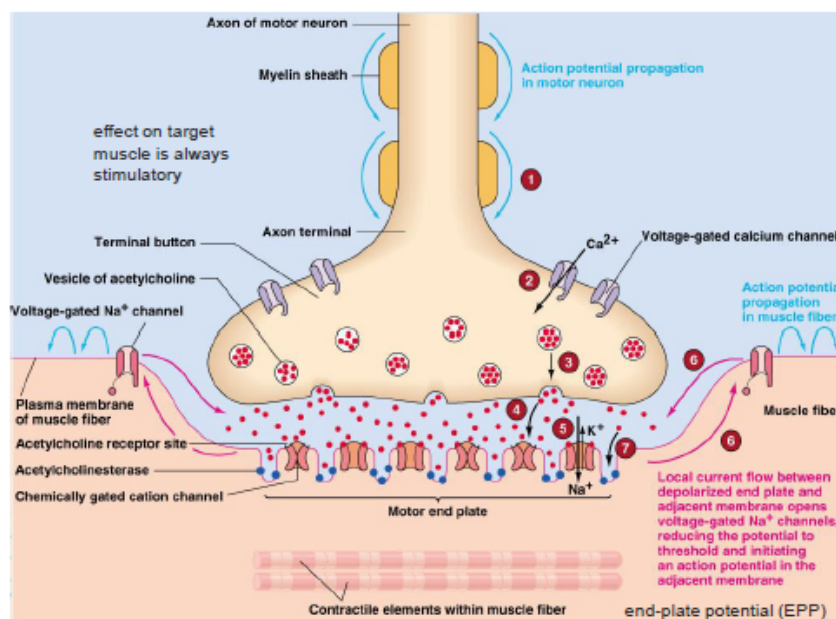


# PHYS30005: MUSCLE AND EXERCISE PHYSIOLOGY NOTES

## LECTURE 1: Functional Properties of Skeletal Muscle I

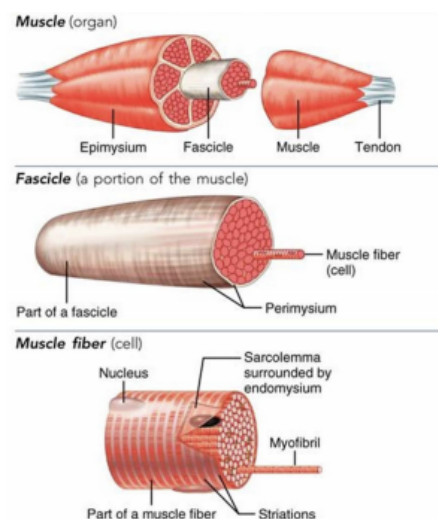
### Neuromuscular Junction

- Action potential travels from the pre-synaptic neuron, through the synaptic cleft then to the post-synaptic channel
- Voltage-gated calcium channels open when membrane reaches threshold and releases calcium into the pre-synaptic neuron
- Acetylcholine (ACh) diffuse across the cleft and binds with receptors on the motor end plate
- Cation channels on the motor end plate opens allowing sodium ( $\text{Na}^+$ ) in and potassium ( $\text{K}^+$ ) out
- Neurotransmitters diffuse across the synapse and bind to specific receptors which elicit a specific response

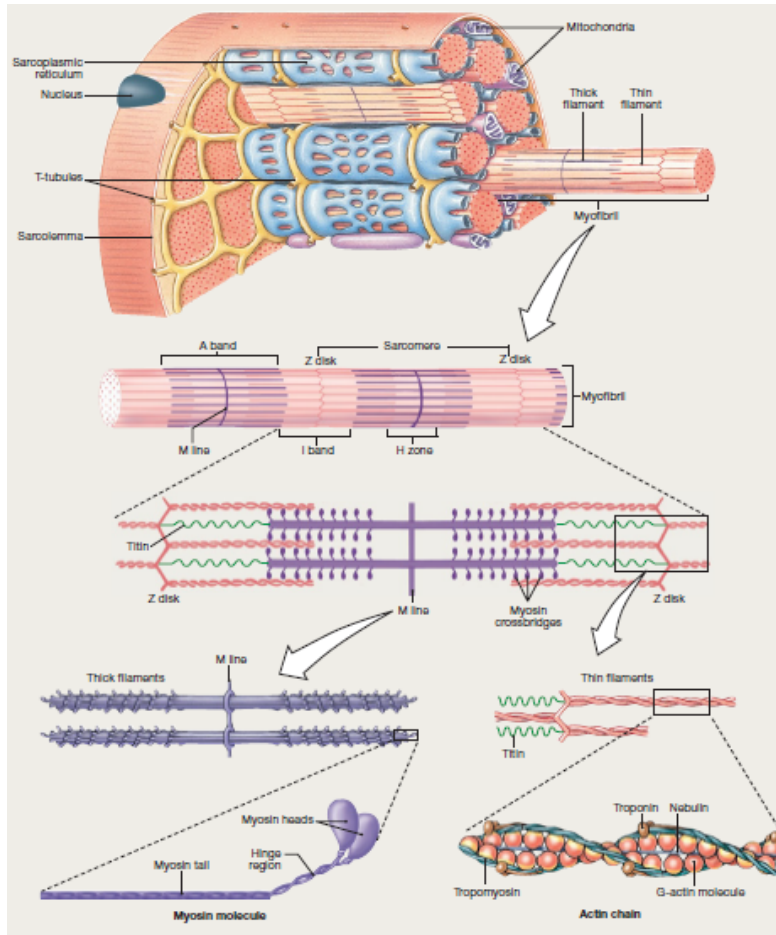


### Muscle Structure

- Muscles have an origin (proximal) and an insertion (distal) ends with tendons on both sides
- Muscle is made up of fascicles of fibres with each fibre made up of many myofibrils
- The myofibrils appear striated due to the dark (A) and light (I) bands in register
- Myofibrils ( $\sim 1\text{-}2\ \mu\text{m}$ ) occupy 80% of the fibre volume with 100s to 1000s of myofibrils that exist in each fibre
- Maintained in transverse register across the cell giving rise to the striation pattern

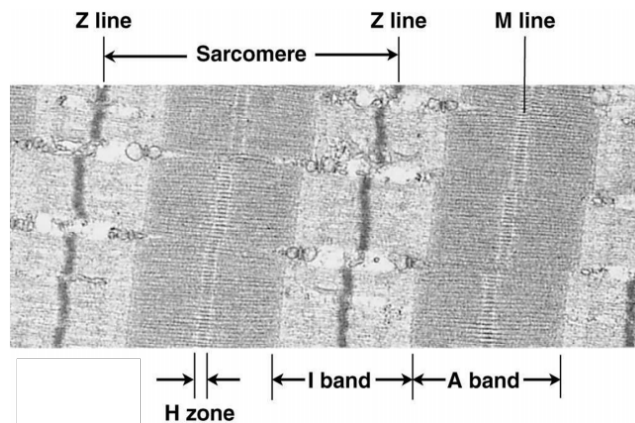


## PHYS30005: MUSCLE AND EXERCISE PHYSIOLOGY NOTES



- Sarcoplasmic reticulum is the internal storage of  $\text{Ca}^{2+}$
- Action potential from the neuromuscular junction travels down the t-tubular network and causes a change in electrical potentials
- This releases  $\text{Ca}^{2+}$  from the internal storage site and allows contraction to occur
- Sarcomeres are the basic contractile unit of the muscle
- Filaments slide within one another during contraction where the length of the filaments does not change

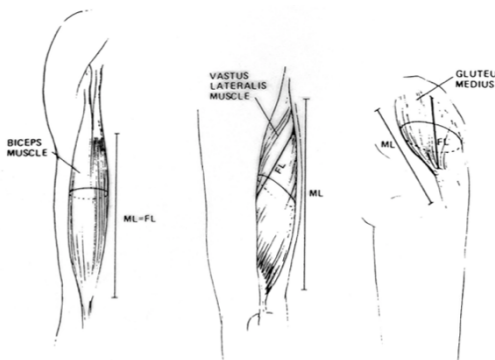
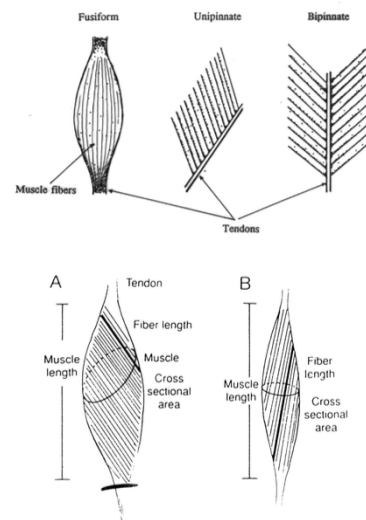
- The Z line will come towards the midline; muscle always have the desire to shorten
- Nebulin helps align actin
- Titin, body's largest protein, provides elasticity and stabilises myosin
- Muscle fibres are held together by a network of cytoskeleton protein that tethers the myofibril into position
- The two headed myosin molecule has two binding sites; one for action, the other for ATP
- The hinge region gives a level of flexibility when a myosin head attaches to an actin filament it can pull sarcomeres towards the midline
- Myosin head cannot bind to the binding sites on the actin molecules at rest due to the presence of troponin binding to tropomyosin
- When  $\text{Ca}^{2+}$  is released it binds to troponin to causes tropomyosin to pull away and allow the myosin head to attach



# PHYS30005: MUSCLE AND EXERCISE PHYSIOLOGY NOTES

## Arrangement of Fibres

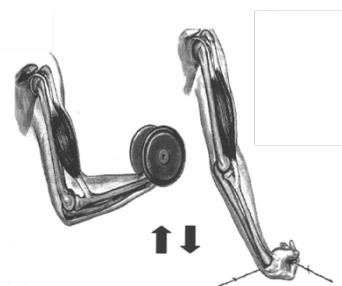
- The arrangement of fibres relative to the axis of force generation
- Fibre length (FL) is rarely the same as muscle length (ML)
- Most fibres insert obliquely into the tendon which represent a feather arrangement are called either pinnation or pennation
- Allows more fibres to be packed in which increases the effective muscle cross-sectional area (CSA)
- Fusiform: Fibres run the length of the muscle (from one tendon to the other)
- Unipennate: A central tendon with fibres running into the midline
- Muscle A is fatter compared to to Muscle B and this more fibres can be packed into that area to produce a greater contraction force
- Force production in proportional the muscle CSA
- Different muscles are required for different tasks; some for dexterity others for force and power



- Biceps: Fusiform; fibre length is almost the size of the muscle length
- Vastus laterallis: Overall muscle length is overall much shorter than the actual muscle length

## Muscle Action

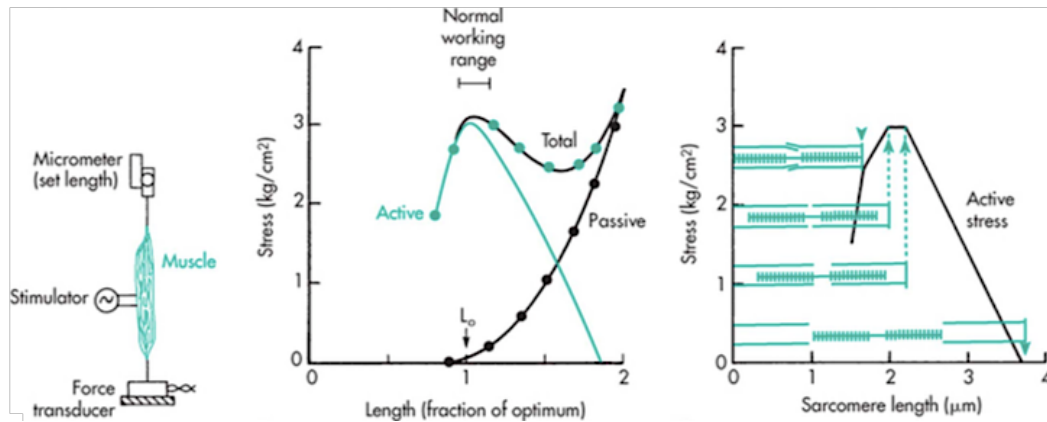
- Isometric Contraction: Any contraction that creates force without moving a load (immovable object)
- Isotonic (concentric) Contraction: Any contraction that creates force and moves a load
  - Every isotonic contraction begins with an isometric phase
  - The force exceeds the weight and overcomes the load, muscle shortens and moves to an isotonic contraction
- Eccentric contraction: Exerting force, muscle still lengthening – load greater than the force exerted by the muscle
- If the force developed by the muscle is greater than the load on the muscle, a shortening or (miometric, concentric) action occurs
- When the force developed by the muscle and the load are equivalent, or the load is immovable, a fixed-end or isometric action occurs
- When the load on the muscle is greater than the force developed by the muscle the muscle is stretched, producing a lengthening (plyometric or eccentric) action



# PHYS30005: MUSCLE AND EXERCISE PHYSIOLOGY NOTES

## Length-Tension Relationship

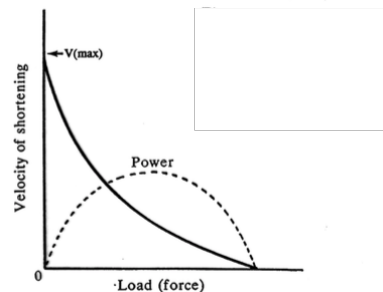
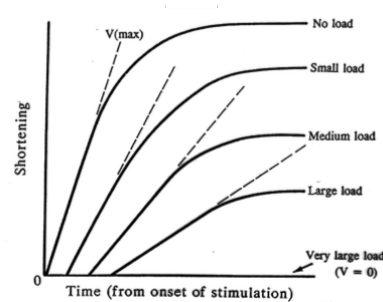
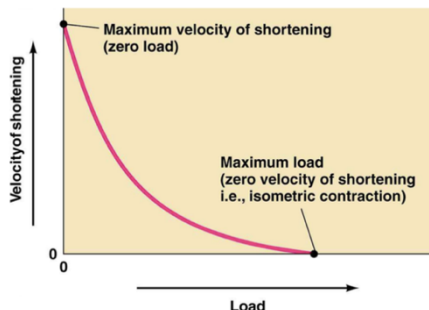
- Relationship between actin and myosin filament overlap and isometric force
- Maximally stimulate muscle at a number of discrete lengths (sarcomere lengths) and the record  $P_0$



- Total: Addition of active and passive
- Ascending limb: Sarcomere length short; overlap not optimal between actin and myosin; maximum active force compromised
- Plateau region: Sarcomere length optimal; overlap optimal between actin and myosin; ideal for maximum force
- Descending limb: Sarcomere length long or stretched; little or no overlap between actin and myosin; maximum force compromised

## Force-Velocity Relationship

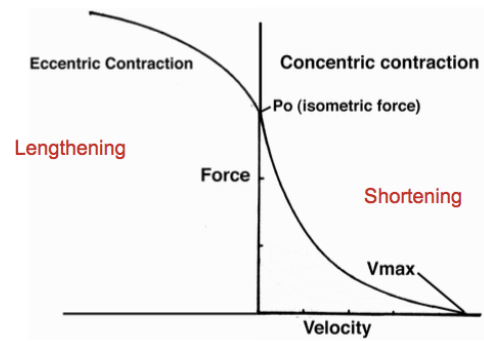
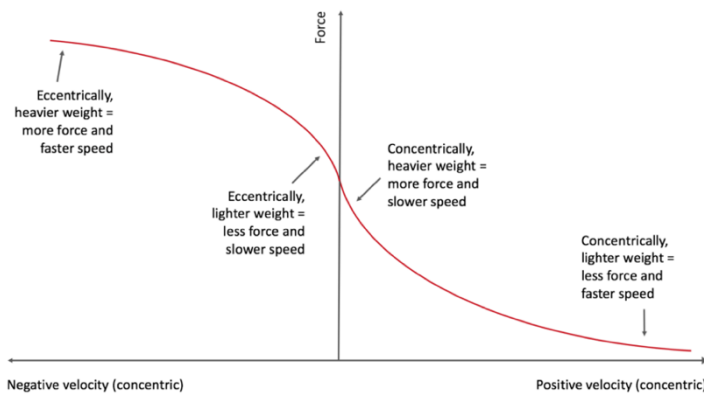
- Muscle allowed to shorten against a constant load and its velocity is plotted against resistive force
  - Heavier the load, slower the lift
- There is a latent period where muscle is contracting isometrically until sufficient tension has been produced to equal the load
- Not only is the rate of shortening reduced for heavy loads, the amount of shortening is decreased



- $V_{max}$ : The maximum velocity of shortening
- Related to muscle fibre type distribution and muscle architecture
- Power = Load x Velocity
- Power will be zero when there is no load on the muscle and when the load is so heavy it cannot be moved at all

# PHYS30005: MUSCLE AND EXERCISE PHYSIOLOGY NOTES

- Success in many sports is dependent upon maximal power output



## Muscle Contraction

Muscle Fibre Activation	
<p>A diagram showing the neuromuscular junction. An axon terminal of a somatic motor neuron (containing vesicles of ACh) is at the top. ACh is released into the motor end plate. An action potential (red arrow) travels down the T-tubule. This activates DHP (Dihydropyridine) receptors, which are coupled to RyR (Ryanodine) receptors on the sarcoplasmic reticulum. This causes Ca<sup>2+</sup> to be released from the sarcoplasmic reticulum into the cytoplasm. Ca<sup>2+</sup> binds to troponin, allowing actin-myosin binding. Labels include: Axon terminal of somatic motor neuron, ACh, Motor end plate, T-tubule, Sarcoplasmic reticulum, Z disk, DHP, RyR, Ca<sup>2+</sup>, Troponin, Myosin head, Actin, Troponin, Myosin thick filament, and M line.</p>	<ul style="list-style-type: none"> <li>- Somatic motor neuron releases ACh at neuromuscular junction</li> <li>- Initiates an action potential along cell membrane</li> <li>- T-tubule are extensions into the surface membrane of muscle fibre</li> <li>- Couples with the T-tubule is the sarcoplasmic reticulum loaded with calcium</li> <li>- Allow both sodium and potassium ions to cross the membrane, however sodium influx exceeds potassium efflux as the electrochemical driving force is greater for sodium</li> </ul>
<p>A diagram showing the sarcoplasmic reticulum releasing Ca<sup>2+</sup> into the cytoplasm. Labels include: Ca<sup>2+</sup> released and Myosin thick filament.</p>	<ul style="list-style-type: none"> <li>- Action potential propagates down the T-tubule</li> <li>- Ryanodine receptors open releasing calcium into the cytoplasm</li> <li>- Calcium binds to troponin allowing actin-myosin binding</li> </ul>
<p>A diagram showing Ca<sup>2+</sup> being pumped back into the sarcoplasmic reticulum by Ca<sup>2+</sup>-ATPase. Labels include: Ca<sup>2+</sup> releases, ATP, and Ca<sup>2+</sup>.</p>	<ul style="list-style-type: none"> <li>- Sarcoplasmic Ca<sup>2+</sup>-ATPase pumps Ca<sup>2+</sup> back into sarcoplasmic reticulum</li> <li>- Decrease in calcium concentration causes calcium to unbind from troponin</li> <li>- Tropomyosin re-covers binding site</li> <li>- Myosin heads return back to relaxed position</li> </ul>