

# *COGS 17 section A02*

neural development



section resources repo  
week 3 guiding questions  
inside folder



## reminders

- midterm exam 1 is **tomorrow** (4/21) during class time 3:30 PM - 4:50 PM
  - open book, online, one shot
  - covers material from week 1 - 3
- homework 3 is due **tonight** 11:59 PM!



# exam game plan

## before the exam

- do the practice homework
  - make sure you don't just memorize the answers and actually understand the concepts
- study all resources provided in the canvas modules
- read the textbook if you have extra time
  - not required just if you would like more detailed concepts
  - i have a pdf of it if you need lmk

## D-0

- please do not pull an all nighter
- pace yourself!
- double check your answers
  - but try to get it right the first time bc you might not have time to revisit them
- have your study notes ready with you



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# embryonic development

3 layers of cells

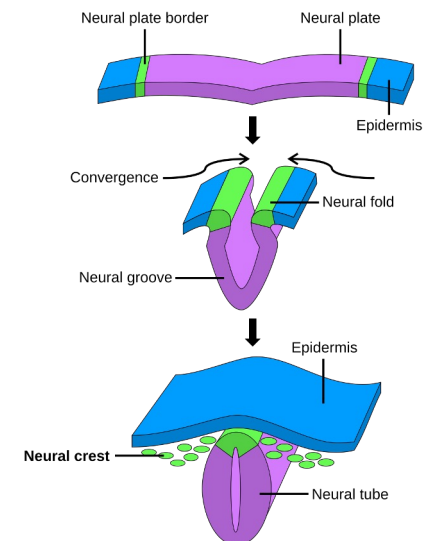
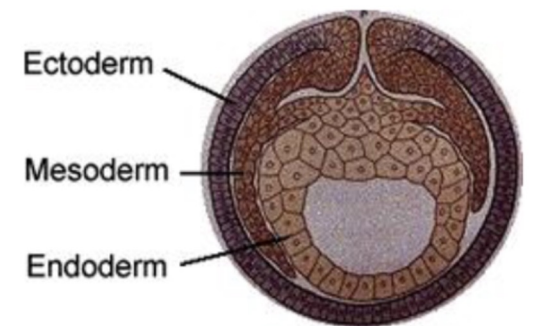
- **ecto(outer)derm** → nervous system and skin
- **meso(middle)derm** → bones, muscles, blood vessels
- **endo(inner)derm** → organs, glands

first 2 weeks

- embryo sphere elongates into a “worm”
  - still 3 layers of cells

neural plate formation

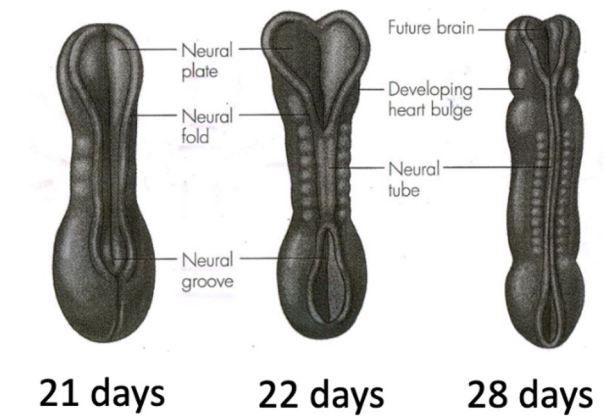
- dorsal ectoderm thickens into neural plate
- edges form neural folds (ridges) that curl up toward each other and fuse
- failed fusion: spina bifida



# embryonic development

week 4: curling and fusing of neural folds complete → neural tube

- becomes central nervous system (CNS)
  - rostral (anterior) end → brain
  - caudal (posterior) end → spinal cord
- surface ridges of neural tube (neural crest) becomes peripheral nervous system
  - hollow center becomes ventricles & central canal

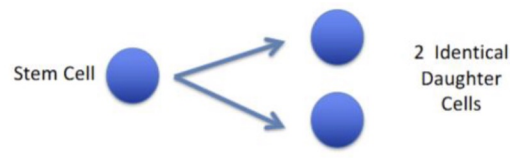


# proliferation (growth of new cells)

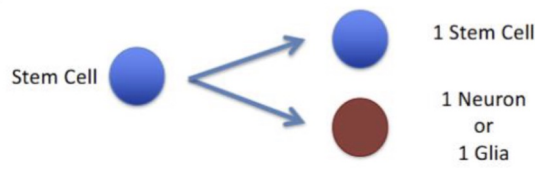
specifically neurons and glia in this context

stem cells: ectodermal cells lining neural tube (ventricular zone)

- early: symmetrical division (increasing size)



- ~week 7: asymmetrical division (produce ~100B neurons in 3 months)



\*each brain region proliferates differently depending on their specialized functions!

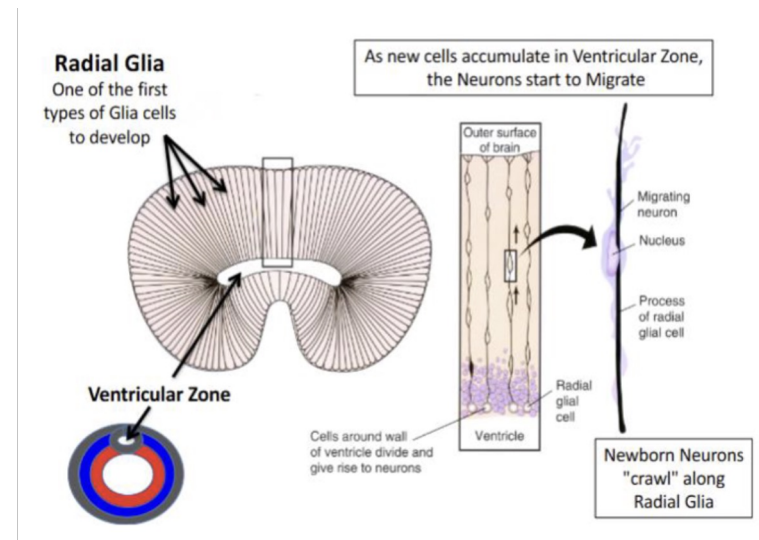
# migration and differentiation of cells

some stem cells become radial glia

- extend fibers outwards from ventricular zone, lengthen as cortex expands
- neurons “crawl” on them, aided by glycoproteins
  - some chemical trails (= neurotrophins) secreted by glia or other neurons

differentiation

- neurons develop specific structures and functions
- driven by
  - cell autonomous factors (genetic factors)
  - induction (local chemical signals / environmental conditions)





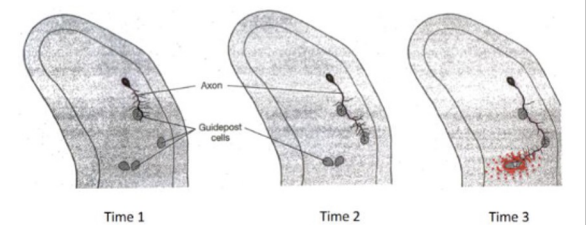
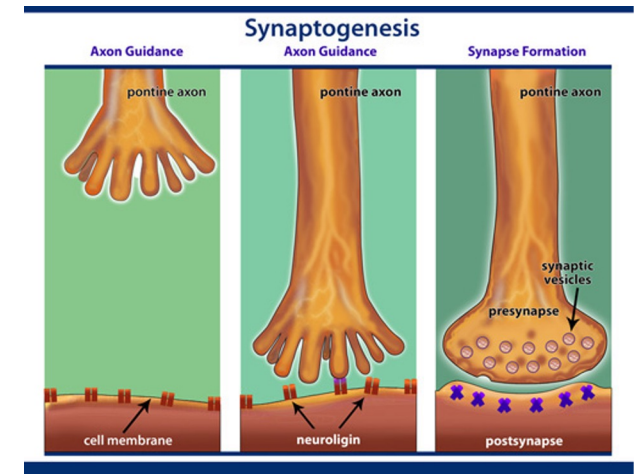
# synaptogenesis

= formation of synapses (cell-to-cell junctions)

- post-migration: neurons settle down; grow axons (outgoing) and dendrites (incoming)
- axons must seek appropriate postsynaptic targets
  - growth cone at axon tip uses filopodia (finger-like extensions) to detect local chemical gradients
  - guided by:
    - guidepost cells (glia): adhere to and direct growing axon to target cell
    - chemical trails (neurotrophins) produced by glia or other migrating neurons/axons

neurotrophins

- attract, repel, and promote neuronal survival & activity
- NGF (Nerve Growth Factor): from muscles/organs → attracts sympathetic NS axons & supports survival
- BDNF (Brain-Derived Neurotrophic Factor): promotes CNS axon survival & branching



# apoptosis

neurons have built-in suicide genes for programmed cell death

- activation depends on brain chemistry & activity patterns
  - eg. abnormal cell growth, failed connections

massive overproduction of neurons during fetal development (~50% more!)

- axons initially branch widely and connect to multiple sites
  - only few sites are strengthened and maintained over time
- neurons compete for connections and neurotrophic factors
  - “losers” (late arrivals, weak connections) undergo apoptosis

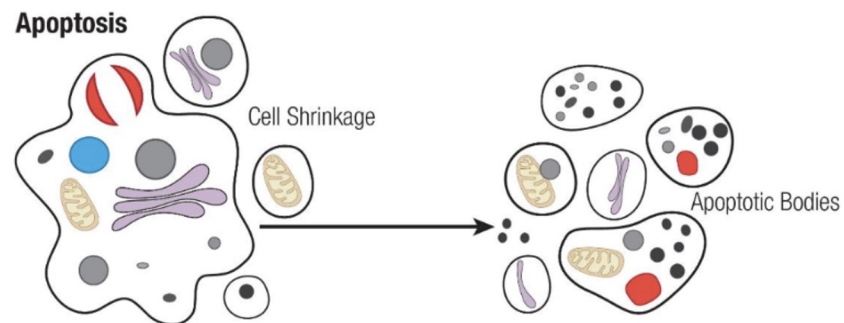
Normal Cells During Apoptosis:



Hey do you want  
to die



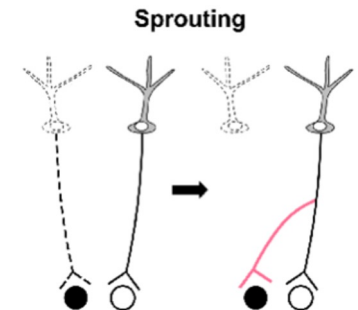
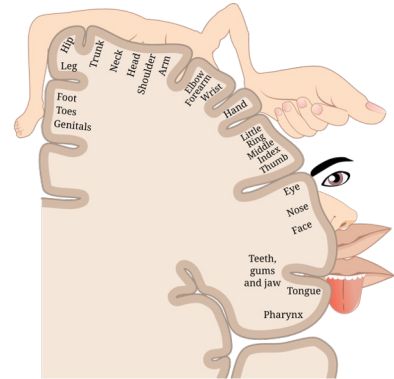
Yes



# patterns of co activity

## **CELLS THAT FIRE TOGETHER, WIRE TOGETHER!**

- coactivity strengthens connections; inactivity dies off connections
  - presynaptic activity triggers postsynaptic neurotrophin release → supports presynaptic cell survival
- neurotrophin release most effective on active presynaptic cells
  - higher correlation of pre/post activity in a pathway (stronger feedback)→stronger pathway
  - cells w/ less correlated activity are targeted for apoptosis
- collateral sprouting: surviving active cells (“winners”) take over losers’ synapses
- adjacent neurons co-activate and form topographic maps
  - spatial relationships along receptor surface preserved in brain



# further development

brain growth continues post-birth

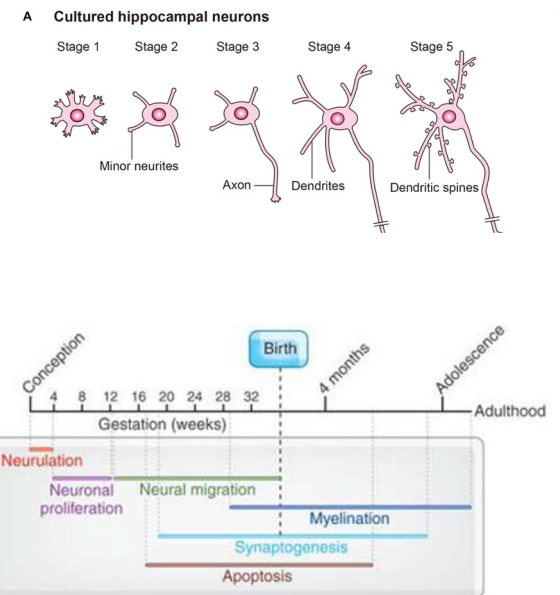
neurons increase in **cell size** and **dendridization** (branching), not in number

- new neurons are rare

dendritic development

- topographic maps formed during fetal development are continuously shaped by experience after birth
- neuroplasticity: learning / enriched environments → more dendritic spines and connections
- early sensory experiences shape circuits
  - eg. kittens exposed to only vertical lines can't perceive horizontal stimuli as cats
  - eg. violin players trained from childhood show expanded somatosensory map (parietal) in left hand fingers

glial development and differentiation continues (eg. myelination cont. through early adulthood)



k a h o o t

<https://play.kahoot.it/v2/?quizId=034733ea-86b4-48d4-bdee-3086fc0faf97&hostId=0889db3c-c5d4-454b-b692-99e48772950b>

go study 🦊 good luck on your midterm



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