

# Assessing Arctic Grayling Relative Abundance and Distribution Through Environmental DNA Analysis

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*We acknowledge the Alaska Native nations located on the traditional lands of the Dena people of the lower Tanana River from which we collected our environmental samples from.*

## Grayling and eDNA

- Arctic Grayling (*Thymallus arcticus*) freshwater species
  - Broad distribution across Alaska
  - Opportunistic feeders; wide diet variety
  - Follow spawning salmonid to feed
  - Not an important species for subsistence harvest
  - Popular for sport fishing; various colors and sizes
- Environmental DNA analysis
  - Potential source of information of distribution, abundance, and ecology
  - qPCR (quantitative Polymerase Chain Reaction) strategy for extracted DNA to determine relative amount of species specific DNA
- **Will we find an abundance of Arctic Grayling DNA in the Chena River during the salmon spawning season?**



Image 1. Greg Beutler fly fishes for Arctic Grayling off Chena Hot Springs Road, close to the North Fork sample site.  
Photo: John Fillmore

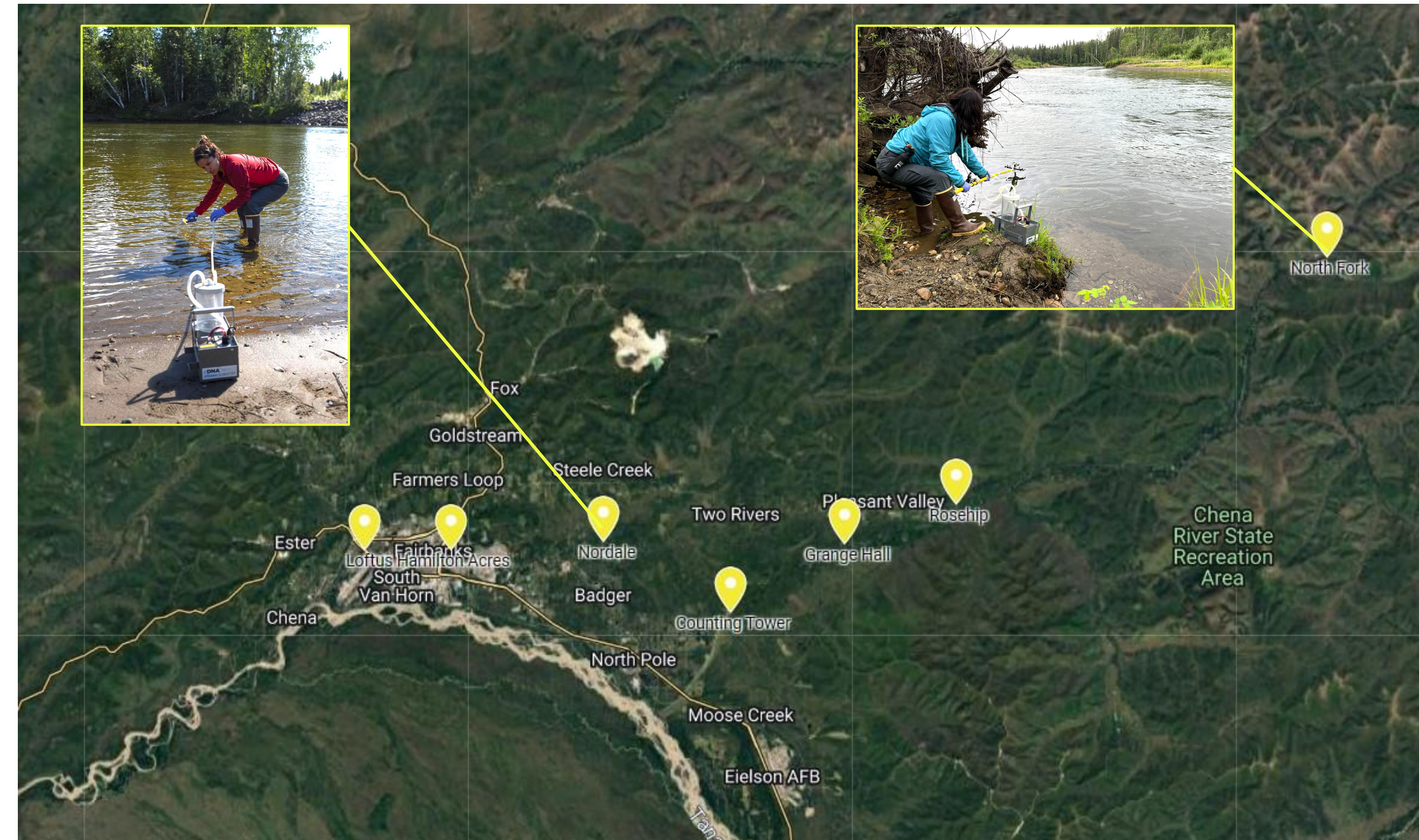


Image 2. Chena River Sample Sites: North Fork, Rosehip, Grange Hall, Moose Creek Counting Tower, Hamilton Acres, Loftus and Nordale.  
Photos: Maggie Harings, UAF



Image 3. Water sample filters before eDNA extraction



Image 4. Extracting DNA from filters  
Photo: Alice Bailey, UAF

Table 1. DNA from tissue samples

<i>Thymallus arcticus</i>	Arctic Grayling
<i>Coregonus autumnalis</i>	Arctic Cisco
<i>Oncorhynchus keta</i>	Chum Salmon
<i>Prosopium cylindraceum</i>	Round Whitefish
<i>Dallia pectoralis</i>	Alaska Blackfish
<i>Cottus cognatus</i>	Slimy Sculpin
<i>Oncorhynchus tshawytscha</i>	Chinook Salmon
<i>Oncorhynchus mykiss</i>	Rainbow Trout/Steelhead
<i>Coregonus sardinella</i>	Least Cisco

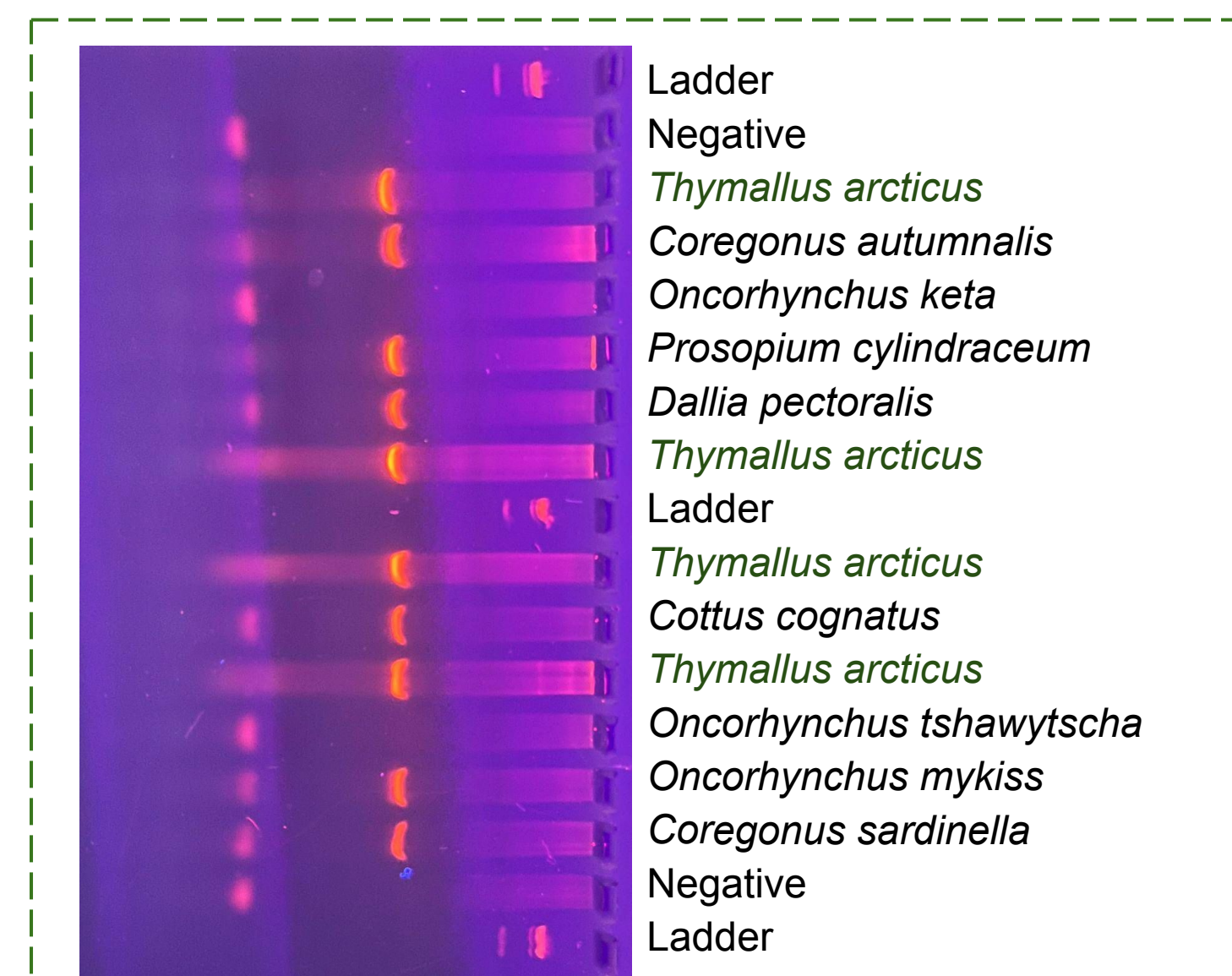


Image 5. Gel electrophoresis from PCR

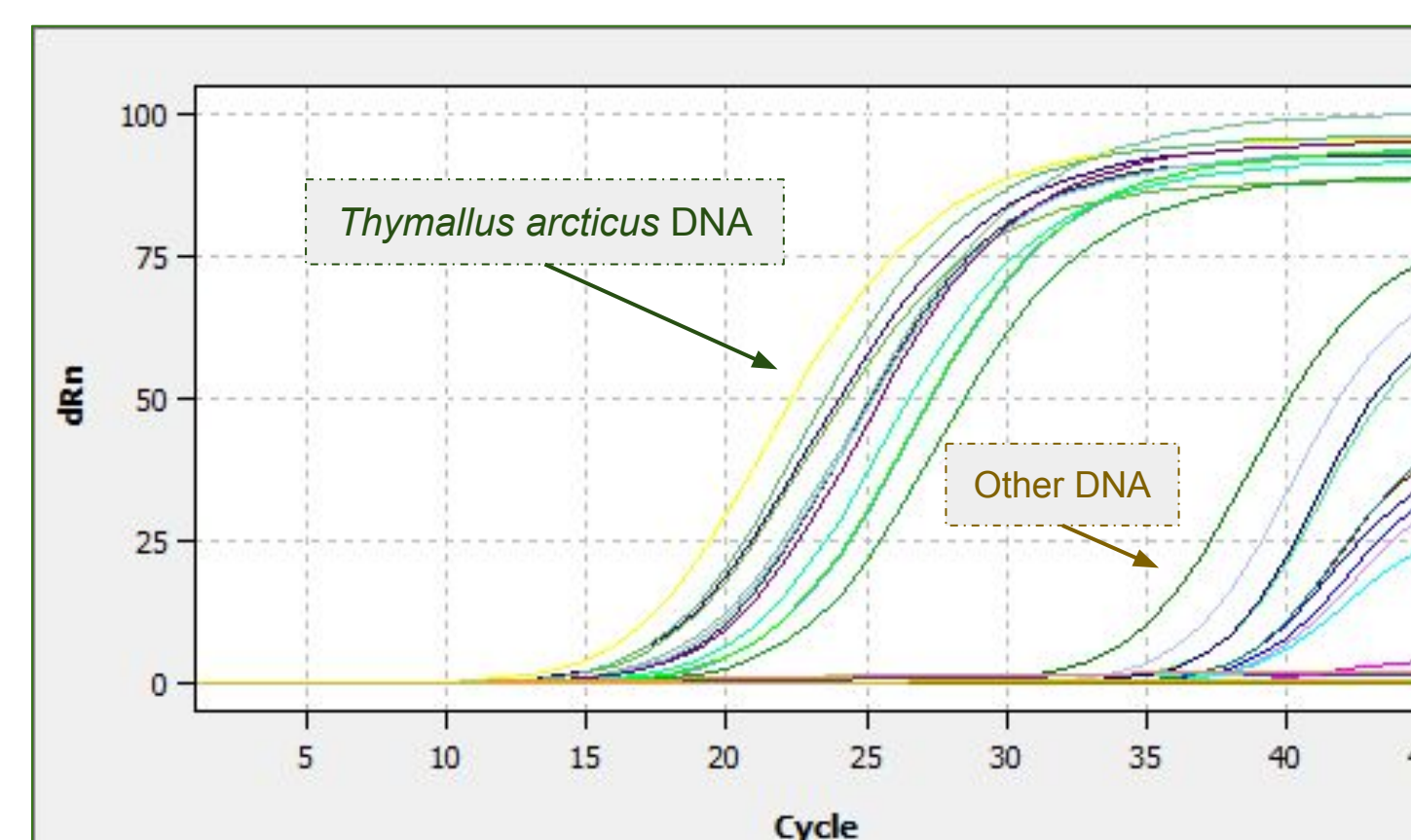


Figure 1. qPCR with tissue sample DNA

Species	Ct Values
<i>Thymallus arcticus</i>	14.97 - 17.31
<i>Oncorhynchus keta</i>	38.1
<i>Oncorhynchus mykiss</i>	38.65
<i>Oncorhynchus tshawytscha</i>	36.52
<i>Prosopium cylindraceum</i>	32.95

## Preliminary Analysis

- Tested assay specificity with tissue samples from Arctic Grayling and closely related/unrelated species (Table 1).
- High DNA concentrations of most salmonids indicate amplification during PCR (Image 5) and qPCR (Figure 1 & Table 2).
- qPCR indicates DNA concentrations of Chena River samples too low to amplify (Figure 2).

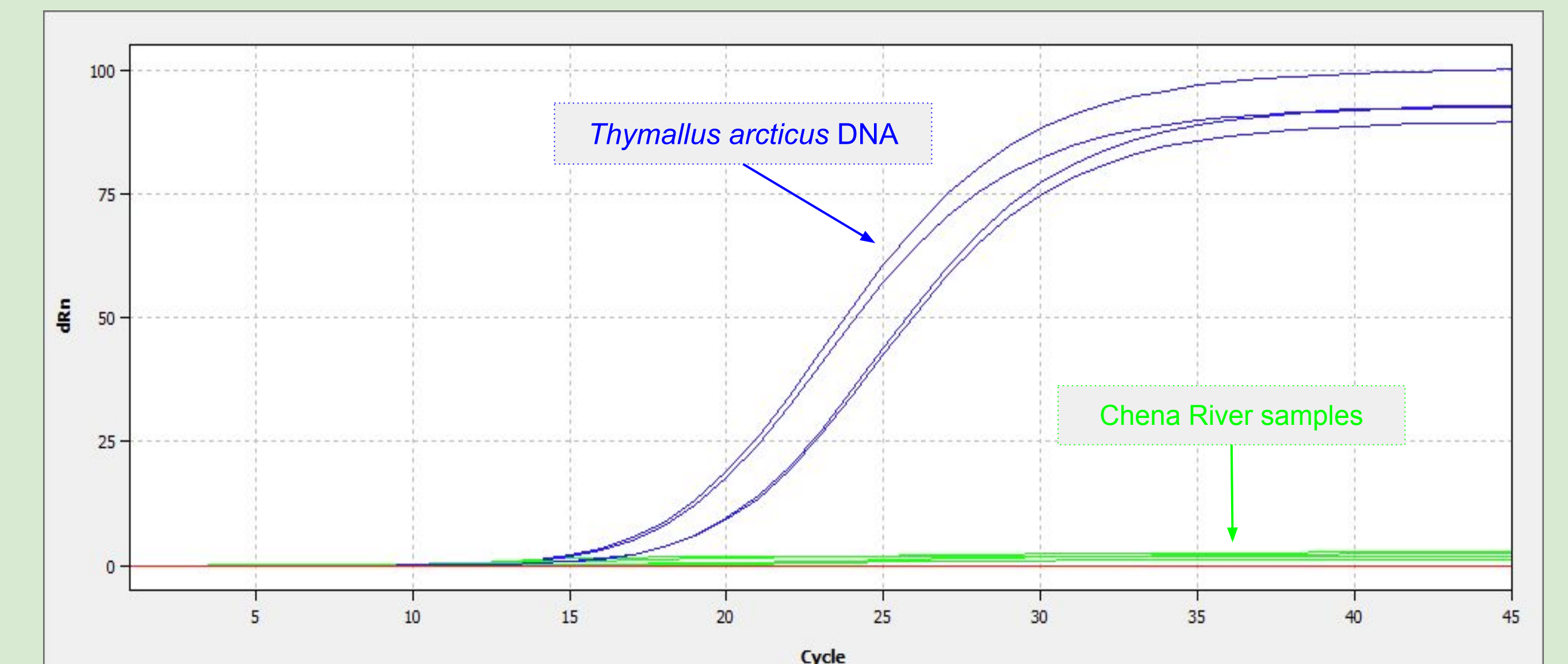


Figure 2. qPCR results of Arctic Grayling vs. Chena River eDNA samples.

## Discussion

- Latest qPCR analysis indicates eDNA sample concentrations are too low to amplify.
- Next Steps:
  - Further testing assay (Rodgers, et al) with known DNA concentrations to develop level of detection for Arctic Grayling.
  - Data configured in relation to Chum and Chinook salmon eDNA spawning periods
- qPCR method may become a complementary technique in the field of assessing relative quantities of species over time and space.

## Sources

Rodgers, T.W., Olson, J.R., Kloubucar, S.L. et al. Conservation Genet Resour (2018) 10: 859. <https://doi.org/10.1007/s12686-017-0883-1>

Schoen, E. R., Sellmer, K. W., Wipfli, M. S., López, J. A., Ivanoff, R., & Meyer, B. E. (2022). Piscine predation on juvenile salmon in sub-arctic Alaskan rivers: Associations with season, habitat, predator size and streamflow. *Ecology of Freshwater Fish*, 31, 243–259. <https://doi.org/10.1111/eff.12626>

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## Methods

- Collected water samples with Citizen Science Sampler
  - 7 locations on 3 separate days: 3 samples each sample site
- Isolated DNA from filters
- Prepared DNA for quantification
- Processed species specific assay with PCR
- Processing samples through qPCR machine (qTOWER3 84)