MARK SCHEME for the October/November 2010 question paper

for the guidance of teachers

9700 BIOLOGY

9700/33

Paper 31 (Advanced Practical Skills 1), maximum raw mark 40

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UNIVERSITY of CAMBRIDGE International Examinations

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Question		Expected	Answers	Additional guidance			
1 (a) (i)	Decide on the concentrations of copper	stigation. [3	I			
	[1]	any 4 or more (volumes/concentrations);					
isions 3	[1]	(highest concentration) 0.3 to 0.15;	(highest concentration) 0.3 to 0.15;				
MMO deci	[1]	 any three consecutive concentrations (incl the same or serial dilution by half or serial dilution by ten; 					
	(ii)	State which variable you will need to co	nples. [1	1			
MMO decision 1	[1]	length or surface area or size or dimensior Allow methylene blue					
	(iii)	Describe how you will control this varial	ble and prepare the samples of plant tiss	ue. [2	I		
sions 2	[1]	(control) measure cut (methylene) rinsing/washing	the same any example of length 3 cm or less/size; excess				
MMO decis	[1]	(prepare samples) use of scalpel/knife or ruler; (methylene blue) water					

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	(iv)	Prepare the space below and	d record your observat	ions.	[5]
2	[1]	 Reject if units for % in body of ta other units e.g. mol dm⁻³ 	ble		
scording		table with all cells drawn	AND heading (top or le percentage conc(entra	ft) tion);	
PDO re	[1]	Rejectif headings/columns for m	nethod/volumes/time 5 n	nins or size/lengths	
		(heading) colour or observations or deso	cription;		
AMO ection 2	[1]	(records clear separate obser after/during 5 min/before mixi	vations/colours) ng	AND after mixing (after/at 5 min);	
n colle	[1]	difference in the strength of colour between the first and last test-tube observations;		Key e.g. + = colour	
MMO decision 1	[1]	5 or more concentrations or observation for water or replicate recorded;			
	(v)	Suggest how copper sulfate	solution affects plant	cell membranes.	[1]
nclusion 1	[1]	In correct context of increasin Idea of damages or destroys or makes more	g or just copper sulfate	it or ((cell) membrane(s)) phospholipid(s) fluid mosaic (model/structure) (fully) permeable	
CE co		denatures		protein	
A		(increases copper sulfate) (decreases copper sulfate)	∫increases ∫decreases	fluidity permeability	
		(increases copper sulfate) (decreases copper sulfate)	}decreases ∫increases	selective permeability;	

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(vi)	Identify three significant sources o	f error in your investigation.	[3]
Reje tem evar any	ect perature pH poration errors which affect all test-tubes equally	/	
Cau	se of error	Error	
[1]	(dependent) qualitative;		
[1] [1]	colour/colour change/observations	difficult judging seeing; qualitative;	
[1]	mixing	more difficult to judge colour/colours the same;	
[1]	(standardised variables) potato or position in potato or age or storage	not same different/variety old;	
[1]	lengths/size/surface areas/volumes Allow mass	not same;	
[1]	staining/washing/handling/forceps	not same loses stain damages potatoes ends not stained or middle more stain;	
[1]	potato/samples (into test-tubes)	time not same/delayed time/not at same time;	max 3

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	(vii)	Suggest how you would make <i>three</i> improvements to this investigation.	[3]
	[1]	same potato or position in same age or storage or fresh use micrometer/cork borer/vernier callipers/ruler with smaller divisions;	
MAX 3	[1]	leave in methylene blue longer/stronger concentration/more than 5 minutes idea of wash more;	
improvements	[1]	more/wider/narrower/different/examples range of concentrations or use burette or graduated pipette or smaller syringe or with smaller divisions;	
ACE	[1]	stagger start or do individually or use more stop clocks or use help;	
	[1]	colorimeter or datalogger with light sensor; Reject c <u>a</u> lorimeter	
	[1]	repeat or replicate;	max 3
		[Total: 18]	

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2 (a	a) (i)	Draw a large plan diagram of a qu	arter of the spec	imen as shown in Fig. 2.1. Label	the endodermis and cortex.	[5]
_	[1]	Rejectif drawn over the print of questi	on			
PDO layout 1		 Reject thick lines-than grid feathery lines 3 'tails' or overlaps or gaps 	AND no shading	AND uses most of space provided;		
		clear, sharp, unbroken lines				
ection	[1]	no additional cells drawn AND (epidermis shows) only the correct quarter;				
0 col 3	[1]	epidermis drawn with two lines 3 mm or closer for most of length;				
MM	[1]	innermost line is wavy/undulating line;				
MO decision 1	[1]	 1] Reject if any label is biologically incorrect e.g. regions belonging to other organs or animals. label within drawn area 		elonging to other organs or		
Σ		correct label with label lines to corte	ex and endodermis	s ;		

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	(ii)	Make a high-power drawing of or circumference. Labels are not requ	ne large xylem v ired.	vessel and the single layer of	cells touching a quarter of the vessel's [5]	
	[1]	Rejectif drawn over the print of questio	n			
PDO layout 1		 Reject thick lines – than on grid feathery lines 4 'tails' or overlaps or gaps if double lines for all cells 1 if single line for any cell 	AND no	AND uses most of space		
		clear, sharp, unbroken lines	shading	provided,		
	[1]	one xylem vessel drawn Ignore band inside	AND only single	e layer of surrounding cells ;		
on 3	[1]	Reject if layer of cells all round xylem vessel If xylem vessel not circular/polygonal				
) collecti		(surrounding cells) (single layer) three to eight cells in a layer only; Allow not touching.				
MMC	[1]	Reject any spaces if single line for cell walls. any gaps between cell walls – floating cells				
		(all cells including xylem vessel) no enclosed spaces more than 1mm	between adjacen	t double cell walls;		
PDO recording 1	[1]	cell walls drawn as double lines with surrounding cells;	middle lamella be	etween three adjacent cells from		

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(b) Prepare the space below so that it is suitable for you to record the observable differences between the specimens on K1 and that in Fig. 2.2.									
PDO recording 1	[1]	orga diag	anise as a table/Venn gram/ruled boxes	AND headed <u>K1</u> and <u>Fig 2.2</u>	AND first difference opposite each other;	ŀ	<u><1</u>	Fig 2.2	
						Ign	ore		
		feature		K1	Fig.2.2	•	 tick and cross without a key ref to non-observable features 		
	[1]	1	epidermis	hairs/trichomes Ignore root	no hairs/trichomes;	•	3D sh	apes	
	[1]			thick(er) or more/2 layers	thin(ner) or few(er);				
	[1]	2	cortex	yes/present/more	no(one)absent/less;				
3	[1]	3	endodermis	yes/present	no(one)/absent;				
tation	[1]	4	pericycle	yes/present	no(one)/absent;				
ACE interpret	[1]	5	vascular bundles) xylem	ring/centre/no(one)/absent/ fewer	scattered/AW/towards edge/yes/present/more;				
		6	thickened cells/	either way round for					
	[1]	sclerenchyma Allow collenchymas		present/absent/under epidermis;					
	[1]		bundle sheath/AW	no(one)/absent	yes/present;				
	[1]	7	pith	yes/present	no(one)/absent;				
	[1]		pith/centre cells	rounded	angular/pentagonal/AW;	-			
	[1] [1]	8	air spaces/lenticels stomata	yes/present no(one)/absent	no(one)/absent; yes/present;	max	nax 3		

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(c) (i)	Plot a chart of the data shown in Table 2.1. MAX 2 for O and S if line graph drawn		[4]
	O [1]	<i>x</i> -axis content(s)	AND <i>y</i> -axis conc(entration in) phloem or sieve tube/element (/) μg cm ⁻³ ;	Must have units
	S	scale as	Reject scale on <i>y</i> -axis any other than 20 to 2 cm.	
	[1]	even widths to 2 cm	AND <i>y</i> -axis <u>20 to 2 cm;</u>	
PDO layout 4	Ρ	 Reject if <i>y</i>-axis scale is awkward if bars arranged differently from order of table if horizontal lines are too thick – 1mm/hal square or not clear Allow bars if scale 20 to 2 cm. even if not 0 25 to 2 cm 	horizontal top line must be clear, sharp and ruled to show plot line must be on horizontal line for sucrose line must be between two lines for all other contents	
	[1]	correct plotting of each bar;		
	L [1]	each bar separate if vertical lines only then must be at least 1 cm apart.	 AND quality – vertical lines no thicker than on grid, not feathery for the complete line; bars – ruled lines Reject irregular thickness labelled clearly with contents – any clear labels e.g. chemical formulae NH₄, Ca, Mg, Na or mixture – underneath, must be directly below correct bar or inside bar or shaded with key. 	Reject solid shading If line shading outside a bar

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	(ii)	Calculate the percentage difference between the conce calcium ions in the phloem sieve tube elements.	entration of calcium ions i	n the xylem vessels and the concentration of [2]
[1]		shows subtraction (190 – 85) divided by 190 multiplied by 100; (190/190 – 85/190) × 100 or (1 – 85/190) × 100		
PDO di	[1]	Reject if no working Allow any answer less than 100 to no more than 3 significant figures 1 decimal place	AND percentage/%;	
(0	d) Sug	ggest why there is 120 μ g cm $^{-3}$ of sucrose in the phloem	sieve tube elements.	[2]
MAX 2	[1]	 (phloem sieve tube elements) (sucrose) transported leaf(ves)/allow type of leaf cell/source to roots/other tissues/sink(s); 		
ACE conclusions	[1]	(detail) <u>load(</u> ed) (in source) or (transported by) mass flow/bulk transport/translocation (sucrose) too large to move out of phloem or sieve tubes or xylem walls impermeable;		
			[Total: 22]	