## (102)輔仁大學碩士班招生考試試題

考試日期:102年3月8日第3節

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## 科目: 分子生物學

系所組:營養科學

- I. Explain the following terms: (6 points for each sentence, 30 points)
  - 1. Shine-Dalgaron sequence
  - 2. RNA splicing
  - 3. Wobble pairing
  - 4. Okazaki fragment
  - 5. Ubiquitination
- II. Question and Answer: (10 points for each question, 30 points)
  - 1. Please detail described the mechanism of post-transcriptional modification of RNA. (10%)
  - 2. What is end-replication problem? How eukaryotic cells to solve this problem? (10%)
  - 3. What is Chromatin immunoprecipitation? Please describe the principle of the methods and how to use this technology. (10%)
- III. Choice questions: (4 points for each question, 40 points)
  - 1. Which of the following is not involved in DNA replication?
    - A. Ligase.
    - B. Helicase.
    - C. SSB.
    - D. Topoisomerase.
  - 2. Which of the following is not involved in the eukaryotic DNA polymerase I function?
    - A. 5' to 3' DNA-Dependent DNA polymerase activity.
    - B. 3' to 5' exonuclease activity.
    - C. 5' to 3' exonuclease activity.
    - D. 3' to 5' DNA-Dependent DNA polymerase activity.
  - 3. Which of the following is not involved in G1 cell cycle regulation?
    - A. CDK1.
    - B. Cyclin D.
    - C. CDK4.
    - D. p21 Waf1/Cip1
  - 4. Which of the following is not correct?
    - A. Okazaki fragments have short RNA sequence.
    - B. DNA polymerase II can remove the RNA from Okazaki fragment.
    - C. DNA ligase I can join the two fragments of DNA.
    - D. Okazaki fragments are synthesized from DNA polymerase III.
  - 5. Which of the following structure is not a DNA-binding domain
    - A. Helix-loop-helix.
    - B. Zinc finger.
    - C. Basic region-leucine zipper.
    - D. Copper finger.
  - 6. Which of the following description is incorrect?
    - A. Bacterial sigma factor can make direct contacts with promoter sequence and ensure transcription begins at the proper position in the DNA.
    - B. Histone modification and chromatin structure must be change before transcription initiation.
    - C. In transcription initiation, transcriptional factors form number of covalent interactions with DNA to ensure the compact interaction.
    - D. The process of transcription can be divided to initiation, elongation and termination.

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7. Which of the following description is correct?

- A. The SCF complex is an ubiquitin ligase that regulates the transition from G1 into S phase by degrading specific proteins that are phosphorylated by G1-S cyclin-Cdks.
- B. Histone acetyltransferases (HATs) and histone deacetylases (HDACs) are two of the chromatin modification enzymes, and HDACs remove the acetyl groups when DNA replication or transcription initiation.
- C. Both of RNA helicase and 5' to 3' exonuclease are included in degradosome.
- D. Stem-loop and poly(U) structures are two specific transcriptional termination structure signals.
- 8. Which of the following description is correct?
  - A. Restriction fragment length polymorphism (RFLP) is a tool to analyze different RNA sequences.
  - B. Polymerase chain reaction (PCR) is to detect RNA expression.
  - C. Electrophoresis mobility shift assay (EMSA) is to analyze the transcriptional factor binding to DNA sequence ability.
  - D. Fluorescence in situ hybridization (FISH) is used to detect the specific RNA sequences on chromosomes.
- 9. Which of the following description is incorrect?
  - A. The key subunit of eukaryotic cells transcription factor IIB is included TBP (TATA binding protein) and TAFs (TBP-associated factors).
  - B. DNA polymerase II has 3'-5' and 5'-3' exonuclease activity and participates in DNA repair system.
  - C. Type IA and B topoisomerases break one of the two strands of DNA and do not require ATP.
  - D. The end replication problem can be solved by telomerase.
- 10. Which of the following description is correct?
  - A. The DNA-binding domain helix-turn-helix using recognition helix (C-terminal helix) to fit in the minor groove of DNA and forms contacts with the base pairs that read the DNA sequence.
  - B. Eukaryotic pre-RNA must undergo 5'-capping by 7-methyl adenosine to protect pre-RNA degradation.
  - C. Eukaryotic cells can utilize RNA interference system to detect and degrade foreign RNAs produced by parasites.
  - D. The function of Uvr B in nucleotide excision repair is to nick the DNA backbone up- and down-stream of the lesion.