



# **A-Level Biology**

## **Immune System**

### **Mark Scheme**

**Time available: 66 minutes**

**Marks available: 51 marks**

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## Mark schemes

1.

- (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,  $9.2 \times 10^{-5}$ ;;

Accept for 1 mark,

0.046 (correct mass injected into 23g mouse)

0.000092 (correct answer but not in standard form)

2

- (d) Mice died

**OR**

Not 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]

- 2.** (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) 'X' written at either or both ends of Y shape; 1
- (d) Joins two (different) polypeptides;  
*Accept holds/attaches*  
*Accept 'prevents polypeptide chains separating'* 1
- [7]**
- 3.** (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*
3. Allows it/attachment protein/virus to bind (to receptors of other species);  
*Accept descriptions of binding eg is complementary* 2 max

(b) For **one** mark, accept any **two** of the following:

- The polymerase chain reaction
- Genetic/DNA fingerprinting
- (Gel) electrophoresis
- DNA/genome sequencing;

*Accept PCR for polymerase chain reaction*

*Accept autoradiography*

*Accept DNA hybridisation*

*Accept compare DNA/base sequence for 'DNA sequencing'*

*Ignore compare mRNA base sequence*

*Ignore compare amino acid sequence*

*Ignore DNA probes*

1

(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

(d) 1. B cell (antibody) binds to (viral) specific/complementary receptor/antigen;

*Accept B cell forms antigen-antibody complex*

2. B cell clones

**OR**

B cell divides by mitosis;

3. Plasma cells release/produce (monoclonal) antibodies (against the virus);

4. (B/plasma cells produce/develop) memory cells;

*Accept B cell undergoes clonal selection/expansion*

3 max

[8]

4.

(a) (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*

1

- (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)

*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]**

**5.**

- (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 er', 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.  $4550 \text{ (cm}^3\text{)} \div 26 \text{ (cm}^3\text{)}$  (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.*

1 max

- (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'.*

3

**[10]**

6.

- (a) 1. Virus can't bind (to receptor)/ can't enter cells;  
2. So can't be replicated/ multiply;  
*Accept can't reproduce*  
3. So, doesn't damage cell(s)/tissues (and cause symptoms);  
*Accept no toxins released*

2 max

- (b) 1. Antigen/glycoprotein on Ebola binds to/stimulates (a specific) B cell;  
*Accept correct reference to stimulation of B cells by T cells*  
2. (Binding causes) replication/cloning of B cell;  
*Accept replication/cloning of plasma cell;*  
3. Plasma cells/B cells release/produce antibodies;

2 max

- (c) 1. Lots of antibodies (against Ebola) in recovered patient;  
2. Transfusion/plasma contains antibodies;  
*Ignore reference to cells*  
3. Antibodies (specific so) will bind with (Ebola) antigen;  
4. (In recipient) virus destroyed/cannot enter cell;  
*Antigen destroyed is insufficient*

3 max

- (d) 1. (High mutation rate leads to) antigens change/antigenic variability;  
*Accept (high mutation rate leads to) changes in base sequence coding for antigen;*  
2. Vaccine contains specific antigen;  
3. Antibodies not complementary to (changed) antigen / won't bind to (changed) antigens;

3

[10]