## Mark scheme – Cell Structure: Microscopes

Qu	iest n	tio		Answer/Ind	icative conte	ent		Marks	Guidance
				laser scanning confocal microscop e	scanning electron microscop e	transmissio n electron microscope			
1		i	maximum resolution						Mark each row
			image appearanc e		3D	2D	~		
			image colour	named colour / coloured	black and white		$\checkmark$		
			larger numbe	er of (named)	organelles √				
		ii	more DNA / I no visible chi	-				2 max	ALLOW twice as much DNA
			nuclear mem	brane preser	nt √				
			Total					4	
2			В					1	
			Total					1	
3	а		(pond comm succession	unity is) final .	/ stable / not s	subject to furth	er	1	<b>IGNORE</b> 'permanent', it is in the rubric.
	b		light microsco graticule (1)	оре (1)				2	
	с	i	urea / uric ac	id				1	ALLOW ammonia, ammonium (ions).
		ii	Nitrosomona nitrite (1) Nitrobacter (1 nitrate (1)					4	
			Total					8	
4		i	3157 <u>µm</u> <sup>3</sup> / 3 <b>ОR</b>	.157 × 10 <sup>3</sup> <u>µ</u>	<u>m</u> ³			3	ALLOW for two marks correctly calculated value not given to 4SF e.g. 3156.55 μm <sup>3</sup> 3157.82 μm <sup>3</sup> (22/7used) 3154.95 μm <sup>3</sup> (3.14 used)

	3155 μm <sup>3</sup> / 3.155 × 10 <sup>3</sup> μm <sup>3</sup> (3.14 used for value of π)		
	OR		OR
			correctly calculated value without units e.g. 3157 / 3.157
	3158 $\mu$ m <sup>3</sup> / 3.158 × 10 <sup>3</sup> $\mu$ m <sup>3</sup> (22/7 used for value of $\pi$ )		OR
			correctly calculated value with inappropriate
	OR		units
	3.157 / 3.155 / 3.158, ×10 <sup>-15</sup> <u>m</u> <sup>3</sup> (answer using SI units)		e.g. 3.157 × 10 <sup>-6</sup> mm <sup>3</sup>
			3.157 × 10 <sup>-9</sup> cm <sup>3</sup>
	イイノ		
			If two or three marks were not awarded for
			the correct answer or calculated value:
			for <b>one mark</b> look for evidence of use of the
			formula:
			$(4/3) \times \pi \times r^3$
			Examiner's Comments
			The necessity to use $\boldsymbol{\pi}$ in the equation for
			Q17(b)(i) generated three different versions of
			the correct answer in order to encompass all the
			various values of $\pi$ e.g. 3.14 or 22/7.
			Candidates could gain credit for a variety of
			responses and many variations that included errors could still gain one or two marks if
			Examiners could clearly see how candidates
			had arrived at their response e.g. one mark was
			awarded for candidates that had used the
			correct equation for calculating the volume of a
			sphere. The most common error was using the
			wrong value for the radius, having confused it
			with the diameter or simply not knowing the formula for a sphere. The equation for
			calculating area of a circle rather than volume of
			a sphere was another common error. Some
			candidates mistakenly squared the value of the
			radius or mis-keyed the correct values into the
			formula and it is recommended that centres
			encourage candidates to repeat calculations to
			check for such errors, provided time allows.
	(transmission) electron (microscope) √		ALLOW TEM DO NOT ALLOW scanning electron microscope
			/ SEM
		2 max	
		∠ max	
	AND ONE of the following:		IGNORE black and white / colour
	2D image √		
<u> </u>	1		1]

		internal details visible $\checkmark$		
		(named) organelles / ultrastructures , visible $\checkmark$		e.g. mitochondria IGNORE nucleus (as visible under a light microscope)
		high <u>magnification</u> √		Examiner's Comments
		high <u>resolution</u> √		For candidates who recognised that an electron microscope was necessary to produce the image of the Kupffer cell, <b>Q17(b)(ii)</b> posed few problems and both marks were often achieved. Some candidates suggested a scanning electron microscope rather than transmission which was not credited but error carried forward was used for the second mark of this question to avoid candidates losing both marks. Error carried forward was also used where candidates had incorrectly stated a light microscope but marks were only gained for the mark points in the mark scheme and not for incorrect statements such as ' <i>no organelles can be seen</i> ' or ' <i>has low resolution</i> '.
		Total	5	
5	i	A = (permanent / temporary) vacuole √ B = <u>nucleolus</u> √	2 (AO1.1)	ALLOW vacule DO NOT ALLOW nucleus Examiner's Comments Most candidates correctly identified A as the vacuole, but only a minority of candidates identified B as the nucleolus, with many identifying it as the nucleus. Other answers seen included xylem and phloem, and air space for vacuole.
	ii	(x)14000 / 1.4 x 10 <sup>4</sup> √√	2 (AO2.8)	If the answer is incorrect, award one mark for a correct calculation not rounded to 2 s.f. (e.g. 0.02 / 0.0000014 = 14285.71429 20000 / 1.4 = 14285.71429) ALLOW 0.019/0.0000014 = 13571.428 or 0.021/0.0000014 = 15000 for 1 mark Examiner's Comments On the whole this question was well answered, with the majority of candidates correctly calculating the magnification. However, some candidates lost a mark for failing to round the answer to 2 significant figures. OCR support

				There are available resources on the 'Maths for Biology' website which can be used to support candidates with the correct use of significant figures: https:/www.ocr.org.uk/subjects/biology/maths- for-biology/handling-data/
				Mark first two improvements described
				e.g. only use outlines
				<b>IGNORE</b> references to labels or annotations and the use of a pencil (because this is mentioned in the question stem)
				<b>IGNORE</b> drawing should take up half a page / no overlapping lines / use continuous lines
				Examiner's Comments
	iii	no, shading / cross hatches / AW ✓ add, a scale / magnification ✓ add a title √	2 max (AO3.4)	Many candidates correctly recognised that the drawing could be improved by removing shading or by adding a scale bar. However, many referred to adding labels or annotations or the use of a sharp pencil which gained no credit, as this was mentioned in the question stem.
				OCR support
				The Biological Drawing handbook can be used as a guidance in this case: https:/www.ocr.org.uk/Images/251799-biology- drawing-skills-handbook.pdf
				Mark as prose IGNORE use forceps / lay sample flat
				ALLOW place stain at side of sample
				ALLOW stated angles given e.g. 45°
	i	place stain at edge of sample (not the centre) $\checkmark$	2 max	ALLOW 'tissue/paper towel' instead of 'blotting
	v	lower cover slip at an angle / use mounted needle $\checkmark$	(AO3.3)	paper' ALLOW ensure stain covers whole sample
		use blotting paper to, remove excess stain / pull stain through $\checkmark$		Examiner's Comments
		use more than one stain (to improve contrast) $\checkmark$		Candidates who had a practical knowledge of slide preparation scored well, mentioning lowering the cover slip at an angle or using blotting paper to remove excess stain, as ways

				to improve the method. However, many candidates wrote about aseptic technique, adding water, wearing gloves, or pressing down on the cover slip to remove air bubbles, which gained no credit.
		Total	8	
6	i	transmission electron (microscope) $\checkmark$	1 (AO2.1)	ALLOW TEM, 'microscopy' for 'microscope'
	ii	M = matrix √ N = crista(e) √	2 (AO1.1)	ALLOW inner membrane for N
		Total	3	
7		В√	1 (AO1.1)	
		Total	1	
8	i	you can now see Golgi body / mitochondria / (smooth / rough) endoplasmic reticulum / ER / RER / SER / ribosomes OR organelles seen in more detail / grana (in chloroplast) / thylakoids (in chloroplast) / nuclear pore / cristae (in mitochondria) / membranes within organelles / double nuclear membrane / (double) nuclear envelope OR resolution is, higher / better √	1	IGNORE clarity IGNORE ref to size of organelles DO NOT ACCEPT chloroplast IGNORE ref to ultrastructure unqualified Examiner's Comments This was answered well with most candidates correctly identifying an organelle which could not be seen with a light microscope, but could then be seen in the second image. Many correctly referred to rough endoplasmic reticulum or mitochondria although the presence of organelles simply being visible was insufficient. Many were able to comment on the higher resolution of cell B but a good proportion failed

					to gain credit due to the use of 'high' rather than 'higher'. Weaker answers often were too vague, simply stating that the 'ultrastructure' or 'detail' of the cell could be seen. <b>ORA</b> for electron microscope needs to be comparative <b>IGNORE</b> colour
		II	LSCM image has lower resolution (than EM) OR can have fluorescent tag OR can see movement (as can be used on living cells) OR can see, different layers / at different depths (of the sample) √	1 max	IGNORE ref to 2D / 3D / depth of field Examiner's Comments The majority of candidates were not confident in answering this question and as a result gave an incomplete or irrelevant answer. Some candidates were awarded the resolution mark but some were clearly confused as to which microscope has the higher resolution or were unsure about the difference between resolution and magnification. Many students responded in terms of the image being either 2D or 3D, some answers stated 'in colour', and several did not state which type of microscope they were referring to. Of the few correct answers seen, the majority referred to different depths of sample or the usefulness of being able to use a fluorescent tag. None referred to the advantage of being able to see movement (in living cells).
			Total	2	
9	а		Ζ;	1	Mark the first answer. If the answer is correct and an additional answer is given that is incorrect or contradicts the correct answer then = 0 marks Examiner's Comments Most students recognised that microscope Z was the transmission electron microscope.
	b		Fig. 3.1(a) (no mark)	max 2	Please place a green blob on paper

		shows surface view; 3D / three dimensional; better <u>resolution</u> (than b);		Do not allow mp 2 if fig 3.1 b selected Do not allow mp 3 if fig 3.1 b selected Must be comparative comment <b>Examiner's Comments</b> Most students recognised that Fig. 3.1(a) was the image from a scanning electron microscope and were able to justify their choice successfully. The most common response was that the image was three dimensional, but many candidates also stated that it was a surface view. Fewer candidates stated that the resolution was higher than in Fig. 3.1(b).
		Total	3	
1 0	i	1.7 mm (1)	1	
	ii	× 50 (1)(1)		ALLOW 1 mark for correct working e.g. 80 / 1.6 ALLOW answer in the range of 48–51
	iii	air spaces give buoyancy (1) supported by (surrounding) water (1)	2	
		Total	5	
1 1		D	1	
		Total	1	
1 2		11.91 µm √√	2	Correct answer = 2 marks (indicated by 2 ticks) even if no working shown ACCEPT 11.06 to 12.77 µm ACCEPT 1.106 x 10 <sup>-5</sup> m to 1.277 x 10 <sup>-5</sup> m [sig figs retained for standard form] Otherwise, Award ONE mark for: correct final answer without (correct) unit OR correct final answer to wrong number of dp or incorrectly rounded OR seeing (one graticule division =) 20 ÷ 2.35 =

					8.51
					0.01
					OR
					seeing (measurement of nucleus = )
					1.3 to 1.5 (graticule / eye piece) units / divisions
					<b>or</b> 1.3 to 1.5 cm
					or
					13 to 15 (graticule / eye piece) units / division
					or
					13 to 15 mm
					OR
					diameter = 110.63 to 127.65 µm
					Examiner's Comments
					A common error was misusing the formula
					linking magnification with image size and actual
					size, so wrongly dividing 2.35 by 20. The correct units were usually used. Students should be
					encouraged to think about what their answers
					actually mean - answers of hundreds of mm or
					cm for a cell nucleus are clearly wrong.
			Total	2	
			use eyepiece graticule $\checkmark$		
			calibrate graticule, using stage micrometer / detail of calibration / calculate the length of one epu $\checkmark$		
1			measure the diameter of the nucleus in, epu / graticule		
1 3			units √	4 max	e.g. of detail: align two scales and record
			take repeat measurements <b>and</b> calculate a mean		number of divisions per graticule unit
			diameter (in epu) √		
			use calibrated epu to calculate diameter (of nucleus) (in $\mu m)$ / described $\checkmark$		
			Total	4	
			sharp blade (should be selected) (1) so slide is thin enough, individual cells are visible / resolution is high (1)		
1		i	method for slicing pieces of tissue (thinly) (1) so slide is thin enough, individual cells are visible /	6	<b>ALLOW</b> any reasonable method (e.g. microtome)
4			resolution is high (1)	Ŭ	
			select thin(nest) slides (1)		ALLOW quantified thickness (e.g. measured
1	1		to ensure maximum light can penetrate sample (1)		with a micrometer)

## Cell Structure - Microscopes

		wet mount (1) prevents dehydration / distortion of tissue (1) squash slide (1) easier to see individual cells / allows light to penetrate tissue more easily (1)		ALLOW description ALLOW description
	ii	contrast is high(er) (1) more (internal) structures visible (1) some (named) organelles / cell components more visible, because they bind to stain (1) clearer image can be obtained (1)	3	
		Total	9	
1 5		D√	1 (AO1.2)	
		Total	1	
1 6		C√	1	
		Total	1	
1 7	i	<ol> <li>to, see / identify, (differences between) cells √</li> <li>to, see / identify, (differences between) organelles √</li> <li>red blood cells visible, anyway / without stain (due to haemoglobin) √</li> <li>ref. contrast √</li> <li>allows, white cells / leucocytes, to be counted √</li> </ol>	max 3 (AO1.1 x5)	<ul> <li>ALLOW so white blood cells / A / C / D can be seen or told apart from RBCs</li> <li>ALLOW named organelles e.g nucleus / cytoplasm</li> <li>ALLOW without stain white cells are, transparent / colourless</li> <li>Examiner's Comments</li> <li>A few candidates gained 2 marks out of the 3 for stating that the staining technique improves visibility of both cells and organelles. Limited vocabulary let candidates down, with frequent references to components or structures rather than 'organelles' or a named organelle. Higher ability candidates sometimes featured the word 'contrast' or made the point that white blood cells needed to be stained to be visible (or to be counted) while red blood cells contained their own pigment.</li> </ul>
	ii	<ul> <li>1 C (is, blue / purple, so) has (more) nucleic acid √</li> <li>2 (C has) (m / t / r) RNA √</li> <li>3 D (is red so) has (more) protein √</li> <li>4 (D has) enzyme / antibody / immunoglobulin √</li> <li>5 idea that different cells have different, roles / (concentrations of) biochemicals / levels of activity √</li> </ul>	max 4 (AO3.2 x5)	IGNORE suggested names for cells IGNORE some / no, protein present 2 DO NOT ALLOW DNA 3 IGNORE some / no, nucleic acid present 4 ALLOW (named) hydrolases / (named) cytokines / perforins / granzymes

				Examiner's Comments
				This question required candidates to use the information about the chemical staining characteristics of haematoxylin and eosin to make deductions about the contents of the cytoplasm of two cell types they observed on Fig. 2. Most creditworthy answers made the link between the colour of the stained cytoplasm and the main biomolecule contained within it. Some candidates went on to suggest that the proteins in cell D were enzymes or antibodies and gained credit. Fewer candidates linked more nucleic acid in the cytoplasm of cell C to more RNA. A pitfall for lower ability students was missing the instruction in the question to consider the cytoplasm of the cells only, not the nuclei (both of which stained purple due to DNA).
		Total	7	
1 8		create / provide / increase contrast; make, cells / (named) component(s), visible OR cells / (named) components, can be, identified / distinguished / differentiated;	2	IGNORE clearer ACCEPT (named) organelle(s) stand out from surroundings ACCEPT regions / parts / AW, of cell Examiner's Comments Most candidates knew that staining made cell components visible and many also understood that the stain increases the contrast.
		Total	2	
1 9		TEM has, better / high <u>er</u> , resolution $\checkmark$ <i>TEM</i> (resolution figure in range) 0.05 - 2 nm $\checkmark$ (shows) image of cell interior $\checkmark$ (shows) ultrastructure / (two named) cell organelles $\checkmark$ <i>SEM</i> (resolution figure in range) 5 - 50 nm $\checkmark$ (shows) 3D / three-dimensional, image $\checkmark$ (shows cell) surface / topography $\checkmark$	4 max (AO1.1)	ALLOW ora SEM has, worse / lower resolution IGNORE magnification ALLOW 0.00005 - 0.002 μm / 50 – 2000 pm 'TEM has resolution of 0.5nm whereas SEM only has resolution of 3-10nm' gets mps 1, 2, 5 (as comparative implied by 'only') ALLOW 0.005 – 0.05 μm ALLOW see depth DO NOT ALLOW organelles in cell unless fracture explained
		Total	4	
2 0		A√	1	
		Total	1	

2			1	Examiner's Comments
1		A √	(AO1.1)	The vast majority of candidates answered this straightforward definition question correctly.
		Total	1	
2 2		С	1	<b>Examiner's Comments</b> This was answered reasonably well. It should be emphasised that questions such as this, where data are presented that do not have definitive values (in this case, varying according to the particular make and model of microscope), that candidates are expected to have an appreciation of the magnitude of the values rather than learning precise figures and expecting to quote those.
		Total	1	
2 3	i	<i>B</i> comment about detail of organelles (1) comment about shapes of cells (1)	2	No Mark for identification of B e.g. light microscope would not allow nuclear pores / ribosomes / endoplasmic reticulum / plasmodesmata to be seen. e.g. sieve tube elements are angular / hexagonal.
	ii	the ability to see more detail / separate two objects (1)	1	
	iii	Nile blue (1) to increase contrast / to make nuclei visible / to show no nuclei in sieve tubes (1)	2	
		Total	5	
2 4		14 000 / 120 = 117 μm;	2	length of line A-B = 14mm / 14000 μm Correct answer = 2 marks. Allow one mark if correct working shown including units for cm & mm e.g. 1.4 cm / 120 14 mm / 120 14000 / 120 If answer = 125 μm allow one mark for correct working but incorrect measurement (15mm instead of 14) Allow one mark if not rounded to nearest micrometre <b>Examiner's Comments</b>

				As ever, many candidates proved incapable of performing a relatively simple calculation. Better candidates did well and gave the correct response or at least showed they knew how to carry out the calculation. However, too many candidates appeared to have little idea of what to do. Commonly the magnification was left out of the calculation and candidates simply converted mm to µm. Another common error was to convert mm to µm by dividing by 1000 rather than multiplying. This is an area in which centres need to improve in light of the increased maths requirements of the new specifications.
2 5	i	Total         A because         nuclei (of white blood cells) are lobed √	<b>2</b> 1	Mark is for the explanation
	ii	(x) 1300 √√	2	If answer is incorrect <b>ALLOW</b> 1 mark for evidence of 0.02 (m) / 0.000015 (m) or equivalent numbers in alternate units
		Total	3	
2 6		C√	1 (AO2.4)	Examiner's Comments Most candidates were able to correctly calculate the real length of the tardigrade and there was often evidence of working shown.
		Total	1	
2 7		12 √ √	2(AO2.2 )	Correct answer = 2 marks even if no working shown. ALLOW 11 / 13 for 2 marks If answer is incorrect then award 1 mark: <i>if answer to</i> >2 <i>s.f.:</i> ALLOW range from 11.2 to 12.8 <i>if answer in mm:</i> 0.011 / 0.012 / 0.013 <i>if answer in cm:</i> 0.0011 / 0.0012 / 0.0013 <i>if answer in m:</i> $1.1 \times 10^{-5} / 1.2 \times 10^{-5} / 1.3 \times 10^{-5}$ for working: $14 \text{ or } 15 \text{ or } 16 \div 1250 \times 1000$ for converting scale bar to $\mu m$ : 15 000 or in range from 14 000 to 16 000

				ECF from mis-measured figure: answer to (x ÷ 1250 x 1000) e.g. 1cm gives an answer of 8 (µm) e.g. 1.5 mm gives an answer of 1.2 (µm) Examiner's Comments Many candidates correctly calculated that the 15mm scale bar represented a length on the magnified image equal to 12µm. Some candidates were given one mark for converting the scale bar length to µm. The use of a scale bar to represent a length on a magnified image confused some candidates A microscopic object such as a cell has a real-life size and a magnified image size. A scale bar's length is as measured on the page but this represents a length on the image that is smaller by a factor of the magnification shown.
		Total	2	
2 8		350, (x / times) √√	2	ALLOW range 333(.33) - 350 x scale bar = 21mm max.1 working mark for: 21mm / 0.06mm or 21 000µm / 60µm Note: candidate may measure size of photo (image size) and calculate size of object using the scale bar. If calculated correctly this gives an answer very close to 350. Allow 2 marks.
		Total	2	