

## Quiz/Self-assessment-II

Q. 1 Shown below is the fragment of an *E. coli* gene (the bold and underlined nucleotides are numbered):

```

      -40      -30      -20      -10      +1
5' - AGGC TTGACA CTTTATGCTT CCGGCTCG TATAAT GTCTGCAAT - 3'
3' - TCCGAACTGT GAAATACGAAG GCCGAGCAT ATTACAGAGCTT - 5'
  
```

- Name the regions highlighted in yellow and green?
- What is the basis of the numbering of the nucleotides?
- Which of the above strands (top strand or bottom strand) is the template strand?

**Answer:**

- Yellow: -35 box                      Green: -10 box**
- +1 is the first nucleotide to be transcribed; sequences in the 5' direction on the non-template strand are negatively numbered.**
- Bottom strand is the template strand (Transcription is from 5' – 3' end)**

Q. 2 Complete the following table:

	Replication	Transcription	Translation
Where does this take place in an eukaryotic cell?	<b>Nucleus</b>	<b>Cytoplasm</b>	<b>Ribosomes</b>
Which enzyme/protein complex carries out this process	<b>DNA polymerase</b>	<b>RNA polymerase</b>	<b>Ribosomes</b>
What is the template that is read during this process?	<b>DNA</b>	<b>DNA</b>	<b>mRNA</b>
Which direction the template is read in?	<b>3' – 5'</b> <b>(Because synthesis is from 5' – 3')</b>	<b>3' – 5'</b> <b>(Because synthesis is from 5' – 3')</b>	<b>5' – 3'</b>
What is the start signal for this process	<b>Origin of replication</b>	<b>Promoter</b>	<b>Ribosome binding site in prokaryotes</b>  <b>5' cap in eukaryotes</b>
What is the product of this process	<b>DNA</b>	<b>RNA</b>	<b>Protein</b>
What are the monomers used in this	<b>Deoxyribonucleotides</b>	<b>Ribonucleotides</b>	<b>Amino acids</b>
What type of bond is formed between the monomers?	<b>Phosphodiester</b>	<b>Phosphodiester</b>	<b>Peptide</b>

Q. 3 Insertion or deletion of one or two base-pairs from the coding region of the gene changes the reading frame of the gene; such mutations are known as .....**frameshift mutations**.....

Q. 4 In the process of translation, each amino acid is coded for by 3 nucleotides—a codon. Why does it have to be at least 3 nucleotides as opposed to 2 or 1 nucleotides coding for an amino acid?

**Answer: As there are only four nucleotides present in RNA, a single nucleotide code can code only for  $4^1 = 4$  amino acids, a two nucleotide code can code only for  $4^2 = 16$  amino acids, and a three nucleotide code can code for  $4^3 = 64$  amino acids. As we have 20 different amino acids in the proteins, the code has to be at least three nucleotides long.**

Q. 5 Codon is the three nucleotide code present in .....**mRNA**.....

Q. 6 How many codon are required for specifying 5 amino acids?

**Answer: Five codons (15 nucleotides)**

Q. 7 What do aminoacyl t-RNA synthetases do?

**Answer: They attach an amino acid to the 3'-end of the tRNA molecules**

Q.8 The 80S eukaryotic ribosome is composed of two subunits of .....**60S**..... and .....**40S**.....

Q. 9 Given below is a hypothetical bacterial gene:

-40                -30                -20                -10                +1

5' -AGGCTTGACACTTTATGCTTCCGGCTCGTATAATGTCTGCAATAGGAGGTGACTATCCTCCAGTGA-3'

3' -TCCGAACTGTGAAATACGAAGCCGAGCATATTACAGAGCTTATCCTCCACTGATAGGAGGTCACT-5'

- (a) Write down the sequence of the RNA molecule that will be synthesized from the above gene.  
(b) What will be the length of the protein/peptide synthesized from this mRNA (Given: AGGAGGU is the ribosome binding site on the mRNA).

**Answer: (a) AAUAGGAGGUGACUAUCCUCCAGUGA**

**(Justification: As bottom strand is the template strand, the other strand is the sequence of the RNA molecule to be synthesized with T replaced by U).**

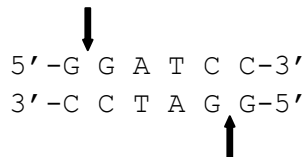
**(b) Zero**

**(Justification: The first amino acid will be coded by AUG (initiation codon), as there is no AUG in the RNA molecule, the length of the peptide/protein to be synthesized is zero).**

Q. 10 Match the columns

A	B
(a) <i>E. coli</i> is grown in a medium having both glucose and lactose	(i) Binds to the operator
(b) Repressor	(ii) Binds to the promoter
(c) RNA polymerase	(iii) Repressor becomes inactive
(d) Bacteria growing in a medium having lactose as the only carbon source	(iv) Catabolite activator protein (CAP) active
	(v) High concentration of cyclic AMP

Q.11 BamHI is a restriction endonuclease with the cleavage sites as shown below:



Which of the following statements is/are correct (tick all correct options):

- (a) The enzyme produces sticky ends with 5' overhangs ✓
- (b) The enzyme produces blunt ends
- (c) The enzyme produces sticky ends with 3' overhangs
- (d) The enzyme produces sticky ends with 3' overhang in one strand and 5' overhang in the other

Q. 12 Mention in one or two sentences is the function of  $\beta$ -galactosidase in *lac* operon?

**Answer:**

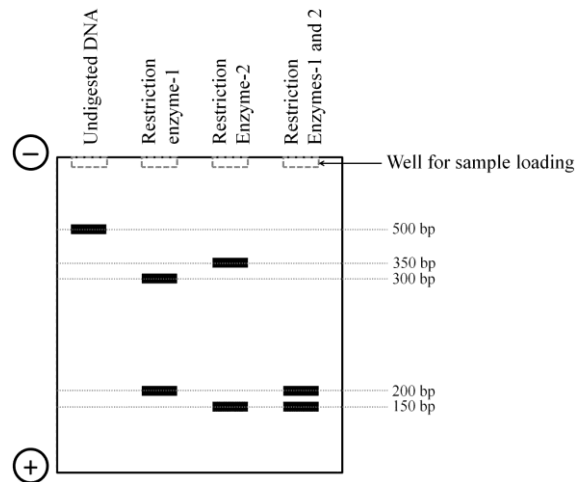
1.  $\beta$ -galactosidase converts lactose into allolactose
2.  $\beta$ -galactosidase converts lactose into glucose and galactose

Q. 13 Name three key steps of polymerase chain reaction.

**Answer:**

1. Denaturation
2. Annealing
3. Extension

Q. 14 Shown below is the gel run following restriction digestion of a given linear DNA molecule with two restriction enzymes. Prepare the restriction map of the enzyme:



**Answer:**

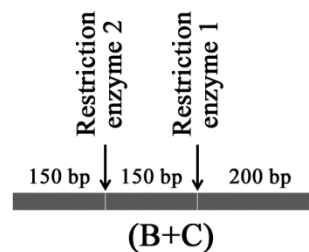
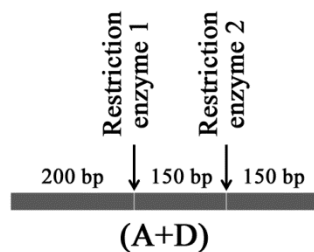
**Digestion with restriction enzyme-1 gives two bands of sizes 200 bp and 300 bp. This results in following two possibilities:**



**Digestion with restriction enzyme-2 gives two bands of sizes 350 bp and 150 bp:**

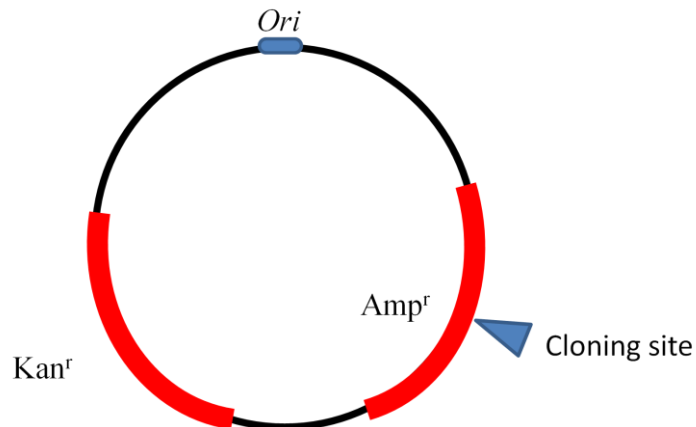


**Digestion with both restriction enzymes-1 and 2 gives two bands of sizes 200 bp and 150 bp. The possible restriction maps from the given data are: (A+D) and (B+C) i.e.**



**The combinations, (A+B), (A+C), (B+D), and (C+D) do not fit the given data.**

Q. 15 You performed cloning of a gene using following vector:



where, Kan<sup>r</sup> and Amp<sup>r</sup> are the Kanamycin resistance and ampicillin resistance genes, respectively. You go ahead like this:

- (a) Cut the plasmid with the appropriate enzyme.
- (b) Cut out your gene of interest from the genome using the same enzyme.
- (c) Mix the cut plasmid and the gene of interest and add a DNA ligase in the reaction. Allow the reaction to go for some time
- (d) Perform bacterial transformation (putting the DNA into bacterial cell) with the above mix.
- (e) Grow these bacteria in a suitable liquid medium

How will you now select the bacteria that have got your gene of interest? Give your answers step-by-step (in not more than 5 sentences).

**Answer:**

- 1. Grow the bacteria after step (e) on a plate containing Kanamycin**
- 2. Only those bacteria that have got the plasmid will grow and produce colonies**
- 3. Take a small amount of bacteria from the plate and try to grow them on the plates containing ampicillin.**
- 4. The bacteria that do not grow on ampicillin have the gene of interest.**