

17. Inheritance

17.4 Monohybrid inheritance

Paper 3 and 4

Marking Scheme

Q1.

(a)(i)	a length of DNA ; that codes for a protein ;	2	
(a)(ii)	nucleus ;	1	A mitochondria
(b)	15 (people) ;	1	
(c)(i)	XX ;	1	
(c)(ii)	4 ;	1	
(c)(iii)	homozygous recessive ;	1	R each additional circle
(d)	parental gametes: A a x a a ; offspring genotypes: Aa (Aa) aa (aa) ; offspring phenotypes: unaffected (unaffected) albinism (albinism) ; ratio 1:1 ;	4	A ecf from previous step A 2:2 / 50:50 / 50%

Q2.

(a)	term	definition	4 one mark for each correct line R each additional line
	dominant	an allele that is expressed if it is present	
	genotype	genetic make-up of an organism	
	heterozygous	having two different alleles of a particular gene	
		having two identical alleles of a particular gene	
	phenotype	observable features of an organism	

Q3.

(b)(i)	1 / one ;	1																
(b)(ii)	D ;	1																
(b)(iii)	3 / three ;	1																
(c)	four correct gamete alleles ; four correct offspring genotypes ; (probability ...) 0 / zero ;	3	<table border="1"> <tr> <td></td><td></td><td colspan="2">male</td></tr> <tr> <td></td><td></td><td>R</td><td>R</td></tr> <tr> <td rowspan="2">female</td><td>R</td><td>RR</td><td>RR</td></tr> <tr> <td>r</td><td>Rr</td><td>Rr</td></tr> </table>			male				R	R	female	R	RR	RR	r	Rr	Rr
		male																
		R	R															
female	R	RR	RR															
	r	Rr	Rr															
(d)	<u>pure</u> ;	1																

Q4.

(a)(i)	gg ; yellow ; yellow ;	3	
(a)(ii)	top row both offspring Gg, bottom row both offspring gg ; 1:1 ;	2	
(b)(i)	2 ;	1	
(b)(ii)	R / S ;	1	
(b)(iii)	0% circled ;	1	

Q5.

(b)(i)	<i>parental genotypes:</i> AA and Aa ; <i>gametes:</i> A A (×) A a ; <i>offspring genotypes:</i> AA, AA, Aa, Aa ; <i>offspring phenotypes:</i> all normal wings ; <i>probability:</i> zero ;	5	ecf for each subsequent row
(b)(ii)	AA ;	1	A homozygous dominant
(b)(iii)	<i>Drosophila ;</i>	1	

Q6.

		4											
	<table><tr><th>number</th><th>genotype</th></tr><tr><td>1</td><td>bb ;</td></tr><tr><td>2</td><td>Bb ;</td></tr><tr><td>4</td><td>Bb ;</td></tr><tr><td>14</td><td>bb ;</td></tr></table>	number	genotype	1	bb ;	2	Bb ;	4	Bb ;	14	bb ;		
number	genotype												
1	bb ;												
2	Bb ;												
4	Bb ;												
14	bb ;												

Q7.

(a)(i)	five / 5 ;	1														
(a)(ii)	<u>black</u> (fur / coat) ;	1														
(a)(iii)	homozygous circled ; dominant circled ;	2														
(b)	discontinuous ;	1														
(c)(i)	(male and female gametes) A and a ; (offspring) AA , Aa , Aa and aa ; (phenotypic ratio) 3 (white) : 1 (black) ;	3	ecf from the step before <table border="1"> <tr> <td colspan="2" rowspan="2"></td><th colspan="2">male</th></tr> <tr> <th>A</th><th>a</th></tr> <tr> <th rowspan="2">female</th><th>A</th><td>AA</td><td>Aa</td></tr> <tr> <th>a</th><td>Aa</td><td>aa</td></tr> </table>			male		A	a	female	A	AA	Aa	a	Aa	aa
		male														
		A	a													
female	A	AA	Aa													
	a	Aa	aa													

Q8.

(b)(i)	individual	genotype	3
	1	Tt ;	
	3	Tt ;	
	4	Tt ;	
(b)(ii)	<i>genotypes:</i> Tt and tt ; <i>genotype ratio:</i> 1 : 1 ;		2

Q9.

(a)	<i>any two from:</i> parent 2, is homozygous recessive / has two alleles (for red-green colour blindness) ; only parent 2 can provide the X chromosome for male children ; males only have one X chromosome and they only need one recessive allele (to be red-green colour blind) ; red-green colour blindness is a sex-linked characteristic ;	2
-----	--	---

(b)	<i>parental genotypes:</i> parent 3 X^bY and parent 4 X^BX^b ; <i>gametes:</i> X^b , Y/0/- and X^B , X^b ; <table><tr><td><i>offspring genotypes</i></td><td>X^BX^b ,</td><td>X^BY</td><td>X^bX^b</td><td>X^bY</td><td>;</td></tr><tr><td><i>offspring phenotypes</i></td><td>normal (colour vision) female</td><td>normal (colour vision) male</td><td>colour-blind female</td><td>colour-blind male</td><td>;</td></tr></table> <i>probability:</i> 0.5 / 50% ;	<i>offspring genotypes</i>	X^BX^b ,	X^BY	X^bX^b	X^bY	;	<i>offspring phenotypes</i>	normal (colour vision) female	normal (colour vision) male	colour-blind female	colour-blind male	;	5
<i>offspring genotypes</i>	X^BX^b ,	X^BY	X^bX^b	X^bY	;									
<i>offspring phenotypes</i>	normal (colour vision) female	normal (colour vision) male	colour-blind female	colour-blind male	;									

Q10.

(a)	4 ; 3 ;	2	A 1, 4, 5 and 7 A 2, 4 and 5
(d)	<i>ref. to I^A and I^B alleles ;</i> <i>I^A and I^B alleles both contribute to phenotype</i> <i>/ ref. to AB blood group ;</i>	2	
(e)	<i>gametes</i> I^A, I^O and I^B, I^O ; <i>genotypes</i> I^AI^O, I^AI^B, I^BI^O, I^OI^O ; <i>phenotypes</i> A, AB, B, O ; <i>probability</i> 0.25 / 25% / 1 in 4 ;	4	ecf from previous stage

Q11.

(e)(i)	parents of 5 and 7 / parents of people with cystic fibrosis / 2 and 3, do not have cystic fibrosis ; parents / 2 and 3 must be, heterozygous / carriers (of the mutant allele) ;	2	
--------	---	---	--

(e)(ii)	<p>1 correct parental genotype aa ;</p> <p>2 correct parental genotype Aa ;</p> <p>3 correct gametes from the parental genotypes ;</p> <p>4 correct offspring genotypes from their gametes / parental genotypes (in any order) ;</p> <p>5 correct offspring phenotypes and correct probability ;</p>	<p>5</p> <p><i>MP1 and MP2 parents may be either way round but following rows must match</i></p> <p>ecf from MP1 <u>and</u> MP2 ecf from MP3 ecf from MP4</p> <p>expected answer:</p> <table> <tr> <td></td> <td>parent 7</td> <td>man who is heterozygous</td> </tr> <tr> <td>parental genotypes</td> <td>aa</td> <td>× Aa</td> </tr> <tr> <td>gametes</td> <td>a a</td> <td>+ A a</td> </tr> <tr> <td>offspring genotypes</td> <td>Aa (Aa)</td> <td>aa (aa)</td> </tr> <tr> <td>offspring phenotypes</td> <td>without cystic fibrosis</td> <td>with cystic fibrosis</td> </tr> <tr> <td>probability</td> <td>50% / 1 in 2 / 0.5</td> <td></td> </tr> </table>		parent 7	man who is heterozygous	parental genotypes	aa	× Aa	gametes	a a	+ A a	offspring genotypes	Aa (Aa)	aa (aa)	offspring phenotypes	without cystic fibrosis	with cystic fibrosis	probability	50% / 1 in 2 / 0.5	
	parent 7	man who is heterozygous																		
parental genotypes	aa	× Aa																		
gametes	a a	+ A a																		
offspring genotypes	Aa (Aa)	aa (aa)																		
offspring phenotypes	without cystic fibrosis	with cystic fibrosis																		
probability	50% / 1 in 2 / 0.5																			

Q12.

(c)(i)	only males, are colour-blind / colour-blindness is more common in males / no females are colour-blind / person 1 and 2 had two male colour-blind children (but no female colour-blind children) ;	1	
(c)(ii)	$X^A X^a$; ;	2	MP1 correct sex chromosomes MP2 correct alleles
(c)(iii)	1 correct genotype for person 3, i.e. $(X)^A(X)^a$; 2 correct genotype for person 4, i.e. $X^A Y$; 3 correct gametes from the parental genotypes ; 4 correct offspring genotypes from their gametes / parental genotypes (in any order) ; 5 correct offspring correct probability ;	5	MP3 ecf from MP1 <u>and</u> MP2 MP4 ecf from MP3 MP5 ecf from their offspring genotype expected answer: <i>genotype</i> (person 3) $X^A X^a$ × (person 4) $X^A Y$ <i>gametes</i> X^A, X^a + X^A, Y <i>offspring genotypes</i> $X^A X^A, X^A X^a, X^A Y, X^a Y$ <i>offspring phenotypes</i> female with normal colour vision female with normal colour vision male with normal colour vision male with colour blindness) <i>probability</i> 0.25 / 25% / 1 in 4 / ¼

Q13.

(a)	selection of suitable letter and case for alleles, e.g. R and r ; <i>parental phenotypes:</i> red fruit × red fruit parental genotypes: Rr ; × Rr ; gametes: R r × R r ; offspring genotypes: (1)RR and (2)Rr and (1)rr ; expected ratio: 3 (red fruit) : 1 (yellow fruit) ;	6	
(b)	(perform a) test cross ;	1	

Q14.

(b)(i)	$p^R p^R$; $p^R p^W$; $p^R p^R, p^R p^W, p^W p^W$;	3	
(b)(ii)	codominance ;	1	

Q15.

(a)	transmission of genetic information from generation to generation ;	1
(b)(i)	1 correct use of X and Y in responses for individual 5 and individual 8 ; 2 correct X allele given for individual 5: X^bY / b ; 3 correct X allele given for individual 8: X^BY / B ;	3
(b)(ii)	<i>any three from:</i> colour blindness is a sex-linked characteristic ; she is, heterozygous for the gene / Bb ; she has, normal allele / B , so has normal colour vision ; but has passed on the, recessive allele / b , to her sons / 5 and 7 ; she has two X chromosomes which have the gene for colour vision ; father / 4, passes on his Y chromosome ;	3

Q16.

(b)(i)	transmission of genetic information from generation to generation ;	1
(b)(ii)	two of the same letter both lower case ;	1
(b)(iii)	$0.25 / 25\% / \frac{1}{4}$;	1
(b)(iv)	<i>any two from:</i> perform a test cross ; by breeding with, homozygous recessive / king cheetah ; if any of the offspring of the test cross are king cheetahs it confirms 17 is heterozygous ; DNA testing ;	2

Q17.

(d)(i)	X^BX^b ;	1	
(d)(ii)	X^bY ;	1	
(d)(iii)	solid shaded square on Fig. 4.2 ;	1	
(d)(iv)	one X chromosome from each parent / an X from father ; mother does not have any colour-blind alleles / father passes on one colour-blind allele ; (all female offspring are) heterozygous / X^BX^b ;	2	

Q18.

(c)	the allele for dwarfism is, recessive / t ; both parents are heterozygous (so do not express the allele) ;	2	
-----	---	---	--

Q19.

(c)(i)	(P) $X^H X^h$; (Q) $X^h Y$; (R) $X^H Y$;	3	
(c)(ii)	0.25 / 25% / 1 in 4 / $\frac{1}{4}$;	1	
(c)(iii)	gene is located on, a sex chromosome / X <u>or</u> Y / X / Y ; characteristic is more common in, males / one sex (than the other) ;	2	