

## Scheme of Work 2020-2021

### Subject: Biology

Year Group: 12 Summer Term 1

Specification: AQA

Lesson No	Topic & Objectives	Big Question – What will students learn?	Key Activities & Specialist Terminology (Do Now Task / Starter/Tasks/Plenary)	Planned Assessment	Homework or flipped learning resources  DODDLE resources	Lit Num SMSC Codes
1 3.4.1 <b>DNA, genes and chromosomes.</b> cholesterol	<p>Eukaryotic cells have chromosomes of linear DNA associated with histones.</p> <p>Prokaryotic cells contain short, circular DNA that is not associated with histones.</p> <p>Mitochondria and chloroplasts contain DNA like that of prokaryotes.</p> <p>A gene is a base sequence of</p>	<ul style="list-style-type: none"> <li>Explain what is meant by the terms chromosome and gene.</li> <li>Compare and contrast DNA in eukaryotes with that in prokaryotes, mitochondria and chloroplasts.</li> <li>Explain what a gene could code for.</li> </ul>	<p><b>Learning activities:</b></p> <ul style="list-style-type: none"> <li>questioning from GCSE about the meaning of key terms like gene, chromosome and allele</li> <li>use animation to show scale of chromosomes in eukaryotic cells and how chromosomes are made of DNA and histones. Introduce the concept of a gene</li> <li>teacher explanation about the difference between the arrangement of DNA in prokaryotic cells and eukaryotic cells</li> <li>students generate a summary table comparing and contrasting prokaryotic and eukaryotic DNA.</li> </ul> <p><b>Skills developed by learning activities:</b></p> <ul style="list-style-type: none"> <li>MS 0.2 – students can be introduced to base pairs/kilobase pairs as a measuring of length when discussing the loci of a gene on a chromosome and convert this from standard to ordinary form</li> </ul> <p>AO1 – development of knowledge and understanding of the arrangement of DNA in eukaryotes and prokaryotes and the</p>		<p><a href="http://yourgenome.org/teachers/zoom.shtml">yourgenome.org/teachers/zoom.shtml</a></p> <p><b>Rich question:</b></p> <p>A textbook stated that “The bacterial chromosome is found in the cytoplasm of the cell”. Evaluate this statement.</p>	C1,C3,Sp2

	DNA that codes for the amino acid sequence of a polypeptide or a functional RNA.		relationship between DNA, genes and chromosomes.			
2	<p>DNA has a triplet code which is universal, non-overlapping and degenerate. Much of eukaryotic DNA does not code for polypeptides. There are non-coding regions of multiple base repeats between genes. There are also introns within genes which separate coding sequences (exons).</p>	<ul style="list-style-type: none"> <li>Explain how the DNA base sequence is able to code for the primary structure of a polypeptide.</li> <li>Explain the terms degenerate, universal and non-overlapping.</li> <li>Explain why much of eukaryotic DNA can be considered as non-coding.</li> </ul> <p>Explain what is meant by an intron and an exon</p>	<p><b>Learning activities:</b></p> <ul style="list-style-type: none"> <li>remind students that there are 20 amino acids and only 4 bases. Ask how many bases would have to code for an amino acid to give sufficient combinations</li> <li>teacher explanation of the triplet code and the fact that there is degeneracy (as well as the fact it is universal and non-overlapping)</li> <li>ask the rich question: how many bases code for a polypeptide of 24 amino acids</li> <li>explain why the answer might in fact be more than 72 as there are introns in the gene. Introduce the idea of introns and also non-coding regions between genes</li> <li>exam questions.</li> </ul> <p><b>Skills developed by learning activities:</b></p> <ul style="list-style-type: none"> <li>MS 0.3 – students could calculate the percentage of human DNA which does code for polypeptides, when supplied with data about the number of coding bases and the total number of bases</li> <li>MS 0.5 – students could work out the possible number of combinations that a triplet code can have (ie 43) to highlight the idea of degeneracy</li> </ul>	<p><b>Past exam paper material:</b></p> <p>BIOL2 June 12 Q5b</p> <p>BIOL2 June 2011 – Q3a</p> <p>BIOL2 Jan 2010 – Q3</p>	<p><a href="http://yourgenome.org/teachers/dnaprotein.shtml">yourgenome.org/teachers/dnaprotein.shtml</a></p> <p><b>Rich questions:</b></p> <ul style="list-style-type: none"> <li>What is meant by the terms: <ul style="list-style-type: none"> <li>degenerate?</li> <li>non-overlapping?</li> <li>universal?</li> </ul> </li> </ul> <p>A polypeptide is made of 24 amino acids. What is the minimum number of bases that the gene coding for it must have had?</p>	C1,C3,Sp2

			AO1 – development of knowledge and understanding of the triplet code and non-coding sections of it.			
3 3.4.2 Protein synthesis.	<p>The concept of the genome and the proteome.</p> <p>The structure of molecules of mRNA.</p> <p>The process of transcription in prokaryotes to produce mRNA.</p> <p>The process of transcription in eukaryotes to produce pre-mRNA which is subsequently spliced..</p>	<ul style="list-style-type: none"> <li>Explain what the terms genome and proteome mean.</li> <li>Describe the structure of mRNA and how it is related to its function (link to 3.1.5.1).</li> <li>Explain the process of transcription in prokaryotes.</li> <li>Explain the process of transcription and splicing in eukaryotes, linking this to knowledge of introns.</li> </ul> <p>Interpret data from experimental work investigating the role of nucleic acids.</p>	<p><b>Learning activities:</b></p> <ul style="list-style-type: none"> <li>questioning to recap knowledge about the role of DNA and RNA from section 3.1.5</li> <li>provide students with data from experimental work investigating the role of nucleic acids eg the Hershey-Chase experiment and ask them to interpret this</li> <li>introduce concept of genome and proteome</li> <li>teacher explanation of the process of transcription and how the structure of mRNA relates to its function of transferring the code to the ribosomes. Use animation to support this.</li> </ul> <p><b>Skills developed by learning activities:</b></p> <ul style="list-style-type: none"> <li>PS 1.2 - apply knowledge of transcription and nucleic acids to explain experimental data from investigations into the role of nucleic acids</li> <li>AO1 – development of knowledge around transcription and the structure and role of mRNA</li> </ul> <p>AO2 – application of knowledge to transcribe a DNA sequence into mRNA</p>	<p><b>Past exam paper material:</b></p> <p>BIOL5 June 2010 – Q2</p> <p>BIOL5 June 2011 – Q1</p>	<p><a href="http://yourgenome.org/teachers/dnaprotein.shtml">yourgenome.org/teachers/dnaprotein.shtml</a></p> <p><b>Rich questions:</b></p> <ul style="list-style-type: none"> <li>What are the advantages of mRNA being used to carry the genetic code to the ribosomes, rather than DNA?</li> <li>Explain how mRNA is adapted to its function.</li> <li>What is the difference between mRNA and pre-mRNA?</li> </ul> <p>Provide students with a DNA code, identify the sense strand and ask students to transcribe it (assuming there are no introns).</p>	So5,Sp2 M2
4	<p>The process of translation.</p> <p>The roles of ribosomes, tRNA and ATP.</p>	<ul style="list-style-type: none"> <li>Explain the process of translation.</li> <li>Explain the specific roles of ribosomes, ATP and tRNA in translation.</li> </ul>	<p><b>Learning activities:</b></p> <ul style="list-style-type: none"> <li>questioning to recap knowledge about transcription, the role of ribosomes from section 3.2.1 and ATP from section 3.1.6</li> <li>teacher explanation of the process of translation and how the structure of tRNA relates to its function in delivering the</li> </ul>	<p>Students could undertake the investigations/questions from the following ISAs:</p> <p>BIO3T P14</p>	<p><a href="http://yourgenome.org/teachers/dnaprotein.shtml">yourgenome.org/teachers/dnaprotein.shtml</a></p> <p><b>Rich questions:</b></p> <ul style="list-style-type: none"> <li>Evaluate the statement “DNA is a</li> </ul>	So5,Sp2 M2

	<p>The structure of molecules of tRNA.</p>	<ul style="list-style-type: none"> <li>Describe the structure of tRNA and how it is related to its function.</li> </ul> <p>Relate the base sequence of nucleic acids to the amino acid sequence of polypeptides, when provided with suitable data about the genetic code.</p>	<p>specific amino acid. Use animation to support this</p> <ul style="list-style-type: none"> <li>exam questions.</li> </ul> <p><b>Skills developed by learning activities:</b></p> <ul style="list-style-type: none"> <li>AO1 – development of knowledge around translation and the structure and role of tRNA</li> </ul> <p>AO2 – application of knowledge to translate a mRNA sequence into a sequence of amino acids.</p>	<p><b>Specimen assessment material:</b></p> <p>A-level Paper 1 (set 1) – Q11.1</p> <p><b>Past exam paper material:</b></p> <p>BIOL5 June 2012 – Q1 (except Q1cii and 1d)</p>	<p>triplet code which instructs the ribosomes how to make amino acids”.</p> <ul style="list-style-type: none"> <li>Explain how the structure of tRNA is adapted for its function.</li> </ul> <p>Provide students with an mRNA code and ask them to translate it into an amino acid sequence (when provided with appropriate information).</p>	
<p>5 3.4.3 <b>Genetic diversity can arise as a result of mutation or during meiosis</b></p>	<p>Gene mutations arise spontaneously during DNA replication and include base deletion and base substitution.</p> <p>The degeneracy of the genetic code means that not all base substitutions cause a change in the</p>	<ul style="list-style-type: none"> <li>Explain what a gene mutation is and how it arises.</li> <li>Explain what is meant by a deletion and substitution mutation and the potential consequences of each (linked to primary protein structure).</li> <li>Interpret base sequences to identify gene mutations and their impact.</li> </ul> <p>Describe what a mutagenic agent is and identify some possible mutagenic agents.</p>	<p><b>Learning activities:</b></p> <ul style="list-style-type: none"> <li>teacher led explanation of how gene mutations arise and mutagenic agents which can increase the risk</li> <li>students work through the transcription and translation activity (linked in resources). Then ask them to repeat the activity twice more but this time putting in a substitution mutation for one and a deletion mutation for another. Compare effects of the two mutations to the original amino acid sequence. Ask students to relate these effects to their knowledge of protein structure</li> <li>teacher explanation of the effects of substitution and deletion mutations and also the possible neutral effects of substitution due to degeneracy.</li> </ul> <p>exam questions</p>	<p><b>Specimen assessment material:</b></p> <p>AS Paper 2 (set 2) – Q3</p> <p><b>Past exam paper material:</b></p> <p>BIOL2 Jan 2013 – Q6a–6</p> <p>BIOL2 June 2013 – Q7b–7c</p> <p>BIOL2 Jan 2012 – Q4</p> <p>BIOL2 June 2011 – Q3b</p> <p>BIOL2 June 2010 – Q3</p>	<p><a href="http://cell-cell-cell.com/wp-content/uploads/CCC_Activity_CrackTheCodon_v01.doc">cell-cell-cell.com/wp-content/uploads/CCC_Activity_CrackTheCodon_v01.doc</a></p> <p><b>Rich questions:</b></p> <ul style="list-style-type: none"> <li>Evaluate this statement: “Sunbathing exposes your body to UV light which causes mutations to occur”.</li> <li>Which type of gene mutation is likely to be the most damaging and why?</li> </ul> <p>A student wrote that UV light increased the</p>	<p>Sp3,Sp2 C1</p>

	<p>amino acid sequence.</p> <p>Mutagenic agents can increase the risk of gene mutation.</p>		<p><b>Skills developed by learning activities:</b></p> <ul style="list-style-type: none"> <li>• AO1 – development of knowledge around gene mutations and their possible consequences</li> </ul> <p>AO2 – application of knowledge of mutation to a model of protein synthesis model to suggest possible effects of gene mutation on the structure of the protein produced.</p>		<p>likelihood of mutations in the protein that the cell made. Why is this not correct?</p>	
6	<p>Meiosis produces genetically unique daughter cells.</p> <p>The process of meiosis involves two nuclear divisions and forms four haploid daughter cells.</p> <p>Independent segregation and crossing over result in genetically different daughter cells.</p>	<ul style="list-style-type: none"> <li>• Explain the different outcome of mitosis and meiosis.</li> <li>• Explain how meiosis results in variation.</li> <li>• Complete diagrams showing the chromosome content of cells after the first and second meiotic division, when given the chromosome content of the parent cell.</li> <li>• Recognise where meiosis occurs when given information about an unfamiliar life cycle.</li> </ul> <p>Explain how random fertilisation of haploid gametes further increases</p>	<p><b>Learning activities:</b></p> <ul style="list-style-type: none"> <li>• introduce the convention of <math>2n</math> and <math>n</math>. Students then calculate the number of possible chromosome combinations (without crossing over)</li> <li>• think, pair, share: there is more variation possible than our calculated number – where does the extra variation come from?</li> <li>• teacher explanation of the process of meiosis, supported by animations and videos</li> <li>• students compare and contrast mitosis and meiosis</li> <li>• students interpret information about unfamiliar life cycles to identify where meiosis and mitosis are occurring.</li> </ul> <p><b>Skills developed by learning activities:</b></p> <ul style="list-style-type: none"> <li>• MS 0.5 – use the expression <math>2n</math> to calculate the possible number of different combinations of chromosomes</li> <li>• MS 0.5 – derive a formula from this to calculate the possible number of different combinations of chromosomes following random fertilisation</li> <li>• 8.4.2.1 and 8.4.2.2</li> </ul>	<p><b>Specimen assessment material:</b></p> <p>A-level Paper 1 (set 1) – Q10; AS Paper 1 (set 1) – Q3.</p> <p><b>Past exam paper material:</b></p> <p>BIOL2 June 2013 – Q1; BIOL2 June 2010 – Q5</p>	<p><a href="http://nuffieldfoundation.org/practical-biology/preparing-anther-squash">nuffieldfoundation.org/practical-biology/preparing-anther-squash</a></p> <p><a href="http://cleapss.org.uk">cleapss.org.uk</a></p> <p><a href="http://highered.mheducation.com/sites/0072495855/student_view0/chapter3/animation_how_meiosis_works.html">highered.mheducation.com/sites/0072495855/student_view0/chapter3/animation_how_meiosis_works.html</a></p> <p><a href="http://highered.mheducation.com/sites/0072495855/student_view0/chapter2/animation_comparison_of_meiosis_and_mitosis_quiz_1.html">highered.mheducation.com/sites/0072495855/student_view0/chapter2/animation_comparison_of_meiosis_and_mitosis_quiz_1.html</a></p> <p><a href="http://sumanasinc.com/webcontent/animations/content/meiosis.html">sumanasinc.com/webcontent/animations/content/meiosis.html</a></p> <p><b>Rich question:</b></p> <p>Compare and contrast the similarities and differences</p>	Sp3,Sp2 C1

			<ul style="list-style-type: none"><li>• AO1 – development of knowledge of meiosis</li></ul> AO2 – application of knowledge to unknown life cycles.		between mitosis and meiosis.	
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