

# Inheritance

## Mark Scheme 2

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

CHEMISTRY ONLINE  
— TUITION —

**Time Allowed:** 58 minutes

**Score:** /48

**Percentage:** /100

1 (a)	the allele that is expressed (if it is present)/AW; always seen in the phenotype; masks (effect of) recessive allele;	max 1	I 'powerful' defines the phenotype defines characteristic(s)
(b) (i)	Parent genotype: Ff , Ff; Parent phenotype: (with) flecks × (with) flecks; Gametes: F , f, F , f; Working shown to derive genotype; Offspring genotype: FF , Ff , ff; linked to correct phenotype	5	ECF on incorrect key usage ECF from each line A Punnett square / criss-cross lines
(ii)	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;	2	A Ff × Ff and Ff × ff ECF on incorrect key usage from (i)  A gene for allele
		[Total: 8]	

2 (a) (i)	<table><tr><td>genetic term</td><td>example used in the passage</td></tr><tr><td>an allele</td><td>Hb<sup>N</sup> / Hb<sup>S</sup>;</td></tr><tr><td>a heterozygous genotype</td><td>Hb<sup>N</sup>Hb<sup>S</sup>;</td></tr><tr><td>a homozygous genotype</td><td>Hb<sup>S</sup> Hb<sup>S</sup>;</td></tr><tr><td>phenotype</td><td>/extreme pain/sickle cell anaemia / mild symptoms;</td></tr></table>	genetic term	example used in the passage	an allele	Hb <sup>N</sup> / Hb <sup>S</sup> ;	a heterozygous genotype	Hb <sup>N</sup> Hb <sup>S</sup> ;	a homozygous genotype	Hb <sup>S</sup> Hb <sup>S</sup> ;	phenotype	/extreme pain/sickle cell anaemia / mild symptoms;	4	<p><b>A</b> N/S, <b>R</b> <b>NS</b> and <b>N</b> × <b>S</b></p> <p><b>A</b> NS</p> <p><b>A</b> SS</p> <p><b>A</b> the disease</p>
	genetic term	example used in the passage											
an allele	Hb <sup>N</sup> / Hb <sup>S</sup> ;												
a heterozygous genotype	Hb <sup>N</sup> Hb <sup>S</sup> ;												
a homozygous genotype	Hb <sup>S</sup> Hb <sup>S</sup> ;												
phenotype	/extreme pain/sickle cell anaemia / mild symptoms;												
(ii)	<p>malara, 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<sup>S</sup> are resistant to malara;</p> <p>Hb<sup>N</sup>Hb<sup>N</sup> / homozygous dominant, susceptible to malara;</p> <p>Hb<sup>N</sup>Hb<sup>N</sup> more likely to die (of malara) before have children (to pass on genes);</p> <p>Hb<sup>N</sup> Hb<sup>S</sup> / sickle cell carriers, do not die from sickle cell anaemia;</p> <p>Hb<sup>N</sup> Hb<sup>S</sup> / sickle cell carriers, have children (and pass on genes);</p> <p>and pass on the (Hb<sup>S</sup>) <u>allele</u>;</p> <p>description of sickle cells are less prone to infection;</p> <p>idea that no advantage of Hb<sup>S</sup> in areas where no malara;</p> <p>AVP;</p>	max 5	<p><b>A</b> reference to selective advantage for MP2</p> <p><b>R</b> immune for resistance (but <b>ECF</b> after first time)</p> <p><b>A</b> carrier for sickle cell trait</p> <p>AVPs: 2 in 4 / ½ , have advantage of resistance to malara; (if Hb<sup>N</sup> Hb<sup>S</sup> × Hb<sup>N</sup> Hb<sup>S</sup> ) 1 in 4 chance of, Hb<sup>S</sup> Hb<sup>S</sup> / 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; <b>ORA</b>  discontinuous variation – no effect of the environment / does not change over (life)time; <b>ORA</b>  discontinuous variation, is discrete / has no intermediates / is qualitative / AW; <b>ORA</b>  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 ( <b>MP1</b> and <b>MP2</b> )  A continuous is measurable
			[Total: 13]	

CHEMISTRY ONLINE  
— TUITION —

	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><b>ignore</b> less haemoglobin</p> <p><b>A</b> carries <u>less</u> oxygen / nutrients...</p> <p><b>A</b> carries <u>less</u> carbon dioxide...</p> <p><b>I</b> 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><b>A</b> just A and S</p> <p><b>A</b> just S and S</p>
(iii)	0.25 / 25 % / $\frac{1}{4}$ / 1 in 4 ;	[1]	<b>I</b> 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</i> / ref to natural selection ;</p> <p>3 <math>H^A H^A</math> / homozygous dominant, susceptible to malaria ;</p> <p>4 <math>H^A H^S</math> / heterozygous, resistant ; <b>A</b> <math>H^S H^S</math> resistant ;</p> <p>5 <math>H^A H^S</math> survive / <math>H^A H^A</math> more likely to die before have children ;</p> <p>6 <math>H^A H^S</math> have children and pass on, the allele / <math>H^S</math> ;</p> <p>7 (if <math>H^A H^S \times H^A H^S</math>) 1 in 4 chance of, <math>H^S H^S</math> / homozygous recessive ;</p> <p>8 2 in 4 / <math>\frac{1}{2}</math> , 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><b>A</b> sickle cell trait / carrier for <math>H^S H^A</math> throughout the answer</p> <p><b>R</b> 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. Thalassaemia</p> <p><b>A</b> mutation described <b>I</b> gene, for allele</p>
	[Total:14]		

Question	Answers	Mark	Additional Guidance
4 (a)	halves the number of chromosomes / diploid to haploid ; <b>ignore</b> halves the genetic material  produces variation / AW ;	[2]	<b>accept</b> produces haploid, nuclei / cells / gametes <b>ignore</b> 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]	<b>R</b> 'from a single cell' but allow ecf for other MPs <b>R</b> egg divides  <b>A</b> same , genetic material / genetic make-up / genome <b>R</b> similar
(c)	increase in, complexity / AW ; ref to specialisation / differentiation ; ref to different types of cells ; ref to, tissues / organs ;	[max 2]	<b>ignore</b> (rapid) growth / change in shape <b>A</b> 'legs / arms / AW, start to grow'
(d)	1. $hY$ ; 2. $HX^h$ ; 3. $HX^H$ ;	[3]	<b>do not accept</b> male genotypes for <b>MP2</b> and <b>MP3</b>

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