The presence of a mycangium in European *Sinodendron cylindricum* (Coleoptera: Lucanidae) and the associated yeast symbionts

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Abstract

Part of the exoskeleton of some wood-inhabiting insects is modified to form a mycangium, which is a specialized organ used to convey fungal spores or yeasts to their offspring. Although most stag beetles (Coleoptera: Lucanidae) are known to have female-specific mycangia and associated yeast symbionts, the evolutionary origin of the mycangium in this group remains unresolved. Here, we report the presence of a mycangium and associated yeast symbionts in the European horned stag beetle *Sinodendron cylindricum* (L.), which belongs to an ancestral clade of the Lucanidae. The mycangium of *S. cylindricum* is shown to be female-specific and have the same developmental origin as that of other stag beetles. A total of five yeast strains were isolated from adult mycangia and larval gut of *S. cylindricum*. Of these, we suggest that SICYAM1 is an undescribed yeast with taxonomic novelty, and have identified SICYLG3 as the xylose-fermenting yeast *Scheffersomyces insectosa* using nuclear ribosomal RNA and ITS sequences. The remaining three yeast strains, SICYAM2, SICYLG1, and SICYLG2, were assigned to the genus *Sugiyamaella*. Yeast density in the adult mycangium was lower than that of the more evolutionarily advanced stag beetles, the European *Lucanus cervus* (L.) and *Dorcus parallelipipedus* (L.), which were also examined in this study. No living yeasts were isolated from the adult guts. However, a third instar larva of *S. cylindricum* harbored 10^4–10^6 living yeasts in each gut region, which suggests that gut yeasts play an important role in these wood-feeding larvae.

Key words: evolution, gut flora, insect–fungal associations, saproxylic insects, vertical transmission

The mycangium is an exoskeletal cavity or saccate structure that harbors symbiotic fungi or yeasts, and can be found in some wood-boring beetles and bark beetles, wood wasps, leaf-rolling weevils, and a bamboo-inhabiting lizard beetle (Batra 1963, Madden and Coutts 1979, Beaver 1989, Six 2003, Kobayashi et al. 2008, Toki et al. 2012, Davis 2015). Recently, a female-specific mycangium was independently discovered in the stag beetle *Lucanus cervus* (L.) (Coleoptera: Lucanidae) (Hawes 2009, 2010, 2013) and 12 genera of Lucanidae in Japan (Tanahashi et al. 2010). Yeast symbions present in the mycangium of *Lucanus* and *Dorcus* species are closely related to the xylose-fermenting yeasts, *Scheffersomyces stipitis* (Pignal) Kurtzman et Suzuki (formerly, *Pichia stipitis*) (Tanahashi et al. 2010, Hawes 2013, Tanahashi and Fremlin 2013). Xylose-fermenting yeasts are commonly found in the digestive tracts and/or feeding tunnels of many xylophagous insects, suggesting an association with wood digestion (Suh et al. 2003, 2006). Stag beetle larvae of some genera feed on subterranean, decaying wood of trees and shrubs, or subterranean woody material of humus-rich soil, while others feed on moist, decaying wood above soil level (Holloway 2007), whereas adults utilize food material that is more nutritive such as fermented tree sap and over-ripe fruits, however a few species are known to exhibit carnivory (Mori and Chiba 2009), and some rarely feed after eclosion (Sadaki et al. 2014). Thus, the biochemical environment of the adult gut is likely to be very different from that of the wood-feeding larvae, where xylose-fermenting yeasts play an essential role. Completely separate from the digestive tract, the saccate mycangium provides a reservoir for yeast symbions (Tanahashi et al. 2010, Fremlin and Tanahashi 2015). Mycangium evolution probably occurred only once in the ancestral stag beetle clade and might have been the starting point for the evolution of the larval and adult dimorphic food habits.
The presence of a mycangium has been reported in nine genera of the subfamily Lucaninae (Dorcus, Prosopocoilus, Aegus, Lucanus, Neolucanus, Pismogonathus, Figulus, Nigidius, and Platycerus), two genera of Aesalinae (Aesalus and Nicagus), and one genus of Sindsinai (Ceruchus). Of the Scarabaeoidea, mycangia have only been found in the family Lucanidae. Consequently, the distribution of mycangia on the phylogenetic tree is markedly similar to the phylogenetic distribution of the family Lucanidae (Krajcik 1868, Arrow 2005), possibly to deter predators, parasitizing the burrow entrance after the female has completed oviposition then packed with wood particles. A male will often continue guarding. Some 20 eggs are laid in tunnel branches, each of which is excavated a burrow in decaying wood. Perhaps uniquely, the male guards the burrow and removes the debris while the female is excavating. Because some supposedly ancestral genera have not been investigated for the presence of a mycangium, together with the phylogenetic ambiguity in the ancestral clades of Lucanidae, the evolutionary origin of the mycangium remains unclear.

The genus Sinodendron is currently placed in the subfamily Syndesinae, which is described as one of the ancestral clades of Lucanidae (Kim and Farrell 2015). Sinodendron species are found in the United Kingdom, Europe, and North America, where they are known as ‘rhinoceros beetles’ or ‘horned stag beetles’ due to the characteristic, prominent horn on the head of the males. Of these, only the species Sinodendron cylindricum (Linneaus, 1758) is present in Europe and the United Kingdom, where its distribution is widespread (Zahradnik and Chvala 1989, Alexander 2002, Sutton 2003, Klausnitzer and Sprecher-Uebersax 2008, GBIF 2016, NBN 2016). Glossy-black, 10–18-mm long and coarsely punctate, the adult beetles are active nocturnally and often found gathering around fermented sap of hardwood trees. Both sexes collaborate to excavate a burrow in decaying wood. Perhaps uniquely, the male guards the burrow and removes the debris while the female is excavating. Some 20 eggs are laid in tunnel branches, each of which is then packed with wood particles. A male will often continue guarding the burrow entrance after the female has completed oviposition (Chapman 1868, Arrow 2005), possibly to deter predators, parasitoids, and/or conspecific competitors. Larvae feed on the decaying wood and pupate within it.

As adult and larval feeding behavior of S. cylindricum is similar to that of other stag beetle species, we hypothesized that (1) S. cylindricum females have a specialized mycangium, similar to that of other stag beetle species, or (2) females and/or males store symbiotic microorganisms in their digestive tracts. In this study, we examined S. cylindricum for (1) presence or absence of a mycangium in females and males, (2) presence or absence of yeast symbionts in the mycangium, (3) presence or absence of yeast symbionts in the adult digestive system, and (4) presence or absence of yeast symbionts in the larval digestive system. In addition, (5) we analyzed the yeasts from the European stag beetles, Dorcus parallelipipededus and L. cervus, which are known to possess a female-specific mycangium and associated xylose-fermenting Scheffersomyces yeasts (Hawes 2013, Tanahashi and Fremlin 2013).

Materials and Methods

Isolation of Yeasts From Adult Insects

Two adult males and two adult females of S. cylindricum (Fig. 1a and b) were collected from decaying logs of Fagus sylvatica L. or Carpinus betulus L. at Grandfontaine, Bas-Rhin, France (48.49°N, 7.16°E) on 4 September 2014. Specimens were anesthetized using gaseous carbon dioxide, then dissected in sterile phosphate-buffered saline (PBS) and examined for the presence or absence of a mycangium, as described in Tanahashi et al. (2010). The mycangium found in both females was removed from its underlying tissue and weighed using an electronic microbalance (MTS, Mettler Toledo). The midgut and hindgut of both sexes were also removed and weighed. These organs were placed separately in 1.5 ml tubes containing 160 µl PBS and then homogenized using pellet pestles (Kimble Kontes, Fischer Science). The total volume was adjusted to 200 µl by adding PBS, and a fivefold dilution series was made by transferring 40 µl of the suspension to a new 1.5 ml tube containing 160 µl of PBS. For quantitative isolation of yeasts, 40 µl of each dilution (i.e., 1/5, 1/25, ... , 1/5² equivalent of the original homogenate) was loaded separately onto five potato dextrose agar (PDA) (BD) plates (9-cm diameter), which contained 50 µg/ml rifampicin, and spread uniformly using a glass spreader. The plates were incubated at 20°C for 4 d.

To compare the density and diversity of yeast symbionts in the mycangium between different insect species, we analyzed three adult female Dorcus parallelipipededus (L.), one of which was collected from sycamore (Acer pseudoplatanus L.) tree stump in Bentley, Ipswich, Suffolk, United Kingdom (51.988°N, 1.076°E, 40 m alt.) on 12 March 2014 and others which were collected from a decaying log in Basel, Switzerland (47.553N, 7.604E, 317 m alt.) on 16 June 2014. A newly emerged female of L. cervus collected in Stutton, Ipswich, Suffolk, United Kingdom (51.970N, 1.134E, 9.5 m alt.) on 10 June 2011 was also added to the analysis.

Isolation of Yeasts From Larvae

Four lucanid-like larvae (Fig. 1c) were collected from a decaying beech log at the northern slope of Lägern, Zurich, Switzerland (47.48°N, 8.39°E, 850 m alt.) on 15 June 2014. As Lägern is a protected area, insect collection was permitted as part of the field survey program in the 8th Symposium on the Conservation of Saproxylic Beetles (Basel, Switzerland, 13–15 June 2014). The larvae were maintained in the same decaying wood at 10–15°C until they were required for examination. For isolation of yeasts, a healthy larva was washed with distilled water and dissected in sterile PBS. The whole gut was removed and washed twice with sterile PBS. The midgut, hindgut and cecum-like sacs were separated from the whole gut, weighed and then homogenized in 1.5-ml tubes. The total volume in each tube was adjusted to 200 µl and a tenfold dilution series made by transferring 20 µl of the suspension to a new 1.5 ml tube containing 180 µl PBS. Finally, 20 µl of each dilution was spread on a PDA plate, as described above. The plates were incubated at 20°C for 4 d. The remaining three larvae were reared to adults for future analysis.

Quantification of Yeasts

The yeast colonies from adults and a larva were roughly classified according to their morphological traits (morphotype), and each morphotype colony was counted on every plate. Sixteen colonies were randomly selected for each morphotype from each isolation source and these were transferred to new PDA plates for later DNA analysis. Although one morphotype is likely to contain multiple species or strains, such pre-classification is helpful in elucidating species diversity, especially when there are minor but morphologically different colonies. Colony forming unit (CFU) was calculated for each morphotype by using the plate where an adequate number of colonies were collected in Stutton, Ipswich, Suffolk, United Kingdom (51.970N, 1.134E, 9.5 m alt.) on 10 June 2011.

DNA Sequencing and Haplotyping of Yeasts

For DNA extraction, small pellets of the isolated yeast colonies were suspended in 50 µl lyticase solution (0.4 U/µl lyticase and 50 mM EDTA) and incubated at 37°C for 2 h. The spheroplast yeast cells were collected by centrifugation and then processed using Wizard (R) Genomic DNA Purification Kit (Promega). For the first DNA screening analysis, internal transcribed spacer (ITS) regions of the
nucleus ribosomal RNA (rRNA) gene were amplified by PCR using the primer pair NS7-NL4 (White et al. 1990), which covers the full length of ITS1, 5.8S, and ITS2. The PCR fragments were sequenced with an internal primer ITS5 to obtain complete nucleotide sequences of those regions. Both ends of the sequences were trimmed to contain only those three ITS regions. The total length of the trimmed sequence varied from 463 to 547 bases. ITS haplotype was determined for each isolate according to the trimmed sequence.

Fig. 1. Adults and a larva of *S. cylindricum*. (a) Male, (b) female, (c) third instar larva, (d) sampling locations in France and Switzerland. (e, f) Male reproductive organs and the absence of mycangium. Arrow indicates the location of a mycangium in a female. (g, h) Female reproductive organs and the presence of mycangium (arrows). Removed hindgut is indicated by dotted lines. (i, j) Larval gut and cecum-like sacs. ag, accessory gland; cs, cecum-like sac; fg, foregut; hg, hindgut; mcg, mycangium; mg, midgut; mpt, Malpighian tubule; ov, ovary; ovd, (main) oviduct; sp, spermatheca; spg, spermathecal gland; ts, testis. Scale bars: (c, i) 5 mm; (d) 100 km; (e–h, j) 1 mm.
the second intensive DNA analysis, a representative isolate was randomly chosen for each ITS haplotype for each insect specimen (i.e., isolates that had the same ITS haplotype from the same specimen were assumed to be the same strain, whereas those from a different specimen were treated as different strains). Approximately 2,900 bp of the nucleus ribosomal RNA genes, including 18S rRNA, ITS1, 5.8S rRNA, ITS2, and 26S rRNA, were amplified by PCR with NS1- 

FS2 and NS7-NL4 primer sets (White et al. 1990), which were designed to have an overlapping region of 170 bases. The PCR fragments were sequenced with multiple sequencing primers; NS1, NS2, NS3, F2, NS7, ITS5, and NLI (White et al. 1990). These short nucleotide sequences (~700 bases) were combined on ContigExpress (Invitrogen, USA) software.

DNA Sequencing of Reference Yeasts

Type strains of five yeast species, S. stipitis JCM 10742, Scheffersomyces segobiiensis (Santa Maria et Garcia) Kurtzman et Suzuki JCM10740, Scheffersomyces shebatae (Buckley et van Uden) Urbina et Blackwell JCM9840, Scheffersomyces insectosa (Kurtzman) Urbina et Blackwell JCM9842, and Sugiyamaella nosakii (Peter et al.) Urbina et Blackwell ATCC 201508, and an identified yeast strain Wickerhamomyces anomalus (Hansen) Kurtzman et al. DBL01s1Shirosato (Toki et al. 2012) were subjected to the sequencing analysis as described above, since there were no continuous, long sequences from 18S to 26S in the GenBank database at that time. Yeasts that had been isolated from the adult mycangium of two Japanese lucanid species D. rectus (Motschulsky) and L. maculifemoratus Motschulsky (Tanahashi et al. 2010) were sequenced in the same way.

Identification of Yeast Species by BLAST

Even though we obtained relatively long, continuous sequences of the rRNA genes of the yeasts, most of the yeast sequences in the GenBank database are short (~600 bases). In this situation, BLAST search usually scoops up longer but weaker-matching sequences (typically, genome sequences of several well-known yeasts), when using a long query sequence with default search parameters. Therefore, we divided our sequence data into three regions, namely, 18S, ITS, and D1/D2, which have been traditionally used for identification and phylogenetic analysis of yeasts. The three regions were then separately subjected to BLAST search (Altschul et al. 1997) using the procedure obtained from the NCBI website (http://blast.ncbi.nlm.nih.gov/Blast.cgi).

Carbon Assimilation Test

The representative isolates were cultured aerobically in 4 ml of yeast nitrogen base (YNB) (BD) containing 0.5% glucose at 25°C for 24 h. The culture media were centrifuged and the cell pellets were suspended in sterile water, in which the OD600 was adjusted to 0.5. 1 µl of the cell suspension was spread onto each of ten nutrient medium plates containing YNB, 1.5% agar and one of the following carbon sources: glucose, galactose, mannose, xylose, arabinose, glucuronic acid, cellobiose, xylan (from oat spelt), glucomannan (from Konjac), and carboxymethyl cellulose. The concentration of each carbon source was 0.5 g/liter, except for insoluble xylan at 1.5 g/liter. The plates were incubated at 25°C for 5 d to determine the growth of each strain.

DNA Sequencing of Host Insects

To identify the species of the lucanid-like larvae from Switzerland, DNA sequencing analysis was conducted on the adult females from France and one of the larvae. Insect DNA was extracted from muscles of the adult prothorax and larval head using a QIAamp (R) DNA Mini Kit (Qiagen) following manufacturer’s instructions. PCR was performed to amplify mitochondrial 16S ribosomal RNA gene with the primers, mt16SB and mt16SC (Hosoya and Araya 2005), and ITS2 region of the nucleus ribosomal RNA gene with the primers, 5.8S3F8 (5'-CGATGAAGAAGCACGCTAATTG-3') and ITS4c0l (5'-TCTCTCCGCTTATGTAATATG-3') (developed in this study). These PCR products were sequenced with mt16SA, mt16SB and mt16SC primers for mt16S, and 5.8S3F8 and ITS4c0l for ITS2 region.

Phylogenetic Analysis of Yeasts and Host Insects

The sequence data were subjected to multiple DNA sequence alignment program ClustalW (Larkin et al. 2007). The alignments were then inspected and corrected manually, from which ambiguously aligned sites were removed. Phylogenetic analyses were conducted using the maximum likelihood (ML) method, program Mega 6 (Tamura et al. 2013). We selected the Tamura-Nei + G + I model for the yeast phylogeny and GTR + I + G model for the insect phylogeny, on the basis of the Akaike criterion estimated by the program Modeltest 3.06. Bootstrap tests were performed with 1,000 replications. Since the yeasts from the three D. parallelipipedus females had an identical sequence, we used that from the United Kingdom.

Results

Species Identification of Insects Using DNA Analysis

There was no difference in the nucleotide sequence of mt16S (977 bases) and ITS2 (999 bases) (accession numbers: LC119078–LC119081) between the adult female of S. cylindricum from France and the unidentified larva from Switzerland. Since other Sinodendron species are not found in Europe, we conclude that the larva has been correctly identified as S. cylindricum.

Morphological Observation of Symbiotic Organs

In adult females of S. cylindricum, a mycangium was found under the eighth tergite (Fig. 1h), at the same location reported for other lucanid species (Tanahashi et al. 2010, Hawes 2013, Tanahashi and Fremlin 2013). The two lateral oviducts of S. cylindricum were expanded, as in other lucanid species, and function as storage organs for mature eggs. Twelve ovarioles were observed at the anterior end of each oviduct and yellow bodies were observed at the posterior end of each oviduct that had produced at least one egg. Female #1 had eggs in the lateral oviducts and yellow bodies were present in all ovarioles (Fig. 1g), which suggests that it had been sexually mature and active the previous summer. Female #2 contained no eggs, or yellow bodies. In contrast, no mycangium-like structure was located in the males (Fig. 1f).

The intestine of the S. cylindricum larva consisted of a short, thin foregut, a long, cylindrical midgut, and an enlarged, segmented hindgut (Fig. 1i). Cecum-like sacs were found at the posterior end of the midgut (Fig. 1j). Although such cecum-like structures are known in most Scarabaeoidea larvae, their function has yet to be fully investigated.

Yeasts From Adult Insects

Two types of yeast colony (M1 and M2) grew on the PDA plates where the mycangium homogenate of female #1 was spread (Fig. 2a), while no yeasts appeared from the mycangium
homogenate of female #2 (Table 1). No living yeasts were found from any of the gut samples. M1 was the dominant colony type, with an estimated CFU of $9.8 \times 10^2$ per mycangium. The growth at 20°C was slow and the surface of the colonies was wrinkled (Fig. 2b). All eight M1 isolates shared the same ITS haplotype ($h1$) and one of these was chosen for the standard strain, namely, SICYAM1 (yeast from *S. cylindricum* adult mycangium, isolate 1). SICYAM1 formed hyphae on the yeast morphology agar after a 5-d incubation period at 25°C (Fig. 2c). The maximum sequence identity values for 18S rRNA, ITS, and 26S rRNA were 96, 91, and 89%, respectively (Table 2). Since values of conspecific strains are usually no less than 99% (i.e., 1% nucleotide difference between two strains) in yeasts (*Kurtzman and Suzuki 2010*), SICYAM1 is thought to be a novel yeast species. M2 appears to be a minor morphotype, with an estimated CFU of $9.1 \times 10^1$ per mycangium. It showed moderate growth and a smooth colony edge and surface on PDA medium (Fig. 2b). All M2 isolates shared the same ITS haplotype ($h2$) however, a minor genetic variant (one nucleotide replacement and one insertion), namely $h2v$, was determined from the midgut (two of eight colonies) and the cecum (one of eight colonies). Therefore, the representative isolates SICYLG1 and SICYLG2 were established from $h2$ and $h2v$ colonies, respectively. There was no sequence difference in 18S or 26S rRNAs between the two isolates. M3 colonies exhibited the fastest growth and smoothness on the PDA plate, and did not produce hyphae on the yeast morphology agar (Fig. 2e). The CFU of M3 was approximately one order of magnitude smaller than M2 in every organ (Table 1). All eight isolates shared the same ITS haplotype ($h3$), thus the representative isolate SICYLG3 was established as a third yeast strain. BLAST search gave 100% identity to the xylose-fermenting yeast, *S. insectosa* (Table 2).

**Yeasts From Other Lucanid Species**

Uniform colonies that were similar to M3 from the *Sinodendron* larva were isolated from the mycangia of *D. parallelipipedus* and *L. cervus* (Table 1), and the strains DOPAAM and LUCEAM with
Table 1 Yeasts isolated from S. cylindricum, D. parallelipipedus, and L. cervus

| Species               | Stage | Sex/specimen | Body weight (mg) | Organ    | Weight (mg) | CFU          |
|-----------------------|-------|--------------|------------------|----------|-------------|--------------|
|                       |       |              |                  |          |             | M1           |
|                       |       |              |                  |          |             | M2           |
|                       |       |              |                  |          |             | M3           |
| S. cylindricum        | Adult | Female #1    | 131.9            | Mycangium| 0.60        | $9.8 \times 10^3$ | $9.1 \times 10^4$ | –         |
|                       |       | female #2    | 156.6            | Mycangium| 0.87        | –            | –          |
|                       |       | male #1      | 115.2            | Mycangium| 0.96        | –            | –          |
|                       |       | male #2      | 160.1            | Mycangium| 0.80        | –            | –          |
|                       |       | larva unknown| 371.7            | Midgut   | 96.3        | $2.5 \times 10^4$ | $1.0 \times 10^4$ | –         |
|                       |       |              |                  | Hindgut  | 31.8        | –            | –          |
|                       |       |              |                  | Cecum    | 24.7        | $2.7 \times 10^5$ | $2.3 \times 10^2$ | –         |
| D. parallelipipedus   | Adult | Females + (n = 3) | 646.0 (± 151.0) | Mycangium| 1.11 (± 0.16) | –            | –          |
|                       |       |              |                  | Mycangium| 19.6 (± 5.6)| –            | –          |
|                       |       |              |                  | Gut      | 1.11 (± 0.16) | –            | –          |
| L. cervus             | Adult | Female      | 2647.0           | Mycangium| 5.20        | –            | $7.6 \times 10^5$ | –         |

M1, M2, and M3 indicate different colony types on PDA plate media.
–, not detected (<10 CFU); n/a, data not available.
aData shown as mean ± SD.

Table 2 Species or taxon of the yeasts isolated from S. cylindricum inferred from NCBI BLASTn search

| Host species     | Morphotype | ITS haplotype | Yeast strain | Accession no. | Maximum partial similarity (%) | Estimated species/taxon |
|------------------|------------|---------------|--------------|---------------|-------------------------------|-------------------------|
| S. cylindricum   | M1         | b1            | Y1028-SICYAM1 | LC119082      | 96                            | Sicyamaella               |
|                  | M2         | b2            | Y1028-SICYAM2 | LC119083      | 99                            | Sugiyamaella sp.          |
|                  | M3         | b3            | Y1053-SICYLG3 | LC119086      | 100                           | S. stipitis               |
| D. parallelipipedus | M3         | b4            | Y719-DOPAAM  | LC120356      | 100                           | Scheffersomyces sp.        |
| L. cervus        | M3         | b5            | Y718-LUCEAM  | LC120355      | 99                            | Sicyamaella               |

Additional notes for Table 2:
- Under ‘Yeast strain’, the leading ‘Y’ and the following group of three or four numbers represent the host specimen (abbreviated in the main text and Table 3).
- The four letters following the hyphen represent the host species, the last two letters indicate the source organ (AM, adult mycangium; LG, larval guts).
- Maximum partial similarity (%) inferred from NCBI BLASTn search.
- Estimated species/taxon following the maximum partial similarity.

Phylogenetic Analysis of Yeasts

SICYAM1, which was dominant in adult mycangia, was not compatible with any clades of previously known Saccharomycocotina yeasts (Fig. 3). It was also far removed from Scheffersomyces yeasts that are known symbionts in lucanids. SICYAM2, SICYLG1, and SICYLG2 were placed in the Sugiyamaella clade, although the ITS sequences indicate that they are different from any known species of Sugiyamaella yeasts.

Carbon Assimilation Test

All five yeast isolates from S. cylindricum were able to utilize glucose, mannose, xylose, and cellobiose (Table 3). SICYAM1 did not assimilate galactose or glucuronic acid. Three Sugiyamaella isolates (SICYAM2, SICYLG1, and SICYLG2) assimilated glucuronic acid (Table 3).

Discussion

In this study, we confirmed the presence of a female-specific mycangium in the genus Sinodendron, which is described as an ancestral clade of the Lucanidae. The mycangium of S. cylindricum originates from the intersegmental membrane that connects the eighth and ninth tergites, as it does in more evolutionarily advanced lucanids (Fremlin and Tanahashi 2015). As far as we know, this homologous structure has never been reported in other Scarabaeoidea families (Tanahashi et al. 2010). Our results therefore strengthen the view that the mycangium is a common and ancient characteristic of the Lucanidae. We obtained a total of five yeast strains from the adult mycangia and larval gut: a supposedly novel yeast from the mycangium, SICYAM1, three conspecific yeast strains that belong to Sugiyamaella clade, SICYAM2, SICYLG1 and SICYLG2, and the yeast identified as S. insectosa, SICYLG3. All of these yeast strains could utilize xylose, and therefore are considered to play a part in the digestion of wood components, especially hardwood.
Hemicelluloses, which consist mainly of xylose and glucose (Sjostrom 1993). Sugiyamaella yeasts are common in the gut of wood-feeding passalids (Coleoptera: Passalidae) (Urbina and Blackwell 2012) and they were present in both adult mycangium and larval guts of S. cylindricum. However, the novel yeast SICYAM1 was found only in the adult mycangium. Since SICYAM1 grew slowly, it might have been overlooked on the larval plate where the fast-growing Sugiyamaella and Scheffersomyces formed their colonies, this being one of the major problems in such a culture-dependent analysis. Males do not possess a mycangium, nor are there any living yeasts in their digestive tract, which suggests that they are unlikely to be involved in the vertical transmission of the yeast symbionts. Moreover, the absence of living yeasts in the female digestive tract indicates that the mycangium alone acts as a storage organ for the yeast symbionts.

Microbial Environment of Adult Mycangium

Although the mycangium of S. cylindricum harbored yeasts, the density of these was low (no more than 10^3 CFU), compared with those found in the mycangium of L. cervus and D. parallelipipedus (more than 10^5 CFU), which often share the same ecological niche as S. cylindricum in Europe and the UK. The mycangium of S. cylindricum was almost transparent and difficult to distinguish from the microbial environment.

Table 3. Growth of the five yeast strains isolated from S. cylindricum on different carbon sources

| Carbon source      | SICYAM1 | SICYAM2 | SICYLG1 | SICYLG2 | SICYLG3 |
|--------------------|---------|---------|---------|---------|---------|
| Glucose            | +       | +       | +       | +       | +       |
| Galactose          | –       | w       | w       | w       | w       |
| Mannose            | +       | +       | +       | +       | +       |
| Xylose             | +       | +       | +       | +       | +       |
| Arabinose          | w       | w       | w       | w       | w       |
| Galacturonic acid  | –       | +       | +       | +       | +       |
| Cellobiose         | +       | +       | +       | +       | +       |
| Xylan              | w       | w       | w       | w       | w       |
| Mannan             | –       | –       | –       | –       | –       |
| Carboxymethyl cellulose | –   | –       | –       | –       | –       |

+, moderate growth; w, weak growth; –, no growth.
surrounding intersegmental membrane, perhaps because of the small amount of mycangial secretion present. The low density of yeasts in the mycangium of *S. cylindricum* is unlikely to be due to the insect’s immaturity as the dissected beetle showed it to be sexually mature. Co-existence of multiple yeast species in the mycangium, contrasts with the single-yeast monoculture in the mycangium of higher lucanids (Tanahashi et al. 2010, Hawes 2013, Fremlin and Tanahashi 2013), and probably reflects a less specialized insect–yeast association in this species. This suggests that the mycangium of *S. cylindricum* represents an early stage in the evolution of insect–yeast symbiosis in the Lucanidae. Bacterial symbionts were not examined in this study; however, bacteria also play important roles in some higher lucanids, which at the same time harbour monoculture xylose-fermenting *Scheffersomyces* yeasts in their mycangia. Kuranouchi et al. (2006), for example, demonstrated nitrogen-fixing chemical activity in the gut of *Dorcas rectus* larvae, suggesting the presence of nitrogen-fixing bacteria, while both Tanahashi et al. (2009) and Tanahashi and Kubota (2013) suggest the probable symbiotic function of *D. rectus* gut bacteria is to assist digestion of fungal mycelia in decaying wood. More recently, Miyashita et al. (2015) discovered antibiotic-producing bacteria from mycangia of some *Dorcas* species. We suggest that the presence and roles of bacterial symbionts in *S. cylindricum* need further investigation.

**Microbial Environment of the Larval Gut**

In lucanid larvae, the midgut forms a straight, cylindrical tube, which contains dark-brown homogenates of ingested wood and digestive fluid. This rather liquid midgut fluid is highly alkaline, ranging from pH 10.2–10.4 in third instar larvae of *D. rectus* (M. Tanahashi, unpublished data). Such high alkalinity is also common in soil- or humus-feeding Scarabaeoidea (Lemke et al. 2003). In contrast, the hindgut of lucanid larvae is expanded when filled with digested food or feces, and usually constricted into two or more sections that will finally form eachecal pellet. The hindgut content has an almost neutral pH (6.7–7.8) in *D. rectus* larvae, and is rather solid and light-colored compared with that of the midgut. In this study, the midgut of the *S. cylindricum* larva contained considerably fewer living yeasts (3.6 × 10^5 CFU/mg) than the hindgut (1.1 × 10^6 CFU/mg), which suggests that the yeasts proliferate in the hindgut.

Similar to other lucanids, cecum-like sacs are located at the junction between midgut and hindgut of *S. cylindricum* larva. Although these seem to be a prominent internal structure, the function of the cecum-like sacs has not been fully investigated. The sacs, which are somewhat harsher than other intestinal organs, were seen to contain fine wood particles and many yeast cells, when observed under the microscope. The inner surface of these sacs seems to be lined with a thick cuticle, suggesting that it originated from ectodermal hindgut. Their yeast content is similar to that of the midgut and hindgut, even though the net capacity of the sacs is small. It is suggested that these sacs act as a refuge for the hindgut symbionts, where they can be retained safely against the gut flow until released into the hindgut. However, gut yeasts in passalid larvae are sometimes found adhering to the gut wall by means of branched filaments (holdfasts), which helps prevent them washing out (Nardi et al. 2006).

**Difference in Microbial Environment Between Adult and Larva**

In an earlier study, Tanahashi et al. (2010) put forward the hypothesis that the evolution of mycangia in Lucanidae is related to the biological difference in food eaten by larvae compared with that eaten by adults. Adult females of most lucanid species deposit eggs on decaying wood, although oviposition preference for decay type varies among species (Araya 1993, 2002). Wood is one of the most indigestible organic materials; it contains high levels of wood polymers such as cellulose, hemicelluloses and lignin, and a very low level of nitrogen (Haack and Slansky 1987). Degradation by wood-decaying fungi may enhance the nutrient value of wood to some extent, but nitrogen level remains low (usually, no more than 0.3%) (Tanahashi and Togashi 2009). Today, microbial symbionts are widely known from many wood-feeding or wood-inhabiting insect groups, and are thought to help digestion of wood polymers such as cellulose and lignin (Breznak and Brune 1994, Geib et al. 2009), detoxify the woody chemicals (Doud 1992), have an ability to fix nitrogen (Benemann 1973) or serve as food for insects (Beaver 1989). Lucanid larvae, including those of *S. cylindricum* referred to in this study, may also benefit from the xylose-utilizing ability of yeast symbionts.

In contrast, adult lucanids feed mainly on fermented tree sap (slime flux). Tree sap is rich in free sugars, amino acids, and vitamins (Kallio et al. 1989, Jiang et al. 2001). Free-living yeasts that are also commonly found in slime flux (Lachance et al. 2001) may improve its nutrient value. This suggests that adult stag beetles do not require specialist yeast symbionts in their food or digestive system, a suggestion supported by *S. cylindricum* adult beetles harboring no living yeasts or fungi, even though some food materials were still present in their digestive system. The same data were obtained from adults of *D. parallelopipedus* and *L. cervus* in this study, and also from Japanese stag beetles *D. rectus* and *L. macluliferatus* when they were examined at least 1 d after feeding (M. Tanahashi, unpublished data). The absence of living yeasts in sap-feeding adults is somewhat puzzling, because their food is likely to contain a large number of living yeasts and moulds. A possible answer to this question is that these yeasts have been completely digested to provide additional nutrient in their passage through the gut. Thus the gut of *S. cylindricum* adults, as well as adults of more evolutionarily advanced lucanids, seem never to act as storage organs for yeast symbionts, which we suggest has resulted in the evolution of a mycangium, an organ that is used for temporary storage of specialist yeast symbionts during the adult phase. In contrast to lucanids, the majority of wood-feeding insects rely on wood or woody tissues of trees throughout their lives, and some develop subsociality, where parents prepare food for their larvae. The Passalidae (Coleoptera) the major wood-feeding family in the Scarabaeoidea exhibits subsociality. Adults and larvae of passalids live in the same piece of decaying wood, which is also used as food by the larvae (Schuster and Schuster 1997, Ento et al. 2008); xylose-fermenting *Scheffersomyces* yeasts are commonly found in the gut of adult beetles (Suh et al. 2003) as well as larvae of this group (M. Tanahashi unpublished). We therefore hypothesize that evolution of the mycangium in the ancestral Lucanidae might have led to the development of adults and larvae using different food sources, sap-feeding in the former and wood-feeding in the latter.

**Limitations and Future Issues**

We are aware that our sample size is small and that these results need to be confirmed. Although *S. cylindricum* is widespread in Europe and the United Kingdom, it is very local and difficult to find, especially as this species spends its life almost entirely within decaying wood.

The evolution of a mycangium is possibly highly associated with the evolution of stag beetles. However, the phylogenetic placement of *Sinodendron* seems to be unresolved. For example, the recent phylogenetic study (Kim and Farrell 2015) showed only weak
branch support values for the ancestral clades of Lucanidae, including Sinodendron. Further studies are required to resolve these issues, and to elucidate the evolutionary origin of the insect-yeast symbiosis in Lucanidae.

Geolocation Information, Bioethics, and Data Availability

Samples of wild organisms used in this study were collected in the United Kingdom, France, Switzerland, and Japan. The biological and statistical analyses were conducted in the United Kingdom and Japan. International agreements were followed, protected areas were respected and bioethical standards met. All nucleotide sequences are available in GenBank (accession nos. LC119078–LC119086 and LC120353–LC120363). Yeast strains are being preserved as frozen stocks or plate cultures by the authors.

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