

Feedback — Module 4 Quiz ****Please Note: No Grace Period****

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Thank you. Your submission for this quiz was received.

You submitted this quiz on **Thu 13 Aug 2015 11:02 PM PDT**. You got a score of **9.00** out of **9.00**.

Question 1

How many lines make up each record contained in a FASTQ file?

Your Answer	Score	Explanation
<input type="radio"/> Two		
<input type="radio"/> Three		
<input type="radio"/> One		
<input checked="" type="radio"/> Four	✓ 1.00	
Total	1.00 / 1.00	

Question 2

Which of the following are components of the FASTQ format?

Your Answer	Score	Explanation
<input type="radio"/> Sequence		
<input checked="" type="radio"/> All of these options	✓ 1.00	
<input type="radio"/> Sequence identifier		

☐ Quality score

Total

1.00 / 1.00

Question 3

Which of the following is NOT a type of encoding of sequence quality scores in a FASTQ file?

Your Answer	Score	Explanation
<input type="radio"/> Sanger		
<input type="radio"/> Solexa		
<input checked="" type="radio"/> Ion torrent	✓ 1.00	
<input type="radio"/> Illumina 1.3+		
Total	1.00 / 1.00	

Question 4

Which of the following operations of quality control can be performed with Galaxy's NGS: QC and manipulation?

Your Answer	Score	Explanation
<input type="radio"/> FastQC: Read Quality reports		
<input checked="" type="radio"/> All of these options	✓ 1.00	
<input type="radio"/> Trim Sequences		
<input type="radio"/> FASTA to FASTQ, and FASTQ to FASTA format conversion, Manipulate FASTQ		
Total	1.00 / 1.00	

Question 5

Which of the following is NOT part of the FastQC report?

Your Answer	Score	Explanation
<input checked="" type="radio"/> Per sequence base quality	✓ 1.00	
<input type="radio"/> Sequence Length distribution		
<input type="radio"/> Per sequence quality score		
<input type="radio"/> Per base sequence quality		
Total	1.00 / 1.00	

Question 6

When do we filter and trim reads? Choose the correct option.

Your Answer	Score	Explanation
<input type="radio"/> Always filter and trim reads, we want high quality equal length reads to gain flexibility of using any tool of choice during downstream analysis.		
<input type="radio"/> Always trim the reads. We want only equal length reads in our data.		
<input checked="" type="radio"/> Depends on tools used during downstream analysis.	✓ 1.00	
<input type="radio"/> Always filter reads. We want only high quality reads in our data.		
Total	1.00 / 1.00	

Question 7

What is ChIP sequencing most commonly used to measure?

Your Answer	Score	Explanation
<input type="radio"/> Expression levels of particular proteins		
<input type="radio"/> Ratios of miRNA to DNA		
<input checked="" type="radio"/> The locations of protein to DNA interaction	✓ 1.00	
<input type="radio"/> Methylation of bases		
Total	1.00 / 1.00	

Question 8

What is MACS used for?

Your Answer	Score	Explanation
<input type="radio"/> Measure RNA levels		
<input checked="" type="radio"/> Peak calling/reconstruction from ChIP-seq data	✓ 1.00	
<input type="radio"/> Detect sequence variants using both qPCR and ChIP-seq data		
<input type="radio"/> Genome assembly using suffix trees and ChIP-seq data		
Total	1.00 / 1.00	

Question 9

What is the advantage of using a control in a ChIP sequencing experiment?

Your Answer	Score	Explanation
<input type="radio"/> Determine background expectation of the number of peaks detected.		
<input checked="" type="radio"/> Both options	✓ 1.00	
<input type="radio"/> Allow MACS to calculate FDR.		
<input type="radio"/> Neither option		
Total	1.00 / 1.00	