

Mutations Worksheet KEY

Original DNA: T A C A C C T T G G C G A C G A C T
Original mRNA: A U G U G G A A C C G C U G C U G A

Original Amino Acid Sequence: Met-Trp-Asn-Arg-Cys-STOP

Mutated mRNA #1: A U G U A G A A C C G C U G C U G A
Mutated AA Sequence: Met-STOP-Asn-Arg-Cys-STOP
Likely Effects: Yes, altering the function of the protein.
Mutation Type: point nonsense mutation

Mutated mRNA #2: A U G C U G G A A A C C G C U G C U G A
Mutated AA Sequence: Met-Leu-Glu-Pro-Leu-Leu
Likely Effects: Yes, creating a mutant protein.
Mutation Type: frameshift insertion mutation

Mutated mRNA #3: A U G U G G A A U C G C U G C U G A
Mutated AA Sequence: Met-Trp-Asn-Arg-Cys-STOP
Likely Effects: none
Mutation Type: silent mutation

Mutated mRNA #4: A U G U G G A A C C G C U G A U G A
Mutated AA Sequence: Met-Trp-Asn-Arg-STOP-STOP
Likely Effects: Yes, prematurely stopping the formation of the protein so it won't be made.
Mutation Type: point nonsense mutation

Mutated mRNA #5: A U G U G G A A C C C U G C U G A
Mutated AA Sequence: Met-Trp-Asn-Pro-Ala-
Likely Effects: Yes, creating a mutant protein
Mutation Type: frameshift deletion mutation

Genetic Code Answer Key

Wenbin Ji



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Beginning with biomolecules the book introduces carbohydrates lipids nucleic acids amino acids and proteins highlighting their classification chemical nature and biological significance The section on bioenergetics explores the concepts of free energy enthalpy and entropy distinguishing between endergonic and exergonic reactions and explaining redox potential energy rich compounds and the significance of ATP and cyclic AMP Carbohydrate metabolism is detailed with pathways such as glycolysis the citric acid cycle HMP shunt gluconeogenesis and glycogen metabolism along with clinical conditions like G6PD deficiency glycogen storage diseases and diabetes mellitus The discussion on biological oxidation emphasizes the electron transport chain oxidative phosphorylation substrate level phosphorylation and the effects of inhibitors and uncouplers Lipid metabolism is explained through the oxidation of fatty acids ketone body metabolism de novo fatty acid synthesis and the biological significance of cholesterol including its conversion into bile acids steroid hormones and vitamin D with attention to disorders such as hypercholesterolemia atherosclerosis fatty liver and obesity Amino acid metabolism covers transamination deamination decarboxylation the urea cycle and its disorders catabolism of phenylalanine and tyrosine and their related diseases such as phenylketonuria albinism alkaptonuria and tyrosinemia It also highlights the synthesis and significance of key biological molecules such as dopamine serotonin 5 HT melatonin noradrenaline and adrenaline alongside heme catabolism hyperbilirubinemia and jaundice The section on nucleic acid metabolism and genetic information transfer describes the biosynthesis and catabolism of purine and pyrimidine nucleotides clinical conditions like gout the structural organization of DNA and RNA DNA replication transcription the genetic code translation and inhibitors of protein synthesis The chapter on enzymes introduces their properties nomenclature and IUB classification followed by enzyme kinetics explained through Michaelis and Lineweaver Burke plots It also addresses enzyme inhibition with examples regulatory mechanisms including induction repression and allosteric regulation and their therapeutic and diagnostic applications Isoenzymes are discussed in medical contexts while the section on coenzymes explains their structures and biochemical functions Altogether this textbook integrates biochemical principles with clinical correlations making it an essential guide for pharmacy medical and life science students bridging molecular understanding with disease mechanisms and therapeutic applications

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