<u>Assembly of Small Microbial</u> <u>Genome</u>

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De novo assembly

- 1. Ideal approach
- 2. Recreate original genome sequence through overlapping sequenced reads
- 3. More accurate
- 4. Not biased towards reference genome unlike mapping-based approach
- 5. Expensive, need a lot of reads
- 6. Time and memory consuming

Paired-end sequencing

- 1. Sequencer starts reading from one end till specified read length
- 2. Reads from the opposite direction again
- 3. improves ability to identify relative read position in genome
- 4. much more effective than single-end seq. in resolving gene insertions, deletions etc.
- 5. improves assembly of repetitive regions

Tools

- sra-tools (fastq-dump)
- 2. fastqc
- 3. trimmomatic
- 4. megahit or other assemblers
- 5. quast

Practical

Let's assemble sequenced SARS-CoV-2 genome

Thank You