Meden School Curriculum Planning							
Subject	Biology Triple	Year Group	11	Sequence No.		Торіс	itance, Variation lution

Retrieval	Core Knowledge	Student Thinking
What do teachers need retrieve from students before they start teaching new content ?	What specific ambitious knowledge do teachers need teach students in this sequence of learning?	What real life examples can be applied to this sequence of learning to development of our students thinking, encouraging them to see the inequalities around them and 'do something about them!'
KS3 Learning	<u>L1: DNA</u>	L2: The human genome
Year 7 <u>Reproduction topic</u> Introduced to the structure of egg and sperm cells.	DNA stands for deoxyribonucleic acid , it's the chemical that all of the genetic material in a cell is made up from. DNA determines what inherited characteristic s you have. DNA is found in the nucleus of animal and plant cells in the form of chromosomes , which are normally in pairs. DNA is a polymer . It's made up of two strands coiled together in the shape of a double helix .	Discuss the importance of understanding the human genome. E.g., the search for genes linked to different types of disease
Year 8 Inheritance topic Introduction to gametes, fertilisation within animals. Understanding of characteristics coming from	A gene is a small section of DNA found on a chromosome. Each gene codes for a particular sequence of amino acids which are put together to make a specific protein . Only 20 amino acids are used, but they make 1000s of different proteins. Genes tell us what order to put the amino acids in. DNA determines what protein a cell produces and so what kind of cell it is.	understanding and treatment of inherited disorders use in tracing human migration patterns from the past.
both genetic and environmental. Selective breeding of animals and cloning of	L2: The Genome	L5: Sex determination
plants. Darwin vs Linnaeus Formation of fossils.	Genome is the entire set of genetic material in an organism. Scientists know the whole human genome. It allows scientists to identify genes that are linked to different types of diseases . This can lead to developing effective treatments. Migration patterns can be traced. All modern humans are descended from a common ancestor who lived in Africa. The human genome is mostly identical in	Students could explore the differences between biological sex and gender to link to LGBTQ+
Introduction to pollination and fertilisation in flowering plants.	individuals, but as different populations migrated away from Africa, they gradually developed tiny differences in their genomes. Scientists can work out when populations split off and what route they	<u>L9: Genetic disorders</u> Students should make informed judgements
<u>Year 9</u>	took. L3: Nucleotides & Protein Synthesis	about the economic, social and ethical issues concerning embryo screening, given appropriate information.
Cell structure topic	Each nucleotide consists of one sugar molecule, one phosphate molecule and one base. The sugar	

The organelles within prokaryotic and eukaryotic cells. Examples of eukaryotic and prokaryotic cells. That prokaryotic cells contain a plasmid.	and phosphate form a backbone and alternate. One of four different bases (A , T , C and G) join each sugar. Each base links to a base on the opposite strand in the helix, A with T and C with G. This is called complementary base pairing . Each amino acid is coded for by a sequence of three bases . There are parts of DNA that don't code for proteins. Some of these non-coding parts switch genes on and off , so they control whether the gene if expressed or not .	<u>L23 & 24 – Antibiotics and Antibiotic resistance</u> Students to explore what can be done to stop drug resistant pathogens emerging. Oracy opportunity of factsheets for 'doctors' or 'public' and then present to the class.
	Proteins are made in the cytoplasm at the ribosomes. They use the code in the DNA which cannot move out of the nucleus as it's too big. mRNA is made by copying the code from DNA. It acts as a messenger between the DNA and ribosomes. The correct amino acids are brought to the ribosomes in the correct order by the carrier molecules.	
	When the chain of proteins has been assembled, it folds into a unique shape which allows the protein to perform the task it's meant to do. Enzymes act as biological catalysts. Hormone s carry messages around the body. Structural proteins are physically strong such as collagen that strengthens connective tissue.	
	L4: Mutations	
	A mutation is a random change. They can be inherited and occur continuously . Mutations can also be spontaneous . The chance is increased when exposed to certain substances or some types of radiation . Mutations change the sequence of the DNA bases in a gene which produces a genetic variant and can lead to changes in the protein coded for. Most mutations have very little or no effect on the protein. An altered protein can have a change in shape which alters its function. If the mutation is in the non-coding DNA, it can alter how genes are expressed.	
	Insertions are when a new base is inserted in the DNA base sequence where it shouldn't be. This changes the way the three bases are read which can change the amino acids they code for. They can change more than one amino acid as they have a knock-on effect. Deletions are when a random base is deleted from the DNA base sequence. They change the way the base sequence is read and have a knock-on effect too. Substitutions are when a random base in the DNA base sequence is changed to a different base.	
	L5: Meiosis	
	Meiosis halves the number of chromosomes in gametes and fertilisation restores the full number of chromosomes.	
	Cells in reproductive organs divide by meiosis to form gametes . When a cell divides to form gametes : copies of the genetic information are made, the cell divides twice to form four gametes , each with a single set of chromosomes all gametes are genetically different from each other.	

Gametes join at fertilisation to restore the normal number of chromosomes. The new cell divides by mitosis. The number of cells increases. As the embryo develops cells differentiate.
Ordinary human body cells contain 23 pairs of chromosomes .22 pairs control characteristics only, but one of the pairs carries the genes that determine sex . In females the sex chromosomes are the same (XX) . In males the chromosomes are different (XY) .
L6: Sexual vs Asexual reproduction
Sexual reproduction involves the joining (fusion) of male and female gametes: sperm and egg cells in animals pollen and egg cells in flowering plants.
In sexual reproduction there is mixing of genetic information which leads to variety in the offspring. The formation of gametes involves meiosis .
Asexual reproduction involves only one parent and no fusion of gametes. There is no mixing of genetic information. This leads to genetically identical offspring (clones). Only mitosis is involved.
L7: Advantages and Disadvantages
Advantages of sexual reproduction:
 produces variation in the offspring
• if the environment changes variation gives a survival advantage by natural
selection
 natural selection can be speeded up by humans in selective breeding to increase food production
food production. Advantages of asexual reproduction:
only one parent needed
 more time and energy efficient as do not need to find a mate
 faster than sexual reproduction many identical offspring can be produced when conditions are favourable
Some organisms reproduce by both methods depending on the circumstances.
Malarial parasites reproduce asexually in the human host, but sexually in the mosquito.
Many fungi reproduce asexually by spores but also reproduce sexually to give variation.

Many plants produce seeds sexually, but also reproduce asexually by runners such as strawberry plants, or bulb division such as daffodils.	
L8: Genetic inheritance	
Some characteristics are controlled by a single gene , such as: fur colour in mice; and red-green colour blindness in humans. Each ge ne may have different forms called alleles .	
The allele s present, or genotype , operate at a molecula r level to develop characteristics that can be expressed as a phenotype .	
A dominant allele is always expressed, even if only one copy is present. A recessive allele is only expressed if two copies are present (therefore no dominant allele present).	
If the two alleles present are the same the organism is homozygous for that trait, but if the alleles are different they are heterozygous .	
Most characteristics are a result of multiple genes interacting, rather than a single gene.	
L9: Punnet Squares	
Students should be able to understand the concept of probability in predicting the results of a single gene cross but recall that most phenotype features are the result of multiple genes rather than single gene inheritance .	
Students should be able to use direct proportion and simple ratios to express the outcome of a genetic cross .	
Students should be able to complete a Punnett square diagram and extract and interpret information from genetic crosses and family trees .	
Disorders are inherited. These disorders are caused by the inheritance of certain alleles. Polydactyly (having extra fingers or toes) is caused by a dominant allele.	
Cystic fibrosis (a disorder of cell membranes) is caused by a recessive allele.	
L10: Screening Embryos	
Embryo screening can help to identify if an offspring will have a genetic disorder. There are economic, social and ethical issues concerning embryo screening to consider with embryo screening.	
L11: Work of Mendel	
In the mid-19 th Centaury Mendel noted how characteristics in plants were passed on from one	

generation to the next. The results were published in 1866 and became the foundation of modern genetics. He showed that the height characteristic in pea plants was determined by separately inherited ('hereditary units') passed on from each parent. The ratios of tall and dwarf plants in the offspring showed that the unit for tall plants 'T' was dominant over the unit for dwarf plants 't'. He determined that characteristics in plants are determined by hereditary units. They are passed on to offspring unchanged from both parents, one unit from each parent. They can also be dominant or recessive.
Mendel's work was cutting edge and new. The background knowledge to understand his findings was not available, they had no idea about genes, DNA and chromosomes. It wasn't until after his death that people realized how significant Mendel's work was. His work was the starting point for the understanding of genes today. In the late 1800s, scientists became familiar with chromosomes, they observed how they behaved during cell division. In the early 20 th century, scientists realized there was similarities in the way chromosomes and 'hereditary units' acted. It was proposed that the 'units' were found on chromones and now we call them genes. In 1953 , the structure of DNA was determined so scientists found out how genes work.
L12: Variation
The genome and its interaction with the environment influence the development of the phenotype of an organism .
Differences in the characteristics of individuals in a population are called variation and may be due to differences in: the genes they have inherited (genetic causes)
, the conditions in which they have developed (environmental causes) or a combination of genes and the environment. There is usually extensive genetic variation within a population of a species recall that all variants arise from mutations and that: most have no effect on the phenotype; some influence phenotype; very few determine phenotype.
Mutations occur continuously. Very rarely a mutation will lead to a new phenotype. If the new phenotype is suited to an environmental change it can lead to a relatively rapid change in the species e.g., the peppered moth.
L13 & L14: Evolution
The theory of evolution by natural selection states that all species of living things have evolved from simple life forms that first developed more than three billion years ago. Evolution occurs through

natural selection of variants that give rise to phenotypes best suited to their environment. The theory of evolution by natural selection is now widely accepted.Evidence for Darwin's theory is now available as it has been shown that characteristics are passed on to offspring in genes. There is further evidence in the fossil record and the knowledge of how resistance to antibiotics evolves in bacteria.Extinctions occur when there are no remaining individuals of a species still alive.Students should be able to describe factors which may contribute to the extinction of a species.L15 & L16: Selective Breeding Selective breeding (artificial selection) is the process by which humans breed plants and animals for desired genetic characteristics. Humans have been doing this for thousands of
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animals for desired genetic characteristics. Humans have been doing this for thousands of
years since they first bred food crops from wild plants and domesticated animals.
Selective breeding involves choosing parents with the desired characteristic from a mixed
population. They are bred together. From the offspring those with the desired characteristic are
bred together. This continues over many generations until all the offspring show the desired characteristic .
The characteristic can be chosen for usefulness or appearance:
The characteristic can be chosen for userumess of appearance.
Disease resistance in food crops.
Animals which produce more meat or milk .
Domestic dogs with a gentle nature.
Large or unusual flowers.
Selective breeding can lead to 'inbreeding' where some breeds are particularly prone to disease or inherited defects.
L17 & L18: Genetic Engineering
Genetic engineering is a process which involves modifying the genome of an organism by
introducing a gene from another organism to give a desired characteristic .
Plant crops have been genetically engineered to be resistant to diseases or to produce bigger, better
fruits.
Bacterial cells have been genetically engineered to produce useful substances such as human insulin to treat diabetes.

(HT only) In genetic engineering:	
 Enzymes are used to isolate the required gene; this gene is inserted into a vector, usually a bacterial plasmid or a virus 	
 The vector is used to insert the gene into the required cells 	
 Genes are transferred to the cells of animals, plants or microorganisms at an early stage in their development so that they develop with desired characteristics. 	
The benefits and risks of genetic engineering in agriculture and in medicine and that some people have objections.	
In genetic engineering , genes from the chromosomes of humans and other organisms can be 'cut out' and transferred to cells of other organisms .	
Crops that have had their genes modified in this way are called genetically modified (GM) crops. GM crops include ones that are resistant to insect attack or to herbicides . GM crops generally show increased yields .	
Concerns about GM crops include the effect on populations of wild flowers and insects . Some people feel the effects of eating GM crop s on human healt h have not been fully explored.	
Modern medical research is exploring the possibility of genetic modification to overcome some inherited disorders .	
 <u>L19: Plant Cloning</u> Tissue culture: using small groups of cells from part of a plant to grow identical new plants. This is important for preserving rare plant species or commercially in nurseries. Cuttings: an older, but simple, method used by gardeners to produce many identical new plants from a parent plant. 	
L20: Animal Cloning Embryo transplants : splitting apart cells from a developing animal embryo before they become specialized, then transplanting the identical embryos into host mothers.	
Adult cell cloning:	

 The nucleus is removed from an unfertilized egg cell. The nucleus from an adult body cell, such as a skin cell, is inserted into the egg cell. An electric shock stimulates the egg cell to divide to form an embryo. These embryo cells contain the same genetic information as the adult skin cell. 	
When the embryo has developed into a ball of cells, it is inserted into the womb of an adult female to continue its development.	
L21: Fossils	
Fossils are the ' remains ' of organisms from millions of years ago, which are found in rocks. Many early forms of life were soft-bodied , which means that they have left few traces behind. What traces there were have been mainly destroyed by geological activity . Therefore, scientists cannot be certain about how life began on Earth. Phylogenic trees can be used to show how different species have evolved over time and who their closest/most distant ancestor species were.	
Fossils may be formed:	
 from parts of organisms that have not decayed because one or more of the conditions needed for decay are absent. 	
 when parts of the organism are replaced by minerals as they decay. as preserved traces of organisms, such as footprints, burrows and rootlet traces. 	
L22: Speciation	
A species is a group of similar organisms that can reproduce to give fertile offspring. Speciation is the development of a new species. It occurs when populations of the same species become so different, they can no longer successfully interbreed to produce fertile offspring. Isolation is where populations of a species are separated. This can happen due to a physical barrier. Conditions on the other side will be slightly different so different characteristics will become more popular in the populations due to natural selection. Alfred Russel Wallace was working at the same time as Charles Darwin, he was one of the early scientists working on speciation (this and warning colours is what he is most famous for). His observations contributed greatly to our understanding today which has developed as more evidence has come available over time. Wallace independently came up with the idea of natural selection and published work on the subject with Darwin in 1858 wish prompted Darwin to publish 'On the Origin of Species' in 1859.	

L23 & 24: Antibiotics and Antibiotic Resistance

Antibiotics, such as **penicillin**, are medicines that help to cure **bacterial disease** by killing **infective bacteria** inside the body. It is important that specific **bacteria** should be treated by specific **antibiotics**.

Bacteria can evolve rapidly because they reproduce at a fast rate.

Mutations of **bacterial pathogens** produce new **strains**. Some **strains** might be **resistant** to **antibiotics**, and so are not killed. They **survive** and **reproduce**, so the population of the **resistant strain rises**. The **resistant strain** will then spread because people are not **immune** to it and there is no effective treatment.

MRSA is resistant to antibiotics.

To reduce the rate of development of **antibiotic resistant** strains:

Doctors should not prescribe **antibiotics** inappropriately, such as treating non-serious or **viral infections.**

Patients should complete their course of **antibiotics** so all **bacteria** are killed and none **survive to mutate** and form **resistant strains.**

The agricultural use of antibiotics should be restricted.

The **development** of new **antibiotics** is costly and slow. It is unlikely to keep up with the emergence of new **resistant** strains

L25 & L26: Classification

In the **1700s Carl Linnaeus** classified organisms into groups according to their characteristics and structures. This was known as the **Linnaean System**. Organism are first divided into **kingdoms**, then subdivided into smaller groups: **phylum**, **class**, **order**, **family**, **genus and species**.

As knowledge of **biochemical processes** taking place inside organisms developed **and microscopes improved** scientists put forward new models of classification. In **1990 Carl Woese** proposed the **threedomain system.** Using RNA analysis technique he found, in some cases, that species thought to be closely related were not. Before kingdom the three domains are **archaea** (primitive bacteria found in extreme places) **bacteria** (true bacteria, they look similar to archaea but there are a lot of biochemical differences between them) and **eukaryota** (broad range of organisms such as fungi, plants, animals and protists).

In the binomial system , every organism has a two-part Latin name . The first name is the genus and the second name is the species . This is used worldwide so scientists from different countries or who speak different languages all refer to a particular species with the same name, avoiding confusion.	
Evolutionary trees show how scientists think species are related to each other. They show common ancestors and relationships between different species. The more recent the common ancestor the more closely related the two species and the more characteristics they are likely to share. Scientists analyze lots of different types of data to work out evolutionary relationships. For currently living organisms they use current classification data (e.g. DNA analysis) and for extinct species they use fossil records .	
L27: Revision	
L28: EoTT	
L29: GPA	