## **BIOLOGICAL PSYCHOLOY**

## Week 1

Somatotopic mapping - mapping of spatial information in the body e.g. rats whiskers body → brain

Structure begets function

Brain has special functions/structures to protect it from its environment and damage: Medically induced comas = protection of the nervous system, enables brain recovery

Ventricles: fluid filled spaces (filled with cerebral spinal fluid). Protection

Choroid plexus: barrier tissue of the CFS from the blood Keep the brains environment protected and highly controlled

The blood brain barrier: helps to control the environment of the brain

Endothelial cells of capillaries are very tightly packed (more tightly packed than the rest of the body).

- The space between them allows nutrients etc. to pass through. The tight junctions prevent many molecules from passing across.
- Poses as a problem in drug discovery
- Some larger molecules e.g. glucose can get through due to active transport
- Bacteria is too big to get through (most of them)

Neurons are so tightly packed and their axons/dendrites so intertwined that looking at unprepared neural tissue reveals almost nothing about them

- Golgi stain: stains each neuron entirely black, making it possible to see individual neurons for the first time only in silhouette)
  - O No indication of no. neurons in an area or the nature of their inner structure
- <u>Nissel stain:</u> uses dye that binds to cell bodies, thus can be used to estimate the number of cell bodies in an area
- <u>Electron microscopy:</u> provides details of neuronal structure. Images are so detailed that it can be difficult to visualise the general aspects of neuroanatomical structure

<u>Anterograde neuroanatomical tracing methods:</u> traces the path of axons projecting <u>away</u> from cell bodies located in a particular area

Retrograde tracing methods: traces the path of axons projecting into a particular area

## Directions in the vertebrate nervous system

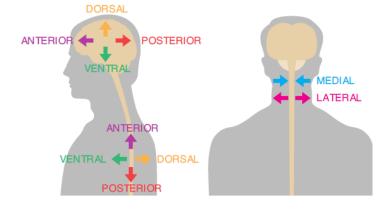
Directions described in relation to the orientation of the spinal cord

<u>Anterior (rostral):</u> towards the nose end. Sounds like 'nostril'

Posterior (caudal): towards the tail end

<u>Dorsal:</u> towards the surface of the back/top of the head. *Superior* in primates

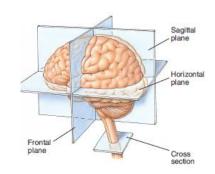
Ventral: towards the surface of the chest/bottom of



#### the head

<u>Medial:</u> towards the midline of the body <u>Lateral:</u> away from the midline toward the body's lateral surfaces

Frontal planes: vertical cuts – parallel to the face
Saggital sections: cut parallel to the side of the brain



## Spinal cord

<u>Gray matter:</u> composed mainly of cell bodies and unmyelinated interneurons White matter: myelinated axons

Pairs of spinal nerves attach to the spinal cord one on the left and one on the right side; axons join to the cord via the dorsal root or the ventral root

<u>Dorsal root axons:</u> sensory (afferent) unipolar neurons, cell bodies group to form the dorsal root ganglia

Ventral root axons: motor (efferent) multipolar neurons with their cell bodies in ventral horns

- Those that are part of the somatic nervous system: project to skeletal muscles
- Those that are part of the autonomic nervous system: project to ganglia, where they synapse onto neurons that in turn project to internal organs

## Five major divisions of the brain

Developing brain in the embryo:

Fluid filled tube  $\rightarrow$  3 swellings occur at the anterior end of this tube  $\rightarrow$  become the adult forebrain, midbrain and hind brain Before birth:

Forebrain swelling grows into  $\Rightarrow$  2 swellings Hindbrain  $\Rightarrow$  2 swellings

From anterior to posterior the 5 swelling are:

- Telencephalon (cerebral hemispheres)
- Diencephalon
- Mesencephalon

Brain stem

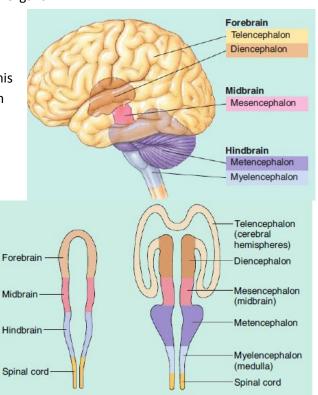
Metencephalon

Myelencephalon (medulla)

# Major structure of the brain

Myelencephalon (medulla) (hind brain):

- Most posterior
- Composed largely of tracts carrying signals between the rest of the brain and the body
- Contains the Reticular formation/RAS: role in arousal, also in sleep, attention, movement



# Developmental psychology/ Developmental Science Week 7

Focuses on research and theory in cognitive, personality and social developmental processes as well as physical changes throughout the lifespan (infancy, childhood, adolescence and adulthood)

<u>Lifespan development:</u> examines patterns of growth, change and stability in behaviour that occur throughout the entire life span

Looks at physical, cognitive, social and personality development

## What affects development?

- *History-graded influences* aka Cohort effects: biological and environmental influences associated with a particular historical movement e.g. WWII, 9/11 attack
- Age-graded influences: biological and environmental influences that are similar for individuals in a particular age group, regardless of when or where they were raised
- Socio-cultural graded influences
- *Non-normative:* specific, a typical events that do not occur in most people's lives e.g. death of a 6 year olds parents

### Pitfalls:

Typical vs atypical Static vs dynamic Nature vs nurture

*Nativist approach:* our DNA alone guides the development of cognitive and social processes and is innately driven

• Cyril Burt - argued intelligence was inherited. Faked his results

*Empiricist approach*: it's through our specific interactions with the environment that we develop cognitive and social processes

- John Locke everyone born with a tabula rasa 'blank slate'
- Watson behaviourism. Albert and the rat trained to fear
  - o 'Albert' had a brain condition hence was not a 'typically developing' infant

Core goal of twin research: is to estimate the heritability of traits - what percentage of variance in the population is due to genes and what's due to the environment e.g. schizophrenia, bipolar

- Environmental constraints hinder conclusions from twin research same environment can never produce twins that are truly identical
- Cyril Burt argued intelligence was inherited. Faked his results

<u>Epigenetics:</u> heritable changes in gene expression that are not caused by changes in DNA sequence but by positive or negative early life experiences

- Our experiences can create a genetic memory without changing the sequence of DNA inherited from out parents. This genetic memory can help build optimal brain development
  - Early environmental experiences e.g. a mothers care or lack of it, can induce chemical changes on the DNA...altering gene expression without changing the underlying DNA sequence
- Attempts to encapsulate the impact of both genes and environmental factors and their combined role in development

## Continuity vs. Discontinuity

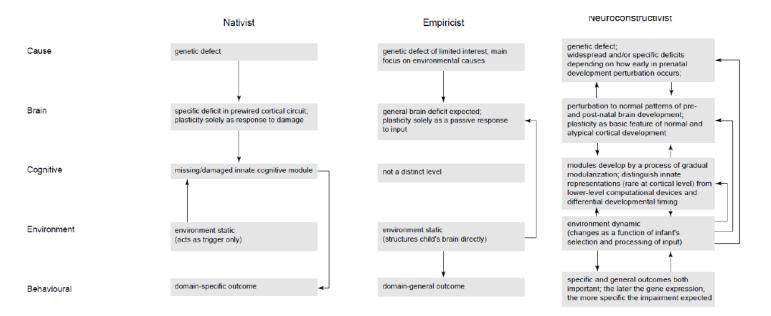
<u>Continuous change:</u> gradual development, achievements at one level building on those of previous levels. Quantitative. Matter of degree, not of kind

<u>Discontinuous change:</u> change occurs in distinct stages. Behaviour and processes are qualitatively different at different stages

<u>Modularity:</u> mind is composed of independent, closed, domain-specific processing modules.

Neuroconstructicism: developed by Annette Karmiloff-Smith

- Looks at the interplay of gene, cognition and environment.
- Not dynamic but static
- Modules develop as a process of graduate modularisation
- Infant brain is highly interconnected and there is gradual modularization across development
- Modules are not innately specified in the infant brain



**Piaget:** development occurs in discrete stages in which knowledge progresses through a progressive sequence (schemas) that adapt and change as the infant becomes more advanced in their motor capabilities

- skills in one stage need to be acquired before moving onto the next stage
- Argued that intellectual development (knowledge) governed all other aspects of development (social, personality, motor etc.)