Fish eDNA

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2024 - 10 - 16

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## Chapter 1

## Our dataset

I was given the following dataset of fastq files which were generated by (1) collecting 18 water samples from rivers and lakes near oil-sands extraction site in the province of Alberta, Canada (see map below), (2) filtering the water samples, (3) extracting the DNA collected on the filer, and (4) sequencing the extracted DNA. For each sample collected, two different marker genes were used to amplify and sequence distinct DNA regions: the **12S** ribosomal RNA gene and the mitochondrial cytochrome oxidase subunit (**COI**).

## Amplified regions details

Marker	Names	Forward primer sequence	Reverse primer sequence	Fragment size (bp)	Source
128	MiFish	-CCGGTAAAACTC	CACAGAGATGC	TOGTAATCCCA	<b>©DDI</b> :TG
	U				%5B10.
					1111/
					mec.
					14395](https
					//
					doi.
					org/
					10.
					1111/
					mec.
					14395)

Marker	Names	Forward primer sequence	Reverse primer sequence	Fragment size (bp)	Source
COI	COI-	CGTATTTGGYGC	CYCHORRACARCOTYRATE		OI:
	PS1				%5B10.
					1098/
					rsos.
					150088](htt
					//
					doi.
					org/
					10.
					1098/
					rsos.
					150088)