- M1.(a) 1. Sugar-phosphate (backbone) / double stranded / helix so provides strength / stability / protects bases / protects hydrogen bonds; Must be a direct link / obvious to get the mark Neutral: reference to histones
 - 2. Long / large molecule **so** can store lots of information;
 - 3. Helix / coiled **so** compact; Accept: can store in a small amount of space for 'compact'
 - 4. Base sequence allows information to be stored / base sequence codes for amino acids / protein; *Accept: base sequence allows transcription*
 - 5. Double stranded **so** replication can occur semi-conservatively / strands can act as templates / complementary base pairing / A-T and G-C so accurate replication / identical copies can be made;
 - (Weak) hydrogen bonds for replication / unzipping / strand separation / many hydrogen bonds so stable / strong; Accept: 'H-bonds' for 'hydrogen bonds'
- 6

- (b) 1. (Mutation) in **E** produces highest risk / 1.78;
 - 2. (Mutation) in **D** produces next highest risk / 1.45;
 - (Mutation) in C produces least risk / 1.30;
 Must be stated directly and not implied
 E > D > C = 3 marks
 Accept: values of 0.78, 0.45 and 0.30 for MP1, MP2 and MP3 respectively
 If no mark is awarded, a principle mark can be given for the idea that all mutant alleles increase the risk

3

(c) **180**;

3.

1

(d) (Similarities):

- 1. Same / similar pattern / both decrease, stay the same then increase;
- 2. Number of cells stays the same for same length of time; Ignore: wrong days stated

(Differences):

(Per unit volume of blood)

 Greater / faster decrease in number of healthy cells / more healthy cells killed / healthy cells killed faster;

> Accept: converse for cancer cells Accept: greater <u>percentage</u> decrease in number of cancer cells / greater <u>proportion</u> of cancer cells killed

4. Greater / faster increase in number of healthy cells / more healthy cells replaced / divide / healthy cells replaced / divide faster;

Accept: converse for cancer cells For **differences**, statements made must be comparative

3 max

- (e) 1. More / too many healthy cells killed;
 - 2. (So) will take time to replace / increase in number; Neutral: will take time to 'repair'
 - 3. Person may die / have side effects;

2 max [15]

1

M2.(a) 250 000;

(b) (i) Loss of 3 bases / triplet = 2 marks;; 'Stop codon / code formed' = 1 mark max unless related to the last amino acid
Loss of base(s) = 1 mark; eg triplet for last amino acid is changed to a stop codon / code = 2 marks 3 bases / triplet forms an intron = 2 marks Accept: descriptions for 'intron' eg non-coding DNA 'Loss of codon' = 2 marks

2

- (ii) 1. Change in tertiary structure / active site; Neutral: change in 3D shape / structure 2. (So) faulty / non-functional protein / enzyme; Accept: reference to examples of loss of function eg fewer E-S complexes formed 2 [5] **M3.**(a) 1. Cell wall not formed / production inhibited; 1. Q Accept: weakened cell wall, but do not accept 'cell wall is broken down' 2. Lower water potential in bacterium; 2. Accept: converse 2. Must be clear that the lower water potential is in the bacterium 3. Water enters and causes lysis / expansion / pressure; 2 max Human cells lack enzyme (B) / have a different enzyme / produce different (b) fatty acids / use different substrates; Neutral: 'human cells do not have cell walls' as out of context 1 (c) 1. Change in base sequence (of DNA / gene) leading to change in amino
 - acid sequence / primary structure (of enzyme);
 - 1. Accept: different amino acids coded for
 - 1. Reject: different amino acids produced
 - 2. Change in hydrogen / ionic / disulphide bonds leading to change in the tertiary structure / active site (of enzyme);
 - 2. Neutral: alters 3D structure / 3D shape
 - 3. Substrate not complementary / cannot bind (to enzyme / active site) / no enzyme-substrate complexes form;

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3

M4.(a) (i) 4;

(ii)	1.	Change in amino acid / (sequence of) amino acids / primary structure; 1. Reject = different amino acids are 'formed'
	2.	Change in hydrogen / ionic / disulphide bonds alters tertiary structure / active site (of enzyme);
		2. Alters 3D structure on its own is not enough for this marking point.
	3.	Substrate not complementary / cannot bind (to enzyme / active site) / no enzyme- substrate complexes form;

3

1

(b) 1. Lack of skin pigment / pale / light skin / albino;

2.	Lack of coordination /	muscles action	affected.
۷.	Lack of coordination /		ancolou,

1

1

1

(c) Founder effect / colonies split off / migration / interbreeding; Allow description of interbreeding e.g. reproduction between individuals from different populations

[7]

M5. (a) Introns;

(b) Ile Gly Val Ser;

(c) (i) Has no effect / same amino acid (sequence) / same primary structure;

	Q Reject same amino acid formed or produced.	1
	Glycine named as same amino acid; It still codes for glycine = two marks.	1
(ii)	 Leu replaces Val / change in amino acid (sequence) / primary structure; Change in hydrogen / ionic bonds which alters tertiary structure / active site; <i>Q</i> Different amino acid formed or produced negates first marking point. Substrate cannot bind / no longer complementary / no enzyme-substrate complexes form; Active site changed must be clear for third marking point but does not need reference to shape. 	
(d) (i)	Interphase / S / synthesis (phase);	3
(ii)	DNA / gene replication / synthesis occurs / longest stage; Allow 'genetic information' = DNA. Allow 'copied' or 'formed' = replication / synthesis	1 [9