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UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS

GCE Advanced Subsidiary Level and GCE Advanced Level

MARK SCHEME for the May/June 2011 question paper for the guidance of teachers

9700 BIOLOGY

9700/52

Paper 5 (Planning, Analysis and Evaluation), maximum raw mark 30

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.

• Cambridge will not enter into discussions or correspondence in connection with these mark schemes.

Cambridge is publishing the mark schemes for the May/June 2011 question papers for most IGCSE, GCE Advanced Level and Advanced Subsidiary Level syllabuses and some Ordinary Level syllabuses.

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	0	GCE AS/A LEVEL – May/June 2011	9700	000
				S
Mark scher	ne abbreviation:	S:		1774
•	separates mark	king points		O.
1	alternative answ	wers for the same point		8
R	reject	·		26
Α	accept (for ans	wers correctly cued by the question, or by	/ extra guidance)	· Ox
AW		ding (where responses vary more than us		7
<u>underline</u>		ven must be used by candidate (grammati		

Mark scheme abbreviations:

max indicates the maximum number of marks that can be given

or reverse argument ora

marking point (with relevant number) mp

error carried forward ecf

ignore

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Page 3	Mark Scheme: Teachers' version GCE AS/A LEVEL – May/June 2011	Syllabus 9700	Pape 52	er dange.
Question	Expected answer		Ext	tra guidance
1. (a)	 8 of: independent variable: ref. to making a range of 0.2, 0.4, 0.6, 0.8, 1.0 mol solution / making separate solutions of 0.2, 0.4, 0.6 from sucrose and water; ref. to using distilled / deionised water (for making 3. ref. to leaving plant tissue for suitable time – minimal dependent variable: ref. to a suitable method of timing the movement of the solution drop); ref. to marking the centre / start point of the solution drop); ref. to how the drop is released; standardising variables (max 3): ref. to using same volume of each solution used for timing ref. to same volume of each solution used for timing ref. to using same number / mass / volume of tissued in the solution ref. to suitable method of keeping constant temperents. ref. to same time for soaking tissue discs; ref. to low risk investigation / any suitable safety pressure for the solution in the solution of the solution in the solution in	dilutions); num of 20 min f the drop; n (to measure king; ng the drop; ne; ature;	3. 4. 5. 6.	allow a general statement of making 5 (min) solutions from 0-1 mol dm³ allow any volumes in correct proportions for making sucrose solutions do not allow if refer to serial dilutions unless it would give the concs. stated by the candidate ignore ref. to 0.0 as a sucrose solution allow in terms of 'long enough for osmotic changes to occur' Ignore keeping in water/solution before using allow stop clock / stop watch / timer allow the idea of using a ruler / graduated test tube e.g. keeping the drop as small as possible / care in releasing the drop ignore same volume 7 and 8. needs to clear what the solution is being used for to award the mark allow surface (area) Ignore size / amount e.g. water bath / incubator . allow room temp. do not allow air conditioning allow 'fixed time' could be subsumed in 3 e.g. cutting away from hands / cutting on a tile, allergy to plants and wearing gloves ignore water and electrics

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	reliability: 13. ref. to at least 2 / several / many replicates and a ref. to increasing number of intermediates / repeat values close to water potential;		Paper 52 13. allow reference to anomalous results 14. allow figures (e.g.0.6 (mol dm ⁻³) to 0.2 (mol dm ⁻³)) allow mp1 if not already given and 5 dilutions are made. [8]
(b) (i)	mm s ⁻¹ / mm / s(ec) / mm per s(ec) / millimetres per se	cond ;	allow cm instead of mm for all versions of the units do not allow min(utes) if several are given, all must be correct [1]
(ii)	(draw (best fit) line through the points to) find intersect the concentration (of sucrose) at which there is little / r drop; (this sucrose) concentration / solution is the water pot of the tissue / cells;	no movement of th	
(c) (i)	independent:concentration / molarity of the sucrose (so dependent: direction and rate of drop / dye moment;		do not allow amount of sucrose / sugar [2]
(ii)	2 of: size / volume of droplet; finding the centre position of the solution or position of measurement; ref. to tissue varying as cutting may not be exact; ref. to source of tissue being different; ref. to removal of pipette;	drop by eye / rule	ignore temperature and pressure ignore number of drops ref. to parallax error must be in the correct context of finding the position of the drop allow as surface (area) / mass / volume ignore size / amount e.g. from different storage organs e.g. being careful not to mix the solution / moving to the side of the tube [2]
(iii)	drop: idea of: larger drops would move at different rate from give false readings); centre of solution: idea of: drops have less / more distance to travel so m inaccurate;	·	allow the idea of introducing variables that cannot be measured and this reducing accuracy allow any ref. to under or overestimating related to a specific variable allow ecf for a correct explanation, related to the investigation, for a variable that could have been controlled equitemperature

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	surface / mass / volume / source of tissue: will change the volume of water movement / density of	f the solution ;	temperature of speed of drop different volume	of diffusion / osr changes kinetic e changes mes of bathing s s changes the wa	energy so olution /	[1]
(d) (i)	0.2 molar – turgid and 0.8 molar – (almost) plasmolysed / flaccid;		turgidity	tions of plasmoly reference to size as of cells		[1]
(ii)	correct ref. to a water / solute potential gradient of either cell ;			water potentia	al	
	correct ref. to the direction of water movement for either	er cell ;		cells/tissue	solution	
			0.2 mol dm ⁻³	lower / more negative / hypertonic	higher / less negative / hypotonic	
			0.8 mol dm ⁻³	higher / less negative / hypotonic	lower / more negative / hypertonic	
			correct. allow ecf if an	re described, bo		
			cells are reve	rsed		[2]
					Total:	[19]

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					Can
2 (a) (i)	to stimulate the growth / development of follicles (cor	taining oocytes);	allow red do not a do not a	creases follicles / oocytes available f. to injection is only possible in vivo allow causes / increases ovulation allow increases the production of / oocytes	[1]
(ii)	ref. to idea of all oocytes starting at the same point fo	r investigation ;	e.g. ood meiosis	ytes are all at the same stage of	[1]
(b)	the closeness of the sample mean to the population of the estimate of the mean;	nean / the reliability	calculate do not a	f. to a measure of the accuracy of the ed mean value allow spread of values around the general references to reliability	[1]
(c) (i)	idea of there is no (significant) difference in the stimulation of meiosis by FF-MAS compared to other compounds;			do not allow the idea of 'does not stimulate meiosis'	
(ii)	(student) <i>t</i> -test; comparing means of (two) populations/ data has a no data is continuous;	ormal distribution /	do not a	Illow 'it is a continuous variable'	[2]
(iii)	1 of: the activator / FF-MAS is causing a change in the sti there (is less than) 0.05 / 5% chance that the differen was caused by chance;		l) allow 'th that the other tha	allow 'the null hypothesis is rejected' here is (more than) 0.95 / 95% chance difference is due to other factors an chance. is not due to chance'	[1]
(d) (i)	No, the (known) LXR alpha receptor activators do not stir	nulate meiosis ;	no without credit e.g. becall the o	ause none of the others are doing	[1]

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(ii) 3 1 2 3 4 5 6	 the stimulation of meiosis; FF-MAS may activate a different receptor / mech not known; none of the other activators / named activator tes meiosis; 22R-HC at 7.0 μmol dm⁻³ may inhibit meiosis; 	more than) doubles	1. 3. 4.	receptor	

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