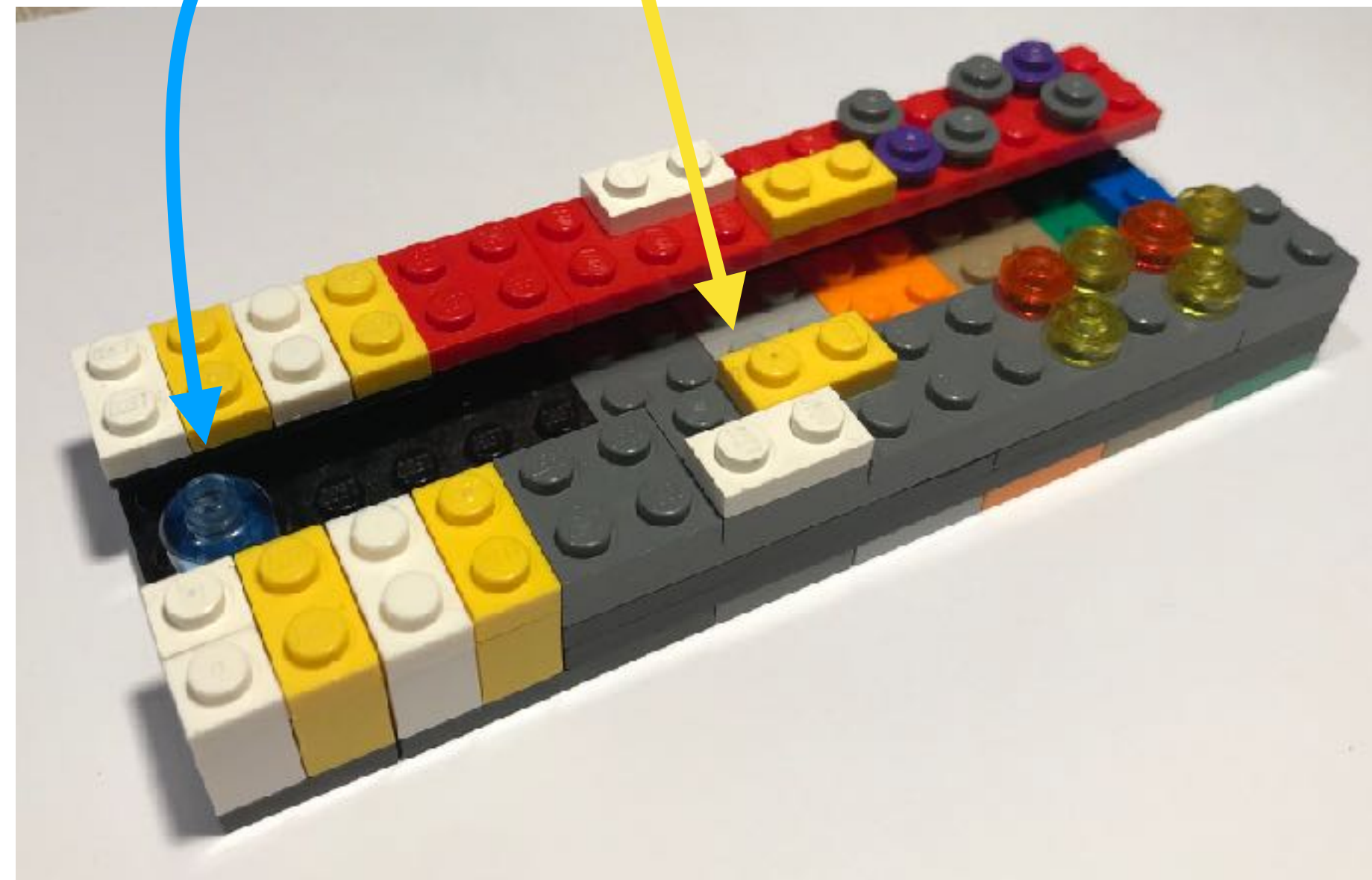
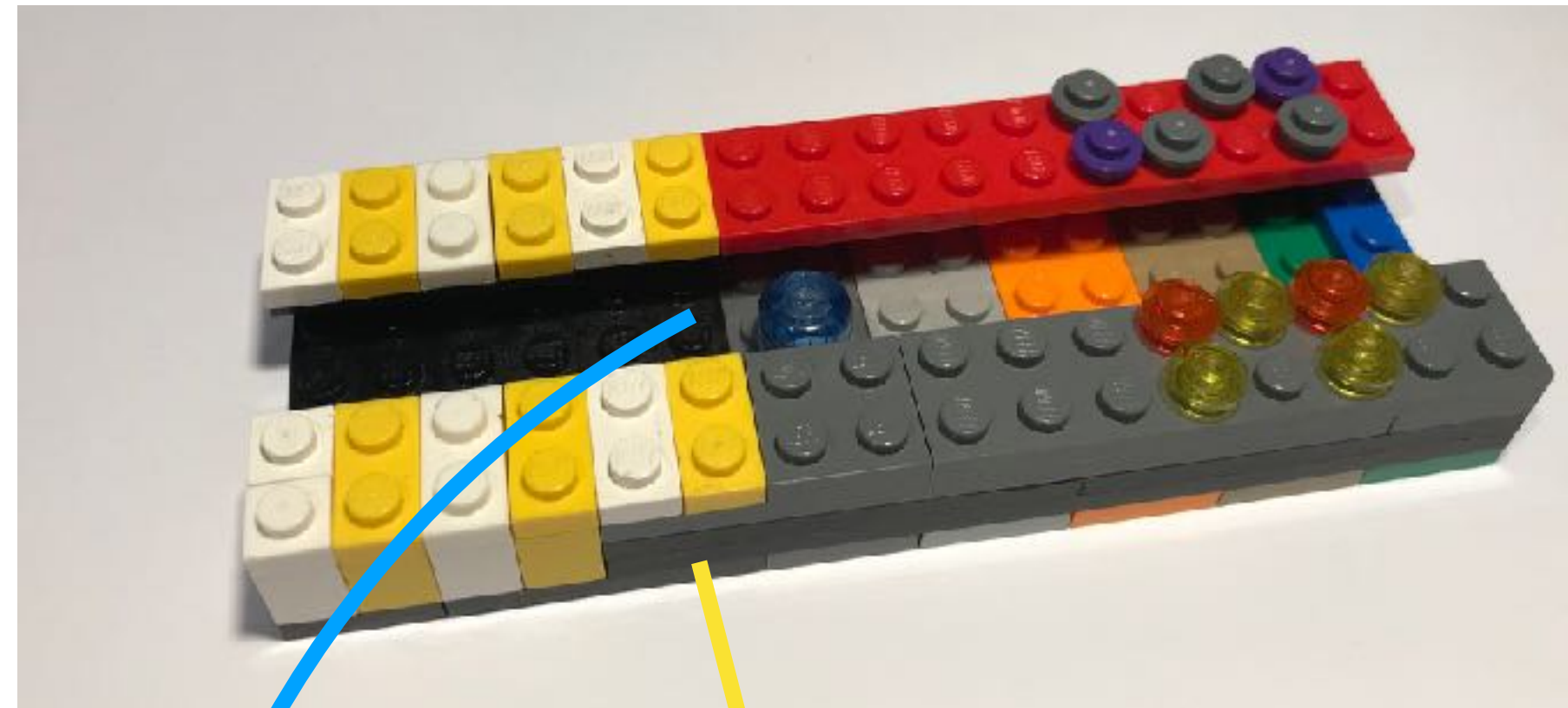
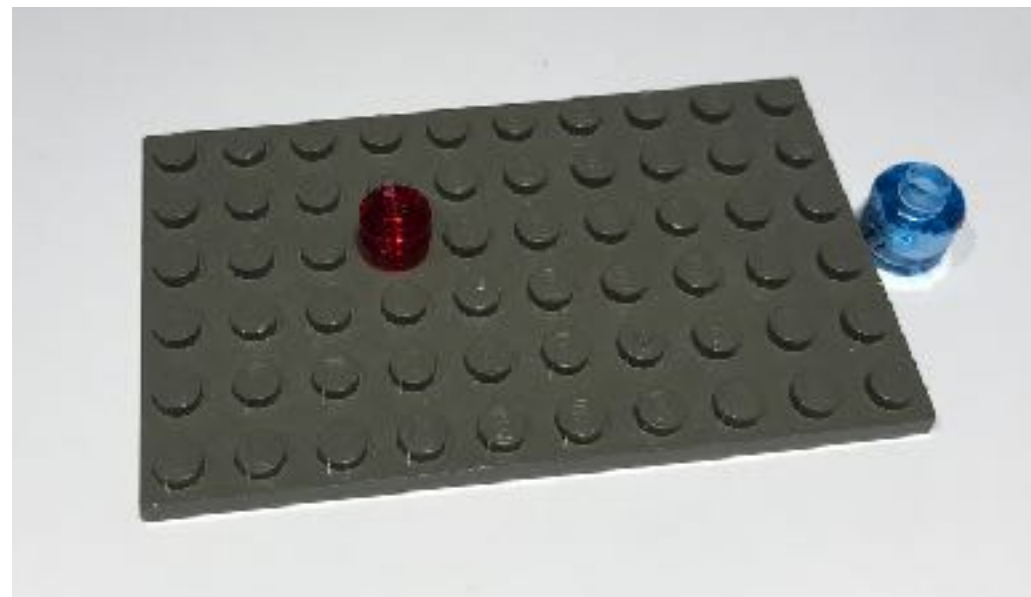




Big questions



Cell migration - finishing touches & completing the cycle for the next generation



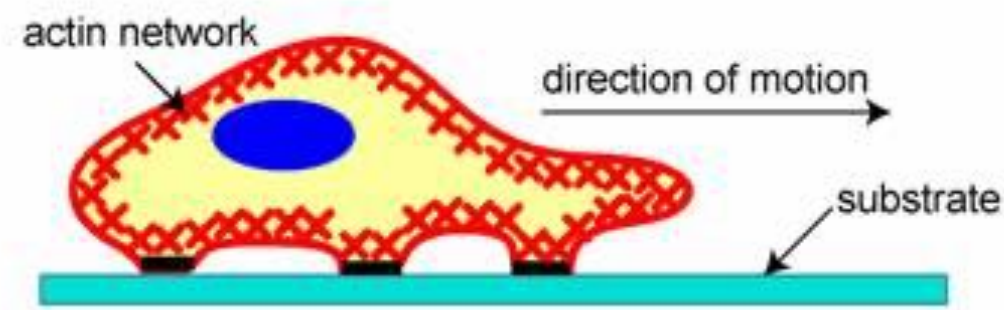
Today's menu

- **Cell migration**
 - *Basic cellular mechanisms*
 - actomyosin + adhesion = traction forces
 - motility + polarization = migration
- **The neural crest and placodes**
 - *The “4th germ layer”*
 - Sculpting face, heart, gut, sensory systems
- **Germ cell migration**
 - *Completing the cycle of life*
 - Wiping and silencing the genome
 - Long Range migration
 - Bi-potency

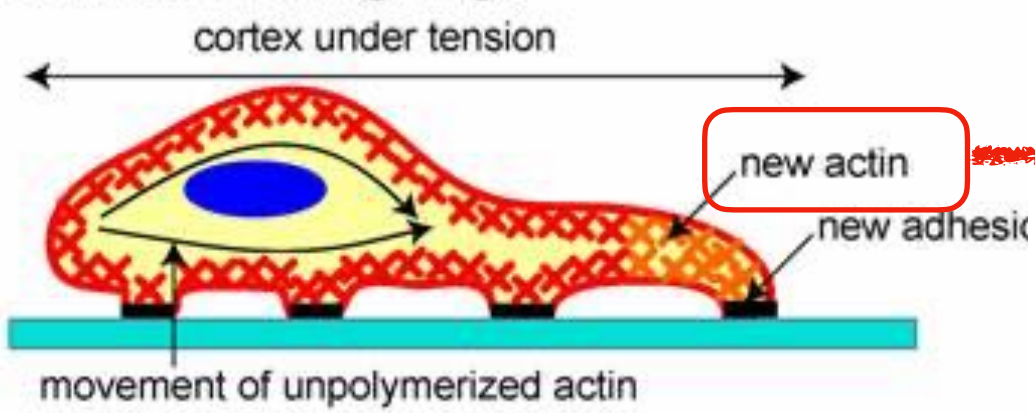
Migration - Basic cellular mechanisms

- Actomyosin + adhesion = traction forces

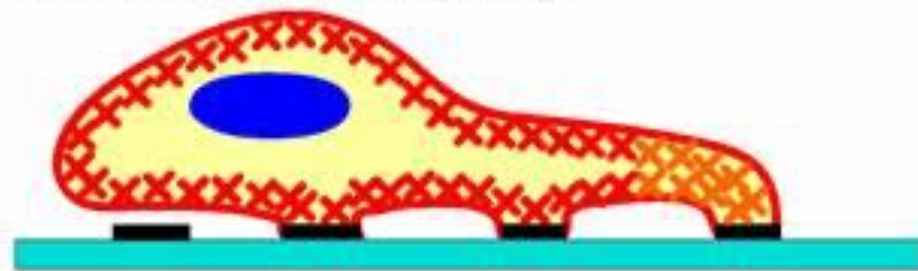
1) Protrusion of the Leading Edge



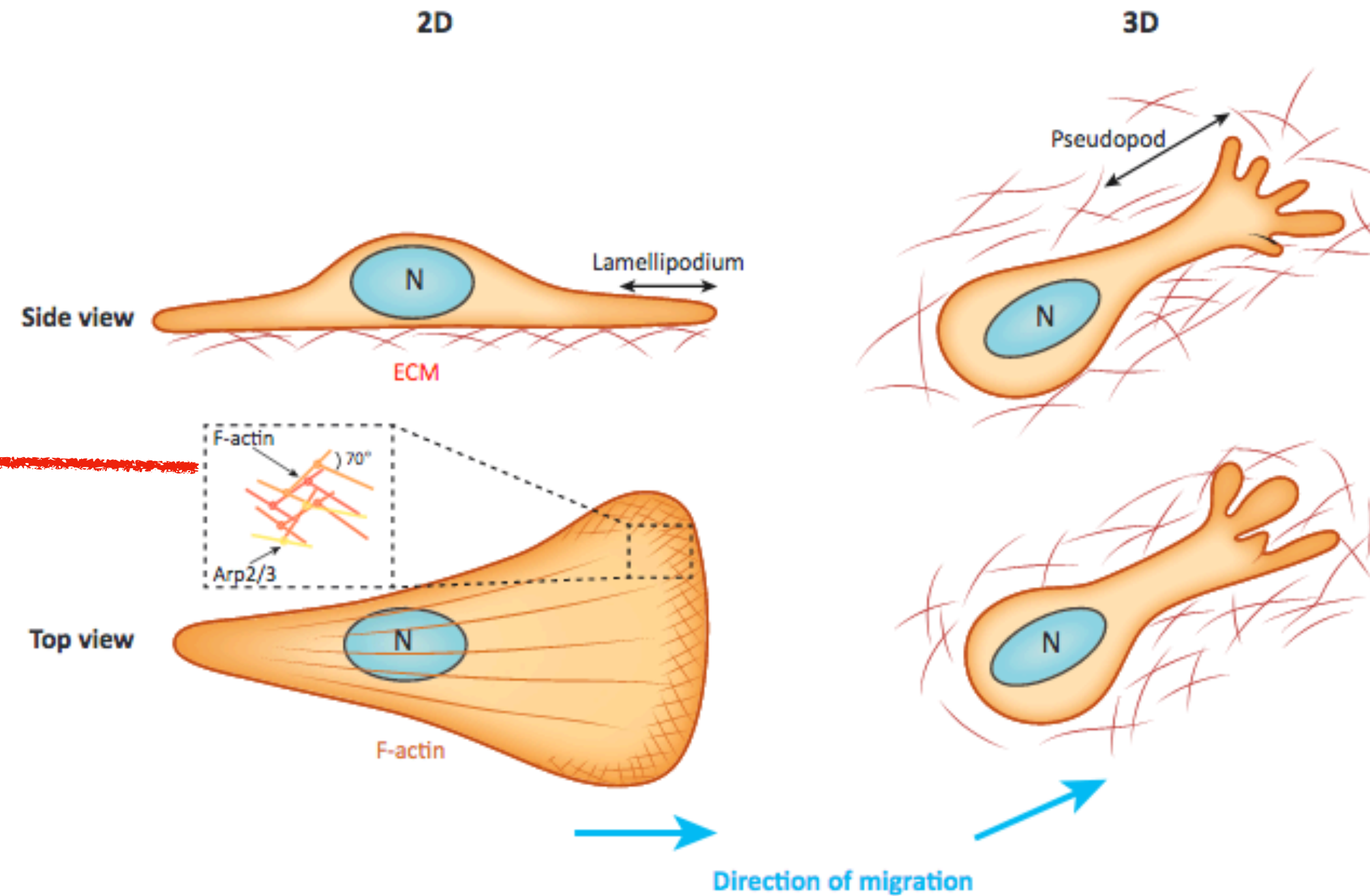
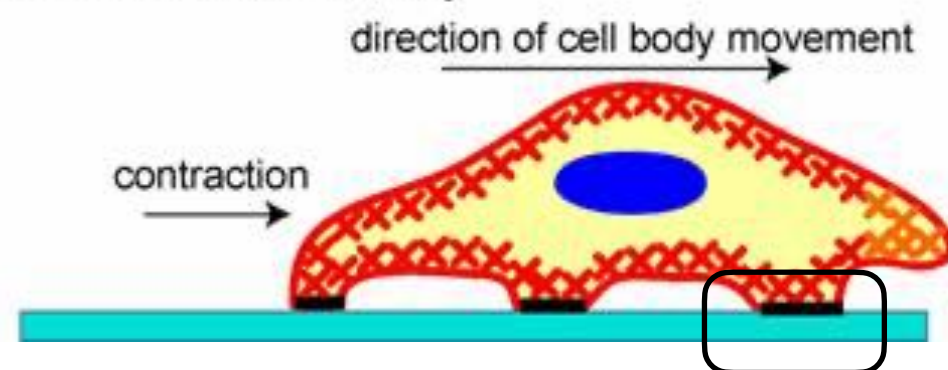
2) Adhesion at the Leading Edge



Deadhesion at the Trailing Edge



3) Movement of the Cell Body



Adhesion receptors: Integrin, Cadherin, etc

Migration - Basic cellular mechanisms

- Actin polymerization and branching at leading edge

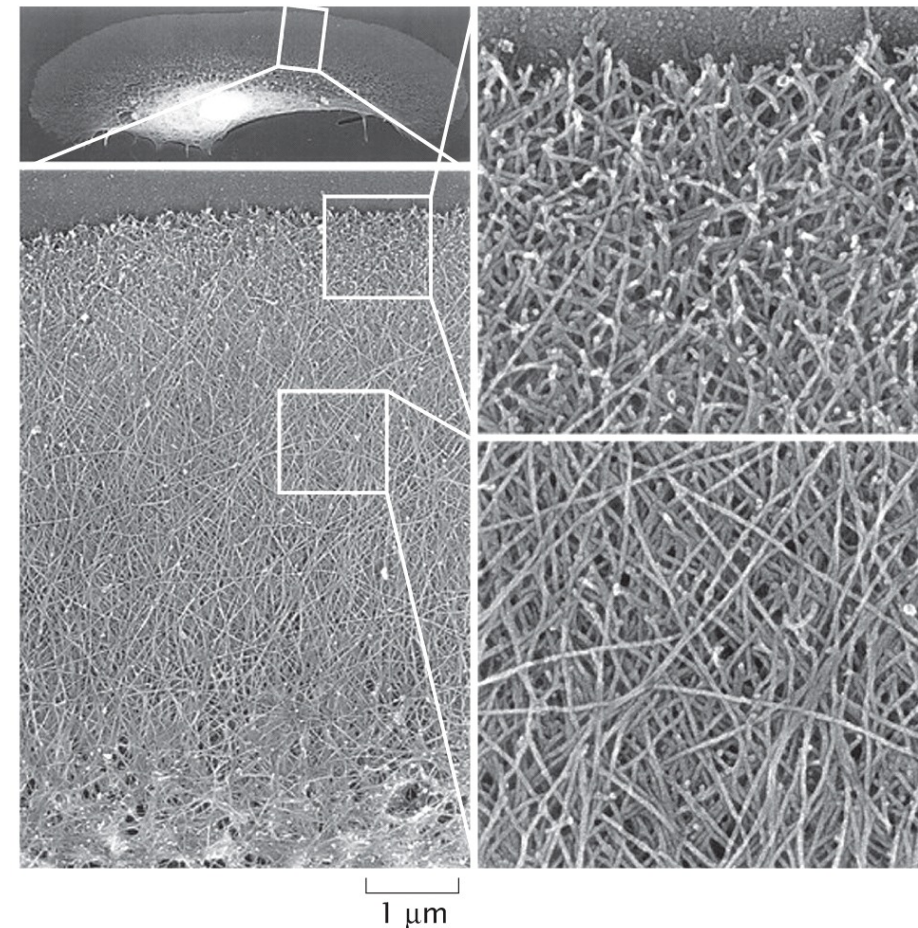


Figure 14.2 Physical Biology of the Cell, 2ed. (© Garland Science 2013)

in vivo polymerization

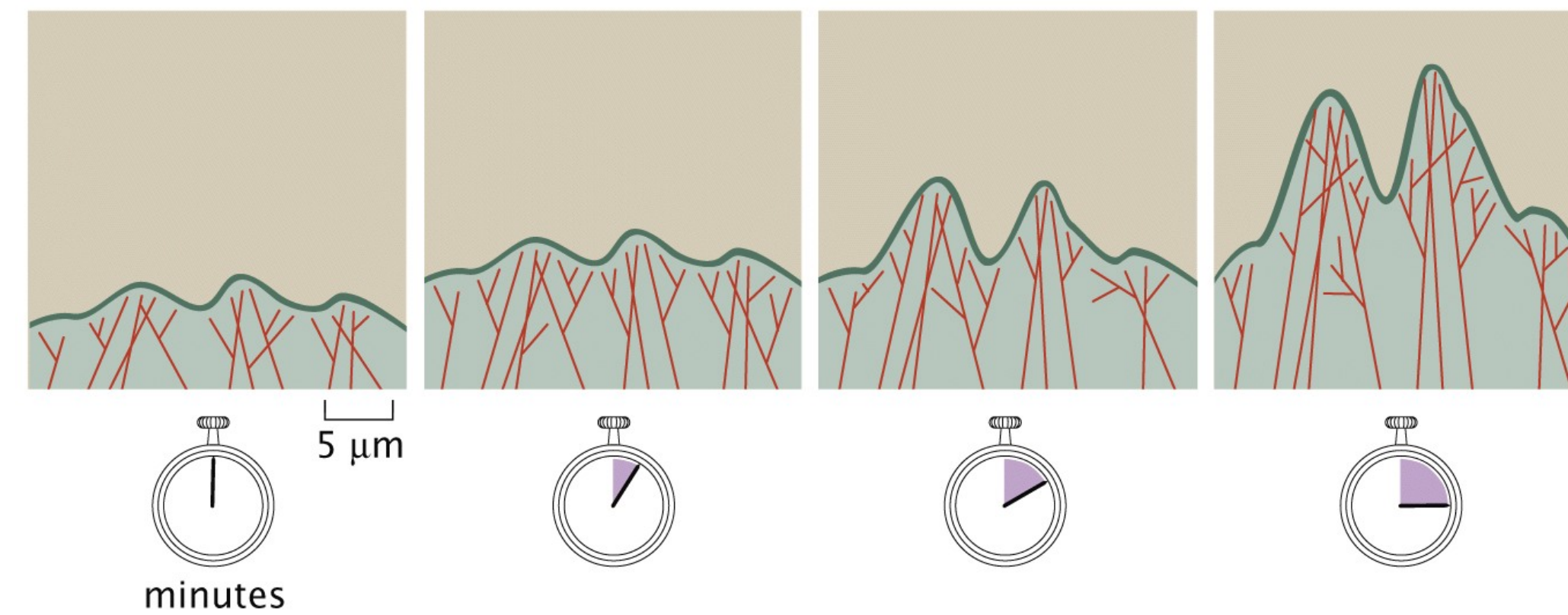
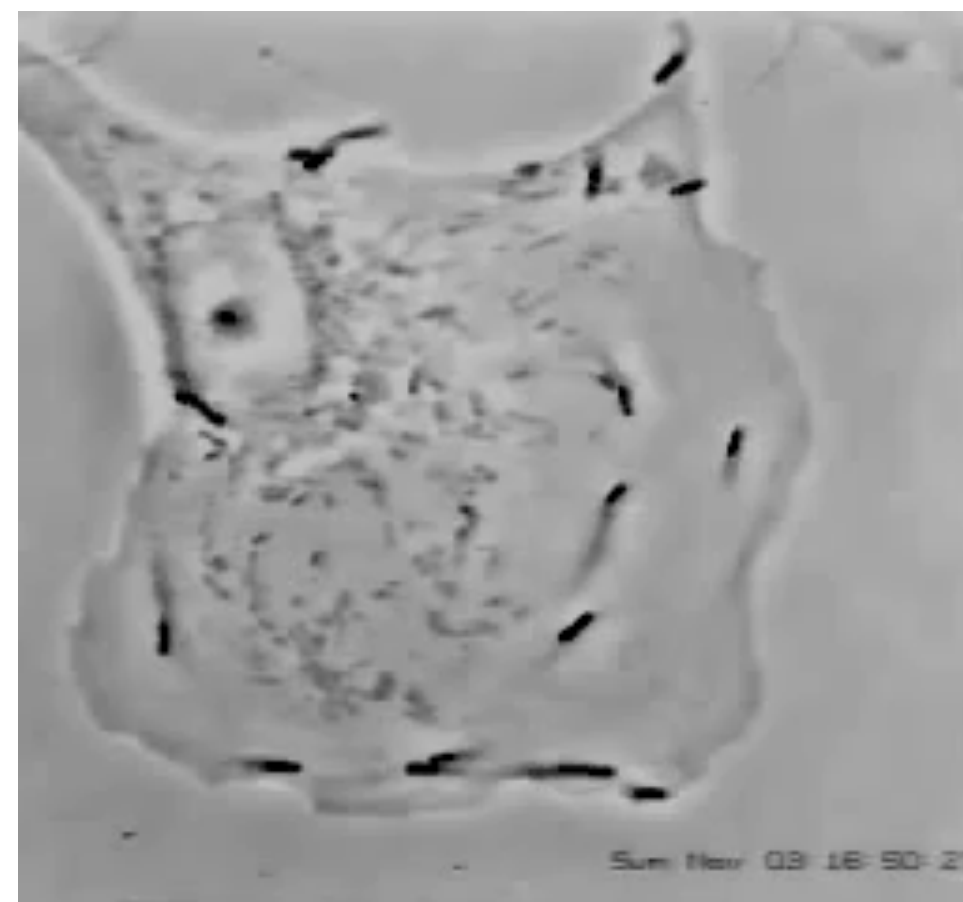
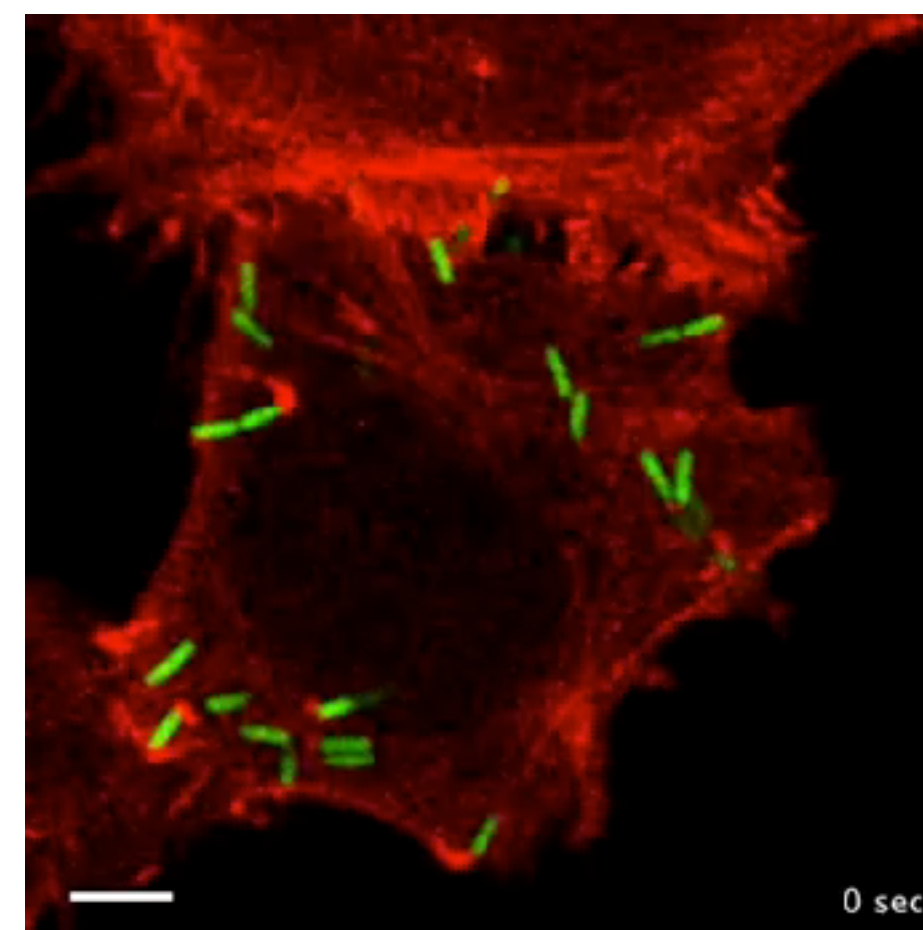


Figure 15.1b Physical Biology of the Cell, 2ed. (© Garland Science 2013)



Listeria propelled inside cell



Listeria (green), actin (red)

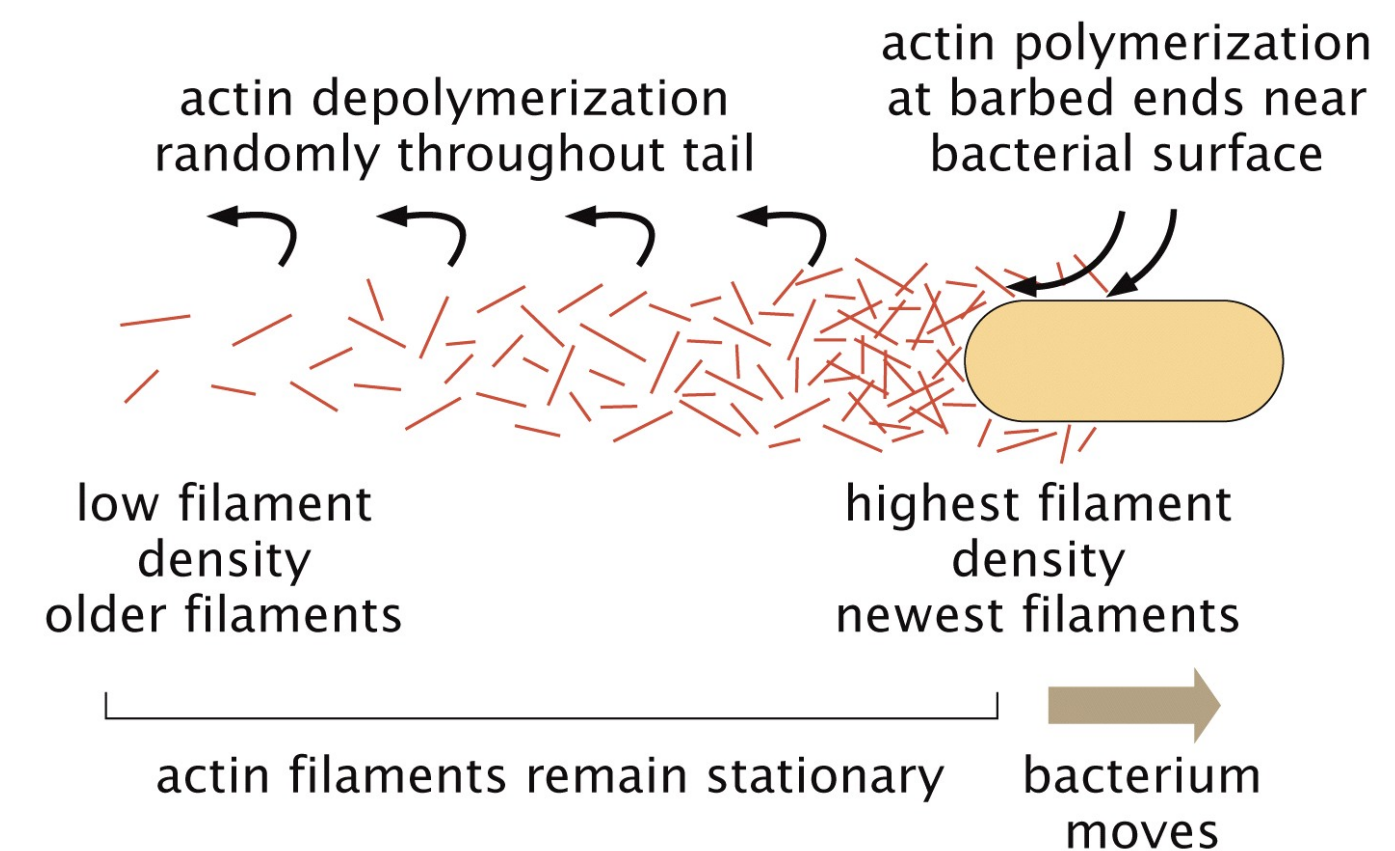
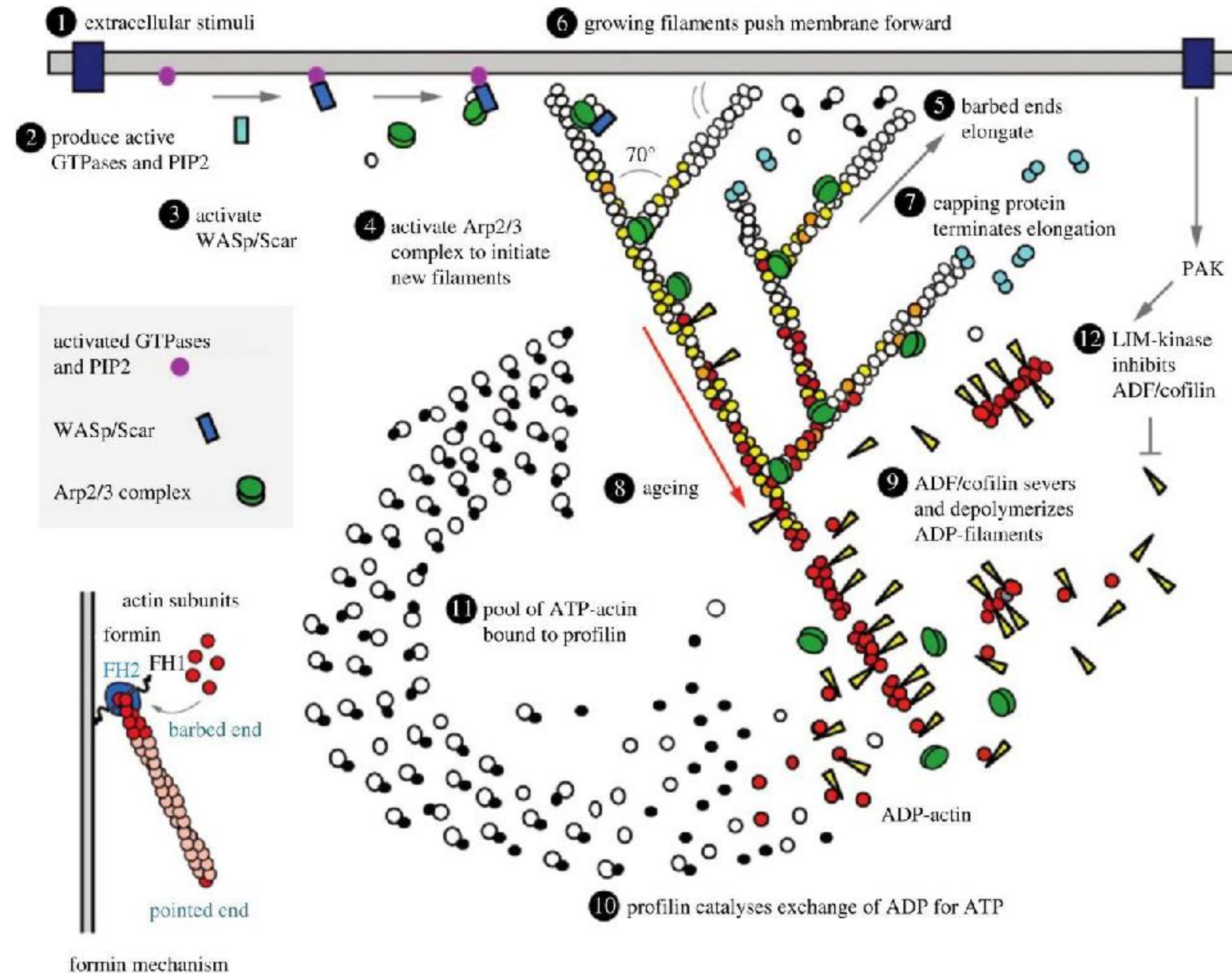


Figure 15.3b Physical Biology of the Cell, 2ed. (© Garland Science 2013)

Migration - Basic cellular mechanisms

- Actin polymerization and branching at leading edge

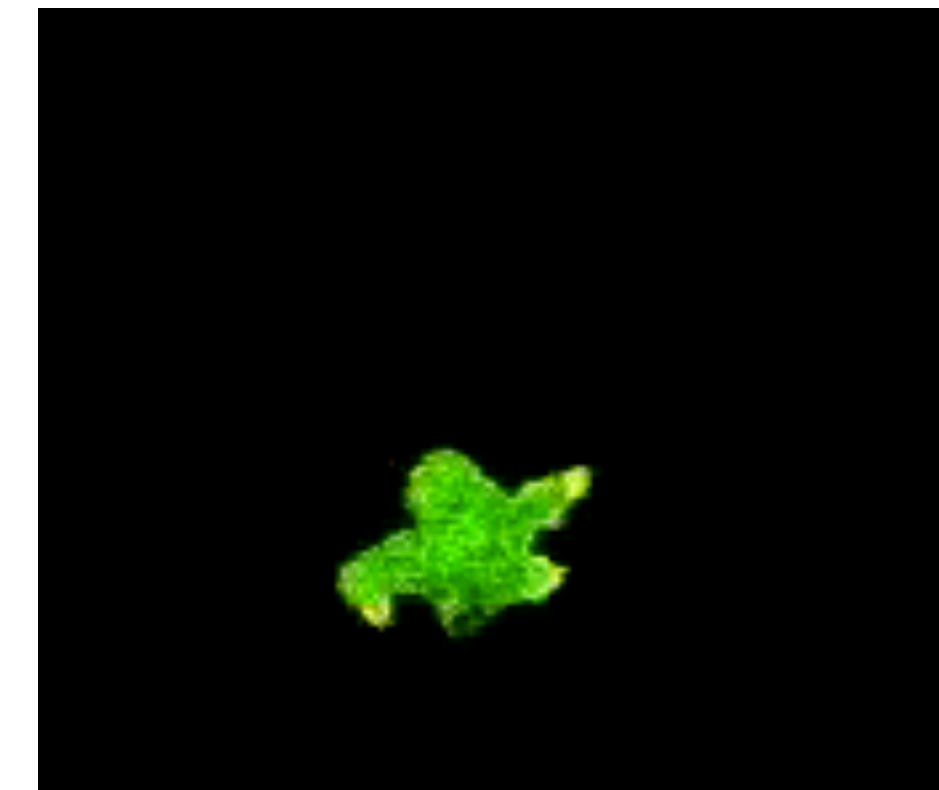
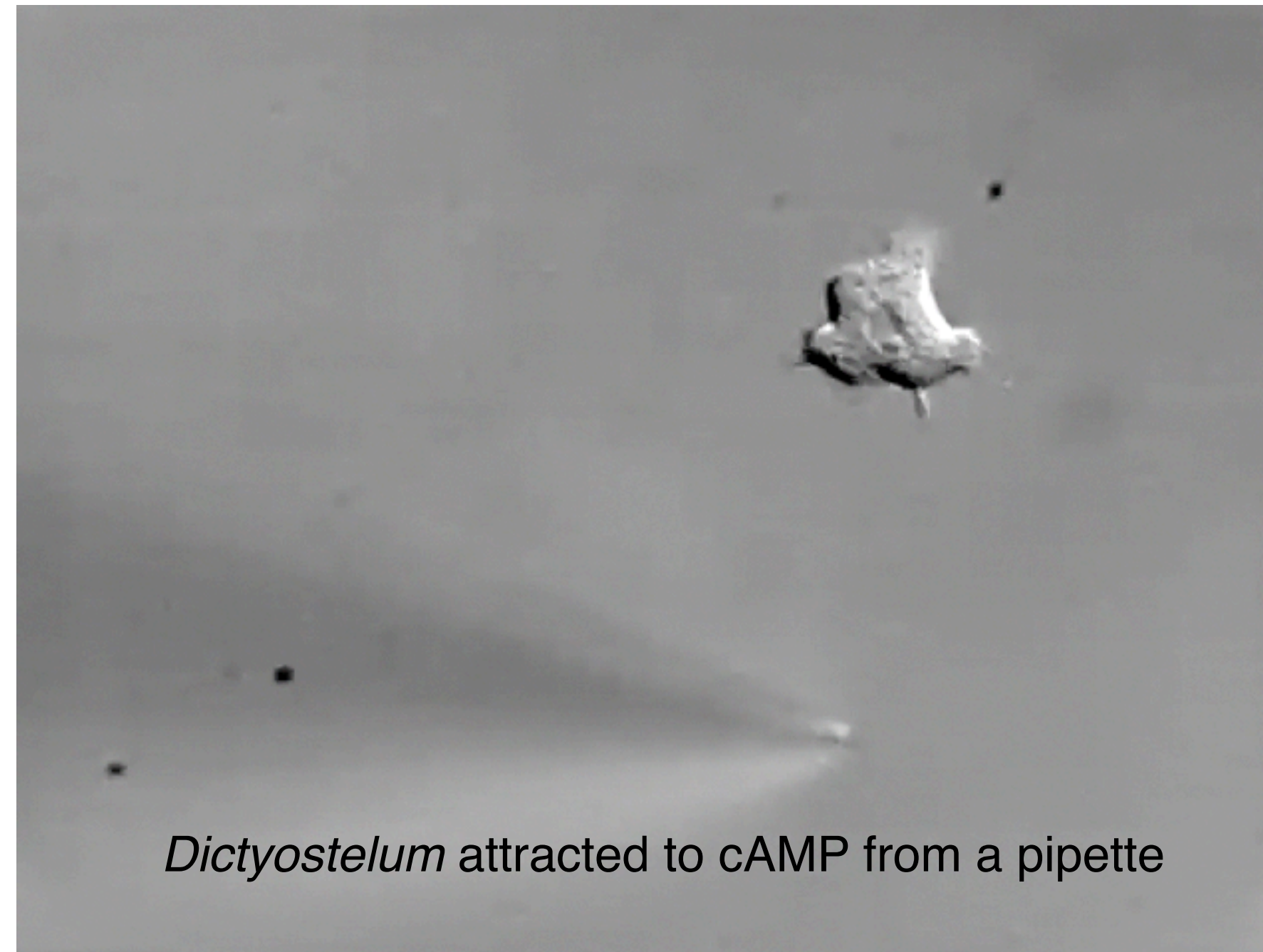


Migration - Basic cellular mechanisms

- motility + polarization = migration



Human neutrophil
chasing bacterium

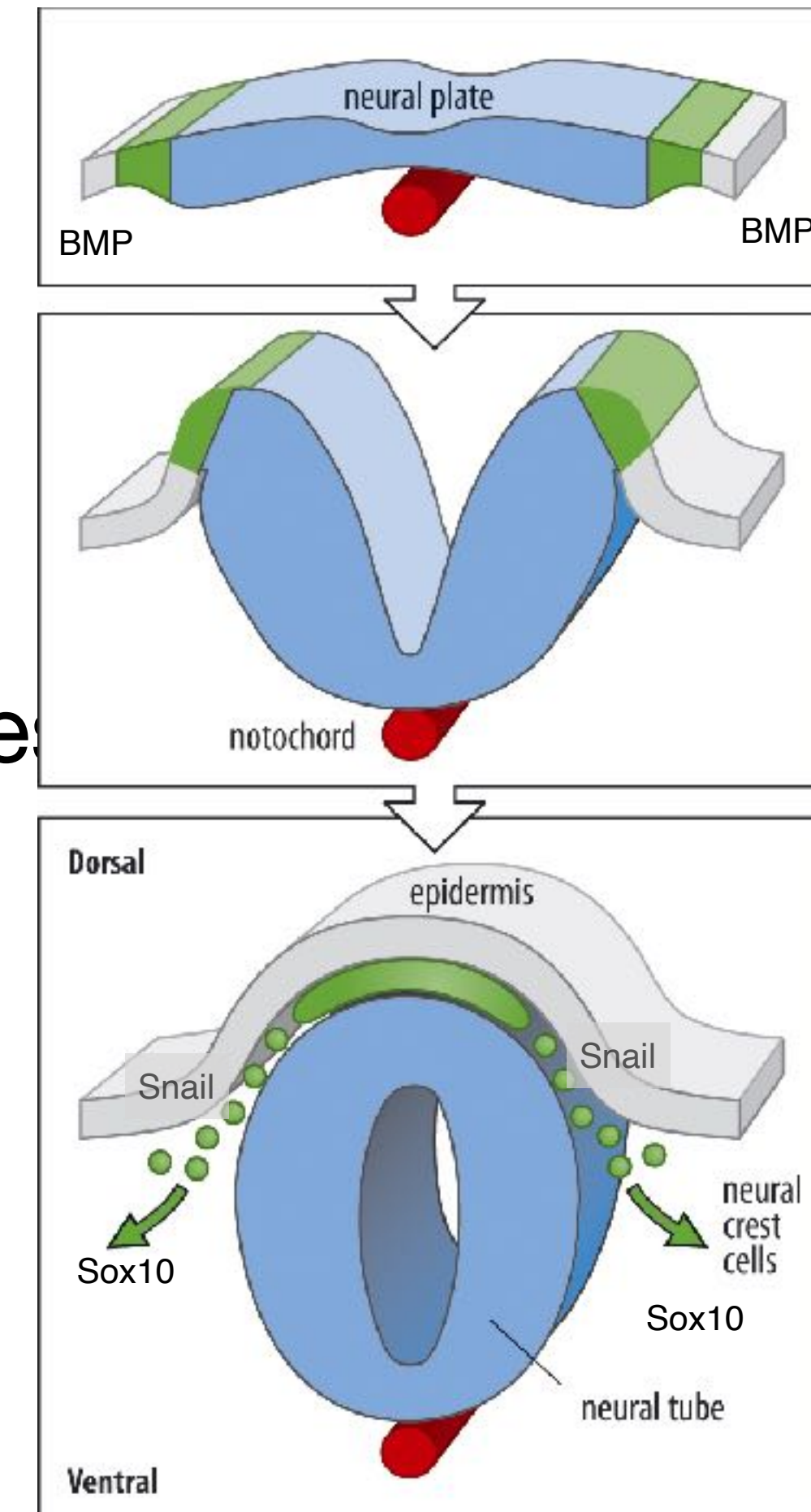


Green = membrane
Red = LimE-RFP = F-actin

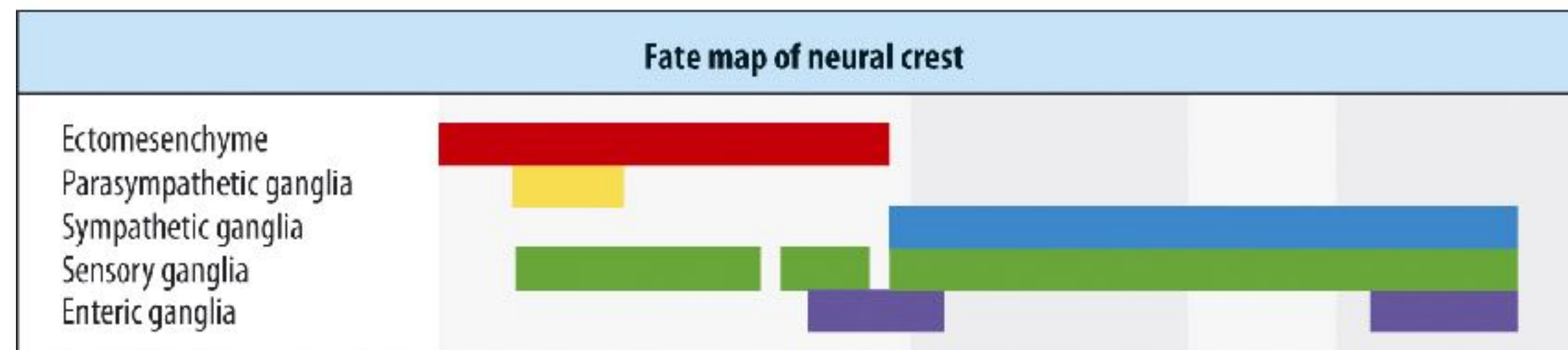
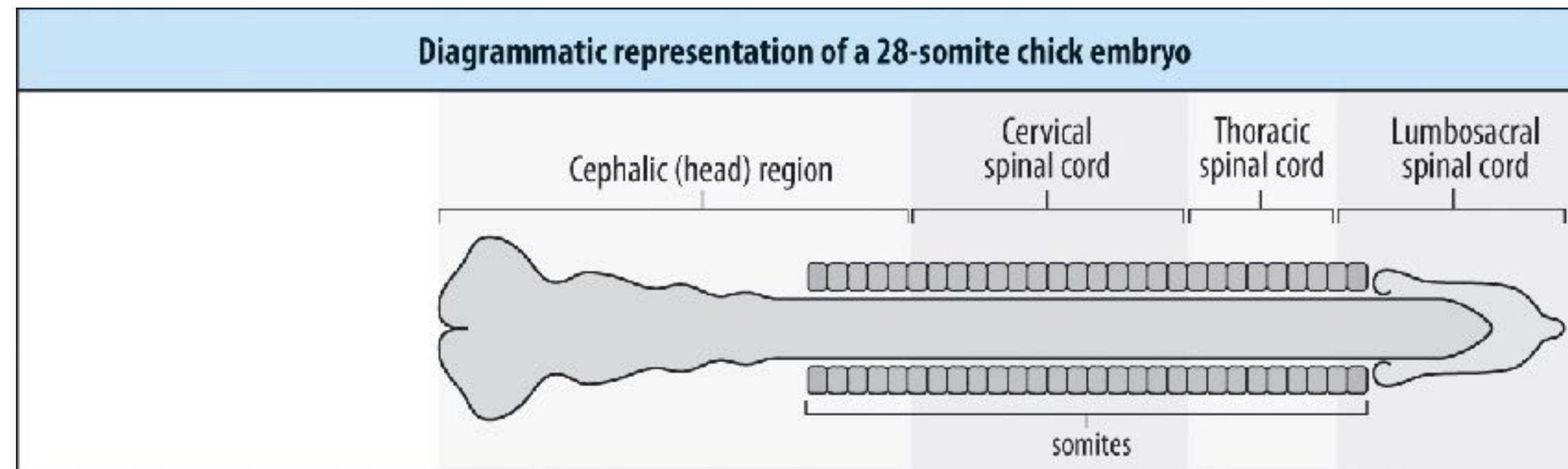
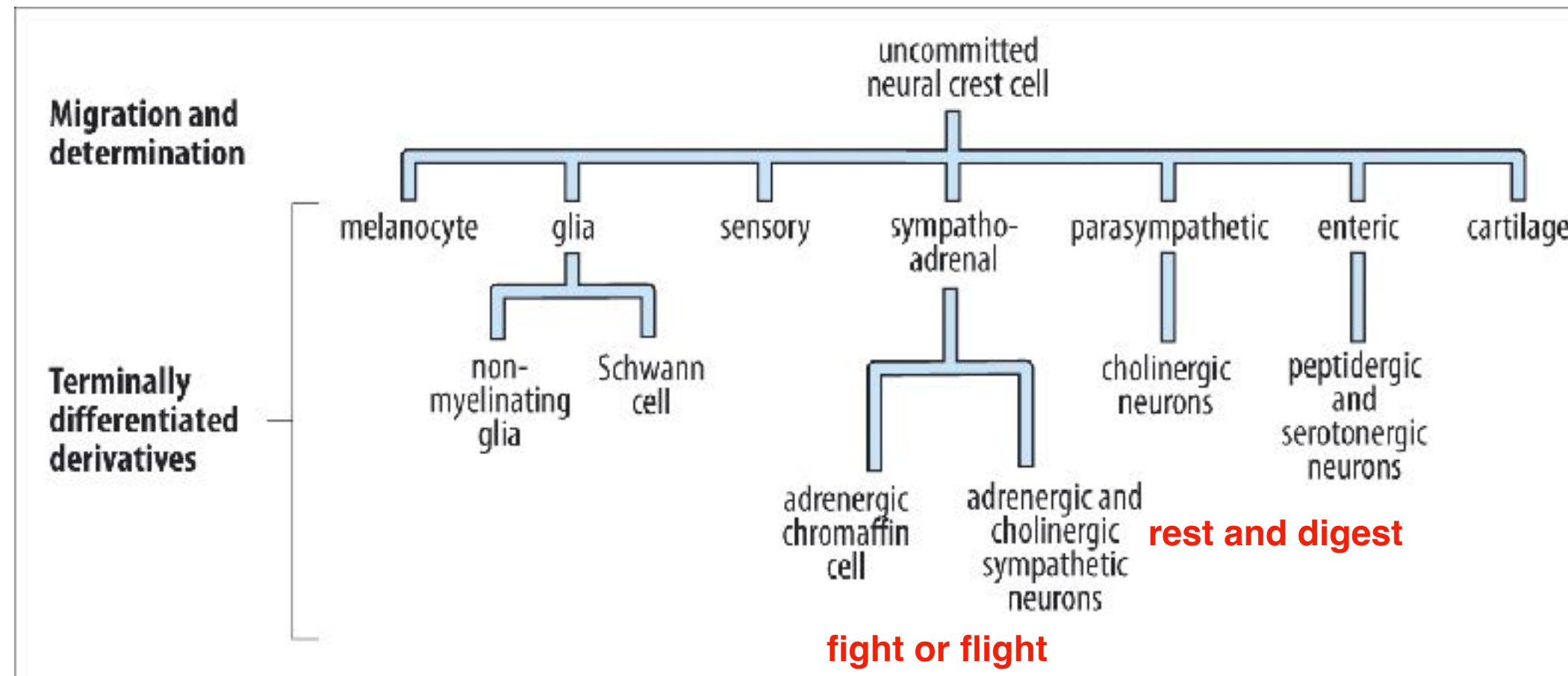
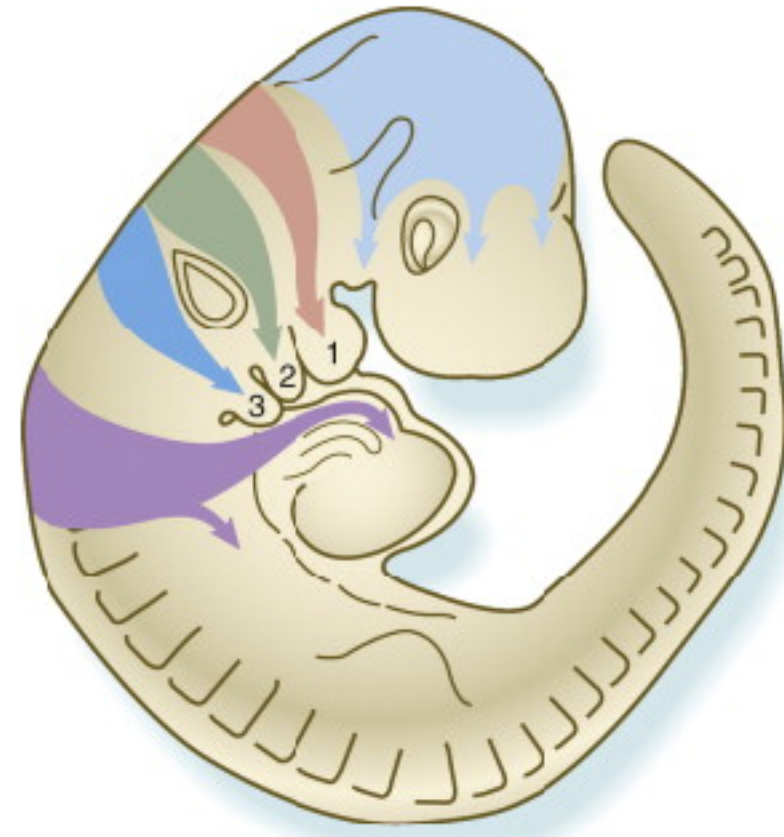
- Polarization - *persistent* directed motion
- Chemoattractant receptors engaged on one side of cell
- Engage Ca^{2+} and small GTP effectors
- Remodel cytoskeleton

The neural crest and sensory placodes

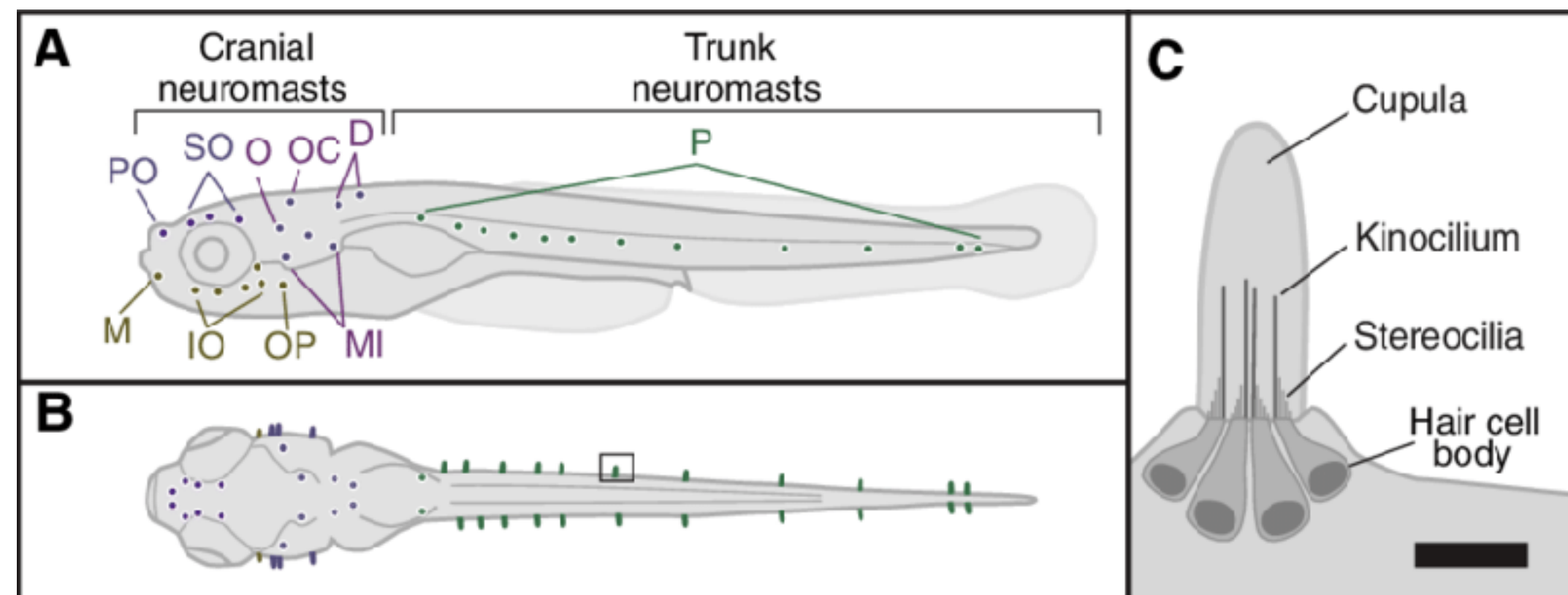
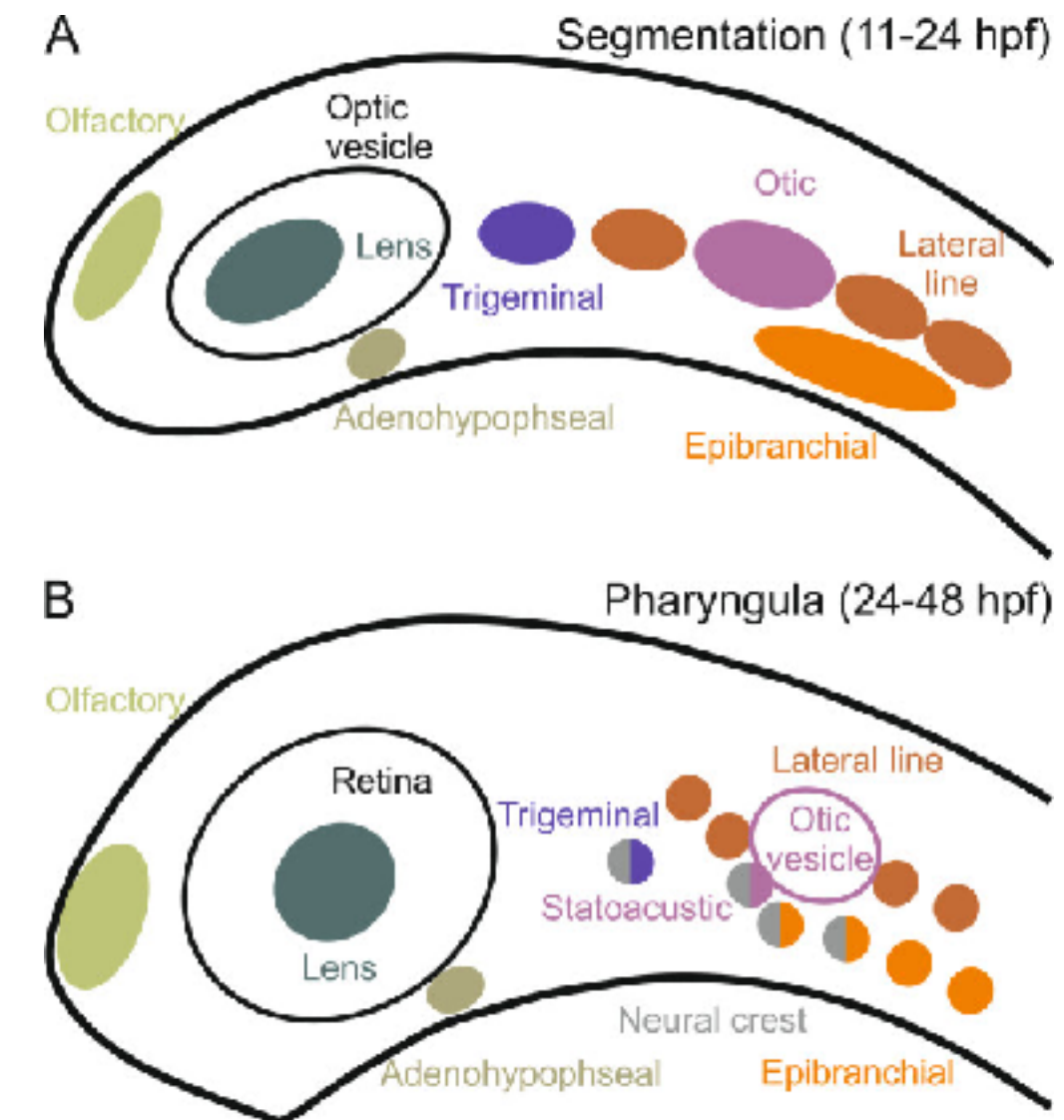
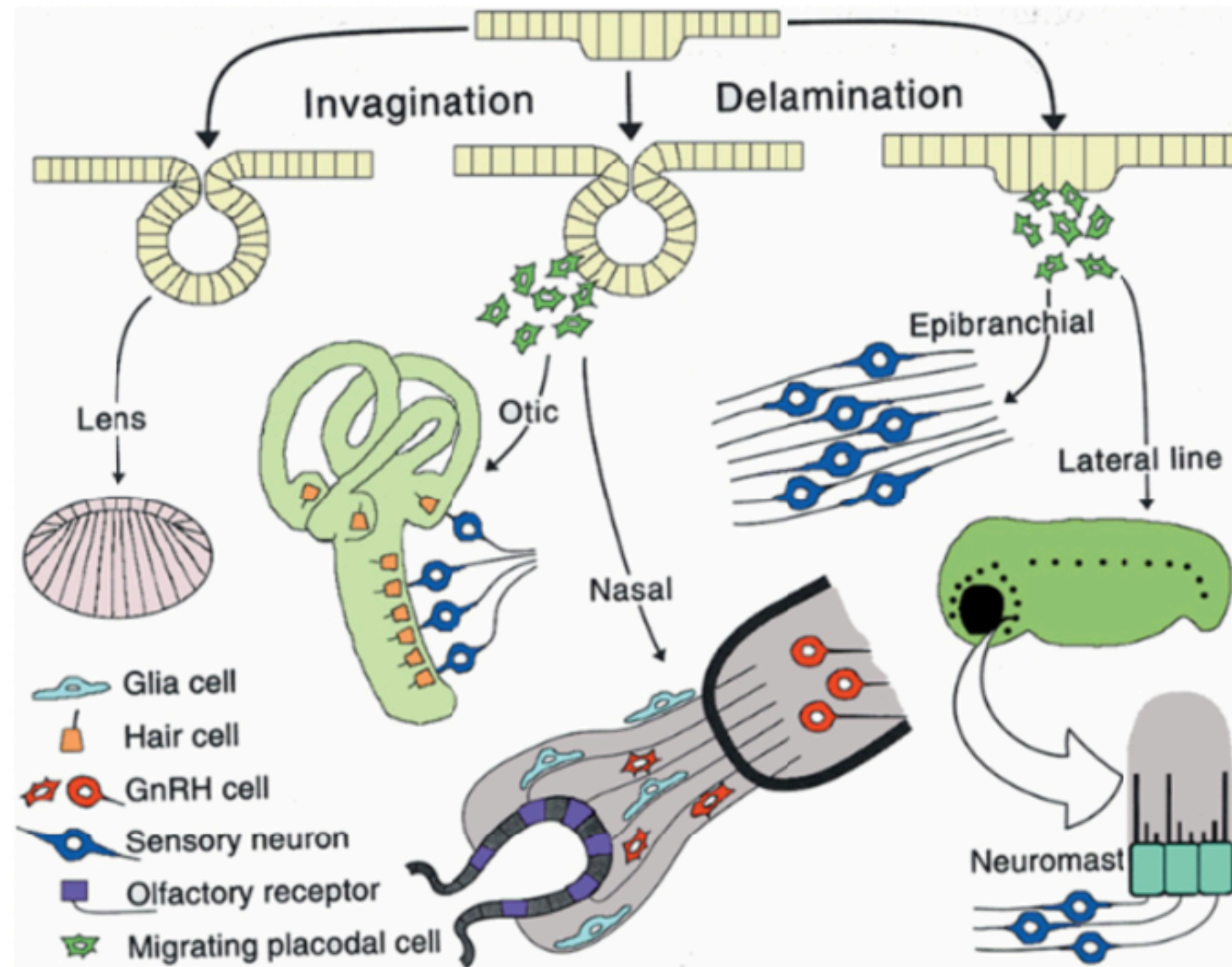
- Derived from neuroepithelium
- NC cells delaminate and migrate away
- Placodal cells deform epithelium + migrate
- Seed existing structures / build new structures



Neural crest-derived structures



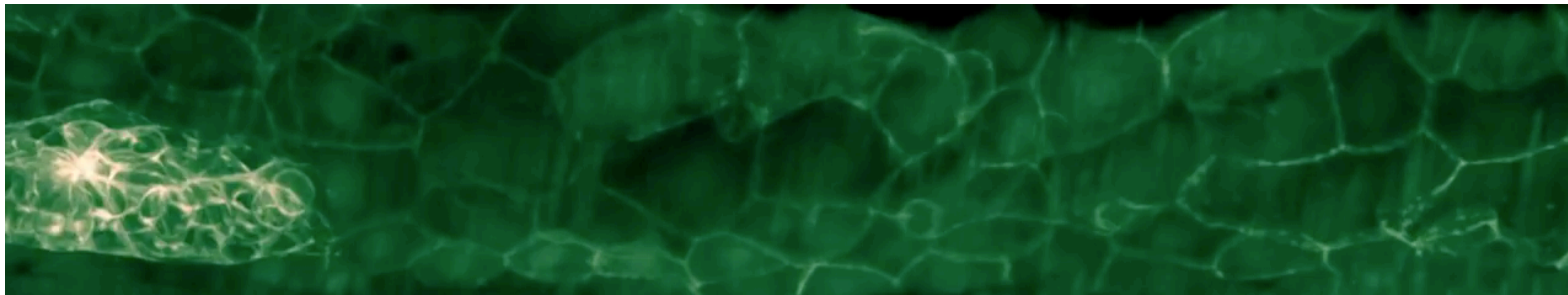
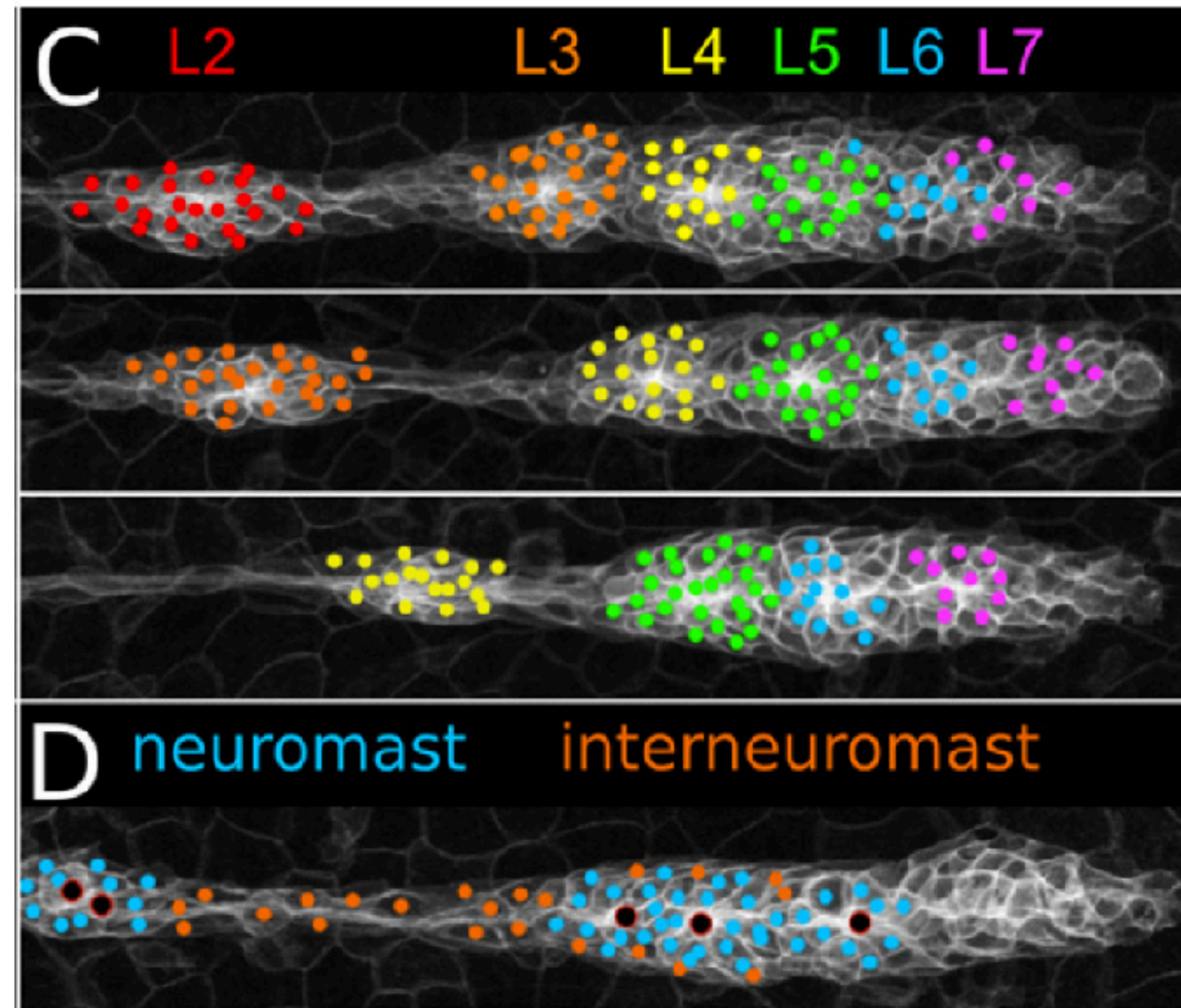
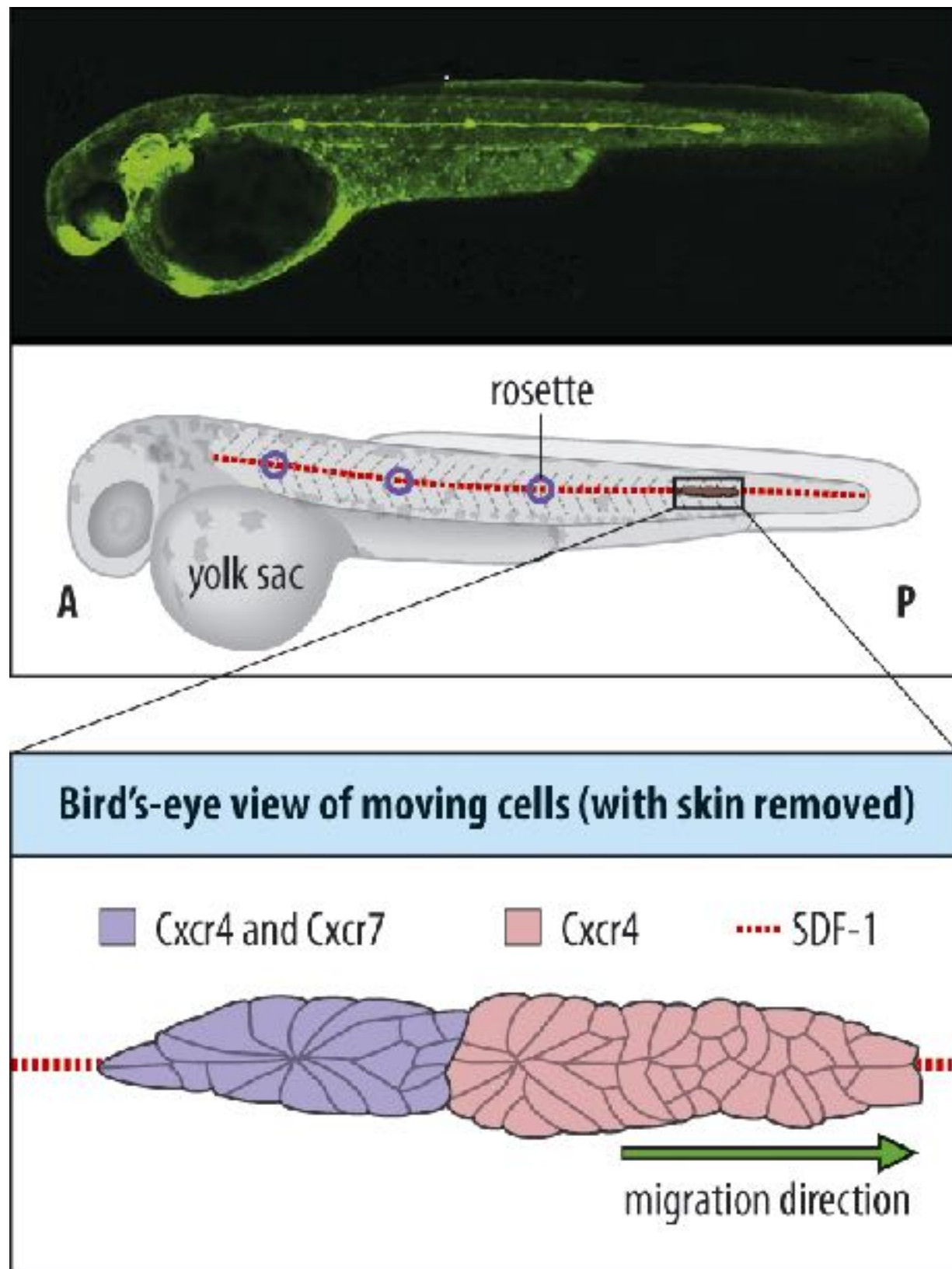
Placodal-derived sensory structures



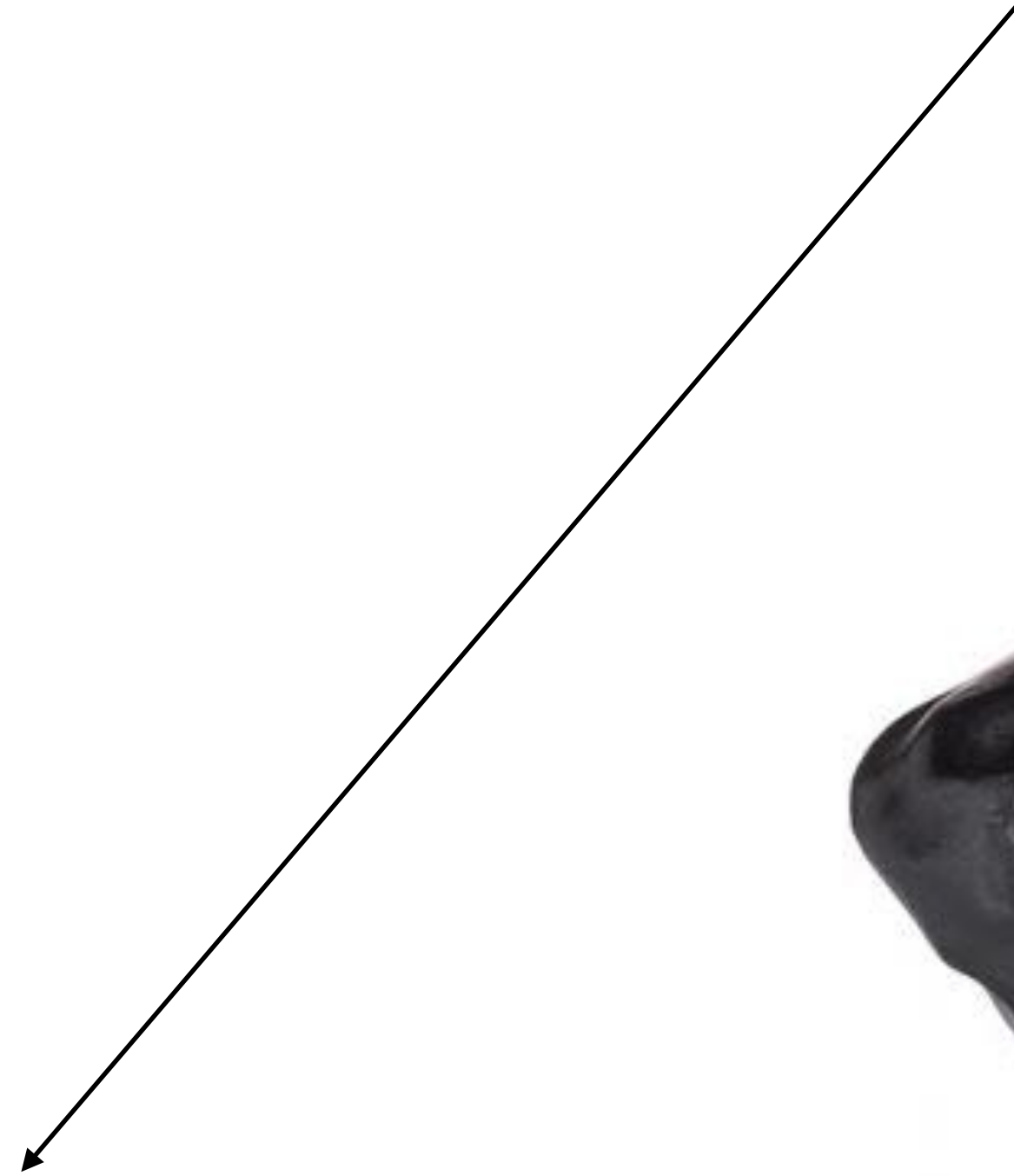
Sensory organ to measure water pressure waves

Blind fish can school

The zebrafish lateral line - collective migration

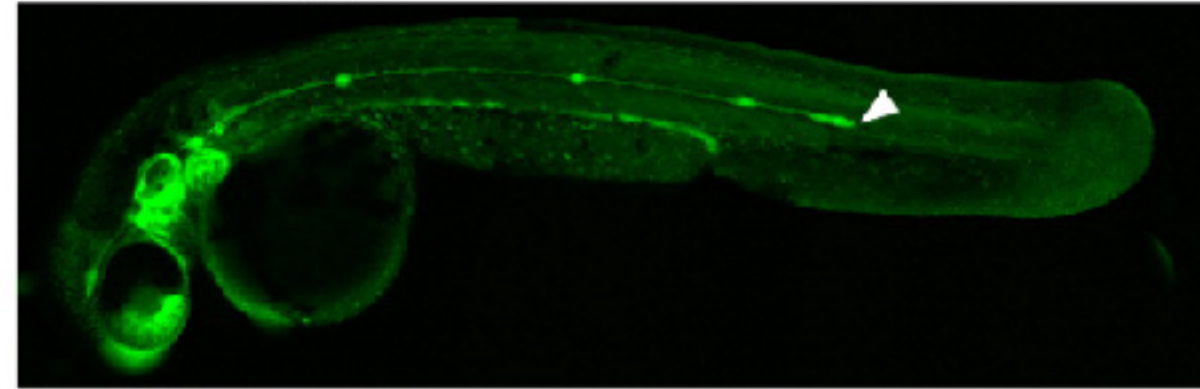


Individual puzzle

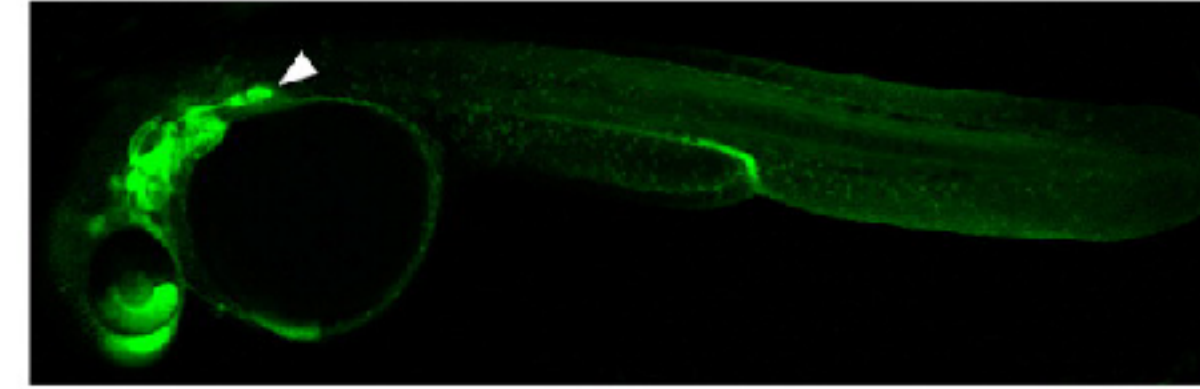


Three genes required for primordium migration

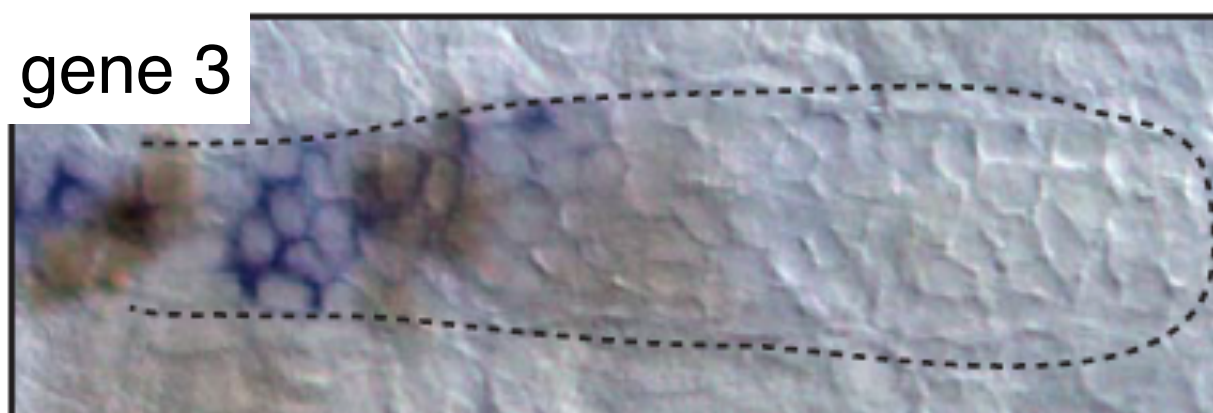
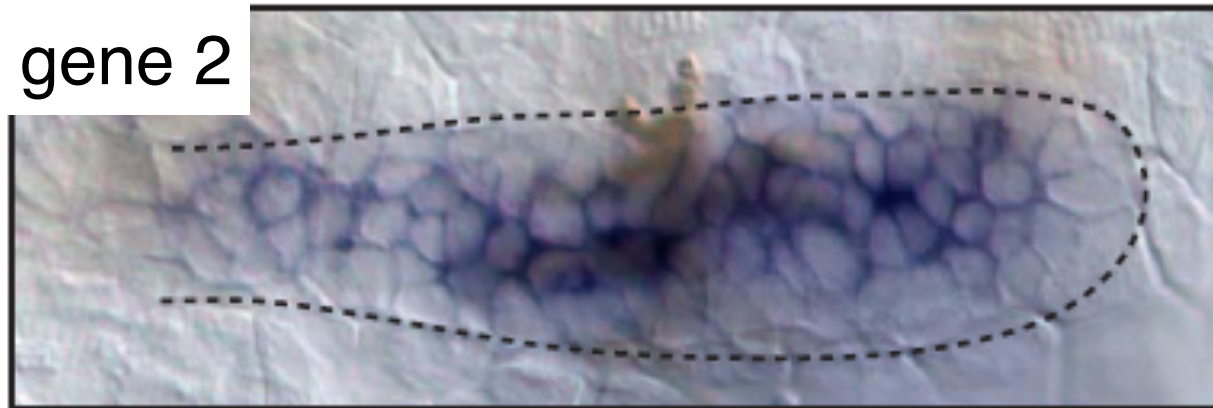
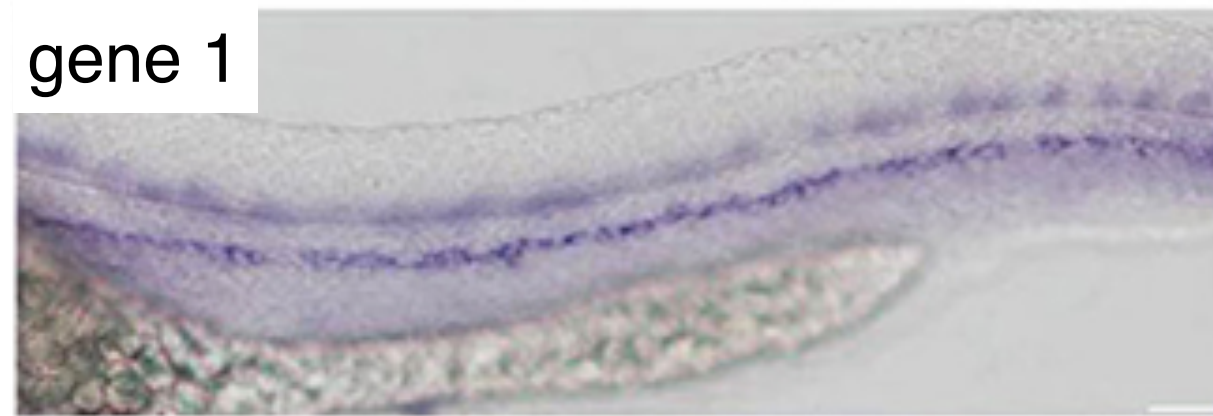
A Wild type



B Mutants' lateral lines



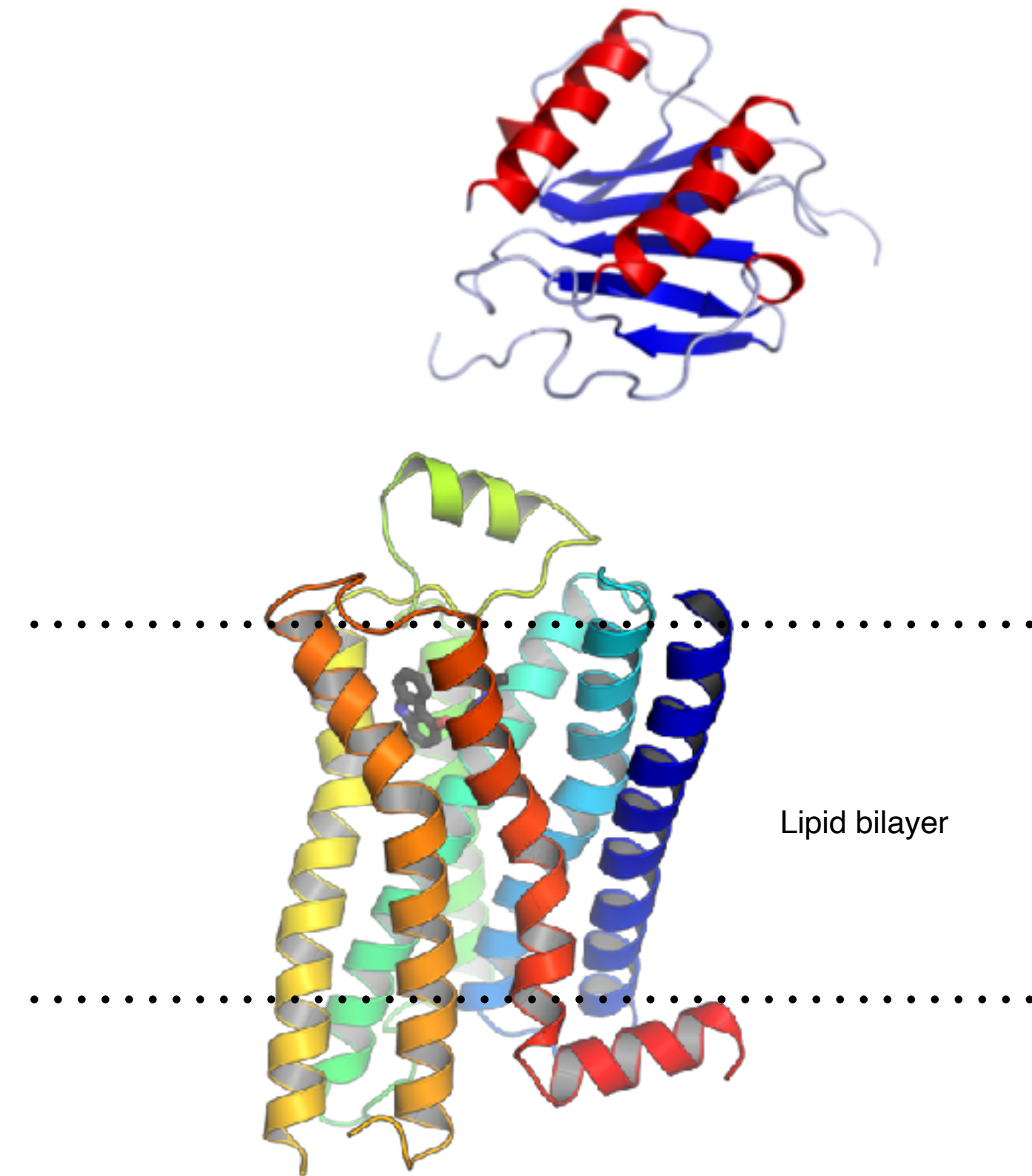
Gene expression patterns



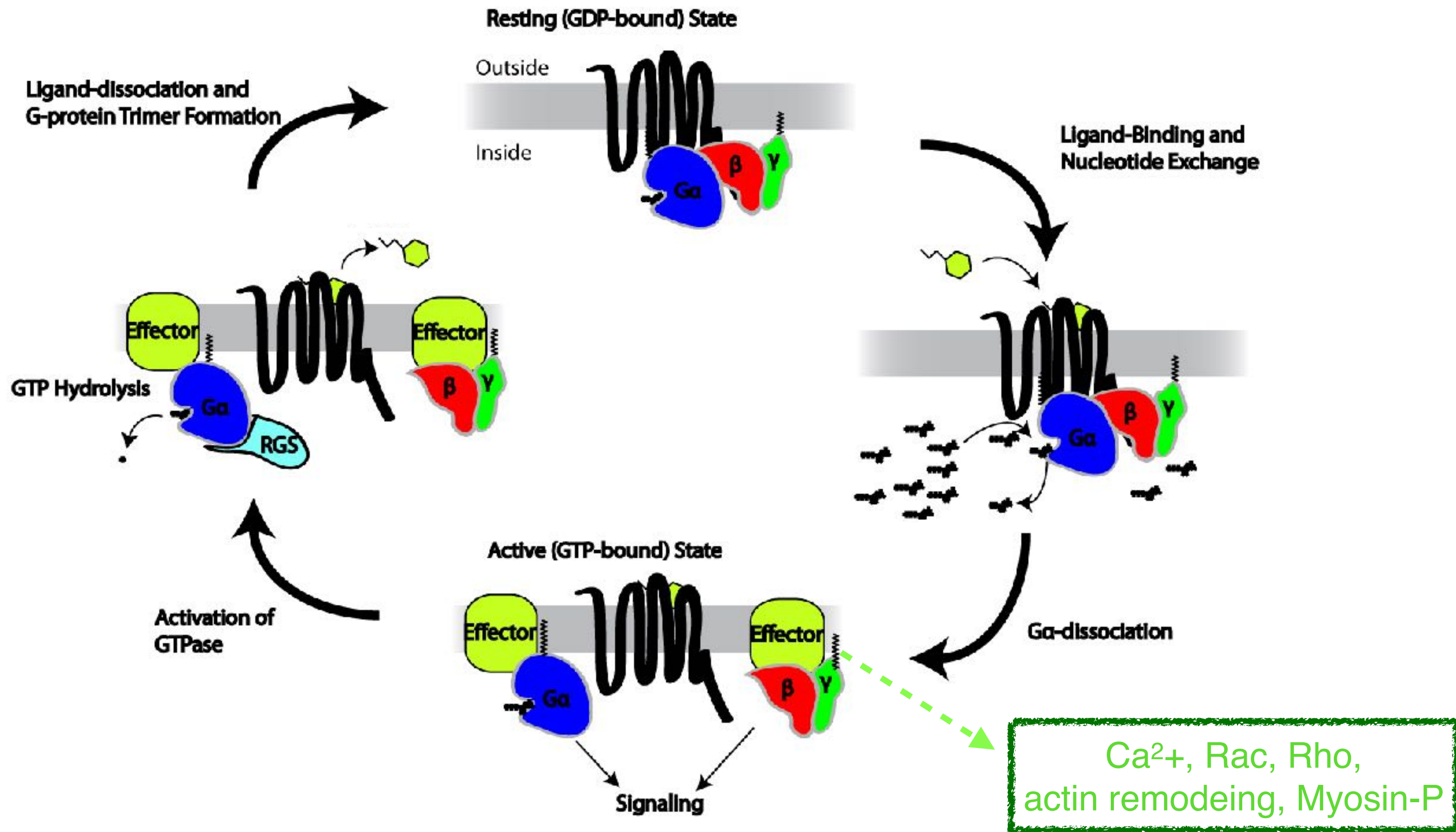
What sort of protein/
function might these
three genes encode?

Chemokine signaling

- **chemotactic cytokines**
- Receptor = CXCR, Ligand = CXCL
- 7-pass trans-membrane receptor
- G-protein coupled
- Discovered in immune cells...

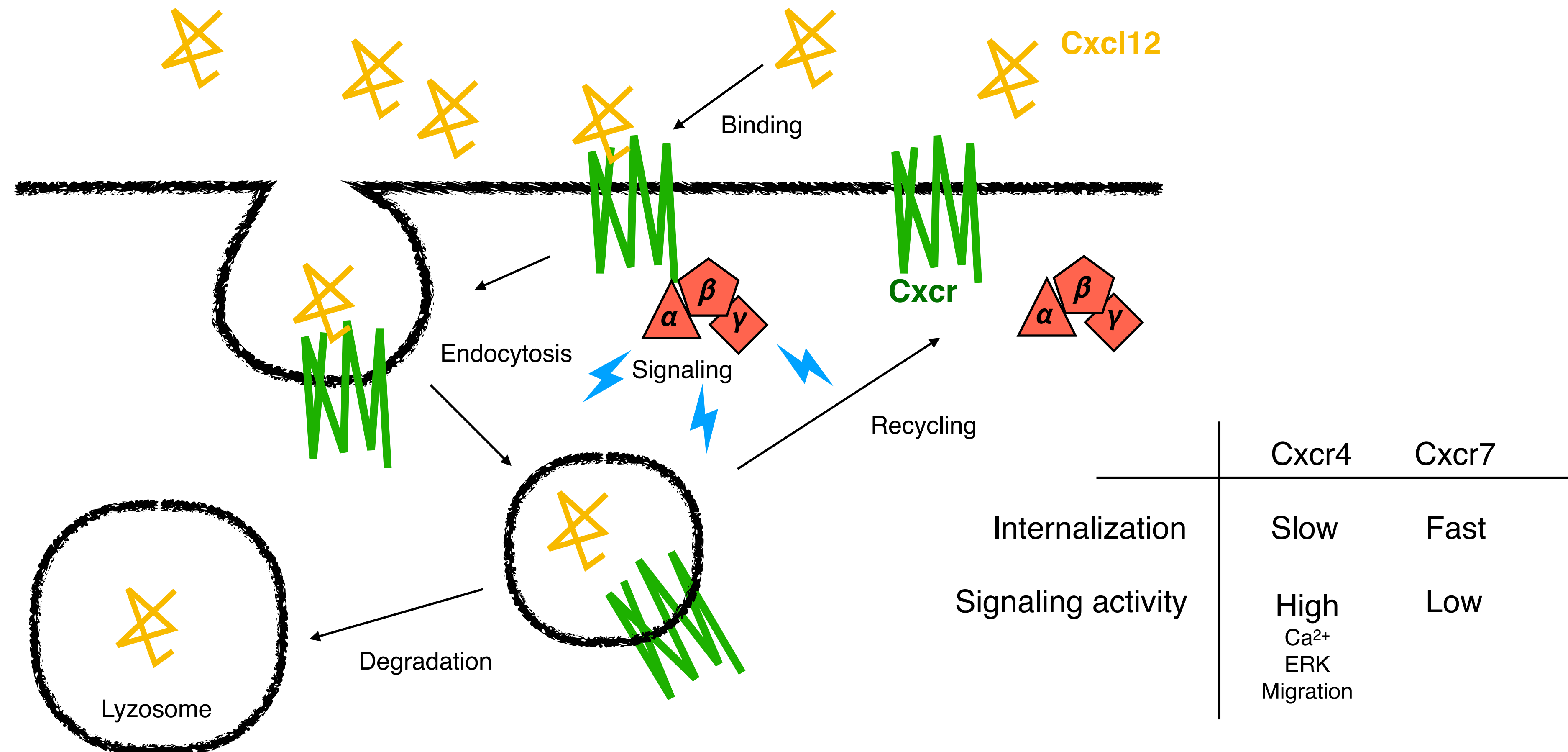


Chemokine signaling cycle



	Plasma Membrane		Gα-subunit (Palmitoylated)		Guanosine diphosphate (GDP)		Agonist
	GPCR		Gβγ-subunit (GPI-anchored)		Guanosine triphosphate (GTP)		Regulator of G-protein Signaling
					PI (Inorganic Phosphate)		Effector (Enzymes, etc)

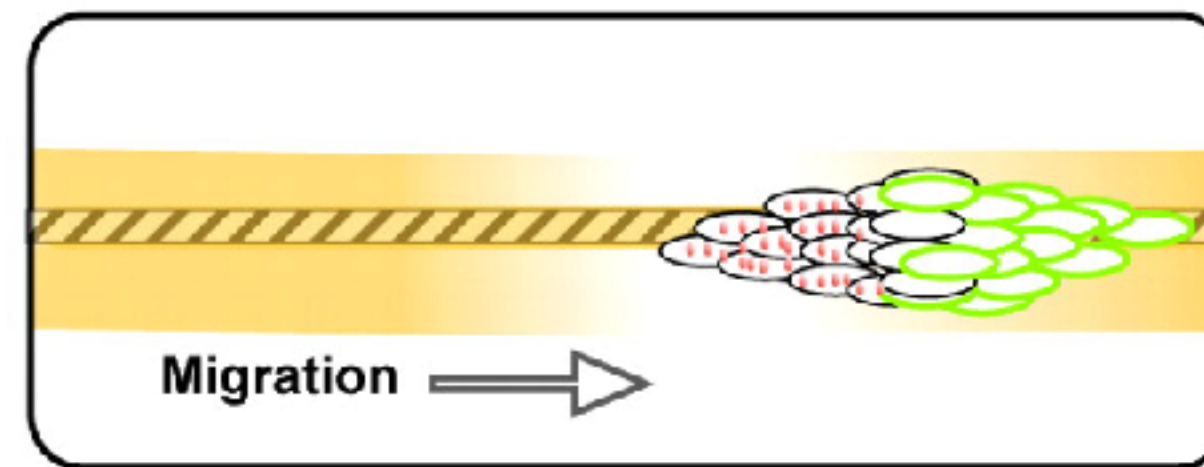
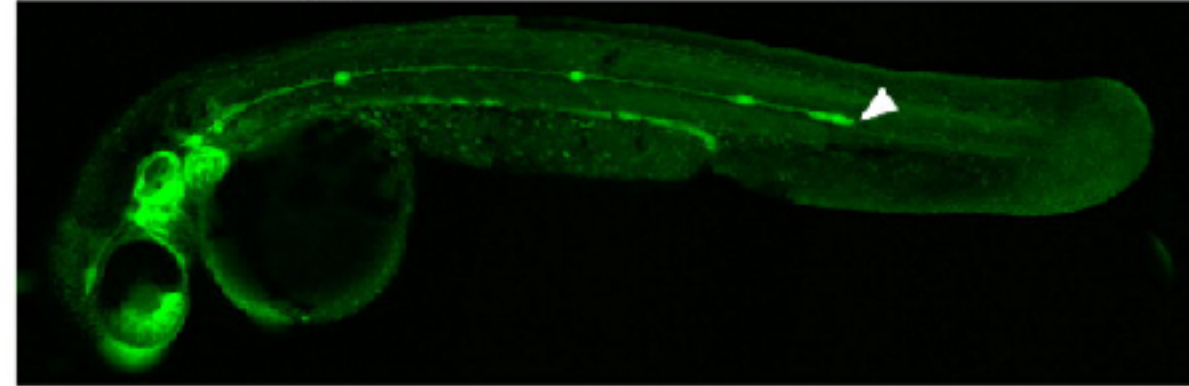
Differential internalisation and activity of Cxcr4 and Cxcr7



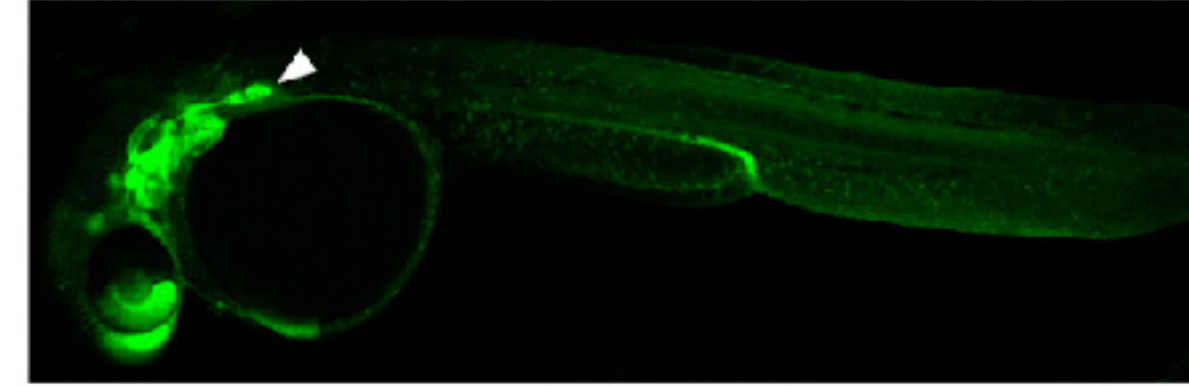
In multiple cell types, Cxcr7 is a “scavenger receptor”

Model: the *primordium* generates the *cxcl12* gradient

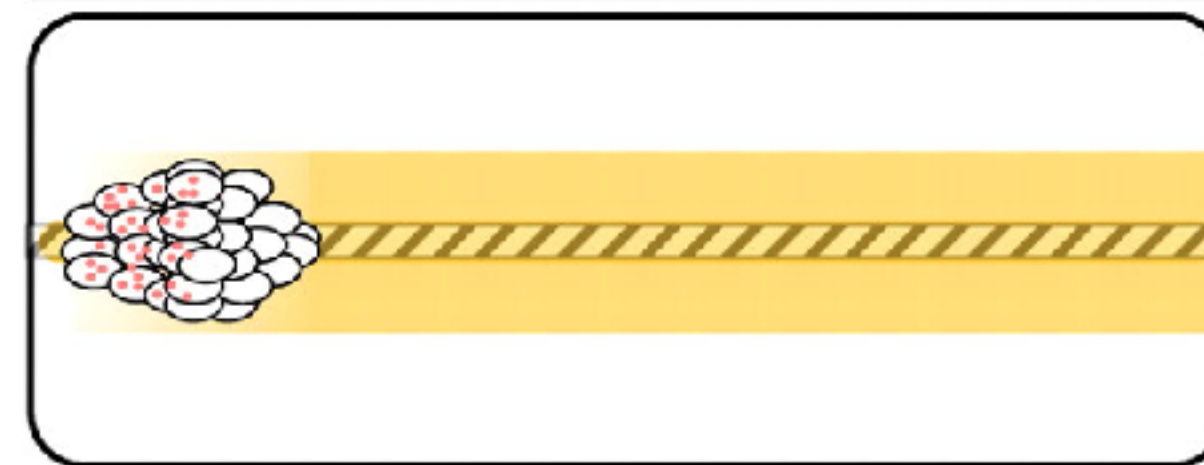
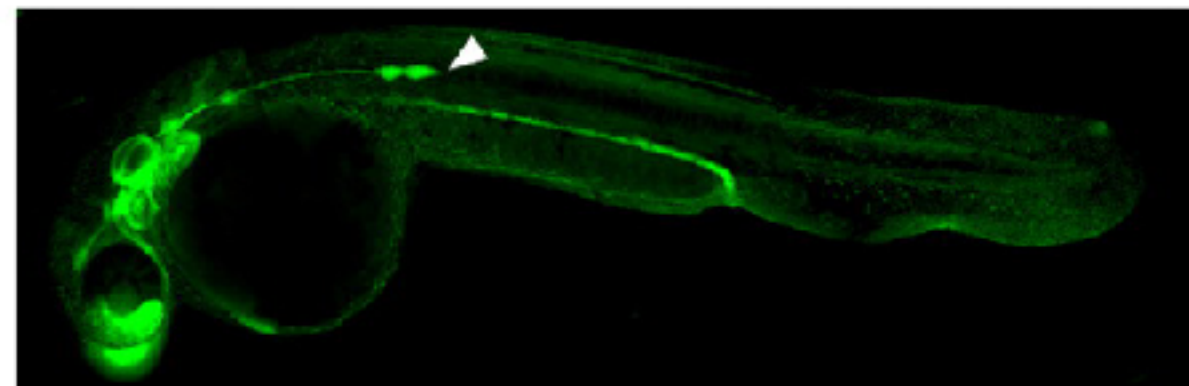
A Wild type



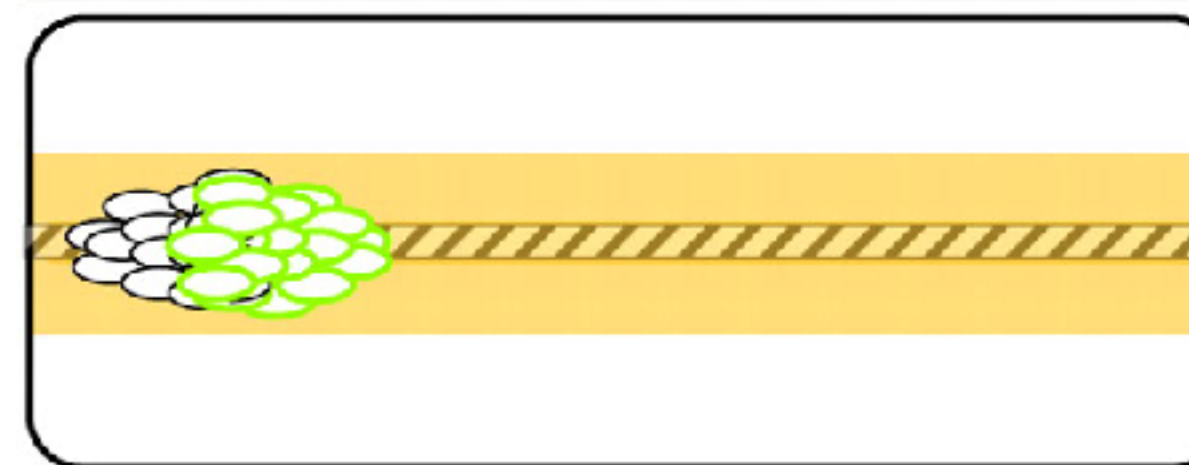
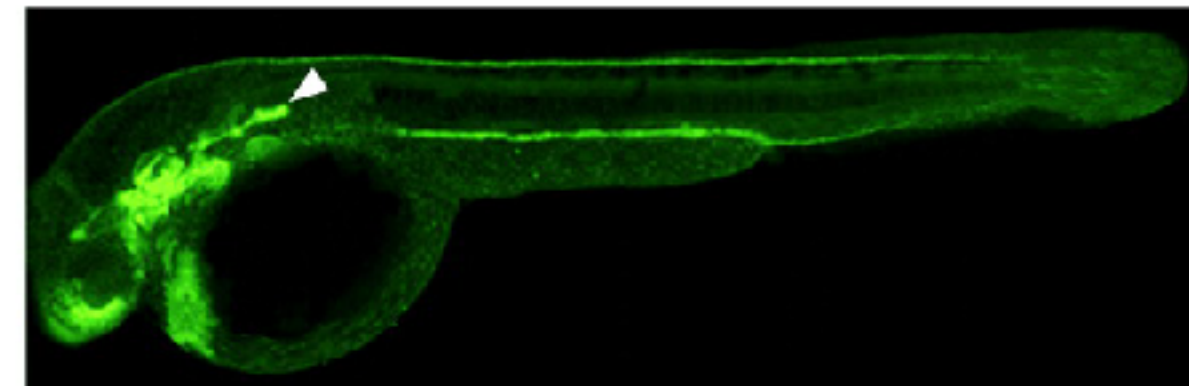
B *cxcl12a*^{-/-}



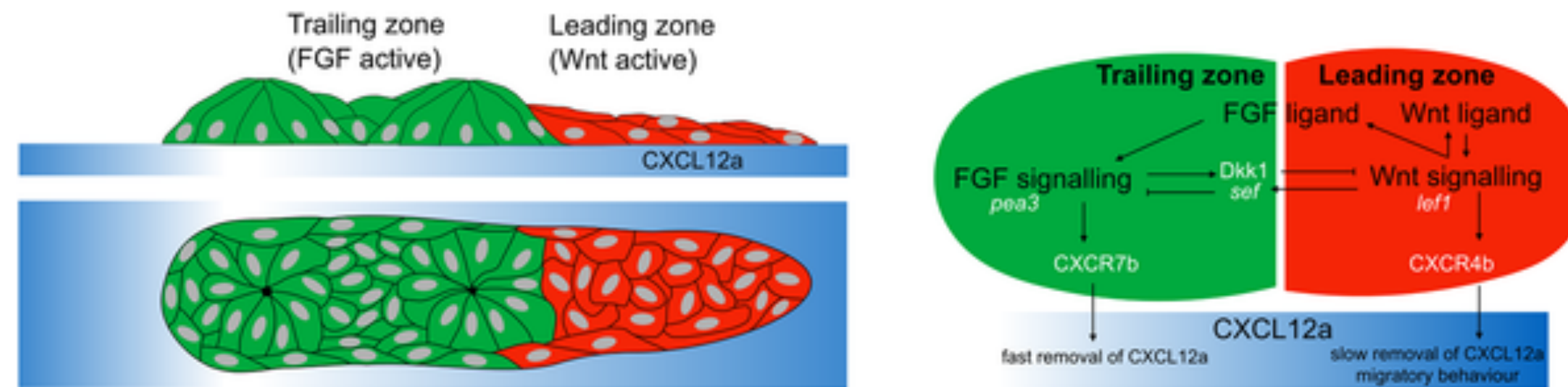
C *cxcr4b*^{-/-}



D *cxcr7b* MO



Bistable Wnt and FGF signaling defines leading and trailing zones & activates CXCRs

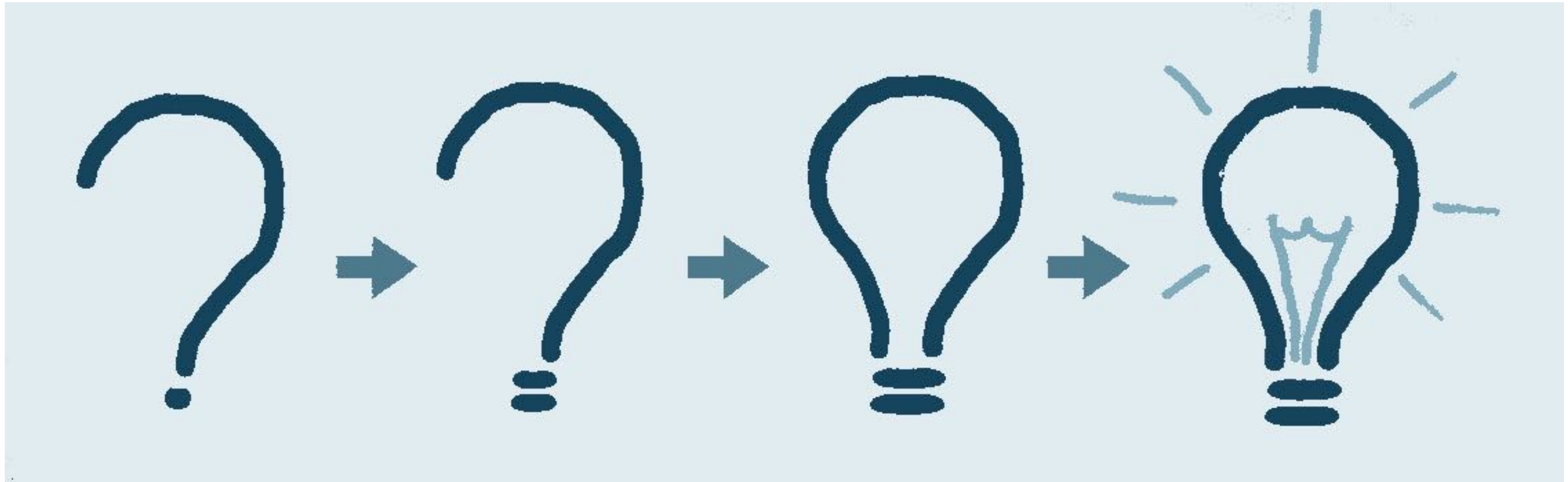


- How is initial Wnt and FGF symmetry broken?
- What are the adhesion systems?
- How is proliferation balanced with rosette deposition?
- What marks the drop-off points?

Summary

- Cell migration results from motility and polarization
- Multiple structures are formed from migratory cells
- Chemokine signaling can direct long-range migration
- Individual and collective migration
 - * We didn't cover this, but also critical in wound healing and metastasis in cancer...
- Collective migration requires internal self-organization

Questions?





Big questions



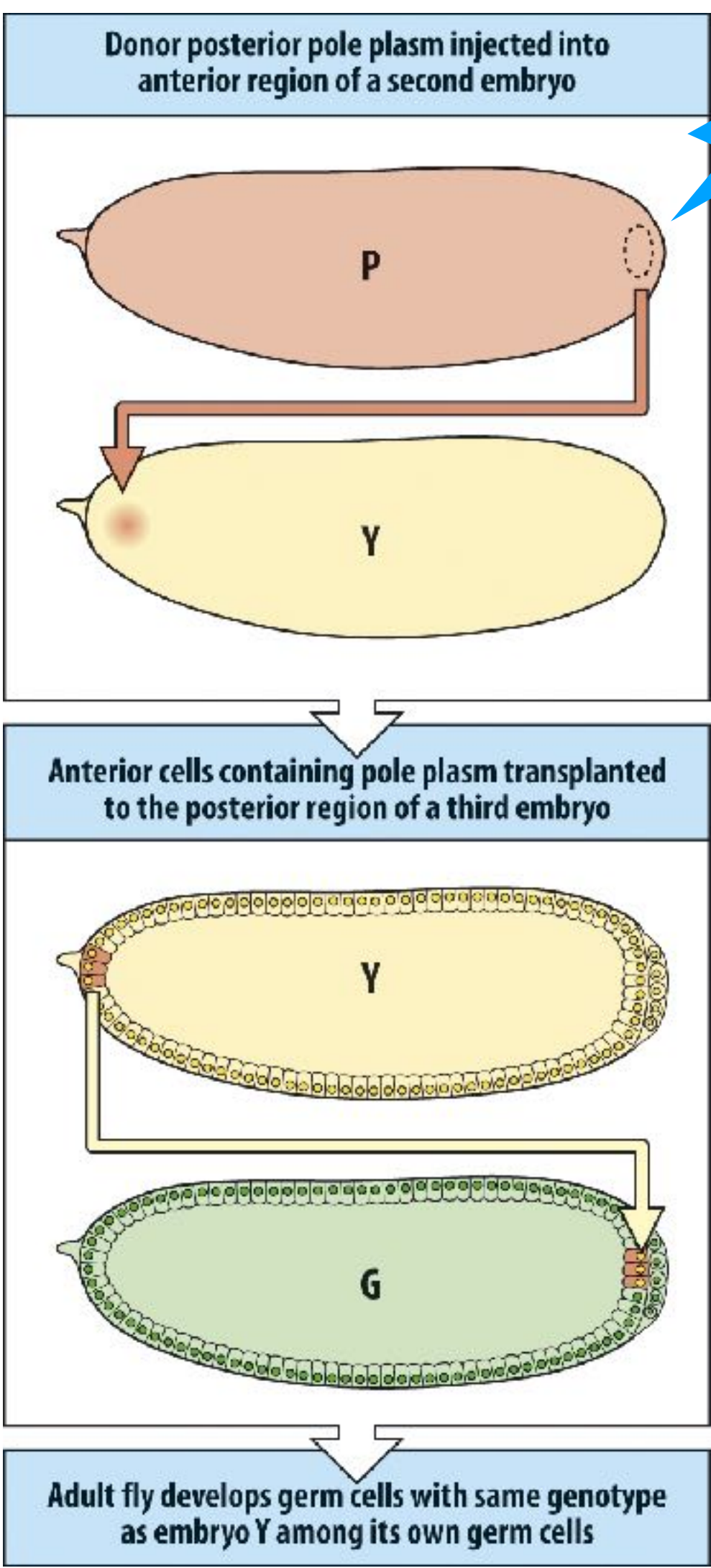
Germ cells - setting aside the next generation

- Specified outside the gonad
- Give rise to gametes - sperm and eggs
- Three roles:
 - Preservation of genetic integrity
 - Generation of genetic diversity (meiosis)
 - Transmission of genetic information
- Maintain pluripotency (immortal...)
- Sperm vs. egg determined by gonad

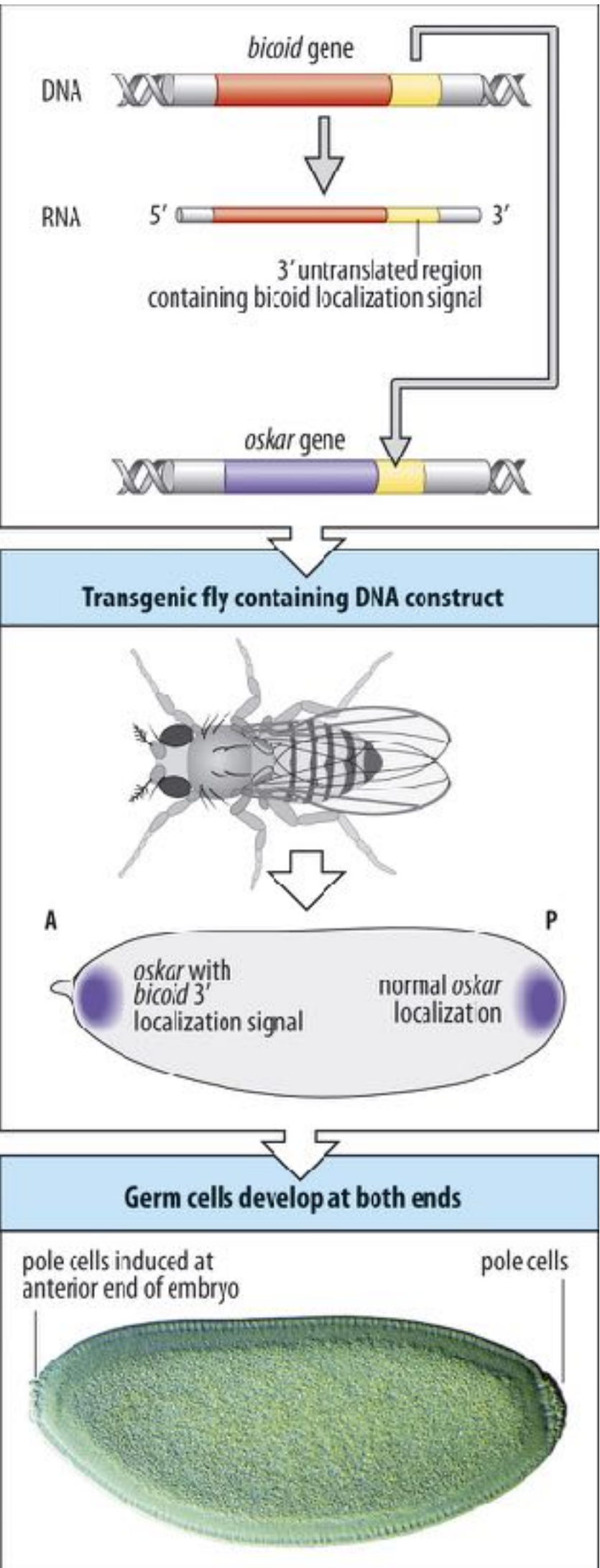
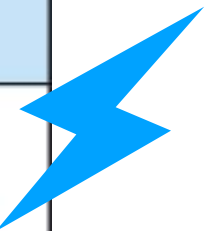
Case I. Maternal determination of germ cells - germ plasm

- Localised in the early embryo
- RNA-binding proteins Vasa, Nanos, Oskar inhibit translation
- Cells that inherit germ plasm become germ cells
- May be distinct organelle...

Localised germ plasm - *D. melanogaster*



UV induced sterility

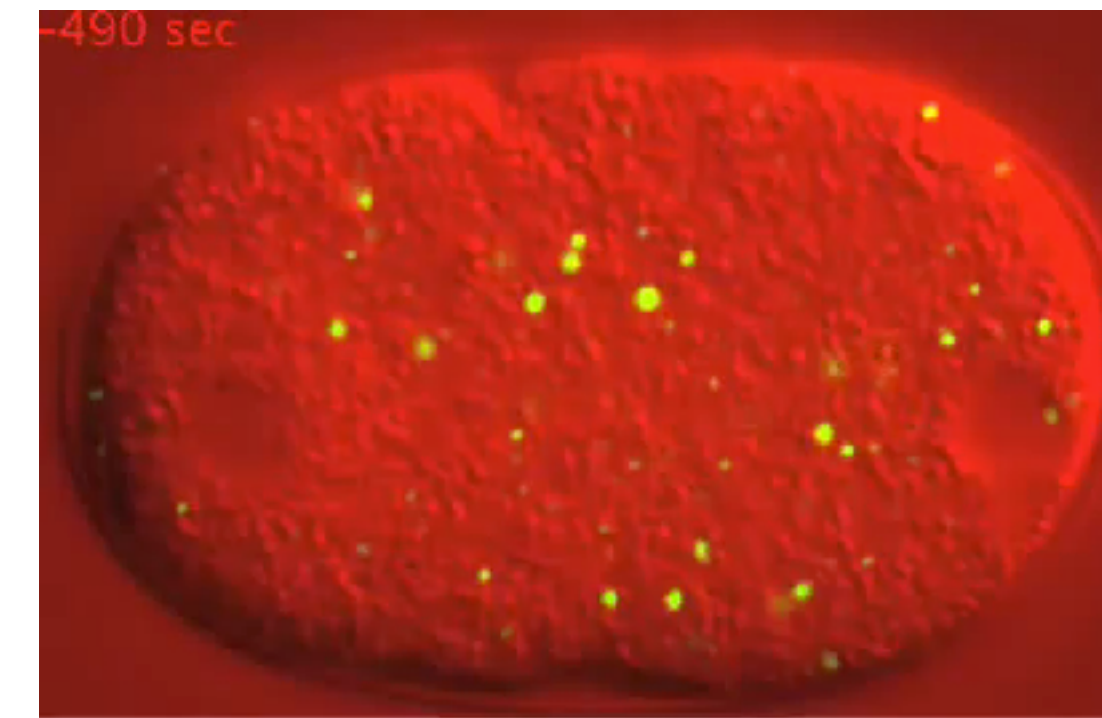
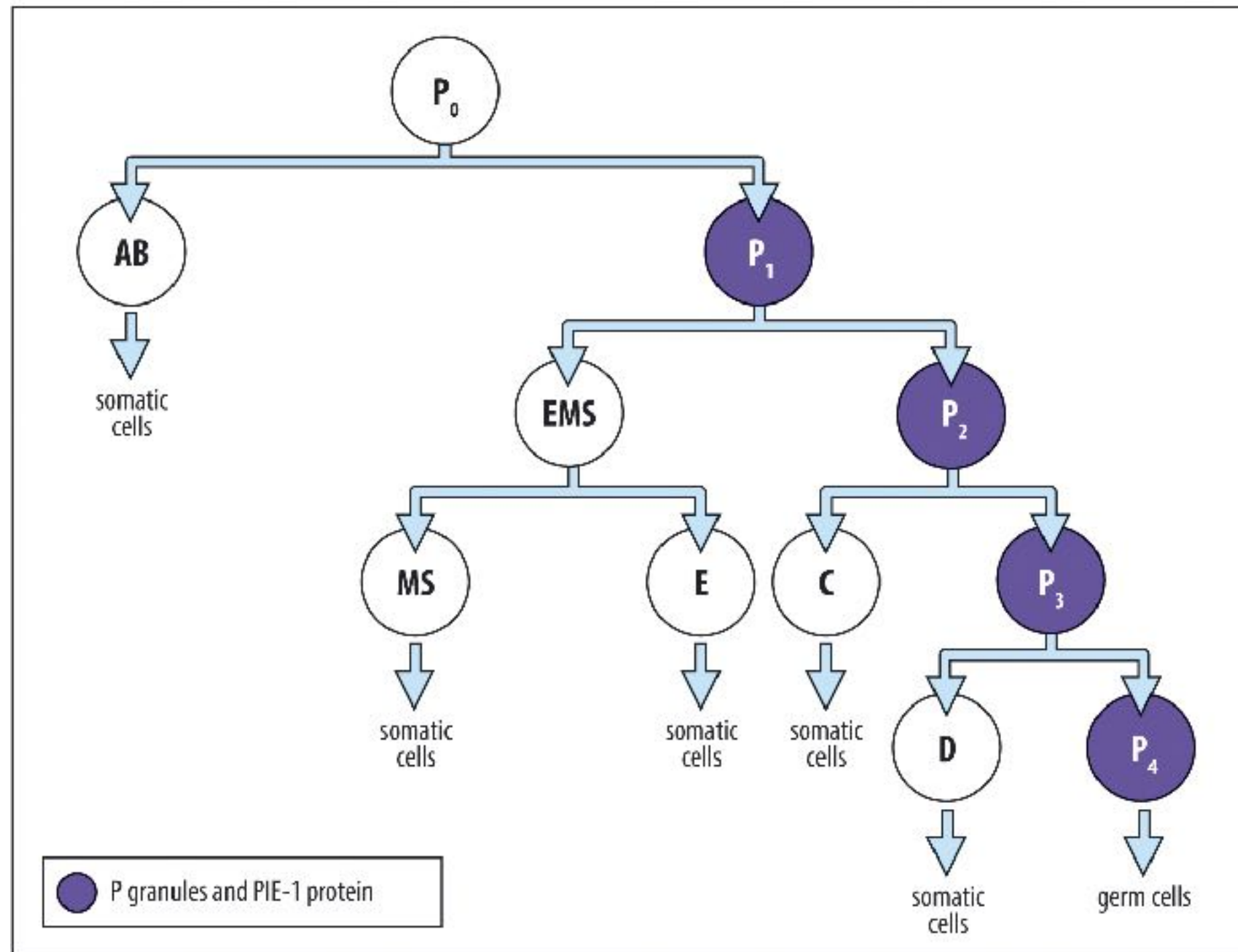
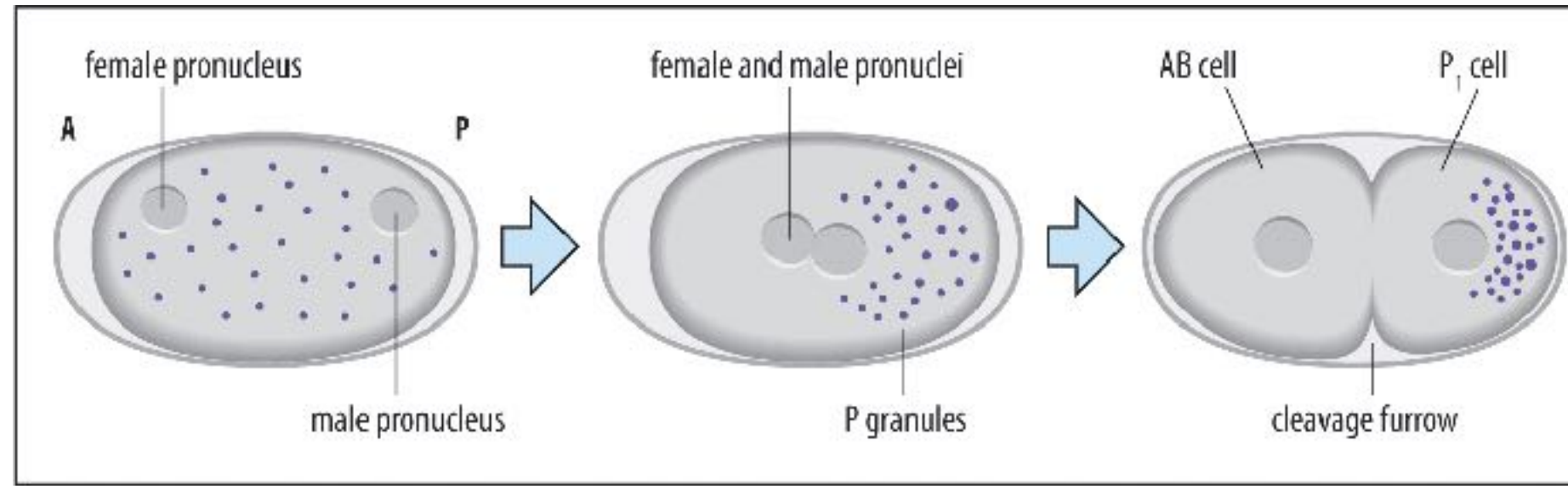


oskar flies are sterile

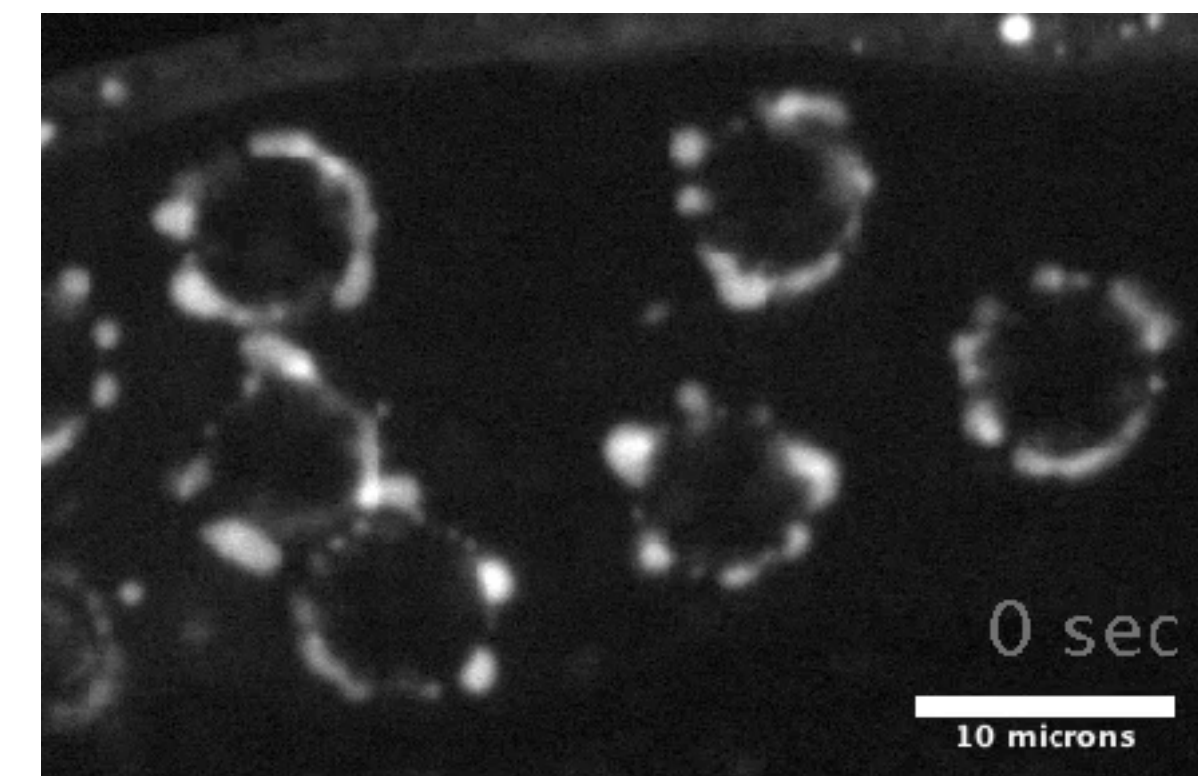
oskar necessary and sufficient for pole cells

Oskar RNA-binding protein nucleates pole granules

Localisation of germ plasm - *C. elegans*



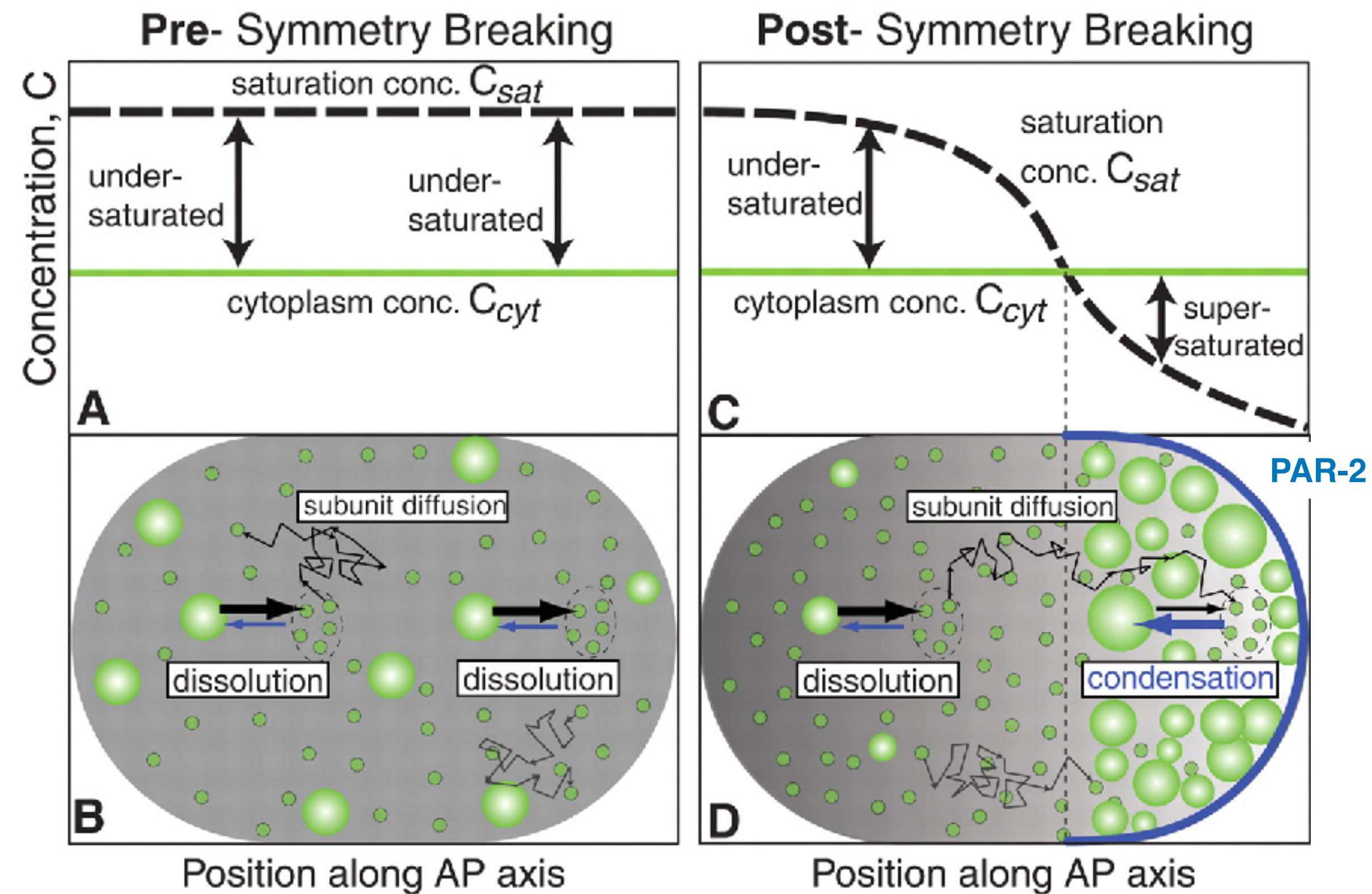
PGL-1-GFP



Syncytial germ cell nuclei in PGL-1-GFP adult

PIE-1 nuclear protein represses transcription - keeps cells silent during early somatic signaling

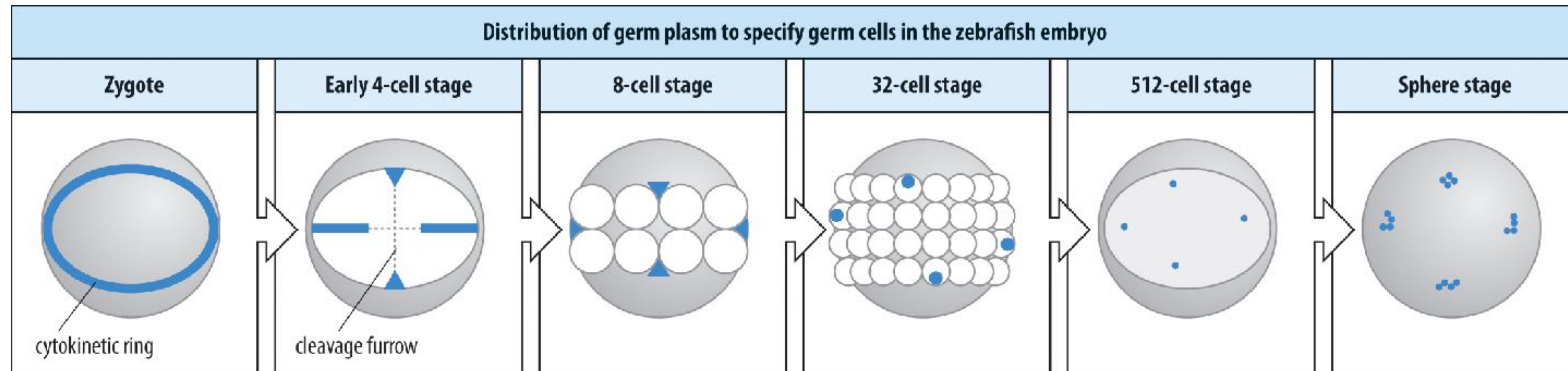
Non-membrane bound organelles: liquid-liquid phase separation



Properties

- RNA-protein, low complexity proteins
- Elevated local concentration
- Buffering of cytoplasm
- Regulation by e.g. phosphorylation

Localisation of germ plasm in zebrafish



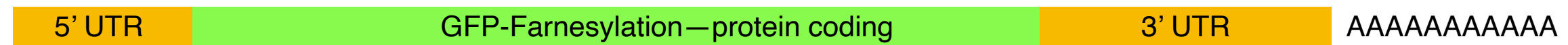
vasa mRNA



nanos mRNA

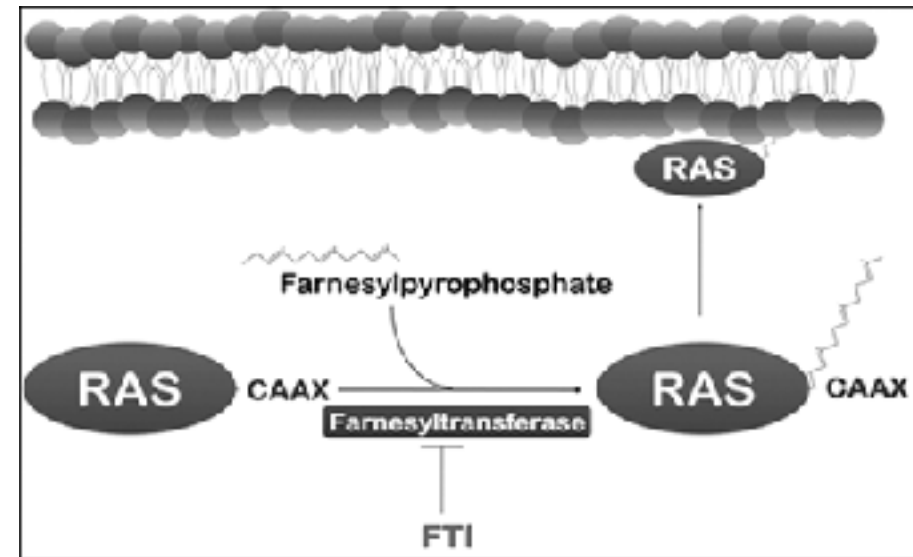


GFP-F-nos 3' UTR mRNA

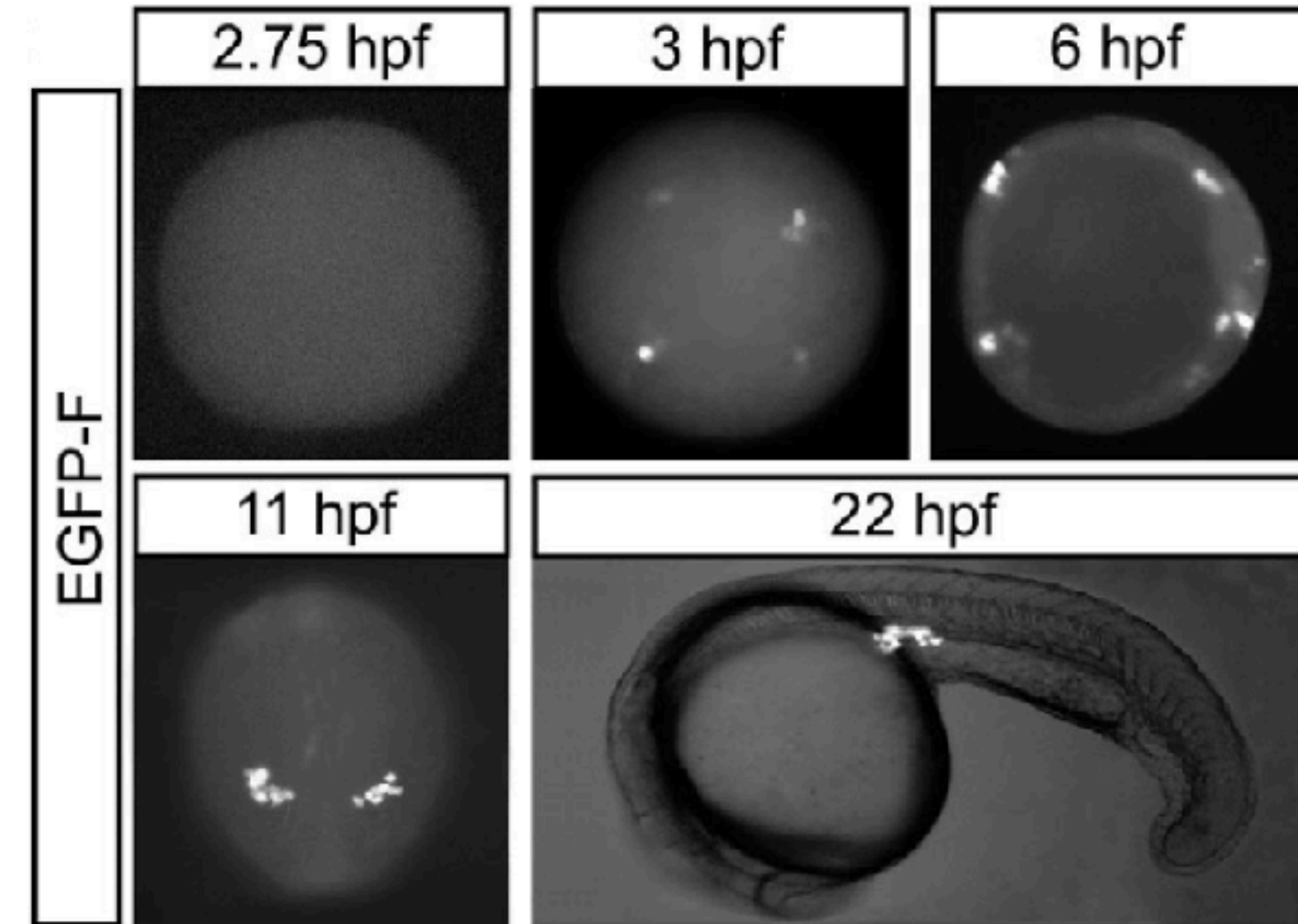
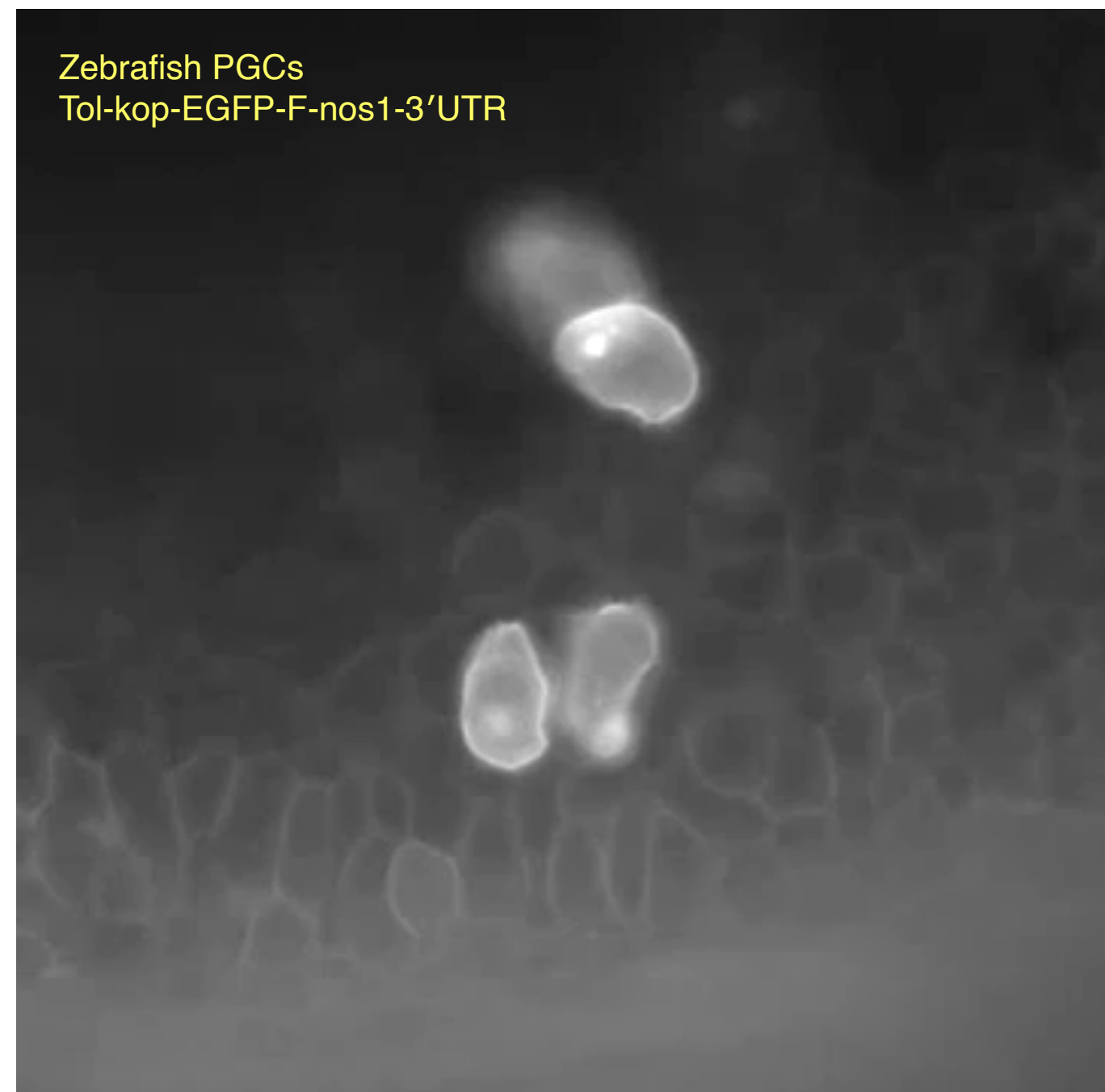


Labeling the PGCs in zebrafish

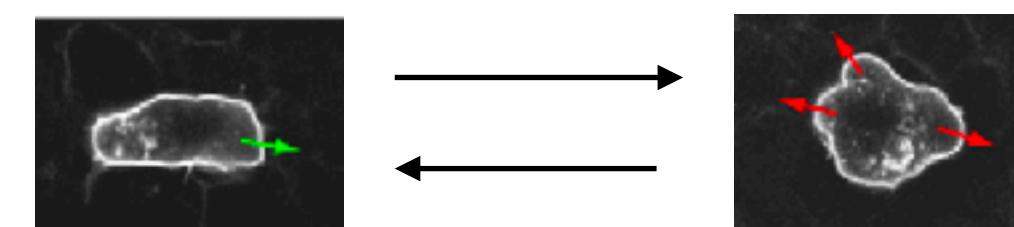
GFP-CAAX—protein coding 3' UTR AAAAAAAAAA



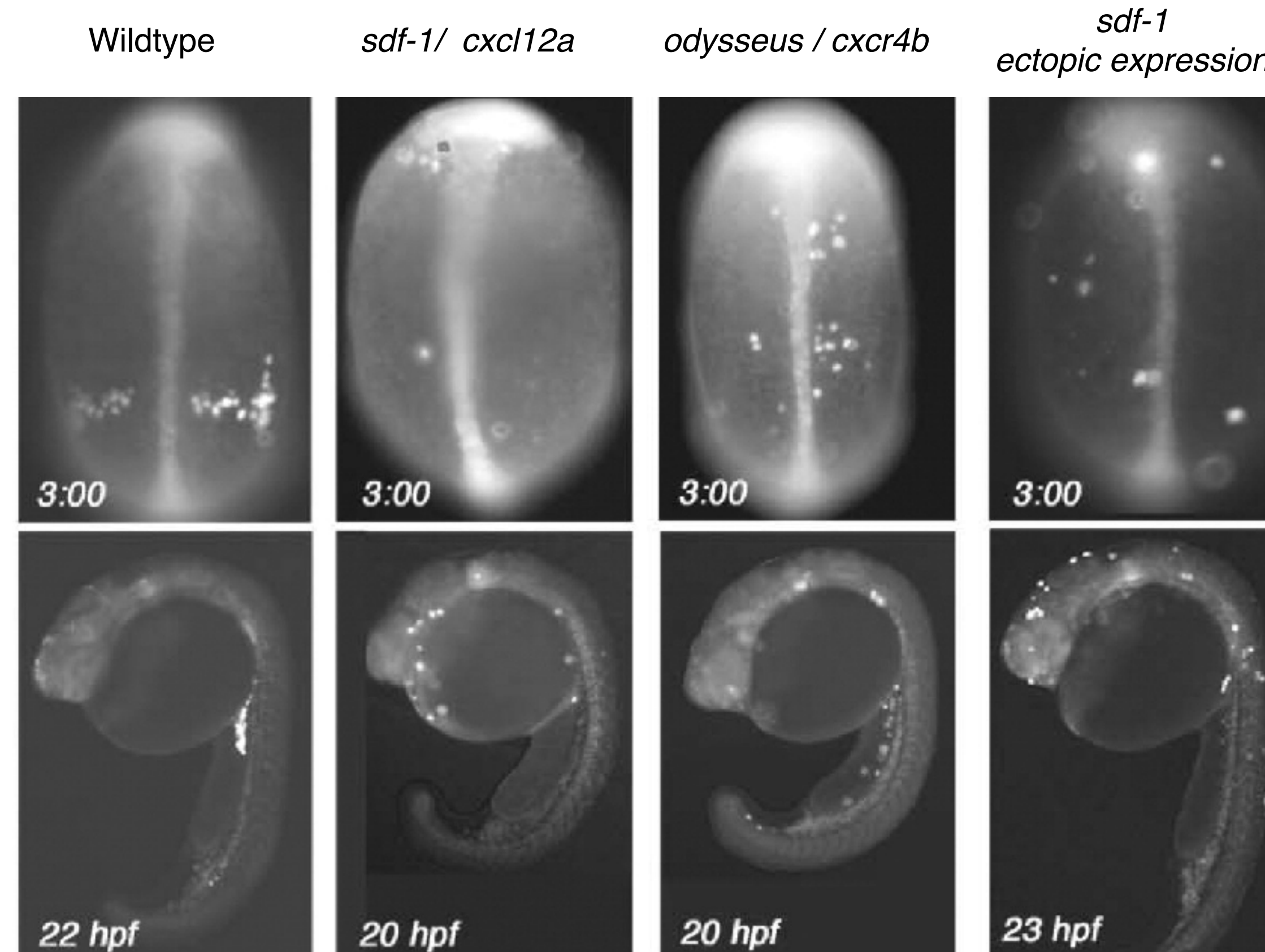
GFP-F-nos 3' UTR mRNA



- PGCs can polarize
- Move by stabilising bleb-based protrusions
- “Run and tumble”
- Require E-cadherin for adhesion
- ~30 PGCs arrive in the gonad region

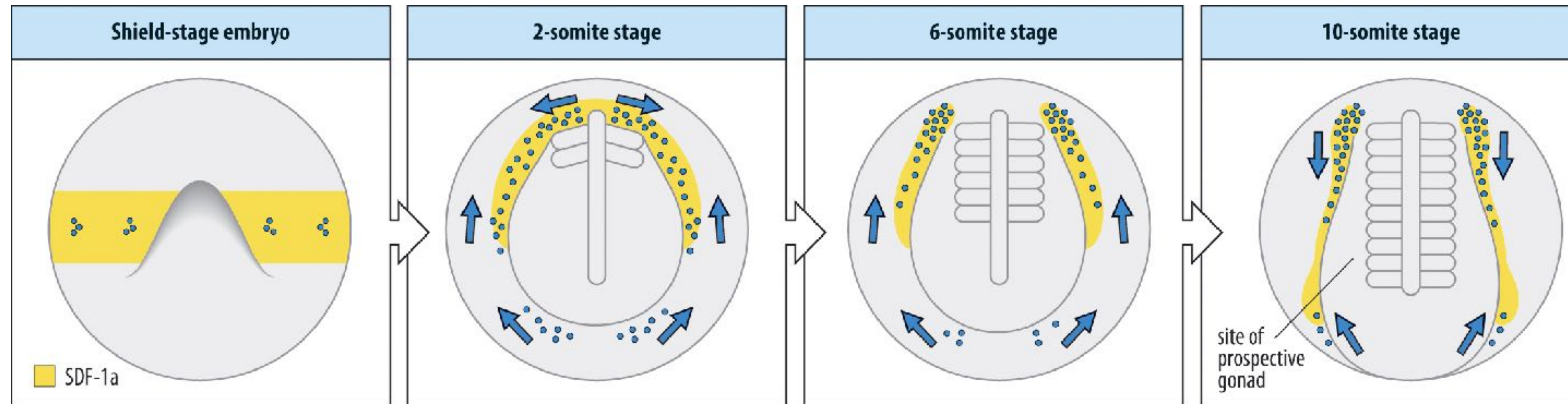


Mutants with ectopic PGCs

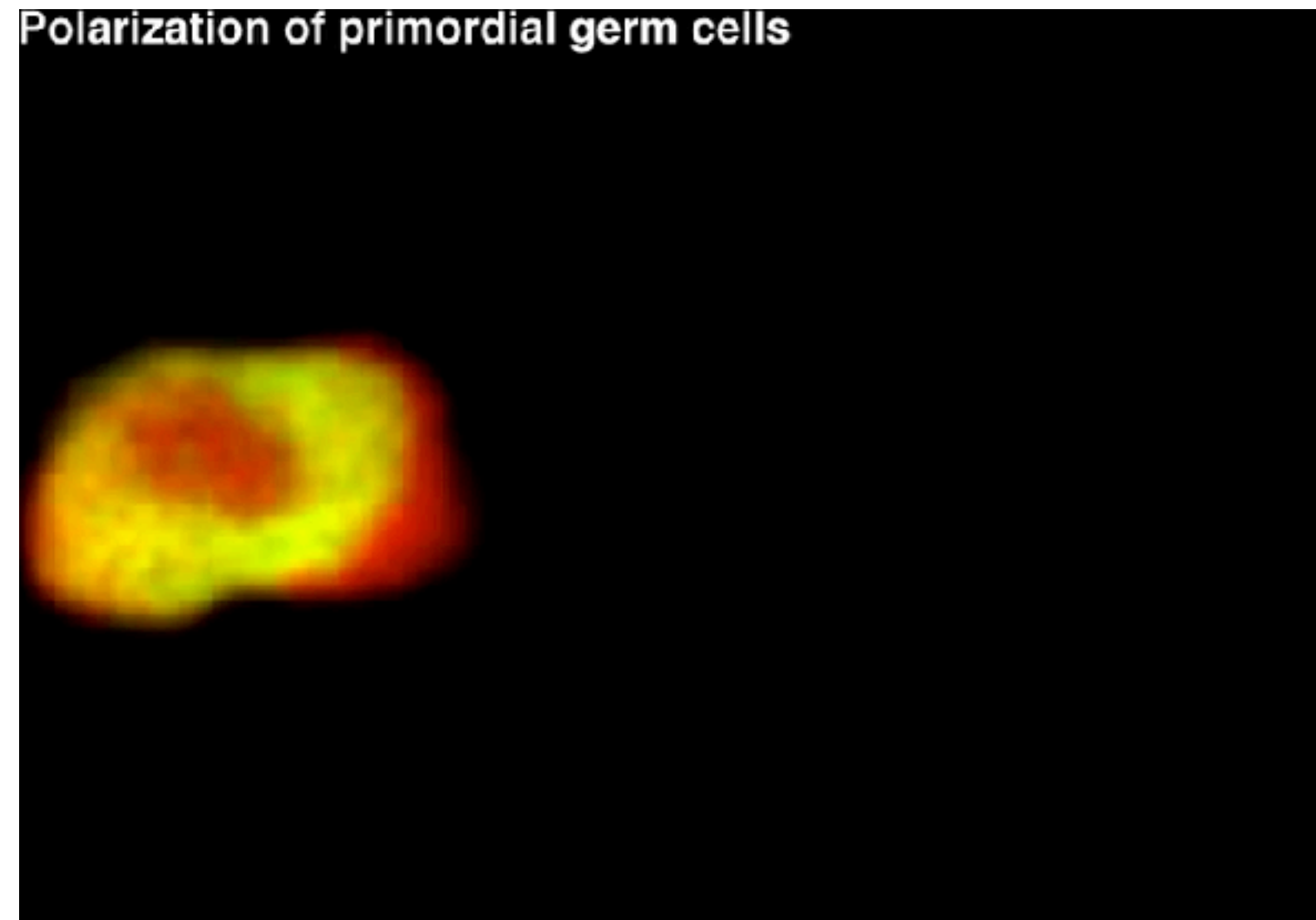


odysseus / cxcr4b expressed in PGCs

Finding the gonads: SDF-1 is attractive



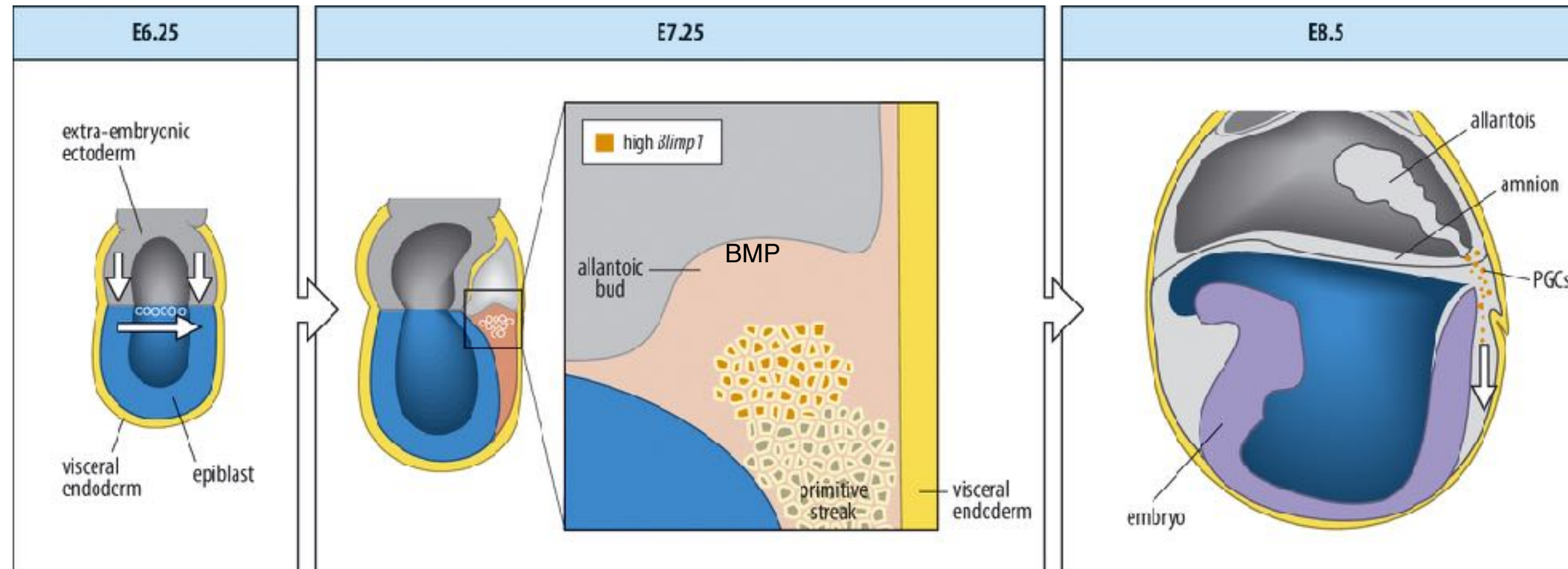
Polarization of primordial germ cells



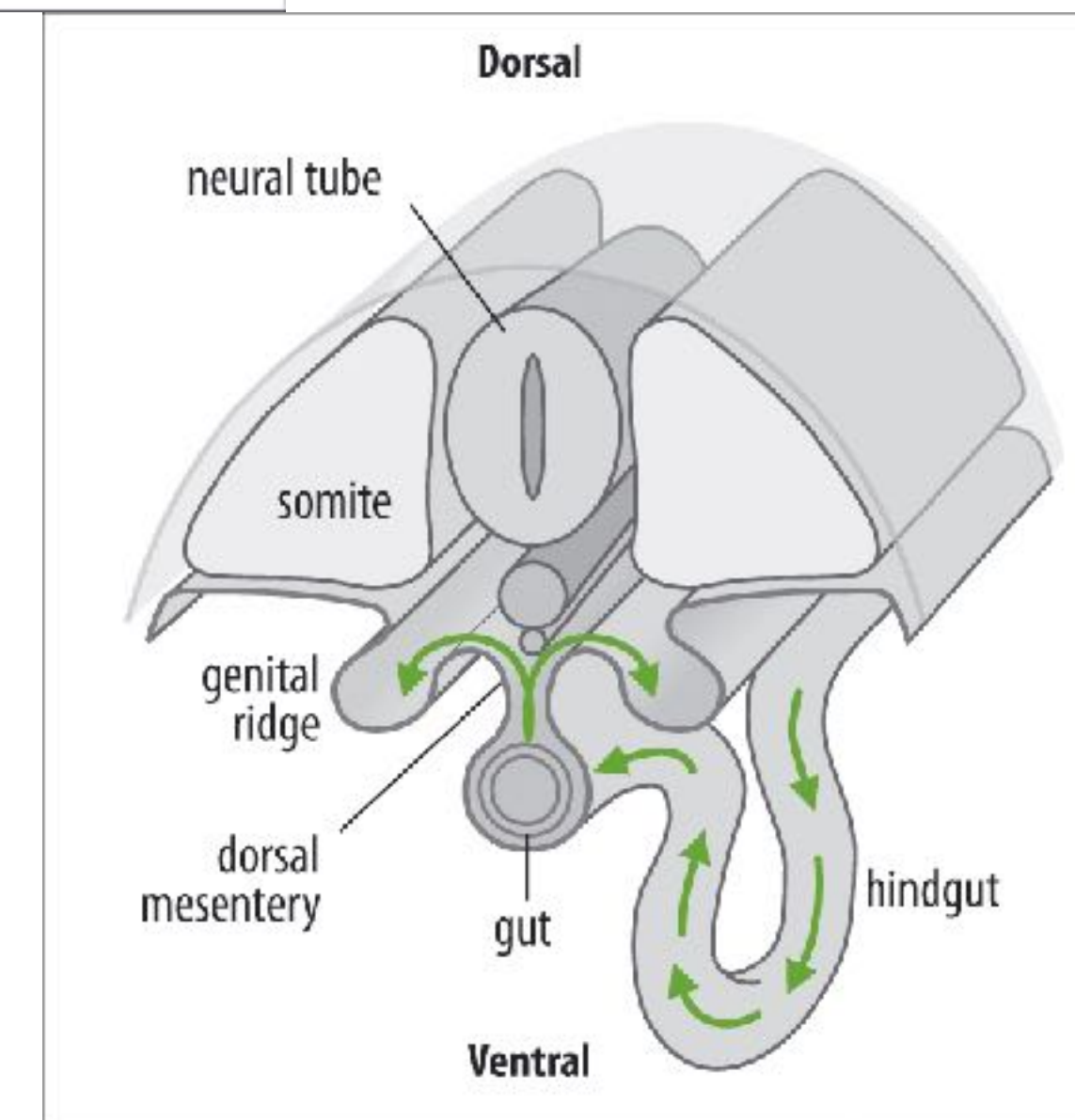
PGCs move towards SDF-1 expressing cells

Transplant of SDF-1-expressing cells

Case 2 - somatic specification mouse PGCs

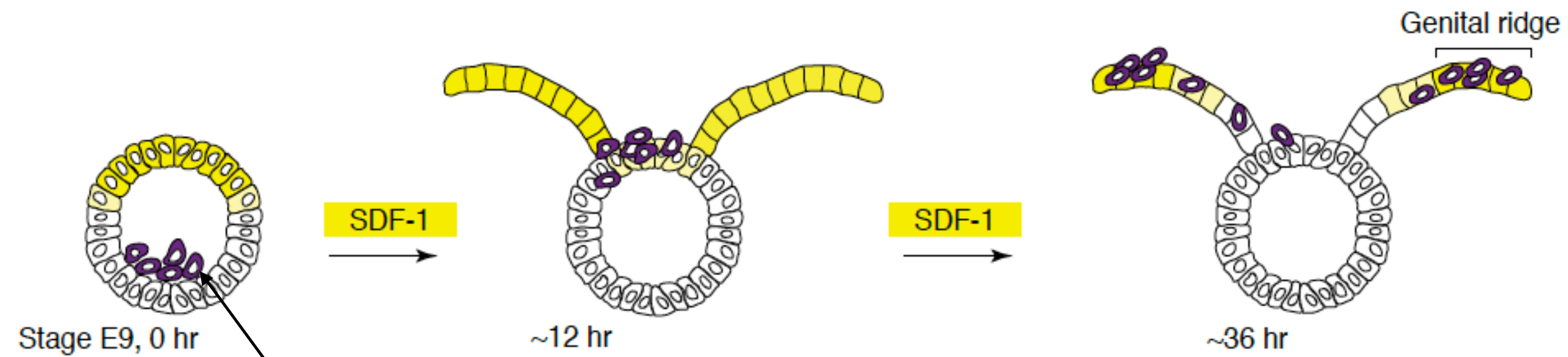


- ~ 40 Blimp1-positive cells at posterior of primitive streak
- Induced by BMP signals
- Migrate into hindgut
- Exit gut into genital ridge



Conservation of guidance mechanisms

Mouse

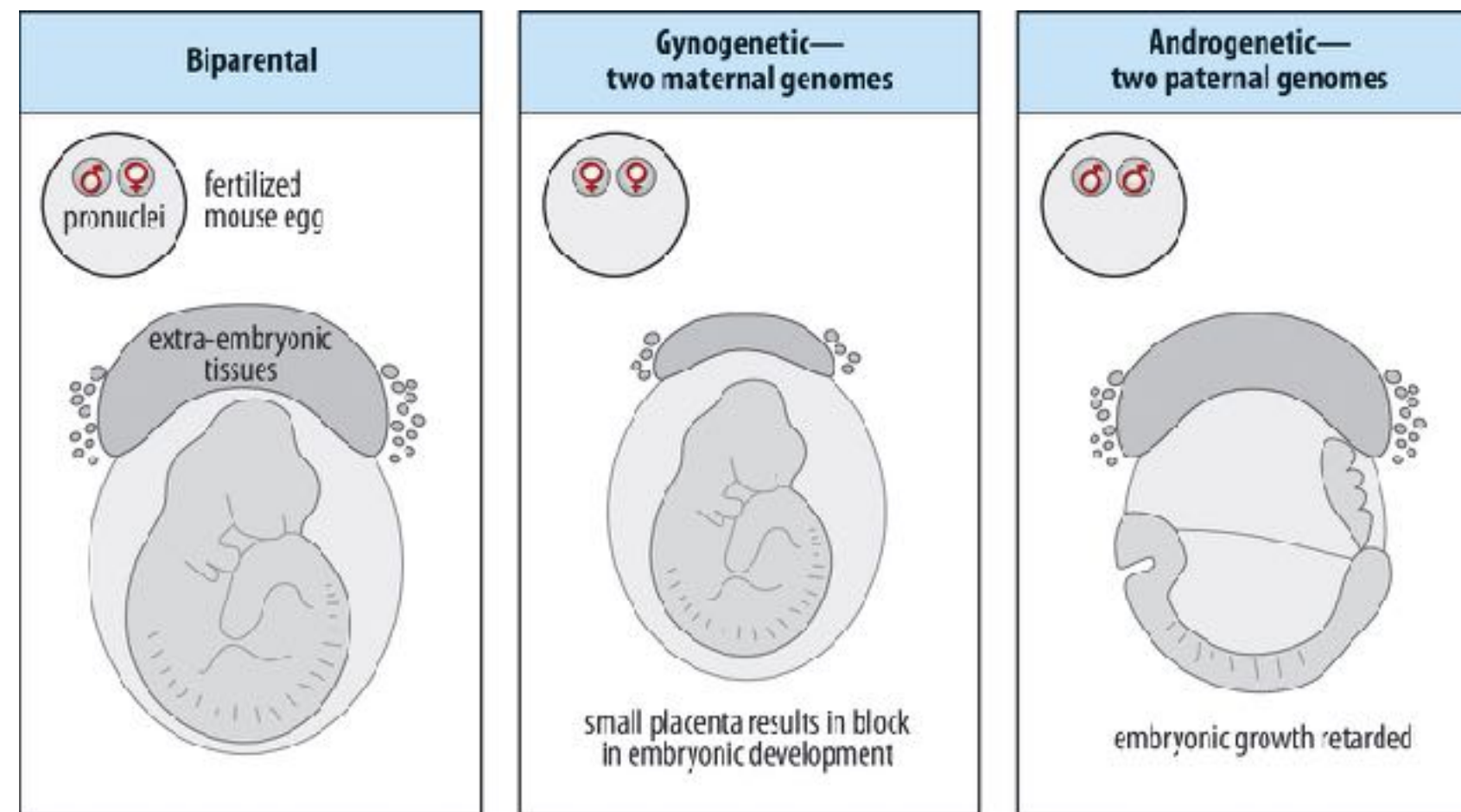


Cxcr4 in PGCs

Zebrafish

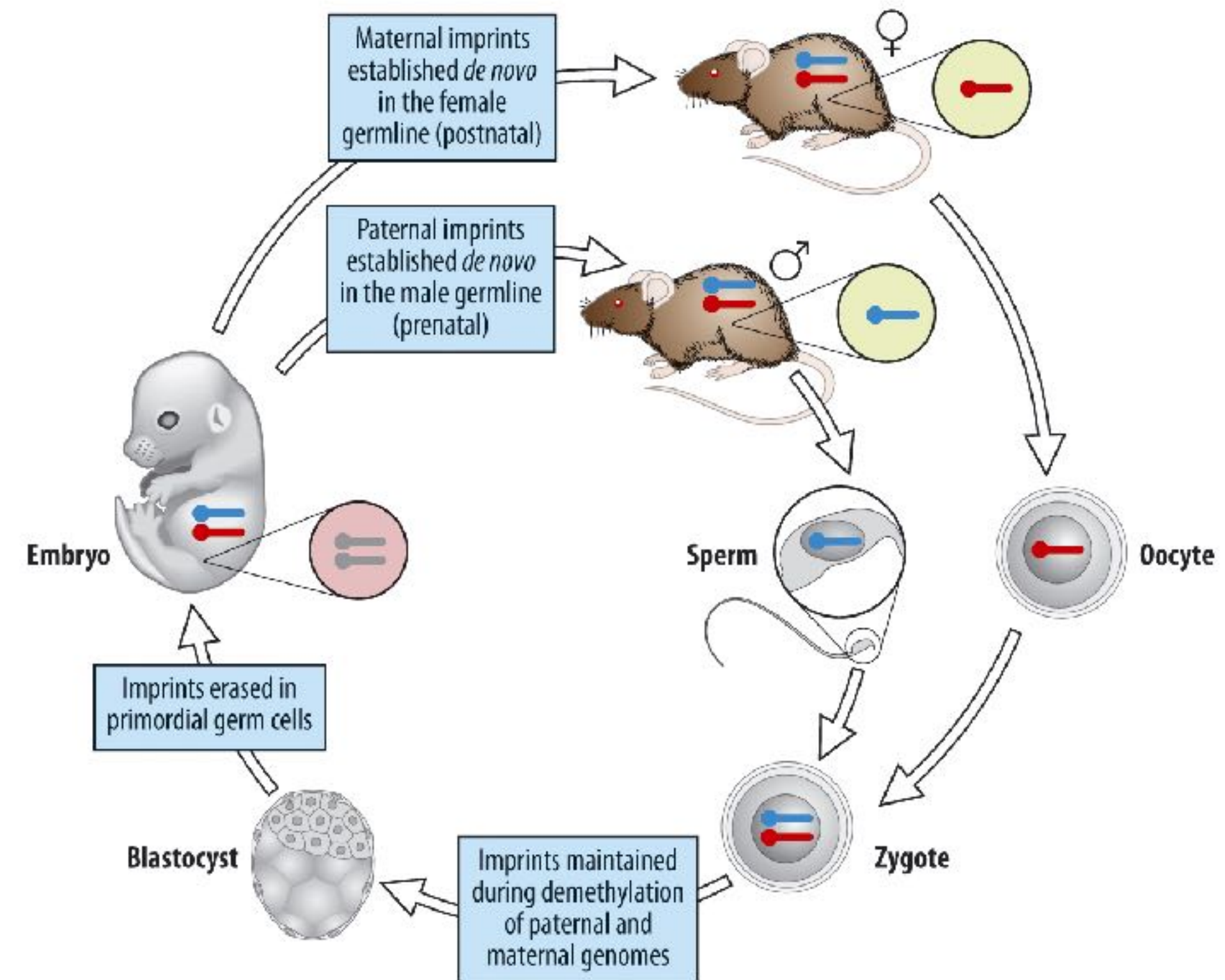


Mammalian gene imprinting - life cycle



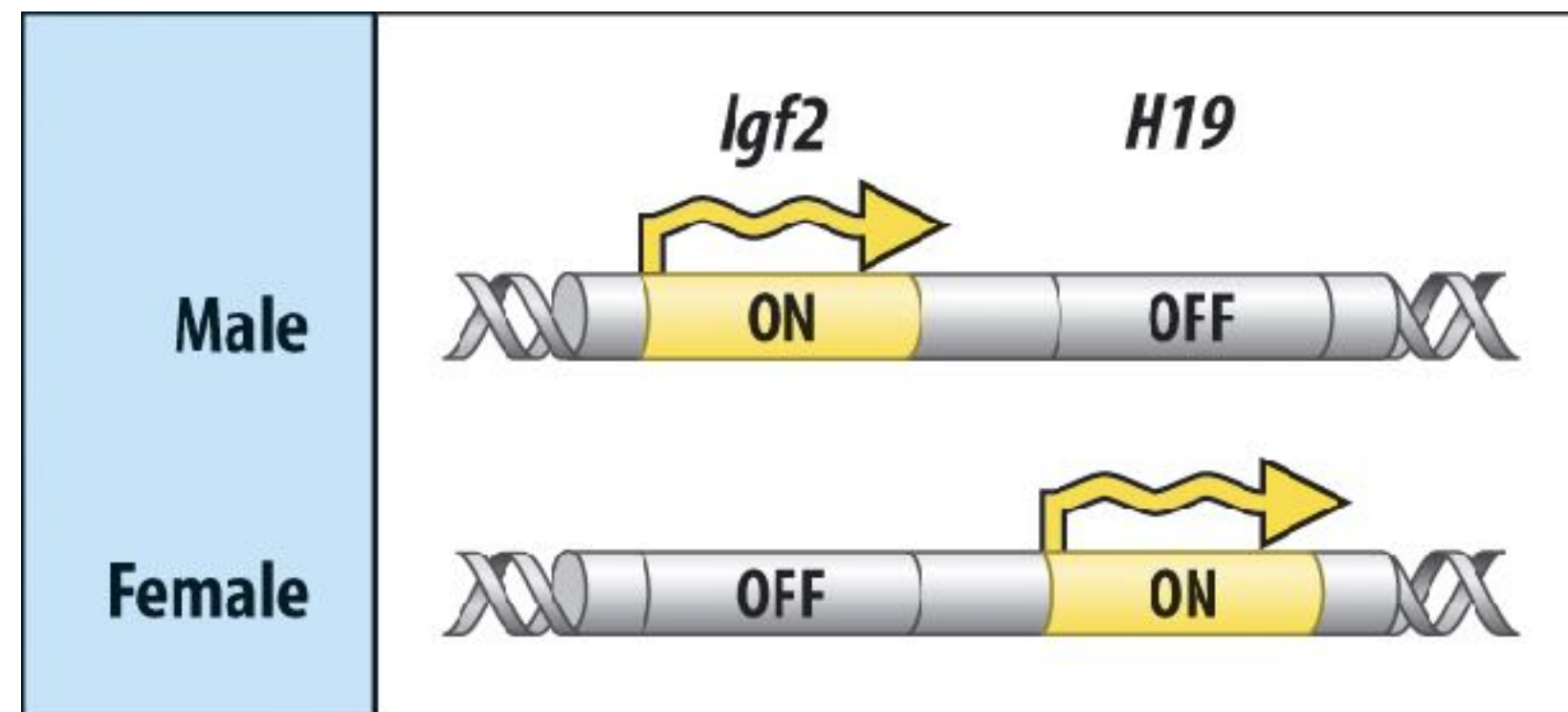
Male and female genomes are not equivalently expressed

A memory of being in sperm or egg



Imprinted genes are “re-set” each generation

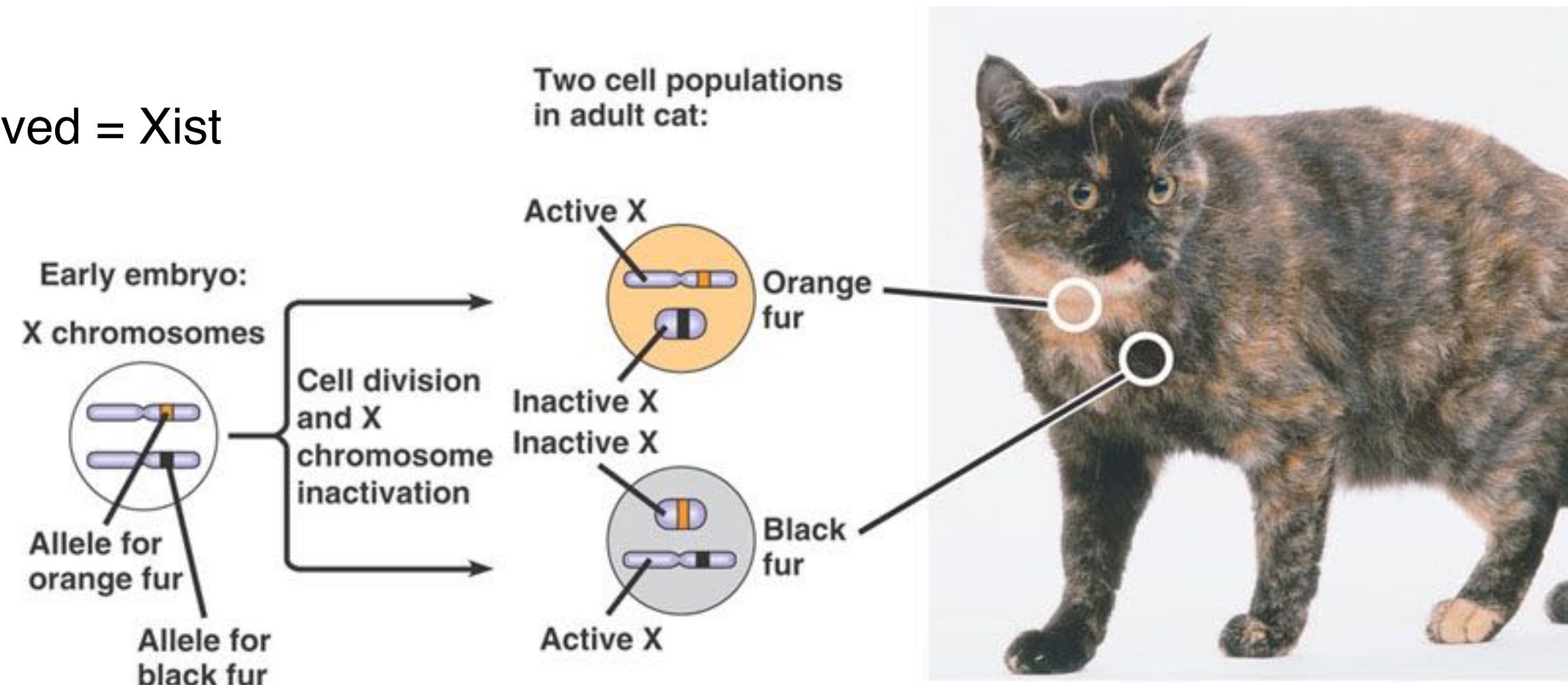
Mammalian gene imprinting - genetics



- 80+ imprinted genes known
- DNA methylation involved
- Many are non-coding RNAs - H19
- IGF2 paternal mutant embryos are small
- Beckwith Wiedemann babies are big
- Parental-conflict theory

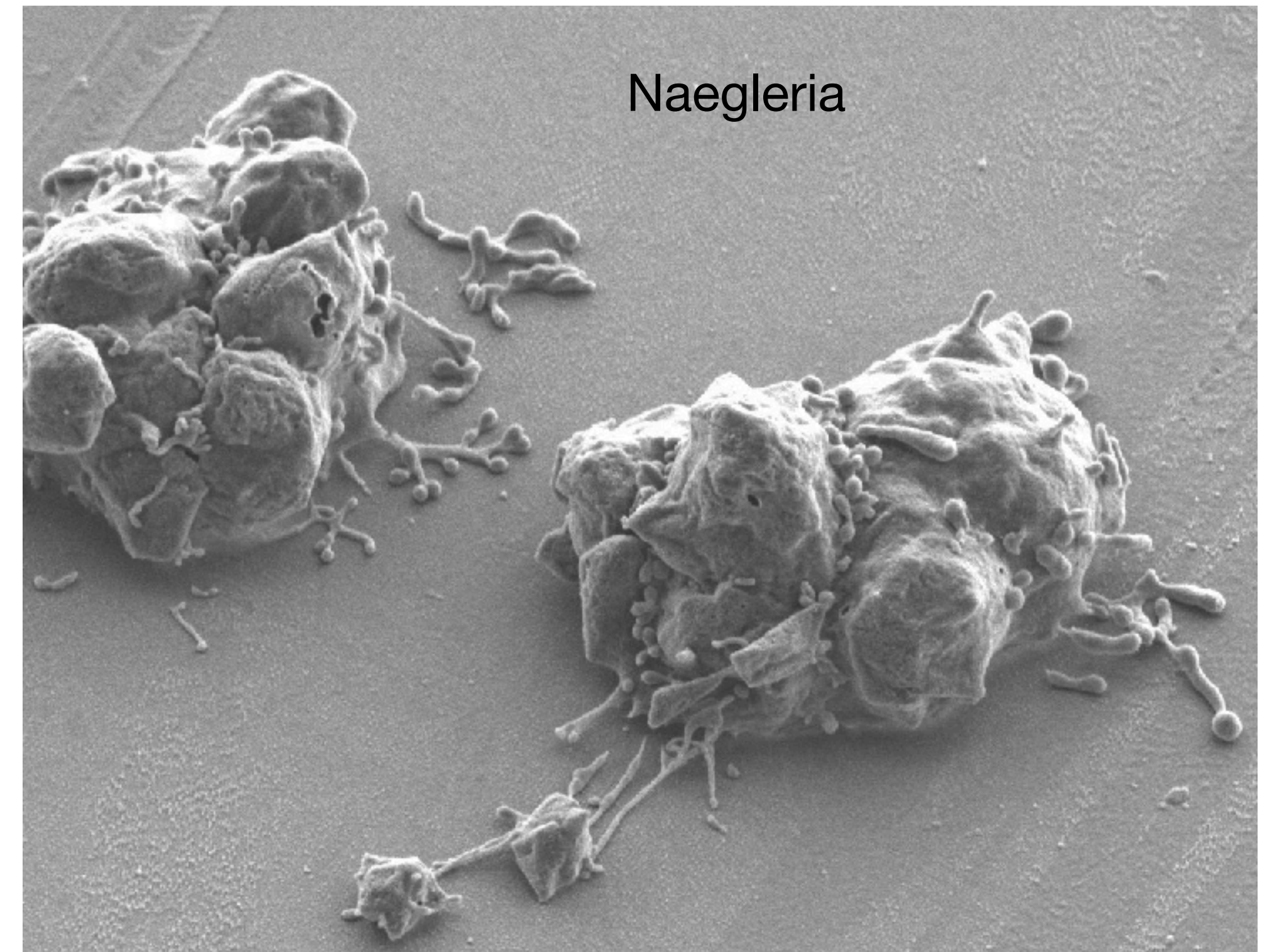
X-chromosome inactivation

- Early, stable
- Noncoding RNA involved = Xist



Summary

- Germ plasm specifies germ cells in many organisms
- RNA binding proteins may have structural and translational function
- CXCR signaling system drives long-range migration of germ cells to gonads
- Germ cells remain bipotent, and harbor pluripotency
- Germ cells wipe most epigenetics



Questions?

