MARK SCHEME for the May/June 2013 series

9700 BIOLOGY

9700/21

Paper 2 (AS Structured Questions), maximum raw mark 60

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 should be read in conjunction with the question paper and the Principal Examiner Report for Teachers.

Cambridge will not enter into discussions about these mark schemes.

Cambridge is publishing the mark schemes for the May/June 2013 series for most IGCSE, GCE Advanced Level and Advanced Subsidiary Level components and some Ordinary Level components.



Page 2	Mark Scheme	Syllabus	Paper
	GCE AS/A LEVEL – May/June 2013	9700	21

Mark scheme abbreviations:

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

P	Page 3 Mark Scheme Syllabus			Syllabus	Paper
	•		GCE AS/A LEVEL – May/June 2013	9700	21
1 (a	1) (i)		endothelial/squamous/epithelial (cell) ; nucleus ;		[2]
	(ii) (:::)	awa awa if no (38–	rd two marks if correct answer given rd one mark if not rounded to nearest whole number rd one mark if given incorrect unit answer given, award one mark if correct measurement 41/3.8–4.1/38000–41000) is divided by 5700		[2]
	(iii)	only	<i>wo marks - one structure and one function two functions = 1 mark two structures = 1 mark</i>		
		2 3 4 5 6 7 8	(capillary) <u>wall</u> is, thin/single layer of cells/one cell thick A endothelium/epithelium for wall short <u>diffusion</u> , pathway/distance/AW; R 'easy' diffusion (many have) endothelial pores/fenestrations/gaps/spa to allow named, substance/cell, to leave the blood; A <i>idea</i> of separation/selection, of named substance(s) small diameter/small lumen/diameter of red blood cells slows down flow of red blood cells/(capillary/blood) clo (capillaries have) large, surface area/surface area to vo <i>idea that</i> allows more exchange; Ignore faster exchange	ces/openings; by size ; se to cells;	[max 2]
(b	o) wh 1 2	(nan A dia high	od cells ned) white blood cells can, leave capillaries/enter tissue apedesis/(suggestion that some) too large to leave the, number in, lymph nodes/thymus/bone marrow/spleen ored/produced	blood/capillarie	S
	g <i>lu</i> 3 4 5	filter take	ll (molecule) ; ed/diffuses/leaves/leaks, from blood/from capillaries/ n up/used, by cells in respiration ; o re supply	into tissue fluid ;	
	p <i>ro</i> 6 7		arge to, leave capillaries/enter lymph/enter tissue fluid /mph / tissue fluid) antibodies/proteins, from/secreted b ;		other [max 5]

Paper
21
only [max 3] Fotal: 14]
[2]
[max 2]
[max 1]
[max 1] [Total: 6]
Γ

Pa	ige 5	Mark Scheme	Syllabus	Paper
		GCE AS/A LEVEL – May/June 2013	9700	21
3 (a)	stomata carbon d ignore re water <u>va</u>	ffusion is a consequence of transpiration open(ings) to allow carbon dioxide in ; ioxide required for photosynthesis ; ef. to oxygen <u>pour diffuses</u> out through stomata ; f evaporation (from mesophyll walls) described as a gas		[3]
(b)	 both ref. 1 both trans both trans both both for n R if 0 com to co or tras 	show, little/low/lowest, transpiration, at night/22.00 to , increase to/peak, at mid day / 12.00 ; o second peak at 16.00 ; , dip/decrease, at 14.00 ; spiration (always) lower for trees at exposed site/ora ; decrease from 16.00 ; <i>nps 1–4 and mp 6, allow a description at one site only</i> contradictory description given for the other site parative data quote to support above marking points ; ; ompare the transpiration rate at two locations at the sam anspiration rate at one location at different times ward data marks arbitrary units (au) must be used at leaf	ne time	o 04.00 ; [max 5]
(c)	 stoma stoma stoma stoma hairs/ low nu ignor thick(a reflect thick(a 	tion of the following features ta close (for longer), during the day/when hot/when dr ta in pits/sunken stomata ; ta only on lower surface of the leaf ; trichomes ; umber of/few(er)/less, stomata (per unit area) ; e 'less open stomata' er) cuticle; tive cuticle (on upper epidermis) ; er) epidermis/more than one layer of epidermal cells ; l/rolled/AW, leaves ;	у;	[max 3]

[Total: 11]

	Page 6		Mark Scheme	Syllabus	Paper
			GCE AS/A LEVEL – May/June 2013	9700	21
4	_	Kg Lh	nymine ; iuanine ; ydrogen bond ; gnore H/H₂ bond		[max 3]
	 (b) 1 zidovudine, <u>competitive</u> inhibitor <u>and</u> efavirenz, <u>non-competitive</u>; 2 zidovudine, <u>complement</u>ary to active site; 3 efavirenz, binds to allosteric site/reference to allostery; 4 efavirenz changes the shape/structure of the active site; 				

- efavirenz changes the, shape/structure, of the active site;
 A denatures/changes tertiary structure so substrate will not fit
 R changes shape unqualified
- 5 either

the effect of zidovudine is, reduced/reversed, by increasing the substrate concentration

or the offer

the effect of efavirenz is not, reduced/reversed, by increasing the substrate concentration;

[max 4]

- (c) 1 virus may be resistant to one or more of the drugs/very low chances that HIV is resistant to all of the drugs;
 R virus immune
 - 2 (resistance due to) change to, active site/allosteric site/tertiary structure, of enzyme;

A drug can no longer fit into active site

- 3 some drugs may be more effective than others/AW;
- 4 reduces risk of drug resistance developing ;
- 5 HIV, has a high mutation rate/changes surface proteins / changes antigens ; A antigen(ic), shift/drift
- 6 person may have mixture of strains of HIV;
- 7 idea that virus will be at different stages in its, life/replication, cycle;
- 8 AVP;
 - e.g. more than one competitive to reduce chances of, ES complexes/AW drugs work better in combination/synergy idea drugs inhibiting two different enzymes, so more effective
- (d) 1 antibiotics are not effective against HIV or viruses/antibiotics are effective against, (named) bacteria/bacterial disease ;
 A fungi/protoctists/protists/malaria
 R antibiotics prevent infection
 - 2 idea that viruses have no, sites / targets, where antibiotics can work ;
 - viruses have no, cell walls/ribosomes/cell membranes;
 A have different enzymes
 - 4 viruses are within cells, *idea that* antibiotics cannot reach them;
 - 5 people with HIV are more susceptible to bacterial infections/reference to immune suppression/weak immune system ; [max 2]

[Total: 11]

[max 2]

GCE AS/A LEVEL – May/June 2013 9700 21 5 (a) P – moves, polar substances/hydrophilic molecules/ions, through membrane/in or out (of cells); A facilitated diffusion/active transport/described Q – receptor/recognition site/cell recognition/binding site; A cell adhesion/'receives' named signal A stabilises membrane (as forms hydrogen bonds with water) R – regulates/AW, fluidity of, membrane/(phospholipid) bilayer; A contributes to hydrophobic layer/impermeability to ions [3] (b) 7.0 nm; [1] (c) fluid idea of phospholipid (and protein) molecules, move about/diffuse (within their monolayer); [1] mosaic to max 1 protein (molecules), interspersed/scattered/not a complete layer/AW; [2]	Page 7	Mark Scheme	Syllabus	Paper
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 (c) <i>fluid</i> <i>idea of</i> <u>phospholipid</u> (and protein) molecules, move about/diffuse (within their monolayer); <i>mosaic to max 1</i> protein (molecules), interspersed/scattered/not a complete layer/AW; different/AW, proteins (molecules); (d) 1 water molecules are polar; R hydrophilic/charged <i>idea that</i> few polar molecules pass through the <u>phospholipid</u> (bilayer); ora for non-polar molecules A none pass/repelled 3 core of membrane/phospholipid tails, are hydrophobic; A hydrophobic core 4 channels (through aquaporins), are hydrophilic; A core of channel proteins/described as R-groups of amino acids 5 (aquaporins) increase <u>permeability of membrane</u> to water; 6 example; e.g. root hairs/small intestine epithelium/nephron 7 role of water in a cell; e.g. solvent/reactant/reaction medium/turgidity <i>or</i> support in plant cell 	out A fa Q – rece A ce A si R – regu	(of cells) ; icilitated diffusion/active transport/described eptor/recognition site/cell recognition/binding site ; ell adhesion/'receives' named signal abilises membrane (as forms hydrogen bonds with wate lates/AW, fluidity of, membrane/(phospholipid) bilayer	er)	
 <i>idea of phospholipid</i> (and protein) molecules, move about/diffuse (within their monolayer); <i>mosaic to max 1</i> protein (molecules), interspersed/scattered/not a complete layer/AW; different/AW, proteins (molecules); (d) 1 water molecules are polar; R hydrophilic/charged <i>idea that</i> few polar molecules pass through the phospholipid (bilayer); ora for non-polar molecules A none pass/repelled 3 core of membrane/phospholipid tails, are hydrophobic; A hydrophobic core 4 channels (through aquaporins), are hydrophilic; A core of channel proteins/described as R-groups of amino acids 5 (aquaporins) increase permeability of membrane to water; 6 example; e.g. root hairs/small intestine epithelium/nephron 7 role of water in a cell; e.g. solvent/reactant/reaction medium/turgidity <i>or</i> support in plant cell 	(b) 7.0 nm ;			[1]
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[Total: 9]	Rh 2 idea ora An 3 core Ah 4 cha A co 5 (aqu 6 exa e.g. 7 role e.g.	ydrophilic/charged a that few polar molecules pass through the <u>phospholipic</u> for non-polar molecules one pass/repelled e of membrane/phospholipid tails, are hydrophobic ; ydrophobic core nnels (through aquaporins), are hydrophilic ; ore of channel proteins/described as R-groups of amino uaporins) increase <u>permeability of membrane</u> to water ; mple ; root hairs/small intestine epithelium/nephron of water in a cell ; solvent/reactant/reaction medium/turgidity <i>or</i> support	o acids in plant cell	

Page 8	Mark Scheme	Syllabus	Paper
	GCE AS/A LEVEL – May/June 2013	9700	21

6 (a) ref. to mutation(s);

in context of initiating uncontrolled mitosis OR as a consequence of uncontrolled mitosis

proto-oncogenes convert to oncogenes/oncogenes switched on/tumour suppressor genes switched off ;

(cell division is by) mitosis;

formation of, tumour/mass of (unspecialised) cells;

no response to (extracellular/intracellular) signals to control mitosis/AW ;

no contact inhibition/AW;

no cell death/no apoptosis;

immune system does not recognise the cells as foreign and destroys them ; A reference to, not non-self/self

metastasis/described;

- (b) R way in which cancer develops/epidemiological evidenceA beagles for dogs
 - 1 tar painted on skin of, mice/rabbits/rats/(small) mammal, led to development of (cancerous/malignant) tumour;
 - 2 dogs that smoked (plain) cigarettes developed, cancer/tumour;
 - <u>dogs</u> that smoked filter-tipped cigarettes did not develop cancer/tumour;
 A developed precancerous changes
 - 4 control group/dogs, which did not smoke and did not develop, cancer/tumour;
 - 5 AVP;
 - e.g. evidence from any other named mammal
 - e.g. inhaling substances from, tar/tobacco

[max 3]

[max 3]

[max 3]

- (c) similarities
 - 1 <u>all</u> (named) countries, increase and decrease / reach a peak and decrease ;

differences

- 2 peaks/AW, have occurred at different years in at least two countries ;
- 3 <u>all maximum mortality rates are different</u>;
- 4 any comparative, data quote/calculation, with units given at least once;
 - e.g. dates and mortality rates for at least two countries
 - e.g. mortality rates for one country at two different dates

accept a range or a single figure within the ranges given

countries	peak mortality rate	year
USA	53–57	1984–1990
Spain	45-48	1993–1997
Finland	69–71	1970–1973
UK	72–75	1970–1975
Hungary	83–87	1996–2000