## **Questions bank**

Genetics-3<sup>rd------Asst</sup> prof-Dr-Kazhal M.Sulaiman

Biology

Q1-What is homologous chromosome, gene and alleles

Q2 - How do genes work? Give an example )

Q3-Discuss this phrase (The genetic code is degeneracy

**Q4**-Which part of chromosome was required for the replication and stability of the chromosome? Explain M)

**Q5-** What are the differences between the followings:

A- Transition and transvertion mutation B- Germinal and somatic mutations

C-Euchromatin and heterochromatin **D**- Acrocentric and telocentric chromosome

**E-** Induced and spontaneous mutations

**Q6-**Illustrate the structure of nucleosome

**Q7-** Recognize different kinds of mutations (missense, non sense, frameshift) and predict the effects on amino acid sequence.

**Q8-** Chromosomes in Eukaryotes and prokaryotes are different, Explain **Q9-**Define the followings: -

A-Chromatin /B- Genotype / C-Alleles / D-Homozygous /

Q10- Explain by figure all types of chromosomes according to centromere position.

Q11- Genetic code where to begin and stop?

Q12-Explain Chromosomes in Eukaryotes

Q13-What is a gene ? How do genes work? Give an example

Q14-Define the followings: -

A-Terminal deletion B- Aneuploidy C-Reciprocal translocation D-Nullisomy

Q15- Explain by figure how thymidine dimer is formed.

**Q16-** Explain chromosomal non disjunction and then illustrate how this case of chromosomal abnormalities happened (XYY)

Q17-What is the functions of the Genetic Material DNA?

Q18- What is the differences between the followings:-

Paracentric and pericentric inversion

A- Chromosomal mutation and gene-point mutation

Q19-Define the followings : -

B- A-Recombination B- Genotype C-Alleles D-Homozygous

**Q20-** More than one codon may specify the same amino acid. What is the name of this phenomena? Explain with example

Q21- Genetic code where to begin and stop

**Q22-** hat is telomerase ?

Q23- What is genome?.

Q24// what is heterochromatin

Q25// What is genetic code ? Where to begin and stop

Q26// Explain by Figure Chromosome Organization & DNA Packaging.

Q27// A-What is the function of genetic material

**B-** Recognize different kinds of mutations (missense, non sense, frameshift) and predict the effects on amino acid sequence

- Q28// Explain chromosomal non disjunction and then illustrate how those cases of chromosomal abnormalities happened(XXX, ,XYY and X0)
  - **Q29-** Explain how thymidine dimmers happened in DNA.
- Q30// More than one codon may specify the same amino acid. Explain this Phenomenon.

Q31// What are the differences between the followings

A- Chromosomal and gene – point mutation

- **B-** Somatic and germinal mutation
- C- Euchromatin and heterochromatin
- **D-** Transition and transversion
- **E-** Paracentric and Pericentric inversion

Q32-. Fill in the blanks:-

1- Detailed analysis of the nucleosome core particles has shown they contain-----

Base pairs of DNA wrapped------times around a histone core consisting of two molecules each of ------,-----,-----.

2- Cri-due-cat syndrome result from----- in a section of ----- on chromosome-

**3-** What are the different syndrome result from Non- disjunction in Oogenesis----

**4-** Move DNA from one chromosome to a non-homologous chromosome called -

Q33.Who would each of the following types of mutations affect the amount of protein that is expressed from a gene:- Non- sence , Missence , Frameshift.

**Q34.** What is genetic code ? Where to begin and stop ?

Q35.Which type of mutation have no effect on the encoded protein ?

Q36. Fill in the blank:-

- 1- Karyotype is------, while idiotype is-----.
- **2-** Euchromatin region that are ----- staining, and contain most of ------, while heterochromatin region that are----- staining, and contain most of ------.
- **3-** The two ends of a chromosome are known as-----, it is required for the ------ of chromosome.
- 4- What are the different syndrome result from non-disjunction in spermatogenesis------
- Q37. What is gene? How do gene work?
- Q38-Explain how thymidine dimmers happened in DNA
- Q39. What are the differences between chromosomal mutations and gene point Mutations?