## 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.

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Page 2	Mark Scheme: Teachers' version	Syllabus	Paper
	GCE AS/A LEVEL – May/June 2011	9700	52

Mark scheme abbreviations:

- ; separates marking points
- *I* alternative answers for the same point
- R reject
- A accept (for answers correctly cued by the question, or by extra guidance)
- **AW** alternative wording (where responses vary more than usual)
- **<u>underline</u>** actual word given must be used by candidate (grammatical variants excepted)
- max indicates the maximum number of marks that can be given
- ora or reverse argument
- **mp** marking point (with relevant number)
- ecf error carried forward
- I ignore

Page 3	ge 3 Mark Scheme: Teachers' version		Paper
	GCE AS/A LEVEL – May/June 2011	9700	52

Expected answer	Extra guidance	Mark
<ul> <li>8 of: <i>independent variable:</i></li> <li>1. ref. to <u>making</u> a range of 0.2, 0.4, 0.6, 0.8, 1.0 mol dm<sup>-3</sup> sucrose solution / making separate solutions of 0.2, 0.4, 0.6, 0.8, 1.0 mol dm<sup>-3</sup> from sucrose and water ;</li> </ul>	1. allow a general statement of making 5 (min) solutions from 0-1 mol dm <sup>-3</sup> 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	
2 ref to using distilled / deionised water (for making dilutions):		
<ol> <li>ref. to leaving plant tissue for suitable time – minimum of 20 min</li> </ol>	<ol> <li>allow in terms of 'long enough for osmotic changes to occur' Ignore keeping in water/solution before</li> </ol>	
dependent variable:	using	
4. ref. to a suitable method of timing the movement of the drop ;	4. allow stop clock / stop watch / timer	
<ol> <li>ref. to marking the centre / start point of the solution (to measure drop);</li> </ol>	<ol> <li>allow the idea of using a ruler / graduated test tube</li> </ol>	
6. ref. to how the drop is released ;	6. e.g. keeping the drop as small as possible / care in releasing the drop	
standardising variables (max 3):	ignore same volume	
7. ref. to using same volume of each solution for soaking ;	7 and 8. needs to clear what the solution	
8. ref. to same volume of each solution used for timing the drop ;	is being used for to award the mark	
9. ref. to using same number / mass / volume of tissue ;	9. allow surface (area) Ignore size / amount	
10. ref. to suitable method of keeping constant temperature ;	10. e.g. water bath / incubator . allow room temp.	
	do not allow air conditioning	
11. ref. to same time for soaking tissue discs ;	11. allow 'fixed time' could be subsumed in 3	
safety:		
12. ref. to low risk investigation / any suitable safety precaution ;	<ul><li>12. e.g. cutting away from hands / cutting on a tile, allergy to plants and wearing gloves ignore water and electrics</li></ul>	
	<ul> <li>8 of: independent variable:</li> <li>1. ref. to making a range of 0.2, 0.4, 0.6, 0.8, 1.0 mol dm<sup>-3</sup> sucrose solution / making separate solutions of 0.2, 0.4, 0.6, 0.8, 1.0 mol dm<sup>-3</sup> from sucrose and water;</li> <li>2. ref. to using distilled / deionised water (for making dilutions);</li> <li>3. ref. to leaving plant tissue for suitable time – minimum of 20 min</li> <li><i>dependent variable:</i></li> <li>4. ref. to a suitable method of timing the movement of the drop;</li> <li>5. ref. to marking the centre / start point of the solution (to measure drop);</li> <li>6. ref. to how the drop is released;</li> <li><i>standardising variables (max 3):</i></li> <li>7. ref. to using same volume of each solution for soaking;</li> <li>8. ref. to same volume of each solution used for timing the drop;</li> <li>9. ref. to using same number / mass / volume of tissue;</li> <li>10. ref. to same time for soaking tissue discs;</li> <li><i>safety:</i></li> </ul>	<ul> <li>8 of: <i>independent variable:</i></li> <li>1. ref. to making a range of 0.2, 0.4, 0.6, 0.8, 1.0 mol dm<sup>3</sup> sucrose solution / making separate solutions of 0.2, 0.4, 0.6, 0.8, 1.0 mol dm<sup>3</sup> from sucrose and water ;</li> <li>2. ref. to using distilled / deionised water (for making dilutions);</li> <li>3. ref. to leaving plant tissue for suitable time – minimum of 20 min <i>dependent variable:</i></li> <li>4. ref. to a suitable method of timing the movement of the drop ;</li> <li>5. ref. to marking the centre / start point of the solution (to measure <i>drop</i>);</li> <li>6. ref. to how the drop is released ;</li> <li><i>standardising variables (max 3):</i></li> <li>7. ref. to using same volume of each solution used for timing the drop ;</li> <li>9. ref. to same volume of each solution used for timing the drop ;</li> <li>9. ref. to same volume of each solution used for timing the drop ;</li> <li>9. ref. to same volume of each solution used for timing the drop ;</li> <li>9. ref. to same volume of each solution used for timing the drop ;</li> <li>9. ref. to same time for soaking tissue discs ;</li> <li>11. ref. to same time for soaking tissue discs ;</li> <li>12. ref. to low risk investigation / any suitable safety precaution ;</li> <li>12. ref. to low risk investigation / any suitable safety precaution ;</li> <li>12. e.g. cutting away from hands / cutting on a tile, alleryy to plants and wearing gloves</li> </ul>

Page 4	Mark Scheme: Teachers' version	Syllabus	Paper	]	
	GCE AS/A LEVEL – May/June 2011	9700	52	]	
	<ul> <li>reliability:</li> <li>13. ref. to at least 2 / several / many replicates and a r</li> <li>14. ref. to increasing number of intermediates / repeat values close to water potential ;</li> </ul>		14. allov (mol allov	w reference to anomalous results w figures (e.g.0.6 (mol dm <sup>-3</sup> ) to 0.2 l dm <sup>-3</sup> )) w mp1 if not already given and 5 ions are made.	[8]
(b) (i)	mm s <sup>-1</sup> / mm / s(ec) / mm per s(ec) / millimetres per se	cond ;	the units do not al	n instead of mm for all versions of llow min(utes) l 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 pote of the tissue / cells ;	o movement of t	he ignore ui allow - w	ine is drawn in Fig. 1.2 nits / ref. to rates vater potential of the tissues / cells is e as that of this (sucrose) ration	[2]
(c) (i)	<i>independent</i> :concentration / molarity of the sucrose (so <i>dependent</i> : direction <b>and</b> rate of drop / dye moment ;	olutions) ;	do not al	llow 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 / rul	er ref. to pa context o allow as ignore si e.g. from e.g. bein	emperature and pressure umber of drops arallax error must be in the correct of finding the position of the drop surface (area) / mass / volume ize / amount n different storage organs og careful not to mix the solution / to the side of the tube	[2]
(iii)	<i>drop</i> : idea of: larger drops would move at different rate from give false readings) ; <i>centre of solution</i> : idea of: drops have less / more distance to travel so m inaccurate ;	·	cannot b accuracy allow any related to allow ect the invest	e idea of introducing variables that be measured and this reducing y ref. to under or overestimating o a specific variable f for a correct explanation, related to stigation, for a variable that could en controlled e.g. temperature	

Page 5	Mark Scheme: Teachers' version	Syllabus	Paper					
	GCE AS/A LEVEL – May/June 2011	9700	52					
	surface / mass / volume / source of tissue: will change the volume of water movement / density of	of the solution ;	changes rate of diffusion / osmosis temperature changes kinetic energy so speed of drop changes different volumes of bathing solution / different times changes the water movement / density		f the solution ; speed of drop changes different volumes of bathing solution / different times changes the water mov		erature changes kinetic energy so d of drop changes ent volumes of bathing solution / ent times changes the water movement	
(d) (i)	0.2 molar – turgid <b>and</b> 0.8 molar – (almost) plasmolysed / flaccid ;		turgidity	reference to size		[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 eith	correct ref. to the direction of water movement for either cell ;		cells/tissue	solution			
			negative / negative /	higher / less negative / hypotonic				
			0.8 mol dm <sup>-3</sup>	higher / less negative / hypotonic	lower / more negative / hypertonic			
			correct.	are described, bo nswer in (i) the ch ersed		[2]		
					Total:	[19]		

Page 6	Mark Scheme: Teachers' version	Syllabus	Paper
	GCE AS/A LEVEL – May/June 2011	9700	52

2 (a) (i)	to stimulate the growth / development of follicles (containing oocytes);	allow increases follicles / oocytes available allow ref. to injection is only possible in vivo do not allow causes / increases ovulation do not allow increases the production of follicles / oocytes	[1]
(ii)	ref. to idea of all oocytes starting at the same point for investigation ;	e.g. oocytes are all at the same stage of meiosis	[1]
(b)	the closeness of the sample mean to the population mean / the reliability of the estimate of the mean ;	allow ref. to a measure of the accuracy of the calculated mean value do not allow spread of values around the mean or 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'	[1]
(ii)	(student) <i>t</i> -test ; comparing means of (two) populations/ data has a normal distribution / data is continuous ;	do not allow 'it is a continuous variable'	[2]
(iii)	1 of: the activator / FF-MAS is causing a change in the stimulation of meiosis ; there (is less than) 0.05 / 5% chance that the difference (from the control) was caused by chance ;	do not allow 'the null hypothesis is rejected' allow 'there is (more than) 0.95 / 95% chance that the difference is due to other factors other than chance. allow 'it is not due to chance'	[1]
(d) (i)	No, the (known) LXR alpha receptor activators do not stimulate meiosis ;	must have a correct reason for their answer, no without an explanation does not gain credit e.g. because the % FF- MAS is higher than all the others e.g. because none of the others are doing any stimulation	[1]

Page 7	Mark Scheme: Teachers' version	Syllabus	Paper
	GCE AS/A LEVEL – May/June 2011	9700	52

(ii)	<ol> <li>3 of :         <ol> <li>FF-MAS above 0.07 at 0.7/at 7.0 μmol dm<sup>-3</sup>/stimulates meiosis ;</li> <li>increasing the concentration of FF-MAS by x10 (more than) doubles the stimulation of meiosis ;</li> </ol> </li> <li>FF-MAS may activate a different receptor / mechanism of action is not known ;</li> </ol>	1.	allow general statement 'as the conc. of FF-MAS increases so does stimulation of meiosis' or 'FF-MAS has the greatest effect on meiosis' allow FF-MAS is not stimulating the LXR receptor	
	<ol> <li>none of the other activators / named activator tested stimulate meiosis ;</li> </ol>	4.	allow in the context of 'little effect as the error bars overlap'	
	5. 22R-HC at 7.0 $\mu$ mol dm <sup>-3</sup> may inhibit meiosis ;		do not allow – the other activators stimulate by the same amount	
	6. ref. to the reliability of the data sets ;	6.	e.g. 0.7 $\mu$ mol dm <sup>-3</sup> data set / 25 HC is most reliable as SM has smallest values e.g. FF-MAS data set is least reliable as SM has largest values	[3]
			Total:	[11]