

GENOME ASSEMBLY WEEK 1

DATA WRANGLING

OVERALL SCHEDULE

- ▶ 1. 4/28: Data wrangling
- ▶ 2. 5/5: Resource allocation and configuration files
- ▶ 3. 5/12: Monitoring assembly progress and troubleshooting errors
- ▶ 4. 5/19: Generating assembly metrics
- ▶ 5. 5/26: Comparing multiple assemblies
- ▶ 6. 6/2: Visualization and next steps

WORKSHOP EXPECTATIONS

- ▶ At the end of the 6 weeks, I hope you:
 - ▶ Feel comfortable with the basic steps of the assembly process and able to perform them on your own.
 - ▶ Are empowered to make decisions about what kind of assembly software you might need for a given project.
 - ▶ Begin to appreciate that this is more of an art than a science!

OUTCOMES

- ▶ You will become versed in three* major assemblers using a mixed bag of public data.
- ▶ We will perform all the steps from raw data to assembly comparison and visualization.

*If everyone becomes super proficient, we can advance to bigger genomes and additional assemblers.

TEST DATASETS

- ▶ NIST (National Institute of Standards and Technology)
 - ▶ *Salmonella enterica* RM 8375
 - ▶ MiSeq
 - ▶ PacBio

WHICH ASSEMBLER DO I NEED?

- ▶ Depends on:
 - ▶ Data (sequencing platform, libraries)
 - ▶ Genome size
 - ▶ Compute resources at your disposal

ASSEMBLERS

- ▶ SPAdes (small genomes, any data)
- ▶ DISCOVAR (Illumina 2X250bp reads only)
- ▶ MaSuRCA (hybrid: pretty much any data, any genome size)

ASSEMBLIES WE WILL RUN

- ▶ Salmonella:
 - ▶ DISCOVAR (Illumina only)
 - ▶ SPAdes (Illumina only)
 - ▶ SPAdes (Illumina and PacBio)
 - ▶ MaSuRCA (Illumina and PacBio)

LET'S MOVE TO THE TUTORIAL

- ▶ https://github.com/SmithsonianWorkshops/GenomeAssembly/blob/master/week1-data_wrangling.md