

The Skin and
MUSCULOSKELETAL
System



PHYSIOLOGY

SLIDES
SHEET

DOCTOR: Dr. M. Khatatbeh
DONE BY:

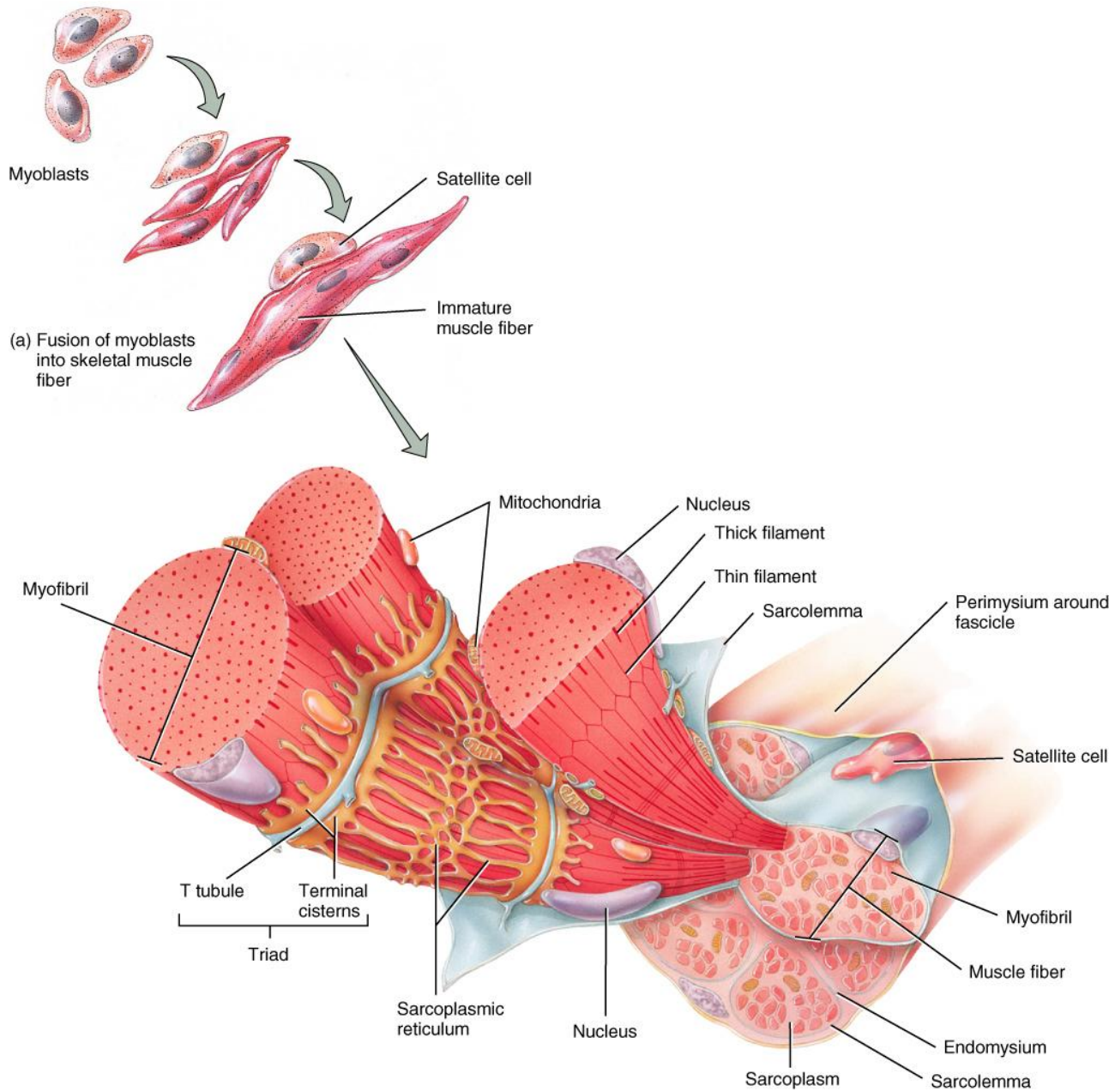
Number # 2-Types of Muscle

Muscle Physiology

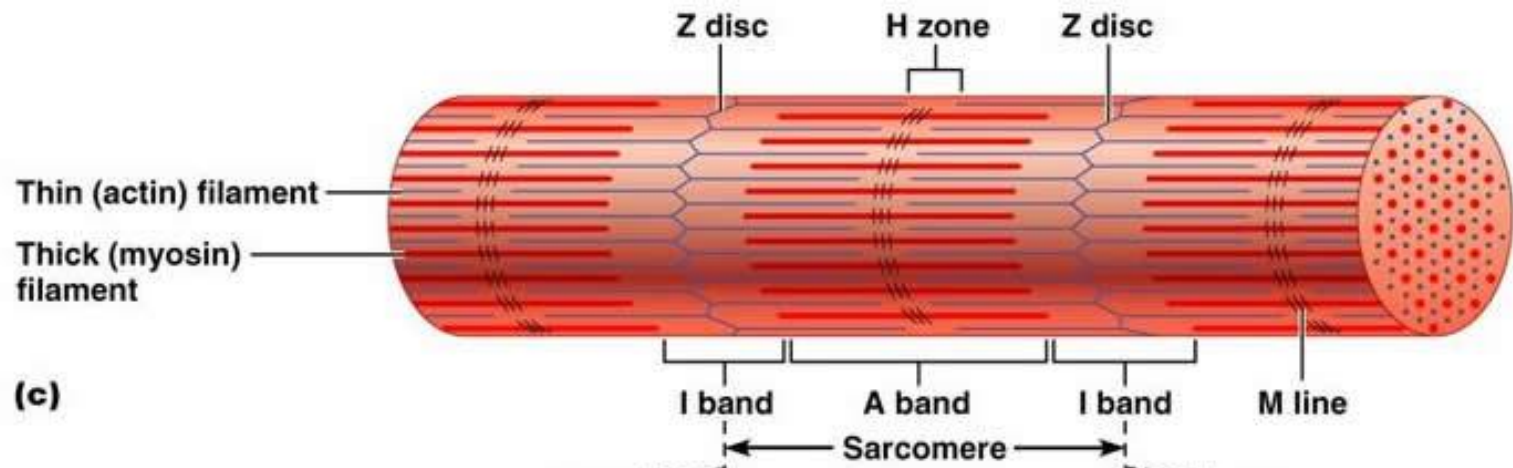
**Ref: Guyton,
chapters: 6,7,8**

Types of Muscle

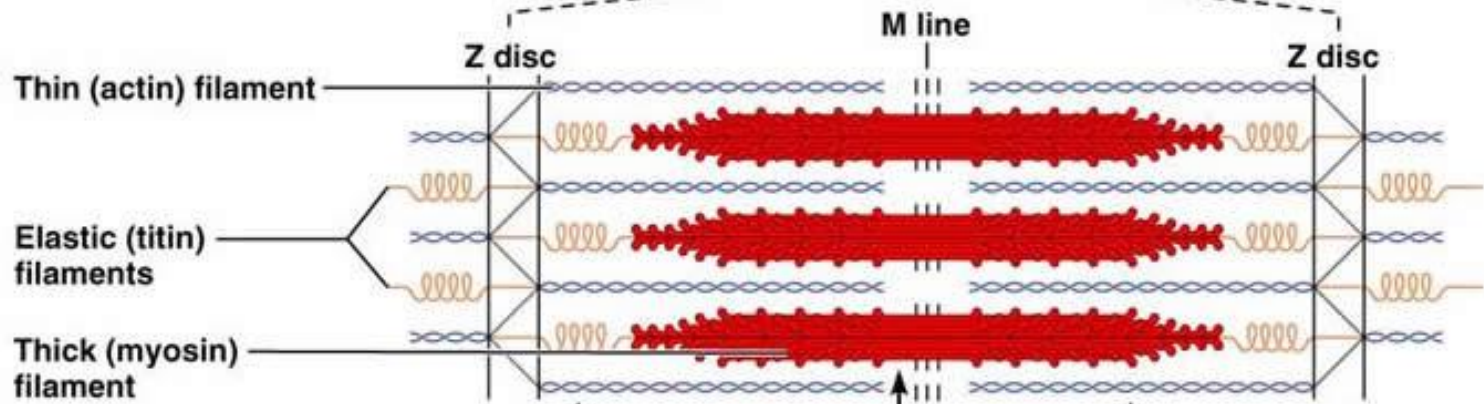
Fig. 10.03



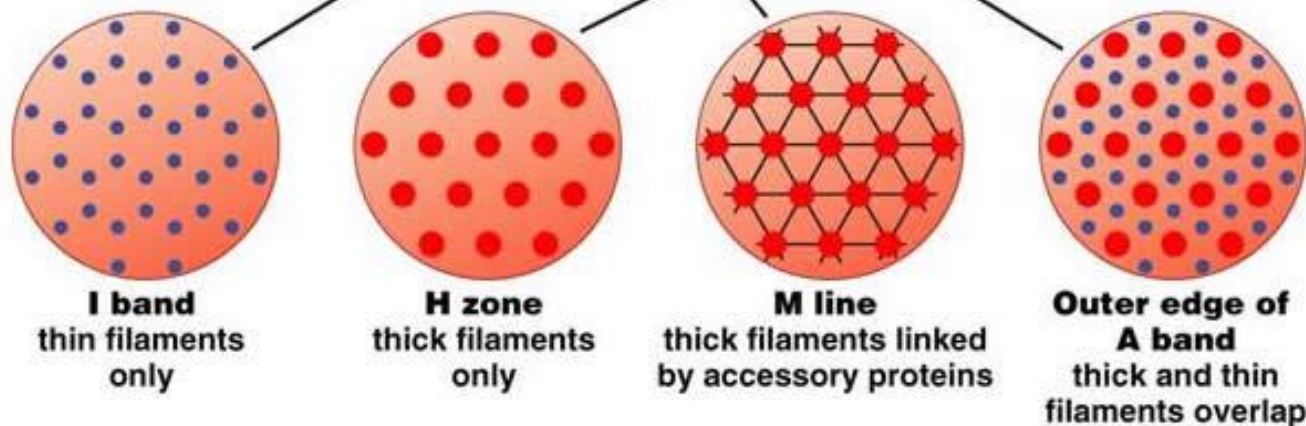
(b) Details of several myofibrils



(c)



(d)



(e)

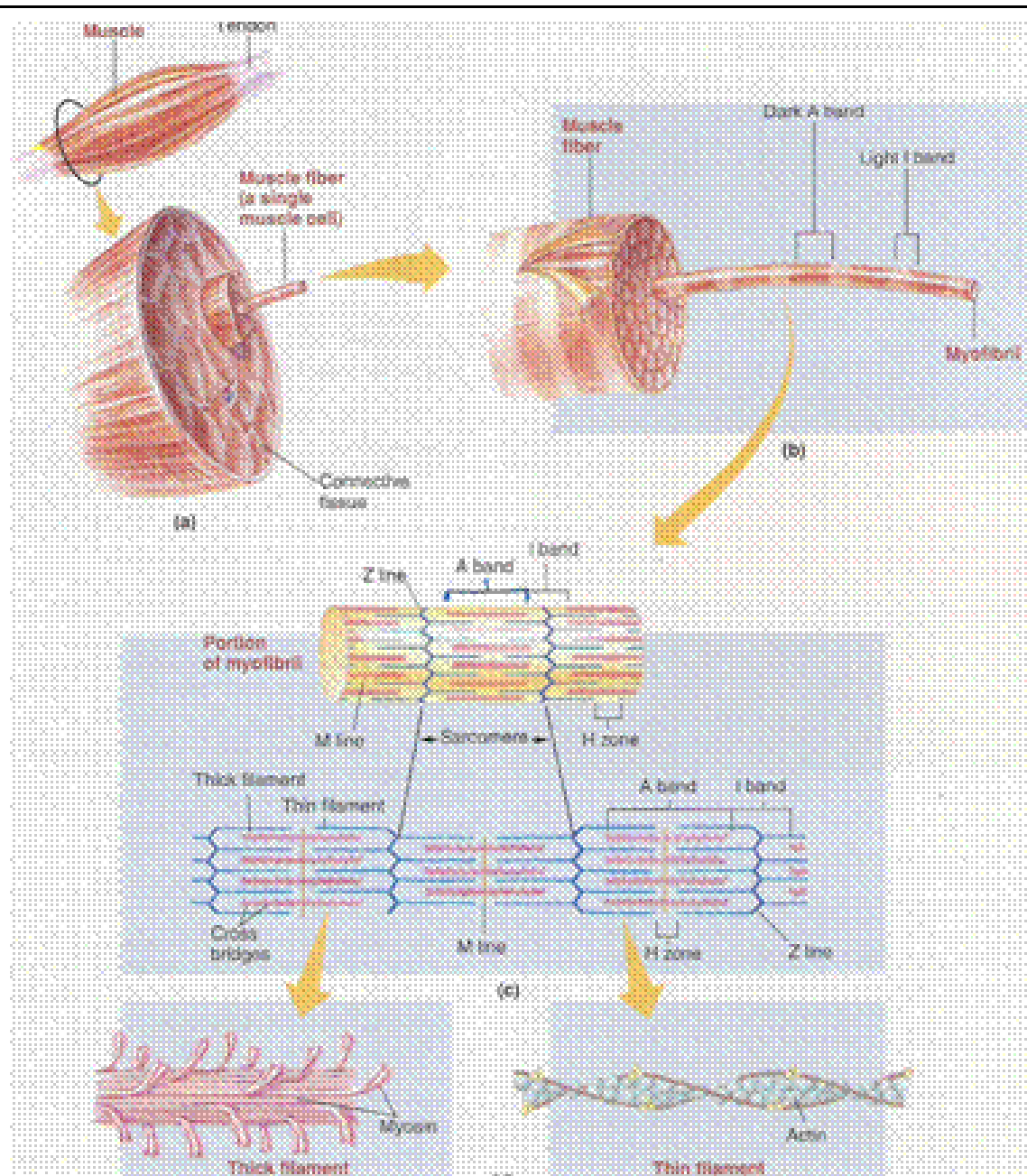


Figure 2. The components of a muscle.

Taken from Sherwood, 2004.

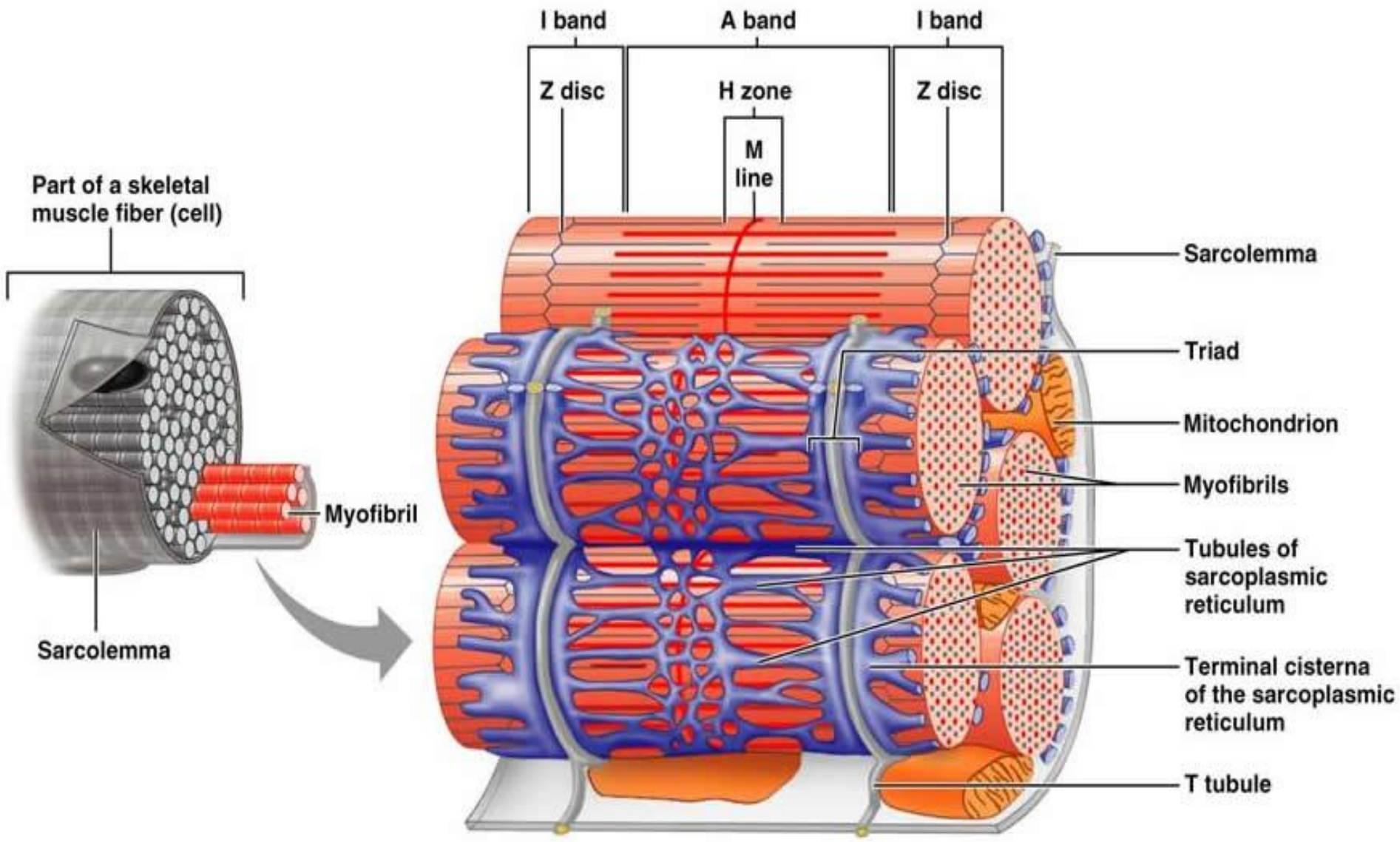
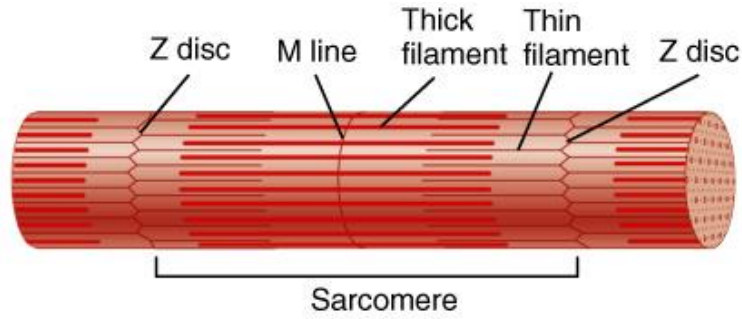
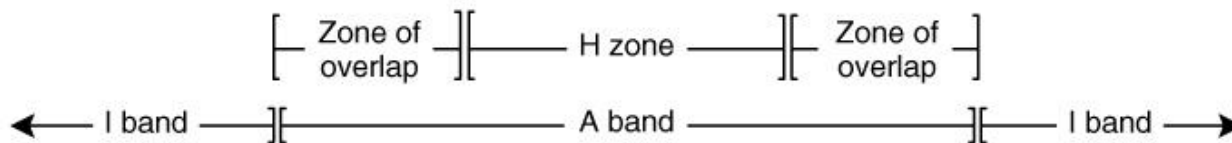
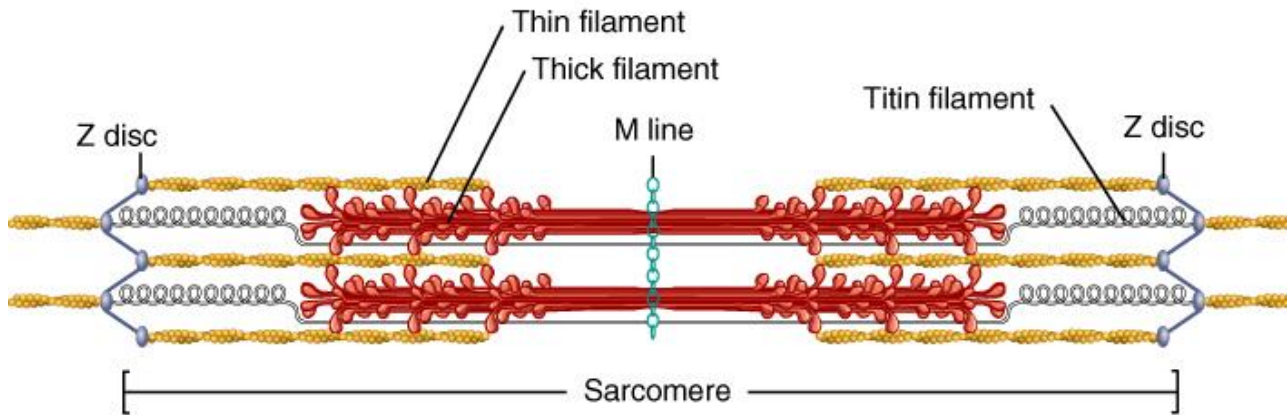


Fig. 10.04



(a) Myofibril



(b) Filaments

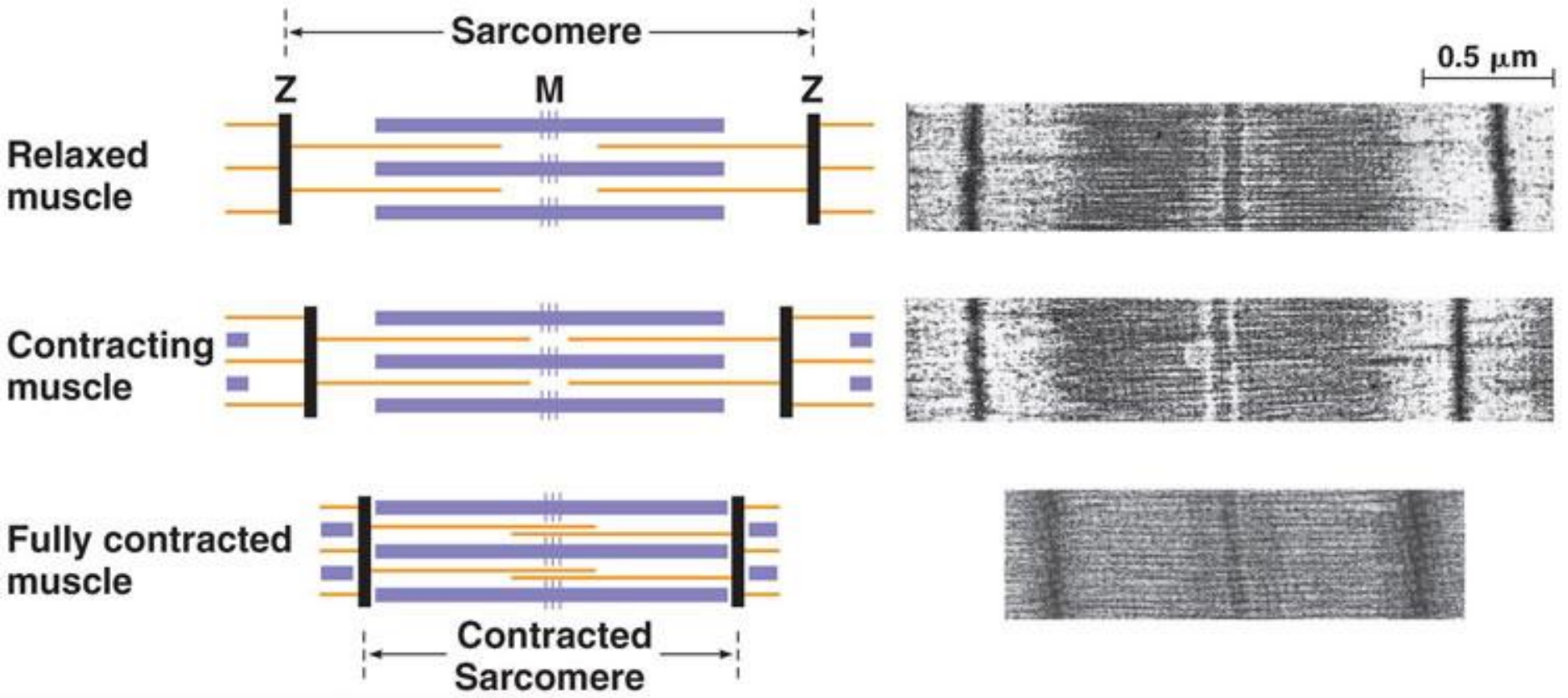
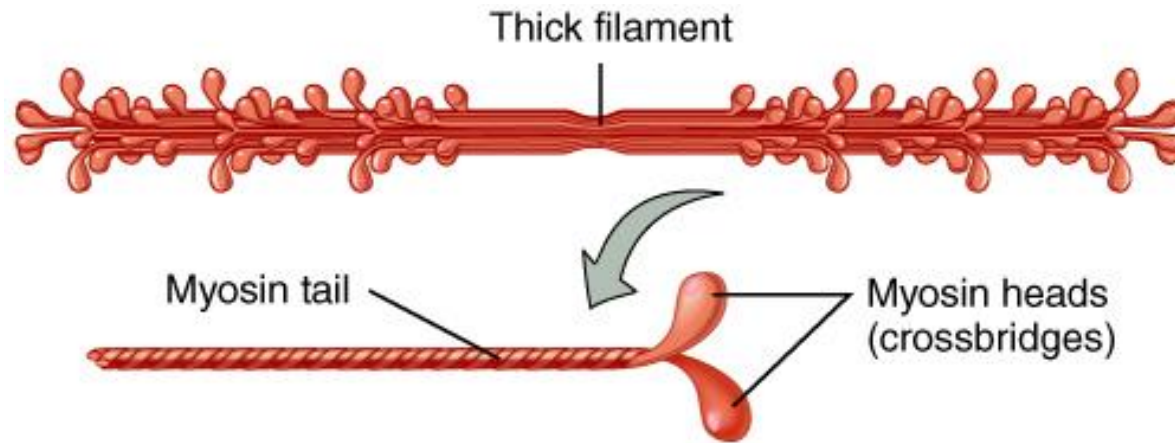
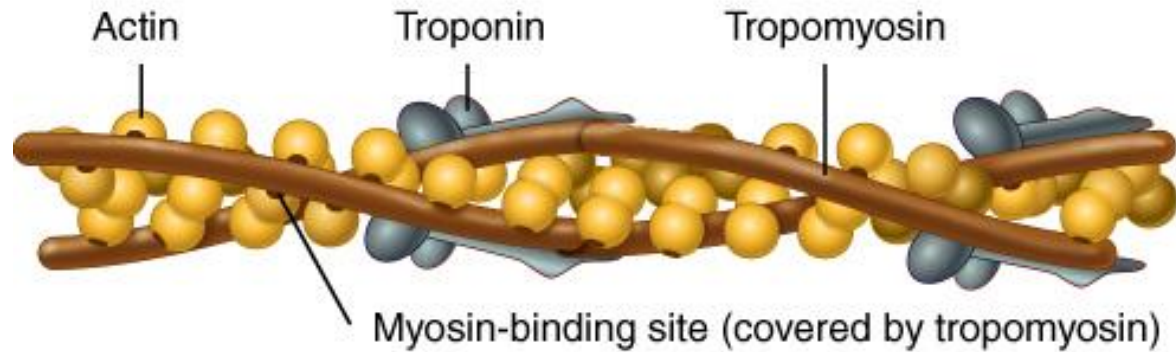


Fig. 10.06

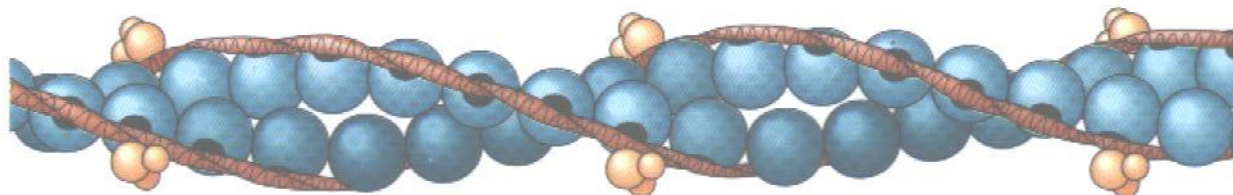
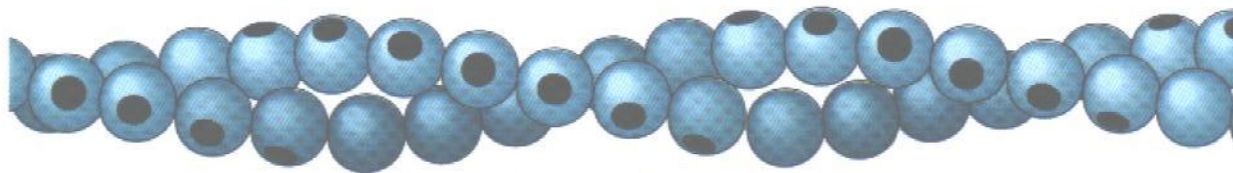


(a) One thick filament (above) and a myosin molecule (below)



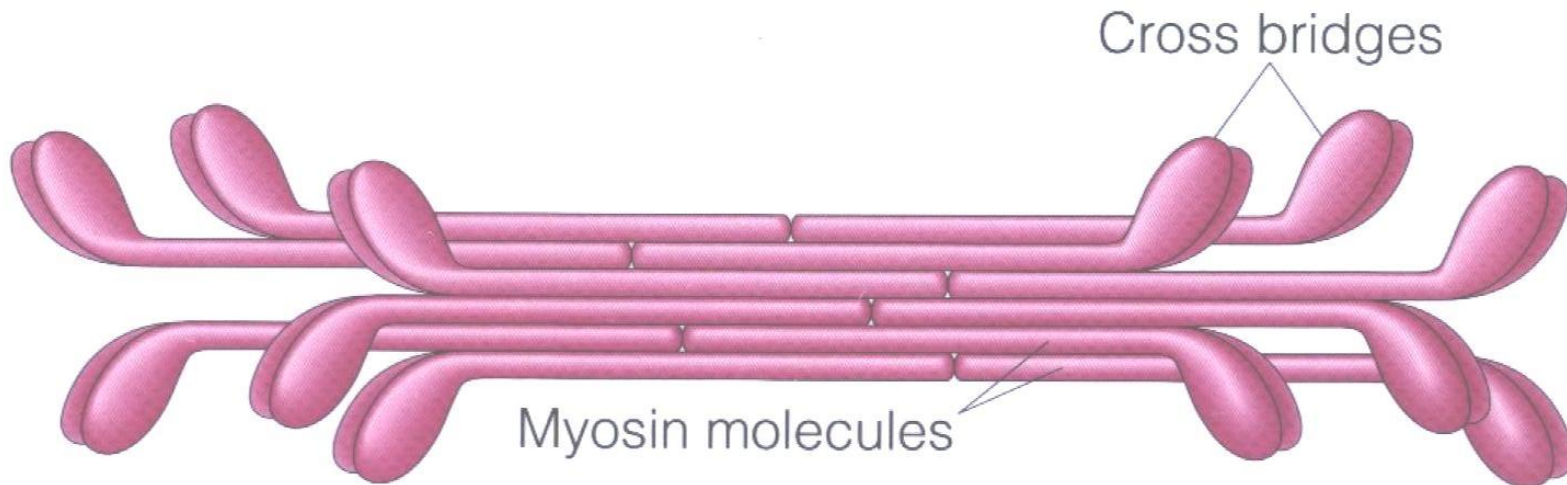
(b) Portion of a thin filament

Composition of Thin Filaments

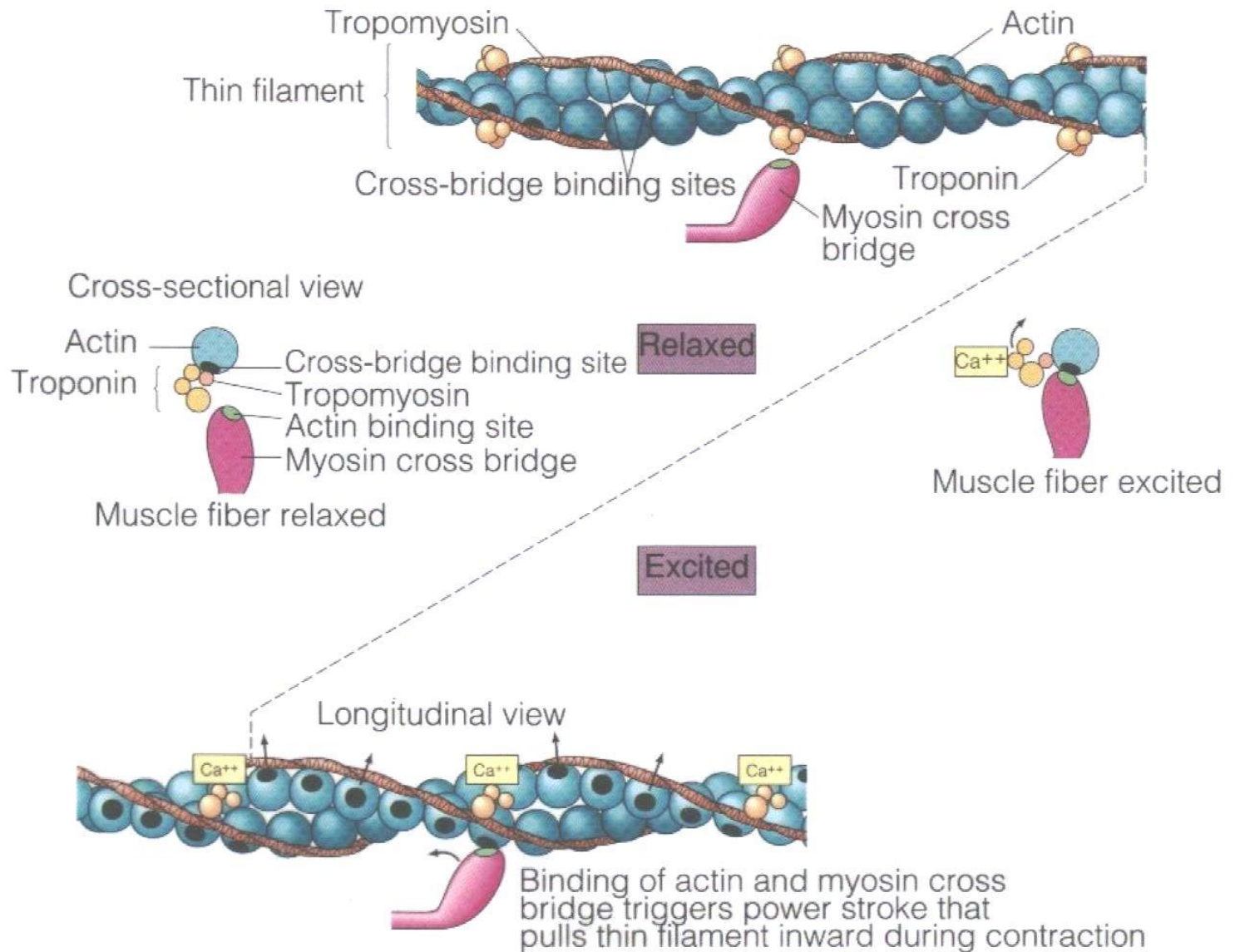


Thin filament

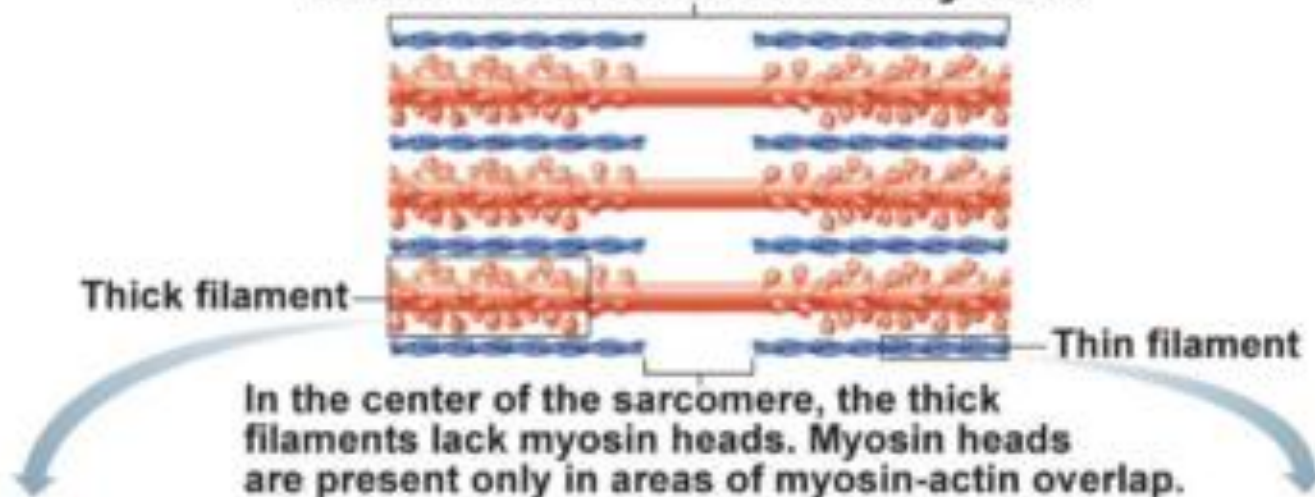
Structure of Myosin Molecules and Their Organization within a Thick Filament



Schematic Representation of Role of Calcium in Turning on Cross Bridges



Longitudinal section of filaments within one sarcomere of a myofibril



Thick filament

Each thick filament consists of many myosin molecules whose heads protrude at opposite ends of the filament.

Portion of a thick filament



Actin-binding sites

ATP-binding site

Heads

Tail

Flexible hinge region

Myosin molecule

Thin filament

A thin filament consists of two strands of actin subunits twisted into a helix plus two types of regulatory proteins (troponin and tropomyosin).

Portion of a thin filament



Actin subunits

Actin subunits

Active sites for myosin attachment

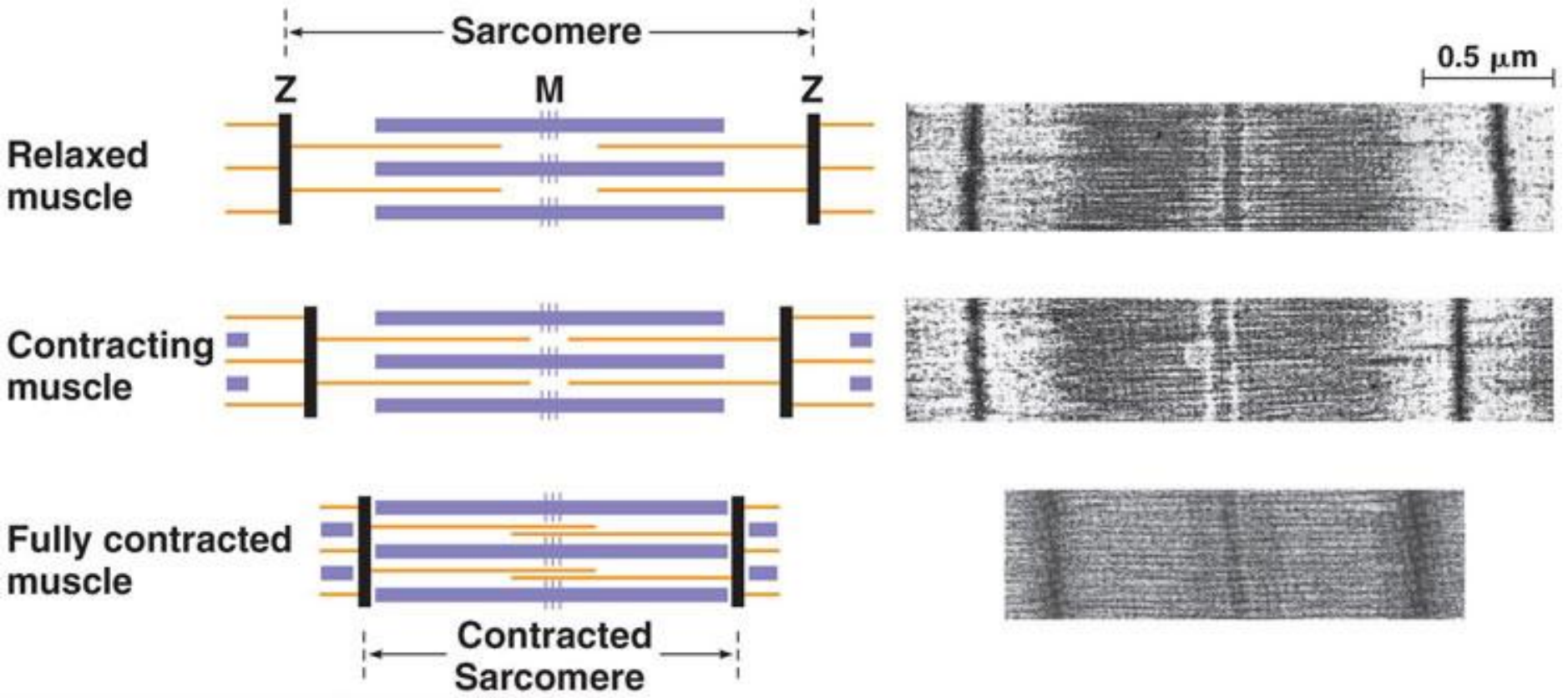


Fig. 10.07

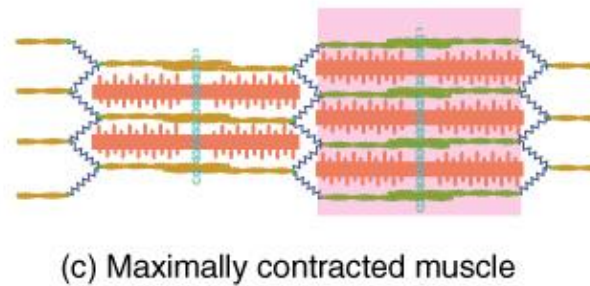
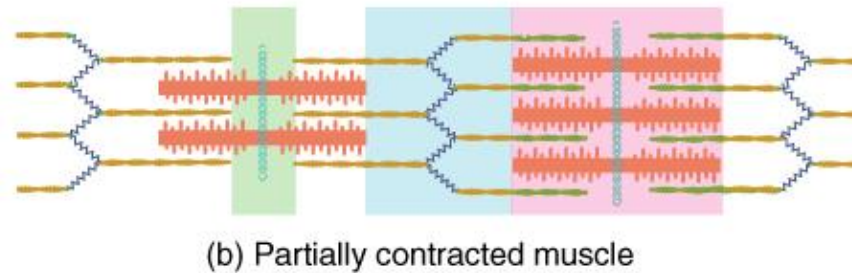
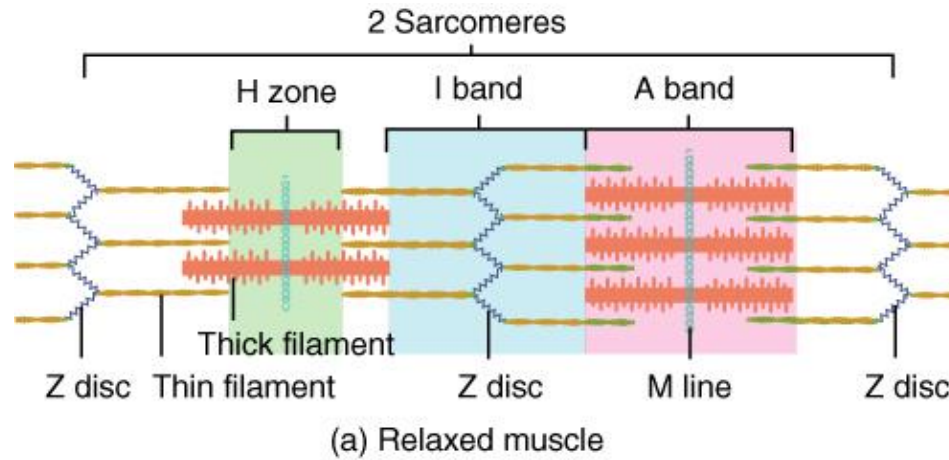
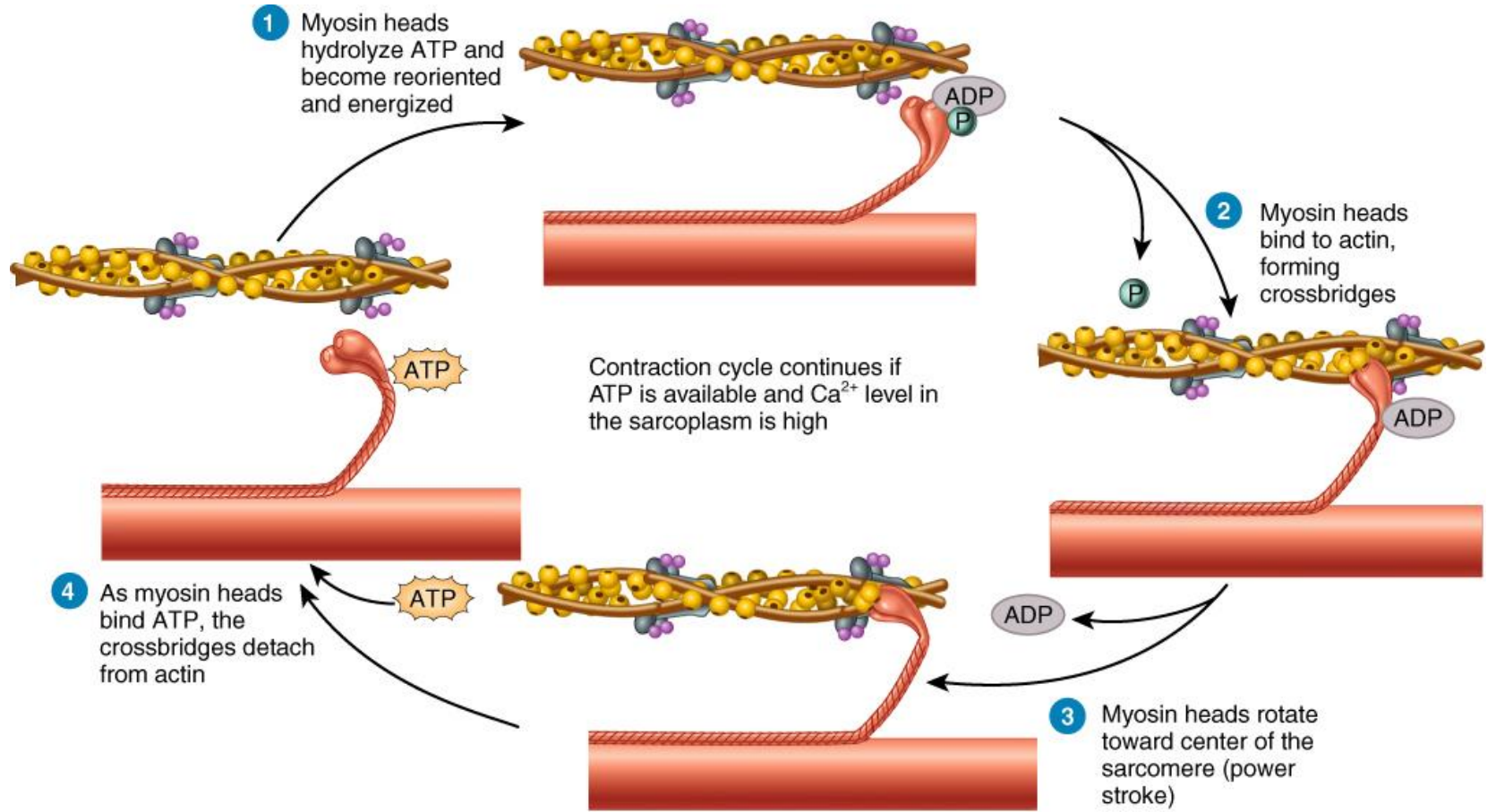
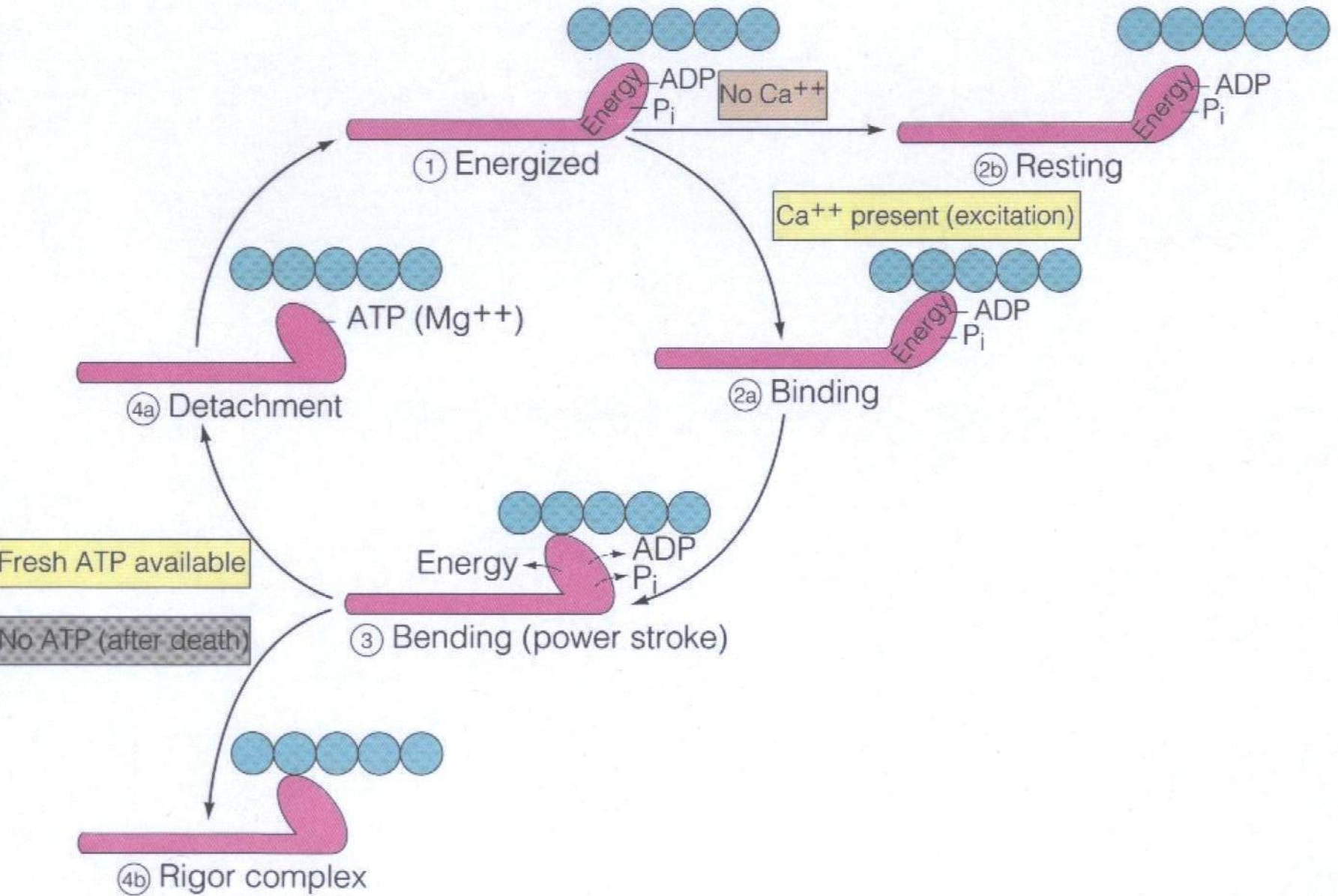


Fig. 10.08



Cross-Bridge Cycle



Sources of energy for muscle contraction

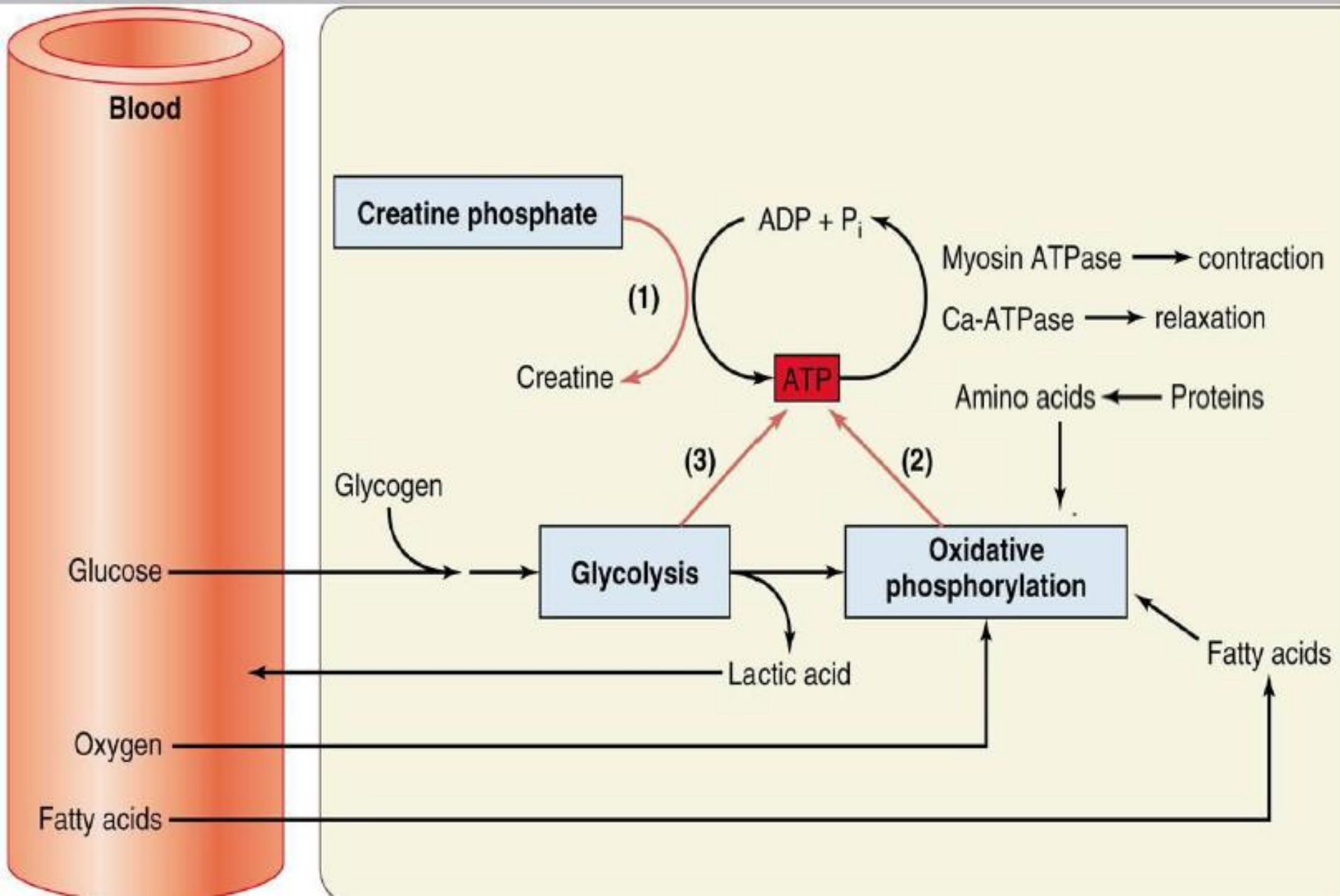
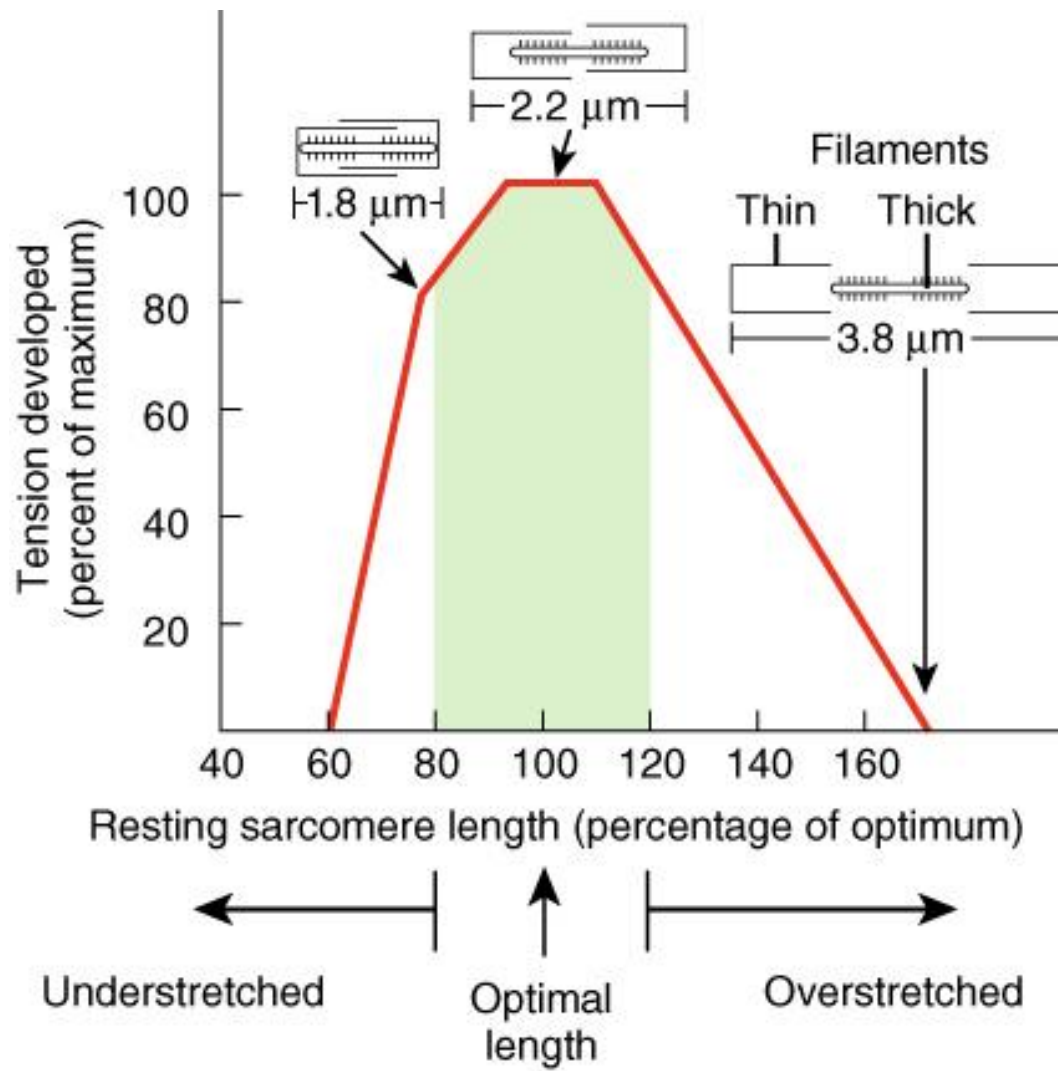
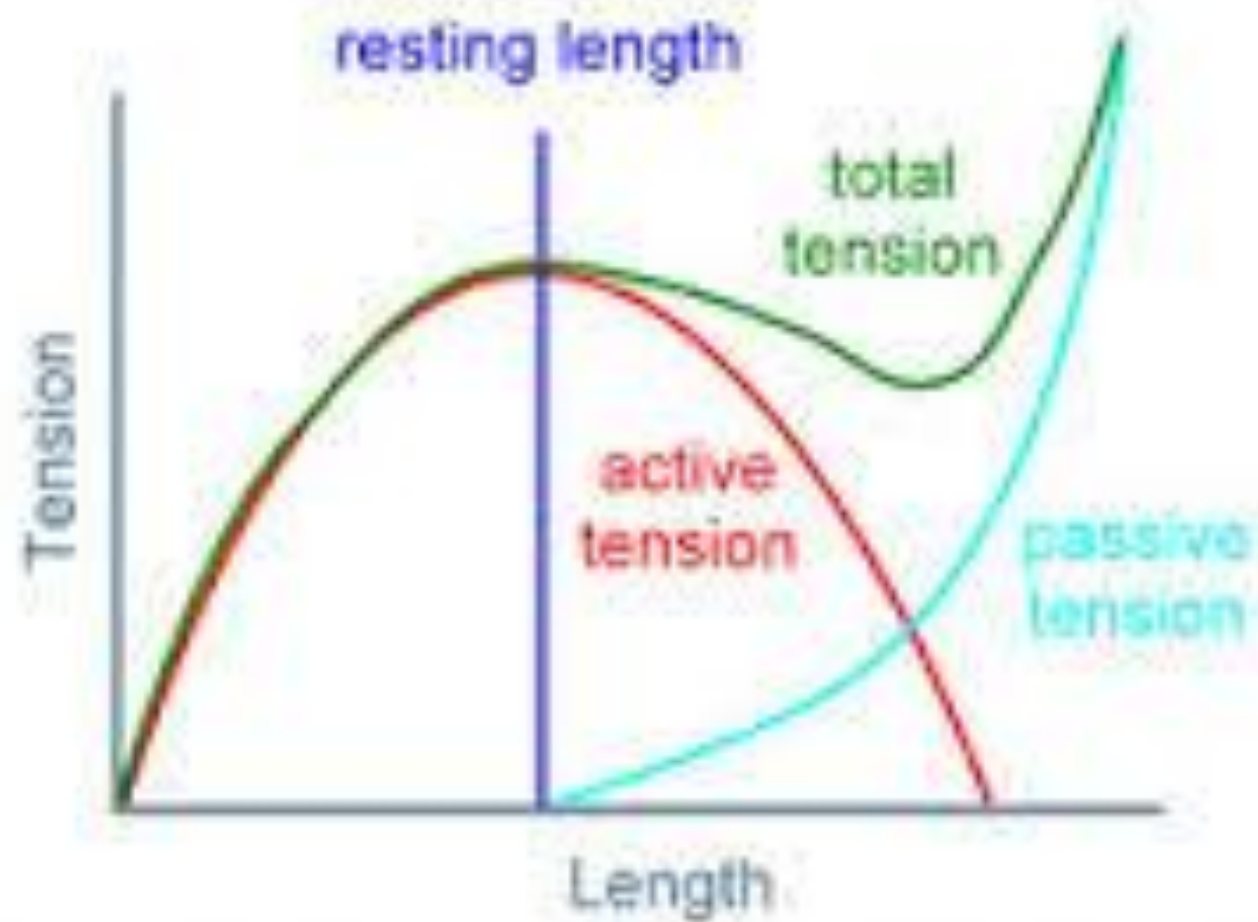


Fig. 10.10





Length-Tension Curve of a Muscle

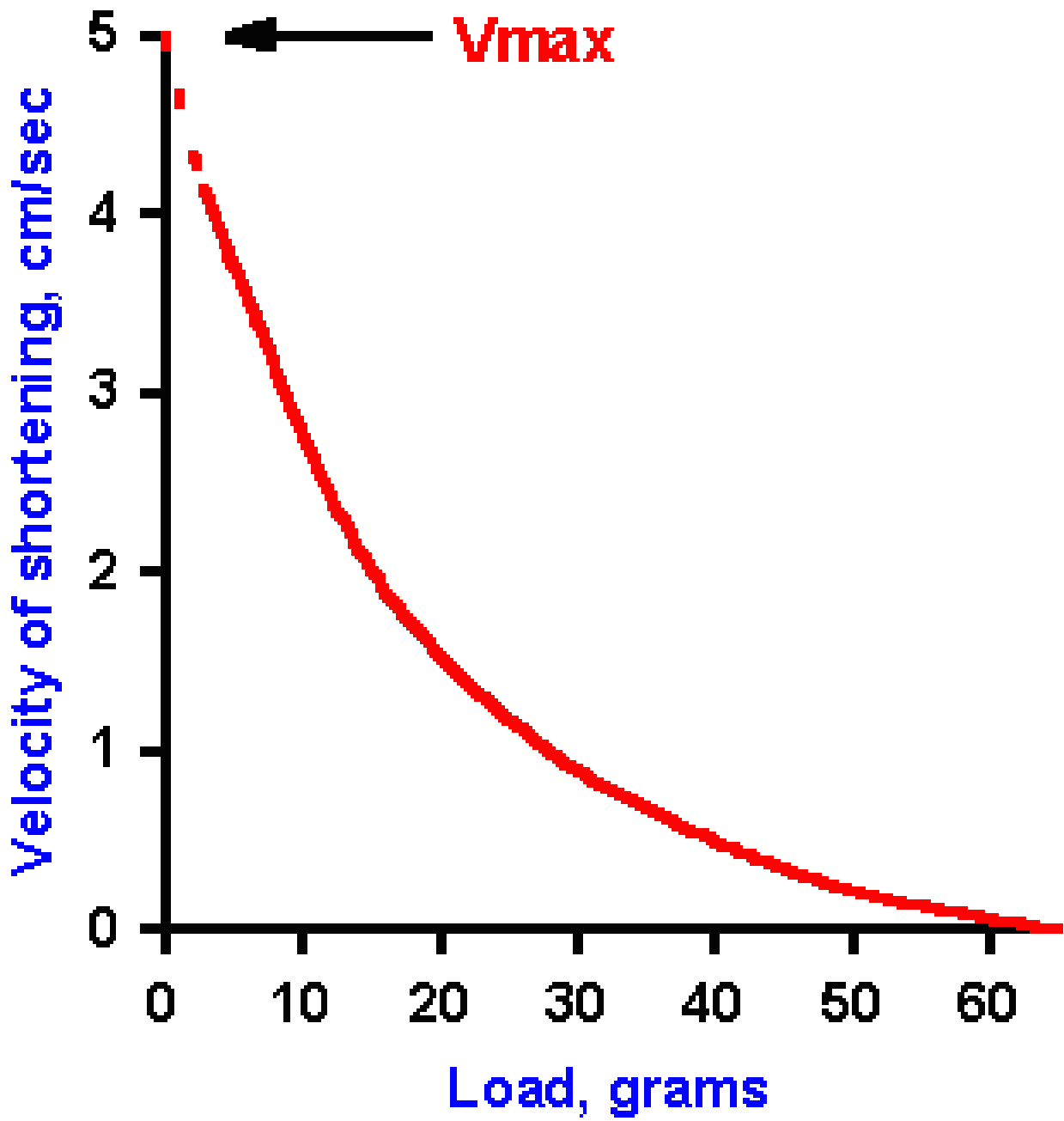
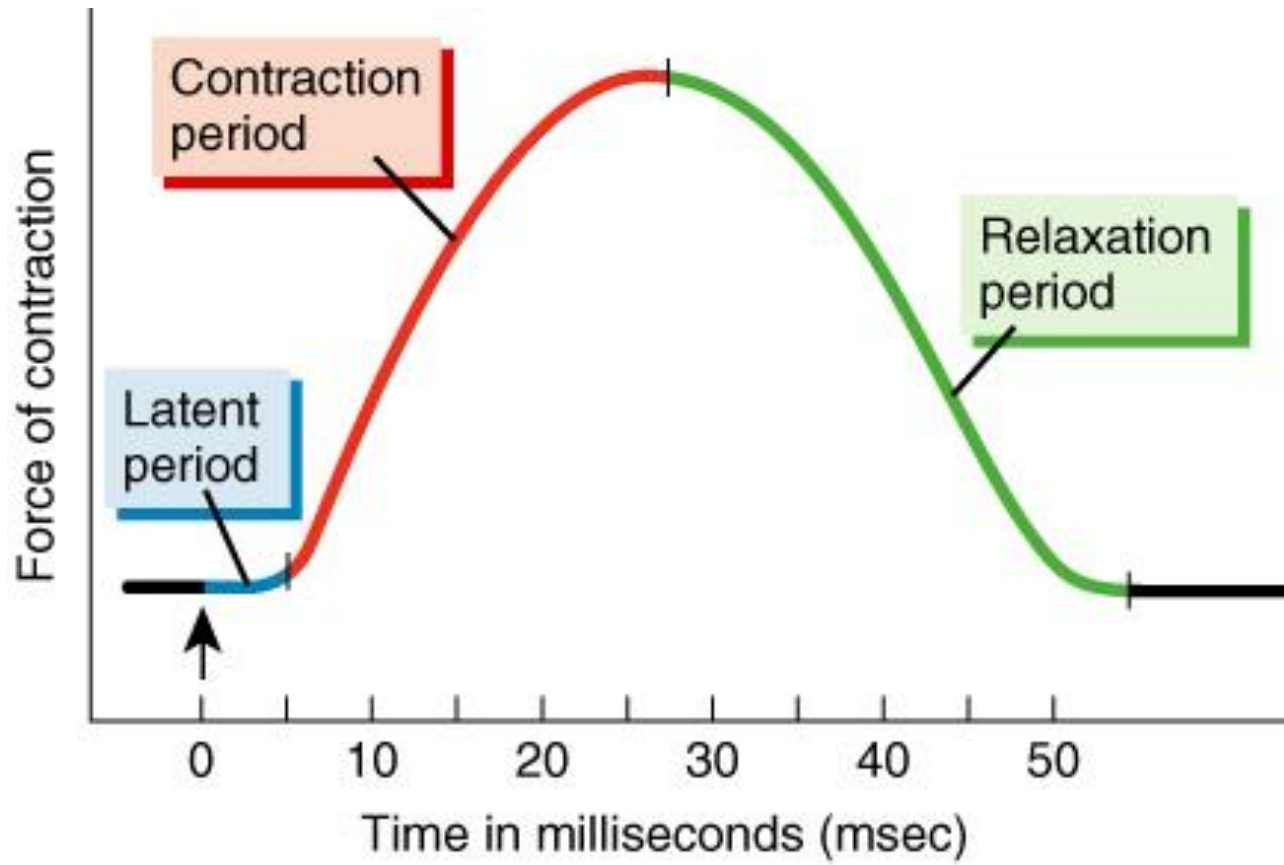
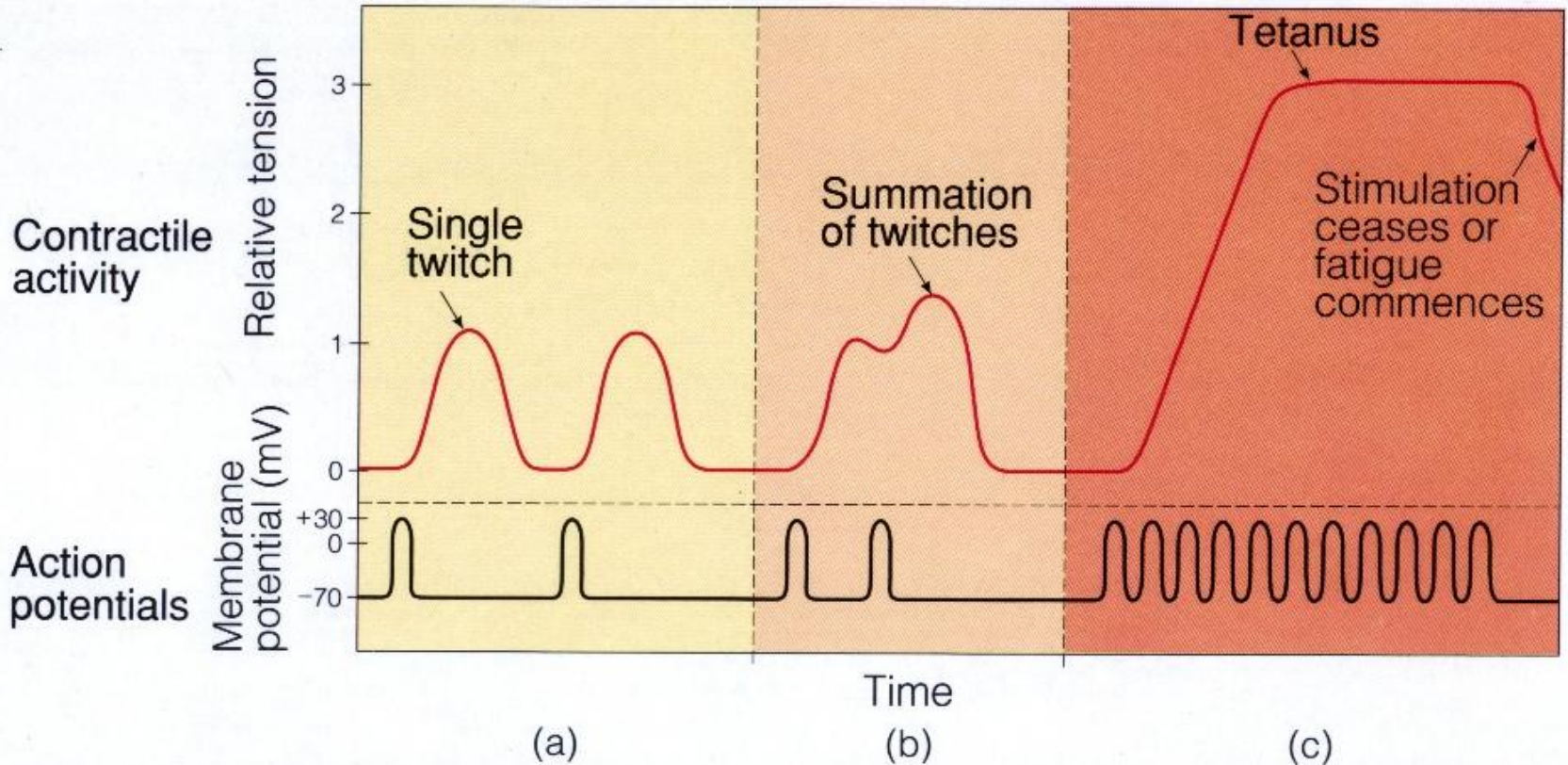


Fig. 10.15



10.15

Summation and Tetanus



The duration of the action potentials is not drawn to scale but is exaggerated.

● FIGURE 8-15

Schematic representation of motor units in a skeletal muscle

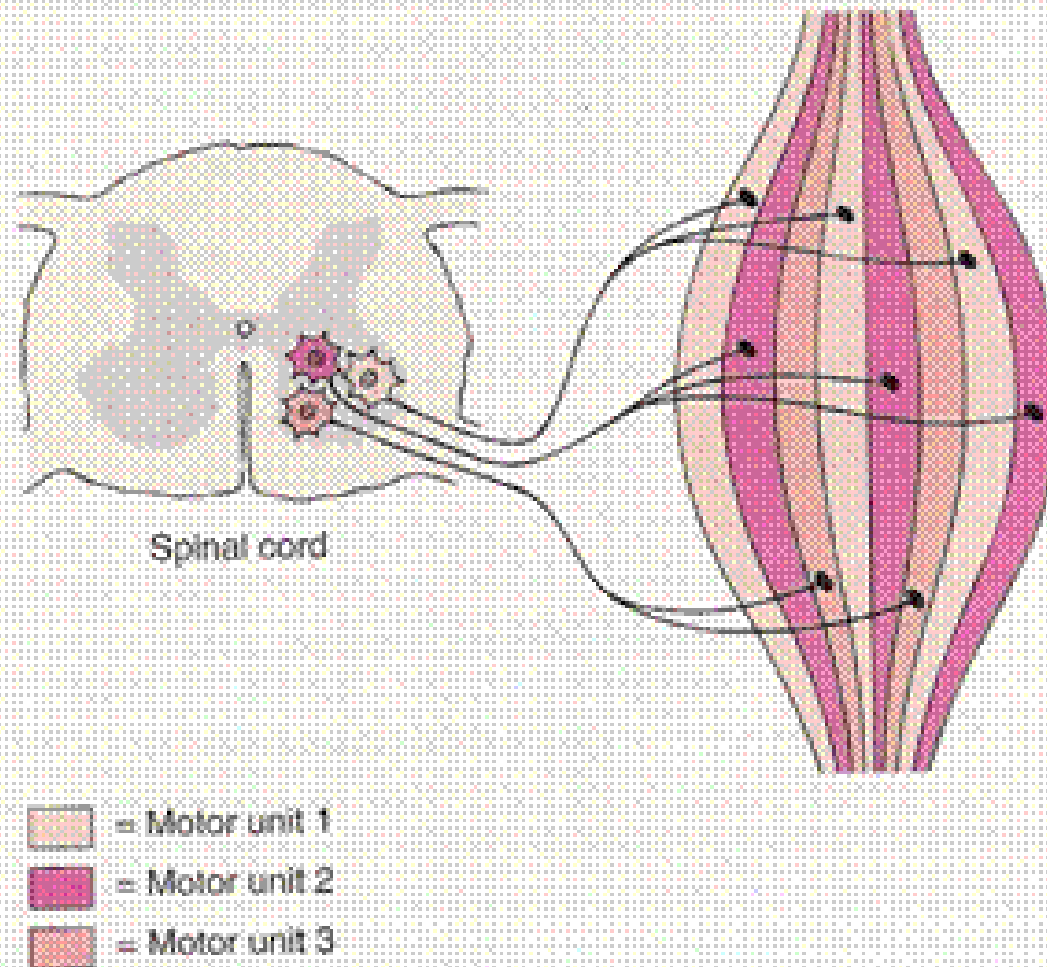
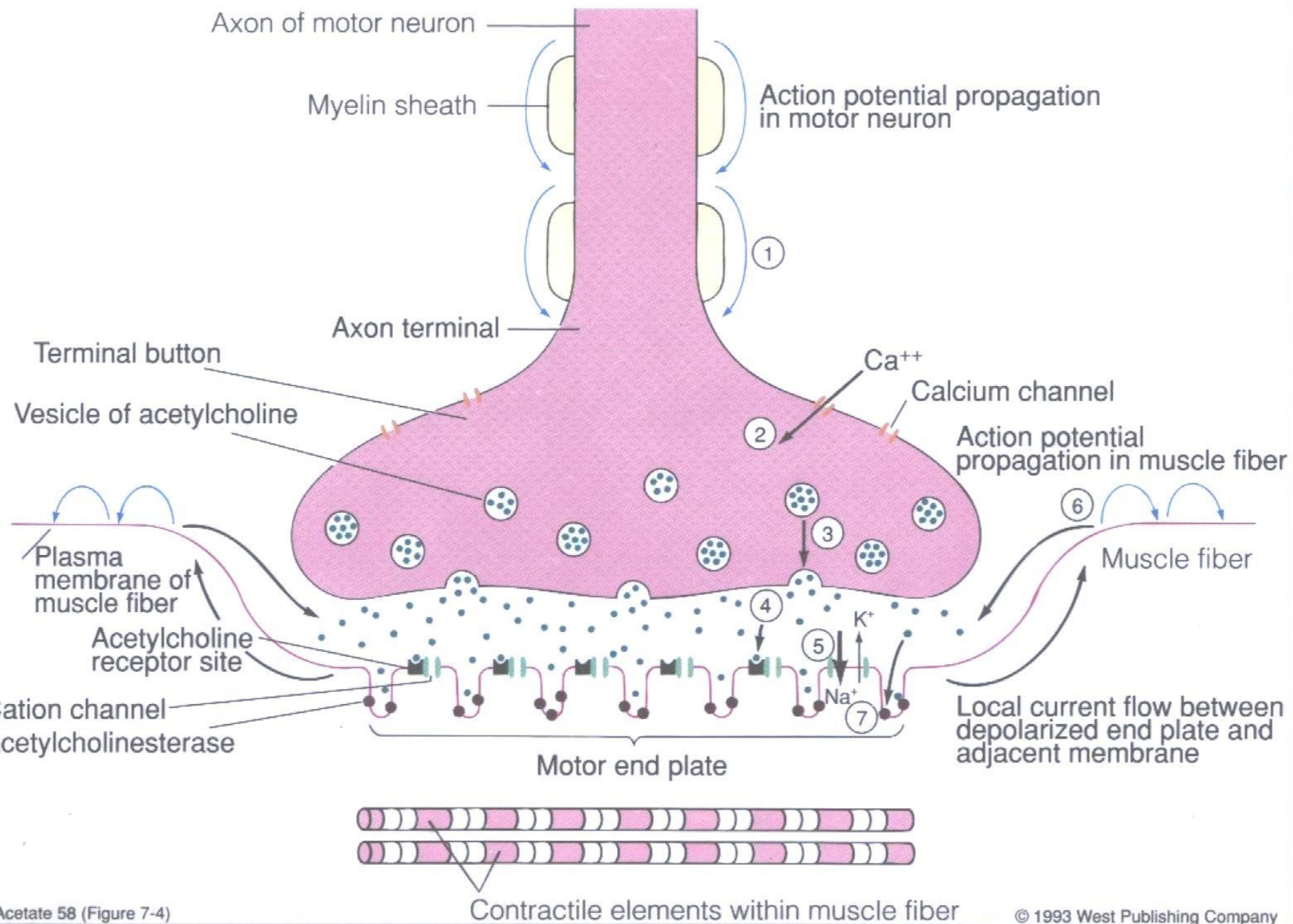
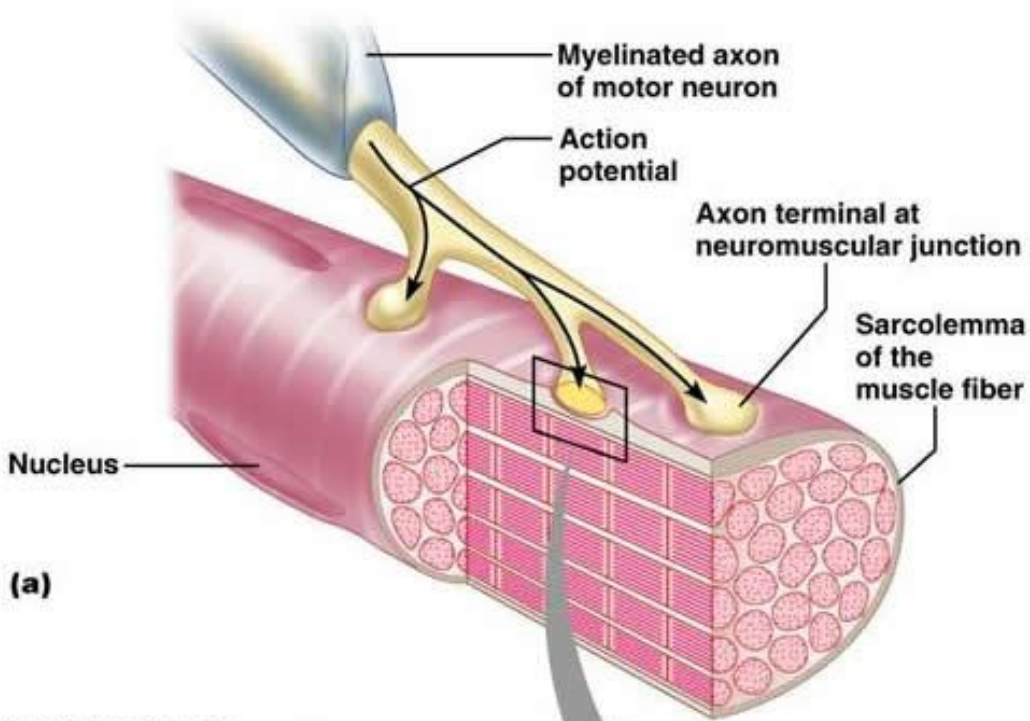


Figure 4. One motor unit can control multiple muscle cells.

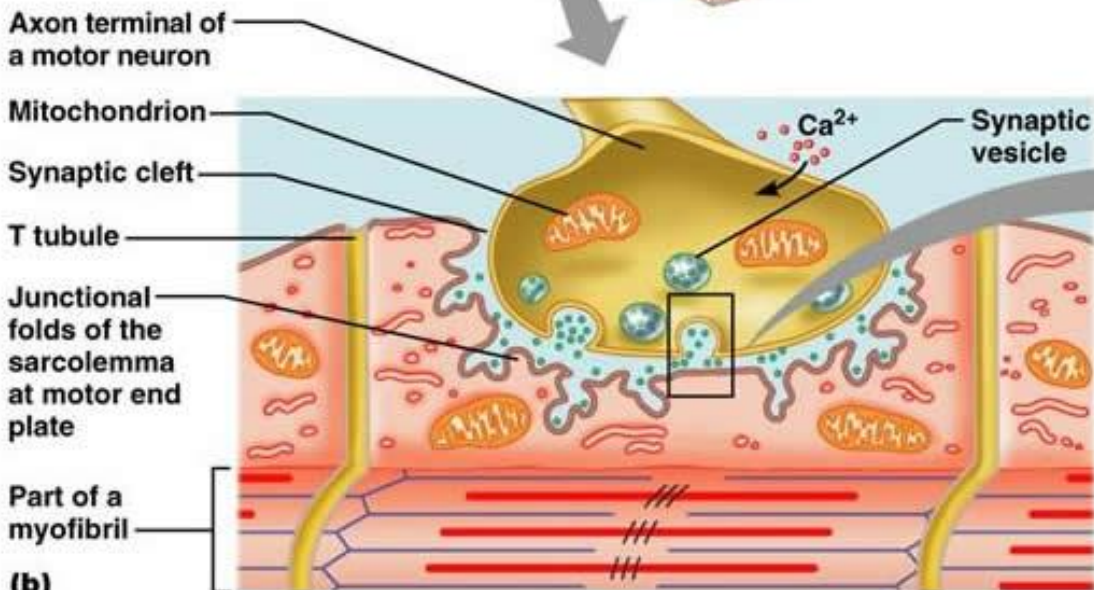
Taken from Sherwood, 2004

Events at a Neuromuscular Junction

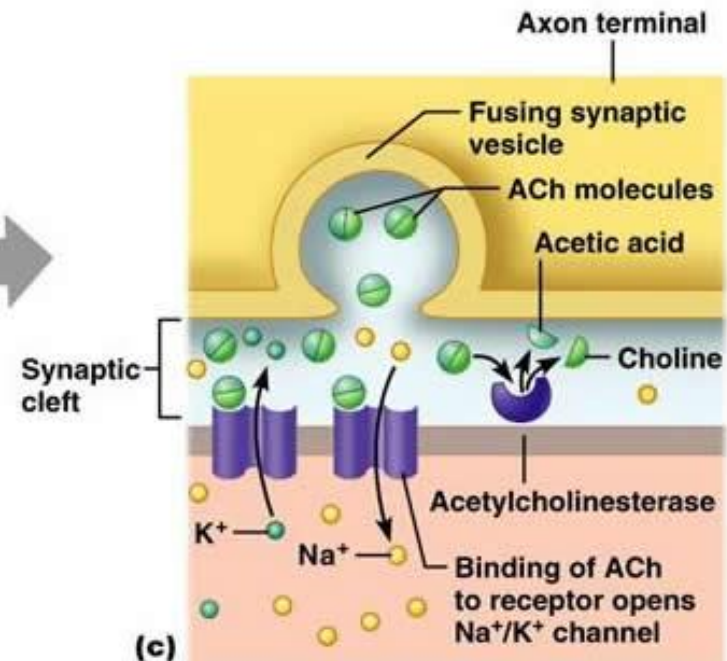




(a)



(b)



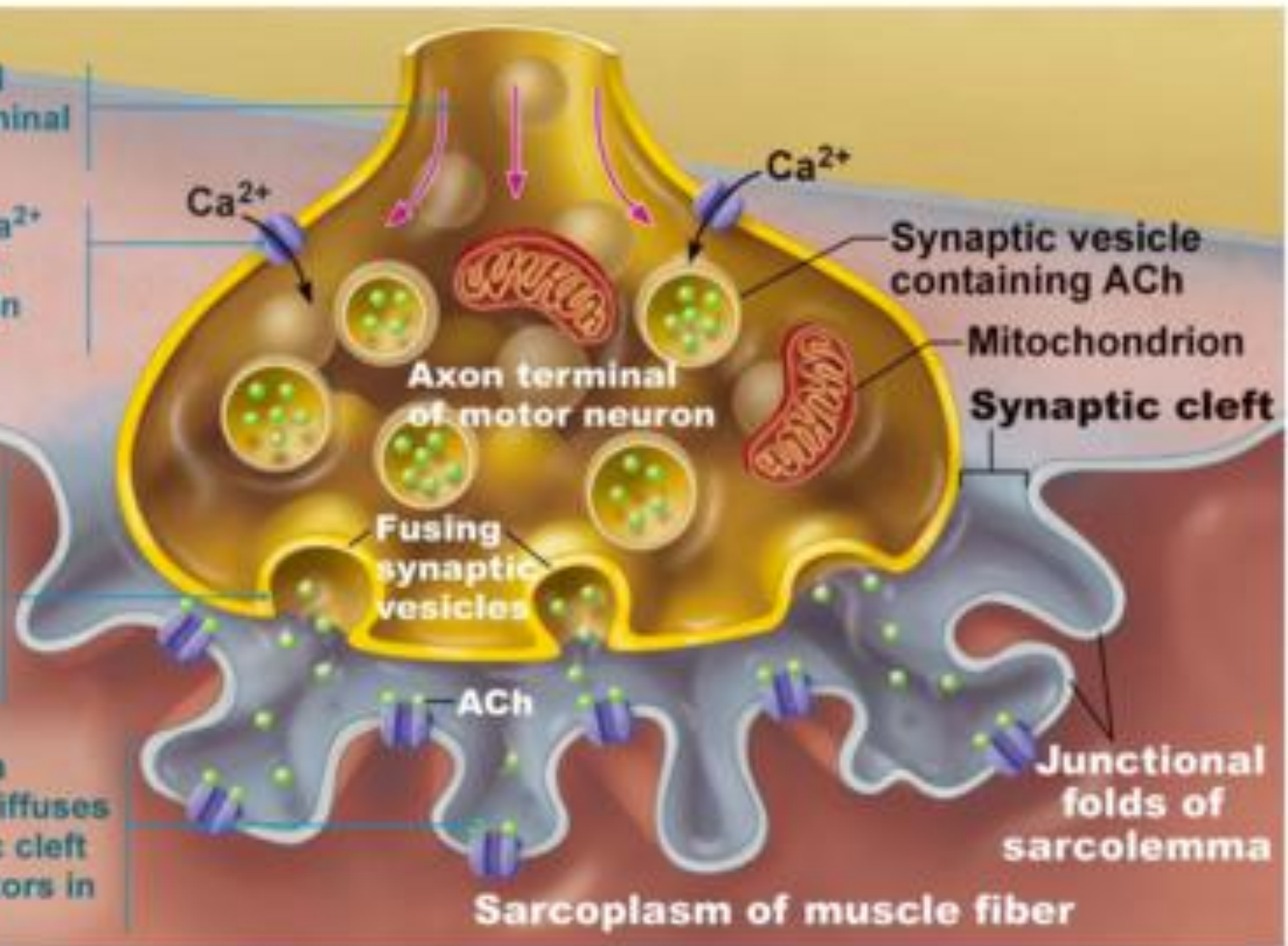
(c)

1 Action potential arrives at axon terminal of motor neuron.

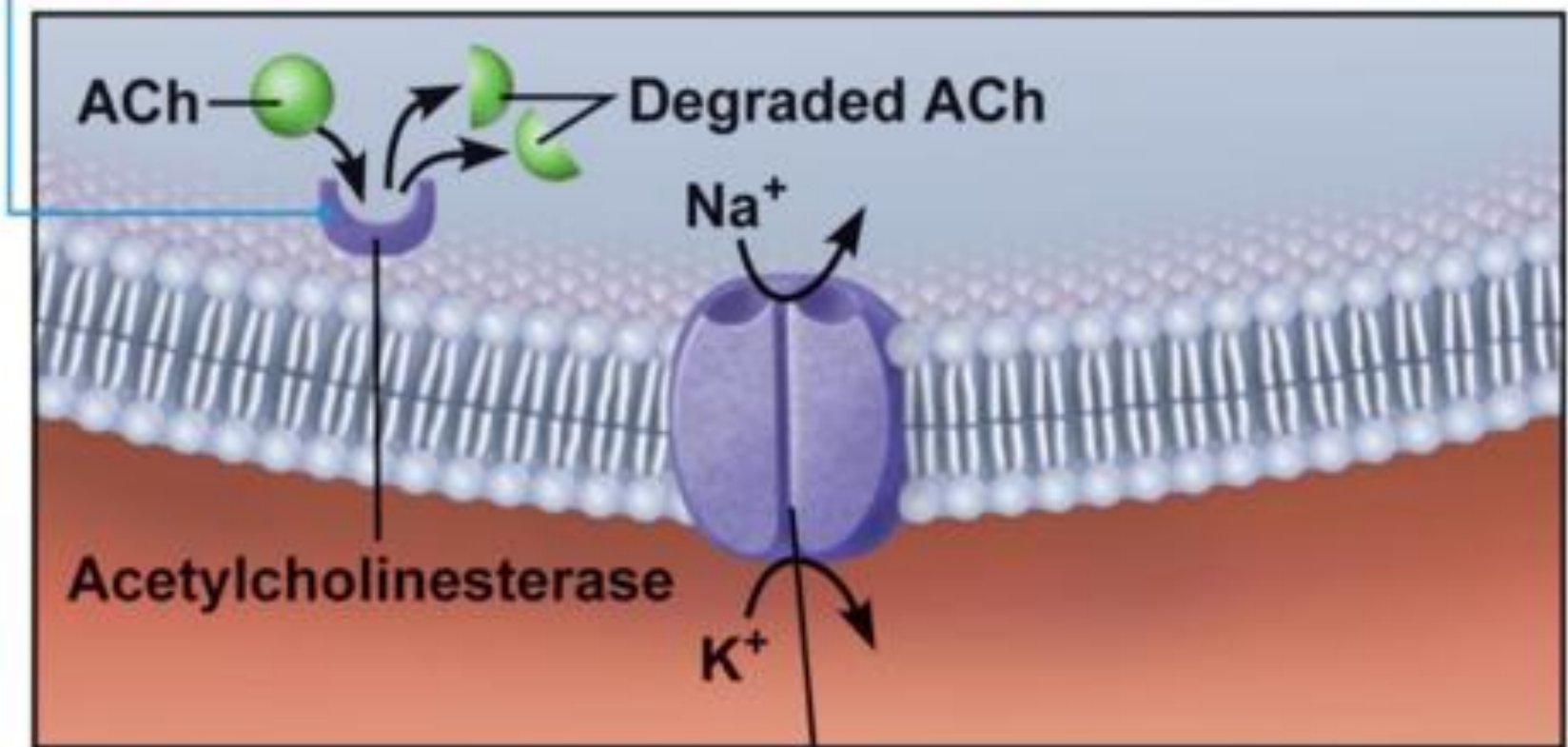
2 Voltage-gated Ca^{2+} channels open and Ca^{2+} enters the axon terminal.

3 Ca^{2+} entry causes some synaptic vesicles to release their contents (acetylcholine) by exocytosis.

4 Acetylcholine, a neurotransmitter, diffuses across the synaptic cleft and binds to receptors in the sarcolemma.

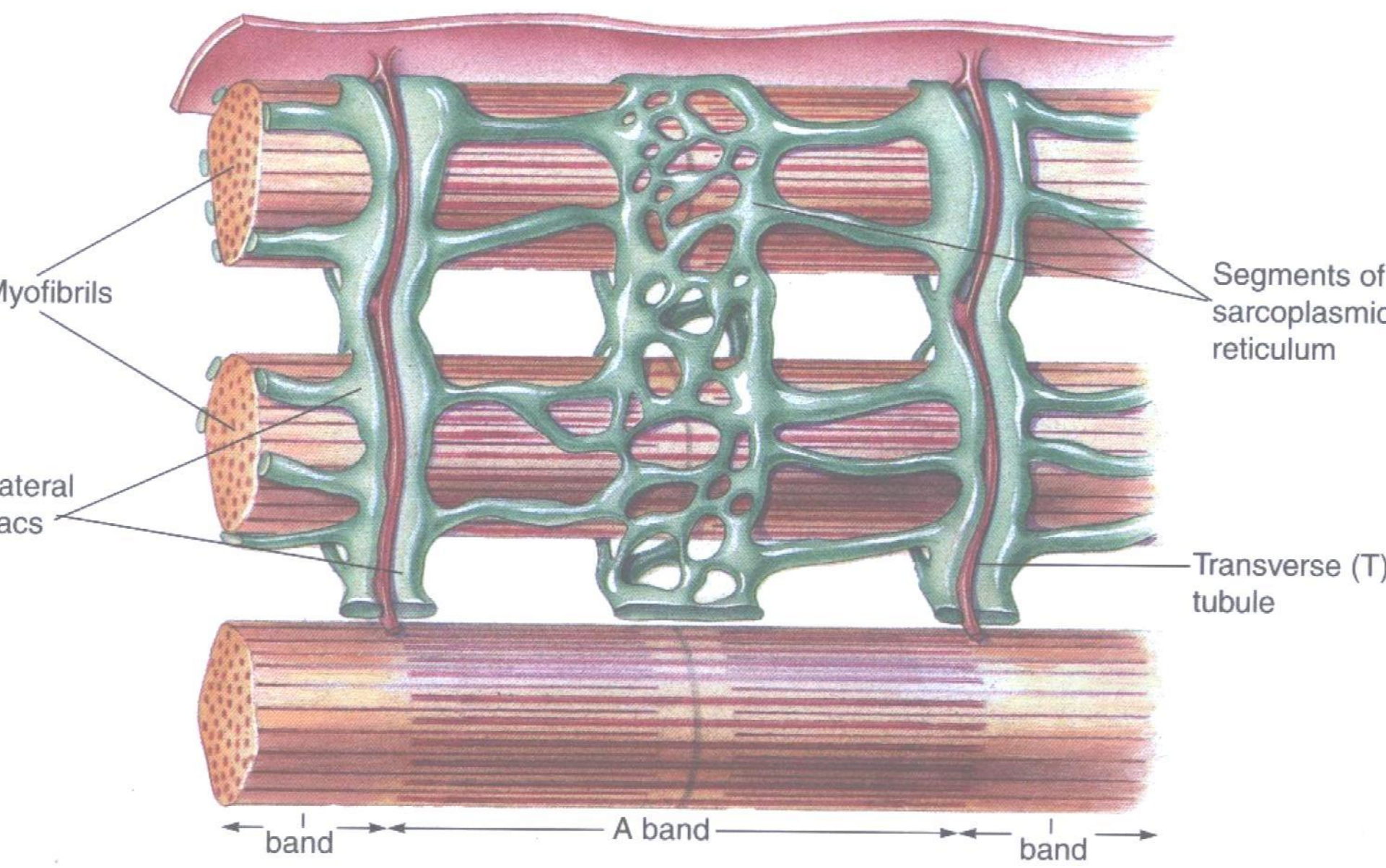


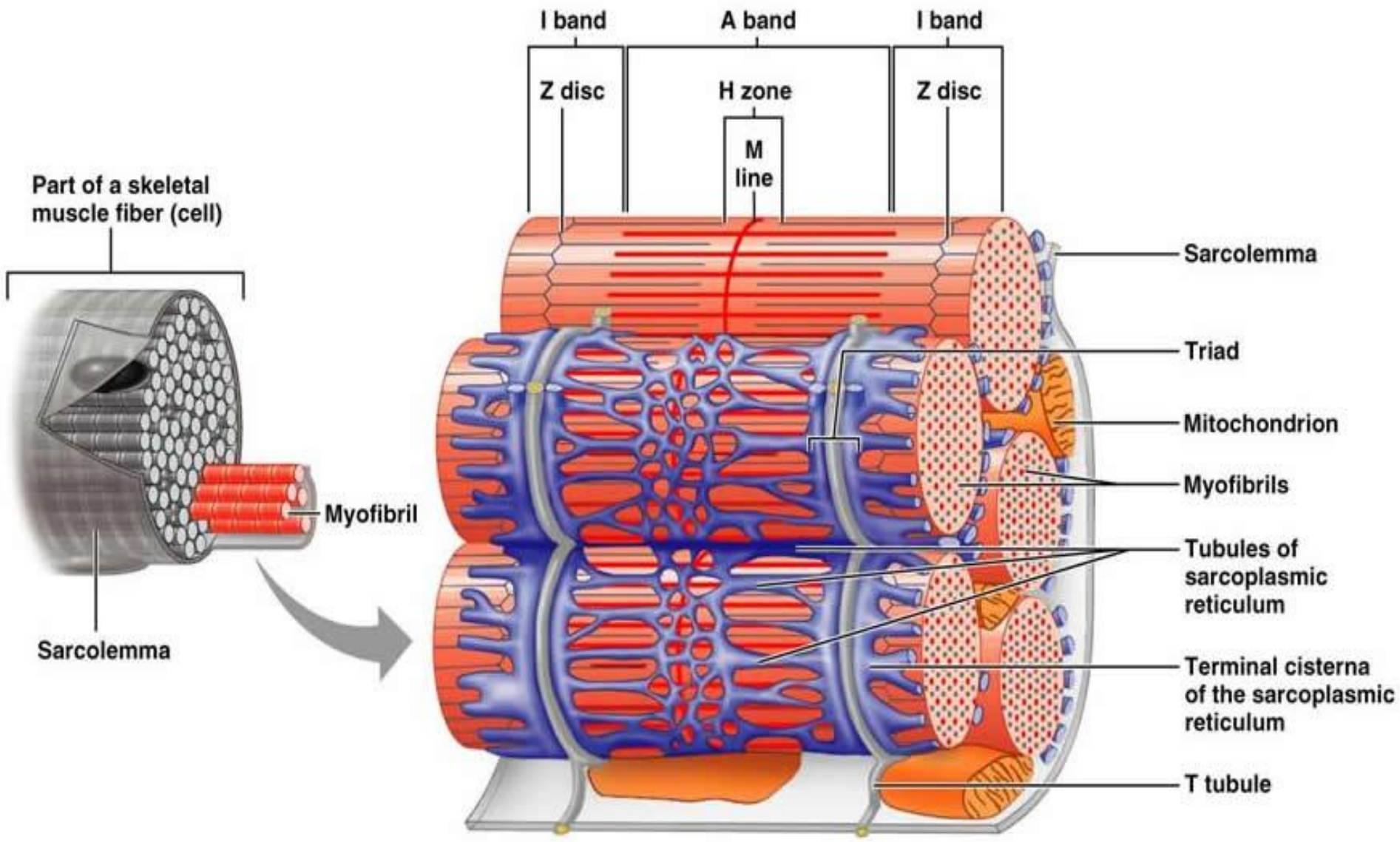
⑥ ACh effects are terminated by its enzymatic breakdown in the synaptic cleft by acetylcholinesterase.

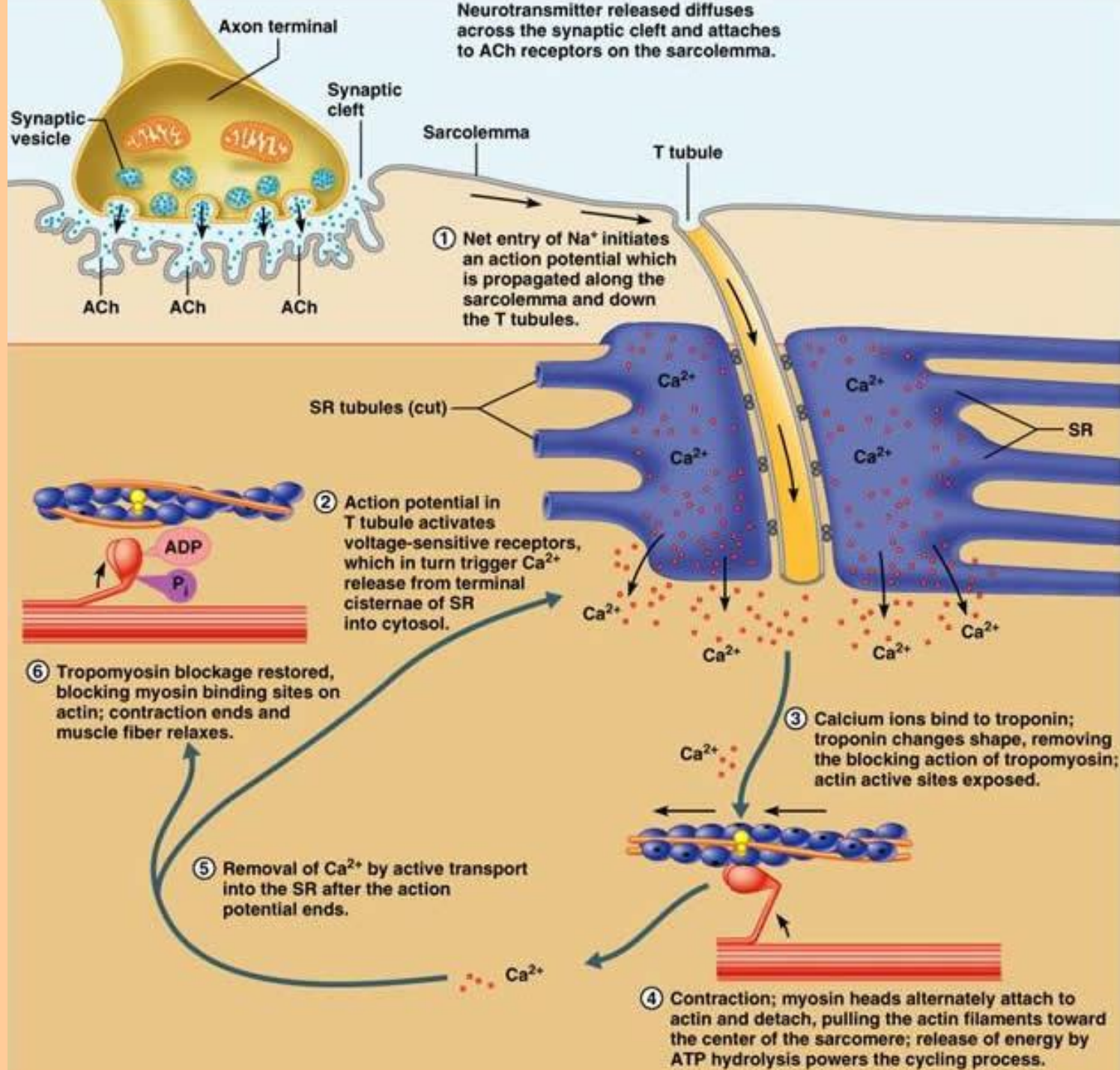


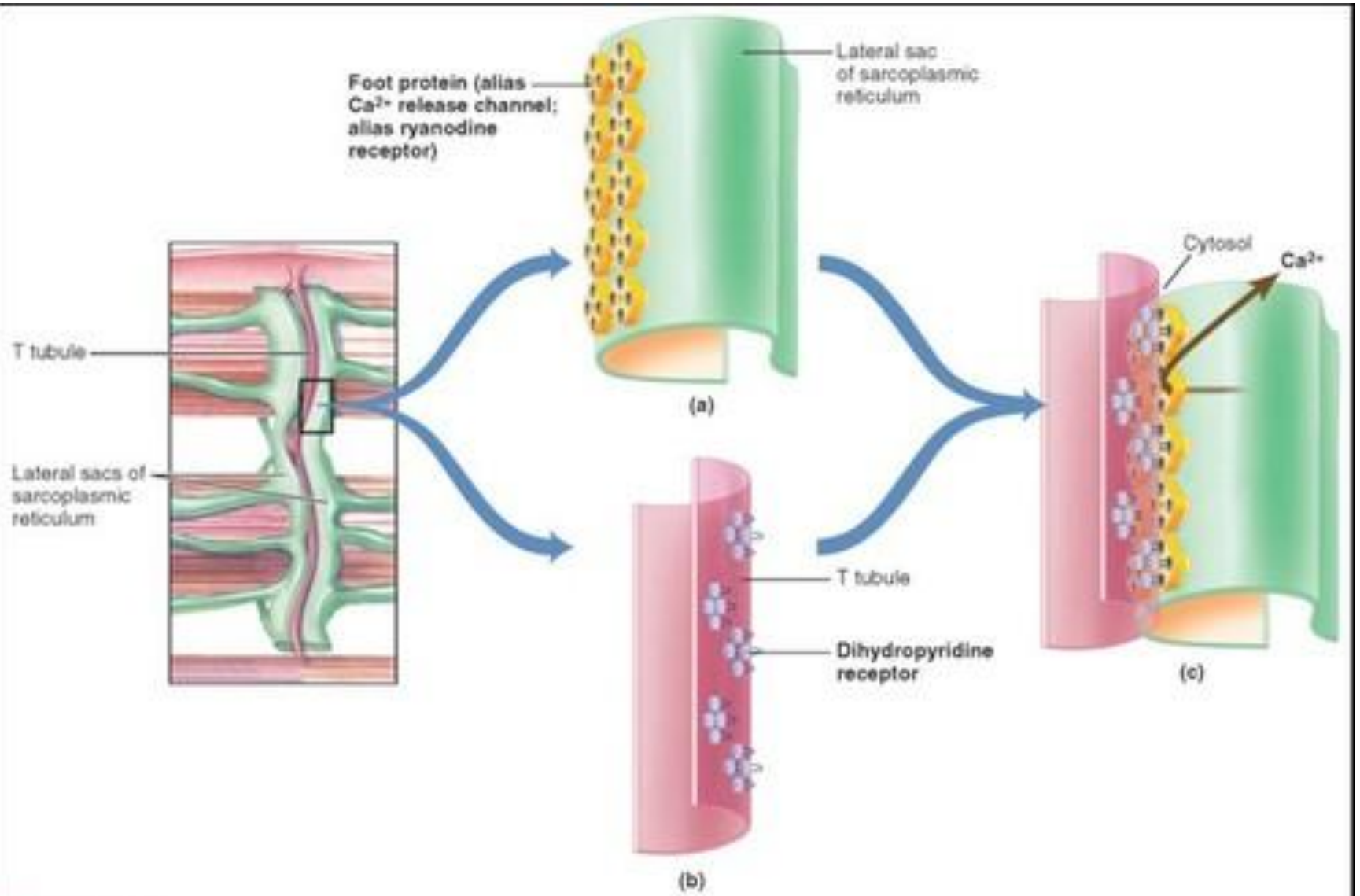
Postsynaptic membrane ion channel closed; ions cannot pass.

Surface membrane of muscle fiber







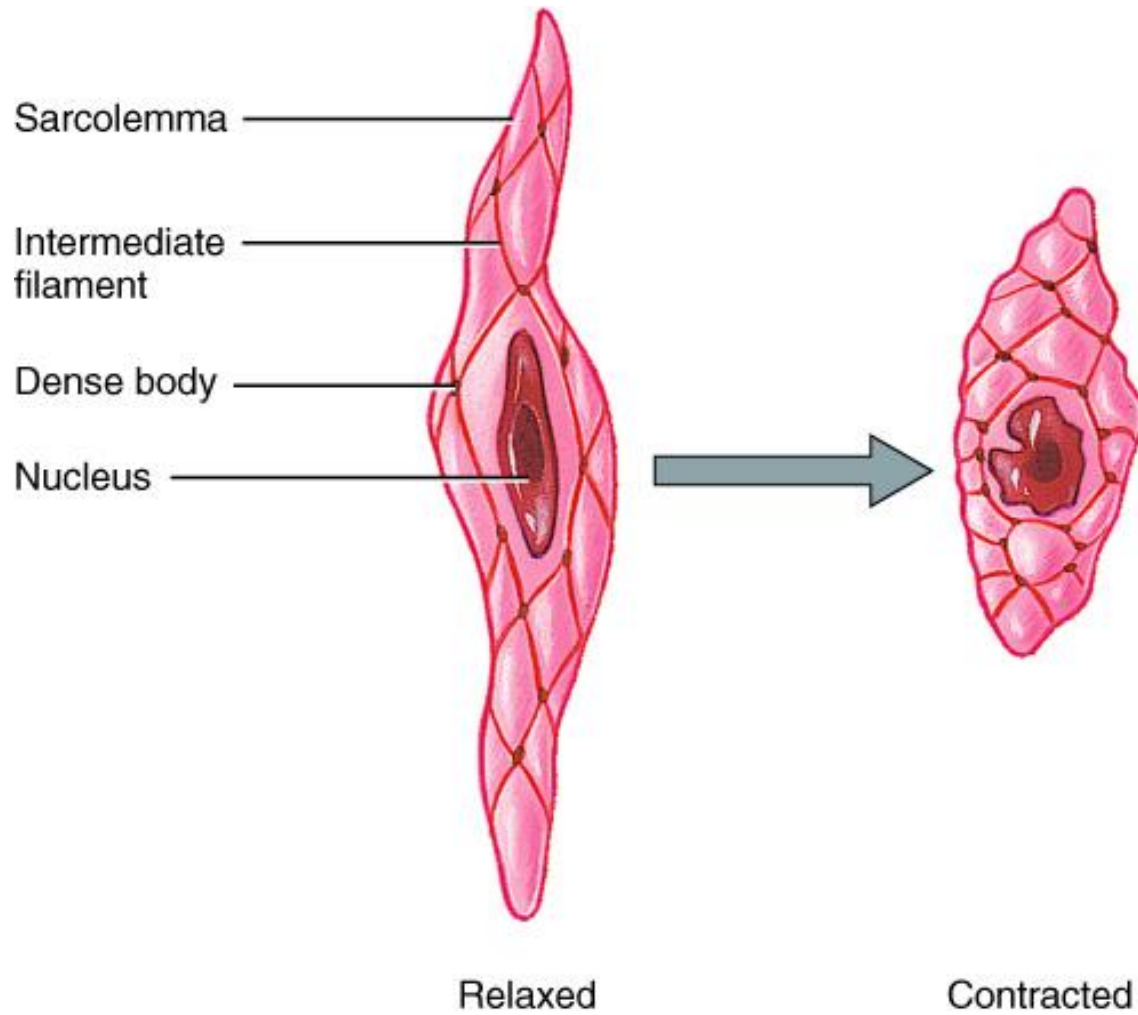


● FIGURE 8-11

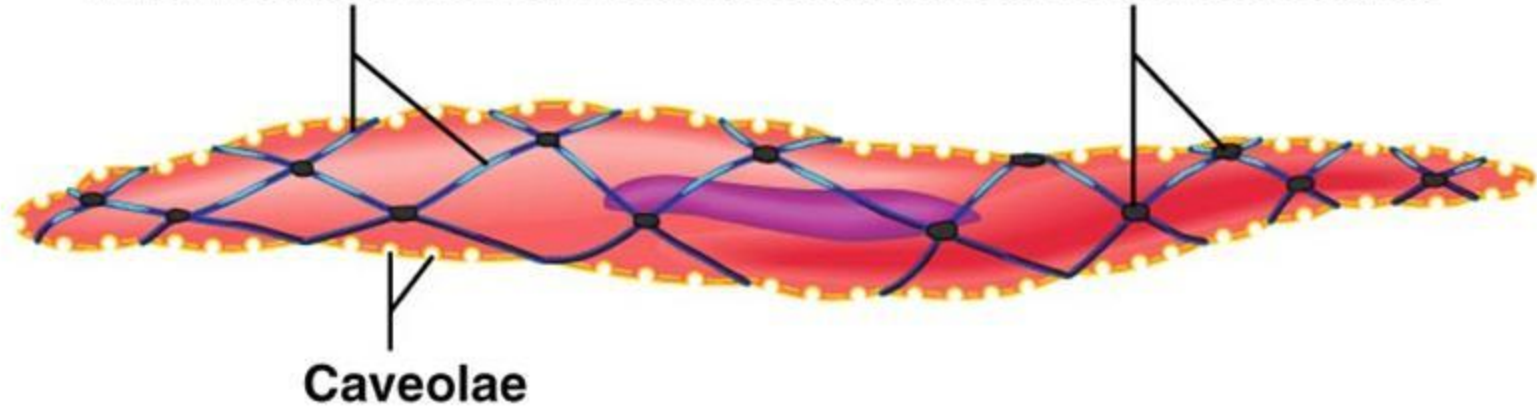
Relationship between a T tubule and the adjacent lateral sacs of the sarcoplasmic reticulum

SMOOTH MUSCLE CELLS

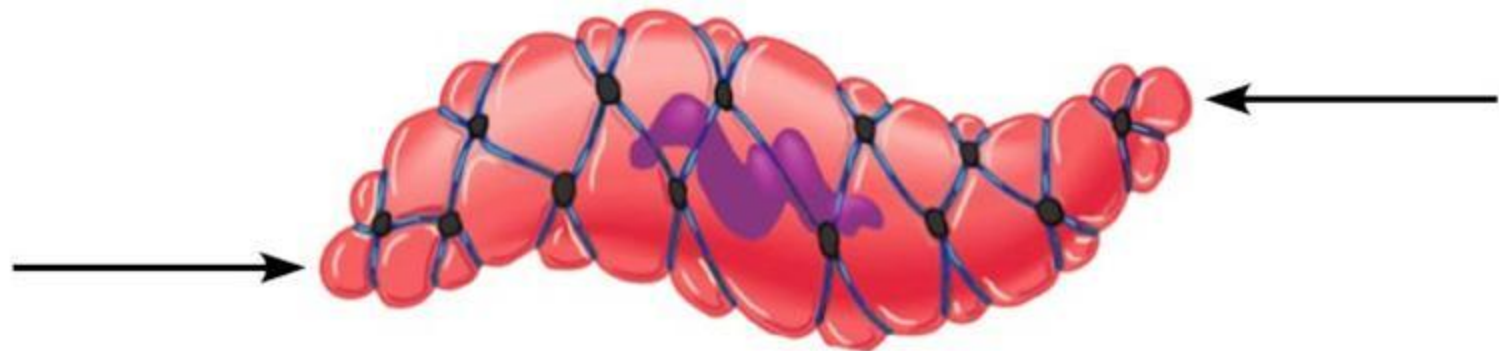
Fig. 10.19



Intermediate filament bundles attached to dense bodies

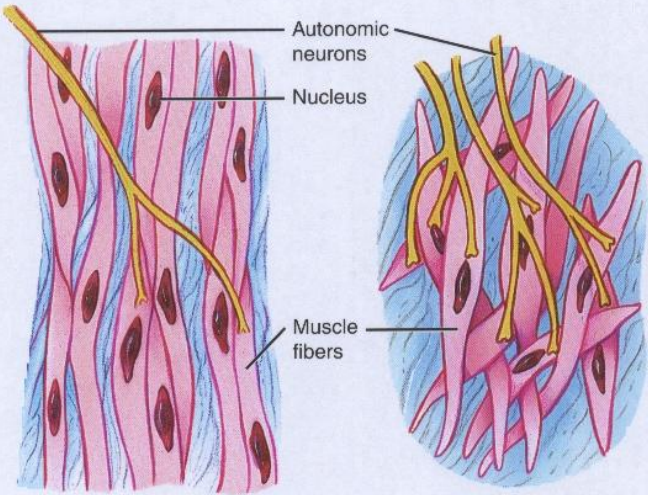


(a) Relaxed smooth muscle cell



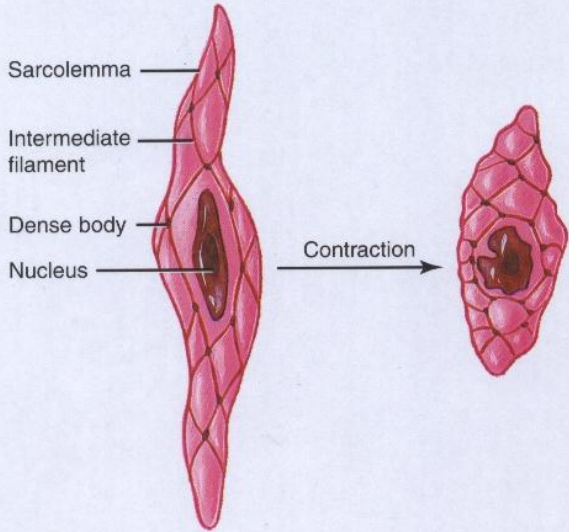
(b) Contracted smooth muscle cell

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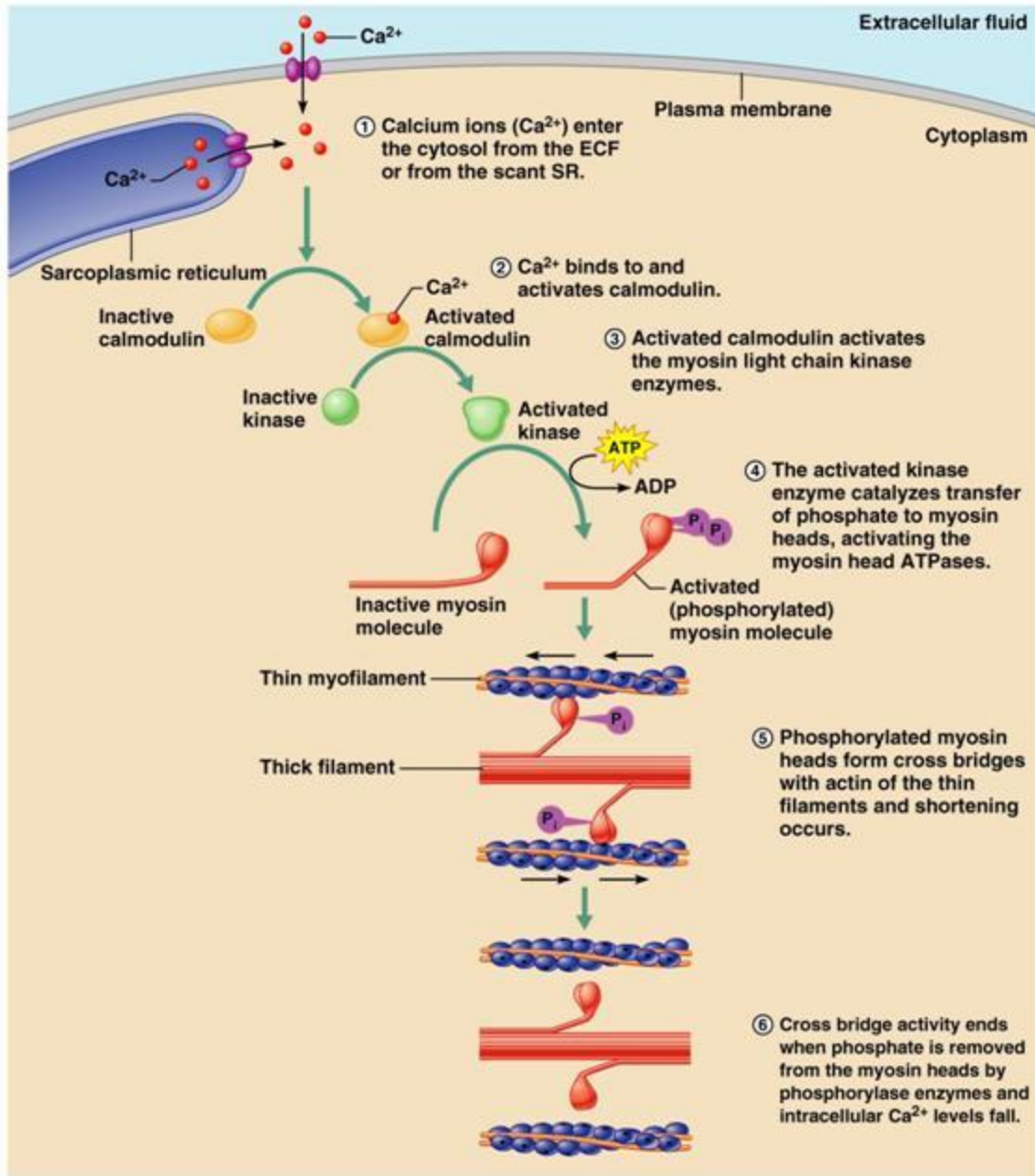


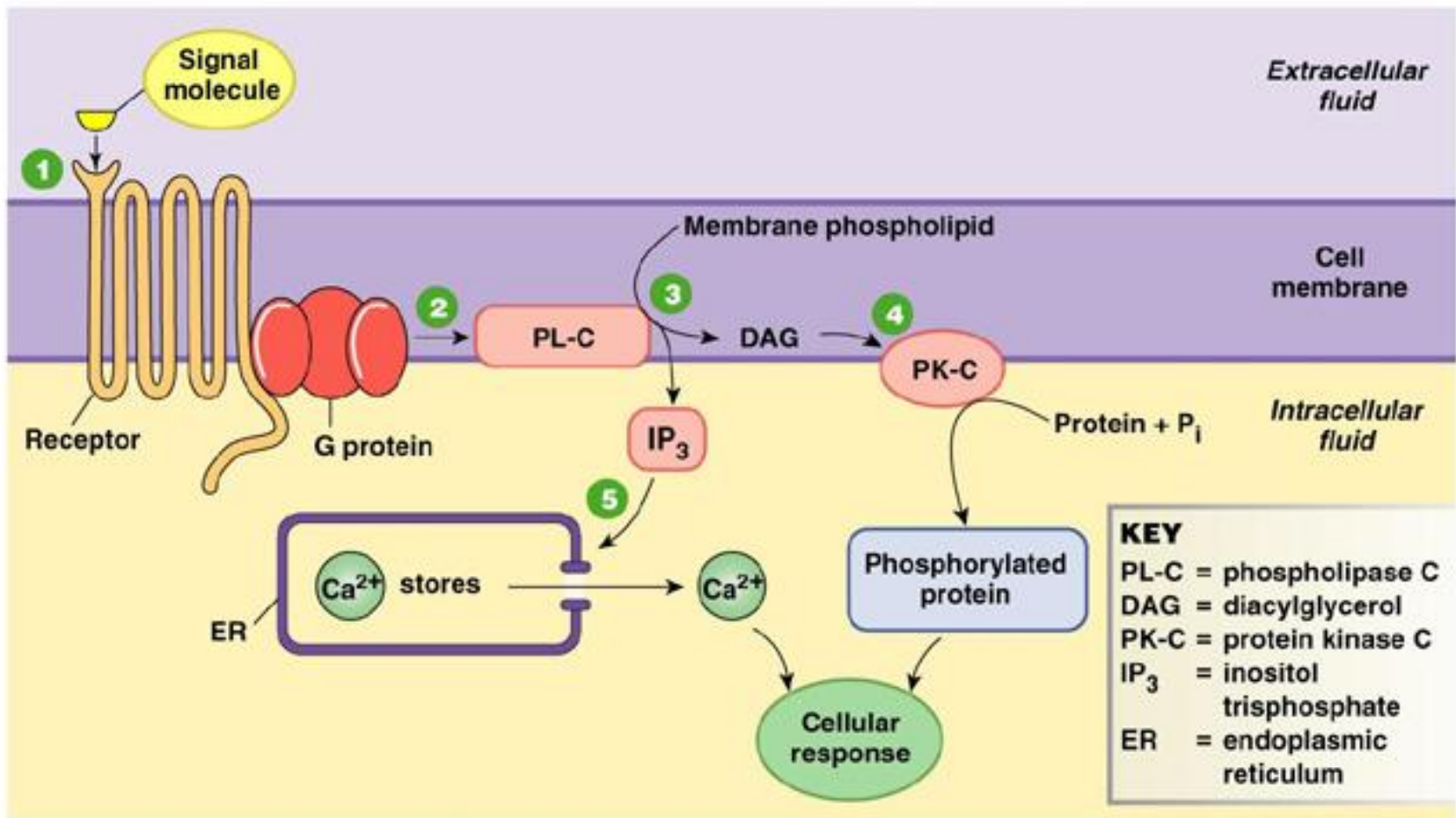
(a) Visceral (single-unit) smooth muscle tissue

(b) Multiunit smooth muscle tissue



(c) Details of a smooth muscle fiber





- 1** Signal molecule activates receptor and associated G protein.
- 2** G protein activates phospholipase C (PL-C), an amplifier enzyme.
- 3** PL-C converts membrane phospholipids into diacylglycerol (DAG), which remains in the membrane, and IP₃, which diffuses into the cytoplasm.
- 4** DAG activates protein kinase C (PK-C), which phosphorylates proteins.
- 5** IP₃ causes release of Ca²⁺ from organelles, creating a Ca²⁺ signal.

