

ME 411 / ME 511

Biological Frameworks for Engineers

Class Organization

- Marita Rodriguez will give Wed lecture
- Homework 4 due on Fri

Class Organization

- *Tiny Workhorse Projects*

Motor Protein	Grad Student
Helicase	Kevin & Ye
Actin-Myosin	Nathan & Kateri
Clathrin	Scott & Spencer
B. Flagella	Brian & Wai
Kinesin-5	Jarrod & Mark

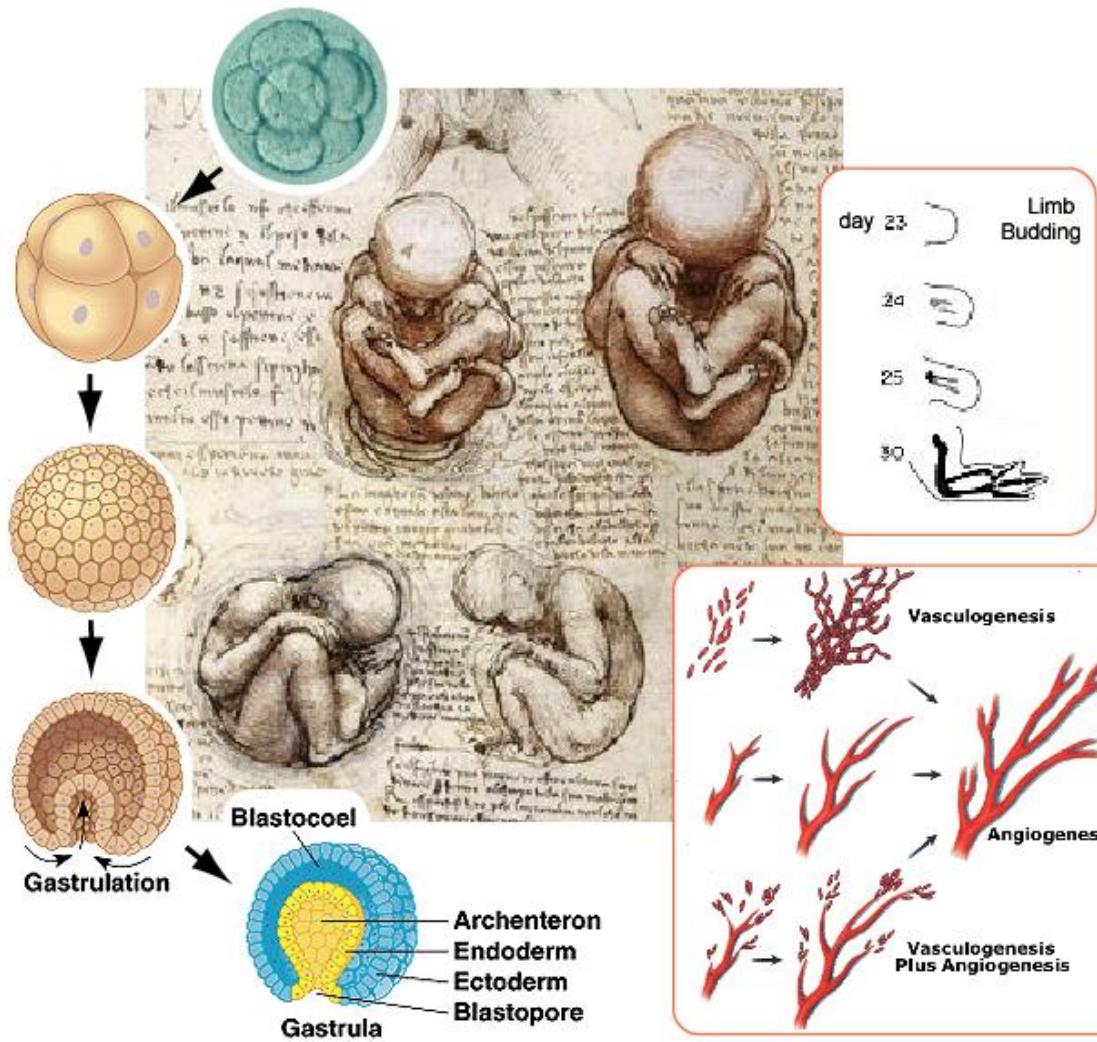


“Git along little doggies!”

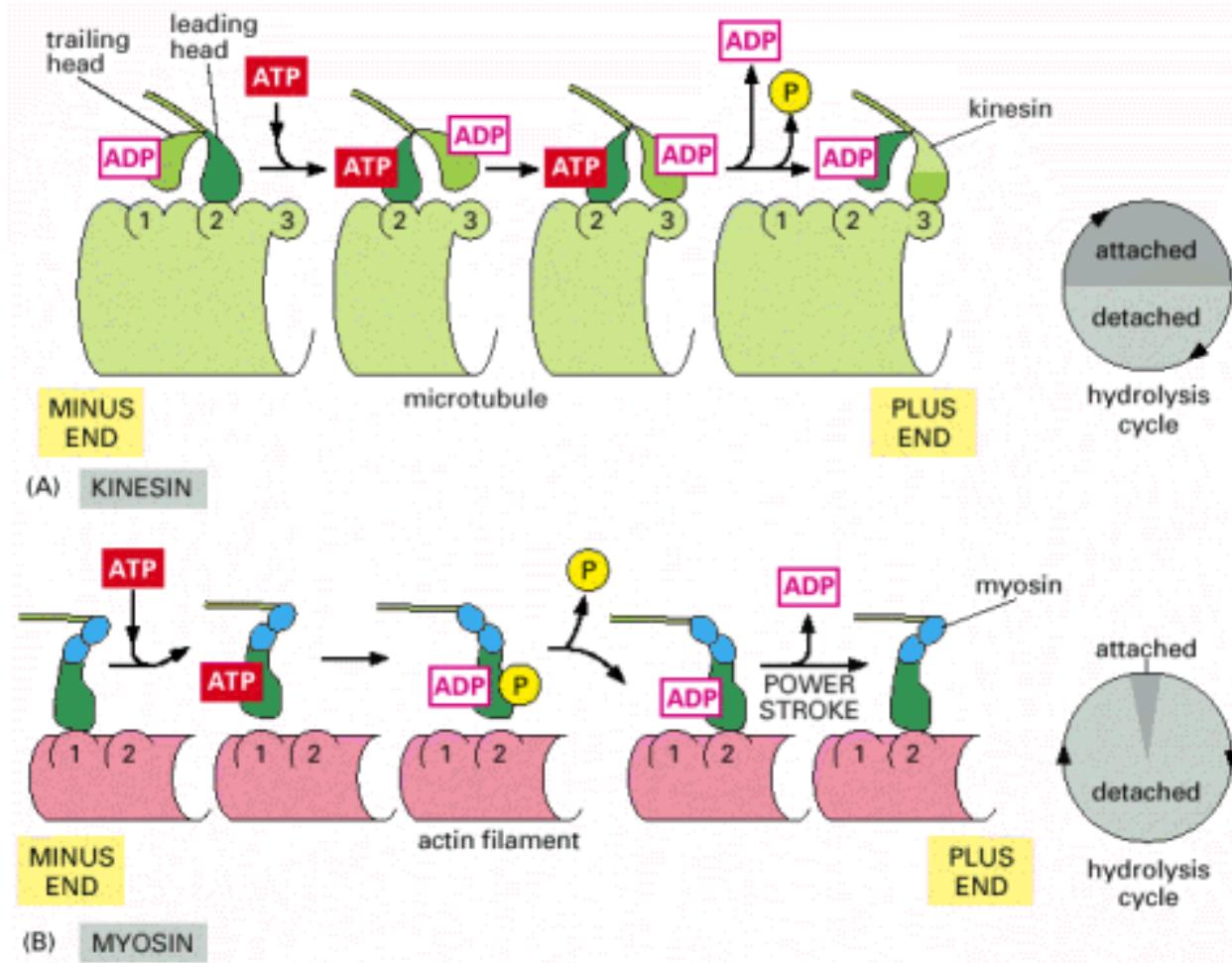
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Cell Movements

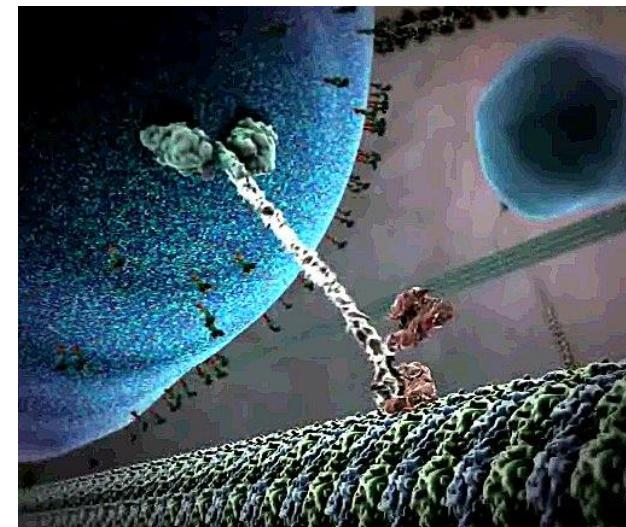
First Movements...



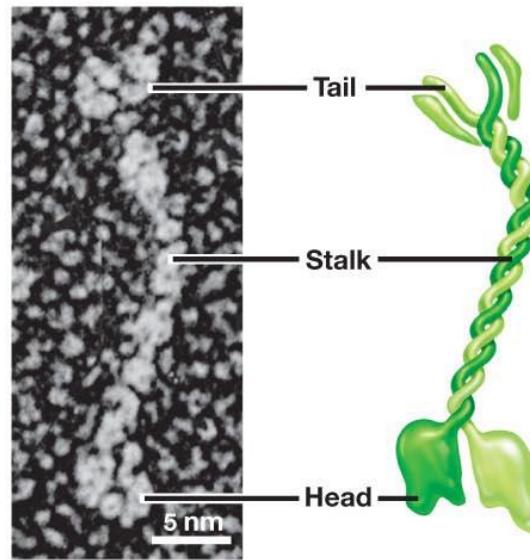
Molecular Motors



Kinesin

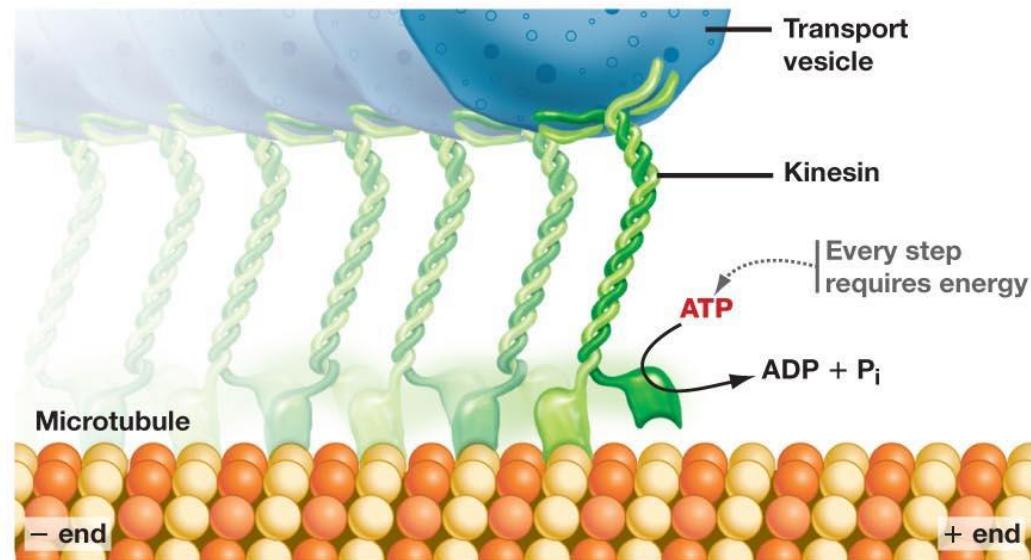


(a) Structure of kinesin

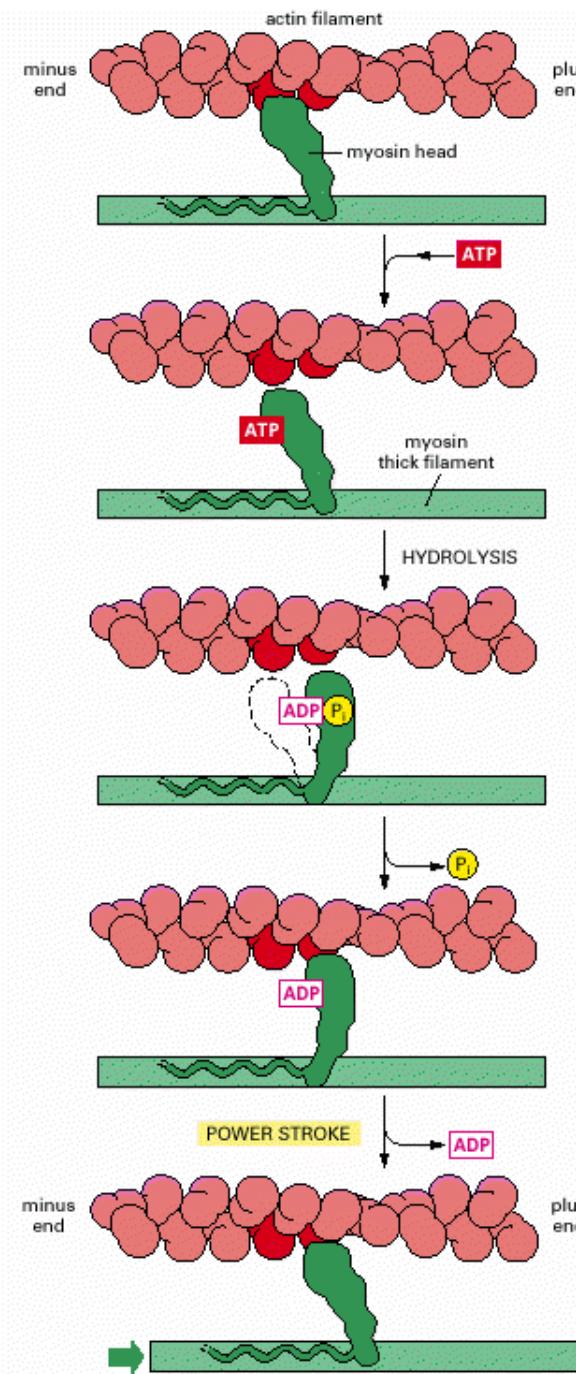
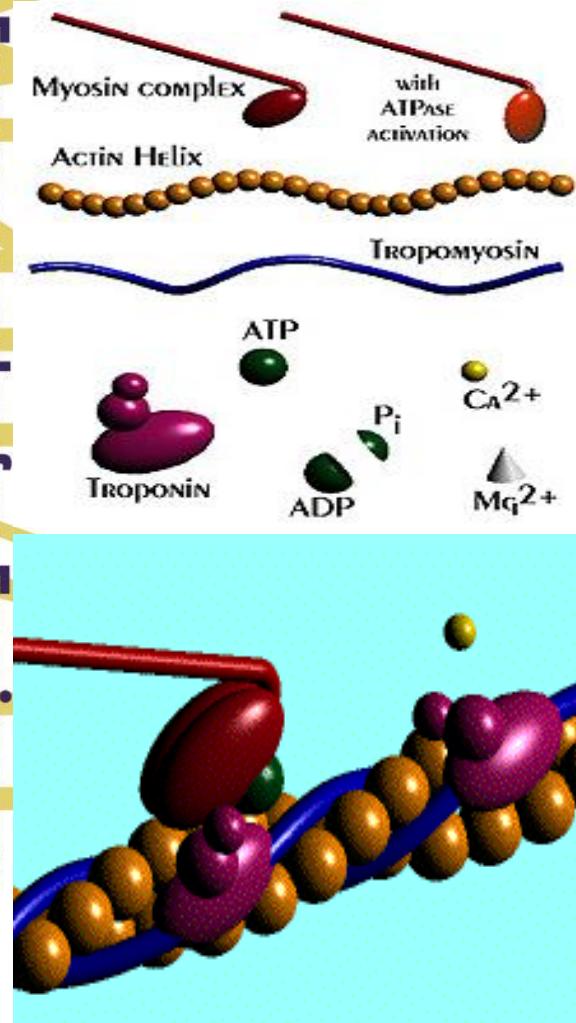


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(b) Kinesin “walks” along a microtubule track.



Myosin



ATTACHED At the start of the cycle shown in this figure, a myosin head lacking a bound nucleotide is locked tightly onto an actin filament in a *rigor* configuration (so named because it is responsible for *rigor mortis*, the rigidity of death). In an actively contracting muscle, this state is very short-lived, being rapidly terminated by the binding of a molecule of ATP.

RELEASED A molecule of ATP binds to the large cleft on the "back" of the head (that is, on the side furthest from the actin filament) and immediately causes a slight change in the conformation of the domains that make up the actin-binding site. This reduces the affinity of the head for actin and allows it to move along the filament. (The space drawn here between the head and actin emphasizes this change, although in reality the head probably remains very close to the actin.)

COCKED The cleft closes like a clam shell around the ATP molecule, triggering a large shape change that causes the head to be displaced along the filament by a distance of about 5 nm. Hydrolysis of ATP occurs, but the ADP and inorganic phosphate (P_i) produced remain tightly bound to the protein.

FORCE-GENERATING A weak binding of the myosin head to a new site on the actin filament triggers release of the inorganic phosphate (P_i) produced by ATP hydrolysis, concomitantly with the tight binding of the head to actin. This release triggers the power stroke—the force-generating change in shape during which the head regains its original conformation. In the course of the power stroke, the head loses its bound ADP, thereby returning to the start of a new cycle.

ATTACHED At the end of the cycle, the myosin head is again locked tightly to the actin filament in a rigor configuration. Note that the head has moved to a new position on the actin filament.

Important Movements

- Cytokinesis
- Migration of Cells
- Contraction of Cells

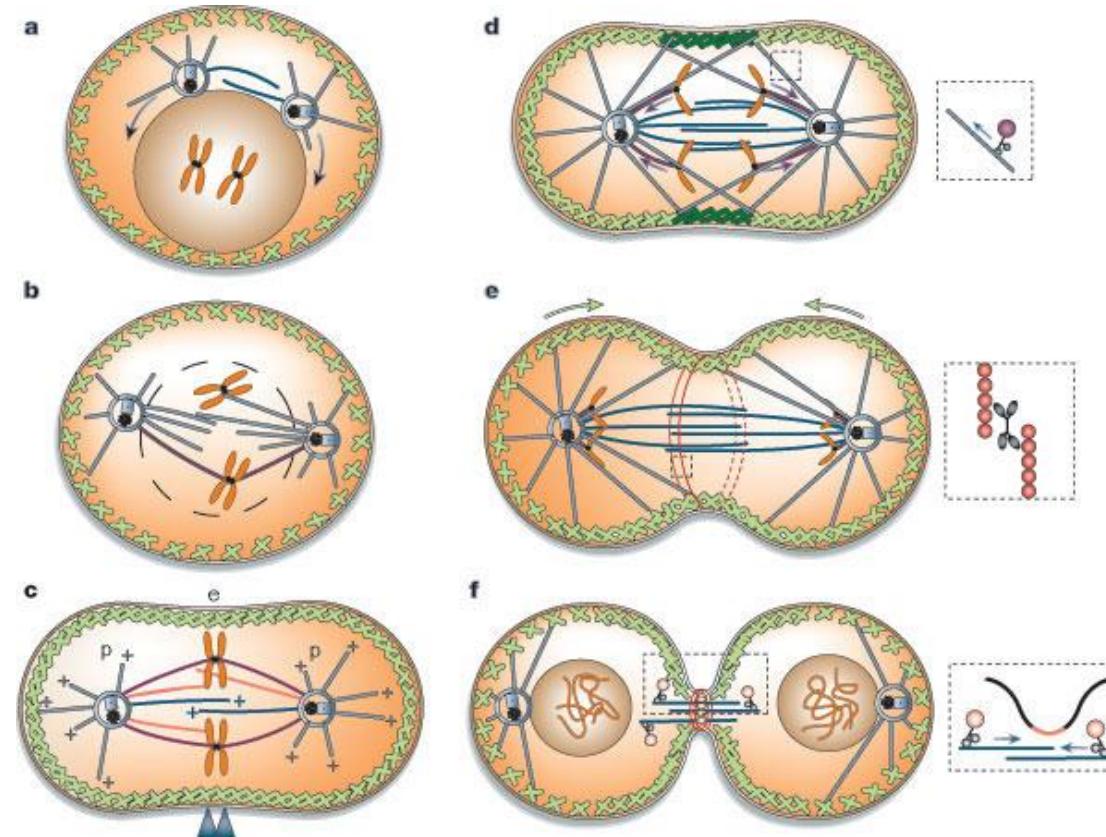
Cytokinesis

Frog Embryo Development



<http://www.youtube.com/watch?v=dXpAbezdOho>

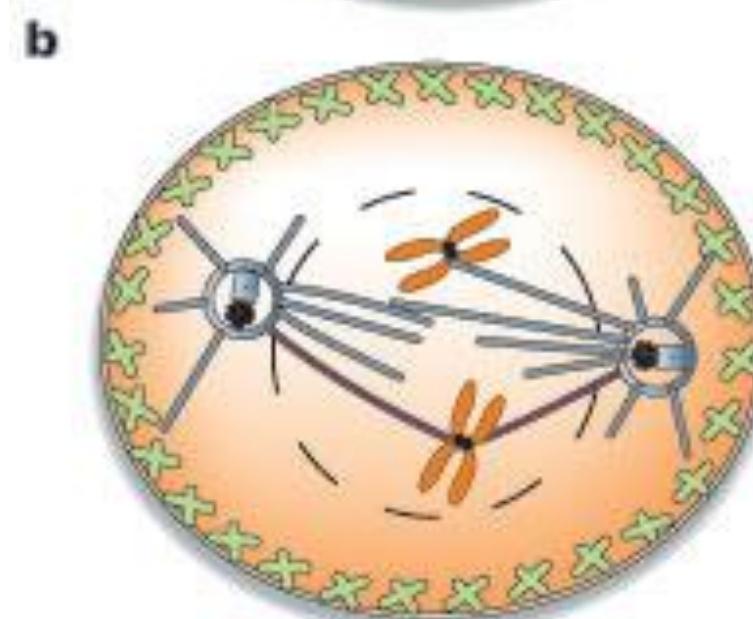
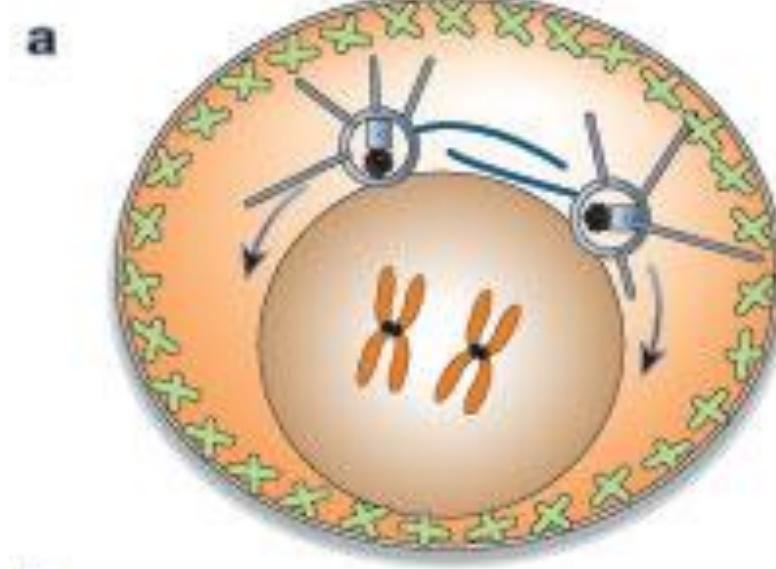
Cytokinesis



Nucleus	Interpolar Microtubule	Chromosomal Microtubule	Contractile Ring	Membrane vesicle
Chromosome	Kinetochore Microtubule	Midbody	Actin filament	Microtubule motor
Centrosome	Astral Microtubule	Cell cortex	Myosin II	

Cytokinesis

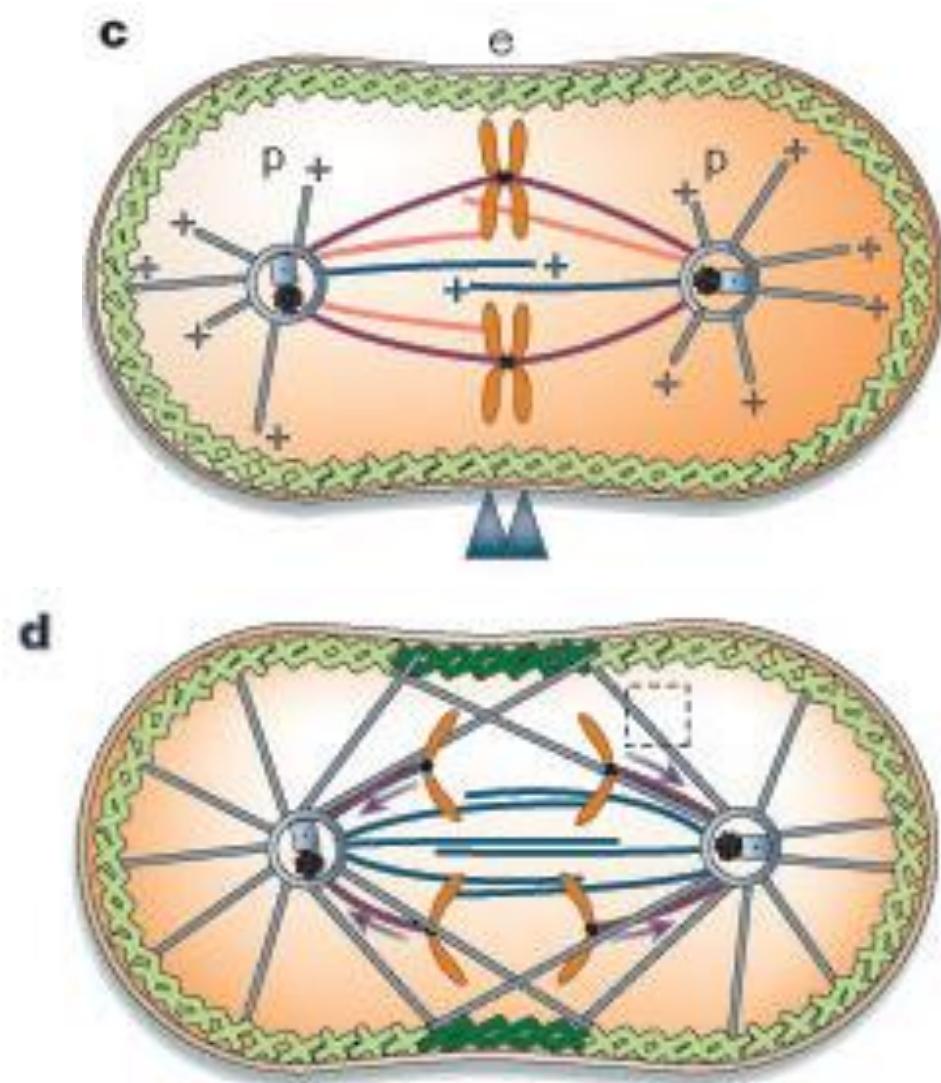
- Prophase
- Prometaphase



Cytokinesis

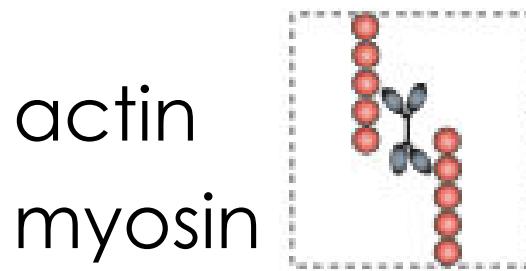
- Metaphase
- Anaphase A

kinesin

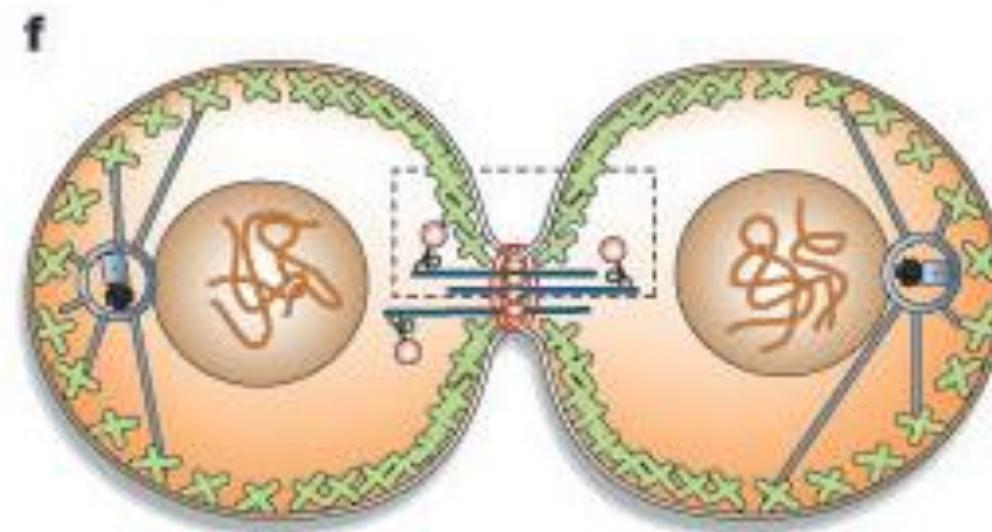
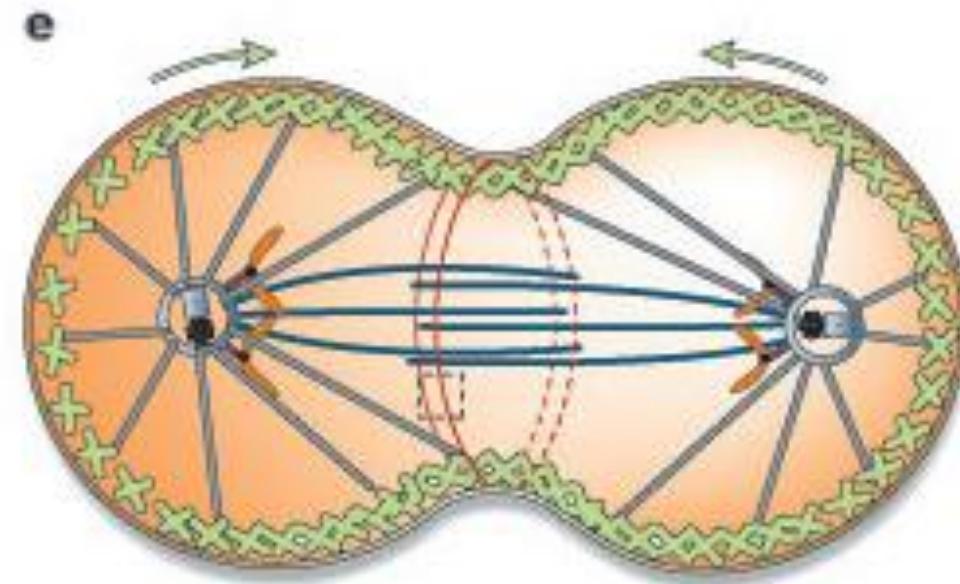
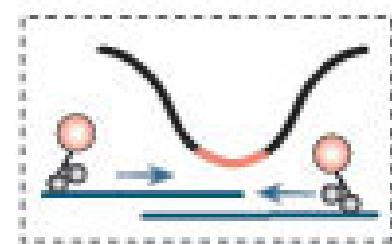


Cytokinesis

- Anaphase B



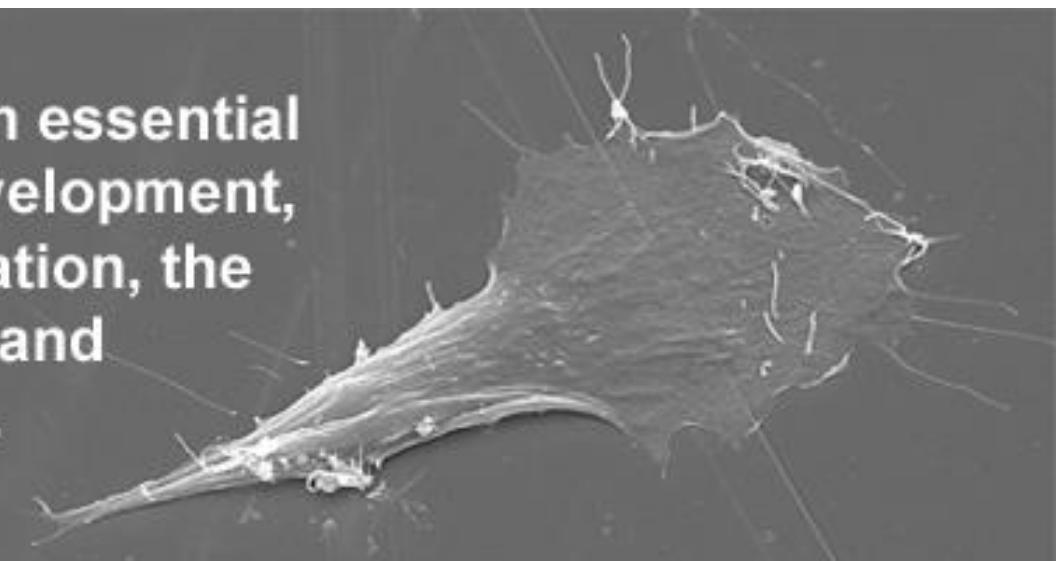
- Telophase abscission



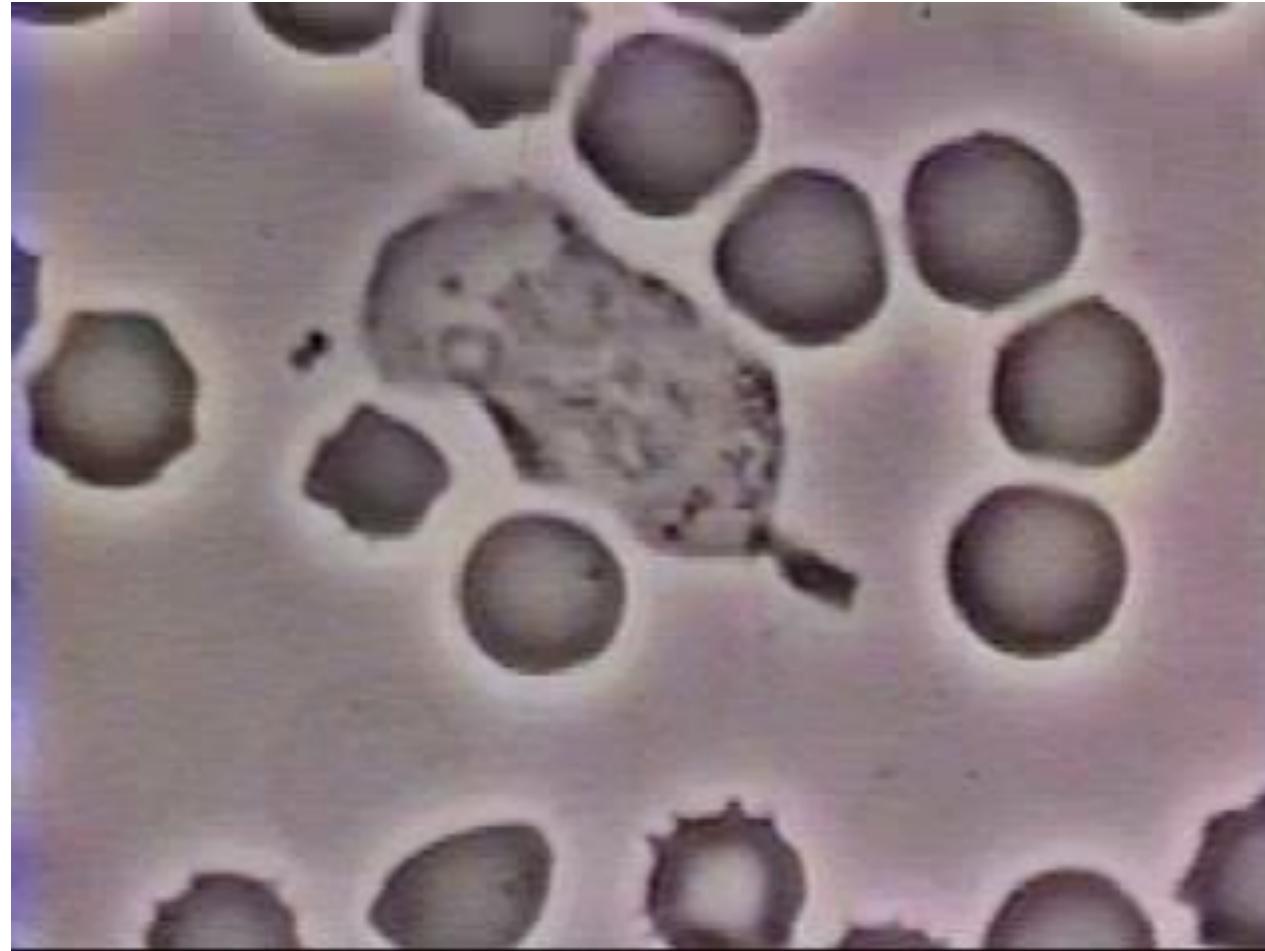
Cell Migration

Cell Migration

Cell migration is an essential part of embryo development, blood vessel formation, the immune response and cancer metastasis.



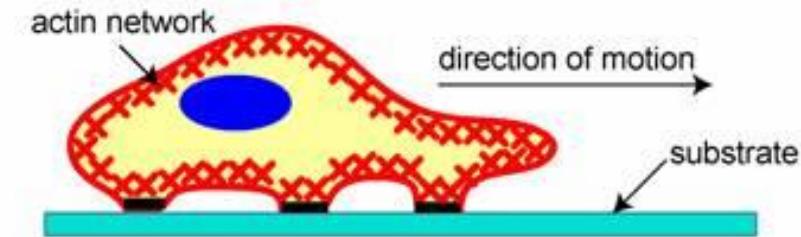
Cell Migration



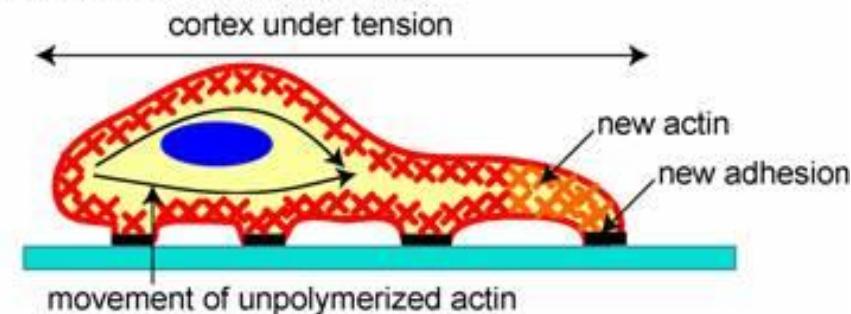
Neutrophil chasing a bacterium

Steps in Cell Migration

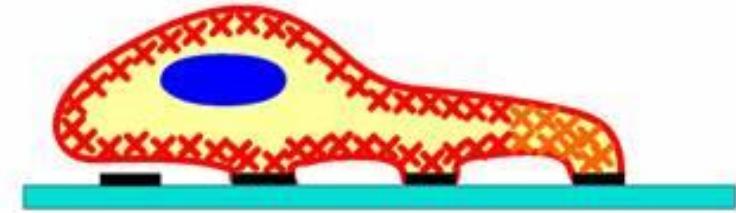
1) Protrusion of the Leading Edge



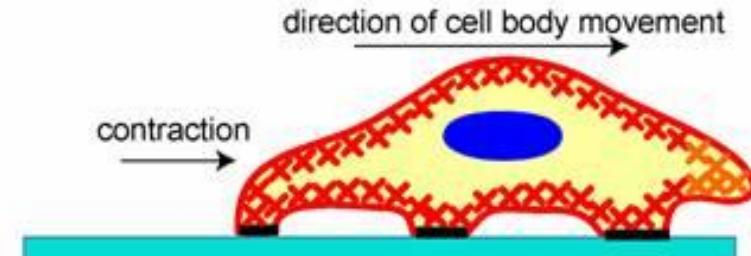
2) Adhesion at the Leading Edge



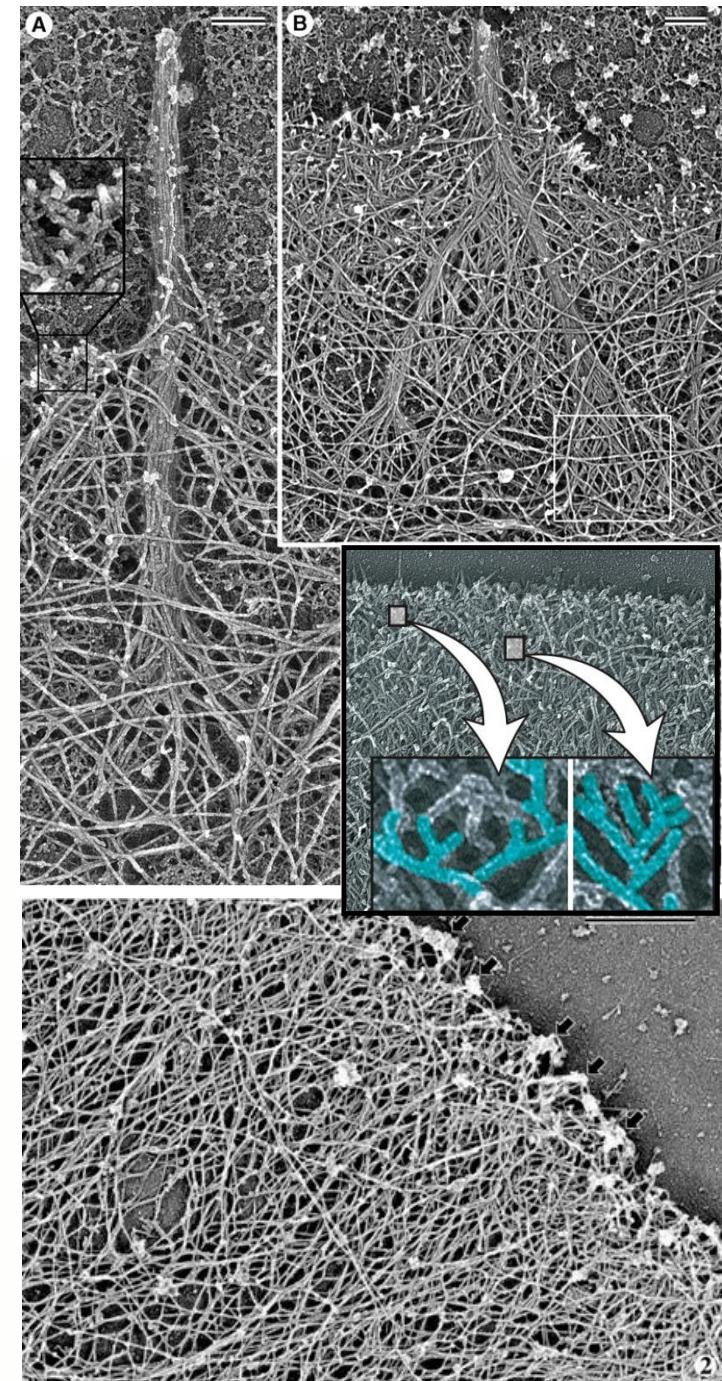
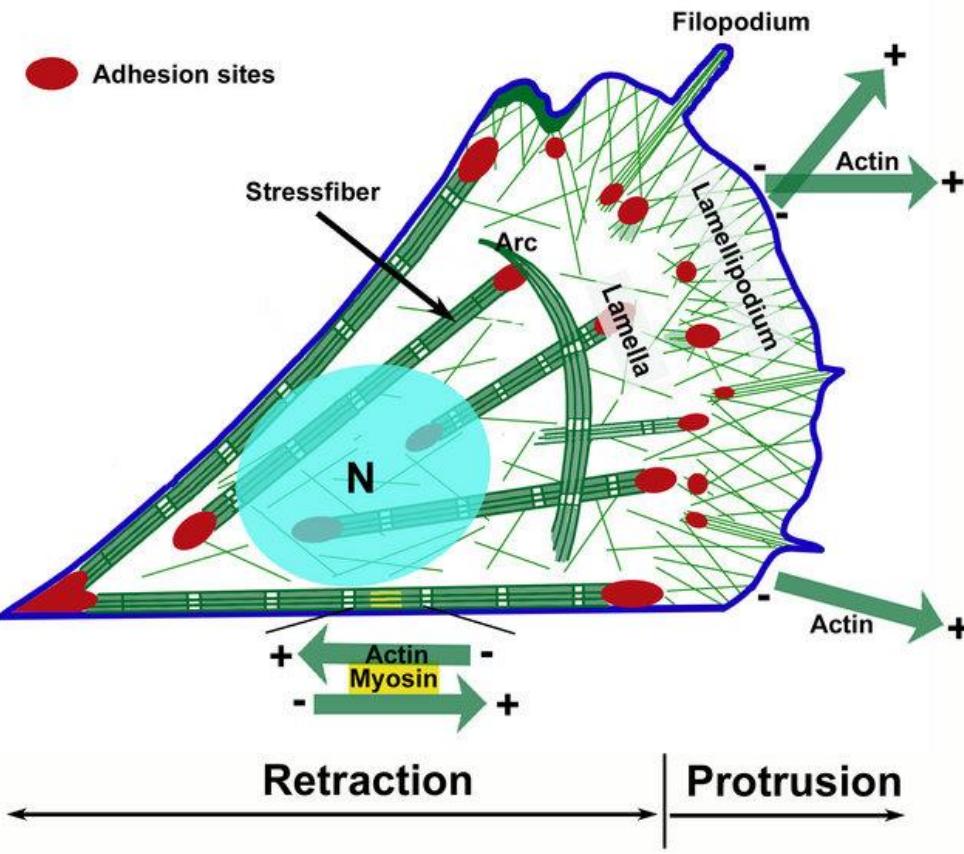
Deadhession at the Trailing Edge



3) Movement of the Cell Body

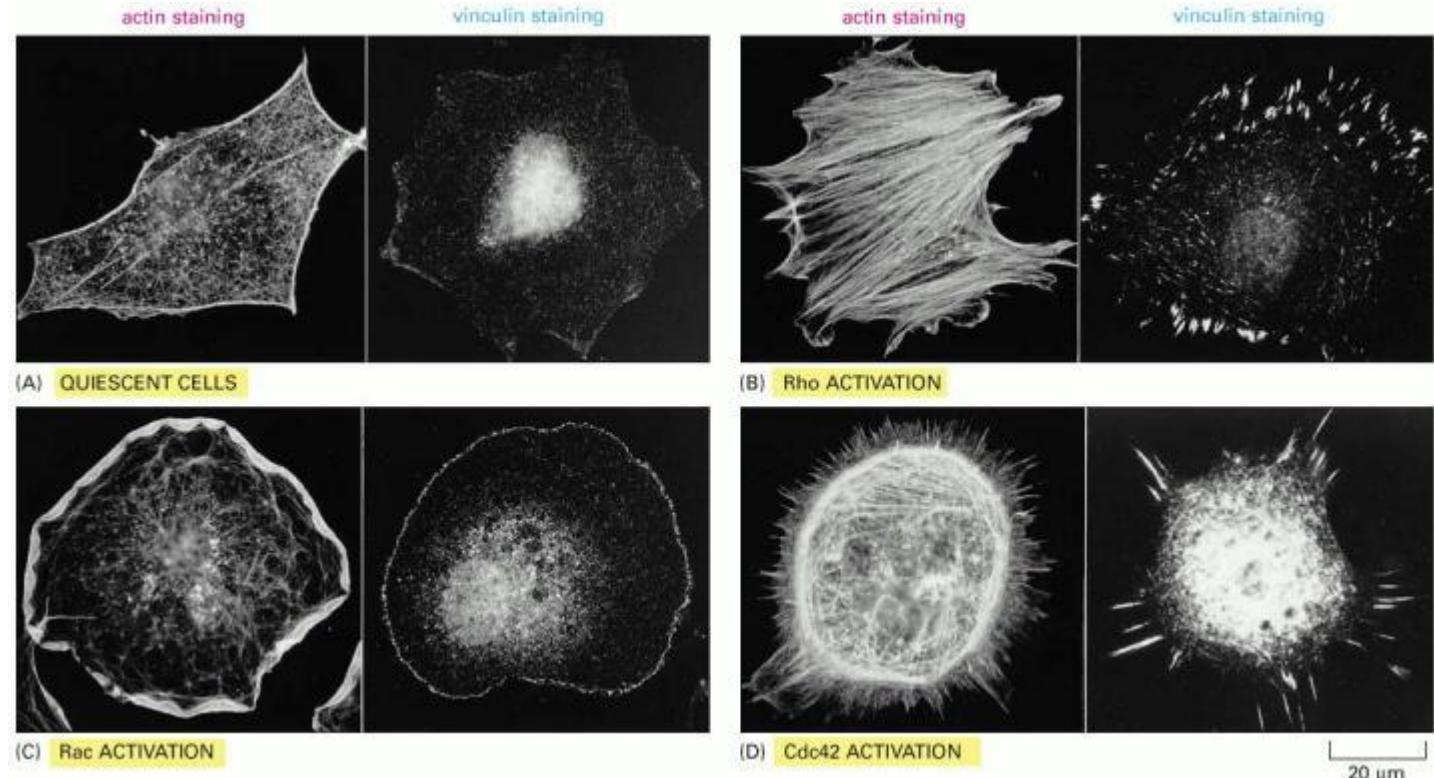
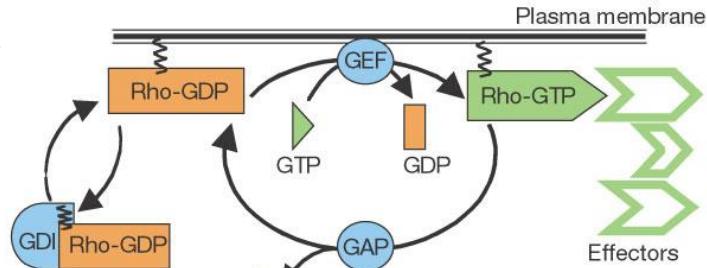


Lamellipodium & Filopodia



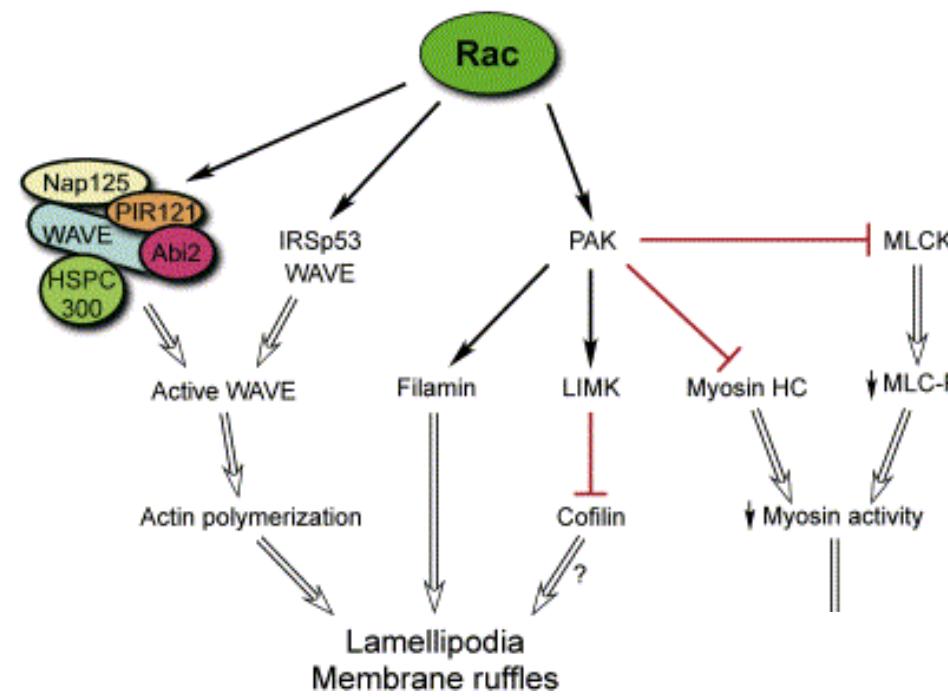
Rho Family GTPases

An interesting discovery by Anne Ridley and Alan Hall while studying the oncogene *ras*...



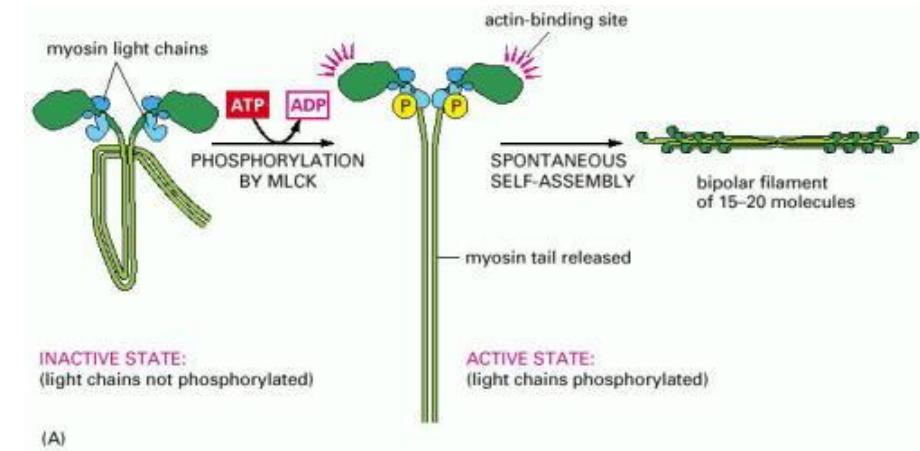
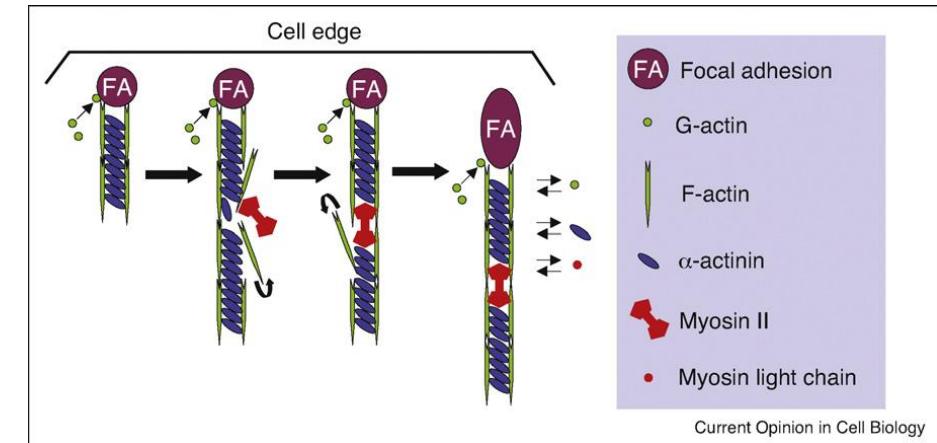
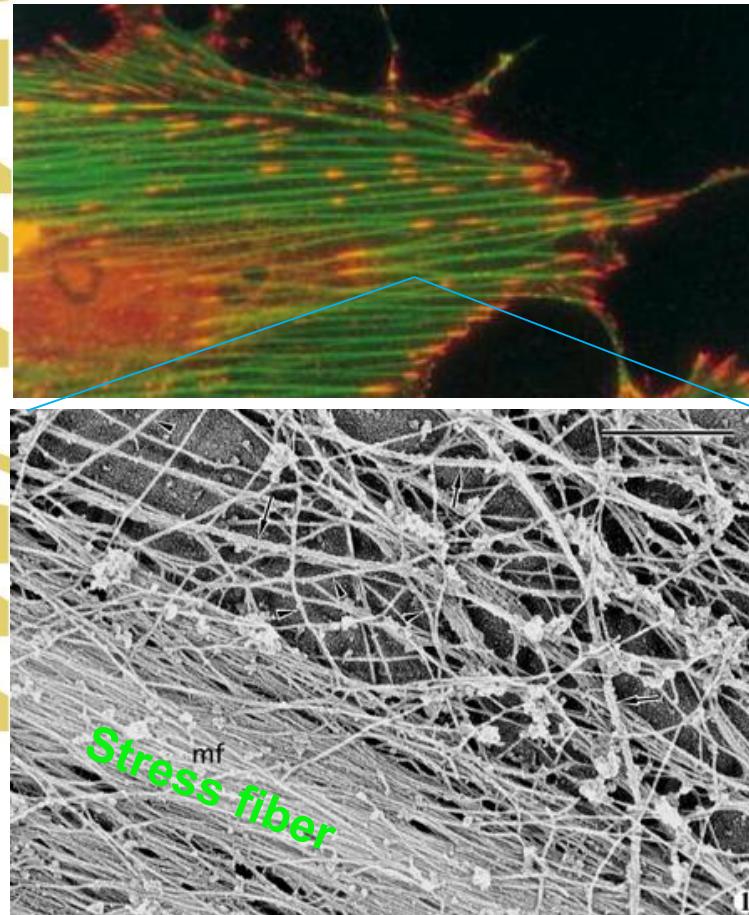
Rac Effectors

- Rac helps migration and cell elongation



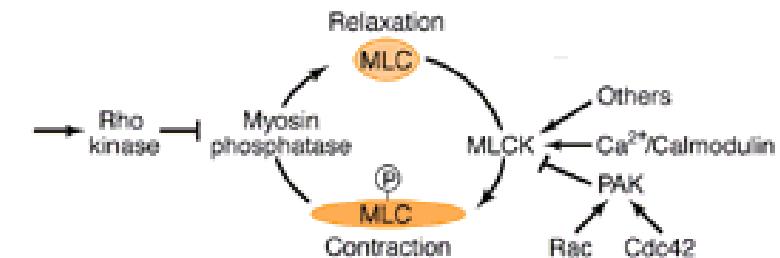
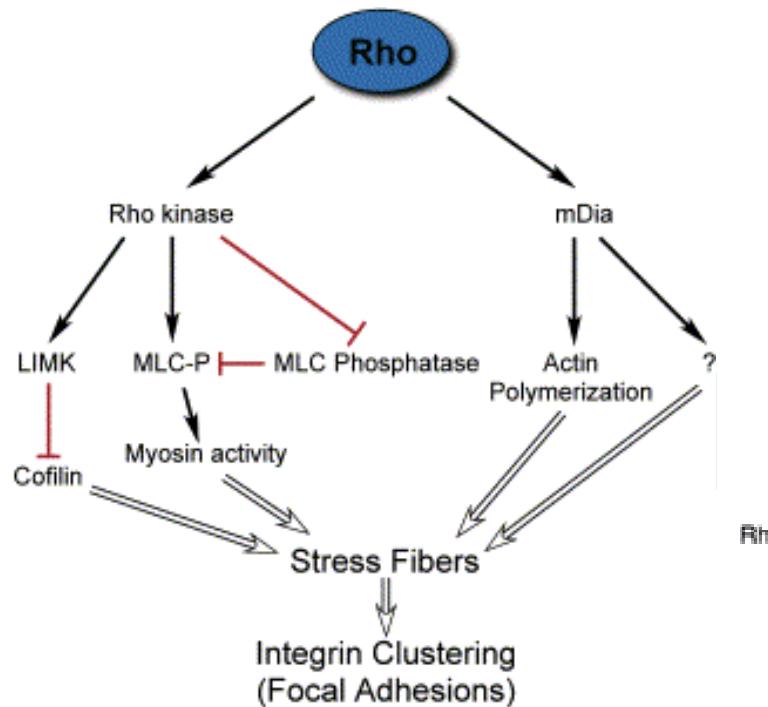
Stress Fibers

- Transient bundled structures

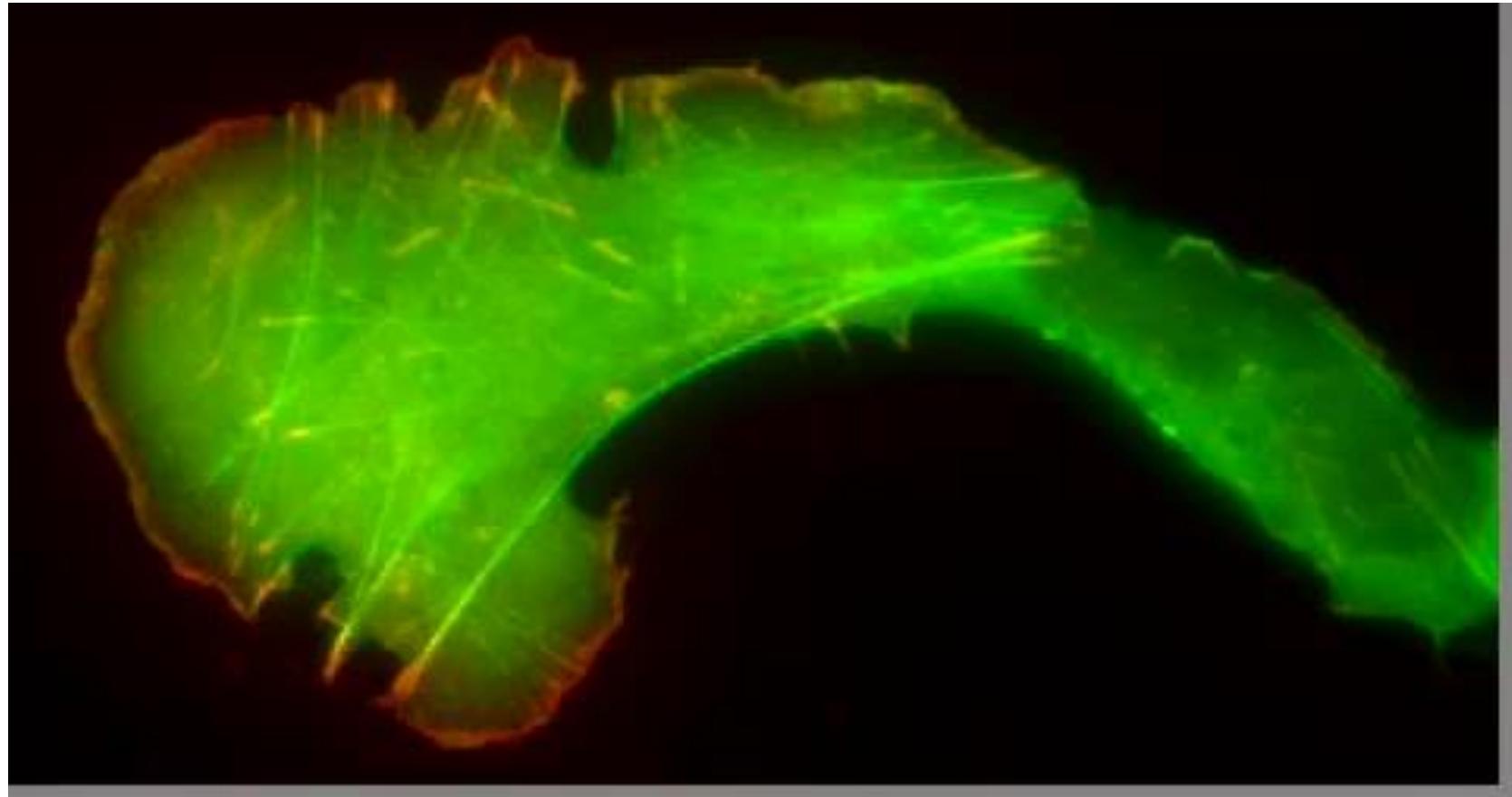


Rho Effectors

- Rho promotes actin-myosin tension and helps migration

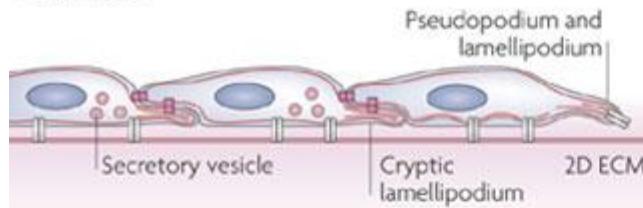


Cell Migration

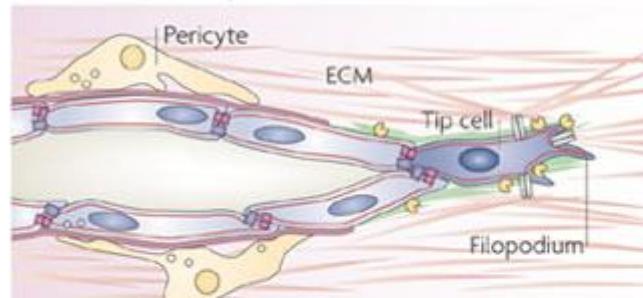


Multicellular Migration

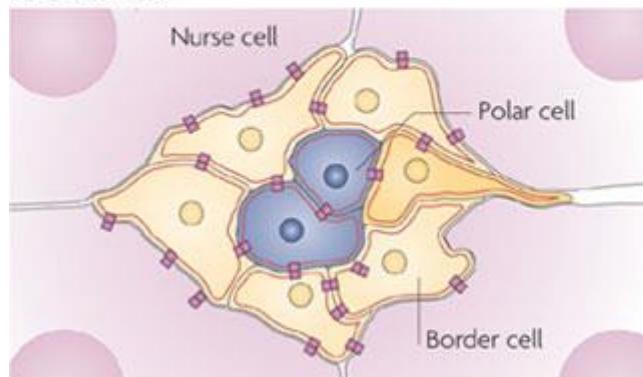
a 2D sheet



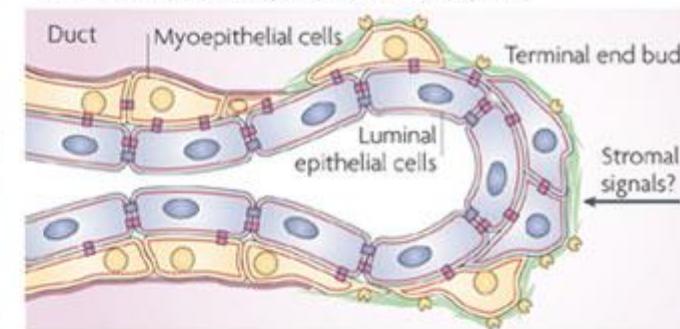
c Vascular sprouting



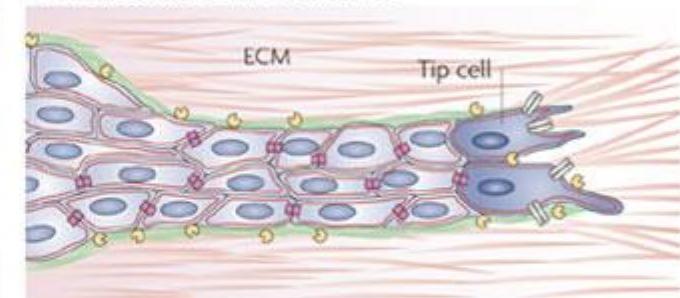
e Border cells



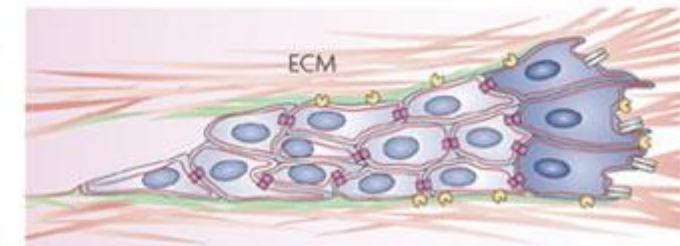
b Branching morphogenesis (mammary gland)



d Multicellular 3D invasion strands



f Detached cluster



— Basement membrane
— Cortical F-actin

■ Adherens junction
■ Tight junction



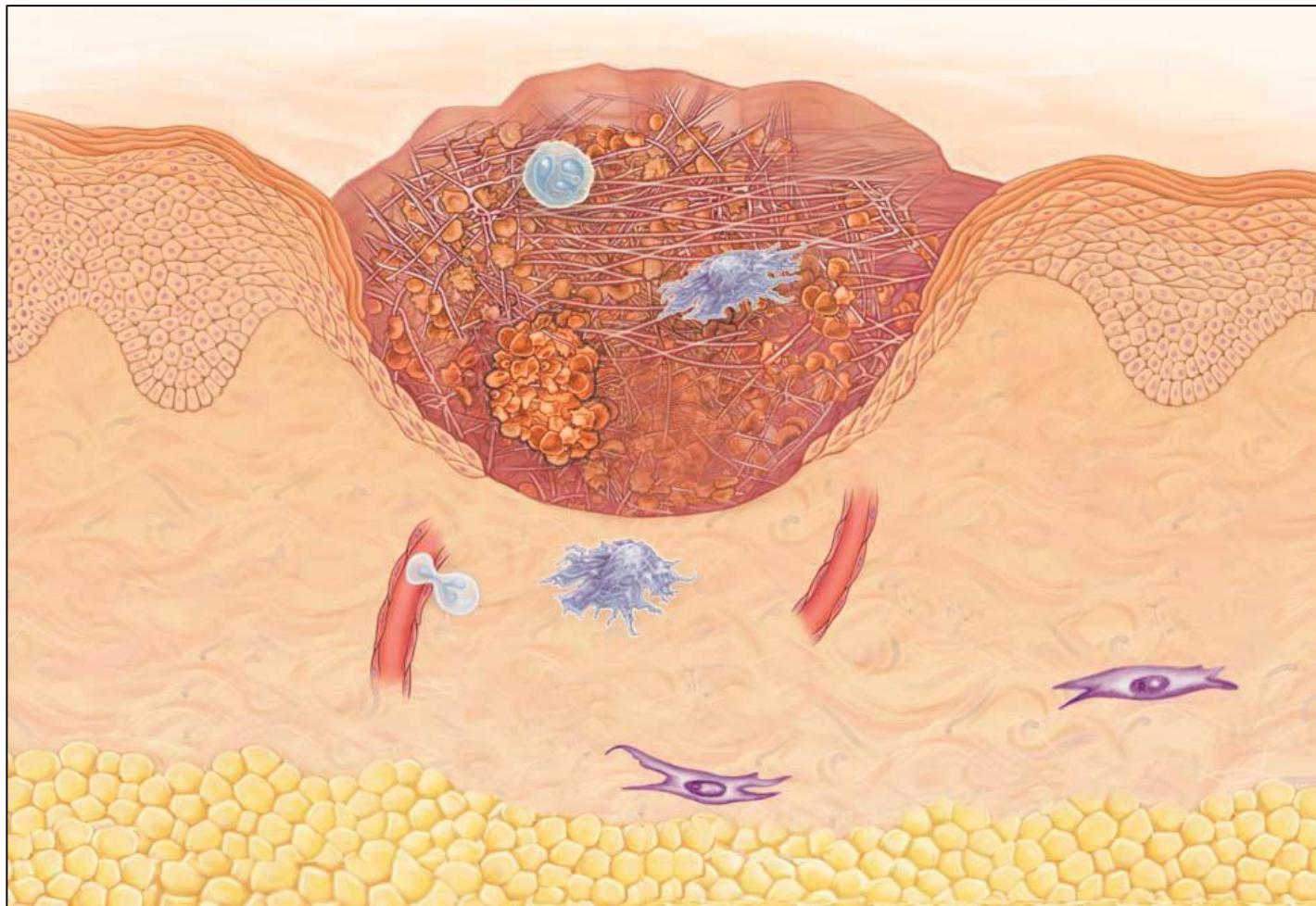
Proteolytically
remodelled ECM

■ Integrins ($\beta 1$ and $\beta 3$)
in focal contacts

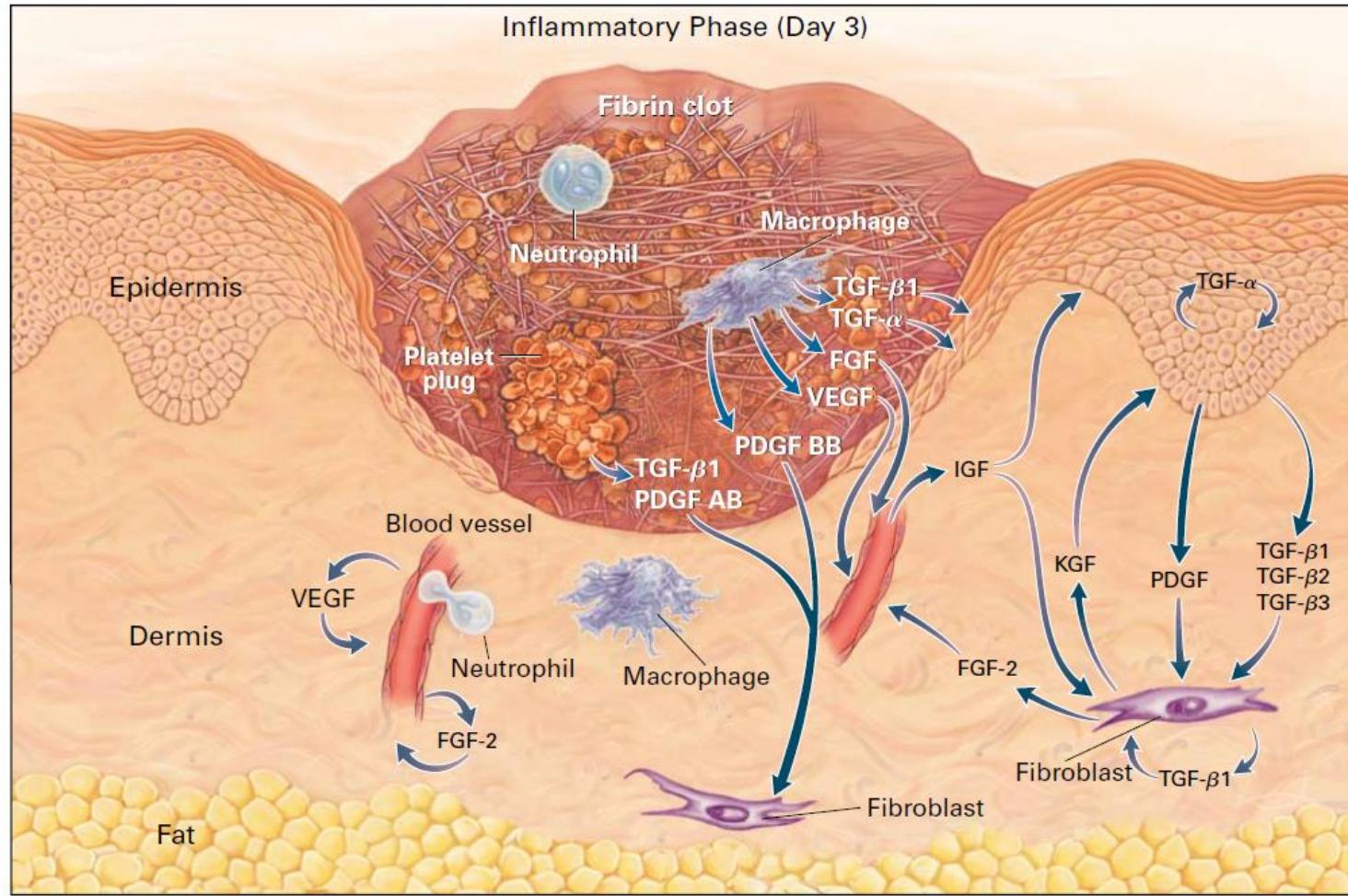
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■ Proteases (MMPs,
MTMMP and UPA)

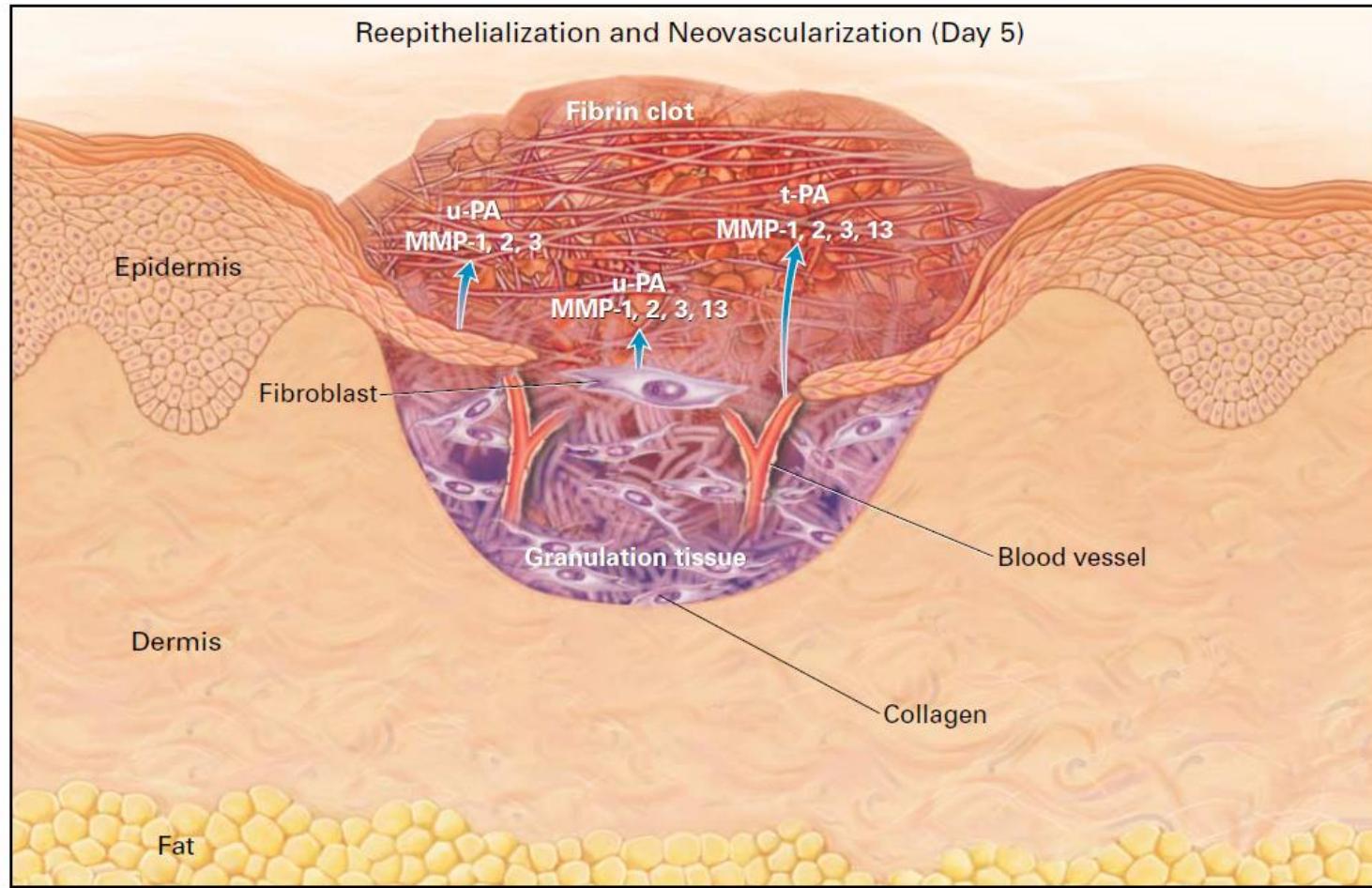
Wound Healing



Cell Migration & Secretion



Healing and Scarring



Cell Contraction

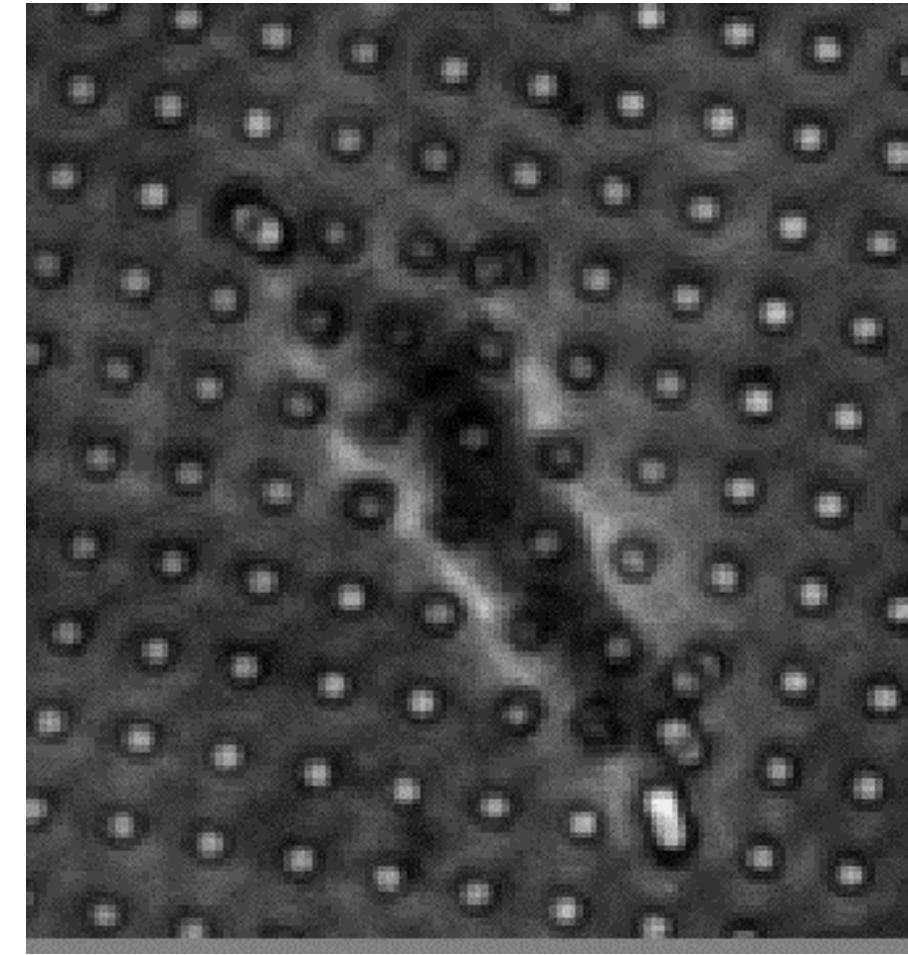
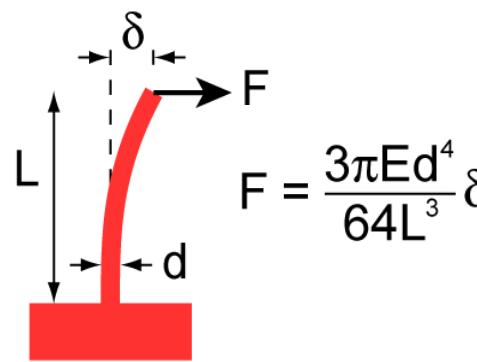
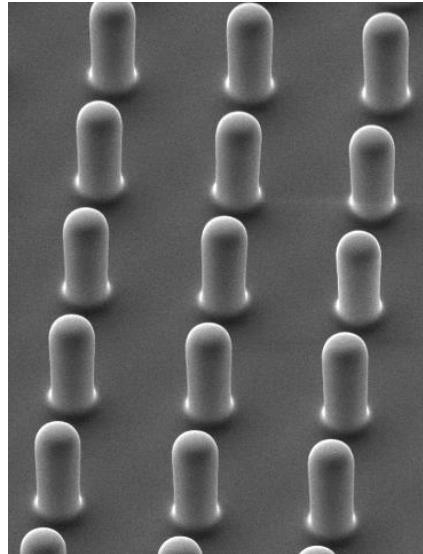
Wrinkling Membranes



Cultured lung myofibroblast,
wrinkling a deformable
silicone substrate

SMA-FP = NH₂-terminal peptide of α -smooth muscle actin inhibits force generation

Microposts

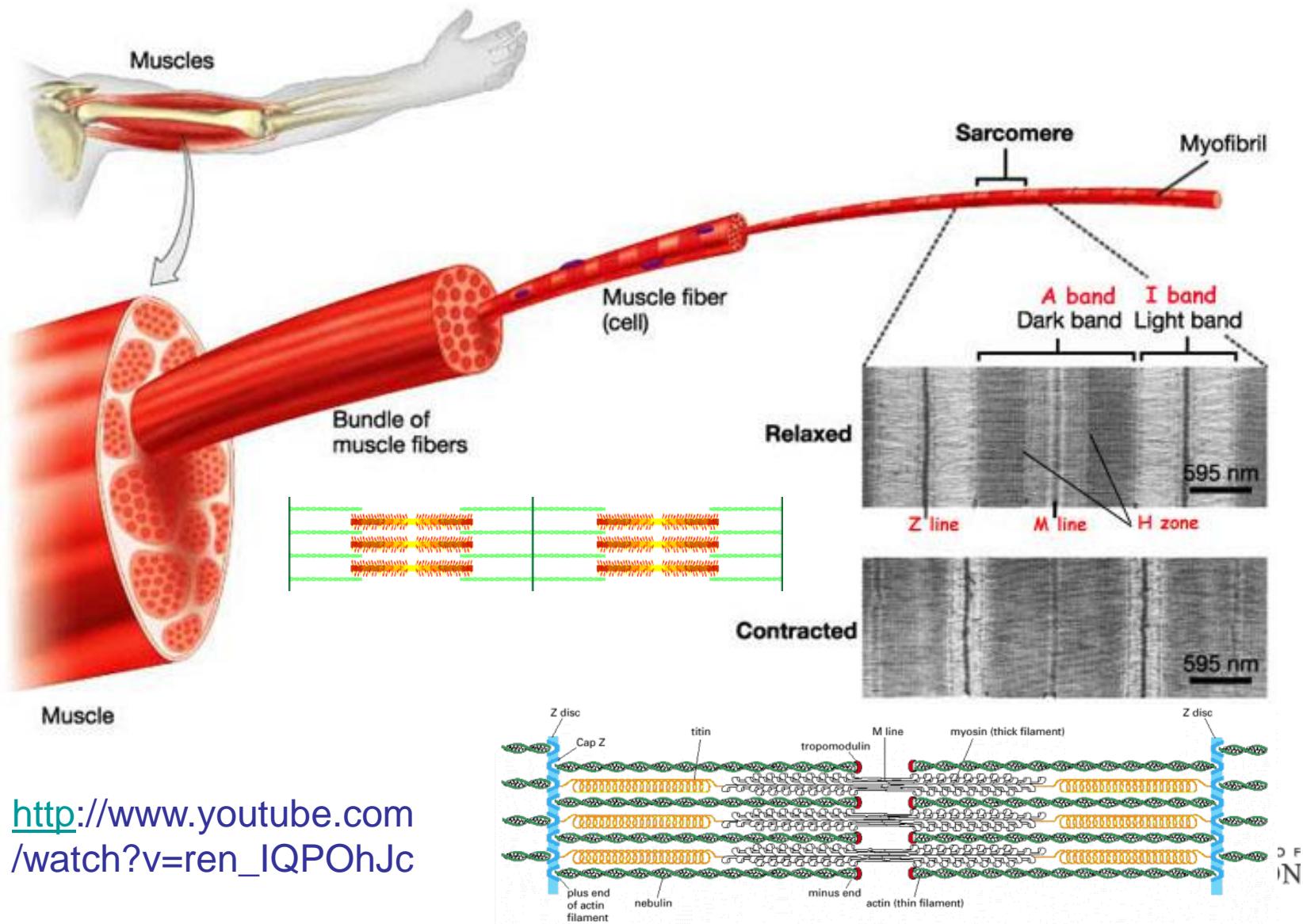


Muscle Cells

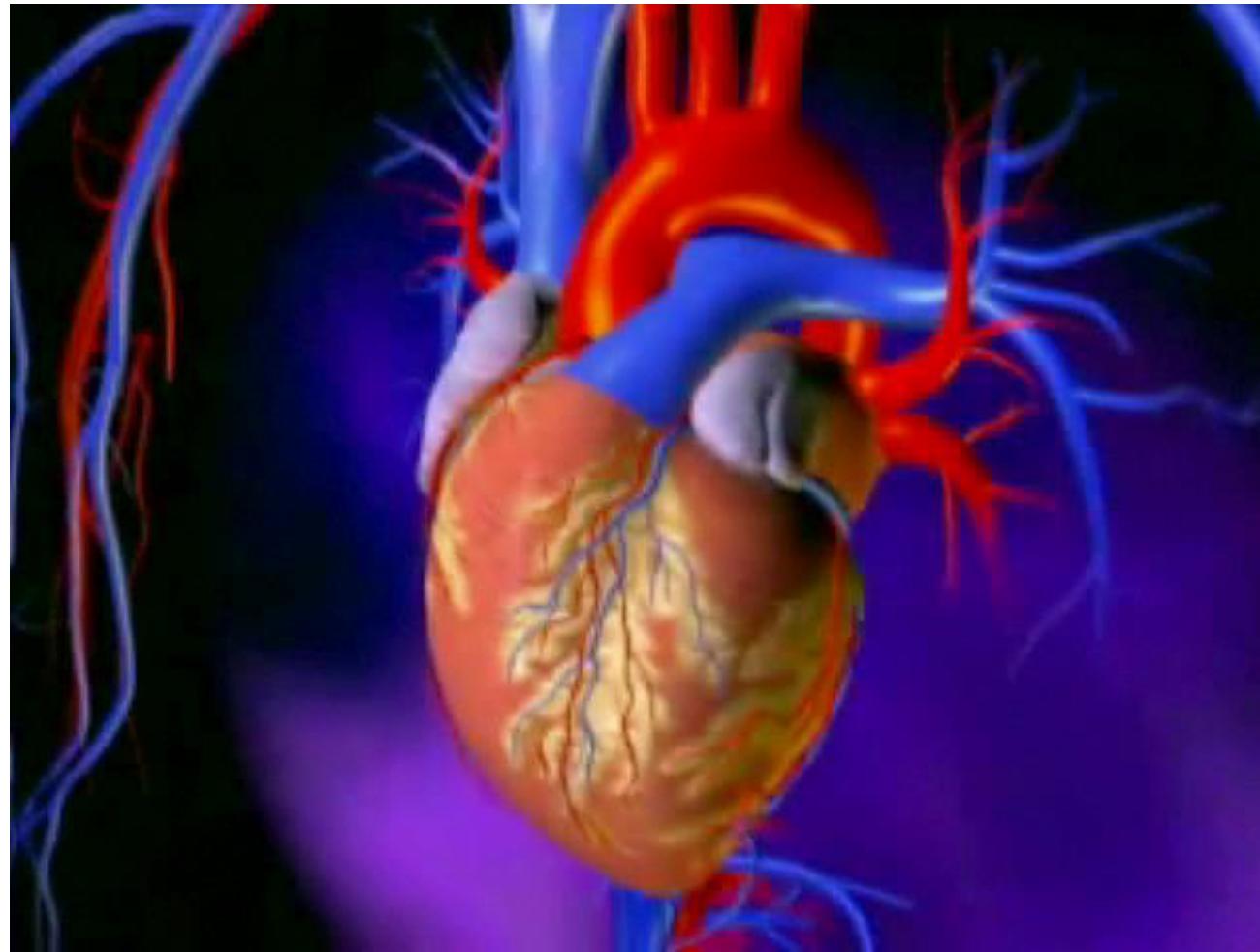


<http://www.youtube.com/watch?v=4j3NW1qrR0k>

Sarcomeres



Cardiac Contraction



Questions?