

Inheritance

Mark Scheme 2

Level	IGCSE
Subject	Biology
Exam Board	CIE
Topic	Inheritance
Paper Type	(Extended) Theory Paper
Booklet	Mark Scheme 2

Time Allowed: 58 minutes

Score: /48

Percentage: /100

<p>1 (a)</p>	<p>the allele that is expressed (if it is present)/AW; always seen in the phenotype; masks (effect of) recessive allele;</p>	<p>max 1</p>	<p>I 'powerful' defines the phenotype defines characteristic(s)</p>
<p>(b) (i)</p>	<p><i>Parent genotype:</i> Ff , Ff; <i>Parent phenotype:</i> (with) flecks × (with) flecks; <i>Gametes:</i> F , f, F , f; Working shown to derive genotype; <i>Offspring genotype:</i> FF , Ff , ff; linked to correct phenotype</p>	<p>5</p>	<p>ECF on incorrect key usage ECF from each line A Punnett square / criss-cross lines</p>
<p>(ii)</p>	<p>ff × ff; parents may be implied as first part of the question asks for parental genotype both parents must have a recessive <u>allele</u> / (if ff × ff) no dominant or F <u>allele</u>, in either parent / (if ff × ff) both parents must be homozygous, recessive / without flecks no parent must be homozygous dominant / presence of (even) one dominant allele in parents could result in flecks;</p>	<p>2</p>	<p>A Ff × Ff and Ff × ff ECF on incorrect key usage from (i) A gene for allele</p>
		<p>[Total: 8]</p>	

<p>2 (a) (i)</p>	<table border="1"> <thead> <tr> <th data-bbox="297 212 548 296">genetic term</th> <th data-bbox="548 212 1189 296">example used in the passage</th> </tr> </thead> <tbody> <tr> <td data-bbox="297 296 548 363">an allele</td> <td data-bbox="548 296 1189 363">Hb^N / Hb^S;</td> </tr> <tr> <td data-bbox="297 363 548 461">a heterozygous genotype</td> <td data-bbox="548 363 1189 461">$Hb^N Hb^S$;</td> </tr> <tr> <td data-bbox="297 461 548 558">a homozygous genotype</td> <td data-bbox="548 461 1189 558">$Hb^S Hb^S$;</td> </tr> <tr> <td data-bbox="297 558 548 703">phenotype</td> <td data-bbox="548 558 1189 703">/ extreme pain / sickle cell anaemia / mild symptoms;</td> </tr> </tbody> </table>	genetic term	example used in the passage	an allele	Hb^N / Hb^S ;	a heterozygous genotype	$Hb^N Hb^S$;	a homozygous genotype	$Hb^S Hb^S$;	phenotype	/ extreme pain / sickle cell anaemia / mild symptoms;	<p>4</p>	<p>A N/S, R NS and N × S</p> <p>A NS</p> <p>A SS</p> <p>A the disease</p>
genetic term	example used in the passage												
an allele	Hb^N / Hb^S ;												
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phenotype	/ extreme pain / sickle cell anaemia / mild symptoms;												
<p>(ii)</p>	<p>malaria, is severe disease / may be fatal;</p> <p>idea that it is the selective agent / ref to (natural) selection;</p> <p>people with sickle cell anaemia / Hb^S are resistant to malaria;</p> <p>$Hb^N Hb^N$ / homozygous dominant, susceptible to malaria;</p> <p>$Hb^N Hb^N$ more likely to die (of malaria) before have children (to pass on genes);</p> <p>$Hb^N Hb^S$ / sickle cell carriers, do not die from sickle cell anaemia;</p> <p>$Hb^N Hb^S$ / sickle cell carriers, have children (and pass on genes);</p> <p>and pass on the (Hb^S) <u>allele</u>;</p> <p>description of sickle cells are less prone to infection;</p> <p>idea that no advantage of Hb^S in areas where no malaria;</p> <p>AVP;</p>	<p>max 5</p>	<p>A reference to selective advantage for MP2</p> <p>R immune for resistance (but ECF after first time)</p> <p>A carrier for sickle cell trait</p> <p>AVPs: 2 in 4 / ½ , have advantage of resistance to malaria; (if $Hb^N Hb^S \times Hb^N Hb^S$) 1 in 4 chance of, $Hb^S Hb^S$ / homozygous recessive;</p>										

2 (b)	(chromosome) mutation; an extra chromosome; non-disjunction / failure during meiosis / translocation;	max 1	A trisomy 21 R more than one chromosome I older mothers, inherited
(c)	discontinuous variation – influenced by genes alone; ORA discontinuous variation – no effect of the environment / does not change over (life)time; ORA discontinuous variation, is discrete / has no intermediates / is qualitative / AW; ORA limited number of <u>phenotypes</u> ;	max 3	assume answer is about discontinuous unless stated otherwise continuous variation influenced by gene and environment = 2 marks (MP1 and MP2) A continuous is measurable
		[Total: 13]	

	Answers	Marks	Guidance for Examiners
3 (a)	<p>1 (red blood cells) get stuck in capillaries / do not flow smoothly / capillaries blocked;</p> <p>2 reduce , supply of, oxygen / nutrients (to tissues / cells / muscles) ;</p> <p>3 reduce , removal of, carbon dioxide / wastes, (from tissues / cells / muscles) ;</p> <p>4 ref to respiration (in tissues) ;</p> <p>5 cause sickle cell crises ;</p> <p>6 pain ;</p> <p>7 increased chance of, thrombosis / blood clotting ;</p> <p>8 death of tissues / cells ;</p> <p>9 AVP ;</p>	[max 4]	<p>ignore less haemoglobin</p> <p>A carries <u>less</u> oxygen / nutrients...</p> <p>A carries <u>less</u> carbon dioxide...</p> <p>I reduced life expectancy</p>
(b) (i)	allele(s) ;	[1]	
(ii)	$H^A, H^S + H^A, H^S$; $(H^A H^A, H^A H^S, H^A H^S) \underline{H^S H^S}$;	[2]	<p>Could be in Punnett square</p> <p>A just A and S</p> <p>A just S and S</p>
(iii)	0.25 / 25% / 1/4 / 1 in 4 ;	[1]	I ratios

	Answers	Marks	Guidance for Examiners
3 (c) (i)	<p>1 malaria, is severe disease / may be fatal ;</p> <p>2 <i>idea that it is the selective agent / ref to natural selection ;</i></p> <p>3 $H^A H^A$ / homozygous dominant, susceptible to malaria ;</p> <p>4 $H^A H^S$ / heterozygous, resistant ; A $H^S H^S$ resistant ;</p> <p>5 $H^A H^S$ survive / $H^A H^A$ more likely to die before have children ;</p> <p>6 $H^A H^S$ have children and pass on, the allele / H^S ;</p> <p>7 (if $H^A H^S \times H^A H^S$) 1 in 4 chance of, $H^S H^S$ / homozygous recessive ;</p> <p>8 2 in 4 / $\frac{1}{2}$, have advantage of resistance to malaria ;</p> <p>9 AVP ; e.g. ref to malarial parasite /</p> <p>10 AVP ; e.g. ref to transmission of malaria</p>	[max 4]	<p>A sickle cell trait / carrier for $H^S H^A$ throughout the answer</p> <p>R immune</p>
(ii)	<p>1 malaria not very serious / not a severe strain of malaria ;</p> <p>2 people have other genetic protection from malaria ;</p> <p>3 malaria has only recently spread to these areas / no malaria before;</p> <p>4 mutation not occurred in populations of these areas ;</p> <p>5 people with mutation / have sickle cell allele , have not migrated here ;</p> <p>6 (majority of) population in Australia has not lived there for long ;</p> <p>7 came from areas where no malaria, is / was, present ;</p> <p>8 AVP ;</p> <p>9 AVP ;</p>	[max 2]	<p>E.g. Thalassemia</p> <p>A mutation described I gene, for allele</p>
		[Total:14]	

Question	E answers	Mark	Additional Guidance
4 (a)	halves the number of chromosomes / diploid to haploid ; ignore halves the genetic material produces variation / AW ;	[2]	accept produces haploid, nuclei / cells / gametes ignore prevents doubling of chromosome number
(b) (i)	question is discounted	[2]	
(ii) 1 2 3 4	(only) one fertilisation / one zygote / one fertilised egg ; zygote / fertilised egg / (cells in) embryo, divides / splits in two ; by <u>mitosis</u> ; into two (groups of) genetically identical cells ;	[2]	R 'from a single cell' but allow ecf for other MPs R egg divides A same , genetic material / genetic make-up / genome R similar
(c)	increase in, complexity / AW ; ref to specialisation / differentiation ; ref to different types of cells ; ref to, tissues / organs ;	[max 2]	ignore (rapid) growth / change in shape A 'legs / arms / AW, start to grow'
(d)	1. hY ; 2. HX^h ; 3. HX^H ;	[3]	do not accept male genotypes for MP2 and MP3

Question	E answers	Mark	Additional Guidance
<p>4 (e) 1</p> <p>2</p> <p>3</p> <p>4</p> <p>5</p> <p>6</p>	<p>mutation / change in DNA ;</p> <p>in the gene, for blood clotting protein / on X chromosome ;</p> <p>in the mother / mother is a carrier / mother is heterozygous ; R parent(s) is / are heterozygous</p> <p>haemophilia is <u>sex linked</u> / shows <u>sex linkage</u> ;</p> <p><i>idea that</i> the mother's egg with the mutant allele fuses with a Y bearing sperm ;</p> <p>e.g. cause of mutation ; ionising radiation / chemical(s)</p>	<p>[max 2]</p>	<p>MP2 can only be awarded if MP1 is awarded</p> <p>MP3 A in context of allele passing down the female line for several / many generations (without being expressed in a male)</p> <p>ignore carried on the X chromosome as this is in the question</p>