



13 May 2021

Our ref: FOI 2021/25

Mr Graeme Sawyer
Protect Country Alliance NT
Via Email:

Dear Mr Sawyer

FREEDOM OF INFORMATION REQUEST

I refer to your email of 12 May 2021, which contained a request, pursuant to the *Freedom of Information Act 1982* (Cth), for access to:

“Correspondence any time during September 2018 to end May 2021 between representatives of GISERA/CSIRO and with representatives of APPEA, Santos and/or Origin Energy containing the terms:

Sulphide reducing bacteria, Sulphate reducing bacteria, Microbially induced concrete corrosion (MICC), sulfur-oxidizing bacteria (SOB), microbial corrosion. Or documents relating to corrosion in petroleum and gas well casings.”

Acknowledgement

CSIRO received your request on **12 May 2021** and the 30 day statutory period for processing your request commenced from the day after that date. You should therefore expect a decision from CSIRO by **11 June 2021**. The period of 30 days may be extended if we need to consult third parties pursuant to sections 27 and 27A of the FOI Act, in relation to their personal information and/or their business affairs contained in the documents you are seeking. CSIRO will advise you if this happens.

Charges

Under the FOI Act charges may be imposed for processing your request pursuant to section 29. These charges are calculated by reference to the time spent in searching and retrieving relevant documents, decision making time, photocopying, postage etc. In the event CSIRO decides that you are liable to pay a charge in respect of processing your request, you will be notified of the preliminary assessment of the charges and you will have the opportunity to contend that the charge should not be imposed or should be reduced.

Exclusion of employee's details

CSIRO is working towards ensuring that all employees have a choice about whether they provide their full name and direct contact details in response to public enquiries. Where such details are included in the scope of a request, this may add to processing time and applicable charges as it may be necessary to consider whether the details are exempt under the FOI Act. On this basis, unless you tell us otherwise, we will assume that these details are out of scope of your request and they will be redacted under section 22 of the FOI Act.

Please note that information released under the FOI Act may later be published online on CSIRO's disclosure log, subject to certain exceptions. For example, personal information will not be published where this would be unreasonable.

CSIRO will contact you using the email address you provided. Please advise if you would prefer us to use an alternative means of contact.

Yours sincerely,

Karen Robinson
FOI Administrative Officer
CSIRO



Attachment B

FOI2021/25 - Document Schedule

Doc No	Date	Addressee	Author	Description	No of Pages	Section of Act	Decision
1.	28 August 2019	CSIRO	Santos	Email (11:12am): Chemical Risk Assessment Attachment: Removed	1	s45, s47, s47E, s47G(1)(a), s22	Part Exempt Exempt in Full
2.	14 June 2019	CSIRO	CSIRO	Email (1:39pm): FW: Beetaloo Basin – Santos Bore Survey Data Attachment: Removed	2	s45, s47, s47E, s47G(1)(a)	Part Exempt Exempt in Full
3.	6 August 2019	Santos	CSIRO	Email (11:15am): GISERA project – Northern Territory – List of chemicals used by onshore gas development that have environmental impacts	2	s45, s47, s47E, s47G(1)(a) s22	Part Exempt
4.	6 August 2019	Origin	CSIRO	Email (11:14am): GISERA project – Northern Territory – List of chemicals used by onshore gas development that have environmental impacts	2	s45, s47, s47E, s47G(1)(a), s22	Part Exempt
5.	6 August 2019	Santos	Santos	Email (11:17am): Re GISERA project – Northern Territory – List of chemicals used by onshore gas development that have environmental impacts	2	s45, s47, s47E, s47G(1)(a), s22	Part Exempt
6.	6 August 2019	Origin	Origin	Email (12:33pm): Re GISERA project – Norther Territory – List of chemicals used by onshore gas development that have environmental impacts	2	s45, s47, s47E, s47G(1)(a), s22	Part Exempt

Doc No	Date	Addressee	Author	Description	No of Pages	Section of Act	Decision
7.	9 August 2019	Santos	Santos	Email (9:39am): Re GISERA project – Northern Territory – List of chemicals used by onshore gas development that have environmental impacts	3	s45, s47, s47E, s47G(1)(a), s22	Part Exempt
8.	12 August 2021	CSIRO	Santos	Email (9:19am): Re GISERA project – Northern Territory – List of chemicals used by onshore gas development that have environmental impacts	3	s45, s47, s47E, s47G(1)(a), s22	Part Exempt
9.	20 August 2021	Santos	CSIRO	Email (1:21pm): Re GISERA project – Northern Territory – List of chemicals used by onshore gas development that have environmental impacts	4	s45, s47, s47G(1)(a), s22	Part Exempt
10.	20 August 2021	Origin	CSIRO	Email (1:27pm): Re GISERA project – Northern Territory – List of chemicals used by onshore gas development that have environmental impacts	2	s45, s47, s47E, s47G(1)(a), s22	Part Exempt
11.	3 July 2021	CSIRO	Santos	Email (2:26pm): Re NT CSIRO GISERA August field trip – Tanumbirini leg Attachment: Removed	10	s45, s47, s47E, s47G(1)(a), s22	Exempt in Full
12.	13 June 2019	Santos & Others	CSIRO	Email (7:40am): With revised agenda – FW: Agenda & meeting papers – NT project fieldwork ad logistics meeting on 14 June 2019 Attachment 1: Removed Attachment 2: Removed Attachment 3: Removed Attachment 4: Removed Attachment 5: Removed	2	s45, s47, s47E, s47G(1)(a), s22	Exempt in Full



12 July 2021

Our ref: FOI 2021/25

Mr Graeme Sawyer
Protect Country Alliance NT
Via Email:

Dear Mr Sawyer

FREEDOM OF INFORMATION REQUEST – DECISION FOI2021/25

Your request

On 12 May 2021, you sought access under the *Freedom of Information Act 1982* (FOI Act) to:

“Correspondence any time during September 2018 to end May 2021 between representatives of GISERA/CSIRO and with representatives of APPEA, Santos and/or Origin Energy containing the terms:

Sulphide reducing bacteria, Sulphate reducing bacteria, Microbially induced concrete corrosion (MICC), sulfur-oxidizing bacteria (SOB), microbial corrosion. Or documents relating to corrosion in petroleum and gas well casings.”

I have identified 12 documents in relation to your FOI request.

Decision maker

I am an authorised decision maker under section 23 of the FOI Act. This letter sets out my decision and reasons for the decision in relation to your application.

Decision

I have decided that exemptions apply to parts of the relevant documents. The relevant provision/exemption provisions include:

Section 45 – Material Obtained in Confidence
Section 47 – Commercially valuable information
Section 47E - Certain operations of agencies
Paragraph 47G(1)(a) -Business and professional affairs
Section 22 – Irrelevant material

Reasons for decision

My findings of fact and reasons for deciding that the exemption provisions apply to a document or part of a document are set out below.

Exemptions Claimed

Section 45 – Material obtained in Confidence

Subsection 45(1) exempts a document if its disclosure would found an action by a person for breach of confidence.

To found an action for breach of confidence a person must be able to:

- (i) specifically identify the information in question;
- (ii) show that the information has the necessary quality of confidentiality (and is not, for example, common or public knowledge);
- (iii) show that the information was communicated in a mutual understanding that the receiver was to keep the information confidential; and
- (iv) show that there is actual or threatened misuse of that information.

I am of the view that disclosure of a document exempted under section 45 would found an action for breach of confidence. CSIRO works closely with industry partners and much of the material communicated between CSIRO and its industry partners is obtained in confidence.

Identified with specificity

The information is able to be identified with specificity as being the information contained in those documents. Moreover, the enforceability of the confidentiality is specifically tied to the material contained within the documents requested.

Confidential in nature

The information is confidential in nature and identified as such within the documents.

Communicated in confidence

I find that the information was communicated within the context of a mutual understanding that both parties would treat it as confidential, and such confidentiality was enforceable.

Disclosure a misuse

I find that disclosure under FOI would constitute a misuse of the information. Such disclosure would be inconsistent with the understanding that CSIRO would keep the information confidential. There is nothing to suggest that the person or body that communicated the information authorised the CSIRO to disclose it under FOI.

Accordingly, I am satisfied the documents are exempt under section 45.

Section 47 - Commercially valuable information

Section 47 provides that a document is exempt from disclosure if its disclosure would disclose a trade secret or, pursuant to para 47(1)(b), the information is has a commercial value “that would be, or could reasonably expected to be, destroyed or diminished if the information was disclosed”.

I find that parts of the documents contain information that has a commercial value and that commercial value would be destroyed or diminished if that information was disclosed.

Commercial value

The parts of the documents exempt under s 47 relate to CSIRO commercial activities, including with private industry partners. Some of the information is of value not only to CSIRO, but more importantly to the third parties to whom CSIRO is contracted with. The information does not have exchange value as it is not a trade secret, nor does it relate to CSIRO’s or third parties’ intellectual property. However, the information relates to the profitability or commercial activities in which CSIRO is involved.

All the information over which s 47 has been applied, is current information and is not out of date.

Diminished or destroyed

I have had regard to whether the value of the information, including the financial information described above, would be diminished or destroyed if disclosed. I am satisfied that the disclosure of the information would result in more than mere criticism or embarrassment for CSIRO and related third parties. I have determined that the intrinsic value of the information would be diminished if it were disclosed. I have had regard to the decision by the Information Commissioner in *McKinnon and Department of Immigration and Citizenship* [2012] AICmr 34 and am satisfied that disclosure of the information would affect more than just the business affairs of the third party.

Having satisfied the two essential criteria of para 47(1)(b), I find that the parts of the documents over which s 47 is claimed, is exempt from disclosure.

Section 47E - Certain operations of agencies

Subsection 47E(3) provides that a document is conditionally exempt if disclosure would or could reasonably be expected to have a substantial adverse effect on the management or assessment of personnel by the Commonwealth or by an agency.

Due to the nature of the material sought under the FOI request, I am of the view that it is appropriate to redact the names of the CSIRO officers from the documents. There have, historically, been incidents where the health and safety of CSIRO officers and their families have been in jeopardy following the release of documents under the FOI Act.

The Office of the Australian Information Commissioner (OAIC) suggest that ‘Specific concerns about the health, safety and wellbeing of staff are most appropriately addressed under the conditional exemption in section 47E(c) of the FOI Act, which is subject to the public interest test. The inclusion of a public interest

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test under section 47E(c) ensures that the public interest in disclosure remains at the forefront of decision making involving this provision.’ (see: [Disclosure of public servants’ name and contact details in response to FOI requests — OAIC](#)).

Further, the OAIC go on to advise:

In certain circumstances, the management of staff and the discharge of the Australian Government’s legal responsibility to ensure the health and safety of its staff may be substantially and adversely affected if public servants’ names and contact details are routinely disclosed in response to FOI requests. Agencies must take all reasonable steps to minimise the risk of harm to staff to be compliant with their statutory obligations under section 19 of the *Work Health and Safety Act 2011*. As discussed, these known risks have evolved over time as a result of the changing digital environment.

I am satisfied, in this instance, that there is a sufficient public interest in the need to protect the identity of CSIRO officers in undertaking their work. It should be noted that this is not a blanket view, but a position that is applied on a case by case basis. I have therefore decided to redact the names of the relevant CSIRO officers from the documents being disclosed.

Section 47G - Business and professional affairs

Section 47G provides that a document is conditionally exempt if disclosure would, or could reasonably be expected to:

‘unreasonably affect that person adversely in respect of his or her lawful business or professional affairs or that organisation or undertaking in respect of its lawful business, commercial or financial affairs.’

Your request concerns documents that contain information concerning the business affairs of a third party (as specified in your request). In determining whether material contained within documents is conditionally exempt under para 47G(1)(a) Part 6 of the Guidelines made by the Office of the Australian Information Commissioner state that consideration of this exemption rests on the effect of disclosure rather than the precise nature of the information. I find that the information in this document could affect the business of the related third party.

I am therefore satisfied that the criteria for conditionally exempting the material in this document under s 47G(1)(a) has been met.

The public interest test: para 47G(1)(a)

Conditionally exempt matter must be released unless, in the circumstances, access to that document would, on balance, be contrary to the public interest under subs 11A(5).

I have considered whether disclosure would be in the public interest. In particular, I have taken into account:

- i. promoting the objects of the FOI Act, particularly in increasing scrutiny, discussion, comment and review of the Government's activities (s 3(2)(b) FOI Act) and in particular:
 - informing debate on a matter of public importance
 - promoting effective oversight of public expenditure

On balance, I do not consider that the public interest in the disclosure of the information is sufficiently strong to override the other considerations, namely that disclosure of the information in this document, if released, would or could reasonably be expected to adversely affect a third party's business affairs.

Accordingly, I am satisfied the documents are part exempt, under section 47G(1)(a).

Section 22 Access to edited copies with exempt or irrelevant matter deleted

Section 22 of the FOI Act provides:

- (1) This section applies if:
 - (a) an agency or Minister decides:
 - (i) to refuse to give access to an exempt document; or
 - (ii) that to give access to a document would disclose information that would reasonably be regarded as irrelevant to the request for access; and
 - (b) it is possible for the agency or Minister to prepare a copy (an edited copy) of the document, modified by deletions, ensuring that:
 - (i) access to the edited copy would be required to be given under section 11A (access to documents on request); and
 - (ii) the edited copy would not disclose any information that would reasonably be regarded as irrelevant to the request; and
 - (c) it is reasonably practicable for the agency or Minister to prepare the edited copy, having regard to:
 - (i) the nature and extent of the modification; and
 - (ii) the resources available to modify the document; and
 - (d) it is not apparent (from the request or from consultation with the applicant) that the applicant would decline access to the edited copy.

Access to edited copy

- (2) The agency or Minister must:
 - (a) prepare the edited copy as mentioned in paragraph (1)(b); and
 - (b) give the applicant access to the edited copy.

I have decided that some of the information in the documents is irrelevant to your request and is therefore removed under section 22. I have decided that I can easily prepare an edited copy of the documents with the irrelevant material deleted and have done so.

Materials taken into account

The materials, information and advice to which I have had reference in making this decision are:

- i. *the terms of your FOI request;*
- ii. *the content of the document in issue;*

- iii. *the relevant provisions of the FOI Act;*
- iv. *guidelines issued by the Office of the Australian Information Commissioner under s 93A of the FOI Act (the Guidelines) and*
- v. *relevant case law.*

Rights of Review

In accordance with section 26(1)(c) of the FOI Act, a statement setting out your rights of review under the Act is at Attachment A.

Yours sincerely

Stephen Jones
Legal Counsel

Attachment A

Review rights

You are entitled to seek review of this decision.

Internal Review

Firstly, under section 54 of the FOI Act, you may apply for an internal review of the decision. Your application must be made by whichever date is the later between:

30 days of you receiving this notice; or 15 days of you receiving the documents to which you have been granted access.

An internal review will be conducted by a different officer from the original decision-maker. No particular form is required to apply for review although it will assist your case to set out in the application the grounds on which you believe that the original decision should be overturned. An application for a review of the decision should be addressed to:

FOI@csiro.au

If you choose to seek an internal review, you will subsequently have a right to apply to the Australian Information Commissioner for a review of the internal review decision.

External review by the Australian Information Commissioner

Alternatively, under 54L of the FOI Act, you may seek review of this decision by the Australian Information Commissioner without first going to internal review. Your application must be made within 60 days of you receiving this notice.

The Information Commissioner is an independent office holder who may review decisions of agencies and Ministers under the FOI Act. More information is available on the Information Commissioner's website www.oaic.gov.au.

You can contact the Information Commissioner to request a review of a decision online or by writing to the Information Commissioner at:

GPO Box 2999
Canberra ACT 2601

Complaints to Ombudsman or Information Commissioner

You may complain to either the Commonwealth Ombudsman or the Information Commissioner about action taken by CSIRO in relation to the application. The Ombudsman will consult with the Information Commissioner before investigating a complaint about the handling of an FOI request.

Your enquiries to the Ombudsman can be directed to:

Phone 1300 362 072 (local call charge)

CSIRO

Australia's National Science Agency

Email ombudsman@ombudsman.gov.au

Your enquiries to the Information Commissioner can be directed to:

Phone 1300 363 992 (local call charge)

Email enquiries@oaic.gov.au

There is no particular form required to make a complaint to the Ombudsman or the Information Commissioner. The request should be in writing and should set out the grounds on which it is considered that the action taken in relation to the request should be investigated and identify CSIRO as the relevant agency.



11 May 2021

Our ref: FOI2021/16

Mr Graeme Sawyer
Protect Country Alliance NT
Via Email:

EEDOM OF INFORMATION REQUEST – DECISION FOI2021/16

Your request

On 12 March 2021, you sought access under the *Freedom of Information Act 1982* (FOI Act) to:

“Correspondence any time during September 2020 to end February 2021 between representatives of GISERA/CSIRO and with representatives of APPEA, Santos and/or Origin Energy regarding the report:

- *Characterisation of the stygofauna and microbial assemblages of the Beetaloo Sub-basin, Northern Territory, (GISERA project number: W18. December 2020)”*

I have identified 17 documents in relation to your FOI request.

Decision maker

I am an authorised decision maker under section 23 of the FOI Act. This letter sets out my decision and reasons for the decision in relation to your application.

Decision

I have decided that exemptions apply to parts of the relevant documents. The relevant provision/exemption provisions include:

Section 22 – Irrelevant material

Reasons for decision

My findings of fact and reasons for deciding that the exemption provision applies to a document or part of a document are set out below.

Exemptions Claimed

Section 22 Access to edited copies with exempt or irrelevant matter deleted

Section 22 of the FOI Act provides:

- (1) *This section applies if:*
 - (a) *an agency or Minister decides:*
 - (i) *to refuse to give access to an exempt document; or*
 - (ii) *that to give access to a document would disclose information that would reasonably be regarded as irrelevant to the request for access; and*
 - (b) *it is possible for the agency or Minister to prepare a copy (an edited copy) of the document, modified by deletions, ensuring that:*
 - (i) *access to the edited copy would be required to be given under section 11A (access to documents on request); and*
 - (ii) *the edited copy would not disclose any information that would reasonably be regarded as irrelevant to the request; and*
 - (c) *it is reasonably practicable for the agency or Minister to prepare the edited copy, having regard to:*
 - (i) *the nature and extent of the modification; and*
 - (ii) *the resources available to modify the document; and*
 - (d) *it is not apparent (from the request or from consultation with the applicant) that the applicant would decline access to the edited copy.*

Access to edited copy

- (2) *The agency or Minister must:*
 - (a) *prepare the edited copy as mentioned in paragraph (1)(b); and*
 - (b) *give the applicant access to the edited copy.*

I have decided that some of the information in the documents is irrelevant to your request and is therefore removed under section 22. I have decided that I can easily prepare an edited copy of the documents with the irrelevant material deleted and have done so.

Materials taken into account

The materials, information and advice to which I have had reference in making this decision are:

- i. the terms of your FOI request;*
- ii. the content of the document in issue;*
- iii. the relevant provisions of the FOI Act;*
- iv. guidelines issued by the Office of the Australian Information Commissioner under s 93A of the FOI Act (the Guidelines) and*
- v. relevant case law.*

Rights of Review

CSIRO

Australia's National Science Agency

In accordance with section 26(1)(c) of the FOI Act, a statement setting out your rights of review under the Act is at **Attachment A**.

Yours sincerely,

Beth Cribb
Senior Legal Counsel
CSIRO

Review rights

You are entitled to seek review of this decision.

Internal Review

Firstly, under section 54 of the FOI Act, you may apply for an internal review of the decision. Your application must be made by whichever date is the later between:

30 days of you receiving this notice; or 15 days of you receiving the documents to which you have been granted access.

An internal review will be conducted by a different officer from the original decision-maker. No particular form is required to apply for review although it will assist your case to set out in the application the grounds on which you believe that the original decision should be overturned. An application for a review of the decision should be addressed to:

FOI@csiro.au

If you choose to seek an internal review, you will subsequently have a right to apply to the Australian Information Commissioner for a review of the internal review decision.

External review by the Australian Information Commissioner

Alternatively, under 54L of the FOI Act, you may seek review of this decision by the Australian Information Commissioner without first going to internal review. Your application must be made within 60 days of you receiving this notice.

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Canberra ACT 2601

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Email ombudsman@ombudsman.gov.au

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Email enquiries@oaic.gov.au

There is no particular form required to make a complaint to the Ombudsman or the Information Commissioner. The request should be in writing and should set out the grounds on which it is considered that the action taken in relation to the request should be investigated and identify CSIRO as the relevant agency.

s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Wednesday, 2 December 2020 12:47 PM
To: s22
Cc: Cunningham, Paul (CorpAffairs, Dutton Park)
Subject: FOR ADVICE: Santos attendees at CSIRO's GISERA knowledge transfer session - NT water project

Hi s22

I hope that you are well.

I am arranging a knowledge transfer session with industry and government to present the key findings from CSIRO's GISERA water project 'Characterisation of the stygofauna and microbial assemblages of the Beetaloo Sub Basin, NT'.

We have earmarked the week commencing 14 December for this session. It is likely to be for 1 hour and will be conducted via WebEx.

Can you please let me know who from Santos should be included in the invite (we were thinking 2-4 people). It is important to note that this will be a closed session and we do try to manage numbers so that we can better facilitate the open discussion part of the session.

I look forward to receiving your earliest response.

Regards
Jizelle

Jizelle Khoury
Executive Officer, CSIRO's Gas Industry Social and Environmental Research Alliance (GISERA)
Energy | CSIRO

s22

Address: PO Box 52, North Ryde NSW 1670, Australia

CSIRO Australia's National Science Agency | csiro.au

s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Wednesday 2 December 2020 12:52 PM
To: s22@origin.com.au
Cc: Cunningham, Paul (CorpAffairs, Dutton Park)
Subject: FOR ADVICE: Origin attendees at CSIRO's GISERA knowledge transfer session - NT water project

Hi s22

I hope that you are well.

I would normally write to s22 as the GISERA NT Regional Research Advisory Committee rep from Origin, but I understand she is currently on leave.

I am arranging a knowledge transfer session with industry and government to present the key findings from CSIRO's GISERA water project 'Characterisation of the stygofauna and microbial assemblages of the Beetaloo Sub-Basin, NT'.

We have earmarked the week commencing 14 December for this session. It is likely to be for 1 hour and will be conducted via WebEx.

Can you please let me know who from Origin should be included in the invite (we were thinking 2-4 people). It is important to note that this will be a closed session and we do try to manage numbers so that we can better facilitate the open discussion part of the session.

I look forward to receiving your earliest response.

Regards
Jizelle

Jizelle Khoury
Executive Officer, CSIRO's Gas Industry Social and Environmental Research Alliance (GISERA)
Energy | CSIRO

s22

Address: PO Box 52, North Ryde NSW 1670, Australia

CSIRO Australia's National Science Agency | csiro.au

s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Tuesday, 8 December 2020 7:39 AM
To: s22
Cc: Cunningham, Paul (CorpAffairs, Dutton Park)
Subject: RE: FOR ADVICE: Santos attendees at CSIRO's GISERA knowledge transfer session - NT water project

Hi s22

I wanted to let you know that we have earmarked Monday, 14 December at 9.30-10.30 NT time (10-11 am QLD) for the knowledge transfer session.

I haven't sent out a calendar invite yet as I'm waiting for the NTG to finalise their participant list. It will go out by Thursday (latest). In the meantime, can I ask that you please hold this timeslot in your diaries.

Many thanks
Jizelle

Jizelle Khoury
Executive Officer, CSIRO's Gas Industry Social and Environmental Research Alliance (GISERA)
Energy | CSIRO

s22
Address: PO Box 52, North Ryde NSW 1670, Australia

CSIRO Australia's National Science Agency | csiro.au

From: s22@santos.com>
Sent: Wednesday, 2 December 2020 1:25 PM
To: Khoury, Jizelle (Energy, North Ryde) s22
Cc: Cunningham, Paul (CorpAffairs, Dutton Park) s22
s22@santos.com>; s22@santos.com>
Subject: RE: FOR ADVICE: Santos attendees at CSIRO's GISERA knowledge transfer session - NT water project

Hi Jizelle,

Can you please invite David Gornall, Mitch Bird (both CC'ed into this email) and myself

Regards,



s22

Santos Limited, 32 Turbot Street, Brisbane QLD 4000

s22



Duplicate Email - Removed

s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Tuesday, 8 December 2020 7:40 AM
To: s22
Cc: Cunningham, Paul (CorpAffairs, Dutton Park)
Subject: RE: FOR ADVICE: Origin attendees at CSIRO's GISERA knowledge transfer session - NT water project

Hi s22

I wanted to let you know that we have earmarked Monday, 14 December at 9.30 10.30 NT time (10 11 am QLD) for the knowledge transfer session.

I haven't sent out a calendar invite yet as I'm waiting for the NTG to finalise their participant list. It will go out by Thursday (latest). In the meantime, can I ask that you please hold this timeslot in your diaries.

Many thanks
Jizelle

Jizelle Khoury
Executive Officer, CSIRO's Gas Industry Social and Environmental Research Alliance (GISERA)
Energy | CSIRO

s22

Address: PO Box 52, North Ryde NSW 1670, Australia

CSIRO Australia's National Science Agency | csiro.au

From: s22@origin.com.au>
Sent: Wednesday, 2 December 2020 1:21 PM
To: Khoury, Jizelle (Energy, North Ryde) s22@origin.com.au>; s22@origin.com.au>
Cc: Cunningham, Paul (CorpAffairs, Dutton Park) s22
Subject: RE: FOR ADVICE: Origin attendees at CSIRO's GISERA knowledge transfer session NT water project

Hi Jizelle,

I remember you from when I was working with Shell; I hope you're well.

At this stage we are planning for Matt and Carl Altmann (copied in) to join the session in w/c 14 December.

Kind regards,

o o o o

s22

Origin
Level 28, 180 Ann Street, Brisbane QLD 4000
s22

Connect with us



originenergy.com.au

I acknowledge the Traditional Owners and Custodians of country throughout Australia and recognise their continuing connection to land, waters and community.

I pay my respects to them and their cultures, and to Elders past, present and future.

Origin shows its commitment to participating in Australia's reconciliation efforts through our [Stretch Reconciliation Action Plan](#).



Duplicate Email - Removed

s22

From: Cunningham, Paul (CorpAffairs, Dutton Park)
Sent: Tuesday, 8 December 2020 4:49 PM
To: s22@aplng.com.au; Bertsch, Paul (L&W, Dutton Park);
s22@santos.com; s22@shell.com;
s22@originenergy.com.au; Niemelae, Marita (Energy, Kensington WA);
David.Lawrence s22 s22@pangaea.net.au
Cc: Barrett, Damian (Energy, Black Mountain); Khoury, Jizelle (Energy, North Ryde);
kathrine.Riley s22 Nathan.Bartlett s22 Bloom, Wendy
(Energy, Clayton); Cunningham, Paul (CorpAffairs, Dutton Park); s22
Subject: CSIRO's GISERA NT project - final report stygofauna and microbial assemblages of
the Beetaloo Sub-basin, NT
Attachments: GISERA Project18 Stygofauna final report 20201208.pdf

Dear all,

Re: GISERA's Northern Territory project [Characterisation of the stygofauna and microbial assemblages of the Beetaloo Sub basin, NT.](#)

Please find attached a copy of the project final report which describes new knowledge about stygofauna and subterranean groundwater dependent ecosystems in the Beetaloo Sub-basin and Roper River system.

This completes project milestone task 7. A knowledge transfer session is scheduled for Government and industry representatives later this month and the final report will be made available on the GISERA web site in early 2021.

Results of this research help address a previously identified knowledge gap regarding stygofauna and subterranean groundwater dependent ecosystems in the Beetaloo Sub-basin and Roper River system, in line with recommendation 7.20 from the Final Report of the Scientific Inquiry into Hydraulic Fracturing in the Northern Territory.

This was a collaborative project between CSIRO and the Research Institute for Environment and Livelihoods, Charles Darwin University. Using a range of sampling devices, researchers collected samples from 26 groundwater bores and two springs in August and October 2019, across a distance of approximately 500 km, from the sub-tropical Mataranka region in the north to the semi-arid Barkly Tablelands (Barkly Stock Route) in the south.

Results confirm Northern Territory aquifers support a diverse range of stygofaunal species. All Beetaloo stygofaunal communities sampled were dominated by crustaceans - shrimps, amphipods, ostracods, copepods and syncarids. This fauna showed little affinity with the stygofauna recorded from more extensively sampled Western Australian aquifers, with new genera and species of crustaceans (amphipods and ostracods) present in the Beetaloo.

Overall, the presence of stygofauna at widely separated sites across the Cambrian Limestone Aquifer is consistent with substantial connectivity within the aquifer. Further work is required to quantify the risk of contamination impacts on stygofauna from possible industrial spill events that takes into account migration pathways and processes including adsorption, dilution and microbial metabolism in soils and aquifers as well as the high connectivity in groundwater systems.

The stygofauna sampling program is part of a suite of GISERA research projects either completed or in progress in the Northern Territory, in line with Scientific Inquiry recommendations. Baseline studies of methane and groundwater characteristics are complete. New projects in progress include microbial degradation of shale gas related chemicals, minimising potential emissions from gas wells through improved leak sealing technologies and well decommissioning practices, development of high quality spatial data to guide land management practices, and assessment of options to offset life-cycle greenhouse gas emissions.

Results of these studies are important for informing appropriate policy and management responses to shale gas development proposals.

If you wish to discuss these results please feel free to contact GISERA Director Dr Damian Barrett.

Paul Cunningham

Communication and Stakeholder Manager
Gas Industry Social and Environmental Research Alliance
Energy | CSIRO

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Australia's National
Science Agency

Characterisation of the stygofauna and microbial assemblages of the Beetaloo Sub-basin, Northern Territory

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GISERA project number: W18. December 2020

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Acknowledgements

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We are grateful to the staff of Origin Energy who assisted with access to bores associated with the Beetaloo Sub basin and the managers and staff of Buchanan Downs, Newcastle Waters and Eva Downs for allowing access to bores on pastoral stations. We acknowledge Steven Tickell (Northern Territory Government) for information on Northern Territory (NT) aquifers, access to NT government bores, and for providing logistical guidance for field support. We also thank Stuart Halse (Bennelongia Environmental Consultants) and John Short (BioAccess Australia) for assisting with taxonomic identifications, Lisa Chandler (Department of Agriculture, Water and the Environment) for reviews of the literature compiled here, Vanessa Solano Rivera (Charles Darwin University) for generating maps, and Matt Northwood (Charles Darwin University) for contributions to the technical aspects of field work preparation.

Executive summary

The Beetaloo Sub basin and Roper River system overlies several major units of the Cambrian Limestone Aquifer (CLA) and is one of the most prospective areas for shale gas in Australia. Groundwater biota, in addition to their biodiversity values, provide an indication of aquatic health of aquifers and are integral to the ecosystem services (the benefits to humans) provided by these systems. As a consequence, protection of subterranean groundwater dependent ecosystems (GDEs) has been recognised at the federal level for over 20 years. Minimal previous sampling of the groundwater ecosystems has been undertaken in this region and knowledge of the subterranean fauna within the region is poor.

The overall objective of this project was:

- To provide new knowledge concerning stygofauna and subterranean groundwater dependent ecosystems in the Beetaloo Sub basin and Roper River system, a critical knowledge gap identified by the Final Report of the Scientific Inquiry into Hydraulic Fracturing in the Northern Territory (2018).

To this end, this project undertook a broad spatial scale pilot survey of bores in the Beetaloo sub Basin to determine the distribution and abundance of stygofauna and characterise stygofauna communities within the subterranean groundwater-dependent ecosystems that might be present.

We sampled 26 groundwater wells (bores) and two springs in August and October 2019, across a distance of ~ 500 km, from the sub tropical Mataranka region in the north to the semi-arid Barkly Tablelands (Barkly Stock Route) in the south. We used a range of sampling devices, including plankton nets and motorised pumps, depending on the type and size of bore hole. All live stygofaunal samples were filtered through a 50 µm mesh sized net, stored in 70% ethanol and subsequently analysed by microscopy in the lab. For DNA analysis, we collected and preserved 300ml of water for subsequent filtering and DNA extraction in the lab. We targeted the cytochrome oxidase I gene (COI) to determine the presence of stygofauna and the 16s ribosomal gene (16sRNA) to gain a fingerprint of all bacteria present. We also extracted genetic material from shrimp tissues for COI barcoding. At every site we measured the depth to the water table, electrical conductivity (EC), pH and water temperature (°C).

Key findings

- Northern Territory aquifers support a diverse range of stygofaunal species.
- All Beetaloo stygofaunal communities sampled were dominated by crustaceans, namely: shrimps, amphipods, ostracods, copepods and syncarids. This fauna showed little affinity with the stygofauna recorded from more extensively sampled Western Australian aquifers, with new genera and species present in the Beetaloo Sub-basin.
- Morphological and genetic (COI and 16S RNA gene) assessment indicates that all atyid specimens (shrimps) comprise a single species, *Parisia unguis*. The presence of this species, ranging across a geographic distance of ~300 km, and the low genetic divergence (maximum 3.9% in COI and 3.29% in 16S RNA gene) among specimens indicate groundwater connectivity in recent times.
- Overall, the presence of stygofauna at widely separated sites across the Cambrian Limestone Aquifer is consistent with substantial connectivity within the aquifer. Further work is required to quantify the risk of contamination impacts on stygofauna from possible spill events that takes into account migration pathways and processes including adsorption, dilution and microbial metabolism in both soil and aquifer as well as the high connectivity in ground water systems.
- eDNA methods were a highly valuable tool in detecting the presence of subterranean biota, in conjunction with traditional sampling methods, but showed particular value where structures prevented using nets to collect samples.
- Diverse microbial communities could be obtained from bore samples, with aerobic heterotrophic bacteria dominating microbial communities.
- Denitrifying bacteria were present in many wells, which is consistent with growth denitrifying bacteria using the high levels of nitrate that have been measured in bore water samples. Similarly, sulfate within the water supported sulfate reducing bacterial populations. These microorganisms are likely to be colonising bore casings etc as part of complex biofilm communities growing on the hard surfaces.

Our study is the first step in the description of the biodiversity and ecological integrity of the subterranean GDEs of the Beetaloo Sub basin. This baseline information will support the development of policy, management and monitoring guidelines for the extraction of shale gas within this region.

Introduction

Groundwater dependent organisms

Groundwater, the water stored beneath the Earth's surface, is an important resource worldwide, and especially so in Australia, where over 70% of the continent is arid or semi-arid, annual rainfall is low (< 500mm) and surface waters are scarce. In addition to supporting terrestrial, aquatic and subterranean ecosystems, known as groundwater dependent ecosystems (GDEs), groundwater supports human settlements and many industries, including agriculture, horticulture, mining and gas and oil extraction. Despite the importance of groundwater across inland Australia, the fact that it is stored underground means that it is often 'out of sight and out of mind'.

Although groundwater biota and groundwater ecology are the subject of an increasing number of studies (e.g. Boulton *et al.* 2008, Humphreys 2009, Tomlinson and Boulton 2010, Nwankwoala 2012) changes in groundwater quality and quantity and their effects on the ecosystem that exists within aquifers remain poorly understood. In addition to micro-organisms (largely Bacteria, Protozoa and algae) and biofilms (an aggregation of micro-organisms), groundwater houses a range of aquatic invertebrates collectively known as stygofauna. These are mainly crustaceans (notably amphipods, copepods and ostracods but also isopods, syncarids and decapods), as well as a range of worms (nematodes, annelids and platyhelminthes), molluscs, mites, beetles and occasionally fish. Stygofaunal communities are considered important as a biodiversity resource, as indicators of groundwater health and as providers of ecosystem goods and services (Glanville *et al.* 2016, Smith *et al.* 2016). Despite the increased focus of research on groundwater systems, there are still many knowledge gaps concerning the diversity, distribution and ecology of stygofauna and the impacts of anthropogenic disturbances and associated toxic chemicals on groundwater communities (Hose *et al.* 2015a, Di Lorenzo *et al.* 2019).

Stygofauna occur in a wide range of groundwater habitats, residing in the pore spaces or fissures of any rock or sediment typically within fresh or saline aquifers, but also in cave systems and springs. The presence of stygofauna is a good indication of a healthy ecological community that also supports other micro-organisms, including bacteria and protozoans (Doody *et al.* 2019). The likelihood of stygofauna occurring in an aquifer is determined by the aquifer type, geology and hydraulic conductivity; groundwater depth and distance from exchange zones; and water quality (Hose *et al.* 2015a). Of these attributes, the availability of large enough pore spaces is key to the type of groundwater organisms that may be supported (Hahn and Fuchs 2009, Hose *et al.* 2015a).

Diversity and abundance of stygofaunal communities is higher within the upper 1–2 m of groundwater, where hydrological exchange between aquifer and surface water is strongest (Danielopol *et al.* 1997, Schmidt *et al.* 2007, Bork *et al.* 2009). Aquifers with a water table < 10m below the surface, and penetrated by phreatophytic tree roots, or influxes of other sources of organic material, are most likely to harbour species-rich communities (Hancock and Boulton 2008, Eberhard and Davies 2011, Chilcott 2013). Stygofauna are rarely found more than 100 m below the ground, where nutrient levels and dissolved oxygen concentrations are low (Hose *et al.* 2015a).

Australian aquifers and stygofauna diversity

In Australia there are three general types of aquifer in which stygofauna have been found—karstic, fractured rock and alluvial (Figure 1). Karstic systems are characterised by sink holes, caves and springs commonly developed in carbonate rocks such as limestone and dolomite. They are found across Australia, including on the Nullarbor Plain, in Cape Range National Park, and throughout northern Australia from the Kimberley to the Barkly Tableland (Tomlinson and Boulton 2008). Fractured rock aquifers occur when fissures or cracks develop in rocks of sedimentary, igneous or metamorphic origin. Groundwater flow follows the fractures but can also permeate the rock matrix, depending on the geology of the system (Tomlinson and Boulton 2008). The Pilbara is a highly diverse region for stygofauna inhabiting fractured rock aquifers (Hose *et al.* 2015a). Alluvial aquifers occur in unconsolidated sediments, often sands and gravels associated with river flood plains and deposits. Stygofauna have been collected from several alluvial aquifers, foremost across eastern Australia, such as the Burdekin River catchment in north Queensland and the Peel and Gwydir river regions in New South Wales (Figure 1) (Hose *et al.* 2015a).

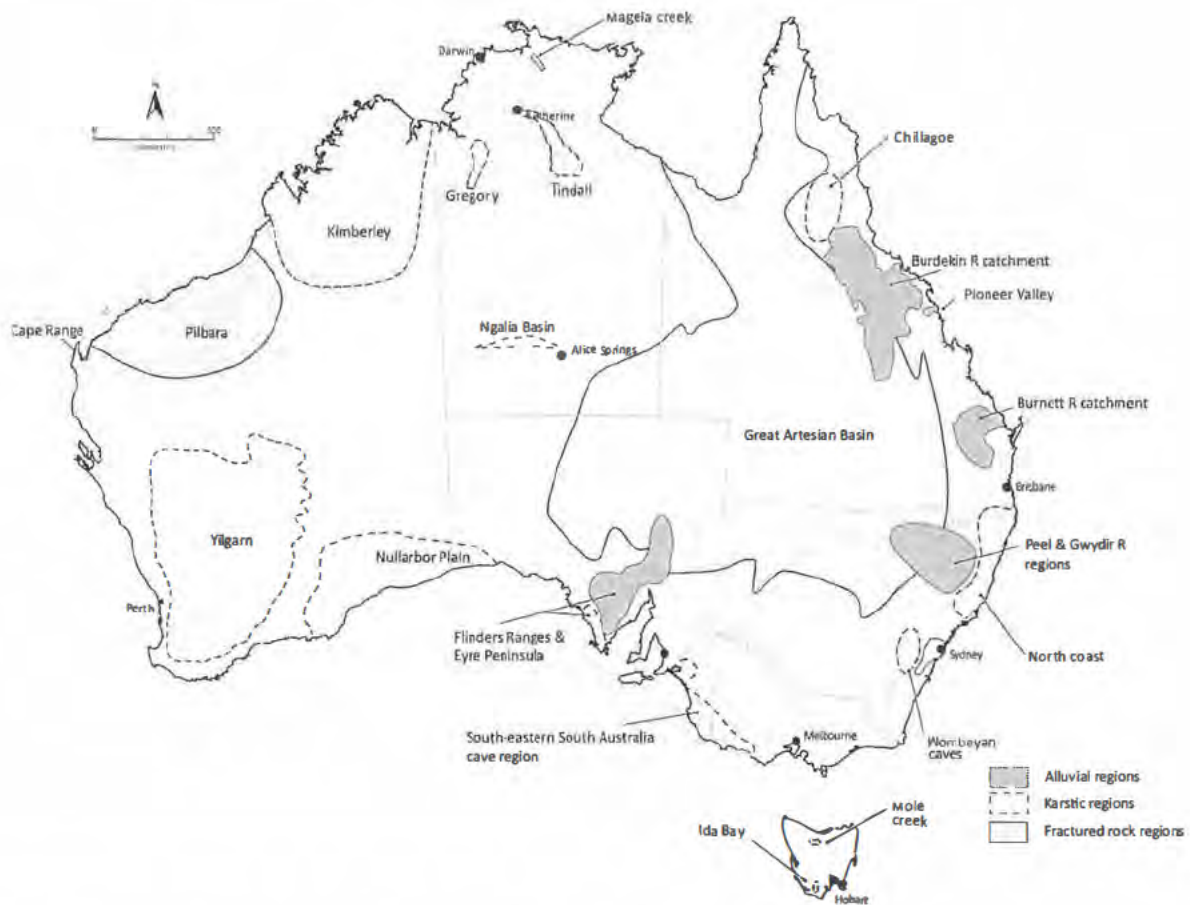


Figure 1. General aquifer types and regions where stygofauna have been found in Australia. Modified from Tomlinson and Boulton (2008) with additional information from Guzik *et al.* (2008), Hose *et al.* (2015a) and Chandler *et al.* (2017) and this study (the Tindall aquifer).

Groundwater biological communities are typically less diverse and abundant than those of surface water environments (Gibert *et al.* 1994). However, the isolation of aquifers and limited dispersal abilities of stygofauna has created a fauna dominated by short range endemic species, so while species diversity may be low locally, diversity across regions is often high (Hose *et al.* 2015a).

The low diversity of stygofauna may be a product of the patchy sampling effort in Australia, and globally, and it is likely that the true extent of biodiversity and distribution of stygofauna is undocumented (Tomlinson and Boulton 2010). Guzik *et al.* (2011) estimated that the reported 770 subterranean taxa (both stygofauna and troglotauna) known from the western half of Australia (from the Kimberley in the north-west through to Eyre Peninsula in the south east) represent only 20% of species, and that much more extensive survey effort is needed to thoroughly assess Australia's stygofauna diversity. Humphreys (2008) noted that, even in its infancy of research,

Australia is a groundwater biodiversity hotspot, the ~750 stygofauna species known to date representing 22% of global totals.

Western Australia

Western Australia represents the forefront of stygofauna research, the first cave-dwelling species recorded as early as the 1940s (Whitley 1945, Holthuis 1960). The 1990s saw the discovery of stygofauna within aquifers (Adams and Humphreys 1993, Humphreys 2000) and led to ongoing surveys of wells especially in association with mineral resources development throughout the Kimberley (e.g. Cho et al. 2005, Karanovic 2005, Rockwater Pty Ltd 2012), Pilbara (e.g. Eberhard et al. 2005, Halse et al. 2014) and Yilgarn (e.g. Leys and Watts 2008, Karanovic et al. 2013) regions. Unlike the rest of Australia, the west encompasses an ancient mineral-rich region that remained largely emerged throughout frequent marine inundations since the Palaeozoic (Guzik et al. 2011). Groundwater systems are largely calcrete karst (Kimberley, Yilgarn) and fractured rock (Pilbara). These have a greater potential to support a diversity of organisms than alluvial sediment because they comprise areas with larger pore spaces (Hose et al. 2015a). Research from Western Australia continues to report a diverse range of species, especially crustaceans and beetles. Only three stygofaunal vertebrates have been recorded in Australia; a blind cave eel, *Ophisternon candidum* Mees 1962, and two blind cave gudgeons, *Milyeringa veritas* and *M. Justitia* Whitley 1945, all from the karst of Cape Range (Humphreys 2006) and Barrow Island (Humphreys et al. 2013). Growing awareness of stygofauna, their rarity and possible fragility, has raised concerns of species loss associated with extensive mining developments throughout the region. To date, 20 stygofauna (and 27 troglifauna) are listed as threatened under the Western Australian Wildlife Conservation Act 1950, and two species—the blind cave eel and a remipedian crustacean, *Lasionectes exleyi* (Yager and Humphreys 1996)—are listed as vulnerable under Commonwealth legislation.

Eastern Australia (Queensland, NSW, Victoria and Tasmania)

Most surveys of stygofauna in eastern Australia have been conducted in alluvial aquifers in northern New South Wales and the southern regions of Queensland. Although the landscapes of eastern Australia are geologically much younger than those of the west and its subterranean fauna is seemingly less diverse (Guzik et al. 2011), it comprises the richest stygofauna recorded from alluvial systems (Hose et al. 2015b). Generally, the same higher taxa known in Western Australia are present, but composition and abundances are quite different. Frequent marine inundations throughout the Cretaceous period are thought to have diminished diversity of some crustacean

taxa, such as amphipods and isopods, which are less frequently recorded in the east (Hose *et al.* 2015a). More commonly encountered are a range of syncarids and copepods (e.g. Cook *et al.* 2012, Schulz *et al.* 2013), and several families of Anaspidacidea, which have not been recorded outside eastern Australia (Hobbs III 2000, Serov 2002). Although rare, dytiscid and elmids beetles are also known (Watts *et al.* 2007, 2008), but comprise distinct lineages to the diverse assemblages recorded from the karstic Yilgarn basin of Western Australian (Leys *et al.* 2003, Watts *et al.* 2008). Small fragments of karstic (e.g. Wombeyan Caves, Ida Bay, Mole Creek) and fractured rock (e.g. Sydney and Hawkesbury regions) regions in New South Wales and Tasmania have yielded over 100 stygobiotic taxa (largely copepods and syncarids; see Thurgate *et al.* 2001), but diversity and abundances are far lower than those known from similar systems in Western Australia (Eberhard *et al.* 2005). Much of eastern Australia remains unexplored and further surveys are needed to better understand the distribution of species. So far, taxa appear spatially limited within aquifers and local endemism is likely high (Asmyhr *et al.* 2014, Little 2014).

South Australia

A diverse range of stygofauna have been recorded in the alluvial aquifers of the Mount Lofty Ranges, Flinders Ranges and Eyre Peninsula in arid South Australia (Leijs and Mitchell 2009). These include chiltoniid amphipods (R. King and R. Leijs pers. comm. in Hose *et al.* 2015a), parabathynellids (Abrams *et al.* 2013) and dytiscid beetles (Leys *et al.* 2010). Some of the dytiscid beetles, such as the genus *Paroster*, show morphological and phylogenetic similarities to species from the Yilgarn area of Western Australia. Although the distribution of stygofauna is highly fragmented throughout Australia, many species appear to be connected to ancient lineages that were once more widespread (Humphreys 2000, Jaume *et al.* 2001). While aquifers are clearly rich in diversity, the distribution, endemism and national significance of the South Australian stygofauna is still too poorly known to develop and implement policies for protecting local groundwater communities (Goonan *et al.* 2015).

Northern Territory

Although there are extensive karstic/carbonate systems across the Top End of the Northern Territory (Tickell 2005), the region's groundwater has rarely been assessed for aquatic biota. There are few published records of stygofauna across the Northern Territory and these appear to be limited to sporadic surveys in five locations. In the arid Ngalia Basin calcrete aquifers, northwest of Alice Springs, taxa similar to those found in Western Australian calcretes have been reported—

dytiscid beetles (Balke and Ribera 2004, Balke *et al.* 2004, Watts and Humphreys 2006, Leys and Watts 2008), *Haloniscus* isopods (S. Taiti, unpub. data in Hose *et al.* 2015a) and a parabathynellid syncarid (Cho *et al.* 2006a). Further north in the Cutta Cutta caves near Katherine, three species of blind atyid shrimps (Williams 1964, Bruce 1992) and a *Mesocyclops* copepod (Dumont and Maas 1983) are known. On the western border of the Northern Territory, in the Judburra/Gregory karst caves, two stygobiont species—an unclassified amphipod and a hydrobiid gastropod—were recorded alongside troglobiont and other caverniferous taxa (Moulds and Bannink 2012). In the monsoonal north, the alluvial aquifers of Magela Creek are known to harbour undescribed hyporheic fauna (Dostine *et al.* 1997), and more recently also stygofauna—cyclopoid and harpacticoid copepods and parabathynellid syncarids (Chandler *et al.* 2017). Zaar (2009) noted the presence of subterranean isopods (Asellota: Protojaniridae) both at Pungalina in the Gulf County and at Gregory National Park in the Victoria River District, and Van Dam *et al.* (2008) of nondescript stygofauna in the karstic aquifers of the Daly River catchment. These records remain unpublished and further investigation is required to assess the presence of stygofauna in these areas.

Study scopes and aims

The importance of understanding groundwater systems and on-shore gas-related threatening processes is becoming increasingly urgent in the Northern Territory. Current stygofauna survey efforts are mainly associated with mining developments, for example, recent surveys of Magela Creek area in Kakadu National Park are associated with rehabilitation of Ranger Uranium Mine (see Chandler *et al.* 2017). Further south, new on-shore gas developments in the Beetaloo Sub-basin and Roper River region have indicated the need to survey groundwater communities, since to date no assessments have been carried out. This region is likely to support stygofauna because much of it contains fractured and karstic aquifers. The connection between Northern Territory aquifers is yet to be fully established but needs to be assessed since hydrological connectivity is an important driver of surface freshwater macroinvertebrate diversity (Davis *et al.* 2018), and could be similarly so for subterranean species.

This study presents a broadscale pilot survey of bores in the Beetaloo Sub-basin and upper Roper River region. It aims to characterise the community structure of the stygofauna and microbial assemblages and determine the environmental variables of the shallow subterranean aquifers of the study area. Such baseline information and ecological understanding of groundwater-

dependent ecosystems in the region is needed to support appropriate policy and resource management decisions in relation to proposed on-shore gas development in the Northern Territory.

Methods

Study location

The Beetaloo Sub-basin lies 180 km southeast of Katherine in the Northern Territory and spans an area of approximately 30,000 km². One of the most prospective areas for shale gas in Australia, it contains an estimated resource of 178,200 petajoules (PJ) of gas. The Beetaloo Sub-basin spans several major aquifers (Figure 2) that consist of either fractured and karstic rocks or fractured and weathered rocks. Of particular importance are those aquifers that contain karstic rocks due to their known importance for stygofauna. The Roper River lies to the north-east of the Beetaloo Sub-basin. It is a perennial river and, with a surface catchment area of more than 80,000 km², is one of the largest river systems in the Katherine region.

Sites/bores

Groundwater was sampled at various locations in the Northern Territory, across a distance of ~500 km from the sub-tropical Mataranka region in the north to the semi-arid Barkly Tablelands (Barkly Stock Route) in the south (Figure 6). This region encompasses the Beetaloo Sub-basin, the upper Roper River and adjacent areas. A total of 28 sites were sampled, which included two springs (Table 2, Table 3). Sites were selected to encompass a broad geographic range, including and adjacent to the Beetaloo Sub-basin, and where sampling was possible. Bore structure and accessibility varied markedly. Bore types included tap-enabled bores (Figure 3A–E); bore openings enclosed in concrete slabs (Figure 3F); bare steel pipes (Figure 3G) or lid-locked steel pipes with inner PVC lining (Figure 3H–I); raised steel pipes with locked screw-cap (Figure 3J); and raised steel pipes enclosed in a telemetered box (Figure 3K). Fig Tree Spring and Botanic Walk Spring in Elsey National Park and Warlock Ponds Spring (on Warlock Ponds Station, adjacent to Elsey National Park) were also sampled, as close as possible to where the spring discharge or groundwater upwelling was visible (Figure 3L). The land tenure of the sites sampled included pastoral leases, local council lands, road-side Northern Territory government registered bores, and Indigenous land trusts.

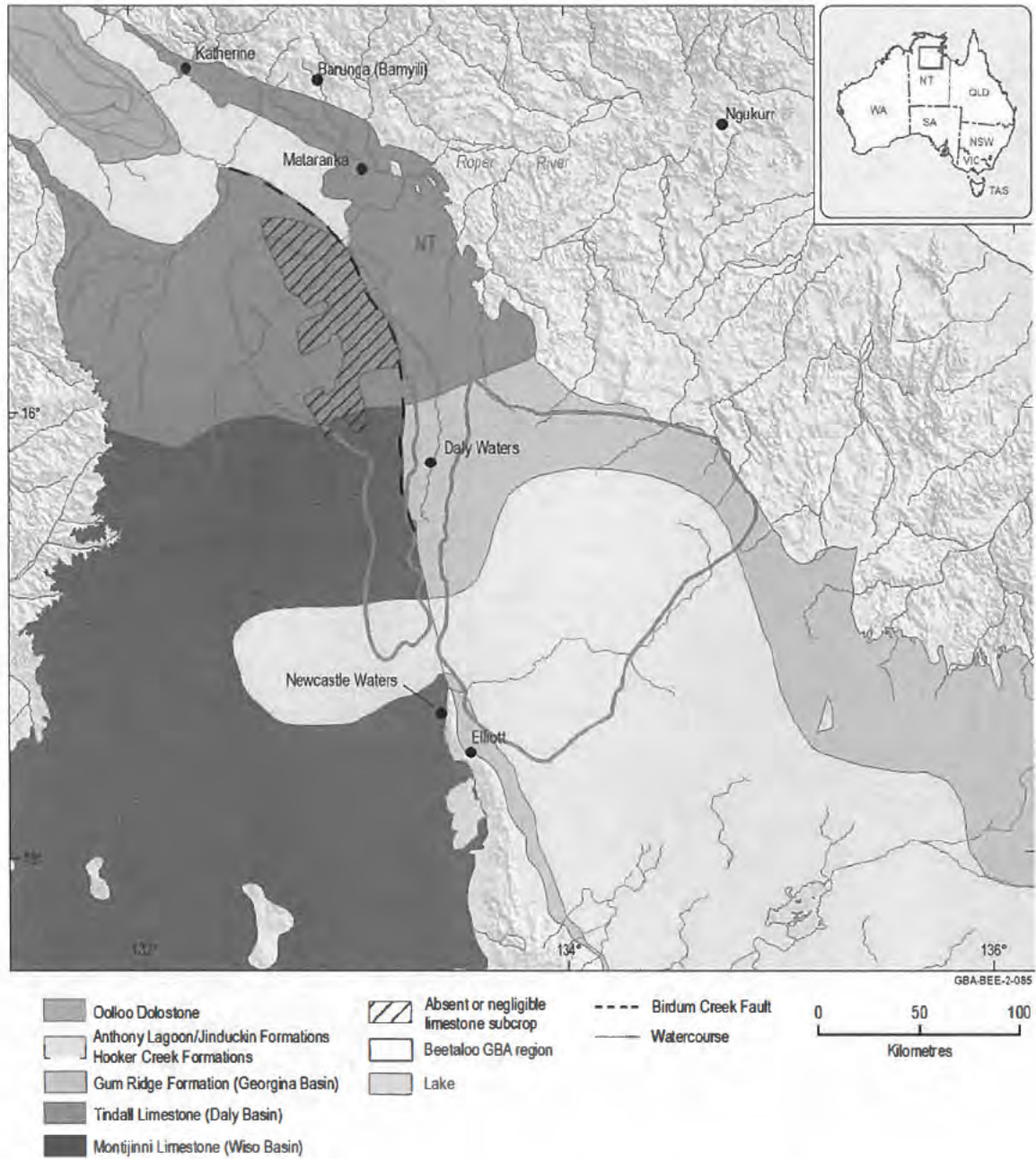


Figure 2. Location of the Beetaloo Sub-basin, marked by the Beetaloo GBA region. The Geological and Bioregional Assessment Program uses the geological boundary of the Beetaloo Sub-basin to delineate the region. Major hydrostratigraphic units that make up the Cambrian Limestone Aquifer are shown. Image source: Evans *et al.* (2020). <https://www.bioregionalassessments.gov.au/assessments/geological-and-bioregional-assessment-program/beetaloo-sub-basin/beetaloo-gba-region-stage-two-report>



Figure 3. Range of bore types sampled; A: Buchanan Downs “Wendy” (RN31243); B: Shenandoah homestead; C: Elliot 8 (RN036781); D: Newcastle Waters (Ferguson Bore); E: Sturt Plains (RN070819); F: Eva Downs 6; G: Carpentaria Highway (RN005942); H: Cutta Cutta South (RN35560); I: Mataranka Homestead (RN35796); J: Tufa (RN RN34032); K: Cave Creek (RN34230); L: Warlock Ponds Spring.

Sampling methods

Sampling was undertaken on two occasions; 5–15 August 2019 and 13–19 October 2019. Sampling devices included motorised pumps and plankton nets, depending on the type and size of the bore hole. Many bores located on pastoral leases had fixed pumps on the bore head (Figure 3A–E) and water samples were collected either directly from pump or via taps. For some bores, on prospective gas industry areas, a large motorised pump was supplied by Origin Energy (Figure 4A),

allowing access to groundwater > 30 m below the surface. Approximately 200–300 L of water was pumped (depending on the pump/tap power and flow rate) and filtered through a large 50 µm mesh size plankton net (Figure 4B). Bores where the water table was higher than 30 m could be accessed by a smaller battery operated pump (Figure 4C–D). At its base the large plankton net (Figure 4B) was fitted with a screw-cap catching jar (Figure 4E) in which organisms were captured and preserved in 70% ethanol. When pumping, bores were not purged to capture the first flow of water. Purging is not recommended for sampling stygofauna because the bore can act as a trap, often containing higher species abundances than aquifers, and so sampling the first-flow is useful for catching specimens (Hahn and Matzke 2005, Roudnew *et al.* 2012, Sorensen *et al.* 2013). Some bore holes were too narrow for pumping. Instead smaller 50 µm mesh-size nets were lowered to the bottom of the bores using a fishing rod (Figure 4F) or manual fishing reel (Figure 4G) and retrieved samples preserved in 70% ethanol. Two sizes of nets were used: a medium-sized (10.5 cm diameter) plankton net with screw cap base (Figure 4G); and a custom made small-sized (2.5 cm diameter) plankton net with a weighted screw-cap steel base (Figure 4H). Where possible, a combination of both pumps and nets were used. One site, Warlock Ponds Springs, was sampled by placing the small-sized plankton net into the spring at the opening chamber of water flux (beneath the water surface; Figure 3L) for a period of 10 minutes.

Groundwater variables

At every site water depth was measured with a water level sounder. Electrical conductivity (EC), pH and water temperature (°C) were measured with a hand-held TPS meter immediately after water was pumped to the surface. To ensure no cross-site contamination, the pump heads, nets and catching jars were thoroughly cleaned and disinfected between sites by washing in a bleach-solution.

A cluster analysis was undertaken, using Euclidean Distance to determine how similar sites were based on the following $\log(x+1)$ transformed variables: depth to the water table, EC and water temperature. pH was not transformed as it is already on a logarithmic scale. All variables were normalised prior to analysis. All multivariate analyses were conducted in PRIMER-E 7 (Clarke and Gorley 2015).

Stygofaunal sample processing

All stygofaunal samples were preserved in 70% ethanol. These samples were subsequently examined using stereo and dark field enabled microscopy at Charles Darwin University Casuarina laboratories. All specimens, or parts thereof, were located, imaged using Leica V4.12 and stored in individual vials. Taxa were identified, based on morphological characters, to the lowest taxonomic level possible. These identifications were subsequently validated by experts: Dr Stuart Halse (Bennelongia Environmental Consultants) and Dr John Short (BioAccess Australia).



Figure 4. Range of sampling methods used; A: large motorised pump; B: filtering pumped water through large plankton net; C: small motorised pump; D: filtering pumped water through large plankton net using small pump; E: catching jar; F: lowering nets down bore hole using fishing rod; G: lowering medium-sized plankton net down bore hole using manual fishing reel; H: custom-made small-sized plankton net.

DNA methods

Cytochrome oxidase I (COI) and 16s RNA gene barcoding shrimp specimens

Genetic material was extracted from small portions of 20 Atyidae specimens, either a leg or tissue from the side of the body, using a DNEasy Blood and Tissue Kit, following the manufacturer's guidelines. A 658 base-pair region of the COI mitochondrial gene was amplified using the Folmer primers (Folmer *et al.* 1994) modified with M13 tails. Primers for 16s RNA gene amplification have been described elsewhere (Page *et al.* 2007). Polymerase Chain Reactions (PCR) were performed on each extraction using 2µl of DNA, 17.5µl of GoTaq® DNA Polymerase, 14.8µl sterile water and 0.35µl of each primer. Cycle conditions for amplification were 1 min at 94 °C; 5 cycles of 1 min at 94 °C, 1.5 min at 45 °C, 1.5 min at 72 °C; 35 cycles of 1 min at 94 °C, 1 min at 50 °C, 1 min at 72 °C; and finally 4 min at 72 °C. PCR products were sequenced at Macrogen Inc. (Seoul, South Korea).

Contiguous DNA sequences were assembled using DNABaser 2.75 (DNABaser 2012) and aligned in MEGA X (Kumar *et al.* 2018) using MUSCLE software (Edgar 2004). Sequences were translated into [invertebrate] proteins to check for stop codons, frame shifts and nuclear paralogues.

Tree inference was conducted using the Neighbour-Joining method (Saitou and Nei 1987), with 500 bootstrap replicates. Included in the tree were sequences (obtained from Genbank) of known Atyidae taxa; the stygal species *Parisia unguis*, *Pycnisia raptor*, *Typhlatya mitchelli* and *Typhlatya pearsei*, and the freshwater species *Caridina steineri*, *Paratya australiensis* and *Halocaridina rubra*.

To explore overall diversity within the samples, intra and inter genetic divergences were calculated in MEGA X method (Kimura 1980), with 500 bootstrap replicates and the 'complete deletion option' (i.e. positions containing gaps and missing data were eliminated).

Water collection and preservation

A dimethyl sulfoxide-ethylenediaminetetraacetic acid-sodium chloride (DEES) stock solution was prepared and used as a preservative for samples collected in the field. Approximately 300ml of the water sample collected at each site (as described previously) was added to the DEES stock solution such that the preserved sample would contain 100ml dimethyl sulfoxide, 33.6 g di-sodium-EDTA and 50g sodium chloride.

eDNA extraction, amplification and sequence analysis

On return to the laboratory, water samples were passed through a 0.1 µm pore size polyvinylidene difluoride filters (Millipore, Bedford, MA, USA) to collect DNA. Where significant sediment was present, samples were centrifuged to avoid clogging the filters and the sediment pellets were combined with their respective water filters. A Qiagen DNeasy Powersoil Pro DNA isolation kit was used to extract total DNA from the pooled samples, according to the manufacturer's instructions. DNA was visualised by gel electrophoresis and a Nanodrop instrument was used to quantify DNA for template DNA calculations and quality scored for protein contamination.

We carried out two metabarcoding assays; the first targeted the cytochrome oxidase I gene (COI) and examined water samples for the presence of invertebrates. The second assay targeted the 16s ribosomal gene (16sRNA) of bacteria and would provide a fingerprint of all bacteria present in water samples. The forward and reverse primer for COI DNA metabarcoding were: (mICOLintF) GGWACWGGWTGAACWGTWTAYCCYCC and (jgHCO2198) TANACYTCNGGRTGNCCRAARAAYCA respectively (Leray *et al.* 2013). For bacterial analysis, the universal 515f forward and 806r reverse primer set were used, which targeted the v4 region of the 16S rRNA gene (Caporaso *et al.* 2011, 2012). Amplicons from both assays were subsequently sequenced using a MiSeq system which was provided by the Ramaciotti Centre for Genomics, University of NSW, Sydney.

A CSIRO in-house automated pipeline (GHAP) was used to manipulate the raw sequence information (<https://data.csiro.au/dap/landingpage?pid=csiro:26534>). GHAP is a hybrid of tools comprising usearch11.0.667, the Ribosomal Database Project classifier and locally written tools for demultiplexing Molecular Operational Taxonomic Unit (MOTU) trimming and produces tables of classified MOTUs. During the pipeline processing, sequence reads were merged, dereplicated, trimmed and clustered at 97% similarity to generate MOTUs. Identification of MOTUs relies on retrieved sequences being the same as those in available databases. In many instances, this is not the case and so a given taxonomic unit will be identified to its best resolution, which can result in different levels of identification across an entire data set. For example, one MOTU may simply be identified to the level of genus (e.g. *Fusarium* sp.), whereas another MOTU may not be resolved better than unidentified arthropod (phylum).

Cytochrome oxidase I gene - data handling

Two levels of data interrogation were carried out. In this first instance, data filtering as part of the DNA read pipeline generated a list of taxa, which is termed the 'full taxonomic list' of organisms detected in bore water samples. Secondly, a conservative approach was adopted to generate a 'concise taxonomic list' that comprised likely subsurface and potential stygofauna. Given the uncertain identity of many of the individuals that had been aggregated as MOTUs at 97% similarity, all MOTUs were aggregated to family level, which could be further aggregated to their respective phyla, given a broad understanding of the types of organisms present in the samples. In generating the concise list, three broad categories of taxa were recognised in the bores: 1) terrestrial organisms whose DNA was detected in bore water through either falling into, or DNA residing within groundwater (e.g. ant species); 2) those taxa that are likely to be subsurface organisms, such as those likely to be associated with soil, plants or water (e.g. fungi), 3), those taxa that are likely, or definitely recognised as being stygofauna (e.g. blind shrimp).

Cytochrome oxidase I gene - multiple sampling Bore RN034032

Given this project was a baseline pilot study, we carried out four separate sampling sessions at bore RN034032 ("Tufa bore") to provide some insight into the rigour of bore water sampling for eDNA. Two sampling sessions were on successive days in August 2019 and a further two sessions were on successive days in October 2019. The aim was to understand more about the diversity of taxa and the number of DNA reads that would be retrieved from replicate sampling of the same bore. For this approach, the number of DNA reads that were retrieved across each sample trip were standardised by converting read number to percentage of the total number of reads, then a 2 stage process was carried out where all those reads contributing less than 0.05% and 0.1% of the total were removed from the data set.

16s ribosomal RNA gene – data handling

Since the intent was to carry out broad examination of the presence of microbial communities in a selection of bores, the pipeline described above was used to generate lists of the microorganisms present in samples from bores. While many DNA barcode sequences have been described for different species of bacteria, online DNA databases for microbes at species level identification can be limited. Therefore all the MOTUs were aggregated to genus.

We used the FAPROTAX clustering routine to aggregate the microbial community according to metabolic functions (Louca *et al* 2016). Taxa identified from the sequence pipeline were aggregated based on their metabolic functions and shows the percent contribution in given samples. In this way we gained some understanding of the different physiological groups present in the ground waters.

We used a cluster analysis approach to visualise similarity of community composition between bores. Taxa lists were transformed to presence/absence on taxa and a similarity matrix was generated based on Bray-Curtis distances. The similarity matrix and cluster analysis was performed using the PRIMER e software package (Quest Research Ltd, Auckland NZ).

Results

Water quality

The depth to the water table varied from very shallow (3-8m in bores in the Mataranka region) to approximately 80m at bores on the Carpentaria Highway and the Hayfield Station (Table 2).

All of the sites sampled within the Beetaloo Sub-basin contained freshwater ($EC < 2,500 \mu S/cm$) i.e. water within the water quality guidelines for drinking water. Seven bores, spanning the entire sampling gradient from north to south, contained very fresh waters ($< 800 \mu S/cm$). The majority of the bore waters samples contained circumneutral water, with pH between 6.5 and 7.5. Eight bores in the northern sampling region were slightly basic (pH between 7.5 and 8). No bores were highly basic ($pH > 8$) or very acidic ($pH < 6.5$). Water temperatures recorded in the bores were between 30 and 40 °C. Surface water sites were slightly lower; 29°C at Fig Tree Springs and 27.9 °C at Little Roper Creek (Table 1, Table 2).

When summarising the water quality parameters in each environmental group, Group A represented surface waters, while the shallow bores tended to form one group (B) and the deeper bores another (C). Those in the Mataranka region were the shallowest and the freshest (group B). Fig Tree Spring formed an outlier, with a higher EC and lower temperature than all other sites (Table 1, Figure 5).

Table 1 Summary of water quality parameters in each environmental group

	Environmental group		
	A	B	C
Average of depth to water table (m)		9.87	69.65
Average of bore depth (m)		74.44	162.33
Average of EC ($\mu S/cm$)	2240	1096	1518
Average of pH	7.4	7.5	6.9
Average of temperature (°C)	29	34.07	35.95

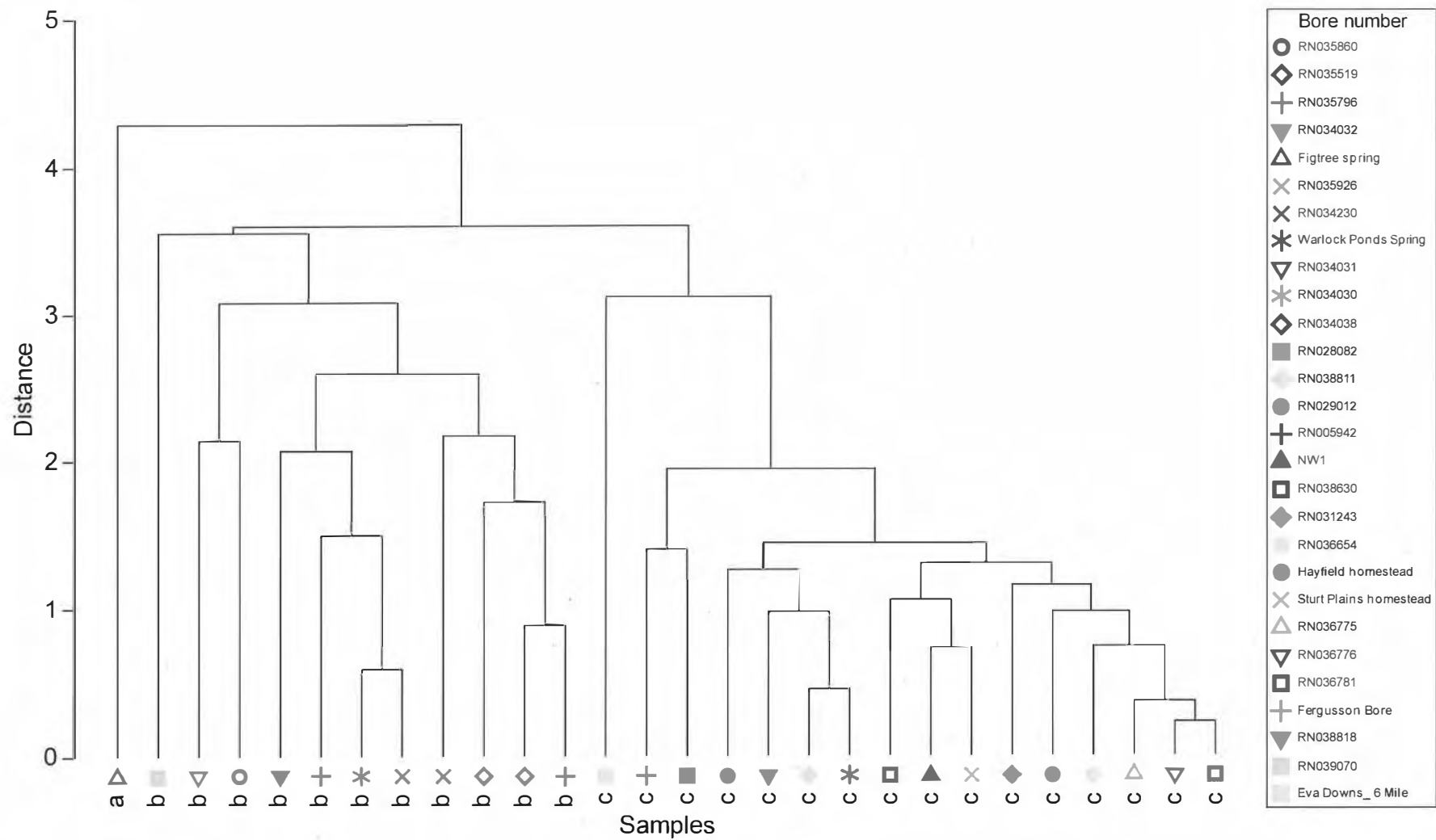


Figure 5. Cluster analysis of bores on the basis of variables measured at the time of sampling (depth to water table, total depth, electrical conductivity, pH and water temperature). Group A = Fig Tree Springs; Group B = Majority of bores in Tindal aquifer, and Group C = majority of bores in regional aquifer.

Stygofauna

Live stygobiotic animals were collected at seven of the 28 sites sampled (Figure 6, Table 3). They were predominantly crustaceans—amphipods, decapods, syncarids, copepods and ostracods—but a stygobiotic annelid worm (*Aeolosoma* sp.; Figure 8O) was also recorded. Mites (Acari) and snails (Gastropoda; Figure 8L) were also collected, but whether these are true stygal taxa is unclear (Table 4). The largest and most impressive record is of the blind atyid shrimp, *Parisia unguis*. (Figure 8A–C). This species was collected on several occasions at five sites (Tufa, Larrimah 1, Larrimah 2, Larrimah 3 and Carpentaria Highway; Table 4) ranging a distance of ~260 km (Figure 7), most notably at bore RN34032 (“Tufa bore”) where both adults and juveniles were captured in a single net haul. Captures from this bore were also rich and abundant in other taxa. One species of melitid amphipod (Figure 8D–E) was collected there on several occasions, as were three species of copepod (Figure 8K, M–N) and two species of ostracod (Figure 8J). Four species of copepod and one species of ostracod (Figure 8I) were also collected at Elliot Bore 6. The same species of ostracod (Candonidae gen. nov. 1 `BOS1374`) was also recorded at Warlock Ponds Springs (Figure 8H). A single syncarid (*Brevisomabathynella* sp.) was collected from the Carpentaria Highway bore (Figure 8G).

Stygofauna were collected by both pumping and netting methods (Table 3). For example, the syncarid recorded from the Carpentaria Highway bore was collected by the large Origin Energy motorised pump and filtering 300 L of water. The ostracods and copepods of Elliot Bore 6 were collected via 200 L filtered water from the bore tap. The three Larrimah bores were accessible by both large motorised pump and net, and both these methods yielded stygofauna. Bore RN34032 (“Tufa bore”) was accessible only by the custom-made small-sized net and it was remarkably successful, capturing animals in each netting haul. Animals captured via pump were often disarticulated, whereas netting samples yielded whole, live specimens that could be seen in the catching jar.

Legend

- Bores sampled in 2019
- Presence of stygofauna indicated by eDNA only
- Presence of stygofauna indicated by both collection of specimens and eDNA

- | | |
|--------------------------|----------------------------------|
| 1. RN035860 | 16. Aumungee NW1 |
| 2. RN035519 | 17. RN038630 |
| 3. RN034230 | 18. RN031243 |
| 4. RN035796 | 19. RN036654 |
| 5. RN034032 | 20. Hayfield Shenandoah Hstd |
| 6. Fig Tree Springs | 21. Sturt Plains Hstd |
| 7. RN035926 | 22. RN036775 |
| 8. RN034030 | 23. RN036776 |
| 9. RN034031 | 24. RN036781 |
| 10. RN034038 | 25. 7 Bore Barkly SR |
| 11. Warlock Ponds Spring | 26. Fergusson Bore nr Lake Woods |
| 12. RN029012 | 27. RN038818 |
| 13. RN038811 | 28. Eva Downs 6 Mile |
| 14. RN028082 | |
| 15. RN005942 | |

Major hydrostratigraphic units

- Ooloo Dolostone
- Tindall Limestone (Daly Basin)
- Gum Ridge Formation (Georgina Basin)
- Montijinni Limestone (Wiso Basin)
- Anthony Lagoon/ Jinduckin Formations

- ★ Main Towns
- Main Roads

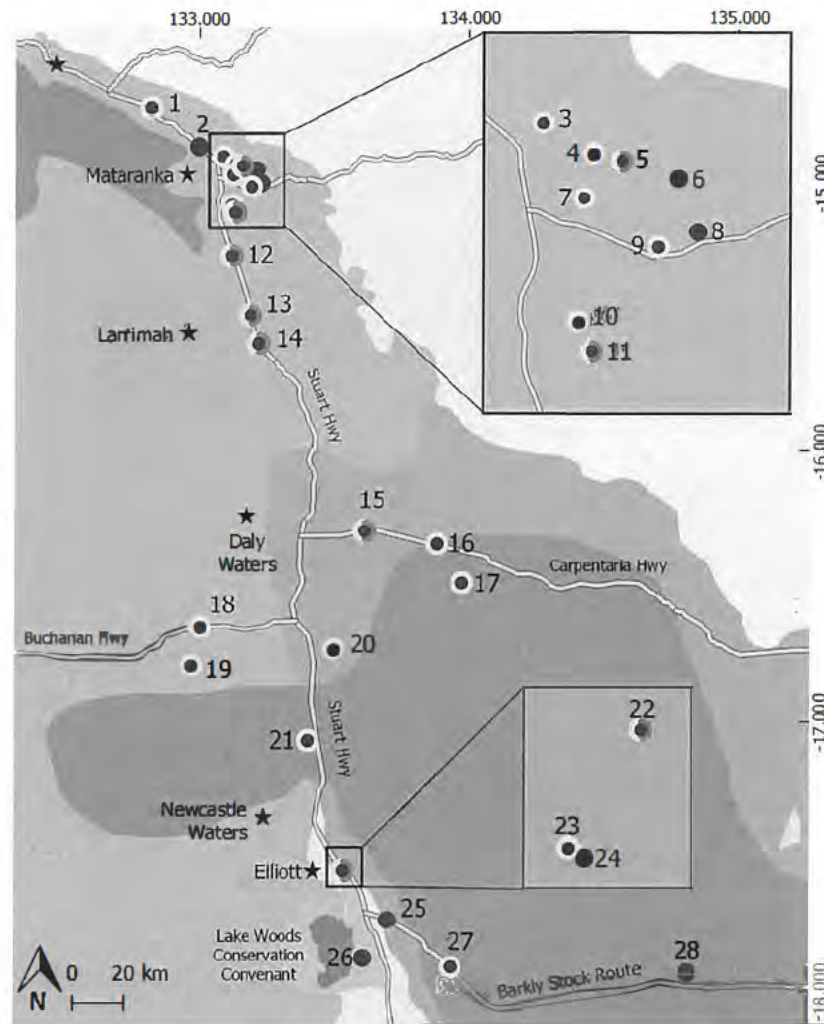


Figure 6. Study area and associated major hydrostratigraphic units of the Cambrian Limestone Aquifer, showing the bores sampled in this study and the presence of live stygofauna and stygofaunal eDNA.

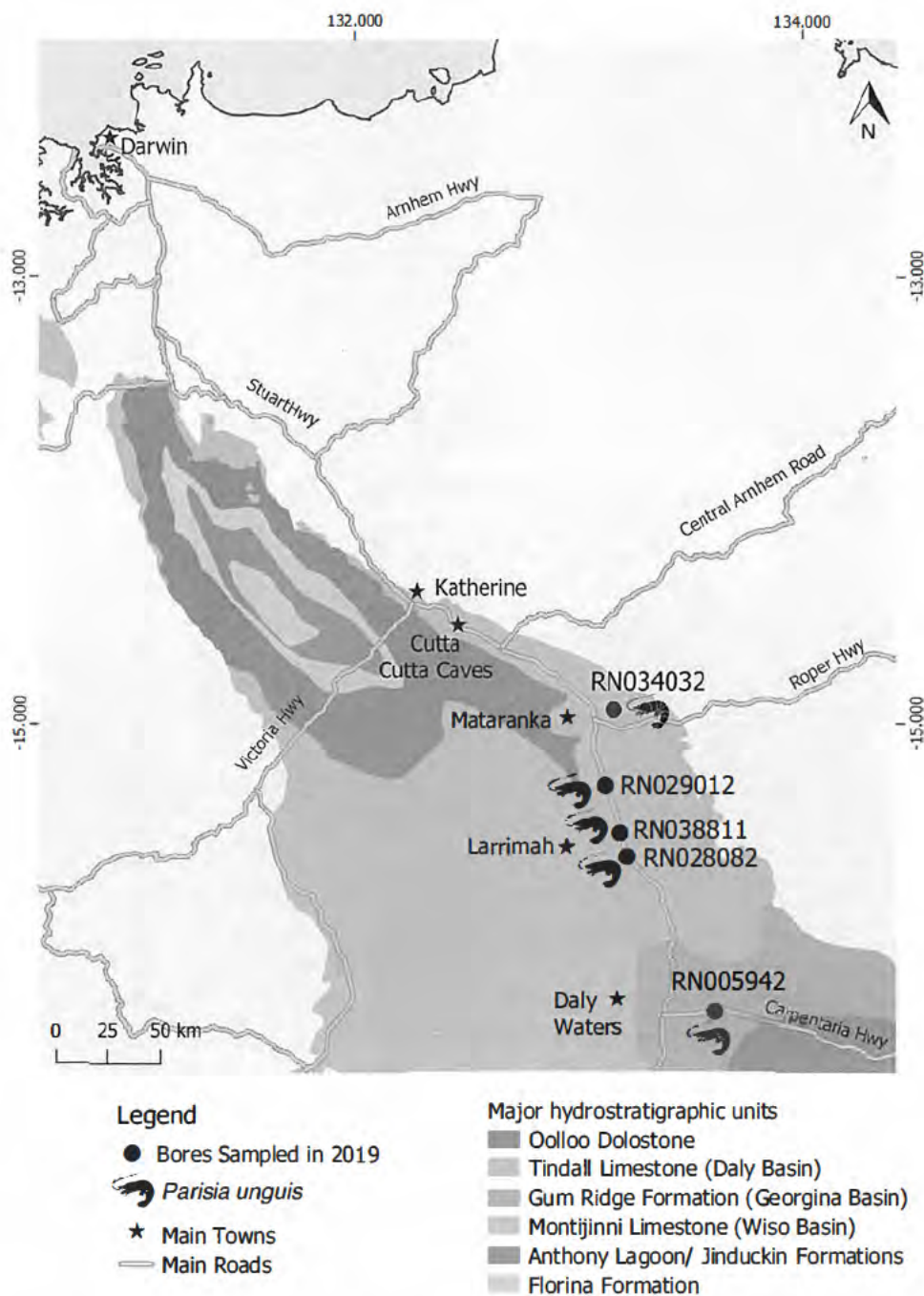


Figure 7. Study area and associated major hydrostratigraphic units of the Cambrian Limestone Aquifer, showing sites at which *Parisia unguis* (Atyidae) was collected via net and/or pump.

Table 2. Location of bores sampled in Beetaloo Sub-basin and water quality parameters.

Locality	RN bore number or Local identifier	Latitude	Longitude	Trip	Depth to water table (m)	Total depth of bore (m)	EC (uS/cm)	pH	Water Temp. (°C)
Sturt highway north of Mataranka	RN035860	-14.72421534	132.823515	August October	22	62	321	7.6	32
	RN035519	14.86837978	133.0023701	August October	8	37	646	7.1	34
Mataranka	RN035796	-14.93191424	133.1381848	August October	5	43	1603	7.6	35
	RN034032	14.938985	133.164316	August October	7	89	2200	7.2	33
	Fig Tree Springs	-14.953311	133.215349	August			2240	7.4	29
	RN035926	-14.97153782	133.1299506	August	2	34	1453	7.4	35
	RN034230	14.903608	133.0929199	October August	3	79	872	7.1	36
	RN034031	-15.01603919	133.1974872	August October					
	RN034030	-15.00225691	133.2331763	October	2	32	1610	7.6	36
	Warlock Ponds Spring	-15.11102	133.13702	October			1678	6.8	35
	RN034038	15.0837	133.1245	October	3	14	775	7.6	35
Larrimah	RN028082	15.59528	133.22612	August October	41	170	1622	6.7	35
	RN038811	-15.489792	133.195027	August October	48	244	1551	6.9	36
	RN029012	15.271076	133.125554	August	37	122	1584	6.9	34

Locality	RN bore number or Local identifier	Latitude	Longitude	Trip	Depth to water table (m)	Total depth of bore (m)	EC (uS/cm)	pH	Water Temp. (°C)
				October					
Carpentaria Highway	RN005942	-16.288648	133.619848	August	85	104	1439	7.0	36
				October					
	NW1	-16.33586	133.8858694	August	115	300	1168	7.0	36
	RN038630	-16.480383	133.97881	August	95	150	1167	7.3	36
Buchanan Highway	RN031243	-16.646094	133.00784	August	103	156	894	6.9	35
	RN036654	-16.792102	132.976861	August	75	106	1482	6.8	34
Sturt Highway north of Elliot	Hayfield Shenandoah Homestead	16.730428	133.501986	August	87		1085	6.8	37
	Sturt Plains Homestead	17.0648	133.41	August			1400	7.1	37
Elliot	RN036775	17.547601	133.541327	August	60	105	1171	6.9	
	RN036776	-17.549002	133.540465	August	61	105	1221	7.1	
	RN036781	-17.549112	133.540662	August	61	101	1197	7.0	
	Fergusson Bore	-17.8717222	133.6174611	October			732	7.3	35
Barkly Stock Route	RN038818	-17.905271	133.939069	October	49	286	1289	7.0	34
	RN039070	-17.963727	134.706142	October	40	298	1020	8.0	35
	Eva Downs_ 6 Mile	17.92678	134.814081	October	57		3490	6.6	

Table 3. Occurrence of stygofauna by direct collection and by detection via eDNA.

Locality	Bore No. or Local identifier	Date	Sample method (net or pump (L))	Stygofauna	'Stygofaunal eDNA'
Sturt Highway north of Mataranka	RN035860	August	Net		✓
		October	Net		
	RN035519	August	Net + 200		
		October			
Mataranka	RN035796	August	200		
		October	Net		✓
	RN034032	August	Net	✓	✓
		October	Net	✓	✓
	Fig Tree Springs RN035926	August	Net		
		August	Net		✓
	RN034230	October	Net		✓
		August	Net		
	RN034031	August	Net		✓
		October	Net		
	RN034030	October	Net		
		October	Net		
Warlock Ponds Spring RN034038	October	Net	✓	✓	
	October	Net		✓	
Larrimah	RN028082	August	300	✓	✓
		October	Net	✓	✓
	RN038811	August	300	✓	✓
		October	Net	✓	No sample
	RN029012	August	300	✓	✓
		October	Net	✓	✓
Carpentaria Highway	RN005942	August	300	✓	✓
		October	Net		
	NW1	August	200		✓
	RN038630	August	200		✓
Buchanan Highway	RN031243	August	300		✓
	RN036654	August	300		✓
Sturt Highway north of Elliott	Hayfield homestead	August	200		✓
	Sturt Plains Homestead	August	200		✓
Elliott	RN036775	6-August	200	✓	✓
	RN036776	6-August	200		✓
	RN036781	6-August	200		
	Fergusson Bore	5-October	Net		
Barkly Stock Route	RN038818	October	Net		✓
	RN039070	4 October	Net		
	Eva Downs_ 6 Mile	4 October	Net		

Table 4. Stygofaunal taxa sampled across bores in the Beetaloo Sub-basin, August–October 2019 (note: a list of non-stygofaunal taxa recorded from the bores is provided in Appendix 1).

Higher taxa	Species	Bore number	Trip
AMOEBOZOA			
Lobosa: Tubulinea			
	Arcellinida: Arcellidae <i>Arcella</i> spp.	Bore 7	August
ANNELIDA			
Aphanoneura			
	Aeolosomatidae <i>Aeolosoma</i> spp.	RN34032	October
ARTHROPODA			
Chelicerata: Arachnida: Acari			
	Mesostigmata <i>Mesostigmata</i> spp.	RN034031	October
Crustacea			
Malacostraca: Eumalacostraca			
	Amphipoda: Melitidae <i>Melitidae</i> unk gen 'BAM177'	RN34032	August
	<i>Melitidae</i> unk gen 'BAM177'	RN34032	October
	Decapoda: Atyidae <i>Parisia unguis</i>	RN34032	August
	<i>Parisia unguis</i>	RN34032	October
	<i>Parisia unguis</i>	RN028082	August
	<i>Parisia unguis</i>	RN028082	October
	<i>Parisia unguis</i>	RN038811	August
	<i>Parisia unguis</i>	RN038811	October
	<i>Parisia unguis</i>	RN020912	August
	<i>Parisia unguis</i>	RN020912	October
	<i>Parisia unguis</i>	RN005942	August
	Syncarida: Bathynellaceae <i>Brevisomabathynella</i> sp.	RN005942	August
Maxillopoda: Copepoda			
	Calanoida <i>Calanoida</i> sp. <i>Centropagi</i>	RN036775	August
	Cyclopoida: Cyclopidae <i>Apocyclops dengizicus</i>	RN036775	August
	Cyclopidae sp.	RN036775	August
	Eucyclopinae ngen 'BCY068'	RN34032	August
	Eucyclopinae ngen 'BCY068'	RN34032	October
	<i>Mesocyclops cuttcuttae</i>	RN34032	August
	<i>Mesocyclops</i> spp.	RN036775	August
	Harpacticoida: Ameiridae <i>Nitakra lacustris</i> s.l.	RN34032	August
	<i>Nitakra lacustris</i> s.l.	RN34032	October
Ostracoda			
	Podocopida: Candonidae <i>Candonidae</i> ngen 1 'BOS1372'	RN34032	August
	<i>Candonidae</i> ngen 1 'BOS1372'	RN34032	October
	<i>Candonidae</i> ngen 1 'BOS1374'	RN036775	August
	<i>Candonidae</i> ngen 1 'BOS1374'	Warlock Ponds Spring	October
	Gastropoda: Caenogastropoda <i>Gastropoda</i> spp.	Warlock Ponds Spring	October
	Hypsogastropoda: Bithynidae <i>Gabbia</i> spp.	RN34032	October

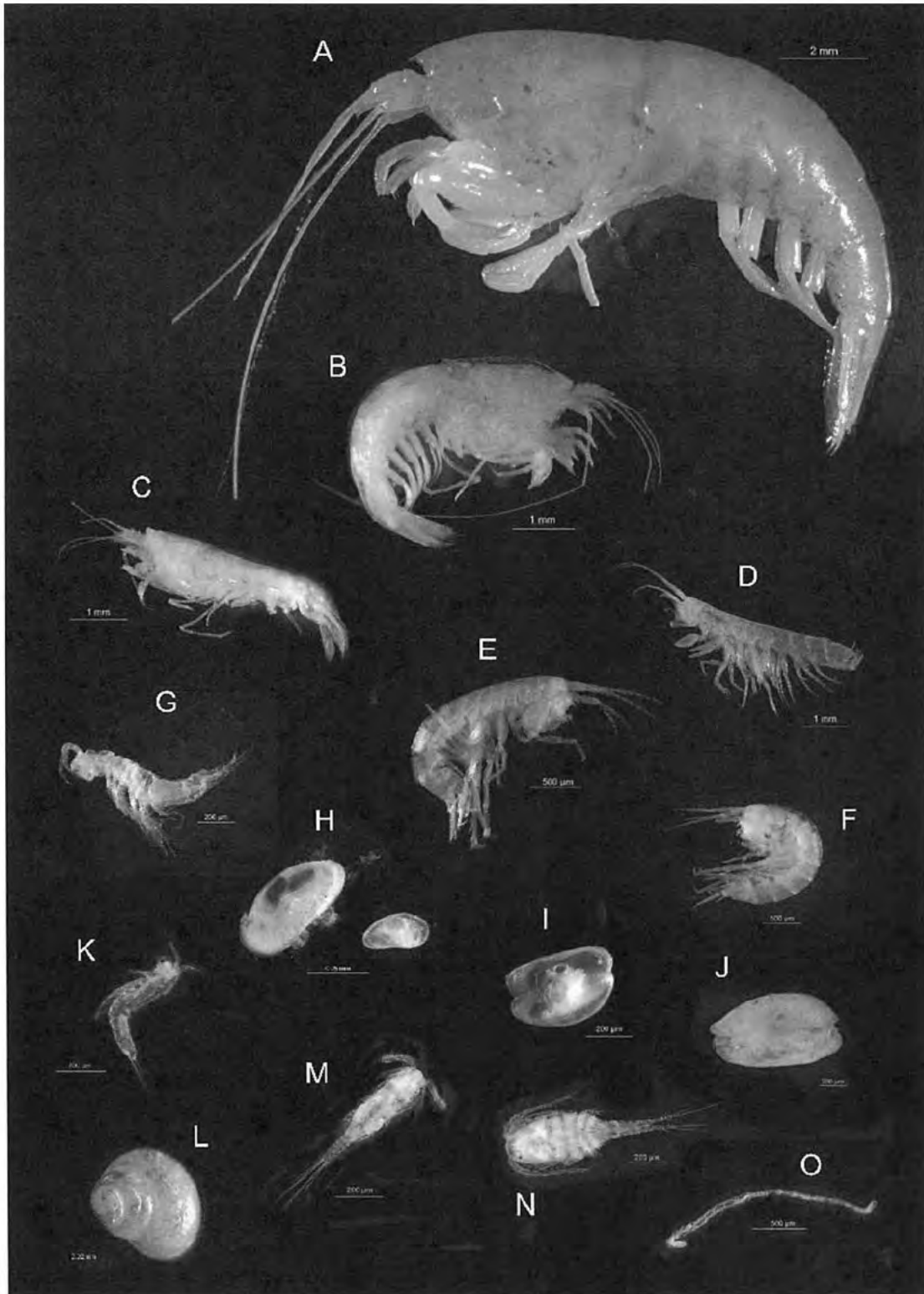


Figure 8. Subterranean fauna collected from Northern Territory aquifers; A–C: Decapoda: Atyidae: *Parisia unguis*. (RN34032); D: Amphipoda: Melitidae Melitidae unk. gen 'BAM177' (RN34032); E: Amphipoda: Melitidae unk. gen 'BAM177' (RN34032); F: Amphipoda Melitidae unk. gen 'BAM177' (RN34032); G: Syncarida: Bathynellaceae: *Brevisomabathynella* sp. (RN005947); H: Ostracoda: Podocopida: Candonidae (Warlock Ponds Spring); I: Ostracoda: Podocopida: Candonidae (RN036775); J: Ostracoda: Podocopida: Candonidae (RN34032); K: Harpacticoida: Ameiridae: *Nitokra lacustris* (RN34032); L: Gastropoda: Caenogastropoda (Warlock Ponds Spring); M: Cyclopoida: Cyclopidae (RN34032); N: Cyclopoida: Cyclopidae (RN34032); O: Annelida: Aeolosomatidae: *Aeolosoma* sp. (RN34032).

COI barcoding and 16S rRNA analysis of shrimp specimens

Thirteen of the twenty Atyidae samples submitted for molecular analysis were successfully amplified. COI barcoding revealed that they formed a well-supported (95% bootstrap support) clade with relatively low intraspecific divergence (Fig. 9a).

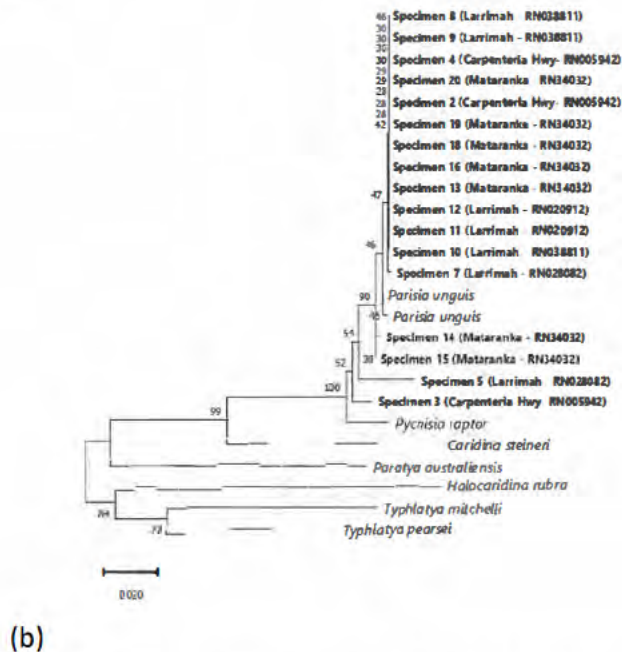
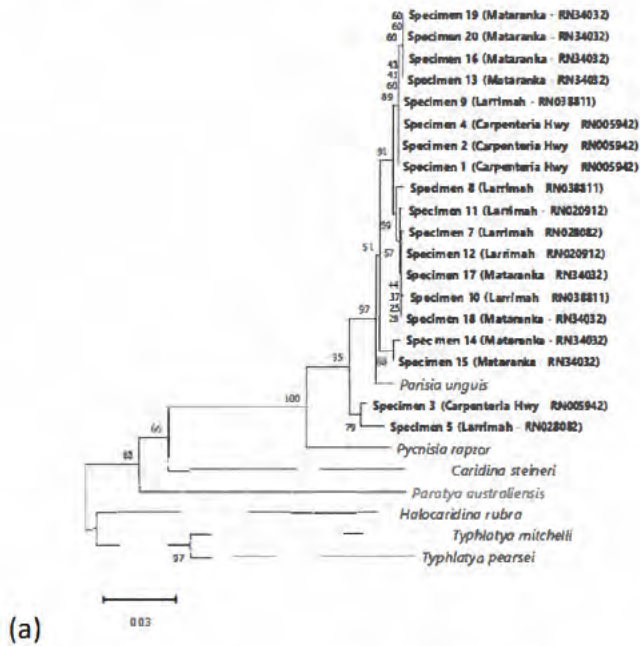


Figure 9 Dendrograms of gene sequences of 13 atyid specimens collected across five sites (RN34032 -Mataranka, RN028082 -Larrimah 1, RN038811 -Larrimah 2, RN020912 -Larrimah 3 and RN005942 -Carpenteria Highway), showing bootstrap support values at the nodes.(a) COI gene tree with selected GenBank sequences; and (b) 16S gene tree with selected GenBank sequences.

The 16S gene tree (Fig.9b) similarly indicates low divergence within the Beetaloo specimens (maximum 3.29%), and high similarity to *Parisia unguis* (minimum genetic diverge 2.35%) and *Pycnisia raptor* (min. 2.58%). Overall, these results suggest that all the specimens belong to a single species, *Parisia unguis*. Although some clade structure is evident, at least one specimen from each of the five localities is included in a strong clade. There little separation between *Pa.unguis* and *Py. raptor* but more sequences for additional specimens of *Py. raptor* are required to determine whether they are the same species.

eDNA analysis

DNA was isolated from all bore water samples, although the yield was highly variable, leading to difficulties in carrying out an extensive set of analyses for bacterial communities. In the first instance, DNA was primarily analysed using COI analysis and where failure to amplify DNA products occurred, a further attempt was carried out to amplify DNA. This process yielded amplicons from 25 bores, which were subsequently used for taxonomic analysis. Successful amplification of microbial DNA also occurred with 25 bores, noting that the bores where we retrieved microbial DNA amplification did not fully mirror those of the COI analysis. Further method modifications were trialled on samples that initially were not successful, but none of those that gave a negative result could be modified to generate microbial DNA for further analysis.

Cytochrome oxidase I - full taxonomic list

The final output from the sequence analysis pipeline generated a list comprising 115 taxa across 25 bores. The average number of taxa in samples ranged from 36 to 76, with an average of 54 per sample. The ability to identify MOTUs was generally very poor and consequently, different MOTU were resolved to differing degrees of taxonomic resolution. The presence of taxa across all bores showed a characteristic pattern whereby 7 taxa were present in all 25 samples, but extending to a 'tail', where some taxa were detected in as few as two samples (Figure 10).

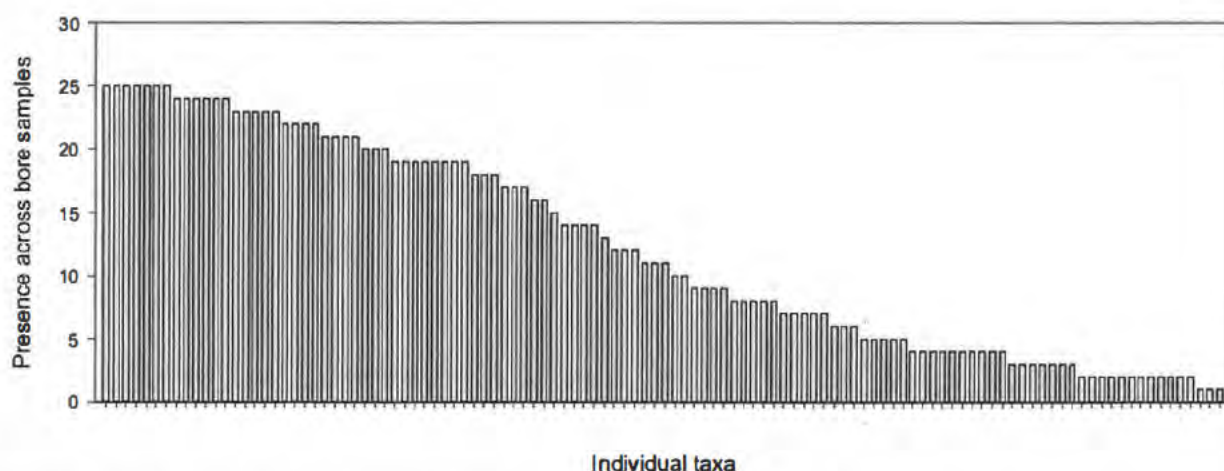


Figure 10. Histogram showing the occurrence of individual taxa across the 21 bores

The full taxonomic list showed the presence of very diverse DNA sequences and highlights the need for a conservative approach to using an eDNA approach to identify organisms in bore water Table 5. The following rationale suggests taxa can be aggregated into those that are clearly contaminants, those that are probable soil organisms that could potentially mix with groundwater and those that are true stygofauna. For example, three groups of algae were identified, which clearly do not exist in groundwater, but either reflect the presence within bores that have exposed openings, thus allowing some colonisation, or the DNA was present by surface entry. A range of fungi were detected that are likely resident in soils, or attached to plants, and therefore their DNA is easily mixed with the groundwater. Similarly, terrestrial invertebrates were detected, such as ant DNA, which supports the visual observations that terrestrial fauna were present in some samples. Rotifer and crustacean DNA were also detected, which supports the observed collection of animals through the netting procedure.

Table 5. The number of different eukaryotic taxa identified within each taxonomic level by COI metabarcoding.

	Phylum	Class	Order	Family	Genus
Animals	Annelidia	3	5	5	5
	Arthropoda	11	20	24	27
	Cnidaria	3	3	3	3
	Echinodermata	1	1	1	1
	Gastrotrichia	1	1	1	1
	Mollusca	2	2	2	2

	Nematoda	5	9	9	9
	Nemertea	2	2	2	2
	Onychophora	1	1	1	1
	Platyhelminthes	1	3	3	3
	Porifera	2	2	2	2
	Rotifera	2	4	6	7
Fungi	Ascomycota	4	7	10	12
	Basidiomycota	3	5	6	6
Protozoa	Amoebozoa	3	4	6	7
Planta	Chlorophyta	2	3	3	3
	Bacillariophyta	2	4	4	4
	Rhodophyta	3	3	3	3
Others	Oomycetes	1	3	3	4
	Stramenopiles	5	7	8	8
	Unknown eukaryotes	5	5	5	5

Cytochrome oxidase I concise taxonomic list

The concise taxonomic list comprised 45 different taxa distributed across 21 bore samples. Of these taxa, none could be identified to resolution below family. In some instances, taxa which could not be identified to level below phylum were detected (Table 6). For example, the organism identified as Amoebozoa1 could not be identified to a taxonomic resolution lower than Amoebozoa. On the other hand, Rotifer3 could be identified as a member of the order Adinetida. The inability to identify organisms to higher resolution reflect absence of suitable DNA sequences in the on-line databases.

Six Amoebozoa, 3 Annelida, 4 Arthropoda, 1 Gastrotricha, 2 Nematoda, 3 Platyhelminthes, 2 Porifera and 6 Rotifera were recognized across all the bores. The frequency of occurrence of different taxa across the bores ranged from present in all samples, to only occurring in one of the bores (Figure 11). The highest frequency of occurrence of stygofauna across the 21 bores were

Amoebazoa⁵, Arthropoda², Rotifera³ and Amoebozoa², in 100, 86, 76 and 62 % of the bores respectively. Amoebazoa⁵ could not be identified to a better taxonomic resolution. Arthropoda² was identified as a member of the class of crustaceans and as noted above, Rotifer³ was identified as a member of the order Adinetida. Amoebozoa² was identified as a member of the class Discosea.

In addition to the dominant four stygofauna, a further five taxa were present in 52 and 48% of the bores. These included two further Amoebozoa, an arthropod belonging to the class Malacostraca, one annelid (worm) of the order Haplotaxida and one member of the phylum Porifera. Some taxa were detected in only one bore. Five ascomycetes were also widely distributed across the bore waters.

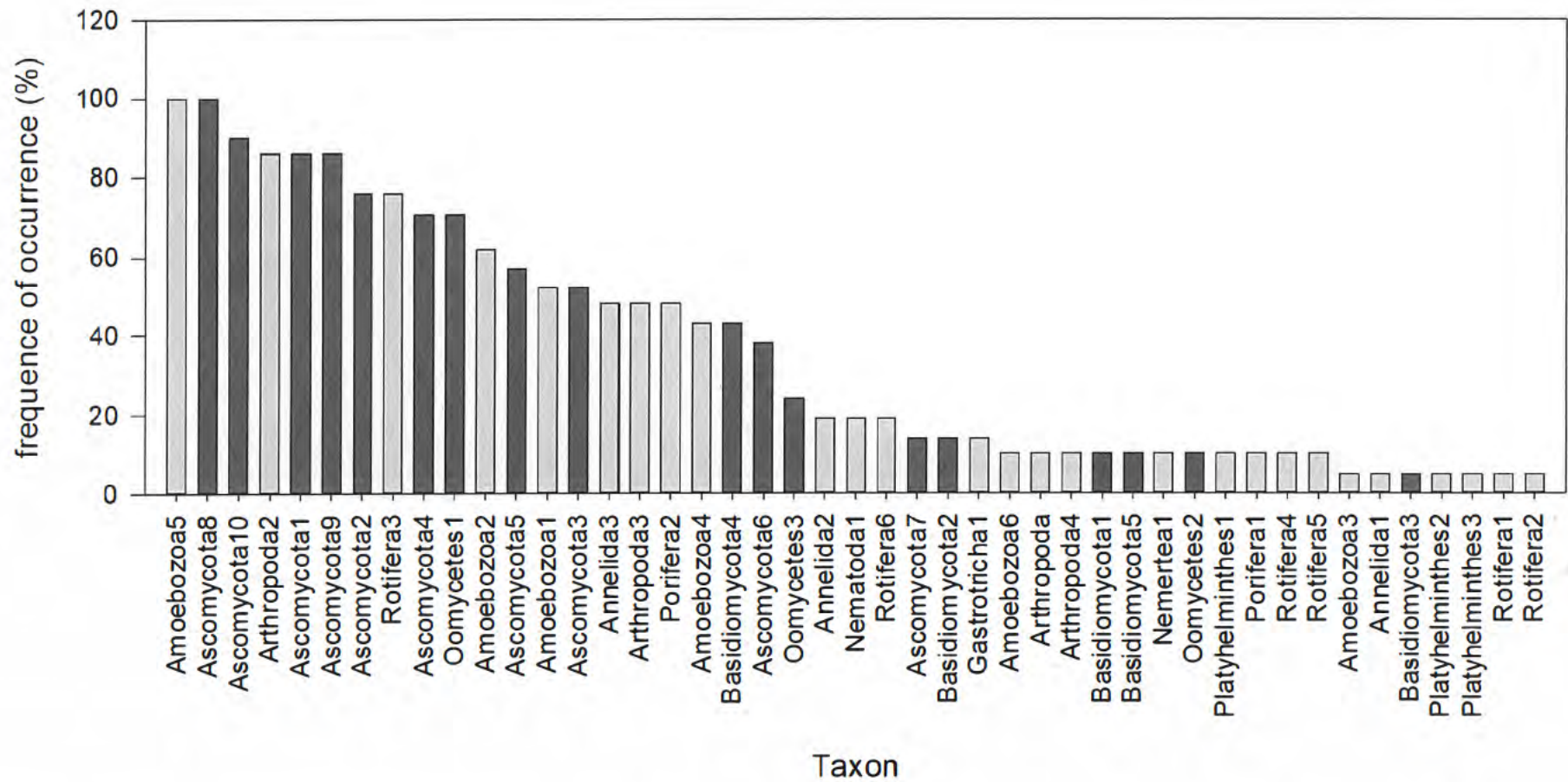


Figure 11. Frequency of occurrence of each taxon across the 21 bore where different biota were detected. Green bars indicate stygofauna. Blue bars indicate subsurface taxa (see methods for definitions)

Table 6. Presence of taxa distributed across bores where DNA could be amplified with cytochrome oxidase I primers. Ticks indicate presence and 0 indicates absence from that bore.

Lowest resolution identification	Highest resolution identification [†]	B1*	B2	B3	B4	B5	B6	B7	B8	B9	B10	B11	B12	B13	B14	B15	B16	B17	B18	B19	B20	B21	
Amoebozoa1	Amoebozoa p	0	✓	0	0	✓	✓	✓	✓	✓	0	✓	✓	0	✓	0	0	0	✓	0	✓	0	
Amoebozoa2	Discosea c	0	0	0	✓	✓	✓	0	✓	✓	0	✓	✓	✓	✓	0	0	✓	✓	0	✓	✓	
Amoebozoa4	Vannelliidae f	0	0	0	0	✓	✓	0	0	0	✓	✓	✓	✓	✓	0	0	0	✓	0	✓	0	
Amoebozoa3	Cochliopodiidae f	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	✓	0	
Amoebozoa5	Himatismenida c	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓
Amoebozoa6	Stemonitidae f	0	0	0	0	0	✓	0	0	0	0	0	0	0	0	0	0	0	0	0	✓	0	
Annelida1	Annelida	0	0	0	0	0	0	0	0	0	0	✓	0	0	0	0	0	0	0	0	0	0	0
Annelida2	Clitellata c	0	0	0	0	0	✓	0	✓	0	0	0	0	✓	0	0	0	✓	0	0	0	0	0
Annelida3	Haplotaxida o	0	0	✓	✓	✓	0	✓	✓	0	✓	✓	0	0	0	✓	0	0	0	✓	✓	0	0
Arthropoda1	Macrotrichidae f	0	0	0	0	0	0	0	0	✓	0	0	0	0	0	0	0	0	0	0	✓	0	0
Arthropoda2	Crustacea c	✓	0	✓	✓	✓	✓	✓	0	✓	✓	✓	✓	✓	✓	✓	0	✓	✓	✓	✓	✓	✓
Arthropoda3	Malacostraca c	0	✓	0	0	✓	0	✓	0	0	0	✓	✓	✓	0	0	✓	0	✓	0	✓	✓	✓
Ascomycota1	Ascomycota	✓	✓	✓	✓	✓	✓	✓	0	✓	✓	✓	✓	✓	✓	0	✓	✓	✓	0	✓	✓	✓
Arthropoda4	Maxillopoda c	0	0	0	0	0	0	0	0	0	0	0	0	✓	0	0	0	0	0	0	✓	0	0
Ascomycota2	Capnodiales o	0	✓	✓	✓	✓	✓	✓	0	✓	0	✓	✓	✓	✓	0	✓	✓	✓	0	✓	✓	✓

Ascomycota3	Dothideomycetes c	0	✓	✓	✓	✓	✓	0	0	0	0	✓	✓	✓	✓	0	0	0	✓	0	0	✓
Ascomycota4	Helotiales_o	0	0	✓	✓	✓	✓	0	0	✓	✓	✓	✓	✓	0	✓	✓	✓	✓	✓	✓	0
Ascomycota5	Leotiomycetes_o	0	✓	0	✓	✓	✓	0	0	0	0	✓	✓	✓	✓	0	✓	✓	✓	0	0	✓
Ascomycota7	Bionectriaceae f	0	✓	0	0	0	0	0	0	0	0	0	0	0	0	✓	0	0	0	0	✓	0
Ascomycota6	Cordycipitaceae_f	0	✓	0	0	✓	✓	0	0	✓	0	✓	✓	✓	0	0	0	0	✓	0	0	0
Ascomycota9	Hypocreales o	✓	✓	✓	✓	✓	✓	0	✓	✓	✓	✓	✓	✓	0	✓	✓	✓	✓	0	✓	✓
Ascomycota8	Hypocreales o	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓
Ascomycota10	Sordariomycetes f	✓	✓	✓	✓	✓	✓	0	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	0	✓	✓
Basidiomycota	Agaricaceae f	0	0	0	0	0	✓	0	0	0	0	0	0	0	0	✓	0	0	0	0	0	0
Basidiomycota2	Psathyrellaceae_f	0	0	0	✓	✓	0	0	0	0	0	✓	0	0	0	0	0	0	0	0	0	0
Basidiomycota3	Agaricomycotina c	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	✓	0
Basidiomycota4	Exobasidiomycetes c	✓	✓	0	0	✓	✓	0	0	0	0	✓	✓	0	✓	0	0	0	✓	0	✓	0
Basidiomycota5	Microstromatales o	0	0	0	0	0	0	0	0	0	0	0	✓	✓	0	0	0	0	0	0	0	0
Gastrotricha	Gastrotricha p	0	0	0	0	0	0	0	0	0	0	✓	0	✓	0	0	0	0	✓	0	0	0
Nemertea	Nemertea_p	0	0	0	0	0	✓	0	0	0	0	✓	0	0	0	0	0	0	0	0	0	0
Nematoda	Nematoda p	0	0	0	0	0	0	0	0	0	0	✓	0	0	0	0	✓	0	✓	0	✓	0
Oomycetes1	Oomycetes_p	✓	✓	✓	0	✓	✓	0	0	✓	0	✓	✓	✓	✓	0	0	✓	✓	✓	✓	✓
Oomycetes2	Pythiaceae_p	0	0	0	0	0	0	0	0	0	0	0	0	✓	0	0	0	0	✓	0	0	0

Oomycetes3	Saprolegniales o	0	0	✓	0	0	0	0	0	✓	0	0	0	0	✓	0	0	✓	0	0	✓	0
Platyhelminthes1	Catenulida o	✓	0	0	0	0	0	0	0	0	0	0	0	0	0	✓	0	0	0	0	0	0
Platyhelminthes2	Platyhelminthes p	0	0	0	0	0	0	0	0	0	0	0	✓	0	0	0	0	0	0	0	0	0
Porifera	Tricladida o	0	0	0	0	0	✓	0	0	0	0	0	0	0	0	0	0	0	0	0	✓	0
Platyhelminthes3	Demospongiae c	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	✓	0
Porifera2	Porifera p	0	0	0	✓	0	✓	0	0	✓	0	✓	✓	✓	✓	0	✓	0	✓	0	✓	0
Rotifera	Adinetida o	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	✓	0	0	0
Rotifera3	Adinetida o	✓	✓	✓	✓	✓	✓	✓	✓	0	✓	✓	✓	✓	0	✓	0	0	✓	✓	✓	0
Rotifera2	Bdelloidea c	0	0	0	0	0	0	0	0	0	0	0	0	0	✓	0	0	0	0	0	0	0
Rotifera4	Monogononta f	0	0	0	0	0	0	0	0	0	0	0	0	0	0	✓	0	0	✓	0	0	0
Rotifera6	Ploima o	0	0	0	0	✓	0	0	✓	0	0	✓	0	0	0	✓	0	0	0	0	0	0
Rotifera5	Ploima o	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	✓	✓

‡ p=phylum, c=class, o=order, f=family. * bore designation.

B1: Amungee NW1, B2: RN038818, B3: RN031243, B4: RN036654, B5: RN005942, B6: RN034230, B7: RN036775, B8: RN036776, B9: RN034038,

B10: Hayfield Shenandoah Homestead, B11: RN028082, B12: RN038811, B13: RN029012, B14: RN035796, B15: RN035926, B16: RN034031

B17: RN038630, B18: RN035860, B19: Stuart Plains-Homestead, B20: RN034032, B21: Warlock Ponds Spring

Return sampling – Bore RN034302

Multiple field trips repeatedly detected organisms in bore RN034032 (Table 7). The degree of diversity differed between samples and reflected some taxa being responsible for a high proportion of the DNA reads. For example, in field trip 1, removing taxa that contributed less than 0.1% of the total reads reduced the overall number of taxa from 68 to 36, indicating a large number of taxa were making only a very small contribution to the overall read number in each sample. This was particularly notable in sample trip 3, where removing taxa contributing less than 0.1% of the total reads reduced the number of taxa from 64 to 4. In the latter case, those 4 taxa, (and their percent contribution) were: Unidentifiable crustacean1 (77.7%), Malacostraca, a crustacean (3.6%), Maxillopoda, a crustacean (5.3%) and a Hymenopteran (ant 12.6%). The latter is notable as a terrestrial contaminant in some bores

Table 7. Taxa present in Bore RN034302 on successive sample trips.

Taxa list	Number of taxa present after filtering			
	Trip 1	Trip 2	Trip 3	Trip 4
Full data set	68	73	64	76
Taxa contributing less than 0.05% of reads removed	38	32	8	25
Taxa contributing less than 0.1% of reads removed	36	26	4	21

Microbial community analysis – community composition

Microbial amplicons were successfully obtained from 20 bore samples. The low levels of DNA that were extracted across samples prevented further exploration of samples where initial failure of DNA amplification occurred. Taxa lists identified an average of 236 genera (range 69 – 362) across the samples. Removing taxa that contributed less than 0.1% of the reads to the overall taxa list, (i.e. rare taxa), reduced the average number of genera to 70 per sample (range 122 – 8) and removing taxa that made up less than 0.5% of the reads reduced the average number of taxa in

samples to 27. This demonstrates that many of the taxa present contributed only a very small contribution to the overall community composition.

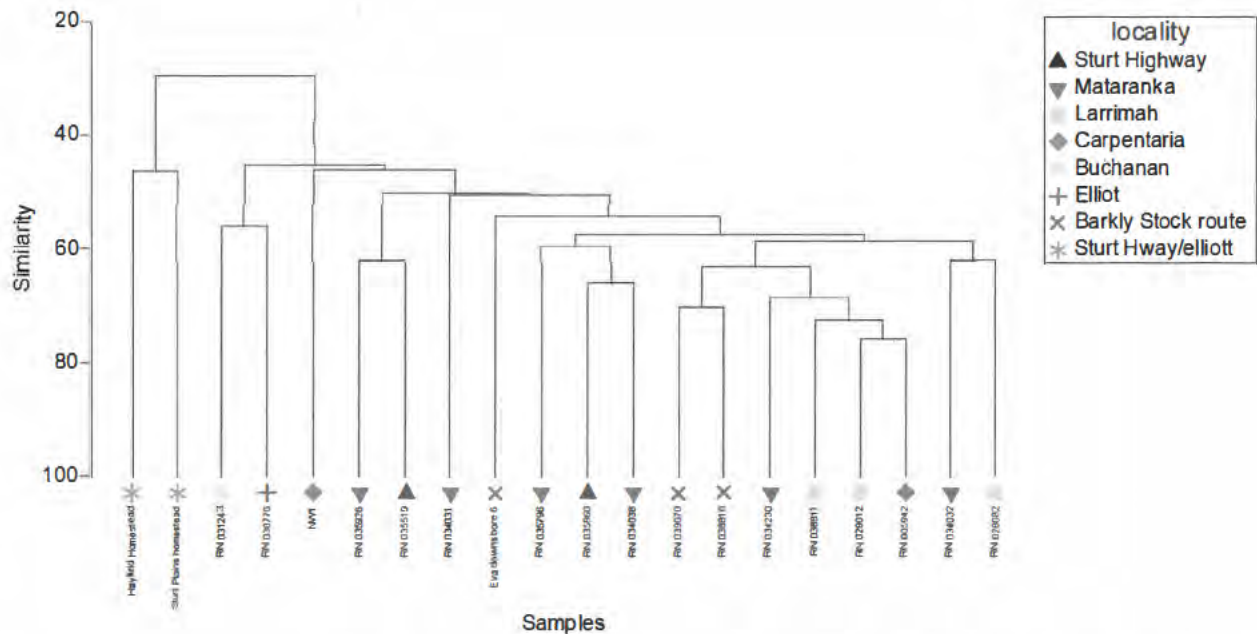


Figure 12. Cluster analysis of the microbial community in bores. Symbols indicate the general locality of each bore as defined in Table 3.

There was only a moderate degree of similarity among the microbial communities present in the bores (Figure 12). A notable grouping occurred with the Hayfield Shenandoah and Sturt Plains homestead samples. While these clustered at approximately 45% similarity with each other, microbial communities were very distinct from all the other bore samples. Two of the bores in the Larrimah locality showed approximately 70% similarity in its composition, the third bore clustered at approximately 50% similarity. A similar result was seen with the bores sampled along the Barkly stock route, with bores RN 39070 and RN 038818 clustering at approximately 70% similarity, and the Eva downs bore #6 some 50% similar to the other two bores sampled along the Barkly stock route (green crosses, Figure 12).

Microbial community analysis – metabolic group analysis

There was a reasonably even spread of aerobic heterotrophic organisms across all the bores, with the heterotrophs comprising between 15 and 20% of the community in most bores (Figure 13). There were three exceptions, (RN 035796, RN 031243 and RN 036776) where the aerobic heterotrophs comprised up to 40% of the community. The presence of microbes that carry out different components of the nitrogen cycle were notable in bores from the more northerly region

of the sampling program. Nitrifying bacteria comprised between 2 and 20% of the population in bores just north and in the vicinity of Mataranka. Denitrifying organism were a dominant group across many of the bores, notably those centred near Mataranka. Sulfate-reducing organisms made up between 3 and 10% of the community in several bores, with their contribution up to 42% in bore NM1. On the other hand, sulfate reducing bacteria were barely detectable in bores RN 03776, Hayfield Shenandoah homestead and Eva Downs bore #6. Methanogenic microorganisms made a major contribution (~20%) of the community in bores RN 035860 and 034038. Microorganisms that carry out fermentative process were well distributed across bores, with notable presence from the Sturt Plains and Hayfield Shenandoah homesteads, and Eva Downs bore #6.

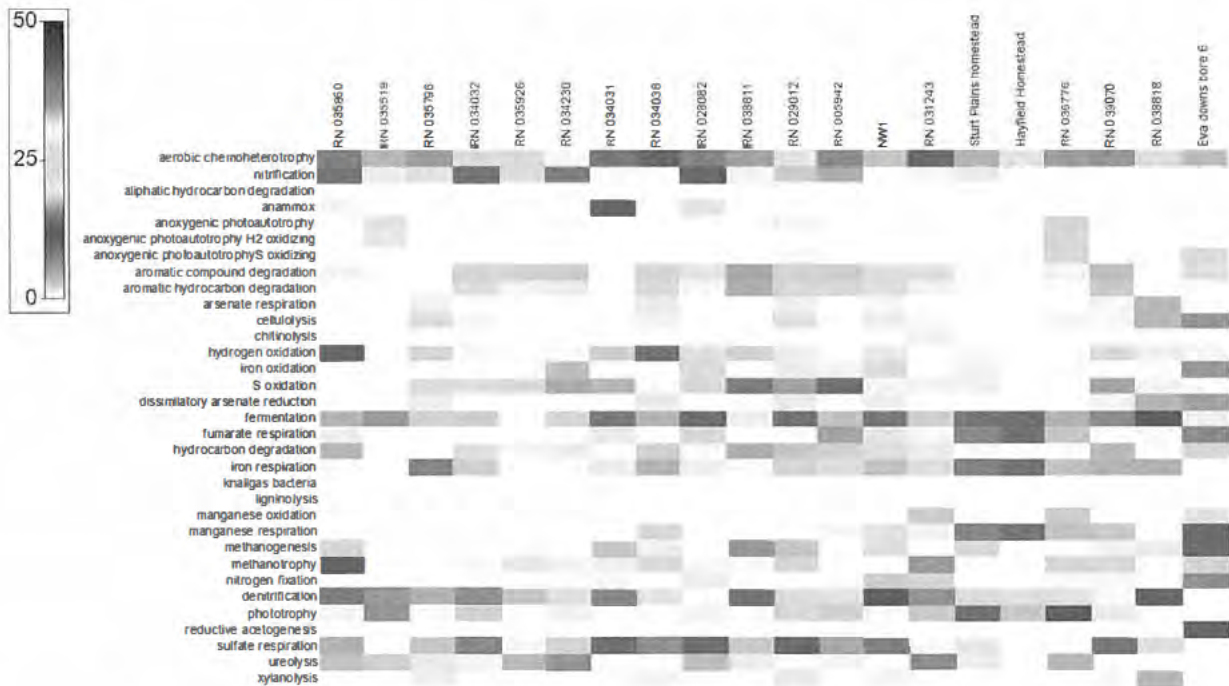


Figure 13. Shade plots showing the contribution of microbial physiological groups to the overall microbial community structure. The graded colour scale shows the percent contribution, based on taxa identified as part of the sequence analysis pipeline.

Discussion

This pilot study has provided the first description of stygofauna in a region of Australia where almost no previous sampling of the subterranean aquatic biota had been undertaken. Accordingly, this study provides baseline information which represents a first step in the conservation of the biodiversity and ecological integrity of the subterranean groundwater dependent ecosystems (GDEs) of the Beetaloo Sub-basin. Knowledge of the subterranean fauna is important because the need to protect subterranean GDEs has been recognised at the federal level (Commonwealth of Australia 1997) for over 20 years.

Diversity and distribution of stygofauna

Across the Beetaloo Sub-basin there were a variety of bore diameters and structures on the bores. For example, groundwater in bores in the Mataranka region was generally accessed via a 5 cm pipe, whereas other bores had fixed pumps. Due to the variation in bores, multiple sampling methods were used to collect stygofauna. The efficiency of the different sampling methods was highly variable, which limits the capacity to make quantitative comparisons between different bores. However, it is clear that all Beetaloo Sub-basin stygofaunal communities are dominated by the arthropod group Crustacea. This includes the shrimps, amphipods, ostracods, copepods and syncarids that were recorded as part of this study (Table 4). Crustacean eDNA was recorded from most bores indicating that, at a broad taxonomic level, crustaceans are distributed throughout the Beetaloo Sub-basin. This corresponds approximately to the distribution of crustaceans that were physically collected with nets. For example, Crustacea (Atyidae) were found in the north of the Sub-basin at Mataranka and Crustacea (*Mesocyclops* spp.) were recorded as far south as Elliott.

The dominance of Crustacea in the taxa recorded here (Table 4) is in accordance with descriptions of stygofaunal assemblages in other regions of Australia (Hose *et al.* 2015a). The most well-described Australian stygofauna are those of Western Australian aquifers, particularly the aquifers of the mineral-rich Pilbara and Yilgarn regions. However, although the general pattern of crustacean dominance holds for the Beetaloo stygofauna, the genera and species recorded show little affinity with the stygofauna recorded from Western Australian aquifers. Dr Stuart Halse (Bennelongia Environmental Consultants) has worked extensively on WA stygofauna. Dr Halse examined all the specimens collected in this study and noted that there are elements of the Beetaloo fauna that do not occur in Western Australian, and these are likely new genera and

species. These include species of Amphipoda: Melitidae, Ostracoda: Candonidae, Cyclopidae: Eucyclopininae, Syncarida: Bathynellaceae and Decapoda: Atyidae (Table 4).

The dominance of Crustacea supports studies undertaken elsewhere in the world that have reported that Crustacea contribute to about 70% of stygofaunal species richness and that Copepoda, Amphipoda and Ostracoda, collectively outnumber all remaining invertebrate groups living in groundwater environments (Stoch and Galassi 2010). The Syncarida occur almost exclusively in groundwaters (i.e. are absent from surface waters) suggesting that their evolutionary history started in subterranean environments. The success of the Crustacea in colonising subterranean aquatic environments is widely attributed to a lack of competition from aquatic insects which are dependent on air for breathing or reproduction (Stoch and Galassi 2010).

Atyidae

A notable feature of the atyid specimens recorded in our study are their large size relative to all other stygofauna collected (Figure 8) and their predatory behaviour, which places them at the top of the truncated foodweb that is characteristic of subterranean aquatic environments (Gibert and Deharveng 2002). Morphologically the atyid specimens closely resemble *Parisia unguis*, a subterranean species recorded from the Cutta Cutta caves near Katherine, Northern Territory (Williams 1964). In agreement with the morphological assessment, COI barcoding and 16S rRNA gene sequencing indicates that all our atyid specimens comprise a single species, *Parisia unguis*, ranging across a geographic distance of ~300 km (from the Cutta Cutta caves down to the Carpentaria Highway). Low genetic divergence (maximum 3.29% and 3.9% for 16S RNA gene and COI respectively) among specimens suggest groundwater connectivity in recent times.

Three species of blind, colourless atyids have been described from the Cutta Cutta caves, *Parisia unguis*, *Parisia gracilis* (Williams 1964) and *Pycnisia raptor* (Bruce 1992). The relationship between these three species requires further assessment. Genetic studies have shown that *Pa. unguis* and *Py. raptor* form a strong clade (Page *et al.* 2007, Page *et al.* 2008), suggesting that *Py. raptor* may be a synonym of *Pa. unguis*.

Amphipoda

Amphipods in the family Melitidae, representing a new genus and species, were recorded from one bore, RN34032, and were potentially present in other bores as represented by the higher

grouping of Arthropoda in the eDNA results. Most members of the Melitidae are considered to be of ancient marine origin and restricted to regions of Australia that were inundated by sea during the Cretaceous (Bradbury and Williams 1999). Amphipods in the families Melitidae, Paramelitidae and Neoniphargidae have been recorded from groundwater in the Pilbara region, Western Australia (Finston *et al.* 2008).

Eberhard (2003) recorded a new, undescribed amphipod species, *Chillagoe* sp., from the karst groundwater system of the Nowranie Caves, near Camooweal, Queensland and noted that it was morphologically similar to, but distinct from, *Chillagoe thea*, a species that inhabits the Chillagoe caves, 600 km to the east of Camooweal (Barnard and Williams 1995, Bradbury and Williams 1997a) and considered that it was probably endemic to the cave system. Eberhard (2003) also noted that the distributions of the Camooweal and Chillagoe species extends the northern distribution of any Australia aquatic (surface or subterranean) amphipods. He considered that only subterranean waters in these regions provide the low temperatures and more stable environmental conditions required to support amphipod populations (Bradbury and Williams 1997b). The melitid species recorded in our study at bore RN34032, in the Mataranka region, represents a further northward extension of the Australian freshwater amphipod fauna.

Ostracoda

Ostracods representing a new genus and species of Candonidae were collected from two locations in the Mataranka region (bore RN34032 in Elosey National Park and a spring on Warlock Ponds Station). Karanovic and McKay (2010) noted that subterranean ostracods mostly belong to the subfamily Candoninae. A total of 84 Candoninae species have been described from Pilbara aquifers, five Candoninae species have been described from Murchison aquifers, three from the Kimberley region, one from Queensland and two from the Perth basin, and almost all genera are considered to be Australian endemics (Karanovic and McKay 2010). Further work is now needed to determine describe the genus and species recorded here.

Copepoda

In accordance with studies undertaken elsewhere in Australia and overseas, cyclopoid copepods were the most numerically diverse species group within the samples we collected. Two described species were recorded, *Apocyclops dengizicus* and *Mesocyclops cuttcuttae* (Table 4). The former species has been recorded from locations in Western Australia (Atlas of Living Australia-

<https://bie.ala.org.au/species>) and the latter has been recorded from the Cutta Cutta caves near Katherine (Dumont and Maas 1983). New genera and species also appear to be present but little further comment can be made on these species without further sampling and more detailed taxonomic investigation. Karanovic (2006) described a rich and interesting subterranean copepod fauna (41 species and subspecies) from the Pilbara region of Western Australia and it seems likely that multiple new species are present in the Beetaloo Sub-basin. The single harpacticoid species recorded, *Nitokra lacustris* s.l. also needs more work to determine its distribution and taxonomic status.

Syncarida

We collected a single syncarid specimen (Bathynellaceae: *Brevisomabathynella* sp.) from a bore (RN005947) along the Carpentaria Highway. It represents a new genus record for the NT; the 12 species comprising *Brevisomabathynella* are all recorded from WA (Cho *et al.* 2006b, Cho and Humphreys 2010). Only one syncarid, *Atopobathynella readi* (Parabathynellidae), has been described from the NT from a bore in the arid Ngalia Basin region (Cho *et al.* 2006a). Parabathynellidae (possibly *Atopobathynella*) and Bathynellidae syncarids have also been collected in northern parts of the NT, from the Magela Creek (Chandler *et al.* 2017; Lisa Chandler pers. com.). Coupled with these northern and southern records, our specimen indicates that syncarids are likely present more broadly throughout the NT and in greater diversity than is currently realised. Australian syncarids are thought to be quite diverse but with highly restricted distributions (Cho *et al.* 2005, Cho *et al.* 2006a, Abrams *et al.* 2013). If this is also the case in the NT, then local species would be of high conservation value.

Aquifer Connectivity

Two studies have been undertaken using environmental tracers to develop an understanding of groundwater flow, and subterranean and surface water connectivity, in the Cambrian Limestone Aquifer (CLA) in the Beetaloo Sub basin (NT). The first (Suckow *et al.* 2018) highlighted the complexity of the groundwater flow system based on the sampling of eight bores in 2017. They raised questions about recharge mechanisms, recharge location, and possible flow from deeper aquifers along fractures, that could not be answered from the limited number of bores sampled in their study. A subsequent study (Deslandes *et al.* 2019) sampled 25 bores between Mataranka and Daly Waters in 2018. They used tracers that included major ions, Rare Earth Elements (REE), the

stable isotopes of water, tritium (^3H), chlorofluorocarbons (CFC-11, CFC-12, CFC-113), sulfur hexafluoride (SF_6), halon-1301 (H1301), radiocarbon (^{14}C & ^{13}C), and noble gases (He, Ne, Ar, Kr, Xe, ^{222}Rn). Their study confirmed the results of the Suckow et al. (2018). They found counter-intuitive tracer patterns and internal contradictions between different tracer types, indicating contributions of both modern water and old water. They concluded that the whole area of the CLA must be regarded as a potential recharge area. They noted that, together with the very high flow velocities typical for a karst aquifer, the whole area of the CLA is at potential risk to possible contamination from surface spills from any source.

The results of our study, primarily the widespread occurrence of the distribution of the blind, colourless shrimp (*Atyidae*), *Parisia unguis*, support the tracer study results that found that the CLA is highly connected. Further work is required to quantify the risk of contamination impacts on stygofauna from possible spill events that takes into account migration pathways and processes including adsorption, dilution and microbial metabolism in both soil and aquifer as well as the high connectivity in groundwater systems.

eDNA analysis – COI analysis

eDNA analysis is rapidly emerging as a useful approach to examining a range of different types of samples for the presence of organisms, particularly in situations where sampling and/or collecting organisms themselves is difficult. In recent times, eDNA has been a target for stygofauna in bore and aquifer waters (Korbel *et al.* 2017, Gibson *et al.* 2019).

While the ability to detect organisms through the presence of their DNA in samples, as compared to their visible detection, is extremely powerful, the method does rely on relevant DNA barcodes for organisms to be present in databases. During the eDNA identification process, eDNA computational analyses compare the DNA sequences present in samples against DNA barcode sequences that are present in public databases. In this way, the identity of DNA can be established. For this process to occur, animals must have been retrieved at some stage from a water sample, their identity established by classical taxonomic identification methods and finally, appropriate DNA barcodes determined for each animal. At present, DNA sequences from the cytochrome oxidase I gene (COI) are a commonly used barcode for invertebrates, although barcodes have been generated for some alternative genes, such as the 16s ribosomal RNA gene. Three recognized three categories of organisms from the eDNA analysis were recognized in this study: terrestrial organisms, subsurface, but likely connected with soils of plant roots (fungi) and

those that are likely resident in the groundwater. The approach was particularly powerful in detecting fauna in the groundwater, particularly small or fragile taxa such as amoebozoans, worms and small arthropods. Taxa were not detected in all our bore water samples which supported the notion that stygofauna were not present across all samples. One bore where animals were collected on multiple occasions was also a source of stygofaunal DNA on each occasion. While this work was a first pilot study, it does show the usefulness of the eDNA approach to sampling groundwater for biota.

Identification of DNA sequences

Given the very limited number of studies on stygofauna, it is therefore not surprising that fine-scale identification of organisms could not be achieved in this study as there are insufficient number of barcodes in DNA databases that matched the DNA we analysed in the study. In further sampling programs, a combined approach that identifies and names all organisms retrieved from bore water samples, as well as carrying out DNA barcoding on specimens is required; the approach we have carried out on the blind shrimp collected during this study. In many instances, the organisms will represent new species, or even genera. Those organisms require a formal description by relevant taxonomic experts.

Current barcoding analyses allow for identification at a coarser taxonomic level. For example, samples from bore RN34032 regularly contained unidentified crustacean DNA. The blind shrimp is now identified as *Parisia unguis*. Once these DNA barcodes are submitted to online databases, reanalysing our current eDNA results against the newly deposited DNA sequences will be able to assign a species identity to the eDNA present in the bore. The shrimp barcode will also be available for future studies examining bore water samples.

eDNA analysis – microbial analysis

Given the small number of bores that yielded DNA that was successfully used as template DNA for microbial analysis, it is not possible to make strong inferences about the overall microbial communities in the bore waters. In general terms, there were diverse microbial communities in the different bores and more widespread sampling, on multiple occasions is required to understand the overall importance of the microbial communities.

First analysis shows that dissolved organic carbon (DOC) could be as high as 4 mg/L in bore water (Table A1). A constant supply of readily bioavailable DOC could support a subsurface microbial community. Analysis of the metabolic groups demonstrated the dominance of aerobic

heterotrophic bacteria, which could use such DOC as a source of carbon and energy. Nitrate concentrations can be very high in bore water samples, providing a source of nitrogen for growth of microbes; phosphorus is below 0.01 mg/L, and may be a limiting factor for growth (Table A1).

The high levels of nitrate clearly support denitrifying bacteria, which were a major proportion of the microbial community. Similarly, the levels of sulfate detected in bores explains the prevalence of sulfate-reducing bacteria in bore water samples.

The source of microbial communities in water samples is an important consideration. Bacteria readily form communities on hard surfaces and those communities can grow to form complex communities of organisms that are referred to as biofilms. The ubiquity of biofilms has meant such communities have been very widely studied (e.g. Branda *et al.* 2005). Components of the biofilm closest to the surface may become depleted in oxygen and colonised by anaerobic bacteria. Sulfate-reducing bacteria are strictly anaerobic bacteria and their role in corrosion of steel while inhabiting the anaerobic zones of biofilms is well documented (Lee *et al.* 1995). Sulfate-reducing bacteria were found widely across the sampling sites in this study and their presence within samples is likely to have been a consequence of materials originally derived from biofilms on the surface casings of the bores. Further work on the microbial communities will need to consider whether organisms retrieved from aquifer samples are resident in the aquifer itself, or through dislodging surface biofilms.

Subterranean biota – a structured foodweb

This pilot scale project demonstrated the presence of stygofaunal communities in bores within the Beetaloo Sub basin. The presence of diverse microbial communities within bores was demonstrated. Blind shrimp are predators, living on other biota present in the bore samples. Given that blind shrimps were isolated on multiple occasions from some bores, their presence provides evidence of a structured food web. It can be presumed that a supply of a basal resource is available to support the numbers of shrimp that were obtained with relative ease. The primary source of carbon for the food web is not known. Either DOC leached through the soil profile, or exudates directly from tree roots that may have close connectivity with aquifers are potential sources of carbon, which would supply DOC for growth of microbes and then become the food source for higher consumers. Further study on the foodweb structure is warranted in order to gain insights into how it would respond to perturbations or potential contamination.

Further work

- Develop a better understanding of the distribution and diversity of stygofauna and aquatic microbial assemblages in the Beetaloo Sub basin by extensive sampling of bores across north south and east-west hydrogeological gradients.
- Develop standard sampling methods to encompass the variety of bore types. Deep bores require large and powerful pumps to access water, but the high flow rates associated with large powerful pumps can damage stygofaunal specimens, making their identification difficult. High flowrate pumps also require certification to operate. Pumps and handheld nets are useful where the groundwater is close to the surface.
- Develop a standard eDNA protocol to ensure that stygofauna can be recorded from bores where the collection of specimens is not possible (i.e. bores with pumps attached or deep bores where handheld nets cannot access the water table).
- Commission taxonomic work and generate DNA barcodes from fauna present in bore water, to ensure that new stygofaunal species are formally described and that their identity can be confirmed by eDNA in subsequent monitoring programs.
- Develop assessment and monitoring protocols and environmental conditions to ensure that stygofauna and subterranean aquatic microbial assemblages are described and protected where on-shore gas extraction may have an impact on groundwater quality and quantity in the Beetaloo Sub basin.

Appendix 1

Table A1. Dissolved organic carbon, nutrients and sulfate concentrations in selected bores. Unpublished data used by permission from Midgely et al GISERA project “Environmental monitoring and microbial degradation of on shore shale gas activity chemicals and fluids in the Northern Territory”.

	Motel bore	RN028082	RN038811	RN029012	RN005942	Amungee NW1	RN038630	RN031243	RN036654	Sturt Plains Homestead	Heyfield/ Shenandoah	RN036775	RN038818
DOC (mg/L)	<1	1	2	2	1	<1	1	2	2	4	2	4	<1
NH ₄ ⁺ -N (mg/L)	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
NO ₃ - N (mg/L)	2.06	0.57	0.82	1.00	0.01	<0.01	<0.01	0.54	2.52	2.37	0.79	2.77	<0.01
Total Nitrogen N (mg/L)	2.2	0.6	0.8	1.0	<0.1	<0.1	<0.1	0.5	2.7	2.6	0.8	3.0	<0.1
Total Phosphorus P (mg/L)	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	0.03
Reactive Phosphorus (mg/L)	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01
SO ₄ ²⁻ (mg/L)	247	151	173	186	109	144	146	27	153	143	70	44	269

Table A2. Non stygofaunal taxa sampled across bores in the Beetaloo Sub-basin, 2019.

Higher taxa	species	Bore number
<u>Arthropoda</u>		
<u>Chelicerata: Arachnida: Acari</u>		
Mesostigmata	Mesostigmata sp.	RN034031
Sarcoptiformes	Oribatida sp. `BAC006`	RN033185
	Oribatida sp. `BAC006`	RN028964
	Oribatida sp. `BAC007`	RN34038
	Oribatida sp. `BAC007`	RN028964
Trombidiformes	Trombidiformes sp.	RN038811
	Acari sp. 1	RN020912
	Acari sp. 2	RN028964
Insecta: Hemiptera	Coccoidea sp.	RN036389

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1300 363 400
+61 3 9545 2176
csiroenquiries@csiro.au
csiro.au

For further information

CSIRO Land and Water
Gavin Rees

s22

s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Tuesday, 8 December 2020 5:05 PM
To: s22
Cc: Cunningham, Paul (CorpAffairs, Dutton Park)
Subject: FW: CSIRO's GISERA NT project - final report - stygofauna and microbial assemblages of the Beetaloo Sub-basin, NT
Attachments: Duplicate Attachment - Removed

Hi s22

I hope that you are well.

It is part of CSIRO's GISERA Communication Protocol to forward the National Research Management Committee a courtesy copy of final reports 10 working days prior to public release. This action does not constitute a form of 'review' by our industry partners but allows our partners to bring to our attention any concerns before the report is made public. We also conduct a knowledge transfer session with industry partners and government where the results are presented and discussed prior to public release (we've already communicated about our upcoming KTS).

The NRMCM have just been emailed a copy of the final report for the NT project [Characterisation of the stygofauna and microbial assemblages of the Beetaloo Sub-basin, NT](#).

I am forwarding this to you as the assigned delegate for Origin's NRMCM representative – Stephanie Stonier.

If you have any questions, please don't hesitate to contact us.

Regards

Jizelle

Jizelle Khoury

Executive Officer, CSIRO's Gas Industry Social and Environmental Research Alliance (GISERA)
Energy | CSIRO

s22

Address: PO Box 52, North Ryde NSW 1670, Australia

CSIRO Australia's National Science Agency | csiro.au

Duplicate Email - Removed

s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Tuesday, 8 December 2020 5:11 PM
To: s22@shell.com
Cc: Cunningham, Paul (CorpAffairs, Dutton Park)
Subject: FW: CSIRO's GISERA NT project final report stygofauna and microbial assemblages of the Beetaloo Sub-basin NT
Attachments: Duplicate Attachment - Removed

Hi s22

I hope that you are well.

It is part of CSIRO's GISERA Communication Protocol to forward the National Research Management Committee (NRMC) a courtesy copy of final reports 10 working days prior to public release. This action does not constitute a form of 'review' by our industry partners but allows our partners to bring to our attention any concerns before the report is made public.

The NRMC (including Patrick McKelvey) have just been emailed a copy of the final report for the NT project [Characterisation of the stygofauna and microbial assemblages of the Beetaloo Sub-basin, NT.](#)

I am forwarding a copy to you also, as I was unsure whether you have officially taken over as Shell/QGC's representative on GISERA's NRMC. If this is the case, it would be great if you could let me know and I'll update our NRMC distribution list.

If you have any questions, please don't hesitate to contact us.

Regards
Jizelle

Jizelle Khoury
Executive Officer, CSIRO's Gas Industry Social and Environmental Research Alliance (GISERA)
Energy | CSIRO

s22

Address: PO Box 52, North Ryde NSW 1670, Australia

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Duplicate Email - Removed

s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Thursday, 10 December 2020 3:58 PM
To: s22
Cc: Cunningham, Paul (CorpAffairs, Dutton Park)
Subject: RE: FOR ADVICE: Origin attendees at CSIRO's GISERA knowledge transfer session NT water project

Hi s22

I am writing to let you know that we have decided to defer the knowledge transfer session to early 2021.

By deferring it, we'll be able to conduct a joint session with the other water project (Environmental monitoring and microbial degradation of onshore shale gas activity chemicals and fluids) which is due to finish shortly.

I'll be in touch in the New Year to confirm a date.

In the meantime, I'd like to wish you all the best for the festive season.

Kind regards
Jizelle

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Tuesday, 8 December 2020 7:40 AM
To: s22@origin.com.au>; s22@origin.com.au>
Cc: Cunningham, Paul (CorpAffairs, Dutton Park) s22
Subject: RE: FOR ADVICE: Origin attendees at CSIRO's GISERA knowledge transfer session NT water project

Hi s22

I wanted to let you know that we have earmarked Monday, 14 December at 9.30 10.30 NT time (10-11 am QLD) for the knowledge transfer session.

I haven't sent out a calendar invite yet as I'm waiting for the NTG to finalise their participant list. It will go out by Thursday (latest). In the meantime, can I ask that you please hold this timeslot in your diaries.

Many thanks
Jizelle

Jizelle Khoury
Executive Officer, CSIRO's Gas Industry Social and Environmental Research Alliance (GISERA)
Energy | CSIRO

s22
Address: PO Box 52, North Ryde NSW 1670, Australia

CSIRO Australia's National Science Agency | csiro.au

From [redacted] s22 [redacted]@origin.com.au>

Sent: Wednesday, 2 December 2020 1:21 PM

To: Khoury, Jizelle (Energy, North Ryde) [redacted] s22 [redacted] s22 [redacted]@origin.com.au>;

[redacted] s22 [redacted]@origin.com.au>

Cc: Cunningham, Paul (CorpAffairs, Dutton Park) [redacted] s22 [redacted]

Subject: RE: FOR ADVICE: Origin attendees at CSIRO's GISERA knowledge transfer session - NT water project

Hi Jizelle,

I remember you from when I was working with Shell; I hope you're well.

At this stage we are planning for Matt and Carl Altmann (copied in) to join the session in w/c 14 December.

Kind regards,

o o o o

[redacted] s22 [redacted]

Origin
Level 28, 180 Ann Street, Brisbane QLD 4000

[redacted] s22 [redacted]

Connect with us



originenergy.com.au

I acknowledge the Traditional Owners and Custodians of country throughout Australia and recognise their continuing connection to land, waters and community.

I pay my respects to them and their cultures, and to Elders past, present and future.

Origin shows its commitment to participating in Australia's reconciliation efforts through our [Stretch Reconciliation Action Plan](#).



Duplicate Email - Removed

s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Thursday, 10 December 2020 3:59 PM
To: s22
Cc: Cunningham, Paul (CorpAffairs, Dutton Park)
Subject: RE: FOR ADVICE: Santos attendees at CSIRO's GISERA knowledge transfer session NT water project

Hi s22

I am writing to let you know that we have decided to defer the knowledge transfer session to early 2021.

By deferring it, we'll be able to conduct a joint session with the other water project (Environmental monitoring and microbial degradation of onshore shale gas activity chemicals and fluids) which is due to finish shortly.

I'll be in touch in the New Year to confirm a date.

In the meantime, I'd like to wish you all the best for the festive season.

Kind regards
 Jizelle

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Tuesday, 8 December 2020 7:39 AM
To: s22@santos.com>; s22@santos.com>; s22@santos.com>
Cc: Cunningham, Paul (CorpAffairs, Dutton Park) s22
Subject: RE: FOR ADVICE: Santos attendees at CSIRO's GISERA knowledge transfer session NT water project

Hi s22

I wanted to let you know that we have earmarked Monday, 14 December at 9.30-10.30 NT time (10-11 am QLD) for the knowledge transfer session.

I haven't sent out a calendar invite yet as I'm waiting for the NTG to finalise their participant list. It will go out by Thursday (latest). In the meantime, can I ask that you please hold this timeslot in your diaries.

Many thanks
 Jizelle

Jizelle Khoury
 Executive Officer, CSIRO's Gas Industry Social and Environmental Research Alliance (GISERA)
 Energy | CSIRO

s22

Address: PO Box 52, North Ryde NSW 1670, Australia

CSIRO Australia's National Science Agency | csiro.au

From [redacted] s22 [redacted]@santos.com>

Sent: Wednesday, 2 December 2020 1:25 PM

To: Khoury, Jizelle (Energy, North Ryde) [redacted] s22 [redacted]

Cc: Cunningham, Paul (CorpAffairs, Dutton Park) [redacted] s22 [redacted]

[redacted] s22 [redacted]@santos.com>; [redacted] s22 [redacted]@santos.com>

Subject: RE: FOR ADVICE: Santos attendees at CSIRO's GISERA knowledge transfer session NT water project

Hi Jizelle,

Can you please invite David Gornall, Mitch Bird (both CC'ed into this email) and myself

Regards,

Santos

[redacted] s22 [redacted]

Santos Limited, 32 Turbot Street, Brisbane QLD 4000

[redacted] s22 [redacted]

   santos.com

Duplicate Email - Removed

s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Thursday, 28 January 2021 10:01 AM
To: s22
Cc: Cunningham, Paul (CorpAffairs, Dutton Park)
Subject: RE: FOR ADVICE: Origin attendees at CSIRO's GISERA knowledge transfer session
NT water project

Hi s22

I hope that you are well.

We've decided to proceed with the KTS for the stygofauna project and not conduct a joint session with the other water project. The other project is still a few weeks away from delivery and I don't want to hold up the public release of the Stygofauna report.

I've just sent through a calendar invite for 11 February.

Regards
Jizelle

Duplicate Email - Removed

s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Thursday, 28 January 2021 10:03 AM
To: s22
Cc: Cunningham, Paul (CorpAffairs, Dutton Park)
Subject: RE: FOR ADVICE: Santos attendees at CSIRO's GISERA knowledge transfer session - NT water project

Hi s22

I hope that you are well.

We've decided to proceed with the KTS for the stygofauna project and not conduct a joint session with the other water project. The other project is still a few weeks away from delivery and I don't want to hold up the public release of the Stygofauna report.

I've just sent through a calendar invite for 11 February.

Regards
Jizelle

Duplicate Email - Removed

s22

Subject: CSIRO's GISERA 'Characterisation of the stygofauna and microbial assemblages of the Beetaloo Sub Basin, NT' project Knowledge Transfer Session

Location: Via WebEx (see inside for dial in details)

Start: Thu 11/02/2021 10:45 AM

End: Thu 11/02/2021 11:45 AM

Show Time As: Tentative

Recurrence: (none)

Organizer: Khoury, Jizelle (Energy, North Ryde)

Dear all

Please join us for a Knowledge Transfer Session where we will be presenting the latest findings from CSIRO GISERA's Characterisation of the stygofauna and microbial assemblages of the Beetaloo Sub Basin, NT project.

Date: Thursday, 11 February 2021

Time: 9.15 10.15 am (Darwin time)

Agenda (NT time):

9.15 am Welcome
 9.20 am Research presentation
 9.50 am Discussion panel with questions from all participants welcome
 10.15 am Finish

Method: The meeting will be hosted by WebEx, please see details below. If you are first time using WebEx please log in 5 minutes prior to the meeting to allow for the program to download.

For any participants unable to connect using the Desktop or Video Conference Room WebEx options, please dial-in using the audio number. I will forward a copy of the PPT slides prior to commencement so that you can follow the presentation.

Welcome to CSIRO Webex Conferencing featuring: audio, video and presentation capabilities. You can join this conference from:

Desktop or Mobile Devices

Once connected to your meeting remember to start your audio and video

Video Conferencing (VC) Rooms

Use the remote control or touch panel and dial the number indicated below:

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 External VC Room

Phones - AUDIO ONLY

Primary Australia
 Quick Dial
 Other Global Numbers

<https://conferencing.csiro.au/Call-in.php>

Meeting Number/Access Code

Password (if prompted)

First time joining a Webex meeting? Watch this short video to get started:

<https://conferencing.csiro.au/videos/videos.php?tab=join>

Need further help? Take a look at the link below for user guides, videos and FAQs:

https://conferencing.csiro.au/index.php?meeting_number=1652631473&site=csiro.webex.com

We look forward to sharing our research with you.

Regards

Jizelle

Jizelle Khoury

Executive Officer, CSIRO's Gas Industry Social and Environmental Research Alliance (GISERA)

Energy | CSIRO

s22

Address: PO Box 52, North Ryde NSW 1670, Australia

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s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Thursday, 11 February 2021 8:42 AM
To: Barrett, Damian (Energy, Black Mountain); Rees, Gavin (L&W, Albury); s22
 Cunningham, Paul (CorpAffairs, Dutton Park); s22
 Carl; s22
 Huddleston-Holmes, Cameron (Energy, Pullenvale)
Subject: CSIRO's GISERA knowledge transfer session Stygofauna project
Attachments: Stygofauna GISERA 11 Feb 2021.pdf

Dear all

You have been invited to participate in this morning's knowledge transfer session for GISERA project Characterisation of the stygofauna and microbial assemblages of the Beetaloo Sub-Basin, NT.

The calendar invite includes the Webex details where you can dial in via video and view the presentation.

For those participants who are not able to connect to Webex, please simply call ph: , followed by meeting id/access code This connects you to audio only. You will then need to open the attached PPT slides and the presenter will instruct you when to move to the next slide.

If you have any questions, please don't hesitate to contact me.

Thanks
Jizelle

Jizelle Khoury
Executive Officer, CSIRO's Gas Industry Social and Environmental Research Alliance (GISERA)
Energy | CSIRO

s22

Address: PO Box 52, North Ryde NSW 1670, Australia

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Characterisation of the stygofauna assemblages of the Beetaloo Sub-basin, Northern Territory

Gavin Rees,

s22

2nd Feb 2021



RIEL

Research Institute for the Environment and Livelihoods



QGC



Santos



Australian Government
Department of Industry, Science,
Energy and Resources



Supported by
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South Australia



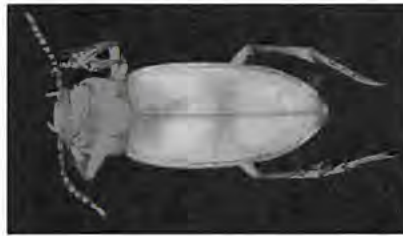
PANGAEA
PANGAEA PTY LTD

Background

- Definition:
 - Ground water fauna, or stygofauna, are animals that live permanently underground in water.
 - Stygofauna live in a range of groundwater habitats—from tiny spaces between sand grains to pools and streams in caves.



Amphipod
(image courtesy Bennelongia)

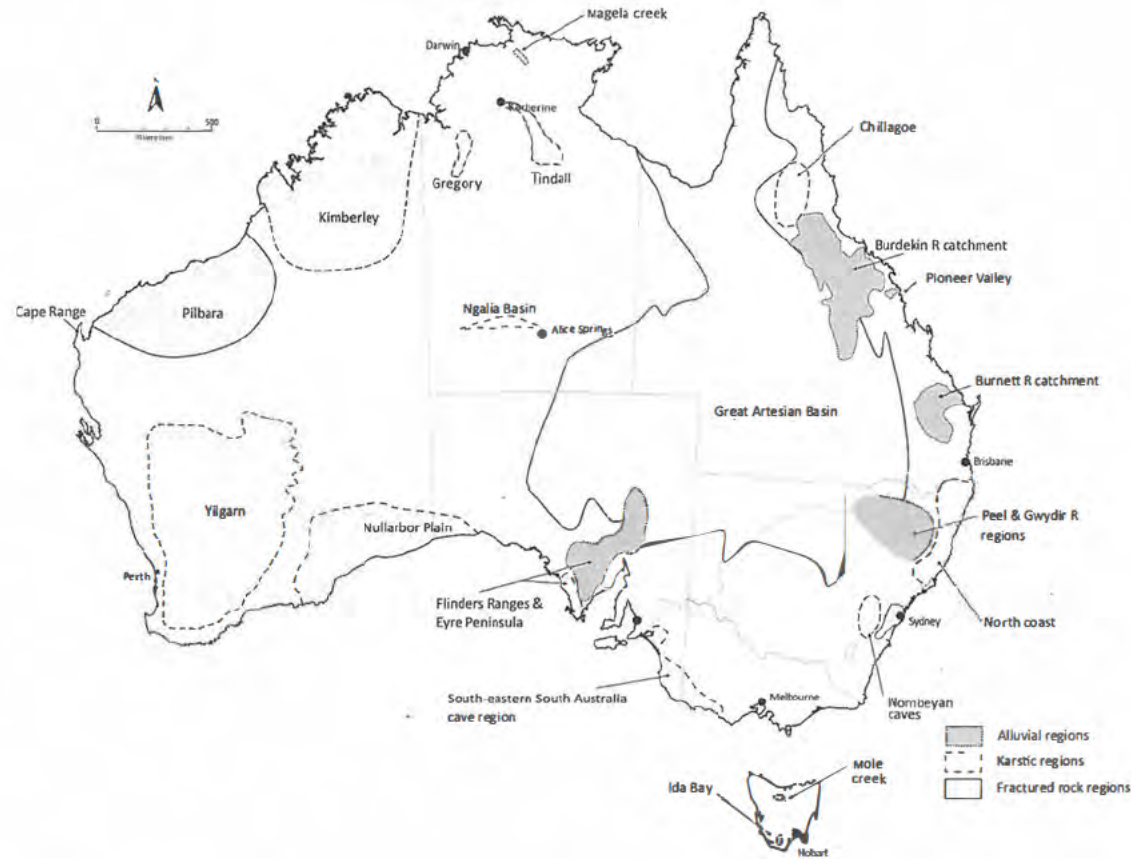


Beetle
(image courtesy Bennelongia)



Blind cave fish

Background - General aquifer types and regions where stygofauna have been found



Modified from Tomlinson and Boulton (2008) with additional information from Guzik *et al.* (2008), Hose *et al.* (2015a) and Chandler *et al.* (2017) and this study (the Tindall aquifer)

Pilot project

Overall project objective.

- To provide new knowledge concerning stygofauna and subterranean groundwater dependent ecosystems in the Beetaloo Sub-basin and Roper River system

Approach.

- Carry out a pilot scale sampling program to examine a limited series of bores/bore water for the presence of stygofauna

Project Team

Gavin Rees (CSIRO)

Daryl Nielsen (CSIRO)

s22 (CDU)

s22 (CDU)

Garth Watson (CSIRO)

s22 (LaTrobe Uni)

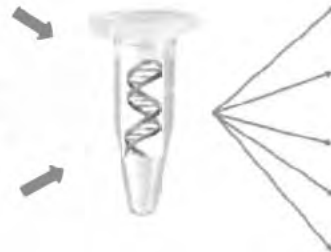


Project – brief approach

- Sample bore water from:
 - 28 sites, including 2 springs,
 - From Mataranka to semi-arid Barkly Tablelands
 - Combination of sites within and outside leases, to sample different types bores
 - Carried out a second sampling trip. Further bores and revisited some earlier bores
- Use a range of bore sampling methods, depending on type of bore
- Preserve and identify any organisms
 - Where relevant, use DNA barcoding to identify organisms
- Use an environmental DNA approach to examine bore water
 - Detecting the DNA from organisms in bore water rather than entire organism



eDNA
metabarcoding



Library of barcodes

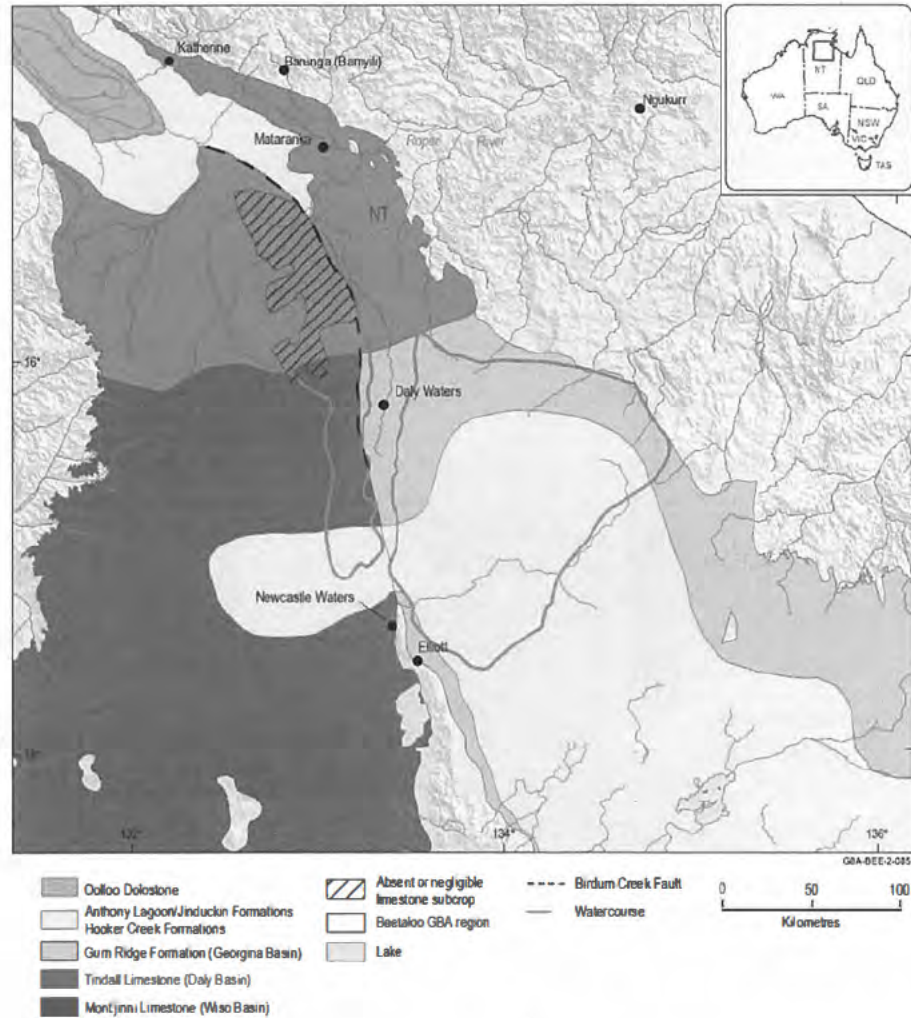
Information on
biodiversity

Fungi

Plants

Animals

Location



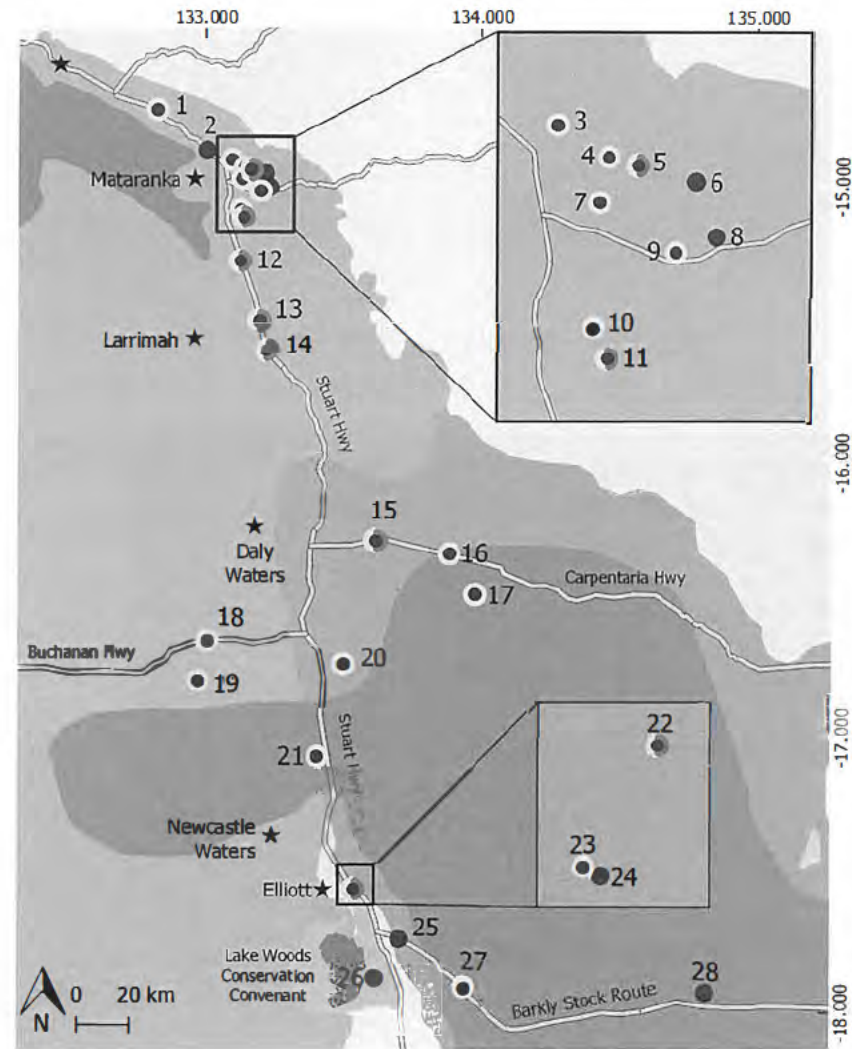
Legend

- Bores sampled in 2019
- Presence of stygofauna indicated by eDNA only
- Presence of stygofauna indicated by both collection of specimens and eDNA

Major hydrostratigraphic units

- Oolloo Dolostone
- Tindall Limestone (Daly Basin)
- Gum Ridge Formation (Georgina Basin)
- Montijinni Limestone (Wiso Basin)
- Anthony Lagoon/ Jinduckin Formations

- ★ Main Towns
- Main Roads



Bores – some examples



Buchanan Downs



Shenandoah homestead

Bores – some examples



Elliot 8 (RN036781)



Mataranka Homestead
(RN35796)

Bores – some examples



Bores – sampling using pumps



Pumped water is passed through an ultra-fine net to collect animals



Bores – hand held nets



Results

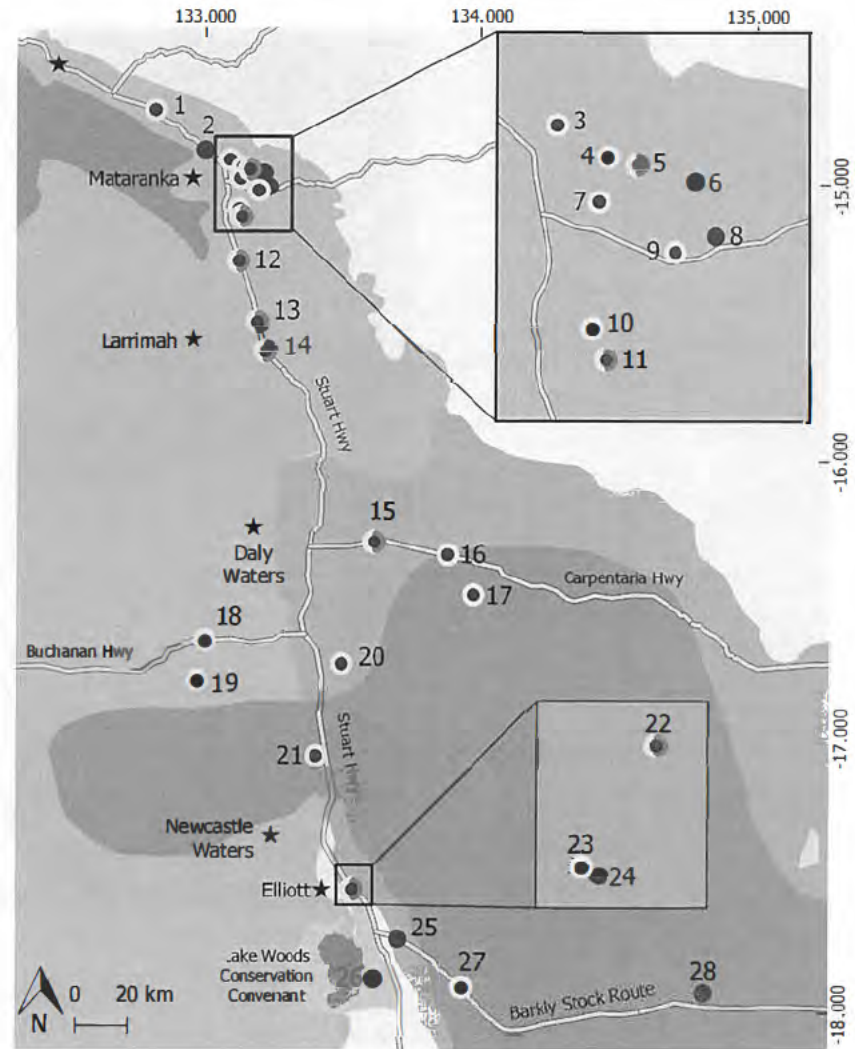
Legend

- Bores sampled in 2019
- Presence of stygofauna indicated by eDNA only
- Presence of stygofauna indicated by both collection of specimens and eDNA

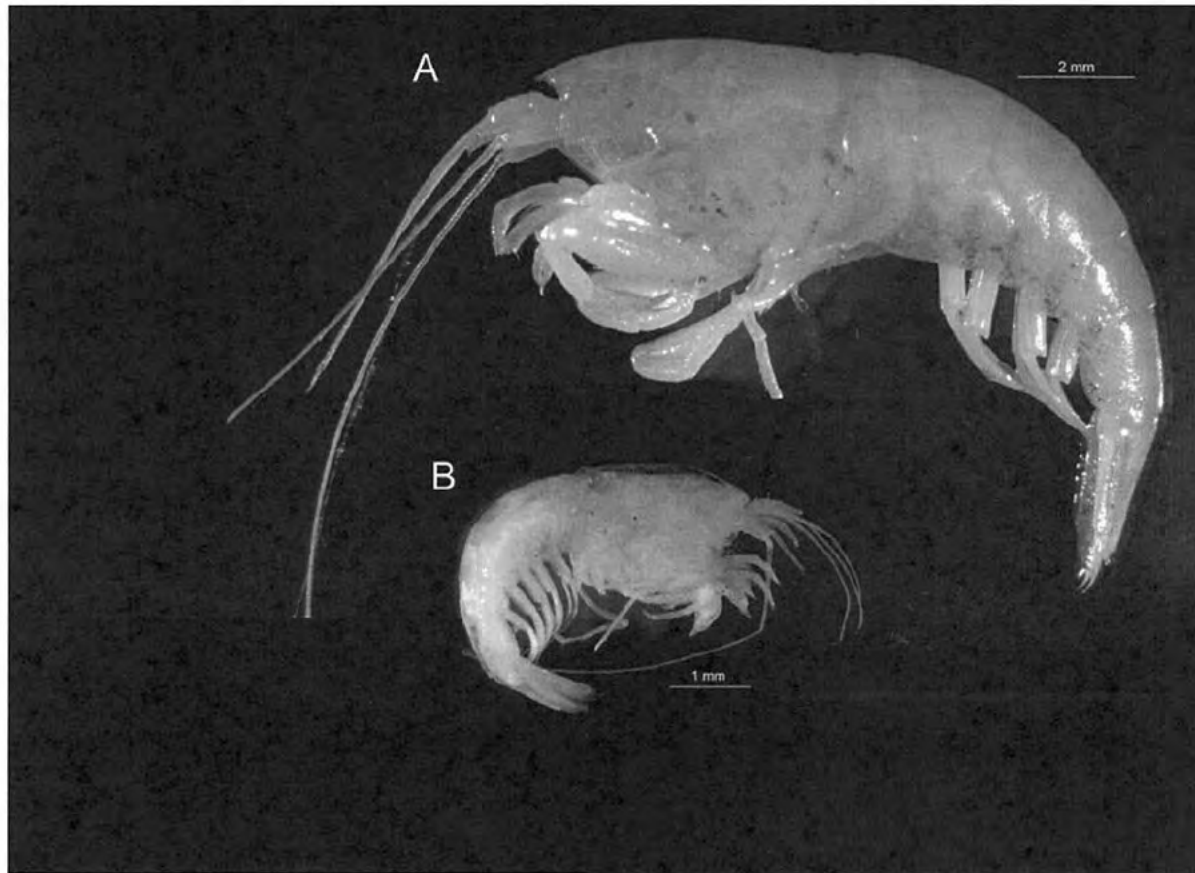
Major hydrostratigraphic units

- Ooloo Dolostone
- Tindall Limestone (Daly Basin)
- Gum Ridge Formation (Georgina Basin)
- Montijinni Limestone (Wiso Basin)
- Anthony Lagoon/ Jinduckin Formations

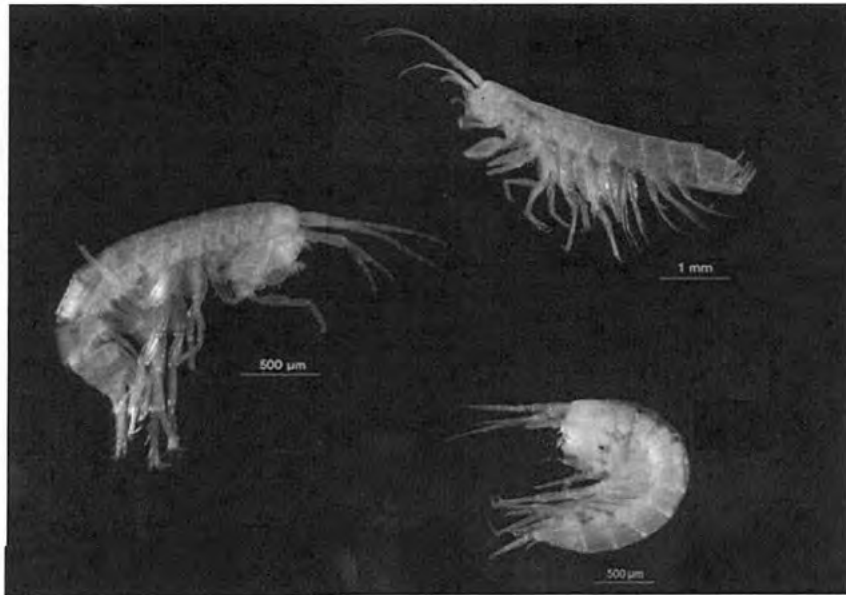
- ★ Main Towns
- Main Roads



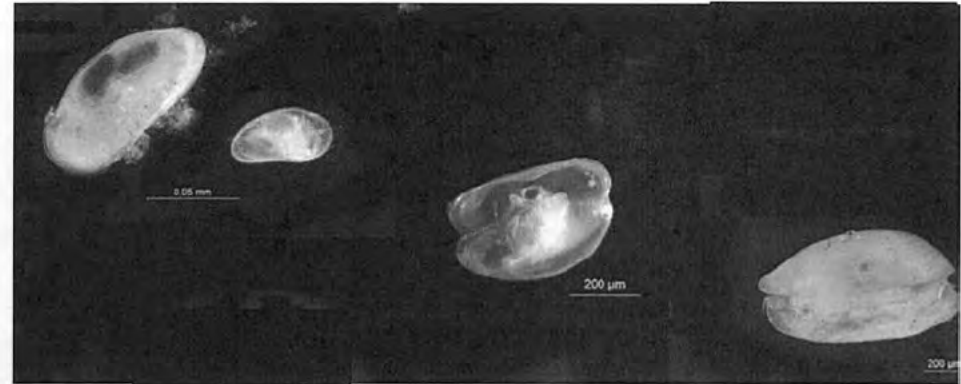
Blind shrimp



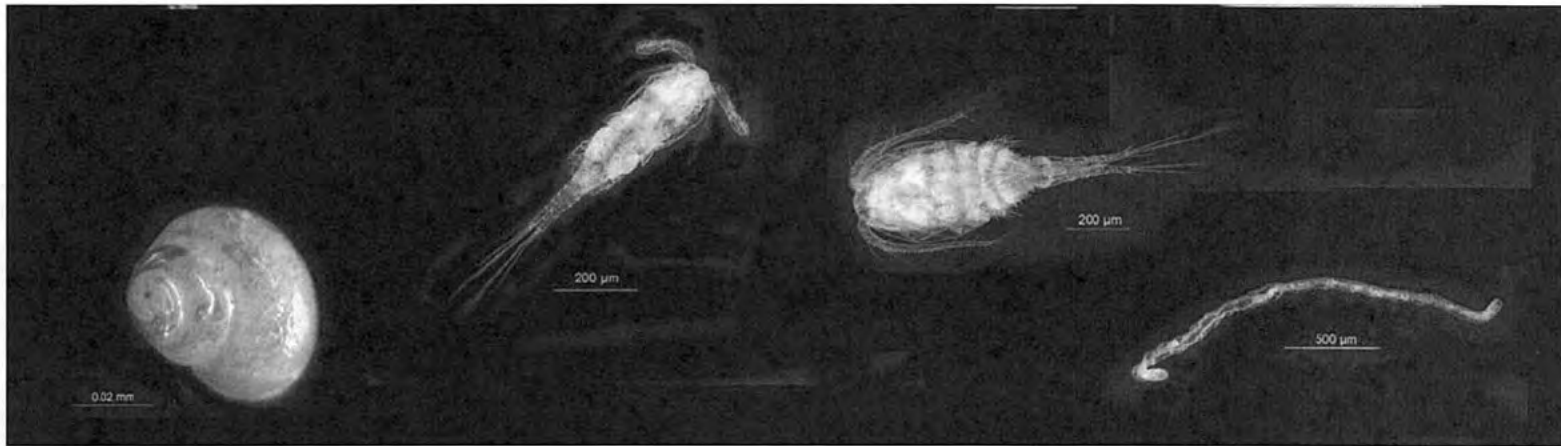
Small crustaceans



Amphipods
A very small crustacean



Ostracods –
Another class of small crustaceans



Snail

Cyclopoids
Tiny crustaceans
(‘zooplankton’)

Worms

eDNA as a detection tool

- Our eDNA recognized three categories of organisms
 - Contaminant terrestrial DNA. Eg, ants
 - Probable soil organisms. Eg, soil fungi,
 - Organisms dwelling in bore waters. Eg, crustaceans detected by netting
- Stygofauna DNA detected across many bores
- Accurate identification of eDNA results requires
 - Animals accurately identified
 - DNA barcodes obtained and have been put into the DNA libraries

Results

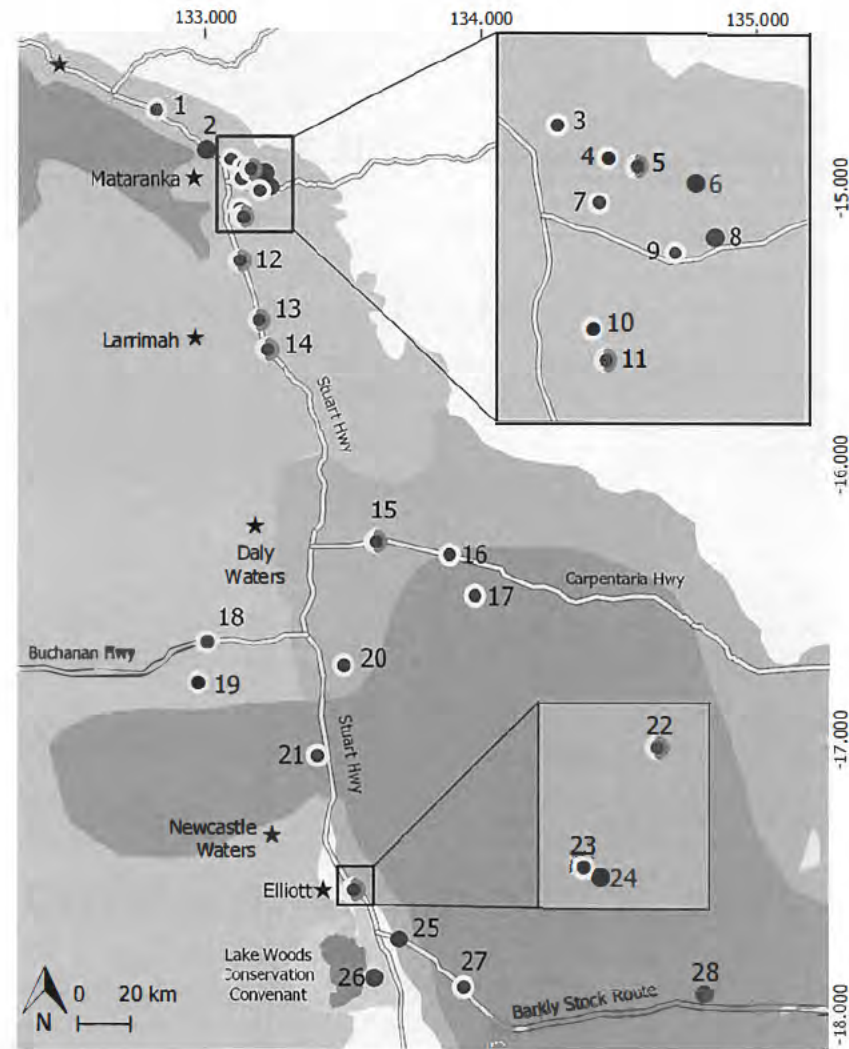
Legend

- Bores sampled in 2019
- Presence of stygofauna indicated by eDNA only
- Presence of stygofauna indicated by both collection of specimens and eDNA

Stygofauna dominated by crustaceans

Organised food web

- shrimp top predator?



Returning to the shrimp

- Three species been described from Cutta Cutta caves near Katherine
 - *Parisia unguis*, *Parisia gracilis*, *Pycnisia raptor*
 - Extremely limited taxonomy (single specimens, pieces of animal), so very low certainty about their true identity
- Our specimens most closely related to *Parisia unguis*
 - Given low genetic diversity of our specimens, this species spread over some 500km

Summary points

- First studies of the aquifers showed stygofaunal communities were dominated by crustaceans
- Showed little affinity with the stygofauna recorded from more extensively sampled Western Australian aquifers
- Highly likely new genera and species present in the Beetaloo Sub-basin
- Evidence of connectivity within the aquifer across our sample sites

Acknowledgements

- Stuart Halse (Bennelongia Environmental Consultants)
- John Short (BioAccess Australia)
- Vanessa Solano Rivera (CDU – for generating maps)



Thank you

Gavin Rees
Principal Research Scientist

t [REDACTED] s22
e [REDACTED]



Australian Government
Department of Industry, Science,
Energy and Resources



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s22

From: Khoury, Jizelle (Energy, North Ryde)
Sent: Thursday, 11 February 2021 9:54 AM
To: s22
Subject: RE: CSIRO's GISERA knowledge transfer session - Stygofauna project

OK thanks for letting me know.

Regards
Jizelle

From: s22@santos.com>
Sent: Thursday, 11 February 2021 9:31 AM
To: Khoury, Jizelle (Energy, North Ryde) s22
Subject: RE: CSIRO's GISERA knowledge transfer session - Stygofauna project

Hi Jizelle, just letting you know I might be 5-10 mins late to this. Dave and Mitch will be there

Regards,

Santos

s22

Santos Limited, 32 Turbot Street, Brisbane QLD 4000

s22

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Duplicate Email - Removed

s22

From: [REDACTED] s22 [REDACTED]@santos.com>
Sent: Thursday, 11 February 2021 12:51 PM
To: Barrett, Damian (Energy, Black Mountain)
Cc: [REDACTED] s22
Subject: Stygo in Beetaloo

Damian

I think the discussion about the effect of a bore on stygo sampling error is somewhat of a red herring. Even if results are valid, contextualising findings will be difficult without a complete risk assessment, or at least some discussion of the possible pathways/mechnisms (what can we discount as a potential hazard e.g. aquifer water extraction), which is beyond the scope of this study.

Re: effect of bores on stygo sampling error this is bigger issue than the Beetaloo Basin, and may not be necessary for the SREBA. However it may be worth acknowledging the uncertainty this introduces, if nothing else but to stimulate the broader research community:

- My observation is that in dry parts of the world, the animals you will find eking out a living in and around a bore can be staggering. It is common to see piles of bones at the bottom of a bore when we run a camera. Pulling pumps out of holes, I've seen large numbers ~10cm frogs attached to a riser where there was no obvious way for them to get into the bore, unless when very small.
- I'm not saying that stygofauna do not exist, but how can stygofauna sampling methods and analysis correct for the effect of the bore on the local groundwater environment.
- Possible research angles, compare results:
 - o How do water quality parameters offer insight for whether the water sampled was bore column or true aquifer water (noting the strict standard for groundwater sampling re: bore purge volumes)
 - o Sampling standing water column method versus purged water methods
 - o Age of bore (new (months) vs old (decades))
 - o Bore condition (is the bore completely open to environment, part-sealed (small gaps), total seal (screw cap/bolted flange))
- eDNA offers a novel research angle using DNA residence times some DNA may only be reasonably explained by surface interaction due to the bore. Acting like a tracer.

Cheers

s22

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s22



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s22

From: Cunningham, Paul (CorpAffairs, Dutton Park)
Sent: Tuesday, 16 February 2021 2:47 PM
To: s22
Cc: Barrett, Damian (Energy, Black Mountain)
Subject: FW: stygofauna release out today

H s22

The Conversation article was initiated by CDU researcher professor s22

Our CSIRO researchers provided feedback to ensure as far as possible The Conversation article reflected the research.

Our own key messages remain tightly focused on the substance of the final report, and our communications material was distributed nationally this morning at 6am.

Here's a link to the CSIRO/GISERA and Charles Darwin University joint [media release and images](#)

CSIRO/GISERA [Fact sheet](#)

CSIRO/GISERA [Final report](#)

[CSIRO News](#)

Happy to discuss further if you have any questions.

Paul Cunningham
Communication and Stakeholder Manager
Gas Industry Social and Environmental Research Alliance
Energy | CSIRO

s22

s22

From: Cunningham, Paul (CorpAffairs, Dutton Park)
Sent: Wednesday, 17 February 2021 5:40 PM
To: s22@chambernt.com.au; s22@ntfarmers.org.au; James.Pratt@s22@originenergy.com.au; Robinson, Cathy (L&W, Dutton Park); Chilcott, Chris (L&W, Darwin); s22@nlc.org.au; s22@santos.com; Dewhurst, David (Energy, Kensington WA); s22@gmail.com; executive.office@s22
Cc: Barrett, Damian (Energy, Black Mountain); Khoury, Jizelle (Energy, North Ryde)
Subject: FW: CSIRO GISERA final report online stygofauna and microbial assemblages of the Beetaloo Sub basin, NT

Dear all

The final report of the CSIRO GISERA project *Characterisation of the stygofauna and microbial assemblages of the Beetaloo Sub-basin, NT* is now online. This project is complete.

This report describes the results of a joint CSIRO GISERA /Charles Darwin University survey of 26 water bores and two groundwater springs in the Beetaloo Sub-basin and Roper River system in the Northern Territory. The survey revealed diverse communities of tiny aquatic animals (stygofauna) in the Cambrian Limestone Aquifer. This study provides the first description of stygofauna in an otherwise little-studied region of Australia and likely includes discovery of new species of crustaceans.

This research responds to recommendations from the Northern Territory Government's *Scientific Inquiry into Hydraulic Fracturing in Northern Territory*. This baseline data is essential for biodiversity conservation and the maintenance of the ecological integrity of high value groundwater dependent ecosystems in the region, and informs appropriate policy and management responses to shale gas development proposals.

Supporting resources: Joint CSIRO GISERA/Charles Darwin University [media release](#) and [factsheet](#) also available on the GISERA web site.

If you would like to discuss these result please contact GISERA Director Damian Barrett.

Paul Cunningham

Communication and Stakeholder Manager
 Gas Industry Social and Environmental Research Alliance
 Energy | CSIRO

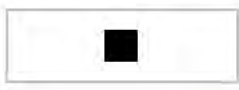
s22

From: s22
To: s22, s47E (Energy, North Ryde)
Cc: (Energy, North Ryde); s22
Subject: Chemical Risk Assessment
Date: Wednesday, 28 August 2019 11:12:59 AM
Attachments: [image001.jpg](#)
[image002.jpg](#)
[image003.jpg](#)
[image004.jpg](#)
s47G(1)(a), s47, s45


Hi s22, s47E



Please find attached s47G(1)(a), s47, s45 in the NT. Let me know if you have any questions.

Regards,
s22



s22



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From: s22, s47E (Energy, Dutton Park)
To: s22, s47E (Energy, North Ryde); s22, s47E (Energy, North Ryde); s22, s47E (L&W, Albury); s22, s47E (L&W, Albury); s22, s47E (Energy, Kensington WA); s22 @cdu.edu.au
Subject: FW: Beetaloo Basin_Santos Bore Survey Data
Date: Friday, 14 June 2019 1:39:52 PM
Attachments: [image001.jpg](#)
[image002.jpg](#)
[image003.jpg](#)
[image004.jpg](#)
s47G(1)(a), s47, s45

FYI

From: s22 [mailto:s22@santos.com]
Sent: Friday, 14 June 2019 12:36 PM
To: s22, s47E (Energy, Dutton Park) s22
Subject: s47G(1)(a), s47, s45

s22,
s47E

See attached s47G(1)(a), s47, s45
s47G(1)(a), s47, s45

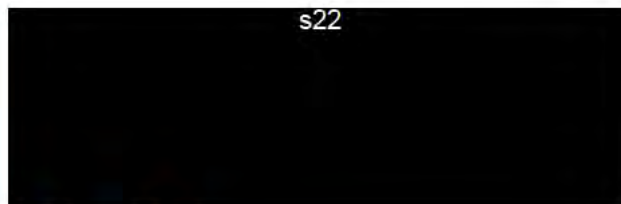
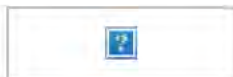
This is provided for the purposes of planning the upcoming GISERA fieldwork for stygofauna and biodegradation studies.

Please get back to me if you have any questions.

We have photos of all the headworks too, but this would be large dataset. Potentially s22, s47E s22, s47E has similar record.

Thanks

s22

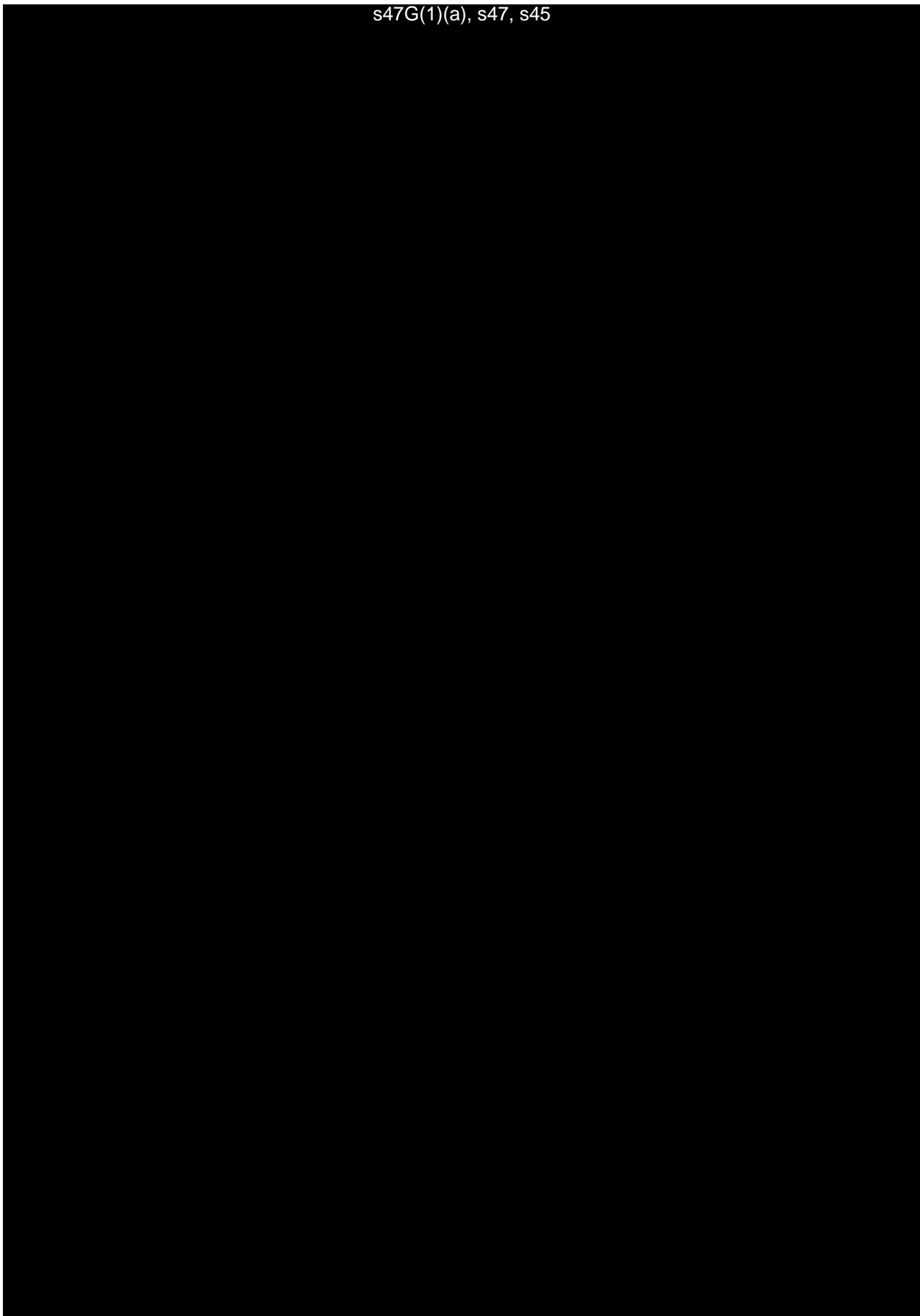


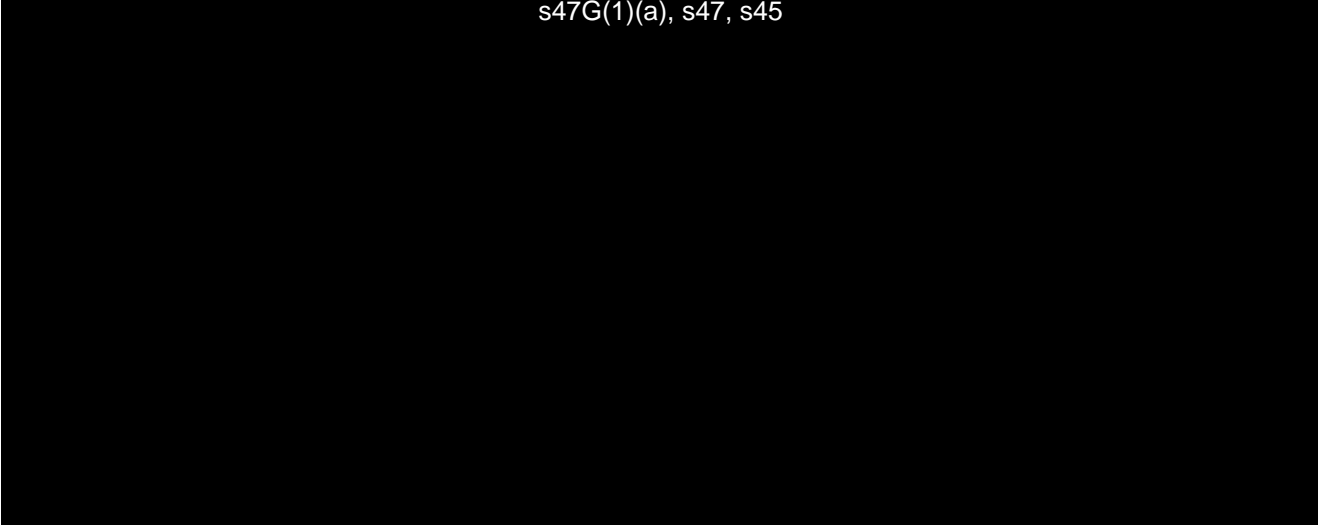
   [santos.com](#)

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From: s22, s47E (Energy, North Ryde)
To: s22
Cc: s22, s47E (Energy, North Ryde)
Subject: GISERA project - Northern Territory - List of chemicals used by onshore gas development that have environmental impacts
Date: Tuesday, 6 August 2019 11:15:00 AM

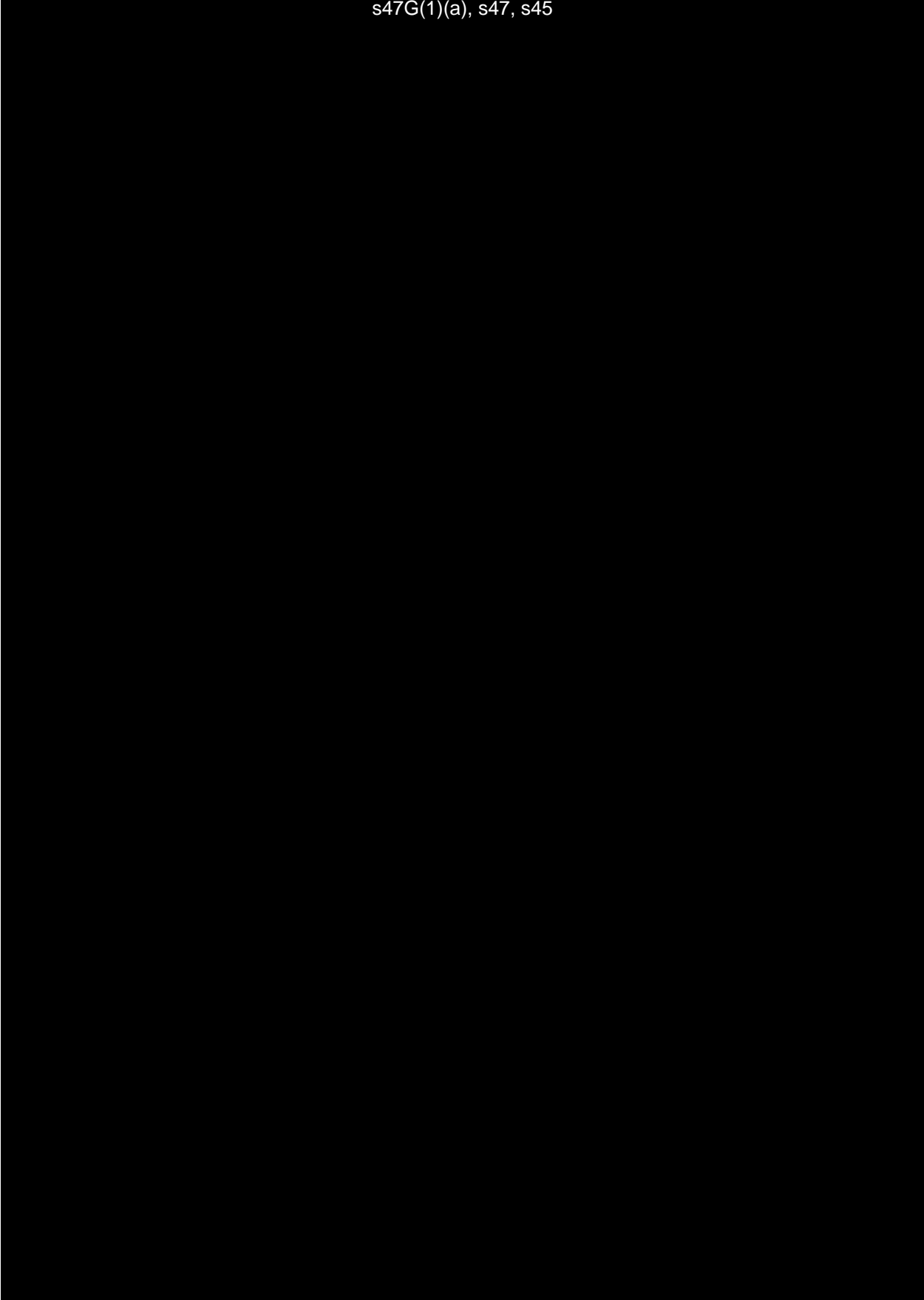
s47G(1)(a), s47, s45





From: [REDACTED] s22, s47E ([Energy, North Ryde](#))
To: [REDACTED] s22
Cc: [REDACTED] s22, s47E ([Energy, North Ryde](#))
Subject: GISERA project - Northern Territory - List of chemicals used by onshore gas development that have environmental impacts
Date: Tuesday, 6 August 2019 11:14:00 AM

s47G(1)(a), s47, s45





From: s22
To: [Redacted]
Cc: s22, s47E (Energy, North Ryde); s22, s47E (Energy, North Ryde)
Subject: RE: GISERA project - Northern Territory - List of chemicals used by onshore gas development that have environmental impacts
Date: Tuesday, 6 August 2019 11:17:44 AM
Attachments: [image001.jpg](#)
[image002.jpg](#)
[image003.jpg](#)
[image004.jpg](#)

Hi s22

Are you able to advise s22, s47E here?

Thanks
s22



Duplicate Email - Removed

From: [REDACTED] s22
To: [REDACTED]
Cc: [REDACTED] s22, s47E ([Energy, North Ryde](#)); [REDACTED] s22, s47E ([Energy, North Ryde](#))
Subject: RE: GISERA project - Northern Territory - List of chemicals used by onshore gas development that have environmental impacts
Date: Tuesday, 6 August 2019 12:33:47 PM
Attachments: [image001.png](#)

Gents,

Can you assist [REDACTED] s22, s47E with her query?

[REDACTED]

[REDACTED] s22

Duplicate Email - Removed

From: [REDACTED] s22
To: [REDACTED]
Cc: s22, s47E (Energy, North Ryde); [REDACTED] s22, s47E (Energy, North Ryde)
Subject: RE: GISERA project - Northern Territory - List of chemicals used by onshore gas development that have environmental impacts
Date: Friday, 9 August 2019 9:39:32 AM
Attachments: [image005.jpg](#)
[image006.jpg](#)
[image007.jpg](#)
[image008.jpg](#)
[image009.jpg](#)
[image010.jpg](#)
[image011.jpg](#)
[image012.jpg](#)

Hi [REDACTED] s22, s47E

Can you please give me a call on my mobile?

This list does not appear to be representative of additives proposed for use in the Beetaloo (maybe 2-3 additives based on a quick scan of the CAS numbers).

Regards,



[REDACTED] s22



Duplicate Email - Removed

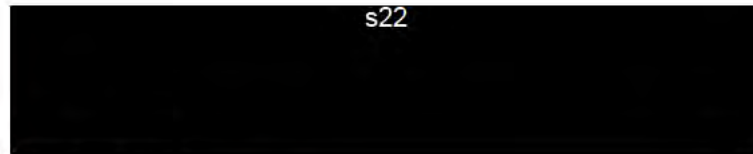


From: s22
To: s22, s47E (Energy, North Ryde); s22
Cc: (Energy, North Ryde)
Subject: RE: GISERA project - Northern Territory - List of chemicals used by onshore gas development that have environmental impacts
Date: Monday, 12 August 2019 9:19:58 AM
Attachments: [image001.jpg](#)
[image002.jpg](#)
[image003.jpg](#)
[image004.jpg](#)
[image005.jpg](#)
[image006.jpg](#)
[image007.jpg](#)
[image008.jpg](#)

Hi s22, s47E

3pm works fine for me

Regards,



From: s22, s47E (Energy, North Ryde) s22
Sent: Saturday, 10 August 2019 4:01 AM
To: s22
Cc: s22, s47E (Energy, North Ryde) s22
Subject: ![EXT]: RE: GISERA project - Northern Territory - List of chemicals used by onshore gas development that have environmental impacts

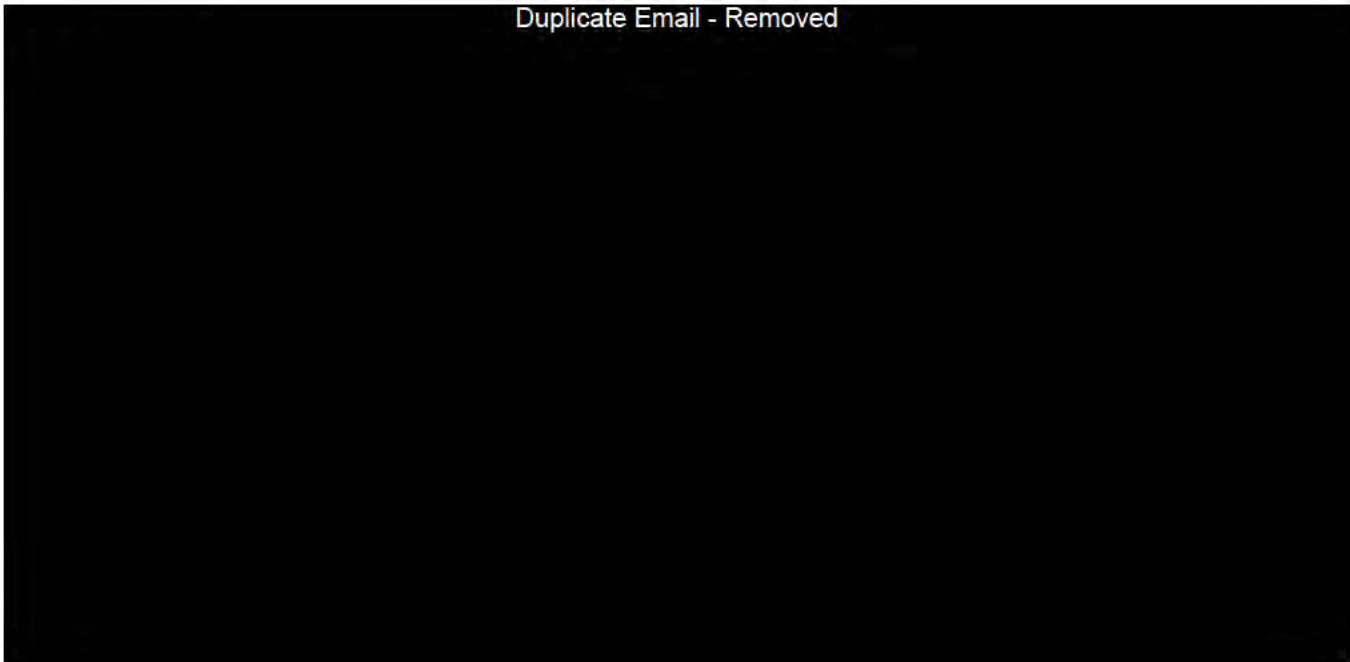
Hi s22

Thank you for your email.

Can we call you on Monday afternoon at 3pm?

Regards,
s22, s47E

Duplicate Email - Removed



From: s22, s47E (Energy, North Ryde)
To: s22
Cc: s22, s47E (Energy, North Ryde)
Subject: RE: GISERA project - Northern Territory - List of chemicals used by onshore gas development that have environmental impacts
Date: Tuesday, 20 August 2019 1:21:44 PM
Attachments: [image001.jpg](#)
[image002.jpg](#)
[image003.jpg](#)
[image004.jpg](#)
[image005.jpg](#)
[image006.jpg](#)
[image007.jpg](#)
[image008.jpg](#)

s47G(1)(a), s47, s45

s22, s47E

Senior Research Scientist | Microbial Ecology & Mycology
Reservoir Characterisation Team Leader

Energy
CSIRO

s22

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Duplicate Email - Removed

From: s22, s47E [Energy, North Ryde]
To: s22
Cc: s22, s47E [Energy, North Ryde]
Subject: RE: GISERA project - Northern Territory - List of chemicals used by onshore gas development that have environmental impacts
Date: Tuesday, 20 August 2019 1:27:00 PM
Attachments: [image001.png](#)

Hi s22

s22, s47E and I were wondering whether you have any response for us regarding the query below.

Regards,

s22, s47E

Duplicate Email - Removed

From: [REDACTED] s22
To: [REDACTED] s22, s47E ([Energy, North Ryde](#))
Cc: [REDACTED] s22, s47E ([Energy, North Ryde](#)); [REDACTED] s22, s47E ([Energy, Dutton Park](#)); [REDACTED] s22, s47E ([Energy, North Ryde](#)); [REDACTED] s22
Subject: RE: NT CSIRO GISERA August field trip - Tanumbirini leg
Date: Wednesday, 3 July 2019 2:26:54 PM
Attachments: [image001.jpg](#)
[image002.jpg](#)
[image003.jpg](#)
[image004.jpg](#)

[REDACTED] s47G(1)(a), s47, s45

[REDACTED] s47G(1)(a), s47, s45

From: [redacted] s22, s47E ([Energy, North Ryde](#))
To: [redacted] s22
[redacted] s22, s47E ([Energy, Dutton Park](#)); [redacted] s22, s47E ([Energy, North Ryde](#)); [redacted] s22, s47E ([L&W, Albury](#)); [redacted] s22, s47E ([Energy, Albury](#)); [redacted] s22
[redacted] s22, s47E ([Energy, Kensington WA](#))
Subject: With revised agenda - FW: Agenda & meeting papers - NT project fieldwork and logistics meeting on 14 June 2019
Date: Thursday, 13 June 2019 7:40:17 AM
Attachments: [redacted] s47G(1)(a), s47, s45

s47G(1)(a), s47, s45

