Enzymes

Mark Scheme 3

Level	International A Level		
Subject	Biology		
Exam Board	CIE		
Topic	Enzymes		
Sub Topic	Enzymes		
Booklet	Theory		
Paper Type	Mark Scheme 3		

Time Allowed: 54 minutes

Score : /45

Percentage : /100

Grade Boundaries:

A*	Α	В	С	D	E	U
>85%	'77.5%	70%	62.5%	57.5%	45%	<45%

1 (a (bacterial urease converts) urea \rightarrow ammonia;

ammonia → nitrite; *Nitrosomonas*; nitrite → to nitrate; *Nitrobacter*;

nitrification;

oxidation / chemosynthesis;

[max 3]

(b) 6

[1] [1]

(ii) ; (iii) ;

[1]

(c) curve starting at 0;but lower;reaches same plateau but at higher concentration of urea;

[2]

(d) inhibition is reversible; enzyme is still active; inhibitor fits into active site temporarily; substrate is broken down (reaction does proceed); same end point; just takes longer / reaction is slower with inhibitor;

[max 2]

[Total: 10]

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— TUITION —

2 (a) Two correct letters required for a mark for each column if list given; mark first 2 letters.

Alcohol	Caffeine	Nicotine	Heroin
U	S	S	C
V	T	T	Υ
Υ	Z	W	W
Z		X	
		Z	

4

- decrease in response to drug/effect of drug becomes less (intense);
 decrease in sensitivity of receptors/more receptors are made;
 drug is metabolised/becomes part of body's metabolism; more
 drug necessary to achieve the same effect/sensation/euphoria; max 2
- (c) award marks from any <u>annotated</u> diagrams
 Either

inhibitor fits site other than active site/allosteric site; tertiary/3D structure or shape changes/any two bonds mentioned break; (ionic, van der Waals, hydrophobic, hydrogen, disulphide, covalent)

active site changes shape;

substrate no longer fits/binds/active site no longer complementary to substrate/E.S. complex not formed;

<u>or</u>

inhibitor fits <u>permanently</u>/irreversibly into active site; substrate can no longer bind/substrate blocked/no E.S. complex formed;

increasing substrate has no effect;

max 3

<u>Either</u> mark scheme as appropriate – <u>do not mix</u> marking points from both mark schemes

[Total 9]

- 3 (a 1 glucose/substrate, is not complementary/is partially complementary, to active site :
 - 2 enzyme/active site, changes shape/moulds around/fits around, when substrate, enters/binds; **R** if substrate/glucose changes
 - 3 stronger binding of substrate to active site;
 - further detail; e.g. becomes complementary to/fits more tightly to, glucose/substrate interaction of, functional groups/R–groups/side-chains formation of (named) bond but not disulfide or peptide bond

[max 3]

- (b) 1 (competitive) inhibitor has, same/similar, shape to substrate;
 - 2 inhibitor does not induce the same change in, 3D shape/tertiary structure/active site (as the substrate);
 - 3 (so inhibitor) less likely to bind (successfully) in active site;
 - 4 idea that because it does not have same functional groups (in same positions)/AW;
 - 5 in lock and key the inhibitor, fits directly into/is complementary to/binds to, active site;

[max 2]

(c) enzymes/hexokinase, denatured; all enzymes molecules are partially denatured/some enzyme molecules are denatured;

changes/disrupts/loss of (specific shape/structure) active site; **A** no longer complementary to, glucose/substrate breakage of, ionic/hydrogen, bonds; **R** disulfide/peptide bonds

idea that loss of structure makes E–S complex formation more difficult/fewer E–S complexes are formed/substrate does not fit into active site;

[max 3]

(d) (i) accept ora

active transport requires, ATP/energy (whereas facilitated diffusion does not);

active transport moves substances against the concentration gradient (whereas facilitated diffusion moves substances down the concentration gradient);

active transport uses only carrier proteins (whereas facilitated diffusion uses both carrier and channel proteins);

A active transport can involve cotransport but facilitated diffusion does not

[max 2]

(ii) too large/too big; R 'it is a big molecule' unqualified

polar/charged, so cannot pass through hydrophobic region of membrane; **A** fatty acid tails for hydrophobic no, specific/AW, protein, in membrane/carrier/channel; e.g. AW = no protein for G–6 AVP; e.g. gated channels are closed

[max 2]



4	(a	(i)	G ;	[1]
		(ii)	B/C;	[1]
		(iii)	A/F;	[1]
		(iv)	В;	[1]
		(v)	D;	[1]
	(b)	1 2	nitrogen and hydrogen/substrates, bind to/AW, active site; enzyme-substrate complex (forms);	
		3	ref. lock and key/induced fit, mechanism;	
		4 5	activation energy of reaction is lowered; example of how activation energy lowered; e.g. strain on (triple) bond of, N_2 / (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/NH ₄ ⁺ , leaves active site ;	
		7 8	ATP, required/used/provided from respiration; 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	4]

- (c) 1 concentration of all the ions is greater in the root tissue than in the solution; ora
 - 2 comparative data quote;

according to these data

- 3 (so) ions will not diffuse into the root tissue;
 - A if (facilitated) diffusion only, initially/till equilibrium reached
- 4 (so) active transport; A active, uptake/pumping I facilitated diffusion
- 5 use ATP; **A** energy **R** ATP energy
- 6 move ions, against concentration gradient/from low to high concentration;
 A diffusion gradient
- 7 use, membrane/integral/intrinsic/transmembrane/transport/carrier, proteins; R channel proteins
- 8 are specific/have specific binding sites;
- 9 involve, conformational/shape, change;
- 10 comparative data quote to suggest that some ions are pumped more than others; e.g. steepest gradients for K^+ and SO $_4^-$
- 11 phospholipid bilayer/hydrophobic core (of cell surface membrane) is impermeable to ions;
- 12 so ions cannot diffuse out/(membrane) proteins only allow inward flow of ions;
- 13 AVP ; e.g. suggestion of differing numbers of specific membrane proteins to explain observation of mp 10 [max 5]

[Total: 14]

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