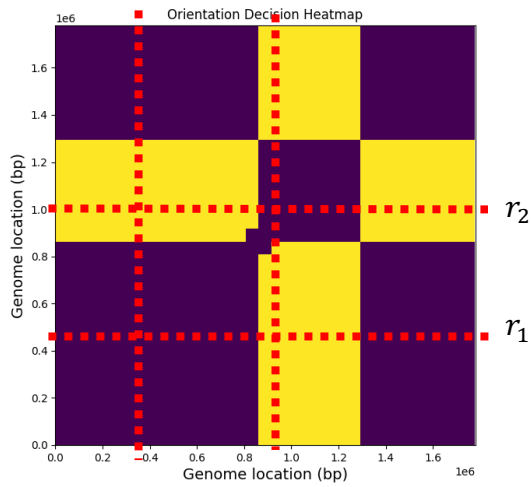
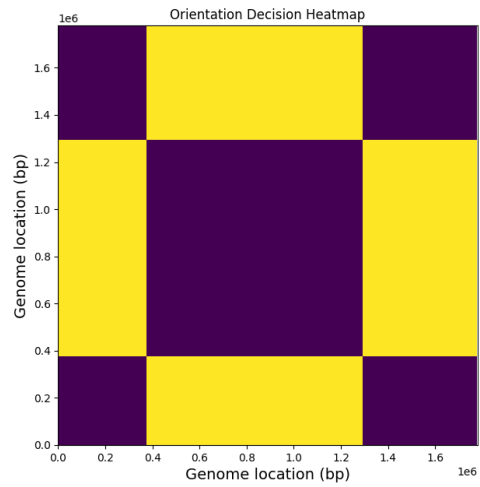


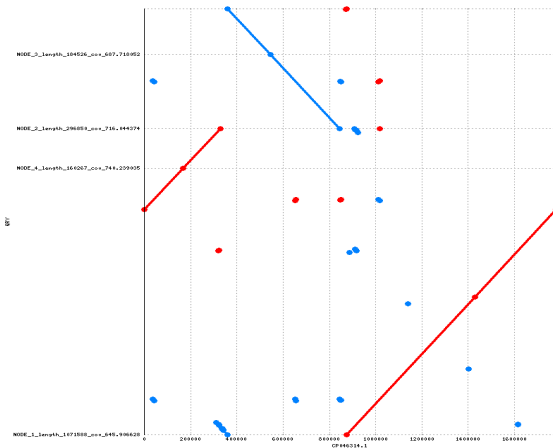
a Original Genome ($bal = 0.45$)



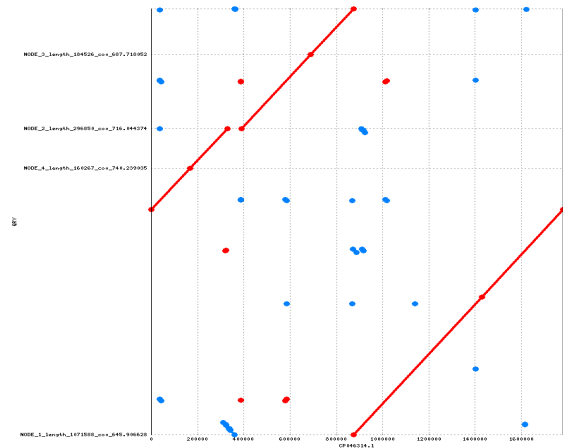
b Corrected genome ($bal = 0.97$)



c Original Genome vs. SPAdes Contigs

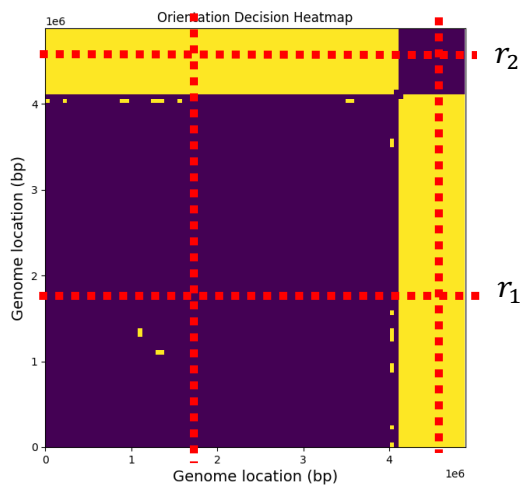


d Corrected Genome vs. SPAdes Contigs

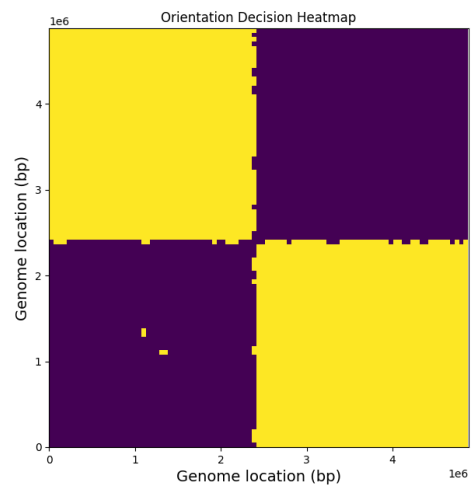


Supplementary Figure S1. Correction of Inverted Misassembly of *G. morbillorum* (strain: FDAARGOS_741; taxID: 29391; assembly accession: GCA_009730315.1; BioSample: SAMN11056456). **(a)** Orientation heatmap of original genome from GenBank is highly unbalanced. Red dashed lines represent the locations of the repeat (r_1, r_2) used to correct the misassembly. **(b)** Heatmap of corrected genome is now balanced. **(c)** Dot plot of original genome against largest four contigs from the re-assembly of the hybrid read data using SPAdes assembler. A large inversion is manually positioned to illustrate the location of the repeat. **(d)** Dot plot of corrected genome against SPAdes re-assembly. The large inversion from (c) is now un-inverted, indicating that the ordering of the contigs chosen in (c,d) is indeed correct.

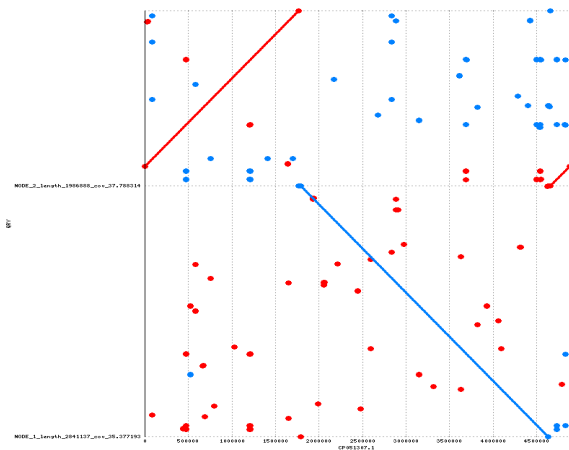
a Original Genome ($bal = 0.34$)



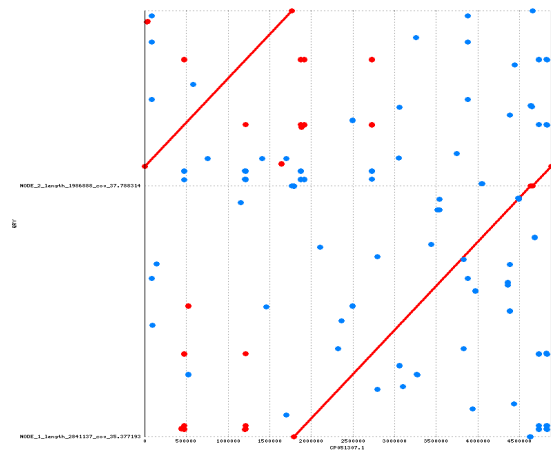
b Corrected genome ($bal = 0.97$)



c Original Genome vs. SPAdes Contigs

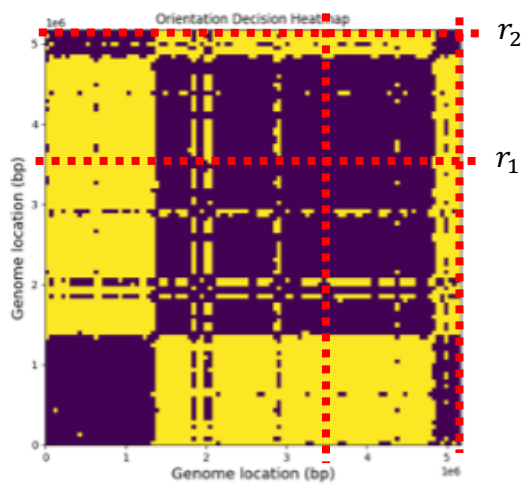


d Corrected Genome vs. SPAdes Contigs

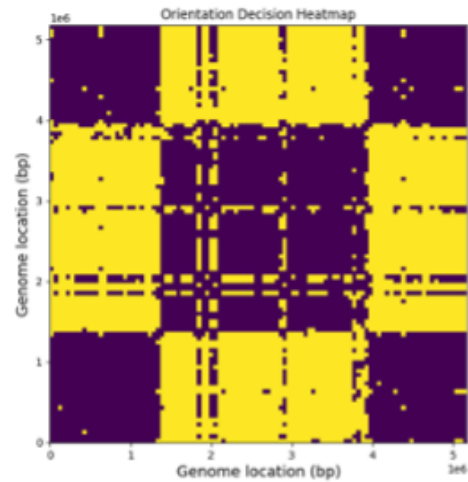


Supplementary Figure S2. Correction of Inverted Misassembly of *S. enterica* (strain: CVM 35189; taxID: 28901; assembly accession: **GCA_016451985.1**; BioSample: SAMN14504941). **(a)** Orientation heatmap of original genome from GenBank is highly unbalanced. Red dashed lines represent the locations of the repeat (r_1, r_2) used to correct the misassembly. **(b)** Heatmap of corrected genome is now balanced. **(c)** Dot plot of original genome against largest four contigs from the re-assembly of the hybrid read data using SPAdes assembler. A large inversion is manually positioned to illustrate the location of the repeat. **(d)** Dot plot of corrected genome against SPAdes re-assembly. The large inversion from (c) is now un-inverted, indicating that the ordering of the contigs chosen in (c,d) is indeed correct.

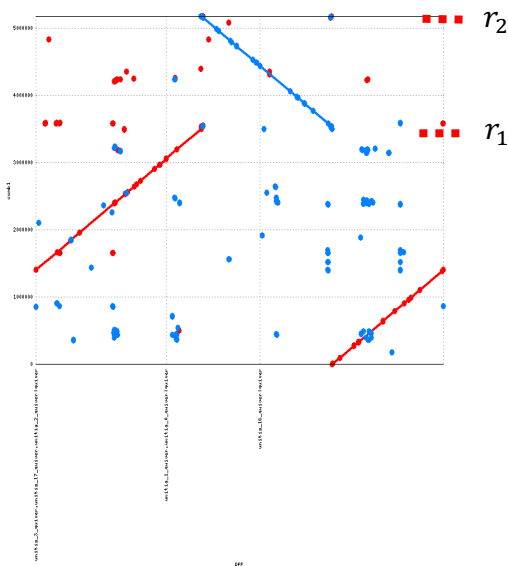
a Traversal 1 ($bal = 0.68$)



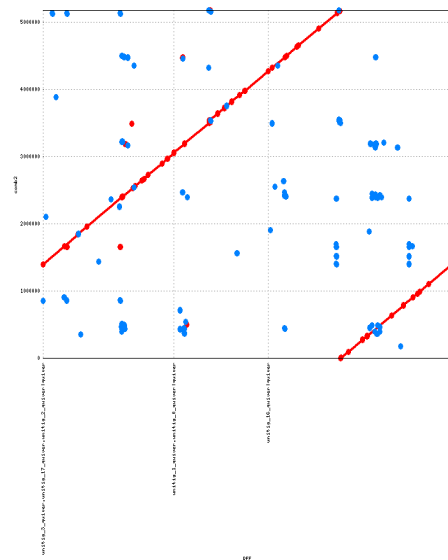
b Traversal 2 ($bal = 0.99$)



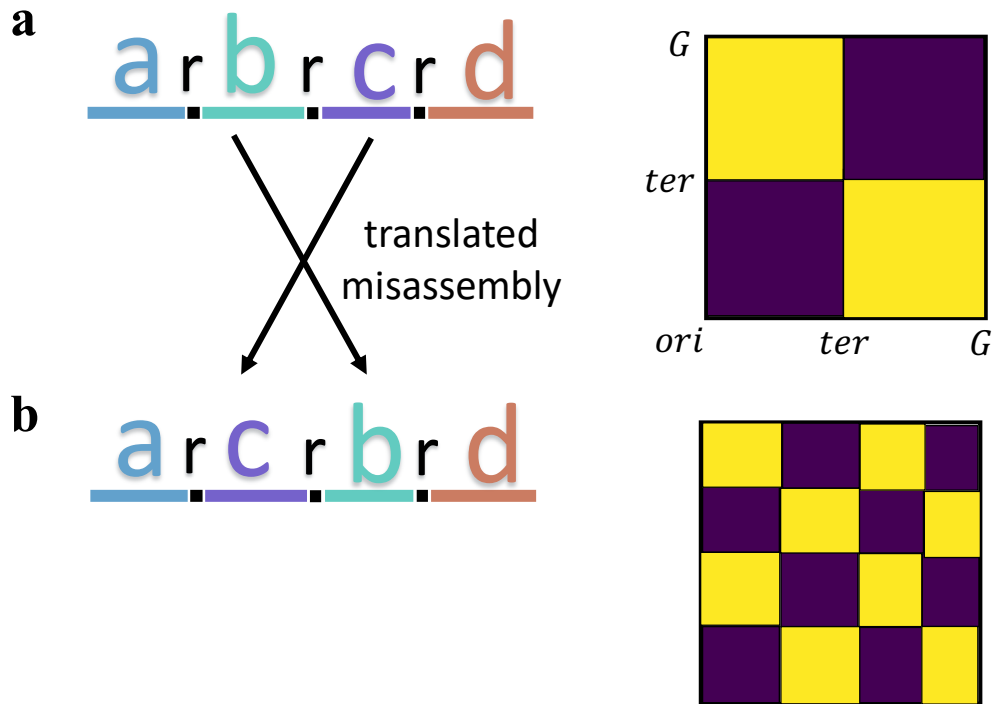
c NCTC Genome vs. Traversal 1



d NCTC Genome vs. Traversal 2



Supplementary Figure S3. Assembly analysis of *E. coli* (strain: NCTC9006; taxID: 562; BioSample: SAMEA3376915). The long read data from NCTC 3000 was assembled using the HINGE assembler, resulting in an assembly graph with two possible traversals. **(a)** Orientation heatmap of Traversal 1 is unbalanced. Red dashed lines represent the locations of the repeat (r_1, r_2) which causes the incomplete assembly. **(b)** Heatmap of Traversal 2 is balanced, suggesting that it is the correct assembly of the genome **(c)** Dot plot of the NCTC genome (containing three contigs) against Traversal 1. A large inversion is present exactly between (r_1, r_2). **(d)** Dot plot of NCTC genome against Traversal 2. The large inversion from (c) is now un-inverted, indicating that the ordering of the contigs chosen in (c,d) is indeed correct.



Supplementary Figure S4. (a) Example of a genome with a triple repeat on the forward strand, along with the corresponding orientation heatmap. **(b)** Depiction of the same genome if an erroneous translation occurred during assembly by switching the locations of segments *b* and *c*. The corresponding orientation heatmap has three distinct transitions in orientation.

assembly_accesion	read_type	sequencer(s)	assembler	misassembly_supported	notes
GCA_005886035.1	hybrid	Illumina HiSeq 4000, PacBio Sequel	SPAdes	yes	Re-assembly agrees with corrected genome
GCA_008065435.1	long	PacBio RS II	HINGE	yes	Two traversals corresponding to original and corrected genome
GCA_009730315.1	hybrid	Illumina HiSeq 4000, PacBio RS	SPAdes	yes	Two orderings of contigs corresponding to original and corrected genome
GCA_016451985.1	hybrid	Illumina MiSeq, PacBio Sequel	SPAdes	yes	Two ordering of contigs corresponding to original and corrected genome
GCA_002012025.1	short	Illumina MiSeq	SPAdes	n/a	Fragmented re-assembly. Possible read data omitted.
GCA_016452025.1	long	PacBio Sequel	HINGE	n/a	Fragmented re-assembly. Possible read data omitted.
GCA_014623225.1	long	PacBio RS II	n/a	n/a	Manually discarded due to poor heatmap quality
GCA_900327275.1	long	PacBio RS	n/a	n/a	Manually discarded due to poor heatmap quality
GCA_003112145.1	short	Illumina MiSeq	n/a	n/a	Some read data omitted. Re-assembly not possible
GCA_003339775.1	long	PacBio RS	n/a	n/a	Some read data omitted. Re-assembly not possible
GCA_000198515.1	none				
GCA_000487155.2	none				
GCA_001562215.1	none				
GCA_001722005.2	none				
GCA_001723625.1	none				

GCA_001936395.1	none				
GCA_001938665.1	none				
GCA_002005165.1	none				
GCA_002441975.1	none				
GCA_004291075.1	none				
GCA_009950475.1	none				
GCA_009951245.1	none				
GCA_011045215.1	none				
GCA_011801475.1	none				
GCA_012934815.1	none				
GCA_013085185.1	none				
GCA_013305705.1	none				
GCA_014168635.1	none				
GCA_014489455.1	none				
GCA_014731795.1	none				
GCA_900324235.1	none				
GCA_900327275.1	none				

Supplementary Table T1. List of misassemblies detected from 5,000 randomly chosen GenBank genomes.