

Q1.

Question	Expected Answers	Marks
4 (a) (i)	<p>A transcription; (ignore mRNA synthesis)</p> <p>B translation;</p> <p>C exocytosis; R secretion</p>	[max 3]
(ii)	<p>D (sub unit of) ribosome</p> <p>E Golgi apparatus/body;</p>	[2]
(iii)	F mRNA;	[1]
(b)	<p>active site;</p> <p>(is) specific shape; A complementary/other amino acids are the wrong shape to fit, R same shape</p> <p>only accepts R groups of these two amino acids; R accepts peptide bond</p>	[2]
(c)	<p>correct bond broken (between C-N);</p> <p>involvement of water molecule in breaking the peptide bond shown clearly;</p> <p><u>two amino acids with free groups as follows</u></p> <p>-COOH/-COO⁻ <u>and</u> -NH₂/-NH₃⁺;</p> <p>A from diagram(s).</p>	[3]
		[Total: 11]

Q2.

(b) (i)	6 ;	[1]
(ii)	5 ;	[1]
(iii)	3 ;	[1]
(c)	<p>curve starting at 0;</p> <p>but lower;</p> <p>reaches same plateau but at higher concentration of urea;</p>	[2]
(d)	<p>inhibition is reversible;</p> <p>enzyme is still active;</p> <p>inhibitor fits into active site temporarily;</p> <p>substrate is broken down (reaction does proceed);</p> <p>same end point;</p> <p>just takes longer / reaction is slower with inhibitor;</p>	[max 2]

Q3.

- 2 (a) catalyst ;
active site ;
complementary (to specific substrate) ;
lock and key/induced fit, correctly described ;
enzyme-substrate complex ; **A** E-S complex
lowers activation energy ; **A** E_a
further detail of active site ; e.g. role of **R** groups in active site/catalytic/
binding, site/mechanism to lower E_a [max. 4]
- (b) (i) (idea of) presence of starch ; [1]
- (ii) control ;
to show, enzyme involved/enzyme catalysed reaction/not spontaneous/**AW** ;
enzyme denatured by boiling ; [max. 2]
- (c) **A** starch, broken down/converted to glucose (1-) phosphate/**AW** ; ora for **B**
- A** at pH 6.5/nearly neutral/**AW**, enzyme is active idea/**AW** ;
e.g. ref to optimum at or near 6.5
- (B)** at pH 2.0/acidic qualified, enzyme is inactive idea/**AW** ;
e.g. well away from optimum
further detail e.g. specific effects of pH / bonds affected by hydrogen ions;
- C** enzyme denatured, by boiling/high temperature ;
ref to bonds broken by high temperature ;
- (D)** glucose phosphate gives, no reaction with iodine/negative
result ; **A** no starch/no substrate added gives, no reaction with iodine/negative result
[max. 4]
- (brackets) denote the letter not required for mark**
- [Total: 11]**

Q4.

- 4 (a) (i) **A** transcription ;
B tRNA / transfer RNA ;
C ribosome ; **A** subunit of ribosome / ribosomal subunit
treat 70S / 80S or small / large as neutral
D anticodon ; [4]

(ii) *similarities*

made of amino acids / amino acid monomers / polymer of amino acids **A** protein / polypeptides
 have quaternary structure / have more than one polypeptide chain ;
 four, sub-units / polypeptides ;
 haem / porphyrin / prosthetic group(s) ; [2 max]

difference

(four) sub-units / polypeptides, are identical ;
or
 haemoglobin has, two different, sub-units / polypeptides ;
or
 haemoglobin has alpha and beta polypeptides ;
 (catalase) has active site(s) ; **A** Hb has (oxygen) binding site [1 max]

- (iii) each, sub-unit / polypeptide, has an active site ; [1 max]
 catalase has four, active sites / haem groups ;

- (b) iodine in potassium iodide solution / iodine in KI solution / I in KI solution ; **A** iodine solution
R iodine

Benedict's, solution / reagent ; **A** Benedict's
A Fehling's solution / NaOH and CuSO₄ [2]

treat refs to colour changes as neutral

[Total: 10]

Q5.

- (c) (i) active site ; **ignore** binding / catalytic [1]

- (ii) 1 (shape of) **U** / active site, gives specificity ; **A** *ecf* from (i)
 2 substrate, fits into / binds with, active site / **U** ; **A** *ecf* from (i)
 3 complementary (shape) / matching shape ;
A 'lock and key' / induced fit **R** 'same shape'
 4 further detail of substrate binding to active site ;
 5 forms, enzyme-substrate / E-S, complex ;
 6 causes stress in substrate / AW ;
 7 lowers activation energy / reactions occur at low(er) temperatures ;
 8 not used up in reaction / remain unchanged / reusable ;
 9 high turnover number / catalyse many reactions per unit time ; [4 max]

Q6.

(c) (i) to check that urea is not hydrolysed/broken down, without enzyme ; ora
A there is no reaction without enzyme [1]

(ii) hydrolysis reduces, substrate/urea, concentration ;
urea, hydrolysed/broken down, more quickly in Tube A than in Tube B ;
A ref. to differences in reaction rates

Tube A enzyme can bind with substrate normally/ES complexes forming (at fast rate) ;
ora Tube B
shape of active site complementary to (shape of) substrate/AW ;

Tube B (competitive) inhibitor, occupying/binding at/AW, active site ;
ref. substrate unable to enter active site/AW ;

correct data quote from either column to illustrate ; [4 max]

Q7.

4 (a) (i) hydrolysis / hydrolysing ; I catabolic / digestive R hydrolysis [1]

(ii) to stop the reaction ; R 'stop it working'
by denaturing, the enzyme / sucrase ; R incorrect context
A 'change shape of active site'
to make the Benedict's solution, react / AW ; [2]

(b) *description to max 2*

rate increases to a, maximum / plateau ; A 'levels off' / remains constant
idea that increase in rate slows ;
11.5 (arbitrary units / au) at 80 - 90, g dm⁻³ ; A range 11.4 – 11.6

explanation to max 4 – accept ora where appropriate

substrate concentration is limiting (factor) ;

(at low concentration) *may be given in terms of increasing concentration*
few collisions between enzyme and substrate ;
few, enzyme-substrate / E-S, complexes formed ;
active sites unoccupied ;

(at high concentration / >80 g dm⁻³)
enzyme concentration is limiting (factor) ;
A 'not enough enzyme for substrate to bind to'
maximum number of enzyme-substrate complexes formed ;
active sites , saturated / always occupied ; A ref to V_{max}

[max 5]

[Total: 8]

Q8.

(c) optimum pH or pH at which, lipase / enzyme, works best ;

[1]

- (d) (i) pH, decreases / AW, over time ;
steep decrease / high rate, in first 5 minutes ; **A** faster
less steep decrease / levels out, correct time ref ; **A** slower
correct, manipulation of data / comparative data quote (ref. to both axes) ;
e.g. pH 8 – 7.3 from 0 – 5 min
pH 7.3 – 6.45 from, 50 / 60, min

[2]

- (ii) triglyceride / oil, hydrolysed / broken down / digested, to produce (fatty) acids ;
increasing, acids / H^+ / hydrogen ions, decreases / AW, pH ;

accept, triglyceride / lipid, for substrate throughout

steep decrease

ref. enzyme has high initial turnover rate or high rate of, collision between enzyme and substrate / ES complex formation ;
(because initially) high concentration of, substrate / triglyceride ;

less steep / levelling / plateau,

substrate, being used up / used up / limiting ;
active sites available or fewer enzyme substrate collisions / fewer ES complexes formed ;
ref. presence of hydrogen ions, partial denaturation (less steep) / denaturation (plateau) ;
A description of denaturation

[4]

Q9.

- 4 (a) *this can be answered in the context of penicillinase*

- 1 complementary shape ;
- 2 substrate, fits into / enters / binds to / with, active site ;
A enzyme-substrate complex / ESC
- 3 ref. to specificity ;
- 4 lock and key / induced fit ; **A** description of induced fit
- 5 ref. to temporary bonds form with, active site / R groups (of amino acid residues) ;

[max 3]

- (b) *shown to max 2*

secondary structure ;
 α / alpha, helix ; **R** 'helix' / helical structure unqualified by alpha
 β pleated sheet ;
tertiary structure / folding ; **ignore** 3D shape or structure
globular ;

not shown to max 2

amino acids / primary structure / sequence of amino acids ;
(types of) R groups ;
bonds / named bonds ; **A** peptide
quaternary structure ;
prosthetic group ;

[max 3]

- (c) (i) one lower peak inside line than uncatalysed ;
start and finish at, dotted lines / same energy levels as uncatalysed ; [2]
- (ii) activation (energy) / (energy of) activation ; [1]
- (d) 1 do not prescribe for viral diseases ;
2 only use when necessary / do not overprescribe ;
3 only available on prescription / not available 'over the counter' ;
4 people must, complete the course / take as instructed ;
R take a long course
5 test to find out which is most appropriate antibiotic to use ;
A use most, appropriate / effective, antibiotics
A use narrow-spectrum antibiotics
6 details of sensitivity test ;
7 rotate / AW, antibiotics / use in combination ; R use many antibiotics
8 do not use same antibiotics for humans and animals ; [max 2]

[Total: 11]

Q10.

- (d) (i) 1 active site has, specific / particular, shape ;
2 complementary to substrate ; A substrate fits into active site
3 ref. to (some enzymes) induced fit mechanism ; A described
4 formation of enzyme-substrate complex ; AW
5 lowering, activation energy / E_a ; A detail of how activation energy lowered
e.g substrates held close together for bond formation
facilitates transfer of electrons
places strain on bond(s) to be broken [max 3]
- (ii) 1 loss of tertiary structure / hydrogen bonds broken / ionic bonds broken ;
R if include disulfide or peptide bonds
2 changes shape / substrate unable to fit, active site; A enzyme changes shape so
alters active site
3 loss of / AW, globular structure ;
4 hydrophobic groups to outside of molecule ;
5 hydrophilic groups no longer interact with water / AW ; [max 2]

(e) penalise once for no units

- 1 with no cryoprotectant, enzyme (remains), inactive / AW ;
A at 0 mmol of cryoprotectant, 0% (of maximum) activity
- 2 for both, increasing concentration increases % (enzyme) activity recovered ;
A comparative data quote with ref. to increase *need units*
- 3 trehalose, steeper curve / AW, up to 10 mmol (cryoprotectant) ; ora **R** rapid
- 4 at all concentrations (below 90 mmol), trehalose has higher percentage of (maximum enzyme) activity
- 5 comparative data quote to support either mps 3 or 4 ;
for mp 3 trehalose from 0 to 80% and glycerol from 0 to 10%
- 6 both cryoprotectants can produce, 100% / maximum, (enzyme, activity / recovery) ;
- 7 trehalose produces, 100% (enzyme) activity / full (enzyme) recovery at,
lower concentrations than glycerol / 30 mmol
compared to, 90-100 (mmol) ; *this is also mp 6*
- 8 trehalose more effective than glycerol (up to 90 – 95 mmol cryoprotectant) ;
A trehalose is a better cryoprotectant (than glycerol)

[max 4]

Q11.

- 4 (a) any one valid ;
e.g.(first) appearance of (brown) colour
use of, colour standards/ colour charts
use of colorimeter
time-lapse photography/ video

[1]

- (b) allow catechol for substrate throughout
rate of reaction 0 au, no substrate to act on / AW ;

at substrate concentrations lower than 5mM
substrate (concentration) is limiting (factor in rate of reaction) ;
presence of free active sites/enzyme is in excess ;
few collisions between enzyme and substrate ;
rate increases with substrate concentration as more, active sites can be occupied/E-S
complexes can form ;
one data quote to support response
 V_{max} reached / rate becomes maximum, at 4.5–5 mM substrate concentration ;
rate constant / levels out / AW, from 4.5–5 mmol substrate concentration ;

at substrate concentrations greater than 5mM
enzyme (concentration) becomes limiting (factor) ;
all active sites, saturated/occupied ;
(so) further increase in substrate concentration does not increase rate ;

[max 5]

- (c) (i) curve always lower than that with no inhibitor ; *must be similar shape*
 curve reaches the maximum ; **A** curve heading to maximum [2]
- (ii) PHBA / inhibitor, similar shape to, substrate / catechol ;
 (so) binds to active site ;
 blocks access to substrate / fewer (successful) enzyme-substrate collisions ;
 reduces rate of, reaction / conversion of substrate to product ;
 AVP ;
 e.g. inhibitor has a greater effect on rate at lower substrate concentrations
 V_{max} reached at higher substrate concentrations
 inhibitor forms same interactions with R-groups in active site [max 2]
- (d) enzymes work in a limited pH range / either side of optimum pH rate decreases ;
 (acid so) presence of H^+ ions, partially denatures / denatures (some), enzymes ;
 further detail ; e.g. ref. to breaking ionic or hydrogen bonds
 change of active site shape means substrate can no longer fit ;
 AVP ; e.g. ref. to antioxidant effect of, lemon juice / citric acid / vitamin C [max 2]

[Total: 12]

Q12.

- 4 (a) active site;
specific shape / configuration / conformation (in ref to active
 site);
 complementary to substrate / exact / perfect fit (between
 substrate and active site);
 combine to form enzyme-substrate / ES complex;
 mould around substrate / substrate alters shape of active site
 (induced fit); R. induced fit unqualified
 ref to temporary bonds / named bond; **3 max**
- (b) (i) EcoR1; **1**
- (ii) sticky ends; **1**
- (c) plasmid DNA cut with same restriction enzyme / endonuclease;
 DNA and plasmid mixed together / AW; R. inserted
 ref complementary / base pairing / C and G on sticky ends pair
 up;
 ref to hydrogen bonding;
 ligase forms bonds between sugar and phosphate /
 phosphodiester bonds; **3 max**

[Total: 8]

Q13.

5 (a) measure

disappearance of substrate; **A** measure conc. of substrate

appearance of product; **A** measure conc. of products

2

(b) active over a wide range of pH/AW e.g. whole range/pH 1-9;

increasing activity as pH increases to, optimum/pH 5;

decreasing activity as pH increases, above optimum/> pH 5;

optimum is, between pH 4 to 5.5/pH 5; **A** any figure between 4-5.5

3

(c) (idea of) some enzymes active/all enzymes partly active;

1

low pH equivalent to high H^+ ion concentration;

(so) enzymes (partly) denatured;

reference to tertiary structure affected;

reference to hydrogen/ionic bonds, disrupted/broken;

(so) active sites changed e.g. no longer complementary to substrate;

(detail) affect on R groups of amino acids (in active site);

(therefore) (few) enzyme-substrate complexes formed;

3 max

(c) curve same shape with same optimum (at pH 5 - between 2.0 and 3.0 units on y axis);

lower (starting at pH 1 and finishing at pH 9 without touching x axis);

2

- (e) similar/same shape to, substrate/organic phosphates;
R similar structure

occupies/binds/combines/fits into, active site; **R** inhibitor competes with substrate for active site

so blocking/preventing, entry of substrate; (therefore) decreased rate of product/
 e-s complex/phosphate, formation (at low substrate concentrations);

inhibitor molecules, not permanently bound to active site/bind briefly;

reference effect of concentration of substrate e.g. inhibitor less effective at high concentrations of substrate

A from sketch graph if given

3 max

[Total 14]

Q14.

- (d) *Penalise once if minutes not used*
- (i) 5 minutes. [1]
- (ii) 10 - 11 minutes. [1]
- (e) Fatty acids are released.; [1]
- (f) Steeper decrease from 5 minutes;
 Levels off at pH 7.0.; [2]

[Total: 11]

Q15.

- 3 (a) *max 2 if no reference to data*
 up to substrate concentration of 24 / 25 g dm⁻³, substrate concentration is limiting ;
 24 / 25 to 30 g dm⁻³, another factor is limiting ;
 enzyme concentration / temperature / pH ;
 active sites, not filled up to 24 / 25 g dm⁻³ / all filled above 24 / 25 g dm⁻³ ;
A enzyme working at maximum rate
 ref to collisions between substrate molecules and enzyme ; [3 max]
- (b) same shape starting at the origin and with plateau starting at 24 / 25 g dm⁻³ ;
 lower ; **A** plateau that starts between 7–12 au [2]

- (c) *either*
 competitive inhibitor / effect described in terms of competition ;
 same shape as protein / substrate / elastin ; **A** complementary shape
 to active site **R** same / similar, structure to active site
 fits into active site ;
 blocking entry of substrate / prevents formation of ES complex ;
- or*
 non-competitive inhibitor / described in terms of not competing ;
 fits into, a site other than active site / allosteric site ;
shape of enzyme changes / shape of active site changes ;
 active site no longer complementary shape to substrate ;
- or*
 combines permanently with, active site / other site on enzyme ;
 e.g. by covalent bonding ;
 blocks access to active site / causes tertiary structure to change ;
 prevents formation of ES complex ; [3 max]
- (d) set up different concentrations of substrate ;
 same concentration of inhibitor ;
 measure rate of reaction ;
- if competitive*
 lower rate at low substrate concentrations, but at high substrate concentration will reach the
 same plateau ;
 increasing substrate concentration reverses inhibition ;
- if non-competitive / irreversible*
 lower rate / no activity / does not reach the same rate at high substrate concentrations ;
 increase substrate concentration does not reverse inhibition ;
- accept sketch graphs to show results* [4 max]
- (e) expands / stretches, during inhalation ;
 recoils during exhalation ;
 forces air out of alveoli ;
 prevents bursting of alveoli ; [2 max]
- (f) emphysema ; **A** chronic obstructive, pulmonary / lung disease
A COPD or COLD [1]

[Total: 15]

Q16.

- 3 (a) (i) tertiary (structure) ; **A** 3° [1]
 (ii) secondary (structure) ; **A** 2° , alpha / α, helix [1]
- (b) active site ; **A** catalytic site [1]

- (d) (i) *reject references to time e.g. rapid, slowly*
 as the concentration of, enzyme / lysozyme, increases the percentage of
 bacteria surviving decreases / AW ; **R** if only 1 named
 steep, decline / decrease, 0 to 10 / first two concentrations, for *E. coli* ;
A large percentage difference in *E.coli* surviving at 0 to 10 / first two concentrations
 less steep / more gradual, decline / decrease, from 10 to 150 for *E. coli* ;
 decline / decrease, shallower / less steep from 0 – ,40 / 60 / 70 / 80, for *S. aureus* ;
A small percentage difference in *S. aureus* surviving from 0 – , 60 / 70 / 80
 decline / decrease, more significant / steeper / more abrupt, from 60 / 70 / 80, up to 150
 for *S. aureus* ; **A** large percentage difference in *S.aureus* surviving from 60 / 70 / 80,
 up to 150
 always more *S. aureus* than *E. coli* ; ora
 all bacteria survive with no lysozyme ;
 lysozyme is more effective, at killing / against, *E. coli* / AW ; **A** ora
 all *E. coli* killed, at 150 pmol dm⁻³ (of lysozyme) / at highest concentration ;
 comparative data quote ; *both axes, both curves*
 comparative data quote ; *penalise once for lack of units in both* [4 max]
 (ii) different, polysaccharides / peptidoglycans, in cell walls ;

S. aureus, does not have / has less, polysaccharides / peptidoglycans, in cell wall ;
 ref to shape of active site ;
 ref to shape of, polysaccharide / peptidoglycan (to fit into active site) ;
S. aureus has a capsule / ora ; **A** protective lipids
 AVP ; e.g. *S. aureus* produces inhibitor [2 max]

Q17.

- 2 (a) *marking points are independent*
 iodine in potassium iodide solution / I in KI solution / iodine solution ;
R iodine / iodine test
A if 'solution' not used, but clear that it is a solution
 positive result = (from yellow / red brown to) blue-black / blue / black ;
R blue-black precipitate [2]
 (b) no activity at pH 2.0 **and** pH 9.0, some activity at pH 3.0 **and** 8.0 ;
 optimum between pH 5.5 and 6.5 ; [2]

(c) *description*

- 1 optimum / peak / described, at pH 6.0 ; *allow ecf from graph*
A 'enzyme works best at' / 'most efficient at'
'rate of reaction / activity, is greatest at...'
- 2 low / no, hydrolysis / activity, with **at least one** correct pH ;
- 3 data quote (from table) using time ;
e.g. within 10 minutes / change within 2 minutes / 1/t

explanation to max 4 accept ora

- 4 at optimum pH, most successful collisions ; **A** alternative wording

greater or less than optimum

- 5 high / low, hydrogen ion concentration ;
- 6 enzyme denatured (fully) at / <pH2 or at / >pH9 ;
- 7 partial denaturation / AW, at other stated value(s) of pH ;

at any pH – optimum or sub-optimum

- 8 ref to, hydrogen bonds / ionic bonds ; **R** if other bonds named
- 9 ref to tertiary structure ; **A** ref to allosteric site
- 10 shape of active site ;
- 11 detail of active site ;
e.g. changes to charge on active site / no longer complementary to substrate forms, no /
fewer, enzyme-substrate complexes [5 max]

[Total: 9]

Q18.

- 2 (a) denature, sucrase / enzyme ; **A** deactivate
stop the reaction (in each tube at the same time) ;

idea that Benedict's test requires a high temperature ;
ref to reducing sugars ;

[2 max]

- (b) starts at, the origin / 5 g dm^{-3} , increases to $45\text{--}55 \text{ g dm}^{-3}$;
constant from 80 to 100 g dm^{-3} ;

[2]

(c) description	conc	rate*
	5	0.0036
	10	0.0069
	15	0.0105
	20	0.0133
	50	0.0213
	100	0.0222

penalise lack of units once only

- 1 increase in rate of hydrolysis to approx 50 g dm⁻³ ;
A decrease in time taken to approx 50 g dm⁻³ / correct rate calculations* to show an increase
 - 2 remains constant / plateaus / levels out / AW, from approx 50 g dm⁻³ to 100 g dm⁻³ ;
- explanation to max 4*
- 3 (sucrase / enzyme) hydrolyses / breaks , glycosidic bonds ;
 - 4 forming, reducing sugars / glucose / fructose ;
 - 5 *idea that* concentration is the limiting factor, at low concentration of, sucrose / substrate ;
 - 6 (at low concentrations) active sites, unoccupied / available ;
A as concentration increases, more active sites are occupied / more enzyme-substrate complexes formed / AW
 - 7 at higher concentrations all active sites, occupied / saturated / AW ;
R enzymes for 'active sites'
 - 8 substrate, in excess / AW ;
 - 9 V_{max} reached / working at maximum rate ;
- idea that*
- 10 at higher concentrations, enzyme / sucrase, is the limiting factor ;

[5 max]

[Total: 9]

Q19.

- 3 (a) spherical / ball-shaped / AW ; **A** round(ed) / circular has tertiary structure ; **R** 3D hydrophilic / polar, (R) group(s), on outside / face to watery exterior ; hydrophobic / non-polar, (R) group(s), in centre ; water soluble ; [max 3]
- (b) (i) *idea that plant cell walls and fungal cell walls have different components* fungal cell walls made of, glucans / chitins / fungal cellulose / different components to plant cell walls ; **A** peptidoglycan / murein **A** plant cell walls contain cellulose, but fungi do not *idea of specificity in context of question* enzymes are specific ; **A** specificity explained e.g. both substrates not complementary / shape of active site specific to one substrate [2]

- (ii) 1 (at optimum pH) maximum / peak, activity ; **A** most efficient / works best
 2 above / below, optimum, activity declines ;
A description / graph sketched with pH and rate / activity
 3 changing pH changes hydrogen ion concentration ;
 4 hydrogen / ionic, bonds (between amino acids), break / disrupted ;
 5 hydrogen / ionic, bonds, important in maintaining shape of, tertiary structure / active site ;
R 4 and 5 if refer to disulfide, hydrophobic interactions, peptide
 at *sub-optimum pH*
 6 active site / tertiary, shape altered ; **A** enzyme denatured
 7 charges at the active site may be affected ;
 8 further detail ; e.g. transfer of electrons may not be possible
 9 the substrate may be altered by pH changes ; **R** cell wall unqualified
 10 (therefore) substrate no longer fits / ES complexes not formed ; [max 3]

Q20.

- 3 (a) (i) glucose and fructose ; *ignore monosaccharides* [1]

- (ii) 1 active site, gives specificity ; **A** specific active site
ignore ref to specific substrate
 2 substrate binds with active site or enzyme-substrate / E-S, complex forms ;
 3 complementary (shape) / substrate fits into active site ; **A** 'lock and key'
A matching shape
R 'same shape'
 4 induced fit / described ;
 5 further detail of substrate and active site ; e.g. binding by hydrogen bonding,
 e.g. transfer of electrons
 6 lowers activation energy / described e.g. causes strain in substrate / AW ;
A Ea
 7 breaks glycosidic bond ;
 8 glucose and fructose / products, no longer fit / AW ; [max 4]

- (iii) non-competitive (inhibition) ;
 irreversible (inhibition) ; [max 1]

- (b) (i) idea of, hydrolysis / product formation / further metabolism, lowering sucrose concentration (in, companion cells / sink cells) ;
 maintains, concentration / diffusion, gradient (between phloem sieve tubes and, companion cells / sink cells) ;
 to remove sucrose from the phloem (sieve tubes) ;
 AVP ; e.g.ref. easier transport of, glucose / fructose, through membranes ; [max 2]

- (ii) ref. facilitated diffusion out / may be lost from cells ;
 products / glucose / fructose, are soluble / AW ;
 (so) will lower the water potential / water potential becomes more negative ;
 causes water to move into cells by osmosis ; **A** osmotic, problems / stress
 reactive / easily metabolised, qualified ; e.g. so interferes with, other metabolic
 processes / cell chemistry **A** more reactive than starch [max 3]

[Total: 11]

Q21.

- 2 (a) (i) tangent drawn on the graph as close as possible to time 0 e.g. 1.6 / 6 ;
0.27 ;

accept
correct volume of gas e.g. $\frac{2.5}{10}$ $\frac{4.3}{20}$ }
 stated time, up to and including 20 secs
 or
 tangent drawn on the graph before 20 secs $\frac{5.8}{20}$ } ;

correct calculation ; e.g. 0.25 (cm³ s⁻¹), 0.22 (cm³ s⁻¹) **A** 0.215
 e.g. 0.29

award one mark if the time is 21–40 s but the calculation is completed correctly [2]

- (ii) *accept hydrogen peroxide or reactant for substrate*
 initially high concentration of substrate so, rate of reaction high / enzyme activity at
 a maximum / AW ;
 (rate slows as) concentration of substrate decreases ; **A** substrate being used up
 no further change in volume / AW, reaction has stopped ;
 correct data quote to support explanation(s) ;

correct ref. to number of (successful) collisions;
 correct ref. to enzyme-substrate complexes / active sites occupied; [max 3]

- (b) 1 (copper ions act as enzyme) inhibitor ; **R** competitive inhibitor
 2 non-competitive (inhibition) ;
 3 (non-competitive) inhibitor / Cu²⁺, combines with enzyme at site other than active
 site ;
 4 active site shape / tertiary structure / 3D shape, changes ;
 5 active site no longer accepts substrate / enzyme-substrate complex not formed /
 AW ;
 6 independent of substrate concentration / increase in substrate concentration has
 no effect / AW ;
 7 comparative rates quoted from Fig. 2.2 ;
 e.g. max, 3.25 cm³ s⁻¹ v 0.22–0.25 cm³ s⁻¹
 8 AVP ; e.g. actual rate depends on the relative concentration of inhibitor / AW
 V_{max} not reached
 effect of ion presence on tertiary structure [max 4]

- (c) enzymes are proteins ;
 ref. transcription ; *accept description* }
 ref. to mRNA ; } *in correct context*
 ref. translation ; *accept description* }
 ref. to further folding / glycosylation / modifying, in, RER / Golgi body ; [max 3]

[Total: 12]

Q22.

- (e) 1 increasing concentration of ara-ATP decreases enzyme activity;
can be comparison between 0 and 5 / 20 or between 5 and 20
A ref. to rate of DNA synthesis for enzyme activity
- 2 ara-ATP acting as an inhibitor;
- 3 substrate unable to bind with active site / fewer enzyme-substrate complexes (formed);
- 4 further detail;
for either competitive
 e.g. competes with substrate for (binding to) the active site / similar, structure / shape, as substrate or complementary shape to active site
or non-competitive inhibition
 e.g. binds to site other than active site / changes shape of active site [max 3]

Q23.

- (b) 1 nitrogen and hydrogen / substrates, bind to / AW, active site ;
- 2 enzyme-substrate complex (forms) ;
- 3 ref. lock and key / induced fit, mechanism ;
- 4 activation energy of reaction is lowered ;
- 5 example of how activation energy lowered ;
 e.g. strain on (triple) bond of, N₂ / (di)nitrogen
A bond broken between nitrogen (atoms)
 nitrogen and hydrogen ions held close together for bond formation
 transfer of electrons
 alternative pathway
- 6 product / NH₄⁺, leaves active site ;
- 7 ATP, required / used / provided from respiration ;
- 8 ref. anaerobic conditions for enzyme action ;
- 9 suggestion as to use of, vanadium / molybdenum, in active site ;
 e.g. act as cofactor / coenzyme
 transfer of, electrons / protons [max 4]

Q24.

(b) (i) 47.5 °C ;

[1]

(ii) *accept activity for relative activity throughout
accept manipulated data quotes and penalise once for, incorrect / no, units*

Fig. 2.2 (relative activity of enzyme at different temperatures)

1 as temperature increases, activity increases up to, optimum / 47.5 °C (*allow ecf from (i), then decreases ;
A peaks (for increase then decrease)*)

2 activity increases from 30 °C to 47.5 °C, then decreases to 70 °C ; *also mp 1
or
increase or decrease, described with comparative data (activity
and temperature compared with another activity and temperature)*

3 at higher temperatures (compared to most others) enzyme still active ;

4 high optimum temperature (compared to most other enzymes) ;

Fig. 2.3 (stability over time for enzyme maintained at different temperatures)

5 enzyme becomes less stable over time ;

A activity decreases over time

A description if at least two temperatures described

6 data quote to support ; *activity at two times for any one temperature
if time 0 or 'start', then assume 100% relative activity
if 100%, assume time 0*

- 7 (over the time period) the lower the temperature, the more stable the enzyme ; ora
A enzyme has higher activity at the lower temperatures
A stated temperatures (at least two) to illustrate the point
 e.g. 28 °C higher activity than 40 °C throughout
A 28 °C, highest activity / enzyme most stable (throughout)
- 8 data quote to support ; *temperatures and (relative) activity (with one time)*

discussion points

- 9 AVP ; ;
- 10 e.g. Fig 2.2
 reason for increasing activity up to optimum / decrease after optimum
 e.g. ref. collisions, kinetic energy increase e.g. denaturation at 60–70 °C
R denaturation at 50 °C (but **A** denaturation begins)
 suggested reason for higher optimum temperature e.g. more bonds

Fig. 2.3

(suggests that) more molecules become, denatured / inactive, as time progresses
 greater stability / higher activity, at 40 °C than 37 °C between 40–50 hours

Fig. 2.2 and 2.3

optimum temperature for activity not most stable temperature
 steep decrease in stability at 60 °C in a short time as (nearly complete) denaturation occurs *allow once only*
 commercial application e.g. if hydrolysis occurs over a longer time period,
 better to use a lower temperature than optimum [max 5]

Q25.

(c) *answers may be general or in the context of phloem transport*

active site (with shape) complementary to substrate ;
A description in terms of lock and key (either way round)
I structure
 induced fit / described ;
 substrate binds to active site / enzyme-substrate complex forms / ESC forms ;
 ref. to specificity of enzymes ;
 activation energy of reaction is lowered ;
 example of how activation energy lowered ;
 e.g. reactants held close together for bond formation
 transfer of electrons
 strain on bonds
 alternative pathway
 holding the substrate in such a way that the bonds needed to be broken are exposed
 product released from, enzyme / active site ;
A enzyme can be used again / enzyme unchanged at end of reaction [max 3]

Q26.

(b) (i) *data quote may help to decide if mp2 is matched*
units must be used at least once in the answer to award mp3

- 1 as retention time increases percentage of cell wall material digested increases / positive correlation ;
A 'time for digestion' / reverse relationship
R directly proportional
- 2 results scattered / not all animals fit the pattern / varying percentages for the same retention time ; *not just a data quote*
- 3 data quote with units (% and h) using both axes ;
e.g. (highest percentage) 65% at 78 hours
(lowest percentage) 35.5 ± 0.5%, 35 hours
- 4 no retention time shorter than 35 hours and none longer than 88 hours ;
A lowest / shortest and highest / longest
A reverse relationship **A** 'time for digestion'
- 5 none of the (24) herbivores can digest the cell wall material completely ;
A no more than 65% is digested
not just a data quote

[max 3]

