COGS 17 WEEK 3 WINTER 2024, A04

ANNOUNCEMENT

- Midterm On Jan 25, 2024 (Tomorrow) 3:30 4:50 pm
- 24 Questions, most of them require multiple responses
- 80 Minutes to complete
- One attempt
- You can revisit and change answers

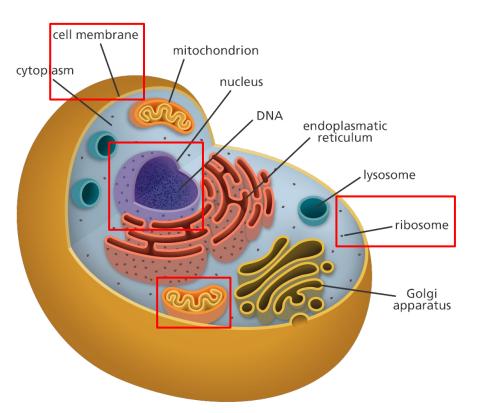
PROBLEM SET FOR REVIEW

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ZWQYAjhzGUhepsGjQWM/edit?usp=sharing



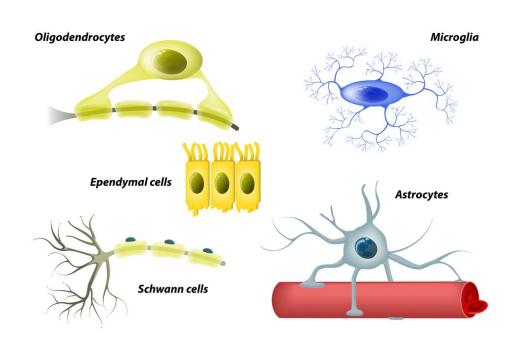
CELLS

CELLS



- Nucleus -- where DNA stored
- Ribosome -- where protein synthesis occurs
- Mitochondria -- Produce ATP, the "powerhouse" of the cell
- Cell membrane -- lipid,
 BILAYER, semi permeable
 membrane

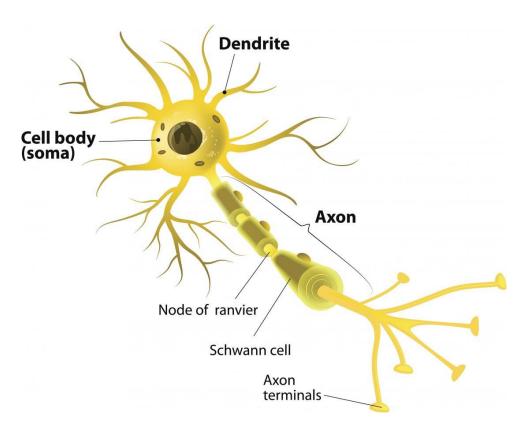
GLIAL CELLS



- Non-neural cells of the nervous system with multiple functions
- Do NOT participate in information transfer
- Much smaller than neurons, but much more numerous
- ~50% of brain by weight

NEURONS

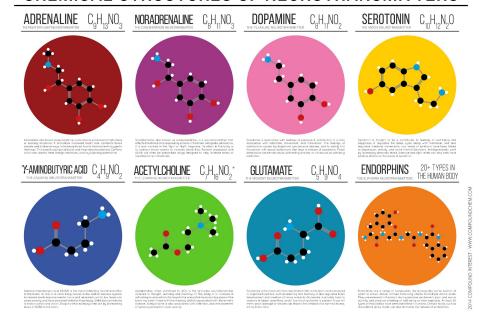
- Cells that are specialized for INFORMATION TRANSFER
- Modified processes (i.e. dendrites & axon)
- Dendrites -- reception of **INCOMING** message via receptors
- Axon -- Site of release of OUTGOING message
- Modified membrane -ion channels



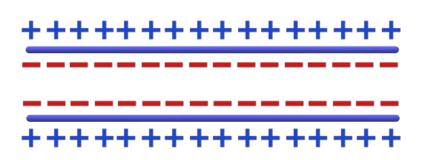
NEUROTRANSMITTERS (NT)

- endogenous chemicals that allow neurons to communicate with each other throughout the body
- E.g. Acetylcholine (ACh),
 GABA, Glutamate, Serotonin
 (5-HT), Norepinephrine,
 etc.
- May have different effects in different parts of body

CHEMICAL STRUCTURES OF NEUROTRANSMITTERS



RESTING POTENTIAL

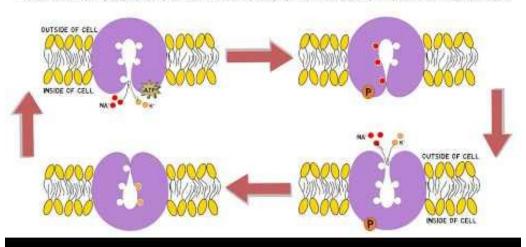


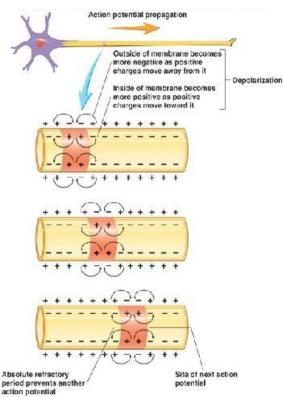
- Generally <u>-70 mV</u>
- More negative inside & more positive outside
- Established in part by Na-K
 pump, which actively transport
 3 Na+ out and 2 K+ in
- Highly POLARIZED -- ready to "fire"

SODIUM-POTASSIUM PUMP

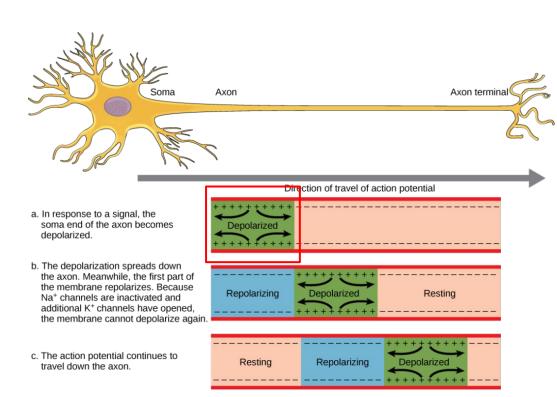
SODIUM-POTASSIUM PUMP CYCLE

-USES ATP TO TRANSPORT SODIUM AND POTASSIUM IONS ACROSS THE CELL MEMBRANE

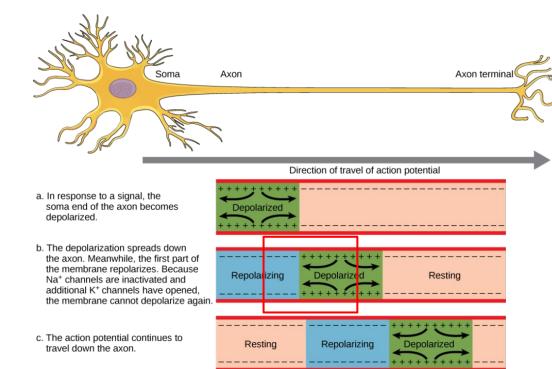




- DEPOLARIZATION of neuron
- Triggered by neurotransmitters from other neuron, electrical stimulation, or other stimuli
- Starting at Axon Hillock



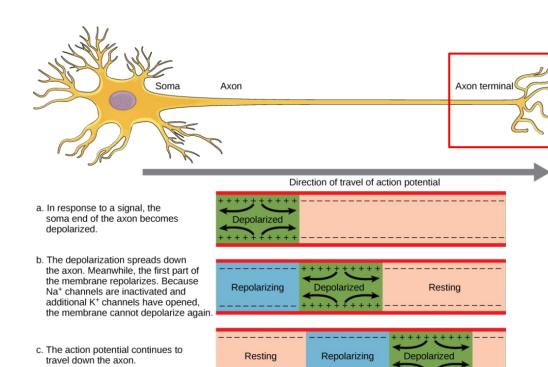
- ●Voltage-activated Na+ gates open -- Na+ in
- •Reverse local polarization to ~ +50 mV
- Na+ influx causes
 adjacent
 Na+ voltage-activated
 gates
 to open & previously
 opened
 gates close
- Propagation of the action potential to the terminal



As previous Na+ gates close, local K+ gates open --K+

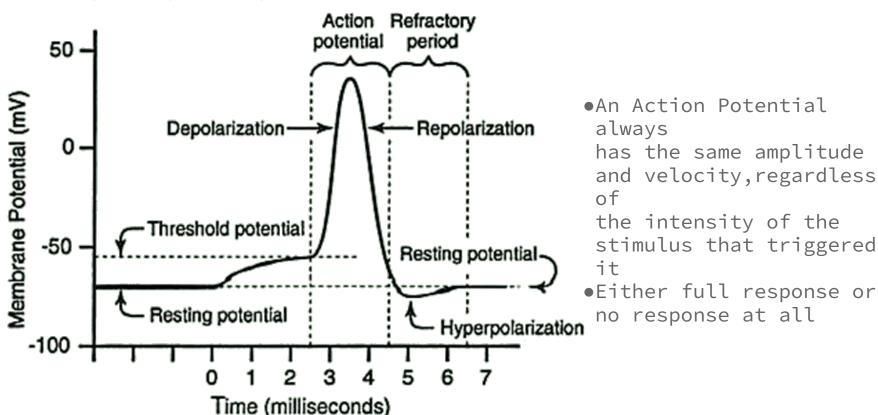
leaves the cell

- Because of opened K+ channels & closed Na+ channels, the membrane cannot depolarize again
- Refractory period -prevents backflow of action potential



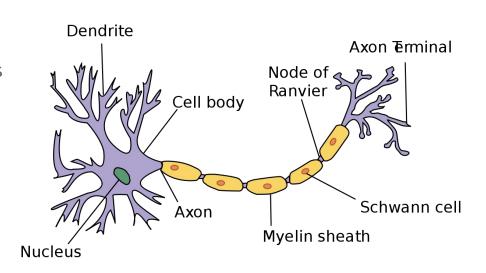
- When reaches terminal,
 Ca2+ enters cell &
 Neurotransmitter
 released
- When outside becomes more positive again,
 K+ channels close
- Na-K pumps restore resting potential to -70 mV
- Ca pump reject Ca2+ from terminal
- Pumps require energy

ALL-OR-NONE LAW

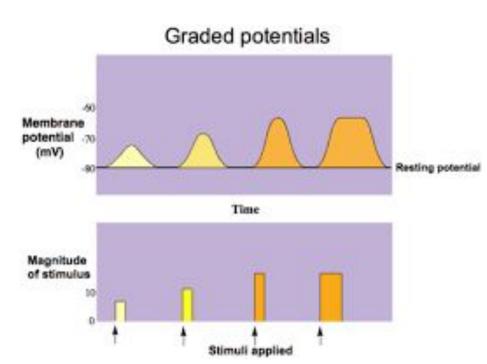


MYELINATION AND SALTATORY CONDUCTION

- Increase the speed of propagation
- Glia cells form insulating sheaths around axons, with small gaps in between sheaths
- Electrical conduction -fast, but degrades as it moves
- Node of Ranvier -- ionic conduction, re-boost the signal to original strength and pass it to next shealth
- Saltatory Conduction "Jumping" of the signal from node to node along axon

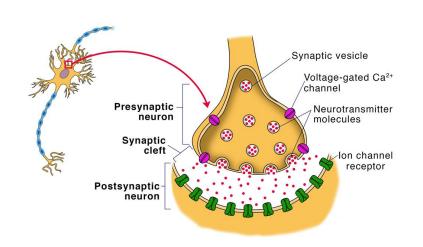


GRADED POTENTIALS



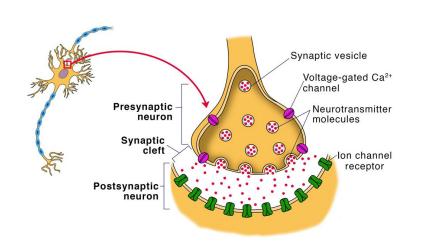
- Releasing NT from a Neuron does
 NOT always requires an Action
 Potential
- •Can vary in amplitude
- •i.e. Strong response due to strong stimulus; weak response due to weak stimulus
- Examples: some receptor cells (e.g. retina), lateral inhibitors, local Neurons, etc.

THE SYNAPSE



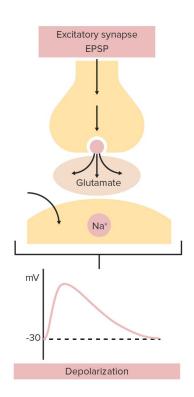
- Presynaptic cell releases NT into synaptic cleft, affects postsynaptic cell via <u>Exocytosis</u> (release of vesicles of NT).
- When membrane depolarization reaches the axon terminal, voltage-gated Ca++ channels open, causes influx of Ca++

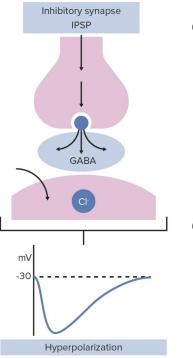
THE SYNAPSE



- NT <u>passively</u> diffuses across synaptic cleft
- NT binds to <u>NT-specific</u> receptor site on (usually) dendrites of postsynaptic cell
- Postsynaptic cell may fire its NT, etc.
- NT soon <u>detaches</u> from receptor
- NT is often then <u>deactivated</u> by enzymes or Glia

EPSP VS. IPSP

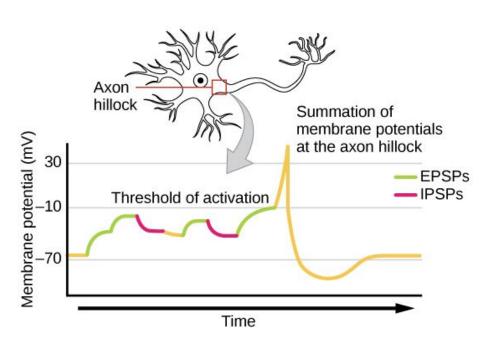




EPSP: <u>Excitatory</u> Postsynaptic
 Potential – postsynaptic
 cells become hypo-polarized.

IPSP: <u>Inhibitory</u> Postsynaptic
Potential - postsynaptic
cells become hyper-polarized.

SUMMATION



- The response of a given neuron is nearly always the product of a <u>summation of EPSPs and/or</u> <u>IPSPs</u>
 - Two types of summation:

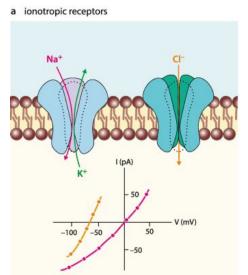
 <u>Spatial</u> summation and
 <u>Temporal</u> summation

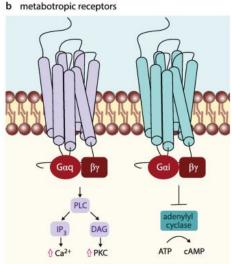
IONOTROPIC VS. METABOTROPIC

- •Ionotropic: NT <u>directly affects</u> <u>ion gates</u>
- -Effects are <u>rapid</u>, <u>short-lived</u>

 Metabotropic: Triggers metabolic changes in Postsynaptic cell

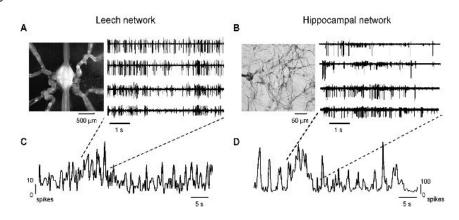
 Activates/triggers production of <u>Secondary Messenger</u>, which binds w/<u>G-Protein</u> to open separate ion gate, requires energy, <u>slower and long-lasting</u>





SPONTANEOUS ACTIVITY

- •Some Neurons show spontaneous Firing, <u>in absence of incoming NT</u>
- •Usually graded potentials
- •Converging NTs increase or decrease such cells' activity, modifying how much NT they will then release



AGONISM VS. ANTAGONISM

 Agonist: chemicals that increase effect of a NT

 Antagonist: chemicals that decrease effect of a NT

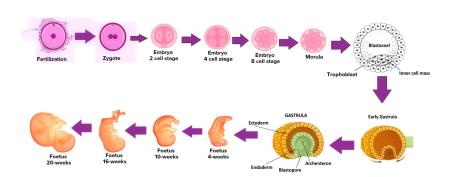
Agonists and Antagonists

Agonists - Drugs that occupy receptors and activate them.

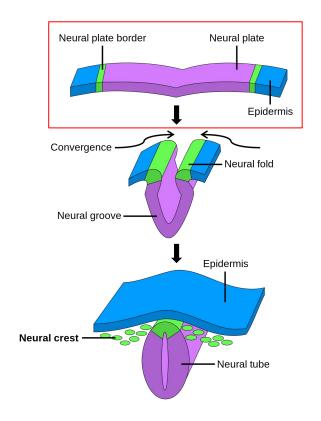
Antagonists - Drugs that occupy receptors but do not activate them Antagonists block receptor activation by agonists.



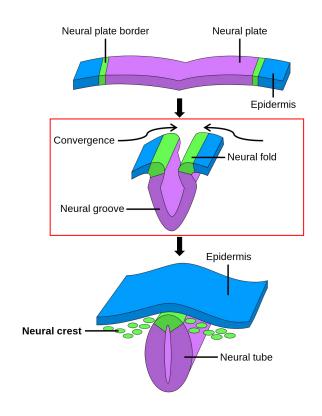
DEVELOPMENT



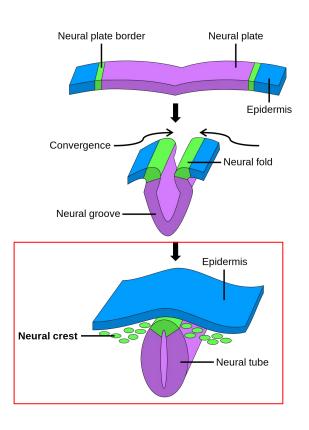
 A new embryo develops three cell layers: Ectoderm (outer layer, becomes nervous system & skin), Mesoderm (middle layer, becomes bones, muscles, blood vessels), endoderm (inner layer, becomes organs, glands)



- Over first 2 weeks, embryo changes from a sphere of cells to an elongated "worm", still 3 layered
- Then dorsal Ectoderm begins to thicken and forms hard Neural Plate



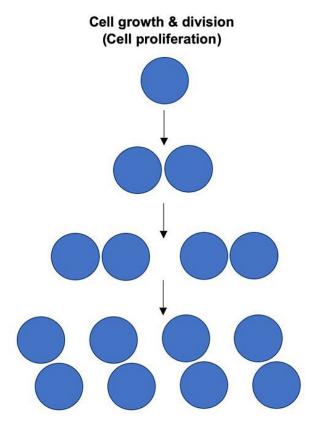
 Edges of plate form ridges (Neural Folds) that curl toward each other along a longitudinal line



- By week 4, edges of Neural Folds have fused, forming Neural Tube lined with Ectoderm, embedded in Mesoderm
- Spina Bifida Neural fold failed to fuse
- Rostral end of Neural Tube >> Brain
- Caudal end >> Spinal Cord
- Surface of ridges (Neural Crest) >> Ganglia of ANS & Peripheral Neurons & Glia

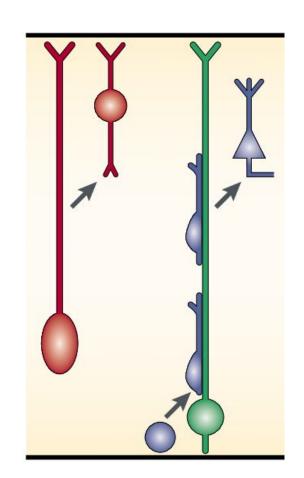
PROLIFERATION

- •Growth of new cells
- •Stem Cells: Ectodermal cells that line the inside of the Neural Tube ogive rise to neurons first through
 - Symmetrical Division
- o∼Week 7, shift to <u>Asymmetrical Division</u>, producing one stem cell + one neuron
- •Stem cells stay to divide, neurons start to migrate

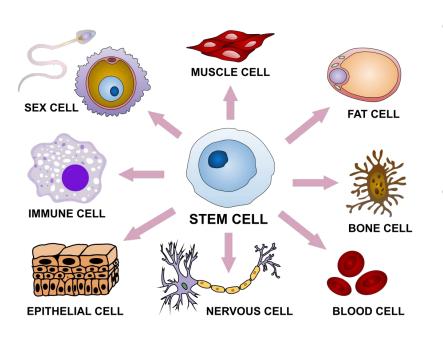


MIGRATION

- Some Neurons migrate by "crawling" along Radial Glia fibers, often aided by Glycoproteins
- Other Neurons may migrate by following chemical trails laid down by Glia Cells or by other Neurons



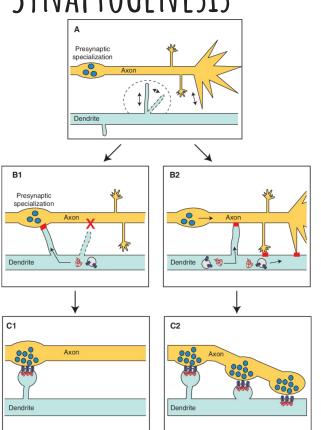
DIFFERENTIATION



•Transition of a cell from one cell type to another

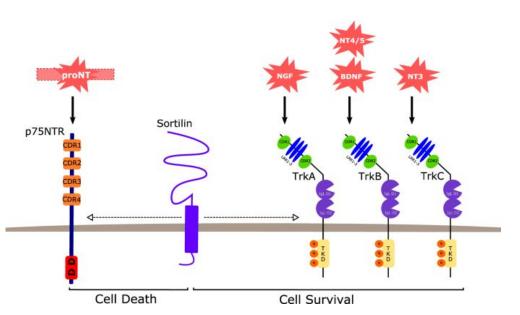
 Per Cell-Autonomous (genetic) and Induction (chemical influences from local environments) factors

SYNAPTOGENESIS



- Developing junctions (Synapses) between cells
- After migration, Neurons grow Axon first and Dendrites later
- Growth Cone at end of elongating Axon has many <u>Filopodia</u> that detect surrounding <u>chemical gradients</u>
- Some Axons are directed by Guidepost Cells
- Others depend on Chemical Trails produced by Glia cells or other migrating Neurons/Axons

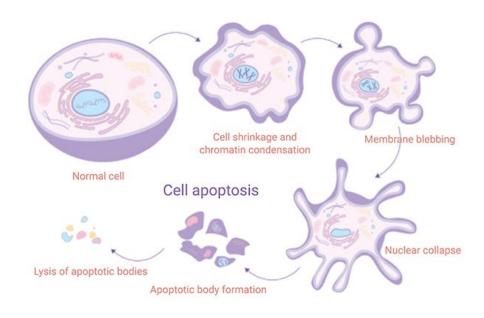
NEUROTROPHINS



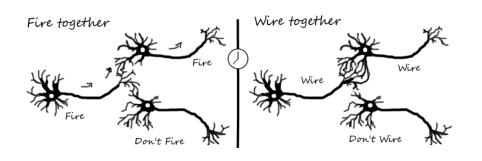
 chemicals that attract/repel and promote survival and activity of Neurons

APOPTOSIS (PROGRAMMED CELL DEATH)

- Triggered by suicide genes
- As cells compete for connections, those who do not have connections die off
- Post-development, most remaining Stem Cells also die by the activation of their suicide genes

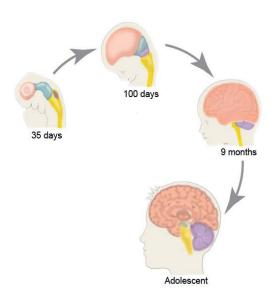


CELLS THAT FIRE TOGETHER, WIRE TOGETHER



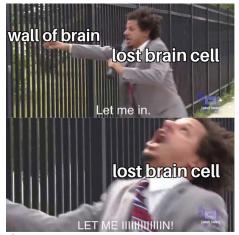
- Patterns of <u>co-activity</u>
 often determine outcome of
 competition
- Adjacent Presynaptic cells tend to correlate their bursts of activity, so tend to develop connections to adjacent Targets

FURTHER DEVELOPMENT

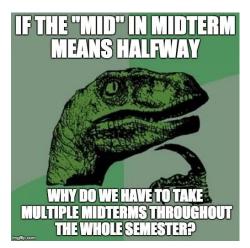


- Brain Growth
- Dendritic branching
- Further Synaptogenesis
- Myelination in neurons





GOOD LUCK ON YOUR MIDTERM!





QUESTIONS?

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Office Hours: Mon 5-6 pm
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To get the section slides:
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https://github.com/JasonC1217/COGS17_A04_Wi24

OR:

