Name:	TOC#
Mutation/ Activity	
Introduction: DNA is genetic material made of nucleotides. Last unit we through transcription (DNA→mRNA) and translation (mRNA→linked as want to see how those processes can "go wrong" and create mutations. In gene mutation in which there is a change in one nucleotide. This can be as where a base pair is "swapped" with the incorrect base. Or it can be an inframeshift mutation. You will see how these changes affect the way the retranslated to protein. Explore the effects of gene mutations.	we saw how proteins were created amino acids). However, in this unit we in this activity you will investigate a a point mutation called a substitution, assertion or deletion: which will cause a
Concepts: By the end of this lab you should	
-be refreshed on transcription and translation	
-understand how a point mutation can alter a gene	
-understand the different types of point mutations	
<b>Background:</b> DNA is an example of a complex biological polymer called small subunits called nucleotides. There are four possible nitrogen bases cytosine (C), and thymine (T). The nitrogen bases will preferentially born base—adenine with thynine and guanine with cytosine. The bonded nitrog DNA is used to complete protein synthesis.	in DNA—adenine (A), guanine (G), and with only one other nitrogenous
During protein synthesis at the ribosome, mRNA sequences are real. Another area of the tRNA transports a specific amino acid. The amino acid enzymes to form proteins.	
Depending on the DNA sequence The 20 amino acids are brough (tRNA). An infinite variety of proteins can be formed from the 20 amino and in any order.	
These amino acids are specified by codons (or three nucleotides is copied incorrectly during transcription a mutation will occur. A <i>gene m</i> of a gene. Point mutations, or <i>single nucleotide polymorphisms</i> (SNPs), is of the three nitrogen bases in a codon. Perform this activity and witness to DNA can have on a resulting protein.	utation is a change in a single base pair involve only one nitrogen base change
Pre-lab Questions  1. This lab shows the effect of point mutations, which will cause a general what is the difference between this and a chromosomal mutation	
<ul><li>2. Describe each of the mutations:</li><li>a. Substitution</li></ul>	
b. Insertion	
c. Deletion	
3. Write the correct type of mutation that occurred in creating the mexamples. Original DNA: ATCCGCTTACG  a. UUGGCGAAUCG:  b. UATGGCGAAUCG:	RNA strand for each of the following
c. UGGCGAAUCG:	

- 4. An insertion and deletion will cause a "frameshift." Explain what this means. Is this more or less likely to cause a noticeable mutation than a substitution?
- 5. Refresh your knowledge on how to use the universal codon chart below

	U	С	A	G	AUG codes for:
	UUU = phe	UCU = ser	UAU = tyr	UGU = cys	AUG codes for.
U	UUC = phe	UCC = ser	UAC = tyr	UGC = cys	
٠	UUA = leu	UCA = ser	UAA = stop	UGA = stop	
	UUG = leu	UCG = ser	UAG = stop	UGG = trp	
	CUU = leu	CCU = pro	CAU = his	CGU = arg	CCG codes for:
С	CUC = leu	CCC = pro	CAC = his	CGC = arg	
ŭ	CUA = leu	CCA = pro	CAA = gln	CGA = arg	
	CUG = leu	CCG = pro	CAG = gln	CGG = arg	
	AUU = ile	ACU = thr	AAU = asn	AGU = ser	GAU codes for:
Α	AUC = ile	ACC = thr	AAC = asn	AGC = ser	OAU codes for.
^	AUA = ile	ACA = thr	AAA = Iys	AGA = arg	
	AUG = met	ACG = thr	AAG = lys	AGG = arg	
	GUU = val	GCU = ala	GAU = asp	GGU = gly	UAA codes for
G	GUC = val	GCC = ala	GAC = asp	GGC = gly	
۳	GUA = val	GCA = ala	GAA = glu	GGA = gly	
	GUG = val	GCG = ala	GAG = glu	GGG = gly	

## **Procedures:**

- 1. Using the DNA strand on the "Mutation Consequences Worksheet" transcribe the DNA into mRNA.
- 2. Using the Universal Codon Chart found in the pre-lab translate the mRNA into its corresponding amino acid sequence.
- 3. Obtain a six pieces of paper labeled 1 through 6 from your teacher. Place them face down on your table.
- 4. The first nucleotide that will be mutated is the 4<sup>th</sup> nucleotide in the strand of DNA.
- 5. Each student will pick a slip of paper and complete the worksheet for the specific mutation they drew, by following the instructions below.
- 6. No student in a group may use the same number.

Number	Corresponding Action
Drawn	
1	Substitute specified nucleotide with an A
2	Substitute specified nucleotide with a C
3	Substitute specified nucleotide with a G
4	Substitute specified nucleotide with a T
5	Delete the nucleotide
6	Insert a nucleotide immediately after the specified nucleotide. Toss the die
	again until you roll 1–4 to determine which nucleotide will be inserted.

- 7. Write the complete DNA sequence with one mutation in nucleotide 4.
- 8. Complete the mRNA sequence from the mutated DNA.
- 9. Translate the amino acid sequence from the mRNA.
- 10. Circle any differences from the original protein produced.
- 11. Repeat this activity again having the mutation occur at nucleotide 6. Each group member must use a new mutation.
- 12. Answer the questions.

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nutate the	e 4 <sup>th</sup>	nuc	leoti	de).	Mal	ke sı	ure (	each	gro	Tria up 1		ıber	is do	oing	a di	ffer	ent 1	muta	ation	ı	
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Nucleotide	T	Α	C	G	Т	G	Т	Т	A	G	T	C	Т	A	Α	G	A	Α	Α	C	Т
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Sequence																					
Number rolle	ed			_																	
Mutated DN	- 1																$\top$				
Sequence	$\overline{}$	_	_	+	+	_				_	$\dashv$	_	_	+	_	+	+	+	+	+	
mRNA from mutated DN	- 1																				
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ember is	doir	ng a	diff6	eren 4	t mu	tati 6	on 7	8	9	doir	ng a	12	13	14	15	16	17	18	19	20	21
ember is	doir 1 T	a g a a a a a a a a a a a a a a a a a a	3 C	4 G	t mu	6 G	7 T	8 T	9 A	doir 10 G	ng a	12 C	13 T	14 A							
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Qı	estions:
1.	What is the significance of choosing a number at random to determine how to mutate the DNA?
2.	Did your mutation in trial 1 cause a change in the amino acid sequence produced? How about during trial 2? Explain why or why not.
3.	Compare your own outcome in trial 1, to your own outcome in trial 2. What do you notice?
4.	Compare the outcomes in trial 1 with your group. Did anyone produce the same exact protein?
5.	Is it possible to have a mutation in nucleotide 4 that would produce the same amino acid?
6.	Could any mutations have occurred in nucleotide 6 that would produce the same amino acid as was produced from the original DNA sequence?