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Results Information

Customer ID: Adrian MI

Project #: P-474

Sample Date Received: 12/20/18 Date of Analysis: 12/20/18 - 12/21/18

Sample Type: 4 Tap Water

Analysis Type: Cyanobacteria ID

Microcystin Toxin Gene ID General Bacteroides ID

Sample Information:

Four tap water samples collected on December 19, 2018 were submitted for analysis. Sample numbers and their corresponding locations are listed below:

Sample 1 –
Sample 2 –
Sample 3 –
Sample 4 –

Analysis Description:

The analysis of the submitted water samples involved determining the presence/absence of Cyanobacteria, the Microcystin producing toxin gene and general Bacteroides bacteria in each of the four samples using real time PCR analysis.

Procedure: DNA Extraction

500 ml of each submitted water sample was vacuum filtered through a 0.45 micron filter membrane that proceeded to capture any microorganisms present in the water sample. Following vacuum filtration, the individual filter membranes were removed from the filter column and microorganism DNA was extracted from each filter membrane using the Generite DNA EZ ST1 DNA extraction kit. The DNA extractions were used in all subsequent analysis.

<u>Procedure: PCR Analysis Cyanobacteria, Microcystin Producing Toxin</u> Gene and General Bacteroides Bacteria

Qualitative PCR analysis to determine the presence/absence of Cyanobacteria, the Microcystin producing toxin gene and general Bacteroides bacteria incorporated oligonucleotide primers that target specific DNA marker sequences. The PCR targets and the location of the specific DNA markers for each target are listed in the table below:

Target

Cyanobacteria Genera Microcystin Toxin Gene General Bacteroides Bacteria

Location of DNA Marker

Cyanobacteria nif Gene Microcystin Synthetase A Gene Bacteroides 16S ribosomal DNA Gene

Real Time PCR analysis was performed using the Stratagene Mx 3005P Real Time PCR System instrument. PCR analysis incorporated oligonucleotide primers that target specific marker DNA sequences unique to Cyanobacteria, the Microcystin producing toxin gene and general Bacteroides bacteria. Appropriate positive and negative control samples were included in PCR experimentation. Replicates of each sample were run for quality assurance purposes and to confirm the observed results.

Following the PCR procedure, the PCR products were run on a 1.5 % agarose gel at 100V for 60 minutes, after which the agarose gel containing the PCR products was visualized under UV light and examined for PCR amplification products that represent the amplification of the target DNA markers.

Results:

Results of the qualitative PCR testing indicate the presence (positive) or absence (negative) of the specific DNA markers for Cyanobacteria, the Microcystin producing toxin gene and general Bacteroides bacteria through PCR amplification of the specific DNA marker sequence.

A sample testing positive for a specific DNA marker confirms the presence of the organism or toxin gene in the sample. A sample testing negative for a specific DNA marker confirms the absence of the organism or toxin gene in the sample.

| <u>Sample</u> | <u>Cyanobacteria</u> | Microcystin Toxin Gene | Bacteroides Bacteria |
|---------------|----------------------|------------------------|-----------------------------|
| 1 – | negative | negative | positive |
| 2 – | positive | negative | positive |
| 3 – | negative | negative | negative |
| 4 – | negative | negative | negative |

These results were verified by Tom Prychitko, Laboratory Director for Helix Biological Laboratory.

For the Contractor (Signature)