



USDA APHIS Identification Technology Program

---

# Practical considerations for seed imaging

Madeline Maher

Botanist, USDA/Colorado State University

<https://idtools.org/>



**United States Department of Agriculture**  
Animal and Plant Health Inspection Service

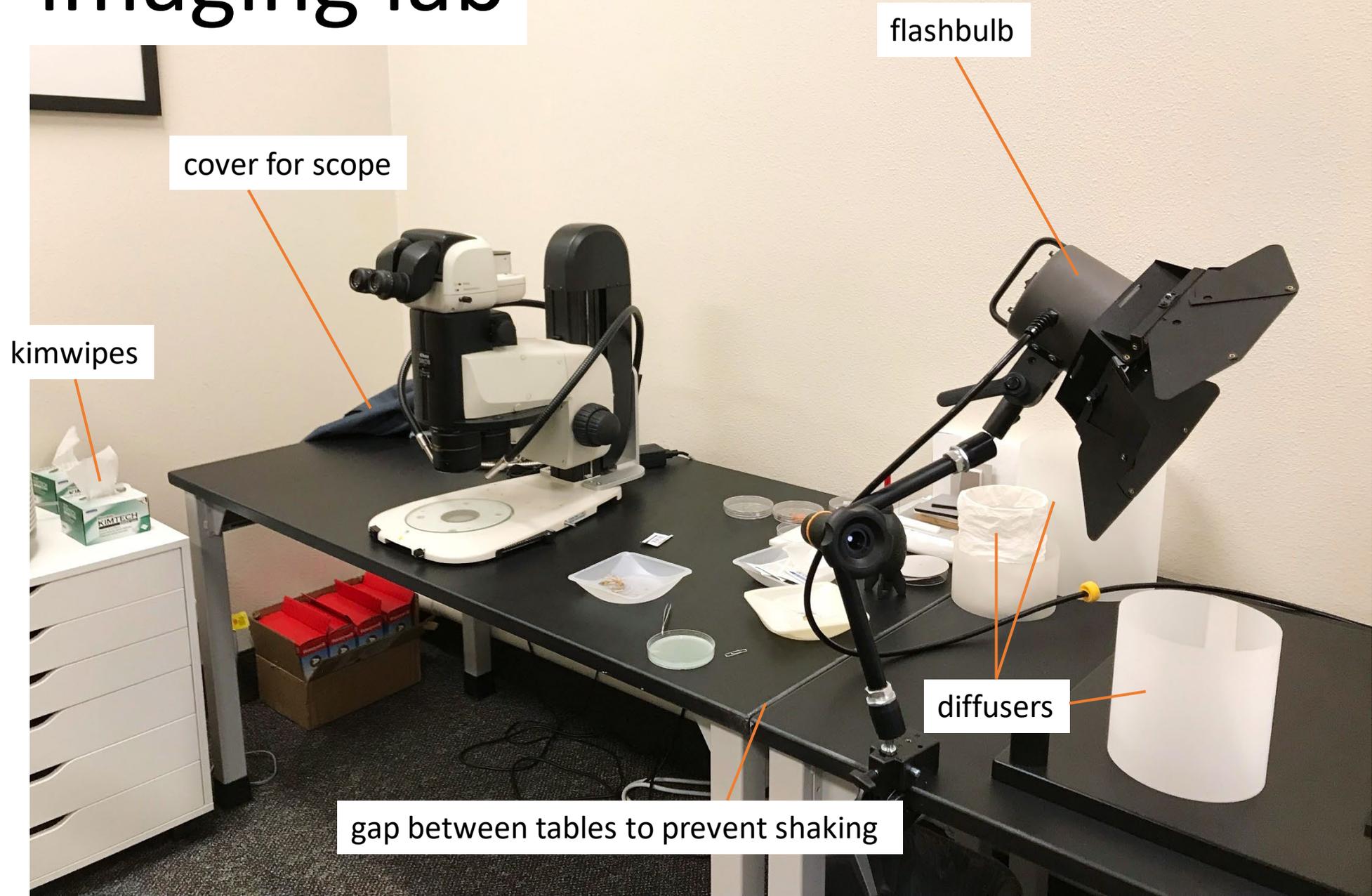
# Key considerations for image quality

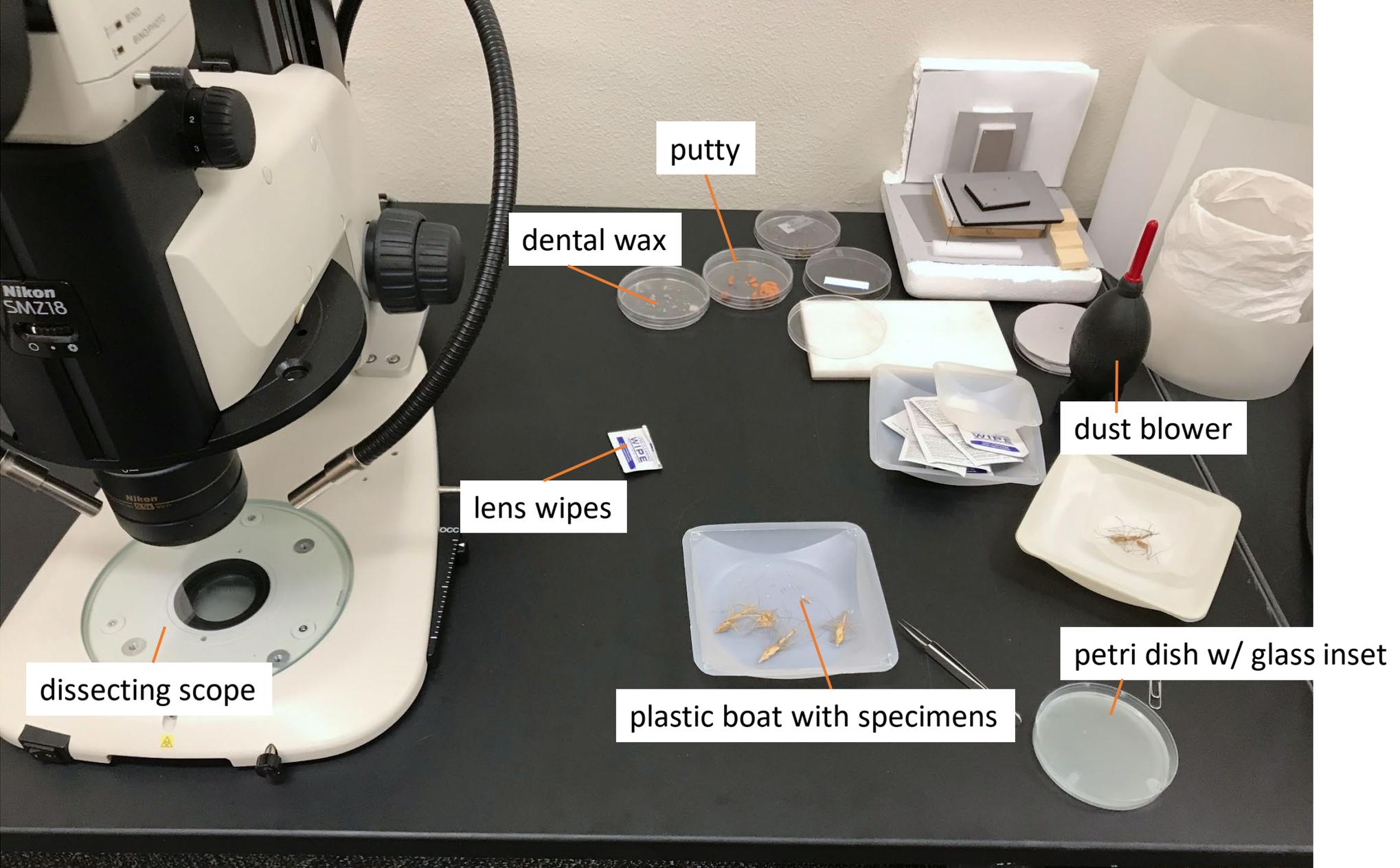
1. Minimize dust/contamination
2. Minimize movement
3. Lighting should be diffuse but bright
4. Camera settings should be appropriate for the circumstances
5. Stacking capabilities helpful for optimal quality, but not always needed

A detailed seed-imaging protocol and guide will be shared on the ISMA website.

Link to follow.

# Our imaging lab





dissecting scope

lens wipes

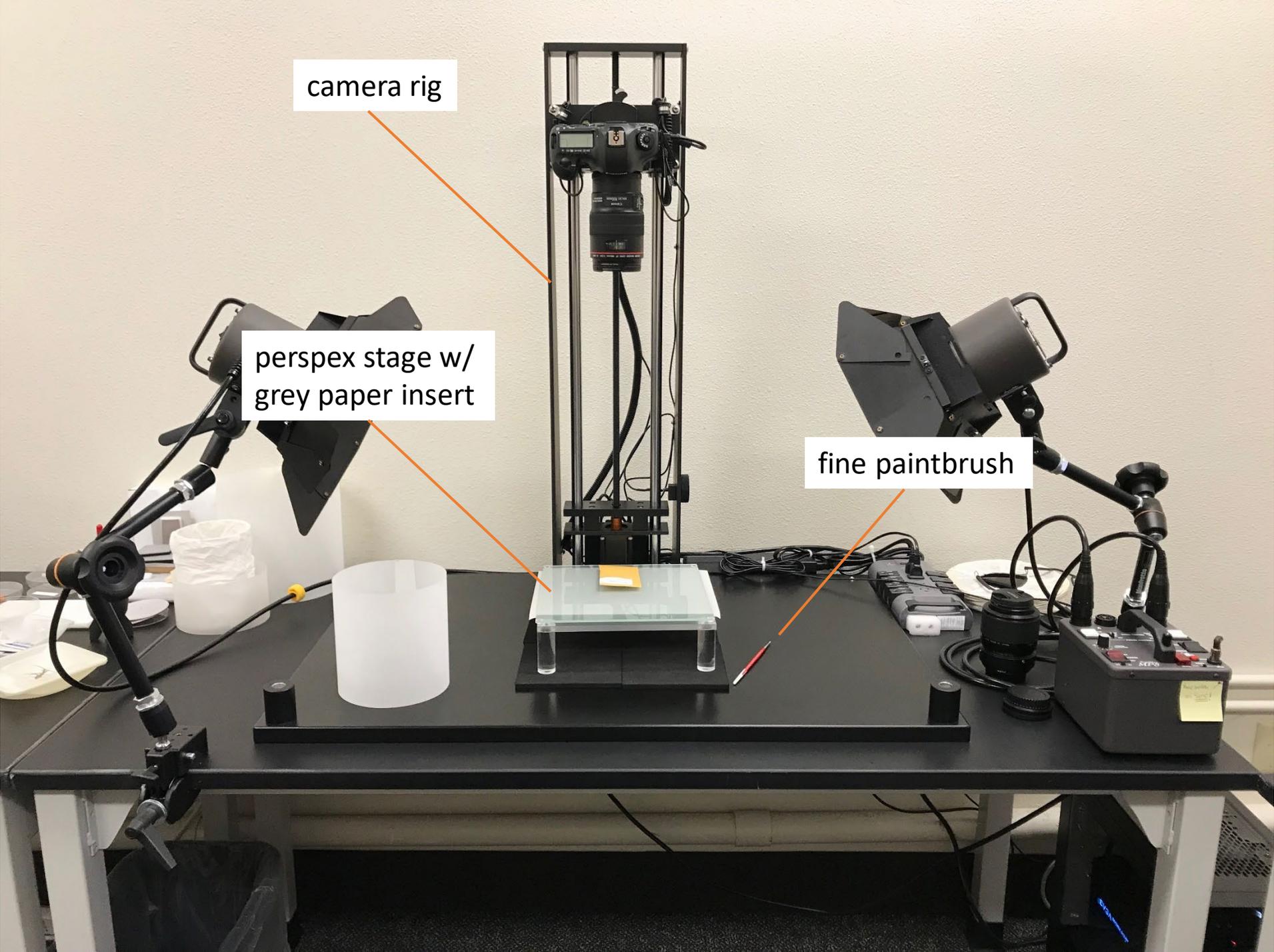
dental wax

putty

plastic boat with specimens

petri dish w/ glass inset

dust blower



camera rig

perspex stage w/  
grey paper insert

fine paintbrush

# Moderately-priced alternatives

Tagarno digital microscope  
(<\$1000 US?)



# Inexpensive alternatives

Phone-camera adapter for dissecting scope (~\$20-300, depending on lens quality)

Gosky (\$)



Labcam (\$\$\$)



# Inexpensive alternatives

Plugable USB microscope (~\$35-50)



**ITP**

USDA APHIS Identification Technology Program

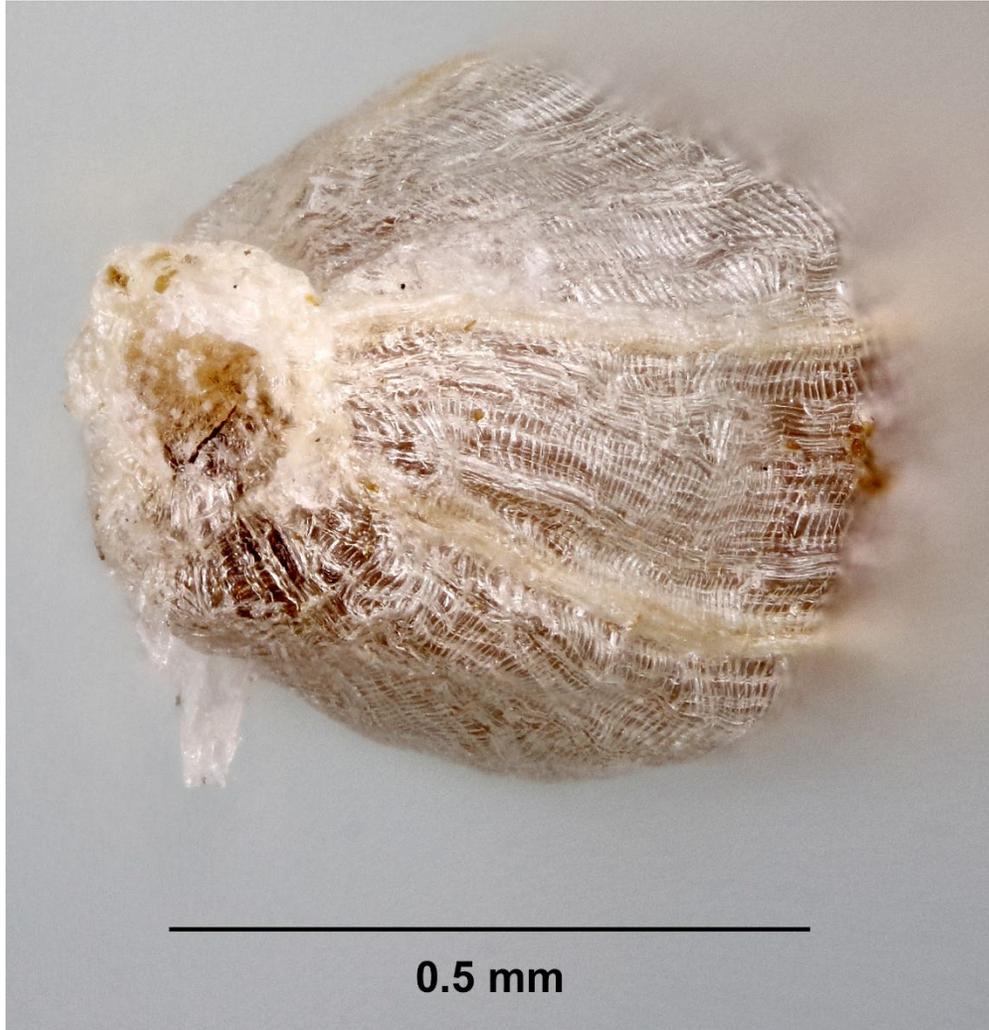
# Imaging with 65mm lens



# Imaging with 65mm lens



# Microscope imaging



# Microscope imaging



# Background color effects



# Defining terms

## ISO

brightness

ISO 100 = crisp image in good light

ISO 400+ = noise in dark parts of image, but can compensate for dark settings

## f-stop

size of aperture

f-stop 4.0 = larger aperture, more light allowed in

f-stop 8.0 = smaller aperture, less light allowed in

## shutter speed

faster (e.g., 1/500, 1/250) = less exposure, less opportunity for blur

slower (e.g., 1/15, 1/30) = more exposure, more opportunity for blur

Good image



No diffuser



# Incorrect white balance



Poor lighting



# Underexposure



# Overexposure



# Specimens moved during imaging



# Correct settings



Final image



# Lessons from our imaging lab

1. A clean environment and clean specimens mean less work later on.
2. Take care to set up the environment to minimize transfer of movement.
3. Time invested in setting up/testing the lighting and diffusion prior to imaging → time saved in post-processing.
4. All of this sounds simple, but it all takes practice to get right.

# Acknowledgements

Images credited to CFIA (Canadian National Seed Herbarium)

- Authors for the imaging protocol published in ISAM website
- \*QUAD Digital Identification Tool Team

Identification Technology Program staff at USDA-APHIS offices in Fort Collins, Colorado

Seed specimens from USDA-ARS labs, Nogales Plant Inspection Station (Dustin Sandberg), California Dept. of Food and Agriculture (Robert Price)

\*Quadrilateral Scientific Collaboration Working Group (the National Plant Protection Organizations (NPPOs) of Australia, Canada, New Zealand, and the United States).



Canadian Food  
Inspection Agency

Agence canadienne  
d'inspection des aliments

Canada



**United States Department of Agriculture**  
Animal and Plant Health Inspection Service

# Thank you!

