Mark schemes

Q1.

(a)

1. A = Attachment protein; Accept gp41 /gp140 /gp120/CD4/ glycoprotein Accept antigen Ignore receptor protein

2. B = Capsid

OR

Capsomere

OR

Protein;

- (b) 1. Attachment proteins attach to receptors on helper T cell/lymphocyte;
 - 2. Nucleic acid/RNA enters cell;
 - 3. Reverse transcriptase converts RNA to DNA;
 - 4. Viral protein/capsid/enzymes produced;
 - 5. Virus (particles) assembled and released (from cell);

4 max

2

[6]

Q2.

(a) 1. Cell ingests/engulfs the antibody/ADC

OR

Cell membrane surrounds the antibody/ADC (to take it inside the cell);

Accept endocytosis for ingest/engulf

- 2. Lysosomes fuse with vesicle/phagosome (containing ADC);
- 3. Lysozymes breakdown/digest the antibody/ADC to release the drug; Accept hydrolytic enzyme for lysozyme

3

- (b) 1. ADC will bind to non-tumour/healthy cells; Reject reference to active site
 - 2. Cause death/damage of non-tumour/healthy cells

OR

		Cause damage to other organs/systems;	2	
(c)	Correct answer for 2 marks, 9.2 x 10 ⁻⁵ ;;			
	Accept for 1 mark,			
	0.046 (correct mass injected into 23g mouse)			
	0.00	0092 (correct answer but not in standard form)	2	
(d)	Mice	died		
	OR			
	Not e	ethical to continue;	1	
(e)	1.	Tested on other mammals to check for safety/side effects; Accept named mammal, eg rat		
	2.	Tested on (healthy) humans to check for safety/side effects; Accept: Tested on (healthy) human tissue/cells to check for no side-effects		
	3.	See if repeat doses stop the tumours regrowing (in Group J);		
	4.	Investigate different concentrations of ADC to find suitable/safe dosage;	2 max	[10]
				[10]
Q3.				
(a)	1.	RNA converted into DNA using reverse transcriptase; Reject 'messenger' or 'm' before RNA		
	2.	DNA incorporated/inserted into (helper T cell) DNA/chromosome/genome/nucleus;		
	3.	DNA transcribed into (HIV m)RNA; Accept descriptions of transcription		
	4.	 (HIV mRNA) translated into (new) HIV/viral proteins (for assembly into viral particles); Accept descriptions of translation Accept named viral protein, eg capsid Reject viral cells 	4	
(b)	<u>For</u>			

1. (There appears to be) no virus/ HIV(-1)/RNA/DNA, so could be a

cure/effective; Max 4 for reasons for or against Ignore virus is killed

2. No CCR5/receptor, so not get HIV(-1) in the future

OR

No CCR5/receptor, so nothing for HIV(-1) to bind to; Reject less CCR5/less HIV(-1) bind

- 3. Only one transplant/BSCT needed (shown by patient Q)
- 4. Would not need (daily) ART (16 months after BSCT);

<u>Against</u>

5. Don't know if chemotherapy/radiotherapy is needed

OR

Do not know if BSCT alone would be effective;

OR

Do not know which treatment is having the effect

OR

Could be due to chemotherapy/radiotherapy;

Accept: chemotherapy/radiotherapy is toxic/harmful/has side-effects

- 6. Only for HIV-1; Accept: Might not work in other types of HIV
- 7. Don't know if it would work in all people

OR

Only worked/tried in 2 cases;

8. Might not be long term

OR

Only 18 months;

- 9. HIV-1 may mutate and be able to bind to a different receptor (on $T_{\rm H}$ cells);
- 10. Might be a lack of (suitable stem cell/BSCT) donors; Accept stem cells/BSCT (might be) rejected

Q4.				
(a)	1.	Engulfs; Accept endocytosis OR Description Ignore 'taken in'		
	2.	Forming vesicle/phagosome and fuses with lysosome;		
	3.	Enzymes digest/hydrolyse; Accept lysozymes for 'enzymes'	3	
(b)	1.	(Cells from) other organisms/transplants;		
	2.	Abnormal/cancer/tumour (cells);		
	3.	(Cells) infected by virus; Accept 'own cells' if autoimmune response suggested Accept APCs Accept non-self		
		2 max	C C	
(c)	'X' v	'X' written at either or both ends of Y shape;		
(d)	Joir	ns two (different) <u>polypeptides;</u> Accept holds/attaches Accept 'prevents polypeptide chains separating' 1		7]
Q5.	1	Less/ne entitledu producedu		
(a)	1.	Less/no antibody produced;		
	2.	(Because HIV) destroys helper T cells; Accept 'reduces number' for 'destroys'		
	3.	(So) few/no B cells activated / stimulated		
		OR		
		(So) few/no B cells undergo mitosis/differentiate/form plasma cells;	3	
(b)	Not	t effective in treating AIDS because		
	1.	Number of T cells < 200 at 4 <u>months;</u> Max 4 if not one of 9. or 10.		

Accept 3.5 - 5 months

Reject day/week only once

2. (So) drug is not effective

OR

AIDS symptoms occur;

3. Does not remove (all) HIV (particles)

OR

Number of HIV (fairly) constant/stable

OR

(Slight) increase in HIV (over 16 months);

- 4. No stats test;
- 5. Only shows (results over) 16 months;
- 6. Only one person;
- 7. Unknown side effects (of drug);
- 8. No control group;

Effective in treating AIDS because

9. Number of T cells > 200 after 5 months

OR

Number of T cells increasing after 4 <u>months;</u> Reject day/week only once Accept any month after 5 months OR 'in the long term'

10. So drug is effective

OR

AIDS symptoms relieved/removed;

5 max

[8]

Q6.

- (a) 1. Mutation in the viral DNA/RNA/genome/genetic material; Accept named examples mutations
 - 2. Altered (tertiary structure of the) viral attachment protein; Accept 'antigen' for 'attachment protein'

Accept causes antigenic variability

Allows it/attachment protein/virus to bind (to receptors of other species);

Accept descriptions of binding eg is complementary

(c) 1. (The scientists) could identify proteins (that derive from the genetic code)

OR

(The scientists) could identify the proteome;

2. (They) could (then) identify potential antigens (to use in the vaccine); Reject if answer suggests vaccine contains antibodies

2

2 max

- (d) 1. B cell (antibody) binds to (viral) specific/complementary receptor/antigen; Accept <u>B cell</u> forms antigen-antibody complex
 - 2. B cell clones

OR

B cell divides by mitosis;

- <u>Plasma cells</u> release/produce (monoclonal) <u>antibodies</u> (against the virus);
- 4. (B/plasma cells produce/develop) memory cells; Accept B cell undergoes clonal selection/expansion

3 max

Q7.

- (Antibodies with the) same tertiary structure
 OR

 (Antibody produced from) identical/cloned plasma cells/B cells/B lymphocytes;
 Accept in context of single plasma/B cell/B lymphocyte
 Reject: genetically identical antibody
- (b) Accept any one suitable use, eg

Targets/binds/carries drug/medicine to specific cells/antigens/receptors **OR**

Block antigens/receptors on cells;

Accept cancer/diseased cells (as a specific cell). Ignore medical diagnosis/pregnancy/ PSA/ELISA test. 1

(c)			
(0)		Ignore mixing of direct or indirect ELISA Accept annotated diagram(s).	
	1.	(First) antibody binds/attaches /complementary (in shape) to antigen;	
	2.	(Second) antibody with enzyme attached is added;	
	3.	(Second) antibody attaches to antigen; Accept (second) antibody attaches to (first) antibody (indirect ELISA test).	
	4.	(Substrate/solution added) and colour changes; Only award if enzyme mentioned. 4	[6]
Q8. (a)	1.	Bind to antigen	
(a)	1.	OR	
		Are markers; Accept opsonin for 'marker'	
		Accept form (antibody-antigen) complexes/are complementary to antigen	
	2.	(Antibodies) cause clumping/agglutination	
		OR Attract phagocytes;	
		Reject clotting	
(b)	Corre	ect answer for 2 marks 110/111/111.1;;	
	Acce	pt for 1 mark, correct readings from graph (5.1 and 2.1) 2	
(c)	1.	Mean (antibody concentration) increases;	
	2.	1 st injection protects some mice/1 mouse/2 mice	
		OR 1 st injection causes primary (immune) response/memory cell production;	
		2. and 3. Accept correct reference to number of unprotected mice	
	3.	2 nd /3 rd injection protects most/all mice OR	
		2 nd /3 rd injection causes secondary (immune) response OR	
		2 nd /3 rd injection uses memory cells;	
	4.	Because antibody at/above protective level/2.1; Accept converse	

[10]

5. Antibody decreased (rapidly after 3rd injection); 6. No mice protected after 180 days OR Injections/vaccine not effective in long term OR Booster required (when antibody below protective level/after 120/180 days); 7. One mouse (after first injection) has big response/already had meningitis/antigen; 4 max (d) Mark as pairs, 1 and 2, 3 and 4 Accept for inject, introduce, give, use 1. Inject vaccine (again)/meningitis antigen/ inactive antigen/dead/living bacteria/ pathogen/use a booster; Must refer to antigen or cell, 'disease' or 'meningitis' is not enough 2. (Memory cells present if) faster/more rapid production/higher concentration antibody (than 1st injection) OR Immune response is quicker (than 1st injection) OR Symptoms do not develop; Accept converse Must be a comparison 3. Add enzyme attached to (second) antibody against memory cell; 4. Colour change shows memory cell present; Ignore to detect (meningitis) antibodies 2

Q9.

(a) 1. (Antivenom/Passive immunity) antibodies bind to the toxin/venom/antigen and (causes) its destruction; For 'bind' accept 'attach', ignore 'attack'. For 'destruction of toxin' accept agglutination or phagocytosis. Ignore reference to antibodies 'neutralising toxin/stopping damage' Reject reference to 'killing' toxin/venom.
2. Active immunity would be too slow/slower; Accept 'passive immunity is fast<u>er</u>', not simply 'passive immunity is fast'.

		2
(b)	1.	May be different form of antigen/toxin (within one species) OR
		Snakes (within one species) may have different mutations/alleles;
	2.	Different antibodies (needed in the antivenom) OR
		(Several) antibodies complementary (to several antigens); No mark points are available for answers related to collecting venom from different species of snake. 2 max
(c)	1.	Horses because more antivenom/antibodies could be collected (as more blood collected);
	2.	4550 (cm ³) v 26 (cm ³) (blood collected);
		Accept 175 rabbits needed to (collect the volume of blood from) one horse.
		2
(d)	1.	(So) the animal does not suffer from the venom/vaccine/toxin;
	2.	(So) the animal does not suffer anaemia/does not suffer as a result of blood collection;
	3.	(So) the animal does not have pathogen that could be transferred to humans;
		Accept 'To fulfil licence/legal requirements'.
		Accept '(So) the animal does not have pathogen that could result in it producing other antibodies (not wanted in the antivenom)'.
		For 'pathogen' accept correct form of pathogen.
(e)	1.	B cells specific to the venom reproduce by mitosis;
		Accept in context of primary or secondary immune response.
		Credit idea of specificity if given once in relation to T or B cell.
		Accept a description for specificity.
		Accept 'clone' for 'reproduce by mitosis'.
		'Clonal selection of B cells' = MP1.
	2.	(B cells produce) plasma cells and memory cells;
	3.	The second dose produces antibodies (in secondary immune response) in higher concentration and quickly OR

The first dose must be small so the animal is not killed; Accept 'a lot of antibody' for 'higher concentration of antibody'.

[10]

Q10.

- (a) 1. Person (infected with HIV) has HIV DNA (in their DNA);
 - 2. New HIV (particles) still made;
 - 3. (AZT) inhibits reverse transcriptase;
 - (AZT) stops these (new HIV particles) from forming new HIV DNA;
 OR

Slows / stops replication of HIV;

- 5. Stops destruction of more / newly infected T cells;
- 6. So immune system continues to work (and AIDS does not develop);
 - 4. Context is important
 - 4. Allow slows / stops (re)production of HIV
 - 4. Reject (AZT) prevents DNA replication

4 max

- (b) 1. Slows / stops the development of AIDS;
 - 2. Because HIV **resistant to AZT** is damaged / destroyed / prevented from replicating (by other drugs);

OR

- 3. AZT continues to work as a drug;
- 4. Because HAART prevents the spread of AZT-resistant HIV to rest of the human population;

OR

- 5. No new HIV particles made;
- 6. Because HAART might interfere with viral protein synthesis;

Mark in pairs.

Do not mix and match.

- 2. Neutral HIV killed
- 2. Accept other drugs prevent HIV resistant to AZT from infecting new / more cells
- Accept blocks transcription / translation / synthesis of lipid envelope / aspect of viral structure

4 max

Q11.

- (a) 1. Phagosome / vesicle fuses with lysosome;
 - 2. (Virus) destroyed by lysozymes / hydrolytic enzymes;
 - 3. Peptides / antigen (from virus) are displayed on the cell <u>membrane;</u>
 - 1. Accept vacuole fuses with lysosome
 - 1. Reject virus fuses with lysosome

3

- (b) 1. Helper T cell / TH cell binds to the antigen (on the antigenpresenting cell / phagocyte);
 - 2. This helper T / TH cell stimulates a specific B cell;
 - 3. B cell clones

(c)	4.	 OR B cell divides by mitosis; (Forms) <u>plasma cells</u> that release antibodies; 1. and 2. 'Helper' is required once only. 2. Accept 'This (helper) T cell stimulates a competent B cell' 'T cell stimulates B cell to undergo clonal selection'. This statement achieves mp2 and mp3. 3 m 	ıax	
(-)	2.	This results in the destruction of the (human) cells / collagen; 2. Ignore 'attacks'	2	[8]
Q12.				
(a)	1.	Foreign protein; Accept glycoprotein / glycolipid / polysaccharide		
	2.	(that) stimulates an immune response / production of antibody;	2	
(b)	1.	A protein / immunoglobulin specific to an antigen;		
	2.	Produced by B cells		
		OR		
		Secreted by plasma cells;	2	
(c)	1750)(%);	1	
(d)	1.	Sample 1 / before vaccination no antibody released because patien not yet encountered vaccine / antigen / virus; Accept 'produced' for 'released'	ts	
	2.	(Sample 2 / primary response / after first dose) activation / clonal selection / expansion of <u>B cells</u> into plasma cells;		
	3.	Plasma cells <u>release</u> antibodies;		
	4.	(Sample 3 / secondary response / after second dose) memory cells produce more antibodies / produce antibodies more quickly;	4	[9]

Q13.

- (a) 1. Antigen / epitope on surface of *N. meninigitidis* / bacterium binds to surface protein / surface receptor on a (specific / single) B cell.
 If answered in context of T cell, allow Antigen binds to (specific / single) T cell
 - 2. (Activated) B cell divides by mitosis / produces clone; If answered in context of T cell, allow (Activated) T cell releases cytokine.
 - 3. (Division) stimulated by cytokines / by T cells; If answered in context of T cell, allow (Cytokine) stimulates production of plasma cells;
 - 4. B cells / plasma cells release antibodies;
 - 5. (Some) B cells become memory cells;
 - 6. Memory cells produce plasma / antibodies faster

6

Q14.

- (a) 1. Vaccine contains antigen from pathogen;
 - 2. Macrophage presents antigen on its surface;
 - 3. T cell with complementary receptor protein binds to antigen;
 - 4. T cell stimulates B cell;
 - 5. (With) complementary antibody on its surface;
 - 6. B cell secretes large amounts of antibody;
 - 7. B cell divides to form clone all secreting / producing same antibody.

5 max

- (b) 1. Active involves memory cells, passive does not;
 - 2. Active involves production of antibody by plasma cells / memory cells;
 - 3. Passive involves antibody introduced into body from outside / named source;
 - 4. Active long term, because antibody produced in response to antigen;
 - 5. Passive short term, because antibody (given) is broken down;
 - 6. Active (can) take time to develop / work, passive fast acting.

5 max

[10]