

# Targeted genome enrichment for efficient purification of endosymbiont DNA from host DNA

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**Abstract** *Wolbachia* endosymbionts are widespread in arthropods and are generally considered reproductive parasites, inducing various phenotypes including cytoplasmic incompatibility, parthenogenesis, feminization and male killing, which serve to promote their spread through populations. In contrast, *Wolbachia* infecting filarial nematodes that cause human diseases, including elephantiasis and river blindness, are obligate mutualists. DNA purification methods for efficient genomic sequencing of these unculturable bacteria have proven difficult using a variety of techniques. To efficiently capture endosymbiont DNA for studies that examine the biology of symbiosis, we devised a parallel strategy to an earlier array-based method by creating a set of SureSelect™ (Agilent) 120-mer target enrichment RNA oligonucleotides (“baits”) for solution hybrid selection. These were designed from *Wolbachia* complete and partial genome sequences in GenBank and were tiled across each genomic sequence with 60 bp overlap. Baits were filtered for homology against host genomes containing *Wolbachia* using BLAT and sequences with significant host homology were removed from the bait pool. Filarial parasite *Brugia*

*malayi* DNA was used as a test case, as the complete sequence of both *Wolbachia* and its host are known. DNA eluted from capture was size selected and sequencing samples were prepared using the NEBNext® Sample Preparation Kit. One-third of a 50 nt paired-end sequencing lane on the HiSeq™ 2000 (Illumina) yielded 53 million reads and the entirety of the *Wolbachia* genome was captured. We then used the baits to isolate more than 97.1 % of the genome of a distantly related *Wolbachia* strain from the crustacean *Armadillidium vulgare*, demonstrating that the method can be used to enrich target DNA from unculturable microbes over large evolutionary distances.

**Keywords** *Wolbachia* · Obligate endosymbiont · Target enrichment · NextGen sequencing · DNA capture · SureSelect™

## 1 Introduction

Whole genome sequencing (WGS) using Next-Generation sequencing (NGS) technologies is becoming a cost efficient and effective procedure for many research laboratories. While metagenomic projects (purification and DNA sequencing of mixed populations of bacteria *en masse*) are becoming common, purification methods for efficient sequencing of selected unculturable bacteria from amongst other DNA populations, such as symbiont hosts, have proven difficult. Purification strategies have generally used a variety of techniques including chemical gradients (Charles and Ishikawa 1999), pulsed-field gel (PFG) purification with or without whole genome amplification, library construction followed by gene walking, etc. (Foster et al. 2005; Iturbe-Ormaetxe et al. 2011; Mavingui et al. 2005;

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Sun et al. 2001). Informatics methods (Kumar and Blaxter 2011) and cell sorting methods (Santos-Garcia et al. 2012) have also been used.

More recently, DNA capture methods using oligonucleotide probes have been used to isolate symbiont or parasitic DNA away from host DNA, providing significant enrichment for more efficient sequencing (Kent et al. 2011; Melnikov et al. 2011). Our goal was to further assess this capture methodology using SureSelect™ (Agilent) technology that could be applied to purify DNA from widely divergent unculturable bacteria (e.g. symbionts) away from other DNA sources as an initiation point for WGS, when some homologous or similar DNA sequence is known. We used the obligate endobacterium *Wolbachia* as the target species for capture, to eliminate host DNA from downstream library construction and subsequent DNA sequencing. *Wolbachia* are obligate alpha-proteobacteria closely related to rickettsial organisms and are present in insects, mites, crustaceans, spiders and parasitic filarial nematodes (Werren et al. 2008; Werren and Windsor 2000; Cordaux et al. 2012; Bouchon et al. 1998). It has been recently estimated that ~40 % of arthropod species are infected with *Wolbachia* (Zug and Hammerstein 2012), making it the most widespread intracellular bacterial species. Phylogenetic analyses using single genes or multilocus sequence typing currently describe seven well-resolved *Wolbachia* groups, designated as supergroup lineages (A to H, no group G) together with a number of additional lineages (Lo et al. 2002; Casiraghi et al. 2005; Casiraghi et al. 2003; Lo and Evans 2007; Baldo and Werren 2007; Bordenstein et al. 2009).

In 1999, the *Wolbachia* Genome Consortium planned to sequence various genomes representing the diversity of *Wolbachia* (Slatko et al. 1999). At the onset of our study, only four *Wolbachia* genomes had been completely sequenced and deposited in GENBANK, *wMel* of *Drosophila melanogaster* (Wu et al. 2004), *wBm* of *Brugia malayi* (Foster et al. 2005), *wPip* of *Culex quinquefasciatus* Pel (Klasson et al. 2008), *wRi* of *Drosophila simulans* (Klasson et al. 2009). Additional genomes are currently being sequenced/annotated (Werren et al. 2008) and partial sequences for several *Wolbachia* strains are available, e.g., <http://www.ncbi.nlm.nih.gov/nuccore/?term=wolbachia>. A common goal of many *Wolbachia* genome projects is to provide comparative genomic information for understanding mechanisms of genome evolution and mechanisms of host phenotype manipulation by *Wolbachia* (Cordaux et al. 2011; Saridaki and Bourtzis 2010; Werren 1997). In addition, the evolutionary analysis of *Wolbachia* prophages is of interest and capturing the symbiont DNA from divergent sources not only provides DNA sequence of *Wolbachia* but also can capture their prophages (Kent et al. 2011).

To test the selectivity of the designed baits, we used the *Wolbachia wBm* strain (supergroup D) as a test case for the efficiency of *Wolbachia* DNA capture. The sequence of the 1.1 Mb *Wolbachia* genome is already known (Foster et al. 2005) and was included in the bait pool. Using these baits, we then attempted to selectively capture *Wolbachia* DNA from a phylogenetically distant strain, the feminizing *wVulC* strain from the isopod *Armadillidium vulgare* (supergroup B) (Cordaux et al. 2004) for which no genomic sequences were included in the bait design.

In the case of *wVulC*, DNA sequencing and comparative analysis from related strains with alternate phenotypes (cytoplasmic incompatibility and feminization) (Bouchon et al. 2008) will aid in the identification of the genetic basis of the phenotypic differences induced by *Wolbachia* within this group of isopod crustaceans. In the case of *wBm*, it has been shown that this endosymbiont is a novel drug target against human filariasis (Slatko et al. 2010), and identification of worldwide variants will be useful. The described DNA capture technique should find application in drug discovery, evolutionary analysis and populational/ecological studies. For example, a similar approach with Roche Nimblegen arrays was utilized by the Bordenstein lab to isolate and analyze the *Wolbachia* genome and WO prophages of the strain *wVitB* from the parasitic wasp *Nasonia vitripennis* (Kent et al. 2011).

## 2 Materials and methods

### 2.1 *Wolbachia* strains

Two *Wolbachia* strains were analyzed in this study: the obligate *wBm* strain from the nematode *B. malayi* (TRS Labs, Georgia, USA) (Foster et al. 2005) and the feminizing *wVulC* strain from the terrestrial isopod *A. vulgare* (maintained in the EES lab) (Cordaux et al. 2004). The 1.1 Mb *wBm* genome sequence is known and the *wVulC* genome is in final assembly steps. The size of the *wVulC* genome has been estimated by PFG electrophoresis at 1.75 Mb (Bouchon et al. 2008) and the current sequence consists of 10 contigs of 1.66 Mb, which agrees with pulsed-field estimates (Liu et al. manuscript in preparation).

### 2.2 RNA bait library design

To design the targeted genome enrichment library, we created a SureSelect™ set of enrichment oligonucleotides for solution hybrid selection. The library of biotinylated complementary RNA baits was designed and synthesized by Agilent (Santa Clara, CA). RNA baits were utilized because of the stability of RNA-DNA hybrids in the selection process and the ease of their removal in subsequent steps. The

120-mer RNA bait library was created based upon compiling 11 *Wolbachia* complete and partial genome sequences found in GenBank (*Wolbachia* infections of *Muscidifurax uniraptor* (NZ\_ACFP00000000.1), *Wuchereria bancrofti* (PRJNA43539), *Onchocerca volvulus* (PRJNA12625), *Drosophila willistoni* (PRJNA16739), *Drosophila simulans gdsi 540* (AAGC01000629.1), *Drosophila simulans* (AAGC00000000.1), *Drosophila melanogaster* (AE017196.1), *Drosophila ananassae gdan 143* (AAGB00000000), *Culex quinquefasciatus pel* (AM999887.1), *Culex quinquefasciatus jhb* (ABZA00000000), *Brugia malayi* (AE017321.1)). Bait sequences were tiled across each genomic sequence with 60 bp overlap and pooled, resulting in approximately 215,000 baits with about 207,000 unique sequences. Baits were then filtered for homology against select host genomes that contain *Wolbachia* (*Brugia malayi* (GCF\_000002995.1), *Onchocerca volvulus* (ADBW00000000.1), *Wuchereria bancrofti* (ADBV00000000.1)) using BLAT, a high speed and more accurate BLAST-Like Alignment Tool with the ability to use an internal set of sequences for assembly and rapidly find high similarity sequences of relatively short length (Kent 2002). 6,000 baits with significant host homology were removed from the bait pool. The final bait count was 201,776 after removing BLAT rejects. Baits were also tested for uniqueness against two nematodes that do not harbor the endosymbiont (*Caenorhabditis elegans* (GCA\_000002985.2), *Loa loa* (GCA\_000183805.1)). This BLAT search did not produce significant hits and thus no additional baits were removed.

### 2.3 DNA extraction and preparation

Total DNA from the nematode *B. malayi* and from the isopod *A. vulgare* were extracted as described (Sambrook and Russell 2001; Bouchon et al. 1998). Quantification of the DNA samples was performed using a Nanodrop 1,000 spectrophotometer (Thermo Scientific) and the Qubit 2.0 fluorimeter (Invitrogen). DNA samples were normalized to 3 µg of DNA for each sequence capture protocol. DNA samples were sheared by sonication to an average length of 200 bp using a Covaris S1 then end repaired, followed by 3'dA addition using the NEBNext® Sample Preparation Kit (New England Biolabs). Adaptors were ligated onto the ends and following purification the DNA was PCR amplified (6 cycles) using indexed PCR primers and the Illumina InPE1.0 forward PCR primer. After purification, quality assays were performed using the Caliper GX (Life Science) and Bioanalyzer (Agilent) to determine the average fragment sizes and concentrations.

### 2.4 Capture and sequencing

*Wolbachia* DNA was captured from the prepared total DNA by hybridization to the biotinylated cRNA baits for 24 h at

65 °C, following the Agilent SureSelect™ protocol, but supplemented with custom blocking oligos complementary to the barcoded adaptors. Bound DNA was recovered using magnetic streptavidin beads, PCR amplified (12 cycles) using Illumina forward and reverse primers and purified. Sequencing samples were prepared using the NEBNext® Sample Preparation Kit (New England Biolabs). The library was paired-end sequenced on the Illumina HiSeq 2000 at HudsonAlpha, Inc.

### 2.5 Bioinformatics analysis

Sequence reads provided by HudsonAlpha were quality controlled using FASTQC (<http://www.bioinformatics.babraham.ac.uk/projects/fastqc/>) and mapped against the complete *wBm* genome and the partial *wVulC* genome using Bowtie (version 0.12.7) (Langmead et al. 2009) and Bowtie2 (version 2.0.0-beta5) (Langmead and Salzberg 2012). Mapping results were processed by SAMtools (Li et al. 2009) and visualized using Artemis (Carver et al. 2012). Assemblies were performed using Velvet assembler (Version 1.2.03) (Zerbino and Birney 2008). Optimization of the assembly was performed by using different k-mers (from 19 to 49 by steps of 2 bases); the optimal assembly was chosen considering the N50, the length of the longest contig and the total bases in the contig.

## 3 Results and discussion

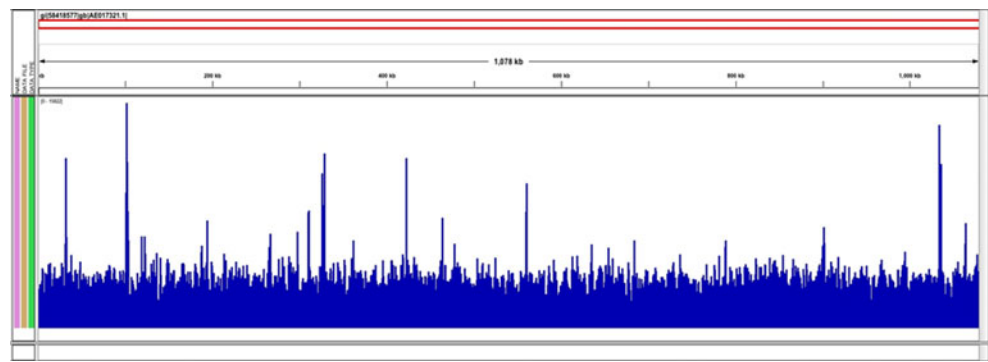
### 3.1 *wBm* sequence capture and sequencing

Nearly ninety-two million (91,724,826) reads from duplicated one third lanes on an Illumina HiSeq 2000 were produced from DNA captured by the oligo bait set. High quality metrics were obtained from the run (Q scores above 30 for bases 1–13 and above 40 for bases 14–50). Sensitive parameters of Bowtie2 were used to map these reads onto the genome sequence of the *Wolbachia* endosymbiont from the TRS strain of *Brugia malayi* (NC\_006833). The entire *Wolbachia* genome was captured at a depth of over 3000×. These results for *wBm* were anticipated as the capture oligos we designed were based on all *Wolbachia* genomic sequences in GenBank or RefSeq including those of *wBm* (Fig. 1).

Depth of coverage was generally uniform but showed spikes over certain regions (Y axis, Fig. 1). These regions could either be repetitive in the genome or could be regions of duplicated lateral gene transfers into the host genome (Dunning Hotopp et al. 2007).

Only 5.20 % of the reads did not map to the *Wolbachia* *wBm* genome. These unmapped reads were assembled by Velvet and the resulting 1,107 contigs were used to query the NCBI database by BLASTn (Fig. 2). 27.8 % of these

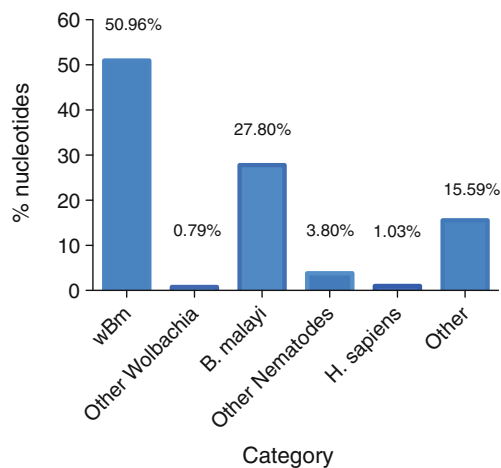
**Fig. 1** The average number of mapped reads per genomic location averaged over a 25 nt window. The X axis represents a linear map of the *Wolbachia* genome from *B. malayi* and the Y axis (1–15,000) represents relative sequence coverage. The count file was generated from a BAM file using IGV Tools (Thorvaldsdottir et al. 2012). 100 % coverage was obtained



~5 % unmapped sequences BLAST to host nematode sequences and 51 % appeared to BLAST to *wBm Wolbachia* sequence. These represent contigs assembled from reads that do not map to the reference genome (reads with more than 3 mismatches were not mapped) but have significant similarity to *wBm* by BLAST. Unknown sequences (15.59 %) were investigated for GC content and they appeared to have an average GC content of 28.1 %, a value similar to *Wolbachia* and the host (Foster et al. 2005; Ghedin et al. 2007). Thus they can not be differentiated as to whether they represent *Wolbachia* or *B. malayi* genes or derive from other organisms with similarly low GC content.

### 3.2 *wVulC* sequence capture and sequencing

Over thirty-six million reads (one-third of a 50 nt paired-end sequencing lane on the Illumina HiSeq™ 2000) were produced from DNA capture by the oligo bait set. As with the



**Fig. 2** Distribution of the 5.2 % reads not mapped to *wBm*. Bar graph denotes % of nucleotides out of the total assembled contigs that show significant BLAST scores to *wBm* genome, other *Wolbachia* the *B. malayi* host, other nematodes or *H. sapiens*. The “Other Nematodes” category corresponds to sequences with a significant BLAST score to other species of nematodes (mainly filarial nematodes). The “*H. sapiens*” category may represent DNA contamination. The “Other” category corresponds to sequences with no significant BLAST similarity to any sequences in the NCBI database

*wBm* mapping set, sequences were quality assayed. The entire dataset was of high quality (Q scores above 30 for bases 1–9 and above 40 for bases 10–50).

Different parameters of Bowtie were tested to improve the mapping of the total paired-end read dataset to the incompletely sequenced *wVulC* genome and the final mapping using Bowtie2 provided a total of 34,524,894 millions reads mapping to the reference genome (94.73 %).

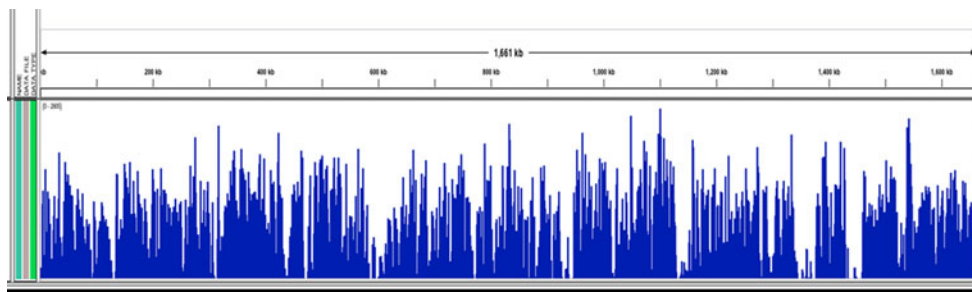
This mapping covered 97.1 % of the partially known *wVulC* sequence (Fig. 3). Several regions appear to be over-represented in the data set, as with the *wBm* mapping. These include the single copy DNA-directed RNA polymerase, beta/beta’ subunit gene as well as the phage major capsid protein E gene, which is likely repetitive in the genome.

The unmapped reads (5.27 % of the total reads) were assembled by Velvet and the resulting 28 contigs were used to query the NCBI database by BLASTn. Of these contigs, 24 (94.78 % of the total unmapped reads) corresponded to other *Wolbachia* sequences. These sequences may represent *wVulC* genes not yet identified in the incomplete genome as the average GC% of these sequences is similar to the GC content of *Wolbachia* strain. None of the remaining 4 contigs had any significant match to any other sequences in the database.

As the *wVulC* genome is still in draft form, it cannot be excluded that there are errors in the preliminary sequence. To correct the sequence, mapping results were used to call variants and to detect errors, which were manually corrected. By this method, more than 300 base substitutions and over 30 indels have been corrected by use of the oligo bait-captured DNA sequence.

## 4 De novo assembly of the *wVulC* genome sequence

A primary goal of this study was to sequence an unknown genome and thus we performed a *de novo* assembly of the oligo-captured *wVulC* sequence data. After optimization of the Velvet parameters (k-mer=45, insert length=170, standard deviation for insert length=63 and minimum contig

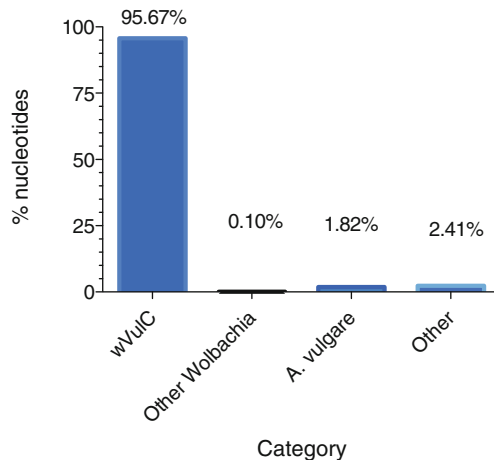


**Fig. 3** The average number of mapped reads per genomic location averaged over a 25 nt window. The X axis represents a linear map of the artificially concatenated *Wolbachia* pseudo-contig from *A. vulgare*

and the Y axis (1–3,000) represents relative sequence coverage. The count file was generated from a BAM file using IGV Tools (Thorvaldsdottir et al. 2012). 97.1 % coverage was obtained

length=200 bases), a manual assembly produced 523 contigs with N50 of 5826, a maximum length of 26,288 bases. This used over 87 % of the oligo-captured reads (31.81/36.45 million reads) providing a total length of 1.32 Mb, representing 75.5 % of the estimated length of the genome and 79.5 % of the partially known sequence.

To ensure that all the contigs built from the oligo-captured DNA were *wVulC* sequences, we aligned and ordered them to the previously sequenced *wVulC* genome using Mauve (Darling et al. 2010) and r2cat (Husemann and Stoye 2010). Overall, 95.7 % of the bases in contigs built from oligo-captured DNA were *wVulC*, which represents 76 % of the 1.66 Mb concatenated reference genome. 61 contigs did not match the incomplete *wVulC* genome (Fig. 4). Several of these contain sequences that show similarities with other *Wolbachia* strain sequences (0.1 %) while a contig of 16 kb appeared to be a region from the *Armadillidium vulgare* 18S ribosomal gene (1.22 %), certainly host DNA contamination (Fig. 4). Since the *wVulC*



**Fig. 4** Capture efficiency of the *wVulC* *Wolbachia* genome. Bar graph shows the distribution of nucleotides out of the total assembled contigs that show significant BLAST scores to the partial *wVulC* genome, other *Wolbachia* or the *A. vulgare* host. The “Other” category corresponds to sequences with no significant BLAST similarity to any sequences in the NCBI database

genome is not finished and gaps remain between the 10 contigs, analysis of the contigs built from oligo-captured DNA that matched other *Wolbachia* strains or didn’t match any known sequence in NCBI (2.41 %) are being used to complete the reference genome sequence as they may represent yet unknown *wVulC* sequences.

Our results confirm and extend the results of Kent et al. (2011) and demonstrate that this method can potentially be used with any *Wolbachia* strain and on any endobacterium with either a reference genome or a highly similar sequence, providing an approach for isolating a significant fraction of symbiont DNA from host DNA for sequencing and comparative genomic analysis. In this context, of interest is the observation that both *wBm* and *wVulC* were successfully purified away from host and mitochondrial DNA with the same bait library even though they are members of phylogenetically distant clades. In filarial nematodes, this approach will enable rapid isolation and analysis of *Wolbachia* strains from worldwide populations to identify polymorphisms related to drug discovery initiatives and evolutionary analysis of *Wolbachia* prevalence and distribution. In isopods, this approach will allow DNA isolation for genomic comparison of *Wolbachia* strains, which induce either various types of feminizing phenotypes or cytoplasmic incompatibility. For example, the isopod *A. vulgare* may harbor another *Wolbachia* strain, *wVulM*, which has a lower feminizing effect (~70 %) than the *wVulC* strain (~80 %) (Cordaux et al. 2004). In the *Porcellionides pruinosus* complex of species, 3 distinct feminizing *Wolbachia* strains have been identified which are present in ~60 % of populations where *Wolbachia* are present only in females or ~90 % in populations where both males and females are infected (Marcadé et al. 1999; Lefebvre and Marcadé 2005), a situation also encountered in populations of *Oniscus asellus* (Rigaud et al. 1999). Further, most *Wolbachia* strains infecting isopods induce a feminizing phenotype (Bouchon et al. 2008), however, 3 of them induce cytoplasmic incompatibility (Legrand et al. 1978; Moret et al. 2001). One of them, *wCon*, which infects *Cylisticus convexus*, is closely related to *wVulC* (Cordaux et al. 2012). Genomic comparisons

will thus be useful in helping decipher the evolution of *Wolbachia* and its various biological manifestations.

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