No part of the candidate evidence in this exemplar material may be presented in an external assessment for the purpose of gaining credits towards an NCEA qualification.



91159



Level 2 Biology, 2016

91159 Demonstrate understanding of gene expression

9.30 a.m. Friday 18 November 2016 Credits: Four

Achievement	Achievement with Merit	Achievement with Excellence
Demonstrate understanding of gene expression.	Demonstrate in-depth understanding of gene expression.	Demonstrate comprehensive understanding of gene expression.

Check that the National Student Number (NSN) on your admission slip is the same as the number at the top of this page.

You should attempt ALL the questions in this booklet.

If you need more space for any answer, use the page(s) provided at the back of this booklet and clearly number the question.

Check that this booklet has pages 2–12 in the correct order and that none of these pages is blank.

YOU MUST HAND THIS BOOKLET TO THE SUPERVISOR AT THE END OF THE EXAMINATION.

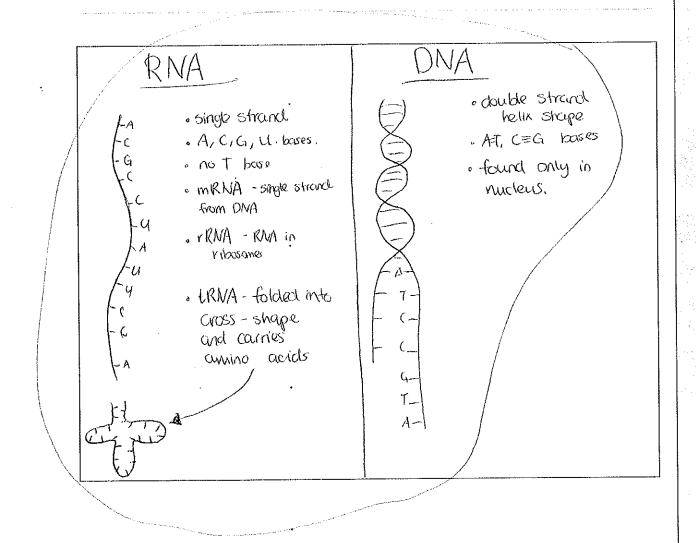
Merit **TOTAL**

(a) Deoxyribonucleic acid (DNA) and ribonucleic acid (RNA) are both involved in protein synthesis.

Describe the structure of DNA and RNA.

You may use diagrams in your answer.

DNA is a double stranded helix. RMI is a single Strander used for transcription and translation which has one more oxygen in the sugar-phosplate Backbone than DNA



(b) DNA, mRNA, and tRNA are all involved in the formation of proteins.

Discuss the significance of these molecules in forming proteins, and why the cell continually makes mRNA molecules, but not DNA molecules, during protein synthesis.

In your answer include:

- an explanation of the function of DNA, mRNA, and tRNA molecules
- an explanation of how mRNA is produced
- a discussion of the significance of DNA, mRNA, and tRNA in forming specific proteins.

DNA provides a base and an order for the p code for a protein which is required. It codes for the protein, mRNA is made corresponding to the bases on the DMA (A=4 and C=G). MRNA a substitute for DM as the DM is too by and too to leave the nucleus. RNA polymerase reads along the DNA and produces the mRNA strand. It has an extra oxygen which allows it to be more sturdy to leave the nucleus. tRNA is the molecule that brings the anti-codon to the coops on the MRNA, and with it the correct cymino acid which forms into a polypeptide chain. DNA codes for a specific protein needed in the organism. The mRNA Carefully copies the code and takes it to the vilossomes where profeins are made. Ribosomes are on the endoplasmic restaulum on the outside of the nucleus. The ribosomes read each three bases of the specific add and match them to its corresponding anti-cooden which tRNA. The HRNA loning a specific amino acid which the mRNA

ASSESSOR'S USE ONLY This page has been deliberately left blank.

The examination continues on the following page.

QUESTION TWO: ENVIRONMENTAL FACTORS AND GENE EXPRESSION

ASSESSOR'S USE ONLY

The honey bee (Apis mellifera) has two female phenotypes.

Female type	Larvae Diet	Adult phenotype	Genotype
Queen bee	royal jelly	increased ovary sizelarge body masslive for 2 years	the same
Worker bee	royal jelly for 3 days, then only pollen and honey	 infertile ovaries smaller body mass live for 3 – 6 weeks 	

www.britannica.com/media/ full/171791/141787

(a) Describe the term gene expression.

Which gene Shows in the pwintype.

(b) Explain why comparing worker and queen honey bee females is ideal for experiments on environmental factors and gene expression.

Their phenotypes are so different, but yet they have the same genotypes. Comparing environmental factors on the different phenotypes can result in large variations. Comparing gene expressions on the two also shows large variation on the same genetypes.

ASSESSOR'S USE ONLY

Discuss the effect the environment has on the expression of the phenotype in honey bee females.

In your answer include:

- a description of the environmental factor that affects honey bee phenotype
- using an example, an explanation of the difference between environmental factor and mutagen
- a discussion of how honey bee phenotype can change without changing the genotype
- a discussion of why the queen bee's phenotype is fully expressed, but the worker bee's phenotype is not.

The environmental factor that affects Phenotype is the availability of nutrients. Their diets one the only thing different that rescuts in different phenotypes for the worker bee and the Queen bee. Environmental factors ane factors do not permanently affect genotype Ovenote ÓY such as apting sunburth-directly affects phenote but will go away once environment Mutagens creete mutations which are permanent, radiation affecting gametes which are passed on to Ottspring. Mutagers have much larger, effects them environmental factors. can change without as the the genotype Queen and worker sufficient availability which affect the environment : S.

A - availability of nutrients.

L - light levels

There is more space for your answer to this question on the following page.

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Queen her phenotype is fully expressed because she has a full availability of nutrients due to her diet of voyal jelly. Rayal jelly is fed to newborn worker bees which is a sign it is full of vital vitamins and nutrients which help growth and strength. As the Queen bee Continues to grow and get stronger, she is able to live longer and produce offspring - fully express genetype phenotype. The worker bees do not have a supply of royal jelly to eat and therefore do not have a ready supply of good nutrients. The worker bees do not grow as much or get as strong as they could because they are not getting the right nutrients to fully express phenotype. As a result the worker bees not produce after 3-6 weeks and do

(a) Describe what a mutation is.

A mutation is a permanent change to the base sequence of a DNA molecule, or a phenotype. Mutations can be point (mutations (one base on a DNA strand) or block mutations (whole genes or parts of chramosomes are changed). H

Question Three continues on the following page.

(b) There are over 1000 mutations that can cause cystic fibrosis. A common mutation is a deletion mutation that results in the absence of one amino acid in the final protein. Another mutation is a substitution mutation that results in a different amino acid in the final protein.

Discuss how these two mutations affect the cystic fibrosis gene's final protein and resulting phenotype.

In your answer include:

- an explanation of why the deletion mutation causes one amino acid to be absent in the final protein, and how this affects protein folding
 - cin folding Chromosome 7

Cystic

fibrosis

ASSESSOR'S USE ONLY

- an explanation of why the substitution mutation causes a different amino acid to be present in the final protein, and how this affects protein folding
- a discussion of why the deletion mutation causes severe cystic fibrosis disease, whereas the substitution mutation causes milder cystic fibrosis disease.

causes there to Veletion mutation missing and therefore with there nitrogen base base too short at the end and the final leaving the is not coded for, be abser one amino acid short and the properly. Proteins told into specific shapes and it everything is different this affects the One armino acid ostitution mutation Whole protein. base with another. not Change an amino acid (due to deopneracy) results ma different other times: agd to be formed. a different amino acid is put into a polypepticle chain, protein will not be made or will not fold properly. Deletion mutation causes because whole amino

may not Stop' or it may stop randomly in the middle, both causing a protein to not be made correctly if at all.

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Substitution mutation causes milder cystic fibrosis. As only one amino acid is changed, there are still enough comino acids on the polypepticle chain to form the right protein, last this protein may not fold entirely correctly, or it. As the protein is still made, but slightly wrong, the cystic fibrosis disease is mild. On the other hand with a deletion frameshiff mutation, the protein is extract unfinished and therefore will not fold properly and can not make the correct protein at all, resulting in severe cystic fibrosis eliseare.

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QUESTION NUMBER	Extra paper if required. Write the question number(s) if applicable.	ASSESS USE OI
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MERIT exemplar for 91159 2016		Total score	Q 5	
Q	Grade score	Annotation		
1	M5 low	The student has good L2 knowledge of the structure of both DNA and RNA. The have described the process of protein synthesis and explained aspects of it, for example that the tRNA molecule will bring in a specific amino acid. More knowledge of the detail of the mRNA synthesis would make this a stronger merit.		
2	M5 low	The student has clear understanding of the difference between an environmental factor and a mutagen in relation to which alters the genotype. The aspect of the environment is correctly identified as diet and while the student knows this is what is having the effect they do not offer biological ideas as to how this could take place.		
3	M5 high	There is clear evidence that the student knows what a mutation is. They link both to base changes. They have knowledge of degeneracy, although they don't clearly describe the term. They link changes briefly to folding of proteins and the severity of CF.		,