

1. Why is whole genome sequencing combined with bioinformatic tools seen as revolutionizing diagnostics and surveillance?

- ☐ Whole genome sequencing combined with bioinformatic tools can only be used by state of the art high profile laboratories
- ☐ Whole genome sequencing combined with bioinformatic tools can only be used for bacteria
- ☐ Whole genome sequencing combined with bioinformatic tools is only for research
- ☒ Whole genome sequencing combined with bioinformatic tools can be used for all organisms

2. What is one of the advances with whole genome sequencing combined with bioinformatic tools?

- ☐ Overall, the time spent from sample to results ( species ID, typing, virulence and relatedness tests) increases and it is less expensive than conventional methodologies combined
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3. What was the aim for Center for Genomic Epidemiology (CGE) project

- ☐ To develop a bioinformatics tool to detect only the bacterial species, virulence properties, antimicrobial resistance genes and relatedness to other strains
- ☐ To develop a commercial available bioinformatics tool for diagnostic
- ☒ To provide a foundation for web based solutions that are easy to use
- ☐ To develop a bioinformatics tool for research

4. Does the Center for Genomic Epidemiology (CGE) have a tool that combines several tools?

- ☐ No, this was not part of the strategy, all tools are single stand-alone tools
- ☒ Yes, It is called the batch upload tool
- ☐ Yes, but it is only used by DTU
- ☐ No, but it is in a development phase

5. Why the urgent need to use whole genome sequencing ?

- ☐ To avoid that developing countries embrace the technology
- ☐ To agree on using only one brand of sequencing platform, globally
- ☒ To initiate and improve standardization and quality control
- ☐ To enhance the resolution beyond the genome level

6. Which sequencing technology produces long read sequences ?

- ☐ Illumina MiSeq
- ☐ Illumina HiSeq
- ☐ Ion Torrent
- ☒ Oxford Nanopore

7. In library preparation for NGS, If you sequence more than one isolate/sample. What part in adapter helps you to distinguish the isolates/samples ?

- ☐ Sequencing primer
- ☐ Amplification primer
- ☒ Barcode
- ☐ None of above

8. What is the format used to store raw read sequences (NGS data) ?

- ☐ FASTA
- ☐ GENBANK
- ☒ FASTQ
- ☐ All of above

9. What is the purpose for 'Trimming' of raw reads ?

- ☒ To get rid of low quality sequences or reads
- ☐ To assemble reads to contigs
- ☐ To transform from FASTQ to FASTA format
- ☐ All of above

10. What is the purpose of mapping reads to a reference genome to identify variance ?

- ☐ To identify core gene
- ☐ To assemble reads to contigs
- ☐ To search for genes
- ☒ To identify SNPs

11. What is the parameter for checking genome quality ?

- ☐ N50
- ☐ Number of contigs
- ☐ Total size of genomes
- ☒ All of above

12. N50 is often used as a quality measure for assemblies. What does it describe?

- ☐ 50% of the total assembly size
- ☐ The total assembly size
- ☐ The size of the longest contig
- ☒ The median of the contig sizes in the assembly

13. What does it mean "to do de-novo assembly"?

- ☒ The assembly is created using only the raw read data.
- ☐ The assembly is done using Velvet.
- ☐ The assembly is partly based upon a reference sequence.
- ☐ The assembly is based upon a reference sequence.

14. When using the CGE Assembler with Illumina data, what assembler will do the assembly?

- ☐ Mira
- ☐ Newbler
- ☐ A combination of Velvet and Newbler
- ☒ Velvet

15. When using the CGE Assembler with Ion Torrent sequencing data, what assembler will do the assembly?

- ☐ Velvet
- ☒ Newbler
- ☐ A combination of Velvet and Newbler
- ☐ Mira