

MUSCULAR SYSTEM

Primary Care Paramedicine

Module: 14

Section: 01c





Excitability

- The ability to receive and respond to stimulus

Contractility

- The ability to contract

Extensibility

- The ability to stretch (opposing pairs)

Elasticity

- The ability to recoil to original shape



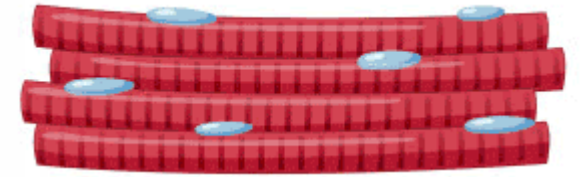
Movement

Posture

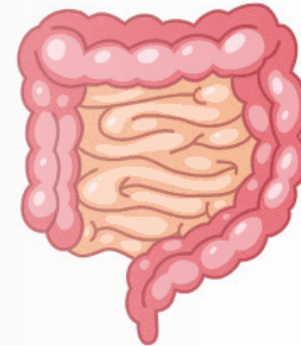
Joint stability

Heat Production

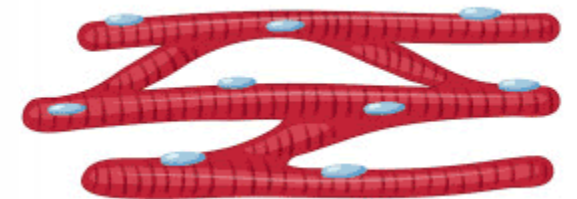
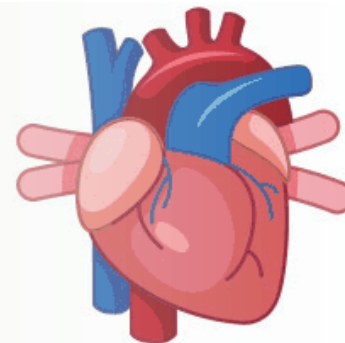
Skeletal Muscle

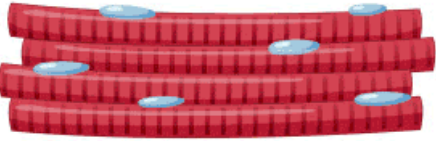
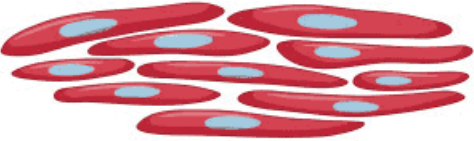
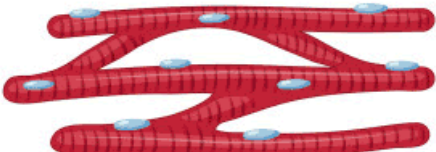


Smooth Muscle



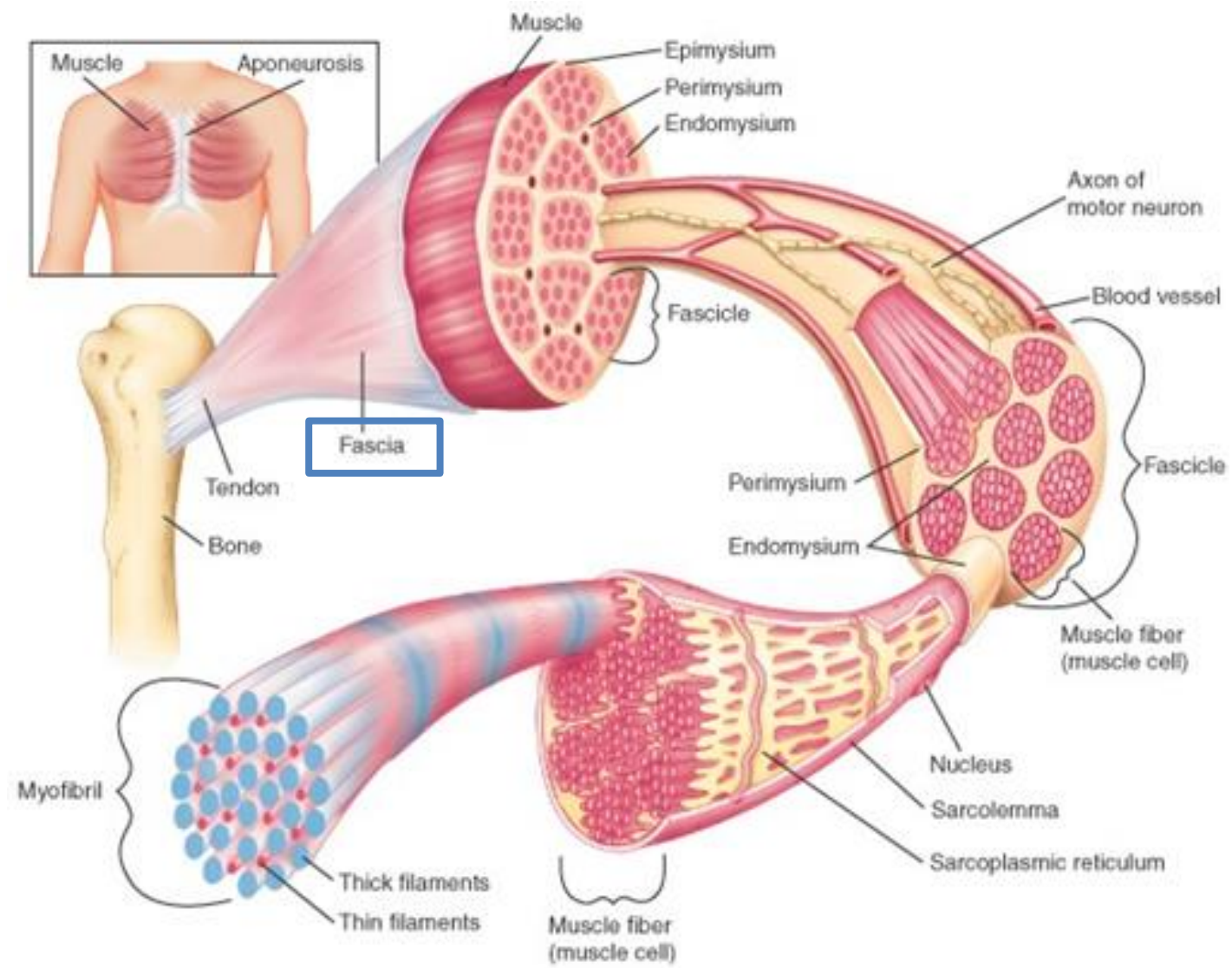
Cardiac Muscle



Type	Location	Function	Appearance	Control
Skeletal 	Skeleton	Movement, heat, posture	<ul style="list-style-type: none"> • Striated • Multi-nucleated (eccentric) • Fibers parallel 	Voluntary
Smooth 	Heart	Pump blood continuously	<ul style="list-style-type: none"> • Striated • One central nucleus 	Involuntary
Cardiac 	G.I. tract, uterus, eye, blood vessels	Peristalsis, blood pressure, pupil size, erects hairs	<ul style="list-style-type: none"> • No striations • One central nucleus 	Involuntary

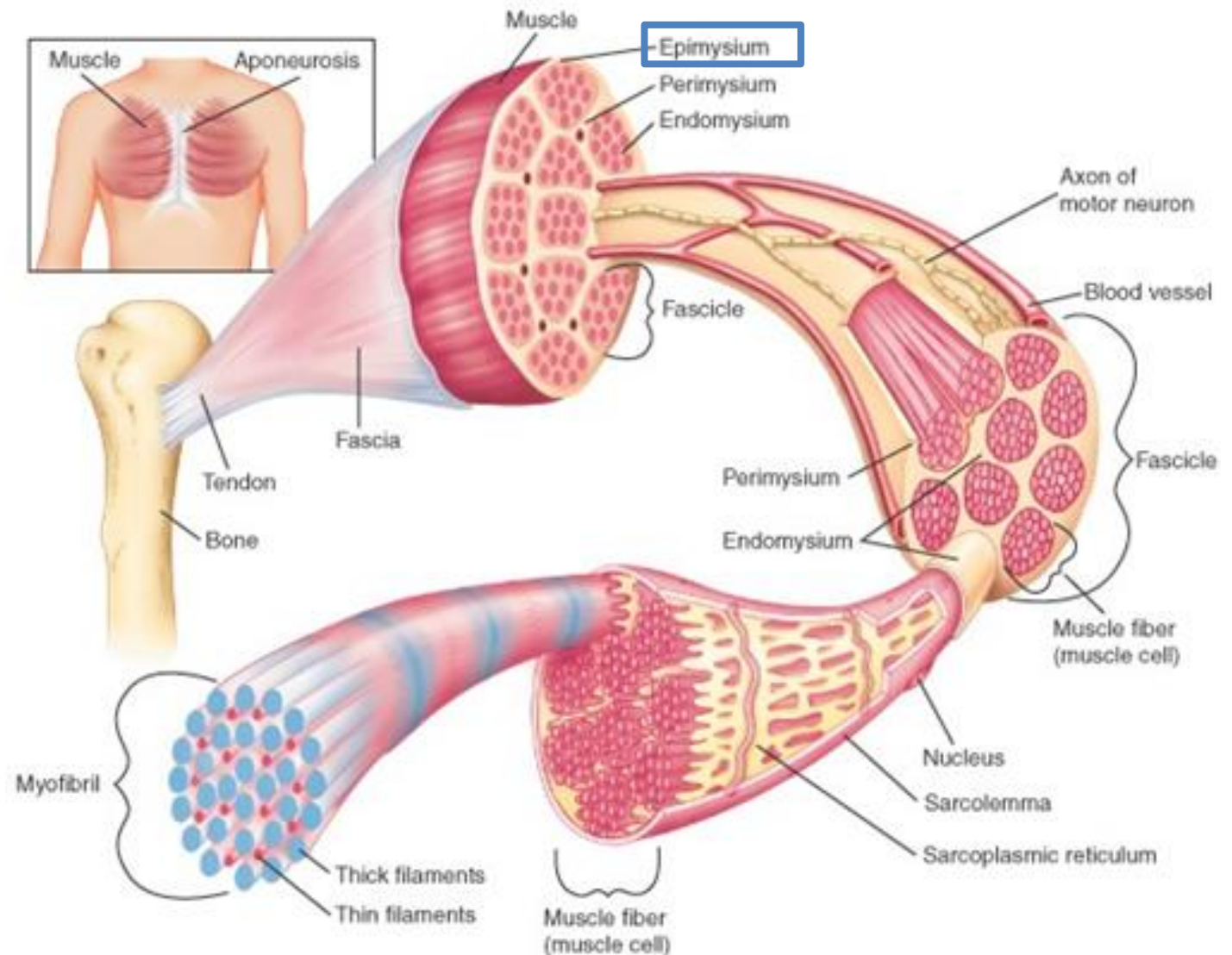
Fascia

- Surrounds and separates each muscle



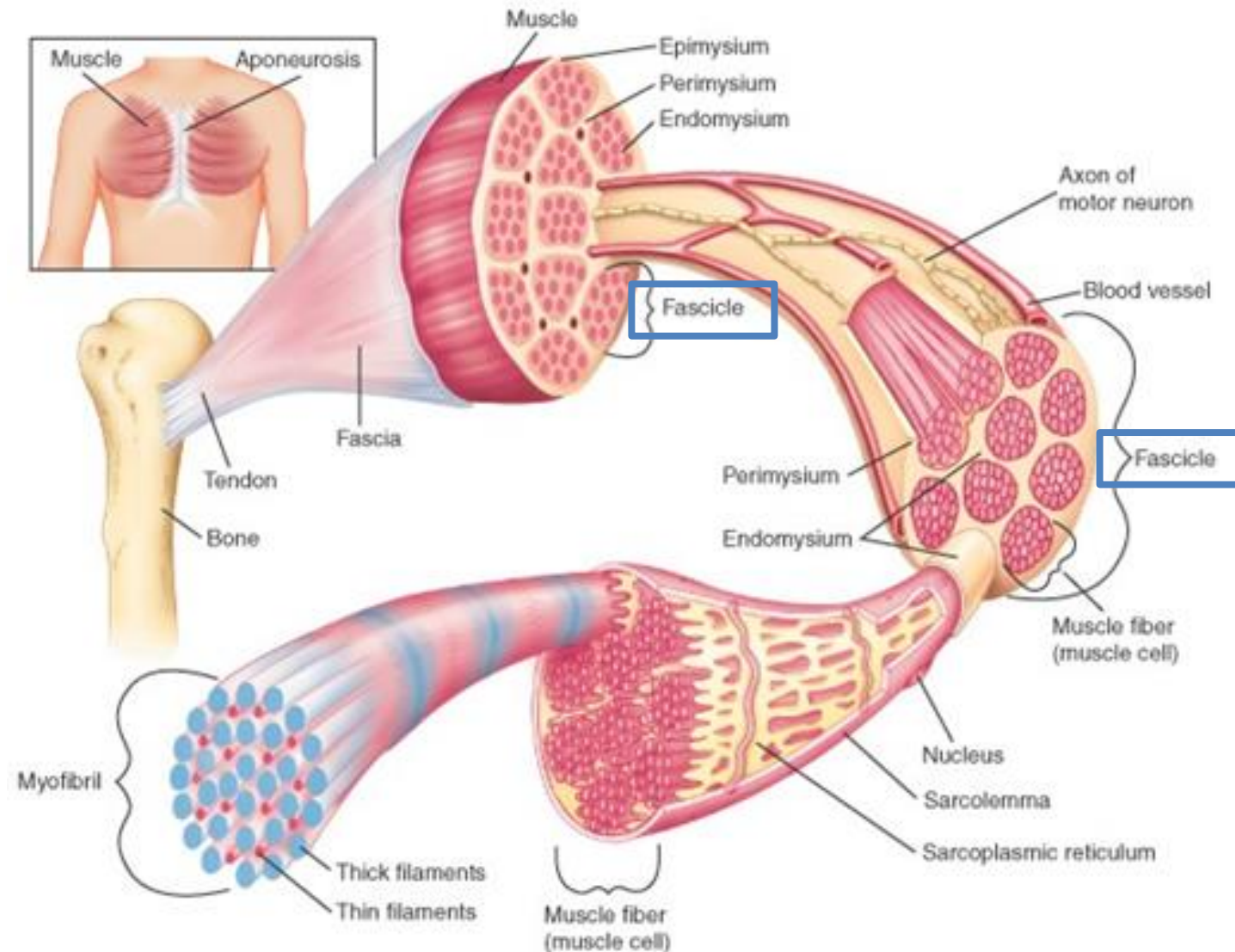
Epimysium

- Surrounds each muscle
- A protective sheath
- Divides the muscle into compartments



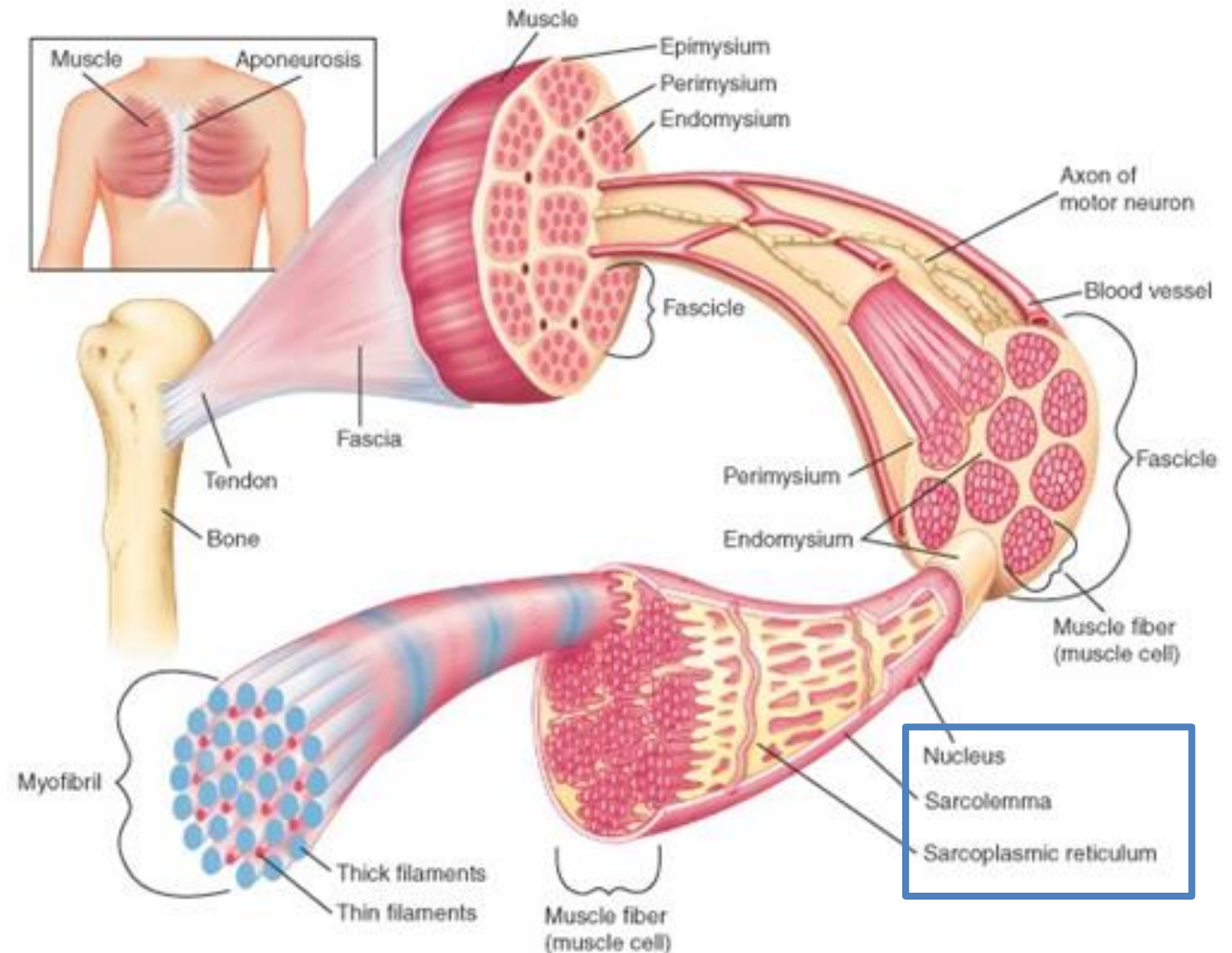
Fasciculus

- Bundle of fibers called a **fasciculus** found in each compartment
- Surrounded by a layer of tissue called **perimysium**
- Each fiber in the **fasciculus** is surrounded by a layer of tissue called the endomysium
- The coverings also contain blood vessels and nerves



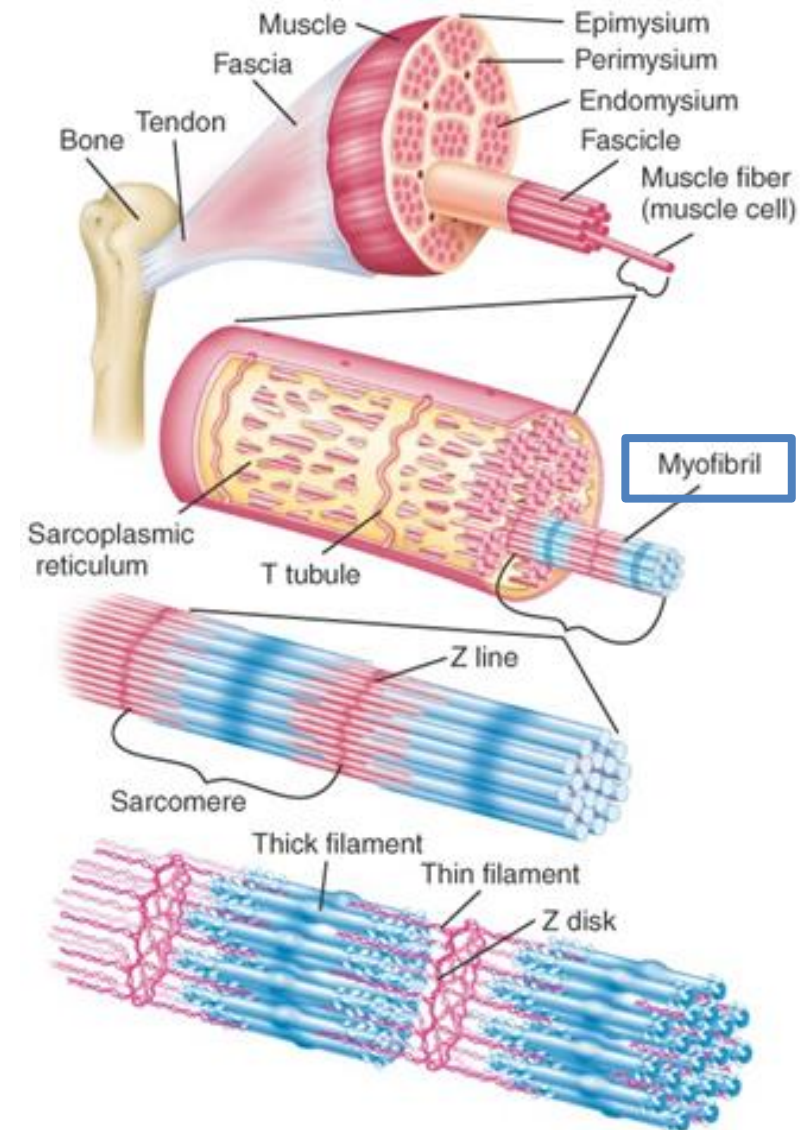
Muscle Fibers

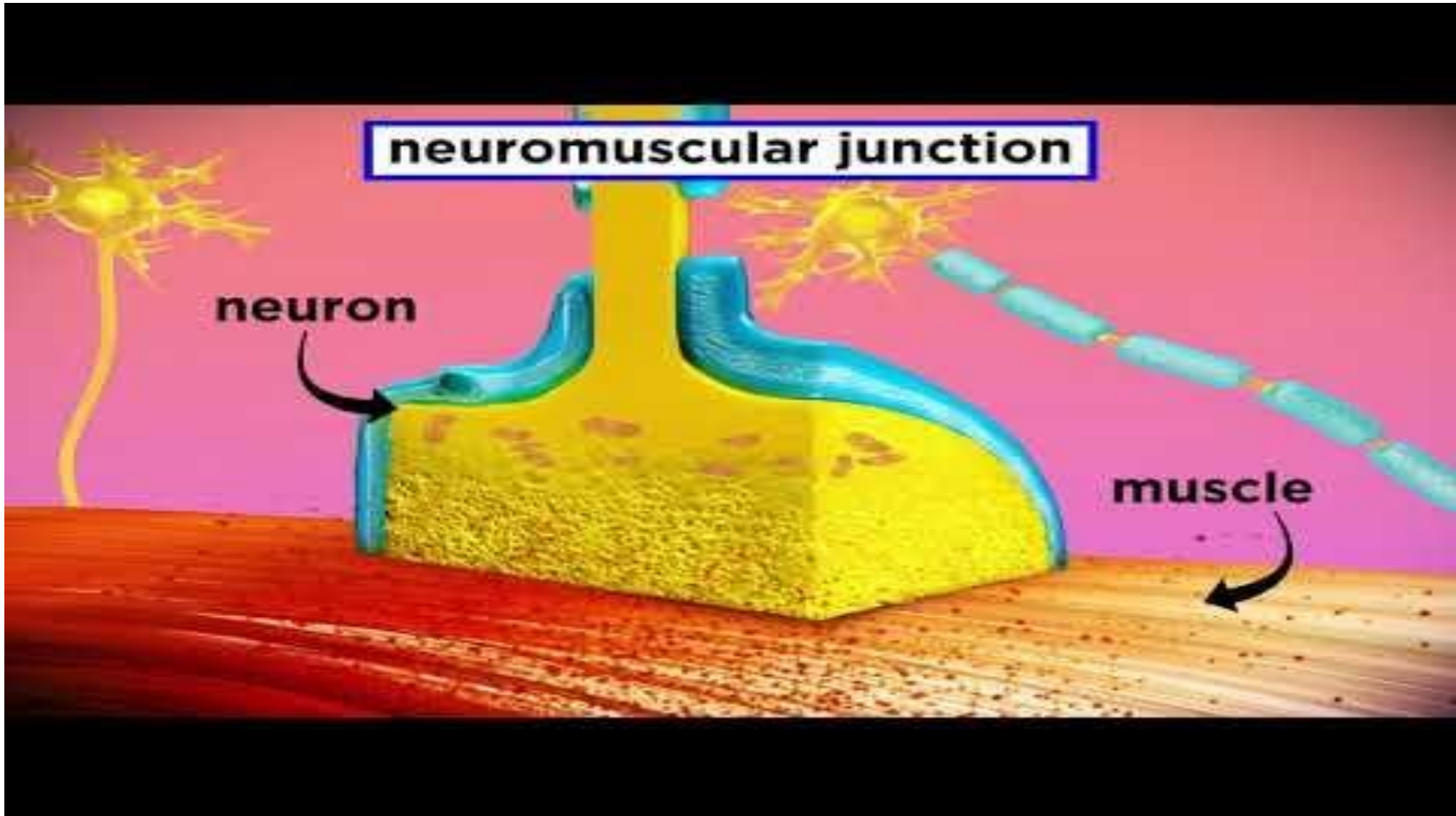
- Each fiber is a cylindrical cell
 - Sarcolemma (Cell membrane)
 - Sarcoplasm (Cytoplasm)
 - Sarcoplasm Reticulum (Endoplasmic Reticulum)
- The sarcolemma has multiple nuclei and mitochondria (for energy production)
- Inward extensions of the sarcolemma are called T-tubules



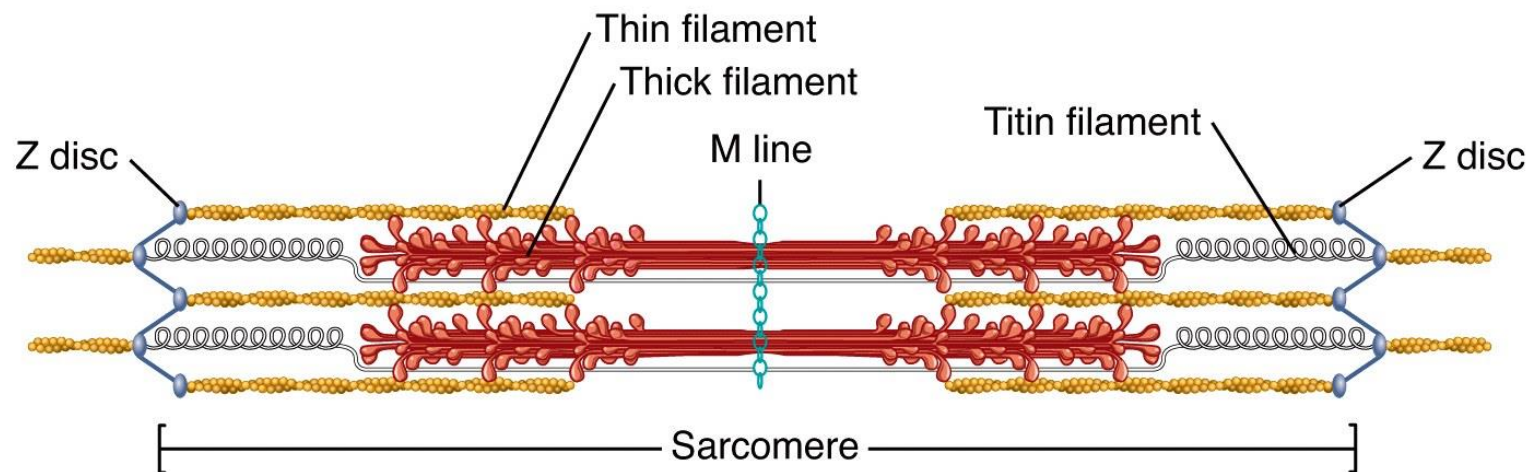
Myofibrils

- Built from three groups of proteins
 - Contractile proteins
 - Generate force during contraction
 - Regulatory proteins
 - Help switch the contraction process on and off
 - Structural proteins
 - Keep the thick and thin filaments in proper alignment and link the myofibrils to the sarcolemma and extracellular matrix





- The thin filaments are comprised mostly of the structural protein actin, and the thick filaments are comprised mostly of the structural protein myosin
- However, in both types of filaments, there are also other structural and regulatory proteins



- In the thin filaments actin proteins are strung together like a bead of pearls



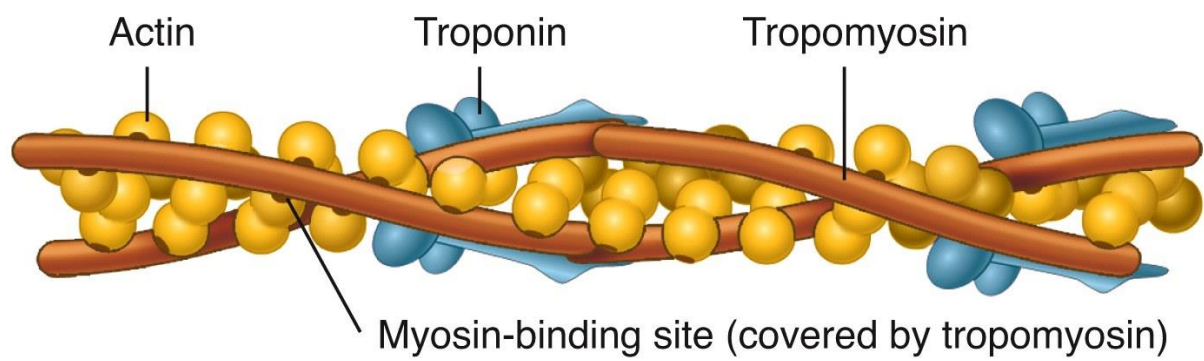
- In the thick filaments myosin proteins look like golf clubs bound together



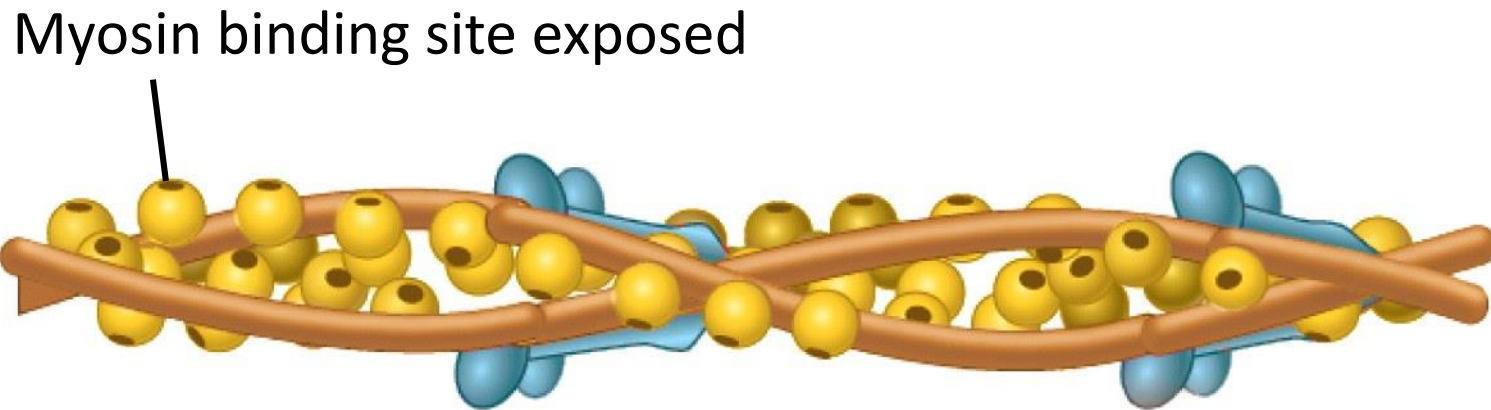
- The myosin binding sites on the actin proteins are readily visible.



- The regulatory proteins troponin and tropomyosin have cover the myosin binding

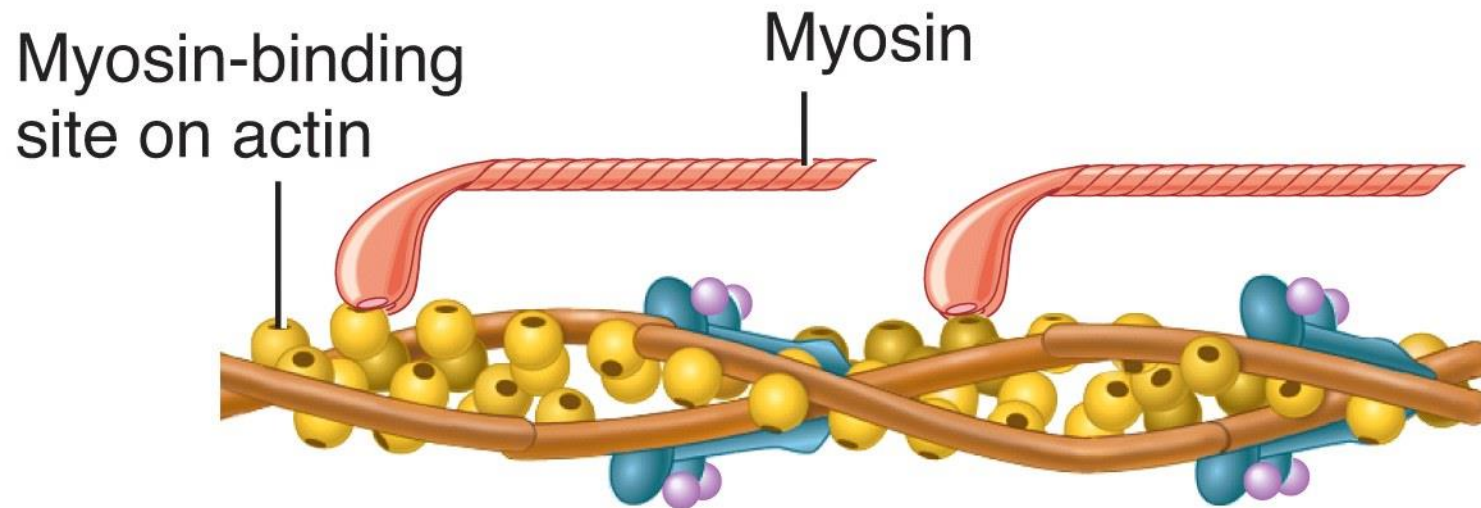


- The troponin-tropomyosin complex has slid down into the “gutters” of the actin molecule unblocking the myosin binding site

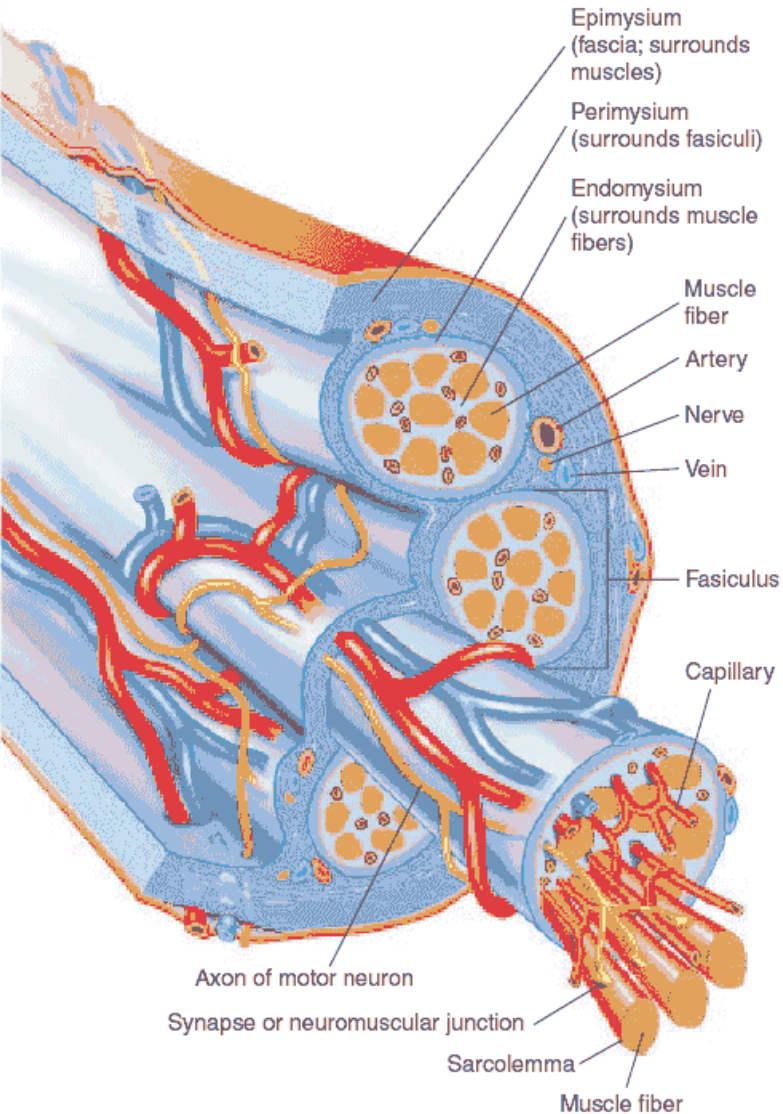


- The troponin-tropomyosin complex can slide back and forth depending on the presence of Ca^{2+}

- Ca^{2+} binds to troponin which changes the shape of the troponin-tropomyosin complex and uncovers the myosin binding sites on actin

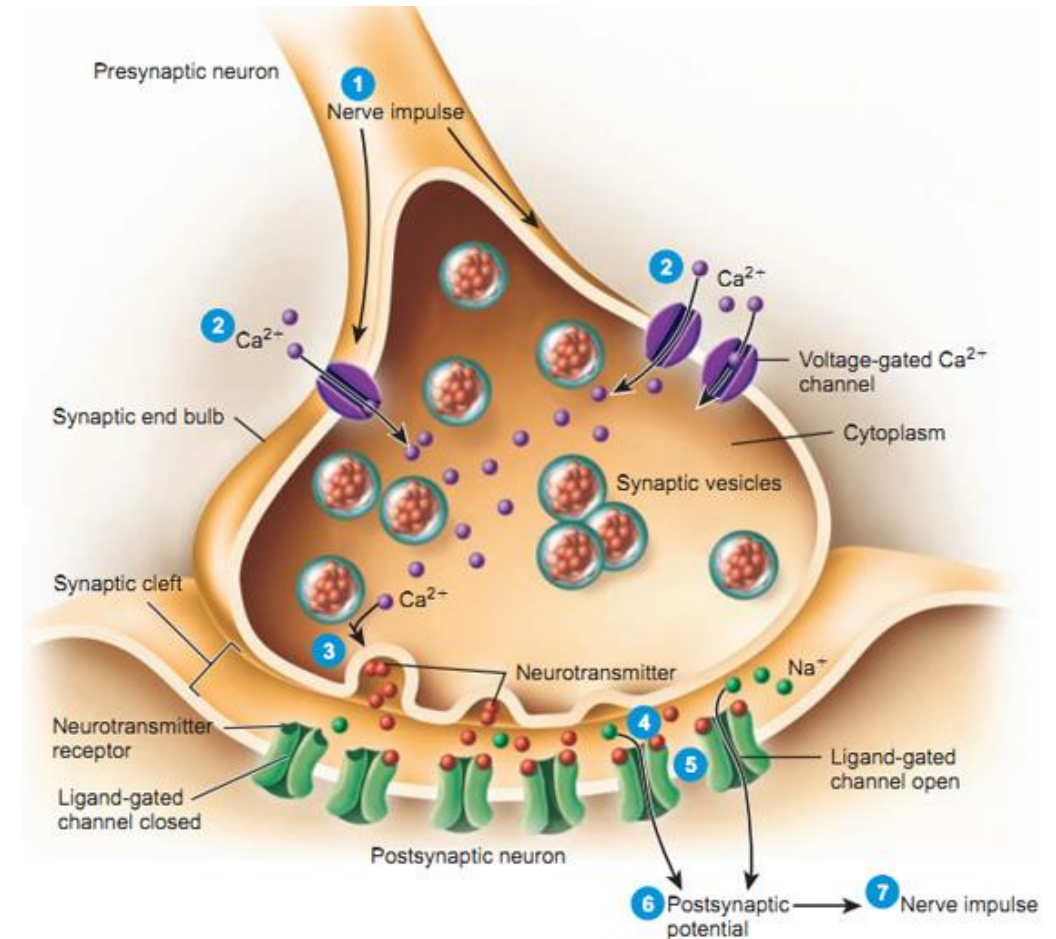


- Besides contractile and regulatory proteins, muscle contains about a dozen structural proteins which contribute to the alignment, stability, elasticity and extensibility of myofibrils
 - Titan is the third most plentiful protein in muscle, after actin and myosin - it extends from the Z disc and accounts for much of the elasticity of myofibrils
 - Dystrophin is discussed later as it relates to the disease of muscular dystrophy



- Have an abundant supply
- Before a muscle can contract it needs a stimulus
- This requires ATP
- Blood supply deliver O_2 and nutrients to produce this and remove the waste products
- One artery and one vein accompany each nerve

- Stimulated by specialized nerve cells called motor neurons
- The motor neuron and muscle(s) is called a motor unit
- Where the axon of the neuron meets the muscle is called the **neuromuscular junction**
- Between the two is a small depression in the muscle membrane called the synaptic cleft

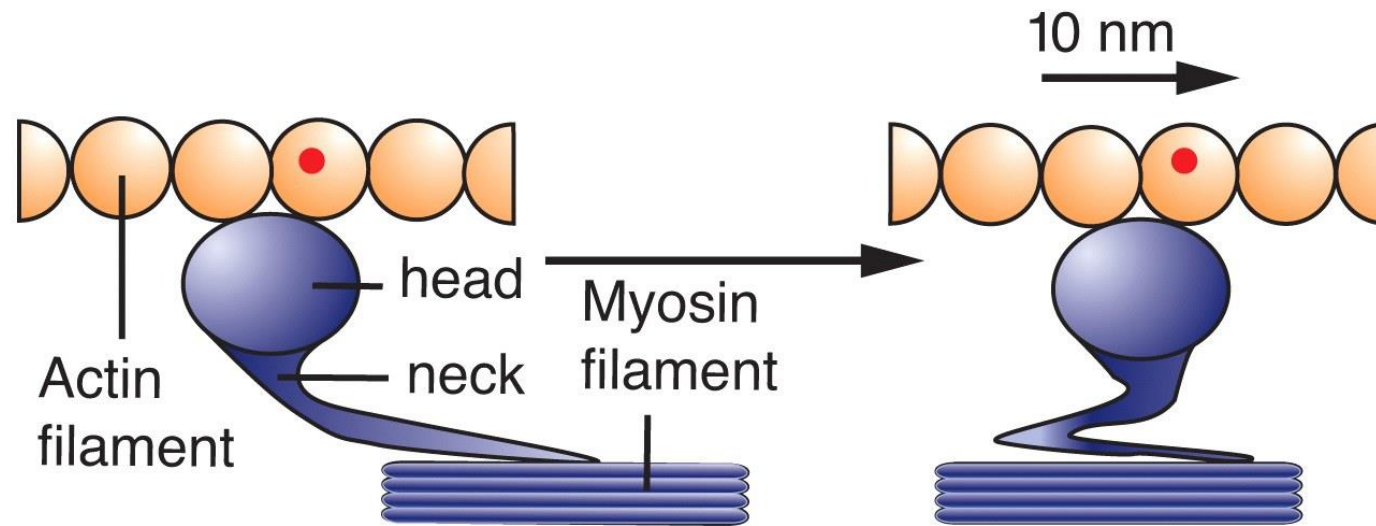


- ACh is contained within the synaptic vesicles of the axon
- Receptors for ACh are in the sarcolemma
- The combination results in a stimulus for contraction (an impulse) which travels along the sarcolemma into the T-tubules where a physiological change occurs causing a contraction
- The enzyme acetylcholinesterase deactivates the ACh at the synaptic cleft

- In a relaxed muscle fiber myosin receptor sites on the actin are inactive
- Heads on the myosin are also inactive and are bound to ATP
- Ca^{2+} is stored in the sarcoplasmic reticulum and has a low concentration in the sarcoplasm
- An impulse into the T-tubule cause release of Ca^{2+} from the SR into the sarcoplasm
- This rapid influx changes configuration of troponin on the actin fibers which exposes receptor sites

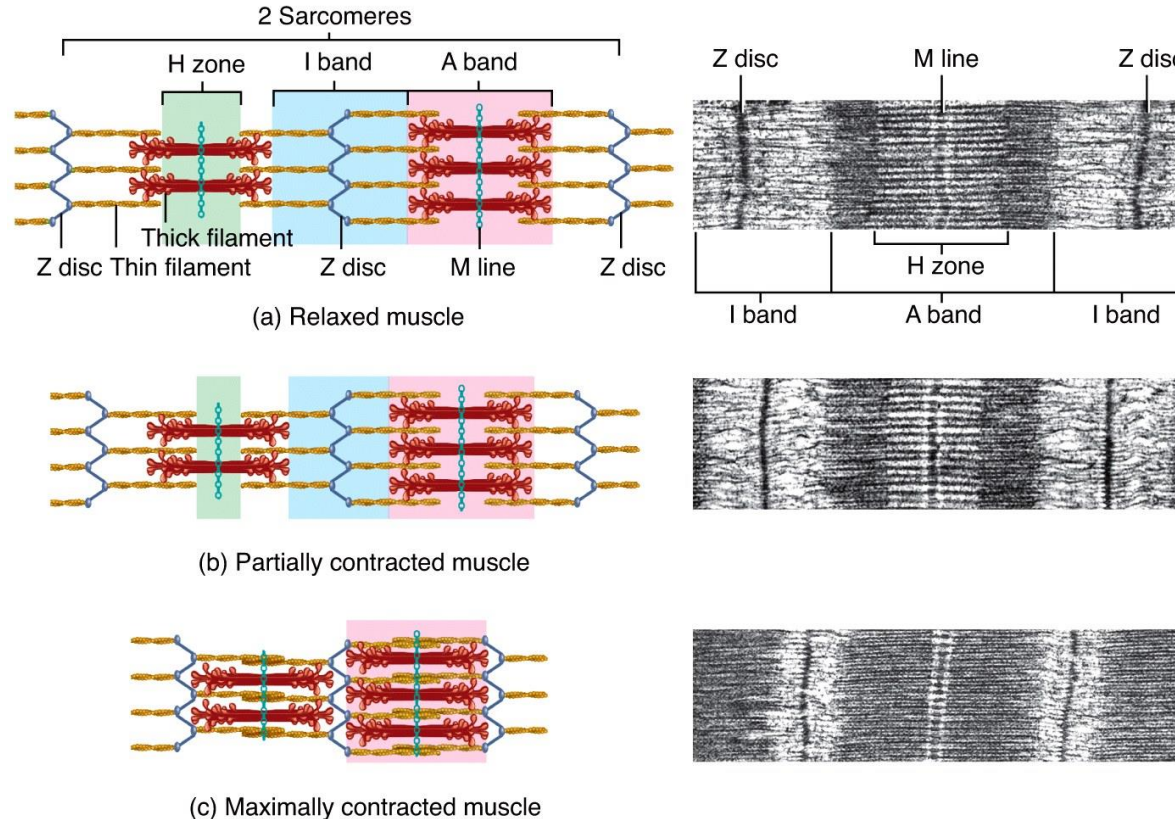
The Sliding-Filament Mechanism

- With exposure of the myosin binding sites on actin (the thin filaments)—in the presence of Ca^{2+} and ATP—the thick and thin filaments “slide” on one another and the sarcomere is shortened



The Sliding-Filament Mechanism

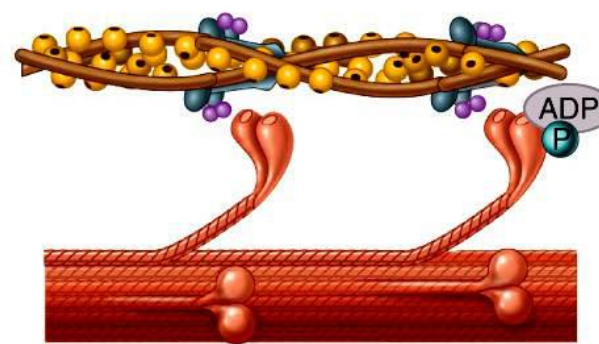
- The “sliding” of actin on myosin (thick filaments on thin filaments) can be broken down into a 4 step process



The Sliding-Filament Mechanism

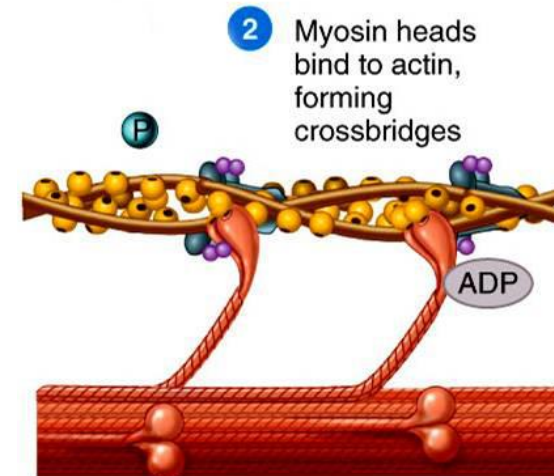
- Step 1: ATP hydrolysis

1 Myosin heads hydrolyze ATP and become reoriented and energized



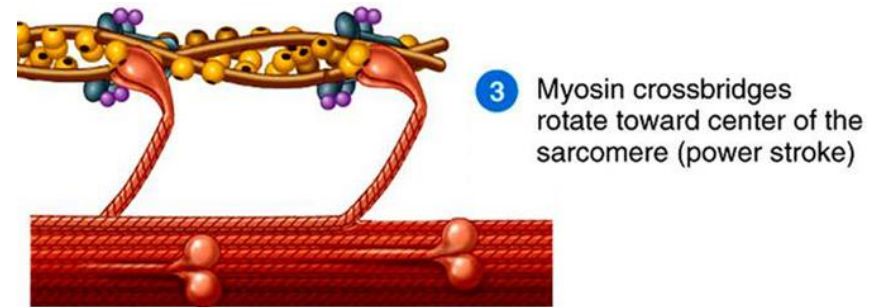
- Step 2: Attachment

2 Myosin heads bind to actin, forming crossbridges

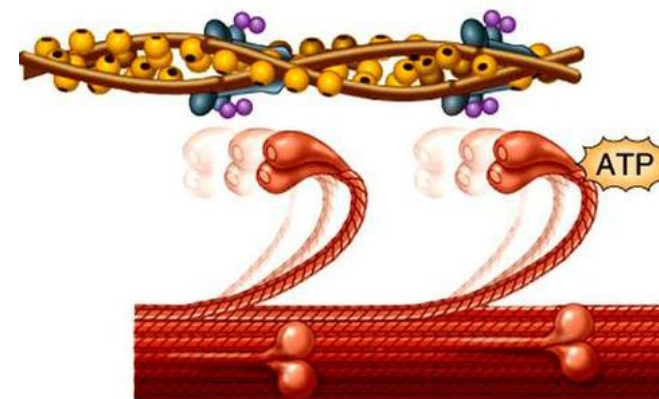


The Sliding-Filament Mechanism

- Step 3: Power Stroke

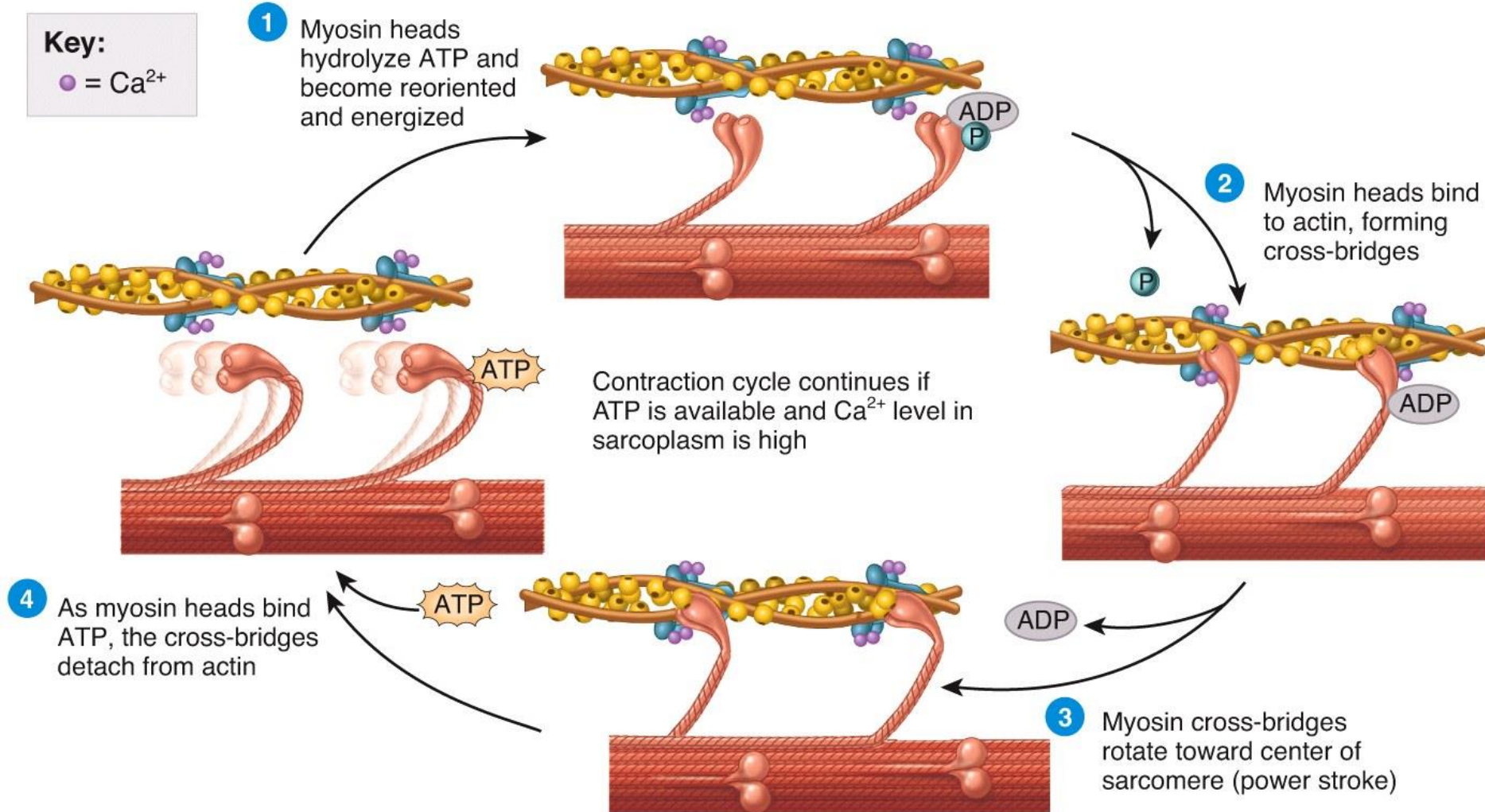


- Step 4: Detachment

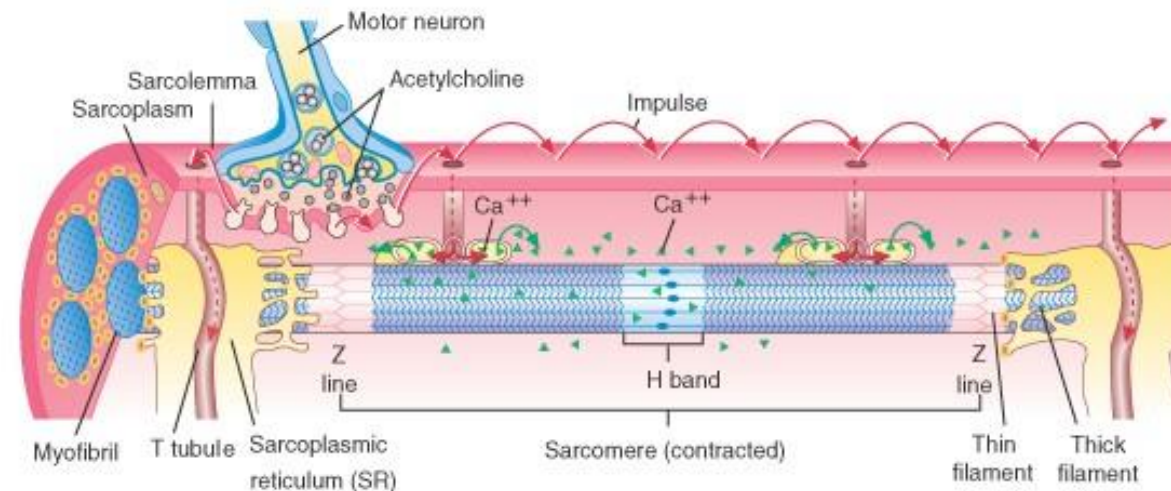


- 4 As myosin heads bind ATP, the crossbridges detach from actin
- ATP
- This diagram illustrates the detachment of a myosin crossbridge. A blue circle with the number '4' is next to the text 'As myosin heads bind ATP, the crossbridges detach from actin'. A yellow starburst labeled 'ATP' is shown next to the heads, indicating that ATP is binding to them. An arrow points from the starburst to the text.

The Sliding-Filament Mechanism

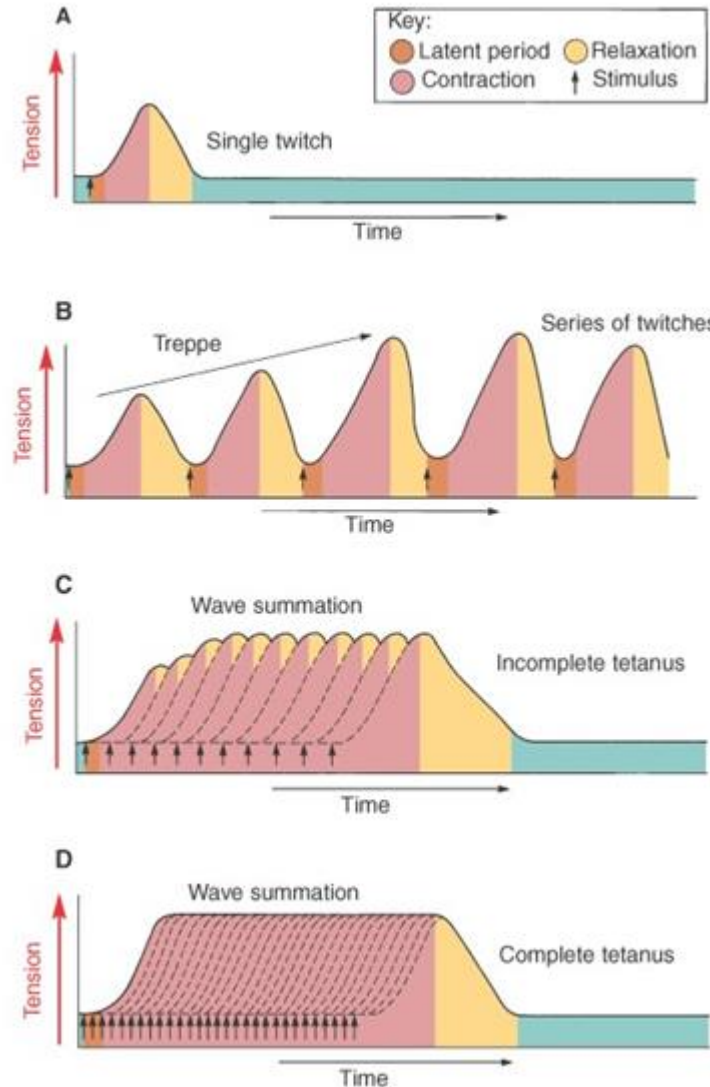


- As a result of the sliding filament mechanism the Z lines are pulled closer together shortening the sarcomere
- This does not shorten the myofilament
- New ATP on myosin reverse the reaction



- When the stimulation ceases, Ca^{2+} is actively transported into the SR
- This causes the receptor sites to close and ceasing the contraction
- Follows the all-or-none principle:
 - A sufficient stimulus is need to cause a contraction (threshold stimulus)
 - A greater stimulus will not produce greater contraction
 - Not enough will elicit no response (sub-threshold stimulus)

- **Does not** follow the All-or-none principle
- Varies due to work load
- Increase contraction is achieved by motor unit summation and wave summation
- A single stimulus causes a twitch (lab setting)
- 3 stages of contraction
 - Lag phase
 - Contraction
 - Relaxation



- A stimulus given during relaxation phase will cause stronger contraction, and continues to build to form a smooth contraction called tetany (multiple wave summation)
- Treppe (staircase) shows an increase in force with a stimulus of same intensity

Whole Muscle Contraction

ISOTONIC
Same tension; changing length

Eccentric
Muscle
lengthens



Concentric
Muscle
shortens

A

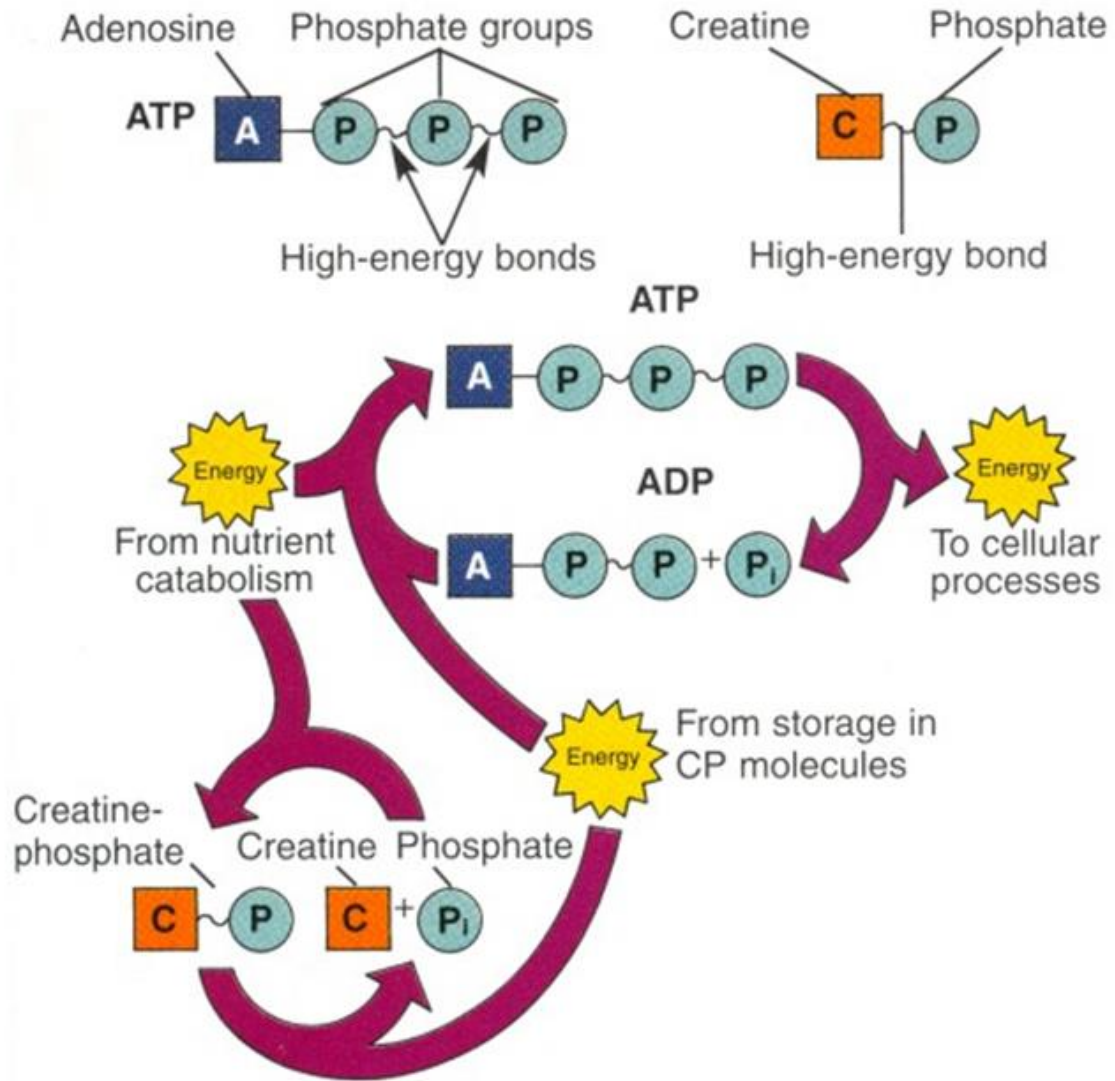
ISOMETRIC
Same length; changing tension

Relaxed

Contracting

B





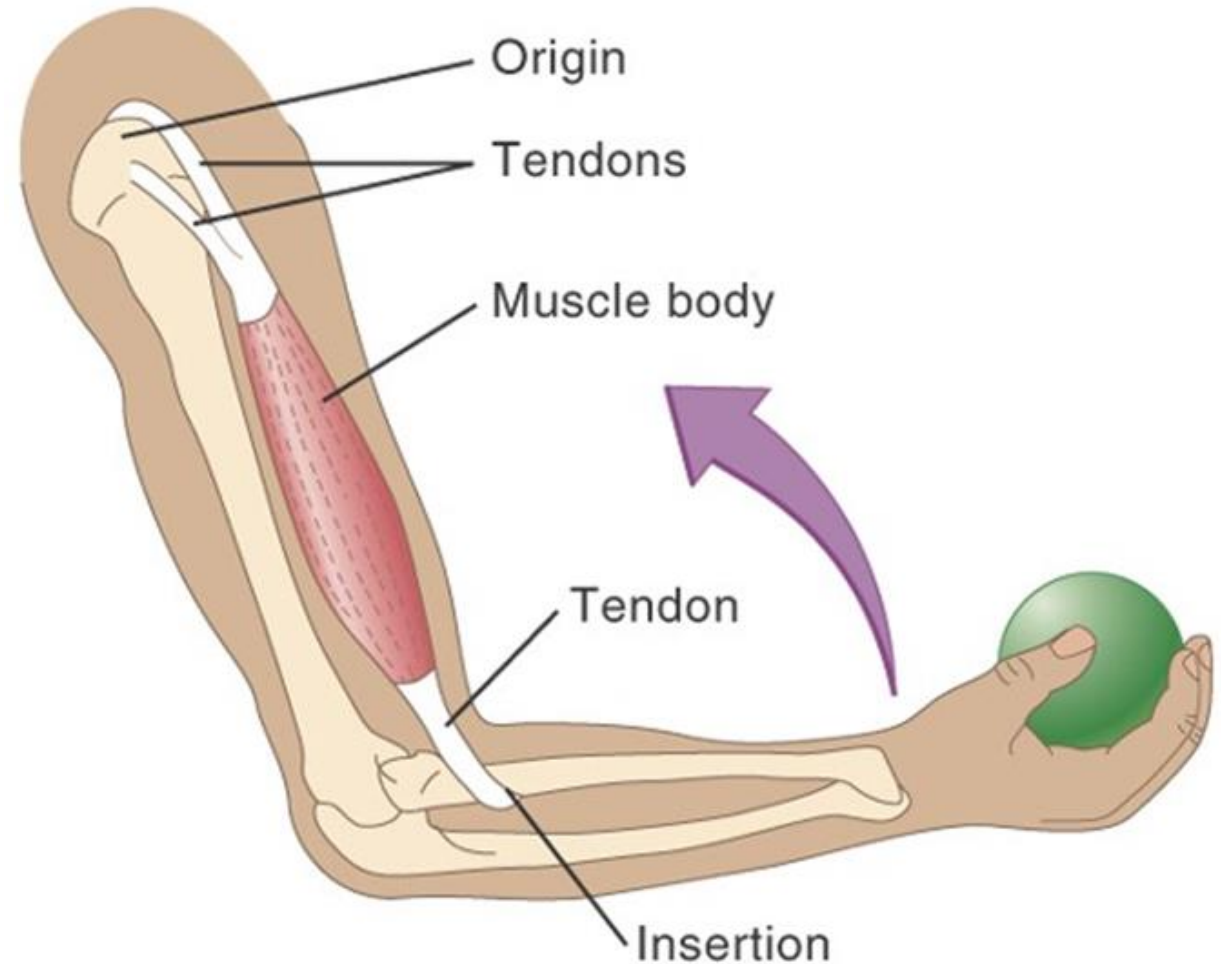
- Initial Source
 - ATP for the cross-bridge and active transport
 - Last only 6 seconds
- Second Source
 - Creatine Phosphate is used to instantaneously give its energy to ADP to synthesis ATP
 - If ATP is in excess it will convert to Creatine phosphate to store for later use
 - Lasts only 10 seconds

- Third Stage
 - Muscles use fatty acids and glucose for energy
 - Fatty acids found in blood
 - Glucose is a derivative of the glycogen found in the muscle
 - If oxygen available then the fats and glucose are broken down with aerobic metabolism (20 times more production)
- Fatty Acids or glucose + O₂ → CO₂ + H₂O + ATP
 - If oxygen is not available then glucose is the primary source of energy (anaerobic metabolism – happens at a faster rate)
- Glucose → lactic acid + ATP

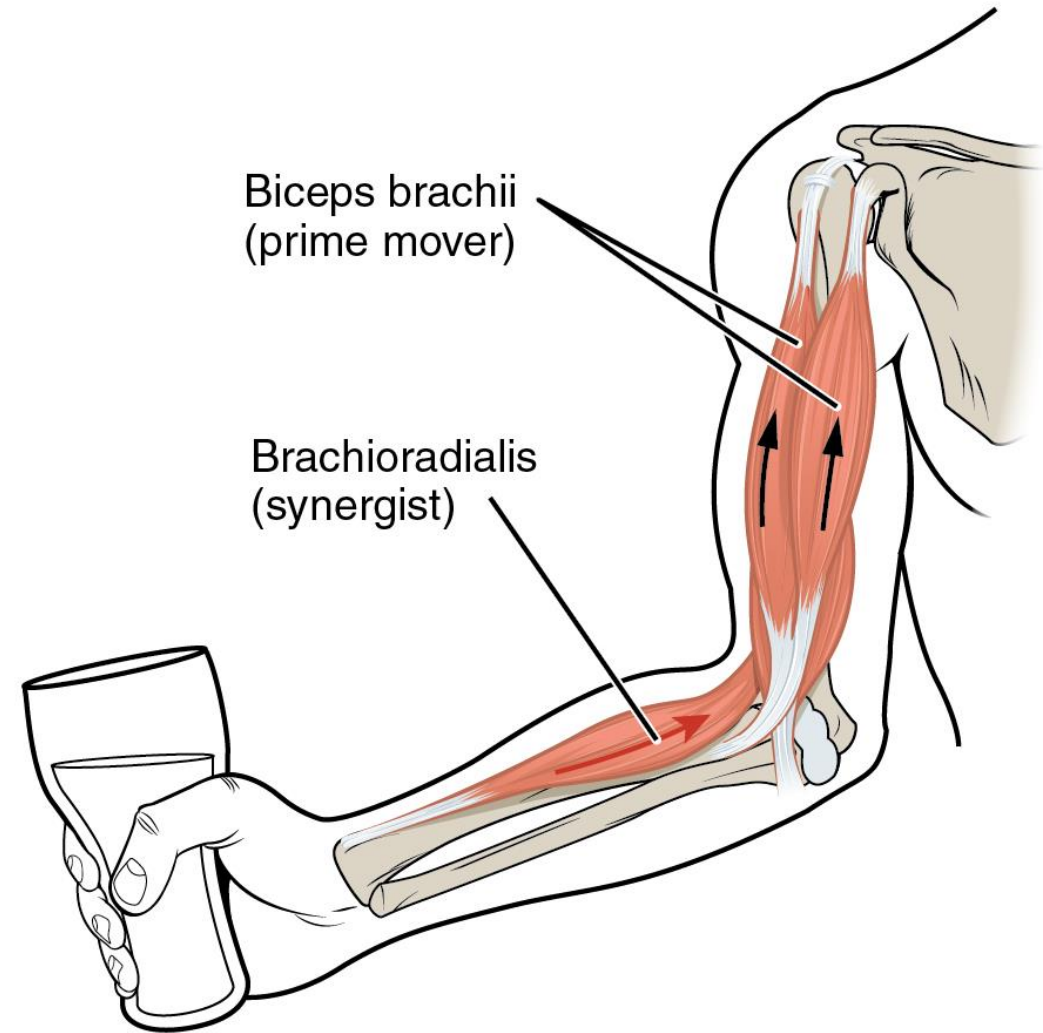
- Oxygen storage
 - Red fibers have myoglobin which has iron to bind with O_2
 - White Fibers do not contain myoglobin
- Lactic Acid
 - Excessive lactic acid is send to the liver when O_2 is available and converted and stored as glycogen
- Oxygen Debt
 - After strenuous exercise using anaerobic metabolism, ATP and creatine phosphate have to be replaced, this requires O_2
 - Is the additional O_2 needed to do this after exercise

- Some may attach to the bone by the epimysium to form a direct attachment
- Most have an extension of the epi, peri and endomysium to form a tendon or a flat sheet-like aponeurosis. This is an indirect attachment
- Muscles typically span a joint
- The attachment that remains relatively fixed is the origin, the other end is the insertion

- Origin
 - Attachment that remains relatively fixed
- Insertion
 - Moves in relation to the origin with a contraction



- Muscles will work in groups to produce a movement
- Prime Mover
 - Muscle has the primary role in the motion
- Synergist
 - Assist prime mover
- Antagonists
 - Muscles that oppose a particular movement



- Size
 - Vastus (huge), maximus (large)
- Shape
 - Deltoid (triangle), teres (round)
- Direction of fibers
 - oblique (diagonal)
- Location
 - Pectoralis (chest)

- # of origins
 - Biceps (2), triceps (3)
- Origin and Insertion
 - sternocleidomastoideous
- Action
 - Abductor, flexor