

# 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	<b>A</b> mitochondria
(b)	15 (people) ;	1	
(c)(i)	XX ;	1	
(c)(ii)	4 ;	1	
(c)(iii)	homozygous recessive ;	1	<b>R</b> each additional circle
(d)	<i>parental gametes: A a x a a ; offspring genotypes: Aa (Aa) aa (aa) ; offspring phenotypes: unaffected (unaffected) albinism (albinism) ; ratio 1:1 ;</i>	4	<b>A</b> ecf from previous step  <b>A</b> 2:2 / 50:50 / 50%

## Q2.

(a)	term	definition	4 one mark for each correct line <b>R</b> 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	
	phenotype	having two identical alleles of a particular gene	
		observable features of an organism	

## Q3.

(b)(i)	1 / one ;	1												
(b)(ii)	<b>D</b> ;	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 colspan="2">male</td> </tr> <tr> <td></td> <td>R</td> <td>R</td> </tr> <tr> <td rowspan="2">female</td> <td>R</td> <td>RR</td> </tr> <tr> <td>r</td> <td>Rr</td> </tr> </table>		male			R	R	female	R	RR	r	Rr
	male													
	R	R												
female	R	RR												
	r	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)	<b>R / S</b> ;	1	
(b)(iii)	0% circled ;	1	

## Q5.

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

## Q6.

	number	genotype	4	
	1	<b>bb</b> ;		
	2	<b>Bb</b> ;		
	4	<b>Bb</b> ;		
	14	<b>bb</b> ;		

## 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) <b>AA</b> , <b>Aa</b> , <b>Aa</b> and <b>aa</b> ; (phenotypic ratio) 3 (white) : 1 (black) ;	3	<p>ecf from the step before</p> <table border="1" style="margin-left: auto; margin-right: auto;"> <tr> <td colspan="2"></td> <td colspan="2">male</td> </tr> <tr> <td colspan="2"></td> <td>A</td> <td>a</td> </tr> <tr> <td rowspan="2" style="background-color: #e0e0e0;">female</td> <td>A</td> <td>AA</td> <td>Aa</td> </tr> <tr> <td>a</td> <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)	<table border="1" style="margin-left: auto; margin-right: auto;"> <tr> <th>individual</th><th>genotype</th></tr> <tr> <td>1</td><td>Tt ;</td></tr> <tr> <td>3</td><td>Tt ;</td></tr> <tr> <td>4</td><td>Tt ;</td></tr> </table>	individual	genotype	1	Tt ;	3	Tt ;	4	Tt ;	3	
individual	genotype										
1	Tt ;										
3	Tt ;										
4	Tt ;										
(b)(ii)	<p><i>genotypes</i>: Tt and tt ;</p> <p><i>genotype ratio</i>: 1 : 1 ;</p>	2									

## Q9.

(a)	<p><i>any two from</i>:</p> <p>parent 2, is homozygous recessive / has two alleles (for red-green colour blindness) ;</p> <p>only parent 2 can provide the X chromosome for male children ;</p> <p>males only have one X chromosome and they only need one recessive allele (to be red-green colour blind) ;</p> <p>red-green colour blindness is a sex-linked characteristic ;</p>	2													
(b)	<p><i>parental genotypes</i>: parent 3 <b>X<sup>b</sup>Y</b> and parent 4 <b>X<sup>B</sup>X<sup>b</sup></b> ;</p> <p><i>gametes</i>: <b>X<sup>b</sup></b> , <b>Y/0/-</b> and <b>X<sup>B</sup></b> , <b>X<sup>b</sup></b> ;</p> <table border="1" style="margin-left: auto; margin-right: auto;"> <tr> <td><i>offspring genotypes</i></td> <td><b>X<sup>B</sup>X<sup>b</sup></b>,</td> <td><b>X<sup>B</sup>Y</b></td> <td><b>X<sup>b</sup>X<sup>b</sup></b></td> <td><b>X<sup>b</sup>Y</b></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> <p><i>probability</i>: 0.5 / 50% ;</p>	<i>offspring genotypes</i>	<b>X<sup>B</sup>X<sup>b</sup></b> ,	<b>X<sup>B</sup>Y</b>	<b>X<sup>b</sup>X<sup>b</sup></b>	<b>X<sup>b</sup>Y</b>	;	<i>offspring phenotypes</i>	normal (colour vision) female	normal (colour vision) male	colour-blind female	colour-blind male	;	5	
<i>offspring genotypes</i>	<b>X<sup>B</sup>X<sup>b</sup></b> ,	<b>X<sup>B</sup>Y</b>	<b>X<sup>b</sup>X<sup>b</sup></b>	<b>X<sup>b</sup>Y</b>	;										
<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)	ref. to $I^A$ and $I^B$ alleles ; $I^A$ and $I^B$ alleles both contribute to phenotype / ref. to AB blood group ;	2	
(e)	gametes $I^A, I^0$ and $I^B, I^0$ ; genotypes $I^A I^0, I^A I^B, I^B I^0, I^0 I^0$ ; phenotypes A, AB, B, O ; probability 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)	1 correct parental genotype aa ; 2 correct parental genotype Aa ; 3 correct gametes from the parental genotypes ; 4 correct offspring genotypes from their gametes / parental genotypes (in any order) ; 5 correct offspring phenotypes and correct probability ;	5	<p>MP1 and MP2 parents may be either way round but following rows must match</p> <p>ecf from MP1 and MP2 ecf from MP3 ecf from MP4</p> <p><b>expected answer:</b></p> <table style="margin-left: 100px;"> <tr> <td>parent 7</td> <td>man who is heterozygous</td> </tr> <tr> <td>parental genotypes</td> <td>aa</td> <td>×</td> <td>Aa</td> </tr> <tr> <td>gametes</td> <td>a</td> <td>a</td> <td>+</td> <td>A</td> <td>a</td> </tr> <tr> <td>offspring genotypes</td> <td colspan="2">Aa (Aa)</td> <td colspan="2">aa (aa)</td> </tr> <tr> <td>offspring phenotypes</td> <td>without cystic fibrosis</td> <td></td> <td>with cystic fibrosis</td> <td></td> </tr> <tr> <td>probability</td> <td>50%</td> <td>/ 1 in 2</td> <td>/ 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)	$X^AX^a$ ;;	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. $X^AY$ ; 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	<p>MP3 <b>ecf</b> from MP1 <u>and</u> MP2 MP4 <b>ecf</b> from MP3</p> <p>MP5 <b>ecf</b> from their offspring genotype</p> <p><b>expected answer:</b> genotype (person 3) <math>X^AX^a</math> <math>\times</math> (person 4) <math>X^AY</math> gametes <math>X^A, X^a</math> + <math>X^A, Y</math> offspring genotypes <math>X^AX^A</math>, <math>X^AX^a</math>, <math>X^AY</math>, <math>X^aY</math> offspring phenotypes female with normal colour vision female with normal colour vision male with normal colour vision male with colour blindness)</p> <p>probability 0.25 / 25% / 1 in 4 / <math>\frac{1}{4}</math></p>

## Q13.

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

## Q14.

(b)(i)	$PPPR$ ; $PPP^W$ ; $PPP^R$ , $P^R P^W$ , $P^W P^W$ ;	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
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**Q19.**

(c)(i)	(P) $X^hX^h$ ; (Q) $X^hY$ ; (R) $X^hY$ ;	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	