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SUMMARY

Genome assemblies can form the basis of comparative analyses fostering insight into the evolutionary genetics of a parasite's pathogenicity, host–pathogen interactions, environmental constraints and invasion biology; however, the length and complexity of many parasite genomes has hampered the development of well-resolved assemblies. In order to improve *Trichinella* genome assemblies, the genome of the sylvatic encapsulated species *Trichinella murrelli* was sequenced using third-generation, long-read technology and, using syntenic comparisons, scaffolded to a reference genome assembly of *Trichinella spiralis*, markedly improving both. A high-quality draft assembly for *T. murrelli* was achieved that totalled 63·2 Mbp, half of which was condensed into 26 contigs each longer than 571 000 bp. When compared with previous assemblies for parasites in the genus, ours required 10-fold fewer contigs, which were five times longer, on average. Better assembly across repetitive regions also enabled resolution of 8 Mbp of previously indeterminate sequence. Furthermore, syntenic comparisons identified widespread scaffold misassemblies in the *T. spiralis* reference genome. The two new assemblies, organized for the first time into three chromosomal scaffolds, will be valuable resources for future studies linking phenotypic traits within each species to their underlying genetic bases.

Key words: Genomics, long-read assembly, synteny, scaffolding, repetitive DNA, *Trichinella*, chromosome, nematode.

INTRODUCTION

Genome evolution influences a parasite's pathogenicity, host–pathogen interactions, environmental constraints and invasion biology (reviewed in Garrett *et al.* 2006; Hoberg and Brooks, 2008; Dittmar, 2009), and genome assemblies form the basis of comparative genomic analyses. These encompass not only nucleotide sequence order and gene content, but also DNA structure. Changes in genomic architecture are known to alter gene regulation affecting phenotypes (Lanctôt *et al.* 2007; Raffaele and Kamoun, 2012), whereas structural conservation generally maintains shared features among related pathogens (Ghedin *et al.* 2004; Peacock *et al.* 2007). Where repetitive elements render assembly of a given genome difficult, shared genome structure (synteny) can enhance assembly efforts (Richter *et al.* 2007; Assefa *et al.* 2009; Husemann and Stoye, 2010). Given that closely related organisms typically maintain appreciable

synteny, the genome assembly of one organism can improve the assembly of a closely related genome.

A number of interesting and important questions concerning differences among species in the parasitic genus *Trichinella* would benefit from high-quality genome assemblies. Nematodes belonging to the genus *Trichinella* are parasites that infect domestic and wild mammals that eat meat. Pork and wild game (including feral pigs, wild boars, bears, dogs, wild cats and walruses) have been implicated as sources of human infections (reviewed in Pozio, 2007). As such, they threaten human health and pose a concern for food safety. As most sylvatic species of *Trichinella* do infect humans but do not infect swine well, understanding the differences in pathogenicity and host range for species within this genus is a subject of considerable interest. In temperate North America, *Trichinella spiralis* and *Trichinella murrelli*, differ in their propensities to infect and persist in domestic swine (Kapel and Gamble, 2000; Kapel *et al.* 2005), although the bases for this difference are not understood. *Trichinella spiralis* has tremendous reproductive capacity in swine, and has historically been the key source of human disease via consumption of infected, undercooked pork products. *Trichinella*

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murrelli, on the other hand, has not been implicated in infections derived from pork, but exhibits a predilection for wild carnivore hosts; human infections have been acquired from bear meat (Hall *et al*. 2012). Experimentally infecting swine with *T. murrelli* produces few muscle larvae; however, it confers some resistance to challenge with *T. spiralis* (D.E. Hill, personal communication, 2016). In North America, *T. murrelli* has been identified as the predominant *Trichinella* species in wild hosts (Zarlenga *et al*. 1991). Consequently, *T. murrelli* threatens the health of those consuming wild game. Genetic investigations might help establish the mechanisms enforcing distinctions in their host affinities.

Genomic comparisons of *T. spiralis* and *T. murrelli* would provide the most comprehensive evaluation of the genetic differences between these two species. Until recently, a high-quality draft genome assembly was only available for *T. spiralis* (Mitreva *et al*. 2011); this assembly has been used by the research community extensively (157 citations since its publication, as of this writing). In 2016, draft genomes were released for every known species of *Trichinella* (Korhonen *et al*. 2016), providing the first opportunity for in-depth interspecific comparisons. Both published *T. spiralis* assemblies were constructed from short-read shotgun sequencing, resulting in average contig and scaffold lengths of 7042 and 5875 bp, respectively. Those data were derived by combining paired-end read information (Mitreva *et al*. 2011; Korhonen *et al*. 2016), fosmid end sequencing (Mitreva *et al*. 2011), with bacterial artificial chromosome end sequencing (Mitreva *et al*. 2011). Further consolidation of contigs into larger scaffolds or chromosomes was hindered by the approximately 20% repetitive DNA content in these genomes and by their AT-rich nature (67% adenines or thymines). Karyotypes indicate that *Trichinella* harbour but two autosomes and an XO sex-determination chromosome (Mutafova *et al*. 1982). Presently, the genomes are fragmented into thousands of sequenced components. Additional measures are needed to condense these genome assemblies towards finished genomes made up of only three chromosomes. Encouragingly, a recent analysis (Foth *et al*. 2014) has enabled each of the 11 longest *T. spiralis* scaffolds to be placed into one of three chromosomes. This was accomplished by analysing single-copy genes with orthologues in the clade I whipworm *Trichuris muris* (Blaxter *et al*. 1998; Holterman *et al*. 2006). Additional consolidation of *T. spiralis* contigs into chromosomes may be possible by comparing syntenic regions among contigs of closely related *Trichinella* species.

The prospects for success in using syntenic relationships among encapsulated species of *Trichinella* to aid in their assembly are favoured, given their relatively close relationships. Direct comparisons of *T. spiralis* and *T. murrelli* mitochondrial DNA revealed complete conservation of gene order, despite an average divergence of 9·5–11·8% at the nucleotide level (Webb and Rosenthal, 2011; Mohandas *et al*. 2014). Single-copy orthologue synteny was found to be appreciable across the nuclear genome (Korhonen *et al*. 2016). The two species are thought to have shared a common ancestor between 7 and 10 million years (MY) ago, based on comparisons of nuclear and mitochondrial gene sequences (Zarlenga *et al*. 2006; Korhonen *et al*. 2016). Furthermore, a higher degree of synteny has been reported among encapsulated trichinellids (including *T. spiralis* and *T. murrelli*) than non-encapsulated species, with a syntenic correlation of 0·118 for genes in the nuclear genomes of *T. spiralis* and *T. murrelli* (Korhonen *et al*. 2016). Therefore, in order to improve *Trichinella* genome assemblies, we used third-generation, long-read technology to sequence through highly repetitive regions of the genome. We sequenced the genome of the North American sylvatic species, *T. murrelli,* using long-read technology and generated a *de novo* assembly which was polished using high-quality base calls derived from Illumina paired-end sequencing. This sequence was exploited to improve the assemblies of both *T. murrelli* and *T. spiralis*. Using syntenic information between the two assemblies, *T. murrelli* contigs were placed into scaffolds representing putative chromosomes. Additionally, *T. spiralis* contigs were added to chromosome groupings based on synteny with the *T. murrelli* assembly. Finally, certain regions of *T. spiralis* scaffolds were reoriented based on putative scaffolding errors revealed by comparisons with longer *T. murrelli* contigs achieved through long-read sequencing. As a result, we achieved a 10-fold improvement in overall genome assemblies of both *T. murrelli* and *T. spiralis*.

MATERIALS AND METHODS

Genome sequencing and assembly

Trichinella murrelli isolate ISS35 was used as the source of genomic DNA for sequencing. This isolate was collected from a black bear (*Ursus americanus*) in North America in 1982, and had been passed through mice approximately twice per year since its isolation. In 2011, Swiss-Webster mice were infected with *T. murrelli* ISS35, and worms were collected from eviscerated mouse carcasses using standard digestion procedures. In order to minimize host DNA contamination, collected worms were pre-treated with DNAse I for 10 min and washed extensively in tap water prior to parasite DNA extraction. Following digestion with proteinase K in 10 mM Tris pH 8.5 with 0.5% sodium dodecyl sulfate (SDS), nucleic acids were collected by phenol–chloroform

extraction and ethanol precipitation. The sample was then treated with a cocktail of RNAse A and RNAse T1. The final genomic DNA preparation was checked for purity using gel electrophoresis and spectrophotometric analysis by NanoDrop. Genomic DNA was quantified using the Stratagene Quantifluor system.

Trichinella murrelli genomic DNA was sequenced using Pacific Biosciences (PacBio) (Menlo Park, CA) long-read technology at the University of Maryland School of Medicine's Institute for Genome Sciences. One PacBio Ultra Long Insert Library was generated. This library was loaded onto four PacBio RS II P6-C4 SMRT-cells in order to generate long-read sequences. Long-read sequencing generated 334 806 reads averaging 9756 bp in length (S.D. = 6637) for a theoretical 50X coverage of the expected 65 Mbp genome.

Long reads were assembled using SMRT Analysis v. 2.3.0, assuming a total assembly size of 65 Mbp (based on the estimated size of the 2011 *T. spiralis* genome assembly). Base calls in the long-read assemblies were improved using Illumina paired-end short reads of 300 bp, obtained using the v3 run capabilities of an Illumina MiSeq, which provided a theoretical 150X coverage prior to quality control of reads. Each Illumina read was trimmed such that no more than two nucleotides within each read had a Q score of <30 (0·1% chance of erroneous base call). Illumina paired-end reads were then aligned to the long-read contigs using the Geneious assembler (Geneious v. 8.0.6, http://geneious.com (Kearse *et al*. 2012)) with medium–low sensitivity and five iterations, resulting in an average of 80X coverage for each base in the SMRT Analysis assembly. To assign nucleotides for the final assembly, consensus base calls were derived from Illumina reads where SMRT assembly contig coverage was greater than 10X. In order to call a base as a heterozygote, minority reads had to account for >10% of all Illumina base calls at that locus. For those sites with less than 10X coverage, the original SMRT Analysis base call was retained.

Repetitive DNA was identified using a combination of *ab initio* repeat finding using RepeatModeler v. 1.0.8 (Smit and Hubley, 2008) and a library-based approach in RepeatMasker v. 4.0.5 (Smit *et al*. 2013). Within RepeatModeler, *ab initio* analysis was conducted in RepeatScout v. 1.0.5 and RECON v. 1.0.8 to produce a library of repeat elements. RepeatMasker was then used to identify each individual repeat in the assembly contigs and classify it as either a simple repeat or as an interspersed repetitive element. RepeatMasker was also used independently, specifying *T. spiralis* repeat elements from Repbase, including the Ginger2 DNA transposon, the R2 family of non-LTR retrotransposons, the Utopia family of non-LTR retrotransposons, and the

TSRP-1 repetitive sequence. Finally, Phobos was used to search for degenerate simple repeats fewer than 5 nt in length, allowing for 20% divergence from the core repeat pattern. Subsequent analyses regarding alignment of *T. murrelli* contigs with *T. spiralis* scaffolds were conducted using the masked DNA file output from RepeatMasker so as to minimize errors and ambiguities that typically ensue when repeats are included.

The new long-read assembly was compared with existing *Trichinella* genome assemblies using QUAST v.4.0 (Gurevich *et al*. 2013) for length metrics and CEGMA v.2.4.010312 (Parra *et al*. 2007) and BUSCO v.2.0 (Simão *et al*. 2015) in order to examine completeness of coding regions in the genome. Default QUAST settings were used to determine the shortest contig or scaffold that was required to encompass 50% (N50) or 90% (N90) of the total length of all contigs or scaffolds for each assembly in the comparison. CEGMA analysis was conducted in the Cyverse.org Discovery Environment (Merchant *et al*. 2016) using the 248 most highly conserved core eukaryotic gene set (CEGs). The new long-read *T. murrelli* assembly, the 2011 *T. spiralis* assembly and the 2016 *T. murrelli* short-read assembly were analyzed for complete, partial and missing CEGs. BUSCO was used to examine single-copy orthologues (BUSCOs) associated specifically with nematodes for both the novel long-read *T. murrelli* assembly and the 2016 *T. murrelli* short-read assembly. The BUSCO.py program was run with the 'nematoda_odb9' lineage in genome mode with species set to 'Caenorhabditis', and complete, duplicated and fragmented BUSCOs were reported.

Synteny and chromosomal scaffolding

ProgressiveMAUVE v. 20150226 (Darling *et al*. 2010) was used to derive an initial overview of synteny between the *T. murrelli* corrected contigs and 2011 *T. spiralis* scaffolds. Using 2011 *T. spiralis* scaffolds as reference sequences, *T. murrelli* contigs were reordered using the Move Contig feature in progressiveMAUVE to achieve optimal alignment. Alignments of long collinear blocks (LCBs) were visualized in the MAUVE viewer, relying on LCBs with a weight >10 000 to assess synteny. LCBs with lesser weight were reserved for MUMmer analysis outlined below.

In order to maximize the organization of both assemblies, 2011 *T. spiralis* scaffolds were divided into putative chromosomes according to single-copy orthologue analyses (Foth *et al*. 2014). Chromosome 1 comprised scaffolds GL622784, GL622787, GL622790, GL623393 and GL624340 of the 2011 *T. spiralis* assembly. Chromosome 2 encompassed scaffolds GL622785, GL622788 and GL623868, while scaffolds GL622791, GL622792 and GL622789 were designated as Chromosome X.

Table 1. *Trichinella* genome assembly sources and statistics

	2011 *T spiralis* assembly	2016 short-read *T. murrelli* Assembly	2017 long-read *T. murrelli* assembly	2017 *T. spiralis* assembly – scaffolding to long-read *T. murrelli* assembly
Description	ISS195	ISS417	ISS35	ISS195
Country of origin	USA	USA	USA	USA
Host	Domestic pig	Coyote	Black bear	Domestic pig
Genome size (bp)	63 521 838	49 039 267	63 190 608	63 521 838
Scaffolds/contigs	6859/9261	5255/6430	543/653	6844
N50 (bp)	6 373 445/76 808	106 482	17 128 578/571 829	17 530 227
N90 (bp)	2049/1512	15 500	27 732/27 652	2049
Max scaffold/contig length	12 041 450/482 167	489 069	20 088 213/3 626 038	20 007 485
Genome GC content (%)	33·9	33·6	33·9	33·9
Repetitive sequences (%)	18	18·78	30·3	18
CEG completeness: complete; partial (%)	93·6; 94·4	96·0; 97·2[a]	94·0; 94·8	93·6; 94·4
BUSCO analysis: complete; duplicated; fragmented (%)		78·2; 0·4; 6·9	88·8; 4·0; 7·3	

[a] As reported by Korhonen *et al*. (2016).

MUMmer v. 3.23 was used to examine local areas of synteny among *T. murrelli* masked contigs and *T. spiralis* putative chromosomes. Using the NUCmer application within MUMmer, maximal unique matches (MUMs) were identified between *T. murrelli* contigs >7000 bp (60% of the total length of all *T. murrelli* contigs) and each putative *T. spiralis* chromosome. Each group of *T. murrelli* contigs was then placed in optimal alignments relative to *T. spiralis* scaffolds using the move contig feature in MAUVE. These optimized *T. murrelli* putative chromosomes were concatenated into scaffolds using 100 indeterminate bases ('N') to signify gaps between contigs that occur within any *T. spiralis* scaffold, and 1000 bases between contig ends that occur between *T. spiralis* scaffolds signifying less confidence in the ordering of the final placement of *T. murrelli* contigs in the chromosomal scaffolds. Subsequently, 2011 *T. spiralis* scaffolds were broken at assembly gaps for reassembly based on synteny with the new *T. murrelli* chromosomes. A custom PerlScript was written to identify stretches of N's that were equal to 10 nucleotides or longer, break the scaffold at each such instance, and place each sub-sequence corresponding to the sequence between adjacent assembly gaps into a new FASTA file, retaining the N's present in the original scaffold. This process was intended to retain as much information as possible from the original scaffolding efforts of Mitreva *et al*. (2011). The resulting sequences are heretofore termed '*T. spiralis* contigs'. After breaking the *T. spiralis* scaffolds into their component parts, these *T. spiralis* contigs were matched to putative *T. murrelli* chromosome sequences using MUMmer. It was expected that this reverse matching would correctly identify not only sequences from the original 11 long *T. spiralis*

scaffolds, but contigs that belong to each chromosomal group that were not hypothesized by Foth *et al*. (2014), thus lengthening chromosomal sequences of the *T. spiralis* assembly.

All statistical analyses were conducted in R. The distribution of contig/scaffold lengths was determined using the density function. Median coverage of *T. murrelli* contigs was determined using Rsamtools, extracting read counts from BAM files of short-reads mapped to *T. murrelli* long-read contigs. Output of median coverage was graphed in Microsoft Excel.

RESULTS

Subjecting an isolate of *T. murrelli* to long-read sequencing using the Pacific Biosystems technology generated 334 806 individual reads averaging 19·6 kbp in length; the longest read extended 51 413 bp. Approximately 3% of the reads corresponded to adapter dimers and inserts <100 bp. Following filtering, nearly 224 000 reads were deemed appropriate for use in *de novo* assembly. These reads averaged 13 786 bp, with an N50 of 19 125, and an average read score (probability of being correct) of 0·85.

After filtering for quality, long reads were assembled into a high-quality draft genome with a total size of 63·2 Mbp (Table 1). Over half of the total assembly's length was derived from just 26 contigs, each of which is longer than 571 000 bp (N50). The N90 is 27 652 bp, and the largest contig is over 3.6 Mbp. Short-read, paired-end sequences were used to verify and correct small errors in the assembled contigs. Paired-end reads of 300 bp were aligned to *T. murrelli* assemblies, providing 80X coverage on average; consensus
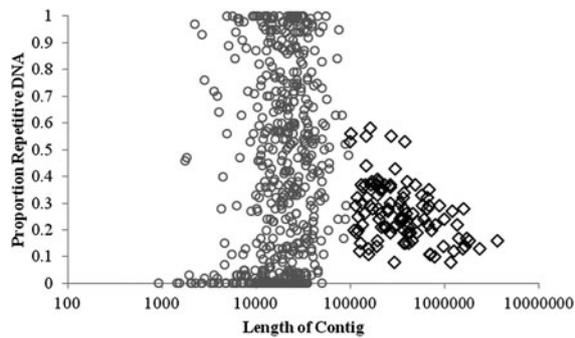
Fig. 1. Proportion of each long-read *T. murrelli* contig that is occupied by repetitive DNA elements. Those contigs greater than 100 000 bp in length (black diamonds) have much more consistent representation of repetitive DNA than smaller fragments (grey circles). The general lower proportion of repetitive sequences may contribute to the ease with which these portions of the genome are assembled. This would lend some greater confidence to these longer contigs when trying to understand reorganization within the genome relative to other closely related genomes.

sequence of the final contigs was determined by those short-read base calls comprising >10% of the coverage. Nearly all of the corrections involved the insertion or deletion of a single nucleotide, following a run of that same nucleotide. Heterozygous bases accounted for 0·5% of all nucleotides or 1 heterozygous base every 200 bp. Attempts at improving the assembly via scaffolding using SSPACE did not further reduce the number of contigs.

Based on Recon and RepeatScout *ab initio* libraries, RepeatMasker identified 13 843 interspersed repeat elements within the final contigs, accounting for 15 795 525 bp (25%) of the *T. murrelli* genome. Additionally, RepeatMasker analysis indicated that 2·7% of the genome could be categorized as short simple repeats. Phobos analysis, allowing for 20% degeneracy in any repeat motif, found 3 355 004 bases (5·3%) within tandem repeats with a 5 nucleotide repeat motif or less (nearly twice as many as detected by RepeatMasker), bringing the total amount of repetitive DNA to 30·3%. Repetitive elements were sub-divided into seven categories. There were 21 SINEs, 929 LINEs, 2856 LTR, 3281 non-LTR DNA elements, 6596 low complexity sequences, 160 satellite elements and 36823 simple repeats. These were spread across all *T. murrelli* contigs, but were at highest concentration in contigs <100 000 bp in length. In order to determine where the repeat sequences were concentrated, we plotted repeat content as a function of contig length. Repeats comprised a smaller proportion of contigs larger than 100 000 bp (Student's *t*-test, $P < 0.0001$); the variance in repeat composition, which ranged from 8 to 58% in contigs greater than 100 000 bp, was reduced when compared with smaller contigs (Fig. 1).

A survey of the core eukaryotic genome (CEGs) was conducted to assess the completeness of the protein coding portions of the assemblies. CEGMA analysis indicated that the 94·0% of CEGs were complete in the new *T. murrelli* long-read assembly, a minor improvement over the 93·6% completeness of the 2011 *T. spiralis* assembly. Using other search parameters, the 2016 short-read assembly of *T. murrelli* (Korhonen *et al*. 2016) had been reported to have 96·0% of CEGs complete. However, subjecting that assembly to search parameters identical to those used above, we found the 2016 *T. murrelli* short-read assembly to have 92·7% of CEGs complete. Similar improvements were found in the new assembly when employing BUSCO to determine the number of nematode-specific single-copy orthologues recovered in each. BUSCO found that 88·8% of nematode single-copy orthologues were complete in the novel long-read assembly whereas the 2016 short-read assembly revealed only 78·2% of single-copy orthologues that were complete (Table 1). Therefore, each assembly was judged successful in assembling areas of the genome carrying coding information, with a 10% improvement in recovering nematode-specific single-copy orthologues in the long-read assembly.

Comparison of T. murrelli *long-read genome assembly with 2011* T. spiralis *assembly and 2016 short-read* T. murrelli *assembly*

The *T. murrelli* assembly presented here represents approximately a 10-fold improvement over the published 2011 *T. spiralis* genome assembly at the contig level. The 2011 assembly of *T. spiralis* in GenBank (Accession # ABIR02·1, accessed 12/9/2015), which was assembled via whole genome shotgun sequencing and hierarchical mapping based on fosmid and BAC end-sequencing, contained 6859 scaffolds comprising 9261 contigs. This *T. spiralis* assembly, with a total length of 63 521 838 bp, had an N50 of 76 808 and an average length of 9261 bp/scaffold. Of those scaffolds, 3008 were smaller than 1000 bp. Similarly, the recently published short-read *T. murrelli* assembly comprised 5255 scaffolds with an N50 of 106 482 bp with 3913 scaffolds <1000 bp in length. Notably, the short-read *T. murrelli* assembly totalled just over 49 Mbp, suggesting that over 14 Mbp were not accounted for in comparison with the 2011 *T. spiralis* assembly or the long-read *T. murrelli* assembly presented here. By contrast, our long-read *T. murrelli* assembly is condensed into nearly 15-fold fewer total contigs (Fig. 2A), and eight times as many of its contigs are larger than 1000 bp (see distribution of scaffold/contig lengths for each assembly in Fig. 2B). Our long-read *T. murrelli* N50 is 7·5 times the size of the 2011 *T. spiralis* N50 and five times longer than the recently published short-
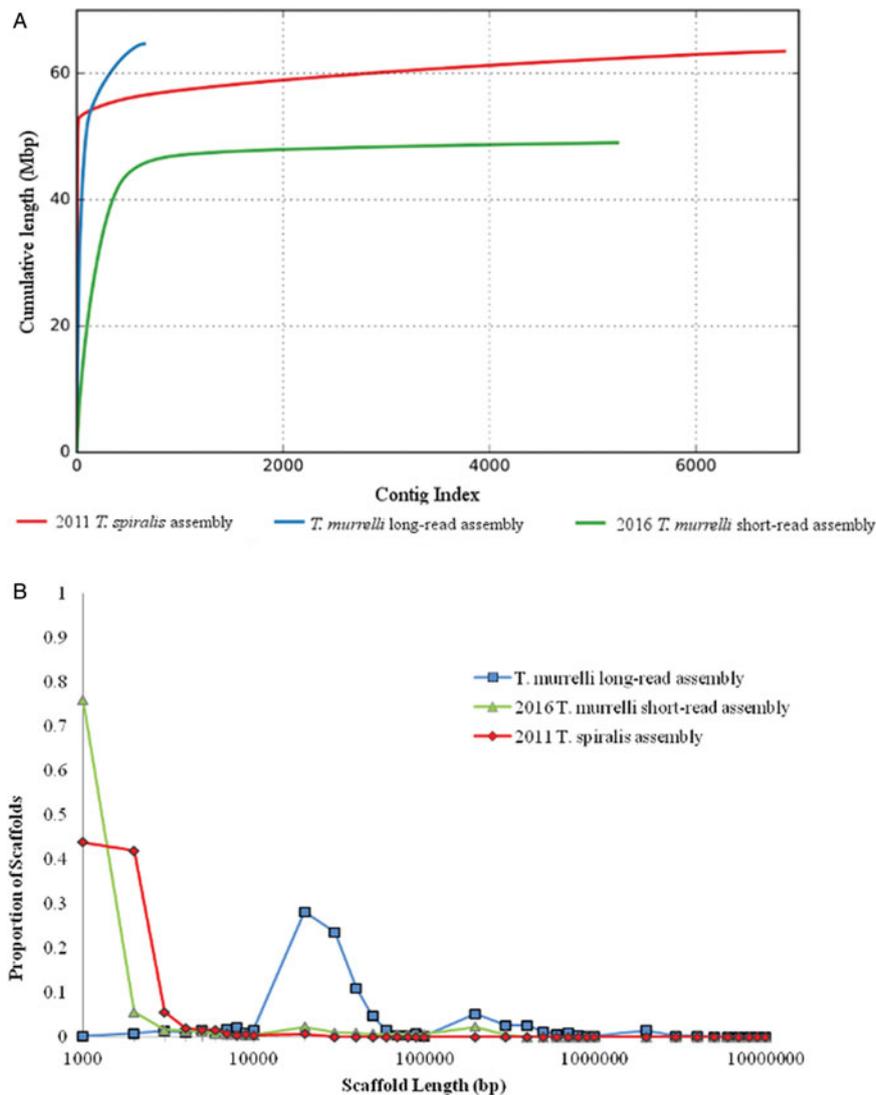
Fig. 2. Comparative summary of *Trichinella* genome assemblies. (A) Cumulative length of contigs or scaffolds when arranged from longest (contig index 1) to shortest for each assembly as implemented in Quast. The *T. murrelli* long-read assembly reaches complete length in many fewer contigs than either the 2011 *T. spiralis* or 2016 *T. murrelli* short-read assembly. (B) For the 2011 *T. spiralis* and 2016 *T. murrelli* short-read assemblies, 91·6% and 83·3% of the contigs are smaller than 3000 bp, respectively; only 2·2% of contigs in the long-read *T. murrelli* assembly are that small. The 63% of all *T. murrelli* long-read contigs fall between 20 000 and 50 000 bp. An additional 10% of long-read *T. murrelli* contigs fall between 200 000 and 500 000 bp. The long-read *T. murrelli* assembly has substantially improved N50 (571 829 bp) (Table 1), despite the fact that the longest *T. spiralis* scaffold (GL622787 = 12·1 Mbp) is over three times the length of the longest *T. murrelli* contig (unitig 1112 = 3·6 Mbp).

read assembly of *T. murrelli*, achieving an average contig length 13·9 times larger for *T. murrelli* than for 2011 *T. spiralis* scaffolds. Almost 8% of positions had been designated as N (indeterminate sequences) within the scaffolds of the 2011 *T. spiralis* assembly, but fewer than 0·1% of the positions in our long-read *T. murrelli* assembly are ambiguous in this way. The recently published short-read *T. murrelli* assembly had small numbers of indeterminate bases (0·6%) but, as noted previously, had 22% less total sequence than the assembly provided here.

Alignment of the *T. murrelli* assembly presented here and the 2011 *T. spiralis* assembly showed remarkable overall synteny. After reordering the

T. murrelli contigs in MAUVE in order to maximize alignment of *T. murrelli* contigs with *T. spiralis* scaffolds, 343 LCBs weighted >10 000 were observed (Fig. 3). At the broadest level, approximately 51·5 Mbp (81·7%) of the assemblies were collinear (ignoring gaps between collinear blocks). Similarly, MUMmer alignment of the 2011 *T. spiralis* assembly and 2016 *T. murrelli* long-read contigs resulted in a total of 51·0 Mbp of aligned sequence, nearly identical to that assessed by alignments in MAUVE. Identity among local MUMmer alignments ranged from 55 to 100% with an overall 92·4% identity across all alignments. Maximal unique matches ranged in size from 67 to 70 998
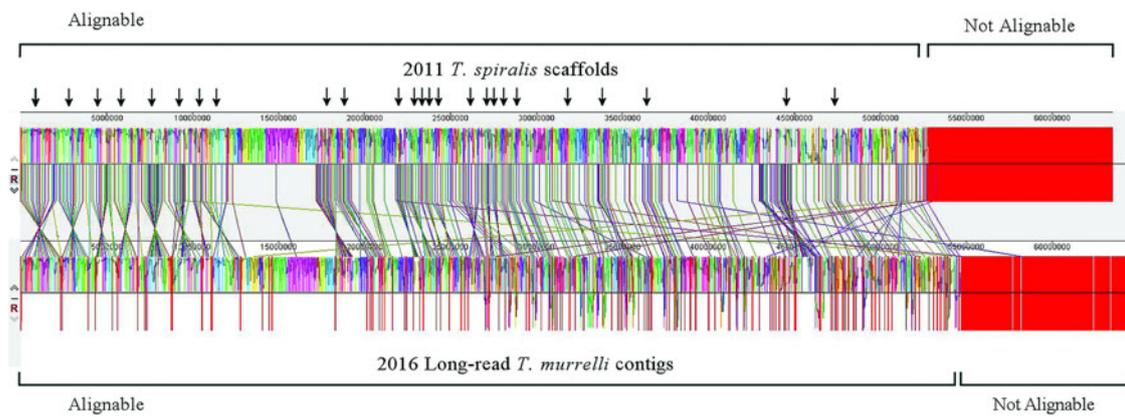
Fig. 3. ProgressiveMAUVE alignment of 2011 *T. spiralis* and *T. murrelli* long-read assemblies using the 'Move Contig' feature to optimize placement of *T. murrelli* contigs relative to the 2011 *T. spiralis* scaffolds. Approximately 51·5 Mbp were located within LCBs, accounting for 82% of the entire assemblies. Arrows indicate the location of local syntenic rearrangements wherein collinear blocks have the same orientation, but have local rearrangements such that each block is precisely reversed in the *T. murrelli* assembly with respect to the *T. spiralis* assembly. Note that each rearrangement occurs within an intact *T. murrelli* contig, denoted by the vertical red lines of the lower assembly. There are 25 different regions of the alignment that follow this pattern.

bp in the masked alignment. For those MUMs larger than 10 000 bp (signifying strong confidence in the correct assignment of orthologous regions in the two genomes), the overall identity was 92·6% and accounted for 48·5% of the total length of aligned sequences. There remained 472 *T. murrelli* contigs that could not be aligned with any confidence; these averaged 20 902 bp in length and were enriched for repetitive sequences. Similarly, 6841 *T. spiralis* scaffolds could not be aligned; these averaged 1561 bp in length. A comparison of the sizes of contigs/scaffolds that could be aligned with those that could not be is shown in Fig. 4. Those contigs and scaffolds that could not be aligned confidently were not included in subsequent chromosomal organization.

The chromosome hypothesis

Foth *et al*. (2014) hypothesized how the largest scaffolds in *T. spiralis* should be grouped into chromosomes, using single-copy orthologues mapped to the whipworm, *T. muris*. In order to assess whether our data supported the organization of contigs into chromosomes, we examined median short-read coverage of *T. murrelli* contigs, expecting coverage of the sex chromosome to be 3/4 the coverage of autosomal sequences (as would be true for an XO sex-determining system in the case where male and female DNA were equally represented in the sequenced template). The coverage of 57 long-read *T. murrelli* contigs ranged between 94 and 111 reads/site, whereas the coverage of another 52 contigs ranged between 72 and 87 reads/site; we supposed that these distinct coverage groupings might correspond, respectively, to the autosomes and to the X chromosome (Fig. 5). Another 545 contigs, mapped fewer than 70 reads/site, were ignored for the purposes of this analysis. Confirming this

supposition, all of the contigs so classified as autosomal also matched sequences aligning with autosomal sequences of *T. spiralis* in the MUMmer analyses. Of the contigs classified as belonging to the X chromosome, 83% matched sequences designated as residing on the X chromosome of *T. spiralis* by Foth *et al*. 2014. The nine exceptions (contigs that matched sex chromosome coverage ratios, but had been judged previously to be autosomal by single-copy orthologue content) merit further analysis to assess their chromosomal assignment. Interestingly, most contigs whose coverage did not match chromosomal expectations are shorter than 100 000 bp and harbour much repetitive DNA, which may compromise attempts at ascertaining their chromosomal location. The evidence from two independent lines justified organizing most of these data into one of three chromosomes.

Employing the chromosomal framework of Foth *et al*. 2014, we assigned scaffolds of *T. spiralis* into their respective chromosomes. We placed homologous *T. murrelli* long-read contigs into corresponding chromosomal files. We then optimized the order of long-read *T. murrelli* contigs using the 'move contig' process of MAUVE, relying on the original *T. spiralis* scaffolds as a baseline for organization; the resulting *T. murrelli* and *T. spiralis* hypothetical chromosomes are depicted in the upper portion of each panel in Fig. 6 and denoted as 'Before', signifying that *T. spiralis* sequences had not yet been optimized relative to long-read *T. murrelli* contigs. While some local inconsistencies persisted among the *T. spiralis* scaffolds and *T. murrelli* contigs, the general agreement between the two assemblies indicates that scaffolding of *T. murrelli* contigs based on *T. spiralis* scaffolds was tenable. This resulted in three *T. murrelli* chromosomal scaffolds with a total size of 52·7
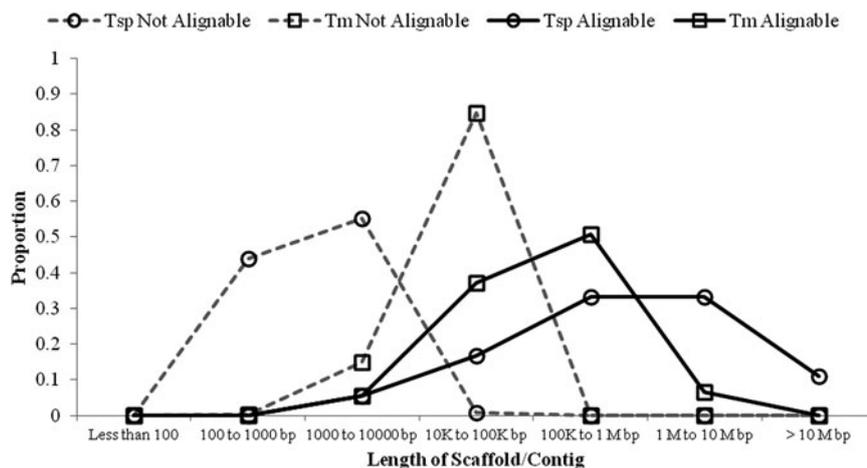
Fig. 4. The distribution of *T. spiralis* (Tsp) scaffold and *T. murrelli* (Tm) contig lengths divided into those that could or could not be confidently aligned in initial MAUVE and MUMmer alignments. For each species, those sequences which could not be aligned were significantly smaller than those that aligned with confidence (Student's *t*-test, *P*<0·01 for both comparisons).

Mbp, and increased the scaffold N50 to 17·1 Mbp. These results represent our current best evidence for chromosome organization for *T. murrelli*.

Certain segments of the MAUVE alignments between *T. murrelli* and the 2011 *T. spiralis* genome revealed a highly consistent criss-cross pattern (highlighted with arrows in Fig. 3) that might be mistaken for large genomic inversions. A simple inversion would have reversed the orientation of those LCBs within such segments, but close inspection of these regions showed them to retain their orientation, despite the reversal of LCB order (Fig. 7). This incongruous pattern always occurred within a single *T. murrelli* contig but among *T. spiralis* contigs that had been connected by 'assembly gaps' in the 2011 *T. spiralis* assembly scaffolding process. If biological in origin, this result would require rampant, local rearrangements in the order of LCBs. Misassembly across assembly gaps in 2011 *T. spiralis* scaffolds would offer a simpler explanation for this pattern.

We therefore reoriented LCB's in the 2011 assembly of *T. spiralis* to conform to the LCB orders indicated by long-read *T. murrelli* contigs. For each set of scaffolds belonging to a putative chromosome (1, 2 or X), we deconstructed *T. spiralis* scaffolds into their component contigs by breaking them at assembly gaps of 10 bp or longer. We then aligned these contigs to *T. murrelli* chromosome scaffolds, and substantially greater agreement resulted. Finally, we reinserted assembly gaps between *T. spiralis* contigs so as to match the total length of the original scaffolds, preserving spacing information gleaned from the 2011 assembly. As a result, we were able to condense the assembly of *T. spiralis* into fewer component pieces with improved confidence in ordering within scaffolds (Fig. 6, 'After').

Pairs of assembled genomes often require transpositions and inversions in order to be fully reconciled; fewer such rearrangements are required to harmonize highly syntenic assemblies. The procedures described above markedly reduced the number of such rearrangements between genome assemblies of two encapsulated members of the genus *Trichinella*. When initially compared, our assembly of *T. murrelli* required 35 inversions and 197 transpositions of LCBs in order to be reconciled with the 2011 assembly of *T. spiralis* (an inversion to transposition rate of 1:5·6). Of these, 24 LCB inversions and 155 LCB transpositions occurred on autosomes, whereas 11 inversions and 42 transpositions on putative sex chromosomes. After identifying and correcting for putative local inconsistencies in the 2011 assembly of *T. spiralis*, the total number of inversions and transpositions dropped to 7 and 14, respectively. All seven of these remaining inversions occurred on autosomal sequences, as did 11 of the 14 transpositions. This resulted in an inversion:transposition ratio of 1:1·6 for autosomes and 0:3 for the X chromosome. Assuming two chromosome breakages for each inversion and three for each translocation as well as 7–10 MY of divergence time, this translates to 0·15 to 0·21 chromosome breakages/Mbp/MY.

DISCUSSION

We sequenced the genome of the sylvatic *Trichinella* species, *T. murrelli*, using third-generation, single-molecule, long-read technology. The final assembly condensed to 653 contigs; half of the assembled genome occurred in contigs longer than 571 031 bp (N50). This represents an approximately 10-fold improvement over the 2011 assembly of the *T. spiralis* genome in terms of length and completeness. The genome assemblies were generally in agreement with 51·5 Mbp of contiguous sequence. *Trichinella murrelli* long-read contigs spanned many gaps in the 2011 *T. spiralis* scaffolds, providing great empirical support for the *T. murrelli* contigs; we therefore
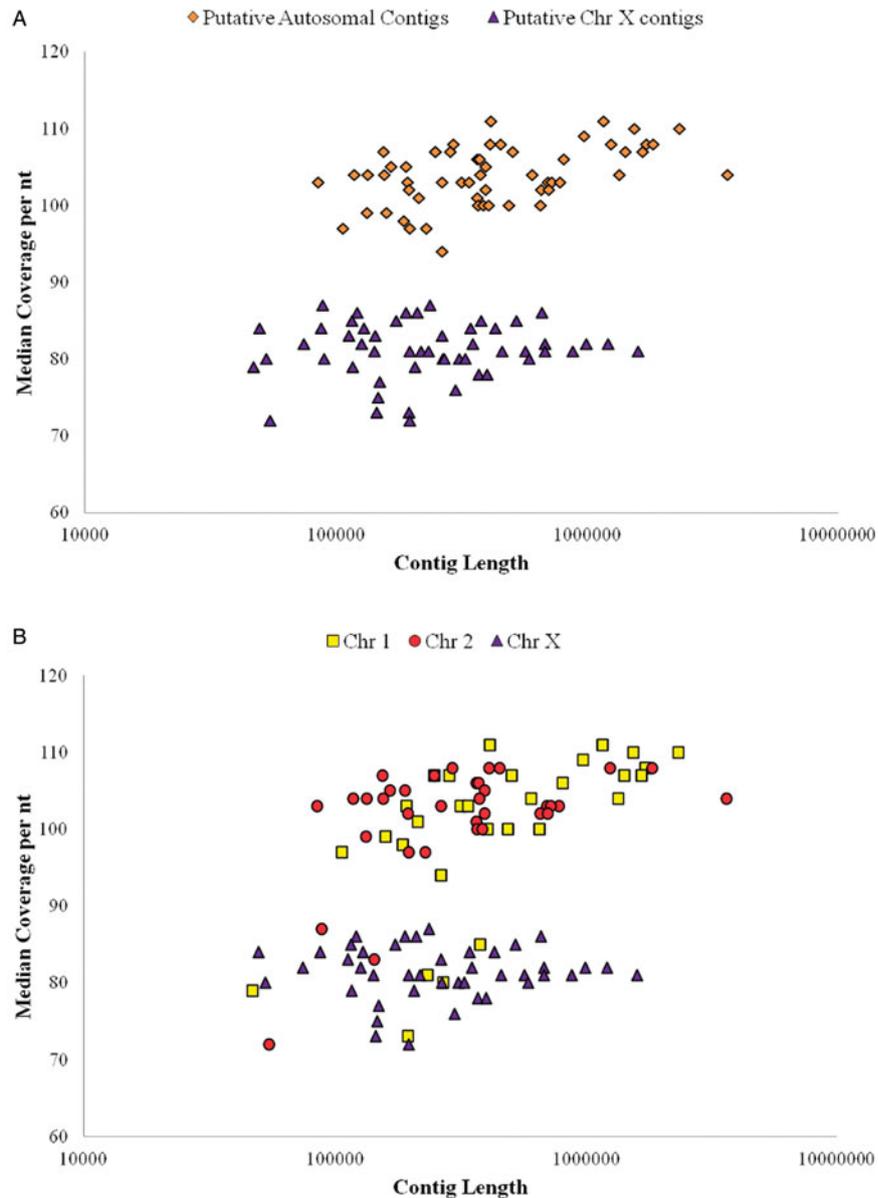
Fig. 5. Median short-read coverage of *T. murrelli* contigs supports hypothesized division of contigs into autosomal and sex chromosome groups. Trichinellids have an XO sexual determination system; if male and female genomes contributed equally to the template used in chromosome sequencing, then reads corresponding to the X chromosome should be 3/4 as frequent as those mapping to either of the two autosomes. (A) The actual distribution of median read coverage for *T. murrelli* contigs by short-read Illumina sequences closely matches this expectation based on the 109 contigs with the highest median coverage, in that read densities either cluster at ~105 reads/base (blue squares) or at ~75 reads/base (purple triangles). Those 544 contigs with median coverage <67 reads/bp (not shown) were considered outside of the normal range required for this analysis. (B) Most contigs (92·7%) assigned as putative autosomal or X chromosome sequences matched their chromosomal assignment by homology to *T. spiralis* scaffolds. However, five contigs matching *T. spiralis* chromosome 1 sequences (yellow squares) and three contigs matching chromosome 2 sequences (red circles) disagreed with homology assignments and require further study for confident placement on chromosomes.

used these to reorder the 2011 *T. spiralis* assembly, resulting in an improved 2016 *T. spiralis* assembly. Furthermore, we were able to use two lines of evidence to situate sequences of *T. murrelli* and *T. spiralis* into their two autosomes and one X chromosome.

The genomes of closely related species often retain syntenic organization. While translocations and inversions occur, chromosomal gene content is often conserved, particularly among nematodes (Guiliano *et al*. 2002; Hillier *et al*. 2007; Simakov *et al*. 2013). Here, two assemblies from species of *Trichinella* were used to improve each other. The 2011 *T. spiralis* scaffolds provided valuable insight into long-range association of contigs, derived from BAC end-sequencing. Furthermore, information from single-copy orthologues allowed scaffolds of *T. spiralis,* and then *T. murrelli,* to be assigned to chromosomes. Conversely, the long *de*
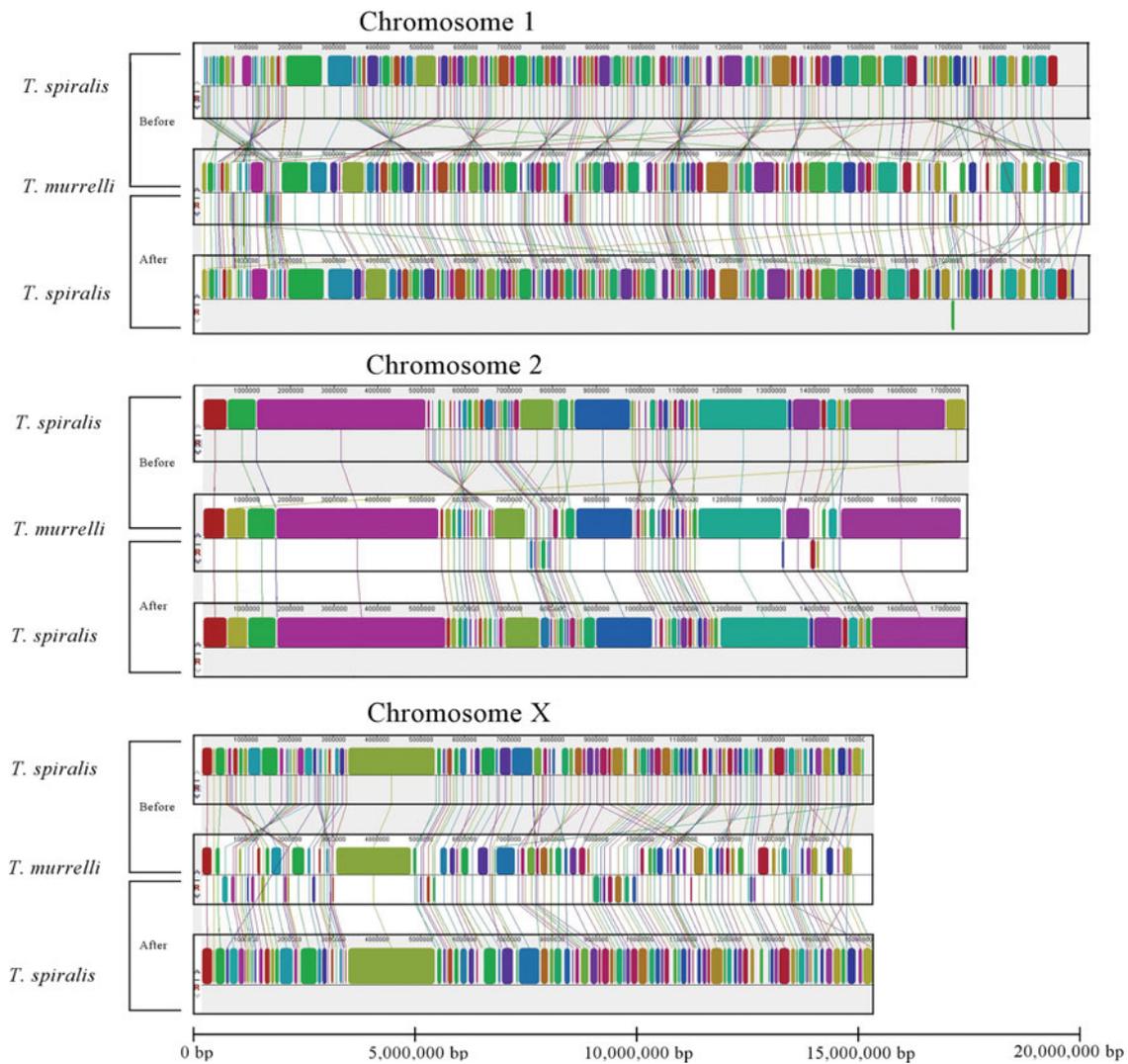
Fig. 6. Depiction of putative chromosomes before and after realignment. In the upper portion of each chromosome panel, *T. murrelli* contigs from each putative chromosome have been moved to optimally match 2011 *T. spiralis* scaffolds predicted to correspond to each chromosome. In the lower part of each chromosomal depiction, *T. spiralis* scaffolds were broken at assembly gaps into component contigs, then contigs were reassembled into putative chromosomes based on MUMmer matches to the newly established *T. murrelli* chromosomes. The decrease in crossing LCB connecting lines is an indication of better syntenic arrangements along putative chromosomes. The final chromosomal scaffolds provide our best estimate of genome sequences for both species based on positional evidence derived from long-read uninterrupted *T. murrelli* contigs and scaffolding information present in the 2011 *T. spiralis* assembly.

novo contigs comprising the *T. murrelli* assembly identified the need to reorder LCBs in *T. spiralis*. These data justified the conclusion that the conspicuous crossing pattern in macrosyntenic alignments had not resulted from rampant inversions, but rather from frequent local misassembly of contigs in the 2011 *T. spiralis* assembly, separated by sequencing gaps. We cannot explain why these blocks appeared in reverse order, but we are confident that the source of conflict was computational, not biological. Thus, the final assembly of *T. murrelli* using long reads allowed assembly limitations imposed by short reads in highly repetitive genomic regions to be overcome. While the ordering of scaffolds within chromosomes still requires refinement, the sequences resulting from this

effort provide a basis for future investigations of inherited linked genes.

Assembly of genomes is most difficult when derived from short reads and when highly repetitive elements force assembly ambiguities; this limitation was largely overcome by obtaining sequence reads far longer repeat intervals. Doing so substantially increased contig length and decreased their number. We achieved an approximately 10-fold improvement over the *T. spiralis* assembly published in 2011 as well as the 2016 short-read *T. murrelli* assembly. We placed the most confidence in contigs >100 000 bp; these exhibited more consistent read coverage and repetitive sequence composition. These longer contigs accounted for 82·2% of the entire assembly length. The longest *T. murrelli* contig spanned 116
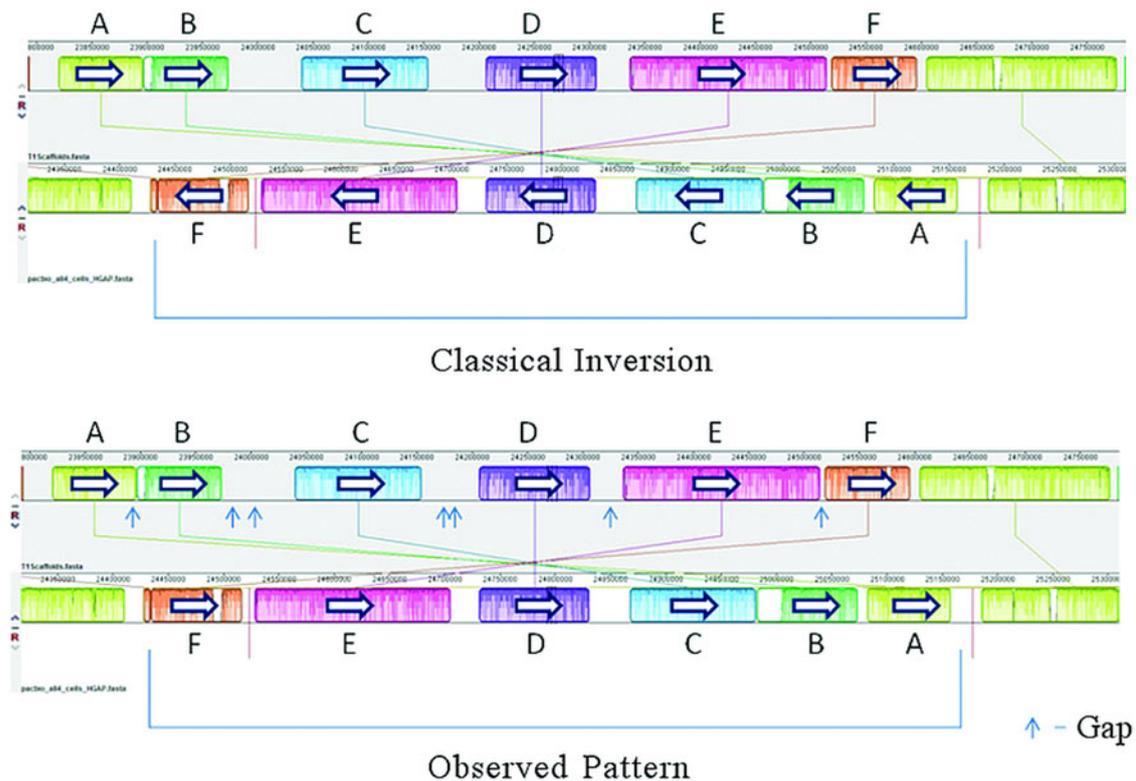
Fig. 7. Graphic showing the difference between the observed LCB alignment pattern and a classic chromosomal inversion. Each coloured block corresponds to an LCB determined by MAUVE. The direction of the arrows indicate the direction of the aligned sequences. In a classical inversion, the segment ABCDEF would be reversed (FEDCBA) and the arrows would point in the opposite direction of the reference sequence. What was observed was that the order of the LCBs was reversed (FEDCBA), but oriented as in the reference sequence, indicating that the LCBs were reordered, but not reversed. Such a rearrangement would have required at least six separate translocations of individual LCBs while retaining regional cohesion within the chromosome. This seems unlikely in closely related species.

gaps in the 2011 *T. spiralis* assembly; these gaps totalled 369 885 bp of the 3·6 Mbp total. In addition, long-read sequencing reduced indeterminate bases 80-fold (from 8 to <0·1%); the longer reads confirmed the scaffolding of some gaps, but revealed many errors in the scaffolding process used previously. By assuming synteny, we were able to leverage these longer contigs in *T. murrelli* to improve scaffolding in the assembly of *T. spiralis*.

Nevertheless, the 2011 *T. spiralis* scaffolds, achieved through hierarchical scaffolding, contain useful proximity information which we used to orient the *T. murrelli* contigs into putative chromosomes, enabling our new scaffolding hypothesis for *T. spiralis* based on alignments between *T. murrelli* contigs and ungapped *T. spiralis* contigs. While further work is required to place the contigs that did not match cleanly with chromosomal sequences, long-read sequencing substantially improved the quality and confidence of both genome assemblies.

We were able to situate our scaffolds into each of two autosomes and one sex-determining X chromosome based on two lines of evidence, single-copy orthologues as per Foth *et al*. 2014, and median depth of short-read coverage of our long-read

assembly. MUMmer was then used to determine which *T. murrelli* contigs aligned with a given *T. spiralis* putative chromosome, and allowed for informed scaffolding into new putative *T. murrelli* chromosomes. Illumina short reads were aligned to the *T. murrelli* contigs and median depth of coverage per base pair was used to test whether putative X chromosome coverage differed from autosomal coverage. Unless only females are sequenced, X chromosomes should be covered by fewer reads than autosomes. The extent of this expected deficit depends on the sex ratio (3/4 for a population comprised of equal numbers of male and female genomes); but the ratio of X:autosome would be 5:6 if, as has been reported, females generally outnumber males by nearly 2:1 (Rappaport, 1943; Gursch, 1949; Boyd and Huston, 1954; Boyd and Huston 1954, Gursch, 1949; Rappaport, 1943). The actual ratio approximated 3:4 based on median coverage, in line with expectations. While nine contigs were classified as having X chromosome coverage but linked to autosome sequence based on gene content, median short-read coverage of contigs provided a reasonable method for preliminary separation of sex chromosome contigs from autosomes.

Long-read assembly of *T. murrelli* substantially refined the estimate of repeat content in *Trichinella* genomes. Our estimate of repetitive DNA content was 12% higher than those made by both Mitreva *et al*. 2011 and Korhonen *et al*. 2016. Conflict in these estimates may derive either from the large proportion of indeterminate sequences present as assembly gaps in the 2011 *T. spiralis* assembly, or from the 14 Mbp of sequence not documented in the 2016 *T. murrelli* short-read assembly. However, because of the long sequence reads presented herein, it is less likely that our assembly would overestimate the amount of repetitive DNA in the genome when compared with assemblies derived from shorter reads.

Long-read sequences were assembled three times, based on three *a priori* assumptions of genome size (60, 65 or 71 Mbp) in order to assess the influence of these assumptions on assembly. Assuming a 60 Mbp genome resulted in an assembly of just under 60 Mbp, but when postulating either 65 or 71 Mb genomes, assemblies of long-reads resulted in 63·2 and 62·7 Mbp total length, suggesting convergence on a stable estimate in spite of differing prior assumptions. MAUVE and MUMmer alignments indicated that the alignable fractions of the *T. spiralis* and *T. murrelli* assemblies were not especially sensitive to differences in the assumed final assembly size. The smaller assembly, derived from the 60 Mbp prior, most likely resulted from excessive condensation of repetitive DNA elements. The difference between *Trichinella* genome assembly sizes presented here and one estimate of genome size made by flow cytometry (71 Mbp) (Mitreva and Jasmer, 2008) may reflect as-yet undetected repetitive DNA. For example, Phobos, which can detect degenerate repeats in the genome, estimated almost twice the number of simple repeats detected by RepeatMasker; this highlights the difficulty of correctly identifying, much less assembling, degenerate repetitive DNA. If there are similar degenerative repeats among interspersed elements, ours is probably an underestimate of the true amount of repetitive DNA. It is likely that this type of repeat remains within the assemblies presented here, despite our best efforts.

The marked improvement in the assembly of *T. spiralis* did not significantly alter our understanding of gene content. CEGMA and BUSCO analyses suggested that all assemblies were nearly complete. Initial predictions by the Augustus and Maker gene prediction programs indicated between 10 400 and 13 500 genes are present in the long-read *T. murrelli* genome assembly, as compared with ~15 000 genes predicted in other studies. The presence of transposons or alternative transcripts did not account for this difference. Our gene prediction parameters may be conservative, as we found nearly 4000 fewer 'hypothetical' genes than were found in the 2011 *T. spiralis* assembly (Mitreva *et al*. 2011) and the 2016 *T. murrelli* short-read assembly (Korhonen *et al*. 2016). These comparisons require further analysis accompanied by RNA sequencing.

Improved organization of contigs is important for understanding genome evolution. The 2011 *T. spiralis* genome assembly implies that 17 genome rearrangements occurred for every Mb of sequence since its divergence from *T. murrelli*. Assuming an estimated 7–10 MY of divergence (Zarlenga *et al*. 2006; Korhonen *et al*. 2016), two chromosome breakages per inversion, and three breakages per translocation, 1·7–2·5 chromosomal breakages/Mbp/MY would have occurred. However, our revised assembly of *T. spiralis* reduces the estimate of genome breakages 10-fold, to 0·15–0·21 breakages/Mbp/MY; a rate more in line with previous estimates for the free-living nematodes *Caenorhabditis elegans* and *Caenorhabditis briggsae* (0·4–1·0/Mbp/MY) (Coghlan and Wolfe, 2002). Furthermore, the improved organization of LCBs within the genome enables better estimates of synteny between these two species, because the syntenic correlation between single-copy orthologues (0·118 as reported by Korhonen *et al*. 2016) was dependent on the 2011 arrangement of scaffolds. However, a revised estimate cannot be made until full annotation of this assembly is complete. Ongoing RNA sequencing will illuminate this issue going forward.

Concluding remarks

Trichinella murrelli and *T. spiralis* maintain remarkable synteny across all three chromosomes. This has enabled us to improve the assembly of each genome. Earlier hierarchical scaffolding of short reads, combined with bacterial artificial chromosomes, provided useful information for linking large numbers of contigs in the 2011 *T. spiralis* assembly. Here, long-read sequencing using PacBio technology substantially improved that scaffolding by spanning repeat regions and allowing more precise ordering of contigs. Together, these refined assemblies of *T. murrelli* and *T. spiralis* provide a resource for future investigations of genome conservation within the genus *Trichinella*, and among clade I nematodes, more generally. RNAseq studies of the gene content of these two genomes will improve comparisons between the two species and provide insight into the gene families that define the genus *Trichinella* and those genes that are dispensable. Furthermore, defined sets of genes may help to indicate why *T. murrelli* infects swine poorly and causes minimal disease whereas *T. spiralis* as adapted well to swine hosts and therefore provides significant danger of zoonotic disease.

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