

2801 Biology Foundation

January 2005

Mark Scheme

ADVICE TO EXAMINERS ON THE ANNOTATION OF SCRIPTS

- 1. Please ensure that you use the **final** version of the Mark Scheme. You are advised to destroy all draft versions.
- 2. Please mark all post-standardisation scripts in red ink. A tick (✓) should be used for each answer judged worthy of a mark. Ticks should be placed as close as possible to the point in the answer where the mark has been awarded. The number of ticks should be the same as the number of marks awarded. If two (or more) responses are required for one mark, use only one tick. Half marks (½) should never be used.
- 3. The following annotations may be used when marking. <u>No comments should be</u> written on scripts unless they relate directly to the mark scheme. Remember that scripts may be returned to Centres.
 - x = incorrect response (errors may also be underlined)
 - ^ = omission mark
 - bod = benefit of the doubt (where professional judgement has been used)
 - ecf = error carried forward (in consequential marking)
 - con = contradiction (in cases where candidates contradict themselves in the same response)
 - sf = error in the number of significant figures
- 4. The marks awarded for each <u>part</u> question should be indicated in the margin provided on the right hand side of the page. The mark <u>total</u> for each question should be ringed at the end of the question, on the right hand side. These totals should be added up to give the final total on the front of the paper.
- 5. In cases where candidates are required to give a specific number of answers, (e.g. 'give three reasons'), mark the first answer(s) given up to the total number required. Strike through the remainder. In specific cases where this rule cannot be applied, the exact procedure to be used is given in the mark scheme.
- 6. Correct answers to calculations should gain full credit even if no working is shown, unless otherwise indicated in the mark scheme. (An instruction on the paper to 'Show your working' is to help candidates, who may then gain partial credit even if their final answer is not correct.)
- 7. Strike through all blank spaces and/or pages in order to give a clear indication that the whole of the script has been considered.
- 8. An element of professional judgement is required in the marking of any written paper, and candidates may not use the exact words that appear in the mark scheme. If the science is correct <u>and</u> answers the question, then the mark(s) should normally be credited. If you are in doubt about the validity of any answer, contact your Team Leader/Principal Examiner for guidance.

Mark Scheme	Unit Code	Session	Year	Version
Page 3 of 9	2801	January	2005	Final

Abbreviations, annotations and conventions used in the Mark Scheme	; = NOT = R = () = ecf = AW = A	words which are not essential to gain credit (underlining) key words which <u>must</u> be used to gain credit error carried forward alternative wording	
---	---	--	--

Question Expected Answers

1

mark two columns separately first. If letter and part of cell both incorrect, look to see if the part of the cell corresponds to this letter. If so, allow 1 mark ecf

function	part of cell	label
controls activities of the cell	nucleus	А
carries out aerobic respiration	mitochondrion / mitochondria ;	D ;
attaches to mRNA in protein synthesis	ribosome(s) / <u>rough</u> ER / <u>R</u> ER ;	С;
produces secretory vesicles	Golgi ;	В;
contains digestive enzymes	lysosome(s);	E;

Marks

[Total: 8]

Mark Scheme Page 4 of 9			Unit Code 2801	Session January	Year 2005		rsion inal
Question Exp		Expected /	Answers				Marks
2 (a)) (i)	polypeptide	; A oligopeptide				1
	(ii)	glycine;	A proline / alanine				1
	(iii)	chain = pol	ver assume that ypeptide groups of 3 polypeptide chains				
		amino acids the small or chains , for	amed amino acid from (ii) but NG s / glycine , small (to allow close ne is , every 3 rd amino acid / at e m a tight coil / lie close to each er by hydrogen bonds ; <i>ignore</i>	epacking) ; every level in the n other ;			
		molecules f covalent bo fibres comp ends of par	between R groups of lysines ; form , fibres / bonds with adjace and between , adjacent molecule posed of parallel molecules ; allel molecules staggered ; he of weakness ;		A fibril S ;		2 max
(b)		cell wall(s) β / beta ; glycosidic ; 180 ; straight ; hydrogen /	 A B NOT glucosidic A polysaccharide / unbranche 	ed / linear			6
						[Total:	10]

Mark Scheme				Year	Version	
Page 5 of 92801January2005						Final
Questi	on	Expected A	Answers			Marks
thin c lack c large prese (mem many		thin cell wal lack of , wa large surfac present in la (membrane many mitoc	terproof layer / cuticle ; ce area ; NOT if cilia / villi / m arge numbers ; c) proteins / carriers / channels			1 ma
	(ii)	active trans A pi	e gives a list or a choice, all mu sport / diffusion / facilitated diffu nocytosis f passive transport / osmosis /	usion / described ;		1
	(iii)	lower <u>water</u> movement through , ch	<u>potential</u> inside / ora ; , down water potential gradien nannel proteins / partially perm permeable ;	t / from high Ψ to lov		2 ma
(b))	U; V; Z;				

s;

4

[Total: 8]

	Downloaded from http://www.thepaperbank.co.uk						
Ма	Mark Scheme		Unit Code Session		Year	Version	
Р	Page 6 of 92801January2005				2005	Final	
Questio	'n	Expected A	Answers			Marks	
4 (a)	(i)	4;				1	
	(ii)	phosphate		A purine / pyrimic	line		
			take a correct base from a lis	t unless that list ir	ncludes uracil	3	
(b)		 2 idention 3 (each 4 1 new 	blecules / helices , (of DNA) prod cal (molecules of DNA produced made up of) 1 , original / parent / strand ; al / parent / old , strands , act as	l) ; t / old , strand ;	bed :		

- 5 original / parent / old , strands , act as template / described ;
- 6 ref to (free DNA) nucleotides ;

[Total: 7]

3 max

				-				
	Mark Scheme			Unit Code	Session	Year	Version	
	Pa	ige 7	' of 9	2801	January	2005	Final	
Qu	estior	ı	Expected	Answers			Marks	
5	(a)	;	2 max					
	(b) (i) (X) 10 / 900% (increase) ; NOT 10% incr ignore 1000%						1	
		(ii)	can e.g.	didates may use information from typical [NOT average] = 20 t threshold = 200 units				
			1 no i	increase , between 0 and 20 unit of radon ;	s / at low levels /	well below three	shold,	
	2 radon			on increasing, from 20 to 200 u	nits / towards thre	shold, increase	s risk ;	
				10X / 900% ; n radon and smoking gives grea	test risk ;			
			5 & 6 othe	er suitable quantitative <u>risk</u> state	ment;;			
			7 con	sequence / relevant effect on ce	H;		2 max	
	(c)		<i>advantage</i> make people aware of risk / let people know that their area is safe / could reduce other risks / other suitable suggestion ;					
			<i>disadvanta</i> worry peop	<i>ge</i> le / lower house prices / migratio	on / other suitable	suggestion ;	1	

Mark Scheme	Unit Code	Session	Year	Version
Page 8 of 9	2801	January	2005	Final

Question Expected Answers

Marks

(d) only award marking points 1, 6, 9, 14 and 16 if descriptions of the stages are correct- do not award simply for identifying the stages - ignore ref to centrioles

- prophase
- 1 C;
- 2 chromosomes / chromatids , condense / coil / shorten and thicken ;
- 3 become visible ;
- 4 consist of two <u>chromatids</u>;
- 5 joined by a centromere ; A kinetochore NOT centrosome

metaphase

- 6 A;
- 7 chromosomes align <u>at</u>, equator / metaphase plate;
- 8 attached to spindle by centromeres ;

anaphase

- **9** B;
- 10 centromere splits ;
- **11** chromatids separate ;
- **12** move to opposite poles ;
- **13** by , contraction / shortening , of spindle ;

telophase

- 14 E;
- 15 chromosomes uncoil;

interphase

- 16 D; A for a description of early prophase
- **17** <u>DNA</u> replication ;
- **18** transcription / formation of mRNA ;
- **19** AVP ; these must relate to behaviour of chromosomes
- 20 AVP; e.g. spindle made of microtubules chromatin becomes chromosomes (in prophase) ora in interphase centromere leads chromatid to pole gene switching during interphase

9 max

1

QWC – clear well organised using specialist terms ; award the QWC mark if three of the following are used in correct context, but Q = 0 if

names of stages of mitosis are used inappropriately

chromatin	equator / metaphase plate
chromatid	DNA replication
centromere	transcription
spindle	

Mark Scheme	Unit Code	Session	Year	Version
Page 9 of 9	2801	January	2005	Final

Qu	estion	Expected Answers	Marks
6	(a)	idea that arachidonate is substrate ; phospholipid source in membrane ; prostaglandin / product , can be , transported / stored ; (S)ER for , lipid / steroid , synthesis / transport ; AVP ; AVP ; AVP ; e.g. separate from other reactions cytoplasm environment not suitable for , reaction / enzyme ora idea that prostaglandin isolated COX does not , damage / use phospholipids from, other membranes	2 max
	(b)	<i>ibuprofen</i> competitive ; ibuprofen blocks / arachidonate cannot enter , channel ; A substrate cannot reach active site ; <i>aspirin</i>	
		non-competitive ; changes shape (of) / blocks ; active site ; AVP ; e.g. <u>allosteric</u>	
		no ESC formed / AW; allow once only	4 max
	(c)	A reverse argument as long as question is answered in terms of low temperature	
		slows, reaction / rate / activity of enzyme / AW;	

ref kinetic energy ; molecules moving , slowly / less ; few collisions / collisions less likely ; few ESC formed / ESC less likely to be formed ; reversible / enzyme not denatured / enzyme still works ; ref activation energy ; ref $Q_{10} = 2$;

4 max

[Total: 10]