- Q1. Dr Semmelweiss collected data about the number of deaths in the two maternity wards in the hospital where he worked.
 - From 1833 to 1838 there were the same number of doctors and midwives delivering babies in both **Ward 1** and **Ward 2**.
 - From 1839 to 1847 medical students and doctors delivered babies in **Ward 1**; midwives delivered babies in **Ward 2**.

Dr Semmelweiss also noticed that doctors often came straight from examining dead bodies to the delivery ward.

The table shows the number of patients and the number of deaths in the two wards.

| Years | Ward | Number of patients | Number of deaths | Death rate as deaths per 1000 patients |
|-----------|--------|--------------------|------------------|--|
| 4022 4020 | Ward 1 | 23 509 | 1505 | 64.0 |
| 1833–1838 | Ward 2 | 13 097 | 731 | 55.8 |
| 4000 4047 | Ward 1 | 20 204 | 1989 | 98.4 |
| 1839–1847 | Ward 2 | 17 791 | 691 | |

(a) (i) Use the formula

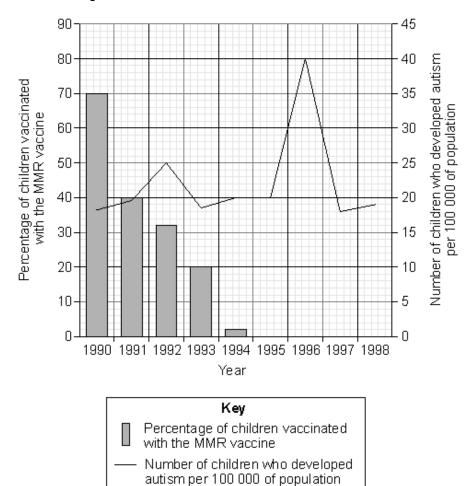
$$death\ rate = \frac{number\ of\ deaths \times 1000}{number\ of\ patients}$$

| | Hamber of patients | |
|------|---|-----|
| | to calculate the death rate for Ward 2 in the years 1839 - 1847. | |
| | | |
| | | |
| | Death rate = deaths per thousand | (2) |
| (ii) | Suggest a hypothesis for the difference in the death rates on Ward 1 and Ward 2 in the years 1839 - 1847. | |
| | | |
| | | |
| | | |
| | | (2) |

| | (D) | Antibiotics are now used in nospitals. | |
|-----|-----|---|------------|
| | | What is an antibiotic, and what does it do? | |
| | | | |
| | | | |
| | | | (2) |
| | | | (2) |
| | (c) | MRSA is causing problems in hospitals. | |
| | | Give one reason why. | |
| | | | |
| | | | (4) |
| | | | (1) |
| | (d) | How can the work of Semmelweiss help to reduce the problems caused by MRSA? | |
| | | | |
| | | | (1) |
| | | (Total 8 ma | |
| | | | |
| | | | |
| Q2. | | Many diseases are caused by viruses. Children are given vaccines to protect them against disease. | |
| | (a) | Complete the following sentences. | |
| | | It is difficult to kill viruses inside the body because | |
| | | viruses | |
| | | The vaccine stimulates the white blood cells to produce | |
| | | | (3) |

(b) In the 1990s many people thought that the MMR vaccine caused autism in some children. This is why the Japanese government stopped using the MMR vaccine.

The graph gives information about the percentage of Japanese children who developed autism during the 1990s.



| | autism. | |
|----|--|---------------|
| | Explain why. | |
| | | |
| | | |
| | | |
| | | |
| | | |
| | | |
| | | |
| | | |
| | | (4) |
| | (Т | otal 7 marks) |
| | | |
| | | |
| F | Read the passage about the use of antibiotics in food production. | |
| Pe | eople do not always agree about the use of antibiotics in food production. | |
| sh | ome farmers put low doses of antibiotics in feed for animals such as cattle and neep. Antibiotics help to keep animals disease-free. Antibiotics also help animals to row. | |
| | ne use of antibiotics in livestock feed means that there is a higher risk of antibiotic- sistant bacteria developing. These could be dangerous to human health. | |
| a) | Explain how a population of antibiotic-resistant bacteria might develop from non-resis bacteria. | stant |
| | | |
| | | |
| | | |
| | | |
| | | (3) |
| | | (3) |

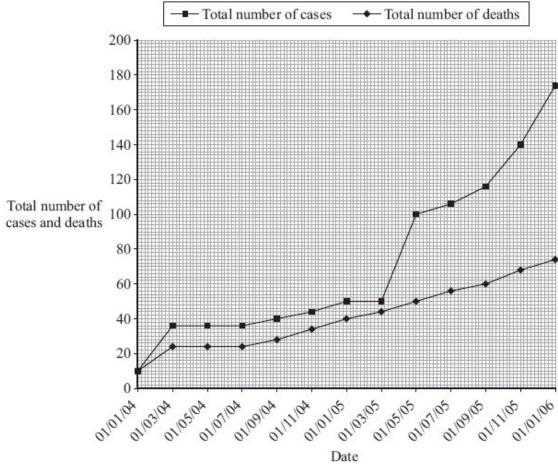
Q3.

(a)

The data in the graph support the view that there is \mathbf{no} link between MMR vaccination and

- Q4. Scientists began to keep records of cases of H5N1 bird flu in humans in January 2004.

The graph shows the total number of cases of bird flu in humans and the total number of deaths up to January 2006.



| | | Date |
|-----|-----|--|
| (a) | (i) | How many people had died from bird flu up to 01/07/05? |
| | | |

(1)

(Total 5 marks)

| | (2) |
|--|-----|
| (b) At present, humans can only catch bird flu from contact with infected birds. The bird flu virus may mutate into a form that can be passed from one human to another. | |
| Explain why millions of people may die if the bird flu virus mutates in this way. | |
| | |
| | |
| | |
| (Total 5 i | (2) |

Q5. Hepatitis B is a liver disease caused by a virus. The virus is found in body fluids such as blood, saliva and urine. Diagram 1 shows the structure of the virus in cross section.

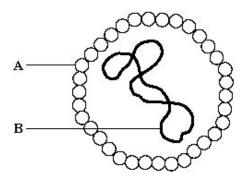


Diagram 1

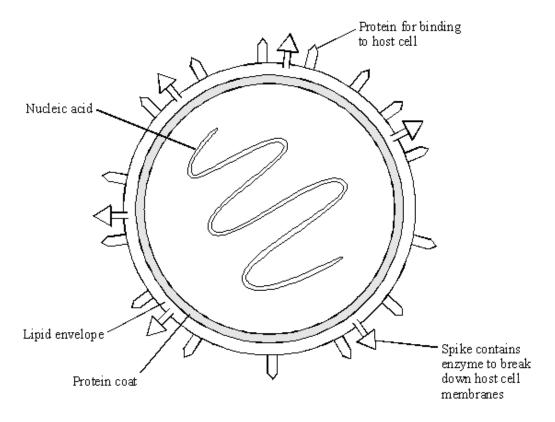
| | | viru | ses from entering the bo | ody. Others act once the viruses | s have entered. | |
|-----|-----|-------|--------------------------|--|-----------------------------|--------------|
| | | (i) | Diagram 2 shows a w | hite blood cell attacking a group | of viruses. | |
| | | | Complete diagram 2 b | y drawing the 2nd stage. | | |
| | | | Viruses | • - | + 8 | |
| | | | 1st stage | 2nd stage | 3rd Stage | |
| | | | | Diagram 2 | | (1) |
| | | | | | | , |
| | | (ii) | What type of chemica | l is released by some white bloo | od cells to attack viruses? | |
| | | | | | | (1) |
| | (b) | | | be spread among people who s n at the beginning of this questi | | (,, |
| | | | | | (Total 4 ma | (2) arks) |
| Q6. | | ghing | g and sneezing spread th | the cells lining the respiratory to the virus. In a method of spreading an infection of spreading an infection of spreading an infection. | | |
| | . , | | | | | |
| | | | | | | (1) |
| | | | | | | |

The human body has several natural defences against viruses. Some of these prevent

(a)

| (b) | sug | n immunisation programme such as that for MMR (Measles, Mumps and Rubella), gest why it is essential for a large proportion of the child population to be vaccinated in er to protect the few individuals who are unable to be vaccinated. | |
|-----|-------|--|-----|
| | | | (1) |
| (c) | virus | ome modern influenza vaccines the protein surface sub-units are separated from the scoat and used for the vaccine. This stimulates an effective immune response in the e way as inactive pathogens. | |
| | (i) | Explain how this immunity is produced in the body following vaccination, and how further illness from the same virus is prevented. | |
| | | | |
| | | | |
| | | | |
| | | | |
| | | | (4) |
| | (ii) | This type of immunity resulting from an influenza injection is described | |
| | | asimmunity. | (1) |
| | | | ` ' |

(d) The diagram shows the structure of an influenza virus.



| Explain why this is so, using the diagram of the influenza virus structure and your knowledge of immunity. | |
|--|---------------|
| | |
| | |
| | |
| | |
| | |
| | |
| | |
| | |
| | |
| | |
| | |
| | (3) |
| (Tot | tal 10 marks) |

Influenza epidemics can arise because the nucleic acid of the virus frequently changes. This results in changes in the virus structure and so a new strain of the virus is formed. A person who has had influenza or who has been vaccinated may not be immune to the new

strain.

| M1. | | (a) | (i) | 38.84 | | |
|-----|-----|------|----------|---|---|------------|
| | | ` , | , | correct answer with or without working gains 2 marks (691 × 1000) / 17 791 gains 1 mark | | |
| | | | | (091 x 1000) / 17 791 gains Tillaik | 2 | |
| | | (ii) | WO | men in Ward 1 infected | 1 | |
| | | | by | pathogens / bacteria / viruses passed on by doctors | 1 | |
| | | | | no have been in contact with dead bodies) | 1 | |
| | (b) | me | dicine | e / drug | | |
| | | (l | c 1-201- | hantada. | 1 | |
| | | tna | t KIIIS | bacteria | 1 | |
| | (c) | res | istant | to / not killed by antibiotics | | |
| | | | | | 1 | |
| | (d) | tou | ich an | weiss showed that infection could be passed on via d so hand washing by doctors / nurses / patients / | | |
| | | VİSİ | itors r | educes the risk of infection | 1 | 701 |
| | | | | | | [8] |
| | | | | | | |
| | | | | | | |
| M2. | | (a) | live i | nside cells | 1 | |
| | | ina | ctive | | 1 | |
| | | ant | ibodie | | 1 | |
| | | anı | ibouie | 5 | 1 | |
| | (b) | the | perce | entage of children vaccinated fell to zero in 1995 | | |
| | | hut | the n | umber of children developing autism rose and fell | 1 | |
| | | | | e period when % vaccinations was falling | 1 | |
| | | nur | mber (| of children developing autism peaked after MMR | - | |
| | | | | on had ceased | 1 | |
| | | whi | ich su | ggests that something other than MMR vaccination | | |
| | | | | sing autism | 1 | |
| | | | | | | [7] |

| IVI3. | | (a) Idea that bacteria mutate or that there is variation in bacteria | 1 | |
|-------|-----|---|---|-----|
| | | leading to bacteria /resistant cells that survive antibiotic | 1 | |
| | | these bacteria (resistant cells) go on to breed | 1 | |
| | | do not allow bacteria get used to antibiotics or idea that antibiotics change the bacteria or bacteria become immune or references to adaptation or evolution | | |
| | (b) | the treated animals do not use energy overcoming illness | 1 | |
| | | an economic reason, eg treated animals do not infect other animals / farm workers | | |
| | | | 1 | [5] |
| | | | | |
| | | | | |
| M4. | | (a) (i) 56 | | |
| | | accept 54 – 58 | 1 | |
| | | (ii) increased | 1 | |
| | | reasonable qualification eg slowly then more quickly or | | |
| | | to 174 / 176 or | | |
| | | by 138 / 140 | 1 | |
| | (h) | any two from: | _ | |
| | (5) | no immunity or antibodies ineffective | | |
| | | accept no resistance | | |
| | | • no vaccines or humans not immunised | | |
| | | idea of large scale contact or large scale travel do not accept passed on | | |
| | | do not accept passed on ignore no cure | 2 | |
| | | | 4 | [5] |
| | | | | |

| M5. | | (a) | (i) | diagram shows extensions of intact cell membrane around viruses | 1 | |
|-----|-----|------|-----------------|---|-------|-----|
| | | (ii) | anti | bodies | | |
| | | ` , | | allow enzymes re (ii) | | |
| | | | | allow interferon ignore antitoxins / proteins | | |
| | | | | ignore antitoxins / proteins | 1 | |
| | 41. | | | | | |
| | (b) | viru | <u>us</u> is tr | ansferred | 1 | |
| | | (vi | rue in\ | blood / body fluids – transfer (via needles) | | |
| | | (VI | ius iii) | blood / body fluids – transfer (via fleedies) | 1 | |
| | | | | | | [4] |
| | | | | | | |
| | | | | | | |
| | | | | | | |
| M6. | | (a) | dropl | et infection or aerosol infection | | |
| | | | | do not accept airborne accept airborne droplets | | |
| | | | | aboopt amborno aropioto | 1 | |
| | (h) | | thara i | is no large group which could establish infection/page on the infection | | |
| | (b) | 50 | uieiei | is no large group which could catch the infection/pass on the infection converse – if large numbers can't pass it on the virus is less likely | | |
| | | | | to reach those few who are susceptible | | |
| | | | | | 1 | |
| | (c) | (i) | any | four of the following points:- | | |
| | . , | ., | - | example of a 3 mark answer: Lymphocytes produce specific | | |
| | | | | antibodies | | |
| | | | con | nment on specificity applied to antibodies or lymphocytes | | |
| | | | (red | cognition by) lymphocytes; | | |
| | | | (wh | ite cells) make antibodies; | | |
| | | | anti | ibodies destroy/neutralise the virus/antigen/protein subunit; do not accept antibodies KILL viruses accept white blood cells replicate | | |
| | | | | accept some white cells form memory cells/live a long time; | | |
| | | | | accept subsequent infection results in very rapid antibody | | |
| | | | | production; | max 4 | |
| | | | | | шал Т | |
| | | (ii) | acti | ve; | 4 | |
| | | | | | 1 | |
| | | | | | | |

(d) any **three** of the following points

Structure change in: protein for binding to host cell;

accept changes in surface proteins (of protein coat)

spike containing enzyme;

changes in antigen

Fit: existing/circulating/old antibodies don't match new virus strain shape/new antigen/new binding protein;

Wrong antibodies: injection does not stimulate antibodies against all strains/different antigens;

accept wrong antibodies for 1 mark

max 3

[10]