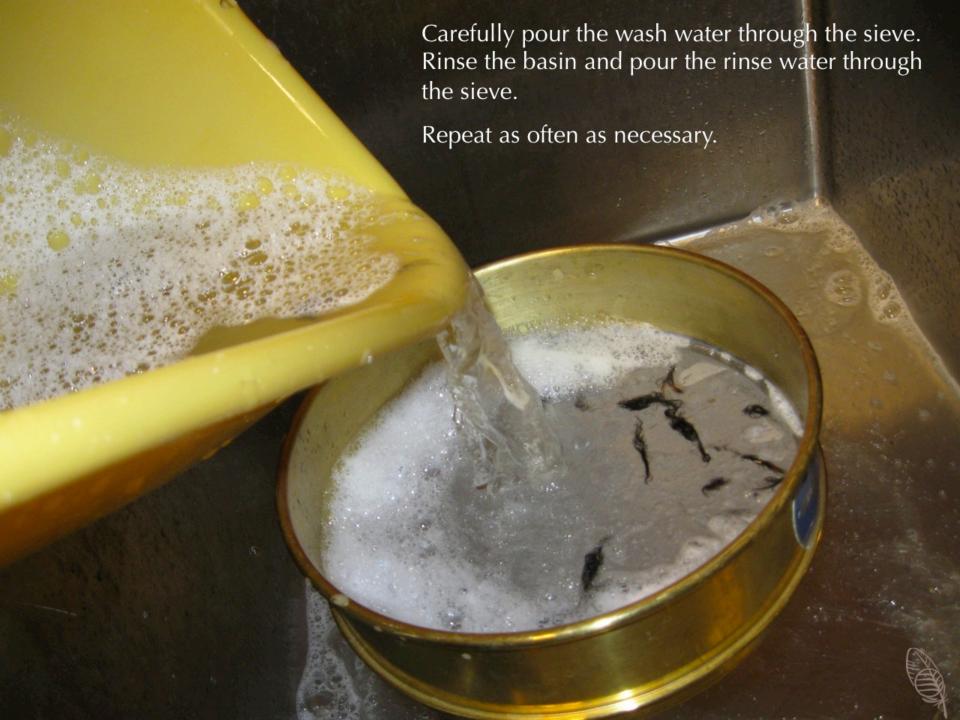






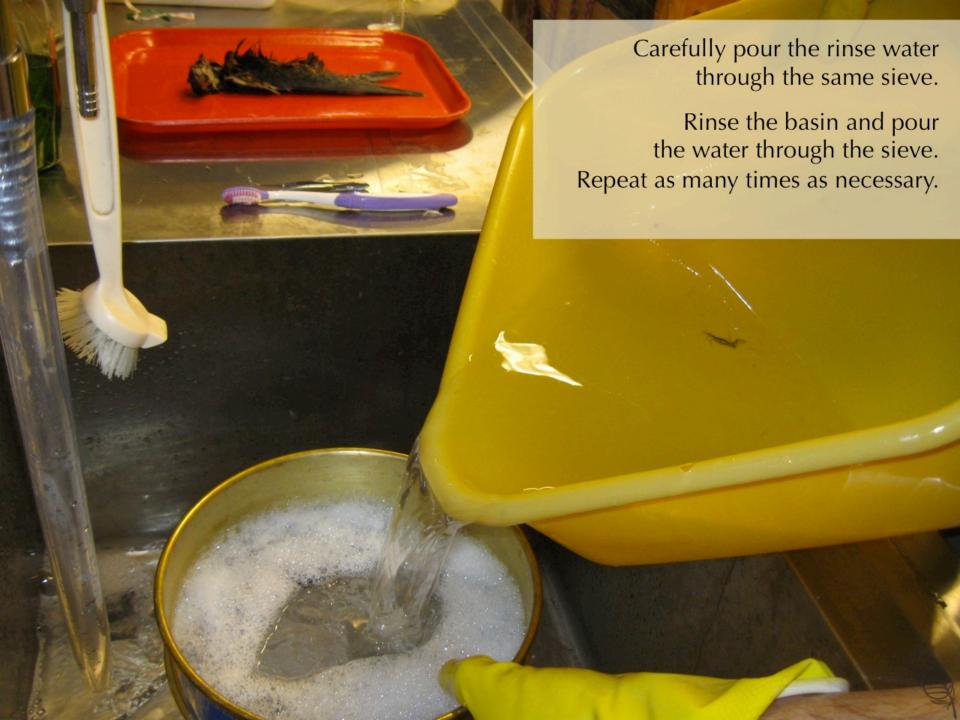


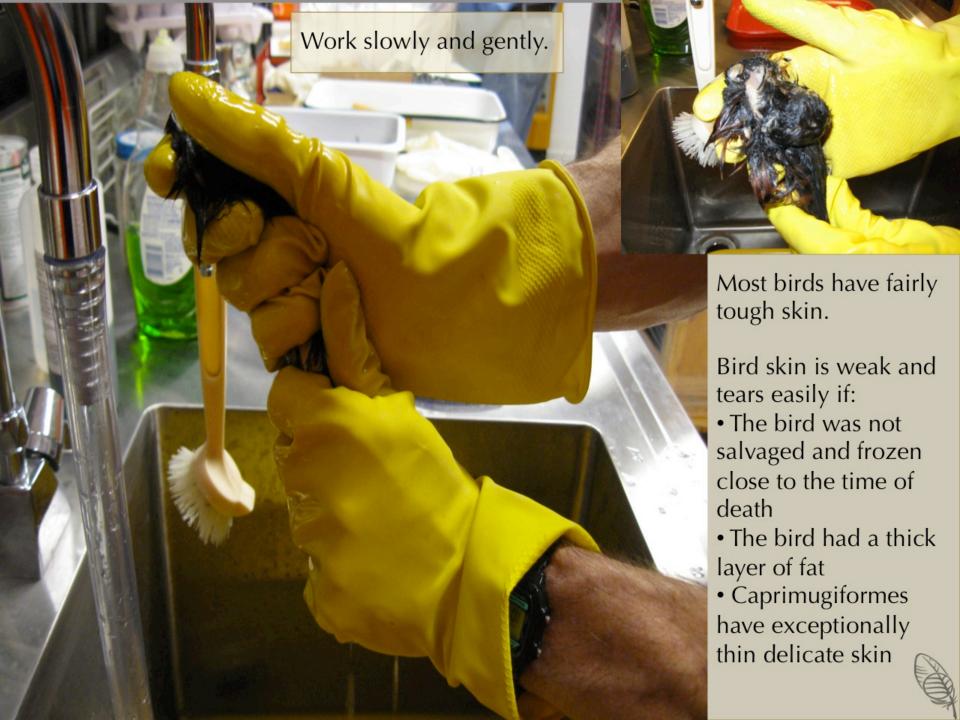












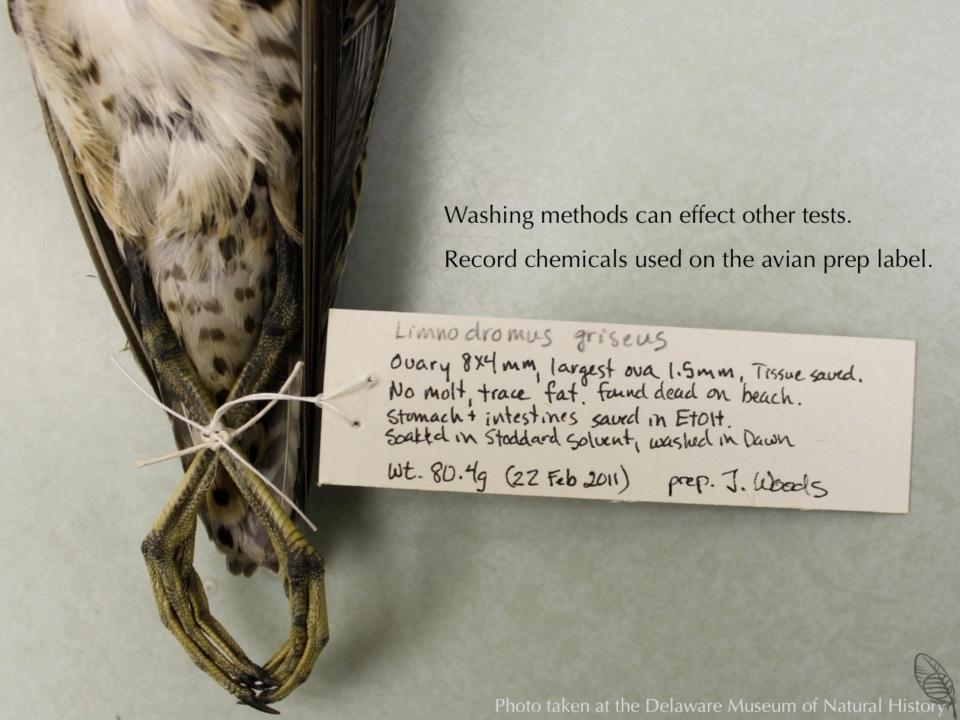


After pouring the final rinse water through the sieve:

- Tip the sieve
- Use a water-filled wash bottle to concentrate the residue in one area of the sieve
- Use a fixative-filled wash bottle to transfer the residue to a vial
- Use a pencil or alcohol resistant pen to write a detailed label on alcohol resistant paper
- Insert label inside the jar
- On the lid, write the prep number, or host species to facilitate vial retrival

Note: Wet feathers dilute the fixative. If possible, use 95% ethanol.





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Part 9b - Drying a washed bird skin

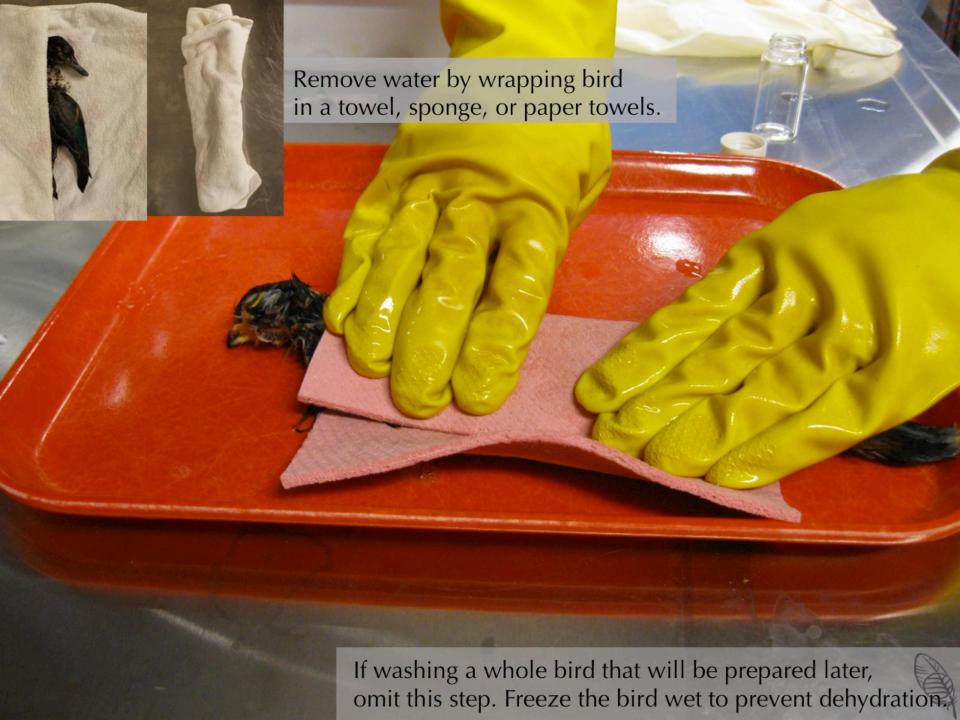


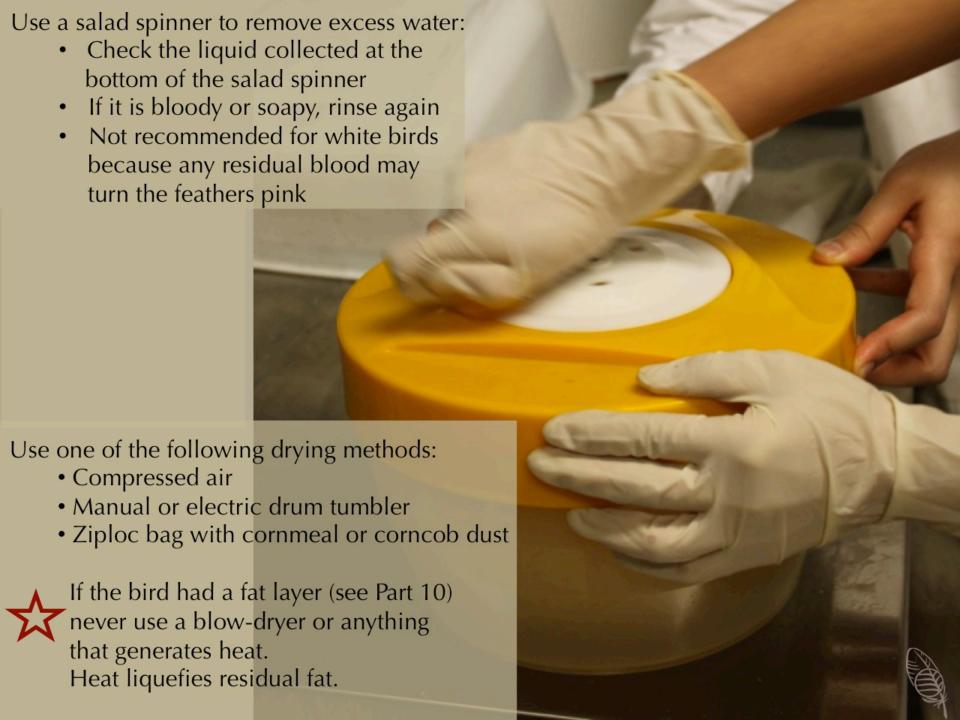
















Create a drying station: (Having both hands free is a real advantage and time saver.)

- Use chemistry stands or the equivalent to hold a hair dryer, or compressed air nozzle
- Commercial hand dryers found in some public washrooms are excellent for drying birds
- A portable fume hood increases the air flow over the bird



Check frequently constantly changing the bird's position.

- If it is a small bird (~20grams) check once a minute
- For a 1 kilo bird, check every five minutes









Drum dryers are available at taxidermy supply stores. They come in several sizes and are either top or side loading. Fill the drum 1/3 full with fine sawdust or corncob dust. An engine spins the drum slowly. Drying time depends on bird size, 10-30 minutes is the norm. This is a messy process.





Going clockwise from the top left, photos taken at the Royal British Columbia Museum, University of Washington Burke Museum, and the Smithsonian Natural History Museum



Use a container with a tightly fitting lid to make a manual tumbler. Wash, towel dry, and submerge in absorbent material (corncob shown below).





Repeat until dry







Be sure to pack the neck from both ends.

In this case, the absorbent is white corn meal and the manual tumbler is a wide mouth white plastic jar with a screw top lid.

In the field, a Ziploc bag is a good alternative.





IN MEMORIUM



DR. REX KENNER

Former Curator of the Cowan Tetrapod Collection who encouraged me to begin this project.

Special thanks to Dr Heather C Proctor for helping out with both this PowerPoint Presentation and for agreeing to be the featured presenter for the Beaty Biodiversity Museum video that was created to complements this presentation.

Without the encouragement of the Spencer Entomology Collection staff ((Karen Needham, Don Griffiths, Robert Curtiss III), this arthropod ectosymbiont presentation would not exists. I also wish to thank Eve Szabo, Hans van Brandwijk, N. John Schmitt, Bud Anderson, Jean Woods, Ellen Paul, and all the wildlife rehabilitators, bird banders, pathologist, museum curators and collection managers who have helped and encouraged me to complete this project. I take full responsibility for any remaining mistakes.

Without the technical assistance of Derek Tan, this project would never have gotten off the drawing board. Dr. Darren Irwin kindly suggested and made the arrangements for this series to be posted on the Beaty Biodiversity Museum website. A huge thank you to the staff and volunteers at the Cowan Tetrapod Collection for providing space and creating a terrific work environment.

Unless otherwise indicted, all pictures were taken by the author at the Cowan Tetrapod Collection, University of British Columbia Beaty Biodiversity Museum.







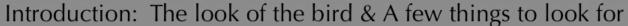








PRESENTATIONS IN THIS SERIES



Part 1 - Spread wings, a good way to start

Part 2 - Skinning your first bird

Part 3 - Other skinning methods

Part 4 - Stuffing your first bird

Part 5 - Other stuffing and pinning methods & Bird parts

Part 6 - Sexing birds using gonads (includes 2 quizzes with answer sheets)

Part 7 - Determining skull pneumatization & Skeleton preparation

Part 8 - DNA tissue sampling & Gut analysis

Part 9 - Washing skins for arthropod ectosymbionts & Drying washed skins

Part 10 - Recording fat levels & Cleaning fatty or stinky skins

Part 11 - Flat skins, shmoos, and other types of study skins

Part 12 - Preserving eggs and shell fragments (in prep)

Part 13 - Determining cause of death

Part 14 - Labelling: the most important step

To download another PowerPoint presentation in this series go to:

http://beatymuseum.ubc.ca/birds









