



Secrets of the Buttermilk Channel

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DNA LEARNING CENTER

Species Richness of Macroinvertebrates in New York Harbor: Buttermilk Channel

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Abstract

The purpose of this research was to learn how to extract DNA from invertebrates, and to discover the species richness of the Buttermilk Channel. Another goal of this research project was to compare the effectiveness of the Folmer and Leroy primers. We found that Buttermilk Channel is not very diverse but with more and more awareness it may become so.

Hypothesis

We hypothesized that there would be little species richness because of the history of dredging and heavy boat traffic in the Buttermilk Channel.

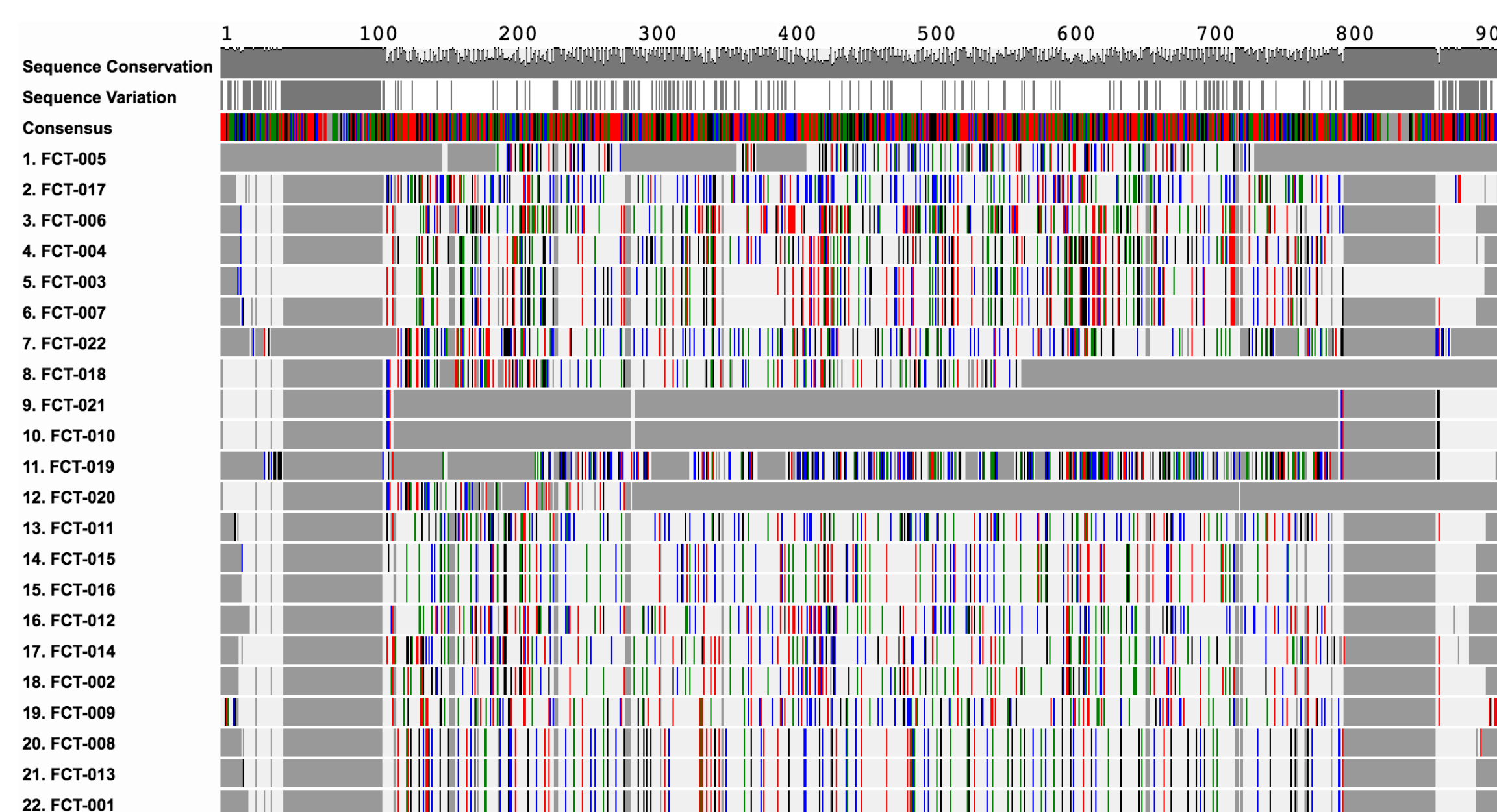
Introduction

Invertebrates are one of the oldest and most diverse group of organisms (Alroy, Aberhan, and Bottjer, 2008). The biodiversity in the Hudson-Raritan estuary is a key factor in creating a healthy and sustainable ecosystem. The biodiversity index is an overarching topic that includes; biodiversity, species richness and species evenness.

Results

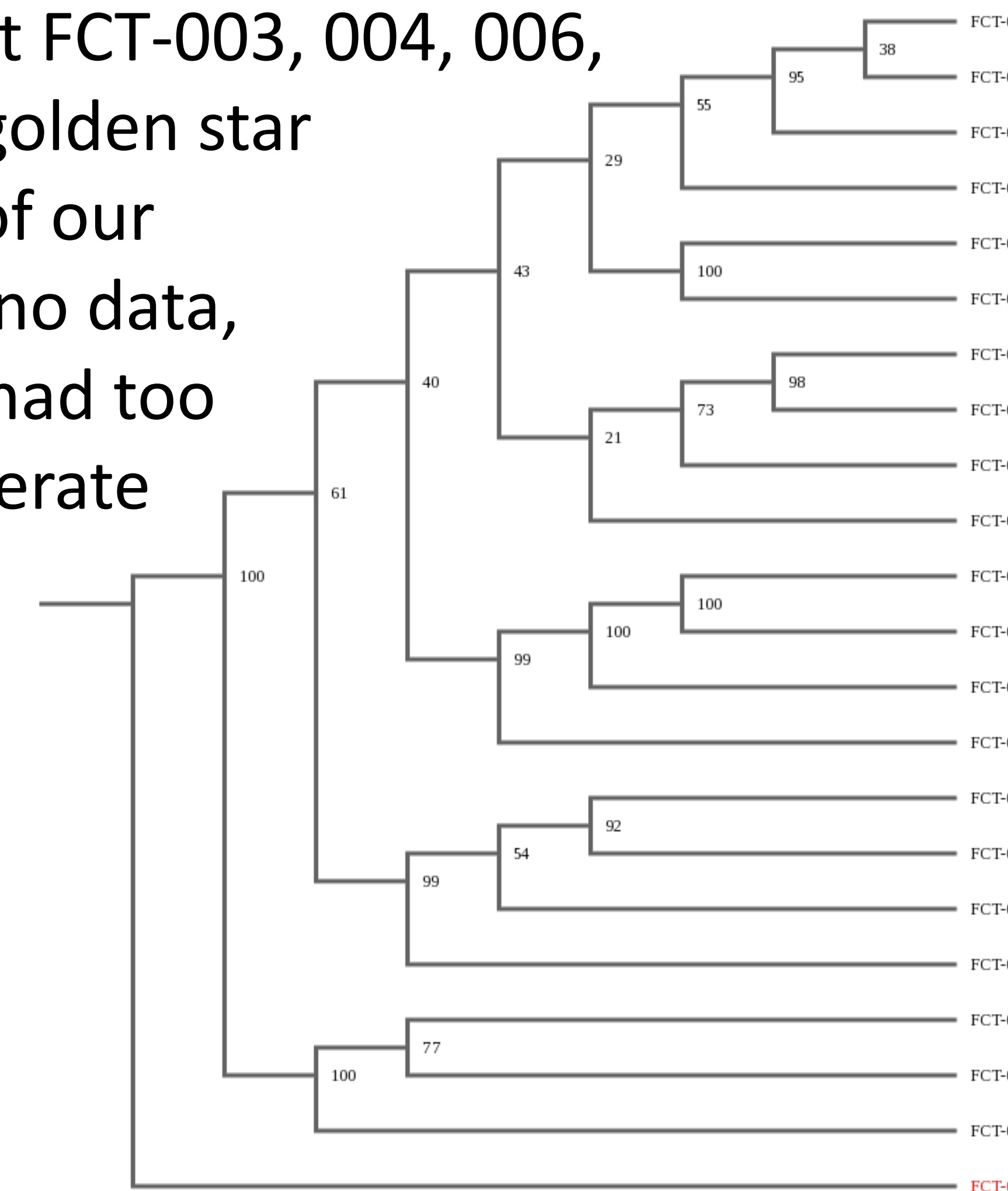
As you can see in the consensus chart, samples with similar base pair patterns were grouped together. We found that FCT-003, 004, 006, and 007 were most likely *Botryllus schlosseri* (golden star tunicate) and closely related species. In many of our samples there are large chunks where there is no data, this is likely due to human error. If the sample had too many missing gaps, the program could not generate a possible species and was left blank, this was the case with FCT-010 and 021.

Anemabuddies Gel 1	Folmer et al Primer 700bp	Comments	Leroy Primer 400pb	Comments
FCT-001	<i>Halichondria panicea</i>	sponge	FCT-001	<i>Halichondria panicea</i>
FCT-002	<i>Urosalpinx cinerea</i>	snail	FCT-002	<i>Urosalpinx cinerea</i>
FCT-003	<i>Botryllus schlosseri</i>	golden star tunicate		no PCR product
FCT-004	<i>Botryllodes vesiculosus</i>	colonial tunicate	FCT-004	<i>Botryllodes vesiculosus</i>
FCT-005	<i>Pione</i> sp./ <i>Stephanaugea nuxilis</i> (?)	many mismatches	FCT-005	<i>Bonnewella regia</i>
FCT-006	<i>Molgula manhattanensis</i>	Sea Grape Tunicate/Tree squirt	FCT-006	<i>Hadra vulgatis</i>
FCT-007	<i>Botryllus schlosseri</i>	golden star tunicate	FCT-007	<i>Botryllus schlosseri</i>
FCT-008	<i>Halichondria panicea</i>	breadcrumb sponge	FCT-008	<i>Cladomeres linata</i>
FCT-009	<i>Ectopleura crocea</i>	In Hydrozoans genus	FCT-009	<i>Botryllodes vesiculosus</i>
FCT-010	No Match	match on GenBank: <i>Cyanea</i> sp. (jellyfish?) questionable	FCT-010	<i>Amphioxe valida</i>
FCT-011	<i>Amphioxe valida</i>		FCT-011	<i>Lepidonotus squamatus</i>
FCT-012	<i>Lepidonotus squamatus</i>		FCT-012	<i>Halichondria panicea</i>
FCT-013	<i>Halichondria panicea</i>	breadcrumb sponge	FCT-013	<i>Crepidula fornicata</i>
FCT-014	<i>Crepidula fornicata</i>		FCT-014	<i>Amphibalanus</i> sp.
FCT-015	<i>Amphibalanus</i> sp.	barnacle		
FCT-016	<i>Amphibalanus</i> sp.	barnacle	FCT-016	<i>Amphibalanus</i> sp.
FCT-017	<i>Parasabella microphthalmia</i>		FCT-017	<i>Parasabella microphthalmia</i>
FCT-018	<i>Jassa maronata</i>	amphipod	FCT-018	<i>Jassa maronata</i>
FCT-019	<i>Stenotrophomonas</i> sp. / vibrio	bacteria- contamination/ often found in brackish saltwater	FCT-019	<i>Halichondria panicea</i>
FCT-020	<i>Lepidonotus squamatus</i>		FCT-020	no seq available
FCT-021	No Match	match on GenBank: match on GenBank: <i>Cyanea</i> sp. (jell)	FCT-021	<i>Zeuxis squicrensis</i>
FCT-022	<i>Monocotylus insidiosus</i>	Amphipodal amphipod	FCT-022	<i>Stenomeres thysax</i>



Methods

First we needed to isolate the DNA, add a series of wash buffers, add the primer, put it through a polymerase chain reaction machine, and send the copied DNA to a bioinformatics lab. After getting the sequences back we used a program called DNA Subway to analyze the results.



As for the Folmer and Leroy primers; the Leroy primer produced species names when the Folmer primer could not, however the same thing happened when the Leroy primer had no answer. We did not have the opportunity to accurately test whether these results were correct.

Discussion

The phylogenetic tree (fig. 1), shows the evolutionary relationships between all our samples and many of them are very closely related. In the consensus chart (fig. 2) the closer a species' sequence is to another, the more likely they are to be related. We had a fairly low rate of biodiversity, which supported our hypothesis, this is likely due to the extreme dredging in the 1900s. In regards to the different primers, the Leroy primer did come up with possible species names where the Folmer primer did not. However, on some samples we knew were amphipods the Leroy gave a result for a sponge (FCT-019). Overall I would recommend using the Leroy primer because it gives more results that appear to be accurate more of the time.

References

- Advances in DNA metabarcoding for food and wildlife forensic species identification, 2016
- A new versatile primer set targeting a short fragment of the mitochondrial COI region for metabarcoding metazoan diversity: application for characterizing coral reef fish gut contents, 2013