



City of Greater Geraldton

Point Moore Residential Onsite Effluent Treatment and Disposal Study

November 2016

Executive Summary

Point Moore accommodates approximately 200 residential dwellings that were built in the 1960s and 1970s on leasehold land within reserves vested with the City of Greater Geraldton. All dwellings use onsite systems (septic tanks with wastewater disposed of via leach drains or soak wells) for wastewater treatment and disposal. Whilst originally established as holiday cottages, many of the Point Moore lessees are now retirees and beach lifestyle enthusiasts that occupy the residences year-round.

The *Point Moore Inundation & Coastal Processes Study* completed in 2015 has highlighted that the area could be severely impacted by coastal inundation and erosion in the future. Given the small size of the residential properties (in the order of 300 m²), the low elevation of the area (existing ground levels in the order of two metres above current sea levels) and predictions that sea levels will rise significantly in the future, the continued viability of onsite wastewater disposal systems in the area has also come into question.

The City of Greater Geraldton (CGG) commissioned GHD Pty Ltd (GHD) to complete a study to review available information and to conduct a series of investigations to assess the performance of the existing onsite systems, their potential impacts on the local environment, and the degree to which they comply with current legislation and guidelines. The fieldwork to locate and to assess the onsite systems at a selection of properties was completed by Sun City Plumbing (SCP).

The following conclusions have resulted from this study:

- 1. Observations made during the field investigation indicate that a significant number of the existing septic tanks and leach drains/soak wells are in a poor condition and require remedial works and in some cases replacement.
- 2. The properties in the study area are significantly smaller than the minimum lot size currently permitted for onsite wastewater disposal (typically 2,000 m²), and many of the onsite systems do not comply with current standards in a number of respects (e.g. sizing, configuration, horizontal setbacks, vertical separation distance to groundwater). For many properties it would not be possible to upgrade the existing onsite systems to meet current standards, or install alternative onsite systems that comply with current standards.
- 3. Though local groundwater is not used for irrigation or any other purpose, it does discharge to the nearby ocean, and residents or others could come into contact with groundwater when undertaking a range of land-based activities. In relation to public health risks:
 - The potential for contact with groundwater when undertaking land based sub-surface activities such as excavation or trenching works is considered to represent a potential health risk to residents and others undertaking such activities in the study area. This risk will increase over time as local groundwater levels increase as a direct consequence of sea level rise.
 - It is considered unlikely that elevated levels of pathogens in groundwater flowing from the study area would pose a significant health risk to persons engaging in primary contact recreation in the ocean near Point Moore given natural purification processes in the aquifer and the high levels of dilution that would typically occur where the groundwater discharges into the ocean. However, under conditions of calm winds and low wave climate, rates of dilution may be greatly reduced, thereby increasing the potential health risk. These conditions typically occur late in the bathing season from March-May.

- It is not possible to discount the possibility that onsite disposal of effluent from the Point Moore residential properties is at least partly responsible for the observed seasonal spikes in *Enterococci* levels at the CGG's local marine water quality monitoring sites.
- 4. In the long term local groundwater levels will rise as sea levels rise, and the magnitude of the rise will severely constrain the potential to dispose of wastewater generated in the study area with the existing conventional onsite septic tank and leach drain/soak well systems approach.
- 5. If residential properties are to remain at Point Moore for the long term then a reticulated wastewater collection system will need to be installed that routs wastewater to the Water Corporation's Geraldton wastewater scheme.
- 6. An indicative cost estimate to design and to construct a conventional reticulated gravity sewer type collection system to serve all properties in the study area is \$6 to 10M. At a unit cost of approximately \$35,000 to \$55,000 per property, this is likely to be prohibitively expensive. Whilst alternative wastewater collection technologies exist that may be able to be implemented at a significantly lower capital cost, ongoing costs for these systems would be higher.
- 7. Whilst nutrient levels in sampled groundwater indicated elevated wastewater-induced contamination above the adopted assessment criteria for all monitoring rounds, given the high levels of dilution that typically occur where groundwater discharges into the ocean it is considered unlikely that elevated levels of nutrients in groundwater flowing from the study area are having any measurable impact on near shore marine ecosystems.

This report is subject to, and must be read in conjunction with, the limitations set out in Section 1.3 and the assumptions and qualifications contained throughout the Report.

Table of Contents

1
1
2
4
4
4
5
7
11
11
11
16
17
17
17
18
19
20
20
20
20
22
22
23
27
33
36
36
36
36
38
39
40
42

Table index

Table 1	Scope of Work	1
Table 2	Key regulatory requirements	18
Table 3	Rationale of GMW Locations	24
Table 4	Summary of Geology Encountered at MW1	27
Table 5	Groundwater Well Survey Data	28
Table 6	Groundwater Well Survey Data	29
Table 7	Groundwater Well Survey Data	30
Table 8	Summary of Laboratory Groundwater Quality Data	34

Figure index

Locality Plan	6
Standing Groundwater Levels at SHP8	7
Near Shore Marine Sampling Points (CGG, 2016)	8
Comparison of Nearshore Marine <i>Enterococci</i> Levels with ANZECC (2000) Recreational Water Quality Guidelines	9
Comparison of Nearshore Marine <i>Enterococci</i> Levels with NHMRC (2008) Recreational Water Quality Categories	10
Coastal Processes Allowance Map (m p rogers, 2015)	12
Inundation Mapping - Present Day Setback (m p rogers, 2015)	13
Inundation Depth – 2030 Setback (20 year ARI Event) (m p rogers, 2015)	14
Combine Coastal Vulnerability Mapping - Present Day (Setback) (m p rogers, 2015)	15
Groundwater Levels Logger Data for MW1, MW2 and MW3	32
Groundwater Contours, August 2016	33
	Locality Plan Standing Groundwater Levels at SHP8 Near Shore Marine Sampling Points (CGG, 2016) Comparison of Nearshore Marine <i>Enterococci</i> Levels with ANZECC (2000) Recreational Water Quality Guidelines Comparison of Nearshore Marine <i>Enterococci</i> Levels with NHMRC (2008) Recreational Water Quality Categories Coastal Processes Allowance Map (m p rogers, 2015) Inundation Mapping - Present Day Setback (m p rogers, 2015) Inundation Depth – 2030 Setback (20 year ARI Event) (m p rogers, 2015) Combine Coastal Vulnerability Mapping - Present Day (Setback) (m p rogers, 2015) Groundwater Levels Logger Data for MW1, MW2 and MW3 Groundwater Contours, August 2016

Appendices

Appendix A – Nearshore Sampling Tabulated Exceedances
Appendix B – Point Moore Inundation and Coastal Processes Study Summary Report
Appendix C – Groundwater Monitoring Well Logs
Appendix D – Groundwater Quality Field Data Sheets
Appendix E – Quality Assurance and Quality Control
Appendix F – Chain of Custody and Laboratory Reports
Appendix G – Tabulated Groundwater Analytical Results
Appendix H – Calibration Certificates
Appendix I – Field Works Photographs

1. Introduction

Point Moore is located in the City of Greater Geraldton (CGG) and is a prominent sandy foreland protected by the Point Moore reefs. There are approximately 200 residential dwellings near the coast built in the 1960s and 1970s on leasehold land within reserves vested in CGG. All dwellings use onsite systems (septic tanks with effluent disposed of via leach drains or soak wells) for wastewater treatment and disposal. Whilst originally established as holiday cottages, many of the Point Moore lessees are now retirees and beach lifestyle enthusiasts that occupy the residences year-round.

Given their age, many of the original septic tanks and soak wells/leach drains may have reached or be approaching their serviceable life and some may have been replaced or upgraded. Due to the low ground levels in the area there is limited vertical separation distance between the soak wells/leach drains and the water table. This separation distance will likely reduce in the future due to rise in sea levels.

M P Rogers & Associates completed the *Point Moore Inundation & Coastal Processes Study*, hereafter termed the Coastal Inundation Study, in 2015 (m p rogers, 2015). The study identified areas of Point Moore which could be impacted by coastal inundation and erosion. Further discussion with regards to this study is provided in Section 3 and the summary report from this m p rogers study is provided in Appendix A.

The CGG commissioned GHD Pty Ltd (GHD) to complete a study to review available information and to conduct a series of investigations to assess the performance of the existing onsite systems and their potential impacts on the local environment and the degree to which they comply with current legislation and guidelines. The fieldwork completed to locate and assess the onsite systems at a selection of properties was completed by Sun City Plumbing (SCP).

1.1 Study Objective

The overall objective of the study is to provide the Point Moore stakeholders with data and documentation on the performance and compliance of the existing onsite wastewater treatment and disposal systems and to assist the CGG in their decision-making process relating to the future status of and potential obligation to Point Moore lessees beyond the current lease expiry dates of 2025 and 2028.

1.2 Scope of Work

The study was completed in two stages, with the key tasks in each stage listed in Table 1.

Phase/Task	Description
Stage 1	
Task 1	Desktop review of available information.
Task 2	Soil and groundwater investigations, including installation of three new monitoring wells.
Task 3	Groundwater level and quality monitoring program (three-month program).
Stage 2	

Table 1Scope of Work

Phase/Task	Description
Task 4	Onsite wastewater system field inspections.
Task 5	Technical assessment and reporting.

This report documents the findings from all five tasks undertaken to complete this study.

1.3 Limitations and Assumptions

This report has been prepared by GHD for the City of Greater Geraldton and may only be used and relied on by the City of Greater Geraldton for the purpose agreed between GHD and the City of Greater Geraldton as set out in Section 1.1 of this report.

GHD otherwise disclaims responsibility to any person other than the City of Greater Geraldton arising in connection with this report. GHD also excludes implied warranties and conditions, to the extent legally permissible.

The services undertaken by GHD in connection with preparing this report were limited to those specifically detailed in the report and are subject to the scope limitations set out in the report.

The opinions, conclusions and any recommendations in this report are based on conditions encountered and information reviewed at the date of preparation of the report. GHD has no responsibility or obligation to update this report to account for events or changes occurring subsequent to the date that the report was prepared.

The opinions, conclusions and any recommendations in this report are based on assumptions made by GHD described in this report. GHD disclaims liability arising from any of the assumptions being incorrect.

GHD has prepared this report on the basis of information provided by the City of Greater Geraldton and others who provided information to GHD (including Government authorities), which GHD has not independently verified or checked beyond the agreed scope of work. GHD does not accept liability in connection with such unverified information, including errors and omissions in the report which were caused by errors or omissions in that information.

The opinions, conclusions and any recommendations in this report are based on information obtained from and testing undertaken at or in connection with, specific sample points. Site conditions at other parts of the site may be different from the site conditions found at the specific sample points.

Investigations undertaken in respect of this report are constrained by the particular site conditions, such as the location of buildings, services and vegetation. As a result, not all relevant site features and conditions may have been identified in this report. At the time of preparing this report, the nature of the proposed redevelopment (and as such the likely magnitude of disturbance across the site) remained unconfirmed.

Site conditions (including the presence of hazardous substances and/or site contamination) may change after the date of this Report. GHD does not accept responsibility arising from, or in connection with, any change to the site conditions. GHD is also not responsible for updating this report if the site conditions change.

GHD has prepared indicative cost estimates using information reasonably available to GHD; and based on assumptions and judgments made by GHD.

The indicative cost estimates have been prepared to provide an order of magnitude indication of the costs to upgrade the existing onsite systems or install new reticulated wastewater infrastructure. They must not be used for any other purpose.

The cost estimates are indicative estimates only. Actual prices, costs and other variables may be different to those used to prepare the cost estimates and may change. Unless as otherwise specified in this report, no detailed quotation has been obtained for actions identified in this report. GHD does not represent, warrant or guarantee that upgrade of the existing onsite systems or installation of new reticulated wastewater infrastructure could be completed at costs which are the same or less than the estimated indicative costs.

Where estimates of potential costs are provided with an indicated level of confidence, notwithstanding the conservatism of the level of confidence selected as the planning level, there remains a chance that the cost will be greater than the planning estimate and any funding would not be adequate. The confidence level considered to be most appropriate for planning purposes will vary depending on the conservatism of the user and the nature of the project. The user should therefore select appropriate confidence levels to suit their particular risk profile.

2. Site Identification and Physical Characteristics

2.1 Site Description

Point Moore is located approximately two kilometres (km) south of the Geraldton business district, and forms the western extremity of the Geraldton tombolo (CGG, 2016). The study area is bounded by the Indian Ocean to the south, east and west, and John Willcock Link to the north. The subject land comprises several land parcels:

- Reserves vested in the City of Greater Geraldton:
 - Reserve R29729 Public Recreation (Pages Beach Coastal Reserve);
 - Reserve R2562 Esplanade and Recreation (Greys Beach, Point Moore and Explosives Coastal Reserve);
 - Reserve R29173 Caravan Park and Tourist Accommodation;
 - Reserve R25459 Recreation and Leasing of Cottages; and
 - Reserve R31658 Parklands.
- Reserves vested in the Australian Maritime Safety Authority:
 - Reserve R44687 Navigation, Communication, Meteorology and Survey (Point Moore Lighthouse).
- Other land:
 - Private land holding 481 Marine Terrace (land and property around Point Moore lighthouse);
 - Vacant Crown Land Part of Point Moore and Greys Beach.

The focus of this study is the residential dwellings located within Reserve R25459. There are approximately 200 residential dwellings in this reserve, typically two to three bedroom dwellings, each on approximately 300 m² of land area. A locality plan is included as Figure 1.

2.2 Geology and Soil

2.2.1 Geology

Regional geological mapping indicates that the surface geology of the study area is dune and beach sand, which is comprised of white calcareous and quartzose sand (Geological Survey WA, 1995).

2.2.2 Soil Landscapes

DAFWA (2016) mapping of the South-West of Western Australia was used to characterise the soil-landscapes present within the study area. The study area is covered by the Quindalup Central 1 urban phase, which is described as follows:

• Urban development on Quindalup 1 coastal dune subsystem on Aeolian calcareous sands and minor limestone in the North Coastal Plain, adjacent to the coast from Jurien Bay to Bluff Point. Man-made, disturbed soils; originally Calcareous deep sand.

2.2.3 Preliminary Acid Sulphate Soil Assessment

A review of the WA Atlas for Department of Environment Regulation (DER) Acid Sulphate Soil risk mapping (Landgate, 2016), indicated that potential ASS has not been mapped within the study area. A search of the CSIRO (2016) database indicated that there was an extremely low probably of ASS occurring with very low confidence.

2.3 Hydrogeology

2.3.1 Aquifers and Groundwater Flow

The Perth Groundwater Atlas indicates that the Site is located above a Superficial (unconfined) Aquifer, Perth – Superficial Swan (DoW, 2016a).

The atlas indicates that groundwater beneath the study area is at approximately three metres below ground level. Groundwater was identified to flow towards the Indian Ocean.

2.3.2 Local Bores

General

Based on advice from CGG, groundwater from the Point Moore area is not used for irrigation or any other purposes in the study area, and the nearest production bores are the CGG bores located approximately 600 m east to north east of the study area near Ocean and Point Streets. These bores are up-gradient of the study area and supply irrigation water for open space areas near the Geraldton city business district.

Department of Water WIN Bore Search

A search of the Department of Water's (DoW) water information reporting (WIN) bore database indicates that three bores are registered within a 2.5 km radius of the study area. The bores are located approximately 2 km north-east and 1.5 km south-east from the study area. No other registered bores occur down-gradient or up gradient of the study area within 1 km during the search.



Figure 1 Locality Plan

2.3.3 Local Groundwater Levels and Quality

Prior to this study there was one groundwater monitoring well (GMW – SHP8) located in the study area, more specifically within Coxswain Park as shown on Figure 11.

Groundwater Quality Data

No data was provided by CGG that could be used to characterise the quality of groundwater in the study area.

Groundwater Depth

The CGG is required under its groundwater licence to report to the DoW the standing water levels in GMW – SHP8 twice a year (Pers. comm. Michael DuFour, 28 July 2016). The standing water levels over the period February 2013 to August 2015 are depicted in Figure 2. There are some data gaps over this period, and no data was provided for the period beyond August 2015.



Figure 2 Standing Groundwater Levels at SHP8

2.4 Nearshore Marine Environment

2.4.1 General

The nearshore bathymetry ranges from 0 to 10 m in depth and comprises a complex system of exposed reefs and deeper channels. Beaches are sandy, and where sheltered they are wide and flat. In exposed areas the beaches are narrower and subject to erosion (CGG, 2016).

2.4.2 Water Quality

The CGG undertakes marine surface water monitoring at designated beach locations within Geraldton and Greenough at sampling locations depicted in Figure 3. GHD has reviewed the laboratory water quality data supplied by CGG for the period November 2012 to April 2016.



Figure 3 Near Shore Marine Sampling Points (CGG, 2016)

Two monitoring locations, Point Moore North (GE3-003) and Pages Beach North (GE3-021), are located close to the study area. The *Enterococci* data at these two locations, and at one location remote from Point Moore (Town Beach Jetty, GE3-001), was assessed in this study. As shown in Figure 4, the *Enterococci* data was compared against the ANZECC (2000) water quality guidelines for primary and secondary contact recreation which are:

- Primary contact: Median of 35 *Enterococci*/100 mL during bathing season with maximum in one sample being 60 to 100 *Enterococcoi*/100 mL;
- Secondary Contact: Median of 230 *Enterococcoi/*100 mL and maximum number in any one sample being 450 to 700 organisms/100 mL.

Based on these water quality guidelines, some reported *Enterococci* results were above the primary contact and in some instances above the secondary contact guideline values. The reported results thus indicate that persons engaging in both primary contact (e.g. swimming, skin diving) and secondary contact (e.g. boating and fishing) recreational activities in the area are exposing themselves to health risks. Of note, as shown Figure 4 there were no exceedances of the ANZECC (2000) guideline values at the Town Beach Jetty monitoring location over the same period.

The *Enterococci* data was also compared against the NHMRC (2008) percentile values for determining water-quality assessment categories for the protection of healthy adult bathers (refer to Appendix A) which are:

- Category A: <40 Enterococci/100 mL (no observed-adverse-effect-level (NOAEL);
- Category B: 41-200 *Enterococci*/100 mL (threshold of illness transmission in most epidemiological studies attempted to define a NOAEL);
- Category C: 200-500 *Enterococci*/100 mL (represents substantial elevation in the probability of all adverse health outcomes); and
- Category D: >500 Enterococci/100 mL (above this level may be a significant risk of high levels of illness transmission of Enterococci/).

As depicted in Figure 5, one reported *Enterococci* result was in Category C, and one result was in Category D, i.e. at levels that pose a significant risk of illness transmission of *Enterococci* and considered to warrant further investigation.



Figure 4 Comparison of Nearshore Marine Enterococci Levels with ANZECC (2000) Recreational Water Quality Guidelines



Figure 5 Comparison of Nearshore Marine Enterococci Levels with NHMRC (2008) Recreational Water Quality Categories

3.1 Study Overview

The Coastal Inundation Study (m p rogers, 2015) identified areas of Point Moore which could be impacted by coastal inundation and erosion. As part of the study the following were completed:

- Cyclone storm surge modelling to determine the potential inundation of these extreme events;
- Analysis of available water level records to determine the potential inundation caused by non-cyclonic events;
- Modelling the potential beach and dune erosion caused by severe events;
- Assessment of historical and potential future shoreline movement caused by natural coastal processes; and
- Assessment of the effects of potential sea level rise (assuming 0.9 metres of sea level rise by year 2110 as required by State Planning Policy No. 2.6 (SPP 2.6)) on the coastal inundation and erosion.

The Coastal Inundation Study Summary Report is provided in Appendix B.

3.2 Maps Depicting Study Results

The Coastal Inundation Study (m p rogers, 2015) included a series of maps to depict the study results and recommendations. Some of the key maps are shown and briefly described next.

Coastal Processes Allowance Plan

The coastal processes allowance plan relates to the extent of coastal erosion over 'planning horizons' including present day, 2030, 2070 and 2110. The recommended present day, 2030, 2070 and 2110 coastal setbacks are shown in Figure 6.

Inundation Mapping Plans

The inundation map (Figure 7) depicts the areas of inundation that could occur in the present, with colours to represent different severity events. The purple shading represents those areas that are predicted to be inundated during a 20-year average recurrence interval (ARI) event, the blue shading demonstrates the additional area that would be inundated during a 100 year ARI event, and the green shading the additional area inundated during a 500 year ARI event. Inundation maps used to establish the 2030, 2070 and 2110 coastal setback are provided in Appendix B.

Inundation Depth Plans

The inundation depth plan provided in Figure 8 shows the potential depth of inundation that is predicted to occur in 2030 under a 20 year average recurrence interval (ARI) event. Other inundation depths maps produced by m p rogers (2015) are included in Appendix B.

Combined Coastal Vulnerability Mapping Plans

The combined coastal vulnerability map provided in Figure 9 depicts the area that would be impacted by inundation or coastal erosion (based on present day setbacks) in a 500 year ARI event. Similar plans generated for the recommended 2030, 2070 and 2110 setbacks (500 year ARI events) are provided in Appendix B.



Figure 6 Coastal Processes Allowance Map (m p rogers, 2015)



Figure 7 Inundation Mapping - Present Day Setback (m p rogers, 2015)



Figure 8 Inundation Depth – 2030 Setback (20 year ARI Event) (m p rogers, 2015)



Figure 9 Combine Coastal Vulnerability Mapping - Present Day (Setback) (m p rogers, 2015)

3.3 Consultation with Regulatory and Planning Bodies

Advice and comments on the Coastal Inundation Study (m p rogers, 2015) provided to CGG by the Department of Health (DoH), Department of Lands (DoL) and Department of Planning (DoP) are summarised next. Note that CGG sought a response from the Department of Environment Regulation (DER), no comment was provided.

3.3.1 Department of Health

In a correspondence letter the DoH advised:

When developing strategies based upon the findings in the report you may consider a risk assessment of each of the respective areas. Such strategies should incorporate issues such as disaster preparedness, emergency shelters and recovery (DoH, 2016a).

3.3.2 Department of Lands

In a correspondence letter the DoL advised:

As indicated at that meeting, the findings of the report indicated that the reserve is at risk from coastal hazards and that the risk is increasing both in extent and depth. The report provides impetus for further discussion with the City on the long term land use planning for the area post the expiry of the current leases on Point Moore Reserve 25459 (DoL, 2016).

3.3.3 Department of Planning

In email correspondence the DoP advised:

The WAPC / DoP position on the West End is contained in the Greater Geraldton Structure Plan June 2011 which identifies it as an area of State Government owned land that provides facilities for tourism, visitor activity. It is severely constrained due to its proximity to the port, a lack of wastewater and vulnerability to storm surges, inundation and long term coastal recession (erosion). I believe the City has just updated its Local Planning Strategy and Scheme, this was the best place to include its intentions for the West End, you should discuss this with your colleague, Murray Connell, the City's Manager Urban and Regional Development. In the LPS on the Geraldton Area Strategy Plan the West End is identified as Strategic Tourism. The LPS also includes at section 4.11 page 27 an action to "ensure land use decision making is based on the best available science regarding coastal processes and the need for adequate setbacks" (DoP, 2016a).

In a correspondence letter the DoP further advised:

DoP.... consider the study to be adequate in demonstrating that much of the current development footprint is already at risk from coastal hazards and that the risk is increasing both in extent and depth. Hence there is sufficient basis and detail in the study to conclude against extension of the current leases when they expire in 2028. It is unlikely that the City or the Department of Lands would wish to fund and maintain coastal protection works for this Crown Reserve and hence leaseholders should not be under any hope of such action (DoP, 2016b).

4. Wastewater Management

4.1 General

Many of the residential properties established on the Point Moore leases were built in the 1960s and 1970s. Originally established as holiday cottages, they have been commonly acquired by retirees and beach lifestyle enthusiasts. Onsite wastewater treatment and disposal systems that were approved by DoH at that time consisted of either:

- combined systems, comprising two concrete septic tanks (1500 mm and 1200 mm diameter) and two standard size (1200 mm diameter) concrete soak wells or one 9 m leach drain; or
- separate systems comprising one tank and one soak well each for toilet waste and another tank and soak well receiving wastewater from separate sewer lines, with the tanks and soak wells typically located on opposite sides of the dwelling (CGG, 2016a).

Having land areas of approximately 300 m², the residential properties are significantly smaller than the minimum lot area currently recommended for houses reliant on onsite wastewater systems. For instance, in its discretionary provisions the Draft Country Sewerage Policy (DoH, 2003) states that lot sizes down to 1,000 m² can be considered under certain conditions (e.g. residential subdivisions in remote areas or towns without sewerage that do not create more than 25 lots). Given that the Point Moore area is within the Geraldton townsite, and that Geraldton is a sewered town, the discretionary provisions of this policy do not apply to Point Moore.

The City's internal health database established in 2003 has two reported cases of failing septic systems; one at Zodiac Lane and another at Bosons Crescent. This compares with 44 other enquiries received from various other locations across the City outside the Point Moore site in relation to poorly functioning septic systems (CGG, 2016).

In 2006 the City received advice from the DoH stating that it "objects to the creation of any new lots and is unsupportive of the extension of the leases, without adequate provision being made for effluent disposal". Accordingly, the City has since issued standard advice on settlement enquires regarding Point Moore properties because of the potential interest at the time from building owners to apply for building extensions; the advice stated;

"The future owner is advised that existing lease lot size may restrict / constrain the available area for future upgrade to onsite effluent disposal system as a result of functional failure or redevelopment in future" (CGG, 2016).

4.2 **Review of Available Information on Existing On-site Systems**

CGG provided GHD with a copy of as-constructed records (stamped block plans) and correspondence that provides information on the existing onsite wastewater treatment and disposal systems for approximately 50% of the houses (dated from 1972 to 2001). GHD's review of this information found that:

- Of the nominal 80 houses for which information was available on the effluent disposal system, approximately 20% of the houses use soak wells (typically two soak wells per house), with the remainder using leach drains (typically 1 x 9 m leach drain).
- In July 1983 the Town of Geraldton's Senior Health Surveyor reported that approximately 59 beach cottage leases were due to expire in 1986, and that of these 30 have septic or disposal systems that do not conform to Health Act legislation. The majority of the nonconformances related to instances where wastewater from adjoining cottages was directed to a single septic tank/soak well installation. These landowners were advised to

effect plumbing amendments to achieve compliance with regulations, and that these amendments need to be completed prior to renewal of the lease agreements.

Since 1989, most household septic tank systems have been installed with either two leach drains or two sets of soak wells (DoH, 2011). These systems are known as alternating systems as they have a diverter box to enable flow to be switched from one leach drain or train of soak wells to the other, which allows the effluent disposal systems to be operated on a cyclic duty:resting regime. Such an operating regime is beneficial (if implemented), as periodic resting of leach drains and soak wells rejuvenates the soil's ability to receive effluent.

4.3 Current Regulatory Requirements

The current requirements for onsite treatment and disposal of wastewater in WA are set out in the Health (Treatment of Sewage and Disposal of Effluent and Liquid Waste) Regulations 1974. In addition to these regulations, the Draft Country Sewerage Policy (Draft from 22 September SOCWM Meeting Amended September 2003) and the Australian Standard AS1547:2012 (Onsite domestic wastewater management) provide guidance that is relevant to the onsite wastewater systems at Point Moore. Some of the key requirements relevant to the performance of the existing onsite systems at Point Moore (sand conditions assumed) are summarised in Table 2.

Item	Requirement/ Guideline	Regulation	Policy or Guideline	Reference ⁽¹⁾
Wastewater disposal area inundation/flooding risk	Probability of such to be less than once every 10 years.		~	В
Soak Well Sizing 2 bedroom dwelling 3 bedroom dwelling 	3 x soak wells ⁽²⁾	√ √		A
 Leach Drain Sizing 2 bedroom dwelling 	$2 \times 6 \text{ m}$ leach drains ⁽³⁾	✓		A
 3 bedroom dwelling 	2 x 8 m leach drains ⁽³⁾	✓		
Minimum Vertical Separation Distance to Groundwater ⁽⁴⁾	1.2 m ⁽⁵⁾ 0.6 - >1.5 m ⁽⁶⁾		\checkmark	B C

Table 2 Key regulatory requirements

Table Notes:

- 1. References:
 - A. Health Regulations 1974
 - B. Draft Country Sewerage Policy
 - C. AS1547:2012
- 2. Each soak well shall have a minimum diameter of 1.2 m and a minimum effective depth of 1.5 m, unless otherwise approved by DoH.
- 3. Assuming standard dimensions of 0.6 m effective depth and 0.4 m internal width.
- 4. Minimum vertical distance between invert of septic tank receptacle (in this case soak well or leach drain) and the highest seasonal water table.
- 5. The policy states that "for existing developed areas or infill areas a depth to highest known groundwater level may be a minimum of 1.2 m from ground level".
- 6. The recommended separation distance is a function of the groundwater pollution hazard

Available as-constructed records (Section 4.2) indicate that existing effluent disposal systems are under-sized with respect to current requirements (Table 2).

In addition to the 'performance requirements' detailed in Table 2, current regulations and guidelines include minimum horizontal clearances that must (unless otherwise approved) be maintained between septic tanks/soak wells//leach drains and buildings, property boundaries etc. The purpose of these clearances is to ensure that septic tanks/leach drains/soak wells can be maintained or replaced without affecting the stability or structural integrity of nearby buildings and the like, and that loadings from buildings do not affect the structural integrity of the wastewater infrastructure. This is further discussed in Section 5.3.

4.4 Estimated Vertical Separation Distance to Groundwater

As shown in Figure 1, although there are some small areas of higher and lower land, the natural surface level across most of the study area is approximately RL 2.0 m. Adopting RL 2.0 m as the typical natural surface level on the residential lots and assuming that the upstream end of the drainage pipework is installed at a depth (to invert) of 0.5 m, and that the typical length and grade of the drain from this point to the septic tank is 10-20 m and 1 in 60, respectively, the typical invert level of the septic tank inlet pipes is estimated to be approximately 0.7-0.9 m below ground level, or RL 1.1 m to RL 1.3 m. Adopting this level, and assuming that soak wells and leach drains have effective depths of 1.5 m and 0.5 m, respectively, the typical invert level of the soak wells are estimated to be approximately RL -0.2 m to RL -0.4 m, and the typical invert level of the leach drains are estimated to be approximately RL 0.6 m to RL 0.8 m.

Noting that the groundwater level currently ranges from approximately RL 0.2-0.5 m (refer to Section 6.3.5), the vertical separation distance between the invert of the leach drains and the water table is estimated to be approximately 0.1 to 0.6 m, and the invert of the soak wells (assuming 1.5 m effective depth as above) is likely to be below water table.

With reference to the current guidelines summarised in Table 2, it is evident that most of the soak wells and many of the leach drains in the study area are unlikely to comply with current vertical separation distance guidelines.

5. Property Assessments

5.1 General

SCP completed field investigations over the week commencing 17 October 2016 to assess the existing wastewater treatment and disposal systems at seventeen properties at Point Moore selected by CGG. Of these fifteen properties had non-invasive investigations undertaken, and two properties had more detailed invasive investigations undertaken.

The scope for the investigations, and the main findings from the investigations, are summarised next. Further detail is provided in SCP's full report.

5.2 Scope of Investigations

The scope of the investigations was to:

- Review available as-constructed information on the existing wastewater systems;
- Liaise with owner/occupiers to gain access to works area, and obtain general advice relevant to this study (e.g. comment on system performance, recent upgrade/maintenance works etc.);
- Take photos of entire property, and measure exterior of house;
- Run a drain camera down the drain (where possible);
- Locate septic tanks by visual inspection or probing, determine if single or dual septic systems, and mark on plan;
- Locate soakwell/leach drains by visual inspection or probing, and mark on plan; and
- Excavate & take photos of the septic tanks and leach drains/soak wells, and then reinstate properties to similar condition to how they were found for the two properties selected for invasive investigations.

5.3 Assessment Findings and Conclusions

The key findings from the investigations were:

- All properties where septic tanks were located had two septic tanks (first tank 1.5 m diameter, second tank 1.2 m diameter). Whilst no markings or stamps were visible stating that the tanks are DoH approved products, they appeared to be proprietary products.
- None of the located septic tanks met all of the standard horizontal separation distance requirements detailed in current regulations. Specifically, the clearance between the tanks (required minimum clearance = 1.2 m), the separation distance between the tanks and buildings (1.8 m required) or property boundaries (1.8 m required) did not comply with current requirements. The same issue applies for the located leach drains.
- Without excavation, the leach drains were difficult to locate. Of those that were located, a number were in poor condition, with some having at least several collapsed segments.
- No twin (alternating) leach drain systems with diverter valves were located, the longest leach drain was 8 m in length, and it was not possible to determine if they are DoH approved products.
- Soil types encountered during the investigations were typically fine grained sands below a topsoil layer, and are expected to have good drainage properties.

Based on the above findings SCP concluded that:

- 1. For the sites where the septic tanks and leach drains were located, none of the homes inspected had onsite systems that comply with relevant Australian Standards or the current DoH regulations.
- 2. Because of the small block sizes at Point Moore, it is not possible to install septic tank and leach drain systems that are fully compliant with current DoH regulations on many of the properties.

6. Soil and Groundwater Investigations

6.1 Guideline Framework for Contamination Assessment

This section of the report summarises the assessment criteria adopted to evaluate the human health and ecological risks posed by the existing septic systems within the study area. Consideration has been given to potential off-site impacts.

The legislation and guidelines that outline the appropriate framework for the investigation include the:

- Contaminated Sites Act 2003 and Contaminated Sites Regulations 2006;
- DER Contaminated Sites Management Series guidelines;
- DER Assessment and management of contaminated sites guideline (DER 2014); and
- National Environmental Protection Council (1999), National Environment Protection (Assessment of Site Contamination) Amendment Measure 2013 (No. 1).

The overarching reference in this assessment was the National Environment Protection (Assessment of Site Contamination) Measure 1999, as amended by the *National Environment Protection (Assessment of Site Contamination) Amendment Measure 2013 (No. 1)*, herein referred to as the NEPM. NEPM Schedule B1 "Guidelines on investigation levels for soil and groundwater" contains investigation and screening levels suitable for the assessment of the chemicals of potential concern in soil and groundwater at the site.

As defined in the NEPM, investigation levels are the concentrations of a contaminant above which further appropriate investigation and evaluation will be required.

6.1.1 Soil assessment criteria

Ecological Assessment Criteria

As defined in the NEPM (NEPC 1999), ecological investigation levels (EILs) have been developed for selected organic substances and are applicable for assessing risk to terrestrial ecosystems. EILs depend on specific soil physicochemical properties and land use scenarios and generally apply to the top 2 metres of soil. The NEPM advises that in arid regions, where the predominant species may have greater root penetration depth, specific consideration may result in their application to 3 m depth. Two metres depth is considered appropriate in this case for the site circumstances.

The following assessment criteria was adopted for a pragmatic consideration of risk to the environment:

• NEPC (2013) EIL for Public Open Space.

6.1.2 Groundwater Assessment Guidelines

The DER has provided criteria for use in the assessment of water (DEC, 2010), which are based upon the "Australian and New Zealand Guidelines for Fresh and Marine Water Quality" (ANZECC, 2000), the "Australian Drinking Water Guidelines" (NHMRC, 2004) and the "Contaminated study areas Reporting Guideline for Chemicals in Groundwater" (DoH, 2006).

Groundwater is likely to discharge into the Indian Ocean, which is located immediately adjacent to Point Moore. This part of the Indian Ocean is generally used for swimming, surfing, windsurfing, kite surfing, diving, fishing and boating.

No abstraction of groundwater takes place at Point Moore (refer to Section 2.3.2). The legislation and guidelines that outline the appropriate framework for the groundwater investigations is summarised next.

Ecological Assessment Criteria

A search was undertaken on the Bureau of Meteorology's (BoM) (2016) Atlas of Groundwater Dependent Ecosystems database. The BoM (2016) reported one groundwater dependent ecosystem reliant on surface expression of groundwater in the vicinity of the study area, which has a low potential for groundwater interaction. This GDE type is *Acacia ligulata*, an open scrub that is situated to north of the study area and is unlikely to interact with groundwater from the study area, and therefore is not considered further in the context of potential impacts from existing Point Moore wastewater practices.

The study area's groundwater is anticipated to flow towards the Indian Ocean. To assess the potential risk to the coastal marine environment and human health of coastal users, the ANZECC (2000) marine inshore default trigger values for South West Australia for physical and chemical stressors and primary recreational contact were adopted, respectively.

Health Assessment Criteria

As noted previously, the groundwater at Point Moore is not used for drinking water or irrigation in the study area, rather all such needs are met with scheme water. The groundwater beneath the study area flows towards the Indian Ocean, where a number of recreational activities are carried out. As a conservative measure the ANZECC (2000) guideline values for primary contact recreation (microbiological water quality limits only) were adopted.

6.2 Methodology

6.2.1 Field Work Preparation

Monitoring Bore Locations

Water Corporation and Telstra assets were identified in the vicinity of the original proposed location for MW1. Consequently, MW1 was relocated beyond these assets and to the western side of Marine Terrace. The rationale for the groundwater monitoring well locations is provided in Table 3, and the GMWs locations are depicted in Figure 11.

Well ID	Location	Rationale
MW1	West side of Marine Terrace, south of Pages Beach carpark	Down-gradient of existing residential properties targeting groundwater flowing in a westerly direction towards the Indian Ocean.
MW2	South side of Captains Crescent, opposite of Helm Way	Down-gradient and south of all residential properties, intersecting groundwater flowing in a southerly direction towards the Indian Ocean.
MW3	Corner of Sextant and Astrolab Lane.	Down-gradient and east of all residential properties, intersecting groundwater flowing in a easterly direction towards the Indian Ocean
SHP8 (previously installed by CGG)	Coxswain Park, situated along Coxswain Crescent	Down-gradient and south of Belair Lifestyle Village, Belair Gardens Caravan Park and small number of residential properties. Up-gradient from older residential properties.

Table 3 Rationale of GMW Locations

6.2.2 Soil Investigation

During the drilling program soils were logged and representative soil samples were collected during the installation of the new GMWs. The soil sampling activities were undertaken with reference to section 7, Schedule B2 of the *NEPM*, and Australian Standard AS 4482.1-2005 *Guide to the Sampling and Investigation of Potentially Contaminated Soil* (Standards, Australia, 2005).

Soil Sampling Methodology

To enable soil sample collection during the drilling program the drilling contractor utilised an air core drilling method for GMW installation. The following was undertaken at the GMW locations:

- Soil samples were collected at 0.5 metre intervals;
- Field observations, including olfactory and visual inspection of the soil, soil lithology, samples, QA samples were recorded on field sheets.

All soil samples were visually inspected and all field observations and subsurface conditions were recorded on field lithological logs. Soil from the borehole was obtained from a collection point on the drill rig and placed on clean sheeting, and samples were subsequently collected from this material.

Soil samples were placed into laboratory prepared containers provided by the primary laboratory and filled to the top to eliminate headspace. Each sample was identified by means of a label showing sample location, depth, date and job number. All soil samples were also identified by the depth at which they have been collected (MW1_0.5). The samples were then transferred to a chilled esky for sample preservation prior to and during shipment to the testing laboratory.

Sample details were entered onto a chain of custody form that accompanied the samples to the laboratory. All samples were transported and handled following chain of custody procedures. A chain of custody form was used for every batch of samples submitted to the laboratory. Delivery of the samples to the laboratory complied with analytical extraction holding times.

All field work was undertaken in the presence of an environmental scientist trained in sampling of contaminated sites. The environmental scientist was present during the services location, supervised the sub-contractors, sampled and completed lithological logs of the soil profile. Field activities were conducted in accordance with accepted industry protocols for environmental sampling.

The study area conditions were photographed during soil sampling and incorporated into the report.

6.2.3 Groundwater Investigation

Groundwater wells were sampled with reference to section 8, Schedule B2 of the *NEPM* and Australian Standard AS5667-11-1998 *Guidance on sampling of groundwaters* (Standards Australia, 1998).

Groundwater well locations were determined based on anticipated hydrogeological characteristics (from the desktop review) including reported depths to groundwater and apparent groundwater flow direction. The locations were confirmed following consultation with the CGG.

Three GMWs were installed in the study area on 16 August 2016 The locations of these GMWs were selected to complement the existing GMW SHP8 located in Coxswain Park (refer to Table 3 for rationale of GMW locations).

Groundwater Monitoring Well Installation

Monitoring wells were drilled to a minimum depth of three metres below the groundwater table and were installed with the following general characteristics:

- 50 mm polyvinyl chloride (PVC) class 18 blank and screened casings;
- Screened casing comprised of machine slotted apertures of approximately 0.5 mm;
- Screened casing extended the full length of the GMWs as the groundwater levels are influenced by tides and may significantly fluctuate. A screen sock extended the full length of the screen casing, to prevent the infiltration of gravel into the GMW;
- Blank and screened PVC casing was attached to each other using flush mounted factorythreaded joints;
- Primary filter pack material used was chemically inert, well rounded material with a high coefficient of uniformity and extend from the base of the borehole annulus to the ground surface;
- Bentonite/cement grout was placed at the ground surface to minimise surface water ingress; and
- Monitoring wells were equipped with flush mounted head-works to protect the wells.

The GMWs logs are provided in Appendix C.

Groundwater Monitoring Well Development

The groundwater monitoring wells were developed immediately following installation with air lifting techniques by the drilling contractor. Monitoring wells were developed until:

- No further noticeable sand or silt is recovered.
- The water was relatively clear when pumped from the well.
- A minimum of four well volumes removed.

Monitoring well development optimises the well efficiency, specific capacity, stabilisation of aquifer material and control of suspended solids.

The newly installed groundwater monitoring wells were allowed to stabilise for a minimum of seven days prior to purging and sampling. The groundwater field sheets are provided in Appendix D.

Groundwater Loggers

Groundwater loggers were deployed from the 16 August 2016 until 30 September 2016 to gauge the depth to groundwater and how this depth varies over time. The loggers were deployed in three GMWs across the network, i.e. MW1, MW2 and MW3. The findings are provided in Section 6.3.5.

Groundwater Sampling Program

The 2016 groundwater sampling program comprised three monitoring rounds across the current groundwater monitoring network within Point Moore.

- GMWs were gauged to determine the standing water level.
- Following gauging, GMWs were purged and samples collected with a low flow pump (less than 1 L/minute) (with reference to the methodology outlined in USEPA 540/5-95/504).

The wells were purged and sampled with low-density poly-ethylene tubing coupled to a Sample-Pro Micro purge ('low flow') pump system. The low flow pump provided an appropriate method for collection of representative samples for the required analytes and is recognised as best practice for groundwater sampling.

Field parameters measured during purging included temperature, pH, conductivity, dissolved oxygen and redox potential. Field parameters were recorded on field data sheets. Wells were purged until field parameters stabilised.

The variance associated with the above mentioned parameters required to establish chemical stabilisation are as follows:

- pH: 0.1 unit
- Temperature: 0.2°C
- Eh (ORP): 10%
- DO: 10%
- SC: 10%

Samples at each bore were collected with new disposable gloves and placed directly into laboratory provided sample bottles. Each sample was identified by a label with the sample location, date, job number and depth. The samples were placed on ice in an insulated container, and kept cold during storage and transportation to the laboratory.

Decontamination Procedure

To ensure samples were collected without the potential presence of cross contamination, all reusable sampling equipment was decontaminated in accordance with the procedure and methods described in AS 4482.1 - 2005. In addition, all samples were handled by field staff with disposable nitrile gloves, which were replaced between each GMW.

Groundwater sampling equipment was decontaminated as follows:

- Washed and scrubbed in tap water;
- Washed and scrubbed in laboratory grade detergent (Decon90); and
- Rinsed in distilled or deionised (Grade 3) water.

Survey

Following installation, the monitoring wells were accurately surveyed to AHD and Geocentric Datum of Australia (GDA94), Map Grid Australia (MGA), Zone 50. As the reduced levels of the GMWs (top of casing levels) were accurately determined, measuring the depth to groundwater in the wells enabled groundwater flow directions beneath the study area to be determined.

6.2.4 Groundwater Laboratory Analysis Program

During each groundwater monitoring event (GME) four primary groundwater samples were submitted for analysis to National Association of Testing Authorities (NATA) accredited laboratories including the Australian Laboratory Services (ALS), SGS Australia (SGS) and mpl Laboratories (mpl). Analyses were selected to ascertain data pertaining the current status of groundwater contamination at Point Moore.

The four primary groundwater samples collected during each GME were submitted for laboratory determination of the following parameters:

- Microbiological quality: Total coliforms, *E. coli*, faecal coliforms and total plate count;
- Nutrients levels: Total phosphorus, ammonia-N, nitrate+nitrite-N, TKN and total nitrogen; and
- Other: Biological oxygen demand and total dissolved solids.

6.3 Results

6.3.1 Soil Investigation

Field Observations

Based on the information gathered during the establishment of the GMWs, the underlying geology is calcareous medium to coarse grained sand. A summary of the geology encountered at MW1 is provided in Table 4, which is typical of the three new monitoring wells. The field logs for all new GMW are provided in Appendix C and the relevant photographs from the works are provided in Appendix I.

Table 4 Summary of Geology Encountered at MW1

Depth (m bgl)	Lithological Description
0 – 2.0	Fine sand, brown with minor loam
2.0 - 6.0	Calcareous coarse sand, grey to white, minor quartz

During the field logging of soil samples (i.e. visually inspection and sampling), there were no olfactory signs (i.e. odours or staining) of contamination that warranted further investigation, so no laboratory analyses of soil samples were undertaken.

6.3.2 Groundwater Investigation: August 2016

The first round of groundwater monitoring at MW1, MW2, MW3 and SHP8 occurred on 24 August 2016.

Field Observations

GMW locations are illustrated in Figure 11, and groundwater monitoring field forms are provided in Appendix D.

To determine groundwater flow direction, groundwater elevations from the newly installed GMWs and the existing GMW SHP8 were calculated from depth to groundwater and survey data. Survey data, depth to groundwater and groundwater elevations are presented in Table 5. Groundwater contours derived from this data are depicted in Figure 11.

ID	Easting	Northing	TOC elevation (m AHD)	Depth to water (m bTOC)	Groundwater Elevation (m AHD)
MW1	263938	6814127	2.608	2.36	0.248
MW2	263709	6813860	2.151	1.78	0.371
MW3	263934	6813674	3.009	2.67	0.339
SHP8	263885	6813915	2.427	2.1	0.327

Table 5 Groundwater Well Survey Data

Groundwater Quality Field Parameters

During purging of the groundwater wells, groundwater quality field parameters were measured with a multi-parameter water quality meter (temperature, pH, specific conductivity (SC), salinity, dissolved oxygen (DO), and oxidation-reduction potential (ORP)). This equipment was calibrated by the equipment supplier (Thermofisher Scientific) prior to use on-site and did not require adjusting for redox measurements. A calibration certificate for the water quality meter is presented in Appendix H. Completed groundwater field sheets including a summary of groundwater quality field parameters can be found in Appendix D.

Groundwater samples were described as follows:

- MW1: Colourless, minor particulate, slight H₂S odour;
- MW2: Colourless, slight H₂S odour;
- MW3: Colourless, minor particulates, slight H₂S odour; and
- SHP8: Colourless, slight H₂S odour.

Field parameters indicate that groundwater beneath the study area is generally brackish to saline, and is alkaline. Groundwater is aerobic and with a reducing condition. No indicators of contamination were noted during the sampling with the exception of organic odour noted in the wells.

Groundwater Analytical Results

The groundwater analytical results are tabulated in Table G1, Appendix G and the Laboratory certificates of analysis are presented in Appendix F. Review of quality control and assurance data completed as part of this GME is provided in Appendix E. Exceedances of the adopted criteria during this monitoring round are discussed next.

Reported Exceedances

The following exceedances were reported during the August 2016 GME:

Ammonia exceeded the adopted EIL (0.91 mg/L) at MW1 (2.68 mg/L);

- Total nitrogen and total phosphorus exceeds the ANZECC (2000) marine inshore default trigger values in all GMWs;
- Nitrate was reported above the ANZECC (2000) Marine Inshore trigger value in MW1 and MW2; and
- Faecal coliforms and *E. coli* results were reported to have exceeded the ANZECC (2000) guideline values for primary contact recreation (150 cfu/100mL) at MW1, MW2 and MW3.

Whilst not an exceedance as ANZECC (2000) does not include guideline values for total coliforms, coliform results (coliforms by membrane filtration) reported in MW1 (250,000 cfu/100 mL) were two orders of magnitude higher than results for up-gradient GMWs including MW2 (1000 cfu/100mL), MW3 (3000 cfu/100mL) and SHP8 (200 cfu/100mL).

6.3.3 Groundwater Investigation: September 2016

The second round of groundwater monitoring at MW1, MW2, MW3 and SHP8 occurred on 22 September 2016.

Field Observations

The depth to groundwater and groundwater elevations are shown in Table 6.

ID	Easting	Northing	TOC elevation (m AHD)	Depth to water (m bTOC)	Groundwater Elevation (m AHD)
MW1	263938	6814127	2.608	2.43	0.178
MW2	263709	6813860	2.151	1.83	0.321
MW3	263934	6813674	3.009	2.68	0.329
SHP8	263885	6813915	2.427	2.08	0.347

Table 6 Groundwater Well Survey Data

Groundwater Quality Field Parameters

During purging of the groundwater wells, groundwater quality field parameters were measured with a multi-parameter water quality meter. A calibration certificate for the water quality meter is presented in Appendix H. The completed groundwater field sheets are reproduced in Appendix D.

Groundwater samples were described as follows:

- MW1: Colourless, particulate, H₂S odour;
- MW2: Colourless, H₂S odour;
- MW3: Colourless, particulates, H₂S odour; and
- SHP8: Colourless, slight H₂S odour.

Field parameters indicate that groundwater beneath the study area is generally brackish to saline, and is acidic to slightly alkaline. The low dissolved oxygen concentrations are indicative of an anaerobic environment with reducing conditions. A hydrogen sulphide odour and high level of particulates was noted during the sampling at all GMW locations. The reported temperatures in the GMWs are likely due to a faulty temperature sensor.

Groundwater Analytical Results

The groundwater analytical results are tabulated in Table G1, Appendix G and the Laboratory certificates of analysis are presented in Appendix F. Review of quality control and assurance data completed as part of this GME is provided in Appendix E. Exceedances of the adopted criteria are discussed next.

Reported Exceedances

The following exceedances were reported during the 22 September 2016 GME:

- Ammonia exceeded the ANZECC (2000) Marine Water 95% (0.91 mg/L) at MW1 and SHP8 (6.28 mg/L and 2.54 mg/L, respectively);
- Total nitrogen and total phosphorus exceeds the ANZECC (2000) default marine inshore trigger values for these parameters in all GMWs;
- Nitrate was reported above the ANZECC (2000) Marine Inshore trigger value in MW1 and 2; and
- Whilst faecal coliforms and *E. coli* were reported above the LOR at two GMWs, i.e. MW2 (1 cfu/100 mL) and MW3 (1 cfu/100 mL), no exceedances occurred.

6.3.4 Groundwater Investigation: October 2016

Field Observations

The third round of groundwater monitoring at MW1, MW2, MW3 and SHP8 occurred on 18 October 2016. The depth to groundwater and groundwater elevations are presented in Table 7.

ID	Easting	Northing	TOC elevation (m AHD)	Depth to water (m bTOC)	Groundwater Elevation (m AHD)
MW1	263938	6814127	2.608	2.41	0.198
MW2	263709	6813860	2.151	1.90	0.251
MW3	263934	6813674	3.009	2.71	0.299
SHP8	263885	6813915	2.427	2.17	0.257

Table 7 Groundwater Well Survey Data

Groundwater Quality Field Parameters

During purging of the groundwater wells, groundwater quality field parameters were measured with a multi-parameter water quality meter. A calibration certificate for the water quality meter is presented in Appendix H. The completed groundwater field sheets are reproduced in Appendix D.
Groundwater samples were described as follows:

- MW1: Colourless, minor suspended solids, H₂S odour;
- MW2: Colourless, H₂S odour, minor suspended solids;
- MW3: Colourless, minor suspended solids, no odour; and
- SHP8: Colourless, slight H₂S odour, minor suspended solids.

Field parameters indicate that groundwater beneath the study area is generally brackish to saline, and is alkaline. Though groundwater had reasonable oxygen levels, it also had reducing conditions. There were no indicators of contamination during sampling with the exception of organic odours. The temperature in MW1 was 0.6 °C higher than other GMWs.

Groundwater Analytical Results

The groundwater analytical results are tabulated in Table G1, Appendix G and the laboratory certificates of analysis are provided in Appendix F. The quality control and assurance report for this GME is provided in Appendix E. Exceedances of the adopted criteria are discussed next.

Reported Exceedances

The following exceedances were reported during the October 2016 GME:

- Ammonia exceeded the ANZECC (2000) Marine Water 95% (0.91 mg/L) at MW1 and MW2 (3.95 mg/L and 1.48 mg/L, respectively);
- Total nitrogen and total phosphorus exceeds the ANZECC (2000) Marine Inshore trigger values for these parameters in all GMWs;
- Nitrate was reported above the ANZECC (2000) Marine Inshore trigger value in MW1 and 2; and
- Whilst not an exceedance, faecal coliforms and *E. coli* were reported above the LOR at all GMWs (all samples reported a count of 1 cfu/mL).

6.3.5 Groundwater Data Loggers

Data loggers were deployed at MW1, MW2 and MW3 for 38 days (16 August to 22 September). This groundwater level data is presented in Figure 10 with the following key findings :

- Water levels of the three MGWs co-varied throughout the 38 day deployment. Hence, groundwater flow directions are seemingly uni-directional as depicted below in Figure 11.
- Groundwater levels across the groundwater network are influenced by coastal tidal variations to varying degrees as follows:
 - MW2 seemingly undergoes daily tidal-induced level variations of 0.05 m during spring tides and 0.01-0.02 m during neap tides. This suggests that connectivity to the marine waters is relatively rapid with concomitant relatively short travel times from the GMW to the marine environment;
 - In contrast, MW3 has much more muted tidal-induced level variations of 0.01-0.02 m during spring tides and no level variations during neap tides. This suggests that connectivity to the marine waters is substantially slower than MW2 with concomitant relatively long travel times from this GMW to the marine environment; and
 - Over the short measurement record of MW1 (only 16-24 August due to subsequent instrument failure), this GWM responds to tidal variations more akin to MW3 in a muted manner.



Figure 10 Groundwater Levels Logger Data for MW1, MW2 and MW3

Calculated Groundwater Flow

The groundwater contours depicted in Figure 11 indicate that groundwater beneath the study area flows towards the Indian Ocean, with groundwater in northern, western and southern portions of the study area flowing to the north, west and south, respectively. The lowest groundwater levels were reported in MW1.



Figure 11 Groundwater Contours, August 2016

6.4 Review of Laboratory Groundwater Quality Data

6.4.1 Summary of Data

A statistical summary of laboratory groundwater quality data obtained over the course of the monitoring program completed as part of this study is presented in Table 8. Trends apparent in the data are discussed in Section 6.4.2.

Table 8 Summary of Laboratory Groundwater Quality Data

		Water Quality Guidelines			Statistical Summary					
						Number of		Concen	tration	
		ANZECC	ANZECC	ANZECC 2000	ANZECC 2000	Results	Minimum	Maximum	Mean	Median
Parameter	Unit	2000 MW	2000 Marine	Primary	Secondary					
		95%	Inshore	Contact	Contact					
				Recreation	Recreation					
Ammonia as N	mg/L	0.91	-	-	-	12	0.13	6.28	1.7	0.675
BOD	mg/L	-	-	-	-	12	3	57	15	5.5
Coliform	cfu/100 mL	-	-	-	-	12	<1	250,000	21,185	5.5
Plate Count (36°C)	CFU/mL	-	-	-	-	12	17	68,000	13,140	4,950
Plate Count (22°C)	CFU/mL	-	-	-	-	12	12	70,000	13,483	3,300
Nitrate (as N)	mg/L	-	0.05	-	-	12	<0.01	13.8	2.6	0.04
Nitrite + Nitrate (as N)	mg/L	-	-	-	-	12	<0.01	14	2.7	0.04
Nitrite (as N)	mg/L	-	-	-	-	12	<0.01	1.85	0.2	0.005
Faecal Coliforms	CFU/100mL	-	-	150	1000	12	<1	400	92	1
E. Coli	cfu/100 mL	-	-	150	1000	12	<1	400	92	1
Total Dissolved Solids	mg/L	-	-			4	1,090	8,890	3,695	2,400
Total Dissolved Solids (Filtered)	mg/L	-	-	-	-	8	1,320	22,300	6,766	2,205
Total Kjeldahl Nitrogen (as N)	mg/L	-	-	-	-	12	0.6	6.7	2.5	2.3
Nitrogen (Total)	mg/L	-	0.23	-	-	12	0.6	17.1	5.2	3.25
Phosphorus (Total)	mg/L	-	0.005	-	-	12	0.06	0.81	0.27	0.245

Legend

ANZECC 2000 MW 95% ANZECC 2000 Marine Inshore ANZECC 2000 Primary Contact Recreation ANZECC 2000 Secondary Contact Recreation Indicates a level is equal to or above the ANZECC (2000) marine waters 95% species protection trigger value Indicates a level is equal to or above the ANZECC (2000) marine inshore trigger value

Indicates a level is equal to or above the ANZECC (2000) guideline value for primary contact recreation (e.g. swimming) Indicates a level is equal to or above the ANZECC (2000) guideline value for secondary contact recreation (e.g. boating, fishing)

6.4.2 Trends

GHD reviewed laboratory water quality data from the three surveys and the following trends were determined:

- Faecal coliforms and *E. coli* results were reported to have exceeded the ANZECC (2000) guideline valves for primary contact recreation (150 cfu/100mL) at MW1, MW2 and MW3 during the August GME. Follow up GMEs September and October reported faecal coliforms and *E. coli* results had declined (1 cfu/100 mL) across the groundwater network, with no exceedances recorded during these two rounds;
- Coliforms (coliform by membrane filter) were highest in the August survey (MW1 -250,000 CFU/mL, MW2 1000 CFU/mL, MW3 3000 CFU/mL and SHP8 200 CFU/mL), whereas the September survey had coliforms ranging from below the LOR (<1) to 10 CFU/mL (MW2, MW3); and the September GME reported that pH levels across the monitoring well network had largely declined from the August GME, however, in the October GME, pH levels inclined;
- In each monitoring round ammonia was reported above the adopted EIL in at least one GMW. Ammonia in MW1 increased from 2.68 mg/L in August to 6.28 mg/L in September, and subsequently declined in October. Ammonia in MW3 and SHP8 also exhibited a similar trend. A single exceedance of the adopted Ammonia EIL was reported at SHP8 during the September GME. Ammonia in MW2 declined between August and September then increased in October. There was a single exceedance of the adopted EIL in October at MW2;
- Nitrate was above the ANZECC (2000) default marine inshore trigger value at MW1 and MW2 for all three surveys. MW1 exhibited an increasing trend over the three surveys, while MW2 exhibited a decreasing trend; and
- Total nitrogen and total phosphorus were reported to have exceeded the ANZECC (2000) default marine inshore trigger values in the three GMEs and at all GMWs. Concentrations were relatively consistent in all GMEs.

With respect to the nutrient exceedances discussed above, Ammonia is the only nutrient species considered a toxicant in marine waters, and as a nutrient may stimulate algal growth and contribute toward eutrophication (Trefey et. Al, 2006). The other nutrients discussed above, whilst not directly toxic to marine life, may also act to stimulate plant growth and contribute towards eutrophication. Phosphorus will adsorb readily onto some mineral phases in soils, and is heavily retarded in some soil types such as the calcareous sands and limestones in the Point Moore area.

The cause of the significantly higher faecal coliform and *E. coli* levels recorded during the August 2016 GME is not clear. Whilst this GME was preceded by more rainfall than was the case for the other GMEs (5.6 mm was recorded at the Bureau of Meteorology's Geraldton Town weather station two days prior to the September GME, 1.6 mm was recorded three days prior to the September 2016 GME and no rainfall was recorded eight days prior to the October GME), it is unlikely that the increased levels would have been caused by the resultant higher rate of infiltration to groundwater. It is also considered unlikely that tidal variations caused this.

7. Discussion

7.1 Asset Condition

As many of the existing onsite systems were installed between 30 and 50 years ago, a significant number of the septic tanks and associated leach drains/soak wells are likely to have reached the end of their serviceable life. Information from the field investigation program verified this, with a number of the septic tanks observed to be in a "fair" condition, one septic tank system with covers that were on the verge of collapse, and several leach drains partially collapsed. During the field investigation there was not any evidence of saturated ground conditions above the leach drains.

An indicative estimate of the cost to construct new septic tanks and 2 x 6 m long leach drains sized for a two-bedroom dwelling, inclusive of the cost of the required plumbing modifications, is \$10,000 per property.

7.2 Compliance with Current Standards

The properties in the study area are significantly smaller than the minimum lot size currently permitted for onsite wastewater disposal (typically 2,000 m²). In addition, based on review of available information and the field investigation for this study:

- Many of the effluent disposal systems are undersized by current standards and are not configured as alternating systems;
- Many of the septic tanks and leach drains/soak wells do not comply with current horizontal setback requirements; and
- The inverts of the leach drains are estimated to be approximately 0.1 to 0.6 m above the water table, and the invert of the soak wells (assuming 1.5 m effective depth as above) is likely to be below water table (nil separation), hence current guidelines on the required vertical separation distance are not met. As natural purification processes are most effective in the aerobic unsaturated zone rather than in the soils below water table (Washington State Department of Health, 1990), the quality of effluent infiltrating to groundwater will be worse than would be the case if there was greater vertical separation between the leach drains/soak wells and groundwater. This situation will worsen as groundwater levels rise in the future (refer to discussion below).

7.3 Health Risks

7.3.1 Current Situation

Local groundwater is not used for irrigation or any other purpose, but does discharge to the nearby ocean. There is also a potential human health risk with regards to residents or others that could come into direct contact with groundwater when undertaking land-based activities. Conservative water quality assessment criteria based on default guideline values for primary contact recreation were used to assess the health risks posed by contact with groundwater. Elevated microorganism levels in sampled groundwater indicated wastewater-induced contamination above the adopted assessment criteria for one of the three monitoring rounds.

The current health risks posted by contact with groundwater are discussed below.

Risks Associated with Recreational Activities in Ocean

Based on studies done elsewhere at locations with similar hydrogeological conditions, e.g. detailed investigations completed at Halls Head (Mandurah) as reported in Toze et.al. (2010),

groundwater quality will improve significantly over time as it flows (sub-surface) to the ocean. With respect to die-off of pathogenic microorganisms:

- The travel time for Point Moore groundwater to reach the inshore marine waters off the coast is uncertain. The groundwater level measurements and seemingly high connectivity to the ocean via tidal variations at GMW MW2 suggests that there are preferential flow paths to rapidly transport groundwater to the ocean for some portions of the study area, with travel times perhaps being in the order of days. In contrast, the muted tidal-induced variations at GMWs MW1 and MW3 suggest that transport times of groundwater to the ocean are likely to be substantially longer. As a conservative measure it has been assumed that the range of transport times from leach drains and soak wells across the Point Moore study area ranges from 2 days (e.g. MW2) to 20 days (e.g. perhaps from the centre of Point Moore). These travel times are likely to be more rapid than actually occurs, which is conservative as it reduces the time for die-off of bacteria and other pathogenic microorganisms (e.g. viruses and protozoa). For example, on the basis of other studies with similar hydrogeological settings, the minimum groundwater travel time from the Point Moore residential properties to the ocean would be estimated to be in the order of 40 days;
- The aquifer residence time for 1 log removal (90% reduction) of bacteria is approximately 2 days (Toze et al, 2010). In the case of bacteria, for every 2 days of aquifer residence time the level of bacteria would reduce by 90%;
- Hence, on the basis of the conservative superficial aquifer residence times adopted in this study (2 days to 20 days), the bacteria will be expected to undergo, from onsite effluent disposal systems to the point of groundwater discharge to the ocean, log reductions ranging between a 1 log (90% reduction in 2 days) and a 15 log (1x10⁻¹³ % remaining) reduction. In short, for those septic systems that are located in preferential flow paths with relatively rapid transport times to the ocean, there is a risk of relatively direct pathways to deliver to the ocean groundwater with elevated pathogens that may pose a risk to human health.

Clearly, one ameliorating factor of the risk of high pathogen levels introduced into the marine environment via preferential groundwater pathways are the high levels of dilution that would typically occur where the groundwater discharges into the ocean. Hence, for the most part, it is considered unlikely that elevated levels of pathogens in groundwater flowing from the study area would pose a significant health risk in terms of primary contact recreation in the ocean near Point Moore. However, under conditions of calm winds and low wave climate, rates of dilution may be greatly reduced, thereby increasing the potential health risk. These conditions typically occur late in the bathing season from March-May.

Based on the above discussion it is not possible to discount the possibility that onsite disposal of effluent from the Point Moore residential properties is at least partly responsible for the observed seasonal spikes in *Enterococci* levels at the CGG's marine water quality monitoring sites near Point Moore (Section 2.4.2).

Other Risks

In addition to the health risk associated with recreational activities in the ocean near Point Moore, there is a risk that residents or others could come into direct contact with groundwater when completing excavation or trenching works, or undertaking dewatering operations, for:

- construction of foundations for new buildings or other structures,
- construction of installation of swimming pools;
- installation of new services, or maintenance of existing services; or

• landscaping/gardening projects.

The risk of such direct contact is a function of several factors, including the depth to groundwater. Whilst the depth to groundwater is currently approximately 1.5m or more across much of the study area, there are some areas (e.g. park north of Sailors Lane which includes land with a surface level below RL1.0m [Figure 1], some residential lots on the south side of Gunners Lane where the surface level is only marginally above RL1.0m [topographic data from CGG Inframaps]) where this depth may be in the order of 0.5m only. The risk of direct contact with groundwater is obviously greatest in such areas.

Whilst elevated faecal coliform and *E. coli* levels above adopted trigger values were only measured at the monitoring bores in one of the three monitoring rounds, it must be noted that testing for viruses and other pathogenic microorganisms (e.g. protozoa such as cryptosporidium and giardia) was not undertaken as part of this study, and that the rate of die-off of these other pathogens in groundwater is significantly slower than faecal coliforms, *E. coli* and other bacteria. In addition, the levels of pathogens in effluent infiltrating to groundwater is likely to be higher than is typically the case because of the limited (or non-existent in the case of soak wells) vertical separation distance between effluent disposal facilities and the water table (refer to Section 7.2). Given this, contact with groundwater in the study area is considered to present a health risk to residents and others in the study area. Whilst for works undertaken by CGG and service utilities it would be possible to mitigate these risks to an acceptable level through use of appropriate personal protective equipment and control measures, it would not be realistic to rely upon such measures to safeguard the health of local residents and others engaging in activities where they come into contact with groundwater.

7.3.2 Long term Future

In the long term, local groundwater levels will rise as sea levels rise. If sea levels rise by 0.9 m in 2110 as predicted, the groundwater levels at that time could be within approximately 0.6 m of the natural surface level at many properties. In some areas groundwater levels could rise above the natural surface level, in which case some areas could become permanently inundated with water containing elevated levels of pathogenic microorganisms.

As groundwater levels rise in the future the likelihood that residents or others could come into direct contact with groundwater will increase, which will in turn increase the public health risks posed by the onsite wastewater systems.

7.4 Environmental Risks

Conservative water quality assessment criteria based on inshore marine default trigger values for nutrients were adopted for the groundwater sampled from the GMWs (three rounds of sampling over two months). Nutrient levels in sampled groundwater indicated elevated wastewater-induced contamination above the adopted assessment criteria for all monitoring rounds.

As discussed in Section 7.3.1, high levels of dilution typically occur where groundwater discharges into the ocean. Given this dilution it is considered unlikely that elevated levels of nutrients in groundwater flowing from the study area are having any measurable impact on near shore marine ecosystems.

7.5 Long Term Wastewater Management

In the long term local groundwater levels will rise as sea levels rise, and the magnitude of the rise will severely constrain the potential to dispose of effluent generated in the study area using conventional onsite septic tank and leach drain/soak well systems. That is:

- If sea levels rise by 0.9 m in 2110 as predicted, the groundwater levels at that time could be within approximately 0.6 m of the natural surface level at many properties.
- If house and ground levels remain as-is, this groundwater level will be too high to enable the onsite disposal systems to function effectively, with potential for odour problems or saturated ground conditions in the vicinity of the onsite disposal systems, and an increased risk of residents or others coming into contact with effluent or contaminated groundwater.

Whilst alternative onsite systems may be able to be used in the long term, the small size of the lots would severely constrain the options are available for effluent disposal (e.g. aerobic treatment units with dedicated irrigation disposal areas could not be installed as insufficient area is available for the irrigation areas). Rather, if residential properties are to remain at Point Moore for the long term it is considered that a reticulated wastewater collection system would need to be installed that discharges wastewater to the Water Corporation's Geraldton wastewater scheme.

Design and construction of a reticulated wastewater collection system to serve properties in the study area would be constrained by the small size of the lots and limited room available to construct sewers along property rear and side boundaries, the relatively flat topography of the area and the shallow depth to groundwater.

An indicative estimate of the cost to design and construct a conventional reticulated gravity sewer type wastewater collection system to serve all properties in the study area, inclusive of the cost of the house connections, as well as pump stations and pressure mains to discharge this wastewater to the Geraldton wastewater system, is \$6M to \$10M. This estimate, which assumes that the collection and conveyance infrastructure is constructed to Water Corporation design standards, equates to a cost of approximately \$35,000 to \$55,000 per property. Additional costs may also apply if the Water Corporation's sewerage scheme needs to be upgraded to cope with the flows from Point Moore.

It would also be possible to serve the properties with a vacuum sewerage system, similar to that operated by the Geraldton Port Authority, or a pressure sewer type collection system. For both alternatives the required depth of the collection pipework would be significantly less than would be the case for a conventional gravity sewer type collection network, which is one of their main advantages. In the case of a vacuum sewerage scheme, wastewater from several properties would gravitate to local pits equipped with proprietary vacuum valves, and from these pits would be conveyed to a single vacuum pump station via vacuum sewers. Conventional wastewater pumps and a pressure main would then deliver the wastewater to the Water Corporation's wastewater scheme. In the case of a pressure sewer system, wastewater from each property would gravitate to small proprietary pump stations equipped with high-head pumps (one pump station per property), and these pumps stations would pump wastewater into a small-diameter interconnected pressure main network that conveys the wastewater direct to the Water Corporation's wastewater scheme, or more likely via a conventional wastewater pump station and pressure main. Whilst the capital cost of installing a vacuum sewerage system or pressure sewer type collection system to serve Point Moore residents may be significantly lower than the cost to install a conventional gravity sewer type collection system, ongoing operations and maintenance costs would be significantly higher for these alternative systems.

8. Conclusions

The overall objective of the study is to provide CGG and Point Moore stakeholders with data and documentation on the performance and compliance of the existing onsite wastewater treatment and disposal systems that assists CGG in the decision-making process relating to the future beyond the current lease expiry dates of 2025 and 2028 and potential obligations it has in relation to Point Moore lessees. Based on work completed in undertaking this study the following conclusions are made:

- 1. Observations made during the field investigation indicate that a significant number of the existing septic tanks and leach drains/soak wells are in a poor condition and require remedial works and in some cases replacement.
- 2. The properties in the study area are significantly smaller than the minimum lot size currently permitted for onsite wastewater disposal (typically 2,000 m²), and many of the onsite systems do not comply with current standards in a number of respects (sizing, configuration, horizontal setbacks, vertical separation distance to groundwater). For many properties it would not be possible to upgrade the existing onsite systems to meet current standards, or install alternative onsite systems that comply with current standards.
- 3. Though local groundwater is not used for irrigation or any other purpose, it does discharge to the nearby ocean, and residents or others could come into contact with groundwater when undertaking a range of land based sub-surface activities. In relation to public health risks:
 - The potential for contact with groundwater when undertaking land based sub-surface activities such as excavation or trenching works is considered to represent a significant public health risk to residents and others undertaking such activities in the study area. This risk will increase over time as groundwater levels increase as a direct consequence of sea level rise.
 - Whilst data from the monitoring program indicates that for some portions of the study area groundwater travel times to the ocean may be short due to preferential groundwater pathways, it is considered unlikely that elevated levels of pathogens in groundwater flowing from the study area would pose a significant health risk to persons engaging in primary contact recreation in the ocean near Point Moore given natural purification processes in the aquifer and the high levels of dilution that would typically occur where the groundwater discharges into the ocean. However, under conditions of calm winds and low wave climate, rates of dilution may be greatly reduced, thereby increasing the potential health risk. These conditions typically occur late in the bathing season from March-May.
 - It is not possible to discount the possibility that onsite disposal of effluent from the Point Moore residential properties is at least partly responsible for the observed seasonal spikes in *Enterococci* levels at the CGG's marine water quality monitoring sites near Point Moore.
- 4. In the long term local groundwater levels will rise as sea levels rise, and the magnitude of the rise will severely constrain the potential to dispose of wastewater generated in the study area with the existing conventional onsite septic tank and leach drain/soak well systems approach.
- 5. If residential properties are to remain at Point Moore for the long term it is considered that a reticulated wastewater collection system would need to be installed that discharges wastewater to the Water Corporation's Geraldton wastewater scheme. An indicative estimate of the cost to design and construct a conventional gravity sewer type wastewater

collection system to serve all properties in the study area, inclusive of the cost of the house connections, as well as the pump stations and pressure mains to discharge this wastewater to the Geraldton wastewater system, is \$6M to \$10M. This equates to a cost of approximately \$35,000 to \$55,000 per property, which is likely to be prohibitively expensive. Whilst alternative wastewater collection technologies exist that may be able to be implemented at a significantly lower capital cost, ongoing costs for these systems would be higher than the ongoing costs associated with a conventional gravity sewer type collection system.

6. Whilst nutrient levels in sampled groundwater indicated elevated wastewater-induced contamination above the adopted assessment criteria for all monitoring rounds, given the high levels of dilution typically occur where groundwater discharges into the ocean it is considered unlikely that elevated levels of nutrients in groundwater flowing from the study area are having any measurable impact on near shore marine ecosystems.

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Appendices

44 | GHD | Report for City of Greater Geraldton - Point Moore, 61/34772

Appendix A – Nearshore Sampling Tabulated Exceedances

Location (Date)	Enterococci (MPN/100mL)	Exceed - ANZECC (2000) Recreational Primary Contact – Lower Limit – 35 cfu/100 mL	Exceed – ANZECC (2000) Recreational Secondary Contact – Upper Limit – 100 cfu/100mL	Exceed –ANZECC (2000) Secondary Contact Lower Limit – 230 cfu/100mL	Exceed – ANZECC (2000) Secondary Contact – Upper Limit – 730 cfu/100mL
Point Moore (9/4/2013)	41	Yes	No	No	No
Pages Beach (9/4/2013)	73	Yes	No	No	No
Point Moore (8/1/2014)	54	Yes	No	No	No
Point Moore (15/1/2015)	84	Yes	No	No	No
Pages Beach (15/04/2015)	170	Yes	Yes	No	No
Pages Beach (12/03/2015)	230	Yes	Yes	Yes	No
Pages Beach (3/02/2016)	910	Yes	Yes	Yes	Yes
Pages Beach (19/04/2016)	97	Yes	No	No	No

Table A1 Nearshore Enterococci Levels, Exceedance of ANZECC (2000) Recreational Water Quality Guidelines

Location (Date)	Enterococci (MPN/100mL)	Exceed – NHMRC Category A – 40 cfu/100 mL	Exceed – NHMRC (2008) Category B Iower limit – 41 cfu/100mL	Exceed –NHMRC (2008) Category B – Upper Limit – 200 cfu/100 mL	Exceed – NHMRC (2008) – Category C – 201 cfu/100 mL	Exceed – NHMRC – Category D - >500 cfu/100mL
Point Moore (9/4/2013)	41	Yes	Yes	No	No	No
Pages Beach (9/4/2013)	73	Yes	Yes	No	No	No
Point Moore (8/1/2014)	54	Yes	Yes	No	No	No
Point Moore (15/1/2015)	84	Yes	Yes	No	No	No
Pages Beach (15/04/2015)	170	Yes	Yes	No	No	No
Pages Beach (12/03/2015)	230	Yes	Yes	Yes	No	No
Pages Beach (3/02/2016)	910	Yes	Yes	Yes	Yes	Yes
Pages Beach (19/04/2016)	97	Yes	Yes	No	No	No

Table A2 Nearshore Enterococci Levels, Exceedance of Criteria that Define NHMRC (2008) Recreational Water Quality Categories

Basis of the derivation of the limits for each category provided in Table A2 are provided in Table A3.

Table A3Basis of derivation of percentile values for determining microbial
water quality categories (NHMRC, 2008)

Category ^a	95 th percentile value for intestinal enterococci/ 100 mL (rounded values)	Basis of derivation	Estimation of probability
A	≤40	This value is below the NOAEL in most epidemiological studies.	GI illness risk: < 1% AFRI risk: < 0.3% The upper 95th percentile value of 40/100 mL relates to an average probability of less than one case of gastroenteritis in every 100 exposures.The AFRI burden would be negligible.
В	41–200	The 200/100 mL value is above the threshold of illness transmission reported in most epidemiological studies that have attempted to define a NOAEL or LOAEL for GI illness and AFRI.	GI illness risk: 1–5% AFRI risk: 0.3–1.9% The upper 95 th percentile value of 200/100 mL relates to an average probability of one case of gastroenteritis in 20 exposures. The AFRI illness rate would be 19 per 1000 exposures or approximately 1 in 50 exposures.
с	201–500	This represents a substantial elevation in the probability of all adverse health outcomes for which dose-response data are available.	GI illness risk: 5–10% AFRI risk: 1.9–3.9% This range of 95 th percentile values represents a probability of 1 in 20 to 1 in 10 risk of gastroenteritis for a single exposure. Exposures in this category also suggest a risk of AFRI in the range of 19–39 per 1000 exposures or a range of approximately 1 in 50 to 1 in 25 exposures.
D	> 501	Above this level there may be a significant risk of high levels of illness transmission.	GI illness risk: > 10% AFRI risk: > 3.9% There is a greater than 10% chance of illness per single exposure. The AFRI illness rate at the guideline value of 500 enterococci per 100 mL would be 39 per 1000 exposures or approximately 1 in 25 exposures.

Appendix B – Point Moore Inundation and Coastal Processes Study Summary Report

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Point Moore Inundation & Coastal Processes Study

In Western Australia, coastal development is guided by State Planning Policy 2.6: The State Coastal Planning Policy (SPP2.6). This policy outlines the general requirements for new development on the coastline in terms of avoiding or managing risks caused by coastal inundation (flooding) or coastal erosion.



SPP2.6 outlines that new development should be safe from coastal inundation caused by an extreme inundation event that has a 0.2% chance of occurring each year. In other words, this event would occur once every 500 years on average. Another way to say this is that the event would have an Average Recurrence Interval (abbreviated to ARI) of 500 years.

For coastal erosion, SPP2.6 states that new development should be safe from an erosion event that has a 1% chance of occurring each year (or would occur once every 100 years on average – i.e. the 100 year ARI event).

In addition to these storm events, the potential impacts of sea level rise and ongoing changes to the shoreline need to be considered when determining appropriate areas for new development.

While the SPP2.6 guidelines relate mainly to the requirements for new development, where existing development does not meet the guidelines there is a general requirement to take action in order to reduce any risks to acceptable levels.

In 2015 a study was completed in order to understand which areas of Point Moore could be impacted by coastal inundation and erosion. This study was completed in accordance with the requirements of SPP2.6. The study involved detailed modelling and assessment of the following items:

- Detailed cyclone storm surge modelling to determine the potential inundation caused by severe cyclones.
- Analysis of available water level records to determine the potential inundation caused by non-cyclonic events.
- Modelling the potential beach and dune erosion caused by severe events. •
- Assessment of historical and potential future shoreline movement caused by the • action of natural coastal processes.
- Assessment of the effects of potential sea level rise (assuming 0.9 metres of sea • level rise by year 2110 as required by SPP2.6) on the coastal inundation and erosion.

The results of this study are summarised on the attached plans. Further details and description of these plans are provided overleaf.

The attached plans show the areas that could be impacted by coastal erosion or inundation for the Present Day, as well as the years 2030, 2070 and 2110. A description of what these plans mean, and how to read them, is provided below.

Coastal Processes Allowance Plan

The Coastal Processes Allowance Plan shows 4 different coloured lines. Each of these lines represents the extent of possible impact of coastal erosion over each planning horizon. The locations of these lines have been determined in accordance with the requirements of SPP2.6. As an example, anything on the ocean side of the **red** line could be vulnerable to coastal erosion by the year 2110.

Coastal Inundation Mapping Plans

The Inundation Mapping Plans show areas that could be inundated by different events for each of the timeframes outlined above. Each of the different plots represents a different timeframe. The different colours represent the potential areas of inundation associated with different event severities. On each of the plots, the area that is shaded **purple** represents the area that would be inundated during the 20 year ARI event; the area that is shaded **blue** represents the <u>additional</u> area that would be inundated during the 100 year ARI event; and the area that is shaded **green** represents the <u>additional</u> area that could be inundated by the 500 year ARI event.

The difference between the plots (as each plot represents a different time), is caused by the potential impact of sea level rise.

Coastal Inundation Depth Plans

The Inundation Depth Plans have been prepared to show the potential depth of inundation caused by the 20, 100 and 500 year ARI events at the year 2030, as well as the 500 year ARI event in 2110. The different colours on these plans show the different inundation depths, as indicated on the legend. For example, anything that is shaded **pink** on the plan would have an inundation depth of between 2.0 and 2.5 metres.

Combined Coastal Vulnerability Mapping Plans

The Combined Coastal Vulnerability Mapping Plans identify the areas that would be impacted by the 500 year ARI inundation event and/or the Coastal Processes Allowance for each timeframe. For example, on the plan depicting the year 2030, the shading depicts the area that would be subject to inundation during the 500 year ARI event as well as the area that would be potentially vulnerable to coastal erosion by 2030 (as shown on the Coastal Processes Allowance Plan).

The significance of these combined plans is that the shaded areas represent the areas that would not be developable under SPP2.6 for each of the different timeframes.

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POINT MOORE COASTAL EROSION AND INUNDATION STUDY INUNDATION MAPPING - PRESENT DAY

TEN UNDATION STUDY





1 111400



<u>NOTES</u>

- INUNDATION EXTENTS BASED ON LIDAR DATA PROVIDED BY THE NORTHERN AGRICULTURAL 1. CATCHMENTS COUNCIL (MARCH 2013). VERTICAL ACCURACY IN THE ORDER OF 0.2m.
- 2. AERIAL PHOTOGRAPH TAKEN NOVEMBER 2014.

scale at a3 1:4,000

JULY 2015

D1242-02-01(B)



coastal and port engineers

t: +61 8 9254 6600 f: +61 8 9254 6699 Suite 1, 128 Main Street Osborne Park 6017 Western Australia admin@coastsandports.com.au INUNDATION MAPPING - 2030 (+0.07 SLR)



LEGEND 20Yr ARI EVENT 100Yr ARI EVENT 500Yr ARI EVENT





<u>NOTES</u>

INUNDATION EXTENTS BASED ON LIDAR DATA PROVIDED BY THE NORTHERN AGRICULTURAL CATCHMENTS COUNCIL (MARCH 2013). VERTICAL ACCURACY IN THE ORDER OF 0.2m. 1.

2. AERIAL PHOTOGRAPH TAKEN NOVEMBER 2014.

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scale at a3 1:4,000

D1242-02-02(B)



INUNDATION MAPPING - 2070 (+0.39 SLR)

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<u>LEGEND</u> 20Yr ARI EVENT 100Yr ARI EVENT 500Yr ARI EVENT





<u>NOTES</u>

INUNDATION EXTENTS BASED ON LIDAR DATA PROVIDED BY THE NORTHERN AGRICULTURAL 1. CATCHMENTS COUNCIL (MARCH 2013). VERTICAL ACCURACY IN THE ORDER OF 0.2m.

2. AERIAL PHOTOGRAPH TAKEN NOVEMBER 2014.

JULY 2015

scale at a3 1:4,000

D1242-02-03(B)

coastal and port engineers

Western Australia

admin@coastsandports.com.au



INUNDATION MAPPING - 2110 (+0.9 SLR)



<u>LEGEND</u> 20Yr ARI EVENT 100Yr ARI EVENT 500Yr ARI EVENT





<u>NOTES</u>

INUNDATION EXTENTS BASED ON LIDAR DATA PROVIDED BY THE NORTHERN AGRICULTURAL CATCHMENTS COUNCIL (MARCH 2013). VERTICAL ACCURACY IN THE ORDER OF 0.2m. 1.

2. AERIAL PHOTOGRAPH TAKEN NOVEMBER 2014.

JULY 2015

scale at a3 1:4,000

D1242-02-04(B)

P:\MRA Paying Jobs\K1242 CGG - Point Moore Inundation Study\MRA DWGs\Innundation\2110\K1242-2110



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POINT MOORE COASTAL EROSION AND INUNDATION STUDY INUNDATION DEPTHS - 2030 20YR ARI EVENT



P:\MRA Paying Jobs\K1242 CGG - Point Moore Inundation Study\MRA DWGs\Innundation\Additional Mapping\2030 20Yr ARI\K1242-2030 20Yr ARI



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POINT MOORE COASTAL EROSION AND INUNDATION STUDY INUNDATION DEPTHS - 2030 100YR ARI EVENT

scale at a3 1:4,000



D1242-03-02(A)



INUNDATION DEPTHS - 2030 500YR ARI EVENT

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Western Australia admin@coastsandports.com.au

coastal and port engineers



SCALE	
AT A3	1:4,000

JULY 2015



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POINT MOORE COASTAL EROSION AND INUNDATION STUDY INUNDATION DEPTHS - 2110 500YR ARI EVENT





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POINT MOORE COASTAL EROSION AND INUNDATION STUDY COMBINED COASTAL VULNERABILITY MAPPING - PRESENT DAY

- 1. INUNDATION EXTENTS BASED ON LIDAR DATA PROVIDED BY THE NORTHERN AGRICULTURAL CATCHMENTS COUNCIL (MARCH 2013). VERTICAL ACCURACY IN THE ORDER OF 0.2m.
- 2. AERIAL PHOTOGRAPH TAKEN NOVEMBER 2014.

DECEMBER 2015

scale at a3 1:4,000

D1242-06-01(B)



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POINT MOORE COASTAL EROSION AND INUNDATION STUDY COMBINED COASTAL VULNERABILITY MAPPING - 2030 (+0.07 SLR)

- 1. INUNDATION EXTENTS BASED ON LIDAR DATA PROVIDED BY THE NORTHERN AGRICULTURAL CATCHMENTS COUNCIL (MARCH 2013). VERTICAL ACCURACY IN THE ORDER OF 0.2m.
- 2. AERIAL PHOTOGRAPH TAKEN NOVEMBER 2014.

DECEMBER 2015

7

scale at a3 1:4,000

D1242-06-02(B)



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POINT MOORE COASTAL EROSION AND INUNDATION STUDY COMBINED COASTAL VULNERABILITY MAPPING - 2070 (+0.39 SLR)

- 1. INUNDATION EXTENTS BASED ON LIDAR DATA PROVIDED BY THE NORTHERN AGRICULTURAL CATCHMENTS COUNCIL (MARCH 2013). VERTICAL ACCURACY IN THE ORDER OF 0.2m.
- 2. AERIAL PHOTOGRAPH TAKEN NOVEMBER 2014.

DECEMBER 2015

7

scale at a3 1:4,000

D1242-06-03(B)



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POINT MOORE COASTAL EROSION AND INUNDATION STUDY COMBINED COASTAL VULNERABILITY MAPPING - 2110 (+0.9 SLR)

- 1. INUNDATION EXTENTS BASED ON LIDAR DATA PROVIDED BY THE NORTHERN AGRICULTURAL CATCHMENTS COUNCIL (MARCH 2013). VERTICAL ACCURACY IN THE ORDER OF 0.2m.
- 2. AERIAL PHOTOGRAPH TAKEN NOVEMBER 2014.

DECEMBER 2015

7

scale at a3 1:4,000

D1242-06-04(B)

Appendix C – Groundwater Monitoring Well Logs

Bore Log

Groundwater Well: MW1
Page: 1 of 3

Client	City of Great	er Geraldton		Drill Co:	Harrington d	Irilling	Easting	263, 938		
Project	Point Moore ROETDS Driller:		James Harrington North		Northing	6,814,127				
Job No.	6134536	4536 Rig Type:		HD1	HD1 Grid Ref					
Location	Point Moore			Total Depth:	6		Elevation	2.608 (top of cas	ing, mAHD)	
Date	15/08/2016			Diameter (mm):	50	1	Logged by	S Petts	Checked by	A. Nagle
Depth	Drilling	Sample ID	Water	Well Details	Graphic	Lithological Description		Consistency	Moisture	CONTAMINANT INDICATORS
(metres)	Method					Soil Type (Classification Group Syn	nbol);			
					Log	Particle ; Size; Colour;				Odours, staining, waste materials,
0.4					Mariana ang ang ang ang ang ang ang ang ang	Secondary/Minor Component			D	separate phase, liquids, imported fill, ash.
0.1		MW1 05		• Inte		Sivi Eine Eine Sand Bro	vn Sub-angular		D	No ouour, no signs of containination
-		10.5		<u>9</u> 🕅 🕅 🤤		Lavor Quartz Wo	wir Sub-angular			
-						Minor loam Loa	n (gravel)			No sampling beyond 3 m inadequate
-				E Source S			ii (gravei)		D	sample amount
0.5									U	sample anount.
-										
-										
F										
F		MW1 1.0								
- 1.0									м	No odours or stains
Γ										
Γ										
Γ										
		MW1_1.5								No odours or stains
1.5										
L										
Ļ										
L										
20		MW1_2.0						_		
- 2.0						SW			M	No odours or stains
-						Coarse Coarse Sand Wh	te Sub-angular			
-				a 🛛 🔜 着		Layer Quartz We	kly Cemented			
-			_	s 🔕 🕺 s						
- 2.5		MW1_2.5	¥	9 🛞 🚫 U						AL
-				i 🕅 🕅					vv	No odours or stains
-										
-										
-		MW1 2 0								
-3.0		101001_5.0							14/	No odours or stains
-									**	
-										
-										
-		MW1 35							w	No odours or stains
- 3.5		WW1_5.5							**	
-										
F										
-										
		ns							w	No odours or stains
-4.0										
Г						1				
Γ										
Ε										
		ns							w	No odours or stains
4.5						1				
L									1	
L						1				
L										
5.0		ns							w	No odours or stains
L	1								1	
L										
F										
F										
- 5.5		ns							w	No odours or stains
F										
F						1				
F										
F										
L	I	I	l	· • • • • • •	1262626262	1		1	W	No odours or stains
Groundwater Well: MW2 Page: 2 of 3

Bore Log

Client	City of Croot	or Coroldton		Deil	1.0	Horrington	Deilling		Fasting	262 700		
Project	Point Moore	ROETDS		Dril	ler:	James Harr	ington		Northing	6.813.860		
Job No.	6134536			Rig	Туре:	HD1			Grid Ref			
Location	Point Moore			Tota	al Depth:		6		Elevation	2.151 (top of cas	ing, mAHD)	
Date	15/08/2016	5	1	Dia	meter (mm):	5	0		Logged by	S Petts	Checked by	A. Nagle
Depth	Drilling	Sample ID	Water	We	ll Details	Graphic	Lithologic	al Description	e 1 1	Consistency	Moisture	CONTAMINANT INDICATORS
(metres)	Method						Soil Type	(Classification Gro	up Symbol);			
						LOg	Minor Co	mpopent	bary/			odours, staining, waste materials
0.1				Ħ,		0000000	SM	mponent			D	No odours or stains
- 0.1				Gro			Fine	Fine Sand	Brown Sub-angular			
[led			Layer	Quartz	Weakly Cemented			
				Sea			Minor Lo	am				
0.5		MW2_0.5									D	No odours or stains
- 0.0		QC01										
-		QC02										
-				IT								
-		MW2 10										No adours or stains
- 1.0		101002_1.0					SM			4	M	
-							Fine	Fine Sand	Brown Sub-angular		IVI	
-							Laver	Quartz	Weakly Less Loam			
Γ									,			
1		MW2_1.5										No odours or stains
- 1.5		_										
							8					
L												
Ļ												
-20		MW2_2.0					å			4		No odours or stains
							SW		brown to		м	
-			-	_			Medium	Medium Sand	White Sub-angular			
-			*	feer			Layer	Quartz	weakly cemented			
-				I Sc								No odours or stains
- 2.5		MW2 2.5		ttec							w	
F				Slo								
-												
Γ												No odours or stains
30		ns										
- 5.0		no sample					8				W	
L		retreivable										
_												
-							8					
- 3.5		MW2_3.5									W	No odours or stains
-												
-												
-												
F		MW2 4.0									w	No odours or stains
-4.0							SW			4		
F							Coarse	Coarse Sand	Grey Sub-angular			
[Layer	Quartz	Weakly Cemented			
							8					
- 4 5		MW2_4.5					8				W	No odours or stains
-							8					
-							3					
-							8					
-							8				14/	
-5.0		WIWZ_5.0					8				vv	
F	1											
F							8					
F		1					1					No odours or stains
5.5		MW2_5.5					8				w	
[^{3.5}							8					
Γ							1					
L							3					
F		1					8					
L		MW2_6.0		•	K K K		8				W	No odours or stains



Groundwater Well: MW3 Page: 3 of 3

Client	City of Great	er Geraldton		Drill Co:	Harrington Dril	ling	Facting	263 934		
Project	Point Moore	ROETDs		Driller:	James Harringt	on	Northing	6,813,674		
Job No.	6134536	-		Rig Type:	HD1		Grid Ref			
Location	Point Moore			Total Depth:	6m		Elevation	3.009 (top of cas	ing, mAHD)	
Date	15/08/2016	i		Diameter (mm):	50		Logged by	S Petts	Checked by	A. Nagle
Depth	Drilling	Sample ID	Water	Well Details	Graphic	Lithological Description		Consistency	Moisture	CONTAMINANT INDICATORS
(metres)	Method				Log	Soil Type (Classification Grou	ıp Symbol);			Odours, staining, waste materials
						Particle; Size; Colour;				
						Secondary/Minor Componer	nt		-	separate phase, liquids, imported fill, ash.
0.1				t t		SP Since Since Cound	Descus Cub secular		D	
-				<u>بة 🗙 🗙 الم</u>		Fine Fine Sand	Brown Sub-angular			No odours or stains
-				i ≩ 🗱 🗮 💥 🗿		very minor loam	weakly cemented			
-		MW3 0 5		9 X 🛛 🖓 🗄		very millior loan			D	
- 0.5		QC03			· [2000000				5	No odours or stains
-		QC04								
[
-10		MW3_1.0								
_									м	No odours or stains
-										
F										
-		M/M/2 1 5								
- 1.5		101003_1.5								No odours or stains
-										
-										
-										
T		MW3_2.0								
2.0						SP			м	No odours or stains
[Fine Medium Sand	White Sub-angular			
L						Layer Quartz	Weakly Cemented			
-										
- 2.5		MM2 25								
-		101003_2.5							vv	NO ODOURS OF Stains
-										
F										
		MW3 3.0	-							No odours or stains
- 3.0		-	· ·						w	
Ε										
[]										
L										
- 3.5		MW3_3.5							w	
-										No odours or stains
-										
-										
-		MW3 4.0							w	
-4.0					00000000	ŚW				No odours or stains
-						Coarse Coarse Sand	Grev Sub-angular			
F						Layer Quartz	Weakly Cemented			
Ε						shell grit				
45		MW3_4.5							w	
										No odours or stains
F										
F										
F		MM2 50							14/	
- 5.0		101003_5.0			83333333	CM .		-	vv	No odours or stains
F .						Sivi Fine Fine Sand	Grev Sub-angular			NO OUDULS OF SLAINS
F						Laver Quartz	Sicy Sub-diiguidi			
F						.,				
55		MW3_5.5							w	No odours or stains
5.5										
L						1				
F										
F								-		
_		MW3_6.0				Limestone	Layer		W	

Appendix D – Groundwater Quality Field Data Sheets



Client: CO	GG									BORE	ID: M	IW1	
Project: P	oint Moore	Groundw	vater Ass	essment						Job No	.: 6134	4772	2
Location:	Marine Te	rrace	0	Casing di	ameter:				50 mm	Total D	epth:	5.99) m
BORE CO	ONSTRUCT	TION											
Head-	🗷 Flush	□Monur	nent					Datu	m 🗷 Cove	er/ 🛛		Dat	tum Elev.:2.608m
works					Casing	Locke	d	Point	Monum	ent Ini Ca	ner sing		Local 🗵 AHD
DUDCING		2								eu	51115		
Mothod	J DETAILS	Datas 20	16				T In	dontal	ron Dru C D	otta	Flore	. Do	tot 0 005 I /S
Method:		Date: 20 Water C	lo olumn· 3	57 m			D.		xell Dy: 5.P		FIOW		
PID reading	ng: N/A	The second secon	orunni. o				Re	q Purş	ge Vol. 1: N/	Α	Actu		ol. Purged:
		Time			Dep LN	th to APL (m)	Dej	pth to	water (m)		Dept	th to	DNAPL (m)
Start		0830			-		2.3	6			-		
Finish		0930			-		2.3	6			-		
PURGINO	G MEASUR	REMENTS	S ²										
Vol. Purged (L)	Elapsed Time (min)	Spec. Cond. (µS/cm)	Act. Cond. (µS/cm)	TDS (ppt)	Temp. (°C)	Salinity (ppt)	p	Н	DO %Sat	DO (mg/L) (m	RP 1V)	Comments – Turbidity, Colour, Odour, etc.
1	5	20,050	18,085	13.03	20.2	12.1	7.	.31	98.45	9.20	-1	64	Colourless, minor
2	10	19,116	17,708	56.2	21.1	11.5	7.	.33	94.20	6.14	-1	99	sulphur type odour
3	12	18,647	17,451	51.2	21.5	11.2	7.	.34	86.14	5.59	-2	.02	
4	15	17,233	18,439	42.1	21.6	11.0	7.	.35	55.64	5.03	-2	.07	
5	17	17,910	16,782	45.3	21.7	10.7	7.	.36	50.96	4.60	-2	10	
		+/-3.0%			+/-3.0%		+/-	- 0.1	+/-10%	+/-3.0%	6 +/- m	- 10 1V	
Comments	s (e.g. condi	ition of he	adworks,	colour,	odour):								
SAMPLIN	IG DETAII	LS			San	ple ID: M	W1						
Date		Time:			Und	ertaken By	: S Pe	etts					
PID reading	g: N/A	Depth to	LNAPL:	N/A	Dep	th to Water	: 2.4	2m	Depth to D	NAPL: N	N/A		
Flow Rate:	0.005 L/s	Vol. Rem	oved: 5L		Sam	pling Meth	od: L	Low Flo	0W				
Temp.: 21.7	∕ °C	DO: 4.60) mg/L		pH:	7.36	Sp	ec. Co	nd: 17,910 uS	S/cm	ORP	: -2	10 mV
DO: 50.9%	Sat	Salinity:	10.7 ppt		TDS	45.3 ppt							
Comments	s (e.g. conta	iners, filti	ration): (QC01 - B	lind, QC	202 – Split,	QC	03 – R	insate, QC04	4 – Field	Blank	K	
CoC Num	ber:240820	15			Che	cked by: S	Steve	n Petts	S	Date	e: 24/0)8/2()16

Bores to be purged dry, until pH, T and EC readings stabilise or a minimum of 3 to 5 times the water column volumes. Water column volumes can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID - 2 L/m; 100 mm ID 8 L/m. Calibration details to be recorded in the instrument –specific calibration book, or in field notes as required by local procedures. 1

2



Client: CO	G									BORE	ID: M	1W2	
Project: P	oint Moor	e Ground	water As	ssessment						Job No	.: 613	4772	2
Location:	Captains (Cres	(Casing di	amete	r:			50 mm	Total D	epth:	: 6.1() m
BORE CO	NSTRUC	TION											
Head-		□Monu	ment				1	Datur	n 🗷 Cove	er/		Dat	tum Elev.:2.151m
works	Flush				Cash	ig Locke	a	Point	Monum	ent Ini Ca	sing		Local 🗵 AHD
PURGING	DETAIL	S											
Method: L	ow Flow	Date: 24	/08/2016				Unc	dertak	en By: S.Pe	etts	Flov	v Ra	te: 0.008 L/S
PID reading	ng: N/A	Water (Column: 4	4.24 m			Req	l Purg	e Vol. ¹ : N/	A	Actu	ual V	ol. Purged:
		Time			D L	epth to NAPL (m)	Dep	oth to y	water (m)		Dep	th to	DNAPL (m)
Start		1000			-		1.78	8			-		
Finish		1100			-		1.78	8			-		
PURGING	MEASU	REMENT	ΓS ²										
Vol. Purged (L)	Elapsed Time (min)	Spec. Cond. (µS/cm)	Act. Cond. (µS/cm)	TDS (ppt)	Temp (°C)	b. Salinity (ppt)	p	Н	DO %Sat	DO (mg/L) (n	nV)	Comments – Turbidity, Colour, Odour, etc.
1	2	15,159	12,459	234.4	15.6	8.88	8.4	42	86.6	8.99	-1	164	Colourless, minor
2	4	2,637	2,274	755.3	17.8	1.37	8.	34	61.7	6.1	-1	62	H ₂ S odour
3	6	2,970	3,283	64.87	20.2	1.73	7.4	47	29.6	2.8	-2	205	
4	8	3,408	3,107	72.1	20.4	1.81	7.	37	24.5	2.3	-2	206	
5	10	3,419	3,125	74.0	20.5	1.81	7.4	43	22.1	2.1	-2	207	
6	12	3,450	3,128	74.2	20.6	1.81	7.4	48	22.5	2.2	-2	207	
7	14	3,500	3,150	75.1	20.6	1.81	7.4	45	22.2	2.1	-2	206	
		+/-3.0%			+/-3.0%	<i>6</i>	+/-	0.1	+/-10%	+/-3.0%	6 +/ 11	/- 10 nV	
Comments	s (e.g. cond	lition of h	eadwork	s, colour	odou	r):							
SAMPLIN	IG DETAI	LS			Sa	ample ID: M	[W2						
Date: 24/08	/2016	Time: 11	:00		U	ndertaken By	: S Pet	tts					
PID reading	g: N/A	Depth to	LNAPL:	N/A	D	epth to Water	: 1.86	m	Depth to D	NAPL: N	N/A		
Flow Rate:	0.008 L/s	Vol. Rem	oved: 7 L		Sa	mpling Meth	od: Lo	ow Flov	w		1		
Temp.: 21.7	′ °C	DO: 2.08	8 mg/L		pł	I: 7.43	Spe	ec. Con	id: 3,419 uS/o	cm	ORF	P: -20	07 mV
DO: 22.1 %	Sat	Salinity:	1.81 ppt		T	DS: 2.22 ppt							
Comments	(e.g. cont	ainers, fil	tration):							<u> </u>			
CoC Num	ber:24082	015 raed dry ur	tilnH To	nd FC read	C	hecked by: S	Stever	1 Petts	5 times the wo	Date	e: 24/(08/20)16 Natar column volumos

Bores to be purged dry, until pH, T and EC readings stabilise or a minimum of 3 to 5 times the water column volumes. Water column volu can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID – 2 L/m; 100 mm ID 8 L/m.

2 Calibration details to be recorded in the instrument -specific calibration book, or in field notes as required by local procedures.



Client: CO	G									BORE	ID: M	1W3	
Project: P	oint Moor	e Ground	water As	sessment	t					Job No	.: 613	4772	
Location:	Astrolabe	Ln	C	Casing di	ameter	•			50 mm	Total D	epth:	5.8 4	l m
BORE CO	NSTRUC	TION											
Head-	x	□Monu	nent					Datu	m 🗷 Cove	er/ 🛛		Dat	tum Elev.:3.009m
works	Flush				Casin	g Locke	ed	Poin	t Monum	ent Ini Ca	ner sing		Local 🗵 AHD
PURCING	L DELAH	S								eu	Sing		
Method · I		Date: 2/	/08/2016				Un	derta	kon Rv. S Pa	otte	Flow	v Ra	te• 0.004 L/S
DID woodin		Water C	Column: 3	.13 m					$\frac{1}{2} \sum_{n=1}^{\infty} \frac{1}{2} \sum_{n=1}^{\infty} \frac{1}$	A	A of the		
FID reading	ig: IN/A	T:			D		Re			A	Actu		oi. Purgeu:
		Time			De LN	pth to NAPL (m)	De	pth to	water (m)		Dept	th to	DNAPL (m)
Start	tart 1100 - 2.67 -												
Finish		1200			-		2.6	7			-		
PURGING	G MEASU	REMENT	TS ²										
Vol. Purged (L)	Elapsed Time (min)	Spec. Cond. (µS/cm)	Act. Cond. (µS/cm)	TDS (ppt)	Temp (°C)	Salinity (ppt)	I	рН	DO %Sat	DO (mg/L) (n	RP nV)	Comments – Turbidity, Colour, Odour, etc.
1	4	2,333	2,259	41.3	23.3	1.21	7	.84	22.6	2.1	-1	92	Minor
2	8	2,415	2,300	42.0	23.4	1.21	7	.83	22.7	2.1	-1	.93	particulates, colourless, H ₂ S
3	12	2,336	2,261	36.4	23.3	1.21	7	.81	22.6	2.05	-1	96	odour
4	16	2,314	2,248	33.1	23.4	1.20	7	.83	17.9	1.69	-2	204	
5	20	2,219	2,233	34.9	23.7	1.19	7	.83	14.2	1.28	-2	211	
		+/-3.0%			+/-3.0%		+/	- 0.1	+/-10%	+/-3.0%	6 +/- n	- 10 nV	
Comments	s (e.g. cond	lition of h	eadworks	s, colour	, odou	·):				1			
SAMPLIN	IG DETAI	LS			Sa	mple ID: M	IW3						
Date: 24/08	/2016	Time: 11	45		Un	dertaken By	: S Pe	etts					
PID reading	g: N/A	Depth to	LNAPL: N	V/A	De	pth to Water	:: 2.7	1 m	Depth to D	NAPL: N	/A		
Flow Rate:	0.04 L/s	Vol. Rem	oved: 5L		Sa	mpling Meth	od: L	low Fl	ow				
Temp.: 23.7	o ₀C	DO: 1.28	8 mg/L		pН	: 7.83	Sp	ec. Co	nd: 2,219 uS/	cm	ORP	P: -2	11 mV
DO: 14.2%	Sat	Salinity:	1.19 ppt		TĽ	S: 34.9 ppt							
Comments	s (e.g. cont	ainers, fil	tration):										
CoC Num	ber:24082	015			Cł	ecked by: S	Steve	n Pett	S	Date	e: 24/0)8/20)16

1 Bores to be purged dry, until pH, T and EC readings stabilise or a minimum of 3 to 5 times the water column volumes. Water column volumes can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID - 2 L/m; 100 mm ID 8 L/m.

2 Calibration details to be recorded in the instrument –specific calibration book, or in field notes as required by local procedures.



Client: CO	GG									BORE	ID: SI	HP8	
Project: P	oint Mo	ore Ground	water As	ssessmen	t					Job No.	.: 6134	4772	
Location:	Coxswa	in Park	(Casing di	ameter	:			50 mm	Total D	epth:	4.72	m
BORE CO	NSTRU	JCTION											
Head-		Monume	nt		×			Datu	m 🗷 Cove	er/ 🛛		Dat	um Elev.:2.427m
works	Flush				Casing	g Locke	ed	Point	Monum	ent Inr Ca	ner sing		local 🗷 AHD
PURGINO	G DETA	ILS									~8		
Method: L	low Flow	v Date: 24	/08/2016				Un	dertak	en By: S.Pe	etts	Flow	v Rat	te: 0.005 L/S
PID reading	ng: N/A	Water C	Column: 2	2.42 m			Re	q Purg	e Vol. ¹ : N/	A	Actu	al V	ol. Purged:
		Time			Der LN	oth to APL (m)	De	pth to	water (m)		Dept	th to	DNAPL (m)
Start		1230			-		2.1				-		
Finish		1330			-		2.1				-		
PURGING	G MEAS	UREMENT	TS ²										
Vol. Purged (L)	Elapse Time (min)	d Spec. Cond. (µS/cm)	Act. Cond. (µS/cm)	TDS (ppt)	Temp. (°C)	Salinity (ppt)]	рН	DO %Sat	DO (mg/L) (m	RP iV)	Sal(ppt) Comments – Turbidity, Colour, Odour, etc.
1	5	6,410	5,917	6.99	20.98	3.54	7	.72	31.2	3.2	-2	21	Colourless, H ₂ S
2	7	6,497	5,987	5.58	20.95	3.57	7	.72	30.2	3.1	-2	22	odour
3	9	6,587	5,974	5.01	20.91	3.58	7	.73	30.1	3.1	-2	21	
4	11	6,588	5,879	5.01	20.91	3.58	7	.73	29.3	3.01	-2	21	
5	13	6,496	5,988	4.22	20.91	3.59	7	.73	29.3	3.01	-2	21	
		+/-3.0%			+/-3.0%		+/	- 0.1	+/-10%	+/-3.0%		- 10 1V	
Comments	s (e.g. co	ondition of h	eadwork	s, colour	, odour)):							
SAMPLIN	IG DET	AILS			San	nple ID: SI	HP8						
Date: 24/08	/2016	Time: 13:20			Und	lertaken By	:SP	etts					
PID reading	g: N/A	Depth to LN	APL: N/A	A	Dep	oth to Water	: 2.3	m	Depth to D	NAPL: N	J/A		
Flow Rate:	-L/s	Vol. Remove	ed: 5 L		San	npling Meth	od: I	.ow Flo	w				
Temp.: 20.9	0°C	DO: 29.32 1	mg/L		pH:	7.73	S	ec. Cor	nd: 6,696 uS/	cm	ORP	: -22	21 mV
DO: 29.3%	Sat	Salinity: 3.5	9 ppt		TDS	S: 24.80 ppt							
Comments	s (e.g. co	ontainers, fil	tration):										
CoC Num	ber:240	82015			Ch	ecked by: S	Steve	n Petts		Date	e: 24/0)8/20	16

 1
 Bores to be purged dry, until pH, T and EC readings stabilise or a minimum of 3 to 5 times the water column volumes. Water column volumes can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID - 2 L/m; 100 mm ID 8 L/m.

2 Calibration details to be recorded in the instrument –specific calibration book, or in field notes as required by local procedures



Client: C	GG												BO	RE II): MW1				
Project: l	Poin	t Moor	e Grou	indwa	ater Ass	essment							Job	No.:	6134772	2			
Location	: Ma	arine Te	errace			Casing	diam	eter:		Locatio	on: M	larine	Tota	al Dej	pth:	5.93			
BORE C	ONS	STRUC	TION																
Head- works	× Flu	sh	⊐Mon	ument	t 🗆 Ca	asing	□ Locke	ed	Da Po	itum int	□ Co Monu	over/ ument	Inr Casin	ner g	Datum Elev.:	I	n 🗵	l Loc] AH	:al D
PURGIN	G D	ETAIL	S																
Method:		Low Flo Pump	ow	Dat	e: 2016				Un S.I	dertak Petts	en By	7 :		Flov	w Rate:]	L/s
PID read	ling:	N/A	Appm	Wa	ter Colu	ımn:		3.5m	Re	q Purg	e Vol	¹ :	-L	Actu	ual Vol.	Purged			-L
		Time			Depth	to LNAP	PL (m	l)	De	pth to v	vater	· (m)		Dep	th to DN	NAPL (n	1)		
Start		09:00			-				2.4	13				-					
Finish		10:15			-				2.4	13				-					
PURGIN	IG N	IEASU	REME	ENTS	2			·											
Vol. Purge (L)	ed	Elapso Time (n	ed nin)	EC (mS/ci	m) Te	mp. (°C)	Sa ()	alinity ppm)		рН	DO	%Sat	DO (p	pm)	Eh (mV)	C Turt C	omme idity,)dour	nts – Colo , etc.	our,
1		10		31.6	5	28.1		-		6.94		-	1.0	8	-282	S	O_2 od	lour,	
2		14		31.7	7	28.1		-		6.97		-	0.3	3	-283	c sust	olour endea	less, 1 soli	ids
3		18		31.7	7	28.1		-		6.96		-	0.3	6	-276				
4		23		31.6	5	28.3		-		6.93		-	0.4	4	-273				
				+/-3.09	% -	+/-3.0%				+/- 0.1	+/	-10%	+/-3.0)%	+/- 10 mV	7			
Commen	ts (e	.g. cond	lition (of hea	dworks	, colour,	odou	r):											
SAMPLI	NG	DETAI	LS							Sample	e ID:	MW1							
Date 2	22/09	/2016		Tim	e:10:00					Undert	aken	By: S Pe	tts						
PID reading	ng:		- ppm	Dep	th to LN	APL:		- 1	m	Depth t	o Wa	ter:	2	.43 m	Depth	to DNAI	Ľ		- m
Flow Rate	e:	0.09L/ s	Vol. I	Remov	ed:		4L	Sampli	ing 1	Method:		Low F	'low						
Temp.:		28.3°C	DO:0	.4		-l r	opm ng/L	pH:		6.93		EC: 31.0	6		mS/cm	Eh:-273]	mV
			DO:-			0	6Sat					Salinity	:		-ppm mg/L				
Commen	ts (e	.g. cont	ainers	, filtra	ation):	QC01_2()1609	22, RB	- Q	C03_20	1609	22, FB	- QC04	4_201	60922				
CoC Nu		er: 20	16092	2 1	T and F	Creadings	ked b	y:	S P	etts	o 5 ti	nos tha m	Da ntor col-	ate:	22/0	9/2016	n vol-	mee	
i Bore	es 10 f	re purged	ury, u	in pH	, i and E	C readings	Stadiff	se or a m	40		u s un	nes the Wa	ater colt	100 IIII	Tumes. W	ater colun	m voit	mes	

can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID - 2 L/m; 100 mm ID 8 L/m.
 Calibration details to be recorded in the instrument –specific calibration book, or in field notes as required by local procedures.



Client: CO	GG	r r											BO	RE II): MW2				
Project: P	Poir	nt Moore	Grou	indw	ater A	Assessment							Job	No.:	6134772	2			
Location:	Ca	aptain Cro	es			Casing	diam	eter:				50 mm	Tota	al Dej	pth:				6.03
BORE CO	ON	STRUCT	ION																
Head- works	⊠ Flı	⊔ush M	onun	nent		Casing]	⊐ Locke	ed	Da Po	atum bint	□ C Mor	Cover/ nument	X Inne Casi	er ing	Datum Elev.:		m	□ L ⊠ A	ocal HD
PURGINO	GΙ	DETAILS																	
Method:		Low Flow Pump	V	Dat	e: 22/	/09/2016			Ur S.I	ndertako Petts	en B	Sy:		Flov	v Rate:			0.	07L/s
PID readi	ing	: N/Ap	opm	Wa	ter C	olumn:		4.2m	Re	eq Purge	e Vo	d. ¹ :	-L	Actu	ual Vol.	Purg	ed:		-L
		Time			Dep	th to LNAP	L (m)	De	epth to v	vate	er (m)		Dep	th to DN	NAPL	(m)		
Start		10:15			-				1.8	83				-					
Finish		11:30			-				1.8	83									
PURGIN	G N	MEASUR	EMF	ENTS	2														
PURGING MEASUREMENTS 2Vol. Purged (L)Elapsed (ms/cm)EC (mS/cm)Temp. (°C) (°C)Salinity (ppm)pHDO %SatDO (ppm)I												Eh (mV)	Тι	Com urbidi Odo	ments ty, Co our, et	5 –)lour, c.			
1		4		2.63	3	25.7		-		7.06		-	1.0	02	-33		Cole	ourles	ss,
2		8		2.6		25.7		-		7.05		-	1.0	02	-30		dour Ispen	less, s ded s	some olids
3		13		2.4		25.8		-		7.04		-	0.9	94	-36				
4		17		2.41		25.7		-		7.06		-	0.9	94	-35				
5		20		2.36	5	25.6		-		7.07		-	0.9	93	-35				
6		24		2.38	3	25.5		-		7.10		-	0.9	91	-37				
7		29		2.38	3	25.4		-		7.11		-	0.3	88	-39				
				+/-3.0	%	+/-3.0%				+/- 0.1		+/-10%	+/-3	.0%	+/- 10 mV	7			
Comment	ts (e	e.g. condit	tion (of hea	dwor	·ks, colour, o	odou	r):											
SAMPLI	NG	DETAIL	'S							Sample	e ID	: MW2							
Date 22	2/0	9/2016		Tim	e:		11	:00		Underta	aken	By: S Pett	s		-				
PID readin	ıg:	-]	ppm	Dep	th to l	LNAPL:		- 1	m	Depth t	o W	ater:	1	.83 m	Depth	to DN	APL		- m
Flow Rate:	:	0.07L/s	Vol. Ren	noved	:		7L	Sampli	ng	Method:		Low Flo)W						
Temp.:		25.4 °C	DO	: 0.88		-p n	pm 1g/L	pH:		6.11		EC:		2.38	mS/cm	Eh:			39mV
			DO	:		-%	Sat					Salinity:			-ppm mg/L				
Comment	ts (e	e.g. contai	iners	, filtr	ation)):		I				I			<u>5</u> , L				
CoC Nu	mb	ber: 201	6092	2		Check	ked b	y:	S P	Petts			Da	ate:	22/0	9/2016			

1 Bores to be purged dry, until pH, T and EC readings stabilise or a minimum of 3 to 5 times the water column volumes. Water column volumes can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID - 2 L/m; 100 mm ID 8 L/m.



Client: CGC	3								BO	RE II): MW3		
Project: Poi	int Moore	Groundw	ater As	sessment					Job	No.:	6134772	2	
Location: A	strolabe L	n		Casing	diam	eter:		50 mm	n Tot	al Dej	pth:		5.90 m
BORE CON	ISTRUCT	ION											
Head- works Fl	l 🗆 lush Me	onument		Casing]	□ Locke	ed	Datum Point	Cover/ Monument	⊠ Inr Casin	ner g	Datum Elev.:	m	□ Local ⊠ AHD
PURGING	DETAILS												
Method:	Low Flow	Da	te: 2016	5			Undertak S.Petts	en By:		Flov	w Rate:		0.09L/s
PID reading	g: N/Ap	pm Wa	ater Col	umn:		m	Req Purg	e Vol. ¹ :	L	Act	ual Vol.	Purged:	L
	Time	•	Depth	n to LNAP	L (m)	Depth to	water (m)		Dep	th to DN	NAPL (m)	
Start	1200		-				2.68			-			
Finish	1240		-				2.68						
PURGING	MEASURI	EMENTS	S ²										
Vol. Purged (L)	Elapsed Time (min	n) EC	C To	emp. (°C)	Sa (j	alinity ppm)	рН	DO %Sat	DO (p	pm)	Eh (mV)	Cor Turbic Od	nments — lity, Colour, our, etc.
1	10	2.8	4	27.9		-	7.42	-	0.4	-2	-170	Colou	urless, SO ₂
2	14	2.8	0	28.0		-	7.39	-	0.2	1	-202	odour,	suspended solids
3	18	2.7	2	28.1		-	7.35	-	0.1	3	-227		
4	22	2.6	2	28.1		-	7.34	-	0.0	9	-240		
5	26	2.5	6	28.1		-	7.34	-	0.0	6	-249		
		+/-3.0)%	+/-3.0%			+/- 0.1	+/-10%	+/-3.0	0%	+/- 10 mV	7	
Comments ((e.g. condit	ion of he	adwork	s, colour, o	odou	r):							
SAMPLING	G DETAIL	S					Sampl	e ID: MW3					
Date 22/0)9/2016	Tir	ne:12:30				Undert	aken By: S Pe	etts				
PID reading:	- I	opm De	pth to LN	NAPL:		- 1	n Depth	to Water:	2	.68 m	Depth	to DNAPL	- m
Flow Rate:	0.09 L/s	Vol. Removed	l:		9L	Sampli	ng Method	: Low I	Flow Pu	mp			
Temp.:	27.8 °C	DO:		-p n	opm ng/L	pH:	5.95	EC:		5.95	mS/cm	Eh:	-10mV
		DO:		71.8%	6Sat			Salinity	:		-ppm mg/L		
Comments ((e.g. contai	ners, filt	ration):								0		
All metals fi	ield filtered	1											
CoC Num	ber: 2010	50922		Check	ked b	y:	S Petts		D	ate:	22/0	9/2016	

Bores to be purged dry, until pH, T and EC readings stabilise or a minimum of 3 to 5 times the water column volumes. Water column volumes can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID - 2 L/m; 100 mm ID 8 L/m.

2 Calibration details to be recorded in the instrument -specific calibration book, or in field notes as required by local procedures.



Client: CO	G											BO	RE II): SHP8		
Project: P	oint Moor	e Grou	undwa	ater As	sessment							Job	No.:	6134772		
Location:	Coxswain	Park			Casing d	liame	eter:			5	50 mm	Tota	al Dej	pth:		4.76 m
BORE CO	NSTRUC	TION														
Head- works	□ I Flush I	⊠ Monun	nent		asing C L] .ocke	ed	Datu Point	m t]	□ Co Monu	over/ ument	Inr Inr Casin	ner g	Datum Elev.:	m	□ Local ⊠ AHD
PURGING	G DETAIL	'S														
Method:	Low Flow	V	Dat	e: 2016				Unde S.Pet	ertake ts	en By	7:		Flov	w Rate:		L/s
PID reading	ng: N/A	ppm	Wa	ter Col	umn:		1.8 m	Req I	Purge	e Vol	· ¹ :	-L	Act	ual Vol.	Purged:	8L
	Time			Depth	to LNAP	L (m)	Dept	h to w	vater	· (m)		Dep	th to DN	NAPL (m)
Start	1310			-				2.08					-			
Finish	1340			-				2.08								
PURGING	G MEASU	REME	ENTS	2												
Vol. Purged (L)	Elapsed T (min)	lime	EC (mS/c	m) Te	emp. (°C)	Sa (j	dinity ppm)	p	н	DO	%Sat	DO (p	pm)	Eh (mV)	Co Turbi Oc	mments – dity, Colour, lour, etc.
1	5		11.4	.9	27.6		-	6.	89			0.6	7	-285	Colo	urless, SO ₂
2	10		8.95	5	27.5			6.	74			0.2	2	301	odoui	, suspended solids
3	15		8.94	4	28			6.	73			0.0	9	-308		
4	17		9.04	4	27.7			6.	67			0.1	6	287		
5	20		9.00	5	27.7			6.	.63			0.0	7	300		
6	23		8.97	7	27.7			6.	58			0.0	5	-306		
			+/-3.09	%	+/-3.0%			+/-	0.1	+/	-10%	+/-3.0)%	+/- 10 mV	7	
Comments	s (e.g. cond	lition o	of hea	dworks	s, colour, o	doui	r):									
SAMPLIN	IG DETAI	LS						Sa	ample	e ID:	SHP8					
Date 22	2/09/2016		Tim	e:		13	30	U	nderta	aken	By: S Pe	tts				
PID reading	g:	- ppm	Dep	th to LN	APL:		- 1	n De	epth to	o Wa	ter:	2	.08 m	Depth	to DNAPI	- m
Flow Rate:	0.06L/s	Vol. I	Remov	ved:		6L	Sampli	ng Me	ethod:							Low Flow
Temp.:	27.7 °C	DO: (0.05		-p) m	pm g/L	pH:	6.	58		EC:		8.97	mS/cm	Eh:	-306mV
		DO:			- %	Sat					Salinity:			-ppm mg/L		
Comments	s (e.g. cont	ainers	, filtr	ation):												
All metals	field filter	ed														
CoC Nui	mber: 20	16031	0 ntil nH	T and F	C readings s	ed b	y: se or a mi	S Petts	s 1 of 3 t	o 5 tir	nes the we	Dater colu	ate:	22/9/16	ater colum	volumes

can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID - 2 L/m; 100 mm ID 8 L/m. 2

Calibration details to be recorded in the instrument -specific calibration book, or in field notes as required by local procedures.



Client: CG	G								BO	RE II	D: MW1		
Project: Po	oint Moor	e Gro	undwat	er Assessment					Job	No.:	6134772		
Location: I	Marine To	errace		Casing	diamet	er:	Locatio	on: Marine	Tot	al De	pth:	5.91	
BORE CO	NSTRUC	TION	[
Head- works	z Flush	□Mor	ument	Casing	□ Locked	I I I	Datum Point	Cover/ Monument	🗷 In Casin	ner Ig	Datum Elev.:	m	□ Local ⊠ AHD
PURGING	DETAIL	S											
Method:	Low Fle Pump	ow	Date:	2016		T S	Undertak S.Petts	en By:		Flo	w Rate:		0.006L/s
PID readin	ng: N/A	Appm	Wate	r Column:	3.5	1 m	Req Purg	e Vol. ¹ :	-L	Act	ual Vol.	Purged:	-L
	Time		1	Depth to LNAP	^P L (m)	I	Depth to v	water (m)		Dep	oth to DN	NAPL (m)	
Start	8:30		-			2	2.41			-			
Finish	1000		-			2	2.41			-			
PURGING	MEASU	REM	ENTS ²										
Vol. Purged (L)	Elaps Time (r	ed nin)	EC (mS/cm)	Temp. (°C)	Turi (N	oidity ΓU)	рН	DO %Sat	D((pp)) m)	Eh (mV)	Con Turbid Od	nments – ity, Colour, our, etc.
1	5		20.7	28.1	6	.7	7.34	7.4	-		-289	S2O o	dour, some
2	8		17.23	27.9	9	.7	7.32	11.4	-		-279	TSS,	colourless
3	10		16.26	27.9	9	.1	7.44	23.6	-		-274		
4	12		16.11	27.9	7	.5	7.46	14.8	-		-273		
5	15		15.89	27.9	8	.4	7.48	15.4	-		-272		
6	18		15.82	27.9	5	.2	7.50	15.4	-		-271		
			+/-3.0%	+/-3.0%			+/- 0.1	+/-10%	+/-3.	0%	+/- 10 mV	7	
Comments	(e.g. cond	dition	of head	works, colour,	odour)	:							
SAMPLIN	G DETAI	ILS					Sampl	e ID: MW1					
Date 18/	/10/2016		Time:	10:00			Undert	aken By: S F	etts				
PID reading	:	- ppm	Depth	to LNAPL:		- m	Depth	to Water:	2	.41 m	Depth	to DNAPL	- m
Flow Rate:	0.006 L/s	Vol.	Removed	l:	6L	Samplin	g Method:	: Low	Flow				
Temp.:	27.9 °C	DO:		-p n	ng/L	pH:	7.50	EC: 15	5.82		mS/cm	Eh:	-271 mV
		DO:		15.4 %	6Sat			Turbid	lity	5	.2 NTU		
Comments	(e.g. cont	ainer	s, filtrat	ion):QC01_201	61018	(blind),	QC02_2	0161018(spl	lit), QC	03_20)161018	(RB)	
QC04_201	61018(FB)											
CoC Nun	nber: 20)16101	8	Chec	ked by	: 5	Petts		D	ate:	18/1	0/2016	
Bores to	be purged d	iry, unt	ирн, Та	nd EC readings sta	idilise or	a minim	um of 3 to 5	b umes the wat	ter colum	n volu	mes. Wate	r column vo	umes

can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID - 2 L/m; 100 mm ID 8 L/m. Calibration details to be recorded in the instrument –specific calibration book, or in field notes as required by local procedures.

2



Client: CG	G								BO	RE ID): MW2		
Project: Po	oint Moore	Grou	ndwat	er Assessment					Job	No.: (6134772		
Location: (Captain Cre	es		Casing	diam	eter:		50 mm	n Tot	al Dep	oth:	6.02	
BORE CO	NSTRUCT	ION											
Head- works F	⊠ □ Flush M	onum	nent	□ Casing	🗆 Locke	ed 1	Datum Point	Cover/ Monument	⊠ Inn Cas	er ing	Datum Elev.:	m	□ Local ⊠ AHD
PURGING	DETAILS												
Method:	Low Flov Pump	V	Date:	: 18/10/2016			Undertak S.Petts	en By:		Flow	v Rate:		0.005L/s
PID readin	ng: N/Ap	pm	Wate	er Column:	4	.12 m	Req Purg	e Vol. ¹ :	-L	Actu	ial Vol.	Purged:	-L
	Time		1	Depth to LNAP	L (m	i)]	Depth to v	water (m)		Dep	th to DN	NAPL (m)	
Start	1015			-			1.90			-			
Finish	1050			-		-	1.90						
PURGING	MEASUR	EME	ENTS ²										
Vol. Purged (L)	Elapsed Time (min	n) (EC (mS/cm)) Temp. (°C)	Tu (1	rbidity NTU)	рН	DO %Sat	D (pj	O pm)	Eh (mV)	Com Turbidit Odor	nents – y, Colour, ır, etc.
1	5		2.17	20.2		-0.8	7.84	17.1		-	-179	S2O	Odour,
2	8		2.16	20.3		-1.7	7.91	20		-	-177	colourle	ess, slight SS
3	11		2.16	20.3		-0.3	8.09	19.0		-	-180		
4	14		2.18	20.3		-0.7	7.97	18.5		-	-180		
5	17		2.17	20.4		-0.8	7.94	18.2		-	-182		
6	20		2.17	20.4		-1.5	7.90	19.0		-	-183		
			+/-3.0	+/-1.0			+/- 0.1	+/-10%	+/-	-1.0	+/- 10 mV	·	
Comments	(e.g. condit	ion o	of head	works, colour,	odou	r):							
SAMPLIN	G DETAIL	S					Sampl	e ID: MW2					
Date 18/	/10/2016		Time:	: 10:40			Undert	aken By: S P	etts				
PID reading	;; - I	opm	Depth	n to LNAPL:		- n	n Depth (to Water:	1	.90 m	Depth	to DNAPL	- m
Flow Rate:	0.005 L/s	Vol. Ren	noved:		6L	Samplir	ng Method	Low 1	Flow				
Temp.:	20.4 °C	DO:		-p n	pm ng/L	pH:	7.84	EC: 2.1	7	1	mS/cm	Eh:-183	mV
		DO:	:	19.0%	Sat			Turbidi	ity	-1.	5 NTU		
Comments	(e.g. contai	ners,	, filtrat	tion):									
				1									
CoC Nun	nber: 201	61018	8	Chec	ked b	y:	S Petts		D	ate:	18/10	0/2016	

Bores to be purged dry, until pH, T and EC readings stabilise or a minimum of 3 to 5 times the water column volumes. Water column volumes can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID - 2 L/m; 100 mm ID 8 L/m.



2

Chent: CG(_															
Chefit: CGG BOKE ID: MW3 Project: Point Moore Groundwater Assessment Job No • 6134772																
Location: Astrolabe Ln Casing diameter:										50 mm	mm Total Denth: 5 87 m					
RODE CONSTRUCTION																
Head- Casing Casing Datum Cover/ Inner Datum Cover/									Local							
works Fl	lush Mo	onume	ent]	Locke	ed	Po	int	Mor	nument	Casin	g	Elev.:	n		AHD
PURGING DETAILS																
Method:	Low Flow	/	Date: 2	2016			Un S.I	dertak Petts	en B	y:		Flov	v Rate:		0.	004 L/s
PID reading	g: N/Ap	pm	Water	Column:	3.	.16 m	Re	q Purg	e Vo	l. ¹ :	L	Actu	ual Vol.	Purged:		L
	Time		De	epth to LNAP	L (m)	De	pth to v	wate	r (m)		Dep	th to DN	APL (n	I)	
Start	1100		-				2.7	'1				-				
Finish	1140		-				2.7	/1				-				
PURGING	MEASURI	EMEN	NTS ²			1										
Vol. Purged (L)	Elapsed Time (min	n) (1	EC uS/cm)	Temp. (°C)	Tur (NT	bidity U)		рН	D	O %Sat	D((pp)) m)	Eh (mV)	Co Turb O	omme idity, dour,	nts – Colour, etc.
1	5		2.24	22.7 21.2		21.2		7.96		17.4	-		-137	N	o odo	urm
2	9		2.22	22.7 17.3		17.3		-		23.6	-		-147	colo	urless SS	, some
3	12		2012	22.8		9.7		-		25.1	-		-145			
4	15		2055	22.7		8.3		7.65		25.9	-		-147			
5	19		2063	22.7		8.5		8.02		25.3	-		-149			
6	23		2062	22.7		6.4		7.96		26.0	-		-151			
		н	+/-3.0%	+/-3.0%				+/- 0.1	-	+/-10%	+/-3.0)%	+/- 10 mV			
Comments	(e.g. condit	ion of	' headw	orks, colour,	odou	r):										
Water meter	issues, pH	readir	ng error													
SAMPLING	G DETAIL	S						Sampl	e ID	: MW3						
Date 18/1	10/2016		Time:		11	:30		Undert	aken	By: S Pet	tts		_			
PID reading:	: - r	opm	Depth t	o LNAPL:		- 1	m	Depth	to W	ater:	2	.71 m	Depth	to DNAP	L	- m
Flow Rate:	0.004 L/s	Vol. Remo	oved:		6 L	Sampli	ing 1	Method	:	Low Fl	ow Pu	mp				
Temp.:	22.7 °C	DO: -ppm mg/L pH:						7.96	96 EC:		2062	uS/cm	Eh:	-	151 mV	
		DO:		26.0 %	6Sat					Turbidit	У	6.	4 NTU			
Comments (e.g. containers, filtration):																
All metals field filtered																
CoC Number: 20161018 Checked by: S Petts Date: 18/10/2016										- 4	Da	ate:	18/1	0/2016	.1	

Bores to be purged dry, until pH, T and EC readings stabilise or a minimum of 3 to 5 times the water column volumes. Water column volumes can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID - 2 L/m; 100 mm ID 8 L/m.

Calibration details to be recorded in the instrument -specific calibration book, or in field notes as required by local procedures.



2

Client: CGG BORE ID: SHP8														
Project: F	I roject: rom moore Groundwater Assessment Job No.: 0154//2 L costion: Covewain Park Cosing diamataw 50 mm Total Danth: 5.95 mm													
Location: Coxswain Park Casing diameter:								50 mm	Tot	al Dej	pth:	5.	85 m	
BORE CO	ONSTRUC	TION	-											
Head- works	□ I Flush I	≭ Monur	nent	□ Casing	□ Locke	ed	Datum Point	□ Cover/ Monument	Inr Casin	ner g	Datum Elev.:	m	□ Local ⊠ AHD	
PURGING DETAILS														
Method: Low Flow Date: 2016				Undertak S.Petts	en By:		Flov	w Rate:		0.004 L/s				
PID reading: N/A Water Column: 3.68 m				.68 m	Req Purg	e Vol. ¹ :	-L	Act	ual Vol. I	Purged:	8L			
	Time			Depth to LNAP	PL (m)	Depth to	water (m)		Dep	oth to DN	APL (m)		
Start	1145						2.17			-				
Finish	1230						2.17							
PURGIN	G MEASU	REMI	ENTS ²	2						I				
Vol. Purged (L)	Elapsed T (min)	Time	EC (mS/cm	Temp. (°C)	Temp. (°C) Turbidity (NTU)		рН	DO %Sat	D((pp)) m)	Eh (mV)	Con Turbid Odo	iments – ity, Colour, our, etc.	
1	5		3.69	20.2	20.2 -3.7		7.63 11.1		-		-226	Colour	less, slight	
2	9		3.71	20.3		-3.2	7.66	13.8	3.8 _		-225	S2O oc	S2O odour, some SS	
3	14		3.74	20.3		-3.2	7.65	13.8 .		224				
4	17		3.77	20.2		-3.2	7.69	7.69 15.6		-				
5	20		3.79	20.2		-3.7	7.70	16.3	-		-222			
			+/-3.0%	+/-3.0%			+/- 0.1	+/- 0.1 +/-10%			+/- 10 mV			
Comment	s (e.g. cond	lition	of head	lworks, colour,	odou	r):		-1 1				1		
SAMPLI	NG DETAI	ILS					Samp	e ID: SHP8						
Date 1	8/10/2016		Time	: 1220			Under	taken By: S Pe	tts					
PID readir	ıg:	- ppm	Dept	h to LNAPL:		- 1	n Depth	to Water:	2	.17 m	Depth (to DNAPL	- m	
Flow Rate:	0.004 L/s	Vol.	Remove	ed:	5L	Sampli	ng Method	: Low F	low					
Temp.:	20.2 °C	DO:		-ppm pH:			7.70	EC:	3.79 m ⁶		mS/cm	Eh:	-222 mV	
DO: 16.3 %Sat Turbidity -3.7 NTU														
Comments (e.g. containers, filtration):														
All metals field filtered														
CoC Nu	mber: 20	16101	8	Chec	ked b	y:	S Petts	-	D	ate:	18/10	/2016		

can be calculated from the following casing volumes per unit length: 40 mm ID - 1 L/m; 50 mm ID - 2 L/m; 100 mm ID 8 L/m. Calibration details to be recorded in the instrument –specific calibration book, or in field notes as required by local procedures.

Appendix E – Quality Assurance and Quality Control

E1 Introduction

The Quality Assurance/Quality Control (QA/QC) procedures are based on the DER Assessment and management of contaminated sites (DER 2014), ASC NEPM (NEPC 1999), AS 4482.1 – 2005 (Standards Australia 2005), AS 4482.2 – 1999 (Standards Australia 1999) and AS 5667 – 1998 (Standards Australia 1998). QA involves all of the actions, procedures, checks and decisions, undertaken to ensure the representativeness and integrity of samples and accuracy and reliability of analytical results. QC involves protocols to monitor and measure the effectiveness of QA procedures.

E2 Field Work Program

E2.1 Field Quality Assurance Procedures

All field work was conducted with reference to the DER (2014) and GHD's Standard Field Operating Procedures, which ensures all environmental samples are collected by a set of uniform and systematic methods, as required by GHD's QA system. The procedures undertaken as part of the soil and groundwater assessments includes:

- Decontamination Procedures: Included the use of new disposable gloves for the collection of each sample, decontamination of the sampling equipment between each sampling event and the use of sampling containers provided by the laboratory;
- Sample Identification Procedures: Collected samples are immediately transferred to sample containers of appropriate composition and preservation for the required laboratory analysis. All sample containers were clearly labelled with a sample number, sample location, sample depth, sample date and sampler's initials. The sample containers were transferred to a chilled esky for sample preservation prior to and during shipment to the testing laboratory;
- Chain of Custody Information Requirements: A chain of custody form was completed and forwarded to the testing laboratory with all sample batches and in some instances emailed separately to the laboratory prior to the Esky's arrival to the laboratories. The purpose of this was to provide the laboratory notification of samples requiring extraction upon arrival; and
- Sample blind duplicate and split duplicate, rinsate blank and field blank frequency.

E2.2 Groundwater Sampling and Analysis Quality Control

The DER Contaminated Sites Management Guidelines outlines the requirements for Quality Control (QC) sampling protocols. The QC samples collected during the investigation are described below and presented in **Table E1**.

Blind (Intra-laboratory) Duplicates

Blind (Intra-laboratory) duplicate samples were used to identify the variation in the analyte concentration between samples collected from the same sampling point and the repeatability of the laboratory's analysis.

Split (Inter-laboratory) Duplicates

Inter-laboratory duplicate samples provide an indication of the repeatability of the results between laboratories.

Rinsate Blanks

Rinsate blank samples are water samples collected from decontaminated, re-used field equipment and used primarily to assess whether the decontamination procedure is effective and if cross-contamination has led to positive observations in subsequent samples.

Rinsate samples were collected by pouring laboratory supplied ultra-high purity rinsate water over the probe of water quality meters and interface probes or running it through the pump and collecting it in laboratory supplied containers. Rinsate samples are then transferred to a chilled esky for sample preservation prior to and during shipment to the testing laboratory.

Field Blanks

Field blank samples were used to estimate contamination of a sample during the collection procedure.

Field blank samples are collected by pouring laboratory supplied rinsate water into laboratory supplied bottles onsite. Field blank samples are then transferred to a chilled esky for sample preservation prior to and during shipment to the testing laboratory.

Table E1 Quality Control Sampling Frequency

Sample	Sample Collection Frequency
Blind (Intra-laboratory) Duplicates	1:20 samples
Split (Inter-laboratory) Duplicates	1:20 samples
Equipment Rinsate Blanks	1 per matrix per day
Field Blanks	1 per matrix per day

E2.3 Relative Percent Difference Calculations

Blind and split duplicate samples were assessed by calculating the relative percentage difference (RPD) between the primary, blind and split samples.

A quantitative measure of the accuracy of the analytical results reported was made by calculating the RPDs between the primary, blind and split results in accordance with the procedure described in AS 4482.1 - 2005. According to AS 4482.1 - 2005 typical RPDs are expected to range between 30% and 50%; however, this may be higher for organics and for low concentrations of analytes. GHD uses 50% as the general assessment criteria.

Where a result was reported below the laboratory limit of reporting (LOR) for one of the duplicate pair samples, the sample will be assigned the concentration of the LOR for RPD calculation purposes.

E3 Laboratory Program

Samples were dispatched to various nominated NATA accredited laboratories for the analytes of concern, with Chain of Custody documentation acknowledging the transference of samples from the Site to the laboratory. The primary and blind duplicate samples were dispatched to ALS Laboratory Group. The split duplicate samples were dispatched to SGS Environmental.

E3.1 Laboratory Analytical Methods

Laboratory methods used by the primary and secondary laboratories are considered suitable for environmental contaminant analysis and are based on established internationally recognised procedures. Each of the laboratories were NATA accredited for the proposed analysis.

Analyte	Method	Limits of Reporting (LOR)	Holding Times
Ammonia	APHA 4500-NH 3 B/C	0.05 mg/L	1 day
Nitrite	APHA 4110/4500-NO ₂	0.05 mg/L	2 days
Nitrate	APHA 4110	0.05 mg/L	2 days
Total Nitrogen	APHA 4500-N	0.05 mg/L	28 days
Total Dissolved Solids	APHA 2540C	10 µg/L	7 days
E. coli (MF)	AS4275:21-2005	1 cfu/100 mL	24 hours

Table E2Laboratory Methods, LORs and Laboratory Holding Times forGroundwater Analysis

E3.2 Laboratory Quality Control Procedures

The following laboratory QC procedures were used during the investigation.

Laboratory Duplicate Samples

Laboratory duplicate sample analysis were the analysis of a laboratory derived duplicate sample from the process batch, at a rate equivalent to one in twenty samples per analytical batch, or one sample per batch if less than twenty samples are analysed in a batch. A laboratory duplicate provides data on the analytical precision and reproducibility of the analytical results.

The permitted ranges for the RPD of Laboratory Duplicates, as specified in ALS Method QWI-EN/38 are dependent on the magnitude of results in comparison to the level of reporting. For RPD results that are less than ten times the limit of reporting (LOR) there are no limits, for results between ten and twenty times the LOR the adopted criteria are 0% - 50% and for results greater than twenty times LOR the criteria are 0% - 20%.

Method Blank Samples

Method or analysis blank sample analysis is the analysis of a sample that is as free as possible of the analytes of interest, but has been prepared the same as the samples under investigation. The analysis is to ascertain if laboratory reagents, glassware and other laboratory consumables contribute to the observed concentration of analytes in the process batch. If below the maximum acceptable method blank (20% of the practical quantitation limit), the contribution is subtracted from the gross analytical signal for each analysis before calculating the sample analyte concentration. GHD notes that the subtraction of the method blank concentration is not appropriate for some organic analytes.

Laboratory Control Spike Samples

Laboratory control spike analysis is the analysis of either a reference material or a control matrix fortified with analytes representative of the analyte class. The purpose of laboratory control spike samples is to monitor method precision and accuracy independent of the sample matrix. Typically, the percent recovery of the laboratory control spike sample is compared to the dynamic recovery limits based on the statistical analysis of the processed laboratory control spike sample analysis.

Matrix Spike Samples

Matrix spike sample analysis is the analysis of one or more replicate portions of samples from the batch, after fortifying the additional portion(s) with known quantities of the analyte(s) of interest. The percent recovery of target analyte(s) from matrix spike samples is used to determine the bias of the method in the specific sample matrix.

Surrogate Spike Samples

Surrogate spike samples are samples with known additions of known amounts of compounds, which are similar to the analytes of interests in terms of extractability, recovery through clean-up procedures and response to chromatographic or other measurement. Surrogate compounds may be alkylated or halogenated analogues or structural isomers of analytes of interest. The purpose of surrogate spikes, which are added immediately before the sample extraction step, is to provide a check for every analysis that no gross processing errors have occurred, which could have led to significant analyte loss or faulty calculation.

Internal Standards

Internal standards are known additions of known amounts of compounds which are not found in real samples, will not interfere with quantification of analytes of interest and may be separately and independently quantified. The purpose of internal standards in instrumental techniques is to provide independent signals, which serve to check the consistency of the analytical step. Internal standards are often used for organic compounds and some inorganic compounds.

E4 Data Management

Laboratory results were reviewed within five working days of receipt from the laboratory.

The individual testing laboratory conducted an assessment of the laboratory QC program, internally, however the results will also be independently reviewed and assessed by GHD, to ensure that no issues exist with the data prior to undertaking any data interpretation. GHD reviewed the Laboratories' quality control certificate and identified any outliers. The outliers identified are discussed in the sections below.

Following receipt of data, the groundwater results were converted to ESDAT format and compared against the adopted criteria provided in Section 6.1 Subsequent to this the ESDAT format was modified to ensure that only test pits location relevant to the Site were presented. All data will be stored in an electronic format as well as produced in hard copy. The hard copies were stored along with chain of custody (CoC) forms in the project file. The CoC forms and laboratory certificates are presented in Appendix F.

E5 Sampling and Analysis Quality Control Results: Groundwater

E5.1 Sample Duplicates

The blind and split duplicated QC samples collected for the 2016 GME is listed in Table E3. The split sample was not analysed for microbes as a result of issues associated with sample receipts department. The split sample was processed outside the 24 hour holding and therefore the laboratory was advised not to proceed with analysing this sample for microbes.

Primary Sample	QC Sample	Description	Date	Analyses
MW1	QC01_20160824	Blind – ALS	24/08/2016	Ammonia, BOD, E. coli, Faecal coliforms, total coliforms, nitrates, nitrite, TDS, TKN, total nitrogen, total phosphorus
MW1	QC02_20160824	Split – SGS	24/08/2016	Ammonia, BOD, nitrates, nitrite, TDS, TKN, total nitrogen, total phosphorus
MW1	QC01_20160923	Blind – ALS	23/09/2016	Ammonia, BOD, E. coli, Faecal coliforms, total coliforms, nitrates, nitrite, TDS, TKN, total nitrogen, total phosphorus
MW1	QC02_20160923	Split – mpl	23/09/2016	Ammonia, BOD, E. coli, Faecal coliforms, total coliforms, nitrates, nitrite, TDS, TKN, total nitrogen, total phosphorus
MW1	QC01_20161018	Blind – ALS	18/10/2016	Ammonia, BOD, E. coli, Faecal coliforms, total coliforms, nitrates, nitrite, TDS, TKN, total nitrogen, total phosphorus
MW1	QC02_20161018	Split – mpl	18/10/2016	Ammonia, BOD, E. coli, Faecal coliforms, total coliforms, nitrates, nitrite, TDS, TKN, total nitrogen, total phosphorus

 Table E3
 Sample Duplicates

Rinsate and Field Blanks

The rinsate and field blanks collected during the groundwater investigation undertaken in 2016 are listed in Table E4.

QC Sample	Description	Date	Analysis
QC03_20160824	Rinsate blank	24/08/2016	Ammonia, BOD, Nitrate, Nitrite, TDS, TKN, total nitrogen and total phosphorus
QC04_20160824	Field blank	24/08/2016	Ammonia, BOD, Nitrate, Nitrite, TDS, TKN, total nitrogen and total phosphorus
QC03_20160922	Rinsate blank	23/09/2016	Ammonia, BOD, Nitrate, Nitrite, TDS, TKN, total nitrogen and total phosphorus
QC04_20160922	Field blank	23/09/2016	Ammonia, BOD, Nitrate, Nitrite, TDS, TKN, total nitrogen and total phosphorus
QC03_20161018	Rinsate blank	18/10/2016	Ammonia, BOD, Nitrate, Nitrite, TDS, TKN, total nitrogen and total phosphorus
QC03_20161018	Field blank	18/10/2016	Ammonia, BOD, Nitrate, Nitrite, TDS, TKN, total nitrogen and total phosphorus

Table E4 Rinsate and Field Blanks

A summary of the reported detects within field and rinsate blank samples is provided:

- QC03_20160824 (rinsate sample) detected BOD above the LOR (5 mg/L). Primary samples reported BOD to range between 4 mg/L and 7 mg/L. Considering two primary samples reported BOD below 5 mg/L, it is unlikely that this detect would impact the overall integrity of the data;
- QC03_20160922 (rinsate sample) detected TDS above the LOR (20 mg/L). Considering the concentrations of Primary samples which reported TDS to range between 1,320 mg/L to 22,400 mg/L, which were reported in three orders of magnitude greater this QC sample; therefore, it is considered that this would no impact on the integrity of the data;
- QC03_20161018 (rinsate sample) and QC04_20161018 (field blank) reported total dissolved solids, ammonia marginally above the LOR. It is likely that the source of this contamination is associated within laboratory supplied water and therefore it is considered that this would no impact on the integrity of the data;

 QC04_20161018 (field blank) reported nitrate and nitrite, the source of this contamination is unknown. Notwithstanding this, the detects are equal to the LOR and considering that two primary samples reported both these CoPC below the LOR. It is therefore considered unlikely that this detect would impact the overall integrity of the data;

Relative Percentage Difference Calculation Results

During the 2016 GME, 79 RPDs were calculated on field samples with six RPDs exceeding the recovery limit of 50%. RPDs for the groundwater investigation are presented in Appendix G and the RPD exceedances are summarised in Table E5.

Table E5 Groundwater RPDs Greater than 50%

Primary/Duplicate Sample Name	Analytes with RPD exceeding 50%	Comment				
MW1/QC01_20160824	Coliform (200%)	The primary sample (250, 000 cfu/100 mL) was reported higher than the blind sample (200 cfu/100 mL). These two samples are in four orders of magnitude and the difference between the two samples is a result of the primary sample having a higher amount of particulate matter than the blind sample.				
	Faecal Coliform (138%)	The blind sample (1100 cfu/100mL) was higher than the primary (200 cuf/100mL). The difference between the primary and blind is a result higher level of particulate, which may be masked by overgrowth of non-				
	<i>E. coli</i> (138%)	target organisms.				
MW1/QC02_20160923	BOD (144%)	The primary sample reported a concentration (31 mg/L) above the LOR, while the split duplicated reported a concentration below the LOR (<1 mg/L). The elevated RPD suggested that the primary laboratory may have over reported the BOD concentration. By considering the primary sample concentration when evaluating the groundwater quality, GHD considers that a conservative approach has been implemented in this regard and secondary laboratory is unlikely to have impacted upon data quality and the overall conclusions drawn.				
	Plate Count 36°C (67%)	The split sample (340 cfu/100 mL) was higher than the primary. The different incubation temperatures used by the primary and secondary laboratory, likely has attributed to the different plate count. With consideration to the above, the elevated RPD value for this duplicate is not considered to have significantly impacted upon the overall data quality.				
	Plate Count 22°C (67%)	The split sample (260 cfu/100 mL) was higher than the primary. The different incubation temperatures used by the primary and secondary laboratory, likely has attributed to the different plate count. With consideration to the above, the elevated RPD value for this duplicate is not considered to have significantly impacted upon the overall data quality.				

Primary/Duplicate Sample Name	Analytes with RPD exceeding 50%	Comment
	Nitrite as N (197%)	The primary sample was reported below the LOR (0.01 mg/L) while the secondary reported a concentration above the LOR (0.006 mg/L). The RPD is a reflection of the different LOR used by the primary and secondary. The secondary utilises a low LOR, as the detected concentration in secondary is below the primary laboratory's LOR. With consideration to the above, the elevated RPD value for this duplicate is not considered to have significantly impacted upon the overall data quality.
	Nitrate as N (195%)	The primary sample reported a concentration (0.05 mg/L) above the LOR, while the split duplicated reported a concentration below the LOR (<0.005 mg/L). The difference between the primary and secondary are the laboratory limit of reporting. Notwithstanding this, the elevated RPD suggested that the primary laboratory may have over reported the BOD concentration. By considering the primary sample concentration when evaluating the groundwater quality, GHD considers that a conservative approach has been implemented in this regard and secondary laboratory is unlikely to have impacted upon data quality and the overall conclusions drawn.
MW1	BOD (81%)	The primary sample (33 mg/L) was higher than the split sample (14 mg/L). The elevated RPD suggested that the primary laboratory may have over reported the BOD concentration. By considering the primary sample concentration when evaluating the groundwater quality, GHD considers that a conservative approach has been implemented in this regard and secondary laboratory is unlikely to have impacted upon data quality and the overall conclusions drawn

E5.2 Laboratory Quality Control Results

Laboratory Duplicates

There were no RPD outliers for laboratory duplicate samples reported by ALS or SGS

Method Blank Samples

There were no method blank recovery value outliers reported by ALS or SGS. GHD considers that the analyses methods used were free from potential laboratory contamination and that the data is suitable for Site soil characterisation.

Laboratory Control Spike Samples

There were no laboratory control outliers reported by ALS or SGS. GHD considers the results from all analyses performed to be sufficiently accurate and independent of the sample matrix and therefore the data is suitable for soil characterisation.

Matrix Spike Samples

There were no matrix spike outliers reported by ALS or SGS for the 2016 groundwater investigation.

Surrogate Spike Samples

There were no surrogate recovery outliers reported by ALS or SGS for the 2016 GME.

E5.3 Review of Sample Management

Laboratory Holding Times

Analysis of samples within analyte specific holding times serves as a measure of QA. The official certificates of analysis and sample receipt notifications obtained from the contracted laboratories reported groundwater analysis was generally completed within specified holding times, with the non-conformance presented in Table E6.

Table E6 Holding Time Non-conformance

Analysis	Samples	Due for Analysis	Date Analysed	Comment		
Nitrate Nitrogen and Nitrite Nitrogen	QC02	28/09/2016	29/09/2016	GHD compared the QC02 results with the primary. The reported		
Total Dissolved Solids	QC02	31/08/2016	1/09/2016	concentrations appear to be in a similar order of magnitude.		

Chain of Custody Forms and Sample Receipts

GHD used Chain of Custody (CoC) forms during the groundwater investigations to establish the traceability of samples and these are attached in Appendix F. A review of the CoC forms by GHD indicates that all prescribed sample transfer, transport and storage protocols were complied with. Sample receipts are presented in Appendix F.

E6 Quality Control and Quality Assurance Summary

The review of the QA/QC results indicates that, overall, the data reported is considered to be of an acceptable quality upon which to draw meaningful conclusions regarding impacts to groundwater at the Site and are within the scope of this investigation for the following reasons:

- **Data Representativeness:** The sampling methodology provided a uniform and systematic approach to the collection of all environmental samples. Laboratory and field QA/QC procedures were carried out to measure data representativeness. Consequently, data representativeness is considered to have been satisfied;
- **Completeness**: It is considered that the field QA/QC procedures carried out, such as the blind and split sample duplicate collection frequencies, and the analytes tested provides completeness in terms of the required number of field duplicate samples. Laboratory QA/QC sample analysis is considered sufficient to provide a complete overview of QA/QC procedures;
- **Precision:** GHD considers that laboratory results are acceptable for interpretation with regards to the suitability of the site for the proposed land use;
- **Accuracy:** Sampling procedures ensured that collection, preservation and laboratory analytical techniques are appropriate for analysis of environmental contaminants.
 - Rinsate blank, field blank and trip blank sample results have been discussed and results indicate that cross-contamination of samples through collection or transport of the samples to the laboratory is unlikely to have occurred and therefore is not considered to have impacted interpretation of contamination at the site.
- **Comparability:** All field work was conducted with reference to the DER Contaminated Sites Management Series Guidelines and GHD Standard Operating Procedures, which ensured all samples were collected by a set of uniform and systematic methods, as required by GHD's QA system. GHD considers that the laboratory data are of a suitable quality for assessing the environmental status of the site.

Consequently, the review of QA procedures and QC results from the primary and secondary laboratories indicates that the analytical data are considered to be of an acceptable quality upon which to draw meaningful conclusions regarding the impacts to soil and groundwater at the site within the scope of this investigation. Any laboratory issues have been listed above and discussed within their relevant sections.

CHAIN OF CUSTORDY RECORD AND



CLIENTS PEOPLE PERFORMANCE GHD House 239 Adelaide Terrace Perth WA

PO Box Y3106 Telephone 08 6222 8222 Facsimile 08 6222 ANALYSIS REQUEST Perth WA 6832 6004 8555 Page <u>1</u> of <u>1</u> COC No. Point Moore Groundwater Assessment Laboratory:MGT Address: Unit 2, 91 Leach Highway, Kewdale WA 6105 Client Job No. 61/34772 City of Greater Geraldton Laboratory Contact: Natalie Container Laboratory Quote No. Turnaround Time Remarks Analyses ≶ Sample Matrix s-soil/ st-sludge/ Water/ A-Air -Vial/Bag/G-Preservative Unpreserv HCI/ H2SO4/HNO3/Other Total Volume (mL) Type B-Bottle/J-Jar/V-Job Manager (Invoice) Email Address (Results) Andrew Nagle andrew.nagle@ghd.com Glass/P-Plastic steven. petts@ghd.com GHD Sample ID Laboratory Sample ID Date Time Hold ۶ X S 250 15/08/2016 NA MW1_0.5 J 1 X MW1 1.0 15/08/2016 250 S NA 1 J X 15/08/2016 250 MW1_1.5 S 1 NA 1 X MW1_2.0 15/08/2016 S NA 1 250 X MW1 2.5 15/08/2016 S NA 1 250 J X MW1_3.0 15/08/2016 S NA 1 250 Л X 15/08/2016 S 250 MW1 3.5 NA 1 J X MW2_0.5 15/08/2016 S NA 1 250 1 MW2_1.0 X MW2 1.5 15/08/2016 S NA 250 J 1 X 15/08/2016 S MW2_2.0 250 NA 1 X MW2 2.5 15/08/2016 S J NA 1 250 X MW2_3.5 15/08/2016 S NA 1 250 1 X MW2_4.0 15/08/2016 S NA 1 250 Л X MW2 4.5 15/08/2016 S NA 1 250 J X MW2_5.0 15/08/2016 S NA 1 250 X 15/08/2016 S NA 250 MW2_5.5 1

				1			
MW2_6.0	15/08/2016	s	1	NA	1	250	
QC01_15082016	15/08/2016	S	J	NA	1	250	
QC02_15082016	15/08/2016	s	1	NA	1	250	
MW3_0.5	15/08/2016	s	J	NA	1	250	
MW3_1.0	15/08/2016	s	1	NA	1	250	
MW3_1.5	15/08/2016	s	J	NA	1	250	
MW3_2.0	15/08/2016	s	J	NA	1	250	
MW3_2.5	15/08/2016	s	J	NA	1	250	
MW3_3.0	15/08/2016	s	J	NA	1	250	
MW3_3.5	15/08/2016	S	J	NA	1	250	
MW3_4.0	15/08/2016	S	J	NA	1	250	
MW3_4.5	15/08/2016	S	J	NA	1	250	
MW3_5.0	15/08/2016	S	J	NA	1	250	
MW3_5.5	15/08/2016	S	J	NA	1	250	
MW3_6.0	15/08/2016	S	J	NA	1	250	
QC03_15082016	15/08/2016	S	J	NA	1	250	
QC04_15082016	15/08/2016	S	J	NA	1	250	
Sampled by: S. Petts		Date/Ti	me: 15/08,	/2016			Relinquished by: S.Petts Date/ Time: 15/08/2016
Received by:			me:				Relinquished by: Date/ Time:
Received by Lab:			me:				Courier/ Transport Company: NA
Sample Conditions:	Sample Conditions: Remarks:						
· · ·							



CERTIFICATE OF ANALYSIS

Work Order	EP1607778	Page	: 1 of 4
Client	: GHD PTY LTD	Laboratory	Environmental Division Perth
Contact	: MR ANDREW NAGLE	Contact	: Lauren Biagioni
Address	: 999 HAY STREET	Address	: 10 Hod Way Malaga WA Australia 6090
	PERTH WA 6000		
Telephone	: +61 08 6222 8222	Telephone	: 08 9209 7655
Project	: 61/34772 Point Moore Groundwater Assessment	Date Samples Received	: 25-Aug-2016 09:20
Order number	:	Date Analysis Commenced	: 25-Aug-2016
C-O-C number	:	Issue Date	31-Aug-2016 16:11
Sampler	: STEVEN PETTS		Hac-MRA NATA
Site	:		
Quote number	:		Accreditation No. 825
No. of samples received	: 7		Accredited for compliance with
No. of samples analysed	: 7		ISO/IEC 17025 - Testing

This report supersedes any previous report(s) with this reference. Results apply to the sample(s) as submitted. This document shall not be reproduced, except in full.

This Certificate of Analysis contains the following information:

- General Comments
- Analytical Results

Additional information pertinent to this report will be found in the following separate attachments: Quality Control Report, QA/QC Compliance Assessment to assist with Quality Review and Sample Receipt Notification.

Signatories

This document has been electronically signed by the authorized signatories below. Electronic signing is carried out in compliance with procedures specified in 21 CFR Part 11.

Signatories	Position	Accreditation Category
Alini Goundar	Senior Analyst	Perth Microbiology, Malaga, WA
Jeremy Truong	Laboratory Manager	Perth Inorganics, Malaga, WA
Tyrone Cole	Inorganics Preparation Supervisor	Perth Inorganics, Malaga, WA



General Comments

The analytical procedures used by the Environmental Division have been developed from established internationally recognized procedures such as those published by the USEPA, APHA, AS and NEPM. In house developed procedures are employed in the absence of documented standards or by client request.

Where moisture determination has been performed, results are reported on a dry weight basis.

Where a reported less than (<) result is higher than the LOR, this may be due to primary sample extract/digestate dilution and/or insufficient sample for analysis.

Where the LOR of a reported result differs from standard LOR, this may be due to high moisture content, insufficient sample (reduced weight employed) or matrix interference.

When sampling time information is not provided by the client, sampling dates are shown without a time component. In these instances, the time component has been assumed by the laboratory for processing purposes.

Where a result is required to meet compliance limits the associated uncertainty must be considered. Refer to the ALS Contact for details.

Key: CAS Number = CAS registry number from database maintained by Chemical Abstracts Services. The Chemical Abstracts Service is a division of the American Chemical Society.

- LOR = Limit of reporting
- ^ = This result is computed from individual analyte detections at or above the level of reporting
- ø = ALS is not NATA accredited for these tests.
- ~ = Indicates an estimated value.
- CFU = colony forming unit
- MF = membrane filtration
- TDS by method EA-015 may bias high due to the presence of fine particulate matter, which may pass through the prescribed GF/C paper.
- MW002: estimate (~) is reported where the growth of bacteria is counted <10cfu and or >300cfu.
- MW006, estimate (~) is reported where the growth of presumptive bacteria on the filtered membrane is counted <10 cfu and/or >100 cfu.
- MW002 is ALS's internal code and is equivalent to AS4276.3.1.
- MW006 is ALS's internal code and is equivalent to AS4276.7.
- MW007 is ALS's internal code and is equivalent to AS4276.5.



Analytical Results

Sub-Matrix: WATER (Matrix: WATER)	Client sample ID			MW1	QC01_20160824	MW2	MW3	SHP8		
	Client sampling date / time			[24-Aug-2016]	[24-Aug-2016]	[24-Aug-2016]	[24-Aug-2016]	[24-Aug-2016]		
Compound	CAS Number	LOR	Unit	EP1607778-001	EP1607778-002	EP1607778-003	EP1607778-004	EP1607778-005		
				Result	Result	Result	Result	Result		
EA015: Total Dissolved Solids dried at 180 ± 5 °C										
Total Dissolved Solids @180°C		10	mg/L	8890	8800	1480	1090	3320		
EK055G: Ammonia as N by Discrete Analyser										
Ammonia as N	7664-41-7	0.01	mg/L	2.68	2.86	0.53	0.57	0.36		
EK057G: Nitrite as N by Discrete Analyser										
Nitrite as N	14797-65-0	0.01	mg/L	<0.01	<0.01	0.20	<0.01	<0.01		
EK058G: Nitrate as N by Discrete Analys	er									
Nitrate as N	14797-55-8	0.01	mg/L	0.19	0.18	13.8	0.02	0.02		
EK059G: Nitrite plus Nitrate as N (NOx) by Discrete Analyser										
Nitrite + Nitrate as N		0.01	mg/L	0.19	0.18	14.0	0.02	0.02		
EK061G: Total Kjeldahl Nitrogen By Discrete Analyser										
Total Kjeldahl Nitrogen as N		0.1	mg/L	3.0	2.9	3.1	0.7	1.5		
EK062G: Total Nitrogen as N (TKN + NOx) by Discrete An	alyser								
^ Total Nitrogen as N		0.1	mg/L	3.2	3.1	17.1	0.7	1.5		
EK067G: Total Phosphorus as P by Discr	ete Analyser									
Total Phosphorus as P		0.01	mg/L	0.29	0.30	0.21	0.06	0.31		
EP030: Biochemical Oxygen Demand (BC	D)									
Biochemical Oxygen Demand		2	mg/L	6	6	7	4	4		
MW002: Heterotrophic Plate Count										
Heterotrophic Plate Count (22°C)		1	CFU/mL	~49000	~60000	~13000	~70000	910		
Heterotrophic Plate Count (36°C)		1	CFU/mL	~46000	~49000	~11000	~68000	110		
MW006: Faecal Coliforms & E.coli by MF										
Faecal Coliforms		1	CFU/100mL	~200	1100	~400	~400	~100		
Escherichia coli		1	CFU/100mL	~200	1100	~400	~400	~100		
MW007: Coliforms by MF										
Coliforms		1	CFU/100mL	250000	~200	~1000	~3000	~200		



Analytical Results

Sub-Matrix: WATER (Matrix: WATER)		Client sample ID		QC03_20160824	QC04_20160824					
	CI	Client sampling date / time		[24-Aug-2016]	[24-Aug-2016]					
Compound	CAS Number	LOR	Unit	EP1607778-006	EP1607778-007					
				Result	Result					
EA015: Total Dissolved Solids dried at 18	80 ± 5 °C									
Total Dissolved Solids @180°C		10	mg/L	<10	<10					
EK055G: Ammonia as N by Discrete Analyser										
Ammonia as N	7664-41-7	0.01	mg/L	<0.01	<0.01					
EK057G: Nitrite as N by Discrete Analyse	er									
Nitrite as N	14797-65-0	0.01	mg/L	<0.01	<0.01					
EK058G: Nitrate as N by Discrete Analys	ser									
Nitrate as N	14797-55-8	0.01	mg/L	<0.01	<0.01					
EK059G: Nitrite plus Nitrate as N (NOx) by Discrete Analyser										
Nitrite + Nitrate as N		0.01	mg/L	<0.01	<0.01					
EK061G: Total Kjeldahl Nitrogen By Disc	rete Analyser									
Total Kjeldahl Nitrogen as N		0.1	mg/L	<0.1	<0.1					
EK062G: Total Nitrogen as N (TKN + NOx	() by Discrete Ar	nalyser								
^ Total Nitrogen as N		0.1	mg/L	<0.1	<0.1					
EK067G: Total Phosphorus as P by Discr	rete Analyser									
Total Phosphorus as P		0.01	mg/L	<0.01	<0.01					
EP030: Biochemical Oxygen Demand (BC)									
Biochemical Oxygen Demand		2	mg/L	5	<2					
MW002: Heterotrophic Plate Count										
Heterotrophic Plate Count (22°C)		1	CFU/mL							
Heterotrophic Plate Count (36°C)		1	CFU/mL							
MW006: Faecal Coliforms & E.coli by MF										
Faecal Coliforms		1	CFU/100mL							
Escherichia coli		1	CFU/100mL							
MW007: Coliforms by MF										
Coliforms		1	CFU/100mL							



QUALITY CONTROL REPORT

Work Order	: EP1607778	Page	: 1 of 4
Client	: GHD PTY LTD	Laboratory	: Environmental Division Perth
Contact	: MR ANDREW NAGLE	Contact	: Lauren Biagioni
Address	999 HAY STREET	Address	: 10 Hod Way Malaga WA Australia 6090
Telephone	PERTH WA 6000 : +61 08 6222 8222	Telephone	: 08 9209 7655
Project	: 61/34772 Point Moore Groundwater Assessment	Date Samples Received	: 25-Aug-2016
Order number	:	Date Analysis Commenced	: 25-Aug-2016
C-O-C number	:	Issue Date	31-Aug-2016
Sampler	: STEVEN PETTS		HACEMRA NATA
Site	:		
Quote number	:		Accreditation No. 825
No. of samples received	: 7		Accredited for compliance with
No. of samples analysed	: 7		ISO/IEC 17025 - Testing

This report supersedes any previous report(s) with this reference. Results apply to the sample(s) as submitted. This document shall not be reproduced, except in full. This Quality Control Report contains the following information:

- Laboratory Duplicate (DUP) Report; Relative Percentage Difference (RPD) and Acceptance Limits
- Method Blank (MB) and Laboratory Control Spike (LCS) Report; Recovery and Acceptance Limits
- Matrix Spike (MS) Report; Recovery and Acceptance Limits

Signatories

This document has been electronically signed by the authorized signatories below. Electronic signing is carried out in compliance with procedures specified in 21 CFR Part 11.

Signatories	Position	Accreditation Category
Alini Goundar	Senior Analyst	Perth Microbiology, Malaga, WA
Jeremy Truong	Laboratory Manager	Perth Inorganics, Malaga, WA
Tyrone Cole	Inorganics Preparation Supervisor	Perth Inorganics, Malaga, WA



General Comments

The analytical procedures used by the Environmental Division have been developed from established internationally recognized procedures such as those published by the USEPA, APHA, AS and NEPM. In house developed procedures are employed in the absence of documented standards or by client request.

Where moisture determination has been performed, results are reported on a dry weight basis.

Where a reported less than (<) result is higher than the LOR, this may be due to primary sample extract/digestate dilution and/or insufficient sample for analysis. Where the LOR of a reported result differs from standard LOR, this may be due to high

Key: Anonymous = Refers to samples which are not specifically part of this work order but formed part of the QC process lot

- CAS Number = CAS registry number from database maintained by Chemical Abstracts Services. The Chemical Abstracts Service is a division of the American Chemical Society.
- LOR = Limit of reporting
- RPD = Relative Percentage Difference
- # = Indicates failed QC

Laboratory Duplicate (DUP) Report

The quality control term Laboratory Duplicate refers to a randomly selected intralaboratory split. Laboratory duplicates provide information regarding method precision and sample heterogeneity. The permitted ranges for the Relative Percent Deviation (RPD) of Laboratory Duplicates are specified in ALS Method QWI-EN/38 and are dependent on the magnitude of results in comparison to the level of reporting: Result < 10 times LOR: No Limit; Result between 10 and 20 times LOR: 0% - 50%; Result > 20 times LOR: 0% - 20%.

Sub-Matrix: WATER				Laboratory Duplicate (DUP) Report						
Laboratory sample ID	Client sample ID	Method: Compound CAS Number			Unit	Original Result	Duplicate Result	RPD (%)	Recovery Limits (%)	
EA015: Total Dissolved Solids dried at 180 ± 5 °C (QC Lot: 564719)										
EP1607760-009	Anonymous	EA015H: Total Dissolved Solids @180°C		10	mg/L	6310	6080	3.81	0% - 20%	
EP1607760-001	Anonymous	EA015H: Total Dissolved Solids @180°C		10	mg/L	6260	6170	1.45	0% - 20%	
EK055G: Ammonia as N by Discrete Analyser (QC Lot: 562898)										
EP1607778-001	MW1	EK055G: Ammonia as N	7664-41-7	0.01	mg/L	2.68	2.72	1.60	0% - 20%	
EP1607791-001	Anonymous	EK055G: Ammonia as N	7664-41-7	0.01	mg/L	0.04	0.09	73.8	No Limit	
EK057G: Nitrite as N by Discrete Analyser (QC Lot: 562895)										
EP1607778-001	MW1	EK057G: Nitrite as N	14797-65-0	0.01	mg/L	<0.01	<0.01	0.00	No Limit	
EK059G: Nitrite plus Nitrate as N (NOx) by Discrete Analyser (QC Lot: 562899)										
EP1607778-001	MW1	EK059G: Nitrite + Nitrate as N		0.01	mg/L	0.19	0.17	10.2	0% - 50%	
EP1607791-001	Anonymous	EK059G: Nitrite + Nitrate as N		0.01	mg/L	0.25	0.22	12.1	0% - 20%	
EK061G: Total Kjeldahl Nitrogen By Discrete Analyser (QC Lot: 566331)										
EP1607775-011	Anonymous	EK061G: Total Kjeldahl Nitrogen as N		0.1	mg/L	2.3	2.4	6.62	0% - 50%	
EP1607778-006	QC03_20160824	EK061G: Total Kjeldahl Nitrogen as N		0.1	mg/L	<0.1	0.1	0.00	No Limit	
EK067G: Total Phosphorus as P by Discrete Analyser (QC Lot: 566330)										
EP1607775-011	Anonymous	EK067G: Total Phosphorus as P		0.01	mg/L	0.22	0.21	0.00	0% - 50%	
EP1607778-006	QC03_20160824	EK067G: Total Phosphorus as P		0.01	mg/L	<0.01	<0.01	0.00	No Limit	
EP030: Biochemical Oxygen Demand (BOD) (QC Lot: 562689)										
EP1607769-001	Anonymous	EP030: Biochemical Oxygen Demand		2	mg/L	4	4	0.00	No Limit	


Method Blank (MB) and Laboratory Control Spike (LCS) Report

The quality control term Method / Laboratory Blank refers to an analyte free matrix to which all reagents are added in the same volumes or proportions as used in standard sample preparation. The purpose of this QC parameter is to monitor potential laboratory contamination. The quality control term Laboratory Control Spike (LCS) refers to a certified reference material, or a known interference free matrix spiked with target analytes. The purpose of this QC parameter is to monitor method precision and accuracy independent of sample matrix. Dynamic Recovery Limits are based on statistical evaluation of processed LCS.

Sub-Matrix: WATER		Method Blank (MB)	Laboratory Control Spike (LCS) Report						
				Report	Spike	Spike Recovery (%)	Recovery	Limits (%)	
Method: Compound	CAS Number	LOR	Unit	Result	Concentration	LCS	Low	High	
EA015: Total Dissolved Solids dried at 180 \pm 5 °C (QC	Lot: 564719)								
EA015H: Total Dissolved Solids @180°C		10	mg/L	<10	2000 mg/L	99.0	83	111	
				<10	293 mg/L	107	70	130	
EK055G: Ammonia as N by Discrete Analyser (QCLot:	562898)								
EK055G: Ammonia as N	7664-41-7	0.01	mg/L	<0.01	1 mg/L	109	87	115	
EK057G: Nitrite as N by Discrete Analyser (QCLot: 56	2895)								
EK057G: Nitrite as N	14797-65-0	0.01	mg/L	<0.01	0.5 mg/L	99.2	86	112	
EK059G: Nitrite plus Nitrate as N (NOx) by Discrete A	nalyser (QCLot: 56	2899)							
EK059G: Nitrite + Nitrate as N		0.01	mg/L	<0.01	0.5 mg/L	112	92	112	
EK061G: Total Kjeldahl Nitrogen By Discrete Analyser	(QCLot: 566331)								
EK061G: Total Kjeldahl Nitrogen as N		0.1	mg/L	<0.1	10 mg/L	84.8	82	110	
EK067G: Total Phosphorus as P by Discrete Analyser	EK067G: Total Phosphorus as P by Discrete Analyser (QCLot: 566330)								
EK067G: Total Phosphorus as P		0.01	mg/L	<0.01	4.42 mg/L	86.8	70	130	
EP030: Biochemical Oxygen Demand (BOD) (QCLot: 562689)									
EP030: Biochemical Oxygen Demand		2	mg/L	<2	198 mg/L	92.8	78	117	

Matrix Spike (MS) Report

The quality control term Matrix Spike (MS) refers to an intralaboratory split sample spiked with a representative set of target analytes. The purpose of this QC parameter is to monitor potential matrix effects on analyte recoveries. Static Recovery Limits as per laboratory Data Quality Objectives (DQOs). Ideal recovery ranges stated may be waived in the event of sample matrix interference.

Sub-Matrix: WATER			Matrix Spike (MS) Report						
				Spike	SpikeRecovery(%)	Recovery Li	nits (%)		
Laboratory sample ID	Client sample ID	Method: Compound	CAS Number	Concentration	MS	Low	High		
EK055G: Ammonia	as N by Discrete Analyser (QCLot: 562898)								
EP1607778-002	QC01_20160824	EK055G: Ammonia as N	7664-41-7	1 mg/L	96.8	70	130		
EK057G: Nitrite as	EK057G: Nitrite as N by Discrete Analyser (QCLot: 562895)								
EP1607778-002	QC01_20160824	EK057G: Nitrite as N	14797-65-0	0.5 mg/L	107	70	130		
EK059G: Nitrite plu	us Nitrate as N (NOx) by Discrete Analyser (QCLot: 562	899)							
EP1607778-002	QC01_20160824	EK059G: Nitrite + Nitrate as N		0.5 mg/L	111	70	130		
EK061G: Total Kjele	dahl Nitrogen By Discrete Analyser (QCLot: 566331)								
EP1607775-011	Anonymous	EK061G: Total Kjeldahl Nitrogen as N		5 mg/L	90.2	70	130		
EK067G: Total Pho	EK067G: Total Phosphorus as P by Discrete Analyser (QCLot: 566330)								
EP1607775-011	Anonymous	EK067G: Total Phosphorus as P		1 mg/L	97.6	70	130		

Page	: 4 of 4
Work Order	: EP1607778
Client	: GHD PTY LTD
Project	: 61/34772 Point Moore Groundwater Assessment





	QA/QC Compliance Asse	ssment to assist with	n Quality Review	
Work Order	: EP1607778	Page	: 1 of 6	
Client		Laboratory	: Environmental Division Perth	
Contact	: MR ANDREW NAGLE	Telephone	: 08 9209 7655	
Project	: 61/34772 Point Moore Groundwater Assessment	Date Samples Received	: 25-Aug-2016	
Site	:	Issue Date	: 31-Aug-2016	
Sampler	: STEVEN PETTS	No. of samples received	: 7	
Order number	:	No. of samples analysed	: 7	

This report is automatically generated by the ALS LIMS through interpretation of the ALS Quality Control Report and several Quality Assurance parameters measured by ALS. This automated reporting highlights any non-conformances, facilitates faster and more accurate data validation and is designed to assist internal expert and external Auditor review. Many components of this report contribute to the overall DQO assessment and reporting for guideline compliance.

Brief method summaries and references are also provided to assist in traceability.

Summary of Outliers

Outliers : Quality Control Samples

This report highlights outliers flagged in the Quality Control (QC) Report.

- <u>NO</u> Method Blank value outliers occur.
- <u>NO</u> Duplicate outliers occur.
- <u>NO</u> Laboratory Control outliers occur.
- <u>NO</u> Matrix Spike outliers occur.
- For all regular sample matrices, <u>NO</u> surrogate recovery outliers occur.

Outliers : Analysis Holding Time Compliance

• NO Analysis Holding Time Outliers exist.

Outliers : Frequency of Quality Control Samples

• <u>NO</u> Quality Control Sample Frequency Outliers exist.



Analysis Holding Time Compliance

If samples are identified below as having been analysed or extracted outside of recommended holding times, this should be taken into consideration when interpreting results.

This report summarizes extraction / preparation and analysis times and compares each with ALS recommended holding times (referencing USEPA SW 846, APHA, AS and NEPM) based on the sample container provided. Dates reported represent first date of extraction or analysis and preclude subsequent dilutions and reruns. A listing of breaches (if any) is provided herein.

Holding time for leachate methods (e.g. TCLP) vary according to the analytes reported. Assessment compares the leach date with the shortest analyte holding time for the equivalent soil method. These are: organics 14 days, mercury 28 days & other metals 180 days. A recorded breach does not guarantee a breach for all non-volatile parameters.

Holding times for <u>VOC in soils</u> vary according to analytes of interest. Vinyl Chloride and Styrene holding time is 7 days; others 14 days. A recorded breach does not guarantee a breach for all VOC analytes and should be verified in case the reported breach is a false positive or Vinyl Chloride and Styrene are not key analytes of interest/concern.

Matrix: WATER					Evaluation	n: × = Holding time	breach ; 🗸 = With	in holding time
Method	Sample Date	Extraction / Preparation			Analysis			
Container / Client Sample ID(s)			Date extracted	Due for extraction	Evaluation	Date analysed	Due for analysis	Evaluation
EA015: Total Dissolved Solids dried at 18	0 ± 5 °C							
Clear Plastic Bottle - Natural (EA015H)								
MW1,	QC01_20160824,	24-Aug-2016				29-Aug-2016	31-Aug-2016	 ✓
MW2,	MW3,							
SHP8,	QC03_20160824,							
QC04_20160824								
EK055G: Ammonia as N by Discrete Analy	yser							
Clear Plastic Bottle - Sulfuric Acid (EK055)	G)							
MW1,	QC01_20160824,	24-Aug-2016				25-Aug-2016	21-Sep-2016	 ✓
MW2,	MW3,							
SHP8,	QC03_20160824,							
QC04_20160824								
EK057G: Nitrite as N by Discrete Analyse	ər 💦							
Clear Plastic Bottle - Natural (EK057G)								
MW1,	QC01_20160824,	24-Aug-2016				25-Aug-2016	26-Aug-2016	✓
MW2,	MW3,							
SHP8,	QC03_20160824,							
QC04_20160824								
EK059G: Nitrite plus Nitrate as N (NOx) b	by Discrete Analyser							
Clear Plastic Bottle - Sulfuric Acid (EK059	G)							
MW1,	QC01_20160824,	24-Aug-2016				25-Aug-2016	21-Sep-2016	 ✓
MW2,	MW3,							
SHP8,	QC03_20160824,							
QC04_20160824								
EK061G: Total Kjeldahl Nitrogen By Discr	rete Analyser							
Clear Plastic Bottle - Sulfuric Acid (EK061)	G)							
MW1,	QC01_20160824,	24-Aug-2016	31-Aug-2016	21-Sep-2016	1	31-Aug-2016	21-Sep-2016	 ✓
MW2,	MW3,							
SHP8,	QC03_20160824,							
QC04 20160824								

Page	: 3 of 6
Work Order	: EP1607778
Client	: GHD PTY LTD
Project	: 61/34772 Point Moore Groundwater Assessment



Matrix: WATER					Evaluatior	n: × = Holding time	breach ; ✓ = With	in holding time	
Method	Sample Date	Extraction / Preparation			Analysis				
Container / Client Sample ID(s)			Date extracted	Due for extraction	Evaluation	Date analysed	Due for analysis	Evaluation	
EK067G: Total Phosphorus as P by Discrete	Analyser								
Clear Plastic Bottle - Sulfuric Acid (EK067G) MW1, MW2, SHP8, QC04_20160824	QC01_20160824, MW3, QC03_20160824,	24-Aug-2016	31-Aug-2016	21-Sep-2016	1	31-Aug-2016	21-Sep-2016	1	
EP030: Biochemical Oxygen Demand (BOD)									
Clear Plastic Bottle - Natural (EP030) MW1, MW2, SHP8, QC04_20160824	QC01_20160824, MW3, QC03_20160824,	24-Aug-2016				25-Aug-2016	26-Aug-2016	~	
MW002: Heterotrophic Plate Count									
Sterile Plastic Bottle - Sodium Thiosulfate (MV MW1, MW2, SHP8	QC01_20160824 , MW3,	24-Aug-2016				25-Aug-2016	25-Aug-2016	1	
MW006: Faecal Coliforms & E.coli by MF									
Sterile Plastic Bottle - Sodium Thiosulfate (MV MW1, MW2, SHP8	V006) QC01_20160824, MW3,	24-Aug-2016				25-Aug-2016	25-Aug-2016	1	
MW007: Coliforms by MF									
Sterile Plastic Bottle - Sodium Thiosulfate (MV MW1, MW2, SHP8	QC01_20160824, MW3,	24-Aug-2016				25-Aug-2016	25-Aug-2016	~	



Quality Control Parameter Frequency Compliance

The following report summarises the frequency of laboratory QC samples analysed within the analytical lot(s) in which the submitted sample(s) was(were) processed. Actual rate should be greater than or equal to the expected rate. A listing of breaches is provided in the Summary of Outliers.

Matrix: WATER				Evaluatio	n: × = Quality Co	ntrol frequency	not within specification ; \checkmark = Quality Control frequency within specification.
Quality Control Sample Type		Co	ount		Rate (%)		Quality Control Specification
Analytical Methods	Method	00	Reaular	Actual	Expected	Evaluation	
Laboratory Duplicates (DUP)							
Ammonia as N by Discrete analyser	EK055G	2	19	10.53	10.00	✓	NEPM 2013 B3 & ALS QC Standard
Biochemical Oxygen Demand (BOD)	EP030	1	9	11.11	10.00	✓	NEPM 2013 B3 & ALS QC Standard
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	2	20	10.00	10.00	✓	NEPM 2013 B3 & ALS QC Standard
Nitrite as N by Discrete Analyser	EK057G	1	7	14.29	10.00	✓	NEPM 2013 B3 & ALS QC Standard
Total Dissolved Solids (High Level)	EA015H	2	20	10.00	10.00	✓	NEPM 2013 B3 & ALS QC Standard
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	2	20	10.00	10.00	1	NEPM 2013 B3 & ALS QC Standard
Total Phosphorus as P By Discrete Analyser	EK067G	2	20	10.00	10.00	1	NEPM 2013 B3 & ALS QC Standard
Laboratory Control Samples (LCS)							
Ammonia as N by Discrete analyser	EK055G	1	19	5.26	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Biochemical Oxygen Demand (BOD)	EP030	1	9	11.11	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	1	20	5.00	5.00	~	NEPM 2013 B3 & ALS QC Standard
Nitrite as N by Discrete Analyser	EK057G	1	7	14.29	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Total Dissolved Solids (High Level)	EA015H	2	20	10.00	10.00	✓	NEPM 2013 B3 & ALS QC Standard
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	1	20	5.00	5.00	1	NEPM 2013 B3 & ALS QC Standard
Total Phosphorus as P By Discrete Analyser	EK067G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Method Blanks (MB)							
Ammonia as N by Discrete analyser	EK055G	1	19	5.26	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Biochemical Oxygen Demand (BOD)	EP030	1	9	11.11	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	1	20	5.00	5.00	~	NEPM 2013 B3 & ALS QC Standard
Nitrite as N by Discrete Analyser	EK057G	1	7	14.29	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Total Dissolved Solids (High Level)	EA015H	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	1	20	5.00	5.00	1	NEPM 2013 B3 & ALS QC Standard
Total Phosphorus as P By Discrete Analyser	EK067G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Matrix Spikes (MS)							
Ammonia as N by Discrete analyser	EK055G	1	19	5.26	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Nitrite as N by Discrete Analyser	EK057G	1	7	14.29	5.00	~	NEPM 2013 B3 & ALS QC Standard
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard
Total Phosphorus as P By Discrete Analyser	EK067G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard



Brief Method Summaries

The analytical procedures used by the Environmental Division have been developed from established internationally recognized procedures such as those published by the US EPA, APHA, AS and NEPM. In house developed procedures are employed in the absence of documented standards or by client request. The following report provides brief descriptions of the analytical procedures employed for results reported in the Certificate of Analysis. Sources from which ALS methods have been developed are provided within the Method Descriptions.

Analytical Methods	Method	Matrix	Method Descriptions
Total Dissolved Solids (High Level)	EA015H	WATER	In house: Referenced to APHA 2540C. A gravimetric procedure that determines the amount of `filterable` residue in an aqueous sample. A well-mixed sample is filtered through a glass fibre filter (1.2um). The filtrate is evaporated to dryness and dried to constant weight at 180+/-5C. This method is compliant with NEPM (2013) Schedule B(3)
Ammonia as N by Discrete analyser	EK055G	WATER	In house: Referenced to APHA 4500-NH3 G Ammonia is determined by direct colorimetry by Discrete Analyser. This method is compliant with NEPM (2013) Schedule B(3)
Nitrite as N by Discrete Analyser	EK057G	WATER	In house: Referenced to APHA 4500-NO2- B. Nitrite is determined by direct colourimetry by Discrete Analyser. This method is compliant with NEPM (2013) Schedule B(3)
Nitrate as N by Discrete Analyser	EK058G	WATER	In house: Referenced to APHA 4500-NO3- F. Nitrate is reduced to nitrite by way of a chemical reduction followed by quantification by Discrete Analyser. Nitrite is determined seperately by direct colourimetry and result for Nitrate calculated as the difference between the two results. This method is compliant with NEPM (2013) Schedule B(3)
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	WATER	In house: Referenced to APHA 4500-NO3- F. Combined oxidised Nitrogen (NO2+NO3) is determined by Chemical Reduction and direct colourimetry by Discrete Analyser. This method is compliant with NEPM (2013) Schedule B(3)
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	WATER	In house: Referenced to APHA 4500-Norg D (In house). An aliquot of sample is digested using a high temperature Kjeldahl digestion to convert nitrogenous compounds to ammonia. Ammonia is determined colorimetrically by discrete analyser. This method is compliant with NEPM (2013) Schedule B(3)
Total Nitrogen as N (TKN + Nox) By Discrete Analyser	EK062G	WATER	In house: Referenced to APHA 4500-Norg / 4500-NO3 This method is compliant with NEPM (2013) Schedule B(3)
Total Phosphorus as P By Discrete Analyser	EK067G	WATER	In house: Referenced to APHA 4500-P H, Jirka et al (1976), Zhang et al (2006). This procedure involves sulphuric acid digestion of a sample aliquot to break phosphorus down to orthophosphate. The orthophosphate reacts with ammonium molybdate and antimony potassium tartrate to form a complex which is then reduced and its concentration measured at 880nm using discrete analyser. This method is compliant with NEPM (2013) Schedule B(3)
Biochemical Oxygen Demand (BOD)	EP030	WATER	In house: Referenced to APHA 5210 B. The 5-Day BOD test provides an empirical measure of the oxygen consumption capacity of a given water. A portion of the sample is diluted into oxygenated, nutrient rich water, and a seed added to begin biological decay. The initial dissolved oxygen content is measured, then the bottle is sealed and incubated for five days. The remaining dissolved oxygen is measured, and from the difference, the demand for oxygen, by biological decay, is determined. This method is compliant with NEPM (2013) Schedule B(3)
Heterotrophic (Total) Plate Count @ 22C and 36C	MW002	WATER	In house: Referenced to AS4276.3.1- 2007
Thermotolerant Coliforms & E.coli by Membrane Filtration	MW006	WATER	In house: Referenced to AS 4276.7 2007
Coliforms by Membrane Filtration	MW007	WATER	In house: Referenced to AS 4276.5 - 2007
Preparation Methods	Method	Matrix	Method Descriptions

Page Work Order Client Project	 6 of 6 EP1607778 GHD PTY LTD 61/34772 Point Moore 	Groundwater Assessment			ALS
Preparation Methods		Method	Matrix	Method Descriptions	
TKN/TP Digestion		EK061/EK067	WATER	In house: Referenced to APHA 4500 Norg - D; APHA 4500 P - H. This method is compliant with NEPM (2013) Schedule B(3)	

Received by:			Sampled by: S. Petts		acoz PEII0063.	GHD Sample ID Laboratory Sam	Job Manager (Invoice) Andrew Nagle		Laboratory Quote No.	City of Greater Geraldton	Client	Point Moore Groundwater Assessmen	CHAIN OF CUSTORDY RECORD AND ANALYSIS REQUEST								
1 2 mm m m m m m m m m m m m m m m m m m	() come)				100	nple ID Date Time	Email Address (Results) andrew.nagle@ghd.com steven.petts@ghd.com	Standard	Turnaround Time	61/34772	Job No.	nt	D CHD SUGAS PROPULATION								
Date/Time: 25/8/16@ 1085	Date/Time: 24/08/2016	Date/Time: 24/08/2016		W BP H29	Sample Matrix S-Soil/ Type B-Bottle/J-Jar/V-Vid Preservative Unprese	/ SL-Sludge/ V al/Bag/G-Glas rved/ HCl/	/-Water/ A-Air s/P-Plastic		Laboratory Contact: Ca	Address: 28 Reid Road,	Laboratory:SGS	GHD House 239 Adelaide Terrace Perth W 6004									
						6	6	. 01	6 R				_		504 3 810	H2504/HN03/Other No Total Volume (mL)			ntainer	rrianne Graham	Perth Airport
Conter/ Hansport	Courier/ Transport	Relinnuished hv-	Relinguished by: S.P		X X	Total Nitrogen, TKN, N NH3, Total Phosphoru Biochemical Oxygen D TDS - Standard Level	lox, NO2, N s remand	03,	-				Telephone 08 6222 8222 8555								
Joinpairy, irea	AM WARNA		etts		X	Total (Heterotrophic P 22oC and 36oC (TPC) Total +Faecal Coliform	late Count YEA s & E. Coli (at MF)					Facsimile 08 6222 Pa								
										Analyses				ge <u>1</u> of <u>1</u>							
				PE11006:	SGS Perth Envi	Uald															
	Date/ Time.	Date/ Time:	Date/ Time: 24/08/2016	3 COC Aug – 2016	ronmental		(Pt 110062	Remarks				COC No.								

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			125mL 1 Plastic F Green
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			one °C



ANALYTICAL REPORT



CLIENT DETAILS		LABORATORY DETAIL	LS
Contact	Andrew Nagle	Manager	Ros Ma
Client	GHD Pty Ltd	Laboratory	SGS Perth Environmental
Address	(PO Box Y3106 PERTH WA 6832) 239 Adelaide Terrace EAST PERTH WA 6004	Address	28 Reid Rd Perth Airport WA 6105
Telephone	08 6222 8222	Telephone	(08) 9373 3500
Facsimile	08 6222 8555	Facsimile	(08) 9373 3556
Email	andrew.nagle@ghd.com	Email	au.environmental.perth@sgs.com
Project	61/34772_Point Moore Groundwater	SGS Reference	PE110063 R0
Order Number	61/34772	Date Received	24 Aug 2016
Samples	1	Date Reported	02 Sep 2016

COMMENTS .

Accredited for compliance with ISO/IEC 17025. NATA accredited laboratory 2562(898/20210).

SIGNATORIES _

Louisettope

Louise Hope Laboratory Technician

welgm

Michael McKay Inorganics and ARD Supervisor

Radel Horsen

Rachel Harrison Inorganics Team Leader

Environment, Health and Safety

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02-September-2016



ANALYTICAL REPORT

	Sar S S	nple Number ample Matrix Sample Date ample Name	PE110063.001 Water 24 Aug 2016 QC02			
Parameter	Units	LOR				
Total Dissolved Solids (TDS) in water Method: AN113 Teste	d: 1/9/2016					
Total Dissolved Solids Dried at 175-185°C	mg/L	10	7500			
CBOD5 Method: AN183 Tested: 26/8/2016						
Biochemical Oxygen Demand (CBOD5)	mg/L	5	<5			
Nitrate Nitrogen and Nitrite Nitrogen (NOx) by FIA Method: AN Nitrate/Nitrite Nitrogen, NOx as N	I258 Testee mg/L	0.05	0.17			
Nitrite, NO ₂ as NO ₂	mg/L	0.2	<0.2			
Nitrate, NO ₃ as NO ₃	mg/L	0.2	0.7			
Ammonia Nitrogen by FIA Method: AN261 Tested: 29/8/2016	5					
Ammonia, NH ₃	mg/L	0.05	2.9			
TKN Kjeldahl Digestion by Discrete Analyser Method: AN281 Tested: 31/8/2016						
Total Kjeldahl Nitrogen	mg/L	0.05	3.1			
Total Phosphorus by Kjeldahl Digestion DA in Water Method:	AN279/AN29	3(Sydney on	ly) Tested: 31/			
Total Phosphorus (Kjeldahl Digestion)	mg/L	0.02	0.19			



ANALYTICAL REPORT

		Sam Sa Sa Sa	ple Numbe mple Matri ample Dat ample Nam	er PE110063.001 x Water e 24 Aug 2016 e QC02
Parameter		Units	LOR	
Calculated Nitrogen Forms - TN, organic N, inorganic N	Metho	od: AN281/292	Teste	d: 2/9/2016
Total Nitrogen (calc)		mg/L	0.05	3.2



QC SUMMARY

PE110063 R0

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.

Ammonia Nitrogen by FIA Method: ME-(AU)-[ENV]AN261

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
Ammonia, NH ₃	LB121813	mg/L	0.05	<0.05	0%	NA

CBOD5 Method: ME-(AU)-[ENV]AN183

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
Biochemical Oxygen Demand (CBOD5)	LB121787	mg/L	5	<5	4%	88 - 90%

Nitrate Nitrogen and Nitrite Nitrogen (NOx) by FIA Method: ME-(AU)-[ENV]AN258

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
Nitrate/Nitrite Nitrogen, NOx as N	LB121813	mg/L	0.05	<0.05	0 - 5%	102 - 107%

TKN Kjeldahl Digestion by Discrete Analyser Method: ME-(AU)-[ENV]AN281

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
Total Kjeldahl Nitrogen	LB121936	mg/L	0.05	<0.05	0 - 6%	105%

Total Dissolved Solids (TDS) in water Method: ME-(AU)-[ENV]AN113

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS	MS	MSD %RPD
	Reference					%Recovery	%Recovery	
Total Dissolved Solids Dried at 175-185°C	LB121986	mg/L	10	<10	1%	104%	102%	2%

Total Phosphorus by Kjeldahl Digestion DA in Water Method: ME-(AU)-[ENV]AN279/AN293(Sydney only)

Parameter	QC	Units	LOR	MB	DUP %RPD	LCS
	Reference					%Recovery
Total Phosphorus (Kjeldahl Digestion)	LB121936	mg/L	0.02	<0.02	0 - 4%	103%



METHOD SUMMARY

METHOD	
- METHOD	METHODOLOGY SUMMARY
	Nitrate and Nitrite by FIA: In an acidic medium, nitrate is reduced quantitatively to nitrite by cadmium metal. This nitrite plus any original nitrite is determined as an intense red-pink azo dye at 540 nm following diazotisation with sulphanilamide and subsequent coupling with N-(1-naphthyl) ethylenediamine dihydrochloride. Without the cadmium reduction only the original nitrite is determined. Reference APHA 4500-NO3- F.
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.
AN113	Total Dissolved Solids: A well-mixed filtered sample of known volume is evaporated to dryness at 180°C and the residue weighed. Approximate methods for correlating chemical analysis with dissolved solids are available. Reference APHA 2540 C.
AN183	BOD: Serial dilutions of the sample are firstly combined with various reagents to aid bacterial growth and the sample is incubated for 5 days at 20°C. The difference between the initial and final oxygen contents of the sample is the amount of oxygen consumed by the bacteria. This is related to the organic loading of the sample therefore cBOD is the measure of the digestibility or bioavailability of organic matter in the sample. Reference APHA 5210 B. Internal Reference AN183
AN261	Ammonia by Continuous Flow Analyser: Ammonium in a basic medium forms ammonia gas, which is separated from the sample matrix by diffusion through a polypropylene membrane. The ammonia is reacted with phenol and hypochlorite to form indophenol blue at an intensity proportional to the ammonia concentration. The blue colour is intensified with sodium nitroprusside and the absorbance measured at 630 nm. The sensitivity of the automated method is 10-20 times that of the macro method. Reference APHA 4500-NH3 H.
AN279/AN293(Sydney)	The sample is digested with Sulphuric acid, K2SO4 and CuSO4. All forms of phosphorus are converted into orthophosphate. The digest is cooled and placed on the discrete analyser for colorimetric analysis.
AN281	An unfiltered water or soil sample is first digested in a block digestor with sulfuric acid, K2SO4 and CuSO4. The ammonia produced following digestion is then measured colourimetrically using the Aquakem 250 Discrete Analyser. A portion of the digested sample is buffered to an alkaline pH, and interfering cations are complexed. The ammonia then reacts with salicylate and hypochlorite to give a blue colour whose absorbance is measured at 660nm and compared with calibration standards. This is proportional to the concentration of Total Kjeldahl Nitrogen in the original sample.
AN281/292	Calculation of total nitrogen and organic nitrogen.



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
- 37 MBq is equivalent to 1 mCi b.

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

This document is issued, on the Client's behalf, by the Company under its General Conditions of Service available on request and accessible at http://www.sgs.com/en/terms-and-conditions. The Client's attention is drawn to the limitation of liability, indemnification and jurisdiction issues defined therein.

Any other holder of this document is advised that information contained hereon reflects the Company's findings at the time of its intervention only and within the limits of Client's instructions, if any. The Company's sole responsibility is to its Client and this document does not exonerate parties to a transaction from exercising all their rights and obligations under the transaction documents.

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STATEMENT OF QA/QC PERFORMANCE

CLIENT DETAILS		LABORATORY DETAIL	.\$
Contact	Andrew Nagle	Manager	Ros Ma
Client	GHD Pty Ltd	Laboratory	SGS Perth Environmental
Address	(PO Box Y3106 PERTH WA 6832) 239 Adelaide Terrace EAST PERTH WA 6004	Address	28 Reid Rd Perth Airport WA 6105
Telephone	08 6222 8222	Telephone	(08) 9373 3500
Facsimile	08 6222 8555	Facsimile	(08) 9373 3556
Email	andrew.nagle@ghd.com	Email	au.environmental.perth@sgs.com
Project	61/34772_Point Moore Groundwater	SGS Reference	PE110063 R0
Order Number	61/34772	Date Received	24 Aug 2016
Samples	1	Date Reported	02 Sep 2016

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 with the exception of the following:

Extraction Date	Nitrate Nitrogen and Nitrite Nitrogen (NOx) by FIA	1 item
	Total Dissolved Solids (TDS) in water	1 item
Analysis Date	Nitrate Nitrogen and Nitrite Nitrogen (NOx) by FIA	1 item
	Total Dissolved Solids (TDS) in water	1 item

- SAMPLE SUMMARY				
Sample counts by matrix	1 water	Type of documentation received	COC	
Date documentation received	26/8/2016	Samples received in good order	Yes	
Samples received without headspace	Yes	Sample temperature upon receipt	12°C	
Sample container provider	ALS	Turnaround time requested	Standard	
Samples received in correct containers	Yes	Sufficient sample for analysis	Yes	
Sample cooling method	Ice Bricks	Samples clearly labelled	Yes	
Complete documentation received	Yes	Number of eskies/boxes received	1	

SGS Australia Pty Ltd ABN 44 000 964 278 Environment, Health and Safety 28 Reid Rd Perth Airport WA 6105 Australia PO Box 32 Welshpool WA 6983 Australia t +61 8 9373 3500 www.sgs.com.au f +61 8 9373 3556

Member of the SGS Group



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.

Ammonia Nitrogen by FIA							Method:	ME-(AU)-[ENV]AN261
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
QC02	PE110063.001	LB121813	24 Aug 2016	24 Aug 2016	21 Sep 2016	29 Aug 2016	21 Sep 2016	29 Aug 2016
CBOD5							Method:	ME-(AU)-[ENV]AN183
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
QC02	PE110063.001	LB121787	24 Aug 2016	24 Aug 2016	26 Aug 2016	26 Aug 2016	02 Sep 2016	31 Aug 2016
Nitrate Nitrogen and Nitrite Nitro	gen (NOx) by FIA						Method:	ME-(AU)-[ENV]AN258
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
QC02	PE110063.001	LB121813	24 Aug 2016	24 Aug 2016	28 Aug 2016	29 Aug 2016†	28 Aug 2016	29 Aug 2016†
TKN Kjeldahl Digestion by Discr	ete Analyser						Method:	ME-(AU)-[ENV]AN281
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
QC02	PE110063.001	LB121936	24 Aug 2016	24 Aug 2016	21 Sep 2016	31 Aug 2016	21 Sep 2016	02 Sep 2016
Total Dissolved Solids (TDS) in v	water						Method:	ME-(AU)-[ENV]AN113
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Due	Analysed
QC02	PE110063.001	LB121986	24 Aug 2016	24 Aug 2016	31 Aug 2016	01 Sep 2016†	31 Aug 2016	01 Sep 2016†
Total Phosphorus by Kjeldahl Di	gestion DA in Water					Metho	d: ME-(AU)-[ENV]AN27	9/AN293(Sydney only)
Sample Name	Sample No.	QC Ref	Sampled	Received	Extraction Due	Extracted	Analysis Du <u>e</u>	Analysed
QC02	PE110063.001	LB121936	24 Aug 2016	24 Aug 2016	21 Sep 2016	31 Aug 2016	21 Sep 2016	02 Sep 2016



SURROGATES

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.



METHOD BLANKS

PE110063 R0

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.

Ammonia Nitrogen by FIA

Ammonia Nitrogen by FIA			Metho	od: ME-(AU)-[ENV]AN261
Sample Number	Parameter	Units	LOR	Result
LB121813.001	Ammonia, NH ₃	mg/L	0.05	<0.05
LB121813.025	Ammonia, NH ₃	mg/L	0.05	<0.05
LB121813.049	Ammonia, NH ₃	mg/L	0.05	<0.05
LB121813.073	Ammonia, NH ₃	mg/L	0.05	<0.05
LB121813.097	Ammonia, NH ₃	mg/L	0.05	<0.05
CBOD5			Metho	od: ME-(AU)-[ENV]AN183
Sample Number	Parameter	Units	LOR	Result
LB121787.001	Biochemical Oxygen Demand (CBOD5)	mg/L	5	<5
LB121787.003	Biochemical Oxygen Demand (CBOD5)	mg/L	5	<5

TKN Kjeldahl Digestion by Discrete Analyser

TKN Kjeldahl Digestion by Discrete Analyser		Me	ethod: ME-(AU)-[ENV]AN281	
Sample Number	Parameter	Units	LOR	Result
LB121936.001	Total Kjeldahl Nitrogen	mg/L	0.05	<0.05

Total Dissolved Solids (TDS) in water

Total Dissolved Solids (TDS) in water				od: ME-(AU)-[ENV]AN113
Sample Number	Parameter	Units	LOR	Result
LB121986.001	Total Dissolved Solids Dried at 175-185°C	mg/L	10	<10

Total Phosphorus by Kjeldahl Digestion	Method:	ME-(AU)-[ENV]A	N279/AN293(Sydney only)	
Sample Number	Parameter	Units	LOR	Result
LB121936.001	Total Phosphorus (Kjeldahl Digestion)	mg/L	0.02	<0.02



Method: ME-(AU)-IENVIAN183

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.

Ammonia Nitrogen by FIA

Ammonia Nitrogen by	/ FIA					Metho	od: ME-(AU)-[ENVJAN261
Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
PE109968.007	LB121813.013	Ammonia, NH ₃	mg/L	0.05	-0.0133571428	30.0218571428	200	0
PE110086.002	LB121813.102	Ammonia, NH ₃	mg/L	0.05	0.0497857142	0.0497857142	115	0

CBOD5

									Î
Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %	
PE110067.001	LB121787.010	Biochemical Oxygen Demand (CBOD5)	mg/L	5	324	337	17	4	1

Nitrate Nitrogen and Nitrite Nitrogen (NOx) by FIA

Nitrate Nitrogen and Nitrite Nitrogen (NOx) by FIA Method: ME-(AU)-[ENV]AN2						ENVJAN258		
Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
PE109968.007	LB121813.013	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	0.219	0.21	38	4
PE109995.001	LB121813.028	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	34.44	33.99	15	1
PE109995.011	LB121813.039	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	57.106	58.26	15	2
PE110020.001	LB121813.054	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	3.24	3.575	16	0
PE110043.001	LB121813.065	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	4.27	4.29	16	0
PE110079.005	LB121813.080	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	14.37	15.169	15	5
PE110079.015	LB121813.091	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	2.902	2.901	17	0
PE110086.002	LB121813.106	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	7.633	7.556	16	1
PE110090.002	LB121813.111	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	1.775	1.775	18	0
TKN Kjeldahl Digesti	on by Discrete Analyser					Meth	od: ME-(AU)-[ENVJAN281
Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
PE109742A.001	LB121936.027	Total Kjeldahl Nitrogen	mg/L	0.05	0.016	0.083	116	50
PE109987.001	LB121936.006	Total Kjeldahl Nitrogen	mg/L	0.05	55.313	58.528	15	6
PE110035.001	LB121936.015	Total Kjeldahl Nitrogen	mg/L	0.05	82.949	82.868	15	0
PE110082.006	LB121936.035	Total Kjeldahl Nitrogen	mg/L	0.05	12	13	15	6
Total Dissolved Solid	s (TDS) in water					Meth	od: ME-(AU)-[ENVJAN113
Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
PE110087.004	LB121986.015	Total Dissolved Solids Dried at 175-185°C	mg/L	10)49.34281404	864.19238780	16	1
PE110088.002	LB121986.023	Total Dissolved Solids Dried at 175-185°C	mg/L	10	00.118098612	208.256652573	16	1

Total Phosphorus	by Kjeldahl Digestion DA in	Water			Method: ME-(AU)-[ENV]AN279/AN293(Sydney only)			
Original	Duplicate	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %
PE109742A.001	LB121936.027	Total Phosphorus (Kjeldahl Digestion)	mg/L	0.02	0.013	0.012	175	0
PE109987.001	LB121936.006	Total Phosphorus (Kjeldahl Digestion)	mg/L	0.02	3.4	3.5	16	4
PE110035.001	LB121936.015	Total Phosphorus (Kjeldahl Digestion)	mg/L	0.02	8.053	7.817	15	3
PE110082.006	LB121936.035	Total Phosphorus (Kieldahl Digestion)	mg/L	0.02	9.3	9.2	15	0



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.

Method: ME-(AU)-[ENV]AN183

CBOD5	M						U)-[ENV]AN18
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB121787.005	Biochemical Oxygen Demand (CBOD5)	mg/L	5	180	198	80 - 120	90
LB121787.007	Biochemical Oxygen Demand (CBOD5)	mg/L	5	180	198	80 - 120	88

Nitrate Nitrogen and Nitrite Nitrogen (NOx) by FIA

Nitrate Nitrogen and Nitrite N	litrogen (NOx) by FIA				l. I	Nethod: ME-(A	U)-[ENV]AN258
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB121813.002	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	4.1	4	85 - 115	102
LB121813.027	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	4.2	4	85 - 115	106
LB121813.052	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	4.2	4	85 - 115	106
LB121813.077	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	4.2	4	85 - 115	106
LB121813.102	Nitrate/Nitrite Nitrogen, NOx as N	mg/L	0.05	4.3	4	85 - 115	107
TKN Kjeldahl Digestion by D	iscrete Analyser				N	Nethod: ME-(A	U)-[ENV]AN281
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB121936.002	Total Kieldahl Nitrogen	ma/L	0.05	1.0	1	80 - 120	105

Total Dissolved Solids (TDS) in water

Total Dissolved Solids (TDS) in water Method: ME-(A								
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %	
LB121986.002	Total Dissolved Solids Dried at 175-185°C	mg/L	10	310	300	80 - 120	104	

Total Phosphorus by Kjeldahl Digestion	IDA In Water			Method	I: ME-(AU)-[EN	V]AN279/AN2	93(Sydney only
Sample Number	Parameter	Units	LOR	Result	Expected	Criteria %	Recovery %
LB121936.002	Total Phosphorus (Kjeldahl Digestion)	mg/L	0.02	0.52	0.5	80 - 120	103



MATRIX SPIKES

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.

Total Dissolved So	olids (TDS) in water					Met	nod: ME-(AU	J)-[ENV]AN113
QC Sample	Sample Number	Parameter	Units	LOR	Result	Original	Spike	Recovery%
PE109998.001	LB121986.004	Total Dissolved Solids Dried at 175-185°C	mg/L	10	1300	270	1000	102



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 identifier when outside suggested criteria. Refer to the footnotes section at the end of this report for failure reasons.

Total Dissolved Solids (TDS) in water Method: ME-(AU)-[EN												
QC Sample	Sample Number	Parameter	Units	LOR	Original	Duplicate	Criteria %	RPD %				
PE109998.001	LB121986.005	Total Dissolved Solids Dried at 175-185°C	mg/L	10	1300	1300	16	2				



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.
- ④ 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).
- 6 LOR was raised due to sample matrix interference.
- ⁽⁷⁾ 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.
- I Low surrogate recovery due to the sample emulsifying during extraction.
- Refer to Analytical Report comments for further information.

This document is issued, on the Client's behalf, by the Company under its General Conditions of Service, available on request and accessible at http://www.sgs.com/en/terms-and-conditions. The Client's attention is drawn to the limitation of liability, indemnification and jurisdiction issues defined therein.

Any other holder of this document is advised that information contained herein reflects the Company's findings at the time of its intervention only and within the limits of Client's instructions, if any. The Company's sole responsibility is to its Client and this document does not exonerate parties to a transaction from exercising all their rights and obligations under the transaction documents.

This test report shall not be reproduced, except in full.

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Point Moore Ground	dwater Assessment			Laborat	ory:ALS												_					1
Client		Job No.		Address	s: 10 Hod \	Nay, Mala	ga 6090															1
City of Greater Gera	ldton	61/34772		Laborat	ory Conta	ct: Lauren	Biagion		SACTOM: NO.	15- 10 ⁻ 10- 10- 10-	1. N. L. 1993 L. 1. AV.		I NA II MATAINA A A A			- Tradition						
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Andrew Nagle		andrew.nagle@ghd.	com	Jac	die/). Tic	tive 3/oth		Ŭ														1
and the second	weight of the state of the stat	steven. petts@ghd.	com	A-Air	B-Bol	HNO		S			Ξ	2						·				1
GHD Sample ID	Laboratory Sample ID	Date	Time	Samp Water/	Type Glass/P	Prese	۶ ۷	Total	NT-8	EP030	EA015	MWD0	MS							Hold		
MW1	1			w	BP	H2SO4	3	810	\square	X	\mathbf{X}	X	$\mathbf{\Sigma}$							E	nvironmental Divis	sior
QC01_20160922	2			w	BP	H2SO4	3	810	$\mathbf{\Sigma}$	X	\mathbf{X}	X	\mathbf{X}		_						Work Order Reference	ж ЛЛ
MW2	3			w	BP	H2SO4	3	810	$\mathbf{\Sigma}$	X	\mathbf{X}	X	\mathbf{X}								LF 10009	+4
MW3	4			w	BP	H2SO4	3	810	\square	X	\square	X	\square						\downarrow			
SHP8	2			w	BP	H2SO4	3	810	\square	X	$\mathbf{\Sigma}$	X	\boxtimes									
QC03_20160922	6			w	BP	H2SO4	2	560	\mathbf{X}	X	$\mathbf{\Sigma}$	X			_							
QC04_20160922	7			w	BP	H2SO4	2	560	\mathbf{X}	X	$\mathbf{\Sigma}$	X								fe	elephone + 61-8-9209 765 *	
																				1	L	
Sampled by: S. Petts	<u> </u>			Date/Ti	me: 22/09	/2016			Relinqu	ished b	y: S.Pe	ts									Date/ Time: 22/09/2016	
Received by:				. n . /==		<u></u>			Relinqu	ished b	iγ:						-				Date/ Time:	1
Received by Lab: V	v Urr"			Date/Ti	me: 231	<u>9116 1</u>	015		Courier	/ Trans	port Co	mpany: N	Α									4
sample conditions:				Remark	5:																	



SAMPLE RECEIPT NOTIFICATION (SRN)

Work Order	: EP1608944						
Client Contact Address	Contact : GHD PTY LTD Contact : MR ANDREW NAGLE E 999 HAY STREET PERTH WA 6000		: Environmental Division Perth : Lauren Biagioni : 10 Hod Way Malaga WA Australia 609				
E-mail Telephone Facsimile	: andrew.nagle@ghd.com : +61 08 6222 8222 : +61 08 9429 6555	E-mail Telephone Facsimile	: Lauren.bia : 08 9209 76 : +61-8-9209	gioni@alsglobal.com 555 9 7600			
Project	: 61/34772 Point Moore Groundwater Assessment	Page	: 1 of 2				
Order number C-O-C number Site Sampler	: : : : STEVEN PETTS	Quote number QC Level	: EP2016GF : NEPM 201	IDSER0029 (EP/919/16 V2) 3 B3 & ALS QC Standard			
Dates Date Samples Receive Client Requested Due Date	d : 23-Sep-2016 10:15 AM : 03-Oct-2016	Issue Date Scheduled Reporting	Date	: 23-Sep-2016 : 03-Oct-2016			
Delivery Details Mode of Delivery No. of coolers/boxes Receipt Detail	Carrier : 7 :	Security Seal Temperature No. of samples receiv	ved / analysed	: Not intact. : 18.9 - Ice Bricks present : 7 / 7			

General Comments

- This report contains the following information:
 - Sample Container(s)/Preservation Non-Compliances
 - Summary of Sample(s) and Requested Analysis
 - Proactive Holding Time Report
 - Requested Deliverables
- Please see scanned COC for sample discrepencies: extra samples , samples not received etc.
- Please direct any queries related to sample condition / numbering / breakages to Sample Receipt (SamplesPerth@alsenviro.com)
- Analytical work for this work order will be conducted at ALS Environmental Perth.
- Please direct any turnaround / technical queries to the laboratory contact designated above.
- Sample Disposal Aqueous (14 days), Solid (60 days) from date of completion of Work Order.
- pH analysis should be conducted within 6 hours of sampling.



en + NO2 + NO3 + NH3 + Total P

1002 (22C and 36C) nic Colony Count (22C & 36C)

Level

- High

Sample Container(s)/Preservation Non-Compliances

All comparisons are made against pretreatment/preservation AS, APHA, USEPA standards.

• No sample container / preservation non-compliance exists.

Summary of Sample(s) and Requested Analysis

Some items described below may be part of a laboratory process necessary for the execution of client requested tasks. Packages may contain additional analyses, such as the determination of moisture content and preparation tasks, that are included in the package.

If no sampling time is provided, the sampling time will default to 15:00 on the date of sampling. If no sampling date is provided, the sampling date will be assumed by the laboratory for processing purposes and will be shown bracketed without a time component.

Matrix: WATER

the sampling dat processing purpos a time component.	client sample ID	ER - EA015H Dissolved Solids - H	ER - EP030	ER - M5 - ES/EB/FM E.coli & TC by MF	ER - MW002 (22C and rotrophic Colony Coun	ER - NT-08 Nitrogen + NO2 + NO
date / time		WAT Total	WAT BOD	WAT FC, F	WAT Hete	WAT
[23-Sep-2016]	MW1	✓	1	✓	✓	1
[23-Sep-2016]	QC01_20160922	1	1	1	✓	1
[23-Sep-2016]	MW2	1	1	1	✓	1
[23-Sep-2016]	MW3	1	1	1	✓	1
[23-Sep-2016]	SHP8	1	1	1	✓	1
[23-Sep-2016]	QC03_20160922	1	1			1
[23-Sep-2016]	QC04_20160922	1	1			1
	the sampling dat processing purpos time component. <i>Client sampling date / time</i> [23-Sep-2016] [23-Sep-2016] [23-Sep-2016] [23-Sep-2016] [23-Sep-2016] [23-Sep-2016] [23-Sep-2016]	the sampling date will be assumed by processing purposes and will be shown time component. Client sampling date / time [23-Sep-2016] MW1 [23-Sep-2016] QC01_20160922 [23-Sep-2016] MW2 [23-Sep-2016] MW3 [23-Sep-2016] SHP8 [23-Sep-2016] SHP8 [23-Sep-2016] QC03_20160922 [23-Sep-2016] QC04_20160922	the sampling date will be assumed by processing purposes and will be shown time component. <i>Client sampling Client sample ID</i> <i>date / time</i> [23-Sep-2016] MW1 \checkmark [23-Sep-2016] QC01_20160922 \checkmark [23-Sep-2016] MW2 \checkmark [23-Sep-2016] MW3 \checkmark [23-Sep-2016] SHP8 \checkmark [23-Sep-2016] SHP8 \checkmark [23-Sep-2016] QC03_20160922 \checkmark	the sampling date will be assumed by processing purposes and will be shown time component.T S S S S Client sample IDS 	the sampling date will be assumed by processing purposes and will be shown time component.T spice poor time component.Client sampling date / timeClient sample ID will be shown verticeV vertice vertice[23-Sep-2016]MW1✓✓[23-Sep-2016]QC01_20160922✓✓[23-Sep-2016]MW2✓✓[23-Sep-2016]MW3✓✓[23-Sep-2016]MW3✓✓[23-Sep-2016]MW3✓✓[23-Sep-2016]MW3✓✓[23-Sep-2016]SHP8✓✓[23-Sep-2016]QC03_20160922✓✓[23-Sep-2016]QC04_20160922✓✓	the sampling date will be assumed by processing purposes and will be shown time component.T spice poor processing purposes and will be shown of the component.T spice poor poor processing purposes and will be shown of the component.T spice poor poor poor processing purposes and will be shown of the component.T spice poor poor poor processing purposes and will be shown processing purposes and will be shown purpose and pu

Proactive Holding Time Report

Sample(s) have been received within the recommended holding times for the requested analysis.

Requested Deliverables

ACCOUNTS PAYABLE (Perth)		
- A4 - AU Tax Invoice (INV)	Email	ap-fss@ghd.com
ANDREW NAGLE		
 *AU Certificate of Analysis - NATA (COA) 	Email	andrew.nagle@ghd.com
 *AU Interpretive QC Report - DEFAULT (Anon QCI Rep) (QCI) 	Email	andrew.nagle@ghd.com
- *AU QC Report - DEFAULT (Anon QC Rep) - NATA (QC)	Email	andrew.nagle@ghd.com
- A4 - AU Sample Receipt Notification - Environmental HT (SRN)	Email	andrew.nagle@ghd.com
- Chain of Custody (CoC) (COC)	Email	andrew.nagle@ghd.com
- EDI Format - ENMRG (ENMRG)	Email	andrew.nagle@ghd.com
- EDI Format - ESDAT (ESDAT)	Email	andrew.nagle@ghd.com
- EDI Format - XTab (XTAB)	Email	andrew.nagle@ghd.com
STEVEN PETTS		
 *AU Certificate of Analysis - NATA (COA) 	Email	steven.petts@ghd.com
- *AU Interpretive QC Report - DEFAULT (Anon QCI Rep) (QCI)	Email	steven.petts@ghd.com
- *AU QC Report - DEFAULT (Anon QC Rep) - NATA (QC)	Email	steven.petts@ghd.com
- A4 - AU Sample Receipt Notification - Environmental HT (SRN)	Email	steven.petts@ghd.com
- Chain of Custody (CoC) (COC)	Email	steven.petts@ghd.com
- EDI Format - ENMRG (ENMRG)	Email	steven.petts@ghd.com
- EDI Format - ESDAT (ESDAT)	Email	steven.petts@ghd.com
- EDI Format - XTab (XTAB)	Email	steven.petts@ahd.com



CERTIFICATE OF ANALYSIS

Work Order	EP1608944	Page	: 1 of 4
Client	: GHD PTY LTD	Laboratory	Environmental Division Perth
Contact	: MR ANDREW NAGLE	Contact	: Lauren Biagioni
Address	: 999 HAY STREET	Address	: 10 Hod Way Malaga WA Australia 6090
	PERTH WA 6000		
Telephone	: +61 08 6222 8222	Telephone	: 08 9209 7655
Project	: 61/34772 Point Moore Groundwater Assessment	Date Samples Received	: 23-Sep-2016 10:15
Order number	:	Date Analysis Commenced	: 23-Sep-2016
C-O-C number	:	Issue Date	: 03-Oct-2016 20:21
Sampler	: STEVEN PETTS		HACEMRA NATA
Site	:		
Quote number	:		Accreditation No. 825
No. of samples received	: 7		Accredited for compliance with
No. of samples analysed	: 7		ISO/IEC 17025 - Testing

This report supersedes any previous report(s) with this reference. Results apply to the sample(s) as submitted. This document shall not be reproduced, except in full.

This Certificate of Analysis contains the following information:

- General Comments
- Analytical Results

Additional information pertinent to this report will be found in the following separate attachments: Quality Control Report, QA/QC Compliance Assessment to assist with Quality Review and Sample Receipt Notification.

Signatories

This document has been electronically signed by the authorized signatories below. Electronic signing is carried out in compliance with procedures specified in 21 CFR Part 11.

Signatories	Position	Accreditation Category
Alini Goundar	Senior Analyst	Perth Microbiology, Malaga, WA
Jeremy Truong	Laboratory Manager	Perth Inorganics, Malaga, WA
Tyrone Cole	Inorganics Preparation Supervisor	Perth Inorganics, Malaga, WA
Vinitha Kesavan	Analyst	Perth Microbiology, Malaga, WA



General Comments

The analytical procedures used by the Environmental Division have been developed from established internationally recognized procedures such as those published by the USEPA, APHA, AS and NEPM. In house developed procedures are employed in the absence of documented standards or by client request.

Where moisture determination has been performed, results are reported on a dry weight basis.

Where a reported less than (<) result is higher than the LOR, this may be due to primary sample extract/digestate dilution and/or insufficient sample for analysis.

Where the LOR of a reported result differs from standard LOR, this may be due to high moisture content, insufficient sample (reduced weight employed) or matrix interference.

When sampling time information is not provided by the client, sampling dates are shown without a time component. In these instances, the time component has been assumed by the laboratory for processing purposes.

Where a result is required to meet compliance limits the associated uncertainty must be considered. Refer to the ALS Contact for details.

Key : CAS Number = CAS registry number from database maintained by Chemical Abstracts Services. The Chemical Abstracts Service is a division of the American Chemical Society.

- LOR = Limit of reporting
- ^ = This result is computed from individual analyte detections at or above the level of reporting
- ø = ALS is not NATA accredited for these tests.
- ~ = Indicates an estimated value.
- CFU = colony forming unit
- MF = membrane filtration
- MW006 and MW007: estimate (~) is reported where the growth of presumptive bacteria on the filtered membrane is counted <10 cfu and/or >100 cfu and where there are many non-target colonies; the typical colonies may be masked by overgrowth of non-target organisms. It may be informative to record this fact.
- MW002 is ALS's internal code and is equivalent to AS4276.3.1.
- MW006 is ALS's internal code and is equivalent to AS4276.7.
- MW007 is ALS's internal code and is equivalent to AS4276.5.



Analytical Results

Sub-Matrix: WATER (Matrix: WATER)		Cli	ent sample ID	MW1	QC01_20160922	MW2	MW3	SHP8
	Cl	ient sampli	ing date / time	[23-Sep-2016]	[23-Sep-2016]	[23-Sep-2016]	[23-Sep-2016]	[23-Sep-2016]
Compound	CAS Number	LOR	Unit	EP1608944-001	EP1608944-002	EP1608944-003	EP1608944-004	EP1608944-005
				Result	Result	Result	Result	Result
EA015: Total Dissolved Solids dried at 1	80 ± 5 °C							
Total Dissolved Solids @180°C		10	mg/L	22300	22400	2120	1320	10300
EK055G: Ammonia as N by Discrete Ana	lyser							
Ammonia as N	7664-41-7	0.01	mg/L	6.28	6.42	0.13	0.78	2.54
EK057G: Nitrite as N by Discrete Analys	er							
Nitrite as N	14797-65-0	0.01	mg/L	<0.01	<0.01	1.85	<0.01	<0.01
EK058G: Nitrate as N by Discrete Analys	ser							
Nitrate as N	14797-55-8	0.01	mg/L	0.05	<0.01	9.95	0.01	0.03
EK059G: Nitrite plus Nitrate as N (NOx)	by Discrete Ana	lyser						
Nitrite + Nitrate as N		0.01	mg/L	0.05	<0.01	11.8	0.01	0.03
EK061G: Total Kjeldahl Nitrogen By Disc	rete Analyser							
Total Kjeldahl Nitrogen as N		0.1	mg/L	6.7	6.7	2.0	0.9	3.3
EK062G: Total Nitrogen as N (TKN + NO)	<) by Discrete Ar	nalyser						
^ Total Nitrogen as N		0.1	mg/L	6.8	6.7	13.8	0.9	3.3
EK067G: Total Phosphorus as P by Disc	rete Analyser							
Total Phosphorus as P		0.01	mg/L	0.81	0.90	0.14	0.08	0.38
EP030: Biochemical Oxygen Demand (B	OD)							
Biochemical Oxygen Demand		2	mg/L	31	22	4	5	22
MW002: Heterotrophic Plate Count								
Heterotrophic Plate Count (22°C)		1	CFU/mL	72	73	1700	2400	12
Heterotrophic Plate Count (36°C)		1	CFU/mL	170	140	12000	2100	17
MW006: Faecal Coliforms & E.coli by MF								
Faecal Coliforms		1	CFU/100mL	<1	<1	~<1	~1	<1
Escherichia coli		1	CFU/100mL	<1	<1	~<1	~1	<1
MW007: Coliforms by MF								
Coliforms		1	CFU/100mL	<1	<1	~10	~10	<1



Analytical Results

Sub-Matrix: WATER (Matrix: WATER)	Client sample ID		QC03_20160922	QC04_20160922				
	Client sampling date / time		[23-Sep-2016]	[23-Sep-2016]				
Compound	CAS Number	LOR	Unit	EP1608944-006	EP1608944-007			
				Result	Result			
EA015: Total Dissolved Solids dried at 1	80 ± 5 °C							
Total Dissolved Solids @180°C		10	mg/L	20	<10			
EK055G: Ammonia as N by Discrete Analyser								
Ammonia as N	7664-41-7	0.01	mg/L	<0.01	<0.01			
EK057G: Nitrite as N by Discrete Analys	ser							
Nitrite as N	14797-65-0	0.01	mg/L	<0.01	<0.01			
EK058G: Nitrate as N by Discrete Analy	ser							
Nitrate as N	14797-55-8	0.01	mg/L	<0.01	<0.01			
EK059G: Nitrite plus Nitrate as N (NOx) by Discrete Analyser								
Nitrite + Nitrate as N		0.01	mg/L	<0.01	<0.01			
EK061G: Total Kjeldahl Nitrogen By Discrete Analyser								
Total Kjeldahl Nitrogen as N		0.1	mg/L	<0.1	<0.1			
EK062G: Total Nitrogen as N (TKN + NO	x) by Discrete Ar	nalyser						
^ Total Nitrogen as N		0.1	mg/L	<0.1	<0.1			
EK067G: Total Phosphorus as P by Disc	rete Analyser							
Total Phosphorus as P		0.01	mg/L	<0.01	<0.01			
EP030: Biochemical Oxygen Demand (B	OD)							
Biochemical Oxygen Demand		2	mg/L	<2	<2			
MW002: Heterotrophic Plate Count								
Heterotrophic Plate Count (22°C)		1	CFU/mL					
Heterotrophic Plate Count (36°C)		1	CFU/mL					
MW006: Faecal Coliforms & E.coli by MF								
Faecal Coliforms		1	CFU/100mL					
Escherichia coli		1	CFU/100mL					
MW007: Coliforms by MF								
Coliforms		1	CFU/100mL					



QUALITY CONTROL REPORT

Work Order	: EP1608944	Page	: 1 of 4
Client	: GHD PTY LTD	Laboratory	: Environmental Division Perth
Contact	: MR ANDREW NAGLE	Contact	: Lauren Biagioni
Address	: 999 HAY STREET	Address	: 10 Hod Way Malaga WA Australia 6090
	PERTH WA 6000		
Telephone	: +61 08 6222 8222	Telephone	: 08 9209 7655
Project	: 61/34772 Point Moore Groundwater Assessment	Date Samples Received	: 23-Sep-2016
Order number	:	Date Analysis Commenced	: 23-Sep-2016
C-O-C number	:	Issue Date	: 03-Oct-2016
Sampler	: STEVEN PETTS		Hac-MRA NATA
Site	:		
Quote number	:		Accreditation No. 825
No. of samples received	: 7		Accredited for compliance with
No. of samples analysed	: 7		ISO/IEC 17025 - Testing

This report supersedes any previous report(s) with this reference. Results apply to the sample(s) as submitted. This document shall not be reproduced, except in full. This Quality Control Report contains the following information:

- Laboratory Duplicate (DUP) Report; Relative Percentage Difference (RPD) and Acceptance Limits
- Method Blank (MB) and Laboratory Control Spike (LCS) Report; Recovery and Acceptance Limits
- Matrix Spike (MS) Report; Recovery and Acceptance Limits

Signatories

This document has been electronically signed by the authorized signatories below. Electronic signing is carried out in compliance with procedures specified in 21 CFR Part 11.

Signatories	Position	Accreditation Category
Alini Goundar	Senior Analyst	Perth Microbiology, Malaga, WA
Jeremy Truong	Laboratory Manager	Perth Inorganics, Malaga, WA
Tyrone Cole	Inorganics Preparation Supervisor	Perth Inorganics, Malaga, WA
Vinitha Kesavan	Analyst	Perth Microbiology, Malaga, WA



General Comments

The analytical procedures used by the Environmental Division have been developed from established internationally recognized procedures such as those published by the USEPA, APHA, AS and NEPM. In house developed procedures are employed in the absence of documented standards or by client request.

Where moisture determination has been performed, results are reported on a dry weight basis.

Where a reported less than (<) result is higher than the LOR, this may be due to primary sample extract/digestate dilution and/or insufficient sample for analysis. Where the LOR of a reported result differs from standard LOR, this may be due to high

Key: Anonymous = Refers to samples which are not specifically part of this work order but formed part of the QC process lot

- CAS Number = CAS registry number from database maintained by Chemical Abstracts Services. The Chemical Abstracts Service is a division of the American Chemical Society.
- LOR = Limit of reporting
- RPD = Relative Percentage Difference

= Indicates failed QC

Laboratory Duplicate (DUP) Report

The quality control term Laboratory Duplicate refers to a randomly selected intralaboratory split. Laboratory duplicates provide information regarding method precision and sample heterogeneity. The permitted ranges for the Relative Percent Deviation (RPD) of Laboratory Duplicates are specified in ALS Method QWI-EN/38 and are dependent on the magnitude of results in comparison to the level of reporting: Result < 10 times LOR: No Limit; Result between 10 and 20 times LOR: 0% - 50%; Result > 20 times LOR: 0% - 20%.

Sub-Matrix: WATER			Laboratory Duplicate (DUP) Report						
Laboratory sample ID	Client sample ID	Method: Compound CAS Number		LOR	Unit	Original Result	Duplicate Result	RPD (%)	Recovery Limits (%)
EA015: Total Dissolv	ed Solids dried at 180 ± 5 $^\circ$	C (QC Lot: 602290)							
EP1608944-001	MW1	EA015H: Total Dissolved Solids @180°C		10	mg/L	22300	21700	2.96	0% - 20%
EP1609004-002	Anonymous	EA015H: Total Dissolved Solids @180°C		10	mg/L	227	222	2.45	0% - 20%
EK055G: Ammonia a	s N by Discrete Analyser(QC Lot: 595630)							
EP1608932-001	Anonymous	EK055G: Ammonia as N	7664-41-7	0.01	mg/L	0.03	0.04	0.00	No Limit
EP1608944-002	QC01_20160922	EK055G: Ammonia as N	7664-41-7	0.01	mg/L	6.42	6.47	0.792	0% - 20%
EK057G: Nitrite as N	by Discrete Analyser (QC	Lot: 595643)							
EP1608944-001	MW1	EK057G: Nitrite as N	14797-65-0	0.01	mg/L	<0.01	<0.01	0.00	No Limit
EP1608946-004	Anonymous	EK057G: Nitrite as N	14797-65-0	0.01	mg/L	<0.01	<0.01	0.00	No Limit
EK059G: Nitrite plus Nitrate as N (NOx) by Discrete Analyser (QC Lot: 595631)									
EP1608932-001	Anonymous	EK059G: Nitrite + Nitrate as N		0.01	mg/L	<0.01	<0.01	0.00	No Limit
EP1608944-002	QC01_20160922	EK059G: Nitrite + Nitrate as N		0.01	mg/L	<0.01	0.02	0.00	No Limit
EK061G: Total Kjeldahl Nitrogen By Discrete Analyser (QC Lot: 602454)									
EP1608944-005	SHP8	EK061G: Total Kjeldahl Nitrogen as N		0.1	mg/L	3.3	3.3	0.00	0% - 20%
EP1608957-004	Anonymous	EK061G: Total Kjeldahl Nitrogen as N		0.1	mg/L	0.5	1.0	61.4	No Limit
EK067G: Total Phosphorus as P by Discrete Analyser (QC Lot: 602453)									
EP1608944-005	SHP8	EK067G: Total Phosphorus as P		0.01	mg/L	0.38	0.39	3.39	0% - 20%
EP1608957-004	Anonymous	EK067G: Total Phosphorus as P		0.01	mg/L	<0.05	0.05	0.00	No Limit
EP030: Biochemical Oxygen Demand (BOD) (QC Lot: 595538)									
EP1608905-002	Anonymous	EP030: Biochemical Oxygen Demand		2	mg/L	245	253	3.13	0% - 20%
EP1608944-004	MW3	EP030: Biochemical Oxygen Demand		2	mg/L	5	4	24.4	No Limit



Method Blank (MB) and Laboratory Control Spike (LCS) Report

The quality control term Method / Laboratory Blank refers to an analyte free matrix to which all reagents are added in the same volumes or proportions as used in standard sample preparation. The purpose of this QC parameter is to monitor potential laboratory contamination. The quality control term Laboratory Control Spike (LCS) refers to a certified reference material, or a known interference free matrix spiked with target analytes. The purpose of this QC parameter is to monitor method precision and accuracy independent of sample matrix. Dynamic Recovery Limits are based on statistical evaluation of processed LCS.

Sub-Matrix: WATER				Method Blank (MB)	Laboratory Control Spike (LCS) Report				
				Report	Spike	Spike Recovery (%)	Recovery	Limits (%)	
Method: Compound	CAS Number	LOR	Unit	Result	Concentration	LCS	Low	High	
EA015: Total Dissolved Solids dried at 180 ± 5 $^\circ$ C (Q	CLot: 602290)								
EA015H: Total Dissolved Solids @180°C		10	mg/L	<10	2000 mg/L	96.8	83	111	
				<10	293 mg/L	112	70	130	
EK055G: Ammonia as N by Discrete Analyser (QCLo	EK055G: Ammonia as N by Discrete Analyser (QCLot: 595630)								
EK055G: Ammonia as N	7664-41-7	0.01	mg/L	<0.01	1 mg/L	107	87	115	
EK057G: Nitrite as N by Discrete Analyser (QCLot: 595643)									
EK057G: Nitrite as N	14797-65-0	0.01	mg/L	<0.01	0.5 mg/L	107	86	112	
EK059G: Nitrite plus Nitrate as N (NOx) by Discrete Analyser (QCLot: 595631)									
EK059G: Nitrite + Nitrate as N		0.01	mg/L	<0.01	0.5 mg/L	110	92	112	
EK061G: Total Kjeldahl Nitrogen By Discrete Analyse	er (QCLot: 602454)								
EK061G: Total Kjeldahl Nitrogen as N		0.1	mg/L	<0.1	10 mg/L	85.4	82	110	
EK067G: Total Phosphorus as P by Discrete Analyser (QCLot: 602453)									
EK067G: Total Phosphorus as P		0.01	mg/L	<0.01	4.42 mg/L	90.6	70	130	
EP030: Biochemical Oxygen Demand (BOD) (QCLot: 595538)									
EP030: Biochemical Oxygen Demand		2	mg/L	<2	198 mg/L	90.4	78	117	

Matrix Spike (MS) Report

The quality control term Matrix Spike (MS) refers to an intralaboratory split sample spiked with a representative set of target analytes. The purpose of this QC parameter is to monitor potential matrix effects on analyte recoveries. Static Recovery Limits as per laboratory Data Quality Objectives (DQOs). Ideal recovery ranges stated may be waived in the event of sample matrix interference.

Sub-Matrix: WATER				Matrix Spike (MS) Report			
				Spike	SpikeRecovery(%)	Recovery Li	mits (%)
Laboratory sample ID	Client sample ID	Method: Compound	CAS Number	Concentration	MS	Low	High
EK055G: Ammonia	as N by Discrete Analyser (QCLot: 595630)						
EP1608932-002	Anonymous	EK055G: Ammonia as N	7664-41-7	1 mg/L	127	70	130
EK057G: Nitrite as	N by Discrete Analyser (QCLot: 595643)						
EP1608944-002	QC01_20160922	EK057G: Nitrite as N	14797-65-0	0.5 mg/L	91.1	70	130
EK059G: Nitrite pl	us Nitrate as N (NOx) by Discrete Analyser(QCLot: 595	631)					
EP1608932-002	Anonymous	EK059G: Nitrite + Nitrate as N		0.5 mg/L	110	70	130
EK061G: Total Kjel	dahl Nitrogen By Discrete Analyser (QCLot: 602454)						
EP1608944-006	QC03_20160922	EK061G: Total Kjeldahl Nitrogen as N		5 mg/L	88.1	70	130
EK067G: Total Pho	sphorus as P by Discrete Analyser (QCLot: 602453)						
EP1608944-006	QC03_20160922	EK067G: Total Phosphorus as P		1 mg/L	95.9	70	130

Page	: 4 of 4
Work Order	: EP1608944
Client	: GHD PTY LTD
Project	: 61/34772 Point Moore Groundwater Assessment




QA/QC Compliance Assessment to assist with Quality Review								
Work Order	: EP1608944	Page	: 1 of 6					
Client		Laboratory	: Environmental Division Perth					
Contact	: MR ANDREW NAGLE	Telephone	: 08 9209 7655					
Project	: 61/34772 Point Moore Groundwater Assessment	Date Samples Received	: 23-Sep-2016					
Site	:	Issue Date	: 03-Oct-2016					
Sampler	: STEVEN PETTS	No. of samples received	: 7					
Order number	:	No. of samples analysed	: 7					

This report is automatically generated by the ALS LIMS through interpretation of the ALS Quality Control Report and several Quality Assurance parameters measured by ALS. This automated reporting highlights any non-conformances, facilitates faster and more accurate data validation and is designed to assist internal expert and external Auditor review. Many components of this report contribute to the overall DQO assessment and reporting for guideline compliance.

Brief method summaries and references are also provided to assist in traceability.

Summary of Outliers

Outliers : Quality Control Samples

This report highlights outliers flagged in the Quality Control (QC) Report.

- <u>NO</u> Method Blank value outliers occur.
- <u>NO</u> Duplicate outliers occur.
- <u>NO</u> Laboratory Control outliers occur.
- <u>NO</u> Matrix Spike outliers occur.
- For all regular sample matrices, <u>NO</u> surrogate recovery outliers occur.

Outliers : Analysis Holding Time Compliance

• <u>NO</u> Analysis Holding Time Outliers exist.

Outliers : Frequency of Quality Control Samples

• <u>NO</u> Quality Control Sample Frequency Outliers exist.



Analysis Holding Time Compliance

If samples are identified below as having been analysed or extracted outside of recommended holding times, this should be taken into consideration when interpreting results.

This report summarizes extraction / preparation and analysis times and compares each with ALS recommended holding times (referencing USEPA SW 846, APHA, AS and NEPM) based on the sample container provided. Dates reported represent first date of extraction or analysis and preclude subsequent dilutions and reruns. A listing of breaches (if any) is provided herein.

Holding time for leachate methods (e.g. TCLP) vary according to the analytes reported. Assessment compares the leach date with the shortest analyte holding time for the equivalent soil method. These are: organics 14 days, mercury 28 days & other metals 180 days. A recorded breach does not guarantee a breach for all non-volatile parameters.

Holding times for <u>VOC in soils</u> vary according to analytes of interest. Vinyl Chloride and Styrene holding time is 7 days; others 14 days. A recorded breach does not guarantee a breach for all VOC analytes and should be verified in case the reported breach is a false positive or Vinyl Chloride and Styrene are not key analytes of interest/concern.

Matrix: WATER					Evaluation	i: × = Holding time	breach ; 🗸 = With	in holding time
Method	Sample Date	Ex	traction / Preparation		Analysis			
Container / Client Sample ID(s)			Date extracted	Due for extraction	Evaluation	Date analysed	Due for analysis	Evaluation
EA015: Total Dissolved Solids dried at 180 ± 5 °C								
Clear Plastic Bottle - Natural (EA015H) MW1, MW2, SHP8, QC04_20160922	QC01_20160922, MW3, QC03_20160922,	23-Sep-2016				30-Sep-2016	30-Sep-2016	~
EK055G: Ammonia as N by Discrete Analyser								
Clear Plastic Bottle - Sulfuric Acid (EK055G) MW1, MW2, SHP8, QC04 20160922	QC01_20160922, MW3, QC03_20160922,	23-Sep-2016				23-Sep-2016	21-Oct-2016	~
EK057G: Nitrite as N by Discrete Analyser							1	
Clear Plastic Bottle - Natural (EK057G) MW1, MW2, SHP8, QC04_20160922	QC01_20160922, MW3, QC03_20160922,	23-Sep-2016				23-Sep-2016	25-Sep-2016	~
EK059G: Nitrite plus Nitrate as N (NOx) by Discr	rete Analyser							
Clear Plastic Bottle - Sulfuric Acid (EK059G) MW1, MW2, SHP8, QC04_20160922	QC01_20160922, MW3, QC03_20160922,	23-Sep-2016				23-Sep-2016	21-Oct-2016	~
EK061G: Total Kjeldahl Nitrogen By Discrete Ana	llyser							
Clear Plastic Bottle - Sulfuric Acid (EK061G) MW1, MW2, SHP8, QC04 20160922	QC01_20160922, MW3, QC03_20160922,	23-Sep-2016	03-Oct-2016	21-Oct-2016	~	03-Oct-2016	21-Oct-2016	~



Matrix: WATER			Evaluation: × = Holding time breach ; ✓ = Within holding time								
Method		Sample Date	E	traction / Preparation			Analysis				
Container / Client Sample ID(s)			Date extracted	Due for extraction	Evaluation	Date analysed	Due for analysis	Evaluation			
EK067G: Total Phosphorus as P by Discrete Analyser											
Clear Plastic Bottle - Sulfuric Acid (EK067G)											
MW1,	QC01_20160922,	23-Sep-2016	03-Oct-2016	21-Oct-2016	1	03-Oct-2016	21-Oct-2016	✓			
MW2,	MW3,										
SHP8,	QC03_20160922,										
QC04_20160922											
EP030: Biochemical Oxygen Demand (BOD)											
Clear Plastic Bottle - Natural (EP030)											
MW1,	QC01_20160922,	23-Sep-2016				23-Sep-2016	25-Sep-2016	✓			
MW2,	MW3,										
SHP8,	QC03_20160922,										
QC04_20160922											
MW002: Heterotrophic Plate Count											
Sterile Plastic Bottle - Sodium Thiosulfate (MW002)											
MW1,	QC01_20160922,	23-Sep-2016				23-Sep-2016	24-Sep-2016	✓			
MW2,	MW3,										
SHP8											
MW006: Faecal Coliforms & E.coli by MF											
Sterile Plastic Bottle - Sodium Thiosulfate (MW006)											
MW1,	QC01_20160922,	23-Sep-2016				23-Sep-2016	24-Sep-2016	✓			
MW2,	MW3,										
SHP8											
MW007: Coliforms by MF											
Sterile Plastic Bottle - Sodium Thiosulfate (MW007)											
MW1,	QC01_20160922,	23-Sep-2016				23-Sep-2016	24-Sep-2016	✓			
MW2,	MW3,										
SHP8											



Quality Control Parameter Frequency Compliance

The following report summarises the frequency of laboratory QC samples analysed within the analytical lot(s) in which the submitted sample(s) was(were) processed. Actual rate should be greater than or equal to the expected rate. A listing of breaches is provided in the Summary of Outliers.

Matrix: WATER			Evaluation: * = Quality Control frequency not within specification ; \checkmark = Quality Control frequency within specification.					
Quality Control Sample Type			Count		Rate (%)		Quality Control Specification	
Analytical Methods	Method	00	Reaular	Actual	Expected	Evaluation		
Laboratory Duplicates (DUP)								
Ammonia as N by Discrete analyser	EK055G	2	20	10.00	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Biochemical Oxygen Demand (BOD)	EP030	2	18	11.11	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	2	20	10.00	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite as N by Discrete Analyser	EK057G	2	17	11.76	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Dissolved Solids (High Level)	EA015H	2	15	13.33	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	2	20	10.00	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Phosphorus as P By Discrete Analyser	EK067G	2	20	10.00	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Laboratory Control Samples (LCS)								
Ammonia as N by Discrete analyser	EK055G	1	20	5.00	5.00	1	NEPM 2013 B3 & ALS QC Standard	
Biochemical Oxygen Demand (BOD)	EP030	1	18	5.56	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	1	20	5.00	5.00	~	NEPM 2013 B3 & ALS QC Standard	
Nitrite as N by Discrete Analyser	EK057G	1	17	5.88	5.00	~	NEPM 2013 B3 & ALS QC Standard	
Total Dissolved Solids (High Level)	EA015H	2	15	13.33	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Phosphorus as P By Discrete Analyser	EK067G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Method Blanks (MB)								
Ammonia as N by Discrete analyser	EK055G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Biochemical Oxygen Demand (BOD)	EP030	1	18	5.56	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	1	20	5.00	5.00	~	NEPM 2013 B3 & ALS QC Standard	
Nitrite as N by Discrete Analyser	EK057G	1	17	5.88	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Dissolved Solids (High Level)	EA015H	1	15	6.67	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Phosphorus as P By Discrete Analyser	EK067G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Matrix Spikes (MS)								
Ammonia as N by Discrete analyser	EK055G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite as N by Discrete Analyser	EK057G	1	17	5.88	5.00	~	NEPM 2013 B3 & ALS QC Standard	
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Phosphorus as P By Discrete Analyser	EK067G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	



Brief Method Summaries

The analytical procedures used by the Environmental Division have been developed from established internationally recognized procedures such as those published by the US EPA, APHA, AS and NEPM. In house developed procedures are employed in the absence of documented standards or by client request. The following report provides brief descriptions of the analytical procedures employed for results reported in the Certificate of Analysis. Sources from which ALS methods have been developed are provided within the Method Descriptions.

Analytical Methods	Method	Matrix	Method Descriptions
Total Dissolved Solids (High Level)	EA015H	WATER	In house: Referenced to APHA 2540C. A gravimetric procedure that determines the amount of `filterable` residue in an aqueous sample. A well-mixed sample is filtered through a glass fibre filter (1.2um). The filtrate is evaporated to dryness and dried to constant weight at 180+/-5C. This method is compliant with NEPM (2013) Schedule B(3)
Ammonia as N by Discrete analyser	EK055G	WATER	In house: Referenced to APHA 4500-NH3 G Ammonia is determined by direct colorimetry by Discrete Analyser. This method is compliant with NEPM (2013) Schedule B(3)
Nitrite as N by Discrete Analyser	EK057G	WATER	In house: Referenced to APHA 4500-NO2- B. Nitrite is determined by direct colourimetry by Discrete Analyser. This method is compliant with NEPM (2013) Schedule B(3)
Nitrate as N by Discrete Analyser	EK058G	WATER	In house: Referenced to APHA 4500-NO3- F. Nitrate is reduced to nitrite by way of a chemical reduction followed by quantification by Discrete Analyser. Nitrite is determined seperately by direct colourimetry and result for Nitrate calculated as the difference between the two results. This method is compliant with NEPM (2013) Schedule B(3)
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	WATER	In house: Referenced to APHA 4500-NO3- F. Combined oxidised Nitrogen (NO2+NO3) is determined by Chemical Reduction and direct colourimetry by Discrete Analyser. This method is compliant with NEPM (2013) Schedule B(3)
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	WATER	In house: Referenced to APHA 4500-Norg D (In house). An aliquot of sample is digested using a high temperature Kjeldahl digestion to convert nitrogenous compounds to ammonia. Ammonia is determined colorimetrically by discrete analyser. This method is compliant with NEPM (2013) Schedule B(3)
Total Nitrogen as N (TKN + Nox) By Discrete Analyser	EK062G	WATER	In house: Referenced to APHA 4500-Norg / 4500-NO3 This method is compliant with NEPM (2013) Schedule B(3)
Total Phosphorus as P By Discrete Analyser	EK067G	WATER	In house: Referenced to APHA 4500-P H, Jirka et al (1976), Zhang et al (2006). This procedure involves sulphuric acid digestion of a sample aliquot to break phosphorus down to orthophosphate. The orthophosphate reacts with ammonium molybdate and antimony potassium tartrate to form a complex which is then reduced and its concentration measured at 880nm using discrete analyser. This method is compliant with NEPM (2013) Schedule B(3)
Biochemical Oxygen Demand (BOD)	EP030	WATER	In house: Referenced to APHA 5210 B. The 5-Day BOD test provides an empirical measure of the oxygen consumption capacity of a given water. A portion of the sample is diluted into oxygenated, nutrient rich water, and a seed added to begin biological decay. The initial dissolved oxygen content is measured, then the bottle is sealed and incubated for five days. The remaining dissolved oxygen is measured, and from the difference, the demand for oxygen, by biological decay, is determined. This method is compliant with NEPM (2013) Schedule B(3)
Heterotrophic (Total) Plate Count @ 22C and 36C	MW002	WATER	In house: Referenced to AS4276.3.1- 2007
Thermotolerant Coliforms & E.coli by Membrane Filtration	MW006	WATER	In house: Referenced to AS 4276.7 2007
Coliforms by Membrane Filtration	MW007	WATER	In house: Referenced to AS 4276.5 - 2007
Preparation Methods	Method	Matrix	Method Descriptions

Page Work Order Client Project	: 6 of 6 : EP1608944 : GHD PTY LTD : 61/34772 Point Moore	Groundwater Assessment			(ALS)
Preparation Methods		Method	Matrix	Method Descriptions	
TKN/TP Digestion		EK061/EK067	WATER	In house: Referenced to APHA 4500 Norg - D; APHA 4500 P - H. This method is compliant with NEPM (2013) Schedule B(3)	

Beint Moore Group	lwater Assessment			Laborato	ry:MPL L	aboratorie	s			1									
Client	Water Assessment	Job No.		Address:	16-18 Ha	yden Cour	t Myar	ee WA 6	154	The Inc.									
City of Greater Gera	dton	61/34772		Laboratory Contact: Kiara Lockerbie															
Laboratory Quote N	0.	Turnaround Time				Contain	ner							Analyses	1			-	Remarks
Standar		Standard		Water/ A-Air	/P-Plastic	04/HNO3/Other			33,				MF)						
Job Manager (Invoid Andrew Nagle	e)	Email Address (Results) andrew.nagle@ghd.com steven.petts@ghd.com		-Sludge/ W-	Bag/G-Glass,	ed/ HCI/ H2S			x, NO2, NC	mand		ite Count a EA	& E. Coli (I						
GHD Sample ID	Laboratory Sample ID	Date	Time	ample Matrix s-soil/ si	YPE B-Bottle/J-Jar/V-Vial/I	reservative Unpreserve	9	otal Volume (mL)	otal Nitrogen, TKN, No: VH3, Total Phosphorus	siochemical Oxygen Dei	rDS - Standard Level	fotal (Heterotrophic Pla 22oC and 36oC (TPC) YI	fotal +Faecal Coliforms					Hold	
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Sampled by: S. Pett	s			Date/Tir	me: 22/09	/2016			Relinc	uished l	y: S.Pe	tts							Date/ Time: 22/9/2016
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16 - 18 Hayden Court, Myaree, Western Australia 6154 PO Box 4023 Myaree BC, Western Australia 6960 Tel: +61 8 9317 2505 / Fax: +61 8 9317 4163 email: laboratory@mpl.com.au www.mpl.com.au Envirolab Services (WA) Pty Ltd ABN 63 140 099 207

SAMPLE RECEIPT ADVICE

Client Details	
Client	GHD Services Pty Ltd
Attention	A Nagle

Sample Login Details	
Your Reference	61/34772 City of Greater Geraldton
Envirolab Reference	186264
Date Sample Received	23/09/2016
Date Instructions Received	23/09/2016
Date Results Expected to be Reported	03/10/2016

Sample Condition								
Samples received in appropriate condition for analysis	YES							
No. of Samples Provided	1 Water							
Turnaround Time Requested	Standard							
Temperature on receipt (°C)	7							
Cooling Method	Ice Pack							
Sampling Date Provided	NO							

Comments

Samples will be held for 1 month for water samples and 2 months for soil samples from date of receipt of samples

Please direct any queries to:

Joshua Lim	Meredith Conroy						
Phone: 08 9317 2505	Phone: 08 9317 2505						
Fax: 08 9317 4163	Fax: 08 9317 4163						
Email: jlim@mpl.com.au	Email: mconroy@mpl.com.au						

Sample and Testing Details on following page

16 - 18 Hayden Court, Myaree, Western Australia 6154 PO Box 4023 Myaree BC, Western Australia 6960 Tel: +61 8 9317 2505 / Fax: +61 8 9317 4163 email: laboratory@mpl.com.au www.mpl.com.au Envirolab Services (WA) Pty Ltd ABN 53 140 099 207





Sample Id	BOD	Total Dissolved Solids (grav)	Microbiological Testing	Nutrients in Water
QC02_20160922	✓	~	1	1



email: lab@mpl.com.au envirolab.com.au

Envirolab Services (WA) Pty Ltd trading as MPL Laboratories | ABN 53 140 099 207

CERTIFICATE OF ANALYSIS 186264

Client: GHD Services Pty Ltd PO Box Y3106 Perth WA 6000

Attention: A Nagle

Sample log in details:

Your Reference: No. of samples: Date samples received: Date completed instructions received: Location:

61/34772 City of Greater Geraldton

1 Water 23/09/2016 23/09/2016

Analysis Details:

Please refer to the following pages for results, methodology summary and quality control data. Samples were analysed as received from the client. Results relate specifically to the samples as received. Results are reported on a dry weight basis for solids and on an as received basis for other matrices. *Please refer to the last pages of this report for any comments relating to the results.*

Report Details:

 Date results requested by:
 3/10/16

 Date of Preliminary Report:
 not issued

 Issue Date:
 30/09/16

 NATA accreditation number 2901. This document shall not be reproduced except in full.

 Accredited for compliance with ISO/IEC 17025 - Testing

 Tests not covered by NATA are denoted with *.

Results Approved By:

Joshua Lim Operations Manager

MPL Reference: Revision No: 186264 R 00

Jessica Miller Micro Team Leader



Page 1 of 8

Client Reference: 61/34772 City of Greater Geraldton

Miscellaneous Inorganics		
Our Reference:	UNITS	186264-1
Your Reference		QC02_20160
		922
Type of sample		Water
Date prepared	-	28/09/2016
Date analysed	-	28/09/2016
BOD	mg/L	<5
Total Dissolved Solids (grav)	mg/L	22,000

Client Reference:

61/34772 City of Greater Geraldton

Microbiological Testing		
Our Reference:	UNITS	186264-1
Your Reference		QC02_20160
		922
Type of sample		Water
Date testing started	-	23/09/2016
Date testing completed	-	28/09/2016
Heterotrophic Plate Count 21C	cfu/mL	340
Heterotrophic Plate Count 35C	cfu/mL	260
Total Coliforms	cfu/100mL	<1
Thermotolerant Coliforms	cfu/100mL	<1
E.coli	cfu/100mL	<1

Client Reference:

61/34772 City of Greater Geraldton

Nutrients in Water		
Our Reference:	UNITS	186264-1
Your Reference		QC02_20160
		922
Type of sample		Water
Date prepared	-	23/09/2016
Date analysed	-	23/09/2016
Total Nitrogen	mg/L	9.2
TKN by Discrete Analyser	mg/L	9.2
NOx as N	mg/L	0.009
Nitrate as N	mg/L	<0.005
Nitrite as N	mg/L	0.006
Ammonia as N	mg/L	5.0
Total Phosphorus	mg/L	0.78

Client Reference: 61/34772 City of Greater Geraldton

MethodID	Methodology Summary
INORG-091	BOD - Analysed in accordance with APHA latest edition 5210 D.
INORG-018	Total Dissolved Solids - determined gravimetrically. The solids are dried at $180\pm5^{\circ}$ C
MICRO-001	Heterotrophic Plate Count: Microbial Water Analysis - in accordance with MICRO-001 (APHA-9215D-2005). Recommended maximums based on NHMRC and ARMC Australian Drinking Water Guidelines.
MICRO-001	Total Coliforms: Microbial Water Analysis - in accordance with MICRO-001 (AS4276.5-2007). Recommended maximums based on NHMRC and ARMC Australian Drinking Water Guidelines.
MICRO-001	Thermotolerant Coliforms: Microbial Water Analysis - in accordance with MICRO-001 (AS4276.7-2007). Recommended maximums based on NHMRC and ARMC Australian Drinking Water Guidelines.
MICRO-001	E. Coli: Microbial Water Analysis - in accordance with MICRO-001 (AS4276.7-2007). Recommended maximums based on NHMRC and ARMC Australian Drinking Water Guidelines.
INORG-055	Total Nitrogen by colourimetric analysis based on APHA 4500-P J, 4500-NO3 F.
INORG-062	TKN by calculation fromTotal Nitrogen and NOx using APHA methodology.
INORG-055	NOx - determined colourimetrically. Soils are analysed from a water extract.
INORG-055	Nitrate - determined colourimetrically. Soils are analysed from a water extract.
INORG-055	Nitrite - determined colourimetrically. Soils are analysed from a water extract.
INORG-057	Ammonia by colourimetric analysis based on APHA latest edition 4500-NH3 F.
METALS-020	Metals in soil and water by ICP-OES.

	Client Reference: 61/34772 City of Greater Geraldton							
QUALITYCONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Miscellaneous Inorganics						Base II Duplicate II % RPD		
Date prepared	-			28/09/ 2016	[NT]	[NT]	LCS-1	28/09/2016
Date analysed	-			28/09/ 2016	[NT]	[NT]	LCS-1	28/09/2016
BOD	mg/L	5	INORG-091	⊲5	[NT]	[NT]	LCS-1	70%
Total Dissolved Solids (grav)	mg/L	5	INORG-018	⊲5	[NT]	[NT]	LCS-1	102%
QUALITYCONTROL	UNITS	PQL	METHOD	Blank				
Microbiological Testing								
Date testing started	-			23/09/				
, , , , , , , , , , , , , , , , , , ,				2016				
Date testing	-			28/09/				
completed				2016				
Heterotrophic Plate Count 21C	cfu/mL	10	MICRO-001	<10				
Heterotrophic Plate Count 35C	cfu/mL	10	MICRO-001	<10				
Total Coliforms	cfu/100 mL	1	MICRO-001	<1				
Thermotolerant Coliforms	cfu/100 mL	1	MICRO-001	<1				
E.coli	cfu/100 mL	1	MICRO-001	<1				
QUALITYCONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Nutrients in Water						Base II Duplicate II % RPD		
Date prepared	-			23/09/ 2016	[NT]	[NT]	LCS-1	23/09/2016
Date analysed	-			23/09/ 2016	[NT]	[NT]	LCS-1	23/09/2016
Total Nitrogen	mg/L	0.1	INORG-055	<0.1	[NT]	[NT]	LCS-1	118%
TKN by Discrete Analyser	mg/L	0.1	INORG-062	<0.1	[NT]	[NT]	[NR]	[NR]
NOx as N	mg/L	0.005	INORG-055	<0.005	[NT]	[NT]	LCS-1	103%
Nitrate as N	mg/L	0.005	INORG-055	<0.005	[NT]	[NT]	LCS-1	103%
Nitrite as N	mg/L	0.005	INORG-055	<0.005	[NT]	[NT]	LCS-1	106%
Ammonia as N	mg/L	0.005	INORG-057	<0.005	[NT]	[NT]	LCS-1	90%
Total Phosphorus	mg/L	0.05	METALS- 020	<0.05	[NT]	[NT]	LCS-1	100%

Report Comments:

Definitions:

NT: Not tested NA: Test not required INS: Insufficient sample for this test PQL: Practical Quantitation Limit <: Less than >: Greater than RPD: Relative Percent Difference LCS: Laboratory Control Sample NS: Not Specified NEPM: National Environmental Protection Measure NR: Not Reported

Australian Drinking Water Guidelines recommend that Thermotolerant Coliform, Faecal Enterococci, & E.Coli levels are less than 1cfu/100mL. The recommended maximums are taken from "Australian Drinking Water Guidelines", published by NHMRC & ARMC 2011

Quality Control Definitions

Blank: This is the component of the analytical signal which is not derived from the sample but from reagents, glassware etc, can be determined by processing solvents and reagents in exactly the same manner as for samples. **Duplicate**: This is the complete duplicate analysis of a sample from the process batch. If possible, the sample selected should be one where the analyte concentration is easily measurable.

Matrix Spike : A portion of the sample is spiked with a known concentration of target analyte. The purpose of the matrix spike is to monitor the performance of the analytical method used and to determine whether matrix interferences exist.

LCS (Laboratory Control Sample) : This comprises either a standard reference material or a control matrix (such as a blank sand or water) fortified with analytes representative of the analyte class. It is simply a check sample.

Surrogate Spike: Surrogates are known additions to each sample, blank, matrix spike and LCS in a batch, of compounds which are similar to the analyte of interest, however are not expected to be found in real samples.

Laboratory Acceptance Criteria

Duplicate sample and matrix spike recoveries may not be reported on smaller jobs, however, were analysed at a frequency to meet or exceed NEPM requirements. All samples are tested in batches of 20. The duplicate sample RPD and matrix spike recoveries for the batch were within the laboratory acceptance criteria.

Filters, swabs, wipes, tubes and badges will not have duplicate data as the whole sample is generally extracted during sample extraction.

Spikes for Physical and Aggregate Tests are not applicable.

For VOCs in water samples, three vials are required for duplicate or spike analysis.

Duplicates: <5xPQL - any RPD is acceptable; >5xPQL - 0-50% RPD is acceptable. Matrix Spikes, LCS and Surrogate recoveries: Generally 70-130% for inorganics/metals; 60-140% for organics (+/-50% surrogates) and 10-140% for labile SVOCs (including labile surrogates), ultra trace organics and speciated phenols is acceptable.

In circumstances where no duplicate and/or sample spike has been reported at 1 in 10 and/or 1 in 20 samples respectively, the sample volume submitted was insufficient in order to satisfy laboratory QA/QC protocols.

When samples are received where certain analytes are outside of recommended technical holding times (THTs), the analysis has proceeded. Where analytes are on the verge of breaching THTs, every effort will be made to analyse within the THT or as soon as practicable.

Where sampling dates are not provided, Envirolab are not in a position to comment on the validity of the analysis where recommended technical holding times may have been breached.

Measurement Uncertainty estimates are available for most tests upon request.

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	w	BP	H2SO4	2	560	\mathbf{X}	\mathbf{X}	\boxtimes											
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Perth Work Order Reference EP1609847

* *



Telephone - 61-8-9209 7655



SAMPLE RECEIPT NOTIFICATION (SRN)

Work Order	: EP1609847						
Client Contact Address	: GHD PTY LTD : MR ANDREW NAGLE : 999 HAY STREET PERTH WA 6000	Laboratory Contact Address	: Environmental Division Perth : Lauren Biagioni : 10 Hod Way Malaga WA Australia 6090				
E-mail Telephone Facsimile	: andrew.nagle@ghd.com : +61 08 6222 8222 : +61 08 9429 6555	E-mail Telephone Facsimile	: Lauren.biagioni@alsglobal.com : 08 9209 7655 : +61-8-9209 7600				
Project	: 61/34772 Point Moore Groundwater Assessment	Page	: 1 of 2				
Order number C-O-C number Site Sampler	: : : : STEVEN PETTS	Quote number: EP2016GHDSER0029 (EP/919)QC Level: NEPM 2013 B3 & ALS QC Stan					
Dates Date Samples Receive Client Requested Due Date	d : 19-Oct-2016 9:25 AM : 26-Oct-2016	Issue Date Scheduled Reporting D	: 19-Oct-2016 Date : 26-Oct-2016				
Delivery Details Mode of Delivery : Carrier No. of coolers/boxes : 1 Receipt Detail :		Security Seal Temperature No. of samples receive	: Intact. : 3.5 - Ice present ed / analysed : 7 / 7				

General Comments

- This report contains the following information:
 - Sample Container(s)/Preservation Non-Compliances
 - Summary of Sample(s) and Requested Analysis
 - Proactive Holding Time Report
 - Requested Deliverables
- Please see scanned COC for sample discrepencies: extra samples , samples not received etc.
- Please direct any queries related to sample condition / numbering / breakages to Sample Receipt (SamplesPerth@alsenviro.com)
- Analytical work for this work order will be conducted at ALS Environmental Perth.
- Please direct any turnaround / technical queries to the laboratory contact designated above.
- Sample Disposal Aqueous (14 days), Solid (60 days) from date of completion of Work Order.
- pH analysis should be conducted within 6 hours of sampling.



en + NO2 + NO3 + NH3 + Total P

1002 (22C and 36C) nic Colony Count (22C & 36C)

Level

- High

Sample Container(s)/Preservation Non-Compliances

All comparisons are made against pretreatment/preservation AS, APHA, USEPA standards.

• No sample container / preservation non-compliance exists.

Summary of Sample(s) and Requested Analysis

Some items described below may be part of a laboratory process necessary for the execution of client requested tasks. Packages may contain additional analyses, such as the determination of moisture content and preparation tasks, that are included in the package.

If no sampling time is provided, the sampling time will default to 15:00 on the date of sampling. If no sampling date is provided, the sampling date will be assumed by the laboratory for processing purposes and will be shown bracketed without a time component.

Matrix: WATER

e sampling dat ocessing purpos e component.	e will be assumed by ses and will be shown	TER - EA015H I Dissolved Solids - H	'ER - EP030	TER - M5 - ES/EB/FM E.coli & TC by MF	ER - MW002 (22C and rotrophic Colony Coun	'ER - NT-08 I Nitrogen + NO2 + NO
date / time	· · · · · · · · · · · · · · · · · · ·	WAT Tota	WAT BOD	WAT FC,	WAT Hete	WAT Tota
[18-Oct-2016]	MW1	✓	1	✓	✓	✓
[18-Oct-2016]	QC01_20160922	✓	1	✓	1	✓
[18-Oct-2016]	MW2	✓	1	✓	✓	1
[18-Oct-2016]	MW3	✓	1	✓	1	1
[18-Oct-2016]	SHP8	1	1	1	✓	✓
[18-Oct-2016]	QC03_20160922	1	1			✓
[18-Oct-2016]	QC04_20160922	1	1			✓
	 Sampling data Sampling data Scessing purpose component. Client sampling date / time 18-Oct-2016] 	 Sampling date will be assumed by pocessing purposes and will be shown a component. Client sampling Client sample ID date / time 18-Oct-2016] MW1 18-Oct-2016] QC01_20160922 18-Oct-2016] MW2 18-Oct-2016] MW3 18-Oct-2016] SHP8 18-Oct-2016] QC03_20160922 18-Oct-2016] QC04_20160922 	 sampling date will be assumed by cleans and will be shown a component. Client sampling client sample ID date / time 18-Oct-2016] MW1 18-Oct-2016] QC01_20160922 18-Oct-2016] MW2 18-Oct-2016] MW3 18-Oct-2016] SHP8 18-Oct-2016] QC03_20160922 18-Oct-2016] QC04_20160922 	 Sampling date will be assumed by cleans and will be shown e component. Client sampling client sample ID date / time 18-Oct-2016] MW1 QC01_20160922 ✓ 18-Oct-2016] MW2 MW3 ✓ 18-Oct-2016] SHP8 QC03_20160922 ✓ 18-Oct-2016] QC04_20160922 ✓ ✓ MW3 ✓ ✓ ✓ ✓ (✓ ✓ ✓	 Sampling date will be assumed by cleans and will be shown e component. Client sampling date vill be assumed by client sample ID Client sampling date / time Client sample ID MW1 V V MW2 V V V V MW3 V <liv< li=""> V <liv< li=""> V V<td>Sampling date will be assumed by coessing purposes and will be shown e component. H Image: Client sample ID Image: Client sample ID</td></liv<></liv<>	Sampling date will be assumed by coessing purposes and will be shown e component. H Image: Client sample ID Image: Client sample ID

Proactive Holding Time Report

Sample(s) have been received within the recommended holding times for the requested analysis.

Requested Deliverables

ANDREW NAGLE

- *AU Certificate of Analysis - NATA (COA)	Email	andrew.nagle@ghd.com
- *AU Interpretive QC Report - DEFAULT (Anon QCI Rep) (QCI)	Email	andrew.nagle@ghd.com
- *AU QC Report - DEFAULT (Anon QC Rep) - NATA (QC)	Email	andrew.nagle@ghd.com
- A4 - AU Sample Receipt Notification - Environmental HT (SRN)	Email	andrew.nagle@ghd.com
- A4 - AU Tax Invoice (INV)	Email	andrew.nagle@ghd.com
- Chain of Custody (CoC) (COC)	Email	andrew.nagle@ghd.com
- EDI Format - ENMRG (ENMRG)	Email	andrew.nagle@ghd.com
- EDI Format - ESDAT (ESDAT)	Email	andrew.nagle@ghd.com
- EDI Format - XTab (XTAB)	Email	andrew.nagle@ghd.com
STEVEN PETTS		
 *AU Certificate of Analysis - NATA (COA) 	Email	steven.petts@ghd.com
- *AU Interpretive QC Report - DEFAULT (Anon QCI Rep) (QCI)	Email	steven.petts@ghd.com
- *AU QC Report - DEFAULT (Anon QC Rep) - NATA (QC)	Email	steven.petts@ghd.com
- A4 - AU Sample Receipt Notification - Environmental HT (SRN)	Email	steven.petts@ghd.com
- Chain of Custody (CoC) (COC)	Email	steven.petts@ghd.com
- EDI Format - ENMRG (ENMRG)	Email	steven.petts@ghd.com
- EDI Format - ESDAT (ESDAT)	Email	steven.petts@ghd.com
- EDI Format - XTab (XTAB)	Email	steven.petts@ghd.com



CERTIFICATE OF ANALYSIS Work Order : EP1609847 Page : 1 of 4 Amendment :1 Client Laboratory GHD PTY LTD : Environmental Division Perth Contact : MR ANDREW NAGLE Contact : Lauren Biagioni Address Address : 10 Hod Way Malaga WA Australia 6090 : 999 HAY STREET PERTH WA 6000 Telephone : +61 08 6222 8222 Telephone : 08 9209 7655 Project : 61/34772 Point Moore Groundwater Assessment **Date Samples Received** : 19-Oct-2016 09:25 Order number Date Analysis Commenced : -----: 19-Oct-2016 C-O-C number Issue Date · 02-Nov-2016 11:53 · ____ Sampler : STEVEN PETTS Site · ----Quote number : -----Accreditation No. 825 No. of samples received : 7 Accredited for compliance with ISO/IEC 17025 - Testing No. of samples analysed : 7

This report supersedes any previous report(s) with this reference. Results apply to the sample(s) as submitted. This document shall not be reproduced, except in full.

This Certificate of Analysis contains the following information:

- General Comments
- Analytical Results

Additional information pertinent to this report will be found in the following separate attachments: Quality Control Report, QA/QC Compliance Assessment to assist with Quality Review and Sample Receipt Notification.

Signatories

This document has been electronically signed by the authorized signatories below. Electronic signing is carried out in compliance with procedures specified in 21 CFR Part 11.

Signatories	Position	Accreditation Category
Jeremy Truong	Laboratory Manager	Perth Inorganics, Malaga, WA
Tyrone Cole	Inorganics Preparation Supervisor	Perth Inorganics, Malaga, WA
Vinitha Kesavan	Analyst	Perth Microbiology, Malaga, WA



General Comments

The analytical procedures used by the Environmental Division have been developed from established internationally recognized procedures such as those published by the USEPA, APHA, AS and NEPM. In house developed procedures are employed in the absence of documented standards or by client request.

Where moisture determination has been performed, results are reported on a dry weight basis.

Where a reported less than (<) result is higher than the LOR, this may be due to primary sample extract/digestate dilution and/or insufficient sample for analysis.

Where the LOR of a reported result differs from standard LOR, this may be due to high moisture content, insufficient sample (reduced weight employed) or matrix interference.

When sampling time information is not provided by the client, sampling dates are shown without a time component. In these instances, the time component has been assumed by the laboratory for processing purposes.

Where a result is required to meet compliance limits the associated uncertainty must be considered. Refer to the ALS Contact for details.

Key: CAS Number = CAS registry number from database maintained by Chemical Abstracts Services. The Chemical Abstracts Service is a division of the American Chemical Society.

LOR = Limit of reporting

^ = This result is computed from individual analyte detections at or above the level of reporting

ø = ALS is not NATA accredited for these tests.

- ~ = Indicates an estimated value.
- CFU = colony forming unit
- MF = membrane filtration
- This report has been amended to update the sample ID's for the QC samples, as per the client's request. There are no changes to the results.
- MW007 and MW006: estimate (~) is reported where there are many non-target colonies; the typical colonies may be masked by overgrowth of non-target organisms. It may be informative to record this fact.
- MW002 is ALS's internal code and is equivalent to AS4276.3.1.
- MW006 is ALS's internal code and is equivalent to AS4276.7.
- MW007 is ALS's internal code and is equivalent to AS4276.5.



Analytical Results

Sub-Matrix: WATER (Matrix: WATER)		Clie	ent sample ID	MW1	QC01_20161018	MW2	MW3	SHP8
	Cli	ent sampli	ng date / time	[18-Oct-2016]	[18-Oct-2016]	[18-Oct-2016]	[18-Oct-2016]	[18-Oct-2016]
Compound	CAS Number	LOR	Unit	EP1609847-001	EP1609847-002	EP1609847-003	EP1609847-004	EP1609847-005
				Result	Result	Result	Result	Result
EA015: Total Dissolved Solids dried at 1	80 ± 5 °C							
Total Dissolved Solids @180°C		10	mg/L	13100	13100	1360	1340	2290
EK055G: Ammonia as N by Discrete Ana	lyser							
Ammonia as N	7664-41-7	0.01	mg/L	3.95	4.00	1.48	0.32	0.46
EK057G: Nitrite as N by Discrete Analys	er							
Nitrite as N	14797-65-0	0.01	mg/L	<0.01	<0.01	0.27	<0.01	<0.01
EK058G: Nitrate as N by Discrete Analys	ser							
Nitrate as N	14797-55-8	0.01	mg/L	0.46	0.46	6.06	<0.01	<0.01
EK059G: Nitrite plus Nitrate as N (NOx)	by Discrete Ana	lyser						
Nitrite + Nitrate as N		0.01	mg/L	0.46	0.46	6.33	<0.01	<0.01
EK061G: Total Kjeldahl Nitrogen By Disc	rete Analyser							
Total Kjeldahl Nitrogen as N		0.1	mg/L	4.2	4.1	2.6	0.6	1.2
EK062G: Total Nitrogen as N (TKN + NO)	k) by Discrete An	alyser						
^ Total Nitrogen as N		0.1	mg/L	4.7	4.6	8.9	0.6	1.2
EK067G: Total Phosphorus as P by Disc	rete Analyser							
Total Phosphorus as P		0.01	mg/L	0.47	0.46	0.18	0.08	0.28
EP030: Biochemical Oxygen Demand (B	OD)							
Biochemical Oxygen Demand		2	mg/L	33	27	3	3	57
MW002: Heterotrophic Plate Count								
Heterotrophic Plate Count (22°C)		1	CFU/mL	4200	4400	4500	14000	2000
Heterotrophic Plate Count (36°C)		1	CFU/mL	4400	3600	7400	5500	980
MW006: Faecal Coliforms & E.coli by MF								
Faecal Coliforms		1	CFU/100mL	~<1	~<1	~<1	~<1	~<1
Escherichia coli		1	CFU/100mL	~<1	~<1	~<1	~<1	~<1
MW007: Coliforms by MF								
Coliforms		1	CFU/100mL	~<1	~<1	~<1	~<1	~<1



Analytical Results

Sub-Matrix: WATER (Matrix: WATER)		Cli	ent sample ID	QC03_20161018	QC04_20161018	 	
	Cli	ient sampli	ng date / time	[18-Oct-2016]	[18-Oct-2016]	 	
Compound	CAS Number	LOR	Unit	EP1609847-006	EP1609847-007	 	
				Result	Result	 	
EA015: Total Dissolved Solids dried at 1	80 ± 5 °C						
Total Dissolved Solids @180°C		10	mg/L	14	16	 	
EK055G: Ammonia as N by Discrete Ana	lyser						
Ammonia as N	7664-41-7	0.01	mg/L	0.06	0.06	 	
EK057G: Nitrite as N by Discrete Analys	er						
Nitrite as N	14797-65-0	0.01	mg/L	<0.01	<0.01	 	
EK058G: Nitrate as N by Discrete Analys	ser						
Nitrate as N	14797-55-8	0.01	mg/L	<0.01	0.01	 	
EK059G: Nitrite plus Nitrate as N (NOx)	by Discrete Ana	lyser					
Nitrite + Nitrate as N		0.01	mg/L	<0.01	0.01	 	
EK061G: Total Kjeldahl Nitrogen By Disc	crete Analyser						
Total Kjeldahl Nitrogen as N		0.1	mg/L	<0.1	<0.1	 	
EK062G: Total Nitrogen as N (TKN + NO)	x) by Discrete An	alyser					
^ Total Nitrogen as N		0.1	mg/L	<0.1	<0.1	 	
EK067G: Total Phosphorus as P by Disc	rete Analyser						
Total Phosphorus as P		0.01	mg/L	<0.01	<0.01	 	
EP030: Biochemical Oxygen Demand (B	OD)						
Biochemical Oxygen Demand		2	mg/L	<2	<2	 	
MW002: Heterotrophic Plate Count							
Heterotrophic Plate Count (22°C)		1	CFU/mL			 	
Heterotrophic Plate Count (36°C)		1	CFU/mL			 	
MW006: Faecal Coliforms & E.coli by MF	;						
Faecal Coliforms		1	CFU/100mL			 	
Escherichia coli		1	CFU/100mL			 	
MW007: Coliforms by MF							
Coliforms		1	CFU/100mL			 	



QUALITY CONTROL REPORT

Work Order	: EP1609847	Page	: 1 of 4
Amendment	: 1		
Client Contact Address	: GHD PTY LTD : MR ANDREW NAGLE : 999 HAY STREET	Laboratory Contact Address	: Environmental Division Perth : Lauren Biagioni : 10 Hod Way Malaga WA Australia 6090
Telephone Project Order number	PERTH WA 6000 : +61 08 6222 8222 : 61/34772 Point Moore Groundwater Assessment :	Telephone Date Samples Received Date Analysis Commenced	: 08 9209 7655 : 19-Oct-2016 : 19-Oct-2016
C-O-C number Sampler Site Quote number No. of samples received No. of samples analysed	: : STEVEN PETTS : : : 7 : 7	Issue Date	: 02-Nov-2016

This report supersedes any previous report(s) with this reference. Results apply to the sample(s) as submitted. This document shall not be reproduced, except in full.

This Quality Control Report contains the following information:

- Laboratory Duplicate (DUP) Report; Relative Percentage Difference (RPD) and Acceptance Limits
- Method Blank (MB) and Laboratory Control Spike (LCS) Report; Recovery and Acceptance Limits
- Matrix Spike (MS) Report; Recovery and Acceptance Limits

Signatories

This document has been electronically signed by the authorized signatories below. Electronic signing is carried out in compliance with procedures specified in 21 CFR Part 11.

Signatories	Position	Accreditation Category
Jeremy Truong	Laboratory Manager	Perth Inorganics, Malaga, WA
Tyrone Cole	Inorganics Preparation Supervisor	Perth Inorganics, Malaga, WA
Vinitha Kesavan	Analyst	Perth Microbiology, Malaga, WA



General Comments

The analytical procedures used by the Environmental Division have been developed from established internationally recognized procedures such as those published by the USEPA, APHA, AS and NEPM. In house developed procedures are employed in the absence of documented standards or by client request.

Where moisture determination has been performed, results are reported on a dry weight basis.

Where a reported less than (<) result is higher than the LOR, this may be due to primary sample extract/digestate dilution and/or insufficient sample for analysis. Where the LOR of a reported result differs from standard LOR, this may be due to high

Key: Anonymous = Refers to samples which are not specifically part of this work order but formed part of the QC process lot

- CAS Number = CAS registry number from database maintained by Chemical Abstracts Services. The Chemical Abstracts Service is a division of the American Chemical Society.
- LOR = Limit of reporting
- RPD = Relative Percentage Difference

= Indicates failed QC

Laboratory Duplicate (DUP) Report

The quality control term Laboratory Duplicate refers to a randomly selected intralaboratory split. Laboratory duplicates provide information regarding method precision and sample heterogeneity. The permitted ranges for the Relative Percent Deviation (RPD) of Laboratory Duplicates are specified in ALS Method QWI-EN/38 and are dependent on the magnitude of results in comparison to the level of reporting: Result < 10 times LOR: No Limit; Result between 10 and 20 times LOR: 0% - 50%; Result > 20 times LOR: 0% - 20%.

Sub-Matrix: WATER				Laboratory Duplicate (DUP) Report					
Laboratory sample ID	Client sample ID	Method: Compound	CAS Number	LOR	Unit	Original Result	Duplicate Result	RPD (%)	Recovery Limits (%)
EA015: Total Dissolv	ed Solids dried at 180 ± 5 °(C (QC Lot: 627327)							
EP1609847-001	MW1	EA015H: Total Dissolved Solids @180°C		10	mg/L	13100	12800	2.00	0% - 20%
EP1609866-001	Anonymous	EA015H: Total Dissolved Solids @180°C		10	mg/L	644	684	6.01	0% - 20%
EK055G: Ammonia as	s N by Discrete Analyser(QC Lot: 623447)							
EP1609847-001	MW1	EK055G: Ammonia as N	7664-41-7	0.01	mg/L	3.95	3.93	0.520	0% - 20%
EP1609870-001	Anonymous	EK055G: Ammonia as N	7664-41-7	0.01	mg/L	0.03	0.03	0.00	No Limit
EK057G: Nitrite as N	by Discrete Analyser (QC	Lot: 623435)							
EP1609882-001	Anonymous	EK057G: Nitrite as N	14797-65-0	0.01	mg/L	0.02	0.02	0.00	No Limit
EP1609888-010	Anonymous	EK057G: Nitrite as N	14797-65-0	0.01	mg/L	0.15	0.14	0.00	0% - 50%
EK059G: Nitrite plus	Nitrate as N (NOx) by Disc	rete Analyser (QC Lot: 623448)							
EP1609847-001	MW1	EK059G: Nitrite + Nitrate as N		0.01	mg/L	0.46	0.46	0.00	0% - 20%
EP1609870-001	Anonymous	EK059G: Nitrite + Nitrate as N		0.01	mg/L	1.56	1.50	3.98	0% - 20%
EK061G: Total Kjelda	hl Nitrogen By Discrete An	alyser (QC Lot: 628160)							
EP1608883-001	Anonymous	EK061G: Total Kjeldahl Nitrogen as N		0.1	mg/L	1.2	1.2	0.00	0% - 50%
EP1609847-003	MW2	EK061G: Total Kjeldahl Nitrogen as N		0.1	mg/L	2.6	2.6	0.00	0% - 50%
EK067G: Total Phosp	horus as P by Discrete Ana	alyser (QC Lot: 628159)							
EP1608883-001	Anonymous	EK067G: Total Phosphorus as P		0.01	mg/L	0.20	0.20	0.00	0% - 20%
EP1609847-003	MW2	EK067G: Total Phosphorus as P		0.01	mg/L	0.18	0.17	8.86	No Limit
EP030: Biochemical	Oxygen Demand (BOD) (Q	C Lot: 625068)							
EP1609847-006	QC03_20161018	EP030: Biochemical Oxygen Demand		2	mg/L	<2	<2	0.00	No Limit
EP1609931-003	Anonymous	EP030: Biochemical Oxygen Demand		2	mg/L	<2	<2	0.00	No Limit



Method Blank (MB) and Laboratory Control Spike (LCS) Report

The quality control term Method / Laboratory Blank refers to an analyte free matrix to which all reagents are added in the same volumes or proportions as used in standard sample preparation. The purpose of this QC parameter is to monitor potential laboratory contamination. The quality control term Laboratory Control Spike (LCS) refers to a certified reference material, or a known interference free matrix spiked with target analytes. The purpose of this QC parameter is to monitor method precision and accuracy independent of sample matrix. Dynamic Recovery Limits are based on statistical evaluation of processed LCS.

Sub-Matrix: WATER			Method Blank (MB)	Laboratory Control Spike (LCS) Report				
				Report	Spike	Spike Recovery (%)	Recovery	Limits (%)
Method: Compound	CAS Number	LOR	Unit	Result	Concentration	LCS	Low	High
EA015: Total Dissolved Solids dried at 180 \pm 5 °C (QC	Lot: 627327)							
EA015H: Total Dissolved Solids @180°C		10	mg/L	<10	2000 mg/L	106	83	111
				<10	1000 mg/L	109	70	130
EK055G: Ammonia as N by Discrete Analyser (QCLot:	623447)							
EK055G: Ammonia as N	7664-41-7	0.01	mg/L	<0.01	1 mg/L	101	87	115
EK057G: Nitrite as N by Discrete Analyser (QCLot: 62	3435)							
EK057G: Nitrite as N	14797-65-0	0.01	mg/L	<0.01	0.5 mg/L	102	86	112
EK059G: Nitrite plus Nitrate as N (NOx) by Discrete A	nalyser (QCLot: 62	3448)						
EK059G: Nitrite + Nitrate as N		0.01	mg/L	<0.01	0.5 mg/L	109	92	112
EK061G: Total Kjeldahl Nitrogen By Discrete Analyser	(QCLot: 628160)							
EK061G: Total Kjeldahl Nitrogen as N		0.1	mg/L	<0.1	10 mg/L	87.9	82	110
EK067G: Total Phosphorus as P by Discrete Analyser	(QCLot: 628159)							
EK067G: Total Phosphorus as P		0.01	mg/L	<0.01	4.42 mg/L	91.4	70	130
EP030: Biochemical Oxygen Demand (BOD) (QCLot: 6	625068)							
EP030: Biochemical Oxygen Demand		2	mg/L	<2	198 mg/L	90.8	78	117

Matrix Spike (MS) Report

The quality control term Matrix Spike (MS) refers to an intralaboratory split sample spiked with a representative set of target analytes. The purpose of this QC parameter is to monitor potential matrix effects on analyte recoveries. Static Recovery Limits as per laboratory Data Quality Objectives (DQOs). Ideal recovery ranges stated may be waived in the event of sample matrix interference.

Sub-Matrix: WATER					trix Spike (MS) Report	t	
				Spike	SpikeRecovery(%)	Recovery Li	nits (%)
Laboratory sample ID	Client sample ID	Method: Compound	CAS Number	Concentration	MS	Low	High
EK055G: Ammonia	as N by Discrete Analyser (QCLot: 623447)						
EP1609847-002	QC01_20161018	EK055G: Ammonia as N	7664-41-7	1 mg/L	85.1	70	130
EK057G: Nitrite as	N by Discrete Analyser (QCLot: 623435)						
EP1609847-001	MW1	EK057G: Nitrite as N	14797-65-0	0.5 mg/L	88.0	70	130
EK059G: Nitrite plu	us Nitrate as N (NOx) by Discrete Analyser (QCLot: 623	448)					
EP1609847-002	QC01_20161018	EK059G: Nitrite + Nitrate as N		0.5 mg/L	105	70	130
EK061G: Total Kjelo	dahl Nitrogen By Discrete Analyser (QCLot: 628160)						
EP1608883-005	Anonymous	EK061G: Total Kjeldahl Nitrogen as N		5 mg/L	96.2	70	130
EK067G: Total Pho	sphorus as P by Discrete Analyser (QCLot: 628159)						
EP1608883-005	Anonymous	EK067G: Total Phosphorus as P		1 mg/L	85.2	70	130





	QA/QC Compliance Asse	ssment to assist with	h Quality Review
Work Order	: EP1609847	Page	: 1 of 6
Amendment	: 1		
Client		Laboratory	: Environmental Division Perth
Contact	: MR ANDREW NAGLE	Telephone	: 08 9209 7655
Project	: 61/34772 Point Moore Groundwater Assessment	Date Samples Received	: 19-Oct-2016
Site	:	Issue Date	: 02-Nov-2016
Sampler	: STEVEN PETTS	No. of samples received	: 7
Order number	:	No. of samples analysed	: 7

This report is automatically generated by the ALS LIMS through interpretation of the ALS Quality Control Report and several Quality Assurance parameters measured by ALS. This automated reporting highlights any non-conformances, facilitates faster and more accurate data validation and is designed to assist internal expert and external Auditor review. Many components of this report contribute to the overall DQO assessment and reporting for guideline compliance.

Brief method summaries and references are also provided to assist in traceability.

Summary of Outliers

Outliers : Quality Control Samples

This report highlights outliers flagged in the Quality Control (QC) Report.

- <u>NO</u> Method Blank value outliers occur.
- <u>NO</u> Duplicate outliers occur.
- <u>NO</u> Laboratory Control outliers occur.
- <u>NO</u> Matrix Spike outliers occur.
- For all regular sample matrices, NO surrogate recovery outliers occur.

Outliers : Analysis Holding Time Compliance

• <u>NO</u> Analysis Holding Time Outliers exist.

Outliers : Frequency of Quality Control Samples

• NO Quality Control Sample Frequency Outliers exist.



Analysis Holding Time Compliance

If samples are identified below as having been analysed or extracted outside of recommended holding times, this should be taken into consideration when interpreting results.

This report summarizes extraction / preparation and analysis times and compares each with ALS recommended holding times (referencing USEPA SW 846, APHA, AS and NEPM) based on the sample container provided. Dates reported represent first date of extraction or analysis and preclude subsequent dilutions and reruns. A listing of breaches (if any) is provided herein.

Holding time for leachate methods (e.g. TCLP) vary according to the analytes reported. Assessment compares the leach date with the shortest analyte holding time for the equivalent soil method. These are: organics 14 days, mercury 28 days & other metals 180 days. A recorded breach does not guarantee a breach for all non-volatile parameters.

Holding times for <u>VOC in soils</u> vary according to analytes of interest. Vinyl Chloride and Styrene holding time is 7 days; others 14 days. A recorded breach does not guarantee a breach for all VOC analytes and should be verified in case the reported breach is a false positive or Vinyl Chloride and Styrene are not key analytes of interest/concern.

Matrix: WATER					Evaluation	i: × = Holding time	breach ; 🗸 = With	in holding time
Method		Sample Date	Ex	traction / Preparation			Analysis	
Container / Client Sample ID(s)			Date extracted	Due for extraction	Evaluation	Date analysed	Due for analysis	Evaluation
EA015: Total Dissolved Solids dried at 180 ± 5 °C								
Clear Plastic Bottle - Natural (EA015H) MW1, MW2, SHP8, QC04_20161018	QC01_20161018, MW3, QC03_20161018,	18-Oct-2016				24-Oct-2016	25-Oct-2016	~
EK055G: Ammonia as N by Discrete Analyser								
Clear Plastic Bottle - Sulfuric Acid (EK055G) MW1, MW2, SHP8, QC04_20161018	QC01_20161018, MW3, QC03_20161018,	18-Oct-2016				19-Oct-2016	15-Nov-2016	~
EK057G: Nitrite as N by Discrete Analyser								
Clear Plastic Bottle - Natural (EK057G) MW1, MW2, SHP8, QC04_20161018	QC01_20161018, MW3, QC03_20161018,	18-Oct-2016				19-Oct-2016	20-Oct-2016	~
EK059G: Nitrite plus Nitrate as N (NOx) by Discre	ete Analyser							
Clear Plastic Bottle - Sulfuric Acid (EK059G) MW1, MW2, SHP8, QC04_20161018	QC01_20161018, MW3, QC03_20161018,	18-Oct-2016				19-Oct-2016	15-Nov-2016	~
EK061G: Total Kjeldahl Nitrogen By Discrete Anal	lyser							
Clear Plastic Bottle - Sulfuric Acid (EK061G) MW1, MW2, SHP8, QC04_20161018	QC01_20161018, MW3, QC03_20161018,	18-Oct-2016	26-Oct-2016	15-Nov-2016	~	26-Oct-2016	15-Nov-2016	~



Matrix: WATER					Evaluation	i: × = Holding time	breach ; 🗸 = Withi	n holding time
Method		Sample Date	Ex	traction / Preparation			Analysis	
Container / Client Sample ID(s)			Date extracted	Due for extraction	Evaluation	Date analysed	Due for analysis	Evaluation
EK067G: Total Phosphorus as P by Discrete Analyser								
Clear Plastic Bottle - Sulfuric Acid (EK067G)								
MW1,	QC01_20161018,	18-Oct-2016	26-Oct-2016	15-Nov-2016	1	26-Oct-2016	15-Nov-2016	✓
MW2,	MW3,							
SHP8,	QC03_20161018,							
QC04_20161018								
EP030: Biochemical Oxygen Demand (BOD)								
Clear Plastic Bottle - Natural (EP030)								
MW1,	QC01_20161018,	18-Oct-2016				20-Oct-2016	20-Oct-2016	✓
MW2,	MW3,							
SHP8,	QC03_20161018,							
QC04_20161018								
MW002: Heterotrophic Plate Count								
Sterile Plastic Bottle - Sodium Thiosulfate (MW002)								
MW1,	QC01_20161018,	18-Oct-2016				19-Oct-2016	19-Oct-2016	✓
MW2,	MW3,							
SHP8								
MW006: Faecal Coliforms & E.coli by MF								
Sterile Plastic Bottle - Sodium Thiosulfate (MW006)								
MW1,	QC01_20161018,	18-Oct-2016				19-Oct-2016	19-Oct-2016	✓
MW2,	MW3,							
SHP8								
MW007: Coliforms by MF								
Sterile Plastic Bottle - Sodium Thiosulfate (MW007)								
MW1,	QC01_20161018,	18-Oct-2016				19-Oct-2016	19-Oct-2016	✓
MW2,	MW3,							
SHP8								



Quality Control Parameter Frequency Compliance

The following report summarises the frequency of laboratory QC samples analysed within the analytical lot(s) in which the submitted sample(s) was(were) processed. Actual rate should be greater than or equal to the expected rate. A listing of breaches is provided in the Summary of Outliers.

Matrix: WATER		Evaluation: 🞽 = Quality Control frequency not within specification ; 🗹 = Quality Control frequency within specificati						
Quality Control Sample Type			ount		Rate (%)		Quality Control Specification	
Analytical Methods	Method	00	Reaular	Actual	Expected	Evaluation		
Laboratory Duplicates (DUP)								
Ammonia as N by Discrete analyser	EK055G	2	12	16.67	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Biochemical Oxygen Demand (BOD)	EP030	2	17	11.76	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	2	13	15.38	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite as N by Discrete Analyser	EK057G	2	20	10.00	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Dissolved Solids (High Level)	EA015H	2	18	11.11	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	2	20	10.00	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Phosphorus as P By Discrete Analyser	EK067G	2	20	10.00	10.00	1	NEPM 2013 B3 & ALS QC Standard	
Laboratory Control Samples (LCS)								
Ammonia as N by Discrete analyser	EK055G	1	12	8.33	5.00	1	NEPM 2013 B3 & ALS QC Standard	
Biochemical Oxygen Demand (BOD)	EP030	1	17	5.88	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	1	13	7.69	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite as N by Discrete Analyser	EK057G	1	20	5.00	5.00	1	NEPM 2013 B3 & ALS QC Standard	
Total Dissolved Solids (High Level)	EA015H	2	18	11.11	10.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Phosphorus as P By Discrete Analyser	EK067G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Method Blanks (MB)								
Ammonia as N by Discrete analyser	EK055G	1	12	8.33	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Biochemical Oxygen Demand (BOD)	EP030	1	17	5.88	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	1	13	7.69	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite as N by Discrete Analyser	EK057G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Dissolved Solids (High Level)	EA015H	1	18	5.56	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Phosphorus as P By Discrete Analyser	EK067G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Matrix Spikes (MS)								
Ammonia as N by Discrete analyser	EK055G	1	12	8.33	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	1	13	7.69	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Nitrite as N by Discrete Analyser	EK057G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	
Total Phosphorus as P By Discrete Analyser	EK067G	1	20	5.00	5.00	✓	NEPM 2013 B3 & ALS QC Standard	



Brief Method Summaries

The analytical procedures used by the Environmental Division have been developed from established internationally recognized procedures such as those published by the US EPA, APHA, AS and NEPM. In house developed procedures are employed in the absence of documented standards or by client request. The following report provides brief descriptions of the analytical procedures employed for results reported in the Certificate of Analysis. Sources from which ALS methods have been developed are provided within the Method Descriptions.

Analytical Methods	Method	Matrix	Method Descriptions
Total Dissolved Solids (High Level)	EA015H	WATER	In house: Referenced to APHA 2540C. A gravimetric procedure that determines the amount of `filterable` residue in an aqueous sample. A well-mixed sample is filtered through a glass fibre filter (1.2um). The filtrate is evaporated to dryness and dried to constant weight at 180+/-5C. This method is compliant with NEPM (2013) Schedule B(3)
Ammonia as N by Discrete analyser	EK055G	WATER	In house: Referenced to APHA 4500-NH3 G Ammonia is determined by direct colorimetry by Discrete Analyser. This method is compliant with NEPM (2013) Schedule B(3)
Nitrite as N by Discrete Analyser	EK057G	WATER	In house: Referenced to APHA 4500-NO2- B. Nitrite is determined by direct colourimetry by Discrete Analyser. This method is compliant with NEPM (2013) Schedule B(3)
Nitrate as N by Discrete Analyser	EK058G	WATER	In house: Referenced to APHA 4500-NO3- F. Nitrate is reduced to nitrite by way of a chemical reduction followed by quantification by Discrete Analyser. Nitrite is determined seperately by direct colourimetry and result for Nitrate calculated as the difference between the two results. This method is compliant with NEPM (2013) Schedule B(3)
Nitrite and Nitrate as N (NOx) by Discrete Analyser	EK059G	WATER	In house: Referenced to APHA 4500-NO3- F. Combined oxidised Nitrogen (NO2+NO3) is determined by Chemical Reduction and direct colourimetry by Discrete Analyser. This method is compliant with NEPM (2013) Schedule B(3)
Total Kjeldahl Nitrogen as N By Discrete Analyser	EK061G	WATER	In house: Referenced to APHA 4500-Norg D (In house). An aliquot of sample is digested using a high temperature Kjeldahl digestion to convert nitrogenous compounds to ammonia. Ammonia is determined colorimetrically by discrete analyser. This method is compliant with NEPM (2013) Schedule B(3)
Total Nitrogen as N (TKN + Nox) By Discrete Analyser	EK062G	WATER	In house: Referenced to APHA 4500-Norg / 4500-NO3 This method is compliant with NEPM (2013) Schedule B(3)
Total Phosphorus as P By Discrete Analyser	EK067G	WATER	In house: Referenced to APHA 4500-P H, Jirka et al (1976), Zhang et al (2006). This procedure involves sulphuric acid digestion of a sample aliquot to break phosphorus down to orthophosphate. The orthophosphate reacts with ammonium molybdate and antimony potassium tartrate to form a complex which is then reduced and its concentration measured at 880nm using discrete analyser. This method is compliant with NEPM (2013) Schedule B(3)
Biochemical Oxygen Demand (BOD)	EP030	WATER	In house: Referenced to APHA 5210 B. The 5-Day BOD test provides an empirical measure of the oxygen consumption capacity of a given water. A portion of the sample is diluted into oxygenated, nutrient rich water, and a seed added to begin biological decay. The initial dissolved oxygen content is measured, then the bottle is sealed and incubated for five days. The remaining dissolved oxygen is measured, and from the difference, the demand for oxygen, by biological decay, is determined. This method is compliant with NEPM (2013) Schedule B(3)
Heterotrophic (Total) Plate Count @ 22C and 36C	MW002	WATER	In house: Referenced to AS4276.3.1- 2007
Thermotolerant Coliforms & E.coli by Membrane Filtration	MW006	WATER	In house: Referenced to AS 4276.7 2007
Coliforms by Membrane Filtration	MW007	WATER	In house: Referenced to AS 4276.5 - 2007
Preparation Methods	Method	Matrix	Method Descriptions

Page	: 6 of 6
Work Order	: EP1609847 Amendment 1
Client	: GHD PTY LTD
Project	: 61/34772 Point Moore Groundwater Assessment



Preparation Methods	Method	Matrix	Method Descriptions
TKN/TP Digestion	EK061/EK067	WATER	In house: Referenced to APHA 4500 Norg - D; APHA 4500 P - H. This method is compliant with NEPM (2013)
			Schedule B(3)

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Date/Time:	18/10/20
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A10 0 0	Date/Time: Date/Time:





16 - 18 Hayden Court, Myaree, Western Australia 6154 PO Box 4023 Myaree BC, Western Australia 6960 Tel: +61 8 9317 2505 / Fax: +61 8 9317 4163 email: laboratory@mpl.com.au www.mpl.com.au Envirolab Services (WA) Pty Ltd ABN 63 140 099 207

SAMPLE RECEIPT ADVICE

Client Details	
Client	GHD Services Pty Ltd
Attention	Andrew Nagle

Sample Login Details	
Your Reference	61/34772
Envirolab Reference	187366
Date Sample Received	19/10/2016
Date Instructions Received	19/10/2016
Date Results Expected to be Reported	26/10/2016

Sample Condition			
Samples received in appropriate condition for analysis	YES		
No. of Samples Provided	1 Water		
Turnaround Time Requested	Standard		
Temperature on receipt (°C)	15		
Cooling Method	Ice Pack		
Sampling Date Provided	Yes		

Comments

Samples will be held for 1 month for water samples and 2 months for soil samples from date of receipt of samples

Please direct any queries to:

Joshua Lim	Meredith Conroy
Phone: 08 9317 2505	Phone: 08 9317 2505
Fax: 08 9317 4163	Fax: 08 9317 4163
Email: jlim@mpl.com.au	Email: mconroy@mpl.com.au

Sample and Testing Details on following page
16 - 18 Hayden Court, Myaree, Western Australia 6154 PO Box 4023 Myaree BC, Western Australia 6960 Tel: +61 8 9317 2505 / Fax: +61 8 9317 4163 email: laboratory@mpl.com.au www.mpl.com.au Envirolab Services (WA) Pty Ltd ABN 53 140 099 207





Sample Id	BOD	Total Dissolved Solids (grav)	Nutrients in Water	Microbiological Testing
QC02_20160922	1	1	1	1



email: lab@mpl.com.au envirolab.com.au

Envirolab Services (WA) Pty Ltd trading as MPL Laboratories | ABN 53 140 099 207

CERTIFICATE OF ANALYSIS 187366

Client: GHD Services Pty Ltd PO Box Y3106 Perth WA 6000

Attention: Andrew Nagle

Sample log in details:

Your Reference:	61/34772
No. of samples:	1 Water
Date samples received:	19/10/2016
Date completed instructions received:	19/10/2016
Location:	

Analysis Details:

Please refer to the following pages for results, methodology summary and quality control data. Samples were analysed as received from the client. Results relate specifically to the samples as received. Results are reported on a dry weight basis for solids and on an as received basis for other matrices. *Please refer to the last pages of this report for any comments relating to the results.*

Report Details:

 Date results requested by:
 26/10/16

 Date of Preliminary Report:
 Not issued

 Issue Date:
 26/10/16

 NATA accreditation number 2901. This document shall not be reproduced except in full.

 Accredited for compliance with ISO/IEC 17025 - Testing

 Tests not covered by NATA are denoted with *.

Results Approved By:

Joshua Lim Operations Manager

MPL Reference: Revision No: 187366 R 00

Jessica Miller Micro Team Leader



Page 1 of 8

Miscellaneous Inorganics		
Our Reference:	UNITS	187366-1
Your Reference		QC02_20160
		922
Date Sampled		18/10/2016
Type of sample		Water
Date prepared	-	19/10/2016
Date analysed	-	19/10/2016
BOD	mg/L	14
Total Dissolved Solids (grav)	mg/L	14,000

Nutrients in Water		
Our Reference:	UNITS	187366-1
Your Reference		QC02_20160
		922
Date Sampled		18/10/2016
Type of sample		Water
Date prepared	-	19/10/2016
Date analysed	-	19/10/2016
Total Nitrogen	mg/L	5.7
TKN by Discrete Analyser	mg/L	5.3
NOx as N	mg/L	0.44
Nitrite as N	mg/L	<0.005
Nitrate as N	mg/L	0.44
Ammonia as N	mg/L	3.2
Total Phosphorus	mg/L	0.57

Microbiological Testing		
Our Reference:	UNITS	187366-1
Your Reference		QC02_20160
		922
Date Sampled		18/10/2016
Type of sample		Water
Date testing started	-	19/10/2016
Date testing completed	-	24/10/2016
Heterotrophic Plate Count 21C	cfu/mL	4,200^
Heterotrophic Plate Count 35C	cfu/mL	3,900^
Total Coliforms	cfu/100mL	3**
Thermotolerant Coliforms	cfu/100mL	<1**
E.coli	cfu/100mL	<1**

MethodID	Methodology Summary
INORG-091	BOD - Analysed in accordance with APHA latest edition 5210 D.
INORG-018	Total Dissolved Solids - determined gravimetrically. The solids are dried at $180\pm5^{\circ}$ C
INORG-055	Total Nitrogen by colourimetric analysis based on APHA 4500-P J, 4500-NO3 F.
INORG-062	TKN by calculation fromTotal Nitrogen and NOx using APHA methodology.
INORG-055	NOx - determined colourimetrically. Soils are analysed from a water extract.
INORG-055	Nitrite - determined colourimetrically. Soils are analysed from a water extract.
INORG-055	Nitrate - determined colourimetrically. Soils are analysed from a water extract.
INORG-057	Ammonia by colourimetric analysis based on APHA latest edition 4500-NH3 F.
METALS-020	Metals in soil and water by ICP-OES.
MICRO-001	Heterotrophic Plate Count: Microbial Water Analysis - in accordance with MICRO-001 (APHA-9215D-2005). Recommended maximums based on NHMRC and ARMC Australian Drinking Water Guidelines.
MICRO-001	Total Coliforms: Microbial Water Analysis - in accordance with MICRO-001 (AS4276.5-2007). Recommended maximums based on NHMRC and ARMC Australian Drinking Water Guidelines.
MICRO-001	Thermotolerant Coliforms: Microbial Water Analysis - in accordance with MICRO-001 (AS4276.7-2007). Recommended maximums based on NHMRC and ARMC Australian Drinking Water Guidelines.
MICRO-001	E. Coli: Microbial Water Analysis - in accordance with MICRO-001 (AS4276.7-2007). Recommended maximums based on NHMRC and ARMC Australian Drinking Water Guidelines.

	Client Reference: 61/34772							
QUALITYCONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Miscellaneous Inorganics						Base II Duplicate II % RPD		
Date prepared	-			19/10/ 2016	[NT]	[NT]	LCS-1	19/10/2016
Date analysed	-			19/10/ 2016	[NT]	[NT]	LCS-1	19/10/2016
BOD	mg/L	5	INORG-091	⊲5	[NT]	[NT]	LCS-1	71%
Total Dissolved Solids (grav)	mg/L	5	INORG-018	45	[NT]	[NT]	LCS-1	96%
QUALITYCONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Nutrients in Water						Base II Duplicate II %RPD		
Date prepared	-			19/10/ 2016	[NT]	[NT]	LCS-1	19/10/2016
Date analysed	-			20/10/ 2016	[NT]	[NT]	LCS-1	19/10/2016
Total Nitrogen	mg/L	0.1	INORG-055	<0.1	[NT]	[NT]	LCS-1	103%
TKN by Discrete Analyser	mg/L	0.1	INORG-062	<0.1	[NT]	[NT]	[NR]	[NR]
NOx as N	mg/L	0.005	INORG-055	<0.005	[NT]	[NT]	LCS-1	110%
Nitrite as N	mg/L	0.005	INORG-055	<0.005	[NT]	[NT]	LCS-1	107%
Nitrate as N	mg/L	0.005	INORG-055	<0.005	[NT]	[NT]	LCS-1	110%
Ammonia as N	mg/L	0.005	INORG-057	<0.005	[NT]	[NT]	LCS-1	108%
Total Phosphorus	mg/L	0.05	METALS- 020	<0.05	[NT]	[NT]	LCS-1	113%
QUALITYCONTROL	UNITS	PQL	METHOD	Blank				
Microbiological Testing								
Date testing started	-			19/10/ 2016				
Date testing completed	-			24/10/ 2016				
Heterotrophic Plate Count 21C	cfu/mL	10	MICRO-001	<10				
Heterotrophic Plate Count 35C	cfu/mL	10	MICRO-001	<10				
Total Coliforms	cfu/100 mL	1	MICRO-001	<1				
Thermotolerant Coliforms	cfu/100 mL	1	MICRO-001	<1				
E.coli	cfu/100 mL	1	MICRO-001	<1				

Report Comments:

^ - Heterotrophic plate count is an estimate.

- ** High background growth of non-coliform bacteria may underestimate Total Coliform Count.
- ** High background growth of non-coliform bacteria may underestimate the Thermotolerant Coliform and E.coli Count.

Definitions:

NT: Not tested NA: Test not required INS: Insufficient sample for this test PQL: Practical Quantitation Limit <: Less than >: Greater than RPD: Relative Percent Difference LCS: Laboratory Control Sample NS: Not Specified NEPM: National Environmental Protection Measure NR: Not Reported

Australian Drinking Water Guidelines recommend that Thermotolerant Coliform, Faecal Enterococci, & E.Coli levels are less than 1cfu/100mL. The recommended maximums are taken from "Australian Drinking Water Guidelines", published by NHMRC & ARMC 2011

Quality Control Definitions

Blank: This is the component of the analytical signal which is not derived from the sample but from reagents, glassware etc, can be determined by processing solvents and reagents in exactly the same manner as for samples. **Duplicate**: This is the complete duplicate analysis of a sample from the process batch. If possible, the sample selected should be one where the analyte concentration is easily measurable.

Matrix Spike : A portion of the sample is spiked with a known concentration of target analyte. The purpose of the matrix spike is to monitor the performance of the analytical method used and to determine whether matrix interferences exist.

LCS (Laboratory Control Sample) : This comprises either a standard reference material or a control matrix (such as a blank sand or water) fortified with analytes representative of the analyte class. It is simply a check sample.

Surrogate Spike: Surrogates are known additions to each sample, blank, matrix spike and LCS in a batch, of compounds which are similar to the analyte of interest, however are not expected to be found in real samples.

Laboratory Acceptance Criteria

Duplicate sample and matrix spike recoveries may not be reported on smaller jobs, however, were analysed at a frequency to meet or exceed NEPM requirements. All samples are tested in batches of 20. The duplicate sample RPD and matrix spike recoveries for the batch were within the laboratory acceptance criteria.

Filters, swabs, wipes, tubes and badges will not have duplicate data as the whole sample is generally extracted during sample extraction.

Spikes for Physical and Aggregate Tests are not applicable.

For VOCs in water samples, three vials are required for duplicate or spike analysis.

Duplicates: <5xPQL - any RPD is acceptable; >5xPQL - 0-50% RPD is acceptable. Matrix Spikes, LCS and Surrogate recoveries: Generally 70-130% for inorganics/metals; 60-140% for organics (+/-50% surrogates) and 10-140% for labile SVOCs (including labile surrogates), ultra trace organics and speciated phenols is acceptable.

In circumstances where no duplicate and/or sample spike has been reported at 1 in 10 and/or 1 in 20 samples respectively, the sample volume submitted was insufficient in order to satisfy laboratory QA/QC protocols.

When samples are received where certain analytes are outside of recommended technical holding times (THTs), the analysis has proceeded. Where analytes are on the verge of breaching THTs, every effort will be made to analyse within the THT or as soon as practicable.

Where sampling dates are not provided, Envirolab are not in a position to comment on the validity of the analysis where recommended technical holding times may have been breached.

Measurement Uncertainty estimates are available for most tests upon request.

Appendix G – Tabulated Groundwater Analytical Results



Appendix G, Table G1 Groundwater Analytical Results

Water Quality Guidelines					6134772 MW1 24/08/2016	6134772 MW1 23/09/2016	6134772 MW1 18/10/2016	6134772 MW2 24/08/2016	6134772 MW2 23/09/2016	6134772 MW2 18/10/2016	6134772 MW3 24/08/2016	6134772 MW3 23/09/2016	
Parameter	Unit	MW 95%	Marine Inshore	ANZECC 2000 Primary Contact Recreation	ANZECC 2000 Secondary Contact Recreation								
Ammonia as N	mg/L	0.91	-	-	-	2.68	6.28	3.95	0.53	0.13	1.48	0.57	0.78
BOD	mg/L	-	-	-	-	6	31	33	7	4	3	4	5
Coliform	cfu/100 ml	-	-	-	-	250,000	<1	1	1000	10	1	3000	10
Plate Count (36°C)	CFU/mL	-	-	-	-	46,000	170	4400	11,000	12,000	7400	68,000	2100
Plate Count (22°C)	CFU/mL	-	-	-	-	49,000	72	4200	13,000	1700	4500	70,000	2400
Nitrate (as N)	mg/L	-	0.05	-	-	0.19	0.05	0.46	13.8	9.95	6.06	0.02	0.01
Nitrite + Nitrate as N	mg/L	-	-	-	-	0.19	0.05	0.46	14	11.8	6.33	0.02	0.01
Nitrite (as N)	mg/L	-	-	-	-	< 0.01	< 0.01	< 0.01	0.2	1.85	0.27	< 0.01	<0.01
Faecal Coliform	CFU/100mL	-	-	150	1000	200	<1	1	400	1	1	400	1
E. Coli	CFU/100 ml	-	-	150	1000	200	<1	1	400	1	1	400	1
Total Dissolved Solids	mg/L	-	-			8890	-	-	1480	-	-	1090	-
Total Dissolved Solids (Filtered)	mg/L	-	-	-	-	-	22,300	13,100	-	2120	1360	-	1320
Total Kjeldahl Nitrogen (as N)	mg/L	-	-	-	-	3	6.7	4.2	3.1	2	2.6	0.7	0.9
Nitrogen (Total)	mg/L	-	0.23	-	-	3.2	6.8	4.7	17.1	13.8	8.9	0.7	0.9
Phosphorus (Total)	mg/L	-	0.005	-	-	0.29	0.81	0.47	0.21	0.14	0.18	0.06	0.08

Legend

ANZECC 2000 MW 95%	Indicates a level is equal to or above the ANZECC (2000) Marine Waters 95% Species Protection Trigger Value
ANZECC 2000 Marine Inshore	Indicates a level is equal to or above the ANZECC (2000) Marine Inshore Trigger Value
ANZECC 2000 Pimary Contact Recreation	Indicates a level is equal to or above the ANZECC (2000) Water Quality Guideline for Primary Contact Recreation (e.g. swimming)
ANZECC 2000 Secondary Contact Recreation	Indicates a level is equal to or above the ANZECC (2000) Water Quality Guideline for Secondary Contact Recreation (e.g. boating, fishing)



Appendix G, Table G1 Groundwater Analytical Results

						6134772	6134772	6134772	6134772					
						MW3	SHP8	SHP8	SHP8					
	Water Quality Guidelines				18/10/2016	24/08/2016	23/09/2016	18/10/2016		Sta	tistical Summa	ry		
		ANZECC 2000	ANZECC 2000	ANZECC 2000	ANZECC 2000					Number of	Minimum	Maximum	Mean	Median
Parameter	Unit	MW 95%	Marine Inshore	Primary Contact	Secondary Contact					Results				
				Recreation	Recreation									
Ammonia as N	mg/L	0.91	-	-	-	0.32	0.36	2.54	0.46	12	0.13	6.28	1.7	0.675
BOD	mg/L	-	-	-	-	3	4	22	57	12	3	57	15	5.5
Coliform	cfu/100 ml	-	-	-	-	1	200	<1	1	12	<1	250,000	21,185	5.5
Plate Count (36°C)	CFU/mL	-	-	-	-	5500	110	17	980	12	17	68,000	13,140	4,950
Plate Count (22°C)	CFU/mL	-	-	-	-	14,000	910	12	2000	12	12	70,000	13,483	3,300
Nitrate (as N)	mg/L	-	0.05	-	-	< 0.01	0.02	0.03	< 0.01	12	<0.01	13.8	2.6	0.04
Nitrite + Nitrate as N	mg/L	-	-	-	-	< 0.01	0.02	0.03	< 0.01	12	<0.01	14	2.7	0.04
Nitrite (as N)	mg/L	-	-	-	-	< 0.01	< 0.01	< 0.01	< 0.01	12	<0.01	1.85	0.2	0.005
Faecal Coliform	CFU/100mL	-	-	150	1000	1	100	<1	1	12	<1	400	92	1
E. Coli	CFU/100 ml	-	-	150	1000	1	100	<1	1	12	<1	400	92	1
Total Dissolved Solids	mg/L	-	-			-	3320	-	-	4	1,090	8,890	3,695	2,400
Total Dissolved Solids (Filtered)	mg/L	-	-	-	-	1340	-	10,300	2290	8	1,320	22,300	6,766	2,205
Total Kjeldahl Nitrogen (as N)	mg/L	-	-	-	-	0.6	1.5	3.3	1.2	12	0.6	6.7	2.5	2.3
Nitrogen (Total)	mg/L	-	0.23	-	-	0.6	1.5	3.3	1.2	12	0.6	17.1	5.2	3.25
Phosphorus (Total)	mg/L	-	0.005	-	-	0.08	0.31	0.38	0.28	12	0.06	0.81	0.27	0.245

Legend

ANZECC 2000 MW 95%	Indicates a level is equal to or above the ANZECC (2000) Marine Waters 95% Species Protection Trigger Value
ANZECC 2000 Marine Inshore	Indicates a level is equal to or above the ANZECC (2000) Marine Inshore Trigger Value
ANZECC 2000 Pimary Contact Recreation	Indicates a level is equal to or above the ANZECC (2000) Water Quality Guideline for Primary Contact Recreation (e.g. swimming)
ANZECC 2000 Secondary Contact Recreation	Indicates a level is equal to or above the ANZECC (2000) Water Quality Guideline for Secondary Contact Recreation (e.g. boating, fishing)

City of Greater Geraldton Point Moore Groundwater Assessment

Appendix G Table G2 Field and Rinsate Blanks

		Rinsate	Field Blank	Rinsate
		QC03_20160824	QC04_20160824	QC03_20160922
ChemName	output	24/08/2016	24/08/2016	23/09/2016
	unit			
Ammonia as N	mg/L	<0.01	<0.01	<0.01
BOD	mg/L	5	<2	<2
Coliform	cfu/100 ml	-	-	-
Plate Count (36°C)	cfu/100 ml	-	-	-
PLATE COUNT 22C	cfu/100 ml	-	-	-
Nitrate (as N)	mg/L	<0.01	<0.01	<0.01
Nitrite + Nitrate as N	mg/L	< 0.01	<0.01	<0.01
Nitrite (as N)	mg/L	<0.01	<0.01	<0.01
Faecal Coliform	cfu/100 ml	-	-	-
E. Coli	cfu/100 ml	-	-	-
Total Dissolved Solids	mg/L	<10	<10	-
Total Dissolved Solids (Filtered)	mg/L	-	-	20
Total Kjeldahl Nitrogen	mg/L	<0.1	<0.1	<0.1
Nitrogen (Total)	mg/L	<0.1	<0.1	<0.1
Phosphorus	mg/L	< 0.01	< 0.01	< 0.01

Appendix G Table G2 Field and Rinsate Blanks

Field Blank	Rinsate	Field Blank
QC04_20160922	QC03_20160922	QC04_20160922
23/09/2016	18/10/2016	18/10/2016
<0.01	0.06	0.06
<2	<2	<2
-	-	-
-	-	-
-	-	-
<0.01	< 0.01	0.01
<0.01	< 0.01	0.01
<0.01	< 0.01	< 0.01
-	-	-
-	-	-
-	-	-
<10	14	16
< 0.1	< 0.1	< 0.1
<0.1	< 0.1	< 0.1
< 0.01	< 0.01	< 0.01

Appendix G Table G3 Blind and Split RPDs

			1					-			-	-	-			-	
		Well	MW1	QC01_20160824	QC02_20160824	RPD	RPD	MW1	QC01_20160922	QC02_20160922	RPD	RPD	MW1	QC01_20161018	QC02_20161018	RPD	RPD
		Date	24/08/2016	24/08/2016	24/08/2016	MW1	MW1	23/09/2016	23/09/2016	23/09/2016	MW1 and	MW1 and	18/10/2016	18/10/2016	18/10/2016	MW1 and	MW1 and
						QC01	QC02				QC01	QC02				QC01	QC02
ChemName	output unit	EQL															
Ammonia as N	mg/L	0.01	2.68	2.86	2.9	6	1	6.28	6.42	5	2	-23	3.95	4	3.2	1	-21
BOD	mg/L	2	6	6	<5	0	-18	31	31	<5	0	-144	33	27	14	-20	-81
Coliform	cfu/100 ml	1	250,000	200	-	200	-	<1	<1	<1	-	-	1	1	<1	0	0
Plate Count (36°C)	CFU/mL	1	46,000	49,000	-	6	-	170	140	340	-19	67	4400	4400	4200	0	-5
PLATE COUNT 22C	CFU/mL	1	49,000	60,000	-	20	-	72	73	260	1	113	4200	3600	3900	-15	-7
Nitrate (as N)	mg/L	0.01	0.19	0.18	0.17	-5	-6	0.05	<0.01	<0.005	-133	-164	0.46	0.46	0.44	0	-4
Nitrite + Nitrate as N	mg/L	0.01	0.19	0.18	0.17	-5	-6	0.05	0.46	-	161	-	0.46	0.46	-	0	-
Nitrite (as N)	mg/L	0.01	<0.01	<0.01	<0.2	-	-	<0.01	< 0.01	0.006	-	-198	< 0.01	< 0.01	<0.005	-	-
Faecal Coliform	CFU/100mL	1	200	1100	-	138	-	<1	<1	<1	0	0	1	1	<1	0	0
E. Coli	cfu/100 ml	1	200	1100	-	138	-	<1	<1	<1	0	0	1	1	<1	0	0
Total Dissolved Solids	mg/L	10	8890	8800	7500	-1	-16	22,300	22,400	22000	0	-1	13,100	13,100	14000	0	7
Total Kjeldahl Nitrogen	mg/L	0.1	3	2.9	3.1	-3	7	6.7	6.7	9.2	0	31	4.2	4.1	5.3	-2	23
Nitrogen (Total)	mg/L	0.1	3.2	3.1	3.2	-3	3	6.8	6.7	9.2	-1	30	4.7	4.6	5.7	-2	19
Phosphorus	mg/L	0.01	0.29	0.3	0.19	3	-45	0.81	0.9	0.78	11	-4	0.47	0.46	0.57	-2	19

Indicates Exceedance of RPD criteria (50%)

City of Greater Geraldton Point Moore Groundwater Assessment

Appendix H – Calibration Certificates

RENTALS

**

Insitu Aqua Troll 600 Profiler Water Quality Meter

This Water Quality Meter has been performance checked / calibrated* as follows:							
рН	□ pH 6.88	₽ pH 7.00	₽ pH 4.00	□ pH 10.00	₽́pH		
Conductivity	🖻 0.0mS/cm	☑ 1413uS/cm	212.88mS/cm	□ 58.6mS/cm	⊯ mS/cm		
TDS	☑ 0.0 ppk	∃ 36 ppk	∠ ppk				
Dissolved Oxygen	2 0.00ppm in Sodiur	n Sulphite		-100% Saturat	ion in Air		
Redox (ORP)**	Electrode operabil	ity test 240mV +,	/- 10%. Actual:	240 mV			
Electrodes cleaned/checked		Temperature					

* Calibration solution traceability information is available upon request.

Date:	22/8	Checked by:	
Signed:	AD		

Please check that the following items are received and that all items are cleaned and decontaminated before return. A minimum \$20 cleaning / service / repair charge may be applied to any unclean or damaged items. Items not returned will be billed for at the full replacement cost.

Sent	Received	Returned	Item
			AquaTroll 600 Unit. Ops check / Battery Voltage @
2			AquaTroll 600 Set to Time: Yes / No
	· 🗌		Aqua Troll 600 Cal Cup
			Troll Cable S/n: Length: m
			USB Cable Connect Comms Cable S/n:
5-			S/steel Sensor Protector Shroud
			Screw Drivers x3
2			Turbidity Wiper
-			PC Software version X.X
F			Aqua Troll 600 Cary Case

Processors Signature/ Initials

Thermo Fisher

SCIENTIFIC

AP

Quote Reference	CP0818	Condition on return	
Customer Ref			
Equipment ID	AT600		
Equipment serial no.			
Return Date	1 1		
Return Time			



RENTALS

Equipment Certification Report - TPS 90FLMV Water Quality Meter

This Water Quality Meter has been performance checked and calibrated as follows:

Sensor	Concentration	Span	1	Span 2		Traceability Lot #	Pass?
pН	pH 7.00 / pH 4.00	7	pН	4	pН	1	
Conductivity	1413uS/cm	1413	mS/cm	12880 1	mS/cm		
TDS	36 ppk	\sim	ppk	36	ppk		
Dissolved Oxygen	Sodium Sulphite / Air	in Sodium	ppm Sulphite	ر <i>©</i> Saturatior	ppm n in Air		
Check only	<u>]</u>]					<u> </u>	1
Redox (ORP) *	Electrode operability test	240n +/- 10	nV D%	240	mV	and a second	
* This meter us mV reading.	ses an Ag/AgCl ORP ele	ctrode. To cor	nvert readings	s to SHE (Stand	dard Hydr	ogen Electrode), add 199	mV to the
Battery Star	tus afety Tag attached (A	_(min 7.2V) S/NZS 3760))	Tempe Electro	erature _ odes Cle	21°C aned and checked	
Tag N	10:						
Valid	to:						
Date: 20/1							
Signed: AD							

Please check that the following items are received and that all items are cleaned and decontaminated before return. A minimum \$30 cleaning / service / repair charge may be applied to any unclean or damaged items. Items not returned will be billed for at the full replacement cost.

Sent	Returned	Item
		90FLMV Unit. Ops check/Battery status: pH sensor with wetting cap, 5m Conductivity/TDS/Temperature K=10 sensor, 5m Dissolved oxygen YSI5739 sensor with wetting cap, 5m Redox (ORP) sensor with wetting cap, 5m Power supply 240V to 12V DC 200mA Instruction Manual Quick Guide Syringe with storage solution for pH and ORP sensors Carry Case Check to confirm electrical safety (tag must be valid)
ate: 00	19	

Date: AD Signed:

Signed:

TFS Reference	CP0818	Return Date: / /
Customer Reference		Return Time:
Equipment ID	90FLMV WA3	Condition on return:
Equipment Serial No.		

"We do more than give you great equipment... We give you great solutions!"

Phone: (Free Call) 13	800 735 295	Fax: (Free Call) 1800 675 123		Email: RentalsAU@Thermofisher.com		
Melbourne Branch	Sydney Branch	Adelaide Branch	Brisbane Branch	Perth Branch		
5 Caribbean Drive,	Level 1, 4 Talavera Road,	27 Beulah Road, Norwood,	Unit 2/5 Ross St	121 Beringarra Ave		
Scoresby 3179	North Ryde 2113	South Australia 5067	Newstead 4006	Malaga WA 6090		



RENTALS

Equipment Certification Report - TPS 90FLT Water Quality Meter

This Water Quality Meter has been performance checked and calibrated as follows:

Sensor	Concen	Concentration Span 1			Spar	12	Traceability Lot #	Pass?	
pН	pH 4.00/	oH 7.00	4	pН	7	pН	1	ď	
Conductivity	12.8	8mS/cm	12.88	,mS/cm	1413	//S/cm		۵,	
TDS		36.0ppk	P	ppk	36	ppk		Ó	
Dissolved Oxygen	Sodium Sulphite / Air		୍ଦ in Sodium S	ppm Sulphite	\∕⊘ Saturat	ppm ion in Air		Ø	
Turbidity	1000	NTU	0	NTU	(00	O NTU		Þ	
Redox (ORP)**240m	V	1	Electrode	e operabil	ty test 240n	nV +/- 10%	. Actual: 220 mV	2	
Battery Sta Electrical	atus Safety Tag	attached	(min 7.2' I (AS/NZS 37	V) 760)		nperature ctrodes C	e°C leaned and checked		
Tag N	No:								
Valid	to:								
Date: 14/	10								
Signed: Af	0								

Please check that the following items are received and that all items are cleaned and decontaminated before return. A minimum \$30 cleaning / service / repair charge may be applied to any unclean or damaged items. Items not returned will be billed for at the full replacement cost.

Sent/	Returned	Item
		90FLMV Unit. Ops check/Battery status:
Δ		pH sensor with wetting cap, 5m
Ď/		Conductivity/TDS/Temperature K=10 sensor, 5m
Ď/		Dissolved oxygen YSI5739 sensor with wetting cap, 5m
2/		Turbidity sensor, 5m
		Power supply 240V to 12V DC 200mA
-		Instruction Manual
		Quick Guide
		Syringe with storage solution for pH and ORP sensors
		Carry Case
2		Check to confirm electrical safety (tag must be valid)

Date: 4/10

Signed: A

TFS Reference	C80818	Return Date: / /
Customer Reference		Return Time:
Equipment ID	90FLT - WAS	Condition on return:
Equipment Serial No.		

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Phone: (Free Call) 1300 735 295		Fax: (Free Call) 1800 675 123	Email: Re	Email: RentalsAU@Thermofisher.com	
Melbourne Branch 5 Caribbean Drive, Scoresby 3179	Sydney Branch Level 1, 4 Talavera Road, North Ryde 2113	Adelaide Branch 27 Beulah Road, Norwood, South Australia 5067	Brisbane Branch Unit 2/5 Ross St Newstead 4006	Perth Branch 121 Beringarra Ave Malaga WA 6090	
Issue 8		Aug 15		G0564	

Appendix I – Field Works Photographs





	MWI-45 MWI-50 MWI-0.5 MWI-0.5 MWI-10 MWI-1.5 MWI-2.0 MWI-2.5 MWI-2.5 MWI-2.5 MWI-2.5 MWI-30 MWI-30 MWI-30 MWI-2.5	
Created by: S Petts	Date	Point Moore Groundwater
Source: GHD (15/08/2016)	16/09/2016	Assessment
Reference: IMG_2221		
File:G:\61\34772\Technical\Photos\	2016	GHD
Description: MW1 – Soil Log 2.5 mbgl		
MWI - 4.5 prw4-50 MWI - 0.5 MWI - 0.5 MWI - 1.0 MWI - 1.5 MWI - 1.20 MWI - 2.20 MWI - 2.5 MWI - 2.5 MWI - 2.5 MWI - 2.5		
Created by: S Petts	Date	Point Moore
Source: GHD (15/08/2016)	16/09/2016	Assessment
Reference: IMG_2222		
File:G:\61\34772\Technical\Photos\	2016	GHD
Description: MW1 – Soil Log 3.0 mbgl		









	MW2- 0.5 NTW 2 - 1.5 MW2 - 2.6 MW2 - 2.6 MW2- 3.5 MW2- 35 MW2- 35 MW2- 35 MW2- 35 MW2- 50 MW2- 50 MW2- 50 MW2- 50	
Created by: S Petts	Date	Point Moore Groundwater
D (10,0002010)	10/03/2010	Assessment
Reference: IMG_2232	2016	CHID
Description: MW2 – Soil Log 2.0 mbg	2010	GHD
	MW2- 05 ptiw 2 - 1 MW2 - 1 MW2 - 2 MW2 - 2 MW2 - 35 MW2- 35 MW2- 35 MW2- 35 MW2- 5 MW2- 5 M	
Created by: S Petts	Date	Point Moore
Source: GHD (15/08/2016)	16/09/2016	Assessment
Reference: IMG_2233	2016	GHD
File:G:\61\34772\Technical\Photos\		
Description: MW1 – Soil Log 2.5 mbgl		




















GHD

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