

Analizis of E.coli strains for outbreak investigation via identification pathogenic genes

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We assemble the genome of deadly E. coli X strain. We analyze strain and find enzymes which lead to pathogenicity.

1 Introduction

2 Methods

To analyze the E. coli X strains, We use the dataset from the TY2482 sample [1]. To estimate genome size, I use Jellyfish [3]. For estimation the genome size, the following formulas is used:

$$N = \frac{M * L}{L - K + 1}, Genome_size = \frac{N}{T},$$

where N — Depth of coverage, M — k-mer peak, K — k-mer-size, L — average read length, T — Total bases).

For assembling the genome the SPAdes tool is used [4]. SPAdes uses information about the distances between reads within read-pairs to combine contigs into ordered collections of adjacent contigs called scaffolds.

We use Prokka for gene prediction and annotation [7].

3 Results

3.1 Data description.

The analyzed data were obtained at Beijing Genome Institute from the isolate of the sample TY2482 using Illumina 1.9. There were three libraries in total - one paired-end (insert size 670 bp) and two mate-pair libraries (insert sizes 2 Kbp and 6 Kbp). The characteristics of each data set are given in Table 1.

ID	Library type	Insert size	Number of reads	Average read length
SRR292678	pair-end	470	5499346	90
SRR292862	mate-pair	2000	5102041	49
SRR292770	mate-pair	6000	5102041	49

Table 1: The sequencing data.

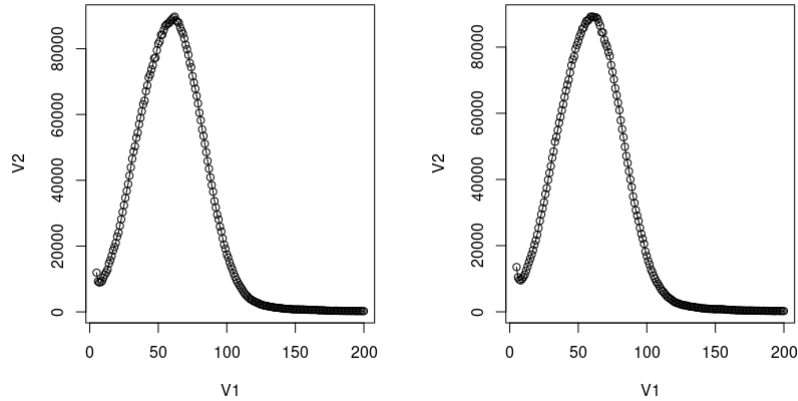


Figure 1: The k-mer distribution in the forward and reverse data among region between 8 and 200

3.2 k-mer profiling using Jellyfish.

The information related to the quality of the resulting assembly after SPAdes usage is available in lab journal. Using Jellyfish we determined k-mer peak bin. For paired-end data the k-mer peak was 125. Depth of coverage was calculated according to formulas described in Methods. To calculate T(total bases number), the number of reads(5499436) was multiplied to average read length(90). These numbers were obtained from FastQC reports(see GitHub repository, as also R script to k-mer counting). As a result, estimated genome size is 5.3 Gb and depth coverage is 188. After reads correction using SPAdes, k-mer profile hasn't changed. There is no significant improvement in k-mer distribution, except first bins. Genome size and depth coverage showed insignificant changes.

3.3 Checking assembly quality using QUAST

In this report, only basic metrics(such as N50,L50 and number of contigs with length more/equal than 500bp) will be reported. You can find the results in Table 2. To sum up, addition of mate-pair libraries improved N50 metric and contigs number, as expected. This fact could be easily explained because addition of mate-pair sequencing

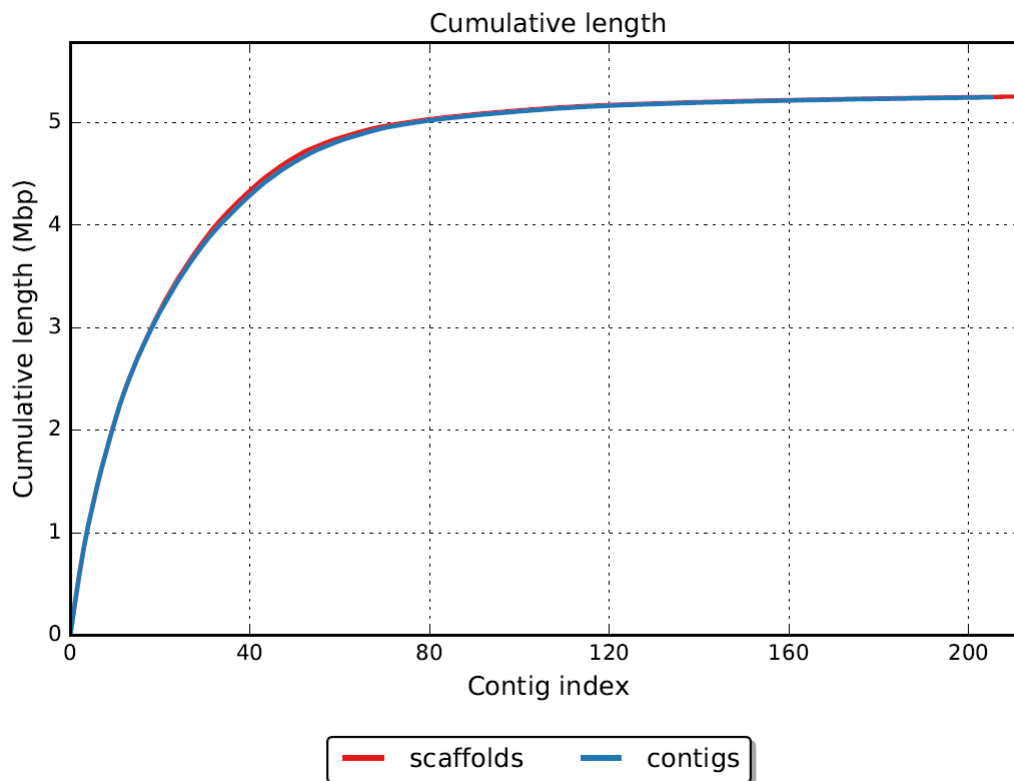


Figure 2: Assessment of the quality of the paired processed data after using SPAdes

	PE	PE+MP
N50	105346	155269
Number of contigs ≥ 50 bp	205	169
Total number of contigs	519	544

Table 2: Quality metrics of SPAdes assemblies.

libraries can negate problems in de novo genome assemblies. Mate pairs cover greater distances because of the increased insert size. As a result, they can cover genome regions that contain repeats. And paired-end reads could fill the gaps between mate-pair reads. This approach (combinig mate-pair/paired-end reads) allows us to obtain better assemblies than just using one library type.

3.4 Genome annotation and 16S rRNA study

Next step was genome annotaion using Prokka. Results are: -5064 CDS -2923 genes -80 tRNAs were found. To find the closest reference genome, 16S rRNA prediction was performed. 16S rRNA is widely used for phylogenetic studies because 16S rRNA gene is highly conserved between different species of bacteria and archaea. 20 rRNA

sequences were found in total, and 8 copies of 16S rRNA were found. Their length is 1536 bp for 7 copies and 406 bp for 1 copy. Using BLAST we successfully identified closest relative strain for our E.coli X strain. This is E.coli strain 55989(*NC_011748.1*). BLAST reporting 100% identity and cover. This newfound strain will be later used as a reference.

3.5 Genome-wide comparison using MSA

To find and see changes between X strain and 55989 strain, multiple sequence alignment using Mauve tool was performed. Using the fact that Shiga toxins can cause HUS, we searched results using Sequence Navigator. As a result, 2 genes (stxA and stxB) were found. These genes encoding A and B subunit of Shiga-like toxin which is responsible for 2011 HUS outbreak. To find how these genes were acquired, regions surrounding stx genes were investigated using Mauve GUI. Several phage-related genes and also mobile element-related genes were found. We can assume that E.coli X contain not only pAA plasmid which has enteroaggregative genes, but also was infected by bacteriophage. We suggest that this phage was a deliverer and source of these toxins.

3.6 Antimicrobial drug resistance analysis

To analyse our reference strain and X strain, ResFinder tool was used. After analyzing both reference and target strains with ResFinder, it was found that the former is resistant to tetracycline due to production of tetB protein which is responsible for the drug efflux and acts as a metal-tetracycline/H⁺ antiporter in the cell. We can definitely say that X strain is almost a superbug because it has multiple resistance—it was reported to be resistant to cefepime, ampicillin, cefotaxime, sulfamethoxazole, trimethoprim and ceftazidime in addition to tetracycline. Most of its resistance was caused by beta-lactamase production. To track the source of this shocking fact Mauve tool was again used, as also the same approach (scanning regions nearby resistance genes). At least 2 enzymes were found: beta-lactamase CTX-M1 and beta-lactamase TEM. And these enzymes were flanked by mobile element-related sequences and/or phage genes responsible for phage replication. So, phage infection could be responsible not only for stxA/B genes, but also for AMR genes. In addition to that, in case of CTX-M1, there was a large insertion that was labeled as Inc11 plasmid transfer proteins. # What a day, huh? Genome of X strain is a mish-mash but thus bacteria is a real danger

4 Discussion

X strain of E.coli is a clear example of horizontal gene transfer. First we found that Shiga-like toxins genes were surrounded by phage proteins. Second, in the case of beta-lactamase CTX-M-1, there is a large insertion containing genes from the Inc11 plasmid. As expected, both groups were absent in the reference genome. However, there is a strong possibility that the mechanisms of obtaining these genes differ. Let's talk about these mechanisms.

In the case of Shiga-like toxins, we could blame the phage integration mechanism. Phage life cycles dictate their role in bacterial and archaeal biology. Three life cycles of phages have been reported: lytic, lysogenic and chronic phages. In general, once a virulent phage (a phage that follows the lytic cycle) has attached to its host cell, the phage's nucleic acid enters the cell and causes the bacterium to produce hundreds of phage copies. This results in the lysis of the cell and the newly formed phages are released into the surrounding environment. Temperate phages (phages that follow the lysogenic cycle) may follow one of two scenarios. The first scenario results in the lysis of the host cell and release of newly formed phages, similar to the lytic life cycle outlined above. In the second scenario, phage DNA may be integrated into the bacterial chromosome. The integrated DNA (prophage) is non-infectious and replicates as part of the bacterial chromosome. Incorporation of the phage DNA into the bacterial chromosome can be beneficial for the evolution of the bacteria as useful genes may be transferred to the bacteria. These prophage-mediated changes have been termed lysogenic conversion. In this state of symbiosis, both phage and the host cell experience an increased level of fitness. Under UV light or certain chemical treatments, the prophage is excised and causes the bacteria to produce phage particles. Figure 2 depicts the lytic and lysogenic lifecycle. The third life cycle is the chronic lifecycle which occurs in archaeal viruses and some filamentous and temperate phages. These viruses do not cause cell disruption or cell death, but instead the newly formed virions are continuously released from the cell. The infected host cells are capable of growing but at a much slower rate. You can read more here [8].

In case of antimicrobial resistance, the situation is a little more complicated. We can assume that E. coli X obtained the TEM beta-lactamase with the help of bacteriophages, as in the case of Shiga-like toxins. But the situation is different in the case of beta-lactamase CTX-M-1. Mauve exploration of neighbouring regions showed that the large insertion containing CTX-M-1 came from the Inc11 plasmid. Several studies show that this plasmid indeed contains such enzyme and can be transferred horizontally by conjugation process [6]. But also this insertion was also surrounded by phage proteins (e.g., phage minor tail protein, phage endopeptidase), according to the

annotation. We can suggest with a strong confidence that E. coli X obtained this gene via Inc11 plasmid since the insertion size is rather big and there are only a few phage genes around. Insertion size is too big to be carried by the phage, but from where these phage genes occur? The exact way of transferring this gene to the strain of interest remains unknown. However, the mechanism of antimicrobial resistance in the case of beta-lactamases is clear. Beta-lactamases are enzymes that are responsible for the hydrolysis of the β -lactam ring of the antibiotic, making it ineffective. Most part of antibiotics that E. coli X is "immune" to compared to the reference strain (cefepime, cefotaxime, ampicillin and ceftazidime) belong to the class of β -lactam antibiotics. But X strain also carries genes of sulfonamide (sul1, sul2) and trimethoprim (dfrA7) resistance which are capable of maintaining their function in presence of these antibiotics, thus making it superbug. HUS syndrome, multiple AMR—I'm concerned.

The simplest way to overcome such resistance is to use a combination of β -lactam antibiotics with beta-lactamase inhibitors (e.g. clavulonic acid). You can check other methods in 1st report of our group in our GitHub repository. Thank you for your attention!

References

- [1] Datasets: (forward and reverse)
SRR292678:
https://d28rh4a8wq0iu5.cloudfront.net/bioinfo/SRR292678sub_S1_L001_R1_001.fastq.gz
https://d28rh4a8wq0iu5.cloudfront.net/bioinfo/SRR292678sub_S1_L001_R2_001.fastq.gz
SRR292862:
https://d28rh4a8wq0iu5.cloudfront.net/bioinfo/SRR292862_S2_L001_R1_001.fastq.gz
https://d28rh4a8wq0iu5.cloudfront.net/bioinfo/SRR292862_S2_L001_R2_001.fastq.gz
SRR292770:
https://d28rh4a8wq0iu5.cloudfront.net/bioinfo/SRR292770_S1_L001_R1_001.fastq.gz
https://d28rh4a8wq0iu5.cloudfront.net/bioinfo/SRR292770_S1_L001_R2_001.fastq.gz
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- [3] Guillaume Marcais and Carl Kingsford, A fast, lock-free approach for efficient parallel counting of occurrences of k-mers. Bioinformatics (2011) 27(6): 764-770 (first published online January 7, 2011) doi:10.1093/bioinformatics/btr011
- [4] SPAdes: <http://cab.spbu.ru/software/spades/>
- [5] QUAST: <http://quast.bioinf.spbau.ru/>

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