

Mark Scheme

Q1.

Question number	Answer	Notes	Marks
(a)	C hypothalamus; A does not control sweating B does not control sweating D does not control sweating		1
(b)	B motor; A transmits impulse from sensory to motor C alternative name for A with same function D transmits impulse to brain		1
(c)	<ul style="list-style-type: none"> • (sweat) evaporates from (the surface of) the skin; • using heat from body/blood; • cools body/reduces body temperature; 		3
(d)	<ul style="list-style-type: none"> • runs in families; • must be faulty/mutated gene/chromosome/allele; • which transfers/passes on information to offspring; 		3
(e)	obesity / high sugar intake / high BMI;		1
(f)	<ul style="list-style-type: none"> • exercise/activity of individual varies; • greater rate of aerobic respiration; • heat given off as by-product/increased body temperature; • external temperature varies; • hotter more sweating/cooler less sweating; • mass of individual varies; • volume of water in the body/fluid intake varies; 	Allow reverse argument for mps 2 and 3	5
			Total 14

Q2.

Question number	Answer	Notes	Marks
(a) (i)	absorbs uv light; reduces/prevents risk of skin cancer;		2
(ii)	eye/ retina/ choroid/ red blood cell/ haemoglobin ;		1
(b) (i)	mother x father Hh hh; gametes H h h; fertilisation Hh hh; phenotype white forelock normal;		4
(ii)	chance of producing a boy is 0.5/½; chance of producing offspring with condition 0.5/½; chance of producing boy with condition is ½ × ½ = ¼;		3
(c)	Any 4 from different (genetic) code produced; different order of amino acids; codes for different proteins/enzymes; enzyme substrate complex not formed; causes change in pigment;		4
Total 14 marks			


Q3.

Question number	Answer	Mark
(a)	A response that makes reference to the following points: <ul style="list-style-type: none"> • form of gene (1) • characteristic (gene) that is expressed (1) 	2

Question number	Answer	Mark
(b)(i)	<p>man Dd (1)</p> <p>wife Dd/dd (1)</p> <p>son dd and son's wife Dd (1)</p> <p>child 1 (boy) dd and child 2 (girl) Dd (1)</p> <pre> graph TD man --- wife man --- son wife --- son son --- son_wife[son's wife] son_wife --- child1[child 1] son_wife --- child2[child 2] </pre> <p>2 marks for setting out as a family tree (minus 1 for each mistake)</p>	6

Question number	Answer	Additional guidance	Mark
(b)(ii)	<p>Process:</p> <p>probability of boy = 1 : 1/1 in 2/0.5/50% (1)</p> <p>probability of polydactyly = 1 : 1/1 in 2/0.5/50% (1)</p> <p>probability of boy with polydactyly is 0.5×0.5</p> <p>= 1 : 3/1 in 4/0.25/25% (1)</p>	allow 3 marks for correct final answer	3

Q4.

Question number	Answer	Notes	Marks								
(a) (i)	chromosomes (1)		1								
(ii)	 (1) (1)		2								
(b) (i)	<table border="1"><tr><td>complementary bases pair up</td><td>3</td></tr><tr><td>DNA double helix unwinds</td><td>1</td></tr><tr><td>strands separate</td><td>2</td></tr><tr><td>two DNA strands form</td><td>4</td></tr></table>	complementary bases pair up	3	DNA double helix unwinds	1	strands separate	2	two DNA strands form	4		1
complementary bases pair up	3										
DNA double helix unwinds	1										
strands separate	2										
two DNA strands form	4										
(c) (i)	thymine/T (1)	R thiamine/thyamine	1								
(ii)	guanine/G/cytosine/C (1)		1								
(c)	any four from <ul style="list-style-type: none">allows formation of gametes (1)haploid (1)allows variation to occur (1)which allows species to evolve (1)allows diploid number to be maintained (1)at fertilisation (1)		4								

Q5.

Question number	Answer	Mark
(a)	A	1

Question number	Answer	Mark
(b)(i)	45 000	1

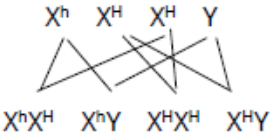
Question number	Answer	Mark
(b)(ii)	A description that makes reference to any two of the following points: <ul style="list-style-type: none"> • large (1) • increase (1) • more than doubled (1) 	2

Question number	Answer	Mark
(c)(i)	One mark for each of the following: <ul style="list-style-type: none"> • abstain from/reduce sexual partners (1) • use condom (1) 	2

Question number	Answer	Mark
(c)(ii)	Antibiotics/named antibiotic	1

Question number	Answer	Mark
(d)	An explanations that makes reference to any two of the following points: <ul style="list-style-type: none"> • lack of barrier (1) • allows fluids to mix (1) • fluids contain bacteria/viruses/fungi (1) 	2

Q6.

Question number	Answer	Notes	Marks
(a) (i)	A (the inheritance of two different alleles, both of which are expressed) B is incorrect as one allele is not expressed in codominant inheritance C is incorrect as the ABO inheritance does not involve the inheritance of multiple alleles D is incorrect as ABO inheritances does not involve the inheritance of multiple alleles		1
(ii)	$I^A I^A$ and $I^A i$		1
(b) (i)	<ul style="list-style-type: none"> blood group O contains no antigens; antigens removed/digested (by enzyme); carbohydrase digest carbohydrates; protease digest proteins; 		Max 3
(ii)	reduces need to find donor/cheaper/blood readily available/less chance of transfusing incorrect blood group/less risk of disease/infection;		1
(c) (i)	parent genotypes $X^h X^H$ and $X^H Y$; gametes and linkage $X^h \quad X^H \quad X^H \quad Y$ <div style="text-align: center;">  </div>	One mark for parent genotypes and one mark for gametes and linkage	1
(ii)	1 in 4/25%/quarter/1/4 /1:3/0.25;		1
Total question = 9 mark			

Q7.

Question number	Answer	Notes	Marks															
(a)	any three from <ul style="list-style-type: none">change in base sequence/named mutation (1)of DNA (1)causing change in genotype/gene (1)incorrect protein produced (1)change in phenotype/characteristics (1)	Allow change in DNA for mps 1 and 2 (1)	3															
(b) (i)	<table><tr><td></td><td colspan="2">father $X^H Y$</td><td colspan="2">mother $X^H X^h$ (1)</td></tr><tr><td></td><td colspan="2">gametes X^H Y</td><td colspan="2">X^H X^h (1)</td></tr><tr><td>F1</td><td>$X^H X^H$ normal female</td><td>$X^H X^h$ carrier female</td><td>$X^H Y$ normal male</td><td>$X^h Y$ (1) haemophiliac(1) male</td></tr></table>		father $X^H Y$		mother $X^H X^h$ (1)			gametes X^H Y		X^H X^h (1)		F1	$X^H X^H$ normal female	$X^H X^h$ carrier female	$X^H Y$ normal male	$X^h Y$ (1) haemophiliac(1) male		4
	father $X^H Y$		mother $X^H X^h$ (1)															
	gametes X^H Y		X^H X^h (1)															
F1	$X^H X^H$ normal female	$X^H X^h$ carrier female	$X^H Y$ normal male	$X^h Y$ (1) haemophiliac(1) male														
(ii)	(chance of first and second child is) $\frac{1}{4}/0.25/25\%$ (1) for both is $\frac{1}{4} \times \frac{1}{4}/0.25 \times 0.25$ (1) = $1/16/0.0625/6.25\%$ (1)		3															
(c)	any five from <ul style="list-style-type: none">males only have one X chromosome (1)two X chromosomes/H <u>and</u> h needed to be a carrier (1)recessive allele/h (1)carried on X chromosome (1)either present or not (1)no chance for dominant allele to be present (1)to mask recessive (1)		5															

Q8.

Question number	Answer	Notes	Marks
(a) (i)	ovulation;		1
(ii)	fertilisation		1
(iii)	Any two from: <ul style="list-style-type: none"> zygote/embryo/cells divides/splits; each half develops into one offspring; 		1 1
(b) (i)	A FSH and LH;		1
(ii)	Any four from: <ul style="list-style-type: none"> removal of egg from female ovary; collect sperm from male; egg fertilised; embryos formed; embryo inserted into uterus/female; 	Allow egg and sperm fuse Allow zygote formed	Max 4
Total question = 9 marks			

Q9.

Question number	Answer	Notes	Marks
(a) (i)	B; (an alternative form of gene) A it is not an allele C it is not an allele D it is not an allele		1
(ii)	only expressed in the homozygous state/not expressed in the heterozygous state;	Allow valid descriptions/examples of homozygous/heterozygous states	1
(b) (i)	<p>grandfather (condition) × grandmother; (no condition)</p> <p>↓</p> <p> father (cond) × mother (no cond) ↓ student (condition) { brother (condition) sister (no condition); } </p>		3
(ii)	<ul style="list-style-type: none"> occurs only in males/not females/more common in males/females are carriers; recessive; probably sex-linked/carried on X chromosome; 		3
			Total 8

Q10.

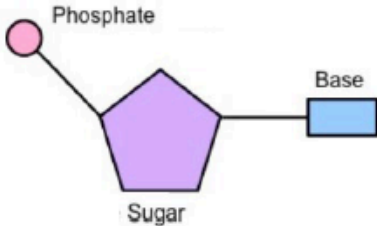
Question number	Answer	Notes	Marks
(a) (i)	<ul style="list-style-type: none"> correct labelling of axes (1) suitable scale (1) X and Y axis correct way round (1) plotting (2) 	minus 1 for each incorrect plot	5
(ii)	Any two from <ul style="list-style-type: none"> few at lower end/150-151/top end/160 (1) majority in middle/154-157 (1) increases to 154-155 then decreases (1) 		2
(b)	Any three from <ul style="list-style-type: none"> nutrition (1) sex (1) age (1) genes/genetically based (1) 		3

Q11.

Question number	Answer	Notes	Marks
(a) (i)	any three from <ul style="list-style-type: none"> prevents excessive loss of blood (1) forms scab over wound (1) prevents entry of bacteria /pathogens/microorganisms(1) so prevents infection (1) allows wound to heal (1) 		3
(ii)	<ul style="list-style-type: none"> (allele) on X/sex chromosome (1) passed on to offspring/inherited by children (1) 		2

(b) (i)	D; (X^hY) A the genotype is not X^hX^h B the genotype is not X^HX^h C the genotype is not X^HY		1
(ii)	B; (X^HX^h) A the genotype is not X^hX^h C the genotype is not X^HY D the genotype is not X^hY		1
(iii)	<ul style="list-style-type: none"> offspring genotypes X^HX^h and X^hY (1) 1 carrier female to 1 haemophiliac male = 50%/0.5/½ (1) 	full marks for correct figures only.	2
(iv)	<ul style="list-style-type: none"> unusual for female to be haemophiliac (1) both alleles have to be affected (1) must have a haemophiliac father and mother with haemophilia/carrier (1) 	any correct reference to excessive blood loss	3
Total for question = 12 marks			

Q12.

Question number	Answer	Notes	Marks
(a) (i)	Any two from: <ul style="list-style-type: none"> (movement of substances/named solute or gas) against a concentration gradient(1) using energy/ATP(1) 		2
(ii)	Any two from: <ul style="list-style-type: none"> parent homozygous recessive/heterozygous / carriers (1) one (recessive) allele inherited from each parent (1) reference to homozygous recessive disorder/caused by recessive alleles (1) 		2
(iii)	<ul style="list-style-type: none"> protein/CFTR is not modified/folded/protein has an incorrect shape(1) protein/CFTR not transported (to its destination)(1) 		2
(iv)	1408 x 3 (1) 4224	Full marks for correct final answer. Max 2 marks ecf for final answer if it is correct from calculation	2
(v)	<ul style="list-style-type: none"> (deoxyribose) sugar(1) joined to a (nitrogenous) base/named base(1) phosphate group attached to sugar (1) 	Allow one mark only if all 3 components just listed	3
(b)	any three from <ul style="list-style-type: none"> deletion(1) of one codon/TTT (1) phenylalanine missing from protein (1) shape of final protein changed/different protein formed (1) 	allow description of deletion	3
Total for Question = 14 Marks			