Enzymatic biosynthesis of B-complex vitamins is supplied by diverse microbiota in the \textit{Rhodnius prolixus} anterior midgut following \textit{Trypanosoma cruzi} infection

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\textbf{A B S T R A C T}

\textit{Trypanosoma cruzi}, the causative agent of Chagas disease, colonizes the gut of triatomine insects, including \textit{Rhodnius prolixus}. It is believed that this colonization upsets the microbiota that are normally present, presumably switching the environment to one more favorable for parasite survival. It was previously thought that one particular bacterium, \textit{Rhodococcus rhodnii}, was essential for insect survival due to its ability to produce vital B-complex vitamins. However, these bacteria are not always identified in great abundance in studies on \textit{R. prolixus} microbiota. Here we sequenced the microbiota of the insect anterior midgut using shotgun metagenomic sequencing in order to obtain a high-resolution snapshot of the microbes inside at two different time points and under two conditions; in the presence or absence of parasite and immediately following infection, or three days post-infection. We identify a total of 217 metagenomic bins, and recovered one metagenome-assembled genome, which we placed in the genus \textit{Rhodococcus}. We show that, despite \textit{Rhodococcus} being present, it is not the only microbe capable of synthesizing B-complex vitamins, with the genes required for biosynthesis present in a number of different microbes. This work helps to gain a new insight into the microbial ecology of \textit{R. prolixus}.

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\section*{1. Introduction}

\textit{Trypanosoma cruzi} is a protozoan parasite that colonizes the digestive tract of triatomine insect vectors (Reduviidae: Triatominae) \cite{1}. The parasite is responsible for causing Chagas disease, an illness that affects several million people worldwide, predominantly in Latin America. The insect hosts are generally nocturnal feeders, that defecate following feeding. The parasite is excreted in the insect faeces and may enter the bloodstream through the bite wound \cite{1,2}. Recent evidence also suggests that food contaminated with faeces or urine is increasingly responsible for infection, where ingested parasite is sufficient to cause an acute phase of disease \cite{3,4}.

Colonization of the triatomines by the parasite results in antimicrobial peptide (AMP) production in as little as two hours post-feeding, ultimately leading to a decreased bacterial load. These changes in the microbiota presumably render the gut ecology more favorable for the parasite to replicate \cite{2,4}. To date, several studies have investigated the community composition of triatomines following parasite colonization. These studies have utilized various methods including denaturing gradient gel electroophoresis \cite{5,6}, RADseq \cite{7} and 16S amplicon sequencing \cite{8–15}. Results from these studies vary, with some suggesting as few as 20 species make up the community composition \cite{6}, while others demonstrate a much higher diversity, despite having clearly dominant members of the microbiota, such as \textit{Pectobacterium} in \textit{R. prolixus} and \textit{Arsenophonus} in the \textit{Triatoma brasiliensis} complex \cite{8}. No consensus has yet been found regarding microbial members of a “core” community. It is therefore difficult to understand the role of the microbiota with respect to \textit{Trypanosoma} infection and triatomines in general. One further obstacle is the fact that the blood meal significantly alters the microbiota in triatomines, a detail further complicated by observed differences between male and female insects \cite{9}.
Next generation sequencing techniques and their application on triatomines have highlighted the diversity of bacteria associated with different species of triatomine and across different labs. In fact, dominant symbionts can include members of the Proteobacteria, Actinomycetes or Bacilli [5–11]. What has yet to be determined is the genetic capabilities that are afforded by the individual microbes, which may account for the observed diversity. Early studies indicated, for example, that B-complex vitamins were an essential contribution of the microbiota and, in Rhodnius prolixus, these vitamins are purportedly supplied by the bacterium, Rhodococcus rhodnii [12]. However, it has not yet been closely investigated whether other associated microbes also contribute to essential metabolic processes.

To address the above supposition, we investigated the changes that occur in the microbiome of the R. prolixus anterior midgut three days post-feeding, by using shotgun metagenomic sequencing. This is the first instance of such a study in triatomines with respect to T. cruzi infection and we provide insight into the coding sequence potential of those organisms that are present following parasitic exposure compared to uninfected controls.

2. Methods

2.1. Ethics statement

All experiments were performed in accordance with guidelines on animal experimentation from FIOCRUZ, adhering to all Brazilian legislation regarding animal welfare. Protocols were based upon procedures set out by The Ministry of Science and Technology (CONCEA/MCT) associated with the American Association for Animal Science (AAAS), the Federation of European Laboratory Animal Science Associations (FELASA), the International Council for Animal Science (ICLAS) and the Association for Assessment and Accreditation of Laboratory Animal Care International (AAALAC). Ethical approval under license number LW-8/17 was approved by the Committee for Ethics in the Use of Animals, CEUA-FIOCRUZ.

2.2. Insect rearing and dissection

Nymphs used in this study came from a colony maintained at a temperature of 25 ± 1 °C, 60 ± 10% relative humidity and natural illumination by the Vector Behaviour and Pathogen Interaction Group, Belo Horizonte, Brazil. Insects are fed monthly with citrated rabbit blood (10% v/v) obtained from the Centro de Criação de Animais de Laboratório (CECAL, Fiocruz, Rio de Janeiro, Brazil) offered in an artificial feeder equipped with a latex membrane and a circulating heated water flow (37 °C), and chicken anesthetized with intraperitoneal injections of a mixture of ketamine (20 mg/kg; Cristália, Brazil) and detomidine (0.3 mg/kg; Syntec, Brazil).

For infection with T. cruzi, a SWR/J mouse was inoculated with 200 μl of triatomine urine containing ~ 5x10^6 metacyclic trypomastigotes/ml from Dm28c strain. The strain was originally isolated from naturally infected opossum and kept in laboratory culture. The parasites were cultured by two weekly passages in liver-infusion tryptose (LIT) supplemented with 15% fetal bovine serum, 100 mg/ml streptomycin and 100 units/ml penicillin. To prevent loss of activity, parasites were submitted to cycles of triatomine-mice infection every six month. Nine days post-infection, the mouse was anesthetized with ketamine (150 mg/kg; Cristália, Brazil) and xylazine (10 mg/kg; Bayer, Brazil) and the presence of parasite was confirmed by light microscopy. The mice were then offered R. prolixus fifth instar nymphs in the early morning, which had fasted for approximately 30 days. Nymphs in the same physiological conditions were fed on an anesthetized healthy mouse and used as a control group. Nymphs were fed for the same amount of time and we proceeded on the assumption that small differences in blood taken would not affect results in a major way since samples from each treatment group were subsequently pooled.

Insect gut sections were segregated using a dissecting microscope (Motic, SMZ-168) and sterile instruments immediately after feeding (approx. 5 to 15 min) and three days post-feeding. This time point was chosen because the transformation of the infectious trypomastigote stages of T. cruzi into epimastigote stages, which only occur in the invertebrate hosts, begins as early as a few hours after infection [13]. Three days after the bloodmeal, the colonization should therefore be well established. In total, twelve triatomines were prepared for DNA extraction. Three anterior midguts from exposed nymphs were combined immediately after feeding as well as three anterior midguts from unexposed insects. This section of the intestine was chosen because there is reportedly a higher antimicrobial activity than in the posterior midgut [4]. The same was done three days post-feeding resulting in the four sample ‘T. cruzi infected blood time point 0’ (TcInfT0), ‘T. cruzi infected blood time point 3’ (TcInfT3), ‘T. cruzi control time point 0’ (TcContT0) and ‘T. cruzi control time point 3’ (TcContT3). By combining the anterior midguts of several individuals, enough intestinal DNA material could be obtained for the shotgun metagenomic sequencing.

2.3. DNA extraction and sequencing

Following dissection of R. prolixus, DNA of the four samples, each with three pooled anterior midguts, was extracted using the AllPrep DNA/RNA minikit (Qiagen) following the manufacturer’s recommendations. Purified DNA was shipped to Novogene and four libraries, one for each sample, were prepared by Novogene. Sample preparation was done using the NEB Ultrall DNA library preparation kit with four amplification cycles. Paired-end (150 bp) sequencing was performed on an Illumina NovaSeq6000 instrument.

2.4. Sequence analysis

Raw reads of all samples were trimmed using Trimmomatic (v0.33) [14] to remove low quality bases and reads lacking a pair using the following settings: ILLUMINACLIP:TruSeq3-PE.fa:2:30:10 LEADING:3 TRAILING:3 SLIDINGWINDOW:4:15 MINLEN:36. For taxonomic classification, reads were uploaded to the Kaiju [15] web server. Results were downloaded and processed according to scripts available at https://github.com/ntobias-85/metagenome-csbj-2020. Trimmed reads were filtered against the R. prolixus genome (GCA_000181055.3) using bowtie2 (v2.2.5) [16] and the --un-conc option to keep only reads not mapping to the reference. Reads not mapping to R. prolixus were used to generate a co-assembly of all four samples with megahit (v1.1.4) [17] with the --kmin-1pass option activated. Individual samples were then independently mapped to the co-assembly to generate bam files using bowtie2 (v2.2.5) and samtools (v1.2) [18]. In this way, sample-specific information such as the relative abundance of sequences from each bin can be determined.

Anvi’o is a bioinformatic package designed to streamline metagenomic workflows. The co-assembly of the four samples was used to generate a contig database in anvi’o (v5.1) [19], which includes annotation of open reading frames using Prodigal (v2.6.3) [20] as a built-in function. Anvi’o was further used to run hidden Markov models to identify single copy gene collections (anvi-run-hmms) and annotate NCBI COGs (anvi-run-ncbi-cogs). COGs (Clusters of Orthologous Groups) represent families of orthologous protein-coding genes which can be used to study microbial phy-
logeny and genome annotation. Taxonomy for genes was added by using centrifuge (v1.0.2) [21] and importing the output into the contig database using the anvi-import-taxonomy-for-genes. Profiles for each single sample were generated, removing all contigs less than 2,500 nucleotides. Finally, the profiles were merged with the anvi-merge command, which includes an automatic binning step with CONCOCT [22]. Amino acid sequences for each gene were exported from the contig database and used as input in the GhostKOALA webserver. The output was parsed back into the contig database using the KEGG-to-anno script, followed by anvi-import of the resulting output. The automatic binning was manually inspected and refined based on predicted taxonomy, including blastx searches of annotated genes, read coverage and GC-content of contigs. The pangenomic analysis [23] was carried out using anvi'o with settings --minbit = 0.5; --mcl-inflation = 10 and --use-ncbi-blast activated.

For phylogenetic analyses, genomes were downloaded from NCBI and converted to contig databases using the 'anvi-script-STA-to-contigs-db' script. The program 'anvi-get-sequences-for-hits' was then used to extract ribosomal RNA sequences common among all downloaded genomes and the respective metagenomic bins. Ribosomal proteins were extracted from the database with --concatenate activated. Concatenated protein sequences were then imported into Geneious v6.1.8 and a maximum likelihood tree was constructed with 100 bootstraps using PhyML [24] (version 3.0).

2.5. KEGG analysis

The output from GhostKOALA was used as input for KEGG_decoyer (Kyoto Encyclopedia of Genes and Genomes) [25] and subsequently KEGG_encoder, which uses hidden markov models to incorporate some further processes. The outputs were then combined with the python script, Decode_and_Expand.py [25], generating a heatmap of all processes identified for each sample.

3. Results and discussion

3.1. Microbial populations of the R. prolixus anterior midgut

Studies on the microbiome of R. prolixus have so far predominantly profiled bacterial species using the 16S rRNA. Results show that there is little intra-individual diversity associated with the microbiota and insects are often dominated by a single group of organisms [8,26]. In the case of Rhodnius spp., these are especially Enterobacteria, such as Pectobacteria and Serratia [6,8,31]. Furthermore, the Actinobacteria Rhodococcus, is found not only in Rhodnius, but also in other triatomine genera such as Triatoma [11,32,33]. One consequence of only investigating 16S rRNA is that interactions between the insect, fungi, viruses and even other parasites that may be present are totally excluded from analysis. Moreover, bacterial species often possess multiple copies of 16S rRNA genes making a reliable quantification difficult [34]. Another drawback is that sequences can only accurately be assigned to higher level taxonomies, with some difficulty to obtain resolution between species. Nevertheless, 16S amplicon sequencing has several advantages. A large and comprehensive reference database is available facilitating the detection of rare taxa. Additionally, it is significantly cheaper than metagenomic shotgun sequencing and the lower data output is easier to process with regard to bioinformatic pipelines and computational capacity. 16S rRNA amplicon sequencing is therefore highly suitable for taxonomic analysis approaches, such as community composition studies.

Given the advances in sequencing technology and the desire to more fully understand the microbial ecology of triatomine midgut at high-resolution, we proceeded to use metagenomic shotgun sequencing to analyze the anterior midgut of R. prolixus under four different conditions (described in methods). Sequencing resulted in a total 145,812,112 clean reads (TcContT0: 34,750,241 (10.4 GB), TcContT3: 47,062,804 (14.1 GB), TcInfT0: 33,499,209 (10.1 GB) and TcInfT3: 30,499,858 (9.2 GB) for further analysis. To begin with, we used Kaiju [15], which directly BLASTs each read pair against the non-redundant nucleotide database, to profile the composition of microbial communities associated with our samples using the paired end input option for each sample (raw data is available from the European Nucleotide Archive under accession number PRJEB33861). This was done initially to determine the number of raw reads that could be directly attributed to any given organism. Our analysis suggests that, apart from bacterial DNA, a large part of the DNA originates from a mixture of fungi, viruses, parasites and other eukaryotic sequences (Supplementary Fig. 1). We also detected an increase in the proportion of bacterial DNA after the blood meal. This is probably due to the increased supply of nutrients and has been observed several times before [4,8]. One key difference to keep in mind between this and other studies is the fact that here, we have used a more natural approach to infection, feeding triatomines on mice infected with triatominne urine (with or without parasite), as opposed to other studies that tend to use a significantly higher parasitic load for infection experiments. One final difference is the way in which our bacterial taxonomy was initially determined. We used kaiju from samples originating from the anterior midgut, where the majority of other studies instead use 16S amplicon sequencing from the hindgut (or directly on faeces). Despite these differences, we do see somewhat similar results to previously published work in that there is a moderate proportion of Actinobacteria, Proteobacteria and Firmicutes identified in the samples (Supplementary Fig. 2) [5,7,11,27–32]. However, the dominant order in our experiment is Chlamydiales, possibly indicative of a dominant microbe in the insect colony used. To more directly compare taxonomic classification of sequencing data with previous works, future experiments should include an additional 16S amplicon sequencing analysis on the same sample used for shotgun sequencing.

3.2. Metagenome assembly and annotation

We were primarily interested in the gene content of the microbiota present in the anterior midgut of R. prolixus. Therefore, initial filtering of our reads was performed against the R. prolixus genome to remove all sequences mapping to the insect DNA. Since the insects originated from the same colony, we predicted that their microbiomes would be relatively similar. We therefore performed a co-assembly with the remaining reads. Co-assembly involves using all available sequence data to get the best assembly possible. The individual samples are then mapped back to this co-assembly to obtain sample specific coverage details. Combining all four samples using megahit, resulted in an assembly containing a total of 480,315 contigs larger than 1,000 bp (max 152,936 bp, average 2,302 bp, N50 2,496 bp). For the annotation, only contigs with 2,500 bp or more were kept (max 152,936 bp, average 6,481 bp, N50 19,857 bp). This cutoff was chosen to ensure a meaningful signal of sequence composition and coverage for subsequent binning steps. The annotation resulted in a total of 357,390 gene calls. Supplementary annotation with the KEGG database allowed us to map the presence or absence of individual pathways to each sample. We then utilized this data to define samples that contained enriched pathways, relative to all other samples. The KEGG decoder and expander script [25] utilizes a hidden Markov database to identify and fill false-negative gaps in pathways that appear incomplete. Based upon this, we screened 162 individual pathways, most of which contained little to no difference between samples. Minor
differences could be identified between time point 0 and time point 3 such as in nitrate reduction pathways (Supplementary Fig. 3), but we expect differences would be more evident when including a greater number of samples.

Annotations were useful in refining the automatic binning output from CONCOCT. Although binning algorithms perform well in most cases, manual refinement is still necessary to ensure the accuracy of contig groupings. In total 217 bins were identified that were unrelated to \textit{R. prolixus}. One draft metagenome-assembled genome (MAG), MAG_NJT_1, was recovered from the genomic bins, which was preliminarily identified as being from the \textit{Dickeya} genus (more details below). Since \textit{Rhodococcus} is reportedly an essential symbiont of \textit{R. prolixus} [33], we also searched for bins containing predominantly \textit{Rhodococcus} sequence (Fig. 1, annotated with the bin name “TcAA_NJT_2”).

Our experiment consisted of DNA samples extracted from either exposed or unexposed insects immediately after feeding and three days post-feeding. We expected that the two samples at time point zero should be similar, while we may see differences between exposed and unexposed samples after three days. For the most part this is true, although the differences associated with three-day time points is limited to just a handful of bins (Fig. 1). We also expected to see changes after three days due to the increase in the expression of various AMPs during this time [2]. However, we saw relatively few differences in the overall relative abundances of microbial profiles between exposed and unexposed (Supplementary Table 1). This may be in part due to the different infection strategy that we utilized (outlined above). The parasite load in this experiment was low, which increases the possibility of a lack of response and may partly explain unexpectedly few differences between the control group and challenged insects. Although further experiments are required to confirm the veracity of our hypothesis, it is possible that microbial changes reported to be associated with parasite entering the insect gut may be, at least in part, due to a higher parasitic load than would naturally occur. Despite this, we observed an increase in sequences derived from MAG_NJT_1, and two other bins (Bin_22_4 and Bin 22_1_1) as well as a decrease in TcAA_NJT_2, which we suspect is \textit{Rhodococcus} (Supplementary Fig. 4). Eichler and Schaub (2002) also identified such a decrease of \textit{Rhodococcus} after the blood meal, but the amount of the bacteria recovered quickly and exceeded the starting level after a few days [34]. These differences could be associated with the differential effects of AMPs against different bacterial species [2], yet still raises the question of the importance of \textit{Rhodococcus} at this time point. Aposymbiotic \textit{Rhodnius prolixus}, which fail to develop beyond the second larval stage, regenerate when fed with blood containing \textit{Rhodococcus} [35]. However, since other bacterial symbionts have never been examined in this way, it cannot be determined whether they would have a similar remediating effect.
3.3. Presence of B-complex vitamin biosynthesis pathways

Early research into *R. prolixus* microbiota provides a narrative that *R. rhodnii* are essential symbionts in triatomine biology [12] and in the absence of these symbionts, the insects fail to moult [36,37]. Deliberation of this point includes arguments that the bacteria are used as a food source themselves, with bacterial lysis supplying required nutrients, as well as strong supporting evidence for the role of B-complex vitamins and their production by the *Rhodococcus* [37]. Many of these studies were performed several years ago (as early as 1936) and include culture-dependent methodologies. In recent years however, several groups have revisited triatomine microbiomes, with 16S amplicon sequencing emerging as the preferred method for investigations of bacteria.

Amplicon sequencing consistently demonstrates that *R. rhodnii* are not the only bacterial species present, nor are they regularly the dominant bacteria. Others have demonstrated that dominant symbionts include Actinobacteria (of which *Rhodococcus* is a member) [11], *Serratia* [6], *Pectobacterium* [8] and *Arsenophonus* [6,26,28], which has already been shown to serve as a source of B-vitamins in whiteflies[38]. Furthermore, in other obligate hematophagous insects, symbioses with various B-vitamin producing bacteria have been detected. The Gammaproteobacteria *Wigglesworthia* spp. is regarded as the main vitamin B provider of the tsetse fly, the aetiological agent of the African trypanosomiasis and *Riesia* spp. seems to be indispensable for *Pediculus humanus humanus*, the human body louse. A lack of these symbionts often results in development retardation and a decrease in fecundity [37,38]. Given this conflicting evidence, we decided to explore the B-vitamin biosynthesis capabilities of the microbiota identified in our study. We extracted all of the gene calls from the co-assembly and uploaded them to GhostKoala to annotate each of our putative coding sequences with KEGG ontology (KO) numbers. Based on the KEGG pathways, we were then able to extract any gene call with a KO number from each of the eight different vitamin B pathways (see Supplementary Table 2 for a list of KO numbers used for each biosynthetic pathway). Unexpectedly, we see that enzymes involved in B-complex vitamin biosynthesis are present in a number of different organisms present within the microbiota (Fig. 1). Therefore, the provision of vitamin B derivatives solely by *Rhodococcus* seems unlikely. Despite this, only two mem-
bers contained enzymes present in all eight vitamin biosynthetic pathways. These were TcAA_NJT_2, which is predicted to be *Rhodococcus*, and have already been shown to possess vitamin B biosynthesis genes including: thiamine (B₁), riboflavin (B₂), niacin (B₃), pantothenate (B₅), pyridoxal (B₆), biotin (B₇), tetrahydrofolate (B₉), and cobalamin (B₁₂) [39]. The second bin containing orthologs in each of the vitamin B biosynthetic pathways was MAG_NJT_1.

3.4. Identification of the metagenome-assembled genome

A closer look at MAG_NJT_1 reveals an 89.21% completion and 0% redundancy estimate based on the presence of single copy genes. To identify a likely taxon, we used BLAST to analyze one of the ribosomal subunits, which suggested the bin originated from *Pectobacterium*. Since some *Pectobacterium* isolates were reclassified into *Brenneria* and Dickeya [40], we used concatenated ribosomal proteins from a selection of these three genera, which placed MAG_NJT_1 into the Dickeya genus (Supplementary Fig. 5). Dickeya are a genus of Gamma-proteobacteria that typically cause disease in plants. Although these phytopathogens are reported to be widespread in insects [41], it is curious that they should be a dominant taxon in a hematophagous insect. However, this is not the first time that a bacterium predominantly associated with plants has been described as a major taxon associated with *R. prolixus*, with *Methyllobacterium* identified as the third most abundant microbe in a North American lab colony of *R. prolixus* [26]. Nevertheless, to check for contamination within this bin, we performed a pan-genome analysis using 15 Dickeya isolates and MAG_NJT_1 (Fig. 2). Excluding MAG_NJT_1, other Dickeya isolates appear to have approximately twice the number of gene clusters and are approximately double the length of MAG_NJT_1. Based on this, we estimate that this bin is in fact approximately 50% complete. Of the single core genes (SCGs) present in all 16 isolates, 50 were randomly selected to create a higher resolution phylogeny (Supplementary Fig. 6, Fig. 2). MAG_NJT_1 clustered closely to Dickeya parasitica (formerly *Brenneria parasitica*).

Although it is still likely that B-complex vitamins are essential to *Rhodinus* development and survival, our data here show that B-vitamins can be contributed to the gut ecology by a number of microbes are (i) present to fulfill the vitamin B biosynthesis and (ii) are capable of doing so.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at https://doi.org/10.1016/j.csbj.2020.10.031.

References


