Methane Production and Methanogenic Archaea in the Digestive Tracts of Millipedes (Diplopoda)

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Abstract

Methane production by intestinal methanogenic Archaea and their community structure were compared among phylogenetic lineages of millipedes. Tropical and temperate millipedes of 35 species and 17 families were investigated. Species that emitted methane were mostly in the juliform orders Julida, Spirobolida, and Spirostreptida. The irregular phylogenetic distribution of methane production correlated with the presence of the methanogen-specific mcrA gene. The study brings the first detailed survey of methanogens’ diversity in the digestive tract of millipedes. Sequences related to Methanosarcinales, Methanobacteriales, Methanomicrobiales and some unclassified Archaea were detected using molecular profiling (DGGE). The differences in substrate preferences of the main lineages of methanogenic Archaea found in different millipede orders indicate that the composition of methanogen communities may reflect the differences in available substrates for methanogenesis or the presence of symbiotic protozoa in the digestive tract. We conclude that differences in methane production in the millipede gut reflect differences in the activity and proliferation of intestinal methanogens rather than an absolute inability of some millipede taxa to host methanogens. This inference was supported by the general presence of methanogenic activity in millipede faecal pellets and the presence of the 16S rRNA gene of methanogens in all tested taxa in the two main groups of millipedes, the Helminthophora and the Pentazonia.

Introduction

About 600 Tg of the greenhouse gas methane is produced on Earth each year [1]. Although methane, which has the second-largest impact on global warming after CO2 [2], is released to the atmosphere from human activities [3] including industry, agriculture, and waste management [4], natural sources account for 37% of global methane emissions. All methane, other than that from industry, is produced by the only known biogenic source - methanogenic Archaea. Methanogenic Archaea are an ancient group of microorganisms that occupy ecological niches with limited oxygen concentrations such as wetlands, rice fields, swamps, marshes, freshwater and marine sediments, agricultural soils, and the digestive tracts of humans and other animals [5,6,7,8,9].

Many authors have estimated methane production from vertebrates [10,11,12,13], and other authors have discussed the global importance of methane originating from invertebrate hosts and its potential contribution to the increase in atmospheric methane [14,15,16,17]. Termites have been recognized as a globally important source of methane in that they are estimated to contribute between 5 and 19% of the global methane emissions [18]. Methane production and the presence of methanogens has also been systematically screened in a wide variety of invertebrate taxa [19,20,21,22,23]; the results indicate that symbiotic associations involving methanogens and methane production are likely to be a characteristic property of the host taxon and that methane production is restricted to millipedes, cockroaches, beetles (Cetonidae), and termites [19].

Using microscopy and specific autofluorescence, Hackstein and Stumm [19] found free-living as well as endosymbiotic methanogens (methanogens living in the cells of Protozoa) in the hindgut of arthropod hosts. Methanogenic Archaea have been detected and their phylogeny analysed in the digestive tracts of methane-producing arthropods other than millipedes by 16S rRNA-based surveys [24,25,26,27,28]. Moreover, phylogenetic analysis in combination with enrichment cultures has suggested the existence of a seventh order of methanogens (the “Methanoplasmatales”) associated with intestinal tracts of both invertebrates and vertebrates [29]. A detailed assessment of methanogenic diversity is still lacking in millipedes, however, because microbial communities in millipede digestive tracts have mainly been studied only by classical cultivation methods and microscopic observation [19,30].

Unlike the digestive tract of most methane-producing arthropods, millipedes have a relatively simple digestive tract that lacks an enlarged pouch in the hindgut [19]. The inner surface of the
millipede hindgut, however, is strongly developed and may facilitate microbial colonization [30].

According to Hopkin and Read [31], there is no evidence that millipedes have a permanent symbiotic microbiota similar to that of termites, although microorganisms are important for millipede digestion. But the review of Byzov [30] demonstrated that the millipede intestinal tract harbours a stable, indigenous microbial community that includes facultative anaerobes that are able to degrade recalcitrant organic polymers. Only a few studies have used molecular techniques to study the microbial community of millipede faeces [32] or millipede intestines [29,33].

Specific autofluorescence microscopy showed that the digestive tract of some millipedes contains free methanogenic Archaea or *Nyctotherus*-type ciliates that host methanogens as endosymbionts [19], van Hoek [34] detected one archaeal sequence obtained from a symbiotic ciliate of an undetermined millipede, while Paul and colleagues [29] detected methanogens related to the “Methanoplasmatales” in the gut of the tropical millipede *Anadenobolus* sp.; no information is available, however, about the differences in intestinal archaean communities among different millipede lineages.

Methane is known to be produced by almost all tropical millipedes but is generally thought not to be produced by millipedes and cockroaches from temperate climates [19]. Sustr and Šimek [23], however, detected methane production in several species of European millipedes. Their results indicated that some groups of European millipedes (mainly Julida) seem to produce methane, while other groups do not (Polydesmida and Glomerida). Because millipedes apparently include producers as well as non-producers of methane, they represent an interesting model group for the determination of factors influencing gut methanogenesis.

In the current study, we compared methane production and the community structure of methanogens among different phylegetic lineages of millipedes. In doing so, we attempted to identify factors that explain the phylegetic differences. To accomplish these objectives, we used gas chromatography in order to measure methane production for a large number of taxonomically diverse millipede species. We verified the presence of methanogens by using PCR to detect the specific genetic marker (*mcrA* gene), and we used molecular profiling (DGGE) to compare the community structure of methanogens in the gut and in faecal pellets of different species of millipedes. Finally, we discuss how methane production relates to millipede phylogeny and millipede characteristics.

**Materials and Methods**

**Animals**

In the current study, most data used to evaluate methane production relative to millipede taxonomic position were obtained by gas chromatography. These data were supplemented by the data published in the recent literature [19,23]. The European millipedes used for our measurements were collected in different habitats in the Czech Republic, Slovakia, Romania, and Greece. The localities of origin of the species included in this study are listed in Table S2.

The permission for the research in Tatra Mountains was issued by the Regional Office of Environment Protection, Prešov, Slovakia (permission No. I/2007/01066-405/KM-R). Millipedes from the Latorica Protected Landscape Area (PLA) were collected with the permission of the Regional Office of Environment Protection, Košice, Slovakia (permission No. 2008/00322), and the research in the Slovak Karst was licensed by the Ministry of the Environment of the Slovak Republic (licence No. 3102/2009-

2.1/jam.). The research in the Bohemian Karst PLA was permitted by the Administration of Český kras PLA, Karlštejn, Czech Republic; in the Blansky les PLA by the Administration of Blansky les PLA, Vysoký, Český Krumlov, Czech Republic; in the Moravian Karst PLA by the Administration of Moravský kras PLA, Blansko, Czech Republic; and in the Pála PLA by the Administration of Pála PLA, Mikulov, Czech Republic. No specific permission was required to collect millipedes in the following five localities in the Czech Republic: the České Budějovice Basin (49°12′0.793″N, 14°24′25.629″E), the Chlěnice (49°06′14.026″N, 14°07′56.443″E), the Českomoravská vrchovina Highlands (49°12′56.878″N, 15°54′55.612″E), the Lanzhôt (48°43′6.003″N, 16°58′11.273″E), and the Žďárský les Highlands (49°04′41.946″N, 16°56′10.599″E). None of the species collected in the Czech Republic are endangered or protected in the Czech Republic. The species collected in Romania are also not endangered or protected. No specific permission was required for the sampling in the locality Mehendiñi Mts. (45°04′14.90″S, 22°45′48.35″E). The species collected in Thessaloniki (40°36′56.37″N, 23°02′08.96″E) is not endangered or protected in Greece, and no specific permission was required for this location.

Specimens of the millipede *Archisprostreptus gigas*, which is naturally distributed in Tanzania, were obtained from a pet shop (M. Kroček, Horni Suchá, Czech Republic). Two species of millipedes were obtained from a laboratory colony maintained at MPI for Terrestrial Microbiology Marburg (Germany); these species were *Glomeris marginata*, which occurs naturally in Germany, and *Orthomorpha coarctata*, which occurs naturally in Southeast Asia. Specimens of *Epibolus pulchripes*, which occurs naturally in Tanzania, were obtained from a breeding colony maintained by the Mikulov grammar school (Czech Republic).

All other localities mentioned in the Table S2 are cited as sites of origin of species used by Hackstein and Stumm [19]. The millipedes were identified to species based on the available literature; before they were assessed for methane production, the millipedes were kept in the laboratory at 10°C for several days to several weeks in plastic boxes containing top soil and litter from the collection site. Appropriate moisture was maintained by regularly moistening the substrate with water, and cuttlefish bone powder was added to the substrate as a source of calcium. Because the diet of most European species of millipedes consists of decomposed leaf litter, the litter in the boxes served as a food source.

As noted earlier, the tropical species *E. pulchripes* and *A. gigas* were obtained from a pet shop and from individual breeders. Species determination of *A. gigas* was verified based on gonopod morphology after Mwabvu et al. [33]. The tropical species were kept at about 25°C on horticultural substrate plus hardwood leaf litter. *E. pulchripes* accepted partly decomposed litter from hardwood trees (a mixture of hazel, maple, and oak leaves) as the only source of food. *A. gigas* consumed large amounts of organic wastes such as pieces of potatoes, cucumbers, and cabbage heads in addition to leaf litter and garden substrate. Coprophagy of millipede faecal pellets was not prevented for either of the tropical species. In total, 35 species of millipedes were maintained and assessed for methane production. Only individuals taken directly from the substrate and with full digestive tracts were used for gut dissection and subsequent microbial analyses.

**Methane production measurement**

Methane production from intact, living millipedes was measured by comparing methane concentrations in closed vessels (250 ml, 5 ml, or 1 ml depending on millipede size) that contained...
or did not contain millipedes – control vessels. The largest species, which were A. gigas, E. pulchripes, and the cockroach Blaptica dubia (B. dubia was used as a known methane producer), were measured in 250-ml glass vessels. Intermediate-sized species such as Julus sandinavicus were measured in 5-ml glass vessels. The 250- and 5-ml vessels were sealed with a rubber stopper and kept at 20°C for 3–4 h (in the case of 250-ml vessels) or for 3–13 h (in the case of 5-ml vessels). A piece of filter paper (2 × 4 cm for 250-ml vessels or 1 × 1 cm for 5-ml vessels) moistened with distilled water was placed in each glass vessel to maintain air humidity during measurement. The control glass vessels without animals were assembled and incubated in the same way. A 0.5-ml sample of the internal gas was collected at the start and at the end of incubation using gas-tight syringes, and the samples were injected into a gas chromatograph (GC) column.

The smallest species were incubated in a 700-µl volume in 1-ml plastic syringes [36]. A circle of moist filter paper (5 mm in diameter) was placed in the syringe to maintain air humidity. Fresh outdoor air was drawn into the syringe immediately before placement of animals. The syringes were incubated at 20°C, and the incubation period ranged from 4 h for the largest species to 24 h for the smallest species in syringes. As controls, the same syringes without animals were assembled, and the mean final concentration of methane in the controls was subtracted from the final concentration in samples. A 500-µl volume of the internal atmosphere in the syringe was injected directly into a GC after the incubation period. The amount of methane was quantified using an HP 5890 Series II gas chromatograph (Hewlett Packard, Palo Alto, CA, USA) equipped with a 2 m Porapak N column at 75°C, and a flame ionization detector using nitrogen as the carrier gas. A standard mixture of 100 ml m⁻³ methane in N₂ was used for calibration purposes [23].

In most cases, one animal was placed in one glass vessel or plastic vessel (syringe) to detect the individual variability in methane production. For some of the smallest species, for which incubation of individuals resulted in non-detectable methane production, several individuals were incubated in one syringe to verify the negative result (for T. costata, for example, three groups of three animals were measured). Animals were weighed at the end of the incubation, and the results were expressed as nl of methane produced per individual per h or as the live-mass specific methane production (nl of methane per g of live mass per h).

The ability to detect methane production depends on the sensitivity of the detection method and on the precision of measurement expressed in the variability of blank controls. If a small amount of methane is generated, the detection limit must be determined to distinguish between negative and positive measurements. We considered a value to be positive if it exceeded the value obtained in five control vessels (glass or plastic vessels without animals) by three standard deviations of controls. In addition, we designated species as non-producers (0% positive samples), accidental producers (at least one but <50% positive samples), facultative producers (from 50% to 99% positive samples), and obligatory producers (100% positive samples).

The distribution of methane-positive species among different taxonomic groups was summarized in contingency tables, and a nonparametric X² test (Statistica v6.0 Nonparametric Statistics, 2 × 2 tables, StatSoft, Inc., USA) was used to test the hypothesis of random distribution of methane production across the millipede phylogenetic tree. We used the previously published taxonomic classification of millipedes [37] along with the phylogenetic tree based on molecular data [39]. The effect of the mean species body mass on the frequency of methane-positive individuals in the species was tested by logit regression in Statistica v6.0 (Advanced Linear/Nonlinear Models: Nonlinear Estimation Analysis).

The faecal pellets of millipedes were anaerobically incubated at 20°C for 7 days to measure their methane production. Fresh pellets were placed in 5-ml glass vessels and moistened with 100 µl of distilled water. The atmosphere in the vessels was changed to argon, and the internal gas was sampled and analysed after 24, 96, and 168 hours by GC as described previously.

Molecular evaluation of the presence and diversity of methanogenic microorganisms in millipedes

Dissected digestive tract contents or freshly released faecal pellets were frozen and stored at −18°C until DNA extraction. Two methods of DNA extraction were tested. A phenol/ chloroform extraction method based on bead-beating homogenization of frozen samples [39] successfully extracted DNA mainly from the faecal pellets or the intestinal tracts of larger animals. For the intestinal tracts of smaller animals (mostly temperate, European millipedes), the NucleoSpin Tissue XS kit (Macherey-Nagel, Germany), based on enzymatic and chemical lyses, enabled the extraction of concentrated DNA from small amounts of starting material (0.025–10 mg). The quantity and quality of extracted DNA was determined with a NanoDrop ND-2000 spectrophotometer (Thermo Scientific, Wilmington, DE, USA).

For detection (expressed as presence or absence) of methanogens in the samples, we used PCR amplification of the mcrA gene, which encodes the methyl coenzyme M reductase β-subunit. The mcrA gene was amplified with two PCR assays, the results of which were combined for detection of positive signals (mcrA-1 and mcrA-2, Table S1). Both reactions were performed in 25-µl reaction volumes containing 1 µl of diluted DNA as template (usually around 5–10 ng), Qiagen Taq DNA polymerase (1.25 U), dNTPs (0.2 mM each), BSA (0.25 µg), Thermoc Scientific, USA), and primers (5 pmols and 8 pmols, respectively) in the presence of MgCl₂ (1.5 and 2.5 mM, respectively) in the PCR buffer recommended by the manufacturer (1x PCR buffer, Qiagen, CA, USA). A 5-µl quantity of the PCR products was analyzed in a 1% agarose gel (Top Vision agarose, Fermentas) that was stained with ethidium bromide (1 mg L⁻¹) for 30 min [40, 41].

Methanogenic community structure was characterized by DGGE using a nested PCR approach that targeted the 16S rRNA gene. In the first round, a stretch of the 16S rRNA gene was amplified using primers Ar109F and Ar915R (both at 25 pmol [42]). In the second round, diluted PCR product (1:100) served as template for amplification of methanogen-specific 16S rRNA using primers MG0357F-gc and MG0691R (both at 25 pmol [43]). Both reactions were carried out with Taq DNA polymerase (Qiagen, CA, USA) and reaction buffer supplemented with 4 µg of BSA (Thermo Scientific, USA). The cycling conditions for each assay are listed in Table S1. DGGE was performed with an Ingeny PhorU system (Ingeny, Leiden, The Netherlands) as follows: 200 ng of PCR product was loaded on an 8% (w/v) polyacrylamide gel with a denaturing gradient range of 30–60% (100% denaturant is equivalent to 7 M urea and 40% deionized formamide). Electrophoresis was run in 0.5×TAE buffer for 16 h at 60°C and 100 V. The DGGE gels were stained with SYBR Green I nucleic acid stain (Lonza Rockland, ME, USA), and molecular profiles were analyzed with GelCompar II software (Applied Maths, Ghent, Belgium). Bands were excised from gels using a sterilized dissecter, eluted in 20 µl of MilliQ water with a freeze-thaw cycle, and re-amplified with primers MG0357F–MG0691R. The resulting PCR products were purified using a MinElute PCR Purification kit (Qiagen, CA, USA) and sequenced using an ABI Prism 3100-Avant Genetic Analyser (Applied
The sequences were edited with Bioedit 7.0.4.1 software [44] and assembled in Geneious version Pro 5.5.6 created by Biomatters (New Zealand, available from http://www.geneious.com/). Sequences were tested against the GenBank database (www.ncbi.nlm.nih.gov) using the Blastn algorithm. The sequences are available in GenBank under accession numbers KF574048–KF574078 and KF739300–KF739307.

The relationship between the results of molecular detection of methanogens and GC detection of methane production was tested by nonparametric statistical methods (Statistica v6.0 Nonparametric Statistics, 2×2 tables). Differences in the complexity of methanogenic communities (numbers of bands detected by DGGE) among different groups of millipedes were tested with a nonparametric Kruskal-Wallis test or Wilcoxon Matched Pairs test (Statistica v6.0, Nonparametric Statistics).

Results

Phylogenetic overview of methane production

Table S2 summarizes the data on the presence or absence of methanogenesis in living and intact millipedes belonging to 46 species and eight orders. Data for 35 species were obtained in the current study, and data for the other 11 species were obtained from the literature [19,23]. With respect to methane production, the species are designated as non-producers (NP), accidental producers (AP), facultative producers (FP), and obligatory producers (OP). Except for Glomeris tetraspica, which was an accidental producer, all species in the order Glomerida were non-producers according to GC analysis (Table S2). Millipedes in the orders Polyzoniida and Chordeumatida (Table S2) were also non-producers. The order Callipodida was represented by only one individual of Callipodella fasciata, which did not produce methane. The order Julida included all categories of methane producers, from non-producers to obligate producers. Of the 18 species tested in the family Julidae, only three were obligatory producers; non-producers, accidental producers, and facultative producers were each represented by five species in the Julidae. Two species were tested in the family Blaniulidae, and neither produced measurable methane. The spirobolid species E. pulchripes was designated an obligatory producer, and the same designation probably applies to the other members of the order Spirostreptida. All A. gigas tested in this study produced large amounts of methane. Among the millipedes belonging to the order Polydesmida that were tested in this study, only the tropical species Orthomorpha coarctata was classified as an obligatory producer (only one individual was tested).

European members of the order Polydesmida belonging to the families Paradoxosomatidae, Polydesmidae, and Trichopolydasmidae did not produce detectable levels of methane except for one of the four tested individuals of Strongylomorpha stigmaticum (Table S2). The data available for this order are biased, however, because most polydesmid methane producers are tropical and large species

Figure 1. The distribution of methane-positive species among millipede orders. The red row of columns (OFP) is based on all tested species classified as either obligatory or facultative producers, and the yellow row (OFAP) includes obligatory, facultative, and accidental producers. doi:10.1371/journal.pone.0102659.g001
(O. coarctata and Pycnotropis acuticollis [19]) but most of the tested polydesmids were European and small species.

The current data and previously published data [19,23] were used to describe the distribution of methane-positive species among the main millipede orders (Fig. 1 and 2). The methane-positive species (obligatory and facultative methane producers) are distributed almost exclusively in the jiliform groups (the orders Julida, Spirobolida, and Spirostreptida). The order Julida, which was the most investigated group, had a higher percentage of accidental producers and non-producers than the Spirobolida and Spirostreptida. In a comparison of the frequencies of all methane-producing species (AP, FP, and OP together) in the jiliform orders Julida, Spirobolida, and Spirostreptida vs. in all other groups, the \(X^2\) test of the 2×2 contingency table indicated that taxonomic position was significantly related to methane production (\(X^2(df = 1) = 13.61, p = 0.0002\)).

Quantitative aspects of methane production

One adult individual of the millipede A. gigas produced more methane than the smaller tropical E. pulchripes in the order Spirobolida and several orders of magnitude more than small European species in the order Julida. The mean individual methane production from A. gigas is comparable to the methane production from the large cockroach, Blaptica dubia. Mean individual methane production (M) by species was positively related to average live body mass (W) by species as indicated by a double logarithmic plot (Fig. 3). The linear regression of log M on log W corresponded to the equation M = 32.494W^{1.569} (\(r^2 = 0.999, p = 0.0001\)). Regression of body mass-specific methane production (M/W) on millipede mass (W) was also positive and significant (\(r^2 = 0.522, p = 0.002\)). It follows that, in addition to taxonomic position, body size may explain differences in methane production among millipede species.

We also analyzed changes in the percentages of methane-positive and methane-negative individuals with decreasing mean body mass in an attempt to detect a general lower limit in body mass for methane production by millipedes. All five species whose body mass was >0.4 g were obligatory producers (OP). The intermediate group in terms of body mass (from 0.06 to 0.30 g) included eight species representing all four categories of methane producers (two species of OP, two species of FP, one species of AP, and three species of NP). All species whose body mass was <0.06 g were non-producers except for Unciger foetidus (the mean body mass of the tested individuals was 0.059 g) and the very small julid species Enantiulus nanus (its mean body mass was 0.006 g); in the case of E. nanus, only one of the 10 tested individuals was positive. In the order Glomerida, the largest species (0.28 g) was classified as NP, and the data set is very small. These data are also scarce for the order Polydesmida, but the existence of some body mass limit for methane production may be expected in this group because all

![Figure 2. Summary of the recent information concerning methane production by millipedes.](https://doi.org/10.1371/journal.pone.0102659.g002)
species that were <0.08 g produced no methane and one species that was >0.4 g produced methane. For the juliform orders (Julida, Spirostreptida, and Spirobolida), a body mass limit of 0.12 g separated the methane-positive and methane-negative species.

Methane production from faecal pellets of millipedes

Excrements of all tested millipede species produced methane under anaerobic conditions (Fig. 4). The time course of the production was similar in all species in that the rate of production quickly increased during the first day of anaerobic incubation and then slowly increased over the next 6 days (Fig. 5).

Methanogenic community structure

DNA was successfully extracted from the faecal pellets of 16 millipede species and from the gut contents of 13 millipede species (Table S2). The extraction was less effective from gut contents than from faecal pellets. Specific amplification of the mcrA gene using two different PCR assays confirmed the presence of methanogens in the gut of three species of millipedes belonging to the orders Julida, Spirostreptida, and Spirobolida. Results were similar for both assays except for Cylindroiulus luridus excrements and Julus scandinavius gut contents (Table S2), which can be influenced by primer sequence bias. The mcrA gene was not found in the gut contents of two species in the family Glomeridae or in the gut contents of the one species in the order Polydesmida (Table S2). A Chi-square test calculated from 2×2 contingency tables indicated that the production of methane as detected by GC was significantly related to the detection of mcrA in the gut contents of the same animal ($X^2(df = 1) = 4.05, p = 0.044, n = 24$). Detection of the mcrA gene in gut contents supported the classification of species based on GC measurement of methane production. All three of the positive gut samples originated from species that were designated FP or OP.

For most species, the detection of mcrA in the gut contents corresponded to mcrA detection in faecal pellets. Discrepancies occurred in only three cases (two in which mcrA was detected only in faecal pellets and one in which mcrA was detected only in the gut contents). All of the tested millipede faecal pellets began to produce methane early during the incubation under anaerobic, moist conditions but the mcrA gene was detected in the faecal pellets of only five of the 16 species that were tested. The detection of mcrA in different millipede lineages is illustrated in Figure 6.

DGGE profiles of the methanogenic microbial communities were obtained from 18 samples of the digestive tracts representing 10 species. This method revealed the presence of methanogens in the digestive tract of members of the orders Glomerida, Julida, Spirobolida, and Spirostreptida and in all categories of methane production except NP. Molecular profiles of the methanogenic microbial communities were also obtained from 30 samples of faecal pellets representing 15 species. In addition, methanogens
were detected in the faecal pellets of Polydesmida members. The
dGGE analysis revealed the presence of the methanogen-specific
16S rRNA gene in all of the tested millipede orders (Fig. 7, Table
S2). dGGE analysis also revealed large variability in the
methanogenic community structure among different gut and
faecal pellet samples in millipedes from the orders Glomerida and
Julida. Clustering of molecular profiles (Fig. 8) showed no relation
of methanogenic community structure to the pre-defined groups
NP, AP, FP, and OP. The results revealed a high diversity of the
methanogenic community among millipede individuals, because
only a few of the molecular profiles were similar (at similarity
cutoff = 70%). The molecular profiles of methanogens from the
faecal pellets of G. hexasticha (MK), G. connexa (MK), M.
unilineatum (TH), and U. foetidus (BL) were similar and
significantly different from that of G. hexasticha (HT) and G.
tetristicha (SK). The methanogenic communities of guts and
excrements of the same species were similar in some cases. In other
species (J. scandinavis and A. gigas) the gut and faecal pellet
communities formed different clusters. The highest dissimilarity
was found between samples of J. scandinavis. Methanogenic
communities of the whole gut content were almost identical among
individuals of G. tetristicha originating from the same
locality, but methanogens in the different gut compartments of A.
gigas had only 40% similarity (Fig. 8).

As indicated by the number of bands in the DGGE profile,
some species harboured rich methanogenic communities while
others did not. Overall, more DGGE bands were obtained in
faecal pellets than in gut contents. This trend was confirmed by
nonparametric statistical tests comparing all DGGE profiles
obtained from guts and faecal pellets (Kruskall-Wallis test, H (1,
n = 48) = 3.99, P = 0.046) and comparing species averages for
pellets and guts (Wilcoxon pair test, n = 9, P = 0.038). The number
of bands in DGGE profiles was greater for the tropical millipedes
(Spirobolida and Spirostreptida), which included the two largest
species in this study, than for the European millipedes (Kruskall-
Wallis test, H (1, n = 48) = 6.95, P = 0.008) (Fig. 8).

It was not possible to re-amplify and characterize all of the
excised bands from each profile, perhaps because of PCR
inhibition or a low concentration of the DNA fragment in the
eluate. A total of 32 bands, representing specific sequences in the
dGGE profiles, were successfully sequenced and identified (Table
S3), and these were assumed to represent the specific or dominant
components of the communities.

In general, diverse methanogens were identified in the gut and
faecal pellets. These included Methanosarcina thermophila, M.
siciliae, Methanoregula boonei, Methanobrevibacter woesei, M.
thermaer, and M. arboriphilus, as well as other species of these
genera that remained unidentified. Methanogenic DGGE profiles
obtained from guts and faecal pellets of millipedes belonging to the
order Julida were dominated by sequences related to Methano-
microbiales and Methanosarcinales. Methanoregula and other

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**Figure 4. Methane production from millipede faecal pellets relative to millipede phylogeny.** Methane production from anaerobically
incubated faecal pellets was detected by gas chromatography (Table S1). Species belonging to framed groups were tested in this study. Red lines
mark the lineages containing some positive species. The phylogenetic tree was used according to Regier et al. [38].
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unclassified Methanomicrobiales or *Methanosarcina* were often identified in these samples.

In samples representing the millipede family Glomeridae, we found Methanomicrobiales, Methanosarcinales, as well as Methanobacteriales (*Methanobacterium* sp. or *Methanobrevibacter* sp.). In faecal pellets and gut content samples from *S. stigmatosum* (Polydesmida), *E. pulchripes* (Spirobolida), and *A. gigas* (Spirostreptida), only *Methanobrevibacter* spp. and related Methanobacteriales were among the dominant bands that were successfully identified.

**Discussion**

The results presented here show that methane production and the *mcrA* gene are unequally distributed among phylogenetic lineages of millipedes (see Fig. 2 and Fig. 6). Previous research also revealed considerable variation in the presence of methane production among different groups of arthropods, and these differences were thought to reflect differences in the ability to host methanogenic microorganisms [19].

Although detection of methane production and the *mcrA* gene indicate a rather restricted distribution of methanogens in millipedes, we found the 16S rRNA genetic marker of methanogens in all tested taxa (members of Julida, Glomerida, Polydesmida, Spirobolida, and Spirostreptida) and potential anaerobic
methane production in all of the tested faecal pellets (see Fig. 7 and Fig. 4). Thus, the methanogens were present in both of the main groups of millipedes: the Helminthophora and Pentazonia. In the current study, the presence of methanogens in the Pentazonia was indicated by methanogenesis in all of their tested faecal pellets, by the detection of methane production from some of living millipedes, and by the detection of methanogen genes. In a previous study, the presence of methanogens in the Pentazonia was supported indirectly by the detection of hydrogen production in the gut of some species of Glomerida [19]. The current results indicate that methanogens may be more broadly distributed in millipedes than previously assumed.

We obtained molecular evidence of methanogens by assessing the ribosomal marker (16S rRNA) and mcrA gene. The results obtained with 16S rRNA frequently differed from those obtained with mcrA. Methane production was better correlated with detection of mcrA than with detection of 16S rRNA. The mcrA-based approach may be less sensitive because its primers are more specific than in the 16S rRNA-based approach. The latter feature has been discussed previously [45] and can be explained by a higher gene copy number of the 16S rRNA gene than of the mcrA gene in the genomes.

The disagreement between detection of methane production and the molecular evidence of methanogens (based on the 16S rRNA gene) can be attributed to the fact that the presence of genetic markers is not an accurate indicator of activity and may represent only the potential for activity. It also suggests that a very small or inactive methanogenic community may be present in the gut of most millipedes. Such communities might be activated when conditions become favourable. This idea is supported by the ability of methanogens to survive unfavourable conditions (drought and aeration) for long periods [46] and to be activated once anoxic conditions return [47,48]. Our results indicate that the striking differences in methane production among different millipede taxa reflect differences in the activity of intestinal methanogens rather than an absolute inability of some taxa to host methanogens.

Hackstein and Stumm [19] considered possible sources of variation in the activity of intestinal methanogens, resulting in differences in methane production among arthropod individuals. The authors discussed body size, environmental temperature or climatic zone of the species distribution, number of intestinal protozoa, and unknown genetically fixed factors.

Among the millipedes investigated in our study, the number of methane-producing animals increased with increasing body size of the species. Moreover, the amount of methane produced was positively correlated with millipede body size in methane-producing species. This effect of body size may be explained by the better aeration in the gut of small vs. large millipedes (the gut of smaller animals has a higher surface-to-volume ratio); the increased aeration in the gut of small animals may prevent the
proliferation of anaerobic methanogenic Archaea [49]. Although the oxygen concentration in the gut of millipedes has rarely been measured, the mean redox potential in the gut of the pill millipede, *G. marginata* (body mass about 0.2 g), was +232 mV in the midgut and +204 mV in the hindgut, and these values correspond to oxidative conditions [50]. This is consistent with our failure to detect methane production in this species.

Thus, differences in body size may explain much of the between-species variability in methane production of millipedes. The absence of methane production in relatively larger members of the Polyzooniida and Callipodida may be caused by their unique feeding habits. Polyzooniida consume moist, soil detritus [51], and Callipodida contains species that have predatory feeding habits and species that consume the dead tissues of animals [52,53].

Hackstein and Stumm [19] speculated that methane production would be more common among tropical than temperate species of arthropods. The effect of climatic origin of species may be masked by the effect of body size because, like other terrestrial invertebrates, millipedes tend to be larger in the tropics than in other regions [54]. Although methane production from individual millipedes decreased with decreasing temperature [23], the inference that high environmental temperatures are required for methanogenesis was disproved by the evidence of methane production from European species of the family Julidae [23] as well as by methane production in permafrost soils [55].

Another possible source of the differences in methane production between tropical and temperate taxa may lie in the symbiotic interaction of large tropical millipedes with ciliate protists related to the genus *Nyctotherus* [19]. The ciliates host considerable numbers of methanogens near hydrogen-producing hydrogenosomes [56,57], where the methanogens consume hydrogen. Therefore, the abundance of ciliates enhances the methanogenic activity and enhances the rate of methane production from cockroaches [17]. It follows that protists may be responsible for quantitative differences in the methane production between tropical spirostreptid millipedes and European Julidae. This is supported by our finding that mainly hydrogenotrophic *Methanobacteriales* (*Methanobrevibacter arborophilus* and other *Methanobrevibacter*-related sequences) dominated the gut of large members of the Spirostreptida (*A. gigas*); hydrogenotrophic Methanobacteriales were previously found in the cytoplasm of ciliates of the genus *Nyctotherus* [17,34]. Moreover, we observed ciliates in the hindgut and fresh faecal pellets of *A. gigas* (unpublished data).

Most methanogenic Archaea lineages previously reported from the insect gut (Methanosarcinales, Methanobacteriales, and Methanomicrobiales; see [39] for a summary) were identified in the present study of millipedes. The list of methanogenic lineages identified here may be completed only by the previously published evidence of the Thermoplasmatales-related methanogens in the...
hindgut of the millipede *Adenobolus* sp. (Spirobolida) [29] and the DNA sequence of an endosymbiotic methanogen (probably related to the Methanobacteriales) isolated from the ciliate *Nycotherus velox* from the gut of a “julid” (probably spirostreptid) millipede [34]. The comparison of methanogenic lineages identified in the digestive tracts of larger tropical orders (Spirostreptida and Spirobolida) vs. smaller temperate orders (Julida, Glomerida) of millipedes generated a pattern similar to that obtained for termites [59]. Methanogens colonizing the hindgut of lower termites (which have gut flagellates) and the gut of Spirostreptida millipedes (which have symbiotic ciliates) belong almost exclusively to the genus *Methanobrevibacter*. Methanogens colonizing the hindgut of higher termites are more diverse than those colonizing lower termites and include Methanomicrobiales, Methanosarcinales, and Methanobacteriales, which were the same orders detected in Julida millipedes.

The main lineages of methanogenic Archaea found in different millipede orders differ in their substrate preference and free vs. endosymbiotic mode of life. Therefore, the composition of methanogenic communities may reflect differences in the habitats and substrates provided by different lineages of millipedes.

*Methanobrevibacter* species isolated from different habitats almost exclusively used H₂ and CO₂ as substrates [39]. Some *Methanobrevibacter* phylotypes may be associated with flagellates [60] or ciliates [57,34]. The metabolism of the strain MP2, which was identified in spirobolids, resembles H₂-requiring methylotrophic methanogens [29]. Regarding Methanosarcinales representatives, rather than finding *Methanomicrococcus blatticola*, which is specialized in hydrogen-dependent reduction of methanol or methylamines to methane [22] and which is found in cockroaches and higher termites [59], we found several *Methanosarcina* lineages in julid millipedes. Methanosarcinales are able to use all three of the known metabolic pathways for methanogenesis (hydrogenotrophic, acetoclastic, and methylotrophic) [61] and thus possess a versatile metabolism that is able to respond to environmental change. Additional detailed studies of selected millipede species are needed to determine whether the metabolic plasticity of different lineages of methanogenic Archaea influences which millipede taxa they inhabit.

In conclusion, the qualitative and quantitative variability in methane production among millipedes seems to reflect the suppression or activation of methanogenic Archaea in the digestive

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Figure 8. Cluster analysis of methanogenic communities in millipedes. DGGE patterns were obtained after PCR amplification of the 16S rRNA gene of the methanogenic communities in the gut contents (Δ) or faecal pellets (●) of individual millipedes from the indicated species. Pearson correlation and UPGMA analysis were used. Abbreviations in brackets indicate the origin of animal samples (see Table S1). mgut = midgut and hgut = hindgut samples. Species belonging to different pre-defined groups of methane producers are marked with different colours: NP – black, AP – green, FP – blue, and OP – red. Asterisks indicate that the samples were from individuals or faecal pellets in which methane production was detected using gas chromatography. Blue line = similarity cutoff 70%.

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tructs of some phylogenetic millipede lineages rather than a fundamental inability of these millipede lineages to host methanogens. Our results also indicate that methane-producing millipede taxa differ in the functional groups of methanogens that they host.

Supporting Information

Table S1 PCR protocols. Primers and PCR conditions used for amplification of the mcrA gene and the 16S rRNA gene for DGGE analysis.

Table S2 Methane production in millipedes and methanogenic microorganisms detected in millipede gut contents and faecal pellets. Methane production was measured by gas chromatography (GC) in the current study or in previous studies, as indicated. The mcrA gene was used as a marker for methanogens in the gut contents and faecal pellets. The presence of methanogens in the gut contents and faecal pellets was also determined by DGGE of the methanogen 16S rRNA gene. Mean values for richness of DGGE profiles are given.

Table S3 The sequenced bands of methanogenic archaeal genes amplified from millipedes. Phylogenetic relationships of the excised and sequenced bands of methanogenic archaeal 16S rRNA genes amplified from millipede faecal pellets or gut contents. Sequences that were not submitted to GenBank were those shorter than 200 bp.

References


