

### **General Certificate of Education**

# Biology 5416

Specification B

BYB2 Genes and Genetic Engineering

## **Mark Scheme**

2007 examination - June series

Mark schemes are prepared by the Principal Examiner and considered, together with the relevant questions, by a panel of subject teachers. This mark scheme includes any amendments made at the standardisation meeting attended by all examiners and is the scheme which was used by them in this examination. The standardisation meeting ensures that the mark scheme covers the candidates' responses to questions and that every examiner understands and applies it in the same correct way. As preparation for the standardisation meeting each examiner analyses a number of candidates' scripts: alternative answers not already covered by the mark scheme are discussed at the meeting and legislated for. If, after this meeting, examiners encounter unusual answers which have not been discussed at the meeting they are required to refer these to the Principal Examiner.

It must be stressed that a mark scheme is a working document, in many cases further developed and expanded on the basis of candidates' reactions to a particular paper. Assumptions about future mark schemes on the basis of one year's document should be avoided; whilst the guiding principles of assessment remain constant, details will change, depending on the content of a particular examination paper.

Further copies of this Mark Scheme are available to download from the AQA Website: www.aqa.org.uk

Copyright © 2007 AQA and its licensors. All rights reserved.

#### COPYRIGHT

AQA retains the copyright on all its publications. However, registered centres for AQA are permitted to copy material from this booklet for their own internal use, with the following important exception: AQA cannot give permission to centres to photocopy any material that is acknowledged to a third party even for internal use within the centre.

Set and published by the Assessment and Qualifications Alliance.

#### Question 1 (a) (i) Stage A is anaphase; Chromatids/chromosomes moving apart/centomere divided; 2 (ii) Stage B is telophase; Chromosomes are uncoiling; 2 (Accept new nuclei forming) (b) Embryo split into separate cells; These (cells) are undifferentiated/totipotent; Each cell grows by mitosis (into new embryo/organism); 2 max Total 6 **Question 2** (a) Heat (DNA) to 95°C, to separate strands/break hydrogen bonds/denature; Cool to 40°C, to allow primers/nucleotides to bind; Primers provide starting point for copying/prevent strands rejoining; Heat to 70°C, optimum for polymerase/enzyme;

(b) Reason with explanation;;

(Accept Taq polymerase)

For example: Easy to insert genes into bacteria; Using a vector/plasmid/virus;

Bacteria reproduce rapidly; Producing many copies of (inserted) gene;

DNA polymerase then joins nucleotides together;

Bacteria can be grown on a large scale/ in industrial fermenters;	
To get a lot of the product of the gene;	2 max

Total 6

4 max

Question 3				
(a)	(i)	<u>Meiosis;</u>		
	(ii)	Reference to homologous/pairs of chromosomes/bivalents; One of each pair goes to each cell/gamete; (In second division) centromere divides/chromatids separate; (Accept crossing over/independent assortment for 1 mark)	3	max
(b)	(i)	F between 'gamete-producing structure and zygote;		
	(ii)	M between '2N' and spores;	2	
(C)	Advan	tage and explanation;;		
	For ex Many s So mo	ample: spores released nearby; re of nutrient source used/used more rapidly;		
	Produc So all v	ces genetically identical fungi; well suited to nutrient source/ environment;	2	max
			Total 7	
Quest	ion 4			
(a)	Plasmi Restric (Comp Use of	id cut; ction enzyme/endonuclease; lementary) sticky ends (however formed); ligase;	3	max
(b)	(Bee) ( So, the	gene will get into/expressed in all cells (of the mosquito); e gene gets passed to future generations;	2	
(c)	To ide So, on Identify	ntify mosquitoes which had taken up the gene; ly these are used; y by looking for those that produce green light;	2	max
(d)	Could Islands If mala malaria	save birds from malaria, so reversing harm done by humans; s isolated, so fewer problems if trial goes wrong; iria parasite becomes resistant to bee protein, resistance not in huma a strain;	an	
	Resist	ance will spread through the mosquito population;	1	max
			Total 8	

#### Question 5

(a)	More radiation, more non-mobile sperm; Radioactive material source of high energy/ionising radiation/ high energy particles/named; Which cause mutations (and abnormal sperm); Detail of how mutation affects sperm movement:	3 max
(b)	Both reduced in high radiation area/correct use of figures for both; Antioxidant 1 most affected;	2
(c)	Reference to <u>sperm</u> unable to fertilise (eggs); Damage to <u>egg</u> DNA (because of lower antioxidants); So non-viable egg/zygote/embryo/offspring; (Mutations) produces individuals with (new) harmful characteristics;	3 max
		Total 8

#### Question 6

(a)	Base sequence different/named mutation; Comparison to functional allele/gene/DNA/mRNA; Leads to different sequence of amino acids; So enzyme/protein with different/non-functional shape;	3 max
(b)	Gene inserted into vector/virus/liposomes; Inhaled/sprayed (into lungs); Vector carries gene into (epithelial) cells; ( <i>Accept plasmids as a vector</i> )	3
(c)	They will have one functional allele/gene; This produces (enough) functional protein;	2

#### Total 8

### Question 7

Quest	tion 7		
(a)	1	Sequence of bases in gene/on DNA determine amino acid sequence;	
	2	Transcription to form mRNA;	
	3	Codons on mRNA;	
	4	Reference to base-pairing;	
	5	A to U, T to A and C to G;	
	6	Specific tRNA for each amino acid;	
	7	Has anticodon to bind to mRNA codon;	
	8	At the ribosome;	
	9	Role of sites on ribosomes in translation.	6 max
(b)	(i)	X is phosphate and Y is pentose/deoxyribose;	
	(ii)	Hydrogen;	2

 (c) (i) AZT binds to adenine/ A (on single DNA strand); But lacks phosphates/ OH; Unable to bind to another nucleotide; So (AZT) enzyme unable to form new strand/DNA; 2 max

Total 10