Bioinformatics Analysis of Whole Genome Sequencing Workshop (Mycobacterium tuberculosis)

Organized by

Faculty of Medicine and Health Sciences



in collaboration with



Saw Swee Hock School of Public Health

• Overview of sequence data processing for mapping reads to bacteria reference genome to identify genetic variants

such as single nucleotide polymorphisms (SNPs)

Date: 8th - 9th October 2018

Time: **9.00 am to 5.00 pm**

Venue: Bioinformatic Lab, Biotechnology Research Institute, Universiti Malaysia Sabah

Seat Limit: 30 participants

Registration fee:

RM 100 (UMS student or staff)

RM150 (Non-UMS)

For more information: **Dr. Chin Kai Ling**bioinformaticsfpsk@gmail.com

For registration:

https://docs.google.com/forms/d/e/1FAIpQ LSfIOSpvufo9aNwBmiD-2mdWnG9DuOICoKiworvGN5uZwJm-Ew/viewform?usp=sf_link

• Use of software programs to infer resistance profile of

bacteria isolates sequenced, and epidemiological clustering

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	8 th October 2018 (Day 1)	9th October 2018 (Day 2)
Session 1 0900 – 1030	Opening Ceremony	Mapping and Variant(s) Detection (Lab Practical)
1030	Introduction to whole-genome sequencing for infectious disease surveillance	• Use of command line software programs to map sequence reads to a reference and identifying genetic variants.
	• Review of sequencing technologies and its applications especially for public health microbiology and surveillance	
Break		
Session 2 1100 – 1230	Sequence data processing for quality control and species identification	Identification and annotation of SNPs for prediction of bacteria drug resistance
	• Overview of sequence data processing for quality control	• Overview of genetic markers associated with bacteria drug resistance
Lunch		
Session 3 1330 – 1500	Sequence data processing for quality control and species identification (Lab Practical)	SNP Phylogenetic tree for inference of bacteria epidemiological clustering
	 Use of command line software programs to assess quality of data generated, removal of poor quality bases/sequences, and rapid identification of species sequenced. 	• Overview of SNP phylogenetic tree for inference of bacteria epidemiological clustering
Break		
Session 4 1530 – 1700	Mapping and Variant(s) Detection	Annotation/Interpretation of SNPs and SNP phylogenetic tree
1.00	• Overview of sequence data processing for manning reads	HCC .