

AQA, OCR, Edexcel

A Level

# A Level Biology

DNA Technology Answers

Name:

**M M E**

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Total Marks:

**M1.(a)** (i) 1. (Tumour suppressor) gene inactivated / not able to control / slow down cell division;

*Ignore: references to growth*

2. Rate of cell division too fast / out of control.

*1 and 2 Accept: mitosis*

*1 and 2 Reject: meiosis* 2

(ii) 1. (Genetic) code degenerate;

*Accept: codon for triplet*

*Accept description of degenerate code, e.g. another triplet codes for the same amino acid*

2. Mutation in intron.

*Accept: mutation in non-coding DNA* 1 max

(b) 1. Antibody has specific tertiary structure / binding site / variable region;

*Do not accept explanations involving undefined antigen*

2. Complementary (shape / fit) to receptor protein / GF / binds to receptor protein / to GF;

*Ignore: same shape as receptor protein / GF*

3. Prevents GF binding (to receptor).

3

[6]

**M2.** (a) (i) Sticky ends / description;

Reference to complementary base-pairing

2

(ii) Ligase;

1

(b) Carrier of DNA / gene; (*context of foreign DNA*)

Into cell / other organism / host;

2

(c) Act as marker gene;

Allows detection of cells containing plasmid / DNA;

2

[7]

**M3.** (a) No cadmium; Other conditions same as cadmium-treated group;

2

(b) (i) As a measure of the effect due to cadmium / to make a comparison;

1

(ii) Becoming more methylated;

*Ignore later slight decrease/no change*

1

- (iii) Production of more methyltransferase enzyme / increased activity of transferase;  
*Extra incorrect relevant information – cancel*  
1

- (c) RNA-polymerase could not bind (to DNA / to promoter); mRNA of p16 could not be made / no transcription of p16 gene;

2

- (d) Any four from: 1. Cadmium causes expression of methyltransferase gene / increased activity transferase (from 2 to 3 weeks in); 2. Methyl groups on to promoter / p16 gene / suppressor (gene); 3. (p16) normally suppresses tumour growth; 4. p16 protein / p16 expression falls after 4 weeks / after methylation; 5. Tumour formation occurs (after 10 weeks) after p16 falls / after suppressor gene activity falls;

4 max

[11]

- M4.(a) 1. Removes (main / largest) source of oestrogen / (different) mice produce different amounts of oestrogen;

*Accept: so oestrogen from ovaries not a confounding variable – idea of.*

2. (Allows) oestrogen to be controlled / oestrogen to be made by aromatase only / only oestrogen made in lungs to be involved.

*Reject: references to injection of aromatase.*

2

- (b) 1. (Anastrozole) prevents / reduces oestrogen production;

2. (Fulvestrant) stops remaining oestrogen binding / less oestrogen binds to receptors.

*Note: brackets around drug names.*

2

- c) (Yes for Group T)

1. Least tumours per animal (from fig. 1);

*Accept: 'mean values' for tumour area.*

2. Lowest (mean) tumour area / size (from fig. 2);

3. Lowest top of range;

(But)

4. Means (tumour area) are similar;

*Where candidates confuse range and standard deviation, do not give credit.*

5. Ranges overlap / share values so differences may not be real / treatments may be just effective in reducing tumour;

*Ignore significance*

6. Range affected by outliers / SD's would be better;

7. Done on mice / not done on women / humans;
8. Only 10 mice used per group / small sample size so may not be representative / reliable;
9. Might be side effects;
10. Only did for 15 weeks so maximum effect of drugs may not have been seen.

5 max

- (d) 1. Tumours may be different depths / area does not take depth into account / tumours are 3-D / are not 2-D;

*Neutral: different sizes*

*Accept: height / thickness for depth*

2. (Measure) tumour volume / mass / weight.

2

- (e) 1. Allows tumours to grow / develop / form;

*Neutral: gives drug more time to work.*

2. (So) can investigate treatment rather than prevention (of tumours) / when tumour / cancer is more advanced.

*Accept: to see whether it can destroy / treat / stop growth of a tumour (that already exists) / to allow / assess treatment of a tumour*

2

- (f) 1. Unethical (not to treat patients) / may increase probability of patients dying / getting more ill;

*Reject: references to giving people tumours*

2. Use normal cancer drugs / treatment.

*Accept: named type of cancer treatment, e.g. chemotherapy*

2

[15]

- M5.** (a) (i) plasmid;

1

- (ii) the bacteria divide / grow, producing many copies of desired gene / plasmid;

OR

the bacteria divide / grow to cover the agar;

1

- (iii) plant tissue that has antibiotic resistance survives; identifies plant tissue which has desired gene / plasmid;

2

- (iv) to clone plants / produce genetically identical plants with gene / characteristic; and produce large numbers / quickly;

2

- (b) (i) (*one reasonable suggestion*),

e.g. toxin present all the time;  
save costs of buying / application of spray;  
no spray drift onto other fields / insects;

1 max

- (ii) (*one reasonable suggestion*),  
e.g. killing of harmless / useful insects that feed on wild plants;  
damage to food chains starting with wild plants;

1 max

[8]

- M6.** (a) only small amounts obtained / PCR increases the amount / mass of DNA;  
so enough DNA available for genetic fingerprinting;

2

- (b) (i) to separate the two strands of the DNA /  
to break the hydrogen bonds;

(Reject "unzip")

1

- (ii) short lengths / fragments of DNA / nucleotides /  
single stranded DNA;

1

- (iii) to mark beginning and / or ends of the part of DNA needed /  
for attachment of enzymes or nucleotides / initiator /  
keeps strands apart;

1

- (iv) would not be denatured;  
must be heated to 95 °C / must withstand high temps;

2

- (c) 1 DNA extracted from sample;  
2 DNA cut / hydrolysed into segments using restriction  
endonucleases;  
3 must leave minisatellites / required core sequences intact;  
4 DNA fragments separated using electrophoresis;  
5 detail of process e.g. mixture put into wells on gel and electric  
current passed through;  
6 immerse gel in alkaline solution / two strands of DNA separated;  
7 Southern blotting / cover with nylon / absorbent paper (to absorb  
DNA);  
8 DNA fixed to nylon / membrane using uv light  
9 radioactive marker / probe added (which is picked up by required  
fragments) / complementary to minisatellites;  
10 (areas with probe) identified using X-ray film / autoradiography;

max 6

- (d) adult 3;  
this is only one which, (with number 1), can provide (all) the DNA  
fragments which children have / all bars match;

(Reject 'genes')

2

[15]

7. (a) a length of DNA;  
that codes for a single protein / polypeptide; 2
- (b) by heating;  
to break the H-bonds (between complementary bases); 2
- (c) (i) to allow the DNA polymerase to attach / start addition of  
nucleotides / mark start and end of sequence to be  
copied / prevents strands re-joining; 1
- (ii) because the sequences at the ends of the target sequence  
are different / one is at the beginning and one at the end; 1
- (d) 8; 1  
*accept 7*
- [7]**

- M8.** (a) 1. DNA is cut;  
2. using restriction enzyme;  
3. electrophoresis;  
4. separates according to length / mass / size;  
5. DNA made single-stranded;  
6. transfer to membrane / Southern blotting;  
7. apply probe;  
8. radioactive / single stranded / detected on film / fluorescent;  
9. reference to tandem repeats / VNTRs / minisatellites;  
10. pattern unique to every individual; 6 max
- (b) cells on toothbrush;  
DNA present in cell; 2
- (c) (i) toothbrush gives small sample of DNA / need more DNA  
for analysis;  
PCR gives many copies; 2
- (ii) uses heat; to separate strands;  
*OR*  
PCR replicates pieces of DNA;  
because DNA has been cut;  
*OR*  
primer added in PCR;  
to initiate replication 2 max

	(d) (i) PCR / amplification needed;	1	
	(ii) other DNA present; need to identify 'required' DNA from rest;	2	
			<b>[15]</b>
<b>M9.</b>	(a) Restriction (enzyme / endonuclease);	1	
	(b) Move towards anode / move because charged; Different rates of movement related to charge / size;	2	
	(c) (i) Piece of DNA; Single stranded; Complementary to / binds to known base sequence / gene;	max 2	
	(ii) DNA invisible on gel / membrane; Allows detection;	2	
			<b>[7]</b>
<b>M10.(a)</b>	1. (If injected into egg), gene gets into all / most of cells of silkworm; 2. So gets into cells that make silk.	2	
	(b) 1. Not all eggs will successfully take up the plasmid; 2. Silkworms that have taken up gene will glow.	2	
	(c) Promoter (region / gene).	1	
	(d) 1. So that protein can be harvested; 2. Fibres in other cells might cause harm.	2	
			<b>[7]</b>
<b>M11.</b>	(a) restriction (enzyme) / endonuclease / named example;	1	
	(b) unpaired bases / sticky ends / staggered; complementary / explained;	2	
	(c) <i>1 mark for each correct outcome</i> plasmid with foreign DNA joined in ring; ring with plasmid only; ring of foreign DNA only; <i>ignore linear structures</i>	3	
			<b>[6]</b>
<b>M12.</b>	(a) Will replace themselves / keep dividing / replicate;  Undifferentiated / can differentiate / develop into other cells / totipotent / multipotent / pluripotent;		

	<i>Accept tissues</i>	2
(b)	Reverse transcriptase; <i>Allow phonetic spelling</i>	1
(c)	(i) Alters base / nucleotide sequence / causes frame shift;  Different sequence of amino acids in polypeptide / protein / primary structure alters the tertiary structure; <i>Accept any reference, such as adding bases, to changing the base sequence of the gene. Reject deletion / substitution.</i> <i>Idea of sequence essential so not makes different amino acids.</i> <i>Accept answers involving stop / start codons and effect on protein.</i>	2
	(ii) Affects tumour suppressor gene;  Inactivates (tumour suppressor) gene;  Rate of cell division increased / tumour cells continue to divide; <i>Ignore answers relating to oncogenes. May gain third point.</i>	2 max
d)	Yes SCID patients unlikely to survive / quality of life poor unless treated; Cancer that develops is treatable / only affects 25% / five children;  No Risk of developing cancer is high / 25%; Cancer may recur / may not be treated successfully in future / only short time scale so more may develop cancer; <i>No mark for yes or no. Marks are for supporting argument based on biological reasoning.</i> <i>Accept any points</i>	2 max
		<b>[9]</b>
13.(a)	RNA polymerase; <i><u>D</u>NA polymerase is incorrect</i> <i>Ignore references to RNA dependent or DNA dependent</i> <i>Allow phonetic spelling</i>	1
(b)	(i) (Receptor / transcription factor) binds to promoter which stimulates RNA polymerase / enzyme X;  Transcribes gene / increase transcription;	2



- (ii) Other cells do not have the / oestrogen / ER $\alpha$  receptors;  
*But do not accept receptors in general.*

1

- c) Similar shape to oestrogen;

Binds receptor / prevents oestrogen binding;

Receptor not activated / will not attach to promoter / no transcription;

*Accept alternative*

*Complementary to oestrogen;*

*Binds to oestrogen;*

*Will not fit receptor;*

2 max

[6]

- M14.**(a) Restriction / endonuclease;

*Ignore specific names of restriction enzymes e.g.*

*EcoR1*

1

- (b) (i) 1. (Acts as a) marker gene to show that the (human) gene has been taken up / expressed;

*1. Accept: gene marker*

2. (Only) implant cells / embryos that show fluorescence / contain the jellyfish gene;

2

- (ii) 1. Factor IX present in / extracted from milk;

2. Gene only expressed in mammary glands / udder / gene not expressed elsewhere;

*2. Ignore references to milk*

*The 'only' aspect is important here.*

3. Do not need to kill sheep (to obtain Factor IX);

2 max

- (c) (i) 1. Mutation / nucleus / chromosomes / DNA may be damaged / disrupts genes;

*1. Neutral: cell may be damaged*

2. May interfere with proteins (produced) / gene expression / translation;

*Ignore references to hormone levels or time of implantation*

**OR**

3. Embryo / antigens foreign;

*3. Neutral: antigens change*

4. Embryo is rejected / attacked by immune system;

4. sNeed idea that the immune system is involved if mark point 3 has not been given

'Embryo foreign so rejected' = 2 marks

'Embryo rejected by immune system' = 1 mark

'Embryo is rejected' = 0 marks

2 max

- (ii) 1. Saves time / money for others;
2. Same work is not repeated / methods can be compared / improved / amended / same errors are not made;

2

[9]

**M15.(a)** 1. No effect at 25°C

*The question only refers to plants with GB*

1. Reject same mass

2. Keeps growing at 30°C and 35°C / up to 35°C (more than without GB);

3. Above 35°C, falls but grows more than plant without GB;

3. Accept at all temperatures above 25°C more growth than without GB

2 max

(b) (i) Significantly different / SEs do not overlap ;  
Accept converse without GB

1

ii) (As temperature increases,)

1. Enzyme activity reduced / (some) enzymes denatured;

2. Less photosynthesis, so fewer sugars formed;

3. Less respiration / less energy / ATP for growth;

4. Less energy for named function associated with growth

4. Eg mitosis, uptake of mineral ions

4

(c) 1. (Rubisco activase attaches to thylakoid and) this changes shape / tertiary structure (of enzyme) / blocks active site / changes active site;

*Note - question states enzyme stops working when it attaches to thylakoid, not before*

1. Accept rubisco in this context

2. (This) prevents substrate / RuBP entering active site / binding;

2. Accept prevents ES complex forming

2. Accept no longer complementary to substrate / RuBP

2

- (d) 1. GB prevents / reduces binding of rubiscoactivase to (thylakoid membrane);  
     1. *Accept enzyme instead of rubiscoactivase.*  
     *Accept rubisco*
2. (Prevents it) up to 35°C;
3. (So) rubiscoactivase / enzyme remains active;
4. (So) photosynthesis / light-independent stage still happens;  
     4. *Accept descriptions of light-independent stage*
5. Above 35°C, some binding still occurs but less than without GB, so less reduction in growth;
- 4 max

- (e) 1. Looked for information / journals, on crop plants that grow at high temperatures;  
     1. *"other research" is minimum accepted*  
     1. *Accept previous experiments research with temperature resistant crops*  
     *Ignore simple references to looking at previous studies / other plants - need to relate to this context*
2. (Crop plants cited in this research) contain / make GB;
3. So assumed making plants produce GB makes them resistant to high temperatures;
- 2 max

[15]

- M16.(a)** 1. Carriers are heterozygous / have one normal copy and one mutant copy of gene / have one recessive allele / don't have the condition;
2. Both have DNA that binds (about) half / 50% amount of probe (that non-carrier does);
3. Probe binds to dominant / healthy allele so only one copy of exon in their DNA / have one copy of gene without exon / base sequence for probe to bind to;  
     3. *Accept normal and gene*  
     3. *Accept have a deletion mutation*

3

- (b) 1. Introns not translated / not in mRNA / (exons) code for amino acids / introns do not code for amino acids;  
     1. *Accept not expressed*  
     1. *Accept polypeptide / protein for amino acids*
2. Mutations of these (exons) affect amino acid sequences (that produce) faulty protein / change tertiary structure of protein;  
     2. *Accept deletion leads to frameshift*  
     2. *In this context, accept affects protein made*

3. So important to know if parents' exons affected, rather than any other part of DNA / introns;

*Accept converse arguments involving - eg introns do not code for amino acids / proteins*

*Reject references to making amino acids, once*

3

- (c) 1. Restriction mapping / described;
2. DNA / base sequencing (of fragments) / description / name of method;

2

[8]

**M17.(a)** Cytosine with Guanine and (Adenine) with Uracil;  
*Ignore G, C and U*

1

- b) Two reasons, with suitable amplification;;

**Q**

Only infected cells have HIV protein on surface;

So carrier only attaches to / specific to these cells / siRNA can only enter these cells;

**OR**

siRNA (base sequence) complementary / specific to one mRNA;  
*Accept idea of specificity*

Only infected cells contain mRNA of HIV / this gene / stops translation of this gene / only binds to this mRNA / destroys this mRNA;

*Accept could not inhibit other / non-HIV mRNA*

4 max

- c) 1. Carrier binds to (protein on) HIV;  
*1. Accept references to HIV membrane*
2. Prevents HIV / it binding to (receptor on human) cell;  
*2. Reject references to binding to HIV protein on human cell*

2

[7]