

Register Number.....

B.TECH. DEGREE EXAMINATIONS: NOV/DEC 2012

Seventh Semester

BIO TECHNOLOGY

BTY211: Genomics and Proteomics

Time: Three Hours

Maximum Marks: 100

Answer All Questions:-

PART A (10 x 1 = 10 Marks)

1. C value paradox is Lack of correlation between
 - A. DNA content and complexity
 - B. mRNA content and complexity
 - C. Protein content and complexity
 - D. Total RNA content and complexity
2. Which of the following model genome study helps in understanding developmental biology
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 - A. *C.elegans*
 - B. *E.coli*
 - C. *Arabidopsis thaliana*
 - D. Yeast
3. Major limitation of using restriction mapping method for preparing genome map of higher organisms is
 - A. large genomes cannot be restricted by RE
 - B. Large fragments cannot be separated by electrophoresis technique.
 - C. Complex technique
 - D. Error rate is high
4. FISH mapping is method
 - A. high resolution physical mapping
 - B. low resolution genetic mapping
 - C. High resolution genetic mapping
 - D. Low resolution physical mapping
5. Physical gaps detected in the genome sequence assembly can be closed by
 - A. Resequencing of already available library clones
 - B. Sequencing of new library clones
 - C. Paired end sequencing of overlapping clones
 - D. Paired end sequencing of non-overlapping clones
6. Draft genome sequence is characterized by
 - A. presence of gaps with unambiguous data
 - B. absence of gaps with ambiguous data
 - C. Presence of gaps with ambiguous data
 - D. Absence of gaps with unambiguous data

(OR)

b) Explain the following restriction mapping methods with suitable illustrations

(i) PFGE

(ii) Optical mapping

23. a) (i) What are the two commonly followed strategies for genome sequencing? (2)

(ii) Elaborate on whole genome shotgun method of genome sequence with its advantages and disadvantages associated with this method. (12)

(OR)

b) Write short notes on the following

(i) Physical gaps (4)

(ii) Sequence gaps (3)

(iii) Draft genome sequence assembly (3)

(iv) Finished genome sequence assembly (4)

24. a) What is the principle of Serial Analysis of Gene expression (SAGE) and elaborate the steps for transcript identification and quantification using SAGE.

(OR)

b) (i) Elaborate on fabrication of microarray using photolithography. (6)

(ii) How will you identify gene expression in two different tissues using microarray hybridization. (8)

25. a) Write the principle of MALDI-ToF Mass Spectrometry and how will you identify the proteins using mass spectrometry?

(OR)

b) How will you exploit iso- electric point and molecular weight of proteins using two dimensional gel electrophoresis?
