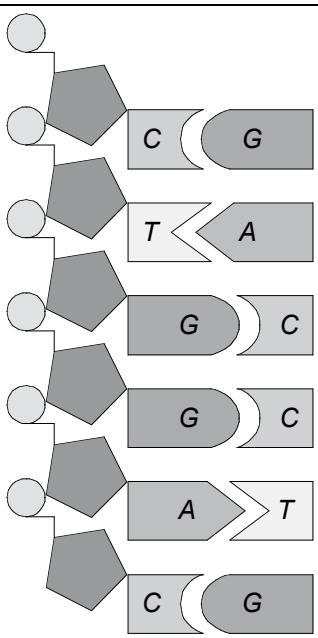


**Assessment Schedule – 2005****Biology: Describe the transfer of genetic information (90163)****Evidence Statement**

| <b>Q</b> | <b>Achievement</b>   | <b>Achievement with Merit</b>   | <b>Achievement with Excellence</b>   |   |   |    |    |   |    |    |  |   |   |   |    |    |   |    |    |  |  |
|----------|--|---|--|---|---|----|----|---|----|----|--|---|---|---|----|----|---|----|----|--|--|
| 1(a)     | Homozygous recessive; homozygous dominant; heterozygous rr; RR; Rr.<br>At least two must be correct, with the corresponding symbols.   |   |  |   |   |    |    |   |    |    |  |   |   |   |    |    |   |    |    |  |  |
| 1(b)     | Correctly completed Punnett squares<br>1.<br><table border="1" style="display: inline-table; vertical-align: middle;"> <tr> <td></td><td style="text-align: center;">R</td><td style="text-align: center;">R</td></tr> <tr> <td style="text-align: center;">r</td><td style="text-align: center;">Rr</td><td style="text-align: center;">Rr</td></tr> <tr> <td style="text-align: center;">r</td><td style="text-align: center;">Rr</td><td style="text-align: center;">Rr</td></tr> </table><br>Zero (0%)<br><br>2.<br><table border="1" style="display: inline-table; vertical-align: middle;"> <tr> <td></td><td style="text-align: center;">R</td><td style="text-align: center;">r</td></tr> <tr> <td style="text-align: center;">r</td><td style="text-align: center;">Rr</td><td style="text-align: center;">rr</td></tr> <tr> <td style="text-align: center;">r</td><td style="text-align: center;">Rr</td><td style="text-align: center;">rr</td></tr> </table><br>1/2 (50%)<br><br>Both Punnett squares and one of the ratios must be correct. |   | R  | R | r | Rr | Rr | r | Rr | Rr |  | R | r | r | Rr | rr | r | Rr | rr |  |  |
|          | R  | R   |  |   |   |    |    |   |    |    |  |   |   |   |    |    |   |    |    |  |  |
| r        | Rr   | Rr  |  |   |   |    |    |   |    |    |  |   |   |   |    |    |   |    |    |  |  |
| r        | Rr   | Rr  |  |   |   |    |    |   |    |    |  |   |   |   |    |    |   |    |    |  |  |
|          | R  | r   |  |   |   |    |    |   |    |    |  |   |   |   |    |    |   |    |    |  |  |
| r        | Rr   | rr  |  |   |   |    |    |   |    |    |  |   |   |   |    |    |   |    |    |  |  |
| r        | Rr   | rr  |  |   |   |    |    |   |    |    |  |   |   |   |    |    |   |    |    |  |  |
| 1(c)     | Different genetic make-ups can give the same flower colour.  | Explanation that whilst an allele may be present (in genotype) it may not be seen (in phenotype) thus altering the ratios.  |  |   |   |    |    |   |    |    |  |   |   |   |    |    |   |    |    |  |  |
| 1(d)     | Cross (breed) with a white flowered plant.<br><br><b>OR</b><br><br>Completing a test / back cross  | Explains that a test cross involves crossing the red flowered plant with a homozygous recessive (white plant).<br><br>Plus gives a reason for cross with a white flowered plant of recessive phenotype.<br>Eg If any white flowered offspring are seen the parent must be heterozygous. | Discussion that involves the need for a <b>back-cross (test-cross)</b> , ie crossing the red flower with homozygous recessive to see if any recessive character is shown (white flower) or not. Discusses the significance of the outcome in identifying the parent genotype.<br>Eg the white offspring show that the unknown genotype of the red plant must have included a white allele which combined with the white plants alleles to create the white plant.<br><b>OR</b> discusses fact that a heterozygous genotype can be determined by a test cross but not a homozygous dominant genotype. |   |   |    |    |   |    |    |  |   |   |   |    |    |   |    |    |  |  |

| Q    | Achievement   | Achievement with Merit   | Achievement with Excellence  |
|------|---|--|--|
| 2(a) |  <p>Diagram completed as above (must be correct pairing) <b>AND</b> any TWO of the three correctly identified: phosphate; sugar; nucleotide.</p> |  |  |
| 2(b) | <p>Describes the importance of the code not changing.<br/>Eg to give exact / identical copies of the DNA.</p>   | <p>Explains how the DNA is replicated ie. the C-G pairs and A-T pairs are vital. (Random pairings would change the code.)<br/><b>OR</b><br/>Why identical replication is necessary.<br/>Eg. A change to the code will result in a mutation.</p>        |  |
| 2(c) | <p>Description of how DNA carries information:<br/>Eg DNA makes up genes.<br/>Recognises DNA as the carrier of the code.</p>  | <p>Explanation of how DNA carries information:<br/>The order of bases affects the gene properties.<br/>Red and white are the result of a different base order / sequence.<br/>Combination of bases makes the characteristics.</p>                      | <p>Discuss the fact that the base sequence is a code for building amino acids / proteins<br/><b>OR</b><br/>That the different sequences give different characteristics such as the flower colour. Red and white are different sequences of bases <b>giving different alleles of the same gene</b>.</p> |
| 3(a) | For growth and repair.  |  |  |
| 3(b) | Mitosis produces the wrong type of cells.   | <p>Explanation eg:<br/>Sperm or egg cells have to be HAPLOID / half the genetic information.<br/><b>OR</b><br/>Mitosis produces cells, DIPLOID / full genetic information, which if combined would give too much information to produce offspring.</p> |  |

| Q    | Achievement  | Achievement with Merit  | Achievement with Excellence  |
|------|--|---|--|
| 3(c) | <p>Describes that the cells divide twice / produce gametes with half the genetic information / variation is needed in order to produce offspring with a greater chance of survival.</p> <p><b>OR</b></p> <p>Describe crossing over or segregation.</p> | <p>Explanation that cell division occurs twice / have half genetic information / are haploid.</p> <p><b>AND</b> that this allows for variety, which contributes to an increased chance of survival for the offspring.</p> | <p>Discussion includes the mention of homologous pairs and exchange of material in <b>crossing over</b></p> <p><b>OR</b></p> <p>the fact that one of each pair of homologous chromosomes goes to a different daughter cell (<b>segregation</b>)</p> <p><b>AND</b> the value of variation to provide offspring with a greater chance of survival in successive generations in a changing environment.</p>                               |
| 4(a) | 2/3rds or (66.66%) or 2 out of 3   |   |  |
| 4(b) | <p>She is heterozygous.</p> <p><b>OR</b></p> <p>She is the carrier of the recessive allele</p>   | <p>She must be heterozygous because some of her offspring are recessive (CF sufferers).</p> <p><b>OR</b></p> <p>If she was homozygous, her offspring would carry the dominant allele (be carriers NOT sufferers).</p>     |  |
| 4(c) | The phenotype of female C is normal / not affected by CF.  | <p>The phenotype of female C is normal / not affected by CF</p> <p><b>PLUS</b></p> <p>Explanation for the appearance of ONE set of children (from normal or carrier male) is correct.</p>                                 | <p>If she has children with B who must be a carrier, then 75% of them would appear normal whilst 25% could show signs of CF. 50% of her children would be carriers but we couldn't tell this from their phenotype.</p> <p>If she has children with a normal male, then all her offspring would appear normal, but 50% of them would be carriers.</p> <p>For male B instead of ratios may use information from Punnett or pedigree.</p> |

### Judgement Statement

| Achievement   | Achievement with Merit   | Achievement with Excellence   |
|---|--|---|
| <p>Total of SEVEN opportunities answered at Achievement or higher.</p> <p><math>7 \times A</math></p> | <p>Total of TEN opportunities answered.</p> <p>THREE at Merit level or higher<br/><i>and</i><br/>SEVEN at Achievement level.</p> <p><math>3 \times M + 7 \times A</math></p> | <p>Total of TWELVE opportunities answered.</p> <p>TWO at Excellence level<br/><i>and</i><br/>THREE at Merit level<br/><i>and</i><br/>SEVEN at Achievement level.</p> <p><math>2 \times E + 3 \times M + 7 \times A</math></p> |