

Name \_\_\_\_\_

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**AP: CHAPTER 17: FROM GENE TO PROTEIN**

1. How did diseases involving metabolic pathways lead to hypotheses about the nature of genes?

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2. Identify some genetic diseases that occur along metabolic pathways.

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3. What was Beadle and Tatum's hypothesis regarding enzymes?

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4. How has that hypothesis been modified?

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5. What occurs during transcription?

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6. What occurs during translation?

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7. How does the protein process differ in prokaryotes and eukaryotes?

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8. Briefly explain how Marshall Nirenberg and Heinrich Matthaei "cracked the genetic code?"

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9. What is the genetic code and why is said to be universal?

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10. List several features about the genetic code.

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11. Give an example of what happens if reading frames are altered?

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12. List the highlights of the three stages of transcription.

a. Initiation \_\_\_\_\_

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b. Elongation \_\_\_\_\_

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c. Termination \_\_\_\_\_

Name \_\_\_\_\_

13. What happens to the transcript RNA before it leaves the nucleus?

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14. What is the advantage of the 5' cap and poly A tail?

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15. Distinguish between exons and introns.

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16. Describe the mechanism for splicing RNA.

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17. What does alternative RNA processing do for cells?

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18. Identify the roles of the players of the translation process.

a. Transfer RNA \_\_\_\_\_

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b. Aminoacyl-tRNA synthetase \_\_\_\_\_

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c. Ribosomes \_\_\_\_\_

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19. Identify and briefly describe the steps of translation. Initiation Elongation Termination

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20. What is the advantage of polyribosomes?

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21. Give an example of how a polypeptide gets into the ER for additional processing.

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22. How does protein synthesis differ between prokaryotes and eukaryotes?

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23. Define point mutations.

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24. Define mutations that are:

a. Missense

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b. Nonsense

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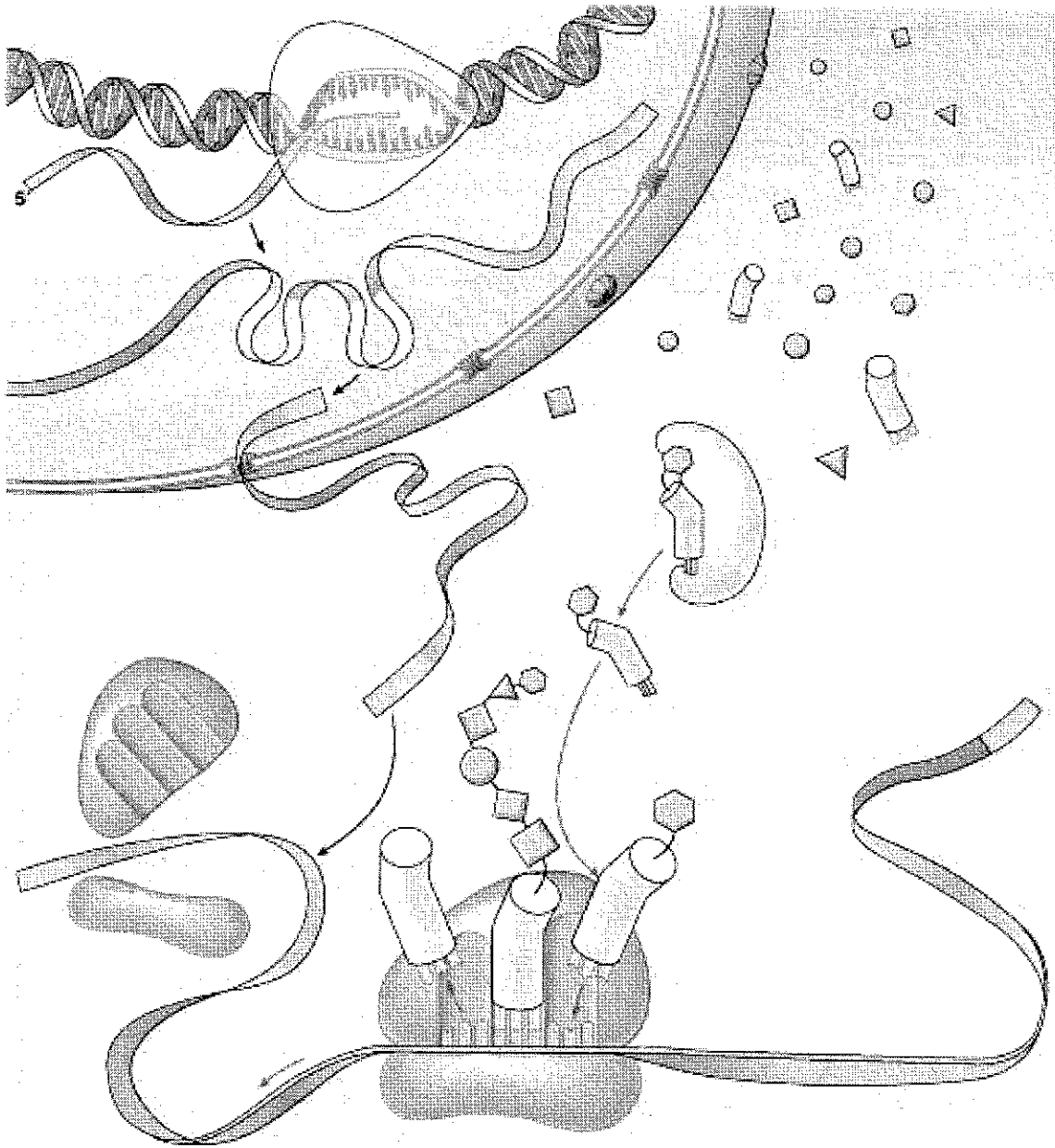
c. Insertion or deletion

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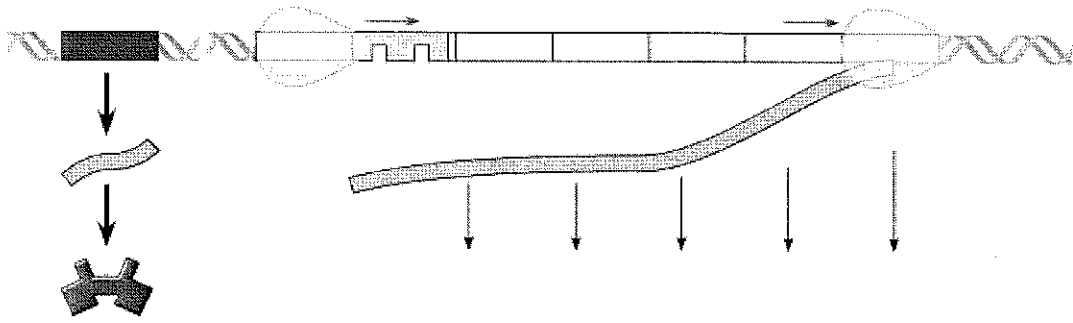
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25. Use the diagram to trace the flow of chemical information from the gene to the protein product.



Name \_\_\_\_\_

25. Use the diagram of the Tryp operon to outline how it regulated tryptophan levels.



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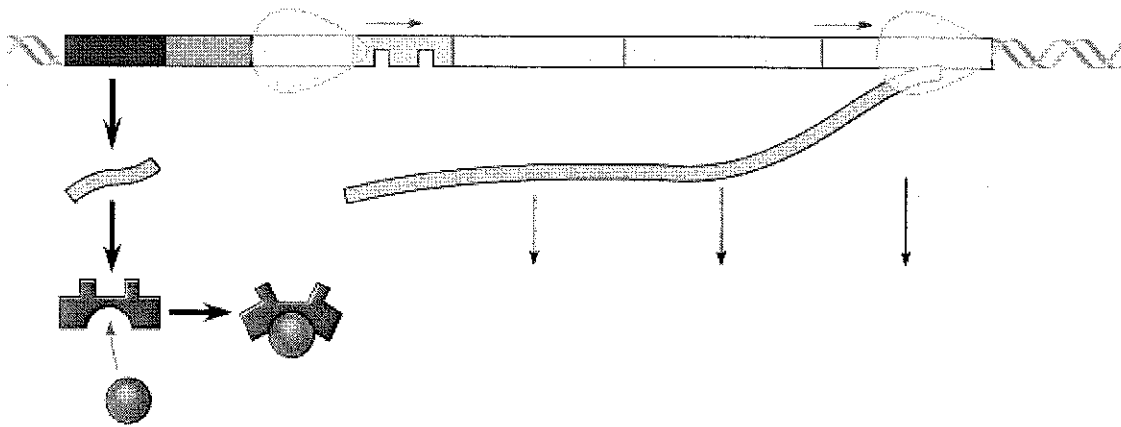
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26. Describe how the trp operon is a repressible operon.

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27. Use the diagram of the lac operon to outline how it regulates glucose levels.



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Name \_\_\_\_\_

28. Does the diagram above represent the condition for the absence or presence of lactose?

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29. Describe what happens when lactose is absent.

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30. How is the lac operon an inducible system?

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31. Summarize how the presence and absence of glucose influences the lac operon.

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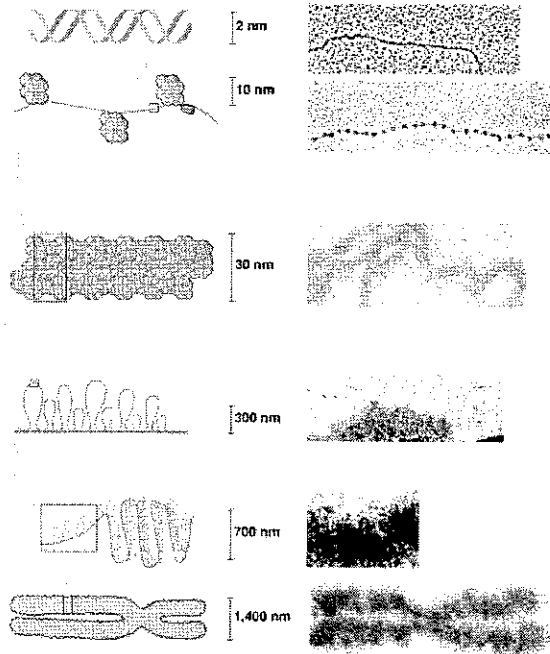
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Period \_\_\_\_\_

Date \_\_\_\_\_

## AP: CHAPTER 19: CONTROL OF EUKARYOTIC GENOME

1. Outline the levels of DNA packing within the eukaryote nucleus.



2. What is the difference between heterochromatin and euchromatin? Which is transcribed?

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3. Which regions of the chromosome will typically be in the form of heterochromatin?

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4. How do the coding regions and genome sizes of prokaryotes and eukaryotes compare?

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5. Much of mammalian non-coding DNA is in the form of \_\_\_\_\_

6. What is the cause of Fragile X?

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7. What is the cause of Huntington's disease?

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8. Discuss an example of interspersed repetitive DNA?

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9. What is a multigene family?

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10. Multigene families are hypothesized to have evolved from...

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11. How is the globulin multigene family an adaptive to mammals?

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12. Explain how gene amplification can regulate gene expression.

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13. How can transposons alter gene expression?

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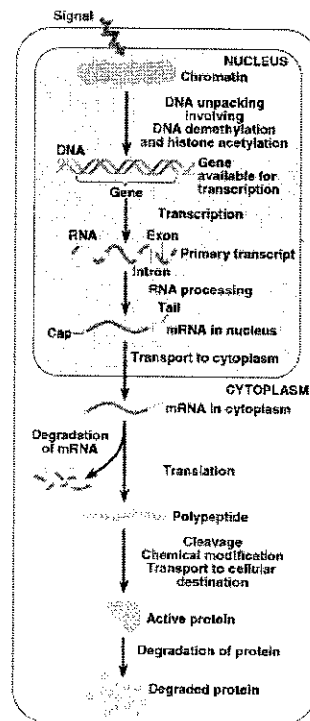
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14. How do immunoglobulin genes code for a seemingly infinite variety of antibodies?

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15. Review the opportunities for gene regulation in eukaryotes in the diagram.



16. Where is the most important step in gene regulation?

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Name \_\_\_\_\_

17. Describe the effect of each of the following control mechanisms.

- a. DNA methylation \_\_\_\_\_
- b. Histone acetylation \_\_\_\_\_
- c. Transcription factors \_\_\_\_\_
- d. Control elements \_\_\_\_\_
- e. Enhancers \_\_\_\_\_
- f. Activators \_\_\_\_\_
- g. DNA-binding domain \_\_\_\_\_

18. How does alternative RNA splicing affect gene expression?

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\_\_\_\_\_

19. How does RNA degradation affect gene expression?

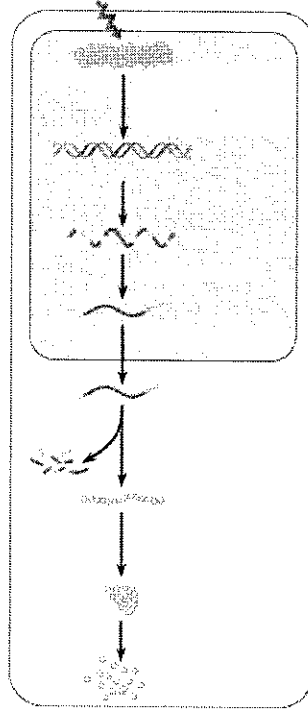
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20. How does protein processing and degradation affect gene expression?

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21. Identify the opportunities to regulate gene expression in eukaryotes.



22. Typically, what happens to cell function when cells become cancerous?

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23. What is a proto-oncogene? What happens to them when cancer occurs?

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24. List the three events that can turn proto-oncogenes into oncogenes.

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25. Identify and describe mutations in specific proteins that can lead to cancer.

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26. What is p53?

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27. Why is it said that cancer formation is a multi-step process?

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