

Skeletal Muscle Structure

Skeletal muscles are connected to at least two bones by tendons (cords of elastic connective tissue)

Muscle fibers have many nuclei b/c they are formed during embryonic life from fusion of cells

Sarcoplasmic reticulum: surrounds each myofibers → activates contraction

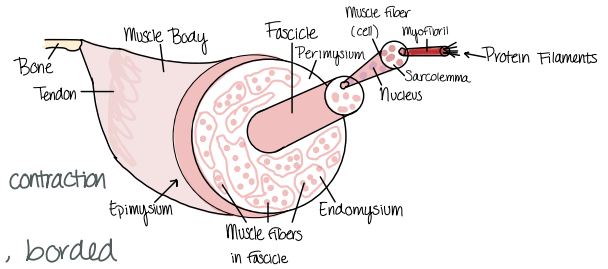
Sarcomere: makes up myofibrils, repeat over + over, bordered

by Z lines ; M lines connect thick filaments (both run perpendicular to the long axis)

↳ Filaments are made up of ^(thin) actin + ^(thick) myosin proteins

- two regulatory proteins that enable contraction: tropomyosin → blocks myosin-binding sites in muscles at rest ; troponin → 3-protein complex

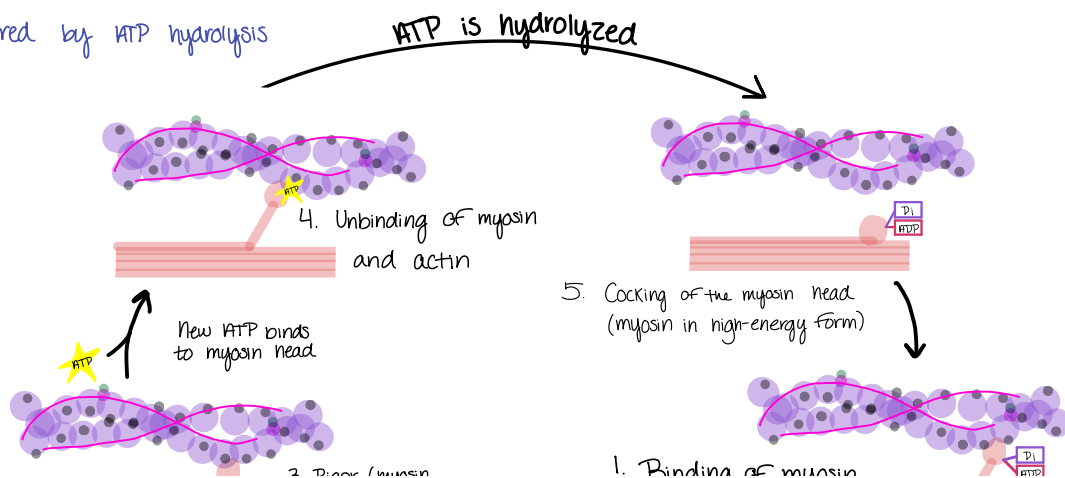
- the head of the myosin molecule actively generates mechanical force → actin-binding site + ATPase binding site (enzymatic activity + hydrolyzes ATP)

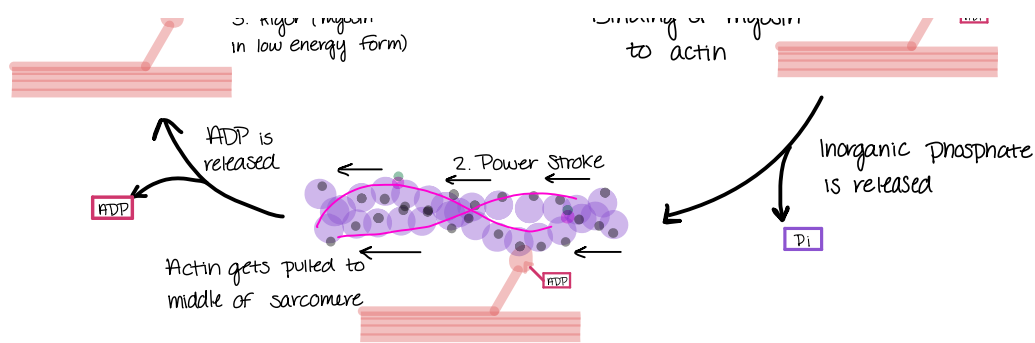


Mechanism of Force

The sliding-filament model: contractions are caused when the I bands and H zones shorten → I band shortening occurs when they slide past thick filaments, deeper into the H zone (decreasing its width) → A bands move closer together (decreasing I band width) → Z lines move closer + sarcomere is shortened

Crossbridge Cycle: what drives thick + thin filaments past one another → oscillating motion of myosin crossbridges, powered by ATP hydrolysis





- **excitation-contraction coupling:** muscle cell receives input from motor neuron $\rightarrow$ cell depolarizes + fires action potential to stimulate contraction
- **neuromuscular junction:** motor neuron's terminal boutons fan out in the sarcolemma, acetylcholine is released, end-plate potential is produced
 - $\hookrightarrow$ if an action potential occurs in a motor neuron, it's always followed by an AP in the muscle cell it innervates

Skeletal muscle Contraction

- **Twitch:** an AP triggers contraction in all the muscle cells connected to a motor neuron (motor unit)
 - latent period $\rightarrow$ delay between AP and contraction ; contraction phase $\rightarrow$ muscular tension/force peaks, cytosolic calcium levels increase ; relaxation phase $\rightarrow$ longest, cytosolic calcium levels and # of active crossbridges decrease
- **Isotonic contraction:** muscle generates constant tension (concentric $\rightarrow$ shortening of the muscle, eccentric $\rightarrow$ lengthening - lowering an object)
- **Isometric contraction:** muscle creates tension but maintains the same length b/c the load is greater than the force exerted by the muscle
- **Contractile component** $\rightarrow$ sarcomeres, **series elastic component** $\rightarrow$ parts of a muscle that don't actively generate force, but passively transmit force to the ends of muscles
- **Factors affecting force**
 - **Frequency of stimulation:** as frequency increases, muscles move from twitch contractions
 - **Treppe:** independent twitches follow each other closely $\rightarrow$ peak tension rises until it reaches constant
 - **summation:** repeated stimulation, twitches are superimposed to yield a force greater than a single twitch

- tetanus: highest frequencies, in unfused tetanus (incomplete), brief periods of relaxation occur; fused tetanus complete
- Fiber diameter: a muscle w/ more sarcomeres arranged in parallel can generate more force $\rightarrow$ fiber's diameter $\propto$ force generating capacity $\propto$ cross-sectional area
- changes in fiber length: optimum length, when its not in that range, force-generating capacity decreases
- ~~Regulation of force~~
- recruitment: an increase in # of active motor units
- size principle: correspondence between size of motor units + order of recruitment
- velocity of shortening: the rate of change of the distance shortened

Skeletal muscle Metabolism

- to ensure steady supply of ATP, muscles use creatine phosphate $\rightarrow$ donates phosphate to ADP $\rightarrow$ ATP
- fast myosin (in fast-twitch fibers) hydrolyzes ATP fast + has a fast ATPase rate $\rightarrow$ can complete more crossbridge cycles / second $\rightarrow$ sarcomeres shorten faster
- Glycolytic Fibers: high cytosolic concentrations of glycolytic enzymes
- Oxidative Fibers: rich in mitochondria + high capacity for producing ATP via oxidative phosphorylation $\rightarrow$ generally smaller diameter + lots of capillaries, contain myoglobin (oxygen-binding protein)

Control of skeletal muscles

- ~~muscle activity across joints~~
- origin $\rightarrow$ where muscle attaches to stationary bone; insertion $\rightarrow$ attachment to moveable bone
- muscle spindles: detect muscle length, consist of intrafusal fibers
- Golgi tendon organs: connective tissue intertwined with collagen fibers of tendons, detect muscle tension
- ~~Smooth muscle~~
- found in internal organs, blood vessels + structures not under voluntary control
- thin + thick filaments, generates force through the crossbridge cycle

↳ dense bodies: points of attachment between filaments + connective tissue inside cells

• regulated by autonomic neurons. Synchronized activity is promoted by Gap Junctions → allow ions to move between cells + electrical signals spread to neighboring cells

• Cardiac Muscle

- connected by gap junctions → action potentials travel throughout the entire cell network

- summation cannot occur so the heart can relax + fill w/ blood