# 國立彰化師範大學 101 學年度碩士班招生考試試題

系所: 生物技術研究所	科目: 分子生物學
☆☆請在答案卷上作答☆☆	共4頁,第1頁
Section I: 單選題,Please choose the best answer for each question. Each 2%	6
<ol> <li>The process for transfer a plasmid into <i>E. coli</i> was called</li> <li>(a) Transfection, (b) Transformation, (c) Transgenic, (d) Transduc</li> </ol>	tion, (e) Transferring
<ul> <li>2. Which of the following is <u>NOT</u> usually found in RNA structure?</li> <li>(a) Adenine, (b) Guanine, (c) Cytosine, (d) Uracil, (e) Thy</li> </ul>	mine
<ul> <li>3. Which of the following is <u>NOT</u> required for cDNA systhesis?</li> <li>(a) DNA, (b) dNTP, (c) reverse transcriptase, (d) oligo(dT),</li> </ul>	(e) none of the above
<ul> <li>4. Which of the following is used for study protein-DNA interaction?</li> <li>(a) SDA-PAGE, (b) DNA microarray, (c) DNA fingerprinting,</li> <li>(d) Two-dimensional gel electrophoresis, (e) DNase footprinting</li> </ul>	
<ul> <li>5. Which of the following belong to prokaryotic promoters?</li> <li>(a) Pribnow box and TATA box, (b) TATA box and CAAT box, (c) -</li> <li>(d) Pribnow box and -35 box, (e) TATA box and -35 box</li> </ul>	-10 box and TATA box,
<ul><li>6. Which of the following play an important role in prokaryotic transcription to</li><li>(a) sigma factor, (b) alpha factor, (c) beta factor, (d) Rho factor,</li></ul>	ermination? (e) none of the above
<ul> <li>7. Which of the following can induce the most gene expression in <i>E.coli lac</i> op (a) Glucose high and lactose high, (b) Glucose low and lactose high, (c) Glucose high and lactose low, (d) Glucose low and lactose low and lactose</li></ul>	beron? (e) all of the above
8. How many RNA polymerase in <i>E.coli</i> ? and How many RNA polymerase in (a) 3 and 3, (b)1 and 3, (c)1 and 1, (d) 3 and 1, (e) none of the a	yeast? above
<ul><li>9. In eukaryotic transcription of class II genes, which transcription factor can s</li><li>(a) TFIID, (b) TFIIE, (c) TFIIS, (d) TFIIB, (e) TFIIF</li></ul>	timulate elongation?
<ul> <li>10. In eukaryotic transcription of class I genes, which are class I transcription fa</li> <li>(a) TFIID and TFIIS,</li> <li>(b) TFID and TFIB,</li> <li>(c) SL1 and UBF,</li> <li>(d) SL1 and TF IIIA,</li> <li>(e) UBF and TFIIIB</li> </ul>	ictors in mammals?
<ul><li>11. Which of the following gene with internal promoter in eukaryotic transcription (a) 5.8S rRNA, (b) 18S rRNA, (c) 5S rRNA, (d) U6 snRNA, (e)</li></ul>	ion? 28S rRNA
<ul><li>12. In <i>E.coli trp</i> operon, what is the function of tryptophan?</li><li>(a) activator, (b) repressor, (c) co-activator, (d) co-repressor,</li></ul>	(e) silencer

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☆☆請在答案卷上作答☆☆ 共4頁,第2頁
13. In prokaryotic translational initiation step, which of the following is <b>NOT</b> involve in 30S initiation complex?
(a) mRNA, (b) GTP, (c) IF2, (d) $23S$ rRNA, (e) $16S$ rRNA.
14. In prokaryotic translational elongation step, how many GTPs are required for each amino acid synthesis?
(a) 1, (b) 2, (c) 3, (d) 4, (e) none of the above
<ul> <li>15. In eukaryotic transcription, 5.8S rRNA is produced by ?</li> <li>(a) RNA polymerase I,</li> <li>(b) RNA polymerase II,</li> <li>(c) RNA polymerase III,</li> <li>(d) DNA polymerase I,</li> <li>(e) RNA polymerase I and RNA polymerase III</li> </ul>
<ul><li>16. The codons in the genetic code that do not specify amino acids are called</li><li>(a) missense codons, (b) start codons, (c) stop codons, (d) promoters, (e) initiator codons</li></ul>
<ul> <li>17. For the DNA strand 5'-TACGATCATAA-3' the correct complementary DNA strand is:</li> <li>(a) 3'-TACGATCATAA-5', (b) 3'-ATGCTAGTATT-5', (c) 3'-AUGCUAGUAUU-5',</li> <li>(d) 3'-GCATATACGCC-5', (e) 3'-AATACTAGCAT-5'</li> </ul>
<ul> <li>18. Which of the following tools of recombinant DNA technology is <u>INCORRECTLY</u> paired with one of its uses?</li> <li>(a) restriction endonuclease - production of DNA fragments for gene cloning</li> <li>(b) DNA polymerase - copies DNA sequences in the polymerase chain reaction</li> <li>(c) reverse transcriptase - production of cDNA from mRNA</li> <li>(d) electrophoresis - RLFP analysis</li> <li>(e) DNA ligase - enzyme that cuts DNA, creating sticky ends</li> </ul>
<ul> <li>19. In homologous DNA recombination via the Holliday model, the Holliday junction is</li> <li>(a) a 2-stranded structure,</li> <li>(b) generated by the action of DNA polymerase I, helicase, SSB and DNA ligase,</li> <li>(c) generated by cutting, exchanging and rejoining (intermolecularly) homologous strands,</li> <li>(d) b and c,</li> <li>(e) none of the above</li> </ul>
<ul> <li>20. Which of the following statements about the 3' poly(A) tail of mRNA is <u>FALSE</u>?</li> <li>(a) It helps align eukaryotic mRNA on the ribosome during translation.</li> <li>(b) It is added to the primary transcript in the nucleus.</li> <li>(c) It is not essential for protein synthesis.</li> <li>(d) It protects mRNA from degradation.</li> <li>(e) It promotes export mRNA from the nucleus and translation.</li> </ul>

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共4頁,第3頁

<u>For questions 21~25</u>, choose the best answer to fill in the blank in the following statement about the process of "**DNA replication**".

The two antiparallel strands are replicated simultaneously in both directions. RNA primers are used to initiate a new strand. The parent strand at the 3' end of the template determines the (21)\_\_\_\_\_\_strand in continuous replication. The parent strand at the 5' end of the template produces the (22)\_\_\_\_\_\_strand as short pieces of DNA (100-200 nucleotides in eukaryotes and longer in prokaryotes). The (23)\_\_\_\_\_\_strand fragments are called Okazaki fragments after their discoverer, Reiji Okazaki. The RNA primers are removed by (24)\_\_\_\_\_and the fragments are joined by (25)\_\_\_\_.

21. (a) leading, (b) lagging, (c) DNA, (d) RNA, (e) none of the above

22. (a) leading, (b) lagging, (c) DNA, (d) RNA, (e) none of the above

- 23. (a) leading, (b) lagging, (c) DNA, (d) RNA, (e) none of the above
- 24. (a) RNase, (b) DNase, (c) DNA polymerase, (d) RNA polymerase, (e) DNA ligase
- 25. (a) RNase, (b) DNase, (c) DNA polymerase, (d) RNA polymerase, (e) DNA ligase

26. How do dideoxynucleoside triphosphates (ddNTPs) terminate a nascent DNA strand?

- (a) They possess a bulky additional group which causes DNA polymerase to dissociate.
- (b) They have no 3' hydroxyl group, so cannot form a phoshodiester bond with the 5' phosphate group of the next nucleotide.
- (c) They form abnormal hydrogen bonds causing the DNA duplex to unwind.
- (d) They form a hair pin structure which stop DNA polymerase activity.
- (e) They form a cross-link between DNA polymerase and the DNA duplex.
- 27. Telomerase is important to eukaryotic cells because \_\_\_\_\_.
  - (a) the leading strand of DNA causes the telomeres to shorten.
  - (b) telomeres attach to MTOC during cell division.
  - (c) telomerase digests telomeres to proper length.
  - (d) telomeres tend to get shortened with each cell division.
  - (e) telomerase activates cyclin B for cell cycle progression.
- 28. Which of these might be an advantage to genetic testing of individuals via microarrays?
  - (a) Many different potential mutations in a single gene could be tested at once.
  - (b) Expression patterns of many different genes can be analyzed simultaneously.
  - (c) Microarray analysis can provide information on relative levels of expression of particular genes.
  - (d) All of the above.
  - (e) None of the above.

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共4頁,第4頁

#### ☆☆請在答案卷上作答☆☆

29. Branch migration is promoted by

(a) recA,
(b) recBCD,
(c) ruvA & ruvB,
(d) recF,
(e) SSB

30. Nuclear mRNA splicing is performed\_\_\_\_\_\_.

(a) in nucleosomes,
(b) in spliceosomes,
(c) in the nucleolus,
(d) by RNA only,
(e) b and c

Section II: 問答題

- Please describe how to make a 250 mL protein extraction buffer, that is 0.5% Nonidet (v/v), 150 mM Tris-HCl, pH 7.5, and 10 mM EDTA, from the given following stock solutions: 100% Nonidet P-40, 1 M Tris-HCl, pH 7.5, and 0.5 M EDTA. (10%)
- 2. Below is a typical eukaryotic gene, its pre-mRNA, and its mature mRNA. The direction of transcription is from left to right. **Label** the diagram with the following structures: start codon(s), stop codon(s), 3'UTR(s), 5'Cap(s), open reading frame(s), promoter(s), poly(A)tail(s), transcription termination signal(s), intron(s), and exon(s). (10%)



- 3. Please diagram and describe the four-step transcription initiation process in *E. coli*. (5%)
- 4. What is iPS and its application? Please give at least one example. (5%)
- 5. What is DNA microarray and its application? Please give at least one example. (5%)
- Please describe detail about how to produce recombinant insulin from *E. coli* BL21(DE3)pLysS. (5%)