



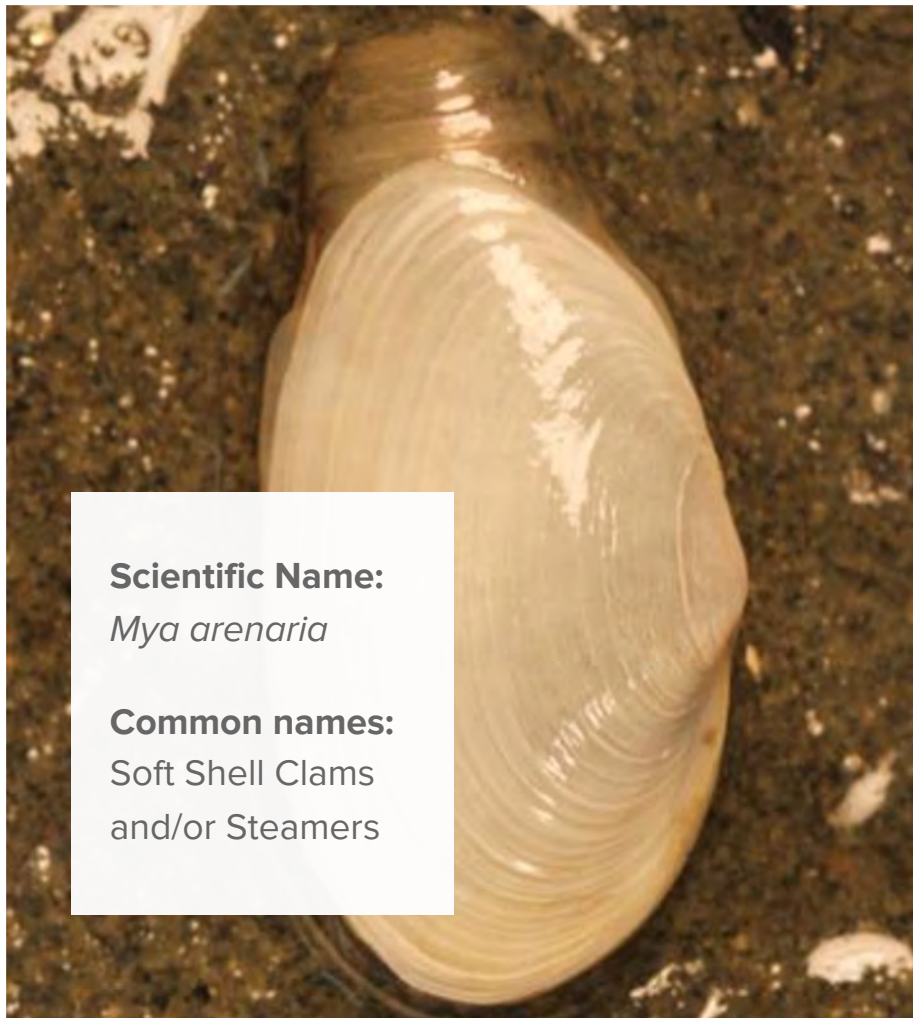
MORGANTM
PATUXENT ENVIRONMENTAL AND
AQUATIC RESEARCH LABORATORY

Pushing the Limits: Heat Tolerance in Soft Shell Clams

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The background of the image shows a close-up, top-down view of a stack of lumber. The wood grain is clearly visible, showing concentric growth rings. The entire image is overlaid with a semi-transparent blue filter. In the bottom-left corner, the word "Background" is written in a large, bold, white sans-serif font.

Background



Scientific Name:

Mya arenaria

Common names:

Soft Shell Clams
and/or Steamers



Distribution:

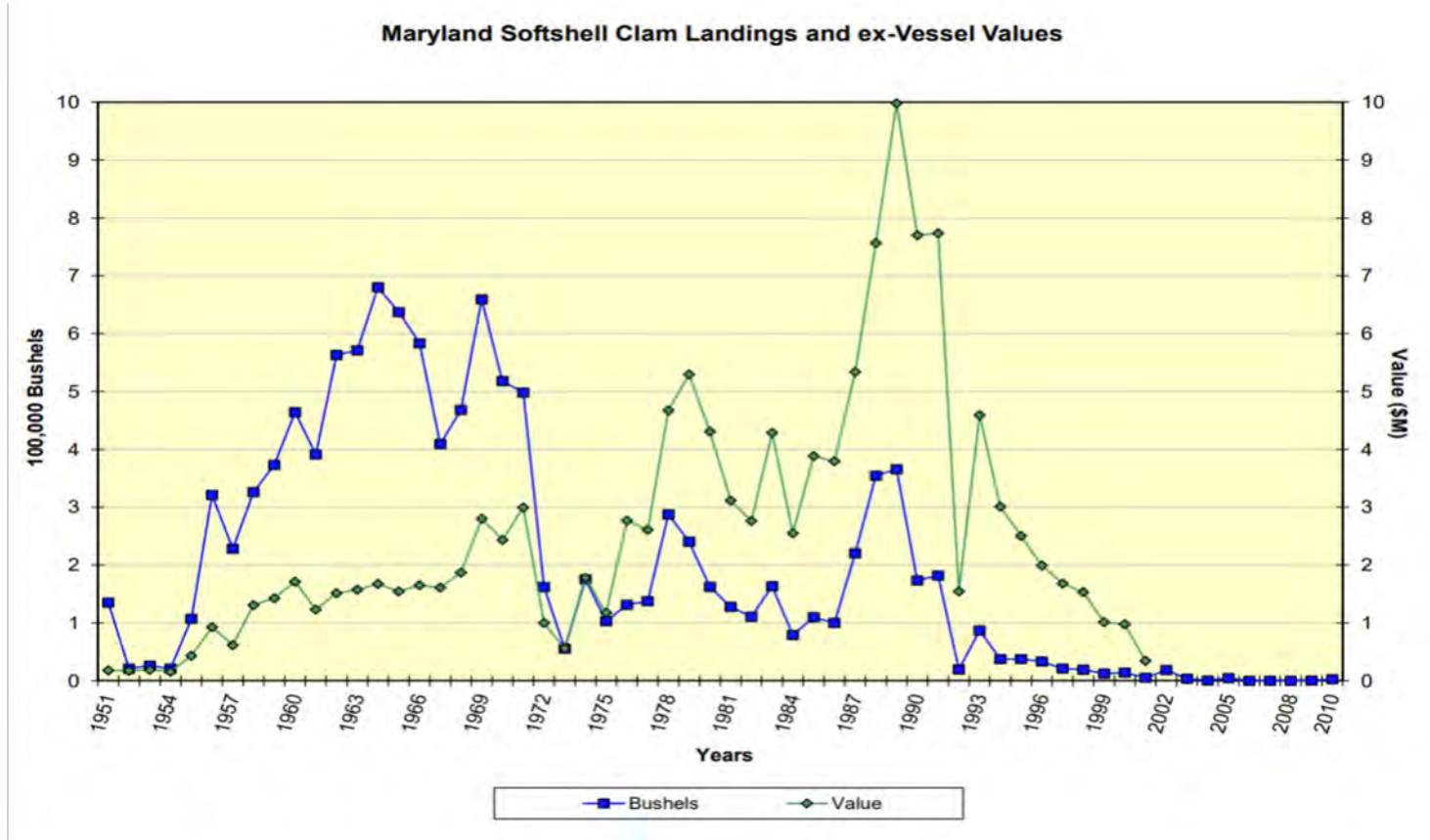
Maine to Virginia

Economic Value

- \$5-10 million annual industry in Maine (Beal, 2002)
- In 2019, the industry was valued at 54 million dollars with wholesalers, distributors, and others (Downeast Institute, 2020)



History in Maryland



Number of bushels and value of Soft Shell Clams in Maryland from 1951-2010

Potential for Maryland Aquaculture

Main species harvested/caught in the Chesapeake Bay are:

Oysters, Blue crabs, Atlantic Menhaden, etc. (Chesapeake Bay Foundation)

Soft Shell Clam Benefits:

- Create a new species in aquaculture
- More tolerant and grow faster in low salinity conditions than oysters
- Reaches commercial size in two years or less (Baker and Mann, 1991)



Number of Bushels of Oysters in the Chesapeake Bay from 1960-2006. This population decline due to overharvesting, disease, and pollution.

Challenges for developing Soft Shell Clam Cultures in MD

1. Seed production
2. Culture method
3. Heat stress

Heat Stress

**Can survive: -2C
to 28C**

**Optimal
temperature: 17C
to 23C**



Objectives

1. **Achieve breeding success: successfully spawned juvenile clams in April and we now have ~468K**
2. **Improve heat tolerance**
3. **Selective breeding based on phenotypic and genotypic breeding methods for heat tolerance**

Methods

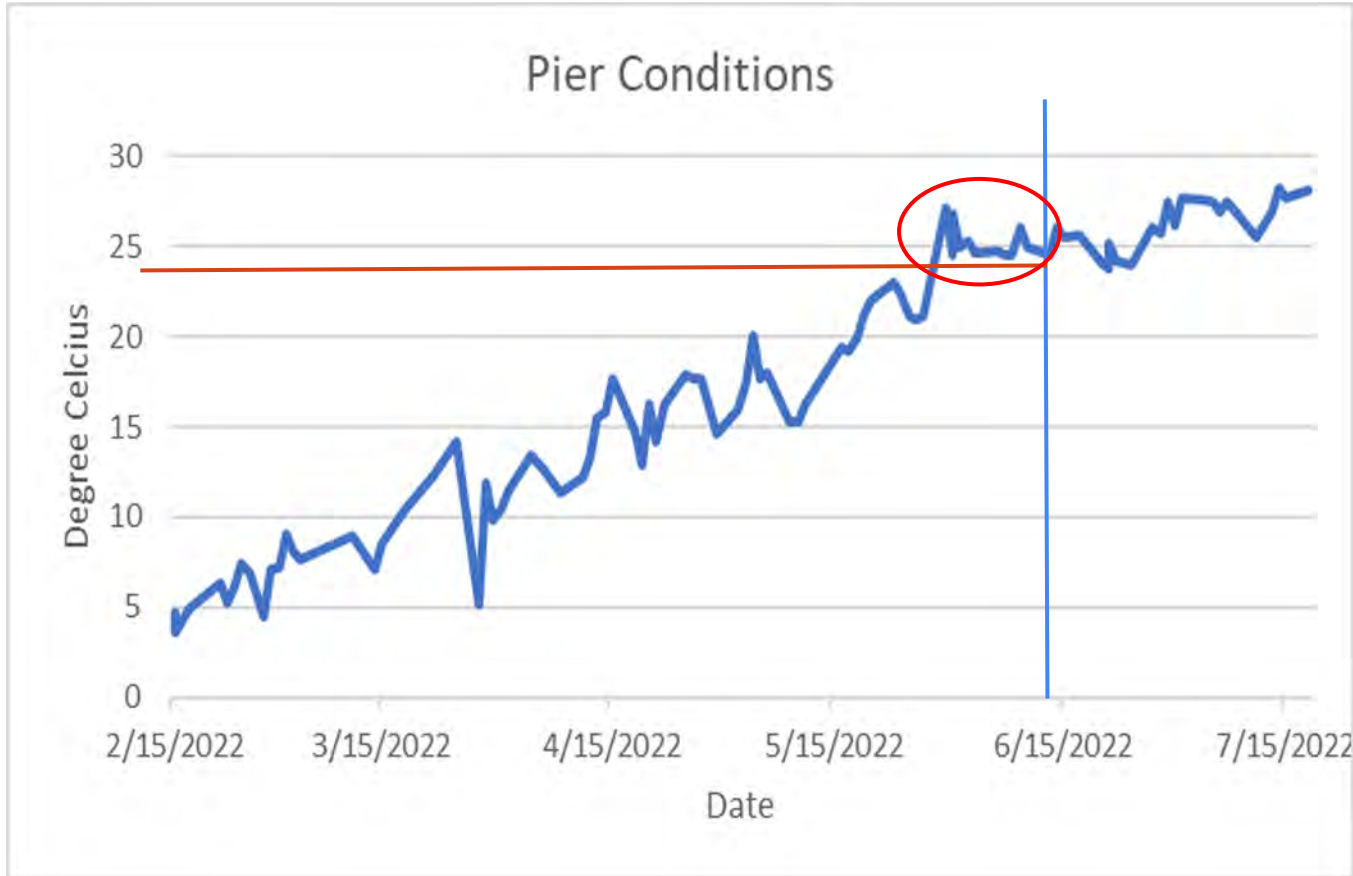
Selective Breeding Strategies

Phenotypic Approach

Genotypic Approach



Phenotypic Approach



Original Plan:
Compare 600 adult and juvenile

However, pier conditions spiked over the 17-23C optimal temperature before they were taken to the lab

Remaining number:

~75 adults
~150 juvenile

Genotypic Approach

Heat shock treatment

Heat shock a smaller amount of clams and sample the alive clams.

RNA sequence

Prepare the rna from the gills to send to a lab to be sequenced.

Identify heat shock genes

Receive sequence from lab and identify genes upregulated or downregulated. Validate these genes using RT PCR. Also, compare gene expression in different organs.

Identify Markers on Genes

Sequence top genes from heat shock or use target region sequencing to find markers of desired genes.

Marker associated selection

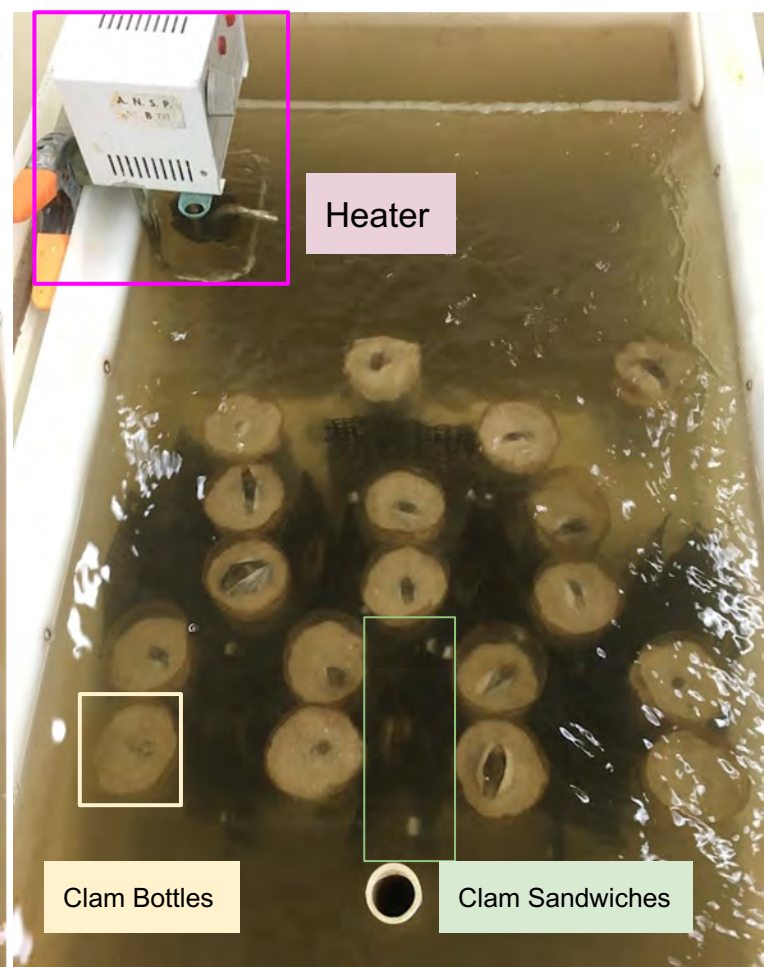
Use the markers to select clams to breed

Results

**For each BS tank:
18 clam bottles
18 sandwiches**



BS tank#1: Control ~19-23C



BS tank #2: Heat Stress ~30C

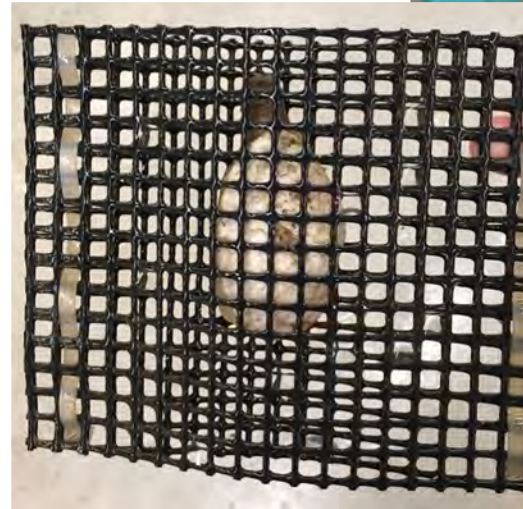
Bottle vs. Sandwich Debate

For Bottle:

- Sand could insulate and protect from challenging conditions
- Mimics the natural environment
- Prevents overcrowding
- Protects against accidents better

For Sandwich:

- Easier to setup and maintain
- Less prep work
- Can hold a higher density of clams in one place
- Do not have to deal with anoxic sand



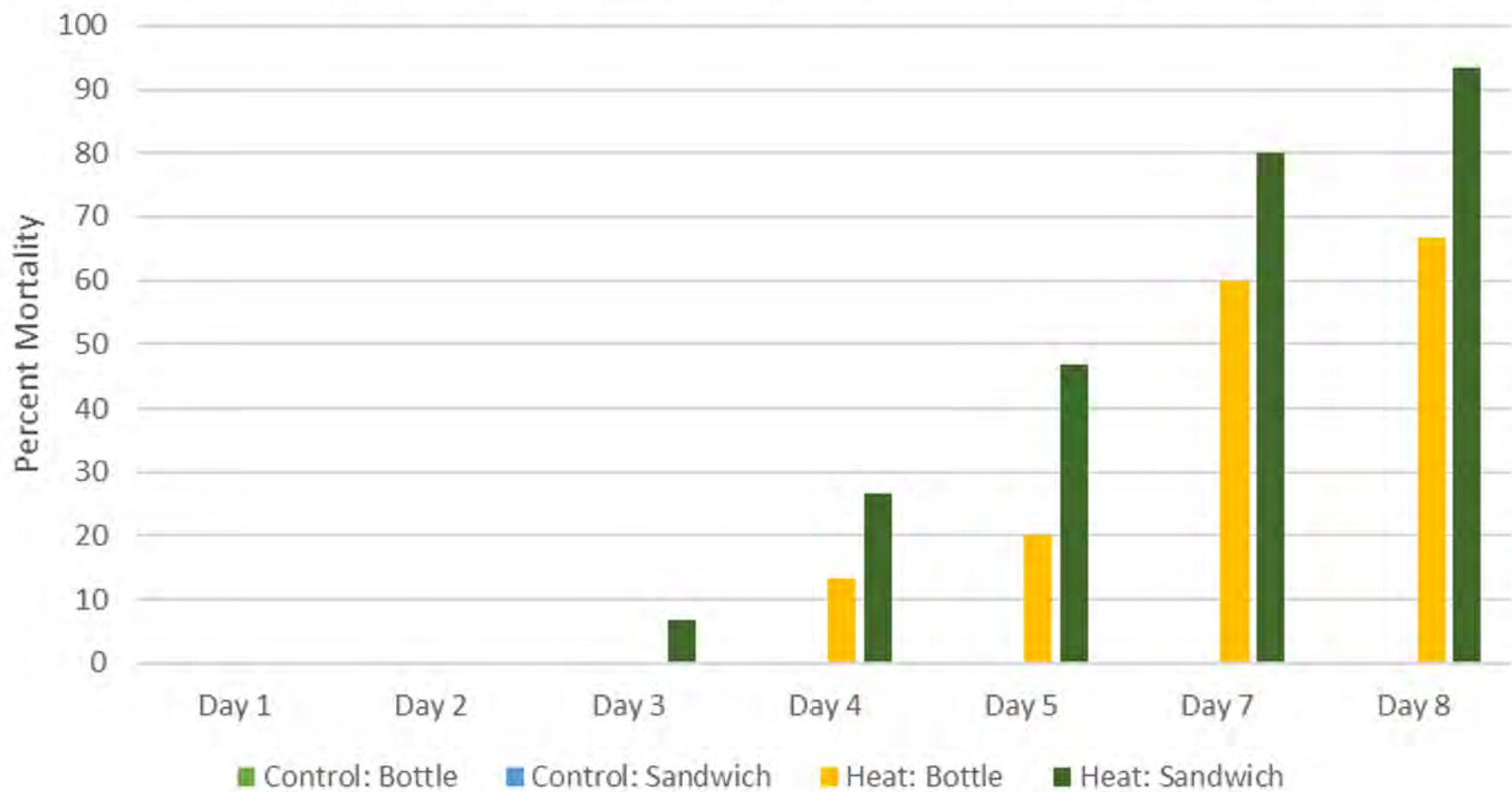
Observations

In Heat Stress Conditions:

1. More siphons were extended
1. Siphons were partially closed in most
1. Siphons responded slower to proding

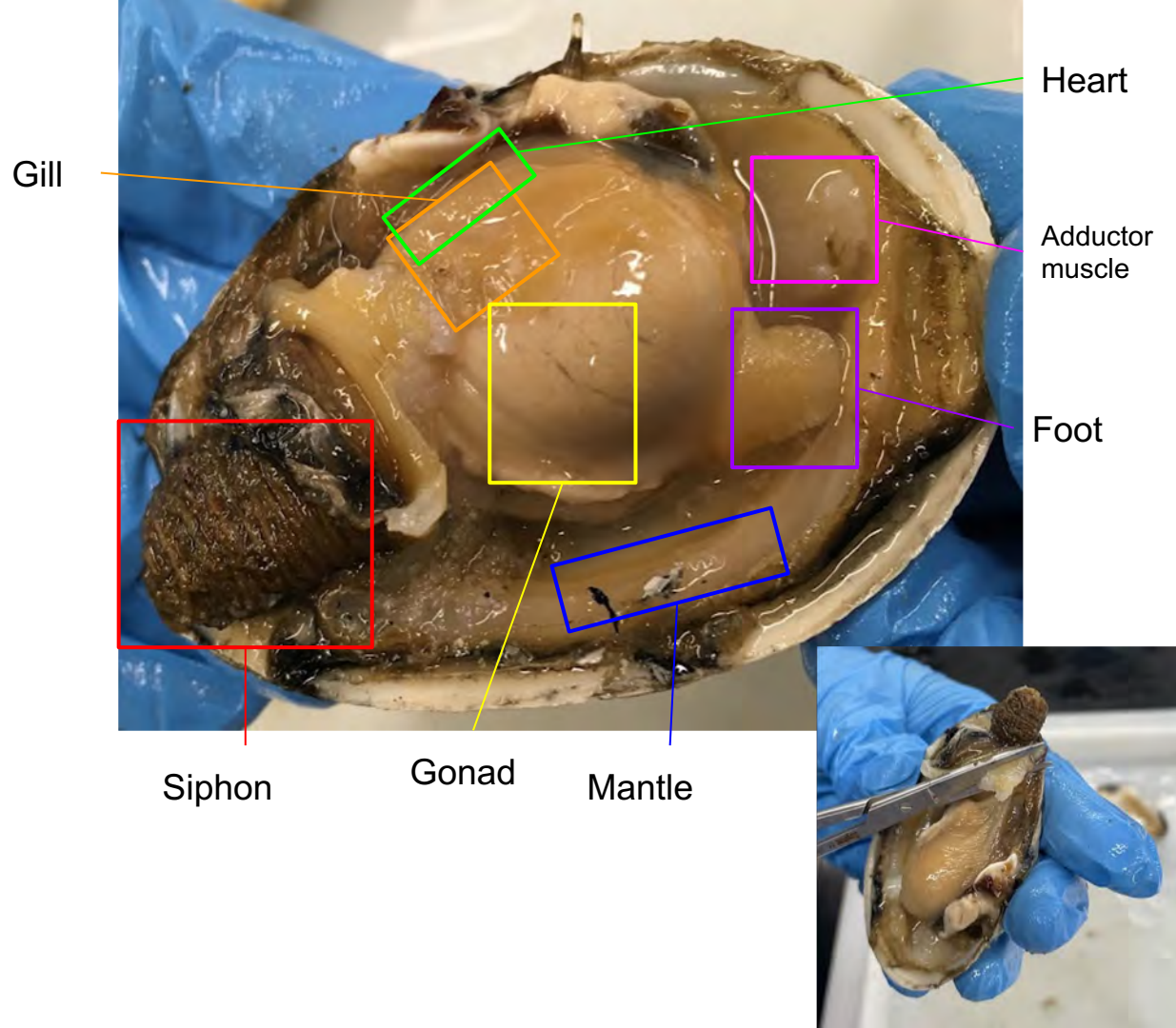


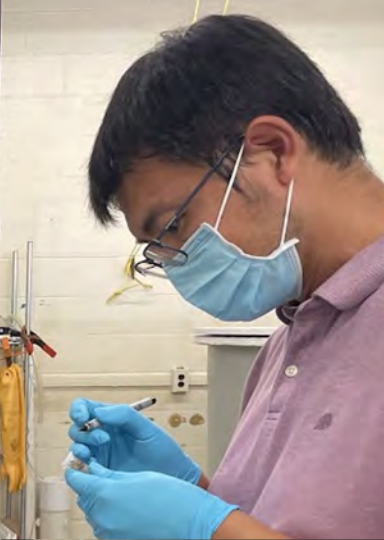
Percent Mortality in Soft Shell Clam Heat Shock Experiment



Sampling

- First death on 7/27/22
- Sampled 3 live clams from each group (total of 12 samples)
- **For Real Time PCR:** took samples from each gill, heart, adductor muscle, foot, mantle, gonad, and siphon
- **For RNA sequencing:** repeated gill samples





Conclusion

- **In Heat Stress:** more clams in sandwiches died
- **In Control:** no clams died, so both methods worked

Future Steps

Selective Breeding:

1. Send gills samples to lab to be RNA sequenced
2. Find these candidate genes and validate them using RT-PCR
3. Perform another heat shock with the 600 clams for phenotypic breeding strategy
4. Sequence the genes from phenotypic breeding group and their genetic markers
5. Use this data to spawn best individuals and run trials on those offspring

***Also continue research on Bottle vs. Sandwich**

Acknowledgements

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Development and Assessment of a Subtidal Grow-out and Method for Culture of Soft-shell Clams (*Mya arenaria*)

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Citations

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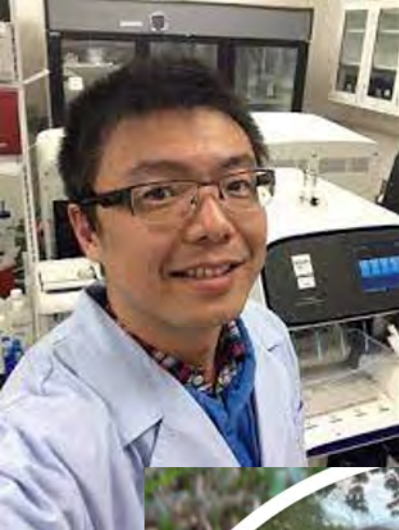
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