

**2007 'A' Level
H2 Biology
Mark Scheme**

PAPER 1 (MCQ)

1	C
2	B
3	B
4	C
5	B

21	B
22	C
23	D
24	A
25	D

6	D
7	B
8	C
9	B
10	C

26	A
27	C
28	D
29	D
30	C

11	D
12	A
13	C
14	A
15	A

31	B
32	D
33	A
34	C
35	D

16	D
17	C
18	C
19	D
20	C

36	A
37	C
38	A
39	D
40	D

PAPER 2 (CORE)**QUESTION 1****(a)**

A – viral RNA

B – reverse transcriptase

C – capsid protein

(b)

- 1 host cell surface membrane surrounds virus particle / (viral) RNA and protein;
- 2 ref. synthesis of surface glycoprotein;
- 3 budding; *R exocytosis*

(c)**(i)**

- 1 block protease active site so that substrate unable to bind;
- 2 viral polyprotein not cut / structural / enzyme protein not formed;
- 3 (newly formed) virus unable to infect further cells;

(ii)

- 1 similar shape to polyprotein / complementary shape to active site;
- 2 inhibitor not hydrolysed so remains in active site;

QUESTION 2**(a)**

- 1 fluid refers to the phospholipids and proteins being free to move;
- 2 phospholipids also being able to move within a layer, transversely;
- 3 mosaic refers to the proteins embedded in the phospholipid bilayer, where they are scattered;

(b)**(i)**

- 1 glucose is a polar molecule and ions are charged;
- 2 unable to pass through hydrophobic region within the membrane;
- 3 ref. to specificity of transport proteins;

(ii)

- 1 high levels of ATP close ATP-sensitive potassium channels;
- 2 build up of potassium ions inside cell;
- 3 depolarize membrane;

(iii)

- 1 insulin release involved exocytosis and therefore did not require a protein channel;
- 2 insulin is a protein;
- 3 too large to pass through membrane;
- 4 ref. to channel large enough to allow insulin through would allow other molecules through;

(c)

- 1 vesicles containing insulin move to cell surface membrane;
- 2 membrane of vesicle fuses with cell surface membrane;
- 3 vesicle contents emptied by exocytosis;

QUESTION 3**(a)**

- 1 DNA strands separate by hydrogen bonds breaking;
- 2 both strands act as templates;
- 3 each new molecule of DNA contains one new and one original strand of DNA;

(b)

- 1 ref. to shape of active site of DNA polymerase;
- 2 needs 3'OH to initiate action of DNA polymerase;
- 3 primers add only at 3';

(c)

- 1 DNA in bacteria occurs in a loop while human DNA is linear;
- 2 bacteria DNA is naked while human DNA is arranged in nucleosomes wrapped around histones;
- 3 ref. to introns only in human DNA;
- 4 ref. to telomeres only in human chromosomes;
- 5 ref. to centromeres only in human chromosomes;
- 6 ref. to operons only in bacteria;

(d)**(i)**

- 1 much longer DNA molecule;
- 2 DNA packaging in nucleosomes with DNA wrapped around histones
- 3 more non-coding DNA in human chromosomes;

(ii)

- 1 multiple sites of replication (on each chromosome);

QUESTION 4**(a)**

- 1 (inherited) change in DNA/nucleotide sequence/base sequence;
- 2 base substitution;
- 3 change in codon/triplet for amino acid/change in R group;
- 4 base addition/deletion;
- 5 ref. to frameshift or described;

(b)

- 1 (if addition/deletion resulting in) frameshift;
- 2 protein coded for no longer synthesized/non-functional;
- 3 (if substitution resulting in) amino acid changed;
- 4 change 3D/tertiary structure of protein/different R group;
- 5 ref. change in binding site *ref. to active site is neutral*

(c)

- 1 both copies of a tumor suppressor gene must be mutated (before a person will develop cancer);
- 2 ref. to effect of mutation being masked/ref. to normal allele being dominant/remaining functional copy of gene;
- 3 sufficient quantities of proteins synthesized to be effective;

(d)

- 1 specific shape of binding site of p53 protein;
- 2 corresponds to/complimentary to shape, part of DNA molecule;
- 3 different sequences of bases have different shapes;
- 4 ref. to major and minor grooves on DNA;

(e)

- 1 change in amino acid in region away from binding area;
- 2 may not affect shape or binding properties;
- 3 ref. idea some amino acids coded for by more than 1 codon/triplet of bases/ref. genetic code/some codons degenerate;
- 4 change in base/codon may not change amino acid/neutral;

QUESTION 5**(a)**

RRPP

RRPp

RrPP

RrPp

(b)

parents RRpp x rrPP;

gametes R_p x r_p

F1

RrPp;

RrPp x RrPp

gametes

RP, Rp, rP, rp

RP, Rp, rP, rp;

F2

walnut

rose

pea

single ;;

RRPP

RRp

rrPP

rrpp

RRPp x 2

Rrpp

rrPp

RrPp x 4

Rrpp

rrPp

Rrpp x 2

(allow on Punnett square, all genotypes must be shown, correspond to correct phenotype)

(c)

RrPP x RRpp

RRPP x Rrpp

RRPP x RRpp

QUESTION 6**(a)**

- P nucleus / euchromatin / nucleoplasm / chromatin; **R:** heterochromatin
Q nuclear envelope;
R cell surface membrane;

(b)

- 1 site of cellular respiration / oxidative phosphorylation / aerobic respiration;
- 2 synthesis / production / release of ATP;
- 3 ref. specific use of ATP (e.g. active transport / vesicle transport / mitosis / macromolecule synthesis, including DNA replication, transcription and translation);
- 4 ref. ATP produced by mitochondria released into cytosol while ATP in chloroplasts not;

(c)**(i)**

- 1 double membrane;
- 2 highly folded / large surface area of inner membrane;
- 3 presence of ETC / proton pumps;
- 4 ATP synthase;
- 5 70S ribosomes;
- 6 DNA loops;

(ii)

- 1 three membranes / double outer membrane in chloroplasts;
- 2 ref. position of stalked particles / chloroplasts "inside out"
- 3 starch grains;
- 4 stack of thylakoid membranes / grana;
- 5 ref. to thylakoids once;
- 6 no cristae in chloroplasts;
- 7 contain pigments / chlorophyll; **A:** other photosynthetic pigments

(d)

- 1 oxidative phosphorylation: final e^- (and H^+) acceptor / combines with e^- and H^+ to form water;
- 2 photophosphorylation: (waste) produce / from photolysis

QUESTION 7

(a) (dot correct answers then ticks bottom right of page)

animal ;
phylum ;
class ;
order ;
Dicrurus ;
species ;

6 correct = 4

5 correct = 3

4 correct = 2

3 correct = 1

(b)

- 1 ref. inherited variation/variation in alleles;
- 2 ref. natural selection;
- 3 differences in selection pressures in different habitats;
- 4 ref. to survive to pass on alleles;
- 5 geographical isolation / ref. allopatric;
- 6 idea that broadleaved woodland not continuous;
- 7 ref. islands;
- 8 prevents interbreeding / gene flow;
- 9 ref. to genetic drift/founder effect;

(c)

- 10 similar (morphological/anatomical/physiological) features;
- 11 capable of interbreeding;
- 12 to produce fertile offspring;

QUESTION 8**(a)**Cellulose

- 1 a polysaccharide;
- 2 (chain of) β /beta glucose;
- 3 1-4 links; (*glycosidic bonds not as important as linking bonds*)
- 4 ref. to microfibrils;
- 5 ref. to bundles of microfibrils/macrofibrils/fibres;

Collagen

- 1 a fibrous protein;
 - 2 (chain of) amino acids;
 - 3 ref. to peptide bonds;
 - 4 ref. to each chain helical;
 - 5 triple helix/tropocollagen;
 - 6 ref. to role/presence of glycine;
 - 7 ref. to staggered ends;
- 8 ref. to H bonds; A once in either

(b)Glycogen

- 1 a polysaccharide;
- 2 (chain of) α /alpha glucose;
- 3 1-4 with (some) 1-6 links/branches;
- 4 insoluble;
- 5 (energy) store;
- 6 stored/produced in muscle/liver;

Glucagon

- 1 a globular protein;
- 2 (chain of) amino acids;
- 3 ref. to peptide bonds;
- 4 soluble in plasma;
- 5 produced by α /alpha cells in pancreas/islets of Langerhans;
- 6 a hormone/regulates/raises blood glucose;

(c)

- 1 ref. energy reserves;
- 2 ref. lipids buoyancy;
- 3 ref. lipids (thermal) insulation;
- 4 ref. lipids more energy per unit mass/higher calorific value;
- 5 ref. to metabolic water;
- 6 ref. protection;

QUESTION 9**(a)**

- 1 length of DNA; (ref) a continuous piece/sequence
- 2 gene/ LacI, coding for repressor/inhibitor protein;
- 3 ref. to CAP (catabolic activator protein) binding site next to promoter;
- 4 promoter (gene);
- 5 RNA polymerase binding site;
- 6 ref. to operator;
- 7 binding site for repressor/inhibitor protein;
- 8 ref. to (3) structural gene;
- 9 Lac Z (galactosidase);
- 10 Lac Y (lactose permease);
- 11 Lac A (lactose transacetylase);
- 12 ref. to termination sequence; 8 max

(b)

- 1 lactose enters (bacterial) cell; *basal level of lac permease*
- 2 ref. to method of entry of lactose / qualified leaky membrane / small amount in;
- 3 allolactose / lactose binds to repressor protein;
- 4 results in a change in shape of repressor protein;
- 5 repressor protein no longer attached (to DNA);
- 6 exposes operator site;
- 7 allows RNA polymerase activity;
- 8 transcription (of genes coding for proteins required to utilise lactose);
- 9 leading to synthesis of proteins;
- 10 ref. named enzymes/proteins;

(c)

- 1 gene switching / majority of genes switched off / for regulation of gene expression;
- 2 so that bacterium only produces enzymes required;
- 3 inefficient / ref. waste of energy/resources **A converse**
- 4 ref. to selective advantage / disadvantage;
- 5 ref. group genes together for control;
- 6 ref. to bacteria able to use a variety of sugars/substrates

Paper 3 (Applications)**QUESTION 1****(a)****(i)**

bacterium/bacteria;

ignore correct qualification (e.g.named)**(ii)**

break phosphodiester bonds within polynucleotide strands/DNA;

(R) cut at, restriction site/gene**(iii)**

protection;

breaks down, foreign/viral/phage, DNA;

allow ecf from (i)**1 max****(b)****(i)**

1 (BamHI + HindIII + EcoRI) all produce, sticky ends/cohesive ends/overhangs/staggered ends;

2 allows, cut DNA/gene, to be, joined/annealed/spliced, to another DNA;
ignore vector

3 using complementary basepairing;

4 AluI + HaeIII require additional procedure to add on, sticky ends/linker DNA/caps;
ora **3 max****(ii)**

1 5' represents end (of strand with nucleotide) with 'free' carbon 5 (on deoxyribose);

2 3' represents end (of strand with nucleotide) with 'free' carbon 3 (on deoxyribose);

3 5' has phosphate/ 3' has hydroxyl;

4 DNA strands are antiparallel;

(c)**(i)****genomic**

1 DNA is broken (into fragments) using Restriction Enzyme(s);

Reject: treated2 fragments/DNA, incorporated into vector;

3 X phage/plasmid/cosmid/BAC/YAC; (BAC: bacteria artificial chromosome, YAC: yeast artificial chromosome)

4 create, further copies/clones, of DNA;

cDNA

5 start with mRNA;

6 to DNA using Reverse Transcriptase;

7 make double-stranded using DNA polymerase;

4max for any section

(ii)

the cDNA Library

- 1 smaller/ more compact;
- 2 contains only, coding sections (of DNA)/ exons;
- 3 introns have been removed/ no junk DNA;
- 4 easier to find desired gene (DNA clone)/no need for gene probes;
- 5 contains genes that are being expressed/reflects proteins actually present (in cell);
- 6 genes are intact;
(R) not been cleaved

credit reverse statements for genomic library**3 max**

QUESTION 2**(a)**Stage 1

- 1 denatures / H bonds broken;
- 2 (2) DNA strand separate;
- 3 by heating to 95⁰C; * **A: 90 – 100⁰C**

Stage 2

- 4 primer / oligonucleotides, bind / join to single (DNA) strands / 3' ends;
- 5 annealing;
- 6 by cooling to 54⁰C; * **A: 30 – 65⁰C**

Stage 3

- 7 new strands (of DNA) are synthesized by, TAQ / DNA polymerase;
- 8 starts at positions of primers / oligonucleotides;
- 9 by heating to 72⁰C; * **A: 60 – 75⁰C**

(b)**(i)**

2²⁶ / 67108864 **Look for 2²⁶ first**

(ii)

- 1 (may only be) very small amount (of original DNA) to work with;
- 2 can, amplify / produce many copies / produce very large amounts of DNA / genetic material;
- 3 (large amounts) make analysis easier;

(c)

- 1 DNA sample / genome (removed from organism) ;
- 2 RFLP / Restriction Fragment Length Polymorphism;
- 3 (sample DNA) cut with restriction enzymes,
- 4 gel electrophoresis / described;
- 5 radioactive probe + X-ray film or ethidium bromide / stain + UV light (to locate bands position)
- 6 wild and captive populations have different genetic profiles;
- 7 genetic fingerprints of captive individuals obtained;
- 8 compare fingerprint / DNA, with that of organism being investigated;
- 9 if bred in captivity there will be a match in, banding / pattern; ora

Comments by examiner:

- Students were weak in identifying how genetic fingerprinting is used to verify what type of organism (wild / bred in captivity) as many elaborated too much on the
- Genetic Fingerprinting method.
- Restriction Enzymes are not just added to the mixture but also used to 'cut' DNA

QUESTION 3**(a)****(i)**

- 1 derived from one parent/not a product of sexual reproduction ; (ignore asexual reproduction)
- 2 products of mitotic division;
- 3 genetically identical/share same DNA/same genotype/same genome/ same alleles; (R) same genes/similar

(ii)

- 1 much larger numbers can be produced/mass production ;
- 2 requires less space;
- 3 (cultures) easy to transport/ref. air freight;
- 4 ensures new plants are disease-free;
- 5 genetic modifications possible;
- 6 not limited by seasons/external conditions/weather; (R) faster techniques

(b)2,4-D

- 1 optimum peak/highest/described, is 0.3mg dm^{-3} with growth rate of 73/74;
- 2 idea that slight change from optimum is critical ;

Cytokinin

- 3 optimum peak/highest/described, is $8(.0)\text{mg dm}^{-3}$ with growth rate of 50;
- 4 not critical/large change from optimum to results in smallest decrease in growth (rate);

Thiamine

- 5 optimum peak/highest/described, is $1(.0)\text{mg dm}^{-3}$ with growth rate of 61/62;
- 6 marked effect/critical from 0.8mg mg dm^{-3} ;
- 7 2,4-D has highest growth rate/cytokinin has lowest growth rate;
- 8 cytokinin required at highest concentration / 2,4-D required at lowest concentration;
- 9 ref. qualified pattern of variation for one component;
- 10 use a mixture of all 3 at optimal levels;
- 11 idea of possible interaction between components;

Comments by examiner:

- Arbitrary units not required in this question but many not be true for all questions.
- Guide students to state the complete units in all questions wherever possible.
- Need to observe scale of graph
- Some form of comparison between graphs wanted by examiner though not very clearly stated in question

(c)**(i)**

- 1 unable to fix nitrogen;
- 2 to produce, amino acids/proteins;
- 3 e.g. of required protein ; (e.g. enzymes/cytochrome/etc)
- 4 DNA/RNA/nucleic acid/nitrogenous base/named base/nucleotide;
- 5 ATP;
- 6 chlorophyll;

(ii)

- 1 ammonium nitrate contains more nitrogen;
- 2 may be, easier to assimilate/easier to absorb/more accessible, ammonium (ion rather than potassium ion);

QUESTION 4**(a)**SCID

- 1 more males affected (than females)
- 2 recessive
- 3 (most common form is) sex-linked (X-chromosome)
- 4 mutation of IL2RG (interleukin-2 receptor gamma) gene (for gamma chain on lymphocyte receptor)
- 5 ADA (adenosine deaminase) deficiency
- 6 mutation of gene on chromosome 20
- 7 prevents formation of , lymphocytes / T cells / B cells
- 8 toxic metabolites not broken down so lymphocytes killed
- 9 so susceptible to opportunistic infections (award only if S7 and S8 awarded)
- 10 children usually die (if not treated)
- 11 pneumonia / PCP / meningitis / chicken pox / measles / other valid

Cystic fibrosis

- 1 autosomal / chromosome 7
- 2 recessive
- 3 mutation of, CFTR / CFTCR gene
- 4 deletion of, 3 base (pairs) / codon
- 5 CFTR / CRTCR, protein has lost phenylalanine
- 6 (channels) no longer able to transport chloride ions
- 7 causing thick mucus
- 8 resulting in stated problem in, lungs / pancreas / reproductive system
- 9 reduces life expectancy

(b)

- 1 gene therapy is a technique for introducing 'normal', allele/gene; *if "therapeutic" need to give further explanation*
- 2 new, allele/gene, generates, functional/correct, protein product; *accept specific example of protein product*
- 3 restores target cell to normal state / corrects function of cell / corrects phenotype;

Viral delivery systems

- 1 viruses are genetically engineered to carry 'normal' human gene;
- 2 adenoviruses/retroviruses/MMLV (for SCID);
- 3 virus releases genetic material into target cells;
- 4 in SCID the gene is inserted (by viruses) into, bone marrow cells / blood stem cells / lymphocytes, *ex vivo / in vitro*;
- 5 in cystic fibrosis the gene is inserted into cells within body;

Non-viral delivery systems

- 1 direct introduction of DNA into target cells (gene gun or other direct method e.g. micropipette);
- 2 creation of, lipid sphere / liposome, which carries DNA;
- 3 liposome can / use of electroporation to, pass through target cell membrane;
must be linked to pt 10
- 4 link DNA to molecule that will bind to receptors in cell membrane;
- 5 DNA is engulfed by membrane;
- 6 possibility of introducing 47th chromosome into cells;

(c)

- 1 difficult to get DNA to integrate into target cell genome;
- 2 results can be short-lived / patients need to be treated on a frequent basis;
- 3 problem with controlling the activity of gene expression;
- 4 risk of stimulating immune / allergic response;
- 5 virus vector may, regain / develop, virulence;
- 6 incorrect insertion of gene may cause, named / described, problem to patient;
- 7 possible problem with finding vector for large gene;
- 8 many genetic diseases are a result of presence of many defective genes / multi-allelic / multigene conditions;
- 9 impossible to introduce many genes at the same time;
- 10 problem dealing with, dominant condition / non-dividing cells;
- 11 ref. problem with liposomes; (toxicity / less efficiency);