MARK SCHEME for the May/June 2012 question paper

for the guidance of teachers

9790 BIOLOGY

9790/04

Paper 4 (Practical), maximum raw mark 70

This mark scheme is published as an aid to teachers and candidates, to indicate the requirements of the examination. It shows the basis on which Examiners were instructed to award marks. It does not indicate the details of the discussions that took place at an Examiners' meeting before marking began, which would have considered the acceptability of alternative answers.

Mark schemes must be read in conjunction with the question papers and the report on the examination.

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Skill	Approximate total marks	Breakdown of the marks	Q.1	Q.2	Q.3	Q.4	Q. 5	Total
Manipulation, measurement and	24	Successful collection of data and observations (MMO collection)	3	11				22
observation		Decisions about measurements or observations (MMO decisions)	6	2				
Presentation of data and	13	Recording data and observations (PDO recording)	2					15
observations		Display of calculation and reasoning (PDO display)	2	4				
		Data layout (PDO layout)	1	3			3	
Planning	16	Defining the problem (P)			6			16
		Methods (M)			10			
Analysis, conclusions and	17	Interpretation of data or observations and identifying sources of error (D)				8	2	17
evaluation		Suggesting improvements and evaluation (E)				2		
		Conclusion (C)					5	
Total	70		14	20	16	10	10	70

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Section A

Question	Sections	Learning outcomes	Indicative material	
			there are 16 marking points available – mark to max 14	
1	MMO decisions	 decide how many tests, measurements or observations to perform make measurements or observations that span the largest possible range within the limits either of the equipment provided or of the instructions given 	 evidence that each solution tested for reducing sugars; evidence that each solution tested for non-reducing sugars; an explanation that reducing sugar test is followed by the non-reducing sugar test on all samples; <i>results as</i> final colour/colour change/time for first appearance of green/precipitate; different colours for any one sample following acid hydrolysis and neutralisation; repeats; 	[6]
	MMO collection	 set up apparatus correctly work out what to do from outline instructions given in the form of written instructions or diagrams use their apparatus to collect an appropriate quantity of data or observations, including subtle differences in colour or other properties of materials 	 7 correct results recorded for solutions C1 to C5; 8 colours/precipitate, recorded unambiguously; 9 no, change in colour/precipitate, with C1 in at least one test; 	[3]

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Question	Sections	Learning outcomes	Indi	cative material	Mark
1	PDO recording PDO layout	 present numerical data, values or observations in a single table of results draw up the table before taking readings/making observations, so that candidates can record directly into the table, to avoid the need to copy up their results include in the table of results, if necessary, columns for raw data, for calculated values and for deductions use column headings that include the quantity and the unit (as appropriate) and that conform to accepted scientific conventions choose a suitable and clear method of presenting the data, e.g. tabulations, chart, graph, drawing or mixture of methods of presentation 	10 11 12 13 14	data for, colours/precipitate/times, and any other observations recorded as a single table ; no evidence of copying up their results ; informative column headings ; e.g. test solution/sample, final colour, precipitate, sugar content, degree of stickiness results linked correctly to relative reducing sugar content ; A ecf more intense colour in C3 with non-reducing sugar test compared with reducing sugar test ;	[5]
	PDO display	 show their working in calculations, and the key steps in their reasoning 	15 16	allow ecf from results deductions about sugar content of fibres ; any deductions about 'stickiness' ;	[2]
					[Total: 14]

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Question	Sections	Learning outcomes	Indicative material	Mark
2 (a)	MMO collection	 set up apparatus correctly use their apparatus to collect an appropriate quantity of data or observations, including subtle differences in colour or other properties of materials 	 correct outline of five seminiferous tubules of appropriate relative sizes with complete lines; layers within the tubule and outline of lumen shown; e.g. connective tissue, germinal epithelium, cells undergoin meiosis, spermatozoa interstitial tissue/cells of Leydig/connective tissue; capillaries/blood vessels; 	
			 <i>labels</i> seminiferous tubule(s); interstitial tissue/cells of Leydig/blood vessel/capillary; two regions within tubule; <i>e.g.</i> connective tissue/germinal epithelium, lumen/sperm 	[6 max]
(b)	MMO collection PDO display PDO layout	 make measurements using millimetre scales, graticules, protractors, stopwatches, balances, measuring cylinders, syringes, thermometers, and other common laboratory apparatus show their working in calculations, and the key steps in their reasoning choose a suitable and clear method of presenting the data, e.g. tabulations, chart, graph, drawing or mixture of methods of presentation 	 measurements of seminiferous tubules given in, graticule units/μm, within acceptable range; calibration shown (within appropriate range); diameters converted to, micrometres/millimetres, correctly; A standard form mean calculated ; standard deviation calculated ;	[1] [4] [2]

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(c)	MMO collectio	on	 use the an ap or observed subtle other 	neir apparatus to collect propriate quantity of data servations, including e differences in colour or properties of materials	1 2 3 4 5 6 7	one Sertoli cell draw large nucleus ; nucleolus ; spermatozoa attache correct shape of hea germinal epithelium cells in stages of div Sertoli cell ;	n, correct relatived to distal surfactorial surfactorial surfactorial surfactorial surfactorial sperm ; sing shown on	ve width and le ace ; lateral surfaces	ngth ; s of [max 4]]
	MMO decisio	ns	 decid meas obser make obser larges the lir equip instru 	e how many tests, urements or vations to perform measurements or vations that span the st possible range within mits either of the ment provided or of the ctions given	8 9 10 11 12	<i>labels</i> sperm(atozoa) ; named part of sperm Sertoli cell + nucleus germinal epithelium spermatogonia/sper	n ; s/nucleolus/cyto ; matocytes/speri	plasm ; matids ;	[max 2]]
	PDO layout		 choose method e.g. ta drawi of pression 	se a suitable and clear od of presenting the data, abulations, chart, graph, ng or mixture of methods esentation	13	cells drawn with clea	ar, complete line	es;	[1]	
						·			[Total: 20)]

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Section B

Planning

P = defining the problem

M = methods

Analysis, conclusions and evaluation

D = interpretation of data or observations and identifying sources of error

C = drawing conclusions

E = suggesting improvements and evaluation

Question 3

Sections	Learning outcomes	Expected answer	Mark
P defining the problem	 identify the dependent and independent variables in the experiment or investigation express the aim in terms of a prediction or hypothesis, and express this in words and in the form of a predicted graph identify the variables that are to be controlled 	 Hypothesis or prediction ; e.g. rate of hydrolysis is faster using free enzyme/quantity of urea hydrolysed over time is greater with free enzyme/immobilised urease catalyses reaction over much longer period of time A ora/null hypothesis Theory to support candidate's hypothesis or prediction ; e.g. refs to accessible active sites/diffusion of substrate into alginate beads/stability of enzyme in alginate beads Outline of strategy ; e.g. method of following the reaction taking samples at intervals and calculating the initial rate Justification/evaluation, of strategy ; e.g. can only alter concentration of immobilised enzyme by changing number of beads/limitations of colour comparison these could be awarded at the end of the plan Method of determining, pH / (the concentration) of ammonium carbonate, at intervals ; e.g. use of pH indicator, to follow colour change At least two control variables ; e.g. temperature, concentration of urea solution, volumes used, number of beads Risk assessment ; ref to hazard <u>and</u> precaution 	[max 6]

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Pmethods		Pre–U describe the method to be used to vary the independent variable, and the means that they will propose to ensure that they have measured its values accurately describe how the dependent variable is to be measured describe how each of the other key variables is to be controlled explain how any control experiments will be used to verify that it is the independent variable that is affecting the dependent variable and not some other factor describe the arrangement of apparatus and the steps in the procedure to be followed suggest appropriate volumes and concentrations of reagents, and explain how different concentrations of reagents, and explain how different concentrations would be prepared assess the risks of their proposed methods describe precautions that should be taken to keep risks to a minimum draw up tables for data that they might wish to record describe how the data might be used in order to reach a conclusion	 May 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 	y/June 2012 use range of concentration use range of concentration to find suitable concentration comparison dilution table(s) included; method to ensure concent in reaction mixtures is the free and immobilised enzy urea solution mixed with p equilibration in water bath mixing, urease/beads, and time = 0; staggered start; samples taken at stated in uncertainty/precision, of re plot results and take grad rate; colour standard set up at time taken (t) to reach coll recorded; rate = 1/t; A 1000/t, etc. colour change followed in repeats/replicates (calcula calculate, standard deviate error; ref to use of <i>t</i> -test to see if significantly different; plot results on appropriated line);	9790 ns of urea ; ns of urease ; <i>tions to make</i> A ratios tration of urease same for both yme ; >H indicator ; ; d urea solution at ntervals ; esults ; ient to give initial known pH ; our standard colorimeter ; ate means) ; ion/standard f rates are e graph (bar or	
						[Total:16]

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Question	Sections	Learning outcome	Expected answer	Mark
4 (a)	ACE interpretation	 describe the patterns and trends shown by tables and graphs describe and summarise the key points of a set of observations 	 general statement, e.g. more males involved, more successful breeding; mean mass of nestlings at day 6 increases the more adults there are to feed them; mean mass increases if males are full-time, not part-time; similar relationships with deaths of nestlings; comparative data quote; ref to any result thought anomalous; 	[max 3]
(b)	ACE interpretation	• use appropriate statistical tests to assess the variability of data or the statistical differences between samples	1 v 3 = 57, not significant/ns ; 4 v 5 = 62, <i>p</i> <0.01 ;	[2]
(c)	ACE evaluation	• use these evaluations and provided information to make informed judgements on the confidence with which conclusions may be drawn	used to show significance or not between similar data sets ; compares mean values ; <i>t</i> -test takes into account the differences in number of nests for each breeding strategy ; <i>t</i> -test may be used with small samples ;	[max 2]
(d)	ACE interpretation	• identify the most significant sources of error in an experiment	 variation in number of nests studied/AW; e.g. 7 v 45; 7 is too small for reliable use of <i>t</i>- test; nestling mass at day 6 may not reflect mass when they, leave the nest/first fly; may not be significant factor in deciding how many survive; mass/number, are not the only indicators of breeding success; another parameter; e.g. number that survive to breed <i>idea of</i> more than one feeding strategy within each reproductive strategy; AVP; ref to SD/only one location 	[max 3]
				[Total:10]

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Question	Sections	Learning outcon	me Expected answer		Mark
5 (a)(i)	ACE conclusion	 make detailed scient explanations of the d and of their conclusio drawing on the skill, knowledge and understanding that th have gained from the studies of the Pre-U syllabus 	ific ata ons, ney eir	cannot make, essential compound(s)/amino acid(s) /histidine ; no enzyme(s) present (for making histidine) ; mutation in gene(s) ;	[max 2]
(ii)	ACE conclusion	make detailed scientific explanations of the data and of their conclusions, drawing on the skill, knowledge and understanding that they have gained from their studies of the Pre-U syllabus		resistant to ampicillin/AW ; detail of mechanism of resistance ; otherwise wild type/AW ; has enzymes for synthesis of tryptophan, histidine and, lactase/ β galactosidase ;	[max 2]
(b)(i)	PDO layout	 select which variable(s) to plot and plot appropriately on clearly labelled <i>x</i>- and <i>y</i>- axes plot all points or bars to an appropriate accuracy follow the IOB recommendations for putting lines on graphs 		<pre>x-axis = time, y-axis = number of colonies, sensible scales and axes labelled appropriately with unit for time; points plotted correctly; curves/straight lines;</pre>	[3]
(ii)	ACE interpretation	 find an unknown value by using co-ordinates or axis intercepts on a graph 		values for first entry into recipient cells read from <i>x</i> -axis correctly (± one small square); genes identified correctly; <i>any four cells correct</i> = 1 <i>mark</i>	[2]
	medium 2 3 4 5	gene time of amp / AW trp / AW his / AW lactase / AW		of gene transfer to recipient cells / min 27 8 17 36	

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(c) ACE conclusions • draw conclusions from an investigati or from interpretati of observations, da and calculated valu		 Pre–U – May/June 2 draw conclusions from an investigation or from interpretations of observations, data and calculated values, 	2012 9790 04 genes mapped in correct order on the chromosome ; apply, mirror image/ecf apply, mirror image/ecf		04
		providing a detailed description of the key features of the observations, data and analyses, and considering whether experimental data support a given hypothesis	lac	try his amp	[1]
					[Total:10]