



CAIRNS SHIPPING DEVELOPMENT PROJECT

Revised Draft Environmental Impact Statement

APPENDIX AK: Groundwater Impact Assessment: Northern Sands DMPA Report (2017)









Cairns Shipping Development Project

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GLOSSARY, ACRONYMS, ABBREVIATIONS

Term	Meaning
AHD	Australian Height Datum
bgl	below ground level
ВОМ	Bureau of Meteorology
COD	chemical oxygen demand
CSD	Cairns Shipping Development
°C	Degrees Celsius
DEHP	Department of Environment and Heritage Protection
DNRM	Department of Natural Resources and Mines
EC	electrical conductivity
EIS	Environmental Impact Statement
FCG	Flanagan Consulting Group
Golder	Golder Associates Pty Ltd
km	Kilometre
m	Metre
mm	Millimetre
bwl	below water level
m/s	metres per second
mm/day	millimetre per day
m ² /day	square metres per day
m ³	cubic metres
m ³ /day	cubic metres per day
μS/cm	micro-Siemens per centimeter
PASS	Potential Acid Sulphate Soils





1.0 INTRODUCTION

Flanagan Consulting Group (FCG) commissioned Golder Associates Pty Ltd (Golder) to provide advice related to soils and groundwater issues as part of the Revised Draft Environmental Impact Statement for the Cairns Shipping Development Project (CSDP).

The CSDP will require the land based placement of approximately 900,000 m³ of soft clays and up to 100,000 m³ of stiff clays at separate Dredge Material Placement Areas (DMPA). The stiff clays are proposed to be placed at Port North's Tingira Street property. The soft clays are proposed to be placed in the Northern Sands void DMPA.

The soft clays are to be dredged via a 5,600 m3 capacity Trailer Suction Hopper Dredge (TSHD) discharging to a temporary floating pump out facility that will be situated between approximately 2.6 and 3.6 km north east of Yorkeys Knob.

Dredge material will be pumped from the pump out facility via a submerged steel pipeline, which will make landfall near the Richters Creek mouth, thence to the Northern Sands DMPA mainly via cane farm headlands. Due to the ~8 km pipeline length that will be required to connect the pump out facility to the Northern Sands DMPA, up to three pipeline booster pumps may be required, depending on TSHD pumping capacity. Tailwater is proposed to be discharged adjacent to the site or pumped to an outfall at the Barron River highway bridge. This report addresses the placement of capital dredged material at the Northern Sands site – an existing void in the Barron Delta as shown on Figure 1. Conceptual placement of dredged materials at Northern Sands would have the following requirements:

- A pumping delivery line along one of the route options shown on Figure 1 including:
 - A floating pump-out facility.
 - The conceptual design for the pipeline includes 1 m diameter welded steel pipe on the seabed and creek bed where appropriate, and flanged steel section above ground on temporary earth pads. Booster stations will be positioned along the pipeline at various locations (possible locations are shown on Figure 1).
- Dredged material will be pumped into the sand pit (which may be enlarged prior to dredging operations) with identified PASS material placed in the deeper portion of the pit and covered by self-neutralising material. PASS will be placed to achieve a final surface at least 1 m below the permanent water table. Other self-neutralising material will also be placed below the permanent water table (subject to final design and approvals).
- Provision for tailwater treatment subject to preliminary concept design.

The aims of the overall studies were to describe the existing groundwater conditions associated with the Northern Sands site and to identify:

- Key groundwater related constraints (and opportunities) to design and construction of the facilities required for placement of the dredged material.
- Potential groundwater related environmental impacts and mitigation/management measures.

2.0 SITE SETTING

2.1 Site Background Information

The Northern Sands site covers an area of about 84 hectares and is located on the Barron River floodplain. The site contains an operating sand pit and is licenced to receive 'inert wastes' and PASS, both of which are placed into the excavated pit below the water table. The current sand pit covers an area of about 25 hectares and is known as "Narelle Lake". It should be noted that the existing pit will be enlarged by extension into an area located to the north of the pit).





2.2 Climate

The climate of the Cairns region and that of the Study Area is tropical with weather patterns consisting of very wet summers and drier winters. Key climatological and weather data was obtained from the nearest weather station, located at the Cairns Airport (Bureau of Meteorology (BOM) Station Number 031011) and is summarised below:

- Mean annual monthly maximum temperature is 29.0° Celsius (°C); with highest temperatures in December and January.
- Mean annual monthly minimum temperature is 20.8°C; with lowest temperatures in July.
- Mean annual rainfall is 1999.7 millimetres (mm) with highest rainfall in January through March.
- Mean number of days of rain greater than or equal to 1 mm is 119.6 days per year.
- Mean annual 9 am humidity is 72%; with February, March, and April having the highest humidity.
- Mean annual 3 pm humidity is 62% with February having the highest humidity.

Mean monthly rainfall and evaporation are presented in Table 1 and the annual rainfall from 2005 to 2015 is presented in Table 2.

Table 1: Mean rainfall and calculated evaporation data at the Cairns Airport (Weather Station Number 031011). Source: Bureau of Meteorology (2016).

Month	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sept	Oct	Nov	Dec
Mean rainfall 1942 to 2016 (mm)	390	448	419	195	92	48	30	27	33	46	94	178
Mean evaporation 1965 to 2016 (mm)	198	164	180	162	152	141	155	174	201	233	225	223

Note: data is rounded to the nearest millimetre

Table 2: Annual total rainfall data between 2005 and 2015 at the Cairns Airport (Weather Station Number 031011). Source: Bureau of Meteorology (2016).

Year	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015*
	2005	2000	2007	2006	2009	2010	2011	2012	2013	2014	2015
Rainfall (mm)	1471	2289	1813	2215	2199	2660	2623	2003	1269	1826	1897

Note: data is rounded to the nearest millimetre

The Cairns region experiences cyclonic storms on a regular basis with extreme rainfall events every two to eight years. Runoff intensity and storm surges from the sea would be expected. Significant rainfalls of 100 mm/day or greater can occur at any time during most of the year. The topography consists of low lying coastal plains which may be influenced by storm surge impacts and large storm run-off or flood events.

2.3 Drainage and Topography

The topography of the Northern Sands site is characterised by alluvial terraces and flat coastal plains of the Barron River Delta extending from the west of the site and east toward the ocean. Ground surface elevations across the site typically range from about 5.5 m to 2 m Australia Height Datum (m AHD).

Narelle Lake is the main surface water feature at the site. Recent hydrographic survey (July 2016 by Ports North) of the lake indicates excavation levels typically range from -1.5 m to -2.5 m AHD across the majority of the pit, apart from the southern and eastern sections of the pit where excavation levels range from -6.5 m to -14.5 m AHD. Contours of the elevation of the base of the lake from the recent hydrographic survey are shown on Figure 2. The water level in the lake varies, with measured water levels generally between -0.5 m



^{*}Not quality controlled by BOM (2016)



AHD and 0.5 m AHD, with higher levels up to approximately RL 1.8 m AHD (refer to Section 3.2 for discussion of lake levels).

The Barron River, Thomatis Creek and Narelle Lake are the dominant drainage features in the site area. The Barron River lies to the immediate south and west of the site, and has an average width of about 150 m with an assumed bed level of around -2 m AHD.

2.4 Regional Geology and Hydrogeology

Surface geology in the area of the site is shown on Figure 3. The Northern Sands site is located on the alluvial fan and delta of the Barron River. Erosion of the bedrock has resulted in the formation of the Barron River floodplain that stretches from Trinity Beach to the north of the site to south of the Cairns Airport. The floodplain is underlain by unconsolidated Quaternary age alluvial deposits of sands, gravels, silts and clays. These sediments reach thicknesses of up to around 90 m, in the area to the immediate north-east of the site, as shown below in Plate 1 (reproduced from QLD Water Resources Commission, 1982).

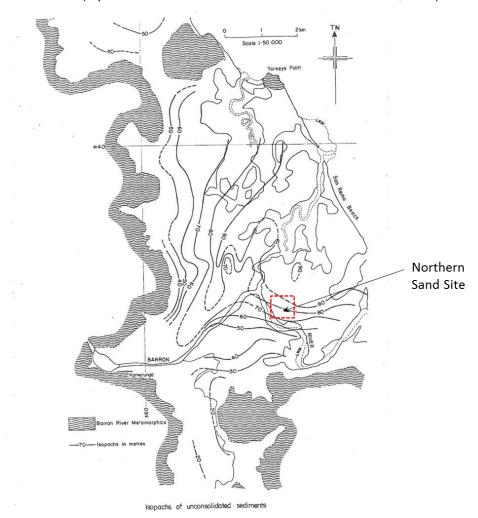


Plate 1 - Thickness of Unconsolidated Alluvium

Bedrock which underlies the unconsolidated sediments is exposed in the Macalister Ranges which outcrop to the west of the delta. Bedrock comprises Silurian/Devonian age metasediments (sedimentary rocks that have undergone some degree of metamorphism) comprising inter-bedded phyllite, schist, quartzite and chert beds which generally strike north to south.





Stratigraphic information available from the Queensland Registered Groundwater database (GWDB) (DNRM, 2016) within 2 km of the site indicates that the Quaternary age sediments are comprised of the following:

- Younger (Holocene age) alluvial deposits that generally range from surface to depths of 90 m. This
 sequence is associated with channel systems and consists of sand, silt and mud deposits, as well as
 minor gravel beds.
- Coastal (Holocene age) deposits in tidal mangrove and supratidal flats. These are located to the west and east of the site and consist of silt, sand and mud. This sequence was deposited in a marine environment with tidal influences from the Coral Sea. During transgressional and regressional periods, changes in sea levels, beach type sand, or silt and clay deposits would inter-tongue with terrestrial deposits.

There are two major aquifers within the unconsolidated sediments of the Barron River delta (QLD Water Resources Commission, 1982):

- An upper, unconfined aquifer varying in thickness from about 2 m to 11 m, which is overlain by up to 5 m of beach ridge deposits or clayey strata.
- A lower, confined or semi-confined aquifer, separated from the upper aquifer by a clay layer of varying thickness from around 3 m to around 25 m. The lower aquifer includes numerous inter-fingering clay layers.

Bore yields of up to 1500 m³/day are reported for these aquifers; however, the potential for use of this water for various beneficial uses is impacted by the salinity of the groundwater. Elevated salinity is observed up to 3 km from the coast (QLD Water Resources Commission, 1982).

Twenty-four registered bores are located within 2 km of the Northern Sands site as shown on Figure 4. Summary details of the registered bores are provided in Appendix A. The range of measured groundwater levels in these registered bores over the period from 1976 to 2016 are also shown on Figure 4. Hydrographs showing the variation in groundwater level over time for the registered groundwater bores are presented on Figure 5.

Figures 4 and 5 indicate that groundwater levels are generally lower in the vicinity of Thomatis Creek. QLD Water Resources (1982) states that "it seems likely that for most of the year, Thomatis/Richter Creek and a major part of the Barron River are effluent streams, acting as a line sink draining water from the aquifer system". This is illustrated in Plate 2 below, reproduced from that report. It is also noted that relatively low groundwater levels have been recorded in registered bores 11000025 and 11000031, which are located in the vicinity of low-lying mangrove areas. It is also noted that relatively low groundwater levels have been recorded in registered bores 11000032 and 11000049 which are located within 300 m of Narelle Lake, towards the east. These lower groundwater levels were recorded between 1977 and 2005. It is not known whether the lower groundwater levels at these bores is associated with the lake, or groundwater extraction.



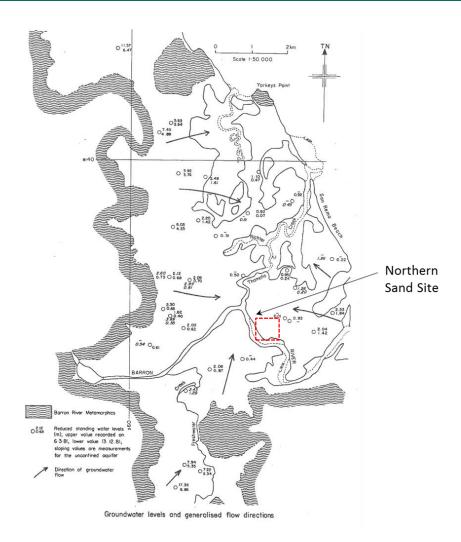


Plate 2 - Groundwater Flow across the Barron Delta

Water quality parameters from the registered bores are shown on Figure 6, and further information on groundwater quality parameters for the registered bores is provided in Appendix B.

Pumping tests carried out for the QLD Water Resources Commission 1982 study indicate transmissivities in the range of 1500 to 6800 square metres per day (m^2 /day) for the upper unconfined aquifer (corresponding to values of hydraulic conductivity in the range of $2x10^{-3}$ to $4x10^{-3}$ m/s), and around 550 to 3900 m²/day for the lower confined aquifer.

2.5 Groundwater Dependant Ecosystems

Groundwater Dependent Ecosystems (GDE) are defined as ecosystems whose ecological processes and biodiversity are wholly, or partially, reliant on groundwater. Examples of GDEs include wetlands, vegetation, mound springs, river base flows, plus saline discharges, springs, mangroves. GDEs may include aquatic ecosystems in rivers and streams that receive groundwater baseflow.

Information on potential groundwater dependent ecosystems is available from the National Atlas of Groundwater Dependent Ecosystems. Based on information from this atlas, the potential for groundwater dependent ecosystems in surface water bodies and for vegetation in the vicinity of the site is shown in Figure 7. Further information on groundwater dependent ecosystems is presented in the Terrestrial Ecology report - Northern Sands (Biotropica, August 2016).

Figure 7 indicates the presence of vegetation with a high potential for groundwater interaction between Narelle Lake and the Barron River along the western and southern boundaries of the lake.





The reaches of Thomatis Creek and Barron River in the vicinity of the Northern Sands site are indicated to have moderate potential for groundwater interaction.

3.0 GROUNDWATER CONDITIONS ON SITE

3.1 Stratigraphy

Previous subsurface investigations carried out on and adjacent to the Northern Sands site include the following:

- Borehole investigation carried out by Probin Pty Ltd in 2007. This work comprised 21 No. boreholes to depths ranging between approximately 15 m to 24 m below ground level (m bgl) on the subject site.
- Borehole investigation carried out by GEO Investigate in 2013. This work comprised 10 No. boreholes within the existing lake to depths ranging from 18 m to 30 m below water level (m bwl).
- Cone Penetrometer Testing (CPT) carried out by GEO Investigate in July 2016. This work comprised 30
 No. CPT's to depths ranging from approximately 8 m to 24 m bgl on the subject site.
- Borehole investigation carried out by Golder in 1995. This work comprised 10 No. boreholes to depths ranging approximately 6 m bgl to the north of the subject site.
- Cone Penetrometer Testing (CPT) carried out by GEO Investigate in July 2016. This work comprised 14
 No. CPT's to depths ranging from approximately 11.6 m to 14.7 m bgl to the north of the subject site.

Additional subsurface investigations carried out for these studies on and adjacent to the Northern Sands site include the following:

- Groundwater monitoring bore installation, sampling and testing by Golder in September 2016 and November 2016. This work comprised 5 No. boreholes to depths ranging from approximately 1.4 m to 12 m bgl on the subject site. Borehole reports are presented in Appendix D.
- Cone Penetrometer Testing (CPT) carried out by Golder in November 2016. This work comprised 7 No. CPT's to depths ranging from approximately 9.8 m to 18.5 m bgl to the east of the subject site. CPT reports are presented in Appendix D.

The investigation locations are shown on Figure 8. Inferred subsurface cross-sections (utilising information from registered groundwater bores as well as the investigations outlined above) are presented in Figures 9 and 10.

The inferred subsurface conditions in the area are broadly consistent with the published geology and generally comprises the following sequence.

- Sandy silty clay typically above -1 m AHD.
- Sand/gravelly sand broadly between 3 m to -7 m AHD.
- Silty clay broadly between 0 m to -16 m AHD and ranging in thickness from around 3 m to 10 m.
 This layer may not be continuous although it is shown to be so in Figures 9 and 10.
- Sand/gravelly sand broadly between -7 m AHD to -20 m AHD.
- Silty clay broadly between 14 m AHD to -26 m AHD, and with a maximum thickness of around 12 m
- Gravelly sand/sandy gravel below -26 m AHD, to a maximum depth of -37 m AHD in registered monitoring bores RN139211 and RN11000053.

The upper two sand/gravelly sand units (and the 3 m to 10 m thick clayey interbed in the site area) are interpreted to represent the upper unconfined regional aquifer. The lower gravelly sand/sandy gravel unit encountered below -26m AHD is interpreted to represent the lower semi-confined/confined regional aquifer.



3.2 Groundwater Levels

Six shallow groundwater monitoring bores (UG1, UG2, BH2, BH3, BH4a and BH5a) have been installed by others at the locations shown on Figure 11. The depth of the monitoring bores ranges between 3.9 m and 8.5 m bgl, however further information regarding the construction of these bores is not available. Hydrographs of groundwater levels and the water levels in the lake between mid-2009 and mid-2016 (information provided by Landline Consulting – environmental consultants to Northern Sands operations) are shown on Figure 12. It was suspected that the recorded lake levels may be approximately 0.5 m above the actual water levels (i.e. actual lake water levels may be approximately 0.5 m below those illustrated in Figure 12). Recent survey of groundwater and lake water levels by Golder confirmed these suspicions in the historical data, however, the data in Figure 12 has not been adjusted from that provided by Landline Consulting.

Notwithstanding the potential error in the lake water levels, the pattern of variation of groundwater levels matches the pattern of variation in the lake level. Measured groundwater levels generally vary between -0.3 m and 1.0 m AHD. It is likely that the groundwater levels in the monitoring bores closest to the Barron River are impacted by tidal fluctuations in the Barron River, and that in particular, measured groundwater levels lower than 0 m AHD are likely to reflect low tide conditions at or close to the time of measurement.

Four shallow groundwater monitoring bores (BH01, BH02, BH03 and GA04) were installed by Golder at the locations shown on Figure 11. The depth of these monitoring bores ranges between 5.5 m and 12 m bgl. Construction information for these monitoring bores are presented on the borehole reports in Appendix D and summarised in Tables 3 & 4. These bore were developed after their installation by purging at least 5 well volumes to remove drilling and filter pack fines and to ensure hydraulic connection between the aquifer and the bore casing. Water quality parameters were monitored during development until parameters stabilised.

Pressure/temperature transducers were installed in BH01 and BH03 in September 2016 to record groundwater levels at one hourly intervals. Hydrographs of groundwater levels at BH01 and BH03 from 29 September to 6 April 2017 are presented in Figure 14. Groundwater level ranges are summarised in Table 4. The hydrographs for BH01 and BH03 show a response to rainfall and tidal influence of approximately 0.10 m.

Table 3: Monitoring bore location and level

Borehole ID	Easting*	Northing*	Elevation (m AHD)	Standpipe Stick up (m)
BH01	363627.63	8136178.11	2.64	0.67
BH02	364376.67	8135447.11	2.13	0.63
BH03	364301.50	8135126.42	1.97	0.67
GA04	364331.65	8135689.89	3.58	0.90

^{*}Coordinates system: GDA 94 zone 55

Table 4: Monitoring bore construction information and water levels

Borehole	Bore depth	Screened	Δαιιιτατ		Water L	evel Range	
ID	(m bgl)	interval (m bgl)	Soil description	type	*(m bgl)	*(m AHD)	
BH01	6.0	3.0 to 6.0	Sand	Unconfined	2.5 to 1.4	0.1 to 1.3	
BH02	6.0	3.0 to 6.0	Clayey sand	Unconfined	2.0 to 1.2	-0.1 to 1.0	
BH03	12.0	6.0 to 12.0	Clayey sand and clay	Unconfined	2.0 to 1.0	-0.06 to 0.97	
GA04	5.5	3.5 to 5.5	Sand	Unconfined	3.4 to 1.8	0.2 to 1.7	

^{*}Where data is not presented in a range, only a single data point exists.





The nearest measurement of river levels in the Barron River is at Cairns Airport, approximately 2 km downstream of the site. It is understood that the tide gauge does not accurately measure low water levels and is not able to read levels below approximately 0 m AHD. Records from this station are interpreted to indicate a long-term average river level of around 0.25 m AHD. Comparison with the groundwater levels illustrated in Figure 12 indicates that the long term average groundwater levels close to the river, and the lake level, are similar to the average river level.

Given the above information, 0 m AHD has been adopted as the lowest permanent water level in the lake.

3.3 Groundwater Quality

Monitoring bores BH2, BH3, BH4a and BH5a have been sampled and tested monthly by others (Landline Consulting) from September 2014 to April 2016 for the following parameters:

A 1		
Alur	ninum	ì

- Copper
- Sulphate

- Arsenic
- EC
- Total Nitrogen

- Cadmium
- Lead
- Total Phosphorus

- Chloride
- Iron
- Zinc

- Chromium
- Mercury
- COD (Chemical oxygen demand)
- pН

Copies of plots showing the results of water quality testing are presented in Appendix C.

Ranges of EC and pH at monitoring bores and regional groundwater bores in the area of Narelle Lake are shown in Figure 13. The EC in Narelle Lake between September 2011 and March 2016 varied between 200 S/cm and 1000 μ S/cm. It is noted that registered groundwater bores 11000049 and 11000033 to the immediate east of Narelle Lake have high recorded EC ranging from 19 000 μ S/cm to 38 000 μ S/cm. The measured EC at these two registered groundwater bores is higher than at all other registered groundwater bores within a 2 km radius where water quality measurements are available.

Golder monitoring bores BH01, BH02, BH03 were sampled on 29 September and 22 November 2016, and GA04 was sampled on 24 November 2016. Groundwater samples from BH01, BH02, BH3 and GA04 were forwarded under chain of custody to SGS (a NATA accredited laboratory) for testing. Samples were analysed for the following parameters. It should be noted that only aluminium, iron and total iron were tested during the 29 September 2016 sampling event.

- Alkalinity
- EC
- Sulphate

- Aluminium
- Iron
- TDS

- Anions
- Total hardness

- Calcium
- Hq
- Total Iron

- Cation
- Potassium

Magnesium

Total Nitrogen

- Chloride
- Sodium

The field data and laboratory results are presented in Appendix E and summarized in Table 5.





Table 5: Field and laboratory pH and salinity results

Monitoring bore	pH range lab results	pH range field results	EC range (μS/cm) lab results	EC range (μS/cm) field results	Salinity
BH01	6.8 to 7.0	6.5 to 7.2	230 to 250	300 to 373	Fresh water
BH02	7.6 to 7.9	6.7 to 8.0	2,900 to 11,000	8,000 to 10,000	Brackish water
BH03	7.9 to 8.0	7.2 to 10.2	8,600 to 24,000	25,500 to 26,540	Brackish water
GA04	6.5	5.74	160	148	Fresh water

3.4 Hydraulic Conductivity

Slug tests were carried out at BH01, BH02, BH03 and GA04 during field investigations in November 2016. Two falling and two rising head test were conducted at each bore location. Data was recorded by a pressure transducer recording water level every second in conjunction with manual water level measurement during the test. Hydraulic conductivities estimated from the falling and rising head tests conducted at each bore are presented in Appendix G and summarised in Table 6.

Talsma tests were carried out within the upper clay layer at the location of GA04 and GA05. Laboratory plasticity and grading tests were carried out to confirm soil classifications at the locations of the Talsma tests. The results of the laboratory testing on the soils are presented in Appendix F. The results of the Talsma tests are presented in Appendix G and are summarised in Table 6.

Table 6: Field hydraulic conductivity testing results

Test Location	Test depth range (m)	Soil description	Method used	Hydraulic conductivity K (m/s)
BH01	3.0 to 6.0	Sand	Hvorslev	2 x 10 ⁻⁴
BH02	3.0 to 6.0	Clayey sand	Hvorslev	9 x 10 ⁻⁷
BH03	6.0 to 12.0	Clayey sand	Hvorslev	3 x 10 ⁻⁴
GA04	3.5 to 5.5	Sand	Hvorslev	2 x 10 ⁻³
GA04	0.3 to1.4	Sandy clay	Talsma equation	7 x 10 ⁻⁷
GA05	0.3 to 1.5	Sandy clay	Talsma equation	1 x 10 ⁻⁷

3.5 Porosity

Samples of the upper sands from GA04 and at GA05 were forwarded for laboratory testing to assess their maximum/minimum densities. Relative density of the sands was inferred from the results of Cone Penetrometer Testing at nearby test locations. The in situ density of the sands was then assessed by comparing the inferred relative density with the maximum/minimum densities achieved in the laboratory testing. Void ratios and porosities for the upper sand formation were calculated based on the interpreted insitu dry densities. Results of the laboratory testing are presented in Appendix F and the assessed porosity results are presented in Table 7.

Table 7: Assessed porosity for upper sand formation.

Sample	Inferred Density Ratio (%) from CPT	Inferred <i>in situ</i> Density (t/m³)	Inferred Dry Density (t/m³)	Moisture Content (%)	Void Ratio (e)	Porosity (n)
GA04 (Brown Sand)	45	1.62	1.27	27.5	1.08	0.39
GA04 (Brown Sand)	96	1.86	1.46	27.5	0.82	0.29
GA05 (Yellow Sand)	45	1.67	1.46	14.0	0.81	0.37
GA05 (Yellow Sand)	96	1.87	1.64	14.0	0.61	0.29

A porosity of 0.35 was adopted for the groundwater modelling for the upper sand layer.





3.6 Conceptual Hydrogeological Model

A conceptual hydrogeological model for the Northern Sands site is illustrated in Figure 15. The following points are noted regarding this conceptual model:

- The Northern Sands site is underlain by an upper unconfined aquifer and a lower confined or semiconfined aquifer. Both of these aquifers extend broadly across the Barron River delta.
- In the area of the Northern Sands site, the upper unconfined aquifer includes a 3 m to 5 m thick clayey interbed, which may or may not be laterally continuous. The upper aquifer is recharged directly by rainfall.
- The overall direction of groundwater flow in the upper unconfined aquifer is towards the Barron River and Thomatis Creek, as discussed in QLD Water Resources Commission (1982). Close to these streams, groundwater exchange will occur as a result of tidal fluctuations in the streams.
- Groundwater flow in the deep confined aquifer is towards the coast. Recharge to this aquifer occurs further to the west where the confining unit is absent in some areas.

In the current condition, the groundwater level in the lake is similar to the groundwater level in the upper unconfined aquifer. When the lake level is raised during the period of dredged material placement along with large volumes of seawater, a hydraulic gradient away from the lake will be created, and saline water will flow away from the lake, primarily through the upper sand layer as a result of the higher permeability of these sediments. Flow will diminish over time as a result of the low permeability of dredged material which will fill the base of the lake and "seal" the upper sand unit.

4.0 CONSTRAINTS AND OPPORTUNITIES

Dredged material will be placed in the sand pit along with large volumes of seawater, within a containment bund that will be constructed around the perimeter of the lake, and across the lake and across the two existing islands that are located within the lake (refer to the drawing included in Appendix H). The northern section of the pit will also be deepened to accommodate the volume of material that will be placed. Water levels in the lake will be raised to between 5 m to 7 m AHD during material placement. As a consequence, the lake level in this area will be at a level above the groundwater level in the surrounding aquifer and above the level in the Barron River for this period. The seepage rate out of the lake during this period will be significantly lower than the flow rate delivered to the lake with the dredged material, and thus the majority of the delivered water will need to be managed through tailwater ponds. The proposed concepts for tailwater management have been developed by BMT JFA and are outlined in their Dredged Material Placement Assessment, 2016.

As noted above, saline water will flow radially away from the lake during the period that the lake level is artificially raised. Along the western boundary of the lake, this water will flow through the upper sandy layer towards the Barron River, with the potential to impact on the salinity of the river. Along the remainder of the perimeter of the lake, water will flow out into the upper unconfined aquifer horizontally through the sandy materials exposed around the edges of the lake with the potential to impact on water quality in this aquifer. There is also the potential for increased salinity in the near surface clay layer, either as a result of direct lateral migration from the lake, or as a result of upward migration of salt from the impacted extent of the upper unconfined aquifer.

The water level in the tailwater treatment ponds will be only slightly above typical lake levels, and will only be elevated for a short period of time. The potential for impact on groundwater is thus significantly less than for the disposal area.

The potential for unacceptable impacts on water quality in the Barron River and in the upper unconfined aquifer, and the potential for increased salinity in near surface soils has been assessed through groundwater modelling as discussed in the following section.





5.0 ASSESSMENT OF IMPACTS

5.1 Groundwater modelling to assess impacts on upper unconfined aquifer and shallow soils, and impacts on Barron River

5.1.1 Description of Groundwater Model

As discussed in Section 3.6, a conceptual hydrogeological model for the Northern Sands site was prepared as shown in Figure 15. The conceptual model is based on the inferred subsurface cross-section shown in Figure 9.

Prior to groundwater modelling all of the additional information on ground conditions from the current studies and from historical investigations in the vicinity was collated, and the ground model for the site area was updated. In order to provide an assessment of saline water flow away from the lake during the period of increased lake water level, two simplified cross-sectional numerical models were developed, based on the inferred subsurface conditions along SSW to NNE and SW to NE oriented cross-sections as shown in Figure 8. The simplified cross-sectional models are shown in Figure 16.

The finite element software SEEP/W was used to develop a variably saturated, density dependent solute transport model. Parameters for the modelling were based on the results of the fieldwork and laboratory testing as appropriate.

During the placement of dredged material, the water level in the lake will be raised and the pit filled with saline water. Modelling has been carried out for maximum lake levels at RL 5 m AHD, RL 5.5 m AHD, RL 6.5 m AHD and RL 7.0 m AHD. Placement of the dredged material has been modelled in 4 sequential steps, with placement of the dredged material in a series of horizontal lifts to raise the dredged material level a maximum level at 1.2 m below the maximum lake level over a period of 80 days. It has been assumed that the dredged material will have a hydraulic conductivity of 1 x 10⁻⁷ m/s.

The adopted boundary conditions are shown in Figure 17. The water level in the lake has been modelled as increasing from 0.25 m AHD to 3 m AHD over the first 7 days of filling, and then being held constant at this level until the level of dredged material in the lake has risen to above the top of the upper sand layer.

5.1.2 Impacts on the upper unconfined aquifer and shallow soils to the north and east of Narelle Lake

Results from the groundwater models for Sections 1 and 2 are presented in Figures 18 to 25. Figures 22 to 25 show contours of the salinity concentration under increased lake levels during the period of placement and for 2 years following filling. Figures 18 to 21 show profiles of the **increase** in salinity concentration above the existing concentration (i.e. prior to placement) with distance away from the lake. The profiles are based on salinity concentrations in the upper sand layer immediately below the near surface clay layer. A summary of the approximate distance from the lake impacted by an increase in salinity is provided in Table 8. These distances are also shown on Figure 26.

The increased distance of salt migration for the lower water levels relates to the time required for the upper sand layer to be covered by the dredged material. The rate of lateral migration reduces once this layer is covered by the low permeability dredged material, and the time required for the dredged material to rise to the level of the top of the sand layer is longer for the models with lower water levels. Note that, as discussed above, the boundary condition representing the water in the lake has been based on the assumption that the water level will be held constant at RL 3 m AHD until the level of dredged material in the lake has risen to above the top of the upper sand layer. This has the effect of limiting the head that drives the lateral migration and will be the primary mitigation measure to limit the extent of lateral migration.





Table 8: Extent of increase in salinity in upper sand layer

Section	Level of water in placement area	Approximate maximum distance to which increased concentration extends
	RL 5m AHD	110m
Section 1	RL 5.5 m AHD	120m
Section	RL 6.5 m AHD	90 m
	RL 7.0 m AHD	80 m
	RL 5.0m AHD	115 m
Section 2	RL 5.5 m AHD	110 m
Section 2	RL 6.5m AHD	105 m
	RL 7 m AHD	105 m

The results of the modelling indicate that lateral migration through the near surface clay layer is significantly less than the extent of migration through the upper sand layer, and that the hydraulic gradient remains downwards throughout the period considered in the modelling (i.e. up to 2 years after the start of placement of dredged material). This downward hydraulic gradient will limit the extent to which salt can migrate upwards into the near surface clay layer and it is assessed that negligible changes in the salinity of the near surface clay will occur.

Within the extent of aquifer that is impacted by outward migration of salt during the period of placement of the dredged material, salt concentrations are likely to remain elevated for a in the long term¹ following the placement. Lateral flow of groundwater will be impacted by the change from the current situation with groundwater flow to the current lake, to groundwater flow around the infilled lake. This will limit the rate at which salt will be flushed from the system, however it is noted that in the area to the north of the placement area, groundwater flow towards Thomatis Creek will be unaffected by the placement of dredged materials.

5.1.3 Impacts on the Barron River

The potential flow rate and solute transport rate between the lake and the Barron River during the period of increased lake water level was preliminarily assessed during previous studies, noting that at that stage a lower lake level was adopted and that a different lake configuration was proposed. The previous estimate of the total seepage volume into the Barron River was based on an assumption that the lake level would be raised across the entire area of the existing lake. The length of river adjacent to the lake will be reduced for the currently proposed approach, which will offset the effect of the higher lake levels within the placement area.

The previous studies indicated that the steady state groundwater seepage from the lake to the river was estimated to be 4800 m³/day as a result of the increase in lake level. The estimated time for breakthrough of the saline water plume from the lake to the river was estimated to be 40 to 80 days after raising the level in the lake. After the breakthrough the salinity of the seepage water would be equal to the concentration in the lake. The previous studies did not account for a progressive diminishing in the rate of seepage as dredged material was placed in the lake. The results of the current modelling indicate that the groundwater seepage from the lake to the river during the period of the raised water level in the lake will briefly reach a rate of 25,000 m³/day with a maximum salt flux of 3500 g/s distributed along the approximately 1.1 km length of the Barron River that is located to the west of the placement area, for a lake level in the placement area of RL 7 m AHD. These seepage and salt flux rates impacts will be of short duration during the period before the dredge material rises above the level of the upper sand layer.

Potential impacts of the seepage on water quality in the Barron River have been addressed by BMT WBM in their Marine Water Quality Impact Assessment Technical Report, 2017.



¹ Refer to Table 10 for definitions of duration of impact that are used in this report.

5.2 Other Impacts

Other impacts which could result from the proposed extension of the lake, construction of the bunds, and increased water level in the lake during placement of dredged material include:

- Seepage beneath the bund wall in areas where the foundation material beneath the bund comprises higher permeability sandy material leading to increased saturation levels/water logging with high salinity at the surface close to the bund wall. Upward migration of water, potentially with elevated salinity, could also occur at locations further from the bund wall as a result of increased groundwater pressures in the upper aquifer, where isolated areas of higher permeability sandy materials are present at surface and are directly connected with the upper aquifer. The potential for such impacts to occur close to the bund wall can be mitigated through appropriate subsurface investigations along the bund, and measures such as the removal of unsuitable material from the foundation. The potential for impacts at locations further from the bund wall can be mitigated through management of groundwater pressures in the upper aquifer. This mitigation will be achieved through controlling the water level in the lake until the level of the low permeability dredged material in the placement area has increased to the level where it limits the direct connection between the aquifer and the water in the placement area.
- If areas of high permeability sandy soils are not detected and addressed in the design and/or construction of the bund wall, the potential exists for piping through such materials, with the potential to impact on the integrity of the bund wall. This could lead to safety risks and risks to adjacent infrastructure, in addition to the environmental risks that would result from the potential release of large volumes of saline water and dredged material. The potential for such impacts to occur can be mitigated through appropriate subsurface investigations along the bund, appropriate design of the bund, and measures such as the removal of unsuitable material from the foundation.

5.3 Summary of Potential Impacts

The potential groundwater impacts associated with the project are summarised as follow:

- Impacts on water quality in the Barron River as a result of seepage from the dredge placement area to the river.
- Impacts on water quality in the upper unconfined aquifer.
- Increased salinity in near surface soils.
- Increased saturation levels/water logging of surface soils as a result of seepage beneath the bund wall.
- Seepage beneath the bund wall adversely impacting on the integrity of the bund wall.
- Upward migration of water, potentially with elevated salinity, at locations further from the bund wall where isolated areas of higher permeability sandy materials are present at surface and are directly connected with the laterally extensive upper aquifer.

Further assessment of soils related impacts is presented in Section 7.0

6.0 IMPACT ASSESSMENT – GROUNDWATER

6.1 Methodology

In order to address the terms of reference, guidelines and other requirements for the currently defined project, the following methodology was adopted:

- Assess impacts (based on the risk assessment format outlined below);
- Provide recommendations for mitigation by design changes; and
- Provide recommendations for mitigation by management.

Flanagan Consulting Group has extracted relevant items from the Queensland Government Terms of Reference and the Commonwealth Government Guidelines for groundwater studies. These items and relevant details are presented in Appendix I.





The initial assessment of impacts utilises a significance table based on that shown in Table 9.

Table 9: Significance criteria

Impact significance / consequence	Description of significance (examples)					
Very High	The impact is considered critical to the decision-making process. Impacts tend to be permanent or irreversible or otherwise long term and can occur over large scale areas. Very high sensitivity of environmental receptors to impact (e.g. permanent loss of groundwater dependent ecosystems).					
High	The impact is considered likely to be important to decision-making. Impacts tend to be permanent or irreversible or otherwise long to medium term. Impacts can occur over large or medium scale areas. High to moderate sensitivity of environmental receptors to impact (e.g. permanent increase in salinity of surface aquifer creating permanent decrease in cane crop yields and reduced health of riparian vegetation).					
Moderate	The effects of the impact are relevant to decision making including the development of environmental mitigation measures Impacts can range from long term to short term in duration Impacts can occur over medium scale areas or otherwise represents a significant impact at the local scale Moderate sensitivity of environmental receptors to impact (e.g. bund failure resulting in discharge of saline waters and dredge material to riparian areas and Barron River and resulting in short term mortality of adjacent cane crops or short term suspended solids loading to the Barron River).).					
Minor	Impacts are recognisable/detectable but acceptable. These impacts are unlikely to be of importance in the decision making process. Nevertheless, they are relevant in the consideration of standard mitigation measures. Impacts tend to be short term or temporary and/or occur at local scale. (e.g short term increase in salinity of surface aquifer creating short term decrease in cane crop yields and reduced health of riparian vegetation).					
Negligible	Minimal change to the existing situation. This could include, for example, impacts which are beneath levels of detection, impacts that are within the normal bounds of variation, or impacts that are within the margin of forecasting error (e.g. minor short term salinity increases in adjacent surface aquifer salinity).					
Beneficial	Impacts have a positive outcome on the existing situation. This could include for example, an improvement in vegetation management or an improvement in air quality as a result of the project.					





The approach to classifying the duration of identified impacts is presented in Table 10.

Table 10: Classifications of the duration of identified impacts

Relative Duration Of Impacts

Temporary	Days to months			
Short Term	Up to one year			
Medium Term	From one to five years			
Long Term	From five to 50 years			
Permanent/Irreversible	In excess of 50 years			

The likelihood of an impact occurring is assessed as per Table 11.

Table 11: Likelihood of impact

Likelihood of Impacts	Risk probability categories
Highly Unlikely	Highly unlikely to occur but theoretically possible
Unlikely	May occur during construction of the project but probability well below 50%; unlikely, but not negligible
Possible	Less likely than not but still appreciable; probability of about 50%
Likely	Likely to occur during construction or during a 12 month timeframe; probability greater than 50%
Almost Certain	Very likely to occur as a result of the proposed project construction and/or operations; could occur multiple times during relevant impacting period

A risk rating is assigned by assessing significance versus likelihood within a risk matrix. Risk is described as the product of likelihood and significance as shown in Table 12.

Table 12: Risk matrix

Likelihood	Significance								
Likelinood	Negligible	Minor	Moderate	High	Very high				
Highly Unlikely/ Rare	Negligible	Negligible	Low	Medium	High				
Unlikely	Negligible	Negligible Low Low Medium		Medium	High				
Possible	Negligible	Low	Medium	Medium	High				
Likely	Negligible	Medium	Medium	High	Extreme				
Almost Certain	Low	Medium	High	Extreme	Extreme				





The rating of risk assessed in the risk matrix is presented in Table 13.

Table 13: Risk Rating Legend

	An issue requiring change in project scope; almost certain to result in a 'significant' impact on a Matter of National or State Environmental Significance
High Risk	An issue requiring further detailed investigation and planning to manage and reduce risk; likely to result in a 'significant' impact on a Matter of National or State Environmental Significance
Medium Risk	An issue requiring project specific controls and procedures to manage
Low Risk	Manageable by standard mitigation and similar operating procedures
Negligible Risk	No additional management required

After assessing the nature and severity of impacts they are summarised under the following categories:

- Adverse/beneficial;
- Consequential;
- Cumulative;
- Short-term/long term;
- Reversible/irreversible; and
- Predictable/unpredictable.

6.2 Results of impact assessment

Potential impacts related to groundwater and possible mitigation strategies have been outlined in Section 5.0.

An assessment of these impacts is presented in Table 15, based on the mitigation measures proposed in Table 14.





Table 14: Summary of Mitigation Measures

Table 14. Suffillary of wildgation measures						
Impacting Processes	Proposed Mitigation Measures					
Seepage from the dredge placement area towards the Barron River causing increases in salinity in the river.						
Lateral migration of saline water away from the dredge placement area causing impacts on water quality in the upper unconfined aquifer.	Limit the water level in the lake until sufficient dredged material has been placed in the lake to create a low permeability barrier between the saline water in the lake, and the surrounding aquifer.					
Lateral migration of saline water away from the dredge placement area causing increased salinity in near surface soils.						
Elevated groundwater pressures in upper unconfined aquifer causing upward migration of potentially saline water, in areas where higher permeability sandy materials are present at surface.						
Seepage beneath the bund causing increased saturation levels/water logging of surface soils close to the bund.	Geotechnical investigation along the alignment of the wall to identify unsuitable foundation materials for the wall, engineering design to take into account foundation					
Seepage beneath the bund wall adversely impacting on the integrity of the bund wall.	materials, and oversight of construction to ensure that the construction is adapted where necessary to ground conditions encountered on site.					





Table 15: Assessment of impacts

Primary impacting processes		ssment with st nitigation meas Place		Residual assessment with additional (proposed) mitigation measures in place			
	Significance Likelihood of of impact impact		Risk rating	Significance of impact	Likelihood of impact	Risk rating	
Seepage from the dredge placement area towards the Barron River causing increases in salinity in the river.	Negligible	Almost certain	Low	Negligible	Almost certain	Low	
Lateral migration of saline water away from the dredge placement area causing impacts on water quality in the upper unconfined aquifer.	Moderate	Likely	Medium	Minor	Likely	Medium	
Lateral migration of saline water away from the dredge placement area causing increased salinity in near surface soils.	Moderate	Unlikely	Low	Moderate	Unlikely	Low	
Seepage beneath the bund causing increased saturation levels/water logging of surface soils close to the bund.	Minor	Possible	Low	Minor	Unlikely	Low	
Seepage beneath the bund wall adversely impacting on the integrity of the bund wall.	Moderate	Possible	Medium	Moderate	Unlikely	Low	
Elevated groundwater pressures in upper unconfined aquifer causing upward migration of potentially saline water, in areas where higher permeability sandy materials are present at surface.	Moderate	Possible	Medium	Moderate	Unlikely	Low	

Based on the above, the risks associated with potential impacts related to groundwater are assessed to be predominantly low, with a likely minor impact on water quality in the upper unconfined aquifer leading to a moderate risk. With reference to the modelling discussed in Section 5.0, these impacts are likely to be limited to a distance of 80 m to 120 m from the placement area. Further assessments of the impacts are presented in Table 16.





Table 16: Summary of assessed impacts

Element	Adverse impact	Beneficial impact	Consequential impact	Cumulative impact	Short term	Long term	Reversible	Irreversible	Predictable	Unpredictable
Ground- water and surface water	Lateral migration of saline water away from placement area causing increased salinity of the Barron River		Impacts on water quality.		X		X		X	
Ground- water	Lateral migration of saline water away from placement area causing increased salinity of upper unconfined aquifer		Limitations on the potential to locate shallow groundwater bores close to the placement area			Х	Х		Х	
Ground- water and soils	Lateral migration of saline water away from placement area causing increased salinity of near surface soils		Decrease in productivity of agricultural land		х		X		X	
Ground- water	Seepage beneath the bund causing increased saturation levels/water logging of surface soils close to the bund.		Poor trafficability in areas close to the bund.		х		X		X	
Ground- water	Seepage beneath the bund wall adversely impacting on the integrity of the bund wall.		Failure of the bund wall with release of saline water and potential acid sulfate soils.		х		х		Х	
Ground- water	Elevated groundwater pressures in upper unconfined aquifer causing upward migration of potentially saline water.		Poor trafficability, impacts on surface infrastructure, decrease in productivity of agricultural land.		х		Х			×

7.0 MONITORING PROGRAM

As identified in Section 6.0 the main potential impacts on groundwater are:

- Localised increase in groundwater level adjacent to lake during dredged material placement; and
- Changes in groundwater quality (salinity) associated with flow of saline water outwards from the lake.

Groundwater monitoring will be carried out to assess changes in water level and water quality parameters, to assess whether such changes are within the expected range. The proposed groundwater monitoring network will make use of some of the existing monitoring bores at the site, and will also include additional bores located around the perimeter of the lake. The location of the existing and proposed monitoring bores is illustrated in Figure 27.





The groundwater monitoring network will be used to collect both groundwater level and water quality data prior to, during, and after placement of dredged material. Pressure/electrical conductivity transducers will be installed in selected bores to enable near real time monitoring of groundwater level, electrical conductivity and pH and to allow a greater understanding of the natural variability of these parameters. Trigger levels for water level and water quality parameters will be set relative to background values established through the pre-dredging period, and based on the predicted changes in water level and salinity.

Table 17 provides details of the proposed monitoring and sampling for different phases of the program.

A more detailed monitoring plan and approach to establishment of baseline values and trigger values will be developed in the detailed design phase.

Table 17: Proposed groundwater monitoring

Monitoring Phase	Parameter	Sampling Frequency
12 months	Water Level	Hourly (data logger) and manually during monthly sampling events
prior to	Electrical Conductivity and pH	Hourly (data logger) and monthly during sampling events
placement of dredged	Field physicochemical parameters (EC, pH, DO, Redox, Temp)	Monthly during sampling events
material	Major Ions	Monthly
	Metals (Total / Dissolved)	Monthly
	Water Level	Hourly (data logger) and manually during monthly sampling events
During	Electrical Conductivity and pH	Hourly (data logger) and monthly during sampling events
placement of dredged material	Field physicochemical parameters (EC, pH, DO, Redox, Temp)	Weekly during sampling events
material	Major Ions	Weekly
	**Metals (Total / Dissolved)	Weekly
24 months	Water Level	Hourly (data logger) and manually during monthly sampling events
after	Electrical Conductivity and pH	Hourly (data logger) and monthly during sampling events
placement of dredged	Field physicochemical parameters (EC, pH, DO, Redox Temp)	Monthly during sampling events
material	Major lons	Monthly
	**Metals (Total / Dissolved)	Monthly

^{**}The need for on-going metal analysis will be assessed based on background concentrations and exceedances observed during filling. pH will be systematically monitoring and should pH values show a decrease to below 6, them metals testing would be recommenced.





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- Golder Associates Pty Ltd (Golder), 2016, *Stage1B-Groundwater Report Northern Sands*, Prepared for Flanagan Consulting Group, Cairns shipping development project.

9.0 IMPORTANT INFORMATION

Your attention is drawn to the document - "Important Information relating to this report", which is included as Appendix J. The statements presented in this document are intended to advise you of what your realistic expectations of this report should be. The document is not intended to reduce the level of responsibility accepted by Golder Associates, but rather to ensure that all parties who may rely on this report are aware of the responsibilities each assumes in so doing.



Report Signature Page

GOLDER ASSOCIATES PTY LTD

Scott Fidler Principal

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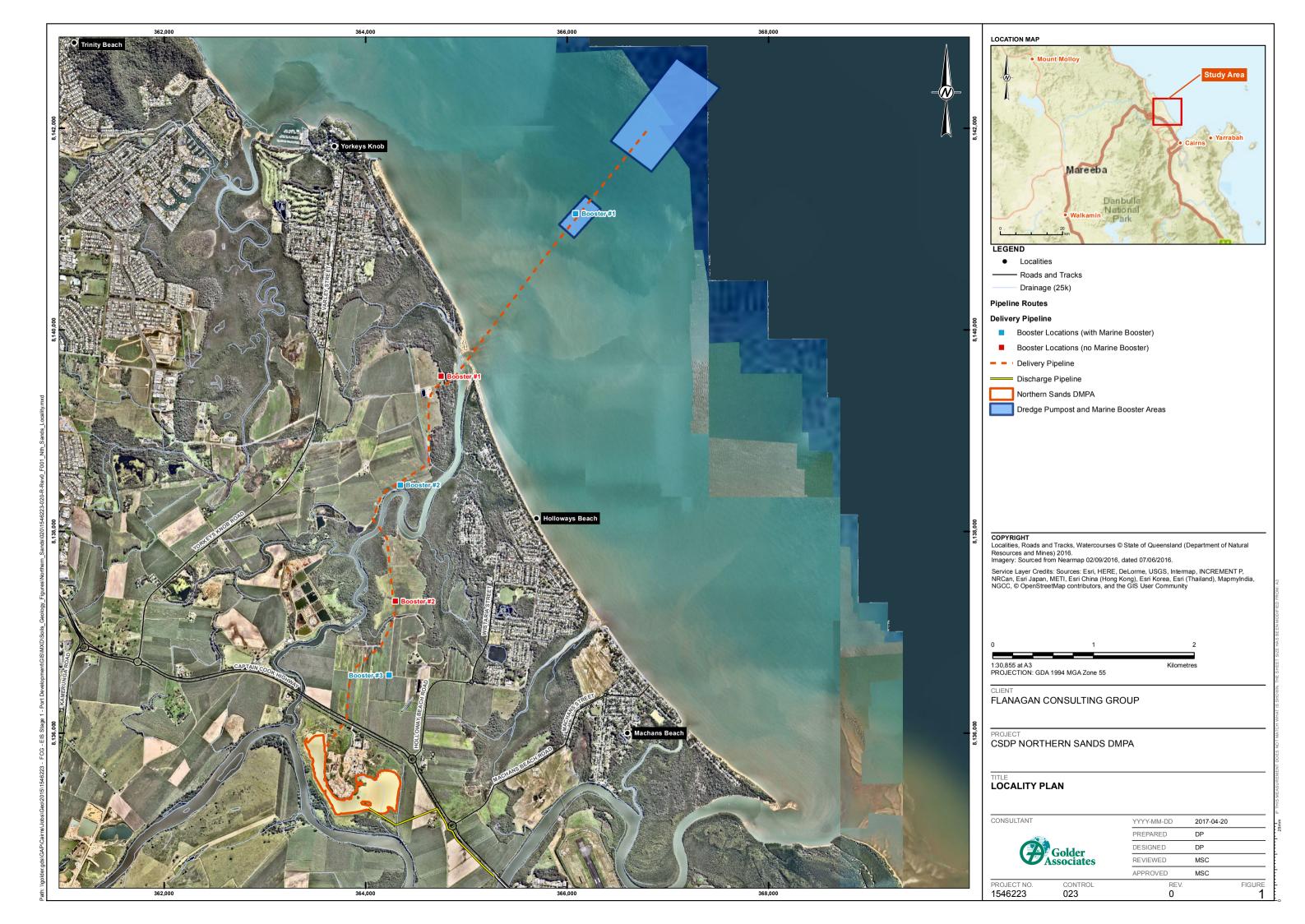
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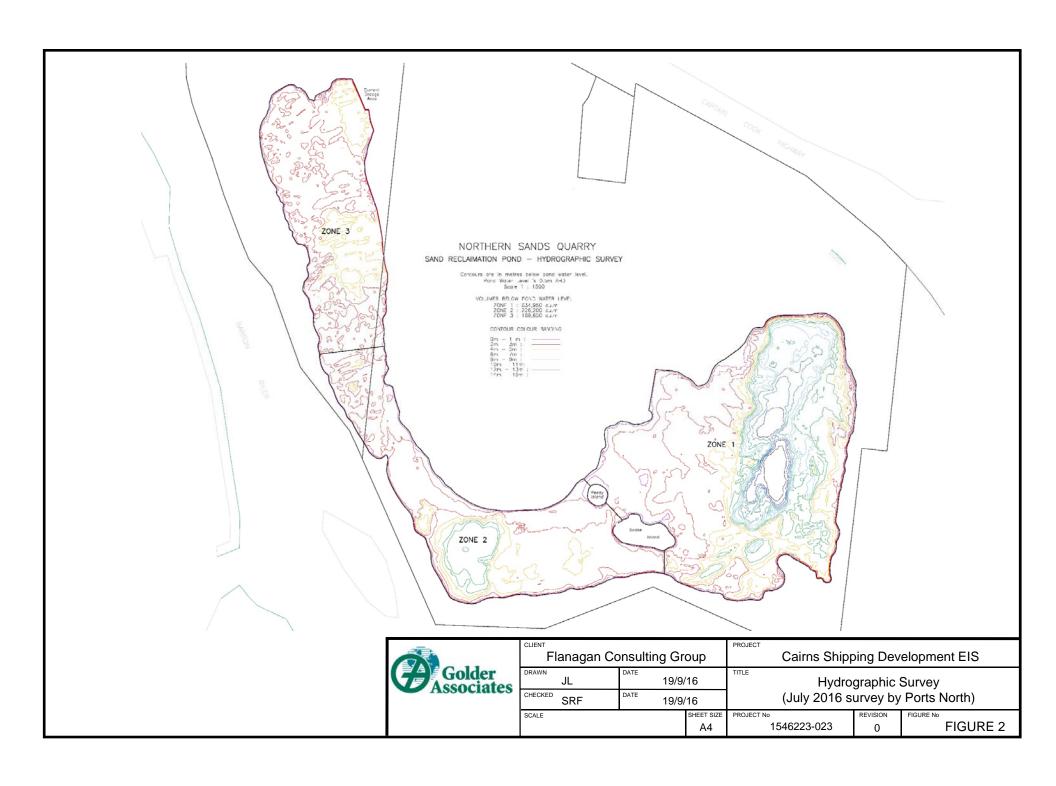


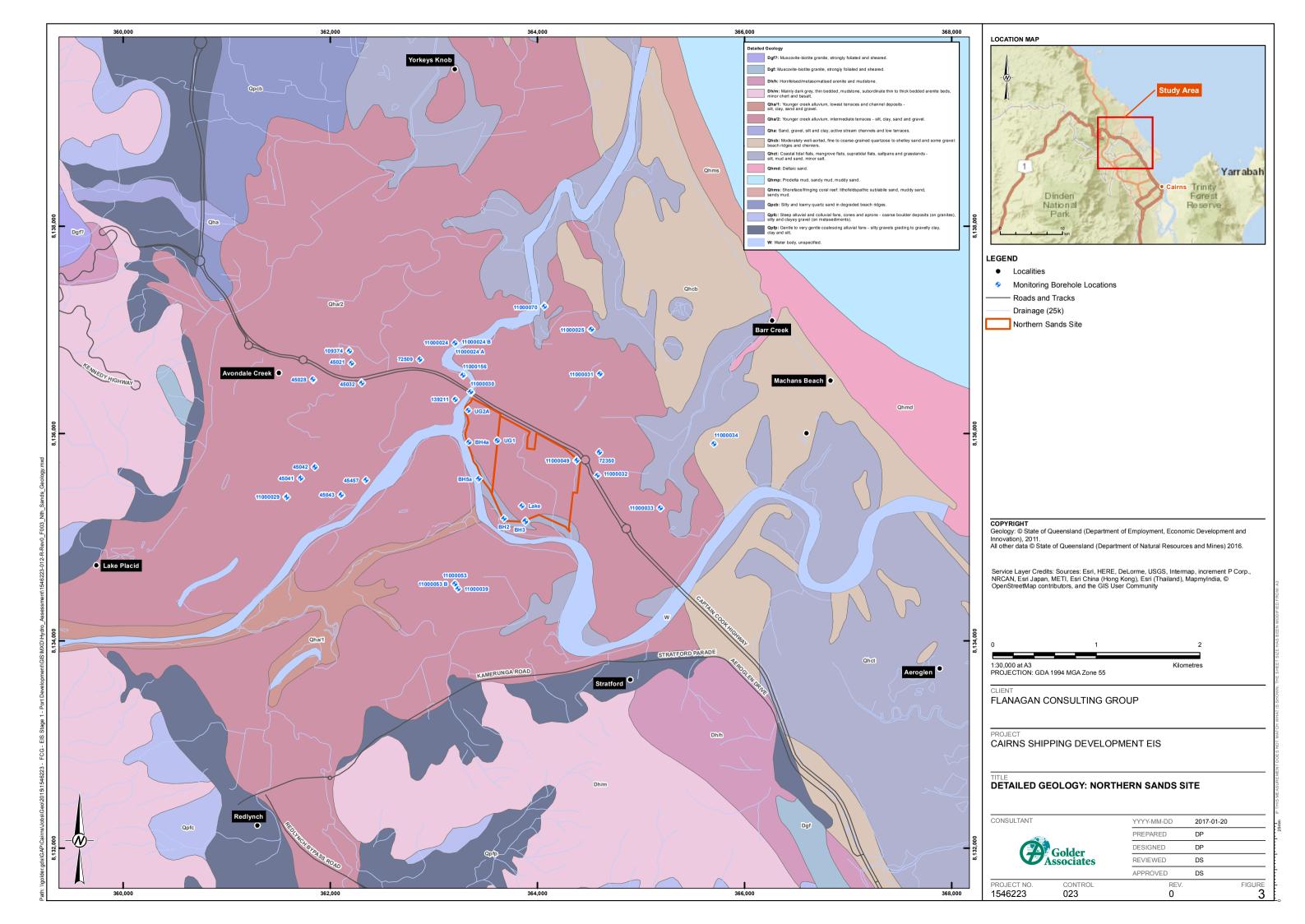


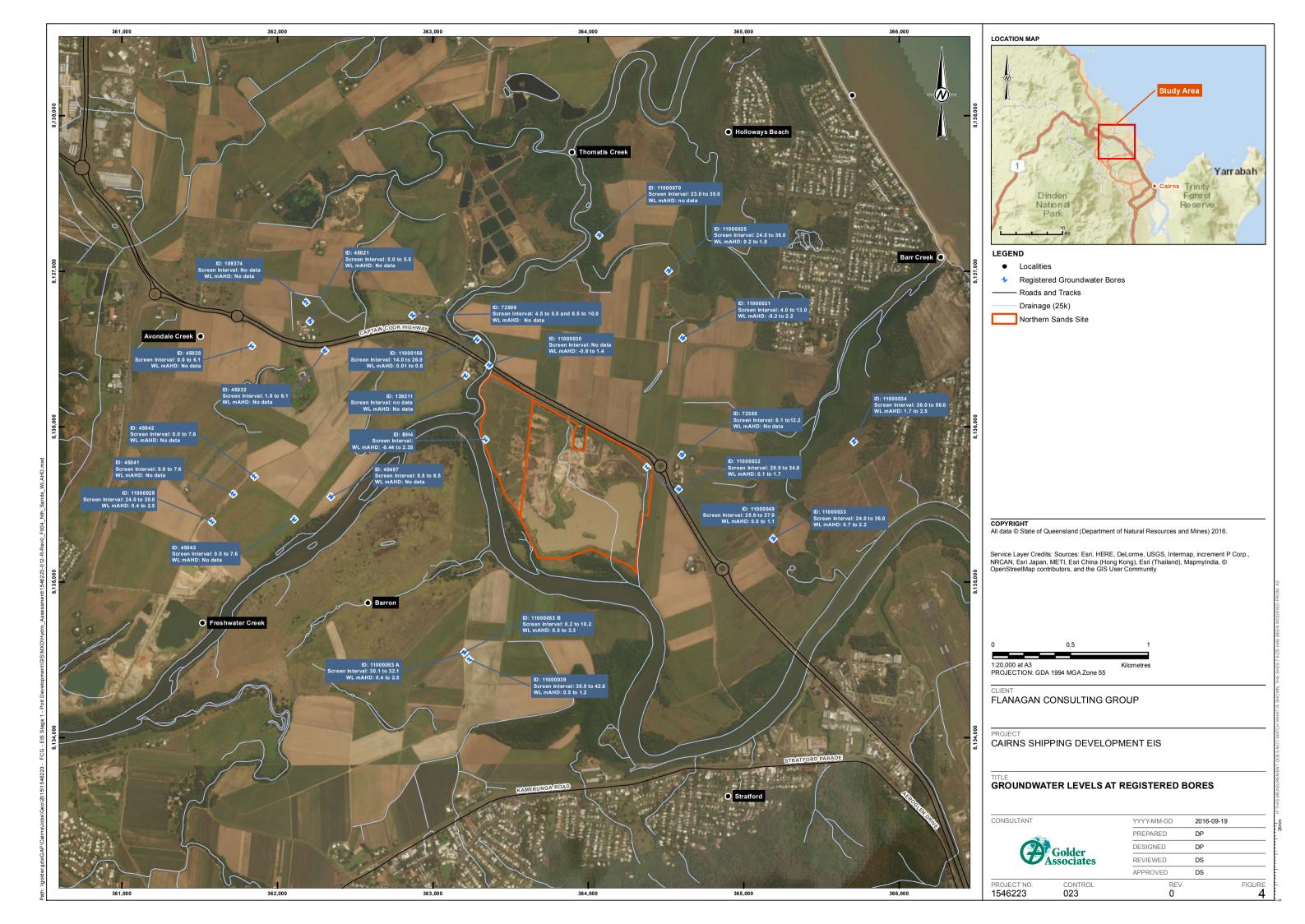
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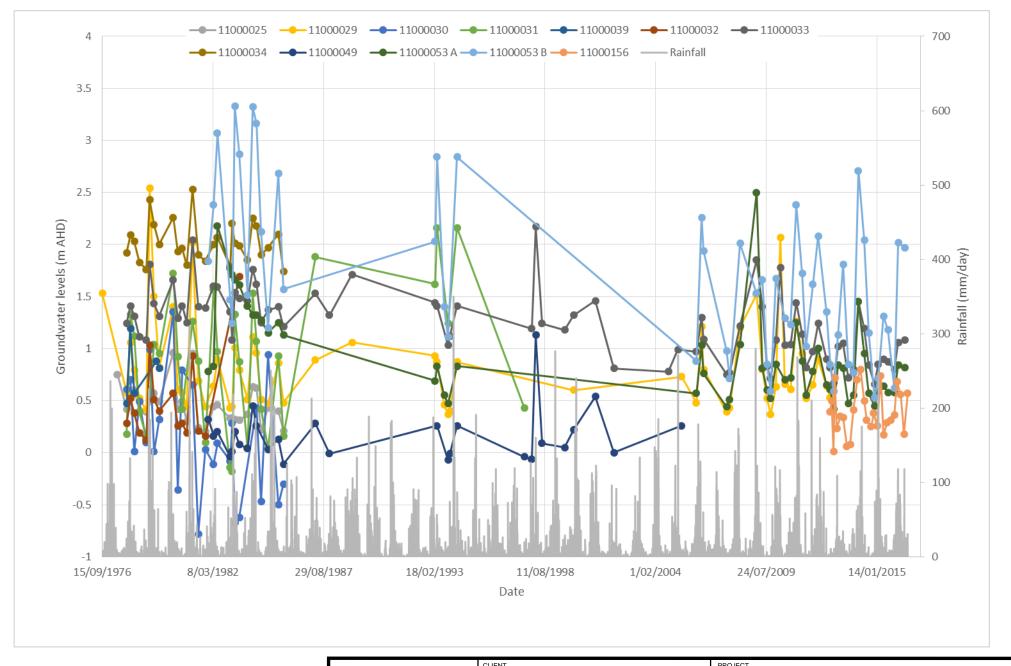




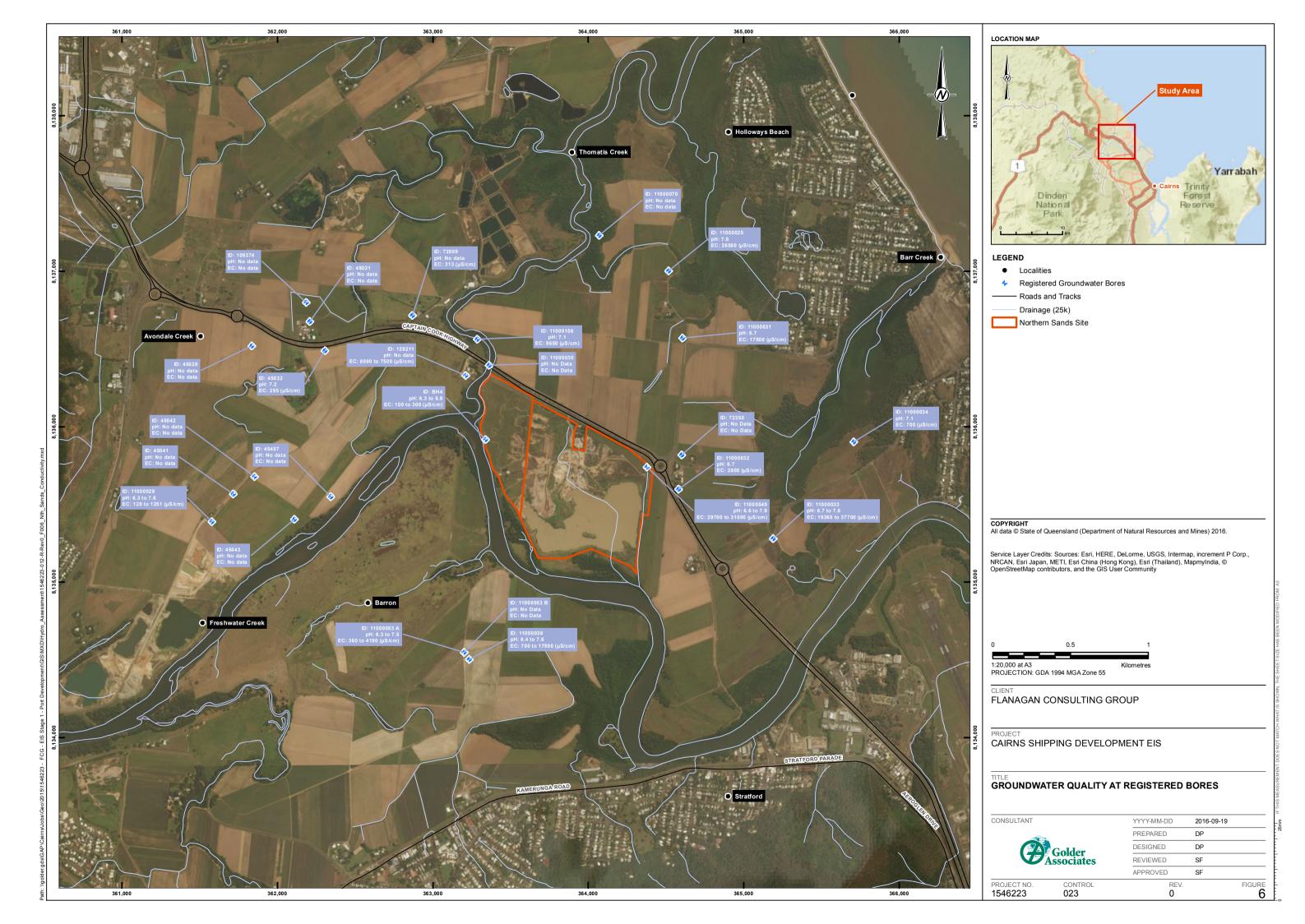


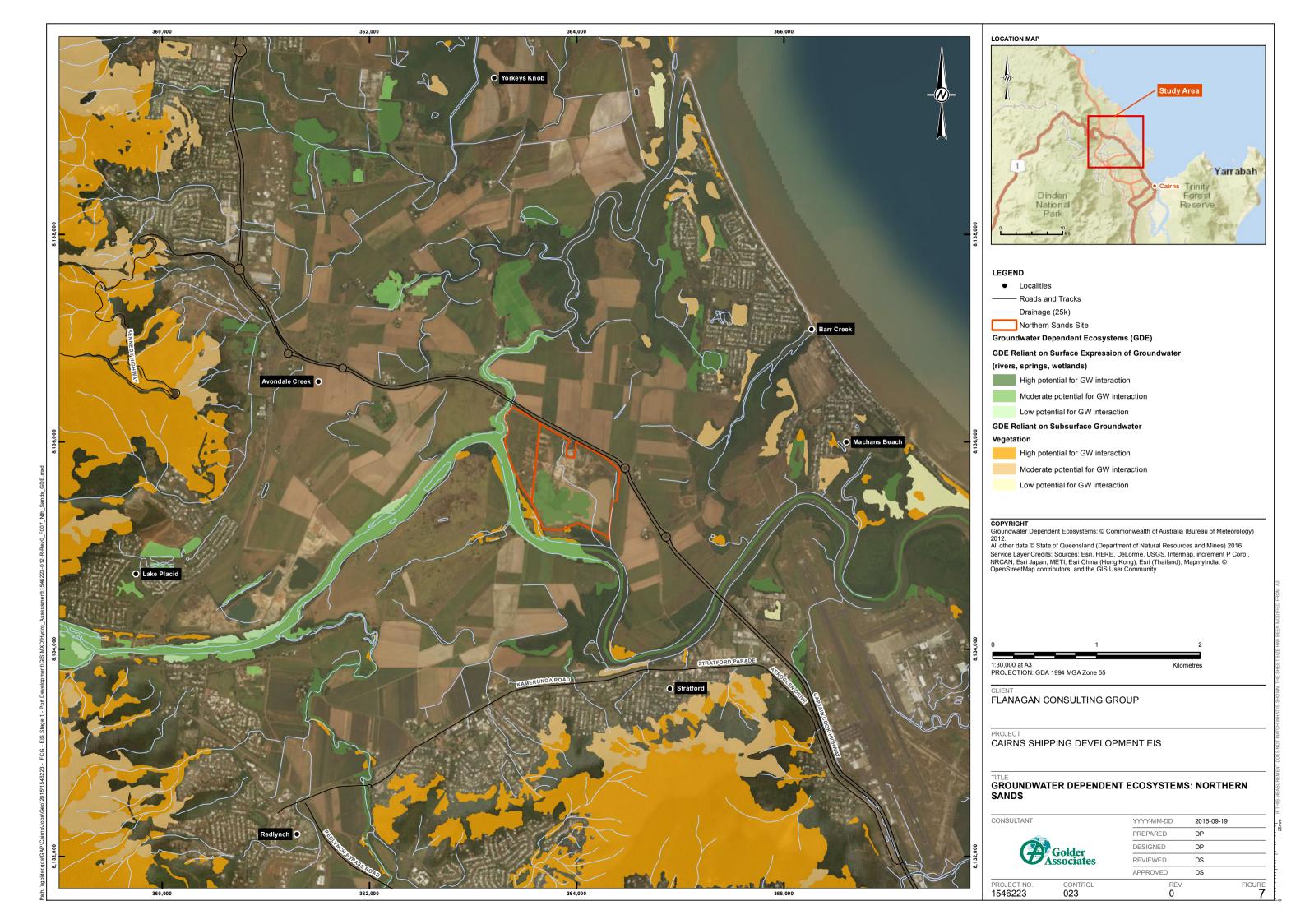


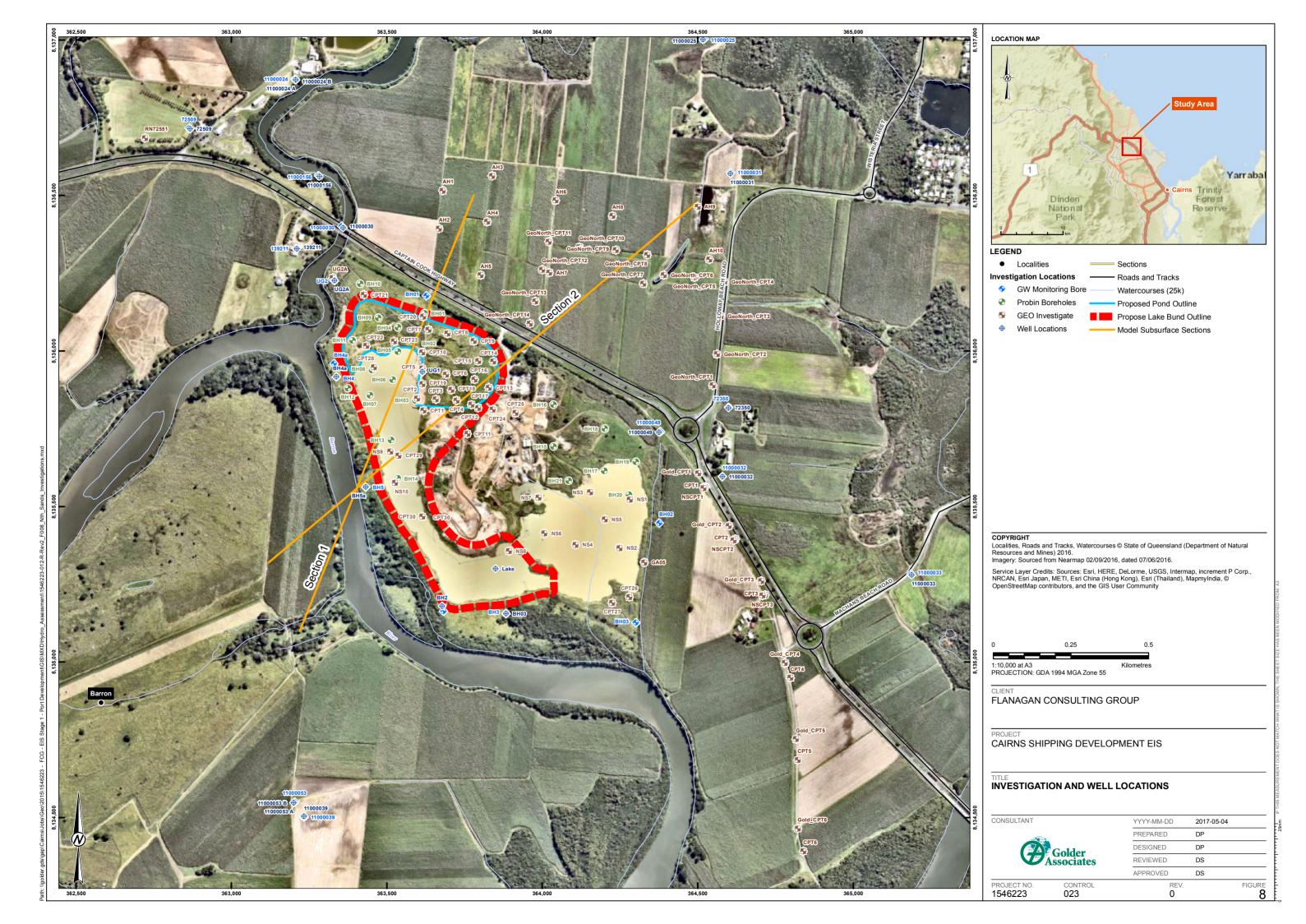


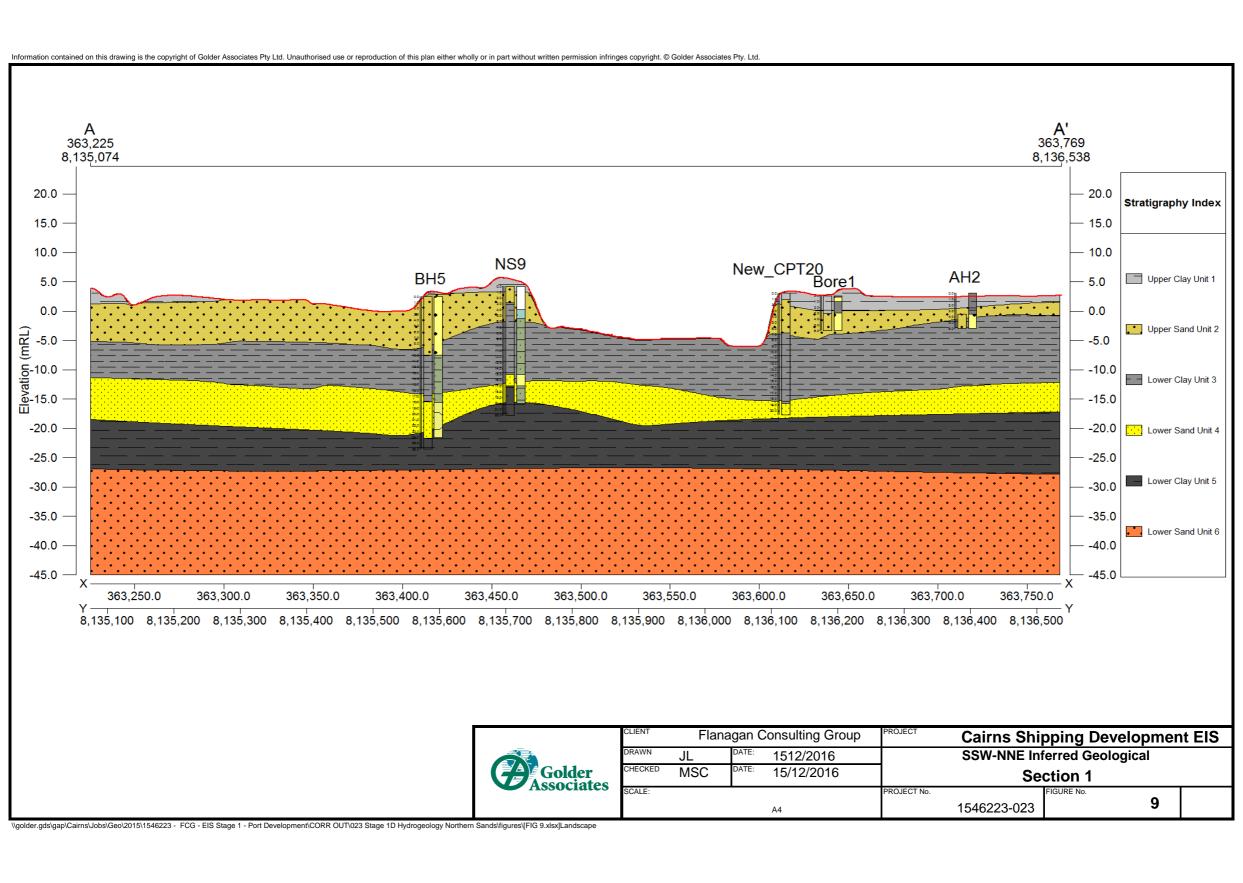


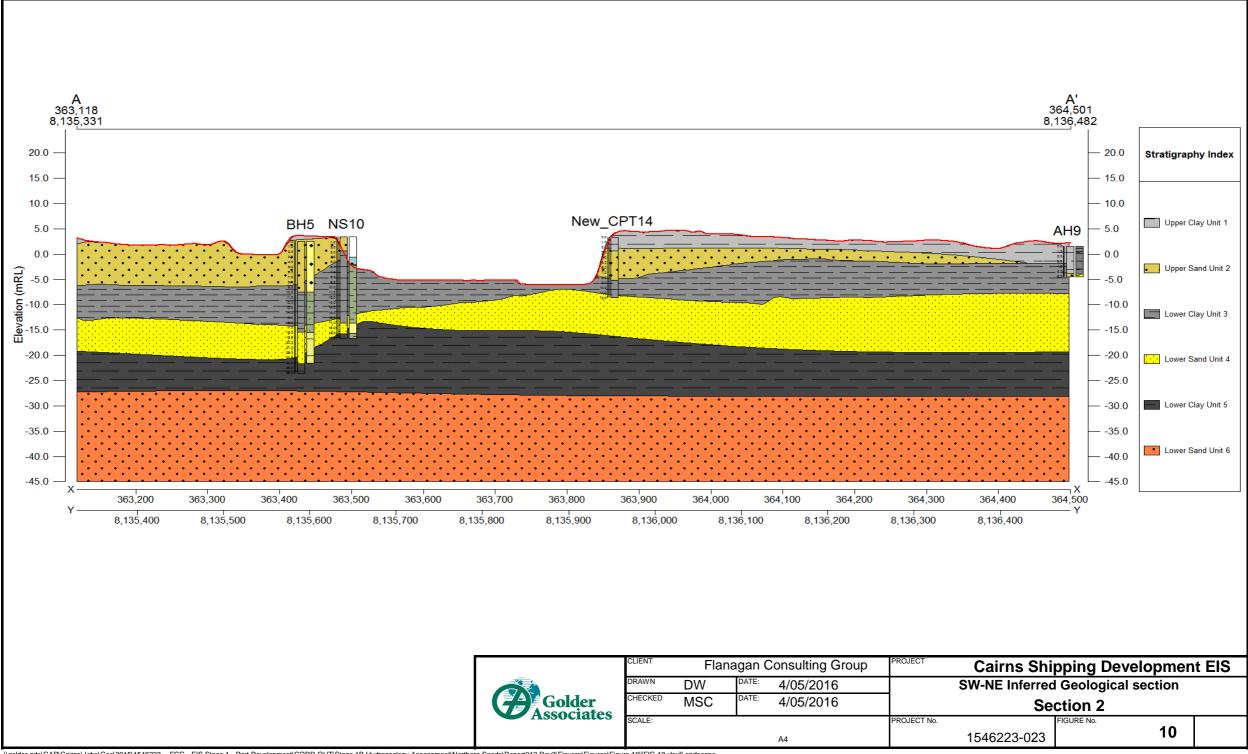
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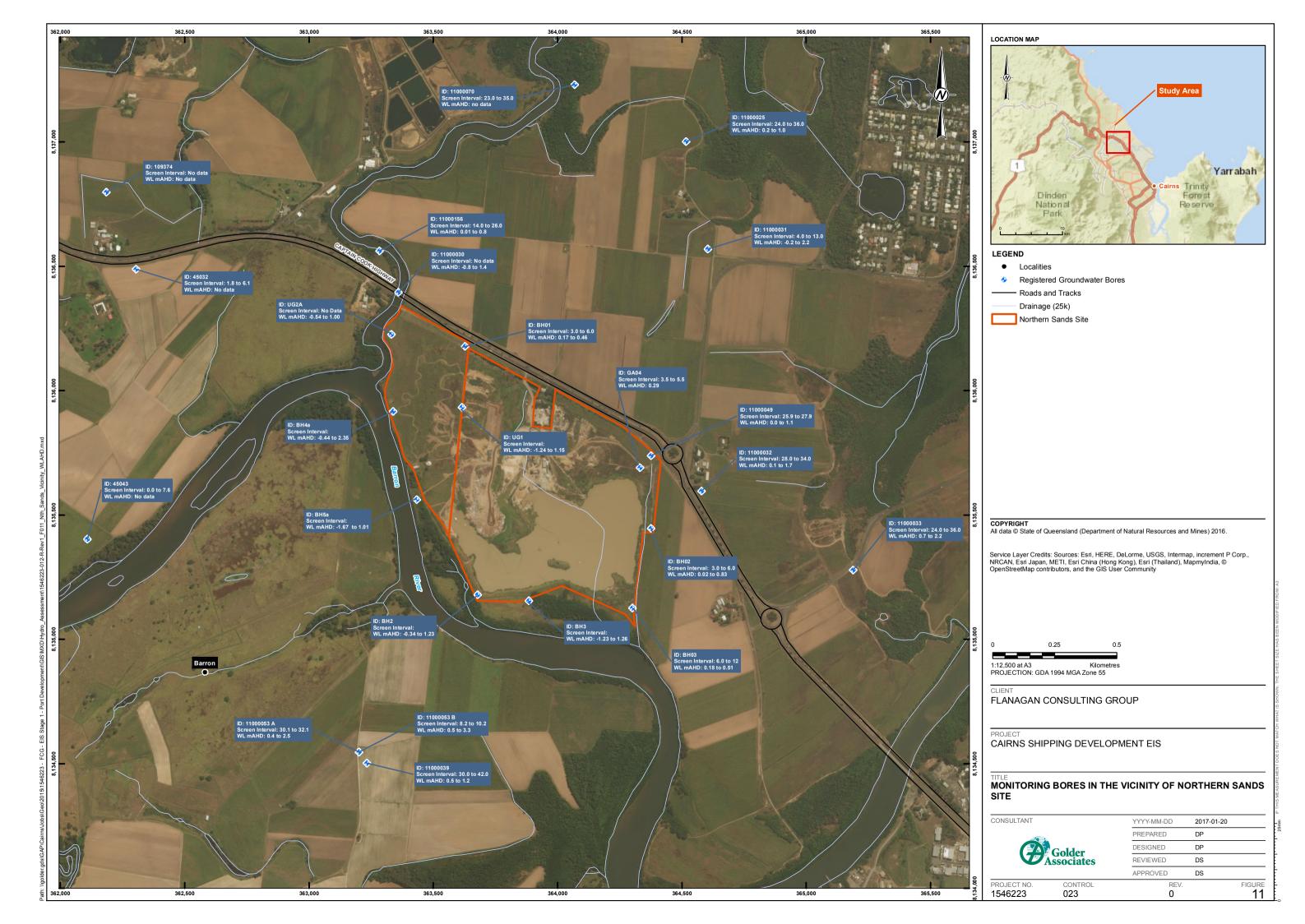


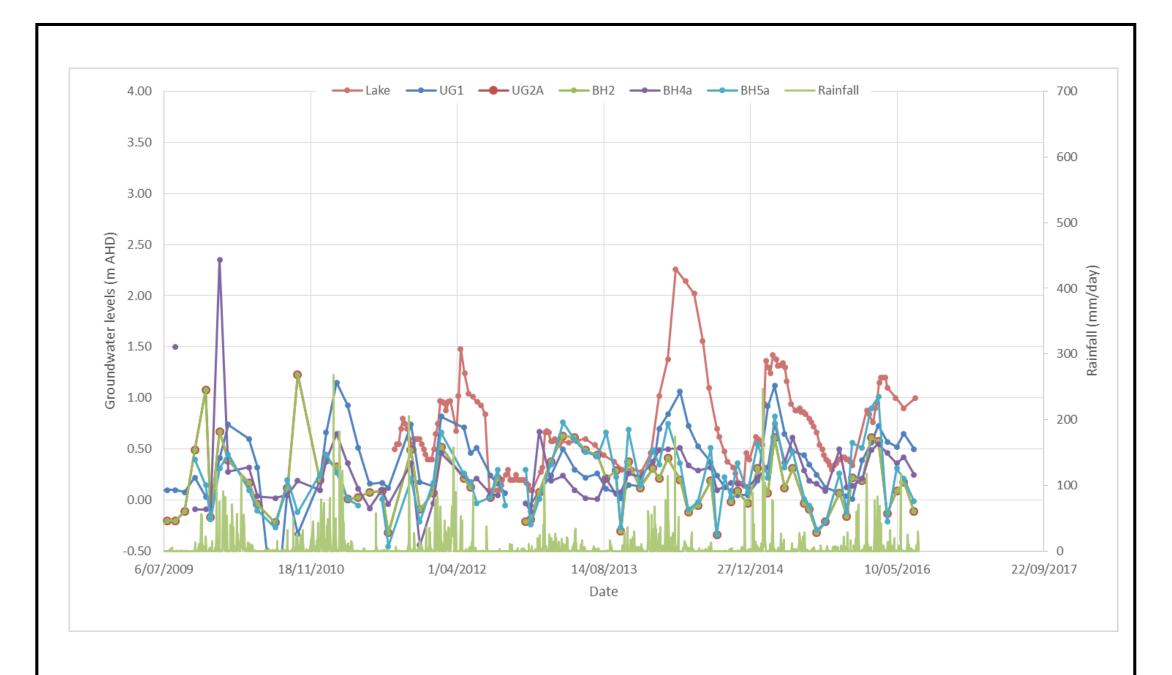




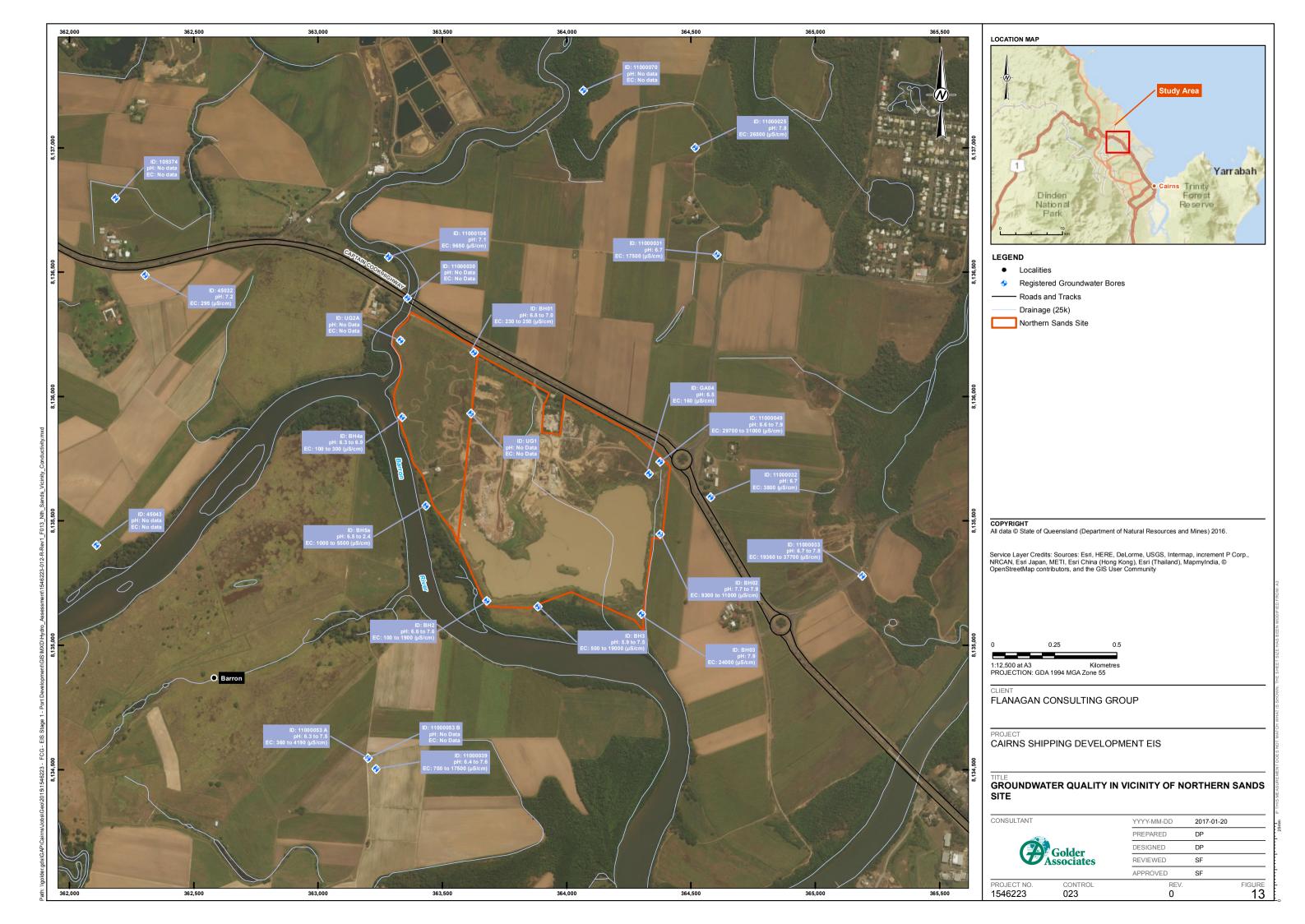


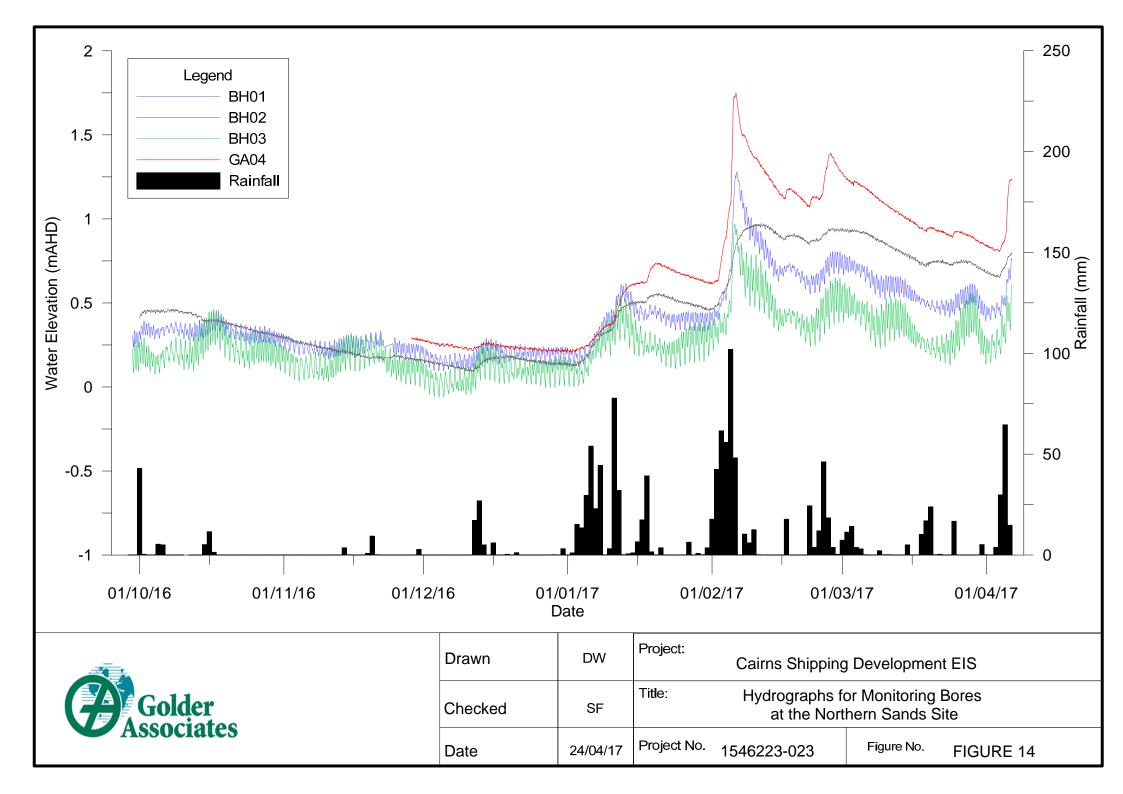






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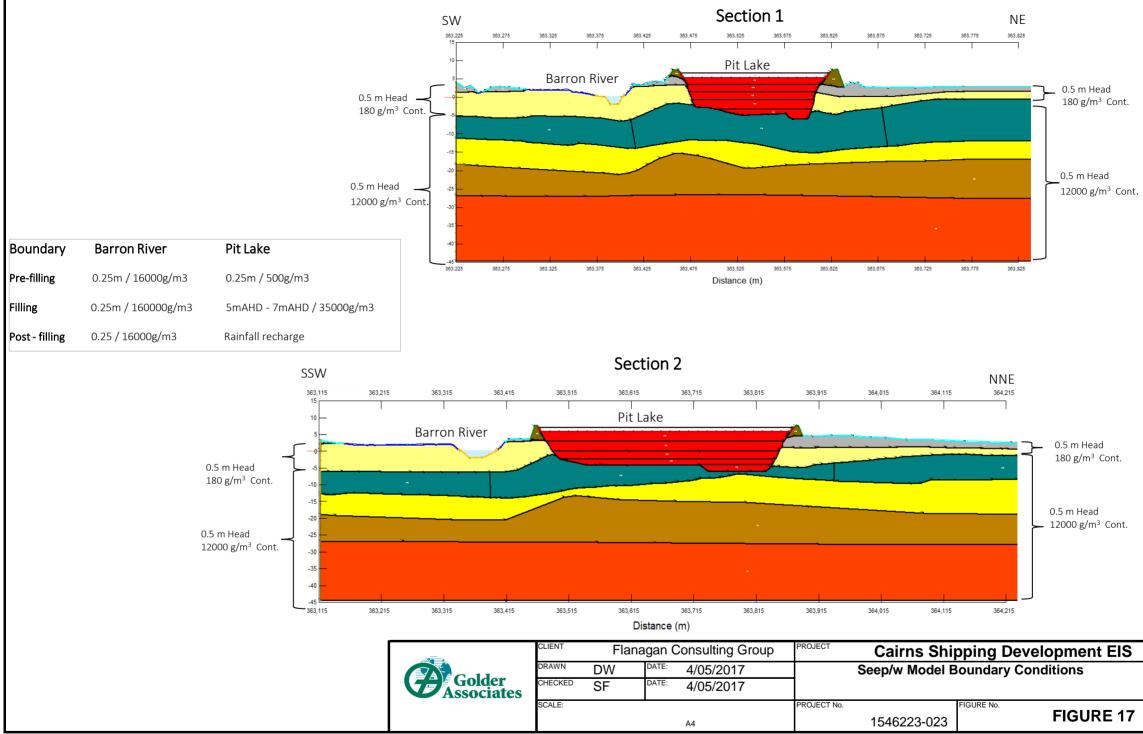




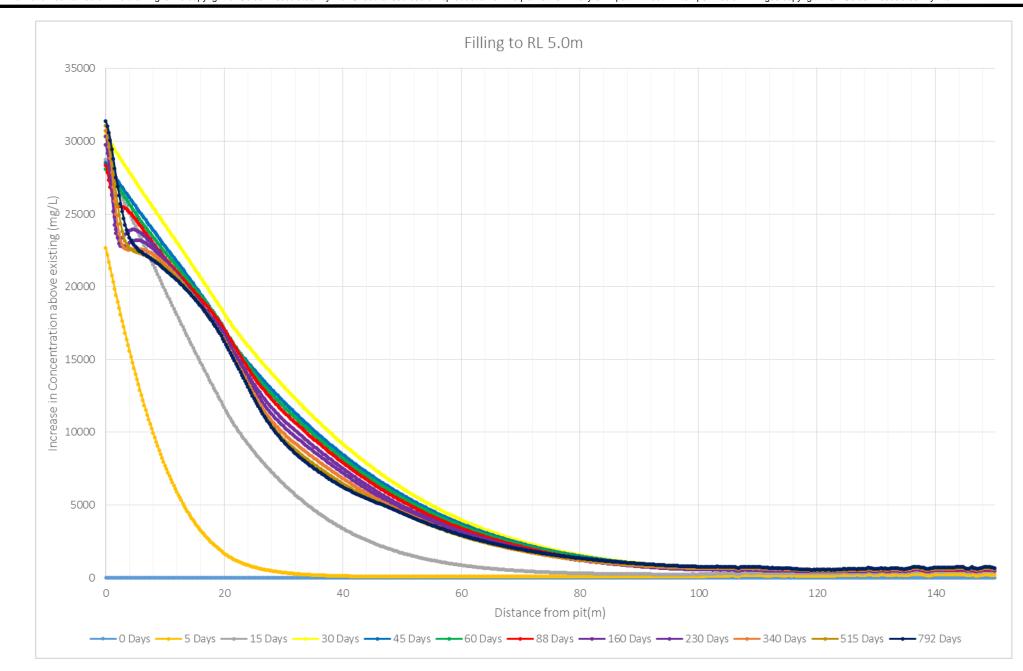
Information contained on this drawing is the copyright of Golder Associates Pty Ltd. Unauthorised use or reproduction of this plan either wholly or in part without written permission infringes copyright. @ Golder Associates Pty. Ltd. NE Section 1 SW 363,225 15 363,475 363,525 363,675 363,725 363,825 Pit Lake Barron River Materials ☐ Upper Sand unit 2 ■ Upper Clay unit 1 Elevation (mRL) Lower Sand unit 4 Lower Clay unit 3 Lower Clay unit 5 Lower Sand unit 6 Dredge material Bund -45⁻ 363,225 363,275 363,325 363,375 363,425 363,475 363,525 363,575 363,625 363,675 363,725 363,775 363,825 Distance (m) SSW NNE Section 2 363,615 364,215 363,115 363,215 363,315 363,415 363,515 363,815 363,915 364,015 364,115 Pit Lake 10 Barron River Elevation (mRL) -25 -30 -35 -45 363,515 363,615 363,715 364,215 363,115 363,215 363,315 363,415 363,815 363,915 364,015 364,115 Distance (m)

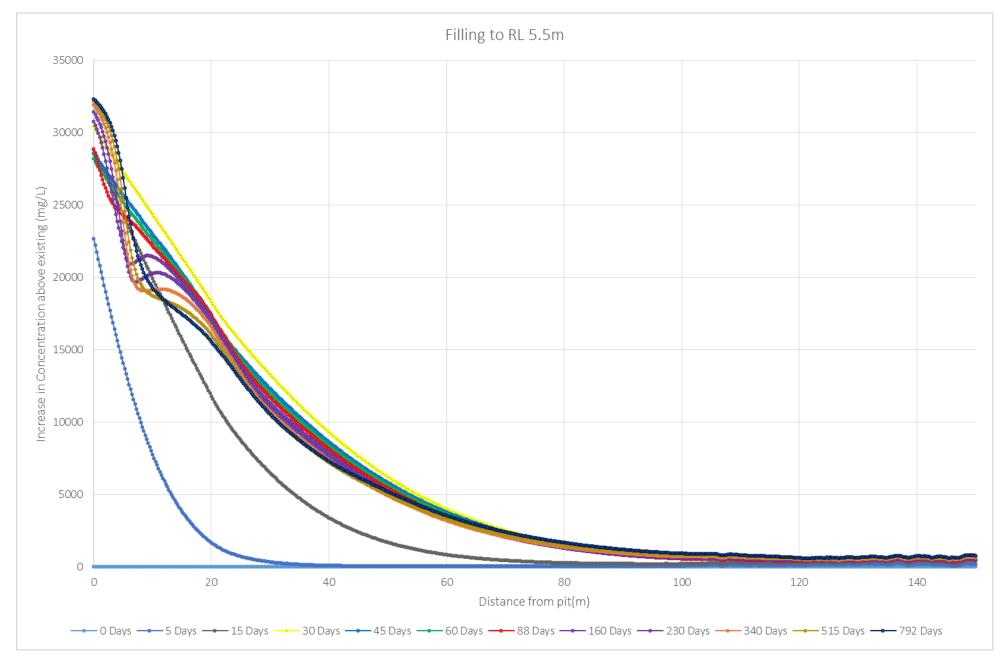
-7 <u>-</u>	CLIENT	Flana	agan C	onsulting Group	PROJECT	Cairns Shi	pping Dev	velopment EIS
Colden	DRAWN	YM	DATE:	1512/2016		Seep/W cross	s-sectional	models
Golder Associates	CHECKED	DB	DATE:	15/12/2016				
	SCALE:				PROJECT No.		FIGURE No.	FIGURE 40
				A4		1546223-023		FIGURE 16

\\golder.gds\GAP\Cairns\Jobs\Geo\2015\1546223 - FCG - EIS Stage 1 - Port Development\CORR OUT\Stage 1B Hydrogeology Assessment\Northern Sands\Report012 Rev3\Figures\Figures\Figures\Figures\Figure 21 seep_W crossection.xlsx]\Landscape



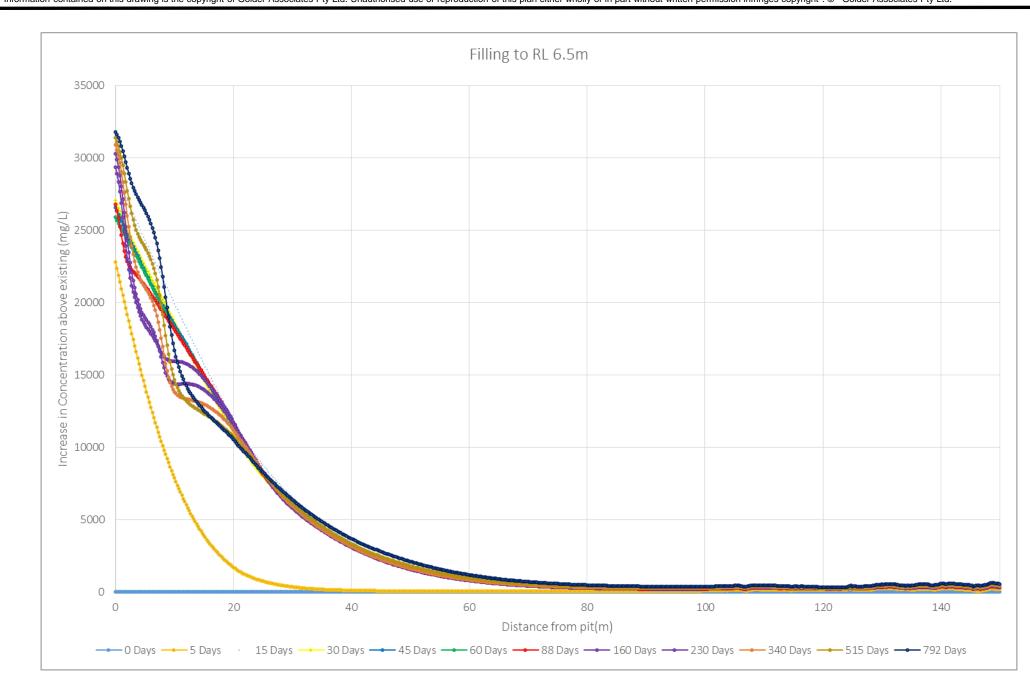
\\golder.gds\\GAP\Cairns\Uobs\Geo\2015\1546223 - FCG - EIS Stage 1 - Port Development\CORR OUT\023 Stage 1D Hydrogeology Northern Sands\figures\\Figures\Figure 17 seep_W Boundary Conditions.xlsx|Landscape

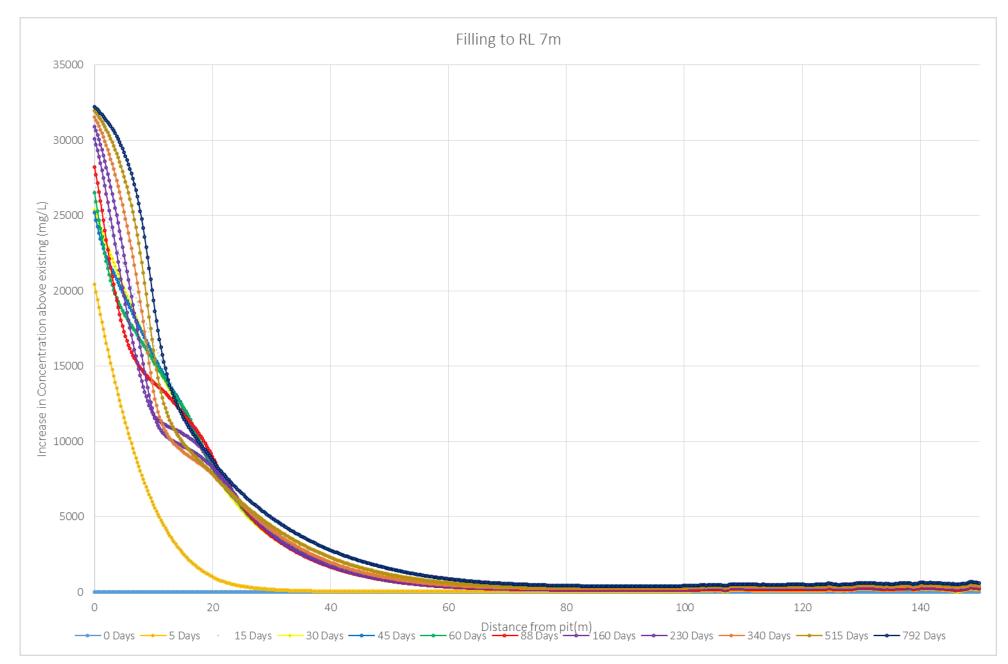




Profiles based on concentrations in the upper sand layer immediately below the upper clay layer

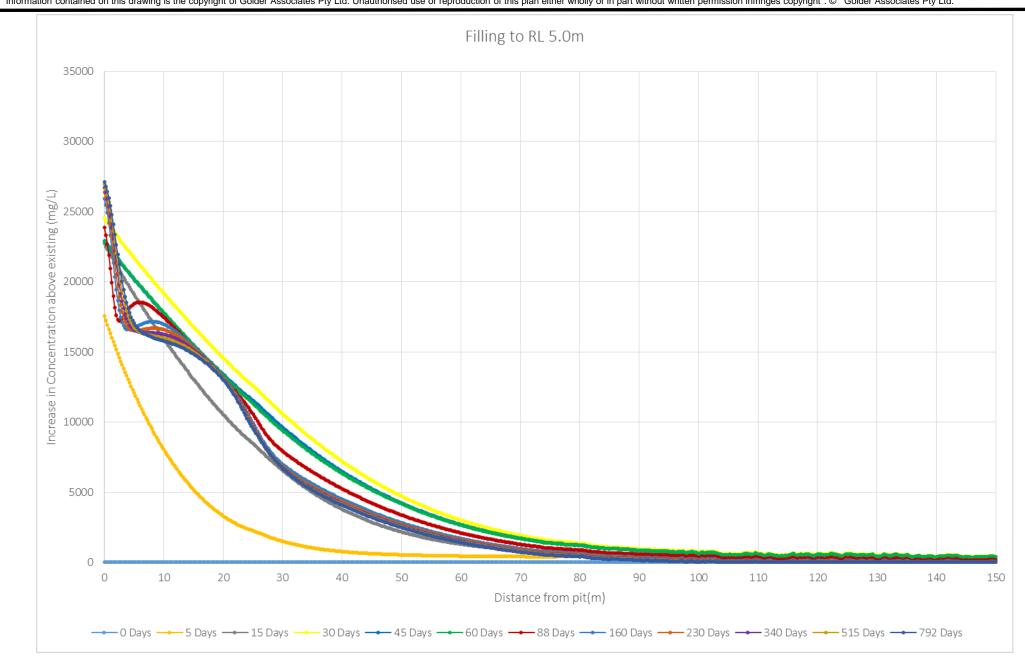
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DRAWN	DW	DATE 04/05/17	TIITLE Profiles of increase in con	centration with	distance
CHECKED	SRF	DATE 04/05/17	from lake – Section	1 (5.0 to 5.5 mF	RL)
SCALE	Not to Sca	le	PROJECT No 1546223-023	FIGURE No 18	nev No A3

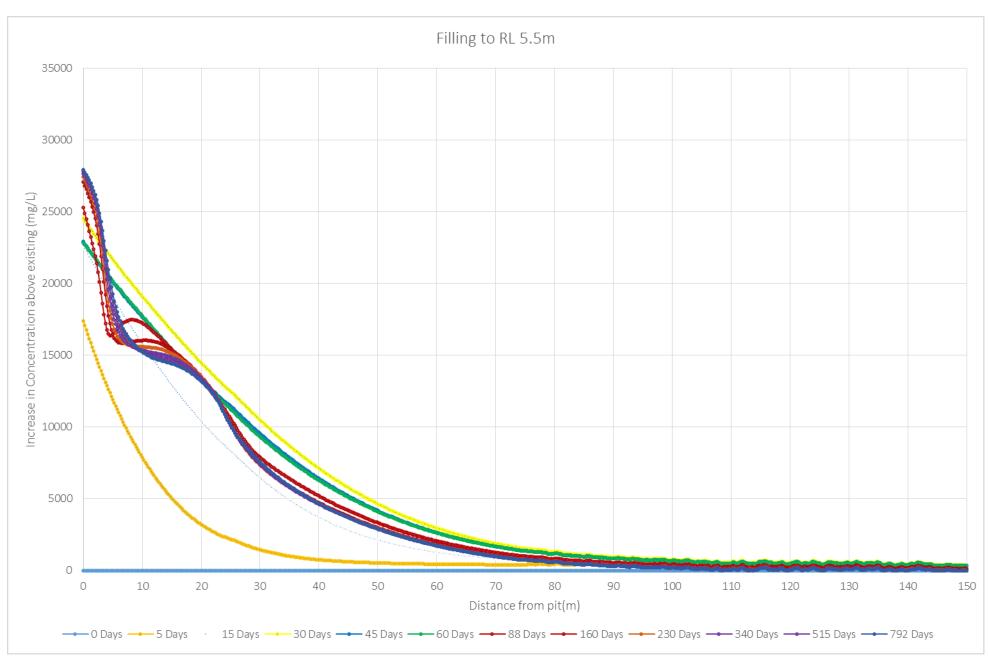




Profiles based on concentrations in the upper sand layer immediately below the upper clay layer

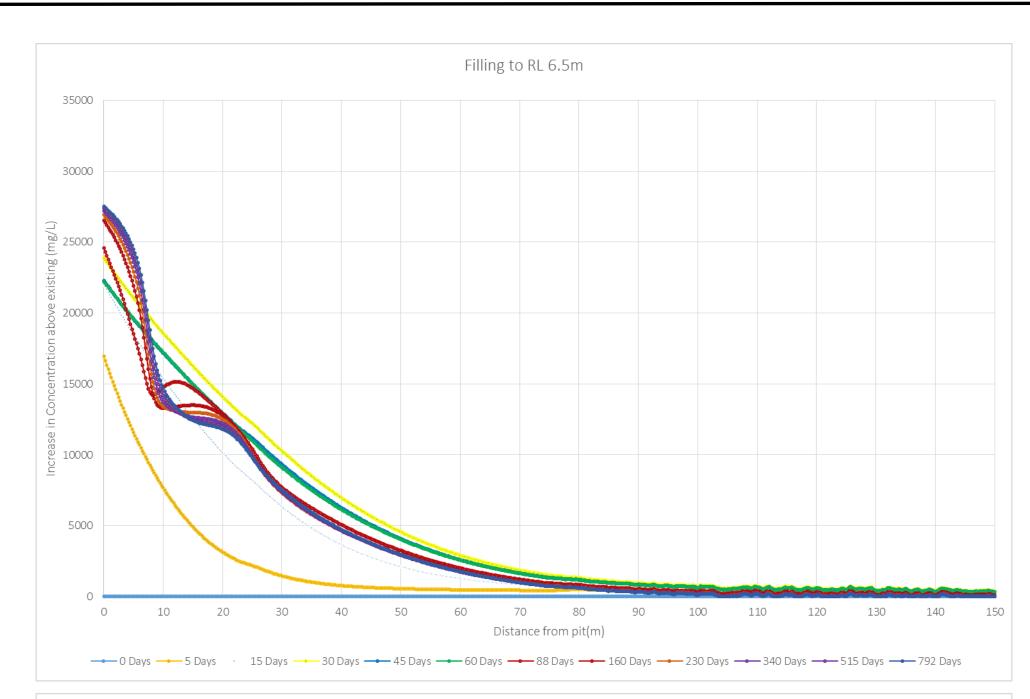
CLIENT FI	anagan Co	nsulting Group	PROJECT Cairns Shipping D	evelopme	nt EIS
DRAWN	DW	DATE 04/05/17	TIITLE Profiles of increase in con	centration	with distance
CHECKED	SRF	DATE 04/05/17	from lake - Section 1	(6.5 to 7.0) mAHD)
SCALE	Not to Sca	le	PROJECT No 1546223-023	FIGURE No	19 REV No A3

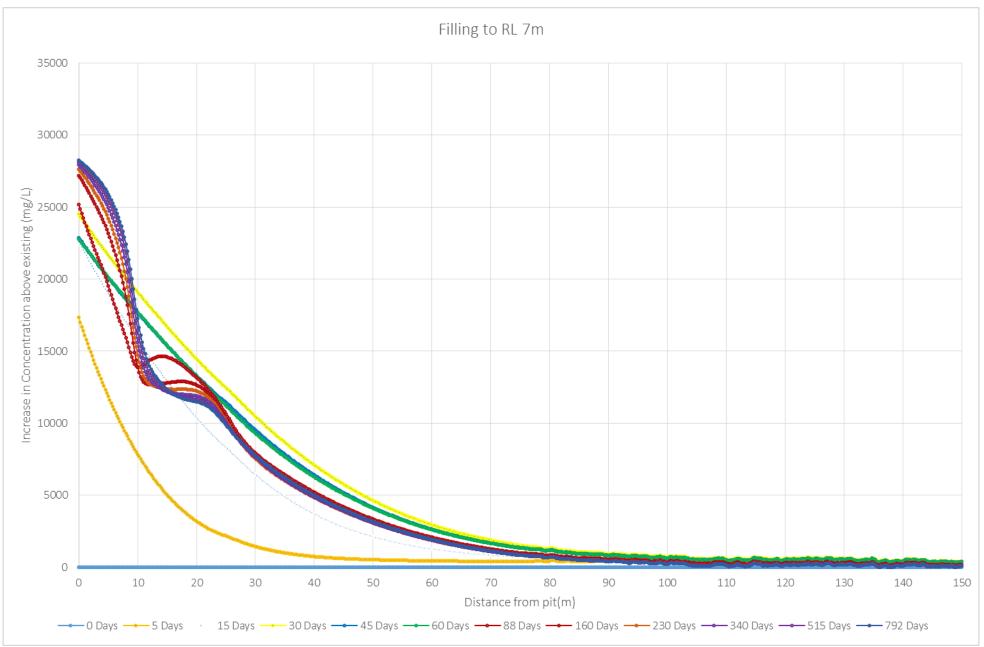




Profiles based on concentrations in the upper sand layer immediately below the upper clay layer

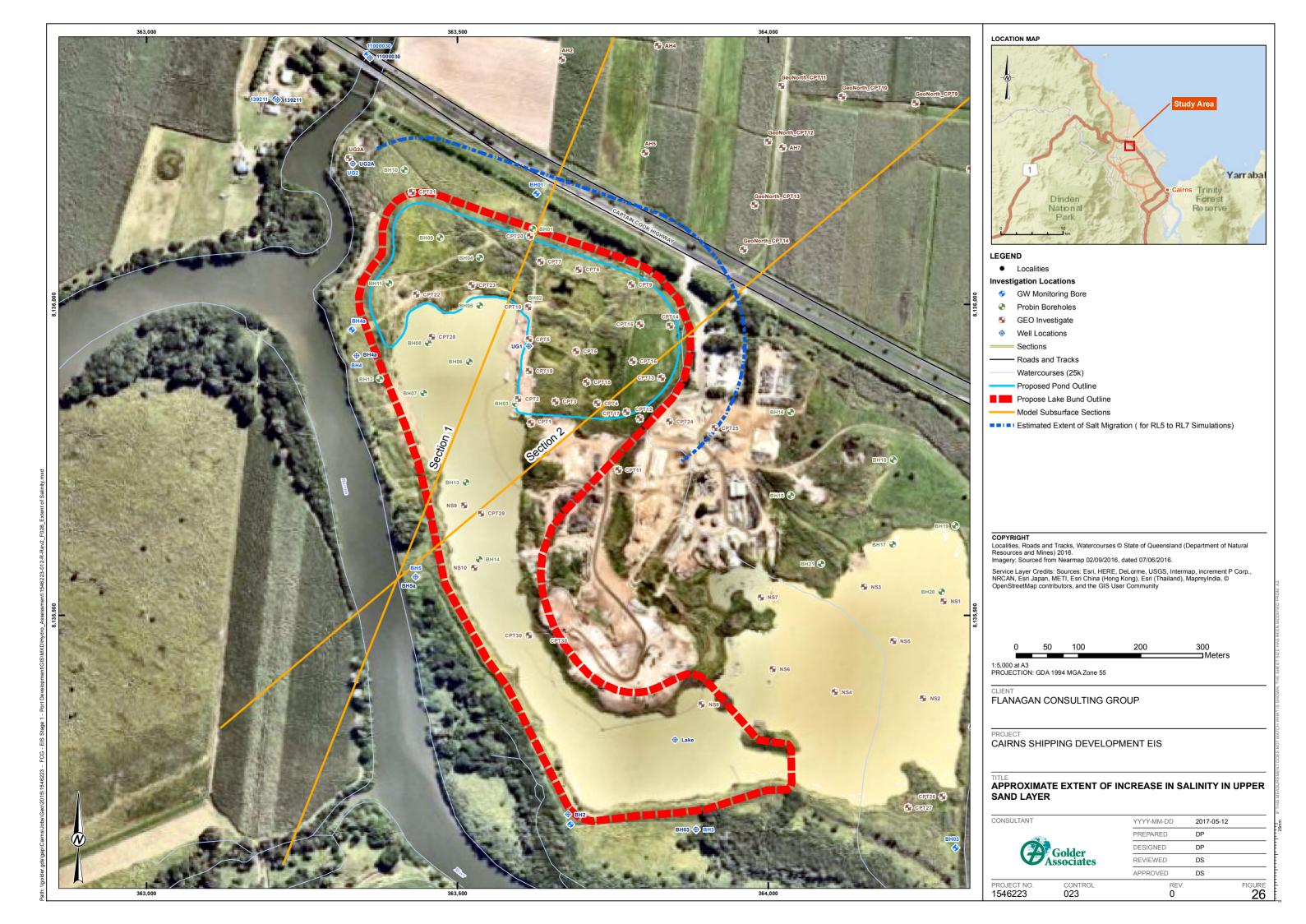
FI	anagan Coi	nsulting Group	Cairns Shipping Development EIS					
DRAWN	DW	04/05/17	TITLE Profiles of increase in con	centration with d	istan	се		
CHECKED	SRF	DATE 04/05/17	from lake - Section 2	2 (5.0 to 5.5 mAHI	D)			
SCALE	Not to Sca	le	PROJECT No 1546223-023	FIGURE No 20	REV No	А3		

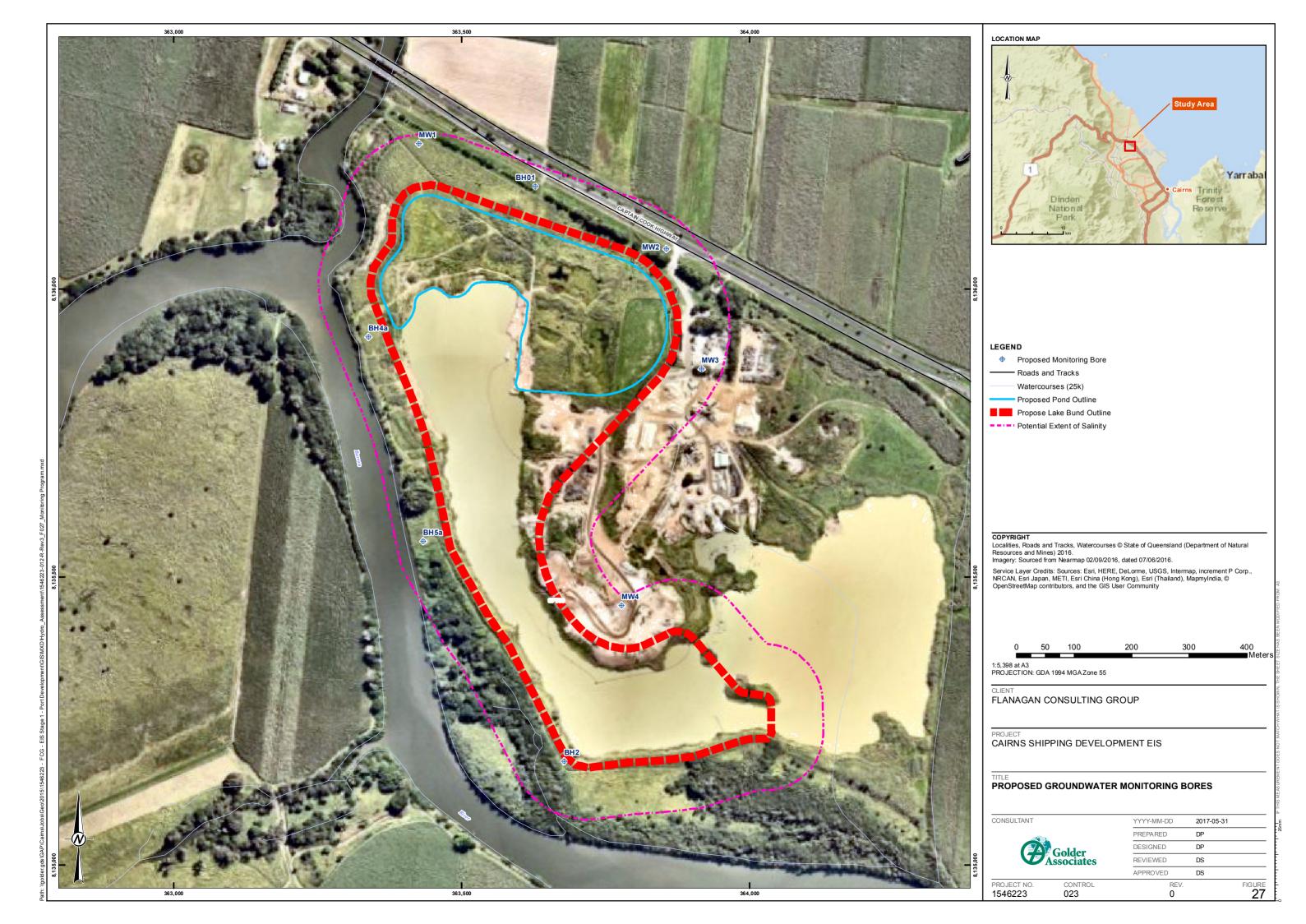




Profiles based on concentrations in the upper sand layer immediately below the upper clay layer

CLIENT	anagan Co	nsulting Group	PROJECT Cairns Shipping D	evelopment EIS	3	
DRAWN	DW	04/05/17	TITLE Profiles of increase in con	centration with c	listan	ce
CHECKED	SRF	DATE 04/051/17	from lake - Section 2	2 (6.5 to 7.0 mAH	D)	
SCALE	Not to Sca	le	PROJECT No 1546223-023	FIGURE No 21	REV No	А3







Appendix A

Summary details of registered groundwater bores





Table A1: Registered bore details.

Registered Borehole ID	Latitude	Longitude	Purpose	Year of installation or drilling	Status
45021	-16.8492	145.7066	Water supply	1974	Existing
45028	-16.8506	145.7031	Water supply	1974	Existing
45032	-16.8509	145.7075	Water supply	1974	Abandoned and destroyed
45041	-16.8592	145.7019	Water supply	1975	Existing
45042	-16.8582	145.7032	Water supply	1975	Existing
45043	-16.8607	145.7056	Water supply	1975	Existing
45457	-16.8594	145.7078	Water supply	1983	Existing
72350	-16.8571	145.729	Water supply	1986	Existing
72509	-16.8489	145.7128	Water supply	1991	Existing
109374	-16.8481	145.7064	Water supply	no data	Existing
139211	-16.8524	145.716	Water supply	2007	Abandoned and destroyed
11000024	-16.8475	145.716	Water resources investigation	1977	Abandoned but still useable
11000025	-16.8464	145.7283	Sub-Artesian monitoring	1977	Abandoned but still useable
11000029	-16.8608	145.7006	Water resources investigation	1976	Existing
11000030	-16.8518	145.7174	Water resources investigation	no data	Abandoned and destroyed
11000031	-16.8503	145.7291	Water resources investigation	1976	Abandoned but still useable
11000032	-16.8591	145.7288	Water resources investigation	1976	Abandoned but still useable
11000033	-16.862	145.7345	Water resources investigation	1976	Existing
11000034	-16.8564	145.7394	Water resources investigation	1977	Abandoned but still useable
11000039	-16.8689	145.7161	Water resources investigation	1977	Abandoned and destroyed
11000049	-16.8578	145.7269	Water resources investigation	1981	Existing
11000053	-16.8685	145.7158	Water resources investigation	1981	Existing
11000070	-16.8443	145.7241	Water resources investigation	1977	Abandoned and destroyed
11000156	-16.8503	145.7167	No data	2010	Existing

Historical groundwater depth information for existing groundwater wells located within 2 km in the surroundings and adjacent to the Northern Sand Site that was available from the DNRM groundwater database is summarised in Table A2.





Table A2: Bore construction details and groundwater levels

	Bore depth	Screened Interval	Screened formation			Standing Water Level Range	
Borehole ID	(m belo w ground level)	(m below ground level)	(m below ground level)	Stratigraphy	Aquifer type	*(m bel ow ground level)	*(m AHD)
45021	5.5	0.0 to 5.5	No data	No data	Shallow	2.4	No data
45028	6.0	0.0 to 6.1	Gravel and sand	Barron river alluvium	Shallow	3.0	No data
45032	6.1	1.8 to 6.1	Gravel and sand	Barron river alluvium	Shallow	3.4	No data
45041	7.6	0.0 to 7.6	No data	No data	Shallow	No data	No data
45042	7.6	0.0 to 7.6	No data	No data	Shallow	No data	No data
45043	7.6	0.0 to 7.6	No data	No data	Shallow	No data	No data
45457	6.5	5.5 to 6.5	Coarse sand	Barron river alluvium	Shallow	2.2	No data
72350	12.2	6.1 to12.2	Coarse sand	Barron river alluvium	Shallow	3.7	No data
72509	11.0	4.5 to 5.5 and 8.5 to 10.0	Mud/coarse sand	Barron river alluvium	Shallow	1.5	No data
109374	No data	No data	No data	No data	NA	No data	No data
139211	66.0	no data	Mud, sand and gravel	Barron river alluvium	Deep	No data	No data
11000024 A	74.0	62.0 to 74.0	Gravel	Quaternary sediment	Deep	1.2 to 4.3	1.0 to 4.1
11000024 B	No data	No data	No data	Quaternary sediment	NA	1.8 to 3.9	1.3 to 3.4
11000025	No data	24.0 to 36.0	Gravel	Quaternary sediment	Deep	1.0 to 1.8	0.2 to 1.0
11000029	30.0	24.0 to 30.0	Gravel	Quaternary sediment	Deep	4.0 to 6.2	0.4 to 2.5
11000030	No data	No data	No data	No data	NA	1.9 to 4.02	-0.8 to 1.4
11000031	85.5	4.0 to 13.0	Gravel	Quaternary sediment	Deep	0.5 to 2.9	-0.2 to 2.2
11000032	88.0	28.0 to 34.0	Gravel	Quaternary sediment	Deep	3.6 to 5.2	0.1 to 1.7
11000033	90.0	24.0 to 36.0	Gravel	Quaternary sediment	Deep	1.3 to 2.5	0.7 to 2.2
11000034	90.0	38.0 to 56.0	Gravel sand and clay	Quaternary sediment	Deep	1.7 to 2.5	1.7 to 2.5



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Borehole ID	Bore depth (m belo w ground level)	Screened Interval (m below ground level)	Screened formation (m below ground level)	Stratigraphy	Aquifer type	Standing Water Level Range	
						*(m bel ow ground level)	*(m AHD)
11000039	72.0	30.0 to 42.0	Gravel	Quaternary sediment	Deep	3.0 to 4.2	0.5 to 1.2
11000049	36.0	25.9 to 27.9	Gravel	Quaternary sediment	Deep	3.2 to 4.4	0.0 to 1.1
11000053 A	42.0	30.1 to 32.1	Gravel	Quaternary sediment	Deep	2.4 to 4.4	0.4 to 2.5
11000053 B	42.0	8.2 to 10.2	Sandy clay and fine to coarse sand	Quaternary sediment	Shallow	1.5 to 4.3	0.5 to 3.3
11000070	73.0	23.0 to 35.0	Sand, gravel and clay	Quaternary sediment	Deep	1.4	no data
11000156	26.0	14.0 to 26.0	Sand, gravel and clay	Barron river alluvium	Deep	2.4 to 4.0	0.01 to 0.8

^{*} Where data is not presented in a range, only a single data point exists.





Appendix B

Water quality data for registered groundwater bores





Water quality parameters for registered bores in the vicinity of the site are available from the Queensland Groundwater Database (GWDB). Bore reports in the database include field parameter measurements for pH and Electrical conductivity (EC), while other analytes are available in some instances where samples have been collected.

Cation and anion concentrations for each groundwater sample were converted to milliequivalents per litre (meq/L) and plotted as percentages of their respective totals in two triangles of the Piper diagram reported in Figure B1 which differentiates groundwater types based on the relative major ion composition.

Table B1 presents the pH, EC and water classification of the registered bores where information is available. Most of the registered bores are classified as a dominant sodium and chloride water types with a mix with sulphate.





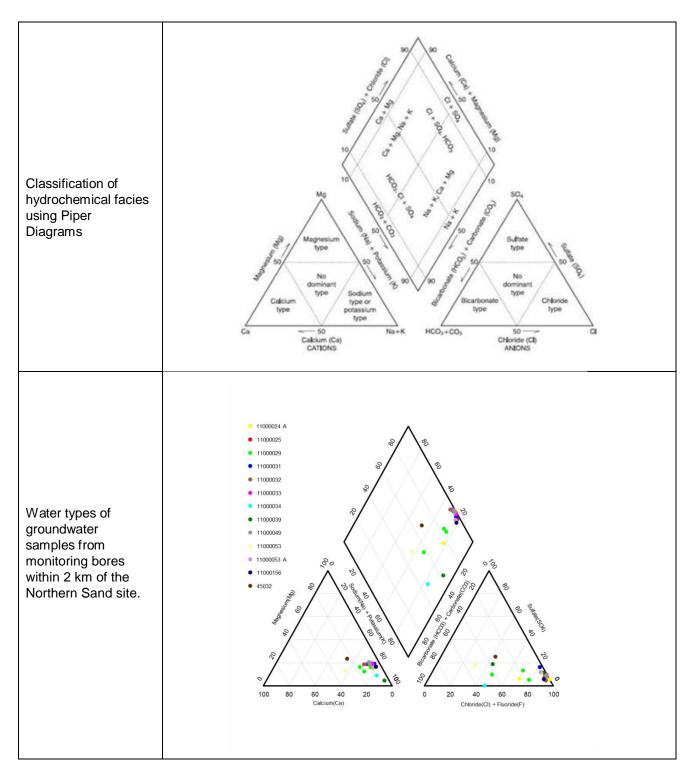


Figure B1: Piper diagram for samples from the Queensland GWDB (DNRM, 2016) within 2 km of the Northern Sand site.





Table B1: pH, EC and water type classification at registered bores

Registered Bore pH range*		EC range (µS/cm)*	Salinity	Water type from piper diagram	
45021	No data	No data	NA		
45028	No data	No data	Potable water	NA	
45032	7.2	295	Neutral and freshwater	CI+S0 ₄ ²⁻ +HC0 ₃ -	
45041	No data	No data	NA	NA	
45042	No data	No data	NA	NA	
45043	No data	No data	NA	NA	
45457	No data	No data	Potable water	NA	
72350	No data	No data	NA	NA	
72509	No data	313	Freshwater	NA	
109374	No data	No data	NA	NA	
139211	No data	6 000 to 7 500	Slightly brackish	NA	
11000024A	6.7 to 7.5	656 to 34 000	Slightly acidic/neutral and freshwater to highly brackish	CI+S0 ₄ ²⁻ +HC03 ⁻	
11000024B	No data	No data	NA	NA	
11000025	7.8	26 500	Neutral, highly brackish	Cl+ Na+ S0 ₄ ²	
11000029	6.3 to 7.6	128 to 1 251	Slightly acidic to neutral, freshwater	Cl+S0₄²-+HC0₃-and Na+K, Ca+Mg	
11000030			Slightly acidic to neutral, freshwater		
11000031	6.7	17 500	Slightly acidic and brackish	Na+K, Ca+Mg	
11000032	6.7	3 800	Slightly acidic and slightly brackish	Cl+ Na+ S0 ₄ ²	
11000033	6.7 to 7.8	19 360 to 37 700	Slightly acidic to slightly alkaline and brackish to highly brackish	Cl+ Na+ S0 ₄ ²	
11000034	7.1	700	Neutral and freshwater	Na+K, Ca+Mg	
11000039	6.4 to 7.6	700 to 17 500	Slightly acidic to slightly alkaline and freshwater to brackish	Cl+S0₄²-+HC0₃-and Na+K, Ca+Mg	
11000049	6.6 to 7.9	29 700 to 31 000	Slightly acidic to slightly alkaline and brackish to highly brackish	CI+S04 ²⁻ +HC03 ⁻	
11000053 A	6.3 to 7.5	360 to 4 190	Slightly acidic to neutral and freshwater to slightly brackish	CI+S0 ₄ ²⁻ +HC0 ₃ -	
11000070	No data	No data	NA	NA	
11000156	7.1	9 650	Neutral and slightly brackish	CI+S04 ²⁻ +HC03 ⁻	





Appendix C

Onsite Groundwater Quality Information





GROUNDWATER REPORT NORTHERN SANDS

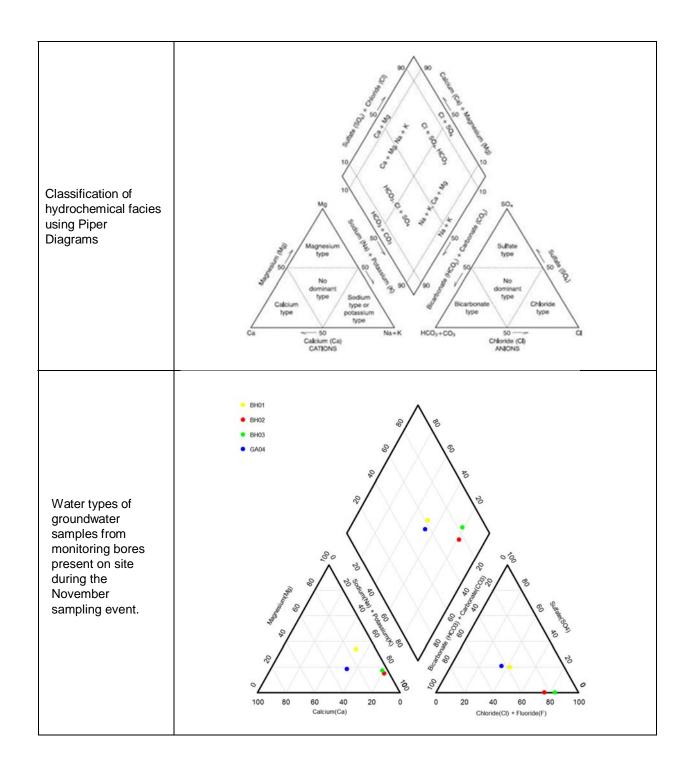


Figure B2: Piper diagram for samples from BH01, BH02, BH03 and GA04.



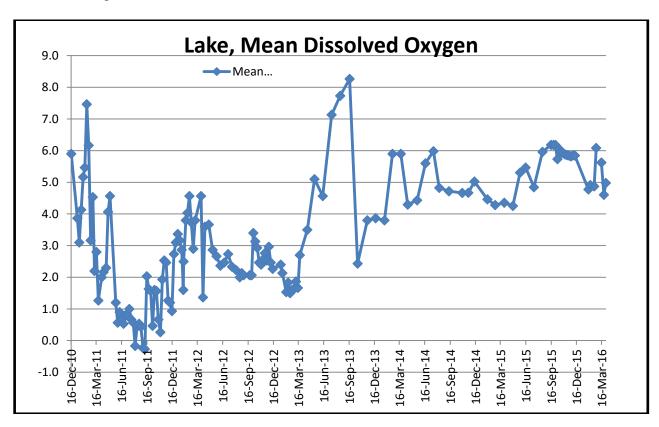
2 June 2017 Report No. 1546223-023-R-Rev0

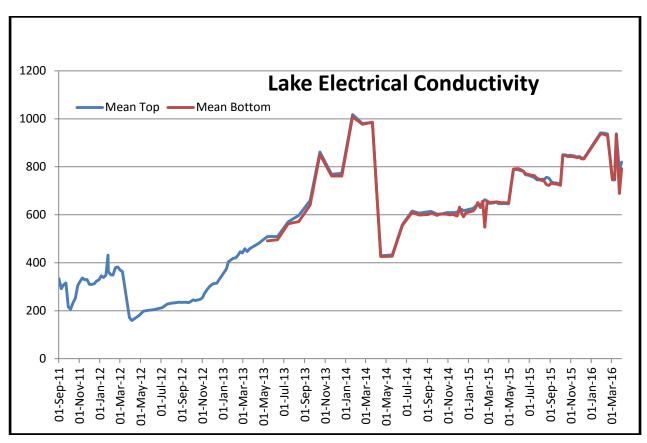
20 July 2016

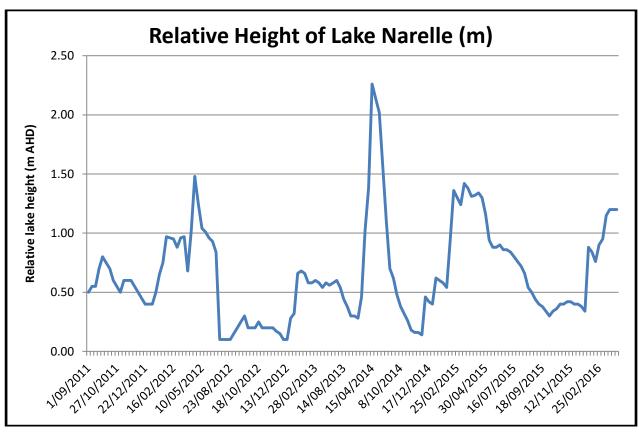
Compiled by Dean Jones of Landline Consulting on request of Tom Hedley for Flanagan Consulting Group representing Ports North.

Northern Sands Lake and Bore Stats.

The first three charts are derived from regular monitoring of the Lake undertaken by Northern Sands using handheld multiparametric water quality metre calibrated by Landline Consulting.

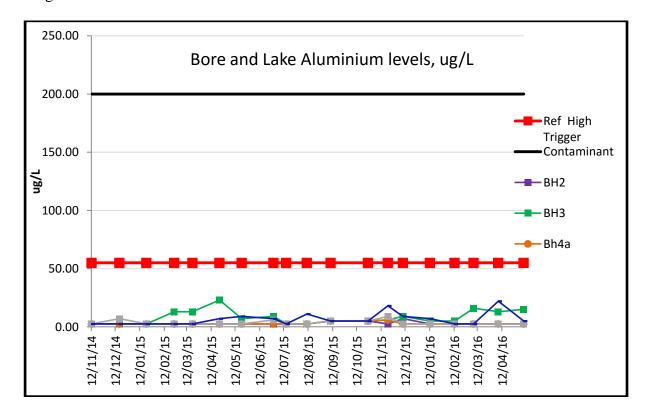


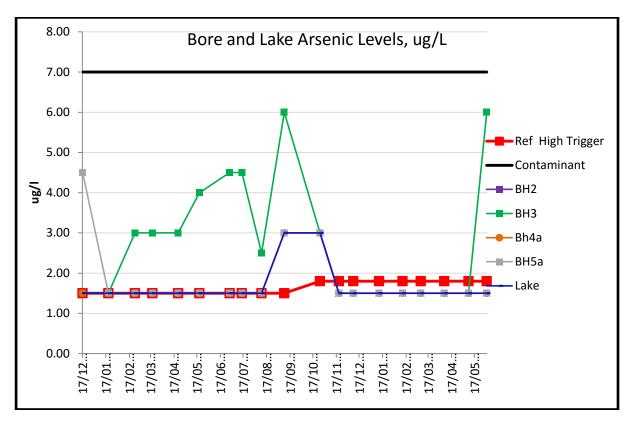


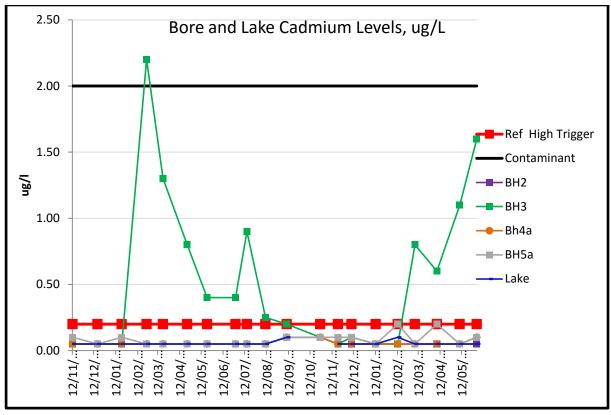


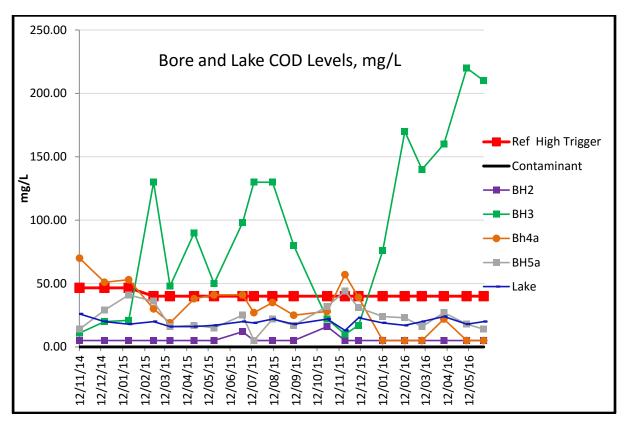
The next group of charts represent samples collected from Northern Sand site bores and lake. Samples are collected monthly and analysed by NATA approved facility. Most charts represent data collected over the last 24 months which is used for examination and interpretation for regular report. Additional historical data, dating back in most cases to 2005, can be provided upon request. However due to the changes in bore numbers over this period it may take some time to collate data.

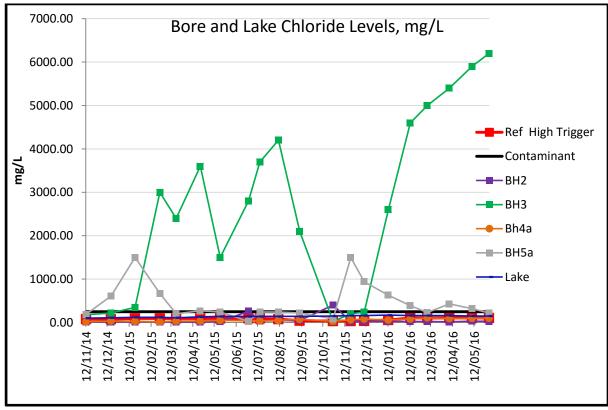
Please note that Bore 3 (BH3) experiences regular exceedences across many parameters. However it's close proximity to the Barron is believed to be responsible for these irregularities and not related to the Lake.

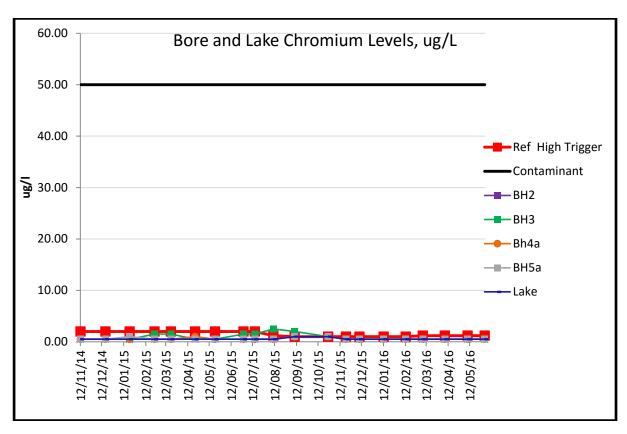


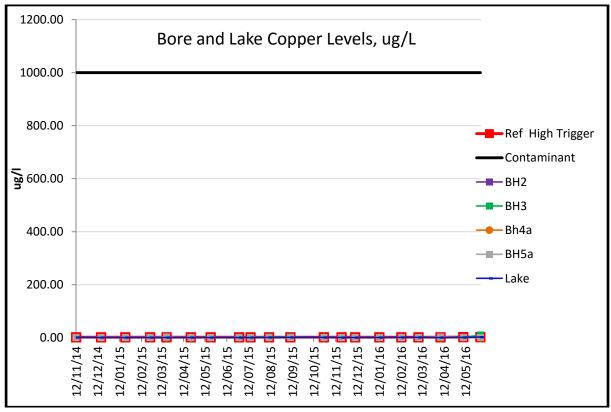


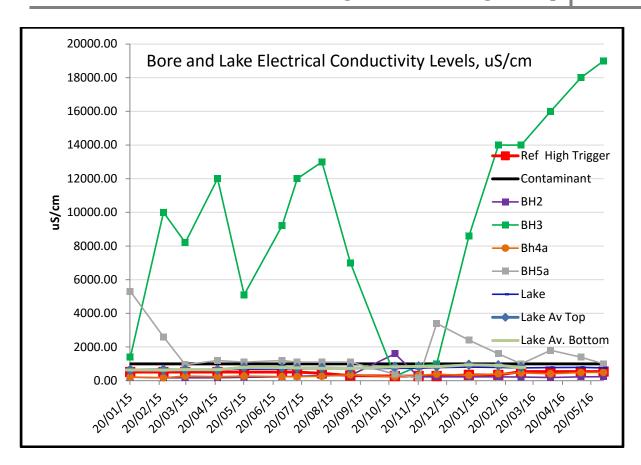


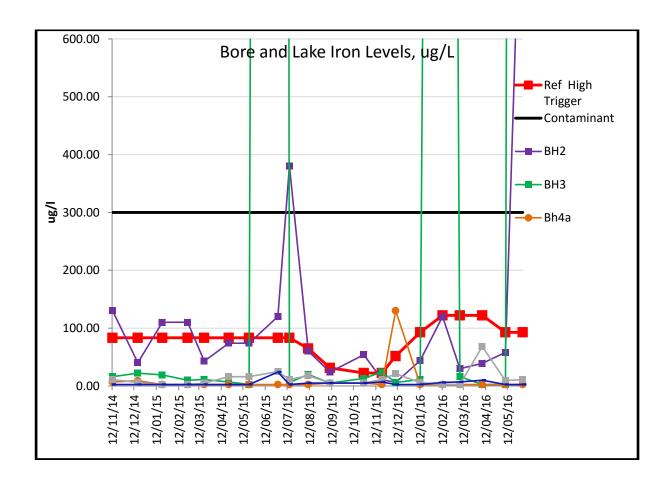


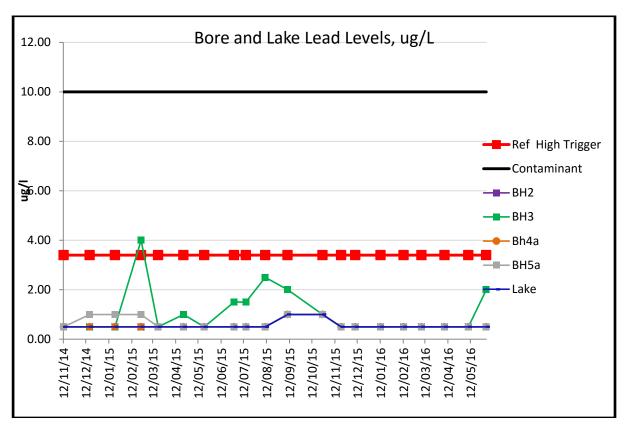


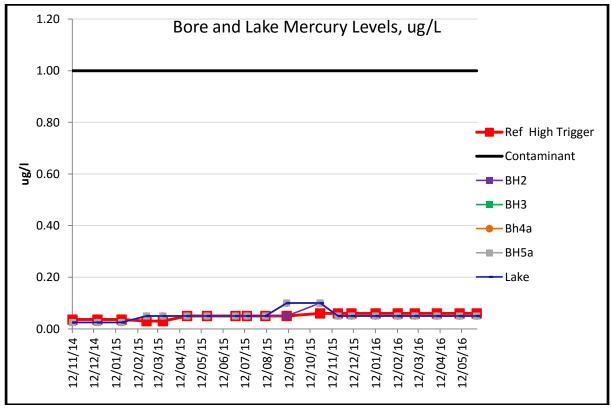


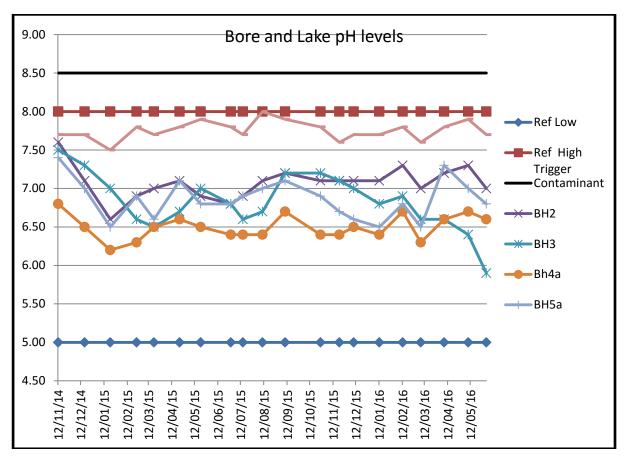


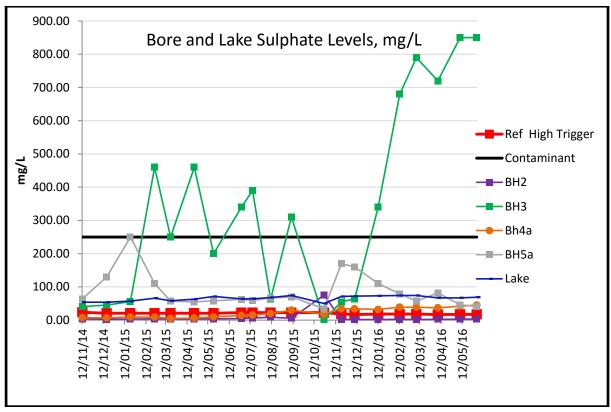


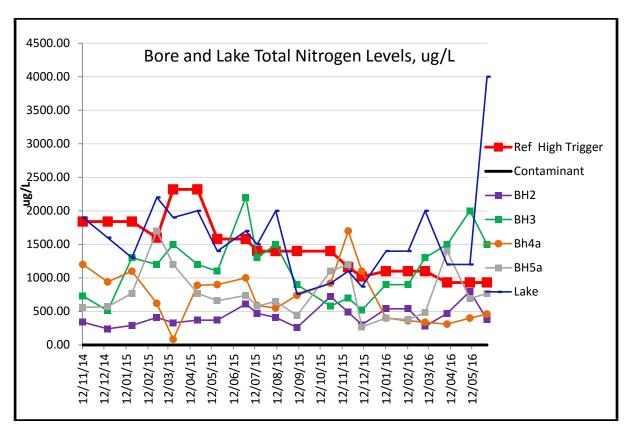


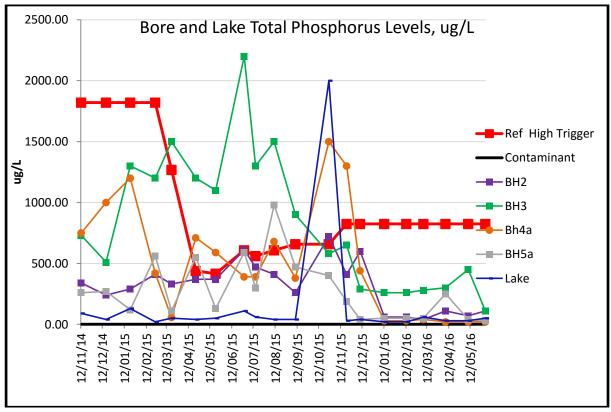


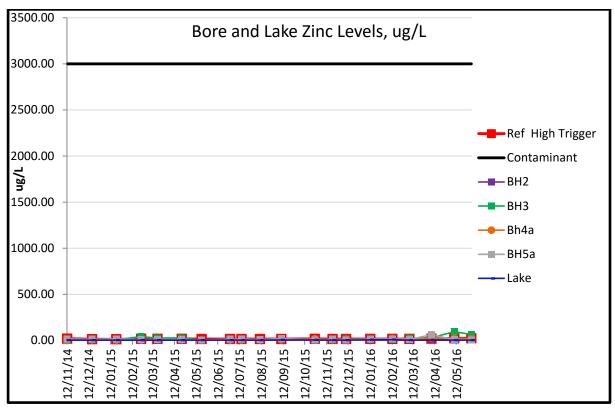


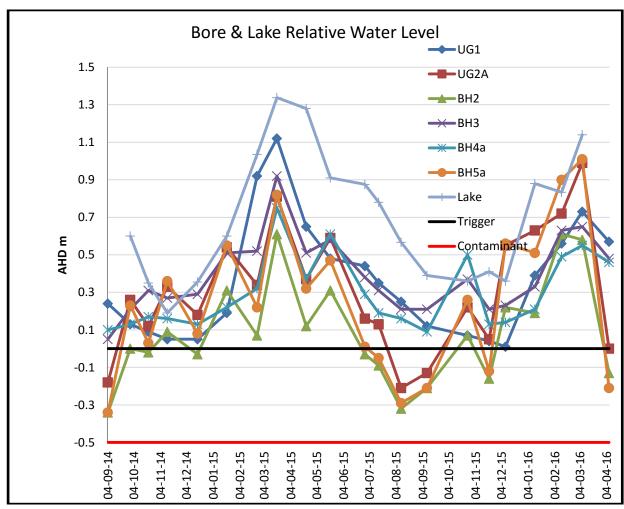


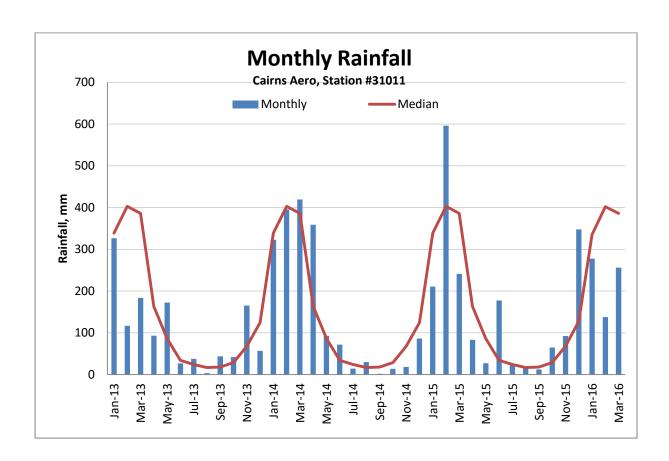














Appendix D

2016 Borehole and CPT Reports





PROJECT: Revised EIA

CLIENT:

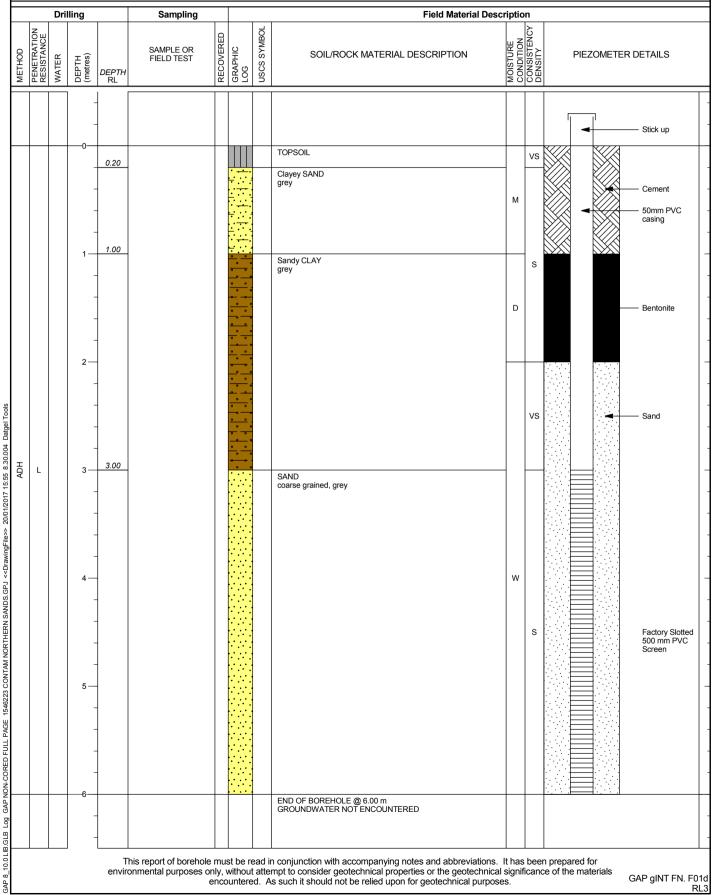
REPORT OF BOREHOLE: BH1

SHEET: 1 OF 1

COORDS: 363630.0 m E 8136179.0 m N MGA94 56 DRILL RIG: HV1

SURFACE RL: DATUM: AHD CONTRACTOR:

LOCATION: Northern Sands INCLINATION: -90° LOGGED: DSS DATE: 20/9/16 JOB NO: 1546223 HOLE DEPTH: 6.00 m CHECKED: MSC DATE: 19/12/16



This report of borehole must be read in conjunction with accompanying notes and abbreviations. It has been prepared for environmental purposes only, without attempt to consider geotechnical properties or the geotechnical significance of the materials encountered. As such it should not be relied upon for geotechnical purposes.

GAP gINT FN. F01d RL3



PROJECT: Revised EIA

CLIENT:

REPORT OF BOREHOLE: BH2

SHEET: 1 OF 1

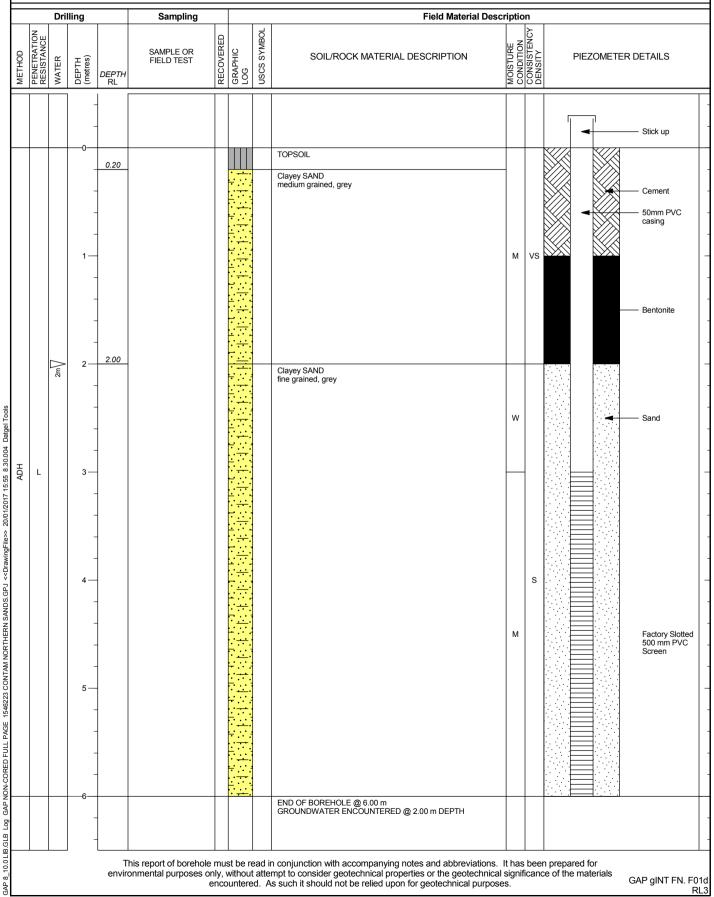
COORDS: 364376.0 m E 8135443.0 m N MGA94 56

DRILL RIG: HV1 CONTRACTOR:

SURFACE RL: DATUM: AHD

LOGGED: DSS DATE: 20/9/16

LOCATION: Northern Sands INCLINATION: -90° JOB NO: 1546223 HOLE DEPTH: 6.00 m CHECKED: MSC DATE: 19/12/16



This report of borehole must be read in conjunction with accompanying notes and abbreviations. It has been prepared for environmental purposes only, without attempt to consider geotechnical properties or the geotechnical significance of the materials encountered. As such it should not be relied upon for geotechnical purposes.

GAP gINT FN. F01d



PROJECT: Revised EIA

CLIENT:

REPORT OF BOREHOLE: BH3

SHEET: 1 OF 1

COORDS: 364302.0 m E 8135127.0 m N MGA94 56 DRILL RIG: HV1

SURFACE RL: DATUM: AHD CONTRACTOR:

LOCATION: Northern Sands INCLINATION: -90° JOB NO: 1546223

LOGGED: DSS DATE: 20/9/16

HOLE DEPTH: 12.00 m CHECKED: MSC DATE: 19/12/16 Drilling Sampling **Field Material Description** MOISTURE CONDITION CONSISTENCY DENSITY **USCS SYMBOL** RECOVERED SAMPLE OR FIELD TEST GRAPHIC LOG SOIL/ROCK MATERIAL DESCRIPTION PIEZOMETER DETAILS WATER DEPTH (metres) DEPTH RL Stick up 0.20 Clayey SAND 1.00 Sandy CLAY medium plasticity, grey М 50mm PVC 2m\ Bentonite W Sand S GAP 8_10.0 LIB.GLB LOg GAP NON-CORED FULL PAGE 1546223 CONTAM NORTHERN SANDS.GPJ <<DrawingFile>> 20/01/2017 15:55 8.30.004 Datgel Tools 6.00 Clayey SAND grey М Factory Slotted 500 mm PVC Screen 9.50 CLAY 10.00 М Sandy CLAY red St D END OF BOREHOLE @ 12.00 m GROUNDWATER ENCOUNTERED @ 2.00 m DEPTH

This report of borehole must be read in conjunction with accompanying notes and abbreviations. It has been prepared for environmental purposes only, without attempt to consider geotechnical properties or the geotechnical significance of the materials encountered. As such it should not be relied upon for geotechnical purposes.

GAP gINT FN. F01d



PROJECT: EIS Stage 1- Port Development

CLIENT:

REPORT OF BOREHOLE: GA04

SHEET: 1 OF 1 DRILL RIG: Ezi Probe

SURFACE RL: 3.58 m DATUM: AHD CONTRACTOR:

LOCATION: Northern Sands INCLINATION: -90° LOGGED: JL DATE: 23/11/16 DATE: 19/12/16 HOLE DEPTH: 6.00 m CHECKED: MSC

COORDS: 364331.6 m E 8135689.9 m N MGA94 56

Drilling Sampling Field Material Description Control of the property of the	DATE: 23/11/16 MSC DATE: 19/12/16
TOPSOIL: Sandy Sitty CLAY high plasticity, brown, fine sand, with some roots <2mm D Sandy Sitty CLAY high plasticity, brown Sandy Sitty CLAY high plasticity, brown Sandy Sitty CLAY high plasticity, brown Sandy Sitty CLAY high plasticity, pale brown, orange, fine to coarse grained sand, trace fine quartz gravel Sitty SAND fine to coarse grained, brown, orange brown, with some fine orange subangular gravel	
1 1.00 2.58 Sandy Silty CLAY high plasticity, brown Sandy Silty CLAY high plasticity, brown Sandy Silty CLAY high plasticity, pale brown, orange, fine to coarse grained sand, trace fine quartz gravel D - M Silty SAND Sandy Silty CLAY high plasticity, pale brown, orange, fine to coarse grained sand, trace fine quartz gravel Silty SAND	PIEZOMETER DETAILS
Sandy Silty CLAY high plasticity, brown 1 1.00 2.58 Sandy Silty CLAY high plasticity, brown Sandy Silty CLAY high plasticity, pale brown, orange, fine to coarse grained sand, trace fine quartz gravel D - M F Silty SAND fine to coarse grained, brown, orange brown, with some fine orange grained subangular gravel	0.9mm Stick up
1 1.00 2.58 Sandy Sitty CLAY high plasticity, brown Sandy Sitty CLAY high plasticity, pale brown, orange, fine to coarse grained sand, trace fine quartz gravel D-M Sitty SAND fine to coarse grained, brown, orange brown, with some fine orange subangular gravel	
Sandy Silty CLAY high plasticity, pale brown, orange, fine to coarse grained sand, trace fine quartz gravel D - M Silty SAND fine to coarse grained, brown, orange brown, with some fine grained, subangular gravel	
2— 2.46 1.12 2.70 Silty SAND fine to coarse grained, brown, orange brown, with some fine grained subangular gravel	50mm UPVC class 18 Blank casing Cement/grout
Silty SAND 2.70 Silty SAND fine to coarse grained, brown, orange brown, with some fine grained, subangular gravel	
grained subangular gravel	
Sandy Silty CLAY high plasticity, brown, grey, fine to coarse grained sand	
Silty SAND fine to coarse grained, grey, brown, orange brown, with some fine quartz gravel SAND SAND	■ Bentonite
SAND medium to coarse grained, orange brown, with some fine to medium quartz gravel	50mm UPVC class 18,
W L	Screen, 1mm aparture size Gravel pack
SAND -1.42 SAND medium to coarse grained, grey, orange brown, with some silt,	
6 -2.42 END OF BOREHOLE @ 6.00 m	
This report of borehole must be read in conjunction with accompanying notes and abbreviations. It has been prepare	red for

This report of borehole must be read in conjunction with accompanying notes and abbreviations. It has been prepared for hydrogeological purposes only, without attempt to assess geotechnical properties or possible contamination. Any reference to geotechnical properties or potential contamination are for information only and do not necessarily indicate the presence or absence of the properties stated.



PROJECT: Port Development ETS S1

1546223

LOCATION: Northern Sands

Flanagan Consulting Group

CLIENT:

JOB NO:

REPORT OF BOREHOLE: GA05

SHEET: 1 OF 1
DRILL RIG: Eziprobe

DRILL RIG: Eziprobe

CONTRACTOR: Golder

 INCLINATION: -90°
 LOGGED: JJP
 DATE: 28/1/16

 HOLE DEPTH: 1.50 m
 CHECKED: MSC
 DATE: 19/12/16

THOSE DEL THE LOOM													
			lling		Sampling				Field Material Desc				
METHOD	PENETRATION RESISTANCE	WATER	DEPTH (metres)	<i>DEPTH</i> RL	SAMPLE OR FIELD TEST	RECOVERED	GRAPHIC LOG	OHANGE SOIL/ROCK MATERIAL DESCRIPTION SOIL/ROCK MATERIAL DESCRIPTION			CONSISTENCY	STRUCTURE AND ADDITIONAL OBSERVATIONS	
			0.0 - - - -	0.45			X X X X X X X X X X X X X X X X X X X	ML	TOPSOIL: Sandy SILT fine to medium, grey brown, with some rootlets and clay	D - N	/I St		
РТ	L	GWNE	0.5	0.10	GA05-DS1 0.60-0.90 m		× · · · × · · · × · · · · × · · · · ×	SM	Sitty SAND fine grained, grey brown and grey, with some clay, (approx. 40% fines content)		MD		
			1.0 —	1.20	GA05-DS2 1.20-1.50 m		× · · · · · · · · · · · · · · · · · · ·	CI	Silty CLAY grey, with trace red brown pockets, with some fine grained sand and organics	М	F		
			 1.5				<u>×</u> ×		END OF BOREHOLE @ 1.50 m TARGET DEPTH GROUNDWATER ENCOUNTERED @ m DEPTH				
			2.0 —										
			- 2.5 — - -										
			3.0 — -										
			3.5 —										
			4.0										
			- - 4.5—										
			- - - 5.0										

COORDS: MGA94 56

SURFACE RL: DATUM: AHD

This report of borehole must be read in conjunction with accompanying notes and abbreviations. It has been prepared for geotechnical purposes only, without attempt to assess possible contamination. Any references to potential contamination are for information only and do not necessarily indicate the presence or absence of soil or groundwater contamination.



LOCATION: Northern Sands

1546223

Flanagan Consulting Group

CLIENT:

JOB NO:

REPORT OF PIEZO-CONE PROBE: CPT1

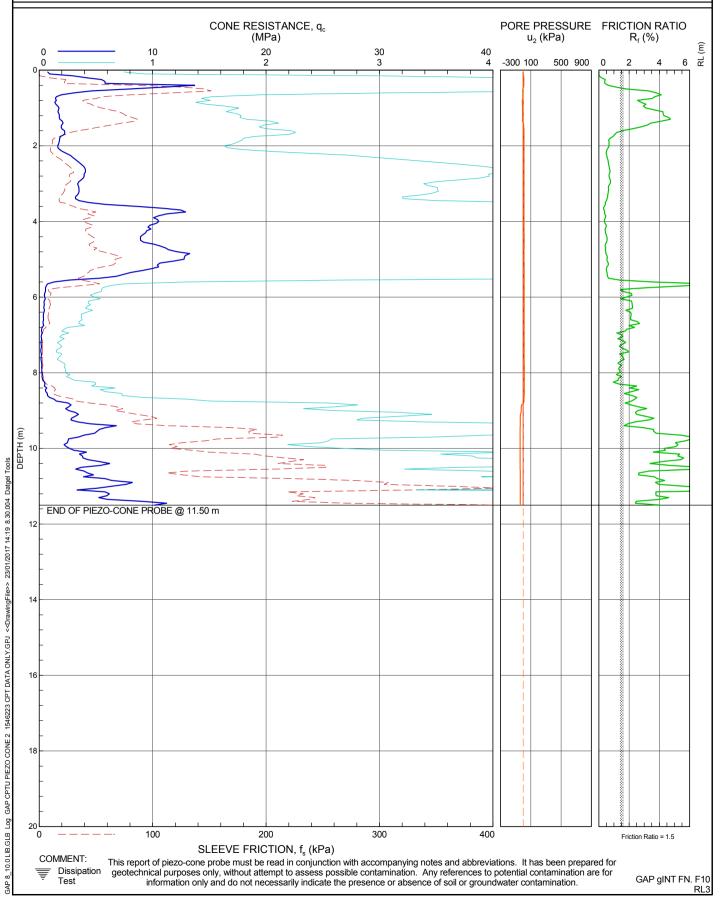
SHEET: 1 OF 1 CONE No.:

RIG:

PROJECT: Cairns Shipping Development SURFACE RL: DATUM: AHD CONTRACTOR: Geo Investigate

COORDS: MGA94 56

INCLINATION: -90° RECORDED: GZL DATE: 31/10/16
HOLE DEPTH: 11.50 m CHECKED: MSC DATE: 31/10/16





REPORT OF PIEZO-CONE PROBE: CPT2

SHEET: 1 OF 1 CONE No.:

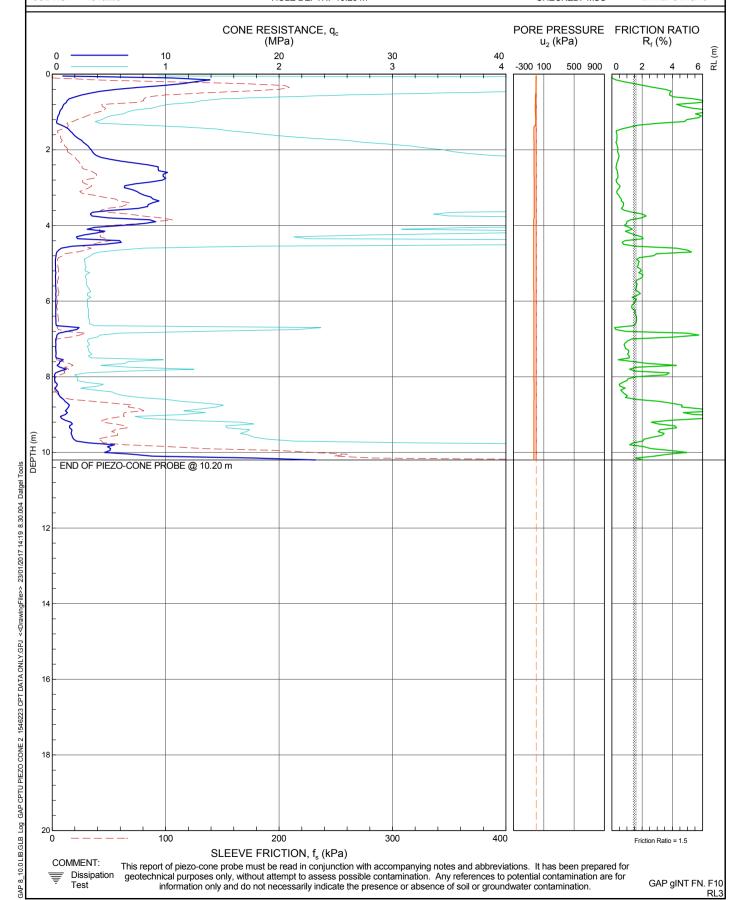
RIG:

PROJECT: Cairns Shipping Development SURFACE RL: DATUM: AHD CONTRACTOR: Geo Investigate

COORDS: MGA94 56

 LOCATION:
 Northern Sands
 INCLINATION: -90°
 RECORDED: GZL
 DATE: 31/10/16

 JOB NO:
 1546223
 HOLE DEPTH: 10.20 m
 CHECKED: MSC
 DATE: 31/10/16





REPORT OF PIEZO-CONE PROBE: CPT3

SHEET: 1 OF 1 CONE No.:

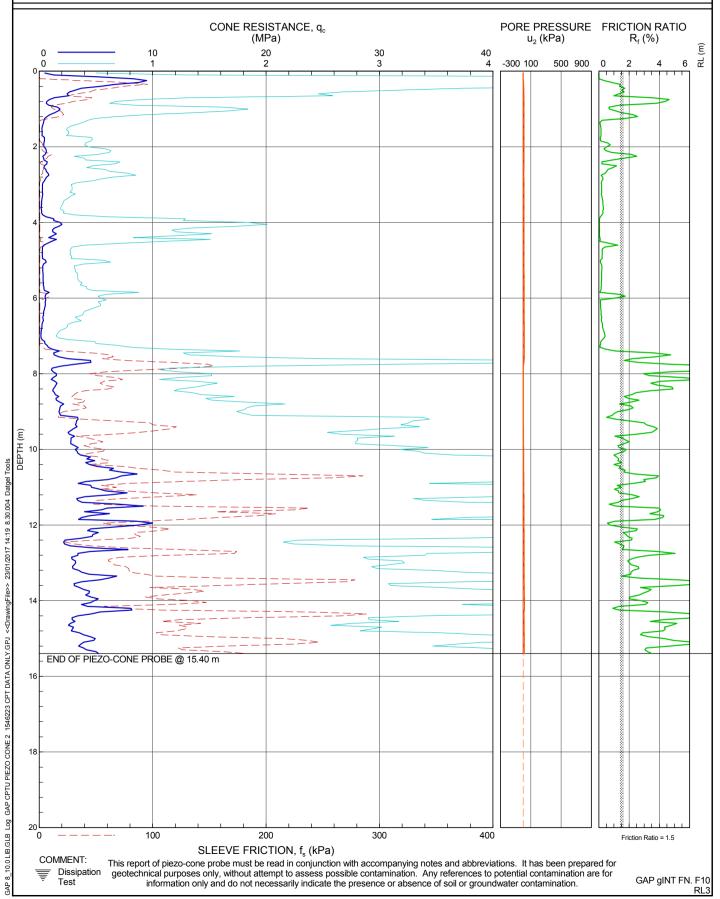
RIG:

PROJECT: Cairns Shipping Development SURFACE RL: DATUM: AHD CONTRACTOR: Geo Investigate

COORDS: MGA94 56

 LOCATION:
 Northern Sands
 INCLINATION: -90°
 RECORDED: GZL
 DATE: 31/10/16

 JOB NO:
 1546223
 HOLE DEPTH: 15.40 m
 CHECKED: MSC
 DATE: 31/10/16





LOCATION: Northern Sands

1546223

Flanagan Consulting Group

CLIENT:

JOB NO:

REPORT OF PIEZO-CONE PROBE: CPT4

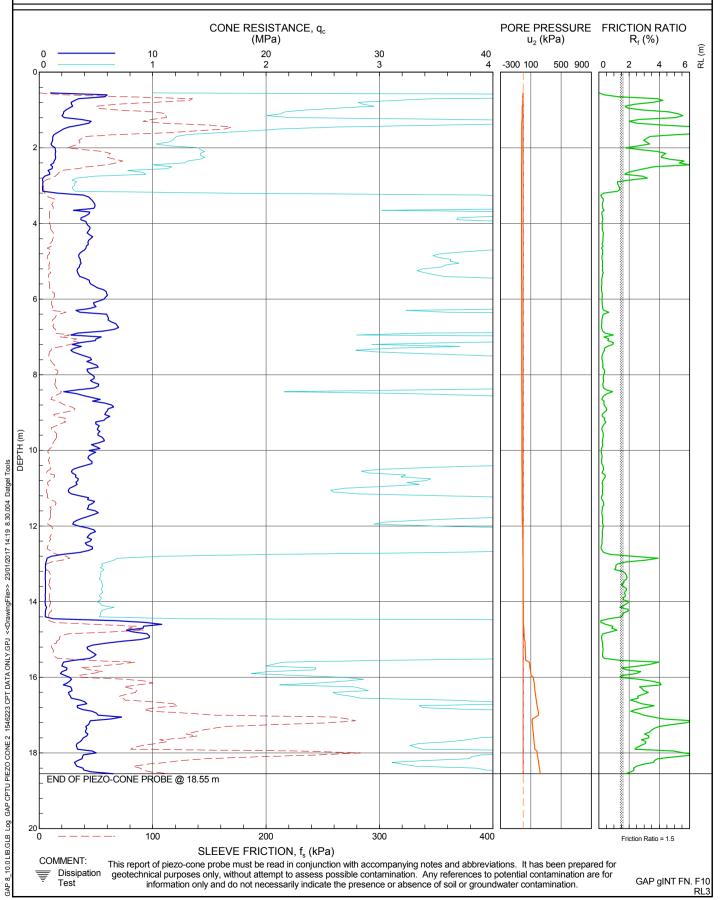
SHEET: 1 OF 1 CONE No.:

RIG:

PROJECT: Cairns Shipping Development SURFACE RL: DATUM: AHD CONTRACTOR: Geo Investigate

COORDS: MGA94 56

INCLINATION: -90° RECORDED: GZL DATE: 28/10/16
HOLE DEPTH: 18.55 m CHECKED: MSC DATE: 28/10/16





CLIENT:

REPORT OF PIEZO-CONE PROBE: CPT5

SHEET: 1 OF 1 CONE No.:

RIG:

CONTRACTOR: Geo Investigate

RECORDED: GZL D.

DATE: 28/10/16

PROJECT: Cairns Shipping Development SURFACE RL: DATUM: AHD LOCATION: Northern Sands INCLINATION: -90°

COORDS: MGA94 56

JOB NO: 1546223 HOLE DEPTH: 13.35 m CHECKED: MSC DATE: 28/10/16 CONE RESISTANCE, q_c PORE PRESSURE FRICTION RATIO (MPa) u2 (kPa) $R_{f}(\%)$ 20 2 10 30 -300 100 牊 500 900 2 6 0 DEPTH (m) END OF PIEZO-CONE PROBE @ 13.35 m 18 400 100 200 Friction Ratio = 1.5 SLEEVE FRICTION, f_s (kPa) COMMENT:

GAP 8.10.0 LIB.GLB Log GAP CPTU PIEZO CONE 2 1546223 CPT DATA ONLY.GPJ <<DrawngFile>> 23/01/2017 14:19 8:30.004 Datgel Tools

Dissipation

This report of piezo-cone probe must be read in conjunction with accompanying notes and abbreviations. It has been prepared for geotechnical purposes only, without attempt to assess possible contamination. Any references to potential contamination are for information only and do not necessarily indicate the presence or absence of soil or groundwater contamination.

GAP gINT FN. F10 RL3



CLIENT:

REPORT OF PIEZO-CONE PROBE: CPT6

SHEET: 1 OF 1 CONE No.:

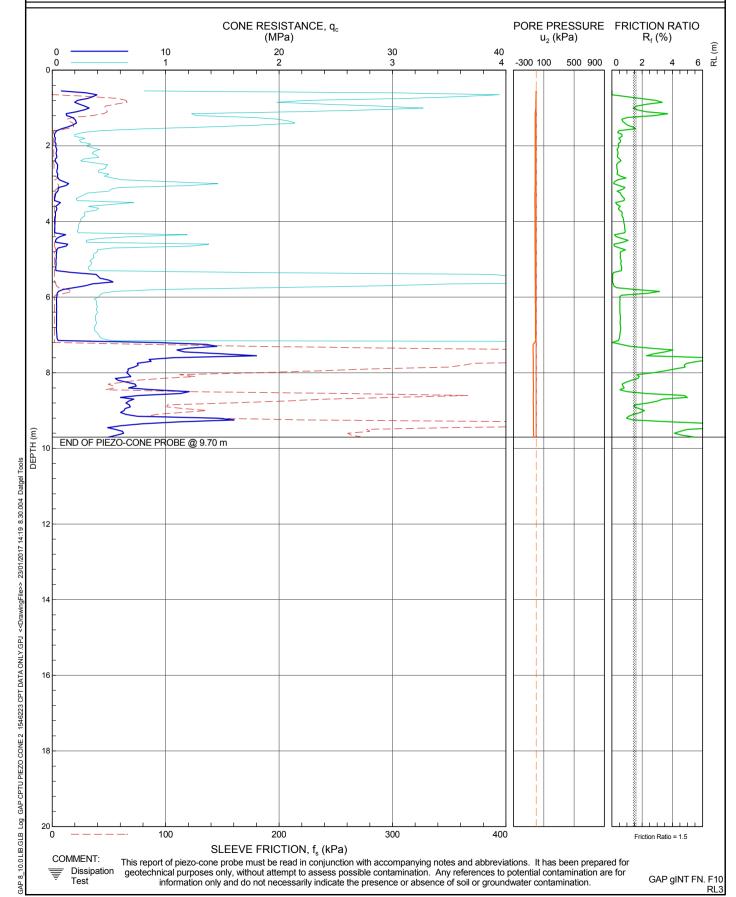
RIG:

PROJECT: Cairns Shipping Development SURFACE RL: DATUM: AHD CONTRACTOR: Geo Investigate

COORDS: MGA94 56

 LOCATION:
 Northern Sands
 INCLINATION: -90°
 RECORDED: GZL
 DATE: 28/10/16

 JOB NO:
 1546223
 HOLE DEPTH: 9.70 m
 CHECKED: MSC
 DATE: 28/10/16





REPORT OF PIEZO-CONE PROBE: CPT7

SHEET: 1 OF 1 CONE No.:

RIG:

PROJECT: Cairns Shipping Development SURFACE RL: DATUM: AHD

CONTRACTOR: Geo Investigate

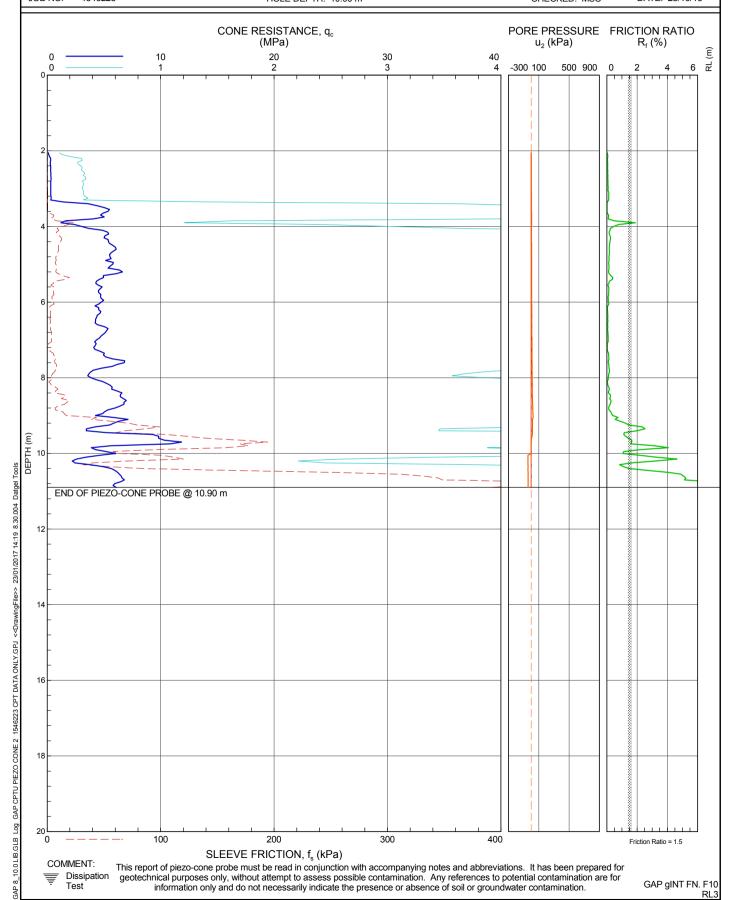
LOCATION: Northern Sands JOB NO: 1546223

INCLINATION: -90° HOLE DEPTH: 10.90 m

COORDS: MGA94 56

RECORDED: GZL CHECKED: MSC DATE: 28/10/16

DATE: 28/10/16





METHOD OF SOIL DESCRIPTION USED ON BOREHOLE AND TEST PIT REPORTS

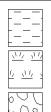
0000

FILL

GRAVEL (GP or GW)

SAND (SP or SW)

SILT (ML or MH)



CLAY (CL, CI or CH)

ORGANIC SOILS (OL or OH or Pt)

COBBLES or BOULDERS

Combinations of these basic symbols may be used to indicate mixed materials such as sandy clay.

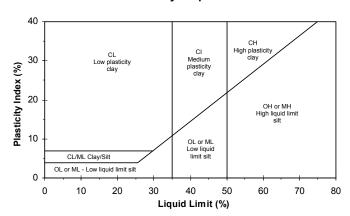
CLASSIFICATION AND INFERRED STRATIGRAPHY

Soil and Rock is classified and described in Reports of Boreholes and Test Pits using the preferred method given in AS1726 – 1993, (Amdt1 – 1994 and Amdt2 – 1994), Appendix A. The material properties are assessed in the field by visual/tactile methods.

Particle Size

Major Division		Sub Division	Particle Size		
В	OULD	> 200 mm			
(СОВВ	LES	63 to 200 mm		
		Coarse	20 to 63 mm		
GRAVEL		Medium	6.0 to 20 mm		
		Fine	2.0 to 6.0 mm		
		Coarse	0.6 to 2.0 mm		
SAND		Medium	0.2 to 0.6 mm		
		Fine	0.075 to 0.2 mm		
	SIL	0.002 to 0.075 mm			
	CLA	ΛΥ	< 0.002 mm		

Plasticity Properties



MOISTURE CONDITION

AS1726 - 1993

		7.01120 1000
Symbol	Term	Description
D	Dry	Sands and gravels are free flowing. Clays & Silts may be brittle or friable and powdery.
М	Moist	Soils are darker than in the dry condition & may feel cool. Sands and gravels tend to cohere.
W	Wet	Soils exude free water. Sands and gravels tend to cohere.

CONSISTENCY AND DENSITY

Undrained Shear Symbol Term Strength VS Very Soft 0 to 12 kPa S Soft 12 to 25 kPa F 25 to 50 kPa Firm St Stiff 50 to 100 kPa VSt Very Stiff 100 to 200 kPa Н Hard Above 200 kPa

AS1726 - 1993

7,61720 1000									
Symbol	Term	Density Index %	SPT "N" #						
VL	Very Loose	Less than 15	0 to 4						
L Loose		Loose 15 to 35							
MD	Medium Dense	35 to 65	10 to 30						
D Dense		65 to 85	30 to 50						
VD Very Dense		Above 85	Above 50						

In the absence of test results, consistency and density may be assessed from correlations with the observed behaviour of the material

SPT correlations are not stated in AS1726 – 1993, and may be subject to corrections for overburden pressure and equipment type.



EXPLANATION OF NOTES, ABBREVIATIONS & TERMS USED ON BOREHOLE AND TEST PIT REPORTS

DRILLING/E	EXCAVATION METHOD				
AS*	Auger Screwing	RD	Rotary blade or drag bit	NQ	Diamond Core - 47 mm
AD*	Auger Drilling	RT	Rotary Tricone bit	NMLC	Diamond Core - 52 mm
*V	V-Bit	RAB	Rotary Air Blast	HQ	Diamond Core - 63 mm
*T	TC-Bit, e.g. ADT	RC	Reverse Circulation	HMLC	Diamond Core – 63mm
HA	Hand Auger	PT	Push Tube	BH	Tractor Mounted Backhoe
ADH	Hollow Auger	CT	Cable Tool Rig	EX	Tracked Hydraulic Excavator
DTC	Diatube Coring	JET	Jetting	EE	Existing Excavation
WB	Washbore or Bailer	NDD	Non-destructive digging	HAND	Excavated by Hand Methods

PENETRATION/EXCAVATION RESISTANCE

- Low resistance. Rapid penetration possible with little effort from the equipment used. L
- M Medium resistance. Excavation/possible at an acceptable rate with moderate effort from the equipment used.
- н High resistance to penetration/excavation. Further penetration is possible at a slow rate and requires significant effort from the equipment.
- R Refusal or Practical Refusal. No further progress possible without the risk of damage or unacceptable wear to the digging implement or machine.

These assessments are subjective and are dependent on many factors including the equipment power, weight, condition of excavation or drilling tools, and the experience of the operator.

V	V.	Δ	Т	F	R

 $\mathbf{\nabla}$ Water level at date shown Partial water loss Water inflow Complete water loss

GROUNDWATER NOT

OBSERVED

The observation of groundwater, whether present or not, was not possible due to drilling water,

surface seepage or cave in of the borehole/test pit.

GROUNDWATER NOT

ENCOUNTERED

The borehole/test pit was dry soon after excavation. However, groundwater could be present in less permeable strata. Inflow may have been observed had the borehole/test pit been left open

for a longer period.

SAMPLING AND TESTING

Standard Penetration Test to AS1289.6.3.1-2004

4,7,11 N=18 4,7,11 = Blows per 150mm.N = Blows per 300mm penetration following 150mm seating Where practical refusal occurs, the blows and penetration for that interval are reported 30/80mm

RW Penetration occurred under the rod weight only

HW Penetration occurred under the hammer and rod weight only

Hammer double bouncing on anvil HB

DS Disturbed sample Bulk disturbed sample **BDS**

G Gas Sample Water Sample W

FP Field permeability test over section noted

F۷ Field vane shear test expressed as uncorrected shear strength (s_v = peak value, s_r = residual value)

PID Photoionisation Detector reading in ppm PM Pressuremeter test over section noted

PP Pocket penetrometer test expressed as instrument reading in kPa

U63 Thin walled tube sample - number indicates nominal sample diameter in millimetres

WPT Water pressure tests

DCP Dynamic cone penetration test **CPT** Static cone penetration test

CPTu Static cone penetration test with pore pressure (u) measurement

Ranking of Visually	Ranking of Visually Observable Contamination and Odour (for specific soil contamination assessment projects)										
R = 0	No visible evidence of contamination	R = A	No non-natural odours identified								
R = 1	Slight evidence of visible contamination	R = B	Slight non-natural odours identified								
R = 2	Visible contamination	R = C	Moderate non-natural odours identified								
R = 3	Significant visible contamination	R = D	Strong non-natural odours identified								

ROCK CORE RECOVERY

TCR = Total Core Recovery (%)

SCR = Solid Core Recovery (%)

RQD = Rock Quality Designation (%)

Length of core recovered × 100 Length of core run

\(\sum_\text{Length of cylindrical core recovered} \) ×100 Length of core run

 \sum Axial lengths of core > 100 mm Length of core run



Appendix E

Water Quality Laboratory Results and Field Data.









CLIENT DETAILS -

Contact

Darcy Simpson

Client

GOLDER ASSOCIATES PTY LTD

Address

PO BOX 5823 216 DRAPER ST CAIRNS QLD 4870

Telephone

07 4054 8200

Facsimile Email

dasimpson@golder.com.au

Project

1546223 Northern Sands

Order Number Samples

07 4054 8201

(Not specified)

LABORATORY DETAILS

Manager

Laboratory Address

Telephone

Facsimile

Date Reported

Email

Jon Dicker

SGS Cairns Environmental

Unit 2, 58 Comport St

Portsmith QLD 4870

+61 07 4035 5111

+61 07 4035 5122

AU.Environmental.Cairns@sgs.com

SGS Reference CE123168 R0 Date Received 29 Sep 2016

11 Oct 2016

COMMENTS

Accredited for compliance with ISO/IEC 17025. NATA accredited laboratory 2562(3146)

SIGNATORIES

Anthony Nilsson

Operations Manager

Jon Dicker

Manager Northern QLD

Leanne Orsmond

Quality & Microbiology Coordinator

Horsmond

Maristela Ganzan Metals Team Leader

SGS Australia Pty Ltd ABN 44 000 964 278

Environment, Health and Safety

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Australia

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www.sgs.com.au



CE123168 R0

	Sa	nple Number ample Matrix Sample Date ample Name	CE123168.001 Water 29 Sep 2016 BH01	CE123168.002 Water 29 Sep 2016 BH02	CE123168.003 Water 29 Sep 2016 BH03
Parameter	Units	LOR			
pH in water Method: AN101 Tested: 29/9/2016					
pH**	pH Units	0.1	7.0	7.7	7.9
Conductivity and TDS by Calculation - Water Method: AN106	Tested: 29/	9/2016			
Conductivity @ 25 C	μS/cm	5	230	11000	24000
Total Dissolved Solids (by calculation)	mg/L	10	140	6300	15000
Alkalinity Method: AN135 Tested: 29/9/2016					
Total Alkalinity as CaCO3	mg/L	5	41	1000	2300
Bicarbonate Alkalinity as CaCO3	mg/L	5	41	1000	2300
Carbonate Alkalinity as CaCO3	mg/L	5	< 5	<5	<5
Hydroxide Alkalinity as CaCO3	mg/L	5	<5	<5	<5
Acidity and Free CO2 Method: AN140 Tested: 29/9/2016					
Acidity to pH 8.3	mg CaCO3/L	5	58	200	250
Chloride by Discrete Analyser in Water Method: AN274 Test	ted: 5/10/201	6	36	3000	7900
			**		
Metals in Water (Dissolved) by ICPOES Method: AN320/AN32	1 Tested: 1	0/10/2016			
Aluminium, Al	mg/L	0.005	<0.005	0.012	0.021
Iron, Fe	mg/L	0.005	0.048	0.31	0.16
Sulphur as Sulphate, SO4	mg/L	0.5	28	14	7.4

11-October-2016 Page 2 of 7



CE123168 R0

	Sa :	nple Number ample Matrix Sample Date ample Name	Water 29 Sep 2016	CE123168.002 Water 29 Sep 2016 BH02	CE123168.003 Water 29 Sep 2016 BH03					
Parameter	Units	LOR								
Metals in Water (Total) by ICPOES Method: AN022/AN320 Tested: 5/10/2016										
Total Iron	mg/L	0.005	10	470	1.8					

11-October-2016 Page 3 of 7



QC SUMMARY

MB blank results are compared to the Limit of Reporting

LCS and MS spike recoveries are measured as the percentage of analyte recovered from the sample compared the the amount of analyte spiked into the sample.

DUP and MSD relative percent differences are measured against their original counterpart samples according to the formula: the absolute difference of the two results divided by the average of the two results as a percentage. Where the DUP RPD is 'NA', the results are less than the LOR and thus the RPD is not applicable.

Acidity and Free CO2 Method: ME-(AU)-[ENV]AN140

Ī	Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
ı		Reference					%Recovery
ı	Acidity to pH 8.3	LB039825	mg CaCO3/L	5	<5	0%	NA

Alkalinity Method: ME-(AU)-[ENV]AN135

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
Total Alkalinity as CaCO3	LB039828	mg/L	5	<5	3 - 11%	103%
Bicarbonate Alkalinity as CaCO3	LB039828	mg/L	5	<5		
Carbonate Alkalinity as CaCO3	LB039828	mg/L	5	<5		
Hydroxide Alkalinity as CaCO3	LB039828	mg/L	5	<5		

Chloride by Discrete Analyser in Water Method: ME-(AU)-[ENV]AN274

ı	Parameter	QC	Units	LOR	МВ	DUP %RPD	LCS
ı		Reference					%Recovery
	Chloride, Cl	LB039925	mg/L	1	<1	1 - 3%	107%

Conductivity and TDS by Calculation - Water Method: ME-(AU)-[ENV]AN106

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
Conductivity @ 25 C	LB039828	μS/cm	5	<5	0 - 1%	99%
Total Dissolved Solids (by calculation)	LB039828	mg/L	10	<10	1%	NA

Metals in Water (Total) by ICPOES Method: ME-(AU)-[ENV]AN022/AN320

Parameter	QC	Units	LOR	LCS
	Reference			%Recovery
Total Iron	LB039917	mg/L	0.005	102%

Metals in Water (Dissolved) by ICPOES Method: ME-(AU)-[ENV]AN320/AN321

D	00	11:4	LOB	MD	DUD W DDD	1.00
Parameter	QC	Units	LOR	МВ	DUP %RPD	LCS
	Reference					%Recovery
Aluminium, Al	LB040030	mg/L	0.005	<0.005	3%	99%
Iron, Fe	LB040030	mg/L	0.005	<0.005	0%	105%
Sulphur as Sulphate, SO4	LB040030	mg/L	0.5	<0.5	3%	NA

11-October-2016 Page 4 of 7





QC SUMMARY

MB blank results are compared to the Limit of Reporting
LCS and MS spike recoveries are measured as the percentage of analyte recovered from the sample compared the the amount of analyte spiked into the sample.

DUP and MSD relative percent differences are measured against their original counterpart samples according to the formula: the absolute difference of the two results divided by the average of the two results as a percentage. Where the DUP RPD is 'NA', the results are less than the LOR and thus the RPD is not applicable.

pH in water Method: ME-(AU)-[ENV]AN101

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
pH**	LB039828	pH Units	0.1	5.5	0%	NA

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METHOD SUMMARY

METLIOD	
METHOD —	METHODOLOGY SUMMARY
AN022/AN320	Total (acid soluble) Metals by ICP-OES: Samples are digested in nitric or nitric and hydrochloric acids prior to analysis for a wide range of metals and some non-metals. This solution is measured by Inductively Coupled Plasma. Solutions are aspirated into an argon plasma at 8000-10000K and emit characteristic energy or light as a result of electron transitions through unique energy levels. The emitted light is focused onto a diffraction grating where it is separated into components.
AN101	pH in Soil Sludge Sediment and Water: pH is measured electrometrically using a combination electrode (glass plus reference electrode) and is calibrated against 3 buffers purchased commercially. For soils, an extract with water is made at a ratio of 1:5 and the pH determined and reported on the extract. Reference APHA 4500-H+.
AN106	Conductivity and TDS by Calculation: Conductivity is measured by meter with temperature compensation and is calibrated against a standard solution of potassium chloride. Conductivity is generally reported as μ mhos/cm or μ S/cm @ 25°C. For soils, an extract with water is made at a ratio of 1:5 and the EC determined and reported on the extract, or calculated back to the as-received sample. Total Dissolved Salts can be estimated from conductivity using a conversion factor, which for natural waters, is in the range 0.55 to 0.75. SGS use 0.6. Reference APHA 2510 B.
AN135	Alkalinity (and forms of) by Titration: The sample is titrated with standard acid to pH 8.3 (P titre) and pH 4.5 (T titre) and permanent and/or total alkalinity calculated. The results are expressed as equivalents of calcium carbonate or recalculated as bicarbonate, carbonate and hydroxide. Reference APHA 2320. Internal Reference AN135
AN140	Acidity by Titration: The water sample is titrated with sodium hydroxide to designated pH end point. In a sample containing only carbon dioxide, bicarbonates and carbonates, titration to pH 8.3 at 25°C corresponds to stoichiometric neutralisation of carbonic acid to bicarbonate. Method reference APHA 2310 B.
AN274	Chloride by Aquakem DA: Chloride reacts with mercuric thiocyanate forming a mercuric chloride complex. In the presence of ferric iron, highly coloured ferric thiocyanate is formed which is proportional to the chloride concentration. Reference APHA 4500CI-
AN320	Photomultipliers or CCDs are used to measure the light intensity at specific wavelengths. This intensity is directly proportional to concentration. Corrections are required to compensate for spectral overlap between elements. Reference APHA 3120 B.
AN320/AN321	Metals by ICP-OES: Samples are preserved with 10% nitric acid for a wide range of metals and some non-metals. This solution is measured by Inductively Coupled Plasma. Solutions are aspirated into an argon plasma at 8000-10000K and emit characteristic energy or light as a result of electron transitions through unique energy levels. The emitted light is focused onto a diffraction grating where it is separated into components.
AN320/AN321	Photomultipliers or CCDs are used to measure the light intensity at specific wavelengths. This intensity is directly proportional to concentration. Corrections are required to compensate for spectral overlap between elements. Reference APHA 3120 B.
Calculation	Free and Total Carbon Dioxide may be calculated using alkalinity forms only when the samples TDS is <500mg/L. If TDS is >500mg/L free or total carbon dioxide cannot be reported. APHA4500CO2 D.

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FOOTNOTES _

IS Insufficient sample for analysis.

LNR Sample listed, but not received.

* NATA accreditation does not cover the performance of this service.

** Indicative data, theoretical holding time exceeded.

LOR Limit of Reporting

↑↓ Raised or Lowered Limit of Reporting
QFH QC result is above the upper tolerance
QFL QC result is below the lower tolerance

- The sample was not analysed for this analyte

NVL Not Validated

Samples analysed as received.

Solid samples expressed on a dry weight basis.

Where "Total" analyte groups are reported (for example, Total PAHs, Total OC Pesticides) the total will be calculated as the sum of the individual analytes, with those analytes that are reported as <LOR being assumed to be zero. The summed (Total) limit of reporting is calcuated by summing the individual analyte LORs and dividing by two. For example, where 16 individual analytes are being summed and each has an LOR of 0.1 mg/kg, the "Totals" LOR will be 1.6 / 2 (0.8 mg/kg). Where only 2 analytes are being summed, the "Total" LOR will be the sum of those two LORs.

Some totals may not appear to add up because the total is rounded after adding up the raw values.

If reported, measurement uncertainty follow the ± sign after the analytical result and is expressed as the expanded uncertainty calculated using a coverage factor of 2, providing a level of confidence of approximately 95%, unless stated otherwise in the comments section of this report.

Results reported for samples tested under test methods with codes starting with ARS-SOP, radionuclide or gross radioactivity concentrations are expressed in becquerel (Bq) per unit of mass or volume or per wipe as stated on the report. Becquerel is the SI unit for activity and equals one nuclear transformation per second.

Note that in terms of units of radioactivity:

- a. 1 Bq is equivalent to 27 pCi
- b. 37 MBq is equivalent to 1 mCi

For results reported for samples tested under test methods with codes starting with ARS-SOP, less than (<) values indicate the detection limit for each radionuclide or parameter for the measurement system used. The respective detection limits have been calculated in accordance with ISO 11929.

The QC criteria are subject to internal review according to the SGS QAQC plan and may be provided on request or alternatively can be found here: http://www.sgs.com.au/~/media/Local/Australia/Documents/Technical%20Documents/MP-AU-ENV-QU-022%20QA%20QC%20Plan.pdf

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LABORATORY DETAILS

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1546223 Northern Sands

Order Number Samples

(Not specified)

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Date Reported

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CE124163 R0

23 Nov 2016 02 Dec 2016

COMMENTS

Accredited for compliance with ISO/IEC 17025. NATA accredited laboratory 2562(3146)

For determination of soluble metals, filtered sample was not received so samples were laboratory filtered on receipt. This may give soluble metals results that do not represent the concentrations present at the time of sampling.

SIGNATORIES

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Operations Manager

Jon Dicker

Manager Northern QLD

Leanne Orsmond

Quality & Microbiology Coordinator

Horsmond

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Anion-Cation Balance

ANALYTICAL REPORT

CE124163 R0

	Sa :	nple Number ample Matrix Sample Date ample Name	Water 22 Nov 2016	CE124163.002 Water 22 Nov 2016 BH02	CE124163.003 Water 22 Nov 2016 BH03
Parameter	Units	LOR			
pH in water Method: AN101 Tested: 23/11/2016					
pH**	pH Units	0.1	6.8	7.9	7.9
Conductivity and TDS by Calculation - Water Method: AN106	Tested: 23/	11/2016			
Conductivity @ 25 C	μS/cm	5	250	9300	24000
Total Dissolved Solids (by calculation)	mg/L	10	150	5600	14000
Alkalinity Method: AN135 Tested: 23/11/2016			,		
Total Alkalinity as CaCO3	mg/L	5	46	1100	2200
Bicarbonate Alkalinity as CaCO3	mg/L	5	46	1100	2200
Carbonate Alkalinity as CaCO3	mg/L	5	<5	<5	
Hydroxide Alkalinity as CaCO3	mg/L	5	<5	<5	<5
Chloride by Discrete Analyser in Water Method: AN274 Test	sted: 25/11/201	1	36	2500	7800
Metals in Water (Dissolved) by ICPOES Method: AN320/AN3		8/11/2016			
Calcium, Ca	mg/L	0.1	6.5	65	200
Magnesium, Mg	mg/L	0.1	9.6	160	580
Potassium, K	mg/L	0.1	3.0	100	270
Sodium, Na	mg/L	0.5	26	1600	4800
Sulphur as Sulphate, SO4	mg/L	0.5	23	11	13
Total Hardness by Calculation	mg CaCO3/L	1	56	810	2900
Calculation of Anion-Cation Balance (SAR Calc) Method: AN	121 Tested:	2/12/2016			
Sum of Cation Milliequivalents*	meq/L	-	2.32	90.2	272
Sum of Anion Milliequivalents*	meq/L	-	2.43	93.5	264

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-100

-2.4

-1.8

1.5



QC SUMMARY

MB blank results are compared to the Limit of Reporting

LCS and MS spike recoveries are measured as the percentage of analyte recovered from the sample compared the the amount of analyte spiked into the sample.

DUP and MSD relative percent differences are measured against their original counterpart samples according to the formula: the absolute difference of the two results divided by the average of the two results as a percentage. Where the DUP RPD is 'NA', the results are less than the LOR and thus the RPD is not applicable.

Alkalinity Method: ME-(AU)-[ENV]AN135

Parameter	QC Reference	Units	LOR	MB	DUP %RPD	LCS %Recovery
Total Alkalinity as CaCO3	LB041247	mg/L	5	<5	0 - 5%	107 - 115%
Bicarbonate Alkalinity as CaCO3	LB041247	mg/L	5	<5		
Carbonate Alkalinity as CaCO3	LB041247	mg/L	5	<5		
Hydroxide Alkalinity as CaCO3	LB041247	mg/L	5	<5		

Chloride by Discrete Analyser in Water Method: ME-(AU)-[ENV]AN274

ı	Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
		Reference					%Recovery
ı	Chloride, Cl	LB041309	mg/L	1	<1	0 - 1%	104%

Conductivity and TDS by Calculation - Water Method: ME-(AU)-[ENV]AN106

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
Conductivity @ 25 C	LB041247	μS/cm	5	<5	0 - 1%	98 - 101%
Total Dissolved Solids (by calculation)	LB041247	mg/L	10	<10	0 - 1%	NA

Metals in Water (Dissolved) by ICPOES Method: ME-(AU)-[ENV]AN320/AN321

Parameter	QC Reference	Units	LOR	MB	DUP %RPD	LCS %Recovery	MS %Recovery
Calcium, Ca	LB041329	mg/L	0.1	<0.1		103%	111%
Magnesium, Mg	LB041329	mg/L	0.1	<0.1		100%	106%
Potassium, K	LB041329	mg/L	0.1	<0.1		103%	113%
Sodium, Na	LB041329	mg/L	0.5	<0.5	0%	96%	100%
Sulphur as Sulphate, SO4	LB041329	mg/L	0.5	<0.5	0%	NA	
Total Hardness by Calculation	LB041329	mg CaCO3/L	1	<1			

pH in water Method: ME-(AU)-[ENV]AN101

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS	
	Reference					%Recovery	
pH**	LB041247	pH Units	0.1	5.6 - 6.2	0 - 4%	NA	

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METHOD SUMMARY

METHOD	METHODOLOGY CHAMADY -
WETTIOD -	METHODOLOGY SUMMARY
AN101	pH in Soil Sludge Sediment and Water: pH is measured electrometrically using a combination electrode (glass plus reference electrode) and is calibrated against 3 buffers purchased commercially. For soils, an extract with water is made at a ratio of 1:5 and the pH determined and reported on the extract. Reference APHA 4500-H+.
AN106	Conductivity and TDS by Calculation: Conductivity is measured by meter with temperature compensation and is calibrated against a standard solution of potassium chloride. Conductivity is generally reported as μ mhos/cm or μ S/cm @ 25°C. For soils, an extract with water is made at a ratio of 1:5 and the EC determined and reported on the extract, or calculated back to the as-received sample. Total Dissolved Salts can be estimated from conductivity using a conversion factor, which for natural waters, is in the range 0.55 to 0.75. SGS use 0.6. Reference APHA 2510 B.
AN121	This method is used to calculation the balance of major Anions and Cations in water samples and converts major ion concentration to milliequivalents and then summed. Anions sum and Cation sum is calculated as a difference and expressed as a percentage.
AN135	Alkalinity (and forms of) by Titration: The sample is titrated with standard acid to pH 8.3 (P titre) and pH 4.5 (T titre) and permanent and/or total alkalinity calculated. The results are expressed as equivalents of calcium carbonate or recalculated as bicarbonate, carbonate and hydroxide. Reference APHA 2320. Internal Reference AN135
AN274	Chloride by Aquakem DA: Chloride reacts with mercuric thiocyanate forming a mercuric chloride complex. In the presence of ferric iron, highly coloured ferric thiocyanate is formed which is proportional to the chloride concentration. Reference APHA 4500CI-
AN320/AN321	Metals by ICP-OES: Samples are preserved with 10% nitric acid for a wide range of metals and some non-metals. This solution is measured by Inductively Coupled Plasma. Solutions are aspirated into an argon plasma at 8000-10000K and emit characteristic energy or light as a result of electron transitions through unique energy levels. The emitted light is focused onto a diffraction grating where it is separated into components.
AN320/AN321	Photomultipliers or CCDs are used to measure the light intensity at specific wavelengths. This intensity is directly proportional to concentration. Corrections are required to compensate for spectral overlap between elements. Reference APHA 3120 B.
Calculation	Free and Total Carbon Dioxide may be calculated using alkalinity forms only when the samples TDS is <500mg/L. If TDS is >500mg/L free or total carbon dioxide cannot be reported. APHA4500CO2 D.

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FOOTNOTES _

IS Insufficient sample for analysis.

LNR Sample listed, but not received.

* NATA accreditation does not cover the performance of this service.

** Indicative data, theoretical holding time exceeded.

LOR Limit of Reporting

↑↓ Raised or Lowered Limit of Reporting
QFH QC result is above the upper tolerance
QFL QC result is below the lower tolerance

- The sample was not analysed for this analyte

NVL Not Validated

Samples analysed as received.

Solid samples expressed on a dry weight basis.

Where "Total" analyte groups are reported (for example, Total PAHs, Total OC Pesticides) the total will be calculated as the sum of the individual analytes, with those analytes that are reported as <LOR being assumed to be zero. The summed (Total) limit of reporting is calcuated by summing the individual analyte LORs and dividing by two. For example, where 16 individual analytes are being summed and each has an LOR of 0.1 mg/kg, the "Totals" LOR will be 1.6 / 2 (0.8 mg/kg). Where only 2 analytes are being summed, the "Total" LOR will be the sum of those two LORs.

Some totals may not appear to add up because the total is rounded after adding up the raw values.

If reported, measurement uncertainty follow the ± sign after the analytical result and is expressed as the expanded uncertainty calculated using a coverage factor of 2, providing a level of confidence of approximately 95%, unless stated otherwise in the comments section of this report.

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Note that in terms of units of radioactivity:

- a. 1 Bq is equivalent to 27 pCi
- b. 37 MBq is equivalent to 1 mCi

For results reported for samples tested under test methods with codes starting with ARS-SOP, less than (<) values indicate the detection limit for each radionuclide or parameter for the measurement system used. The respective detection limits have been calculated in accordance with ISO 11929.

The QC criteria are subject to internal review according to the SGS QAQC plan and may be provided on request or alternatively can be found here: http://www.sgs.com.au/~/media/Local/Australia/Documents/Technical%20Documents/MP-AU-ENV-QU-022%20QA%20QC%20Plan.pdf

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STATEMENT OF QA/QC **PERFORMANCE**

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1546223 Northern Sands CE124163 R0 SGS Reference Project

(Not specified) 23 Nov 2016 Date Received Order Number 02 Dec 2016 Date Reported Samples

COMMENTS

All the laboratory data for each environmental matrix was compared to SGS' stated Data Quality Objectives (DQO). Comments arising from the comparison were made and are reported below.

The data relating to sampling was taken from the Chain of Custody document and was supplied by the Client.

This QA/QC Statement must be read in conjunction with the referenced Analytical Report.

The Statement and the Analytical Report must not be reproduced except in full.

All Data Quality Objectives were met (within the SGS Cairns Environmental laboratory).

SAMPLE SUMMARY

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HOLDING TIME SUMMARY

SGS holding time criteria are drawn from current regulations and are highly dependent on sample container preservation as specified in the SGS "Field Sampling Guide for Containers and Holding Time" (ref: GU-(AU)-ENV.001). Soil samples guidelines are derived from NEPM "Schedule B(3) Guideline on Laboratory Analysis of Potentially Contaminated Soils". Water sample guidelines are derived from "AS/NZS 5667.1 : 1998 Water Quality - sampling part 1" and APHA "Standard Methods for the Examination of Water and Wastewater" 21st edition 2005.

Extraction and analysis holding time due dates listed are calculated from the date sampled, although holding times may be extended after laboratory extraction for some analytes. The due dates are the suggested dates that samples may be held before extraction or analysis and still be considered valid.

Extraction and analysis dates are shown in Green when within suggested criteria or Red with an appended dagger symbol (†) when outside suggested criteria. If the sampled date is not supplied then compliance with criteria cannot be determined. If the received date is after one or both due dates then holding time will fail by default.

Alkalinity							Method:	ME-(AU)-[ENV]AN13
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
BH01	CE124163.001	LB041247	22 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016
BH02	CE124163.002	LB041247	22 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016
BH03	CE124163.003	LB041247	22 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016
Chloride by Discrete Ana	llyser in Water						Method:	ME-(AU)-[ENV]AN2
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
BH01	CE124163.001	LB041309	22 Nov 2016	23 Nov 2016	20 Dec 2016	25 Nov 2016	20 Dec 2016	28 Nov 2016
BH02	CE124163.002	LB041309	22 Nov 2016	23 Nov 2016	20 Dec 2016	25 Nov 2016	20 Dec 2016	28 Nov 2016
BH03	CE124163.003	LB041309	22 Nov 2016	23 Nov 2016	20 Dec 2016	25 Nov 2016	20 Dec 2016	28 Nov 2016
Conductivity and TDS by	Calculation - Water						Method:	ME-(AU)-[ENV]AN1
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
BH01	CE124163.001	LB041247	22 Nov 2016	23 Nov 2016	20 Dec 2016	23 Nov 2016	20 Dec 2016	23 Nov 2016
BH02	CE124163.002	LB041247	22 Nov 2016	23 Nov 2016	20 Dec 2016	23 Nov 2016	20 Dec 2016	23 Nov 2016
BH03	CE124163.003	LB041247	22 Nov 2016	23 Nov 2016	20 Dec 2016	23 Nov 2016	20 Dec 2016	23 Nov 2016
Metals in Water (Dissolve	ed) by ICPOES						Method: ME-(AU)-[ENV]AN320/AN3
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
BH01	CE124163.001	LB041329	22 Nov 2016	23 Nov 2016	21 May 2017	28 Nov 2016	21 May 2017	28 Nov 2016
BH02	CE124163.002	LB041329	22 Nov 2016	23 Nov 2016	21 May 2017	28 Nov 2016	21 May 2017	28 Nov 2016
BH03	CE124163.003	LB041329	22 Nov 2016	23 Nov 2016	21 May 2017	28 Nov 2016	21 May 2017	28 Nov 2016
pH in water							Method:	ME-(AU)-[ENV]AN1
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
BH01	CE124163.001	LB041247	22 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016
BH02	CE124163.002	LB041247	22 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016
BH03	CE124163.003	LB041247	22 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016	23 Nov 2016

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SURROGATES

CE124163 R0

Surrogate results are evaluated against upper and lower limit criteria established in the SGS QA/QC plan (Ref: MP-(AU)-[ENV]QU-022). At least two of three routine level soil sample surrogate spike recoveries for BTEX/VOC are to be within 70-130% where control charts have not been developed and within the established control limits for charted surrogates. Matrix effects may void this as an acceptance criterion. Water sample surrogate spike recoveries are to be within 40-130%. The presence of emulsions, surfactants and particulates may void this as an acceptance criterion.

Result is shown in Green when within suggested criteria or Red with an appended reason identifier when outside suggested criteria. Refer to the footnotes section at the end of this report for failure reasons.

No surrogates were required for this job.

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METHOD BLANKS

Blank results are evaluated against the limit of reporting (LOR), for the chosen method and its associated instrumentation, typically 2.5 times the statistically determined method detection limit (MDL).

Result is shown in Green when within suggested criteria or Red with an appended dagger symbol (†) when outside suggested criteria.

Alkalinity	Method: ME-(AU)-[ENV]AN135
------------	----------------------------

Sample Number	Parameter	Units	LOR	Result
LB041247.001	Total Alkalinity as CaCO3	mg/L	5	<5
LB041247.028	Total Alkalinity as CaCO3	mg/L	5	<5
LB041247.055	Total Alkalinity as CaCO3	mg/L	5	<5
LB041247.082	Total Alkalinity as CaCO3	mg/L	5	<5

Chloride by Discrete Analyser in Water

Chloride by Discrete Analyser in Water	•		Metho	od: ME-(AU)-[ENV]AN27
Sample Number	Parameter	Units	LOR	Result
LB041309.001	Chloride, Cl	mg/L	1	<1
LB041309.024	Chloride, Cl	ma/L	1	<1

Conductivity and TDS by Calculation - Water

Method: ME-(AU)-[ENV]AN106

Sample Number	Parameter	Units	LOR	Result
LB041247.001	Conductivity @ 25 C	μS/cm	5	<5
	Total Dissolved Solids (by calculation)	mg/L	10	<10
LB041247.028	Conductivity @ 25 C	μS/cm	5	<5
	Total Dissolved Solids (by calculation)	mg/L	10	<10
LB041247.055	Conductivity @ 25 C	μS/cm	5	<5
	Total Dissolved Solids (by calculation)	mg/L	10	<10
LB041247.082	Conductivity @ 25 C	μS/cm	5	<5
	Total Dissolved Solids (by calculation)	mg/L	10	<10

Metals in Water (Dissolved) by ICPOES

Method:	MF-/ALI	LIENV	AN320/AN321

Sample Number	Parameter	Units	LOR	Result
Sample Number LB041329.001	Calcium, Ca	mg/L	0.1	<0.1
	Magnesium, Mg	mg/L	0.1	<0.1
	Potassium, K	mg/L	0.1	<0.1
	Sodium, Na	ma/L	0.5	<0.5

pH in water

Method: ME-(AU)-[ENV]AN101

Sample Number	Parameter	Units	LOR	Result
LB041247.001	pH**	pH Units	0.1	5.6
LB041247.028	pH**	pH Units	0.1	6.2
LB041247.055	pH**	pH Units	0.1	5.8
LB041247.082	pH**	pH Units	0.1	5.8

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DUPLICATES

Duplicates are calculated as Relative Percentage Difference (RPD) using the formula: RPD = | OriginalResult - ReplicateResult | x 100 / Mean

The RPD is evaluated against the Maximum Allowable Difference (MAD) criteria and can be graphically represented by a curve calculated from the Statistical Detection Limit (SDL) and Limiting Repeatability (LR) using the formula: MAD = 100 x SDL / Mean + LR

Where the Maximum Allowable Difference evaluates to a number larger than 200 it is displayed as 200.

RPD is shown in Green when within suggested criteria or Red with an appended reason identifer when outside suggested criteria. Refer to the footnotes section at the end of this report for failure reasons.

Alkalinity Method: ME-(AU)-[ENV]AN135

Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
CE124106.001	LB041247.123	Total Alkalinity as CaCO3	mg/L	5	81	80	21	2
	LB041247.132	Total Alkalinity as CaCO3	mg/L	5	81	80	21	2
CE124111.001	LB041247.125	Total Alkalinity as CaCO3	mg/L	5	380	380	16	0
CE124122.001	LB041247.126	Total Alkalinity as CaCO3	mg/L	5	5	6	105	5
CE124125.001	LB041247.127	Total Alkalinity as CaCO3	mg/L	5	200	200	17	1
CE124133.001	LB041247.128	Total Alkalinity as CaCO3	mg/L	5	120	130	19	1
CE124141.001	LB041247.129	Total Alkalinity as CaCO3	mg/L	5	20	17	42	15
CE124142.001	LB041247.130	Total Alkalinity as CaCO3	mg/L	5	240	240	17	0
CE124145.001	LB041247.134	Total Alkalinity as CaCO3	mg/L	5	510	500	16	1
CE124145.011	LB041247.135	Total Alkalinity as CaCO3	mg/L	5	830	850	16	2
CE124146.001	LB041247.137	Total Alkalinity as CaCO3	mg/L	5	14	14	50	1
CE124163.001	LB041247.140	Total Alkalinity as CaCO3	mg/L	5	46	46	26	0

Chloride by Discrete Analyser in Water

Method: ME-(AU)-[ENV]AN274

Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
CE124143.001	LB041309.005	Chloride, Cl	mg/L	1	33	32	18	1
CE124145.006	LB041309.016	Chloride, Cl	mg/L	1	450	450	15	0
CE124146.005	LB041309.030	Chloride, Cl	mg/L	1	17	17	21	0

Conductivity and TDS by Calculation - Water

Method: ME-(AU)-[ENV]AN106

Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
CE124106.001	LB041247.123	Conductivity @ 25 C	μS/cm	5	150	150	16	1
		Total Dissolved Solids (by calculation)	mg/L	10	88	88	17	1
	LB041247.132	Conductivity @ 25 C	μS/cm	5	150	150	16	1
		Total Dissolved Solids (by calculation)	mg/L	10	88	88	17	1
CE124111.001	LB041247.125	Conductivity @ 25 C	μS/cm	5	1000	1100	15	1
CE124125.001	LB041247.127	Conductivity @ 25 C	μS/cm	5	770	780	15	0
CE124133.001	LB041247.128	Conductivity @ 25 C	μS/cm	5	530	530	15	0
		Total Dissolved Solids (by calculation)	mg/L	10	320	320	16	0
CE124141.001	LB041247.129	Conductivity @ 25 C	μS/cm	5	11.944000244	1 1.4734611511	20	1
CE124142.001	LB041247.130	Conductivity @ 25 C	μS/cm	5	680	680	15	1
	LB041247.131	Conductivity @ 25 C	μS/cm	5	680	680	15	1
CE124146.001	LB041247.144	Conductivity @ 25 C	μS/cm	5	89	88	17	1
		Total Dissolved Solids (by calculation)	mg/L	10	53	53	19	1
CE124163.001	LB041247.147	Conductivity @ 25 C	μS/cm	5	250	250	16	1
		Total Dissolved Solids (by calculation)	mg/L	10	150	150	16	1

Metals in Water (Dissolved) by ICPOES

Method: ME-(AU)-[ENV]AN320/AN321

Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
CE124171.001	LB041329.014	Sodium, Na	mg/L	0.5	35	35	16	0
		Sulphur as Sulphate, SO4	mg/L	0.5	28	28	17	0

pH in water

Method: ME-(AU)-[ENV]AN101

Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
CE124106.001	LB041247.123	pH**	pH Units	0.1	7.7	7.8	16	0
	LB041247.132	pH**	pH Units	0.1	7.7	7.8	16	0
CE124111.001	LB041247.125	pH**	pH Units	0.1	7.4	7.3	16	2
	LB041247.134	pH**	pH Units	0.1	7.4	7.3	16	2
CE124122.001	LB041247.126	pH**	pH Units	0.1	6.4	6.6	17	3
	LB041247.135	pH**	pH Units	0.1	6.4	6.6	17	3
CE124125.001	LB041247.127	pH**	pH Units	0.1	7.9	7.9	16	0
	LB041247.136	pH**	pH Units	0.1	7.9	7.9	16	0
CE124133.001	LB041247.128	pH**	pH Units	0.1	8.0	8.1	16	1
	LB041247.137	pH**	pH Units	0.1	8.0	8.1	16	1
CE124141.001	LB041247.129	pH**	pH Units	0.1	6.8	6.6	16	2
	LB041247.138	pH**	pH Units	0.1	6.8	6.6	16	2
CE124142.001	LB041247.130	pH**	pH Units	0.1	7.5	7.5	16	0
	LB041247.131	pH**	pH Units	0.1	7.5	7.8	16	4
	LB041247.139	pH**	pH Units	0.1	7.5	7.5	16	0
	LB041247.140	pH**	pH Units	0.1	7.5	7.8	16	4
CE124145.001	LB041247.141	pH**	pH Units	0.1	7.9	7.9	16	0

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DUPLICATES

CE124163 R0

Duplicates are calculated as Relative Percentage Difference (RPD) using the formula: RPD = | OriginalResult - ReplicateResult | x 100 / Mean

The RPD is evaluated against the Maximum Allowable Difference (MAD) criteria and can be graphically represented by a curve calculated from the Statistical Detection Limit (SDL) and Limiting Repeatability (LR) using the formula: MAD = 100 x SDL / Mean + LR

Where the Maximum Allowable Difference evaluates to a number larger than 200 it is displayed as 200.

RPD is shown in Green when within suggested criteria or Red with an appended reason identifer when outside suggested criteria. Refer to the footnotes section at the end of this report for failure reasons.

pH in water (continued)

Method: ME-(AU)-[ENV]AN101

Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
CE124145.011	LB041247.142	pH**	pH Units	0.1	7.9	8.0	16	1
CE124146.001	LB041247.144	pH**	pH Units	0.1	7.1	7.0	16	1
CE124157.001	LB041247.145	pH**	pH Units	0.1	6.8	6.9	16	2
CE124157.011	LB041247.146	pH**	pH Units	0.1	7.0	7.0	16	0
CE124163.001	LB041247.147	pH**	pH Units	0.1	6.8	6.8	16	0

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7.415

98 - 102

100



LB041247.087

pH**

LABORATORY CONTROL SAMPLES

Laboratory Control Standard (LCS) results are evaluated against an expected result, typically the concentration of analyte spiked into the control during the sample preparation stage, producing a percentage recovery. The criteria applied to the percentage recovery is established in the SGS QA /QC plan (Ref: MP-(AU)-[ENV]QU-022). For more information refer to the footnotes in the concluding page of this report.

Recovery is shown in Green when within suggested criteria or Red with an appended dagger symbol (†) when outside suggested criteria.

Alkalinity					1	Method: ME-(A	.U)-[ENV]AN13
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB041247.002	Total Alkalinity as CaCO3	mg/L	5	64	59.5	80 - 120	107
LB041247.003	Total Alkalinity as CaCO3	mg/L	5	240	229	80 - 120	104
LB041247.029	Total Alkalinity as CaCO3	mg/L	5	69	59.5	80 - 120	115
LB041247.030	Total Alkalinity as CaCO3	mg/L	5	240	229	80 - 120	103
LB041247.056	Total Alkalinity as CaCO3	mg/L	5	67	59.5	80 - 120	113
LB041247.057	Total Alkalinity as CaCO3	mg/L	5	240	229	80 - 120	104
LB041247.083	Total Alkalinity as CaCO3	mg/L	5	65	59.5	80 - 120	108
LB041247.084	Total Alkalinity as CaCO3	mg/L	5	240	229	80 - 120	103
Chloride by Discrete Analyse	er in Water					Method: ME-(A	U)-[ENV]AN27
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB041309.002	Chloride, Cl	mg/L	1	130	125	80 - 120	103
LB041309.003	Chloride, Cl	mg/L	1	10	10	80 - 120	104
LB041309.025	Chloride, Cl	mg/L	1	130	125	80 - 120	103
LB041309.026	Chloride, Cl	mg/L	1	10	10	80 - 120	104
Conductivity and TDS by Cal	culation - Water					Method: ME-(A	U)-[ENV]AN10
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB041247.004	Conductivity @ 25 C	μS/cm	5	300	303	90 - 110	98
LB041247.005	Conductivity @ 25 C	μS/cm	5	59000	58670	90 - 110	100
LB041247.031	Conductivity @ 25 C	μS/cm	5	290	303	90 - 110	97
LB041247.032	Conductivity @ 25 C	μS/cm	5	59000	58670	90 - 110	100
LB041247.058	Conductivity @ 25 C	μS/cm	5	310	303	90 - 110	102
LB041247.059	Conductivity @ 25 C	μS/cm	5	59000	58670	90 - 110	100
LB041247.085	Conductivity @ 25 C	μS/cm	5	310	303	90 - 110	101
LB041247.086	Conductivity @ 25 C	μS/cm	5	59000	58670	90 - 110	100
Metals in Water (Dissolved)	by ICPOES				Method	: ME-(AU)-[EN	V]AN320/AN32
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB041329.002	Calcium, Ca	mg/L	0.1	21	20	80 - 120	103
	Magnesium, Mg	mg/L	0.1	20	20	80 - 120	100
	Potassium, K	mg/L	0.1	21	20	80 - 120	103
	Sodium, Na	mg/L	0.5	19	20	80 - 120	96
pH in water						Method: ME-(A	U)-[ENV]AN10
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB041247.006	pH**	pH Units	0.1	7.4	7.415	98 - 102	100
LB041247.033	pH**	pH Units	0.1	7.4	7.415	98 - 102	100
LB041247.060	pH**	pH Units	0.1	7.4	7.415	98 - 102	100

pH Units

0.1

7.4

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MATRIX SPIKES

CE124163 R0

Matrix Spike (MS) results are evaluated as the percentage recovery of an expected result, typically the concentration of analyte spiked into a field sub-sample during the sample preparation stage. The original sample's result is subtracted from the sub-sample result before determining the percentage recovery. The criteria applied to the percentage recovery is established in the SGS QA/QC plan (ref: MP-(AU)-[ENV]QU-022). For more information refer to the footnotes in the concluding page of this report.

Recovery is shown in Green when within suggested criteria or Red with an appended reason identifer when outside suggested criteria. Refer to the footnotes section at the end of this report for failure reasons.

Metals in Water (Dissolved) by ICPOES

Method: ME-(AU)-[ENV]AN320/AN321

QC Sample	Sample Number	Parameter	Units	LOR	Result	Original	Spike	Recovery%
CE124146.001	LB041329.004	Calcium, Ca	mg/L	0.1	57	1.8	50	111
		Magnesium, Mg	mg/L	0.1	55	1.6	50	106
		Potassium, K	mg/L	0.1	58	1.3	50	113
		Sodium, Na	mg/L	0.5	62	13	50	100

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MATRIX SPIKE DUPLICATES

CE124163 R0

Matrix spike duplicates are calculated as Relative Percent Difference (RPD) using the formula: RPD = | OriginalResult - ReplicateResult | x 100 / Mean

The original result is the analyte concentration of the matrix spike. The Duplicate result is the analyte concentration of the matrix spike duplicate.

The RPD is evaluated against the Maximum Allowable Difference (MAD) criteria and can be graphically represented by a curve calculated from the Statistical Detection Limit (SDL) and Limiting Repeatability (LR) using the formula: MAD = 100 x SDL / Mean + LR

Where the Maximum Allowable Difference evaluates to a number larger than 200 it is displayed as 200.

RPD is shown in Green when within suggested criteria or Red with an appended reason identifer when outside suggested criteria. Refer to the footnotes section at the end of this report for failure reasons.

No matrix spike duplicates were required for this job.

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FOOTNOTES



Samples analysed as received.

Solid samples expressed on a dry weight basis.

QC criteria are subject to internal review according to the SGS QA/QC plan and may be provided on request or alternatively can be found here: http://www.sgs.com.au/~/media/Local/Australia/Documents/Technical Documents/MP-AU-ENV-QU-022 QA QC Plan.pdf

- * NATA accreditation does not cover the performance of this service.
- Sample not analysed for this analyte.

IS Insufficient sample for analysis.

LNR Sample listed, but not received.

LOR Limit of reporting.

QFH QC result is above the upper tolerance.

QFL QC result is below the lower tolerance.

- ① At least 2 of 3 surrogates are within acceptance criteria.
- 2 RPD failed acceptance criteria due to sample heterogeneity.
- 3 Results less than 5 times LOR preclude acceptance criteria for RPD.
- Recovery failed acceptance criteria due to matrix interference.
- ® Recovery failed acceptance criteria due to the presence of significant concentration of analyte (i.e. the concentration of analyte exceeds the spike level).
- © LOR was raised due to sample matrix interference.
- ① LOR was raised due to dilution of significantly high concentration of analyte in sample.
- ® Reanalysis of sample in duplicate confirmed sample heterogeneity and inconsistency of results.
- Low surrogate recovery due to the sample emulsifying during extraction.
- © Legionella Test Result <10 cfu/mL</p>

Control Strategy (1)

Maintain Monthly Program or at least 3-monthly monitoring. Maintain water treatment program

(f) Legionella Test Result <1000 cfu/mL

Control Strategy (2)

Investigate problem. Review water treatment program. Take necessary remedial action (including immediate online disinfection)

and undertake control strategy 3.

© Control Strategy (3)

Retest water within 3 to 7 days of plant operation. If not detected, continue to retest water every 3 to 7 days until 2 consecutive samples

return readings of 'not detected', then repeat control strategy (1).

If detected at <100 cfu/mL, repeat control strategy (2)

If detected at >100 cfu/mL, investigate the problem and review water treatment program,

and immediately carry out online decontamination.

If detected at >1000 cfu/mL, undertake control strategy (4).

Legionella Test Result >1000 cfu/mL

Control Strategy (4)

Investigate problem. Review water treatment program.

Take necessary remedial action (including immediate online decontamination)

and undertake control strategy (5).

(5) Control Strategy

Retest water within 3 to 7 days of plant operation. If not detected, continue to retest water every 3 to 7 days until 2 consecutive samples

return readings of 'not detected', then repeat control strategy (1).

If detected at <100 cfu/mL, repeat control strategy (1)

If detected at >100 and <1000 cfu/mL, investigate the problem and review water treatment program, immediately carry out online decontamination.

and repeat control strategy (5).

If detected at >1000 cfu/mL, investigate and review the water treatment program,

immediately carry out system decontamination

and repeat control strategy (5).

(5) HPC - Test Result < 100 000 cfu/mL

Control Strategy (1)

Maintain Monthly Program. Maintain water treatment program.

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FOOTNOTES

Samples analysed as received.

Solid samples expressed on a dry weight basis.

QC criteria are subject to internal review according to the SGS QA/QC plan and may be provided on request or alternatively can be found here: http://www.sgs.com.au/~/media/Local/Australia/Documents/Technical Documents/MP-AU-ENV-QU-022 QA QC Plan.pdf

6 HPC Test Result > 100 000 cfu/mL <5 000 000 cfu/mL</p>

Control Strategy (2)

Investigate problem. Review water treatment program. Take necessary remedial action (including immediate online disinfection)

and undertake control strategy 3.

① Control Strategy (3)

Retest water within 3 to 7 days of plant operation. If the test result is <100 000 cfu/mL, repeat control strategy (1) If the test result is >100 000 cfu/mL but <5 000 000 cfu/mL, undertake control strategy (2).

If the test result is >5 000 000 cfu/mL, undertake control strategy (4).

® HPC Test Result >5 000 000 cfu/mL

Control Strategy (4)

Investigate problem. Review water treatment program. Take necessary remedial action (including immediate online disinfection)

and undertake control strategy (5).

(5) Control Strategy

Retest water within 3 to 7 days of plant operation. If the test result is <100 000 cfu/mL, repeat control strategy (1)

If the test result is >100 000 cfu/mL but <5 000 000 cfu/mL, undertake control strategy (4).

If the test result is >5 000 000 cfu/mL, investigate the problem.

review the water treatment program, and carry out immediate online decontamination.

Enterococci - Median result should not exceed 230 cfu/100mL

(maximum number in any one sample: 450-700 cfu/100mL) Sourced from NHMRC (National Health and Medical Research Council)

& NWQMS (National Water Quality Management Strategy)

-Australian Guidelines for Recreational Use of Water. Version Oct 2000.

† Refer to Analytical Report comments for further information.

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07 Dec 2016

Jon Dicker

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SGS Reference CE124214 R0 Date Received 25 Nov 2016

Date Reported

COMMENTS

Accredited for compliance with ISO/IEC 17025. NATA accredited laboratory 2562(3146)

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Quality & Microbiology Coordinator

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CE124214 R0

Sample Number Cl Sample Matrix Sample Date 2 Sample Name

Parameter Units LOR

pH in water Method: AN101 Tested: 25/11/2016

pH** pH Units 0.1 **6.5**

Conductivity and TDS by Calculation - Water Method: AN106 Tested: 25/11/2016

Conductivity @ 25 C	μS/cm	5	160
Total Dissolved Solids (by calculation)	mg/L	10	93

Alkalinity Method: AN135 Tested: 25/11/2016

Total Alkalinity as CaCO3	mg/L	5	50
Bicarbonate Alkalinity as CaCO3	mg/L	5	50
Carbonate Alkalinity as CaCO3	mg/L	5	<5
Hydroxide Alkalinity as CaCO3	mg/L	5	<5

Chloride by Discrete Analyser in Water Method: AN274 Tested: 30/11/2016

Metals in Water (Dissolved) by ICPOES Method: AN320/AN321 Tested: 5/12/2016

Calcium, Ca	mg/L	0.1	8.0
Magnesium, Mg	mg/L	0.1	3.2
Potassium, K	mg/L	0.1	0.8
Sodium, Na	mg/L	0.5	17
Sulphur as Sulphate, SO4	mg/L	0.5	23
Total Hardness by Calculation	mg CaCO3/L	1	33

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CE124214 R0

		ample Number Sample Matrix Sample Date Sample Name	CE124214.001 Water 25 Nov 2016 GA04
Parameter	Units	LOR	

Calculation of Anion-C	ation Balance (SAR C	Ic) Method: AN121	Tested: 6/12/2016

Sum of Cation Milliequivalents*	meq/L	-	1.43
Sum of Anion Milliequivalents*	meq/L	-	2.27
Anion-Cation Balance	%	-100	-23

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QC SUMMARY

MB blank results are compared to the Limit of Reporting

LCS and MS spike recoveries are measured as the percentage of analyte recovered from the sample compared the the amount of analyte spiked into the sample.

DUP and MSD relative percent differences are measured against their original counterpart samples according to the formula: the absolute difference of the two results divided by the average of the two results as a percentage. Where the DUP RPD is 'NA', the results are less than the LOR and thus the RPD is not applicable.

Alkalinity Method: ME-(AU)-[ENV]AN135

Parameter	QC Reference	Units	LOR	МВ	LCS %Recovery
Total Alkalinity as CaCO3	LB041324	mg/L	5	<5	110 - 119%
Bicarbonate Alkalinity as CaCO3	LB041324	mg/L	5	<5	
Carbonate Alkalinity as CaCO3	LB041324	mg/L	5	<5	
Hydroxide Alkalinity as CaCO3	LB041324	mg/L	5	<5	

Chloride by Discrete Analyser in Water Method: ME-(AU)-[ENV]AN274

ı	Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
ı		Reference					%Recovery
ı	Chloride, Cl	LB041417	mg/L	1	<1	0 - 2%	104%

Conductivity and TDS by Calculation - Water Method: ME-(AU)-[ENV]AN106

Parameter	QC	Units	LOR	MB	LCS
	Reference				%Recovery
Conductivity @ 25 C	LB041324	μS/cm	5	<5	99 - 100%
Total Dissolved Solids (by calculation)	LB041324	mg/L	10	<10	NA

Metals in Water (Dissolved) by ICPOES Method: ME-(AU)-[ENV]AN320/AN321

Parameter	QC Reference	Units	LOR	MB	DUP %RPD	LCS %Recovery	MS %Recovery
Calcium, Ca	LB041532	mg/L	0.1	<0.1		103%	NVL
Magnesium, Mg	LB041532	mg/L	0.1	<0.1		101%	NVL
Potassium, K	LB041532	mg/L	0.1	<0.1		107%	NVL
Sodium, Na	LB041532	mg/L	0.5	<0.5		97%	NVL
Sulphur as Sulphate, SO4	LB041532	mg/L	0.5	<0.5	NVL	NA	
Total Hardness by Calculation	LB041532	mg CaCO3/L	1	<1			

pH in water Method: ME-(AU)-[ENV]AN101

ı	Parameter	QC	Units	LOR	MB	LCS	
1		Reference				%Recovery	
ı	pH**	LB041324	pH Units	0.1	5.6 - 5.7	100%	

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METHOD SUMMARY

METHOD	METHODOLOGY SUMMARY
ETTTOD	WETHODOLOGT SUNIWART
AN101	pH in Soil Sludge Sediment and Water: pH is measured electrometrically using a combination electrode (glass plus reference electrode) and is calibrated against 3 buffers purchased commercially. For soils, an extract with water is made at a ratio of 1:5 and the pH determined and reported on the extract. Reference APHA 4500-H+.
AN106	Conductivity and TDS by Calculation: Conductivity is measured by meter with temperature compensation and is calibrated against a standard solution of potassium chloride. Conductivity is generally reported as μ mhos/cm or μ S/cm @ 25°C. For soils, an extract with water is made at a ratio of 1:5 and the EC determined and reported on the extract, or calculated back to the as-received sample. Total Dissolved Salts can be estimated from conductivity using a conversion factor, which for natural waters, is in the range 0.55 to 0.75. SGS use 0.6. Reference APHA 2510 B.
AN121	This method is used to calculation the balance of major Anions and Cations in water samples and converts major ion concentration to milliequivalents and then summed. Anions sum and Cation sum is calculated as a difference and expressed as a percentage.
AN135	Alkalinity (and forms of) by Titration: The sample is titrated with standard acid to pH 8.3 (P titre) and pH 4.5 (T titre) and permanent and/or total alkalinity calculated. The results are expressed as equivalents of calcium carbonate or recalculated as bicarbonate, carbonate and hydroxide. Reference APHA 2320. Internal Reference AN135
AN274	Chloride by Aquakem DA: Chloride reacts with mercuric thiocyanate forming a mercuric chloride complex. In the presence of ferric iron, highly coloured ferric thiocyanate is formed which is proportional to the chloride concentration. Reference APHA 4500Cl-
AN320/AN321	Metals by ICP-OES: Samples are preserved with 10% nitric acid for a wide range of metals and some non-metals. This solution is measured by Inductively Coupled Plasma. Solutions are aspirated into an argon plasma at 8000-10000K and emit characteristic energy or light as a result of electron transitions through unique energy levels. The emitted light is focused onto a diffraction grating where it is separated into components.
AN320/AN321	Photomultipliers or CCDs are used to measure the light intensity at specific wavelengths. This intensity is directly proportional to concentration. Corrections are required to compensate for spectral overlap between elements. Reference APHA 3120 B.
Calculation	Free and Total Carbon Dioxide may be calculated using alkalinity forms only when the samples TDS is <500mg/L. If TDS is >500mg/L free or total carbon dioxide cannot be reported. APHA4500CO2 D.

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FOOTNOTES _

IS Insufficient sample for analysis.

LNR Sample listed, but not received.

* NATA accreditation does not cover the performance of this service.

** Indicative data, theoretical holding time exceeded.

LOR Limit of Reporting

↑↓ Raised or Lowered Limit of Reporting QFH QC result is above the upper tolerance QFL QC result is below the lower tolerance

- The sample was not analysed for this analyte

NVL Not Validated

Samples analysed as received.

Solid samples expressed on a dry weight basis.

Where "Total" analyte groups are reported (for example, Total PAHs, Total OC Pesticides) the total will be calculated as the sum of the individual analytes, with those analytes that are reported as <LOR being assumed to be zero. The summed (Total) limit of reporting is calcuated by summing the individual analyte LORs and dividing by two. For example, where 16 individual analytes are being summed and each has an LOR of 0.1 mg/kg, the "Totals" LOR will be 1.6 / 2 (0.8 mg/kg). Where only 2 analytes are being summed, the "Total" LOR will be the sum of those two LORs.

Some totals may not appear to add up because the total is rounded after adding up the raw values.

If reported, measurement uncertainty follow the ± sign after the analytical result and is expressed as the expanded uncertainty calculated using a coverage factor of 2, providing a level of confidence of approximately 95%, unless stated otherwise in the comments section of this report.

Results reported for samples tested under test methods with codes starting with ARS-SOP, radionuclide or gross radioactivity concentrations are expressed in becquerel (Bq) per unit of mass or volume or per wipe as stated on the report. Becquerel is the SI unit for activity and equals one nuclear transformation per second.

Note that in terms of units of radioactivity:

- a. 1 Bq is equivalent to 27 pCi
- b. 37 MBq is equivalent to 1 mCi

For results reported for samples tested under test methods with codes starting with ARS-SOP, less than (<) values indicate the detection limit for each radionuclide or parameter for the measurement system used. The respective detection limits have been calculated in accordance with ISO 11929.

The QC criteria are subject to internal review according to the SGS QAQC plan and may be provided on request or alternatively can be found here: http://www.sgs.com.au/~/media/Local/Australia/Documents/Technical%20Documents/MP-AU-ENV-QU-022%20QA%20QC%20Plan.pdf

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STATEMENT OF QA/QC **PERFORMANCE**

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1546223 Northern Sands CE124214 R0 SGS Reference Project 25 Nov 2016 (Not specified) Order Number Date Received 07 Dec 2016 Date Reported

COMMENTS

Samples

All the laboratory data for each environmental matrix was compared to SGS' stated Data Quality Objectives (DQO). Comments arising from the comparison were made and are reported below.

The data relating to sampling was taken from the Chain of Custody document and was supplied by the Client.

This QA/QC Statement must be read in conjunction with the referenced Analytical Report.

The Statement and the Analytical Report must not be reproduced except in full.

All Data Quality Objectives were met (within the SGS Cairns Environmental laboratory).

SAMPLE SUMMARY

Samples clearly labelled Yes Complete documentation received Yes Sample container provider SGS Sample cooling method Ice Samples received in correct containers Yes Sample counts by matrix 1 water 25/11/2016 COC Date documentation received Type of documentation received Number of eskies/boxes received Samples received in good order Yes Samples received without headspace Yes Sample temperature upon receipt Chilled Sufficient sample for analysis Turnaround time requested Yes standard

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Alkalinity

GA04

Sample Na

Sample Nan

GA04

HOLDING TIME SUMMARY

CE124214 R0

Method: ME-(AU)-[ENV]AN135

25 Nov 2016

25 Nov 2016

Analysis Due Analysed

26 Nov 2016

Analysis Due

26 Nov 2016

SGS holding time criteria are drawn from current regulations and are highly dependent on sample container preservation as specified in the SGS "Field Sampling Guide for Containers and Holding Time" (ref: GU-(AU)-ENV.001). Soil samples guidelines are derived from NEPM "Schedule B(3) Guideline on Laboratory Analysis of Potentially Contaminated Soils". Water sample guidelines are derived from "AS/NZS 5667.1 : 1998 Water Quality - sampling part 1" and APHA "Standard Methods for the Examination of Water and Wastewater" 21st edition 2005.

Extraction and analysis holding time due dates listed are calculated from the date sampled, although holding times may be extended after laboratory extraction for some analytes. The due dates are the suggested dates that samples may be held before extraction or analysis and still be considered valid.

Extraction and analysis dates are shown in Green when within suggested criteria or Red with an appended dagger symbol (†) when outside suggested criteria. If the sampled date is not supplied then compliance with criteria cannot be determined. If the received date is after one or both due dates then holding time will fail by default.

25 Nov 2016

Extraction Due

26 Nov 2016

Extraction Due

26 Nov 2016

Extracted

25 Nov 2016

25 Nov 2016

QC Ref

LB041324

LB041324

25 Nov 2016

25 Nov 2016

CE124214.001

CE124214.001

Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
GA04	CE124214.001	LB041417	25 Nov 2016	25 Nov 2016	23 Dec 2016	30 Nov 2016	23 Dec 2016	01 Dec 2016
Conductivity and TDS by	Calculation - Water						Method:	ME-(AU)-[ENV]AN
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
GA04	CE124214.001	LB041324	25 Nov 2016	25 Nov 2016	23 Dec 2016	25 Nov 2016	23 Dec 2016	25 Nov 2016
letals in Water (Dissolve	ed) by ICPOES						Method: ME-(AU)-[ENV]AN320/AN
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
GA04	CE124214.001	LB041532	25 Nov 2016	25 Nov 2016	24 May 2017	05 Dec 2016	24 May 2017	06 Dec 2016

25 Nov 2016

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SURROGATES

CE124214 R0

Surrogate results are evaluated against upper and lower limit criteria established in the SGS QA/QC plan (Ref: MP-(AU)-[ENV]QU-022). At least two of three routine level soil sample surrogate spike recoveries for BTEX/VOC are to be within 70-130% where control charts have not been developed and within the established control limits for charted surrogates. Matrix effects may void this as an acceptance criterion. Water sample surrogate spike recoveries are to be within 40-130%. The presence of emulsions, surfactants and particulates may void this as an acceptance criterion.

Result is shown in Green when within suggested criteria or Red with an appended reason identifier when outside suggested criteria. Refer to the footnotes section at the end of this report for failure reasons.

No surrogates were required for this job.

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Method: ME-(AU)-[ENV]AN274



Chloride by Discrete Analyser in Water

METHOD BLANKS

Blank results are evaluated against the limit of reporting (LOR), for the chosen method and its associated instrumentation, typically 2.5 times the statistically determined method detection limit (MDL).

Result is shown in Green when within suggested criteria or Red with an appended dagger symbol (†) when outside suggested criteria.

Alkalinity	Meth	Method: ME-(AU)-[ENV]AN135		
Sample Number	Parameter	Units	LOR	Result
LB041324.001	Total Alkalinity as CaCO3	mg/L	5	<5
LB041324.028	Total Alkalinity as CaCO3	mg/L	5	<5

Sample Number	Parameter	Units	LOR	Result
LB041417.001	Chloride, Cl	mg/L	1	<1
LB041417.024	Chloride, Cl	mg/L	1	<1
LB041417.047	Chloride, Cl	mg/L	1	<1
LB041417.070	Chloride, Cl	mg/L	1	<1
Conductivity and TDS by Calculation -	Water		Metho	od: ME-(AU)-[ENV]AN10
Sample Number	Parameter	Units	LOR	Result
LB041324.001	Conductivity @ 25 C	μS/cm	5	<5

Parameter	Units	LOR	Result
Conductivity @ 25 C	μS/cm	5	<5
Total Dissolved Solids (by calculation)	mg/L	10	<10
Conductivity @ 25 C	μS/cm	5	<5
Total Dissolved Solids (by calculation)	mg/L	10	<10
	Conductivity @ 25 C Total Dissolved Solids (by calculation) Conductivity @ 25 C	Conductivity @ 25 C μS/cm Total Dissolved Solids (by calculation) mg/L Conductivity @ 25 C μS/cm	Conductivity @ 25 C μS/cm 5 Total Dissolved Solids (by calculation) mg/L 10 Conductivity @ 25 C μS/cm 5

Metals in Water (Dissolved) by ICPOES				Method: ME-(AU)-[ENV]AN320/AN321		
Sample Number	Parameter	Units	LOR	Result		
LB041532.001	Calcium, Ca	mg/L	0.1	<0.1		
	Magnesium, Mg	mg/L	0.1	<0.1		
	Potassium, K	mg/L	0.1	<0.1		
	Sodium, Na	mg/L	0.5	<0.5		

pH in water			Meth	od: ME-(AU)-[ENV]AN101
Sample Number	Parameter	Units	LOR	Result
LB041324.001	pH**	pH Units	0.1	5.7
LB041324.028	pH**	pH Units	0.1	5.6

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DUPLICATES

CE124214 R0

Duplicates are calculated as Relative Percentage Difference (RPD) using the formula: RPD = | OriginalResult - ReplicateResult | x 100 / Mean

The RPD is evaluated against the Maximum Allowable Difference (MAD) criteria and can be graphically represented by a curve calculated from the Statistical Detection Limit (SDL) and Limiting Repeatability (LR) using the formula: MAD = 100 x SDL / Mean + LR

Where the Maximum Allowable Difference evaluates to a number larger than 200 it is displayed as 200.

RPD is shown in Green when within suggested criteria or Red with an appended reason identifer when outside suggested criteria. Refer to the footnotes section at the end of this report for failure reasons.

Chloride by Discrete Analyser in Water

Method: ME-(AU)-[ENV]AN274

Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
CE124193.001	LB041417.005	Chloride, Cl	mg/L	1	590	590	15	1
CE124202.010	LB041417.016	Chloride, Cl	mg/L	1	167.414	168.798	16	1
CE124202.030	LB041417.041	Chloride, Cl	mg/L	1	173.93	169.787	16	2
CE124202.040	LB041417.055	Chloride, Cl	mg/L	1	731.413	722.563	15	1
CE124241.002	LB041417.066	Chloride, Cl	mg/L	1	16974.816	16908.926	15	0
CE124241.012	LB041417.080	Chloride, Cl	mg/L	1	544.711	536.859	15	1
CE124242.001	LB041417.091	Chloride, Cl	mg/L	1	210	210	15	0

Metals in Water (Dissolved) by ICPOES

Method: ME-(AU)-[ENV]AN320/AN321

Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
CE124298.006	LB041532.014	Sulphur as Sulphate, SO4	mg/L	0.5	44.9664	45.0432	16	0
CE124298.009	LB041532.018	Sulphur as Sulphate, SO4	mg/L	0.5	0.2559888	0.2457522	200	0

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pH in water

LB041324.006

LB041324.033

Sample Number

pH**

pH**

LABORATORY CONTROL SAMPLES

CE124214 R0

Method: ME-(AU)-[ENV]AN101

100

100

98 - 102

98 - 102

LOR Result Expected Criteria % Recovery %

7.415

7.415

Laboratory Control Standard (LCS) results are evaluated against an expected result, typically the concentration of analyte spiked into the control during the sample preparation stage, producing a percentage recovery. The criteria applied to the percentage recovery is established in the SGS QA /QC plan (Ref: MP-(AU)-[ENV]QU-022). For more information refer to the footnotes in the concluding page of this report.

Recovery is shown in Green when within suggested criteria or Red with an appended dagger symbol (†) when outside suggested criteria.

Alkalinity						Method: ME-(A	.U)-[ENV]AN13
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB041324.002	Total Alkalinity as CaCO3	mg/L	5	71	59.5	80 - 120	119
LB041324.003	Total Alkalinity as CaCO3	mg/L	5	230	229	80 - 120	102
LB041324.029	Total Alkalinity as CaCO3	mg/L	5	66	59.5	80 - 120	110
LB041324.030	Total Alkalinity as CaCO3	mg/L	5	230	229	80 - 120	102
Chloride by Discrete Analyse	er in Water				1	Method: ME-(A	U)-[ENV]AN27
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB041417.002	Chloride, Cl	mg/L	1	130	125	80 - 120	104
LB041417.003	Chloride, Cl	mg/L	1	10	10	80 - 120	104
LB041417.025	Chloride, Cl	mg/L	1	130	125	80 - 120	104
LB041417.026	Chloride, Cl	mg/L	1	10	10	80 - 120	104
LB041417.048	Chloride, Cl	mg/L	1	130	125	80 - 120	104
LB041417.049	Chloride, Cl	mg/L	1	10	10	80 - 120	104
LB041417.071	Chloride, Cl	mg/L	1	130	125	80 - 120	104
LB041417.072	Chloride, Cl	mg/L	1	10	10	80 - 120	104
Conductivity and TDS by Ca	Iculation - Water				1	Method: ME-(A	U)-[ENV]AN10
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB041324.004	Conductivity @ 25 C	μS/cm	5	310	303	90 - 110	102
LB041324.005	Conductivity @ 25 C	μS/cm	5	58000	58670	90 - 110	100
LB041324.031	Conductivity @ 25 C	μS/cm	5	310	303	90 - 110	102
LB041324.032	Conductivity @ 25 C	μS/cm	5	58000	58670	90 - 110	99
Metals in Water (Dissolved)	by ICPOES				Method:	: ME-(AU)-[EN	V]AN320/AN32
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB041532.002	Calcium, Ca	mg/L	0.1	21	20	80 - 120	103
	Magnesium, Mg	mg/L	0.1	20	20	80 - 120	101
	Potassium, K	mg/L	0.1	21	20	80 - 120	107
	Sodium, Na	mg/L	0.5	19	20	80 - 120	97

pH Units

pH Units

0.1

0.1

7.4

7.4

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MATRIX SPIKES

CE124214 R0

Matrix Spike (MS) results are evaluated as the percentage recovery of an expected result, typically the concentration of analyte spiked into a field sub-sample during the sample preparation stage. The original sample's result is subtracted from the sub-sample result before determining the percentage recovery. The criteria applied to the percentage recovery is established in the SGS QA/QC plan (ref: MP-(AU)-[ENV]QU-022). For more information refer to the footnotes in the concluding page of this report.

Recovery is shown in Green when within suggested criteria or Red with an appended reason identifer when outside suggested criteria. Refer to the footnotes section at the end of this report for failure reasons.

Metals in Water (Dissolved) by ICPOES

Method: ME-(AU)-[ENV]AN320/AN321

								*
QC Sample	Sample Number	Parameter	Units	LOR	Result	Original	Spike	Recovery%
CE124214.001	LB041532.004	Calcium, Ca	mg/L	0.1	59	8.0	50	103
		Magnesium, Mg	mg/L	0.1	52	3.2	50	98
		Potassium, K	mg/L	0.1	54	0.8	50	106
		Sodium, Na	mg/L	0.5	63	17	50	93

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MATRIX SPIKE DUPLICATES

CE124214 R0

Matrix spike duplicates are calculated as Relative Percent Difference (RPD) using the formula: RPD = | OriginalResult - ReplicateResult | x 100 / Mean

The original result is the analyte concentration of the matrix spike. The Duplicate result is the analyte concentration of the matrix spike duplicate.

The RPD is evaluated against the Maximum Allowable Difference (MAD) criteria and can be graphically represented by a curve calculated from the Statistical Detection Limit (SDL) and Limiting Repeatability (LR) using the formula: MAD = 100 x SDL / Mean + LR

Where the Maximum Allowable Difference evaluates to a number larger than 200 it is displayed as 200.

RPD is shown in Green when within suggested criteria or Red with an appended reason identifer when outside suggested criteria. Refer to the footnotes section at the end of this report for failure reasons.

No matrix spike duplicates were required for this job.

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FOOTNOTES

Samples analysed as received.

Solid samples expressed on a dry weight basis.

QC criteria are subject to internal review according to the SGS QA/QC plan and may be provided on request or alternatively can be found here: http://www.sgs.com.au/~/media/Local/Australia/Documents/Technical Documents/MP-AU-ENV-QU-022 QA QC Plan.pdf

- * NATA accreditation does not cover the performance of this service.
- Sample not analysed for this analyte.

IS Insufficient sample for analysis.

LNR Sample listed, but not received.

LOR Limit of reporting.

QFH QC result is above the upper tolerance.
QFL QC result is below the lower tolerance.

- ① At least 2 of 3 surrogates are within acceptance criteria.
- 2 RPD failed acceptance criteria due to sample heterogeneity.
- 3 Results less than 5 times LOR preclude acceptance criteria for RPD.
- Recovery failed acceptance criteria due to matrix interference.
- ® Recovery failed acceptance criteria due to the presence of significant concentration of analyte (i.e. the concentration of analyte exceeds the spike level).
- © LOR was raised due to sample matrix interference.
- ① LOR was raised due to dilution of significantly high concentration of analyte in sample.
- ® Reanalysis of sample in duplicate confirmed sample heterogeneity and inconsistency of results.
- Low surrogate recovery due to the sample emulsifying during extraction.
- © Legionella Test Result <10 cfu/mL</p>

Control Strategy (1)

Maintain Monthly Program or at least 3-monthly monitoring. Maintain water treatment program

(f) Legionella Test Result <1000 cfu/mL

Control Strategy (2)

Investigate problem. Review water treatment program. Take necessary remedial action (including immediate online disinfection)

and undertake control strategy 3.

© Control Strategy (3)

Retest water within 3 to 7 days of plant operation. If not detected, continue to retest water every 3 to 7 days until 2 consecutive samples

return readings of 'not detected', then repeat control strategy (1).

If detected at <100 cfu/mL, repeat control strategy (2).

If detected at >100 cfu/mL, investigate the problem and review water treatment program,

and immediately carry out online decontamination.

If detected at >1000 cfu/mL, undertake control strategy (4).

Legionella Test Result >1000 cfu/mL

Control Strategy (4)

Investigate problem. Review water treatment program.

Take necessary remedial action (including immediate online decontamination)

and undertake control strategy (5).

(5) Control Strategy

Retest water within 3 to 7 days of plant operation. If not detected, continue to retest water every 3 to 7 days until 2 consecutive samples

return readings of 'not detected', then repeat control strategy (1).

If detected at <100 cfu/mL, repeat control strategy (1)

If detected at >100 and <1000 cfu/mL, investigate the problem and review water treatment program, immediately carry out online decontamination.

and repeat control strategy (5).

If detected at >1000 cfu/mL, investigate and review the water treatment program,

immediately carry out system decontamination

and repeat control strategy (5).

(5) HPC - Test Result < 100 000 cfu/mL

Control Strategy (1)

Maintain Monthly Program. Maintain water treatment program.

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Samples analysed as received.

Solid samples expressed on a dry weight basis.

QC criteria are subject to internal review according to the SGS QA/QC plan and may be provided on request or alternatively can be found here: http://www.sgs.com.au/~/media/Local/Australia/Documents/Technical Documents/MP-AU-ENV-QU-022 QA QC Plan.pdf

(6) HPC Test Result > 100 000 cfu/mL <5 000 000 cfu/mL

Control Strategy (2)

Investigate problem. Review water treatment program. Take necessary remedial action (including immediate online disinfection)

and undertake control strategy 3.

① Control Strategy (3)

Retest water within 3 to 7 days of plant operation. If the test result is <100 000 cfu/mL, repeat control strategy (1) If the test result is >100 000 cfu/mL but <5 000 000 cfu/mL, undertake control strategy (2).

If the test result is >5 000 000 cfu/mL, undertake control strategy (4).

® HPC Test Result >5 000 000 cfu/mL

Control Strategy (4)

Investigate problem. Review water treatment program. Take necessary remedial action (including immediate online disinfection)

and undertake control strategy (5).

(5) Control Strategy

Retest water within 3 to 7 days of plant operation. If the test result is <100 000 cfu/mL, repeat control strategy (1)

If the test result is >100 000 cfu/mL but <5 000 000 cfu/mL, undertake control strategy (4).

If the test result is >5 000 000 cfu/mL, investigate the problem.

review the water treatment program, and carry out immediate online decontamination.

Enterococci - Median result should not exceed 230 cfu/100mL

(maximum number in any one sample: 450-700 cfu/100mL)

Sourced from NHMRC (National Health and Medical Research Council)

& NWQMS (National Water Quality Management Strategy)

-Australian Guidelines for Recreational Use of Water. Version Oct 2000.

† Refer to Analytical Report comments for further information.

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LABORATORY DETAILS

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2

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SGS Reference CE126723 R0 06 Apr 2017 Date Received

18 Apr 2017

COMMENTS

Accredited for compliance with ISO/IEC 17025-Testing. NATA accredited laboratory 2562(3146)

For determination of soluble metals, filtered sample was not received so samples were laboratory filtered on receipt. This may give soluble metals results that do not represent the concentrations present at the time of sampling.

SIGNATORIES

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Anion-Cation Balance

ANALYTICAL REPORT

CE126723 R0

	Sample Number Sample Matrix Sample Date Sample Name		CE126723.001 Water 06 Apr 2017 BH02	CE126723.002 Water 06 Apr 2017 BH03
Parameter	Units	LOR		
pH in water Method: AN101/MA1490(Melb) Tested: 6/4/2017	•			
pH**	pH Units	0.1	7.6	8.0
Conductivity and TDS by Calculation - Water Method: AN106/l	MA1489(Melb	Tested:	6/4/2017	
Conductivity @ 25 C	μS/cm	5	2900	8600
Alkalinity Method: AN135/MA1127(Melb) Tested: 6/4/2017				
Total Alkalinity as CaCO3	mg/L	5	430	670
Bicarbonate Alkalinity as CaCO3	mg/L	5	430	670
Carbonate Alkalinity as CaCO3	mg/L	5	<5	<5
Hydroxide Alkalinity as CaCO3	mg/L	5	<5	<5
Chloride by Discrete Analyser in Water Method: AN274 Test	ted: 10/4/2017			
Chloride, Cl	mg/L	1	660	2400
Metals in Water (Dissolved) by ICPOES Method: AN320/AN32	1 Tested: 1	0/4/2017		
Calcium, Ca	mg/L	0.1	20	68
Magnesium, Mg	mg/L	0.1	39	190
Potassium, K	mg/L	0.1	40	99
Sodium, Na	mg/L	0.5	570	1500
Sulphur as Sulphate, SO4	mg/L	0.5	7.3	3.5
Total Hardness by Calculation	mg CaCO3/L	1	210	940
Calculation of Anion-Cation Balance (SAR Calc) Method: AN1	21 Tested:	18/4/2017		
Sum of Cation Milliequivalents*	meq/L	-	30.0	86.0
Sum of Anion Milliequivalents*	meq/L	-	27.3	81.8

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4.6

2.5



QC SUMMARY

MB blank results are compared to the Limit of Reporting

LCS and MS spike recoveries are measured as the percentage of analyte recovered from the sample compared the the amount of analyte spiked into the sample.

DUP and MSD relative percent differences are measured against their original counterpart samples according to the formula: the absolute difference of the two results divided by the average of the two results as a percentage. Where the DUP RPD is 'NA', the results are less than the LOR and thus the RPD is not applicable.

Alkalinity Method: ME-(AU)-[ENV]AN135/MA1127(Melb)

Parameter	QC Reference	Units	LOR	МВ	DUP %RPD	LCS %Recovery
Total Alkalinity as CaCO3	LB045094	mg/L	5	<5	2%	98 - 112%
Bicarbonate Alkalinity as CaCO3	LB045094	mg/L	5	<5		
Carbonate Alkalinity as CaCO3	LB045094	mg/L	5	<5		
Hydroxide Alkalinity as CaCO3	LB045094	mg/L	5	<5		

Chloride by Discrete Analyser in Water Method: ME-(AU)-[ENV]AN274

	Parameter	QC	Units	LOR	МВ	DUP %RPD	LCS
ı		Reference					%Recovery
	Chloride, Cl	LB045153	mg/L	1	<1	0 - 1%	107 - 108%

Conductivity and TDS by Calculation - Water Method: ME-(AU)-[ENV]AN106/MA1489(Melb)

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
Conductivity @ 25 C	LB045094	μS/cm	5	<5	0 - 1%	98 - 99%

Metals in Water (Dissolved) by ICPOES Method: ME-(AU)-[ENV]AN320/AN321

Parameter	QC Reference	Units	LOR	MB	DUP %RPD	LCS %Recovery	MS %Recovery
Calcium, Ca	LB045144	mg/L	0.1	<0.1	0%	102%	103%
Magnesium, Mg	LB045144	mg/L	0.1	<0.1	0%	103%	97%
Potassium, K	LB045144	mg/L	0.1	<0.1	0%	97%	117%
Sodium, Na	LB045144	mg/L	0.5	<0.5	0%	97%	
Sulphur as Sulphate, SO4	LB045144	mg/L	0.5	<0.5	0%	NA	
Total Hardness by Calculation	LB045144	mg CaCO3/L	1	<1			

pH in water Method: ME-(AU)-[ENV]AN101/MA1490(Melb)

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
pH**	LB045094	pH Units	0.1	5.5 - 5.7	0 - 3%	NA

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METHOD SUMMARY

METHOD	METHODOLOGY SUMMARY
AN101/MA1490(Melb)	pH in Soil Sludge Sediment and Water: pH is measured electrometrically using a combination electrode (glass plus reference electrode) and is calibrated against 3 buffers purchased commercially. For soils, an extract with water is made at a ratio of 1:5 and the pH determined and reported on the extract. Reference APHA 4500-H+.
AN106/MA1489(Melb)	Conductivity and TDS by Calculation: Conductivity is measured by meter with temperature compensation and is calibrated against a standard solution of potassium chloride. Conductivity is generally reported as µmhos/cm or µS/cm @ 25°C. For soils, an extract with water is made at a ratio of 1:5 and the EC determined and reported on the extract, or calculated back to the as-received sample. Total Dissolved Salts can be estimated from conductivity using a conversion factor, which for natural waters, is in the range 0.55 to 0.75. SGS use 0.6. Reference APHA 2510 B.
AN121	This method is used to calculation the balance of major Anions and Cations in water samples and converts major ion concentration to milliequivalents and then summed. Anions sum and Cation sum is calculated as a difference and expressed as a percentage.
AN135/MA1127(Melb)	Alkalinity (and forms of) by Titration: The sample is titrated with standard acid to pH 8.3 (P titre) and pH 4.5 (T titre) and permanent and/or total alkalinity calculated. The results are expressed as equivalents of calcium carbonate or recalculated as bicarbonate, carbonate and hydroxide. Reference APHA 2320. Internal Reference AN135
AN274	Chloride by Aquakem DA: Chloride reacts with mercuric thiocyanate forming a mercuric chloride complex. In the presence of ferric iron, highly coloured ferric thiocyanate is formed which is proportional to the chloride concentration. Reference APHA 4500CI-
AN320/AN321	Metals by ICP-OES: Samples are preserved with 10% nitric acid for a wide range of metals and some non-metals. This solution is measured by Inductively Coupled Plasma. Solutions are aspirated into an argon plasma at 8000-10000K and emit characteristic energy or light as a result of electron transitions through unique energy levels. The emitted light is focused onto a diffraction grating where it is separated into components.
AN320/AN321	Photomultipliers or CCDs are used to measure the light intensity at specific wavelengths. This intensity is directly proportional to concentration. Corrections are required to compensate for spectral overlap between elements. Reference APHA 3120 B.
Calculation	Free and Total Carbon Dioxide may be calculated using alkalinity forms only when the samples TDS is <500mg/L. If TDS is >500mg/L free or total carbon dioxide cannot be reported. APHA4500CO2 D.

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FOOTNOTES

IS Insufficient sample for analysis.

LNR Sample listed, but not received.

* NATA accreditation does not cover the

performance of this service.

** Indicative data, theoretical holding time exceeded.

LOR Limit of Reporting

↑↓ Raised or Lowered Limit of Reporting
QFH QC result is above the upper tolerance
QFL QC result is below the lower tolerance
The sample was not analysed for this analyte

NVL Not Validated

Samples analysed as received.
Solid samples expressed on a dry weight basis.

Where "Total" analyte groups are reported (for example, Total PAHs, Total OC Pesticides) the total will be calculated as the sum of the individual analytes, with those analytes that are reported as <LOR being assumed to be zero. The summed (Total) limit of reporting is calcuated by summing the individual analyte LORs and dividing by two. For example, where 16 individual analytes are being summed and each has an LOR of 0.1 mg/kg, the "Totals" LOR will be 1.6 / 2 (0.8 mg/kg). Where only 2 analytes are being summed, the "Total" LOR will be the sum of those two LORs.

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Results reported for samples tested under test methods with codes starting with ARS-SOP, radionuclide or gross radioactivity concentrations are expressed in becquerel (Bq) per unit of mass or volume or per wipe as stated on the report. Becquerel is the SI unit for activity and equals one nuclear transformation per second.

Note that in terms of units of radioactivity:

- a. 1 Bq is equivalent to 27 pCi
- b. 37 MBq is equivalent to 1 mCi

For results reported for samples tested under test methods with codes starting with ARS-SOP, less than (<) values indicate the detection limit for each radionuclide or parameter for the measurement system used. The respective detection limits have been calculated in accordance with ISO 11929.

The QC criteria are subject to internal review according to the SGS QAQC plan and may be provided on request or alternatively can be found here: http://www.sgs.com.au/~/media/Local/Australia/Documents/Technical%20Documents/MP-AU-ENV-QU-022%20QA%20QC%20Plan.pdf

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18-April-2017 Page 5 of 5







CLIENT DETAILS -

LABORATORY DETAILS

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Email

dasimpson@golder.com.au

Project

1546223 Northern Sands

Order Number

(Not specified)

Samples

Jon Dicker Manager

Laboratory

Telephone

Date Reported

SGS Cairns Environmental

Address

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Portsmith QLD 4870

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Facsimile +61 07 4035 5122

AU.Environmental.Cairns@sgs.com Fmail

CE126058 R0

SGS Reference Date Received

03 Mar 2017

15 Mar 2017

COMMENTS

Accredited for compliance with ISO/IEC 17025. NATA accredited laboratory 2562(3146)

For determination of soluble metals, filtered sample was not received so samples were laboratory filtered on receipt. This may give soluble metals results that do not represent the concentrations present at the time of sampling.

SIGNATORIES

Alyson Bergamo

Senior Laboratory Technician

Anthony Nilsson **Operations Manager** Jon Dicker

Manager Northern QLD

Maristela Ganzan Metals Team Leader

SGS Australia Pty Ltd ABN 44 000 964 278

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Anion-Cation Balance

ANALYTICAL REPORT

CE126058 R0

	Sample Number Sample Matrix Sample Date Sample Name		CE126058.001 Water 03 Mar 2017 BH01	CE126058.002 Water 03 Mar 2017 BH04
Parameter	Units	LOR		
pH in water Method: AN101 Tested: 3/3/2017				
pH**	pH Units	0.1	6.9	6.2
Conductivity and TDS by Calculation - Water Method: AN106	Tested: 3/3	/2017		
Conductivity @ 25 C	μS/cm	5	250	140
Alkalinity Method: AN135 Tested: 3/3/2017				
Total Alkalinity as CaCO3	mg/L	5	41	15
Bicarbonate Alkalinity as CaCO3	mg/L	5	41	15
Carbonate Alkalinity as CaCO3	mg/L	5	<5	< 5
Hydroxide Alkalinity as CaCO3	mg/L	5	<5	<5
Chloride by Discrete Analyser in Water Method: AN274 Test	ted: 6/3/2017			
Chloride, Cl	mg/L	1	39	17
Metals in Water (Dissolved) by ICPOES Method: AN320/AN32	1 Tested: 9	0/3/2017		
Calcium, Ca	mg/L	0.1	5.7	4.1
Magnesium, Mg	mg/L	0.1	8.2	2.8
Potassium, K	mg/L	0.1	1.8	0.5
Sodium, Na	mg/L	0.5	28	19
Sulphur as Sulphate, SO4	mg/L	0.5	22	24
Total Hardness by Calculation	mg CaCO3/L	1	48	22
Calculation of Anion-Cation Balance (SAR Calc) Method: AN1	21 Tested:	15/3/2017		
Sum of Cation Milliequivalents*	meq/L	-	2.23	1.26
Sum of Anion Milliequivalents*	meq/L	-	2.36	1.28

15-March-2017 Page 2 of 5

-100

-2.9

-0.8



QC SUMMARY

MB blank results are compared to the Limit of Reporting

LCS and MS spike recoveries are measured as the percentage of analyte recovered from the sample compared the the amount of analyte spiked into the sample.

DUP and MSD relative percent differences are measured against their original counterpart samples according to the formula: the absolute difference of the two results divided by the average of the two results as a percentage. Where the DUP RPD is 'NA', the results are less than the LOR and thus the RPD is not applicable.

Alkalinity Method: ME-(AU)-[ENV]AN135

Parameter	QC Reference	Units	LOR	MB	DUP %RPD	LCS %Recovery
Total Alkalinity as CaCO3	LB044113	mg/L	5	<5	0%	112%
Bicarbonate Alkalinity as CaCO3	LB044113	mg/L	5	<5		
Carbonate Alkalinity as CaCO3	LB044113	mg/L	5	<5		
Hydroxide Alkalinity as CaCO3	LB044113	mg/L	5	<5		

Chloride by Discrete Analyser in Water Method: ME-(AU)-[ENV]AN274

ı	Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
ı		Reference					%Recovery
ı	Chloride, Cl	LB044132	mg/L	1	<1	0 - 1%	109%

Conductivity and TDS by Calculation - Water Method: ME-(AU)-[ENV]AN106

	Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
ı		Reference					%Recovery
	Conductivity @ 25 C	LB044113	μS/cm	5	<5	1%	101%

Metals in Water (Dissolved) by ICPOES Method: ME-(AU)-[ENV]AN320/AN321

Parameter	QC Reference	Units	LOR	MB	DUP %RPD	LCS %Recovery	MS %Recovery
Calcium, Ca	LB044236	mg/L	0.1	<0.1	1%	110%	99%
Magnesium, Mg	LB044236	mg/L	0.1	<0.1	1 - 2%	107%	104%
Potassium, K	LB044236	mg/L	0.1	<0.1	1 - 3%	105%	111%
Sodium, Na	LB044236	mg/L	0.5	<0.5	0 - 1%	97%	96%
Sulphur as Sulphate, SO4	LB044236	mg/L	0.5	<0.5	1%	NA	
Total Hardness by Calculation	LB044236	mg CaCO3/L	1	<1			

pH in water Method: ME-(AU)-[ENV]AN101

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
pH**	LB044113	pH Units	0.1	5.8	1%	NA

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SGS

METHOD SUMMARY

METHOD	METHODOLOGY CHAMADY
WILTHOU -	METHODOLOGY SUMMARY
AN101	pH in Soil Sludge Sediment and Water: pH is measured electrometrically using a combination electrode (glass plus reference electrode) and is calibrated against 3 buffers purchased commercially. For soils, an extract with water is made at a ratio of 1:5 and the pH determined and reported on the extract. Reference APHA 4500-H+.
AN106	Conductivity and TDS by Calculation: Conductivity is measured by meter with temperature compensation and is calibrated against a standard solution of potassium chloride. Conductivity is generally reported as μ mhos/cm or μ S/cm @ 25°C. For soils, an extract with water is made at a ratio of 1:5 and the EC determined and reported on the extract, or calculated back to the as-received sample. Total Dissolved Salts can be estimated from conductivity using a conversion factor, which for natural waters, is in the range 0.55 to 0.75. SGS use 0.6. Reference APHA 2510 B.
AN121	This method is used to calculation the balance of major Anions and Cations in water samples and converts major ion concentration to milliequivalents and then summed. Anions sum and Cation sum is calculated as a difference and expressed as a percentage.
AN135	Alkalinity (and forms of) by Titration: The sample is titrated with standard acid to pH 8.3 (P titre) and pH 4.5 (T titre) and permanent and/or total alkalinity calculated. The results are expressed as equivalents of calcium carbonate or recalculated as bicarbonate, carbonate and hydroxide. Reference APHA 2320. Internal Reference AN135
AN274	Chloride by Aquakem DA: Chloride reacts with mercuric thiocyanate forming a mercuric chloride complex. In the presence of ferric iron, highly coloured ferric thiocyanate is formed which is proportional to the chloride concentration. Reference APHA 4500CI-
AN320/AN321	Metals by ICP-OES: Samples are preserved with 10% nitric acid for a wide range of metals and some non-metals. This solution is measured by Inductively Coupled Plasma. Solutions are aspirated into an argon plasma at 8000-10000K and emit characteristic energy or light as a result of electron transitions through unique energy levels. The emitted light is focused onto a diffraction grating where it is separated into components.
AN320/AN321	Photomultipliers or CCDs are used to measure the light intensity at specific wavelengths. This intensity is directly proportional to concentration. Corrections are required to compensate for spectral overlap between elements. Reference APHA 3120 B.
Calculation	Free and Total Carbon Dioxide may be calculated using alkalinity forms only when the samples TDS is <500mg/L. If TDS is >500mg/L free or total carbon dioxide cannot be reported. APHA4500CO2 D.

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FOOTNOTES _

IS Insufficient sample for analysis.

LNR Sample listed, but not received.

* NATA accreditation does not cover the performance of this service.

** Indicative data, theoretical holding time exceeded.

LOR Limit of Reporting

↑↓ Raised or Lowered Limit of Reporting QFH QC result is above the upper tolerance QFL QC result is below the lower tolerance

- The sample was not analysed for this analyte

NVL Not Validated

Samples analysed as received.

Solid samples expressed on a dry weight basis.

Where "Total" analyte groups are reported (for example, Total PAHs, Total OC Pesticides) the total will be calculated as the sum of the individual analytes, with those analytes that are reported as <LOR being assumed to be zero. The summed (Total) limit of reporting is calcuated by summing the individual analyte LORs and dividing by two. For example, where 16 individual analytes are being summed and each has an LOR of 0.1 mg/kg, the "Totals" LOR will be 1.6 / 2 (0.8 mg/kg). Where only 2 analytes are being summed, the "Total" LOR will be the sum of those two LORs.

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15-March-2017 Page 5 of 5

GROUNDWATER SAMPLING RECORD FORM



FOR THR	EE WELL V	OLUME METHO	DD				55 A	ssociates
PROJECT	INFORMATIO	NC						
Project Number: 1546 223 Client: Site Location: Walkern Sand Pro					Tech	Date Sampled By		(3)16
GROUNDW	ATER WELL				0		Programme and the second	
Information	4.5	13:25	2 -> .	087			DODE ID	
Surveyed refe		13.0	3 5.0	DOT	1, 2×15	-6C Note? o(i)	BORE ID	1300
Depth of well			61	m	6x3-	Note -1	(0,0	BHO
	vell (inc filter pa	ack) (mm)	0		OKJ	10000000	20	
Height of filter						Standard refere	ence point is top	of PVC standpipe
Information I	recorded on s	ite			•	mbRP - metres	below top of re	ference point
	tandpipe (mm)	1	501	Com				•
Standpipe stic	ck up (m)	-						
Time	/ I DD: 00	0-	1-2	-3		Interface probe		YES / NO
Depth to wate		MBIOC	/			Depth to produc		
Total depth of		The light of	60	Jr.		Depth to water		-
	er in standpipe	ase of well (m)	9	180		Thickness of pro	oduci (m)	
Well volume (V/	3	101				
		, well volume wihtin 5	0mm standpip	e is 2 litres/m				
Estimation of St	tanding Well Vol	lumes for 50 mm dian	35///		water column wit	thin filter pack		
Well diameter (standpipe plus fi	filter pack)	100mm	115mm	120mm	125mm	150mm	200mm
ures per lineal	meter of water i	in bore	3.7	4.5	4.8	5.1	6.7	10.8
PURGING R	ECORD						Arte X	
Time	Volume Purged (L)	Conductivity (mS or (µS)	Temp (°C)	pН	Redox Potential	Dissolved Oxygen (mg/L)		earance eidity, odour, etc)
1.31	1	27.0	26 -16	710	(mV)	2	AND O	1001110
1032-5	015	350	00:04	8.4	256	20	Manny	uch lours
12.5	200	217	26.01	9,92	293	25		-//
135	0.05	257	9-03	221	297	0.5		11
1.36	130	298	20.91	7.27	202	01		//
1037	240	297	1602 BA	706-8	216	0.6		1/
2-40	. '	373	12.85	7.18	263	-/		r
lotal vo	ol. purged (L)		No. bor	re vol. purged		Purging	Time (minutes)	
	at end of purg	ing (mbRP)		_			×	Somethin
urging Method	d:	luaro	ma po	Smo				goving in
AMPLING F	RECORD			9				whomby
VEN 112 NO. 12 N	an and an annual control of the second	. 4	and the second s	THE PERSON NAMED IN COLUMN TWO IS NOT THE PERSON NAMED IN COLUMN TWO IS NAMED IN COLUMN TWI	Lander Colored (1995)		P Uni	A della
ampling meth ime sampled:		a pera pi	mp		Samples 64a	Sample ID red for metals?:	Yes /(No)	•
ample Appear		196	V	3	Samples like	red for metals?.	res / No	
Colour	10	shr may			Turbidity	Low Media	um / High	
Odour_		J. No.		Hydroc	arbon sheen?	100	,	
ample Contai	iner and Prese	ervation:	n A			ken? / Dup ID.:		
	1	grean t	x +160	4				
		0	J.1 ~	194000 VIII				
offer phosphilare variety	Same in the same		100 m	TOTAL SAFETON		FIGT OF THE WITTER STATES		
BSERVATIO	ONS		jer Lend L					1 1
Weather	Conditions:	Temperature:	30			WC1. 139	3.08.	Defene de
	10	Precipitation :	1	10		Jan	2.084	alla
Notes:	1:1	4 : Ja	Ne ou	r daka	19904	es co	500/	- yin
_	7	00 ;	PUT 1	Deck.	davo	laggers		
	-	_	1	The state of the s				

GAP Form No. 37 RL 0

GROUNDWATER SAMPLING RECORD FORM FOR THREE WELL VOLUME METHOD



PRUJELI	INFORMATI	ON			和"生"为"利"。		TO A TO LET BE	ssociates
a comment of the state of the s	roject Numbe	er: 1546 (223			Date Sampled By	-661	121/16
	Site Location	n: Nul	hern Sas	ne more	V	,	10	
GROUNDY	VATER WEL			4 0				
Information	from file					21 (BORE ID	I D Us S
Surveyed ref	ference point		1	14	7, - (BOKE ID	1 RAOS
Depth of well			61	M		Note:		
	well (inc filter p	ack) (mm)		0-1	-	Note.		
leight of filte				- 1		Standard refere	nce point is ton	of PVC standpipe
	recorded on s	site			7	mbRP - metres		
iameter of s	standpipe (mm)	500	mm	7	more - metres	below top of fer	erence point
tandpipe sti					49			
ime			7.71)	1	Interface probe	used?	YES / NO
epth to wate	er (mbRP)	maroc	765	2.3		Depth to produc		120 / 110
otal depth of	f well (mbRP)	mBack	hom			Depth to water (
nickness of	sediment on ba	ase of well (m)	0.4		47	Thickness of pro		
	er in standpipe					L. Fridances of pro	7.2301 (111)	
ell volume ((L)	4			7.3			
ater column a	above filter pack	, well volume wihtin s	50mm standnir	e is 2 litres/m	,			
		lumes for 50 mm dia			water column w	ithin filter nack		
ell diameter (standpipe plus f	ilter pack)	100mm	115mm	120mm	125mm	150mm	200mm
res per lineal	meter of water	in bore	3.7	4.5	4.8	5.1	6.7	10.8
JRGING R	ECORD							
ne	Volume	Conductivity (mS	Temp	рН	Redox	Dissolved	Anne	earance
	Purged (L)	OP ITS)	(°C)	P.1.	Potential	Oxygen (mg/L)		dity, odour, etc)
2.211	95	Xooo	Valet.	0 2	(mV)		Pod	11 0 0 0 0
200	010	7000	2000	7.0	77	0.8	Brown	Highrah
549	280	800	7639	3-86	39	0.8		//
25	2	7800	16.31	400	30	0.6	/	
139	2	3000	86.08	7:65	200	-0		
1.5	200	9000	19 XC	7560	40	-0.7	11	- ACTA
901	934 33 8000 76.0		6.91	7.00	Sta	-0./	10101211	
9.33	33	Dago		(20 ())	74	-02-1		C
70 X	37	8000	76.054	to und pure d		5 .	/	
Total vo	ol. purged (L)	8000	(1) (-) (4)	re vol. purged		Purging	Time (minutes)	
pth to water	ol. purged (L)	ing (mbRP)	No. bor	610	العيار ال	Purging	Time (minutes)	
pth to water	ol. purged (L)	ing (mbRP)	No. bor	e vol. purged		Purging	Time (minutes)	
pth to water	ol. purged (L) at end of purg	ing (mbRP)	No. bor	610),	Purging	Time (minutes)	
pth to water rging Method	at end of purg	ing (mbRP)	No. boi	610	Charles mil		Time (minutes)	
pth to water rging Method MPLING R	at end of purg d: RECORD	ing (mbRP)	No. boi	610)	Sample ID.	BH02	
pth to water rging Method MPLING R mpling method re sampled:	at end of purg d: RECORD	ing (mbRP)	No. boi	610	Samples filte		BH 2 Yes /(No)	
pth to water rging Method MPLING R mpling method re sampled: mple Appear	at end of purg d: RECORD od:	wa Vene	No. boi	610		Sample ID red for metals?:	BH o Z	
pth to water rging Method MPLING R mpling method to sampled: mple Appear Colour	at end of purg d: RECORD od:	ing (mbRP) Lua Noua 1.40	No. bor	e pump	Turbidity	Sample ID.	BH o Z	
pth to water rging Method MPLING R mpling method e sampled: nple Appear Colour Odour	at end of purg d: RECORD od:	Cua Vena	No. bor	6 76 La guirm	Turbidity arbon sheen?	Sample ID. red for metals?:	BH o Z	
pth to water rging Method MPLING R mpling method the sampled: nple Appear Colour Odour	at end of purg d: RECORD od:	Cua Vena	No. bor	6 76 La guirm	Turbidity arbon sheen?	Sample ID red for metals?:	BH o Z	
pth to water rging Method MPLING R mpling method he sampled: mple Appear Colour Odour	at end of purg d: RECORD od:	Cua Vena	No. bor	6 76 La guirm	Turbidity arbon sheen?	Sample ID. red for metals?:	BH o Z	
pth to water rging Method MPLING R mpling method ne sampled: mple Appear Colour Odour	at end of purg d: RECORD od:	Cua Vena	No. bor	6 76 La guirm	Turbidity arbon sheen?	Sample ID. red for metals?:	BH o Z	
ppth to water rging Method MPLING R mpling method ne sampled: mple Appear Colour Odour	at end of purg d: RECORD od:	Cua Vena	No. bor	6 76 La guirm	Turbidity arbon sheen?	Sample ID. red for metals?:	BH o Z	
epth to water irging Method AMPLING R impling method ne sampled: mple Appear Colour Odour mple Contain	at end of purg d: RECORD od: rance:	Cua Vena	No. bor	6 76 La guirm	Turbidity arbon sheen?	Sample ID. red for metals?:	BH o Z	
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pth to water rging Method MPLING R mpling method ne sampled: mple Appear Colour Odour mple Contain	at end of purg d: RECORD od: rance:	Cua Vena	No. bor	6 76 La guirm	Turbidity arbon sheen?	Sample ID. red for metals?:	BH o Z	
pth to water rging Method MPLING R mpling method ne sampled: mple Appear Colour Odour mple Contain	at end of purg d: RECORD od: rance:	ing (mbRP) Lyo Moun Lyo Pown Lyo Lyo Lyo Lyo Lyo Lyo Lyo Ly	No. bor	6 76 La guirm	Turbidity arbon sheen?	Sample ID. red for metals?:	BH o Z	
pth to water rging Method MPLING R mpling method es sampled: nple Appear Colour Odour nple Contain SERVATIO	at end of purg d: RECORD od: rance:	ing (mbRP) County Co	No. bor	6 76 La guirm	Turbidity arbon sheen?	Sample ID. red for metals?:	BH o Z	

GROUNDWATER SAMPLING RECORD FORM FOR THREE WELL VOLUME METHOD



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PROJECT INFORMATION						
Project Number: 154 792	1			Date	. 271	1111/
Client:)	_		Sampled By		4110
	in Sent	proxect			- 36	
GROUNDWATER WELL DATA		4. "Q.		7.7	The second of	***
Consideration of the control of the second of the constant of the control of the		经现场和现在 。		1912年1月1日	DODE ID	1 (2/112)
Information from file Surveyed reference point			1		BORE ID	BHOS
Depth of well (from log)			1	Note:		
Diameter of well (inc filter pack) (mm)			1	. 1010.		
Height of filter pack (m)		9	1	Standard refere	nce point is top	of PVC standpipe
Information recorded on site			_	mbRP - metres	below top of ref	erence point
Diameter of standpipe (mm)]			
Standpipe stick up (m)]			
Time	3.10]	Interface probe	used?	YES / NO
Depth to water (mbRP) MPS CC	7.50			Depth to produc	t (mbRP)	
Total depth of well (mbRP) AMB TOC	11.31	212m		Depth to water (
Thickness of sediment on base of well (m)				Thickness of pro	oduct (m)	
Height of water in standpipe (m)			2 1/6	alde		~
Well volume (L)			Lollar Ja	elitersta	3 nomol	e
Water column above filter pack, well volume wihtin 50			wellbildon			
Estimation of Standing Well Volumes for 50 mm diam Well diameter (standpipe plus filter pack)	100mm	es, for height of	mater column with 120mm	125mm	150mm	200mm
Litres per lineal meter of water in bore	3.7	4.5	4.8	5.1	6.7	10.8
PURGING RECORD						
Time Volume Conductivity mS	Temp	рН	Redox	Dissolved	Appe	earance
Purged (L) or μS)	(°C)		Potential	Oxygen (mg/L)		idity, odour, etc)
	0.00		(mV)			1 1 2
5.14 10 25.46	2594	8.86	165	0.5	Roun	MOSHOO
315 20 755	25.0	4°52	105	//		-
X11 20 255	25.83	8.47	144	067	- 1	
1800 40 85.428	2	800	148	0.0	1	
300 43 65-49	85.03	6,0	13 t	6.3		
97 XX 977	18/04	3-211	13.9	6-2		
Total vol. purged (L)	No ho	re vol. purged	146		Time (minutes)	T T
3.76 60	140. 50	70/23	11	1/	Time (Himates)	L
Depth to water at end of purging (mbRP)						
Purging Method:	ena (pung				
SAMPLING RECORD						
Sampling method:	a Q1	Sw		Sample ID.	R	103
Time sampled: 2, 26	4	The state of the s	Samples filte	red for metals?:	Yes / No	
Sample Appearance:						
Colour pange shown			Turbidity	Low / Medi	um / High	
Odour			carbon sheen?	NO		
Sample Container and Preservation:	1	100 Dup	licate sample ta	aken? / Dup ID.:	ND	
Lons	pan r	atte				
		N N				
				Nation Transfer	The state of the s	
DBSERVATIONS	.Hr	0 -	المناه العاملات	A		
Weather Conditions: Temperature:		135				
Precipitation :		, No			2 -	1
Notes:	cul	av 3	10,	1966	X 2.	(
1			1 27	2 051	suche	
		(wt: 2.4	8 por	xeel	



GROUNDWATER SAMPLING RECORD FORM FIELD PARAMETER MEASUREMENTS

Depth to water (m TOC)

Thickness of product (mm)

Yes) No

PROJECT INFORMATION

Client: FCC Project: Northern Sends Location: NS GROUNDWATER BORE DATA	Project No: 154622 Date of Sampling: 25.3.16 Sampled By: 58
Diameter of Column (mm)	BORE
Diameter of Bore (mm) Standing Water Level (m TOC) 50	Interface probe used?
Total Depth of Bore (m TOC) Depth of Water in Column (m)	Depth to product (m TOC)

PURGING RECORD

eli Diametei (Casing pius nite

Litres per lineal meter of water in well

Standpipe stick up (m)

Volume Purged (L)	Dissolved Oxygen (mg/L)	Temperature (C)	TDS (ppm / ppt)	рН	Conductivity	Redox Potential	Turbidity (NTU)	Other
10	0.9	22.37	0.15	9.23	0.30	99.	600	-10.0
15	0.4	22.25	0.18	6.52	0.30	142	196.	-10-
30	0.2	22.25	0.15	6.49	0.30	170	175	10.9
	-				-			
Tatal walva	no munua d (1.)							
	ne purged (L)	Supments for pH you		umes purged	3	Purging	Time (minutes)	9

(Prior to sampling consecutive measurements for pH vaule should be within 0.1 pH units, for conductivity, salinity and dissolved oxygen should be within 10% and temperature should be within 0.5 °C.)

	SAMPLING RECORD							
	Samples Taken	~				Container:	Preservation:	
	Duplicate sample taken	YES (NO				Vial 🔲	H ₂ SO ₄ / other:	
	Samples filtered?	YES / W	for Metals / ot	her:		500ml Glass	none / other;	
	Filter Method:	0.45 mm filter 8	& syringe / other	:		500 ml Plastic	none / other.	
	Water Quality Meter type:					1 l Plastic	H ₂ SO ₄ / other:	
	Water Dipper type:					200 ml Plastic	NaOH-Fother: ACI	'w
	Pumping Method:	Submersible Pu	mp / Disposable	Bailer / ot	her:	200 mL Glass	HNO ₃ / other:	
-	OBSERVATIONS				000			
	Samples:	Colour:	CLEAR	~ 6	Turbidity	Low / Medium / H	ligh	
		Odour:		Hydro	carbon sheen?	Yes / No		
	Weather Conditions:	Sampling Day	Rain		Temperature	31		
		Previous Week	Rain		Temperature			
	Notes:							
	ene							
R	eference Information: Estimation of S	landing Well Volumes f	or 50mm diameter We	dle				

5.1

150mm

Sampling procedures are based on AS/NZS 5667.11:1998 Water Quality - Sampling Part 11: Guidance on sampling of groundwaters

Note: Typical

Eziprobe well diameter =75mm

200mm

2.7



GROUNDWATER SAMPLING RECORD FORM FIELD PARAMETER MEASUREMENTS

PROJECT INFORMATION Project No: 1546223 Client: MORTHERY SAMO Project: Date of Sampling: Location: Sampled By: **GROUNDWATER BORE DATA** Diameter of Column (mm) BORE ID Diameter of Bore (mm) Standing Water Level (m TOC) Interface probe used? Yes / No Total Depth of Bore (m TOC) Depth to product (m TOC) Depth of Water in Column (m) Depth to water (m TOC) Standpipe stick up (m) Thickness of product (mm) **PURGING RECORD** Volume Dissolved Temperature (C) TDS Conductivity Redox Turbidity (NTU) Purged (L) ||Oxygen (mg/L) (ppm / ppt) (US, mS) **Potential** 0.0 10.54 23.03 HEAVY 10.08 Total volume purged (L) No. bore volumes purged Purging Time (minutes) (Prior to sampling consecutive measuements for pH vaule should be within 0.1 pH units, for conductivity, salinity and dissolved oxygen should be within 10% and temperature should be within 0.5 °C.) SAMPLING RECORD Samples Taken? Container: Preservation: Duplicate sample taken? Vial 🔲 H₂SO₄ / other: Samples filtered? YES / NO) for Metals / other: 500ml Glass none / other: Filter Method: 0.45 mm filter & syringe / other: 500 ml Plastic 🗖 none other: Water Quality Meter type: YEOKAL H₂SO₄ / other: 1 | Plastic NaOH / other: HNO3 / other: Nonthaco Water Dipper type: 200 ml Plastic Pumping Method: Submersible Pump / Disposable Bailer / other: 200 mL Glass **OBSERVATIONS**

Reference Information: Estimation of Standing Well Volumes for 50mm diameter Wells

Previous Week

Samples:

Weather Conditions: Sampling Day

Notes:

Note: Typical 75mm Litres per lineal meter of water in well Eziprobe well diameter =75mm

Rain

Sampling procedures are based on AS/NZS 5667.11:1998 Water Quality - Sampling Part 11: Guidance on sampling of groundwaters

Colour: GREY



Turbidity

Hydrocarbon sheen?

Temperature

Temperature

Low / Medium ,



GROUNDWATER SAMPLING RECORD FORM FIELD PARAMETER MEASUREMENTS

Project:	NORT MORT TER BOR Jumn (mm) Te (mm) Level (m TO	E DATA	125 50 2:41 11-28	HYDRO Da	Project N te of Samplin Sampled B	g: 29	BORE ID be used?	(Yes)/No	
Standpipe stick	up (m)		0.65			Thickness of	product (mm)		
PURGING REC									
	issolved gen (mg/L)	Temperature (C)	TDS (ppm / ppt)	pΗ	Conductivity	Redox Potential	Turbidity (NTU)	Other	
5	0.0	22.32	16-11	8.52	26.36	65	600	RED/BROWN	الر
25	0-0	22.28	16-19	9.58	26.45	7	600	CLÉARING	
50 0	0.0	25.28	16.23	10.21	26.54	-3	600	11 0	
									4
									4
Total values as	unned (I.)								
(Prior to sampling cons	secutive mea	suements for pH vac	ule should be with	lumes purged hin 0.1 pH units,	for conductivity, s	Purging Salinity and dissolu	g Time (minutes) ved oxygen		
should be within 10% a		ure should be within	0.5 °C.)					2.4	۵۲
Samples Duplicate sample Samples fi Filter M	taken? Itered?	YES / NO YES / NO YES / NO for the second of	or Metals / oth yringe / other :		E	Container: Vial 500ml Glass 500 ml Plastic	Preserval H ₂ SO ₄ / o none oth	tion: ther: ner:	
Water Quality Met	er type:	YEOKA				1 Plastic	H ₂ \$O ₄ / ot		
	Water Dipper type:								
OBSERVATIONS					2	00 mL Glass	HNO ₃ / ot/	ier;	
Sar	mples:	Colour: _	PIL		Turbidity bon sheen?	Low / Medio Yes /	- (°)		
Weather Condi			Rain _		mperature	31			
N	Pre	vious Week	Rain	MPPED	mperature	ER S	ANPC/A	16	
Reference Information: Estim	nation of Standi		0mm diameter Wel	ls					
itres per lineal meter of water		75mm 2.7	3.7	125mm 5.1	150mm 6.7	10.8 Ez	ote: Typical ciprobe well ameter =75mm		

Sampling procedures are based on AS/NZS 5667.11:1998 Water Quality - Sampling Part 11: Guidance on sampling of groundwaters

Well	DTW	DTL
B403	7-41	10.0
R1402	2.294	513
BHOL	2.99	5.3

GROUNDWATER SAMPLING RECORD FORM



FOR THE	REE WELL \	OLUME METH	OD					Golder
PROJECT	INFORMATI	ОН						
		r: 15462	23	and the second second	AND	Date	25	C1.18
	Clier Site Location		HODA	SANDS		Sampled By	r:	45
GROUND	NATER WEL		TIERRY	- 2V+ 100				
a so a which is a second	and the same and the			Con De Artiste		A LOS TOWNS		LINES OF STREET
Information	ference point		1 1	20	1		BORE ID	
				.4	-	NI-4		17Aol
Depth of we	well (inc filter p	ack) (mm)			-	Note:		<u></u>
Height of filt		iack) (IIIII)		00 t-3	-{	Otom dond refer		5000
	recorded on s	rito	1	+-3	1			of PVC standpipe
	standpipe (mm		50 B	1-01	1	mbRP - metres	below top of re	terence point
Standpipe st		/	70 8	a	-			~
Time	ion up (iii)		6-	7.20	1	Interface probe	usod?	VEQUID
Depth to wat	er (mbRP)		C	171		Depth to produc		YES / NO
	of well (mbRP)	T	- T	. //		Depth to water		185 1-71
		ase of well (m)	-	× +				44.71
	ter in standpipe			2,3		Thickness of pro-	oduct (m)	
Well volume		(111)	4	7				
		k, well volume wihtin s	Omm standain	e in O literatur	21L			
		olumes for 50 mm dia				ista etter ment		
Well diameter	(standpipe plus t	filter pack)	100mm	115mm	vater column wit	125mm	150mm	200mm
Litres per linea	meter of water	in bore	3.7	4.5	4.8	5.1	6.7	10.8
PURGING F	RECORD		orkean and					
Time	Volume	Conductivity (mS	Temp	pH	Redox	Dissolved	Ann	earance
	Purged (L)	or μS)	(°C)	.	Potential (mV)	Oxygen (mg/L)		idity, odour, etc)
0730	7	142	22-56	5.67	453	-0.9	BROW X	HI / GOOM
0733	14	140	22.76	5-70	55.450	~0.9	~	/ //
0736	21	148	22-68	5-74	457	-0.4	4	4
	•							
Total v	ol. purged (L)		No. bor	e vol. purged		Purging	Time (minutes)	
Denth to water	at end of purg	ing (mhPP)	4.1	7				
Purging Metho		WATERRA		nell	VOL			
SAMPLING	RECORD						engineratura (n. 17. gr. gr. gr. gr. 18. february (n. 17. gr. gr. gr. gr. gr. gr.	
	21.	CIPPRA	2	rell us			1183	Z
Sampling meth		410.10	39	rell u	91	Sample ID.	GAU	7
ime sampled		1 40			Samples filter	ed for metals?:	Yes / 160	
Sample Appea	rance:	m 1 1				3	0	00
Colour	00.	econo			Turbidity	Low / Mediu	ım /(High	
Odour				-	arbon sheen?_			
ample Conta	iner and Pres	ervation:	VII	Dupli	cate sample tal	ken? / Dup ID.: _		1.
	DVO	general management of Ac-	5 . 47	to Torrigo	a no handle a men v	्रांग् । गांस्यु १५५० स्ट ार हो स्टब्स्टकारः	The Trial of Straight of	
BSERVATIO	CMC			-			ر په خوموني چې امريکون	
Weather	Conditions:	Temperature:	H	01				
		Precipitation :						
Notes:								
								1050
_								



Appendix F

Soil Laboratory Test Results





ABN: 74 128 806 735

Earlville QLD 4870

Shed 3, 5 Commercial Place

Laborator Cairns Laboratory

0740337815 Fax: 0740546632 Phone:

Email: cnz.lab@cardno.com.au

MINIMUM / MAXIMUM DENSITY REPORT

Client: Golder Associates Pty Ltd

Client Address: 216, Draper Street, Cairns

Project: General Testing

Location: North Queensland Component: Compliance Testing

Area Description:

Date Tested

Report Number:

11512/R/11062-1

Project Number: 11512/P/592

BH1-002 Lot Number:

Internal Test Request: 11512/T/6454

Client Reference/s: Baron River Delta

Report Date / Page: Page 1 of 2 1/12/2016

Test Procedures: AS1289.5.5.1 GA04

Sample Number 11512/S/31833 **Brown Sand**

Sampling Method Tested As Received

Date Sampled 25/11/2016 Sampled By Client Sampled

Material Source Supplied Samples 30/11/2016 Material Type **QAA Sample**

Maximum Dry Density Result					
Size of Mould Used (Litres) 1.0					
Maximum Dry Density (t/m³)	1.88				

Minimum Dry Density Result					
Size of Mould Used (Litres) 1.0					
Minimum Dry Density (t/m³)	1.46				

Remarks



e results of the tests, calibrations and/or measurements included in document are traceable to Australian/national standards. Accredited for compliance with ISO/IEC 17025

1986 Accreditation Number: Corporate Site Number: 11512

approved Signatory: Peter Gode Form ID: W19Rep Rev 1



ABN: 74 128 806 735

Earlville QLD 4870

Address: Shed 3, 5 Commercial Place

Laborator Cairns Laboratory

0740337815 Phone:

Email: cnz.lab@cardno.com.au

MINIMUM / MAXIMUM DENSITY REPORT

Client: Golder Associates Pty Ltd

Client Address: 216, Draper Street, Cairns

Project: General Testing Location: North Queensland

Component: Compliance Testing

Area Description:

Date Tested

Report Number:

11512/R/11062-1

Project Number: 11512/P/592

BH1-002 Lot Number:

Internal Test Request: 11512/T/6454

Fax: 0740546632

Client Reference/s: Baron River Delta

Report Date / Page: 1/12/2016 Page 2 of 2

Test Procedures: AS1289.5.5.1 GA05

Sample Number 11512/S/31834 Yellow Sand

Sampling Method Tested As Received

Date Sampled 25/11/2016 Sampled By Client Sampled

Material Source Supplied Samples 29/11/2016 Material Type **QAA Sample**

Maximum Dry Density Result					
Size of Mould Used (Litres) 1.0					
Maximum Dry Density (t/m³)	1.89				

Minimum Dry Density Result				
Size of Mould Used (Litres)	1.0			
Minimum Dry Density (t/m³)	1.52			

Remarks



e results of the tests, calibrations and/or measurements included in document are traceable to Australian/national standards. Accredited for compliance with ISO/IEC 17025

1986 Accreditation Number: Corporate Site Number: 11512

approved Signatory: Peter Gode Form ID: W19Rep Rev 1



ABN: 74 128 806 735

Laborator Cairns Laboratory

0740337815 **Fax:** 0740546632 Phone:

Email: cnz.lab@cardno.com.au

Shed 3, 5 Commercial Place Earlville QLD 4870

MOISTURE CONTENT REPORT

Client: Golder Associates Pty Ltd

Client Address: 216, Draper Street, Cairns

Project: General Testing

Location: North Queensland

Compliance Testing Component:

Area Description:

11512/R/11061-1 Report Number:

11512/P/592 Project Number:

BH1-002 Lot Number:

Internal Test Request: 11512/T/6454

Client Reference/s: Baron River Delta

Report Date / Page: 1/12/2016 Page 1 of 1

Alea Description.			teport Date / Fage.	1/12/2010	- 1 ago 1 01 1
Test Procedures:	AS1289.2.1.1				
Sample Number	11512/S/31833	11512/S/31834			
ID / Client ID	-	-			
Lot Number	BH1-002	BH1-002			
Date / Time Sampled	25/11/2016	25/11/2016			
Sampling Method	Tested As Received	Tested As Received			
Date Tested	29/11/2016	29/11/2016			
Material Source	Supplied Samples	Supplied Samples			
Material Type	QAA Sample	QAA Sample			
	GA04	GA05			
	Brown Sand	Yellow Sand			
Moisture Content (%)	27.5	14.0			
Sample Number					
ID / Client ID					
Lot Number					
Date / Time Sampled					
Sampling Method					
Date Tested					
Material Source					
Material Type					
Moisture Content (%)					

Remar	ks
-------	----



e results of the tests, calibrations and/or measurements included in document are traceable to Australian/national standards. Accredited for compliance with ISO/IEC 17025

Accreditation Number: 1986 Corporate Site Number: 11512

pproved Signatory: Peter Gode Form ID: W20Rep Rev 1



ABN: 74 128 806 735

Shed 3, 5 Commercial Place Earlville QLD 4870

Laborator Cairns Laboratory

0740337815 Fax: 0740546632 Phone:

Email: cnz.lab@cardno.com.au

ATTERBERG LIMITS REPORT

Client: Golder Associates Pty Ltd

Client Address: 216, Draper Street, Cairns

Project: General Testing

Location: North Queensland

Component: Compliance Testing

Area Description:

11512/R/11063-1 Report Number:

Project Number: 11512/P/592

BH1-002 Lot Number:

Internal Test Request: 11512/T/6454

Client Reference/s: Baron River Delta

Report Date / Page: Page 1 of 2 1/12/2016

Brown Sand

AS1289.3.1.2, AS 1289.3.3.1, AS1289.3.2.1, AS1289.3.4.1 Test Procedures:

Sample Number 11512/S/31833 Sample Location Sampling Method Tested As Received GA04

Date Sampled 25/11/2016 Sampled By Client Sampled **Date Tested** 30/11/2016

Material Source Supplied Samples Att. Drying Method Atterberg Preparation **QAA Sample** Material Type

Material Description

Atterberg Limits Results					
Atterberg Limit	Specification Minimum	Test Result	Specification Maximum		
Liquid Limit (%)		Not Obtainable			
Plastic Limit (%)		Not Obtainable			
Plasticity Index (%)		Non Plastic			
Linear Shrinkage (%)		-			
Linear Shrinkage Defects:	-				

Remarks



e results of the tests, calibrations and/or measurements included in document are traceable to Australian/national standards. Accredited for compliance with ISO/IEC 17025

Accreditation Number: 1986 Corporate Site Number: 11512

approved Signatory: Peter Gode Form ID: W11bRep Rev 1



ABN: 74 128 806 735

Address:

Earlville QLD 4870

Shed 3, 5 Commercial Place

Laborator Cairns Laboratory

0740337815 Fax: 0740546632 Phone:

Email: cnz.lab@cardno.com.au

ATTERBERG LIMITS REPORT

Client: Golder Associates Pty Ltd

Client Address: 216, Draper Street, Cairns

Project: General Testing

Location: North Queensland

Component: Compliance Testing

Area Description:

11512/R/11063-1 Report Number:

Project Number: 11512/P/592

Lot Number: BH1-002

Internal Test Request: 11512/T/6454

Client Reference/s: Baron River Delta

Report Date / Page: Page 2 of 2 1/12/2016

AS1289.3.1.2, AS 1289.3.3.1, AS1289.3.2.1, AS1289.3.4.1 Test Procedures:

Sample Number 11512/S/31834 Sample Location

Sampling Method Tested As Received GA05 Date Sampled 25/11/2016 Yellow Sand

Sampled By Client Sampled **Date Tested** 1/12/2016

Material Source Supplied Samples Att. Drying Method Atterberg Preparation -**QAA Sample** Material Type

Material Description

Atterberg Limits Results					
Atterberg Limit	Specification Minimum	Test Result	Specification Maximum		
Liquid Limit (%)		Not Obtainable			
Plastic Limit (%)		Not Obtainable			
Plasticity Index (%)		Non Plastic			
Linear Shrinkage (%)		-			
Linear Shrinkage Mould Length / Defects:	Mould Length: 150,0mm / -				

Remarks



e results of the tests, calibrations and/or measurements included in document are traceable to Australian/national standards. Accredited for compliance with ISO/IEC 17025

Accreditation Number: 1986 Corporate Site Number: 11512

approved Signatory: Peter Gode

Form ID: W11bRep Rev 1



BN: 74 128 806 735

Address:

Shed 3, 5 Commercial Place Earlville QLD 4870 Laboratory: Cairns Laboratory

Phone: 0740337815 **Fax:**

Email: cnz.lab@cardno.com.au

QUALITY OF MATERIALS REPORT

Client: Golder Associates Pty Ltd

Client Address: 216, Draper Street, Cairns

Project: Project No.: 1546223 Contact: Joseph Parisi

Location: Cairns

Component: Soil Testing

Area Description: Submitted samples

Report Number: 11512/R/11246-1

0740546632

Project Number: 11512/P/677

Lot Number:

Internal Test Request: 11512/T/6529

Client Reference/s: PO: Q003655 | Submitted 02-12-16

Report Date / Page: 9/12/2016 Page 1 of 4

Depth: 0.5-0.8m

Test Procedures AS1289.3.6.1, AS1289.3.1.2, AS1289.3.2.1, AS1289.3.4.1, AS1289.2.1.1, AS 1289.3.3.1

Sample Number 11512/S/32143 GA04-DS1

Sampling Method Tested As Received

Date Sampled 2/12/2016

Sampled By Client Sampled
Date Tested 6/12/2016

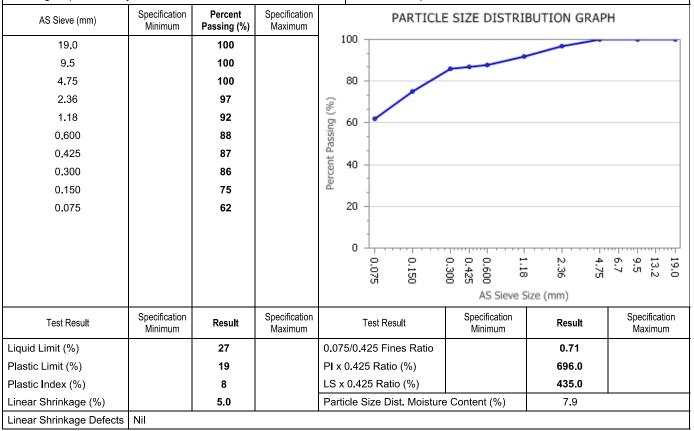
Att. Drying Method Oven Dried

Atterberg Preparation Dry Sieved

Material Source Not Supplied

Material Type Not Supplied (Not Supplied)

Material Description -



Remarks



The results of the tests, calibrations and/or measurements included in this document are traceable to Australian/national standards.

Accredited for compliance with ISO/IEC 17025

Accreditation Number: 1986 Corporate Site Number: 11512





BN: 74 128 806 735

Address:

Shed 3, 5 Commercial Place Earlville QLD 4870

Sciences Laboratory: Cairns Laboratory

Phone: 0740337815 **Fax:**

Email: cnz.lab@cardno.com.au

QUALITY OF MATERIALS REPORT

Client: Golder Associates Pty Ltd

Client Address: 216, Draper Street, Cairns

Project: Project No.: 1546223 Contact: Joseph Parisi

Location: Cairns

Sampling Method

Component: Soil Testing

Area Description: Submitted samples

Report Number: 11512/R/11246-1

0740546632

Project Number: 11512/P/677

Lot Number:

Internal Test Request: 11512/T/6529

Client Reference/s: PO: Q003655 | Submitted 02-12-16

Page 2 of 4

Report Date / Page: 9/12/2016

Test Procedures AS1289.3.6.1, AS1289.3.1.2, AS1289.3.2.1, AS1289.3.4.1, AS1289.2.1.1, AS 1289.3.3.1

Sample Number 11512/S/32144 GA04-DS2

Tested As Received Depth: 1.2-1.5m

Date Sampled 2/12/2016
Sampled By Client Sampled

Date Tested 6/12/2016

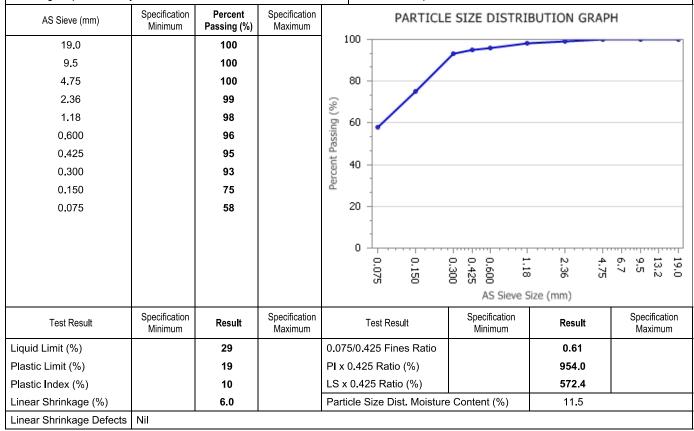
Att. Drying Method Oven Dried

Atterberg Preparation Dry Sieved

Material Source Not Supplied

Material Type Not Supplied (Not Supplied)

Material Description -



Remarks



The results of the tests, calibrations and/or measurements included in this document are traceable to Australian/national standards.

Accredited for compliance with ISO/IEC 17025

Accreditation Number: 1986 Corporate Site Number: 11512





74 128 806 735

Address:

Shed 3 5 Commercial Place Earlville QLD 4870

Laboratory: Cairns Laboratory

0740337815 Phone: Fax:

Email: cnz.lab@cardno.com.au

QUALITY OF MATERIALS REPORT

Client: Golder Associates Pty Ltd

Client Address: 216, Draper Street, Cairns

Project: Project No.: 1546223 Contact: Joseph Parisi

Location: Cairns

Component: Soil Testing

Area Description: Submitted samples Report Number: 11512/R/11246-1

0740546632

Project Number: 11512/P/677

Lot Number:

Internal Test Request: 11512/T/6529

Client Reference/s: PO: Q003655 | Submitted 02-12-16

Depth: 0.6-0.9m

Page 3 of 4

Report Date / Page: 9/12/2016

AS1289.3.6.1, AS1289.3.1.2, AS1289.3.2.1, AS1289.3.4.1, AS1289.2.1.1, AS 1289.3.3.1 **Test Procedures**

Sample Number 11512/S/32145 GA05-DS1

Sampling Method Tested As Received

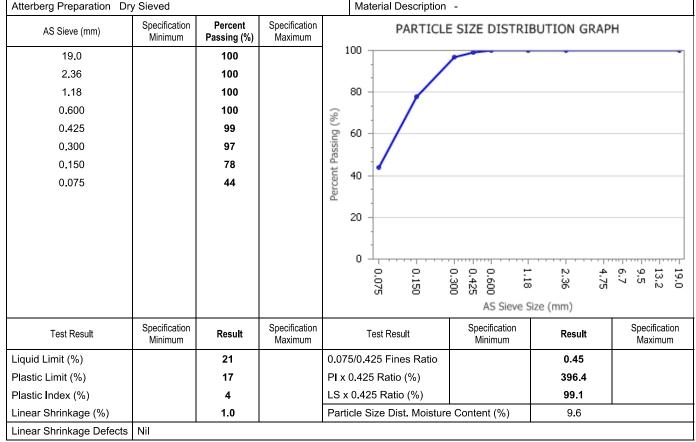
Date Sampled 2/12/2016 Sampled By Client Sampled

Date Tested 6/12/2016

Att. Drying Method Oven Dried Material Source Not Supplied

Material Type Not Supplied (Not Supplied)

Material Description -



Remarks



The results of the tests, calibrations and/or measurements included in this document are traceable to Australian/national standards. Accredited for compliance with ISO/IEC 17025

Accreditation Number: 1986 11512 Corporate Site Number:





74 128 806 735

Address:

Shed 3 5 Commercial Place Earlville QLD 4870

Laboratory: Cairns Laboratory 0740337815

Phone:

Email: cnz.lab@cardno.com.au

QUALITY OF MATERIALS REPORT

Client: Golder Associates Pty Ltd

Client Address: 216, Draper Street, Cairns

Project: Project No.: 1546223 Contact: Joseph Parisi

Location: Cairns

Sampling Method

Component: Soil Testing

Area Description: Submitted samples Report Number: 11512/R/11246-1

0740546632

Fax:

Project Number: 11512/P/677

Lot Number:

Internal Test Request: 11512/T/6529

Client Reference/s: PO: Q003655 | Submitted 02-12-16

Page 4 of 4

Report Date / Page: 9/12/2016

AS1289.3.6.1, AS1289.3.1.2, AS1289.3.2.1, AS1289.3.4.1, AS1289.2.1.1, AS 1289.3.3.1 **Test Procedures**

Sample Number 11512/S/32146 GA05-DS2

> Tested As Received Depth: 1.2-1.5m

Date Sampled 2/12/2016 Sampled By Client Sampled

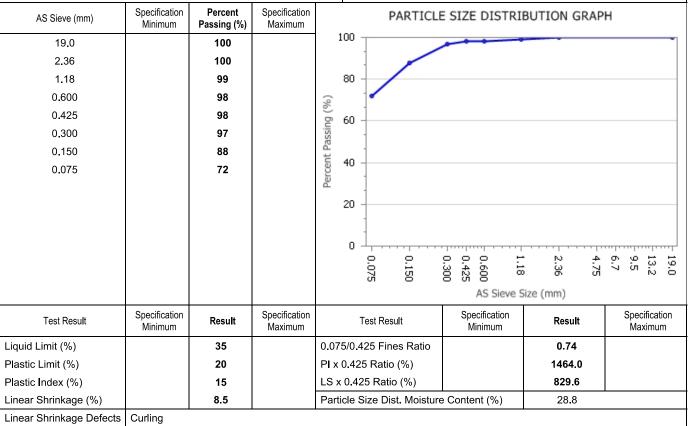
Date Tested 6/12/2016

Att. Drying Method Oven Dried

Atterberg Preparation Dry Sieved Material Source Not Supplied

Material Type Not Supplied (Not Supplied)

Material Description -



Remarks



The results of the tests, calibrations and/or measurements included in this document are traceable to Australian/national standards. Accredited for compliance with ISO/IEC 17025

Accreditation Number: 1986 11512 Corporate Site Number:

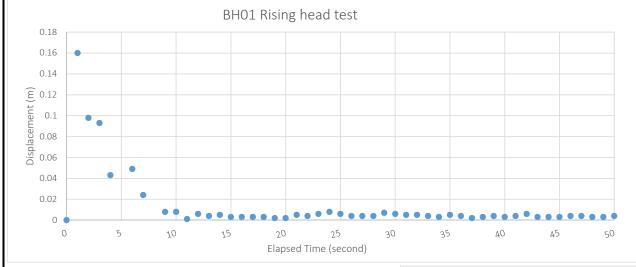




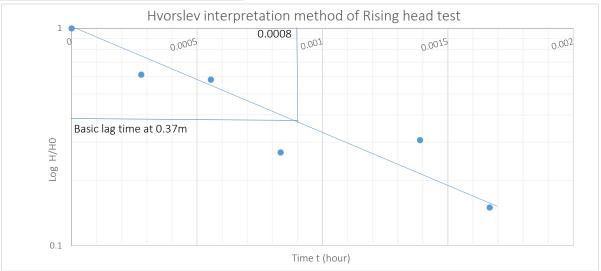
Appendix G

Permeability Test Results



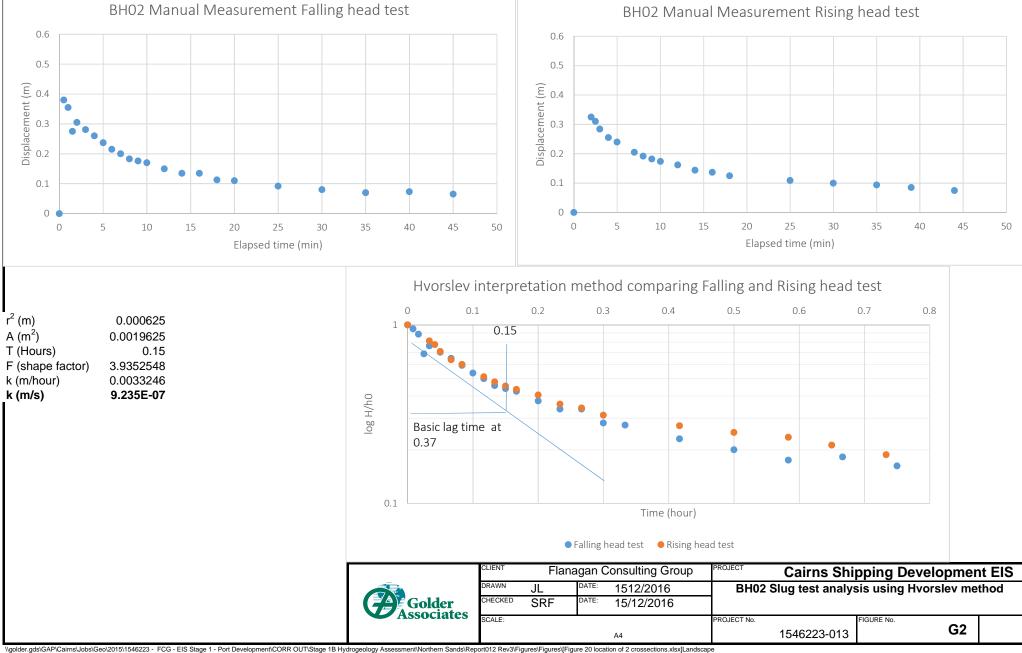


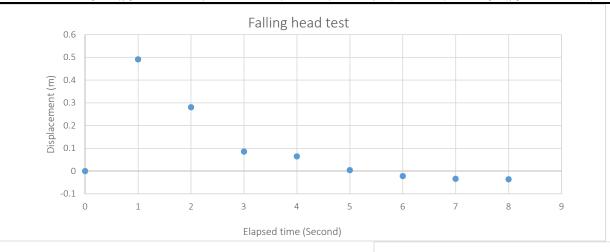
r² (m) 0.000625 A (m²) 0.0019625 T (hour) 0.0008 F (Shape factor) 3.935254829 k (m/hour) 0.623371321 k (m/s) 1.73E-04





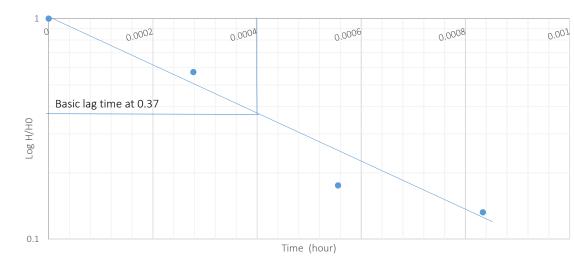
	CLIENT Flanagan Consulting Group				PROJECT	Cairns Shi	pping Dev	elopmen	t EIS
	DRAWN	JL	DATE:	1512/2016	BH01	Slug test analys	sis using H	orslev me	thod
,	CHECKED	SRF	DATE:	15/12/2016					
•	SCALE:			A4	PROJECT No.	1546223-013	FIGURE No.	G1	
\Eia	uroc\[Eiguro 20	location of 2 gross	octions vis	vII andecano					





r² (m) 0.000625 A (m²) 0.0019625 T (Hours) 0.0004 F (shape factor) 4.447913896 k (m/hour) 1.103045184 **k (m/s) 3.06E-04**

Hvorslev interpretation method of falling head test





CLIENT	Flana	agan C	onsulting Group	PROJECT Cair	ns Shipping D	evelopment EIS
DRAWN	¹ JL	DATE:	1512/2016	BH03 Slug tes	st analysis using	Hvorslev method
CHECK	ED SRF	DATE:	15/12/2016			
SCALE:		•		PROJECT No.	FIGURE No.	00
			A4	15462	23-013	G3

0.01

	CLIENT	Flana	agan C	Consulting Group	PROJECT Cairns Shi	pping Developmer	nt EIS
	DRAWN	JL	DATE:	1512/2016	GA04 Slug test analy	sis using Hvorslev me	thod
Golder Associates	CHECKED	SRF	DATE:	15/12/2016			
Associates	SCALE:				PROJECT No.	FIGURE No.	
				A4	1546223-013	G4	

Time t (hour)

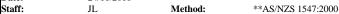
Hydraulic Conductivity Test - Constant Head Test (Talsma-Method)

Site: Northern Sands

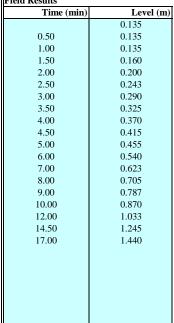
Project Number: 1546223 GA04 **Test Location:**

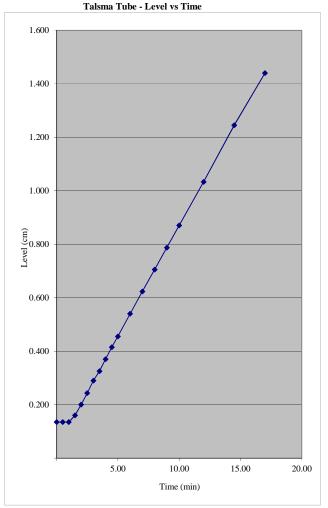
Date: 24/11/2016

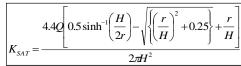
Talsma-Hallam method for determining the permeability of soil using a constantly maintained head of water.



Field Results







Where

Ksat = saturated hydraulic conductivity of soil in cm/min

= correction factor for a systematic under-estimate of soil permeability in the mathematical derivation of the equation.

Q = rate of loss of water from the reservoir in cm³/min

Н = depth to water in test hole in cm.

= radius of the test hole in cm.

Talsma outflow / minute = 8.21 cm/min

Permeameter cross-sec area = 10.86 cm²

 $Q = 89.14 \text{ cm}^3/\text{min}$

H = 133.00 cm

r = 5.00 cm

Correction factor = 4.4

Apply Steady State (See graph above)

Ksat =	4.15E-03	cm/min
=	6.92E-07	m/sec
=	5.98E-02	m/day

Input By: JJP Date: 28/11/2016 Reviewed By: MSC Date: 28/11/2016

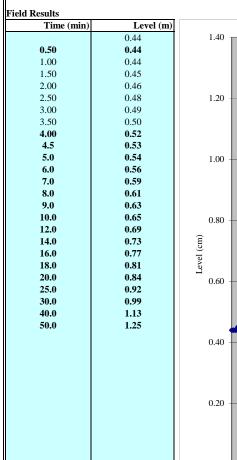
Hydraulic Conductivity Test - Constant Head Test (Talsma-Method)

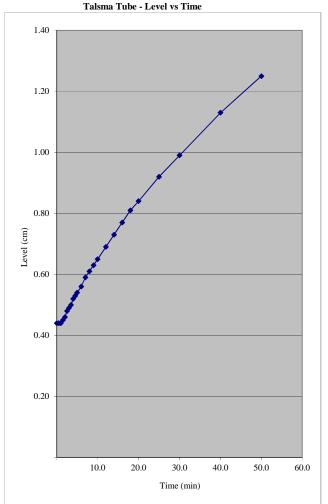
Site: Northern Sands

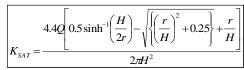
Project Number: 1546223 **Test Location:**

GA05 older Date: 24/11/2016 Method: Staff: **AS/NZS 1547:2000

Talsma-Hallam method for determining the permeability of soil using a constantly maintained head of water.







Where

Ksat = saturated hydraulic conductivity of soil in cm/min

= correction factor for a systematic under-estimate of soil permeability in the mathematical derivation of the equation.

Q = rate of loss of water from the reservoir in cm³/min

Н = depth to water in test hole in cm.

= radius of the test hole in cm.

Talsma outflow / minute = 1.32 cm/min

Permeameter cross-sec area = 10.86 cm²

 $Q = 14.34 \text{ cm}^3/\text{min}$

H = 143.00 cm

r = 3.75 cmCorrection factor = 4.4

Apply Steady State (See graph above)

Ksat =	6.61E-04	cm/min
=	1.10E-07	m/sec
=	9.52E-03	m/day

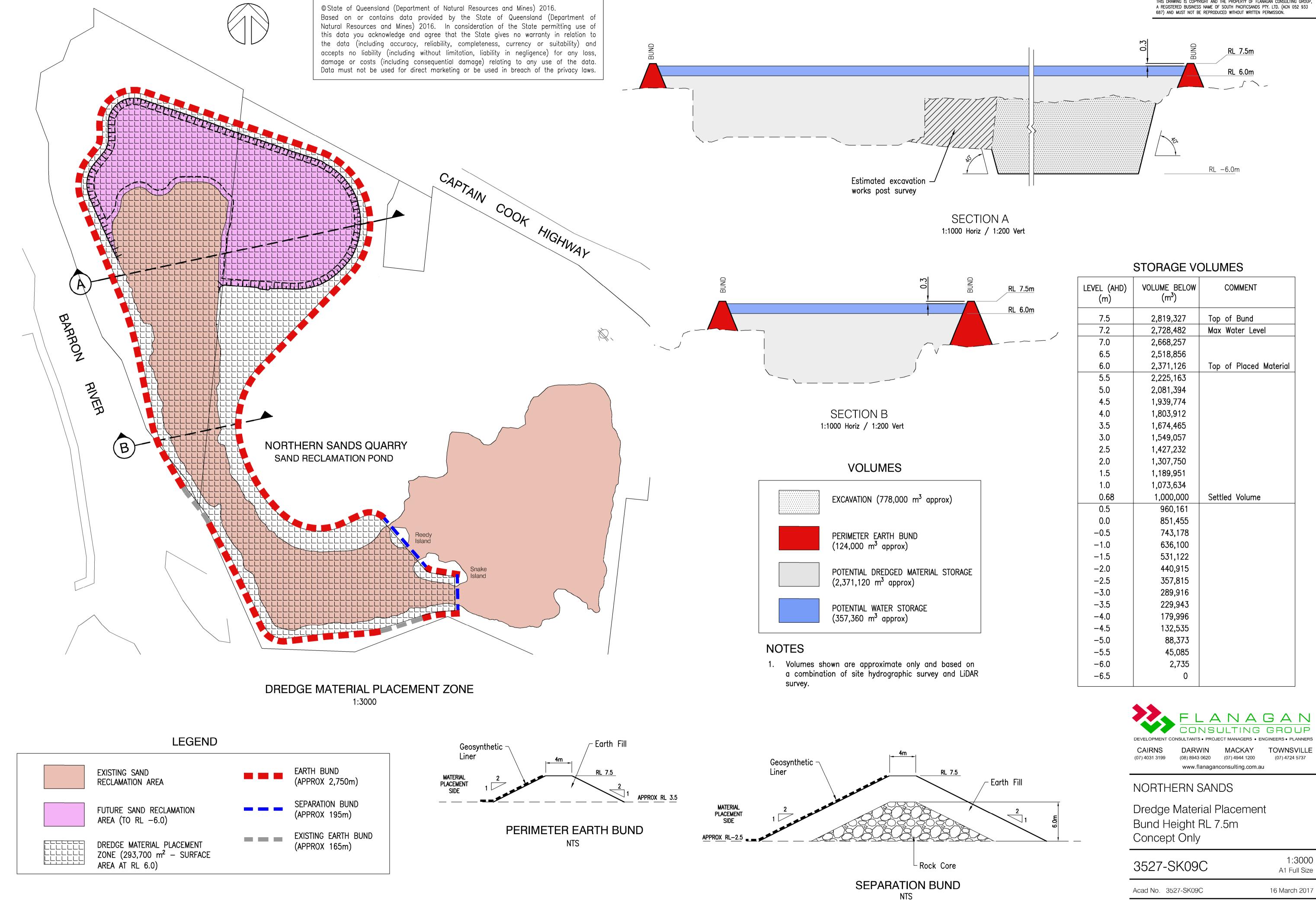
Input By:	JJP	Date:	25/11/2016	
Reviewed By:	MSC	Date:	25/11/2016	



Appendix H

Proposed Expansion of Lake for Placement of Dredge Spoil (Drawing 3527-SK09C)







Appendix I

Queensland Government Terms of Reference Commonwealth Guidelines





Queensland Government Terms of Reference

Flanagan Consulting Group has extracted relevant items from the Queensland Government Terms of Reference for soils and groundwater studies. These items and relevant details are presented below. Items that are not in scope are shaded grey in the table. The section of this report where each item is discussed is also shown in the table.

Relevant Queensland Government ToR - Groundwater

ToR	Title	Details	Comments
5.2	Land	Detail the existing land environment values for all areas associated with the project.	Not addressed in this report.
5.2.1	Land use and tenure	· declared water storage catchments	Not addressed in this report
5.3.2	Water quality	groundwater quality for likely discharge, runoff or seepage waters from any dredge spoil structures using the ANZECC Guidelines (Australian and New Zealand	Section 2.4 Section 3.3
5.5.1	Water resources	Provide an overview of the quality and quantity of any water resources in the vicinity of the project area and the relevance to the project.	Section 2.4 Section 3.2 Section 3.3 Section 3.4
5.5.2	Water resources	potential impacts on the flow and the quality of surface and groundwater from all phases of the project, with reference to their suitability for the current and potential downstream uses and discharge licences	Section 4 Section 5
5.5.2	Water resources	an assessment of all likely impacts on groundwater depletion or recharge regimes	Not addressed in this report
5.5.2	Water resources	where placement of dredge spoil to land is being considered, the use of the ANZECC Guidelines methodology is required to derive water quality trigger values and investigate surface and groundwater quality of the receiving environment that are likely to receive discharge, runoff or seepage waters from any dredge spoil	Not addressed in this report
5.5.2	Water resources	an assessment of the potential to contaminate surface and groundwater resources and measures to prevent, mitigate and remediate such contamination	Section 5
5.5.2	Water resources	details of a monitoring program for the groundwater resources, using existing deep bores, to establish the base line yield and water quality of the supply from those bores	Section 3.3

Commonwealth Guidelines





Flanagan Consulting Group has extracted relevant items from the Commonwealth Guidelines for soils and groundwater studies. These items and relevant details are presented below. Items that are not in scope are shaded grey in the table. The section of this report where each item is discussed is also shown in the table.

Relevant Commonwealth Government Guidelines

Guideline	Title	Details	Comments
5.9	The Existing Environment	This section must provide a description of the project area including baseline condition and trends of coastal, terrestrial and marine environments, including hydrology, sediment characteristics, sediment flows, geography, flora and fauna, cultural and heritage values, and all relevant socioeconomic considerations. This section must link to the proposal description, potential impacts, and proposed avoidance, mitigation, adaptive management framework and/or offset measures throughout the life of the project including preconstruction, construction, operation, and any decommissioning. This section is to also identify and reference any relevant (published and unpublished) studies undertaken in the area which will assist in describing patterns and trends in the environment. Acute and chronic) from geotechnical activities (such as blasting (such as blasting (such as blasting (such as blasting increased marine underwater marine species, including the impacts from noise at varying from each project component (considering the variables e.g. depth, wave height, bottom profile); impacts from proposal on air quality impacts; dredged material and impacts from increased shipping	Section 3
5.9	The Existing Environment	The section must include a description of the environment of the proposal site and the surrounding areas that may be affected by the action. This must include the following information:	Section 2 Section 3
5.9	The Existing Environment	b) A detailed assessment of the nature, extent, likelihood and consequence of the likely short-term and long-term impacts including but not limited to: description of the risks and potential impacts placement impacts	Section 5 Section 6
5.9	The Existing Environment	e) Any technical data, including modelling, and other information used or needed to make a detailed assessment of the relevant impacts;	Section 3 Section 5
5.9	The Existing Environment	h) Consideration of potential impacts throughout the life of the proposal – from preconstruction, construction through to operation and any decommissioning;	Section 5 Section 6
5.9	The Existing Environment	I) Impacts on the existing use of the area and nearby areas that may be affected by the proposed action;	Section 5 Section 6



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Guideline	Title	Details	Comments
5.10	Relevant Impact of the Proposed Action	The EIS must include a description of all of the relevant impacts of the action. Relevant impacts (both direct and indirect) are impacts that the action will have or is likely to have on a matter protected by a controlling provision (as listed in the preamble of this document). This section must provide clear linkages with the existing environmental values described in section 5.9 and proposed avoidance, safeguards, management and mitigation measures described in section 5.11. Impacts during all phases of the project must be addressed. This section must include:	Section 5 Section 6
5.10	Relevant Impact of the Proposed Action	a) A description of the framework used to assess impacts, including risk assessment processes based on an approved standard;	Section 6
5.10	Relevant Impact of the Proposed Action	b) A detailed assessment of the nature, extent, likelihood and consequence of the likely short-term and long-term impacts including but not limited to: description of the risks and potential impacts (acute and chronic) from geotechnical activities (such as blasting and pile driving), impacts of increased marine underwater noise on marine species, including the impacts from noise at varying distances from each project component (considering the environmental variables e.g. depth, wave height, bottom profile); impacts from the proposal on air quality impacts; dredging and dredged material placement impacts and impacts from increased shipping;	Section 5 Section 6
5.10	Relevant Impact of the Proposed Action	c) A statement whether any relevant impacts are likely to be unknown, unpredictable, irreversible or sub-lethal (reversible over time) and what confidence level is placed on the predictions of relevant impacts;	Section 6.2
5.10	Relevant Impact of the Proposed Action	d) Analysis of the significance of the impacts;	Section 6.2
5.10	Relevant Impact of the Proposed Action	e) Any technical data, including modelling, and other information used or needed to make a detailed assessment of the relevant impacts;	Section 2 Section 3 Section 5
5.10	Relevant Impact of the Proposed Action	f) Consideration of potential impacts throughout the life of the proposal – from preconstruction, construction through to operation and any decommissioning;	Section 6
5.10.9	Dredging and Dredged Material	h) Assessment of the risk and potential impacts of acid sulfate soils (ASS) and potential acid sulfate soils (PASS);	Not considered in this report





Guideline	Title	Details	Comments
5.11	Proposed avoidance, safeguards, management and mitigation measures	The EIS must provide information on proposed avoidance, safeguards and mitigation measures to deal with the impacts of the action. Specific and detailed descriptions of proposed measures must be provided and substantiated, based on best available practices/standards and must include the following elements.	Section 5
5.11	Proposed avoidance, safeguards, management and mitigation measures	a) Identify the level of risk associated with potential impacts already identified and those that require mitigation, monitoring or management to avoid or reduce impacts to an acceptable level;	Section 6.2
5.11	Proposed avoidance, safeguards, management and mitigation measures	b) A consolidated list of measures proposed to be undertaken to avoid, prevent, minimise or compensate (in priority order) for the impacts of the action (as specified in section 5.10), including: i. A description of proposed avoidance, safeguards and mitigation measures to deal with impacts of the action, including measures proposed to be taken by State governments, local governments or the proponent; ii. Assessment of the expected or predicted effectiveness of the measures; iii. Any statutory or policy basis for the mitigation measures; iv. The cost of the mitigation measures; and v. The resulting risk level for that impact post- avoidance, mitigation and/or management.	Section 5 Section 6
5.11	Proposed avoidance, safeguards, management and mitigation	c) Particular focus must be given to: i. Determining factors in the planning of the proposal so as to avoic damage to the environment;	Section 5





Appendix J

Important information relating to this document





IMPORTANT INFORMATION RELATING TO THIS REPORT

The document ("Report") to which this page is attached and which this page forms a part of, has been issued by Golder Associates Pty Ltd ("Golder") subject to the important limitations and other qualifications set out below.

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Having regard to the matters referred to in the previous paragraphs on this page in particular, carrying out the Services has allowed Golder to form no more than an opinion as to the actual conditions at any relevant location. That opinion is necessarily constrained by the extent of the information collected by Golder or otherwise made available to Golder. Further, the passage of time may affect the accuracy, applicability or usefulness of the opinions, assessments or other information in this Report. This Report is based upon the information and other circumstances that existed and were known to Golder when the Services were performed and this Report was prepared. Golder has not considered the effect of any possible future developments including physical changes to any relevant location or changes to any laws or regulations relevant to such location.

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