		Riology	Ι.Δ.	Novel - ve	nar 12	EOY Exam	Sequencing and Progression	
KAA Curriculum Overview Biology			A level - year 13					
Rationale Following on from year 12, year 13 students begin learning about energy transfers within and between organisms. They then study the nervous system and how organisms respond to stimuli. Pupils study DNA and inheritance before they move on to gene expression and DNA technology. Pupils are provided with opportunities to apply their knowledge of biological molecules and processes learnt in Y12 to various contexts in Y13. It is essential pupils have an understanding of how the sun's energy sustains life on earth so pupils begin the year with a focus on bioenergetics. As the year progresses, pupils gain an in depth look at genetics before ending the year focussing on fundamental disciplinary knowledge within the study of genetics and how it can be applied in real life contexts such as genetic fingerprinting. As with Y12, required practicals are taught in conjunction with the theory within lessons. Pupils will develop independent thinking and problem solving skills using practical experience gained in Y12. Pupils will ultimately be assessed by teachers against Common Practical Assessment Criteria (CPAC) in order to obtain a practical endorsement by AQA.					All three assessment points across the year assess AO1, AO2 and AO3 skills. The end of year A level papers are mock papers 1 to 3, covering all AS and A level content. Roughly 10% of marks will be attributed to mathematical skills and 15% to practical skills. The 2O22 mocks can be found here: https://drive.google.com/drive/folders/1AGWkikNOKJU3ntChpgLE9lm1nwU1fHB	The content studied in year 12 forms the basis for understanding units 5 to 8 in year 13. Direct links are made between AS and A2 topics - for example, between biological molecules and bioenergetics.	Studying A level biology develops resilience, and builds students' practical, analytical and evaluative skills. These competencies will well equip them for anything they decide to undertake after year 13.	
Term	Autumn 1		Autumn 2		Spring 1	Spring 2	Sum 1	Sum 2
Link to MTP Overview								
Topic studied & Fertile Question	recap, respi ecosystems	photosynthesis ration & energy in (unit 5) inherited change	Teacher 1 - response to st (unit 6) Teacher 2 - populations, evolution and ecosystems 7)		Teacher 1 - nervous coordination and muscles (unit 6) Teacher 2 - gene expression (unit 8)	Teacher 1 - homeostasis (unit 6) Teacher 2 - DNA technology (unit 8)	Revision	Revision
Adjustments following last assessments / evaluation.	None		None		None	None	None	None
Key knowledge and skills students need to have gained by the end of the unit	sym pho - Expl of p - Deso in th reac - Deso in th reac - Deso effe facto pho - Deso	esis e the word and bol equation for tosynthesis. lain the importance hotosynthesis. cribe what happens he light dependent ction. cribe what happens he light independent ction. cribe and explain the ct of environmental ors on the rate of tosynthesis. cribe some cultural process used	Response to stimuli Define taxis and e some examples in organisms. Define kinesis and explain some examin organisms. Define tropism an explain some examin plants. State the role of laregulating growth plants and describit works. Describe the structure the nervous syste. Describe the reflerand explain the importance of ref	d amples amples IAA on th in the how acture of em. ex arc	Knowledge Nervous coordination and muscles - Describe structure of a myelinated motor neurone. - Describe how the resting potential is established Describe how action potentials are generated Compare the passage of an action potential along unmyelinated and myelinated axons Describe the refractory period and explain its importance Describe and explain the factors affecting the speed of conductance:	Knowledge Homeostasis Define homeostasis. Explain the importance of maintaining body temperature, blood pH, blood glucose and blood water potential. Define negative feedback and give examples. State the role of the liver in glycogenesis, glycogenolysis and gluconeogenesis. Describe the actions of insulin, glucagon and adrenaline in maintaining blood glucose levels. Describe the second messenger model of		

to overcome these limiting factors.

Respiration

- State the word and symbol equations for aerobic and anaerobic respiration.
- Explain the importance of respiration.
- Describe what happens in glycolysis, the Link reaction, the Krebs cycle and oxidative phosphorylation.
- Describe the process of anaerobic respiration in detail and the products formed, and compare this to the process of aerobic respiration.

Energy in ecosystems

- Define biomass and describe how to calculate the chemical energy stored in dry biomass.
- Define GPP and NPP.
- Describe how energy is lost through trophic levels of a food chain.
- Appreciate the ways in which production is affected by farming practices designed to increase the efficiency of energy transfer.
- Students must be able to describe the stages of the nitrogen and phosphorus cycles.
- Describe the phosphorus cycle.
- Describe the nitrogen cycle.
- Describe the role of microorganisms in the nitrogen and phosphorus cycle.
- Explain why farmers use fertilisers.
- Explain the
 environmental issues
 arising from the use of
 fertilisers, such as
 leaching and
 eutrophication.
- Inherited change

- Describe the basic structure of a Pacinian corpuscle and label the parts on diagrams.
- Describe how deformation of stretch-mediated sodium ion channels in a Pacinian corpuscle leads to the establishment of a generator potential.
- Describe the distribution of rods and cones in the retina.
- Identify the pigments in rod and cone cells.
- Explain how rod cells' visual acuity, sensitivity to light and sensitivity to colour are accounted for by the presence of rhodopsin and connections to the optic nerve.
- Explain how cone cells' visual acuity, sensitivity to light and sensitivity to colour are accounted for by the presence of different forms of iodopsin and connections to the optic nerve.
- Explain the importance of constriction and dilation of the pupil in response to changes in light.
- Describe the role of the radial and circular muscles in accommodation.
- Describe the location of, and the role played by, chemoreceptors and pressure receptors involved in detecting changes which lead to changes in heart rate.
- Explain what is meant by the sympathetic and parasympathetic nervous system and explain their role in controlling heart rate. Describe the role of the medulla oblongata.

- myelination and saltatory conduction; axon diameter; temperature.
- Describe the detailed structure of a synapse and of a neuromuscular
- junction.
- Describe transmission across a cholinergic synapse.
- Compare transmission across a cholinergic synapse and across a neuromuscular junction.
- Predict and explain the effects of specific drugs on a synapse.
- Describe the structure of skeletal muscle and a myofibril.
- Describe the roles of actin, myosin, calcium ions, tropomyosin, troponin and ATP in myofibril contraction.
- Compare slow and fast twitch skeletal muscle fibres.

Gene expression

- Define mutation and explain each type.
- Describe and explain the effects of mutations on proteins.
- Define a mutagen and give an example.
- Compare totipotent, pluripotent and unipotent stem cells, including what they can do, and when and where they are produced.
- Describe how induced pluripotent stem cells are produced.
- Describe and explain the role of transcription factors in gene expression.
- Describe the role of the steroid hormone, oestrogen, in initiating transcription.
- Define epigenetics.
- Explain the effect of increased methylation of the DND and decreased

- adrenaline and glucagon action, involving adenylate cyclase, cyclic AMP (cAMP) and protein kinase.
- State the causes of types

 I and II diabetes and their
 control by insulin and/or
 manipulation of the diet.

DNA technology

- State what recombinant DNA is.
- Describe the three different ways DNA fragments can be produced.
- Describe the ways that DNA fragments can be amplified - in vivo and in vitro (PCR).
- Describe how DNA is inserted into a vector.
- Describe the transformation of DNA into a host.
- Describe how to identify successful hosts.
- Describe how to grow/clone host cells.
- Describe how labelled DNA probes and DNA hybridisation can be used to locate specific alleles of genes.
- Describe how labelled DNA probes can be used to screen patients for heritable conditions, drug responses or health risks.
- Define VNTR.
- Explain the biological principles that underpin genetic fingerprinting techniques.
- Describe the use of genetic fingerprinting in the fields of forensic science, medical diagnosis, animal and plant breeding.

Skills

Homeostasis

 Interpret information relating to examples of

- Explain the meaning of the key terms: gene; allele; genotype; phenotype; homozygous; and heterozygous.
- Draw labelled genetic diagrams for monohybrid crosses.
- Explain dihybrid inheritance.
- Draw labelled genetic diagrams for dihybrid crosses.
- Explain how codominance affects the inheritance of characteristics.
- Explain how multiple alleles affect inheritance.
- Explain how blood groups in humans are inherited.
- Explain how sex is determined genetically.
- State what is meant by sex-linkage.
- Explain how sex-linked diseases such as haemophilia are inherited.
- Describe autosomal linkage.
- Explain how autosomal linkage affects the combinations of alleles in gametes.
- Explain what is meant by epistasis and explain the effects of epistasis.

Skills

Photosynthesis

- Evaluate data relating to common agricultural practices used to overcome the effect limiting factors of photosynthesis.
- Required practical 7 Use of chromatography to investigate the pigments isolated from leaves of different plants, eg, leaves from shade-tolerant and shade-intolerant plants

 Understand the roles of the sinoatrial node (SAN), atrioventricular node (AVN) and Purkyne tissue in the bundle of His in the control of heart rate.

Populations, evolution and ecosystems

- Define species, population, gene pool and allelic frequency.
- Explain why individuals within a population of a species may show a wide range of variation in phenotype.
- Describe, in detail, the process of natural selection and evolution.
- Describe the effects of stabilising, directional and disruptive selection.
- Describe the process of speciation and how this arises.
- Define allopatric and sympatric speciation.
- Describe genetic drift and its importance in causing changes in allele frequency in small populations.
- Explain how evolutionary change over a long period of time has resulted in a great diversity of species.
- Define community, ecosystem and niche.
- Define biotic and abiotic factors and give examples.
- Define interspecific and intraspecific competition and predation.
- Describe the use of quadrats and the mark-release-recapture method to estimate the size of a population.
- Describe the process of succession.
- Show understanding of the need to manage the conflict between human needs and conservation

- acetylation of associated histones on transcription.
- Explain the relevance of epigenetics on the development and treatment of disease, especially cancer.
- Describe how translation of the mRNA produced from target genes can be inhibited by RNA interference (RNAi).
- Compare malignant and benign tumours.
- Describe the role of tumour suppressor genes and oncogenes.
- Explain how abnormal methylation and increased oestrogen concentrations can cause tumour development.

Skills

Nervous coordination and muscles

Use appropriate units when calculating the maximum frequency of impulse conduction given the refractory period of a neurone.

Gene expression

- Evaluate the use of stem cells in treating human disorders.
- Interpret data provided from investigations into gene expression.
- Evaluate appropriate data for the relative influences of genetic and environmental factors on phenotype.
- Evaluate evidence showing correlations between genetic and environmental factors and various forms of cancer.

- negative and positive feedback.
- Required practical 11 Production of a dilution series of a glucose solution and use of colorimetric techniques to produce a calibration curve with which to identify the concentration of glucose in an unknown 'urine' sample.
- Evaluate the positions of health advisers and the food industry in relation to the increased incidence of type II diabetes.
- Describe the roles of the hypothalamus, posterior pituitary and antidiuretic hormone (ADH) in osmoregulation.
- Describe the structure of the nephron.
- Describe the role of the nephron in urine formation.

DNA technology

- Interpret information relating to the use of recombinant DNA technology.
- Evaluate the ethical, financial and social issues associated with the use and ownership of recombinant DNA technology in agriculture, in industry and in medicine.
- Balance the humanitarian aspects of recombinant DNA technology with the opposition from environmentalists and anti-globalisation activists.
- Evaluate information relating to screening individuals for genetically determined conditions and drug responses.
- Interpret data showing the results of gel

or leaves of different	in order to maintain the	electrophoresis to	
colours.	sustainability of natural	separate DNA fragments.	
- Required practical 8 -	resources.		
Investigation into the			
effect of a named factor	Skills		
on the rate of	Response to stimuli		
dehydrogenase activity in	- Required practical 10:		
extracts of chloroplasts.	Investigation into the		
Respiration	effect of an		
- Required practical 9 -	environmental variable		
Investigation into the	on the movement of an		
effect of a named	animal using either a		
variable on the rate of	choice chamber or a		
respiration of cultures of	maze.		
•	- Use values of heart rate		
single-celled organisms.			
Energy in ecosystems	(R) and stroke volume (V)		
- Calculate the NPP of	to calculate cardiac		
producers given values	output (CO), using the		
for GPP and respiratory	formula CO = R × V		
losses. Derive the	Danulations of the		
appropriate units.	Populations, evolution and		
- Calculate the NPP of	ecosystems		
consumers given values	- Evaluate evidence and		
for energy stored in food,	data concerning issues		
energy lost in urine and	relating to the		
faeces and respiratory	conservation of species		
losses.	and habitats and		
- Calculate the efficiency	consider conflicting		
of energy transfers	evidence.		
within ecosystems.	- Use given data to		
- Calculate percentage	calculate the size of a		
yields.	population estimated		
Inherited change	using the		
- Explain why results of	mark-release-recapture		
genetic crosses often	method.		
differ from predicted	- Required practical 12 -		
results.	Investigation into the		
- Use information to	effect of a named		
represent phenotypic	environmental factor on		
ratios in monohybrid and	the distribution of a		
dihybrid crosses.	given species.		
- Explain what the	- Investigate the		
chi-squared test is.	distribution of organisms		
- Calculate values for	in a named habitat using		
chi-squared and use it to	randomly placed frame		
compare the goodness of fit of observed	quadrats, or a belt		
	transect.		
phenotypic ratios with	- Use both percentage		
expected ratios. - Demonstrate how the	cover and frequency as		
	measures of abundance of a sessile species.		
chi-squared test is used	oi a sessile species.		
in genetics.			
- Calculate the frequency			
of alleles, genotypes and			
phenotypes in a			
population using			
appropriate data and the			

	Hardy Weinberg equation.							
at the end of the unit?	45 minute topic test at the end of each unit (2 per half term), covering AO1, AO3. 15 minute mini tests per fortnight. CPACs assessed during required practicals 7-12. Mock exams at the end of SPR2.							