

Molecular analysis of *Oryctes rhinoceros* collected from Philippines (January 2017)

Sean Marshall and Aubrey Moore

May 2017



Report for University of Guam

Client Report Number: TBD

This report has been prepared for Univeristy of Guam, and is confidential to University of Guam and AgResearch Ltd. No part of this report may be copied, used, modified or disclosed by any means without their consent.

Every effort has been made to ensure this Report is accurate. However scientific research and development can involve extrapolation and interpretation of uncertain data, and can produce uncertain results. Neither AgResearch Ltd nor any person involved in this Report shall be responsible for any error or omission in this Report or for any use of or reliance on this Report unless specifically agreed otherwise in writing. To the extent permitted by law, AgResearch Ltd excludes all liability in relation to this Report, whether under contract, tort (including negligence), equity, legislation or otherwise unless specifically agreed otherwise in writing.

CONTENTS

Molecular analysis of *Oryctes rhinoceros* collected from Philippines (January 2017)

1. EXECUTIVE SUMMARY	4
2. BACKGROUND	5
3. METHODS	6
3.1 Collection and DNA extraction of CRB tissue	6
3.2 PCR-RFLP detection of CRB-G haplotype	6
3.3 PCR detection of OrNV infection	7
3.3 Histopathology detection of OrNV infection	7
4. RESULTS AND DISCUSSION	7
5. RECOMMENDATIONS	8
6. ACKNOWLEDGEMENTS	8
7. REFERENCES	9
8. APPENDICES	9

1. EXECUTIVE SUMMARY

Oryctes rhinoceros nudivirus (OrNV) has been effective in reducing *Oryctes rhinoceros* (coconut rhinoceros beetle; CRB) population levels and keeping them at low levels elsewhere in the Pacific for over 40 years. However, to date the CRB population that has invaded Guam (CRB-G haplotype) has so far proven recalcitrant to currently available OrNV isolates that are able to cause disease in other CRB populations. OrNV is known to be widespread across the Philippines, where CRB is considered as native, and the CRB-G haplotype has been identified from here. Native habitats provide a good opportunity to identify candidate OrNV isolates (and other potential biocontrol agents).

The objective of this project was to begin searching for new OrNV isolates that could be effective against the CRB-G haplotype. Identification of an OrNV isolate with good efficacy against CRB-G would assist in preventing further mortality and damage to coconut palms on CRB-G infested Pacific islands (such as Guam), and slow or halt its spread into other areas of the Pacific region.

CRB tissue samples were obtained from live CRB adults collected using both pheromone trapping and hand collection from the Philippines. Gross observation and molecular DNA analyses were used to determine the OrNV infection status and haplotype of individual specimens. The pheromone trapping results were disappointing, with only a single CRB adult caught; however, hand collection enabled a number of adults and larvae to be collected. From the CRB specimens collected in the Philippines, an OrNV infected CRB adult was identified. Isolation of OrNV from infected CRB-G tissue will be undertaken to prepare inoculum for testing efficacy against CRB-G.

Recommendations:

- Isolate live OrNV from the infected CRB-G specimen and begin characterizing its potential efficacy as a biocontrol agent CRB-G.
- It is still important to continue searching for OrNV isolates (and other potential biocontrol agents) in regions known to harbor CRB-G populations, particularly where CRB caused palm damage is not reported as an issue.
- As the current CRB lure does not appear to be efficient at attracting CRB-G, new lures that can attractant CRB-G needs to be identified.

2. BACKGROUND

The coconut rhinoceros beetle (CRB), *Oryctes rhinoceros*, is a major pest of coconut palm, oil palm and other palm species. Palms are damaged when adult beetles bore into the crowns of palms to feed on sap. Tree mortality occurs when beetles destroy the growing tip (meristem). Immature beetles (grubs) do no damage. They feed on dead, decaying vegetation in breeding sites. Preferred breeding sites are dead, standing coconut stems, and piles of decaying vegetation such those left behind by typhoons or after replanting of oil palm plantations. If a CRB population is not suppressed, it is possible for a positive feedback cycle to initiate whereby adult beetles kill massive numbers of palms, thereby generating more food for even more grubs which turn into adults which kill even more palms. An outbreak following this scenario occurred in the Palau Islands during the late 1940s resulting in about 50% coconut palms being killed by CRB throughout the archipelago and 100% mortality on some of the smaller islands (Gressitt, 1953).

Following 40 years of no geographical range expansion, CRB is on the move in the Pacific. CRB has been detected for the first time at several Pacific Island locations including Saipan (2006), Guam (2007), Port Moresby, Papua New Guinea (2010), Oahu, Hawaii (2013), and Honiara, Solomon Islands (2015). Eradication of CRB is extremely difficult, having been achieved only once, on Niuaupolu (Keppel) Island, an island with an area of only 16 km² belonging to the Kingdom of Tonga (Catley, 1969). Failing eradication, the usual response to CRB infestations during the second half of the 20th century was introduction of OrNV, the biological control agent of choice for this pest. OrNV attacks only *Oryctes rhinoceros* beetles, typically reducing CRB damage by up to 90% with population suppression lasting indefinitely (Bedford, 2013). OrNV is auto disseminated, meaning the pathogen is carried between breeding sites by CRB adults. Like many biocontrol agents, OrNV is density dependent, working best at high population densities. Highly pathogenic OrNV isolates suppress population growth to levels, which result in only minor damage.

Current invasions of Pacific Islands by CRB involve a new invasive biotype (CRB-G) that has escaped from biological control by OrNV (Marshall et al.). Discovery of *Oryctes rhinoceros* nudivirus in the 1960s enabled the successful management of populations in Pacific Island Countries. Augmentative release of OrNV continues to be an important mechanism for CRB management in both coconut and oil palm growing regions. For 40 years after adoption of this biocontrol strategy, no new outbreaks of CRB were reported from uninfested palm growing islands in the Pacific ensuring continuity of palm based village economies. However, the situation has recently changed. For the first time in 40 years, CRB invasion into completely new areas has been reported. Additionally, Pacific areas with established CRB populations (e.g. Palau) have reported increased severity and frequency of CRB damage. Common to all these areas is the high incidence of severe palm damage not seen since the introduction of OrNV. Initial attempts to introduce OrNV into the Guam CRB population (CRB-G) were unexpectedly unsuccessful, raising the possibility that the CRB-G population that invaded Guam become tolerant or resistant to the commonly applied OrNV isolates. Analysis of several CRB populations has demonstrated that, in addition to Guam, the CRB-G biotype is also found in Hawaii, Palau, and most recently in Port Moresby (PNG) and Honiara (Solomon Islands) during 2015.

Uncontrolled infestations of CRB may kill most palms within a few years, as it currently being observed. A worse case scenario can be triggered by a massive outbreak of adult CRB emerging from abundant breeding sites made by large amounts of decaying vegetation left in the wake of a typhoon. This is the situation in Guam, which was visited by Typhoon Dolphin in May, 2015. Very high feeding activity will kill mature coconut palms, leaving standing dead coconut trunks that are ideal breeding sites for subsequent generations of beetles. During a CRB outbreak, there will be an increased risk of further spread to uninfested islands throughout the Pacific. Palms are important on Pacific Islands for various reasons: as a cash crop for nuts, oil and lumber, as an ornamental tree appreciated by residents and tourists. On some of the smaller, more traditional islands the coconut palm is referred to as the tree of life. Here, this species is an essential natural resource

providing income, housing, food, oil, soap, clothing, mats, baskets, and other containers. The smaller, poorer Pacific islands will suffer the most if the spread of CRB-G cannot be controlled.

The objective of this project was to begin searching for new OrNV isolates that could be effective against the CRB-G haplotype. Native habitats provide a good opportunity to identify candidate OrNV isolates (and other potential biocontrol agents). OrNV is known to be widespread across the Philippines, where CRB is considered as native, and the CRB-G haplotype has been identified from here. Identification of an OrNV isolate with good efficacy against CRB-G would assist in preventing further mortality and damage to coconut palms on CRB-G infested Pacific islands (such as Guam), and slow or halt its spread into other areas of the Pacific region.

3. METHODS

3.1 Collection and DNA extraction of CRB tissue

CRB specimens were collected using both hand collection (larvae and adults) and 'DeFence Traps' (hanging Tekken fish gill netting) fitted with the CRB attractant, Oryctalure (www.chemtica.com). Gut tissue samples from live CRB specimens were dissected and used for DNA extraction because decaying tissue is often of limited value in terms of DNA quality. To ensure DNA quality was maintained, a 0.5 - 1 cm piece of the midgut tissue was submerged in monopropylene glycol (PPG), and stored at -20°C.

DNA was extracted from CRB tissue using a ZR Genomic DNA Tissue MiniPrep (Zymo Research) kits. DNA elution was carried out using 100 µl of elution buffer from the appropriate kit, and aliquots of eluted DNA samples were subsequently used for further analyses.

3.2 PCR-RFLP detection of CRB-G haplotype

The primers used to amplify a 523 base pair (bp) fragment of the CRB COI gene were C1-J-1718Oryctes (5'- GGAGGTTTCGGAAATTGACTTGTTCC -3') and C1-N-2191Oryctes (5'- CCAGGTAGAATTAATAATRTATACCTC -3') (Marshall et al.). Each 25 µl PCR reaction constituted 0.125 µl i-StarTaq DNA Polymerase (iNtRON Biotechnology), 2.5 µl 10x PCR buffer (iNtRON Biotechnology), 0.5 µl dNTP mixture (10 mM), 0.5 µl C1-J-1718Oryctes (10 µM), 0.5 µl C1-N-2191Oryctes (10 µM), 2 µl diluted (1 in 50) DNA template, and 18.75 µl water. PCR amplifications were performed in a C2100 (BioRad) thermocycler with a cycling profile of 35 cycles of 94°C denaturation (30 s), 50°C annealing (45 s), 72°C extension (1 min) with an initial denaturation of 3 min at 94°C and a final extension of 5 min at 72°C. A 5 µl aliquot of each PCR reaction was checked by agarose gel electrophoresis (1%, 0.5xTBE), stained with RedSafe (iNtRON Biotechnology) and fluorescence visualized over UV light. Photographs were recorded using an UVIdoc HD2 gel doc (UVItech). For RFLP analysis, successfully amplified COI PCR products (8 µl) were each combined with 0.2 µl Mse1 (10U/µl; New England BioLabs, NEB), 1 µl 10x NEB Buffer#4, 0.1 µl 100x NEB BSA and 10.7 µl water, and incubated at 37°C for 3 h. Digested samples (10 µl) were mixed with DNA loading dye, loaded onto on a 2% agarose gel in 0.5xTBE buffer. The gel was electrophoresed using 60 V for 1.5 h, stained with RedSafe and DNA fluorescence detected over UV light. Photographs were taken using an UVIdoc HD2 gel doc.

3.3 PCR detection of OrNV infection

The PCR protocol for detection of OrNV was based on that described in Richards et al. (Richards et al., 1999), and has been subsequently modified to distinguish infection from mere presence (e.g. incidental contact contamination). The primer pairs used to amplify a 945 base pair (bp) fragment of the OrNV genome were OrNV15a (5'-ATTACGTCGTAGAGGCAATC-3') and OrNV15b (5'-ATGATCGATTCGTCTATGG-3')(Richards et al., 1999). Each 25 µl PCR reaction contained 0.2 µl i-StarTaq DNA Polymerase (iNtRON Biotechnology), 2.5 µl 10x PCR buffer (iNtRON Biotechnology), 0.5 µl dNTP mixture (10 mM), 0.5 µl OrNV15a (10 µM), 0.5 µl OrNV15b (10 µM), 1 µl diluted DNA (paired reactions of 1 in 100 and 1 in 5000), and 19.8 µl water. PCR amplifications were performed in a C2100 (BioRad) thermocycler with a cycling profile of 35 cycles of 94°C denaturation (30 s), 50°C annealing (45 s), 72°C extension (1 min) with an initial denaturation of 3 min at 94°C and a final extension of 5 min at 72°C. An 8 µl aliquot of each PCR reaction was separated by agarose gel electrophoresis (1%, 0.5xTBE), stained with RedSafe (iNtRON Biotechnology) and fluorescence visualized over UV light. Photographs were recorded using an UVIdoc HD2 gel doc (UVItech). Detection of OrNV PCR product in the 1 to 5000 dilution was considered indicative of OrNV infection (unpublished data; validated in combination with gross visual inspection and histological analysis based on pathology description (Huger, 2005)).

3.3 Histopathology detection of OrNV infection

Histopathology studies of tissue displaying symptoms of OrNV infection. Dissected midgut tissue was fixed for in a 10% neutral buffered formalin solution. Standard paraffin embedding, serial sectioning and hematoxylin and eosin staining methods were carried out on samples by Gribbles Veterinary Services (Christchurch, New Zealand). Stained slides of the alimentary tract were examined under brightfield and differential interference contrast (DIC) optics using an Olympus BX50 upright microscope and photographed with an Olympus DP-72 digital camera.

4. RESULTS AND DISCUSSION

A total of 214 CRB larvae and adults were collected from within the Dumagate - Dauin region of Negros Island, Philippines. Tissue from 94 individuals was preserved for further analysis (see Appendix 1 for specimen details). Only one specimen (a hand caught adult female; Dug-42) displayed the symptoms of OrNV infection (thick, milky coloured gut; see Huger, 2005). From this pool of preserved tissue specimens, 16 representative samples (including Dug-42) were selected for molecular analysis to determine haplotype and OrNV infection status. A summary of the results is presented in Table 1. Amplification of the partial COI gene failed for specimens Dug-55, -56, and -57; therefore, the CRB-G haplotype status could not be determined from these specimens. As COI is present in all host cells, this meant that the OrNV infection status could also not be resolved, despite the observation of OrNV DNA in specimens Dug-55 and -. For the remaining 13 specimens, all were of the CRB-G haplotype, while only Dug-42 was positive for OrNV infection. Histopathology examination of Dug-42 gut tissue revealed diagnostic indicators of OrNV infection as described in Huger (Huger, 2005).

Since three separate methods indicated OrNV presence, we conclude that the Dug-42 tissue was infected with OrNV. Isolation of OrNV from Dug-42 tissue has been undertaken, and early results from testing this material on the OrNV permissive cell line DSIR-Ha-1179 have shown the expected cytopathic effects (data not shown). If the early cell culture results are confirmed, the next step will be to conduct a pathogen challenge assay using the Dug-42 OrNV isolate against CRB-G.

Table 1: Summary of haplotype and OrNV infection status for CRB collected from Negros Island, Philippines.

Specimen	COI PCR	OrNV PCR	Virus present	Haplotype
Dug-1	+	-	No	CRB-G
Dug-2	+	-	No	CRB-G
Dug-16	+	-	No	CRB-G
Dug-17	+	-	No	CRB-G
Dug-18	+	-	No	CRB-G
Dug-19	+	-	No	CRB-G
Dug-42	+	+	Yes	CRB-G
Dug-43	+	-	No	CRB-G
Dug-44	+	-	No	CRB-G
Dug-45	+	-	No	CRB-G
Dug-55	-	+	Undetermined	Undetermined
Dug-56	-	+	Undetermined	Undetermined
Dug-57	-	-	Undetermined	Undetermined
Dug-62	+	-	No	CRB-G
Dug-63	+	-	No	CRB-G
Dug-69	+	-	No	CRB-G

5. RECOMMENDATIONS

- Isolate live OrNV from the infected CRB-G specimen and begin characterizing its potential efficacy as a biocontrol agent CRB-G.
- It is still important to continue searching for OrNV isolates (and other potential biocontrol agents) in regions known to harbor CRB-G populations, particularly where CRB caused palm damage is not reported as an issue.
- As the current CRB lure does not appear to be efficient at attracting CRB-G, new lures that can attractant CRB-G needs to be identified.

6. ACKNOWLEDGEMENTS

We thank Ian Iriarte for his assistance in collection of CRB specimens and dissection of tissue. We also thank Bonito for sharing his invaluable local knowledge and in helping with the collection of CRB specimens. This project work was funded through a USDA Farm Bill grant.

7. REFERENCES

- Bedford, G. O., 2013. Biology and management of palm dynastid beetles: Recent advances. Annual Review of Entomology. 58, 353-372.
- Catley, A., 1969. The Coconut Rhinoceros Beetle *Oryctes rhinoceros* (L)[Coleoptera: Scarabaeidae: Dynastinae]. Pest Articles & News Summaries. 15, 18-30.
- Gressitt, J. L., 1953. The Coconut Rhinoceros Beetle (*Oryctes rhinoceros*) with particular reference to the Palau Islands. Bernice P. Bishop Museum Bulletin. 212, 1-157.
- Huger, A. M., 2005. The *Oryctes* virus: Its detection, identification, and implementation in biological control of the coconut palm rhinoceros beetle, *Oryctes rhinoceros* (Coleoptera: Scarabaeidae). Journal of Invertebrate Pathology. 89, 78-84.
- Marshall, S. D. G., Moore, A., Vaqalo, M., Noble, A., Jackson, T. A., Under Review. A new haplotype of the coconut rhinoceros beetle, *Oryctes rhinoceros*, has escaped biological control by *Oryctes rhinoceros* nudivirus and is invading Pacific islands. Journal of Invertebrate Pathology.
- Richards, N. K., Glare, T. R., Aloali'i, I., Jackson, T. A., 1999. Primers for the detection of *Oryctes* virus from Scarabaeidae (Coleoptera). Molecular Ecology. 8, 1552-1553.

8. APPENDICES

Appendix 1: Details of CRB tissue specimens collected from Philippines in January 2017.

Date	Specimen ID	Site	Collection Method	Stage/Gender	Condition	PPG	FAA	Live
26/01/2017	Du1	Farm log2	Fallen log By hand	L3 -large	slow, reddish colour	1	1	1
26/01/2017	Du2	Farm log2	Fallen log By hand	L3 -large	slow, red all over (almost purple)	1	1	1
26/01/2017	Du3	Farm log2	Fallen log By hand	L3 -large	slow, reddish colour, stripped, a little purple	1	1	1
26/01/2017	Du4	Farm log2	Fallen log By hand	L3 -large	slow, reddish colour	1	1	1
26/01/2017	Du5	Farm log2	Fallen log By hand	L3 -small	slow, reddish colour	1	1	1
26/01/2017	Du6	Farm log2	Fallen log By hand	L3 -small	normal	1	0	0
26/01/2017	Du7	Farm log2	Fallen log By hand	L3 -small	normal	1	0	0
26/01/2017	Du8	Farm log2	Fallen log By hand	L2	normal	1	0	0
26/01/2017	Du9	Farm log2	Fallen log By hand	L2	normal	1	0	0
26/01/2017	Du10	Farm log2	Fallen log By hand	L3 -small	normal	1	0	0
26/01/2017	Du11	Farm log2	Fallen log By hand	L3 -small	normal	1	0	0
26/01/2017	Du12	Farm log2	Fallen log By hand	L3 -small	normal	1	0	0
26/01/2017	Du13	Farm log2	Fallen log By hand	L3 -small	normal	1	0	0

26/01/2017	Du14	Farm log2	Fallen log By hand	L3 -small	normal	1	0	0
26/01/2017	Du15	Farm log2	Fallen log By hand	L3 -small	normal	1	0	0
26/01/2017	Du16	Lumber yard, Sibulan	Sawdust By hand	Male	vigorous, active, fatty, brown medium gut	1	1	1
26/01/2017	Du17	Lumber yard, Sibulan	Sawdust By hand	Male	reasonable active, not fatty, thin light brown gut	1	1	1
26/01/2017	Du18	Farm log1	Fallen log By hand	L3	normal	1	0	0
26/01/2017	Du19	Farm log1	Fallen log By hand	L3	normal	1	0	0
26/01/2017	Du20	Farm log1	Fallen log By hand	L3	normal	1	0	0
26/01/2017	Du21	Farm log1	Fallen log By hand	L3	normal	1	0	0
26/01/2017	Du22	Farm log1	Fallen log By hand	L3	normal	1	0	0
26/01/2017	Du23	Farm log1	Fallen log By hand	L3	normal	1	0	0
26/01/2017	Du24	Farm log1	Fallen log By hand	L3	normal	1	0	0
26/01/2017	Du25	Farm log1	Fallen log By hand	L3	normal	1	0	0
26/01/2017	Du26	Lumber yard, Sibulan	Sawdust By hand	L3 -small	flacid	1	1	1
26/01/2017	Du27	Lumber yard, Sibulan	Sawdust By hand	L3 -small	flacid	1	1	1
26/01/2017	Du28	Lumber yard, Sibulan	Sawdust By hand	prepupa	flacid, very fatty, lots liquid	1	1	1
26/01/2017	Du29	Lumber yard, Sibulan	Sawdust By hand	prepupa	very fatty	1	1	1
26/01/2017	Du30	Lumber yard, Sibulan	Sawdust By hand	L3 -large	soft flacid posterior, firm anterior (white tissue)	1	1	1
26/01/2017	Du31	Lumber yard, Sibulan	Sawdust By hand	prepupa	very fatty	1	1	1
26/01/2017	Du32	Lumber yard, Sibulan	Sawdust By hand	L3 -large	normal	1	1	1
26/01/2017	Du33	Lumber yard, Sibulan	Sawdust By hand	L3 -large	normal	1	1	1
26/01/2017	Du34	Lumber yard, Sibulan	Sawdust By hand	L3 -small	normal	1	0	0
26/01/2017	Du35	Lumber yard, Sibulan	Sawdust By hand	L3 -large	normal	1	0	0
26/01/2017	Du36	Lumber yard, Sibulan	Sawdust By hand	L3 -large	normal	1	0	0
26/01/2017	Du37	Lumber yard, Sibulan	Sawdust By hand	L3 -large	normal	1	0	0
26/01/2017	Du38	Lumber yard, Sibulan	Sawdust By hand	L3 -large	normal	1	0	0

26/01/2017	Du39	Lumber yard, Sibulan	Sawdust By hand	L3 -small	normal	1	0	0
26/01/2017	Du40	Lumber yard, Sibulan	Sawdust By hand	L3 -large	normal	1	0	0
26/01/2017	Du41	Lumber yard, Sibulan	Sawdust By hand	L3 -large	normal	1	0	0
28/01/2017	Du42	Grandma home	window by hand	Female	live, fatty, thick white gut	1	1	1
28/01/2017	Du43	Grandma home	window by hand	Male	live, thin brown gut (kept in same bottle as #42 for 2 days)	1	1	1
28/01/2017	Du44	Bascofadco coop, Bacong	By hand	Male	live, thin brown gut	1	1	1
28/01/2017	Du45	Bascofadco coop, Bacong	By hand	Male	live, thin brown gut	1	1	1
28/01/2017	Du46	Bascofadco coop, Bacong	By hand	Male	live, thin brown gut	1	1	1
30/01/2017	Du47	Ton home, Dauin	stumps/logs By hand	Male	live, fatty, thick brown gut	1	1	1
30/01/2017	Du48	Ton home, Dauin	stumps/logs By hand	Teneral Male	live, fatty, thick brown gut	1	1	1
30/01/2017	Du49	Ton home, Dauin	stumps/logs By hand	Male	live, thick brown gut	1	1	1
30/01/2017	Du50	Ton home, Dauin	stumps/logs By hand	Teneral Male	live, fatty, thick brown gut	1	0	1
30/01/2017	Du51	Ton home, Dauin	stumps/logs By hand	Male	live, thin brown gut	1	0	1
30/01/2017	Du52	Ton home, Dauin	stumps/logs By hand	Female	live, thin brown gut	1	0	1
30/01/2017	Du53	Ton home, Dauin	stumps/logs By hand	Male	live, thick brown gut	1	0	1
30/01/2017	Du54	Ton home, Dauin	stumps/logs By hand	Female	live, thin brown gut	1	0	1
30/01/2017	Du55	Ton home, Dauin	stumps/logs By hand	L3	found dead in breeding site log	1	0	1
30/01/2017	Du56	Ton home, Dauin	stumps/logs By hand	L2 (x6)	composite sample	1	0	1
30/01/2017	Du57	Ton home, Dauin	stumps/logs By hand	L3	found dead in log	1	0	1
28/01/2017	Du58	Bascofadco coop, Bacong	Fallen log by hand	L3	Healthy looking	1	0	0
28/01/2017	Du59	Bascofadco coop, Bacong	Fallen log by hand	L3	Healthy looking	1	0	0
28/01/2017	Du60	Bascofadco coop, Bacong	Fallen log by hand	L3	Healthy looking	1	0	0
28/01/2017	Du61	Bascofadco coop, Bacong	Fallen log by hand	L3	Healthy looking	1	0	0
1/02/2017	Du62	Aubrey's, Lunga	balcony light	Female	live, thin brown gut	1	1	1
1/02/2017	Du63	Aubrey's, Lunga	Pheromone trap	Male	live, thick brown gut	1	1	1

31/01/2017	Du64	palm plantation, San Jose	stumps/logs By hand	Female	live, thin brown gut	1	1	1
31/01/2017	Du65	palm plantation, San Jose	stumps/logs By hand	Female	live, thin brown gut	1	1	1
31/01/2017	Du66	palm plantation, San Jose	stumps/logs By hand	Male	live, thin brown gut, very fatty	1	1	1
31/01/2017	Du67	palm plantation, San Jose	stumps/logs By hand	Male	live, thin brown gut	1	1	1
31/01/2017	Du68	palm plantation, San Jose	stumps/logs By hand	Male	live, thin brown gut	1	1	1
31/01/2017	Du69	palm plantation, San Jose	stumps/logs By hand	Female	live, thin light brown gut ,	1	1	1
31/01/2017	Du70	palm plantation, San Jose	stumps/logs By hand	Male	live, thin brown gut, fatty	1	1	1
31/01/2017	Du71	palm plantation, San Jose	stumps/logs By hand	Male	live, thin brown gut	1	1	1
31/01/2017	Du72	palm plantation, San Jose	stumps/logs By hand	Female	live, thin brown gut	1	1	1
31/01/2017	Du73	palm plantation, San Jose	stumps/logs By hand	Male	live, thin brown gut	1	1	1
31/01/2017	Du74	palm plantation, San Jose	stumps/logs By hand	Female	live, thin brown gut	1	1	1
31/01/2017	Du75	palm plantation, San Jose	stumps/logs By hand	Male	live, thin brown gut	1	1	1
31/01/2017	Du76	palm plantation, San Jose	stumps/logs By hand	Male	live, medium brown gut	1	1	1
31/01/2017	Du77	palm plantation, San Jose	stumps/logs By hand	Female	live, thin dark brown gut	1	1	1
31/01/2017	Du78	palm plantation, San Jose	stumps/logs By hand	Female	live, thin dark brown gut	1	1	1
31/01/2017	Du79	palm plantation, San Jose	stumps/logs By hand	Female	live, thin dark brown gut	1	1	1
31/01/2017	Du80	palm plantation, San Jose	stumps/logs By hand	Female	live, thin brown gut	1	1	1
31/01/2017	Du81	palm plantation, San Jose	stumps/logs By hand	Male	live, thin brown gut	1	1	1
31/01/2017	Du82	palm plantation, San Jose	stumps/logs By hand	Female	live, thin brown gut	1	1	1
31/01/2017	Du83	palm plantation, San Jose	stumps/logs By hand	Male	live, thin brown gut , fatty	1	1	1
1/02/2017	Du84	Ton home, Dauin	stumps/logs By hand	Male	live, thin dark brown gut	1	1	1
1/02/2017	Du85	Ton home, Dauin	stumps/logs By hand	Female	live, thin brown gut	1	1	1
1/02/2017	Du86	Ton home, Dauin	stumps/logs By hand	Female	live, thin dark brown gut	1	1	1
1/02/2017	Du87	Ton home, Dauin	stumps/logs By hand	Female	live, thin brown gut	1	1	1

1/02/2017	Du88	Ton home, Dauin	stumps/logs By hand	Male	live, thin brown gut	1	1	1
1/02/2017	Du89	Ton home, Dauin	stumps/logs By hand	Female	live, thin dark brown gut	1	1	1
1/02/2017	Du90	Ton home, Dauin	stumps/logs By hand	Male	live, thin brown gut	1	1	1
1/02/2017	Du91	Ton home, Dauin	stumps/logs By hand	Female	live, thin brown gut , fatty	1	1	1
1/02/2017	Du92	Ton home, Dauin	stumps/logs By hand	Male	live, thin brown gut	1	1	1
1/02/2017	Du93	Ton home, Dauin	stumps/logs By hand	Female	live, thin light brown gut	1	1	1
1/02/2017	Du94	Ton home, Dauin	stumps/logs By hand	pupa (male)	all white (messy inards)	1	1	1