

### Certificate of Analysis

Laboratory Reference: 240112-017

<b>Attention:</b> News .	<b>Final Report:</b> 536307-0
<b>Client:</b> VEOLIA WATER	<b>Report Issue Date:</b> 16-Jan-2024
<b>Address:</b> PO Box 761, Thames, 3540	<b>Received Date:</b> 13-Jan-2024
<b>Client Reference:</b> Hahei Estuary	<b>Laboratory Activity Dates:</b> 13-Jan-2024 - 15-Jan-2024
<b>Purchase Order:</b> 7300164040	<b>Quote Reference :</b> 5949

#### Sample Details

<b>Lab Sample ID:</b>	240112-017-1
<b>Client Sample ID:</b>	
<b>Sample Date/Time</b>	12/01/2024 14:45
<b>Description:</b>	Estuary

#### General Testing

Ammoniacal Nitrogen (as N)	mg/L	0.58
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#### Microbiology

##### Enterococci by Membrane Filtration

Enterococci	cfu/100 mL	31
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##### Escherichia coli by Membrane Filtration

Escherichia coli	cfu/100 mL	150
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*Results marked with \* are not accredited to International Accreditation New Zealand. A dash indicates no test performed.*

*Where samples have been supplied by the client, they are tested as received.*

*The results of analysis contained in this report relate only to the sample(s) tested. Where sample collection was performed by the laboratory, the results of analysis contained in this report relate only to the sample(s) collected.*

#### Reference Methods

The sample(s) referred to in this report were analysed by the following method(s)

Analyte	Method Reference	MDL	Samples	Location
<b>General Testing</b>				
Ammoniacal Nitrogen (as N) by Flow Analysis	APHA (online edition) 4500-NH3 H	0.005 mg/L	All	Auckland
<b>Microbiology</b>				
<b>Enterococci by Membrane Filtration</b>				
Enterococci	APHA (online edition) 9230 C	2 cfu/100 mL	All	Auckland
<b>Escherichia coli by Membrane Filtration</b>				
Escherichia coli	USEPA Method 1603	1 cfu/100 mL	All	Auckland

*The method detection limit (MDL) listed is the limit attainable in a relatively clean matrix. If dilutions are required for analysis the detection limit may be higher.  
 For more information please contact the Compliance and Projects Manager.*

Samples, with suitable preservation and stability of analytes, will be held by the laboratory for a period of two weeks after results have been reported, unless otherwise advised by the submitter.

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