

Eastland Port Matawhero Logyard Sampling Report – August 2015

For Eastland Port Limited

September 2015

REPORT INFORMATION AND QUALITY CONTROL

Prepared for

Eastland Port Limited

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1 INTRODUCTION

Under consents DW-2011-104235-01, WP-2011-104234-01, and WP-2011-104292-01 stormwater and groundwater monitoring is required.

For the stormwater monitoring two monthly sampling is undertaken at three locations; the stormwater retention ponds culvert outlets, the Awapuni Drain 10 metres downstream of the confluence with the tributary drain, and 10 metres upstream of the confluence with the tributary drain. For the groundwater monitoring, six monthly sampling, during February and August, is undertaken at three locations; two groundwater monitoring bores and the sump tile drainage outlet. Refer to Appendix A for a plan of the sample sites.

Both the stormwater and groundwater sampling was undertaken on 7 August 2015. The stormwater samples represent the July/August sampling round. The groundwater samples represent the August sampling round.

Sampling was undertaken in accordance with the Sampling Protocols and Standard Operating Procedures prepared by 4Sight Consulting (formerly known as Andrew.Stewart Ltd). The sampling was undertaken by Logic Forest Solutions.

This report has been prepared for Gisborne District Council and provides the results and analysis of the Matawhero Logyard August 2015 stormwater and groundwater sampling rounds.

This is the third sampling report for stormwater sampling. The last report, titled "Eastland Port Matawhero Logyard Sampling Report – July 2015", was prepared for the May/June sampling round.

This is the second sampling report for groundwater sampling. The first report, titled "Eastland Port Sampling Results Report", was prepared for the February sampling round undertaken in March 2015. The first report also included sampling results for Southern Logyard and Wairakaia Bark Disposal site.

2 SAMPLING DETAILS

Table 1: Stormwater Sample time and dates

Location	Date	Time
Stormwater retention ponds culvert outlets on the river side of the railway track	07/08/2015	8:45am
Awapuni Drain 10m downstream	07/08/2015	8:29am
Awapuni Drain 10m upstream	07/08/2015	8:08am

Table 2: Groundwater Sample time and dates

Location	Date	Time
Monitoring Bore 1 (MLY GW01)	07/08/2015	9:40am
Monitoring Bore 2 (MLY GW02)	07/08/2015	10:32am
Sump tile drainage outlet	07/08/2015	8:59am



2.1 Relevant Site Information

Table 3: Stormwater Sample Information

Location	Rainfall event	Number of dry days prior to sampling	Discharge/water colour	Obvious or visual features
Stormwater retention ponds culvert outlets on the river side of the railway track	16.8mm	0	Clear	No debris or scums/foams present
Awapuni Drain 10m downstream	16.8mm	0	Slightly yellow	No debris or scums/foams present
Awapuni Drain 10m upstream	16.8mm	0	Slightly yellow	No debris or scums/foams present

Table 4: Groundwater Sample Information

Location	Rainfall event	Number of dry days prior to sampling	Discharge/water colour	Obvious or visual features
Monitoring Bore 1 (MLY GW01)	16.8mm	0	Clear/slightly cloudy	No debris or scums/foams present
Monitoring Bore 2 (MLY GW02)	16.8mm	0	Clear	Some sediment
Sump tile drainage outlet	16.8mm	0	Slightly cloudy	No debris or scums/foams present



3 ANALYSIS OF LABORATORY RESULTS

3.1 Stormwater Results

Table 5 shows the results of the 7 August 2015 stormwater sample round. The Awapuni Drain 10 metres downstream sample (MLYSW Site 2) is the mixing zone boundary and the compliance point. Exceedances of the consent trigger limits at this location and also the background site are highlighted in purple.

Parameter	Units Consent trigger limits		Stormwater retention ponds culvert outlets (MLYSW Site 1)	Awapuni Drain 10m downstream (MLYSW Site 2)	Awapuni Drain 10m upstream (MLYSW Site 3)	
рН	-LOG(H+)	6.5 – 8.5	6.52	7.41	7.29	
Total Suspended Solids	nded g/m ³ 100 g/m ³ above background site ¹		9	18	18	
BOD ₅	OD ₅ g/m ³ 20		8	6	5	
Total Petroleum Hydrocarbons			<0.7	<0.7	<0.7	
Total Nitrogen	g/m ³ 0.4		1.36	3.6	4.1	
Total Tannins	g/m³	Indicator test only	<0.2	<1.0	<1.0	
Dissolved Oxygen	Dissolved Oxygen Total Not less than 80% saturation		36.5	54.5	64.4	
Conductivity	mS/cm	0.3	0.877	9.605	9.664	
Total Resin Acids	otal Resin Acids g/m ³ 0.06		<0.0001	<0.0001	<0.0001	

Table 5: August 2015 stormwater sample results

3.2 Findings on Consent Condition Compliance – Stormwater

At all locations, pH levels are within the consent range. Total petroleum hydrocarbons and total resin acids were less than detection limits. Biological oxygen demand (BOD) was below the consent trigger limits.

Total suspended solids (TSS) concentration at the downstream site is the same as the upstream site and is therefore compliant with the consent trigger limit $(100g/m^3 above background site)$. TSS concentrations in the Awapuni Drain and the pond outlet were low (the pond outlet is $9g/m^3$).

This pond outlet concentration is significantly less than the July result of 2100g/m³. We consider that result to be anomalous and due to likely substrate disturbance during sample collection and other factors at the discharge outlet on the river side of the railway line.

This location which is shown in Figure 1 below, was inspected by Mark Poynter of 4Sight on the afternoon of 26 August 2015. This location is not an appropriate sampling point as it is beyond the site and susceptible to multiple potential influences not related to the log storage yard operation. These influences which are evident in the July photo, include backflooding into the outlet route from the Awapuni drain as well as pugging and sediment generation due to stock activity along the drain edge. Going forward, the outflow samples from the retention pond will be collected from the left hand side culvert upstand within the stormwater pond. A photograph of this location is shown in Figure 2.

¹ Background site is the Awapuni Drain 10 metres upstream



To minimise the likelihood of contamination during sample collection, debris and weed growth will be kept away from the sampling point.



Figure 1: Culvert outlets sampling point July 2015

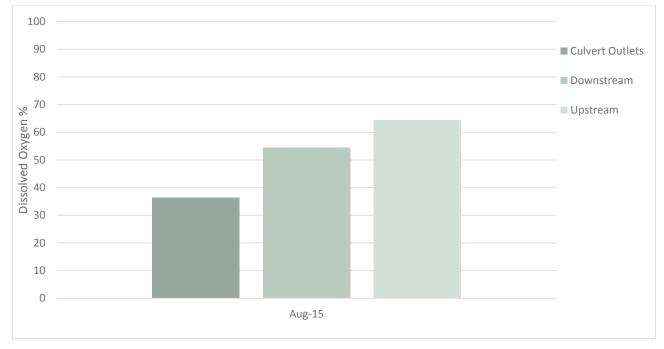


Figure 2: Culvert upstands within stormwater pond

Dissolved oxygen concentration is low in the Awapuni Drain. At the downstream site the August sampling showed a concentration of 54.5% and an upstream concentration of 64.4%. This suggests some reduction (by 9.9%) in downstream concentration due to the influence of the discharge which was at 36.5% concentration. Given the low TSS concentration, the low discharge oxygen value may be due to seasonal die off and breakdown of vegetation within the pond rather than sourced to run off from the log storage activity per see. As noted above, other factors may also be involved and influence dissolved oxygen concentration in the outlet flow once it passes beyond the Matawhero site.



It is suggested that with the next round of sampling, if the dissolved oxygen concentration in the retention pond outlet sample is low (below 80%), and if the downstream site concentration is both below the background concentration and also below the 80% trigger threshold, then further meter readings of dissolved oxygen concentration in the Awapuni Drain should be made at 20m and 30m below the outlet confluence. This will establish the physical extent of oxygen suppression that might be attributable in whole or part to the discharge from the log yard retention pond.





Conductivity results confirm that the Awapuni Drain at the time of sampling is tidally influenced and the discharge is not a significant influence on the receiving environment conductivity. Specific conductivity at the Awapuni Drain sites may depend on the particular state of the tide at the time of sampling. Results of conductivity since March 2015 are shown in Figure 4.

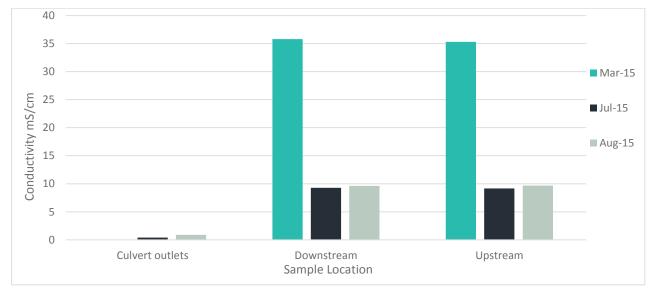


Figure 4: Conductivity results for 2015

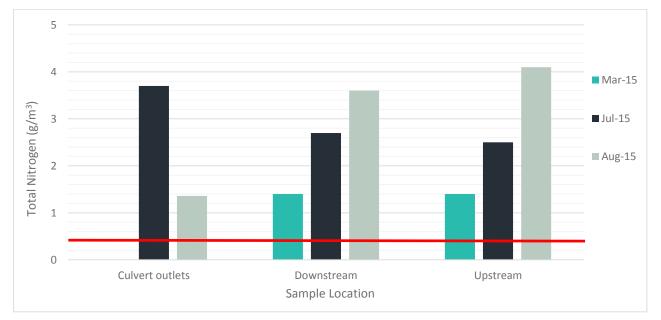


Total nitrogen (TN) concentration at the downstream site was below the background concentration. This may be suggesting a diluting influence from the discharge in respect of TN notwithstanding that the Awapuni Drain is highly enriched and all concentrations are well above the trigger level. This is likely to be a very localised effect as it is unlikely that the downstream sampling is representative of the full body of water across the width of the river. The downstream sample is collected close to the true left bank and the discharge itself is likely to disperse close to the bank edge as it is moved downstream.

Background total nitrogen has increased compared to the previous sample rounds. This may reflect seasonal influences with greater runoff and loss of nitrogen to waterbodies as the winter progresses.

In response to the July report, Paul Murphy of GDC asked what could be contributing to the total nitrogen result. There are no significant sources of nitrogen on site that could influence surface water or subsurface drainage (tile drainage). The retention pond discharge TN concentration may also be influenced by the state of the heavy crop of emergent vegetation that has colonised the pond. Die off of this vegetation in the winter may release nutrients and decrease uptake by way of plant growth resulting in higher discharge concentrations.

The constituents of total nitrogen for the pond outlet July sample also shows that total kjeldahl nitrogen (TKN) was $3g/m^3$ while nitrate-N + nitrite-N was $0.66g/m^3$. TKN, which represents ammonia and protein nitrogen, also dominates at the upstream and downstream sites for all three 2015 sample rounds. A potential source of TKN in the Awapuni Drain could be dairy or stock farming. The high proportion of TKN in the pond outlet sample is potentially more difficult to understand but may also reflect other 'farming' influences at the particular sampling location used as discussed above. For example it was apparent on the field inspection carried out on the 26 August that sheep had been grazing along this bank edge.



TN results for 2015 are shown in Figure 5.

Figure 5: Total nitrogen results for 2015 – consent trigger limit shown as red line



3.3 Groundwater results

Table 6 shows the results of the 7 August 2015 groundwater sample round. The sump tile drainage outlet is the groundwater quality compliance point. Exceedances of the consent trigger limits at this location and the monitoring bores are highlighted in purple.

Parameter	Units	Limit from consent condition	Limit from background samples at the sump tile drainage outlet ²	Monitoring Bore 1 (MLY GW01)	Monitoring Bore 2 (MLY GW02)	Sump tile drainage outlet
рН	-LOG(H+)	6.5 – 8.5	6.5 - 8.5	6.64	6.52	6.52
Conductivity	umho/cm	0.3 above background	866.3	891	935	804
Total Petroleum Hydrocarbons	g/m³	15	15	<0.7	<0.7	<0.7
Total Resin Acids	g/m ³	0.06 above background	n/a³	<0.0001	<0.0001	<0.0001
Total Nitrogen	g/m³	0.6 above background	1.49	0.26	3.6	1.67

Table 6: August 2015 groundwater sample results

3.4 Findings on Consent Condition Compliance – Groundwater

At the sump tile drainage outlet pH was within the consent range. Conductivity was below the consent trigger limit. Total petroleum hydrocarbons and total resin acids were less than detection limits.

Total nitrogen was above the consent trigger limit of $1.49g/m^3$ by 1.1 times. This exceedance is not considered significant, especially with Monitoring Bore 2, a background bore, having a total nitrogen result of 3.6 g/m³. This shows that groundwater flowing into the site already has high total nitrogen level, which is not increased further by activities on the logyard.

3.5 Further Actions Required in Light of Findings

As this is the first round of sampling with an exceedance of the trigger limit for dissolved oxygen, additional in situ measurements are recommended for the next round of sampling on the basis described above.

Taking into consideration background concentrations, no significant exceedences of the consent trigger limits for other parameters occurred so no further action is required. The next round of stormwater sampling (September/October round) is scheduled to taken by the end of October if an appropriate rain event occurs. The next groundwater sampling is scheduled for February.

² Two samples were collected in October 2010 as background samples. The results of these samples have been used to determine the trigger limits where required, the average of the results has been used.

³ Background samples were not tested for total resin acids.



4 CONCLUSIONS

4.1 Stormwater

- The Awapuni Drain is tidally influenced which explains high receiving environment conductivity and preempts any concern about discharge conductivity.
- Dissolved oxygen may be slightly suppressed close to the discharge relative to background concentration which is also low. The frequency and physical extent of this effect will be investigated further but is expected to be highly localised.
- The discharge is not adversely affecting receiving environment Total Nitrogen concentration.
- All other sample results are not notable and are were within the consent trigger limits.
- The next round of sampling is scheduled to be taken by the end of October.

4.2 Groundwater

- Total nitrogen at the sump tile drainage outlet exceeded the consent trigger level by 1.1 times. This result is not considered a significant exceedance.
- All other results were within the consent trigger limits.
- The next round of sampling is scheduled for February.



Appendix A:

Sampling Locations



Author: Paul Sorensen Checked: Christine Oakey Approved: Christine Oakey



either in whole or in part by any means without

prior consent of 4Sight Consulting Ltd.

0

25

50

75

100 m



Appendix B:

Laboratory Analysis Reports



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NALYSIS REPOR 7

Client:	4SIGHT Consulting Limited
Contact:	Kim Wepasnick
	C/- 4SIGHT Consulting Limited
	PO Box 25356
	Featherston Street
	WELLINGTON 6146

Lab No:	1460366 SPv1
Date Registered:	08-Aug-2015
Date Reported:	14-Aug-2015
Quote No:	66824
Order No:	
Client Reference:	Eastland Port-Dunstan Rd Surface Water
Submitted By:	Kim Wepasnick

Sampl	e Name:	MLYSW Site 1	MLWSW Site 2	MLYSW Site 3 07-Aug-2015 8:08		
		am	am	am		
Lab	Number:	1460366.1	1460366.2	1460366.3		
Individual Tests						
Total Suspended Solids	g/m ³	9	18	18	-	-
Total Nitrogen	g/m³	1.36	3.6	4.1	-	-
Nitrate-N + Nitrite-N	g/m³	1.01	0.157	0.182	-	-
Total Kjeldahl Nitrogen (TKN)	g/m³	0.35	3.4	3.9	-	-
Carbonaceous Biochemical Oxygen Demand (cBOD ₅)	g O ₂ /m ³	8	6	5	-	-
Tannin	g/m³	< 0.2 #1	< 1.0 ^{#1}	< 1.0 #1	-	-
Total Petroleum Hydrocarbons in Wat	er		I	I I		
C7 - C9	g/m³	< 0.10	< 0.10	< 0.10	-	-
C10 - C14	g/m³	< 0.2	< 0.2	< 0.2	-	-
C15 - C36	g/m³	< 0.4	< 0.4	< 0.4	-	-
Total hydrocarbons (C7 - C36)	g/m ³	< 0.7	< 0.7	< 0.7	-	-

Analyst's Comments

^{#1} Severe matrix interferences required that a dilution be performed prior to analysis of this sample, resulting in a detection limit higher than that normally achieved for the Tannin analysis.

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The following table(s) gives a brief description of the methods used to conduct the analyses for this job. The detection limits given below are those attainable in a relatively clean matrix. Detection limits may be higher for individual samples should insufficient sample be available, or if the matrix requires that dilutions be performed during analysis.

Sample Type: Aqueous			
Test	Method Description	Default Detection Limit	Sample No
Total Petroleum Hydrocarbons in Water	Hexane extraction, GC-FID analysis US EPA 8015B/MfE Petroleum Industry Guidelines [KBIs:2803,10734]	0.10 - 0.7 g/m ³	1-3
Filtration, Unpreserved	Sample filtration through 0.45µm membrane filter.	-	1-3
Total Kjeldahl Digestion	Sulphuric acid digestion with copper sulphate catalyst.	-	1-3
Total Suspended Solids	Filtration using Whatman 934 AH, Advantec GC-50 or equivalent filters (nominal pore size 1.2 - 1.5µm), gravimetric determination. APHA 2540 D 22 nd ed. 2012.	3 g/m ³	1-3
Total Nitrogen	Calculation: TKN + Nitrate-N + Nitrite-N. Please note: The Default Detection Limit of 0.05 g/m ³ is only attainable when the TKN has been determined using a trace method utilising duplicate analyses. In cases where the Detection Limit for TKN is 0.10 g/m ³ , the Default Detection Limit for Total Nitrogen will be 0.11 g/m ³ .	0.05 g/m ³	1-3
Nitrate-N + Nitrite-N	Total oxidised nitrogen. Automated cadmium reduction, flow injection analyser. APHA 4500-NO ₃ ⁻ I 22 nd ed. 2012 (modified).	0.002 g/m ³	1-3
Total Kjeldahl Nitrogen (TKN)	Total Kjeldahl digestion, phenol/hypochlorite colorimetry. Discrete Analyser. APHA 4500-N _{org} D. (modified) 4500 NH ₃ F (modified) 22 nd ed. 2012.	0.10 g/m ³	1-3





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tests marked *, which are not accredited.

Sample Type: Aqueous			
Test	Method Description	Default Detection Limit	Sample No
Carbonaceous Biochemical Oxygen Demand (cBOD₅)	Incubation 5 days, DO meter, nitrification inhibitor added, dilutions, seeded. Analysed at Hill Laboratories - Microbiology; 1 Clow Place, Hamilton. APHA 5210 B (modified) 22 nd ed. 2012.	2 g O ₂ /m ³	1-3
Tannin	Colorimetric with Folin phenol reagent, tannic acid used for calibration. APHA 5550 B 22 nd ed. 2012.	0.10 g/m ³	1-3

These samples were collected by yourselves (or your agent) and analysed as received at the laboratory.

Samples are held at the laboratory after reporting for a length of time depending on the preservation used and the stability of the analytes being tested. Once the storage period is completed the samples are discarded unless otherwise advised by the client.

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Carole Kester-Canoll

Carole Rodgers-Carroll BA, NZCS Client Services Manager - Environmental Division



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NALYSIS REPOR T

Client:	4SIGHT Consulting Limited
Contact:	Kim Wepasnick
	C/- 4SIGHT Consulting Limited
	PO Box 25356
	Featherston Street
	WELLINGTON 6146

Lab No:	1460416 SPv1
Date Registered:	08-Aug-2015
Date Reported:	18-Aug-2015
Quote No:	66825
Order No:	
Client Reference:	Eastland Port - Feb & Aug
Submitted By:	Kim Wepasnick

Sample Type: Aqueous						
Ś	Sample Name:	MLYGW 01	MLYGW 02	MLY STD01	WR GW01	WR GW02
	-	07-Aug-2015	07-Aug-2015	-	07-Aug-2015 1:11	-
		10:12 am	11:35 am 1460416.2	am	pm	pm 1460416.5
Individual Tests	Lab Number:	1460416.1	1460416.2	1460416.3	1460416.4	1400416.5
						1
pH	pH Units	-	-	7.1	-	-
Electrical Conductivity (EC)	mS/m	-	-	88.9	-	-
Dissolved Mercury	g/m ³	-	-	-	< 0.00008	< 0.00008
Total Mercury	g/m ³	-	-	-	< 0.0008	< 0.0008
Total Nitrogen	g/m³	0.26	3.6	1.67	-	-
Total Ammoniacal-N	g/m³	-	-	-	0.167	< 0.010
Nitrite-N	g/m³	-	-	-	0.020	< 0.002
Nitrate-N	g/m³	-	-	-	0.016	0.22
Nitrate-N + Nitrite-N	g/m³	0.046	0.135	1.14	0.036	0.22
Total Kjeldahl Nitrogen (TKN)	g/m³	0.21	3.4	0.53	-	-
Heavy metals, dissolved, trace	As,Cd,Cr,Cu,Ni,Pl	b,Zn				
Dissolved Arsenic	g/m³	-	-	-	0.0082	0.0015
Dissolved Cadmium	g/m³	-	-	-	< 0.00005	< 0.00005
Dissolved Chromium	g/m³	-	-	-	< 0.0005	< 0.0005
Dissolved Copper	g/m³	-	-	-	0.0007	0.0099
Dissolved Lead	g/m³	-	-	-	< 0.00010	< 0.00010
Dissolved Nickel	g/m³	-	-	-	< 0.0005	0.0067
Dissolved Zinc	g/m³	-	-	-	0.0113	0.063 #1
Heavy metals, totals, trace As,	Cd,Cr,Cu,Ni,Pb,Zr	1				
Total Arsenic	g/m³	-	-	-	0.0151	0.0022
Total Cadmium	g/m³	-	-	-	< 0.000053	< 0.000053
Total Chromium	g/m³	-	-	-	0.00077	< 0.00053
Total Copper	g/m³	-	-	-	0.0116	0.0117
Total Lead	g/m³	-	-	-	0.00127	0.00039
Total Nickel	g/m³	-	-	-	0.0027	0.0074
Total Zinc	g/m³	-	-	-	0.0128	0.059 #1
Total Petroleum Hydrocarbons	in Water			1	1	1
C7 - C9	g/m ³	< 0.10	< 0.10	< 0.10	< 0.10	< 0.10
C10 - C14	g/m ³	< 0.2	< 0.2	< 0.2	< 0.2	< 0.2
C15 - C36	g/m ³	< 0.4	< 0.4	< 0.4	< 0.4	< 0.4
Total hydrocarbons (C7 - C36)	g/m ³	< 0.7	< 0.7	< 0.7	< 0.7	< 0.7
Analyst's Comments			1	1		

Analyst's Comments

^{#1} It has been noted that the result for the dissolved fraction was greater than that for the total fraction, but within analytical variation of the methods.





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(ILAC-MRA) this accreditation is internationally recognised. The tests reported herein have been performed in accordance with the terms of accreditation, with the exception of tests marked *, which are not accredited.

SUMMARY OF METHODS

The following table(s) gives a brief description of the methods used to conduct the analyses for this job. The detection limits given below are those attainable in a relatively clean matrix. Detection limits may be higher for individual samples should insufficient sample be available, or if the matrix requires that dilutions be performed during analysis.

Sample Type: Aqueous			
Test	Method Description	Default Detection Limit	Sample No
Heavy metals, dissolved, trace As,Cd,Cr,Cu,Ni,Pb,Zn	0.45µm filtration, ICP-MS, trace level. APHA 3125 B 21st ed. 2005.	0.00005 - 0.0010 g/m ³	4-5
Heavy metals, totals, trace As,Cd,Cr,Cu,Ni,Pb,Zn	Nitric acid digestion, ICP-MS, trace level	0.000053 - 0.0011 g/m ³	4-5
Total Petroleum Hydrocarbons in Water	Hexane extraction, GC-FID analysis US EPA 8015B/MfE Petroleum Industry Guidelines [KBIs:2803,10734]	0.10 - 0.7 g/m ³	1-5
Filtration, Unpreserved	Sample filtration through 0.45µm membrane filter.	-	1-5
Total Digestion	Boiling nitric acid digestion. APHA 3030 E 22 nd ed. 2012 (modified).	-	4-5
Total Kjeldahl Digestion	Sulphuric acid digestion with copper sulphate catalyst.	-	1-3
рН	pH meter. APHA 4500-H ⁺ B 22 nd ed. 2012. Note: It is not possible to achieve the APHA Maximum Storage Recommendation for this test (15 min) when samples are analysed upon receipt at the laboratory, and not in the field.	0.1 pH Units	3
Electrical Conductivity (EC)	Conductivity meter, 25°C. APHA 2510 B 22 nd ed. 2012.	0.1 mS/m	3
Dissolved Mercury	0.45µm filtration, bromine oxidation followed by atomic fluorescence. US EPA Method 245.7, Feb 2005.	0.00008 g/m ³	4-5
Total Mercury	Bromine Oxidation followed by Atomic Fluorescence. US EPA Method 245.7, Feb 2005.	0.00008 g/m ³	4-5
Total Nitrogen	Calculation: TKN + Nitrate-N + Nitrite-N. Please note: The Default Detection Limit of 0.05 g/m ³ is only attainable when the TKN has been determined using a trace method utilising duplicate analyses. In cases where the Detection Limit for TKN is 0.10 g/m ³ , the Default Detection Limit for Total Nitrogen will be 0.11 g/m ³ .	0.05 g/m ³	1-3
Total Ammoniacal-N	Filtered sample. Phenol/hypochlorite colorimetry. Discrete Analyser. (NH ₄ -N = NH ₄ +·N + NH ₃ -N). APHA 4500-NH ₃ F (modified from manual analysis) 22 nd ed. 2012.	0.010 g/m ³	4-5
Nitrite-N	Automated Azo dye colorimetry, Flow injection analyser. APHA 4500 -NO ₃ I 22^{nd} ed. 2012 (modified).	0.002 g/m ³	4-5
Nitrate-N	Calculation: (Nitrate-N + Nitrite-N) - NO2N. In-House.	0.0010 g/m ³	4-5
Nitrate-N + Nitrite-N	Total oxidised nitrogen. Automated cadmium reduction, flow injection analyser. APHA 4500-NO ₃ ⁻ I 22 nd ed. 2012 (modified).	0.002 g/m ³	1-5
Total Kjeldahl Nitrogen (TKN)	Total Kjeldahl digestion, phenol/hypochlorite colorimetry. Discrete Analyser. APHA 4500- N_{org} D. (modified) 4500 NH ₃ F (modified) 22 nd ed. 2012.	0.10 g/m ³	1-3

These samples were collected by yourselves (or your agent) and analysed as received at the laboratory.

Samples are held at the laboratory after reporting for a length of time depending on the preservation used and the stability of the analytes being tested. Once the storage period is completed the samples are discarded unless otherwise advised by the client.

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Tuesday, 25 August 2015

Kaiti Beach Road PO Box 1048 Gisborne 4040

Attn: Martin Bayley

TRACE RESIN ACID ANALYSIS:

Resin Acids (incl. DHA) analysis for Marty Bayley (Eastland Port Ltd) – August 2015.

CLIENT'S ORDER NUMBER:

AA 1146 EPL Compliance Programme - Eastland Port

WORK PERFORMED BY:

WORK CHECKED BY:

APPROVED BY:

DATE OF ISSUE:

DISTRIBUTION:

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(SIGN)

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25 August 2015

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New Zealand Forest Research Institute Limited – A Crown Research Institute of New Zealand

DATE SAMPLES RECEIVED 10.08.2015

SAMPLE DESCRIPTIONEight samples in 1L glass bottles (450°C muffled Scion
sample bottles) – samples sent by James Isaac (Logic
Forest Solutions Ltd).

SAMPLE IDENTIFICATION

MLY GW01 MLY GW02 MLY STD01 MLY SW Site 1 MLY SW Site 2 MLY SW Site 3 WR GW01 WR GW02

SAMPLING PROCEDURE

This report relates only to the items tested as received and therefore does not necessarily represent the sample from which it was taken.

DATE OF TESTING 12.08.2015

METHODS

In-house method, involving unfiltered pH9 liquid/liquid extraction with dichloromethane (DCM), followed by gas chromatography - mass spectrometry (GC/MS) analysis.

RESULTS:

RESIN ACIDS (µg/L)

Sample name	MLY GW 01	MLY GW 02	MLY STD 01	MLY SW Site 1	MLY SW Site 2	MLY SW Site 3	WR GW 01	WR GW 02
Pimaric acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Sandaracopimaric acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Isopimaric acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Palustric acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Levopimaric Acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Dehydroabietic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Abietic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Neoabietic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Pimarenic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Sandaracopimarenic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Isopimarenic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
13-Abietenic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Pimaranic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Isopimaranic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Abietanic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Seco-1-dehydroabietic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Seco-2-dehydroabietic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
12-Chlorodehydroabietic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
14-Chlorodehydroabietic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
12,14-Dichlorodehydroabietic	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
7-Oxodehydroabietic acid	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.
Total Resin Acids	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.	n.d.

n.d. = not detected, method detection limit is $0.1 \mu g/L$ All results presented are from duplicate sample analysis and concentrations are in $\mu g/L$. Compounds are quantified if they have a response 2.5 times higher than the average blank.



Appendix C:

Field Form

Produced by Andrew.Stewart Ltd for Eastland Port Ltd. © 2015

		т	Field Quality Control Checks						
Was pre-cleaning sampling equipment used for these samples?	\bigcirc	z	N Were gas bubbles present in vials at time of collection?	~	z	NA)	Consistent with COC form?	3	2
Was pre-cleaning sampling equipment properly protected from contamination? 🕜 N Was sample filtered for metals prior to preservations?	\bigcirc	z	Nas sample filtered for metals prior to preservations?	Y	Z	NA	COC Filled out?	Θ	2
									L

ľ

		Additional Comments:	

	Sample	Sample Details		-	Vater Qual	Water Quality Parameters	U)		Q	Observations		
Sample ID	Sample Time	Approx. Depth (m)	Approx. Stream Flow Rate	Temp. (°C)	DO (%)	EC (µS/cm)	рH	Water colour	Debris	Foams / scums	Sediment plumes observable?	Photos Taken?
Muy Sw B.45cm 0.15	8.4sem	0.15	10L	8.11	36-5	440	6.52	11.8 36.5 877 6.52 Clear.	Z	Z	7	\prec
MLY SW 8.29mm 0.6	8.29am	0.6	101	11.8	2:4.2	14.2 S096 S.45 8.11	14.4	Slightly	2	Z	Z	~
MLYSW 8.08an 0.5	8.0800	.0.5		11.8	64.4	64.4 9664 7.29		Slightly	Z	Z	Z	~
MLYSTD 8. Sam 025	8. Stan		1205	11.7	36.0	11-7 36.0 80H 6.52	6.52	Slightly	Z	Z	Z	\prec
								C				

Surface Water Sampling Form

	Job Information	Equi	iquipment
Date: 7/8/15	Time: Arrive: 8 and Depart:	Water quality equipment description:	Calibration Records Filed?
Project Name: E PL Outsoured Conflictne	Project Number:	Interface Probe Number: 12 CI Co 9.01	Calibration Records Filed?
Site Location: MLLY	Operator: James Isear.	Sampling Equipment Type: Ruckat	
Weather:	_	Event Rainfall Depth: 1 6 . 8 m. M.	Number of Dry days Prior to Sampling:
Reason for sampling: Standard Compliance Prog	Reason for sampling: Standard Compliance Programme (Circle frequency: (2 Monthly / Quarterly) or	Additional Monitoring (describe):	

ANDREW.STEWART Eastland Port



ANDREW.STEWART

Groundwater Well Sampling Form (with criteria specific to A National Protocol for State of the Environment Groundwater Sampling in New Zealand, MfE, 2006)

Job Info	ormation
Date: 7/8/15	Time: Arrive: 9:15an Depart: 10:25an
Project Name: EPL Outsourced Compliance Programme	Project Number: AA1146
Site Location: MLLY	Operator: T. T. M. M.
Well ID: MLY QWOI	Weather: Fine

Equip	ment
Water quality equipment description: 13 C 100 791	Calibration Records Filed? (V) N
Interface Probe Number: 577506	Calibration Records Filed? (Y) N
Purging Equipment Type: (Please circle) Bailer Type: Plastic Teflon Pump Typ	e: Peristaltic Submersible Micro-purge Other:

			We	I Gauging	and Purge	e Volume C	alculation	s	
Casing Diameter	25mm	50mm	50m	50mm	50mm	100mm	100mm	100mm	Volume of water in a well
Bore Diameter	50mm	100mm	125mm	150mm	200mm	125mm	200mm	250mm	$V = \pi x r^2 x h$
Conversion Factor (volume L/m)	0.93	3.73	5.06	6.68	10.8	10.8	14.2	20.2	V = Volume in litres π = 3.142
Total Well Depth (-)	Water Lev	vel (=) Wat	ter Column						r = radius in m h = Height of water column in m
			er Column 821 m	(x) Convers	sion Factor	(=) Litres p		olume L	

Water G	Quality Pa	rameters							Low Flow Sar	npling
Beginni	ng Purging	g Time: 9	:40am	End Pur	ging Time	10 4	12 am	Fill Time:	-	Discharge Time: -
Litres	Time	DO (mg/L)	Cond. (µS/cm)	рН	Redox (mV)	Temp °C	DTW (mbTOC)		Commer	its
0	9:42	2.60	544	6.88	226.1	15.5	1.724	Clea	r water	-
0.5	9:52	1.92	901	6.62	191.5	15.4	1.728	Stigi	n Her ils	uder
1	9:58	1.92	888	6.63	1770	15.3	1.723	No	odeur.	0
1.5	10:03	1.94	887	6.64	164.6	15.3	1.725			
2	10:08	1.84	891	6.64	152	15.4	1.722			
2.5	10:12	1.92	891	6.64	1576	15.5	1-721			
-		_								
						4				
									11	
							<i>n</i>			4
Stabil	sation	±10%1	±3% or ±5% if <100*	±0.1*	± 10mV ¹	±0.1*	Example C odour / slig	omments: clear / s ht odour / strong od	slightly cloudy / turbid lour / describe odour	/ very turbid / colour / no (hydrocarbon/solvent/organic)
Crit	eria						Sampling in	NZ, 2006, ¹ Based	on Vic EPA (Australia	
2:5	1		ell Volume						Did field parame	eters stabilise? <u> N</u> N
20	~	Actual am	ount of water rem	oved prior to	sampling				Was the well dr	y purged? Y N

Field Quality Control Che	ecks			
Was pre-cleaning sampling equipment used for these samples?	\heartsuit	N	-	Consistent with COC form? (Y) N
Was pre-cleaning sampling equipment properly protected from contamination?	V	Ν	-	COC Filled out? Y N
Were gas bubbles present in vials at time of collection?	Y		NA	Time Sample Collected: 10 12 cm
Was sample filtered for metals prior to preservations?	Y	N	(NA)	Sample ID:
Analytes: Ammonia, Nitrate, Nitrite, Total Nitrogen, Heavy Metals (As, Cd, Cr, Cu, Pb, Hg, Ni, Zn), TP	H, Total	Resin A	cids	MINGWOD



Groundwater Well Sampling Form (with criteria specific to A National Protocol for State of the Environment Groundwater Sampling in New Zealand, MfE, 2006)

Job In	formation
Date: 718115	Time: Arrive: 10:30 an Depart: 11:40 an
Project Name: EPL Outsourced Compliance Programme	Project Number: AA1146
Site Location: MLN	Operator: J.T. M.M.
Well ID: MLY GWOZ	Weather: Fine

Equipment	
Water quality equipment description: $1 3 0 7 9 1$	Calibration Records Filed? 💋 N
Interface Probe Number: 5 7 5 0 6	Calibration Records Filed? 🕐 N
Purging Equipment Type: Bailer Type: Plastic Teflon Pump Type: Peristaltic Submersible	Micro-purge Other:

		a seed on a seed of the second se	We	II Gauging	and Purg	e Volume C	alculation	S	
Casing Diameter	25mm	50mm	50m	50mm	50mm	100mm	100mm	100mm	Volume of water in a well
Bore Diameter	50mm	100mm	125mm	150mm	200mm	125mm	200mm	250mm	$V = \pi x r^2 x h$
Conversion Factor (volume L/m)	0.93	3.73	5.06	6.68	10.8	10.8	14.2	20.2	V = Volume in litres π = 3.142
Total Well Depth (-)	Water Le	vel (=) Wat	er Column						r = radius in m h = Height of water column in m
			er Column 659 m	(x) Convers	sion Factor	(=) Litres p	er 1 Well V	olume _ L	

Water O	Quality Pa	rameters							Low Flow Sa	ampling	
Beginni	ng Purging) Time: 丨	0:32	End Pur	ging Time	: 11 :	21	Fill Time:	-	Discharge Tim	ie: -
Litres	Time	DO (mg/L)	Cond. (µS/cm)	pН	Redox (mV)	Temp °C	DTW (mbTOC)		Comme	ents	
0	10:40	7.5	928	6.52	90.1	15.2	1-587	ileer	butsome	sedimen	t.
0.5	10:47	2.5	930	6.52	84.1	15-2	1.921	0 -	ganic / R	o Hing ode	ur.
1	10:56	3.2	929	6.52	81.6	15.5	2.02		0	4	
1.5	11:07	2.6	933	6.52	78.3	169	1.95	- Left +	o rechard	ge For Q 1	nins
2	11:13	2.2	933	6.52	78.1	15.6	2.04				
2.5	11:28	2.6	935	6.52	74.2	15.6	2.140	well,	-e charge	ed ver	1
								slowle	ч.		0
									0		
						_				2	
Stabil	isation	±10% ¹	±3% or ±5% if <100*	±0.1*	± 10mV ¹	±0.1*				id / very turbid / colo ir (hydrocarbon/solve	
Cri	teria								on Vic EPA (Austra onsecutive readings	lia) 669. (either 3 min or 0.5l	. apart)
2	51		ell Volume		-				Did field paran	neters stabilise?	N
2.	ット	Actual am	ount of water rem	loved prior to	sampling			Non-Arrow Address of Malace Science and Science and Science	Was the well c	Iry purged?	Y N

Field Quality Control Che	cks		and the set	
Was pre-cleaning sampling equipment used for these samples?	V	N	-	Consistent with COC form? (V) N
Was pre-cleaning sampling equipment properly protected from contamination?	N	Ν	-	COC Filled out? Y N
Were gas bubbles present in vials at time of collection?	Y	N	NA	Time Sample Collected: 11: 35 and
Was sample filtered for metals prior to preservations?	Y	N	(NA)	Sample ID:
Analytes: Ammonia, Nitrate, Nitrite, Total Nitrogen, Heavy Metals (As, Cd, Cr, Cu, Pb, Hg, Ni, Zn), TP	H, Total	Resin A	cids	MLY GWOZ



Groundwater Gauging Form



Site Name: matanhero Logyard	Project Number: AA 1146
Operator: T. I	Date: 7/8/15
Interface Probe: 57506	Weather Conditions: Fair

Well ID	Time	Depth to LNAPL (mb TOC)	DTW (mb TOC)	LNAPL Thickness (m)	Well Depth (mbTOC)	Comments
MLYGW 01	9:42	NA	1.722	NA	9.543	
nevan 02	10:400		1-587	NA	4.246	
		FT.				
1						
		-				
						•
						low top of casing, m = metres.

Version 1.0 February 2015



Daily Summary Sheet



Site Name:	Natanhero Logizard, Project Number: AA 1146.										
Operator: 50	mes Isaar/Mat Main. Date: 7/8/15										
Reason for Vis	it: Surface and Ground Weather Conditions: Fine										
Time	Comments										
8.am	Arrived at MLY and proceeded to take										
	Surface vater samples, as previous days rainfall ensured that there was dis- charge from MLY SW site 1 into Anapuni drain.										
8:0800	Took sample for MLY SN Site 3.										
8:29am	Took sample for MLYSW Site 2.										
8:4 Sam	Took sample for MLYSW site 1										
8:59 an	Took sample for MLY STO 01.										
	Field analysis (using 451 meter) and photos taken at each site.										
9:15 am	After SW samples had been										
	organized into a chilly bin										
	ue moved to MLMAWOI. to.										
	start Ground water sampling. We encounted another well enroute										
	which could possibly be MLYGWOI										
	(see photos) hovever ne sampled										
	the well that we did in March.										



Daily Summary Sheet



Site Name:	nata where Logyard Project Number: AA1146									
Operator: Jo	mes Isace / matthein Date: 7/8/15									
	it: Surface / aprod water sampling Weather Conditions: Fine									
Time	Comments									
10:300	After collecting samples for MLYGWOI									
	Le moveel to MLYGWOZ.									
	Field analysis and photos were									
	taken for both tites,									
11:40am	Departed MLY to take									
	Ground water samples at where-									
	rata road,									

Page 2 of 2



Appendix D:

Matawhero Logyard Stormwater Results



D.1 Matawhero Logyard Stormwater Results

Summary table of results starting in March 2015. Exceedance of the consent trigger limits at the downstream site are shown in purple.

Parameter		Consent trigger limits	March/April 2 March 2015				May/June		July/August 7 August 2015		
	Units						9 July 2015				
			Stormwater retention ponds culvert outlets (MLYSW Site 1) ¹	Awapuni Drain 10m downstream (MLYSW Site 2)	Awapuni Drain 10m upstream (MLYSW Site 3)	Stormwater retention ponds culvert outlets (MLYSW Site 1)	Awapuni Drain 10m downstream (MLYSW Site 2)	Awapuni Drain 10m upstream (MLYSW Site 3)	Stormwater retention ponds culvert outlets (MLYSW Site 1)	Awapuni Drain 10m downstream (MLYSW Site 2)	Awapuni Drain 10m upstream (MLYSW Site 3)
рН	-LOG(H+)	6.5 – 8.5	n/a	8.17	8.17	7.31	7.77	7.54	6.52	7.41	7.29
Suspended Solids	g/m³	100 mg/L above background site ²	n/a	20	22	2100	18	19	9	18	18
BOD ₅	g/m³	20	n/a	7	7	7	7	7	8	6	5
Total Petroleum Hydrocarbons	g/m³	15	n/a	< 0.7	< 0.7	<1.4	<0.7	<0.7	<0.7	<0.7	<0.7
Total Nitrogen	g/m³	0.4	n/a	1.4	1.4	3.7	2.7	2.5	1.36	3.6	4.1
Total Tannins	g/m³	Indicator test only	n/a	1.5	1.3	<5	1.2	1.1	<0.2	<1.0	<1.0
Dissolved Oxygen	Total saturation	Not less than 80%	n/a	n/a	n/a	82.4	81	53.9	36.5	54.5	64.4
Conductivity	mS/cm	0.3	n/a	35.81	35.329	0.40	9.28	9.15	0.877	9.605	9.664
Total Resin Acids	g/m ³	0.06	n/a	<0.0001	<0.0001	<0.0001	<0.0001	n/a³	<0.0001	<0.0001	<0.0001

1 - No sample was collected as no discharge from the ponds was occurring

2 - No sample was collected as no discharge from the ponds was occurring

3 - Sample bottle broke during transport to the lab so no analysis for this site was able to be collected



Appendix E:

Matawhero Logyard Groundwater Results



E.1 Matawhero Logyard Groundwater Results

Summary table of results starting in March 2015. Exceedance of the consent trigger limits at the sump tile drainage outlet are shown in purple.

Parameter	Units	Consent trigger limits	Limit from		February	1	August			
			background samples at the sump tile drainage outlet		2 March 20)15	7 August 2015			
				MLY GW 01	MLY GW 02	Sump Tile Drainage	MLY GW 01	MLY GW 02	Sump Tile Drainage	
рН	-LOG(H+)	6.5 – 8.5	6.5 - 8.5	6.81	6.88	7.91	6.64	6.52	6.52	
Conductivity	umho/cm	0.3 above background	866.3	896	942	2043	891	935	804	
Total Petroleum Hydrocarbons	g/m³	15	15	<0.1	<0.1	<0.7	<0.7	<0.7	<0.7	
Total Resin Acids	g/m³	0.06 above background	n/a	<0.0001	<0.0001	<0.0001				
Total Nitrogen	g/m³	0.6 above background	1.49	0.35	0.23	1.46	0.26	3.6	1.67	

1 - Two samples were collected in October 2010 as background samples. The results of these samples have been used to determine the trigger limits where required, the average of the results has been used.

2 - Background samples were not tested for total resin acids.

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