## UNIVERSITY OF CAMBRIDGE INTERNATIONAL EXAMINATIONS GCE Advanced Subsidiary Level and GCE Advanced Level

## MARK SCHEME for the May/June 2008 question paper

## 9700 BIOLOGY

9700/32

Paper 32 (Advanced Practical Skills 2), maximum raw mark 40

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.

All Examiners are instructed that alternative correct answers and unexpected approaches in candidates' scripts must be given marks that fairly reflect the relevant knowledge and skills demonstrated.

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

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Skill	Total marks	Breakdown of marks		Question 1	Question 2	
Manipulation, measurement and	16 marks	successful collection of data and observations	8 marks	2	6	
observation		nature of measurements or observations	8 marks	5	3	
Presentation of data and	12 marks	recording data and observations	4 marks	2	2	
observations		display of calculation and reasoning	2 marks	1	1	
		data layout	6 marks	3	3	
Analysis, conclusions and evaluation	12 marks	interpretation of data or observations and identifying sources of error	6 marks	3	3	
		drawing conclusions	3 marks	2	1	
		suggesting improvements	3 marks	3	0	

MMO = Manipulation, measurement and observation Collection = successful collection of data and observations Decisions = decisions relating to measurements or observations

- PDO = Presentation of data and observations Recording = recording data and observations Display = display of calculation and reasoning Layout = data layout
- ACE = Analysis, conclusions and evaluation Interpretation = interpretation of data or observations and identifying sources of error Conclusions = drawing conclusions Improvements = suggesting improvements
- ecf = error carried forward
- AW = alternative wording
- ora = or reverse argument

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Q		Expected Answers	Additional Guidance		Marks
1	(a)	How solutions made, concentration, measurements and observations.	2 PDO display, 2 MMO co	Check Supervisor's report for their results but mark according to mark scheme. Reject point 3 if heading units different from those recorded.	[5]
		<ol> <li>single table, all cells drawn;</li> <li>column headings: concentration/mol dm<sup>-3</sup> to left/across top, any one of length/mm/change in length/mm/volume cm<sup>3</sup> or ml;</li> <li>in table changes shown, 0.6, gets smaller/–ve, water, gets no change/longer/+ve;</li> <li>volume kept the same;</li> <li>at least two of 0.4/0.3/0.2/0.15;</li> </ol>	Mark first table, ignore other tables/further writing. No outer boundary needed. M or molar or mole(s)/l or per litre. Allow strength. No units in table. Must have units. Read from final length results in table. Allow if not shown in table. Mark for volumes anywhere. allow 0.48/0.42/0.36/0.24/0.18/ 0.12/0.06	report for their results but mark according to mark scheme. Reject point 3 if heading units different from those	
	(b)	Estimate the concentration of sucrose in X1.	ACE interpretation	1	[1]
		correct concentration from results, mol/dm <sup>3</sup> or mol dm <sup>-3</sup> ;	Match any one correct. O values but no made up es units M or molar or mole(s	timate and must have	

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(c)	Des	scribe and explain the results from all the solutions that you made.	3 MMO decisions		[3]
	con lenç	ncentration from results table, correct increase/decrease/no change in gth;	Ignore <b>X1</b> . Allow for their results even if wrong. Allow general statement.	If no changes in length recorded, allow 20mm/2cm – final length.	
	gra	rect ref to direction of water movement, correct ref to water potential dient; <u>osmosis;</u>	If no change then water moving in both directions/no net water movement. Higher or high to low/er WP ora.		
(d)	(i)	Describe and explain the results from all the solutions that you made.	2 ACE interpretation		[2]
		changes are small/accuracy of measuring/ruler +/-1mm/parallax error; widths are not standard; syringe qualified with ref. to uncertainty/large size vs coarse measurement; evaporation of solution; strips float/not immersed; no/not enough repeats/replicates; times different/not possible to add/remove at same time;	Mark any correct. Reject bubbles in syringe. Reject if just syringe not accurate. Must have +/–		
	(ii)		ACE improvements		[3 max]
		more concentrations/examples (at least 2) any between those given; smaller syringe/syringe with lower uncertainty/finer divisions; method/how to remove bubbles in syringe; longer strips; cover solution; repeat/use more strips; use burette/graduated pipette; use vernier callipers; do one experiment at a time/timed sequence/AW;			
(e)	(i)	Complete the Table 1.1 by calculating the missing value.	PDO display	1	[1]
		<u>5.6, -1.4;</u>	Only these answers.		



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(ii)	When the student performed this investigation, the diameter of cell 2 in 0.20 mol dm <sup>-3</sup> solution was 3.1 $\mu$ m. Explain why the student discarded this result and repeated the experiment with freshly made solutions.			[1]
	reading should have been more/too low/reading anomalous/not reliable/does not fit trend;	Allow idea of e.g. reading is odd/completely different.	Ignore just some error.	

0	x-axis conc.(sucrose)/mol $dm^{-3}$ or mol/ $dm^{3}$ ;	M or molar or mole(s)/ <i>l</i> or per litre
S/P	scale <i>x</i> -axis 0.2 to 2cm and <i>y</i> -axis uses half or more plotting crosses or dot in circle ONLY, correct for correct graph i.e. mean diameter of red blood cells only; Allow ecf from their figures/ crosses in circles	Ignore if 0 not at origin but if scale starts at 0.2 then must have 0.2. Reject if <b>mean change.</b> Reject 1 mm or more blobs in circles. Ignore shape of circles.
L	either ruled/straight lines joining each point/ ruled line of best fit with 2 one side and one the other/one on the line and two each side of the line;	If join the dots, then allow only 1mm/half square extension beyond first and last point. If line of best fit then allow 2mm/one square at 0.8/high concentration and to the axis for 0.05/low concentration end Reject line if more than 1mm thick

	reater the concentration of sucrose solution, the greater the eter of red blood cells.	ACE conclusions		[2]
as the	hesis wrong/incorrect; e concentration increases, the diameter of cells decreases/inversely ortional; ORA	Needs clear statement. Idea of correct relationship may quote figures to get same idea.	Reject idea that not totally wrong.	



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2 (a) (i)	Draw a LARGE, LOW-POWER plan diagram of part of specimen K TS Ranunculus.	1 MMO collection, 3 PDO	layout	[4]
	(quarter that is) 4 to 7 vascular bundles drawn, sharp, clear unbroken lines; no cells, larger than 6 cms in any one direction; distance of outer epidermis to top of nearest vascular bundle less than top of vascular bundle to pith; largest vascular bundle at least twice size of smallest vascular bundle;	VA Xey OX C(	Ignore labels. If more than a quarter drawn, then max1 point 2 only.	
		Ring 3 errors on simple diagram then reject. Use acetate square, accept if any line outside.	Reject if no pith drawn or a dotted line.	



		Page 7	М	ark Scheme		Syllabus	Paper		
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(ii)	Use this in bundle.	formation to ca	Iculate the actual v	vidth of the vascular	2 MMO o	collection, 1 PDC	) recording, 1	PDO display	[4]
	<u>Step 1</u> :	k the calculation.	W · X Y Z	✓ (1) s (Z/Y × X × W) ×1000			any 1 deci OR betwee	veen 2 and 3.5 mal place en 20 and 35.5 .5 no other	
	prrectly to 1 ace only. up or down. eck where th			Answer wrong. Go back to working MUST show Z/Y × X × W – all fig must be there and o the correct thing in whatever order OR calculated it correct (ignore decimal figured AND Look for ×1000 or × Must be shown not	loing have ly res). ✓ 10 <sup>3</sup> .				



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(iii)	Suggest how an error in measuring the vascular bundle could occur.	ACE interpretation	[1]
	thickness of lines/difficult to line up scales/depth of focus/not knowing where edge of vascular bundle is;	Allow focussing problem/blurring/aligning. Ignore can't count/do a mean/average/other excuses. Reject scale not calibrated correctly. Ignore parallax.	
(iv)	Make a LARGE, HIGH-POWER drawing of 5 cells to include a companion cell.	1 MMO collection, 3 MMO decisions	[4]
	<u>5 complete</u> cells, all touching (one group of five cells); sharp, clear unbroken lines, larger than 6 cm in any one direction;	Reject if more than 6 cells/no marks for textbook diagrams.	
	(companion cell) smallest cell drawn less than a quarter of the largest cell (phloem sieve tube), no more than one nucleus in a smaller cell, no sieve plates;	O $O$ $X$ $($ $($	
	all cells have single line walls, no shaded walls, no intercellular spaces;	A rings for errors reject touching cell wall lines/incomplete cell walls. Use acetate square. Reject if drawing not like slide e.g. too much detail drawn sieve plate and pores between cells or nuclei in cells.	



PMT

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2 (b) (i) Suggest the of stem.	he purpose of the region Y (	collenchyma cells) in corner	ACE conclusion		[1]
support ste	em/keep shape of stem/makes	corner;	Allow strengthening		
(ii) Compare a	pare and contrast TS Ranunculus with Fig 2.2.		2 MMO collection, 1 PDO recording, 2 ACE interpretation		[3 max]
organise as a table/venn diagram/ruled connected boxes, correctly labelled; (1) comparative statements opposite each other/in one sentence; (1) THREE from:		YFirst table ignore additional e.g. similarities separate table minimum.Allow for one statement as comparison.	statement as		
	K1 (Ranunculus)	Fig 2.4	Allow similarities vs differences as column	Reject two separate lists.	
(pith)	space in middle/no pith,	no space/has pith;	headings.		
(shape)	circular/no corners,	square/has (4) corners			
	no protrusions/AW/Y,	protrusions/AW/has Y;		Reject spherical.	
(vascular bundles)	lots/many,	few/not as many;			
	all round/evenly spaced,	mainly in corners/square;			
	rounded shape/oval,	shape described/curved;			
	big and small,	big and small;			
	arranged around sides,	arranged round sides;			
AVP;	no trichomes/AW,	trichomes;			

