A level Biology Personal Learning Checklist

<u>Paper 1</u> = Topics 1, 2, 3, 4, 5 and 6

Paper 2 = Topics 1, 2, 3, 4, 7 and 8

<u>Paper 3</u> = Practicals and scientific article

Topic 1 Lifestyle, Health and Risk

Objective	Confidence (R/A/G)		
	1	2	3
1.1 Understand why many animals have a heart and circulation (mass			
transport to overcome limitations of diffusion in meeting the			
requirements of organisms).			
1.2 Understand the importance of water as a solvent in transport,			
including its dipole nature.			
1.3 Understand how the structures of blood vessels (capillaries, arteries			
and veins) relate to their functions.			
1.4 i) Know the cardiac cycle (atrial systole, ventricular systole and cardiac			
diastole) and relate the structure and operation of the mammalian heart,			
including the major blood vessels, to its function.			
ii) Know how the relationship between heart structure and function can			
be investigated practically.			
1.5 Understand the course of events that leads to atherosclerosis			
(endothelial dysfunction, inflammatory response, plaque formation,			
raised blood pressure).			
1.6 Understand the blood-clotting process (thromboplastin release,			
conversion of prothrombin to thrombin and fibrinogen to fibrin) and its			
role in cardiovascular disease (CVD).			
1.7 Know how factors such as genetics, diet, age, gender, high blood			
pressure, smoking and inactivity increase the risk of cardiovascular			
disease (CVD).			
1.8 Be able to analyse and interpret quantitative data on illness and			_
mortality rates to determine health risks, including distinguishing			
between correlation and causation and recognising conflicting evidence.			

risk factors, including sample selection and sample size used to collect data that is both valid and reliable. 1.10 Understand why people's perceptions of risks are often different from the actual risks, including underestimating and overestimating the risks due to diet and other lifestyle factors in the development of heart disease. 1.11 i) Be able to analyse data on energy budgets and diet. ii) Understand the consequences of energy imbalance, including weight loss, weight gain, and development of obesity. 1.12 i) Know the difference between monosaccharides, disaccharides and polysaccharides, including glycogen and starch (amylose and amylopectin). ii) Be able to relate the structures of monosaccharides, disaccharides and polysaccharides to their roles in providing and storing energy (β-glucose and cellulose are not required in this topic). 1.13 Know how monosaccharides join to form disaccharides (sucrose, lactose and maltose) and polysaccharides (glycogen and amylose) through condensation reactions forming glycosidic bonds, and how these can be split through hydrolysis reactions. 1.14 i) Know how a triglyceride is synthesised by the formation of ester bonds during condensation reactions between glycerol and three fatty acids. ii) Know the differences between saturated and unsaturated lipids. 1.15 i) Be able to analyse and interpret data on the possible significance for health of blood cholesterol levels and levels of high-density lipoproteins (HDLs) and low-density lipoproteins (LDLs). ii) Know the evidence for a causal relationship between blood cholesterol levels (total cholesterol and LDL cholesterol) and cardiovascular disease (CVD). 1.16 Understand how people use scientific knowledge about the effects of diet, including obesity indicators, body mass index and waist-to-hip ratio, exercise and smoking to reduce their risk of coronary heart disease. CORE PRACTICAL 2: Investigate the effect of caffeine on heart rate in Daphnia. 1.17 Be able discuss the potential ethical issues regarding the use of i	1.9 Be able to evaluate the design of studies used to determine health		
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	CORE PRACTICAL 2:		
1.18 Know the benefits and risks of treatments for cardiovascular disease	Investigate the vitamin C content of food and drink.		
	1.18 Know the benefits and risks of treatments for cardiovascular disease		

Topic 2 Genes and Health

Objective	Confidence (R/A/G)		e
	1	2	3
2.1 i) Know the properties of gas exchange surfaces in living organisms			
(large surface area to volume ratio, thickness of surface, difference in			
concentration).			
ii) Understand how the rate of diffusion is dependent on these properties			
and can be calculated using Fick's Law of Diffusion.			
iii) Understand how the structure of the mammalian lung is adapted for			
rapid gaseous exchange.			
2.2 i) Know the structure and properties of cell membranes.			
ii) Understand how models such as the fluid mosaic model of cell			
membranes are interpretations of data used to develop scientific			
explanations of the structure and properties of cell membranes.			
CORE PRACTICAL 3:			
Investigate membrane structure, including the effect of alcohol			
concentration or temperature on membrane permeability.			
2.3 Understand what is meant by osmosis in terms of the movement of			
free water molecules through a partially permeable membrane			
(consideration of water potential is not required).			
2.4 i) Understand what is meant by passive transport (diffusion, facilitated			
diffusion), active transport (including the role of ATP as an immediate			
source of energy), endocytosis and exocytosis.			
ii) Understand the involvement of carrier and channel proteins in			
membrane transport.			
2.5 i) Know the basic structure of mononucleotides (deoxyribose or ribose			
linked to a phosphate and a base, including thymine, uracil, cytosine,			
adenine or guanine) and the structures of DNA and RNA (polynucleotides			
composed of mononucleotides linked through condensation reactions).			
ii) Know how complementary base pairing and the hydrogen bonding			
between two complementary strands are involved in the formation of the			
DNA double helix.			
2.6 i) Understand the process of protein synthesis (transcription)			
including the role of RNA polymerase, translation, messenger RNA,			
transfer RNA, ribosomes and the role of start and stop codons.			
ii) Understand the roles of the DNA template (antisense) strand in			
transcription, codons on messenger RNA and anticodons on transfer RNA.			

overlapping and degenerate). 2.8 Know that a gene is a sequence of bases on a DNA molecule that codes for a sequence of amino acids in a polypeptide chain. 2.9 i) Know the basic structure of an amino acid (structures of specific amino acids are not required). ii) Understand the formation of polypeptides and proteins (amino acid monomers linked by peptide bonds in condensation reactions). iii) Understand the significance of a protein's primary structure in determining its three-dimensional structure and properties (globular and fibrous proteins and the types of bonds involved in its three-dimensional structure). iv) Know the molecular structure of a globular protein and a fibrous protein and understand how their structures relate to their functions (including haemoglobin and collagen). 2.10 i) Understand the mechanism of action and the specificity of enzymes in terms of their three-dimensional structure. ii) Understand that enzymes are biological catalysts that reduce activation energy. iii) Know that there are intracellular enzymes catalysing reactions inside cells and extracellular enzymes produced by cells catalysing reactions outside of cells. CORE PRACTICAL 4: Investigate the effect of enzyme and substrate concentrations on the initial rates of reactions. 2.11 i) Understand the process of DNA replication, including the role of DNA polymerase. ii) Understand how Meselson and Stahl's classic experiment provided new data that supported the accepted theory of replication of DNA and refuted competing theories. 2.12 i) Understand how errors in DNA replication can give rise to mutations. ii) Understand how errors in DNA replication one of a number of possible gene mutations. ii) Understand how cystic fibrosis results from one of a number of possible gene mutations.	2.7 Understand the nature of the genetic code (triplet code, non-		
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2.14 Understand how the expression of a gene mutation in people with		
cystic fibrosis impairs the functioning of the gaseous exchange, digestive		
and reproductive systems.		
2.15 i) Understand the uses of genetic screening, including the		
identification of carriers, pre-implantation genetic diagnosis (PGD) and		
prenatal testing, including amniocentesis and chorionic villus sampling.		
ii) Understand the implications of prenatal genetic screening.		
2.16 Be able to identify and discuss the social and ethical issues related to		
genetic screening from a range of ethical viewpoints.		

Topic 3 Voice of the Genome

Objective	Confidence (R/A/G)		:e
	1	2	3
3.1 Know that all living organisms are made of cells, sharing some			
common features.			
3.2 Know the ultrastructure of eukaryotic cells, including nucleus,			
nucleolus, ribosomes, rough and smooth endoplasmic reticulum,			
mitochondria, centrioles, lysosomes, and Golgi apparatus.			
3.3 Understand the role of the rough endoplasmic reticulum (rER) and the			
Golgi apparatus in protein transport within cells, including their role in			
the formation of extracellular enzymes.			
3.4 Know the ultrastructure of prokaryotic cells, including cell wall,			
capsule, plasmid, flagellum, pili, ribosomes, mesosomes and circular DNA.			
3.5 Be able to recognise the organelles in 3.2 from electron microscope			
(EM) images.			l
3.6 Understand how mammalian gametes are specialised for their			
functions (including the acrosome in sperm and the zona pellucida in the			İ
egg).			1
3.7 Know the process of fertilisation in mammals, including the acrosome			
reaction, the cortical reaction and the fusion of nuclei.			
3.8 i) Know that a locus (plural = loci) is the location of genes on a			
chromosome.			1
ii) Understand the linkage of genes on a chromosome and sex linkage.			1
3.9 Understand the role of meiosis in ensuring genetic variation through			
the production of non-identical gametes as a consequence of			İ
independent			
assortment of chromosomes and crossing over of alleles between			
chromatids (details of the stages of meiosis are not required).			

3.10 Understand the role of mitosis and the cell cycle in producing		
identical daughter cells for growth and asexual reproduction.		
CORE PRACTICAL 5:		
Prepare and stain a root tip squash to observe the stages of mitosis.		
3.11 i) Understand what is meant by the terms 'stem cell, pluripotency		
and totipotency'.		
ii) Be able to discuss the way society uses scientific knowledge to make		
decisions about the use of stem cells in medical therapies.		
3.12 Understand how cells become specialised through differential gene		
expression, producing active mRNA leading to synthesis of proteins,		
which in turn control cell processes or determine cell structure in animals		
and plants, including the lac operon.		
3.13 Understand how the cells of multicellular organisms are organised		
into tissues, tissues into organs and organs into systems.		
3.14 i) Understand how phenotype is the result of an interaction between		
genotype and the environment.		
ii) Know how epigenetic changes, including DNA methylation and histone		
modification, can modify the activation of certain genes.		
iii) Understand how epigenetic changes can be passed on following cell		
division.		
3.15 Understand how some phenotypes are affected by multiple alleles		
for the same gene at many loci (polygenic inheritance) as well as the		
environment and how this can give rise to phenotypes that show		
continuous variation.		

Topic 4 Biodiversity and Natural Resources

Objective	Confidence (R/A/G)		ce
	1	2	3
4.1 Know that over time the variety of life has become extensive but is			
now being threatened by human activity.			
4.2 i) Understand the terms biodiversity and endemism.			
ii) Know how biodiversity can be measured within a habitat using species			
richness and within a species using genetic diversity by calculating the			
heterozygosity index (H)			
iii) Understand how biodiversity can be compared in different habitats			
using a formula to calculate an index of diversity (D)			

4.3 Understand the concept of niche and be able to discuss examples of	
adaptation of organisms to their environment (behavioural, physiological	
and anatomical).	
4.4 Understand how natural selection can lead to adaptation and	
evolution.	
4.5 i) Understand how the Hardy-Weinberg equation can be used to see	
whether a change in allele frequency is occurring in a population over	
time.	
ii) Understand that reproductive isolation can lead to accumulation of	
different genetic information in populations, potentially leading to the	
formation of new species.	
4.6 i) Understand that classification is a means of organising the variety of	
life based on relationships between organisms using differences and	
similarities in phenotypes and in genotypes, and is built around the	
species concept.	
ii) Understand the process and importance of critical evaluation of new	
data by the scientific community, which leads to new taxonomic	
groupings, including the three domains of life based on molecular	
phylogeny, which are Bacteria, Archaea, Eukaryota.	
4.7 Know the ultrastructure of plant cells (cell walls, chloroplasts,	
amyloplasts, vacuole, tonoplast, plasmodesmata, pits and middle lamella)	
and be able to compare it with animal cells.	
4.8 Be able to recognise the organelles in 4.7 from electron microscope	
(EM) images.	
4.9 Understand the structure and function of the polysaccharides starch	
and cellulose, including the role of hydrogen bonds between β-glucose	
molecules in the formation of cellulose microfibrils.	
4.10 Understand how the arrangement of cellulose microfibrils and	
secondary thickening in plant cell walls contributes to the physical	
properties of xylem vessels and sclerenchyma fibres in plant fibres that	
can be exploited by humans.	
CORE PRACTICAL 6:	
Identify sclerenchyma fibres, phloem sieve tubes and xylem vessels and	
their location within stems through a light microscope.	
4.11 Know the similarities and differences between the structures,	
position in the stem and function of sclerenchyma fibres (support), xylem	
vessels (support and transport of water and mineral ions) and phloem	
(translocation of organic solutes).	
4.12 Understand the importance of water and inorganic ions (nitrate,	
calcium ions and magnesium ions) to plants.	
CORE PRACTICAL 7:	

Investigate plant mineral deficiencies.		
CORE PRACTICAL 8:		
Determine the tensile strength of plant fibres.		
4.13 Understand the development of drug testing from historic to		
contemporary protocols, including William Withering's digitalis soup,		
double blind trials, placebo, three-phased testing.		
4.14 Understand the conditions required for bacterial growth.		
CORE PRACTICAL 9:		
Investigate the antimicrobial properties of plants, including aseptic		
techniques for the safe handling of bacteria.		
4.15 Understand how the uses of plant fibres and starch may contribute		
to sustainability, including plant-based products to replace oil-based		
plastics.		
4.16 Be able to evaluate the methods used by zoos and seed banks in the		
conservation of endangered species and their genetic diversity, including		
scientific research, captive breeding programmes, reintroduction		
programmes and education.		

Topic 5 On the Wild Side

Objective	Confidence (R/A/G)		ce
	1	2	3
5.1 Understand the terms ecosystem, community, population and			
habitat.			
5.2 Understand that the numbers and distribution of organisms in a			
habitat are controlled by biotic and abiotic factors.			
5.3 Understand how the concept of niche accounts for distribution and			
abundance of organisms in a habitat.			
CORE PRACTICAL 10:			
Carry out a study on the ecology of a habitat, such as using quadrats and			
transects to determine distribution and abundance of organisms, and			
measuring abiotic factors appropriate to the habitat.			
5.4 Understand the stages of succession from colonisation to a climax			
community.			
5.5 Understand the overall reaction of photosynthesis as requiring energy			
from light to split apart the strong bonds in water molecules, storing the			
hydrogen in a fuel (glucose) by combining it with carbon dioxide and			
releasing oxygen into the atmosphere.			

5.6 Understand how phosphorylation of ADP requires energy and that		
hydrolysis of ATP provides an immediate supply of energy for biological		
processes.		
5.7 Understand the light-dependent reactions of photosynthesis including		
how light energy is trapped by exciting electrons in chlorophyll and the		
role of these electrons in generating ATP, reducing NADP in		
photophosphorylation and producing oxygen through photolysis of water.		
5.8 i) Understand the light-independent reactions as reduction of carbon		
dioxide using the products of the light-dependent reactions (carbon		
fixation in the Calvin cycle, the role of GP, GALP, RuBP and RUBISCO).		
ii) Know that the products are simple sugars that are used by plants,		
animals and other organisms in respiration and the synthesis of new		
biological molecules (polysaccharides, amino acids, lipids and nucleic		
acids).		
CORE PRACTICAL 11:		
Investigate photosynthesis using isolated chloroplasts (the Hill reaction).		
5.9 Understand the structure of chloroplasts in relation to their role in		
photosynthesis.		
5.10 i) Be able to calculate net primary productivity.		
ii) Understand the relationship between gross primary productivity, net		
primary productivity and plant respiration.		
5.11 Know how to calculate the efficiency of biomass and energy		
transfers between trophic levels.		
5.12 Understand the different types of evidence for climate change and		
its causes (including records of carbon dioxide levels, temperature		
records, pollen in peat bogs and dendrochronology), recognising		
correlations and causal relationships.		
5.13 Understand the causes of anthropogenic climate change, including		
the role of greenhouse gases (carbon dioxide and methane) in the		
greenhouse effect.		
5.14 i) Understand that data can be extrapolated to make predictions and		
that these are used in models of future climate change.		
ii) Understand that models for climate change have limitations.		
5.15 Understand the effects of climate change (changing rainfall patterns		
and changes in seasonal cycles) on plants and animals (distribution of		
species, development and life cycles).		
5.16 Understand the effect of temperature on the rate of enzyme activity		
and its impact on plants, animals and microorganisms.		
5.17 Understand how evolution (a change in the allele frequency) can		
come about through gene mutation and natural selection.		
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5.18 Understand the role of the scientific community (scientific journals,		
the peer review process, scientific conferences) in validating new		
evidence, including proteomics and genomics, that supports the accepted		
scientific theory of evolution.		
5.19 Understand how isolation reduces gene flow between populations,		
leading to allopatric or sympatric speciation.		
CORE PRACTICAL 12:		
Investigate the effect of temperature on the initial rate of an enzyme-		
catalysed reaction, to include Q10.		
CORE PRACTICAL 13:		
Investigate the effects of temperature on the development of organisms		
(such as seedling growth rate, brine shrimp hatch rates).		
5.20 Understand the way in which scientific conclusions about		
controversial issues, such as what actions should be taken to reduce		
climate change or the degree to which humans are affecting climate		
change, can sometimes depend on who is reaching the conclusions.		
5.21 Understand how knowledge of the carbon cycle can be applied to		
methods to reduce atmospheric levels of carbon dioxide.		
5.22 Understand how reforestation and the use of sustainable resources,		
including biofuels, are examples of the effective management of the		
conflict between human needs and conservation.		

Topic 6 Immunity, Infection and Forensics

Objective	Co	nfidenc	e
	((R/A/G)	
	1	2	3
6.1 Understand how to determine the time of death of a mammal by			
examining the extent of decomposition, stage of succession, forensic			
entomology, body temperature and degree of muscle contraction.			
6.2 Know the role of micro-organisms in the decomposition of organic			
matter and the recycling of carbon.			
6.3 Know how DNA profiling is used for identification and determining			
genetic relationships between organisms (plants and animals).			
6.4 Know how DNA can be amplified using the polymerase chain reaction			
(PCR).			
CORE PRACTICAL 14:			
Use gel electrophoresis to separate DNA fragments of different length.			
6.5 Be able to compare the structure of bacteria and viruses.			
6.6 Understand how Mycobacterium tuberculosis (TB) and Human			

Immunodeficiency Virus (HIV) infect human cells, causing a sequence of		
symptoms that may result in death.		
6.7 Understand the non-specific responses of the body to infection,		
including inflammation, lysozyme action, interferon, and phagocytosis.		
6.8 Understand the roles of antigens and antibodies in the body's		
immune response including the involvement of plasma cells,		
macrophages and antigen-presenting cells.		
6.9 Understand the differences between the roles of B cells (B memory		
and B effector cells) and T cells (T helper, T killer and T memory cells) in		
the body's immune response.		
6.10 Understand how one gene can give rise to more than one protein		
through posttranscriptional changes to messenger RNA (mRNA).		
6.11 i) Know the major routes pathogens may take when entering the		
body.		
ii) Understand the role of barriers in protecting the body from infection,		
including skin, stomach acid, and gut and skin flora.		
6.12 Understand how individuals may develop immunity (natural,		
artificial, active, passive).		
6.13 Understand how the theory of an 'evolutionary race' between		
pathogens and their hosts is supported by the evasion mechanisms		
shown by pathogens.		
6.14 Understand the difference between bacteriostatic and bactericidal		
antibiotics.		
CORE PRACTICAL 15:		
Investigate the effect of different antibiotics on bacteria.		
6.15 Know how an understanding of the contributory causes of hospital		
acquired infections have led to codes of practice regarding antibiotic		
prescription and hospital practice that relate to infection prevention and		
control.		
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Topic 7 Run for your Life

Objective	Confidence (R/A/G)		ce
	1	2	3
7.1 Know the way in which muscles, tendons, the skeleton and ligaments			
interact to enable movement, including antagonistic muscle pairs,			
extensors and flexors.			

the sliding filament theory, including the role of actin, myosin, troponin, tropomyosin, calcium ions (Ca2+), ATP and ATPase. 7.3 i) Understand the overall reaction of aerobic respiration as splitting of the respiratory substrate, to release carbon dioxide as a waste product and reuniting of hydrogen with atmospheric oxygen with the release of a large amount of energy. ii) Understand that respiration is a many-stepped process with each step controlled and catalysed by a specific intracellular enzyme. 7.4 Understand the roles of glycolysis in aerobic and anaerobic respiration, including the phosphorylation of hexoses, the production of ATP, reduced coenzyme, pyruvate and lactate (details of intermediate stages and compounds are not required). 7.5 Understand the role of the link reaction and the Krebs cycle in the complete oxidation of glucose and formation of carbon dioxide (CO2), ATP, reduced NAD and reduced FAD (names of other compounds are not required) and why these steps take place in the mitochondria, unlike glycolysis which occurs in the cytoplasm. 7.6 Understand how ATP is synthesised by oxidative phosphorylation associated with the electron transport chain in mitochondria, including the role of chemiosmosis and ATP synthase. 7.7 Understand what happens to lactate after a period of anaerobic respiration in animals. CORE PRACTICAL 16: Investigate rate of respiration. 7.8 i) Know the myogenic nature of cardiac muscle. ii) Understand how the normal electrical activity of the heart coordinates the heart beat, including the roles of the sinoatrial node (SAN), the atrioventricular node (AVN), the bundle of His and the Purkyne fibres. iii) Understand how the use of electrocardiograms (ECGs) can aid the diagnosis of cardiovascular disease (CVD) and other heart conditions. 7.9 i) Be able to calculate cardiac output. iii) Understand how variations in ventilation and cardiac output enable rapid delivery of oxygen to tissues and the removal of carbon dioxide	7.2 Understand the process of contraction of skeletal muscle in terms of		
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from them, including how the heart rate and ventilation rate are	from them, including how the heart rate and ventilation rate are		
controlled and the roles of the cardiovascular control centre and the	controlled and the roles of the cardiovascular control centre and the		
ventilation centre in the medulla oblongata.	ventilation centre in the medulla oblongata.		
CORE PRACTICAL 17:	CORE PRACTICAL 17:		
Investigate the effects of exercise on tidal volume, breathing rate,	Investigate the effects of exercise on tidal volume, breathing rate,		
respiratory minute ventilation and oxygen consumption using data from	respiratory minute ventilation and oxygen consumption using data from		
spirometer traces.	spirometer traces.		

7.10 i) Know the structure of a muscle fibre.		
ii) Understand the structural and physiological differences between fast		
and slow twitch muscle fibres.		
7.11 i) Understand what is meant by negative feedback and positive		
feedback control.		
ii) Understand the principle of negative feedback in maintaining systems		
within narrow limits.		
7.12 Understand homeostasis and its importance in maintaining the body		
in a state of dynamic equilibrium during exercise, including the role of the		
hypothalamus and the mechanisms of thermoregulation.		
7.13 Understand the analysis and interpretation of data relating to		
possible disadvantages of exercising too much (wear and tear on joints,		
suppression of the immune system) and exercising too little (increased		
risk of obesity, cardiovascular disease (CVD) and diabetes), recognising		
correlation and causal relationships.		
7.14 Understand how medical technology, including the use of keyhole		
surgery and prostheses, is enabling those with injuries and disabilities to		
participate in sports.		
7.15 Be able to discuss different ethical positions relating to whether the		
use of performance-enhancing substances by athletes is acceptable.		
7.16 Understand how genes can be switched on and off by DNA		
transcription factors including hormones.		

Topic 8 Grey Matter

Objective	Confidence		ce
		(R/A/G)	
	1	2	3
8.1 Know the structure and function of sensory, relay and motor			
neurones including the role of Schwann cells and myelination.			
8.2 i) Understand how the nervous systems of organisms can cause			
effectors to respond to a stimulus.			
ii) Understand how the pupil dilates and contracts.			
8.3 Understand how a nerve impulse (action potential) is conducted along			
an axon including changes in membrane permeability to sodium and			
potassium ions and the role of the myelination in saltatory conduction.			
8.4 Know the structure and function of synapses in nerve impulse			
transmission, including the role of neurotransmitters, including			
acetylcholine.			

8.5 Understand how the nervous systems of organisms can detect stimuli with reference to rods in the retina of mammals, the roles of rhodopsin, opsin, retinal, sodium ions, cation channels and hyperpolarisation of rod cells in forming action potentials in the optic neurones. 8.6 Understand how phytochrome and IAA bring about responses in plants to environmental cues, including their effects on transcription. 8.7 Understand how co-ordination is brought about through nervous and hormonal control in animals. 8.8 Know the location and functions of the cerebral hemispheres, hypothalamus, cerebellum and medulla oblongata in the human brain. 8.9 Understand how magnetic resonance imaging (MRI), positron emission tomography (PET) and computed tomography (CT) scans are used in medical diagnosis and the investigation of brain structure and function. 8.10 Understand what happens during the critical period so that mammals can develop their visual capacities to the full. 8.11 Understand the role animal models have played in the research into human brain development and function, including Hubel and Wiesel's experiments with monkeys and kittens. 8.12 Be able to discuss moral and ethical issues relating to the use of animals in medical research from two ethical standpoints. 8.13 Understand how animals, including humans, can learn by habituation. CORE PRACTICAL 18: Investigate habituation to a stimulus. 8.14 Understand how imbalances in certain, naturally occurring brain chemicals can contribute to ill health, including dopamine in Parkinson's disease and serotonin in depression, and to the development of new drugs. 8.15 Understand the effects of drugs on synaptic transmissions, including the use of L-Dopa in the treatment of Parkinson's disease and the action of MDMA in Ecstasy. 8.15 Understand how the outcomes of genome sequencing projects are being used in the development of personalised medicine and the social, moral and ethical issues this raises. 8.17 Know how drugs can be produced using genetically modified			
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abilities of new-born babies, animal experiments, studies of individuals		
with damaged brain areas, twin studies and cross-cultural studies.		

Key skills

Objec	pjective		onfidend (R/A/G)	ce
		1	2	3
ation	Express numbers in decimal form.			
Comput	Express numbers in standard form.			
Arithmetic and Numerical Computation	Use ratios, fractions and percentages.			
tic and N	Make estimates of the results of simple calculations.			
Arithme	Use calculators to find and use power, exponential and logarithmic functions.			
	Use an appropriate number of significant figures/			
o o	Calculate the mean.			
Handling Data	Understand the terms mean, mode and median.			
ndlin	Make order of magnitude calculations.			
На	Select and use a statistical test.			
	Understand measures of dispersion, including standard deviation and range.			
_	Understand and use the symbols: =, <, <<, >>, >, α , $^{\sim}$			
Algebra	Change the subject of an equation.			
₹	Substitute numerical values into equations using appropriate units.			
	Understand that $y = mx + c$ represents a linear relationship.			
	Plot a line graph from experimental data, including drawing a line of best fit.			
Draw a tangent to a curve of the rate of change.	Determine the gradient and intercept of a linear graph.			
	Draw a tangent to a curve and calculate its gradient as a measure of the rate of change.			
	Identify a correlation from a graph.			
Ge	Visualise and represent 2D and 3D forms.			

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	Calculate areas of triangles and rectangles.			
	Calculate surface areas and volumes of cubes.			
lysis	Describe overall trends and significant changes from data in tables and graphs.			
Data analysis	Manipulate the data to emphasise trends observed.			
Data	Draw inferences and conclusions by linking the data to your own biological knowledge.			
	Identify the meanings of the key words of an exam question.			
	Select the relevant information given in a question to answer it.			
Literacy	Answer extended questions strategically.			
Lit	Link the relevant subject knowledge to answer questions on text such as the article in paper 3.			
	Identify the range of the independent variable required.			
	Identify the equipment required and what will be recorded for the dependent variable.			
	Identify the controlled variables and how they will be kept the same/monitored to ensure validity.			
s	Identify how to make the experiment repeatable.			
Practicals	Identify any ethical issues.			
	Identify all risks and precautions.			
	Apply your knowledge of the method to new contexts and scenarios.			
	Apply your knowledge of the practical method to comment on			
	and improve the validity and repeatability of a similar method. Evaluate results and draw conclusions with reference to measurement uncertainties and errors.			