## We can now align the viral reads to a known reference sequence.

Sequence of the reference strain (already known).

Quality Viral Reads, from the previous step.

• Use the "bwa" program to map the viral reads to the known reference assembly.

Reference Start with quality reads and a trimmed trimmed genome, reference genome. R1 pair R2 pair indexed for Use bwa mem to align the reads ed.fastq ed.fastq use with bwa to the reference. Save the output in a .sam file, which links the read to a Mapping Name of Name of files to location in the reference algorithm reference. process genome where the read aligns. bwa mem ebola\_ref trimmedR1\_paired.fastq trimmedR2\_paired.fastq \ > Prelim\_alignment.sam Save output in a .sam file. Prelim alig Continue Reference nment.sam **Based Assembly**