

nerve system structure & function

skull & spinal cord

- skull: protect the brain from measurable amount of force.
- spinal cord: protected by the spinal column
 - ↳ individual components are vertebrae.
 - ↳ sc and brain are connected.
- CSF flows between the arachnoid and pial layers.

Layers of Meninges:

(outer to inner layer):

- ① Dura
- ② Arachnoid
- ③ Pia

↳ made up of collagen fibers (elastic).

cerebrospinal fluid

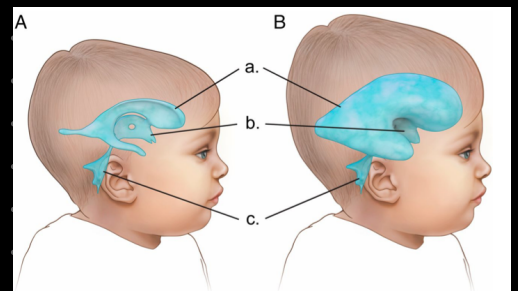
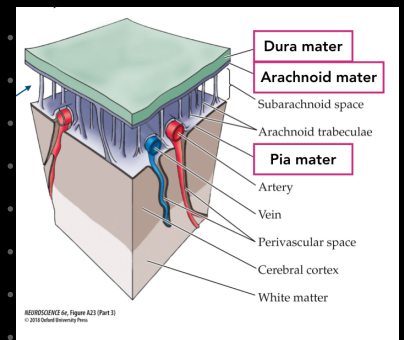
- CSF: balanced ionic concentration
- CSF movement is via the pulsation of the blood vessels.
 - drainage into the sagittal sinus to join back into the bloodstream.
- rate of flow: under circadian rhythm
 - ↳ fastest when asleep.
- The CSF = made via the ependymal cells in the choroid plexus of the ventricles.
- CSF production in mice = increased under anaesthetic & decreased under AD gene mice.

FUNCTIONS

- ① cushion the brain from dmg
- ② maintains homeostatic environm.
- ③ provide neurotrophic factor for the growth of axons.
- ④ waste removal.

WHAT HAPPENS WHEN CSF BUILDS UP?

- in children: hydrocephalus → the skull bones are not fused yet, so the brain starts to swell.
 - neurological deficits.
- in adults: the skull bones are all fused together, so the ICP would increase.
 - resolved with trephination (hole in the brain)



NEURON DOCTRINE

neuron doctrine: original argument

= the neurons are **reticulum** or **discrete cells**.

observations that it's **reticulum**:

- golgi stain = only 5-10% of cells were stained.
- revealed **neurites** (chainy processes)
- the neurites were growing out of a cell body.

observations that it's **discrete**:

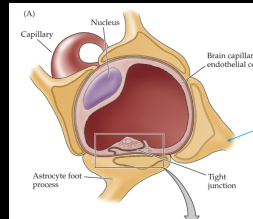
- the sub-cellular processes: fibril structures were discontinuous
- proven by **EM**: the neuron's neurites were able to touch but the membranes were non-continuous.

the longer you wait after the axonal death, the longer the axon would grow back.

Neurons have to have:

- ① **Transmission of information** → the movement of information from the dendrite of the neuron → the axon.
- ② The neurons have to be able to move information from one neuron to another:
passing information

BLOOD BRAIN BARRIER



BBB = made via: the glial cell processes → the **astrocyte** not process wrap around the **blood vessel**.

Astroglia are the support cells that are contributing to the maintenance of the BBB.

↳ **locally controls the blood vessel diameter.**

molecular transport: **transporter proteins** are required to move the **glc** across the membrane.

↳ use **ATP** to move essential molecules.

Drugs to penetrate the BBB: **hydrophilic** to move thru blood.

* need specialized transport proteins for each molecule & **hydrophobic** to move thru membrane.

Neuron Doctrine

- ① The basis of the nervous system: the neuron.
- ② The neurons are distinct and non continuous.

DIRECTIONAL FLOW

directional flow was determined via the dendrite and axon.

- the dendrites in the sensory neurons are always farthest from CNS, and the axons are closest into the CNS.

Synapses

info: transmitted from one cell → next via synapse.

- 2 main kinds of synapse: **electrical** and **chemical synapse**.

Chemical Synapse

- no specialized structure to move chemical from one neuron \rightarrow next.
 - \hookrightarrow the NT are released into the synapse (the gap) and diffuse into the next neuron.

Electrical currents are metabolically more efficient.

Electrical Synapse

- specialized junction for the movement of electrical signals from one neuron \rightarrow next.
 - \hookrightarrow the electrical signals move directly from the prev to the next neuron.
- current flows directly into the next neuron.

Experiment: Frog Hearts

signal modification

(signals can be changed plastically).

signal amplification

Advantage of chemical synapse

computation: can

invert the signal from $+$ \rightarrow $-ve$.

the HR was measured by Heart tension (\downarrow T \rightarrow \downarrow HR when chemical used).



1. stimulate donor heart to decrease HR via CNIO.



2. ECF w/ chemicals from donor heart: given to the recipient and the recipient heart also had \downarrow HR.

- \rightarrow the electrical stimulation resulted in chemical changes that altered the HR on another heart w/o the vagus nerve.
- \rightarrow chemical stim could work independently of the electrical stimulation.

neuron diversity

3 main kinds of neurons

sensory neuron

- brings in info from outside of the CNS

interneuron

- connects the sensory MN.

motor neuron

- brings the info out into the periphery.

4 kinds of neuronal diversity

① **Structural**: number + arrangement of neurites.

② **Connectivity**: convergent/diverge

③ **Chemical**: diff NT for diff neurons

④ **Electrical**: diff AP frequencies.

neuronal classification:

- modern method: RNA sequencing, via separating the neurons and testing their RNA sequences.
- previous categories of neurons can be split into 27 individual neurons b/c of the RNA sequencing.

glial cells

- glial cells are non-excitabile cells that support the neurons.

\hookrightarrow glial cell: neuron ratio $\sim 1:1$

Maintaining polarity:
packing, sorting, mvmt of the proteins

site and transport

Problems w/ diversity

the use of metabolic energy: very high.
- need to make / rebuild NT.

switching from chemical / electrical signals fast and reliably.

Neurons overcome this problem via the use of support cells.

- Astrocytes**
- most abundant cell type involved:
 - EC concentration [ions]
 - growth regulation
 - homeostatic regulation
 - neurotrophic factors

- Myelinating cell**
- myelinating cell: cells that make the myelin (which makes outgrowths & wraps around axons).

↳ oligodendrocyte CNS, Schwann cell PNS.



- ependymal cells**
- involved in secretion of fluid that becomes CSF
 - permissive form of BBB.
 - in the choroid plexus of the ventricles.

- Microglia**
- macrophage descendants
 - involved in: inflammation, neuroplasticity of axon growth, scavenging.

Nervous system organization

Complete circuit: one sensory and one motor neuron.

When you have more neurons in the circuit → the neurons are more specialized.

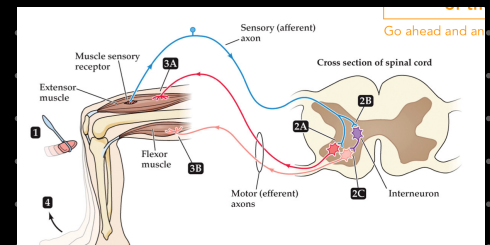
Signal integration: occurs @ the soma, not the axon.

↳ the signals are generated @ axon but not integrated.

vertebrate reflex arc:

ipsilateral flexion, inhibit extension.

Stimulus (on sensory neuron) → IN → the same sensory neuron can diverge: IN can send signal across midline of SC. → contralateral extension, inhibit flexion.



Hub & Spoke NS

- CNS / PNS distinction: CNS & PNS (size, cell #).
- CNS → PNS connection, not PNS - PNS.
- unidirectional info flow
- specialized neurons
- ↳ distinct afferent/efferent/ IN.

Neural Net NS.

- no CNS / PNS distinction.
- no central ganglia: info just flows everywhere.
- bidirectional information flow.
- no specialization of cell/neuron function.
- ↳ efferent/afferent are all in one neuron.

AP VS. PSP

- AP: non decaying, travels @ one magnitude (binary signal) → electrical mechanism of signal transduction.
- ↳ detectable via the extracellular recording.
- PSP: decaying, travels @ graded potentials → current due to the NT activating receptor.
- ↳ detected only via intracellular recording.

Types of PSP:

- ① **Excitatory PSP** → they cause the cell to be more likely to fire.
- ② **Inhibitory PSP** → cause cell to be less likely to fire.
- ③ **Modulatory** → they alter the synapses around the cell, making it altered.
↳ not distinctly excitatory/inhibitory.

Recurrent inhibition / Excitation



Recurrent inhibition

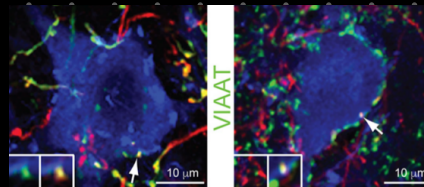
(-ve feedback, activation means turning off the neuron that initiated it)



Recurrent excitation

(+ve feedback, the activation of the cell continuously activates itself)

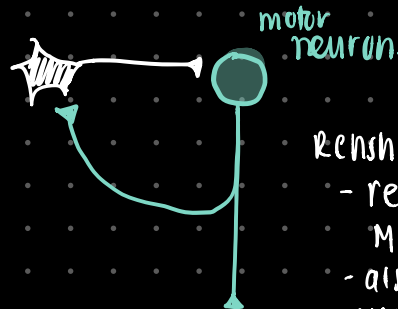
↓
need external source to turn off



Renshaw cell

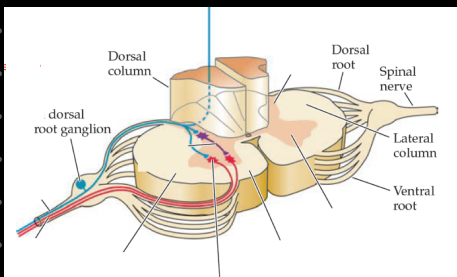
determined by staining for protein:

- a) **calbindin** → exclusive to the Renshaw cell in SC.
- b) **AChR** → bIC R cell get input from the motor axons.



Renshaw cell: inhibitory.
- receives collateral from the MN
- also has one extension of axon back to MN.

spinal cord



- white matter: myelinated axons.
- Gray Matter: **neuropil**, soma.

↓
axon terminal, dendrite, synapse.

WM > GM in cervical SC: b/c there's more axons going to from brain to diff regions.

↳ basis of localization of Fcn.

SC = mapped **topographically**: mapped by fcn.

Somatotopy: mapping of the diff body parts along vertical axis of SC.

- **Distal muscle**: innervated by **lateral GM**.
- **Proximal muscle**: innervated by **medial GM**.

Exp: changing sensory - Motor neuron connection.

muscle in ventral horn of GM = matched to the soma via injecting dye
↳ the injection of dye from terminal → goes to soma.

deletion of FoxP1 gene: MN oriented diff.

↓
loss of fine motor scale; movements are nonsmooth.
mapping of MN.

↓
more distal muscle = more ventral soma.

ANS

- sympa/parasympa NS.
- uses 2 neurons to get to organ → CNS uses 1.
 - still have 1 sensory 1 motor.
 - ↳ pre/post-ganglionic neurons.
- innervates the visceral organs.
- sympa & parasympa (time): b/c the syma
 - = used to get the body ready for fight/flight.
 - ↓
 - doesn't actually innervate the muscles to move them.
- metabolic balance of the 2 maintains the metabolic functioning of the body.

ENS

- can act independently of the CNS.
- - mainly involved in the innerv. of the GI.



the parasympa would synapse on the ENS neurons (they're found in plexuses of GI) rather than the non-neuronal GI cells.