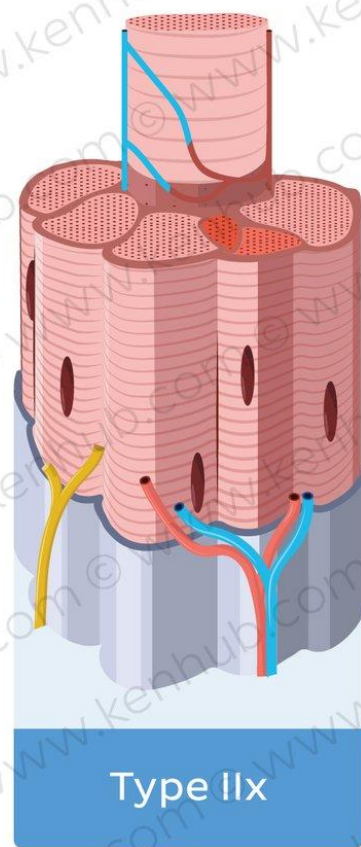
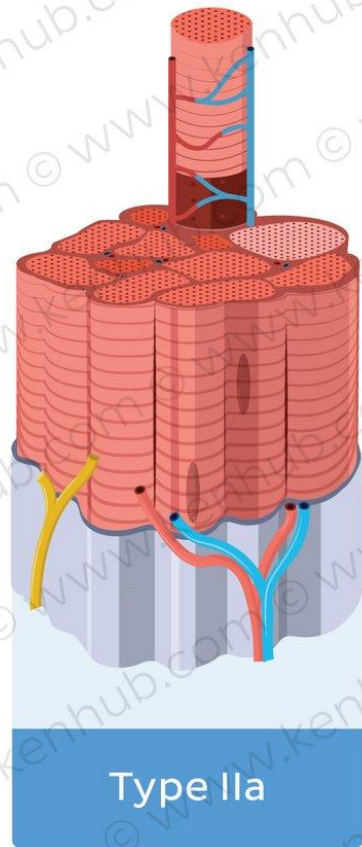
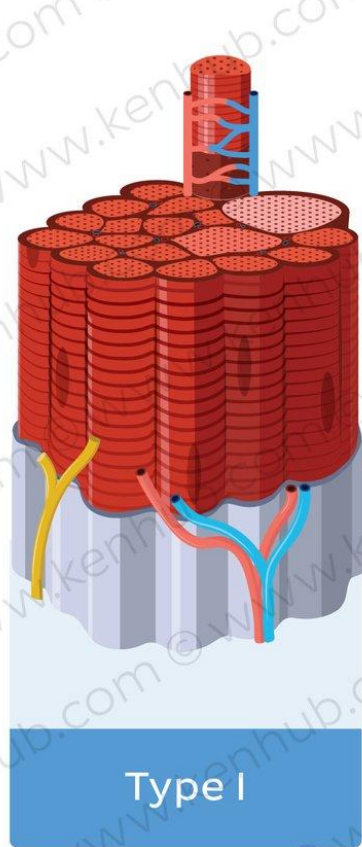


Medical Biology

Classification of skeletal muscle fibers

- Type I or slow, red oxidative fibers:
- Type IIa or fast, intermediate oxidative-glycolytic fibers:
- Type IIb or fast, white glycolytic fibers:

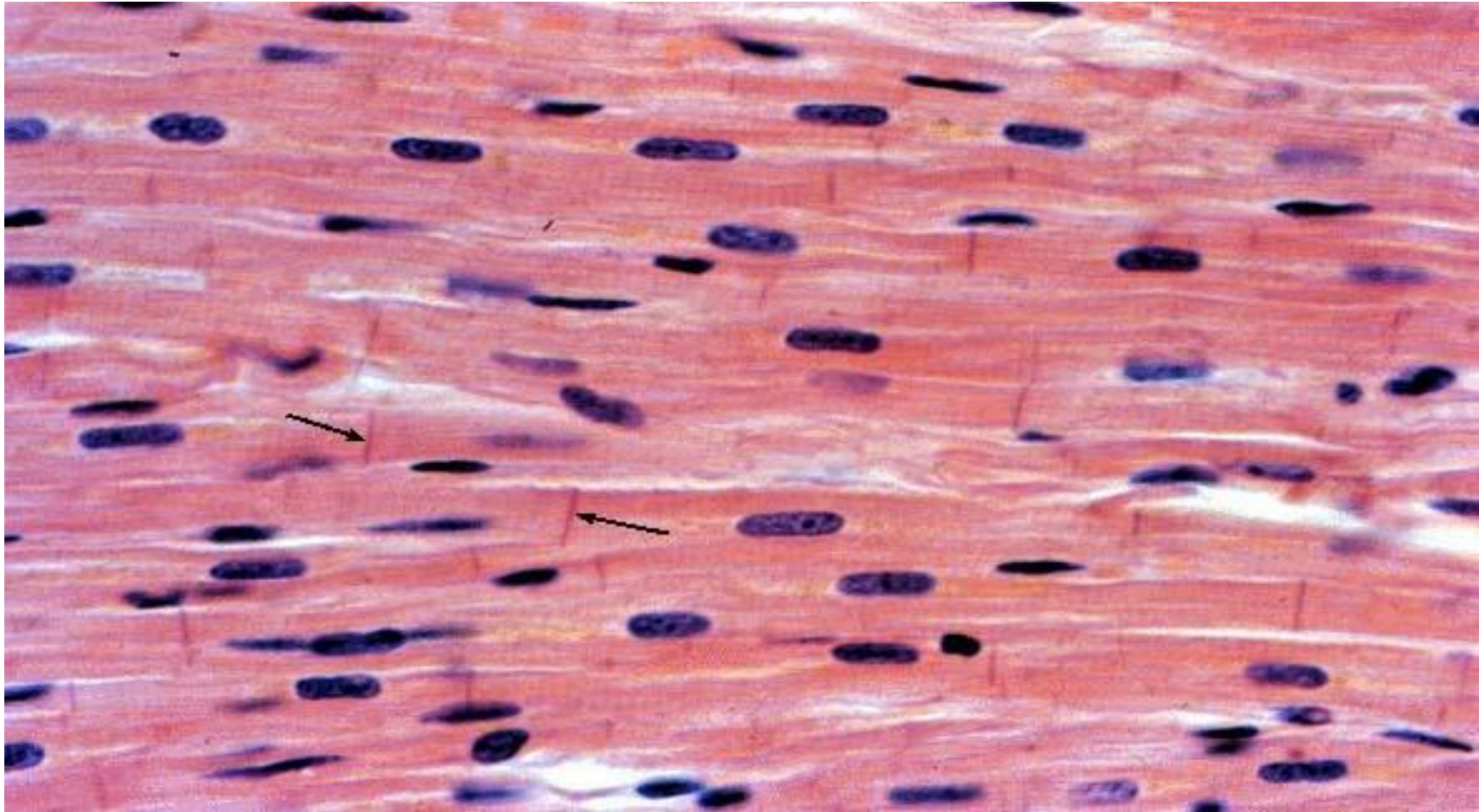


Atypical Striated Muscle

- cremaster muscle (near the spermatic cord).
- esophageal striated muscle, external urethral sphincter, external anal sphincter.

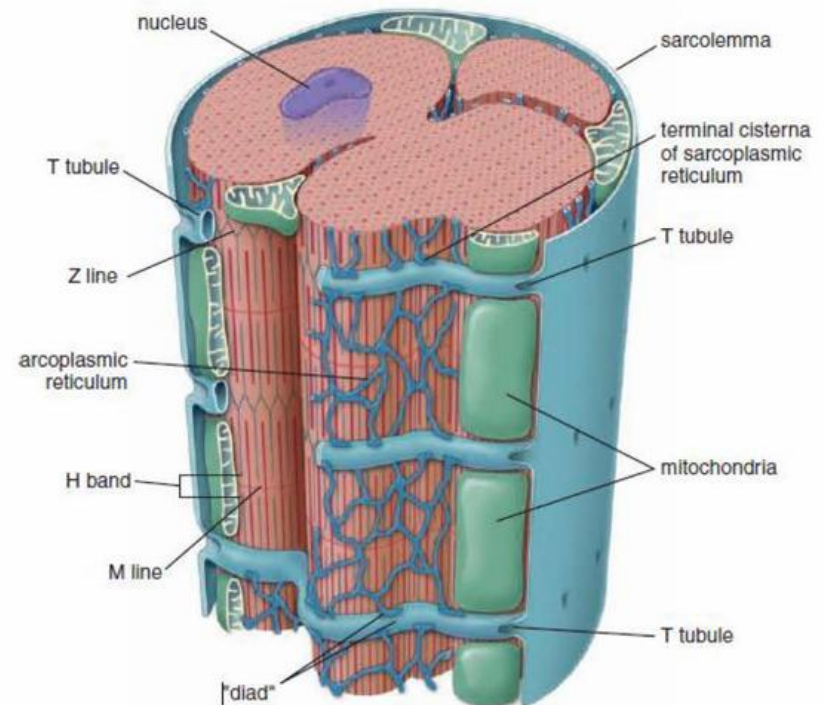
CARDIAC MUSCLE

- The muscle **fibers branch** (bifurcate) and are arranged in series to form an anastomosing network.
- Each myocyte has one or two **central nuclei** (unlike the many peripheral nuclei of syncytia of skeletal muscle fibers).
- The fibers have more sarcoplasm.
- The mitochondria are larger and better developed.
- **All the fibers are Type I** (red fibers, with abundant myoglobin).
- **Glycogen** may also present.
- The myocytes have specialized areas of contact - the **intercalated disks**.
- **Contractions are rhythmic, spontaneous and involuntary.**
- Delicate sheath of **endomysium** with a rich capillary network.
- A thicker **perimysium** separates bundles and layers of muscle fibers.

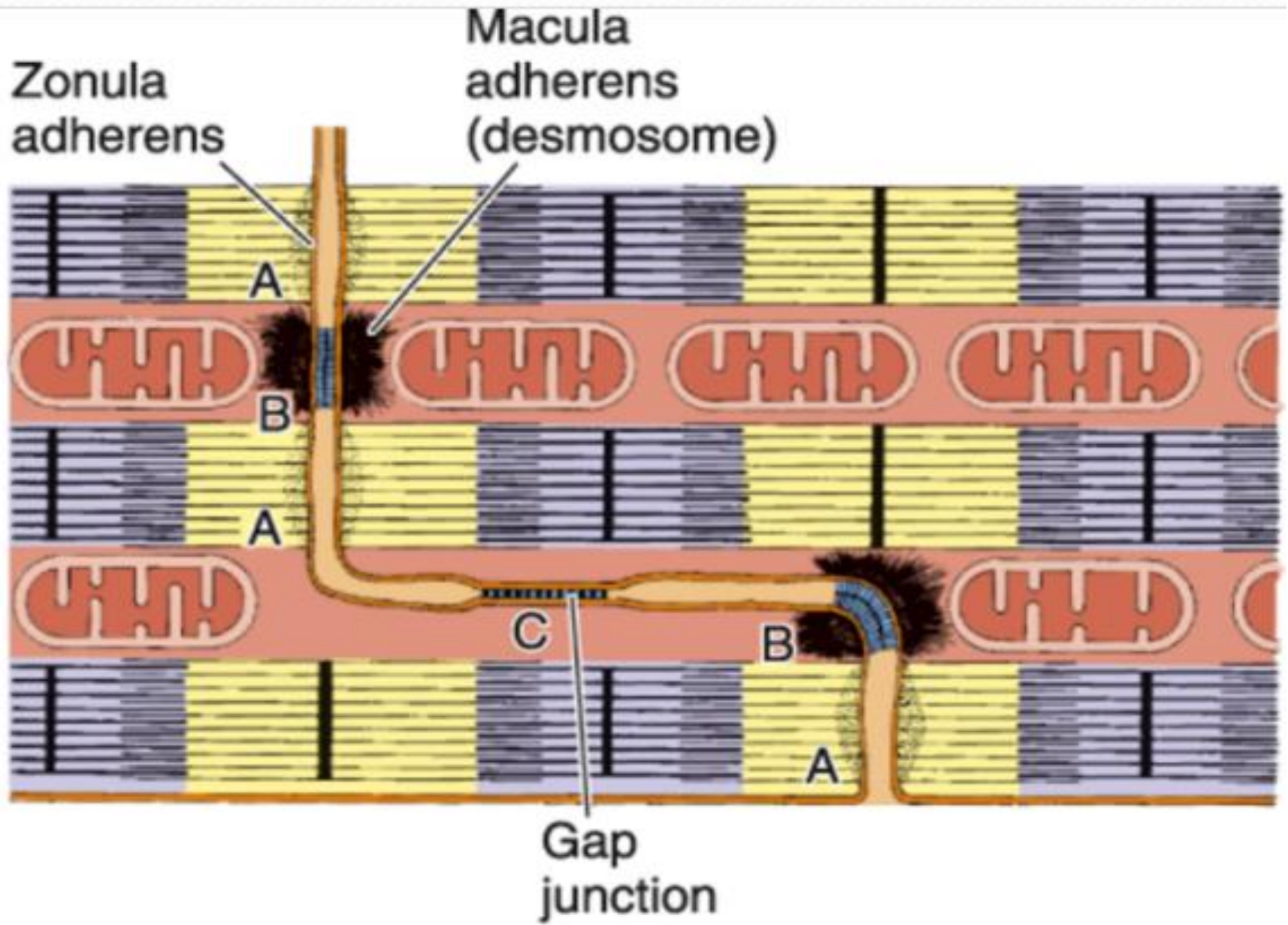


Cardiac muscle cells

- The **T tubule system** and **sarcoplasmic reticulum** are not as regularly arranged in the cardiac myocytes.
- **Diads end near Z disc**
- sarcomere
- Lipofuscin pigment granules (aging pigment)



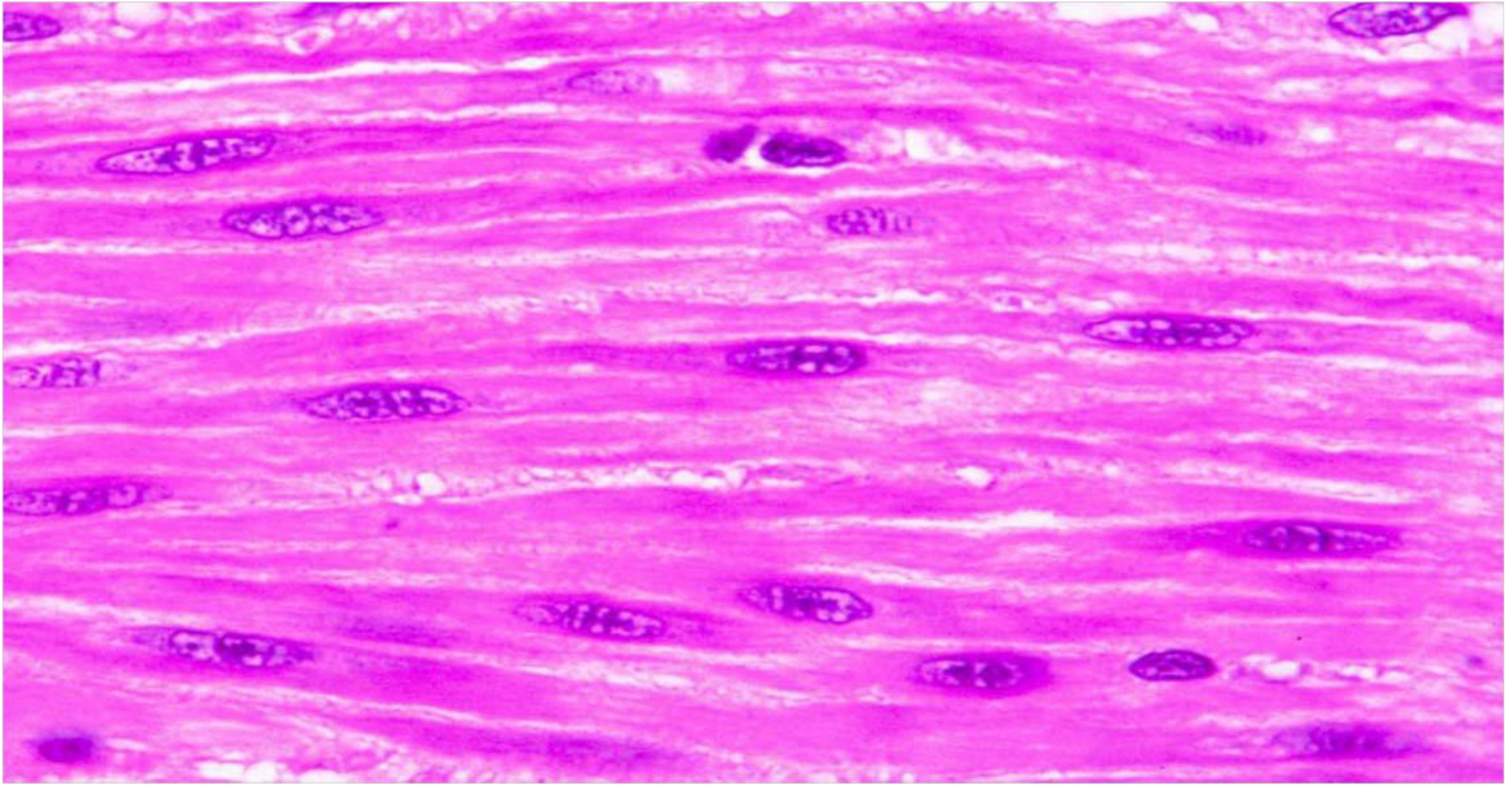
Intercalated disks



SMOOTH MUSCLE

**innervated by the autonomic nervous
system**

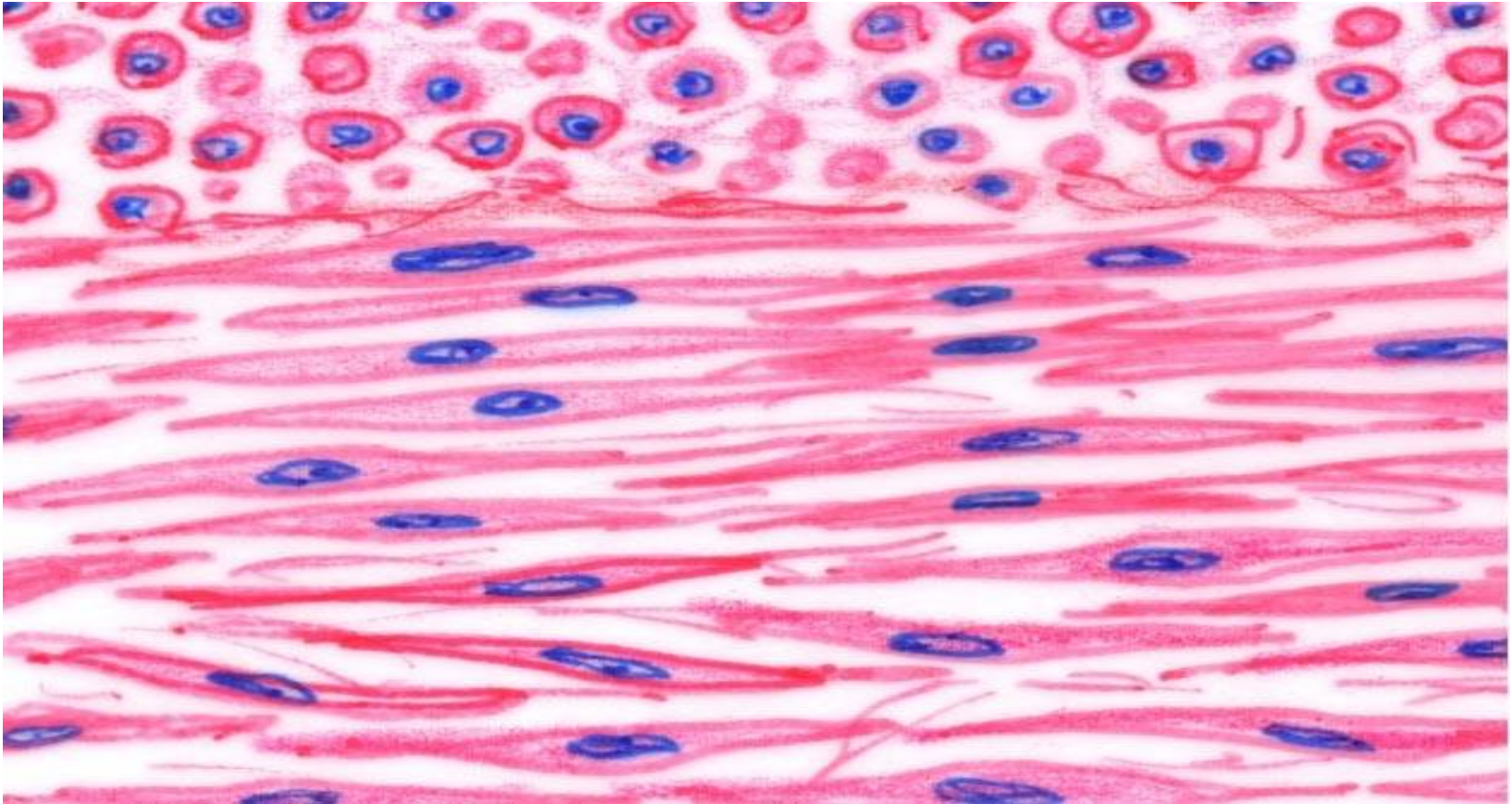
involuntary muscle



Location of smooth muscle

- Smooth muscle is found in the **walls of the hollow internal organs**
- **walls of blood vessels** (vascular smooth muscle, especially in arterial vessels).
- Smooth muscle is found in the **dermis of the skin** (arrector pili).
- Smooth muscle is found in the eye (**iris diaphragm**, controlling the amount of light reaching the retina).

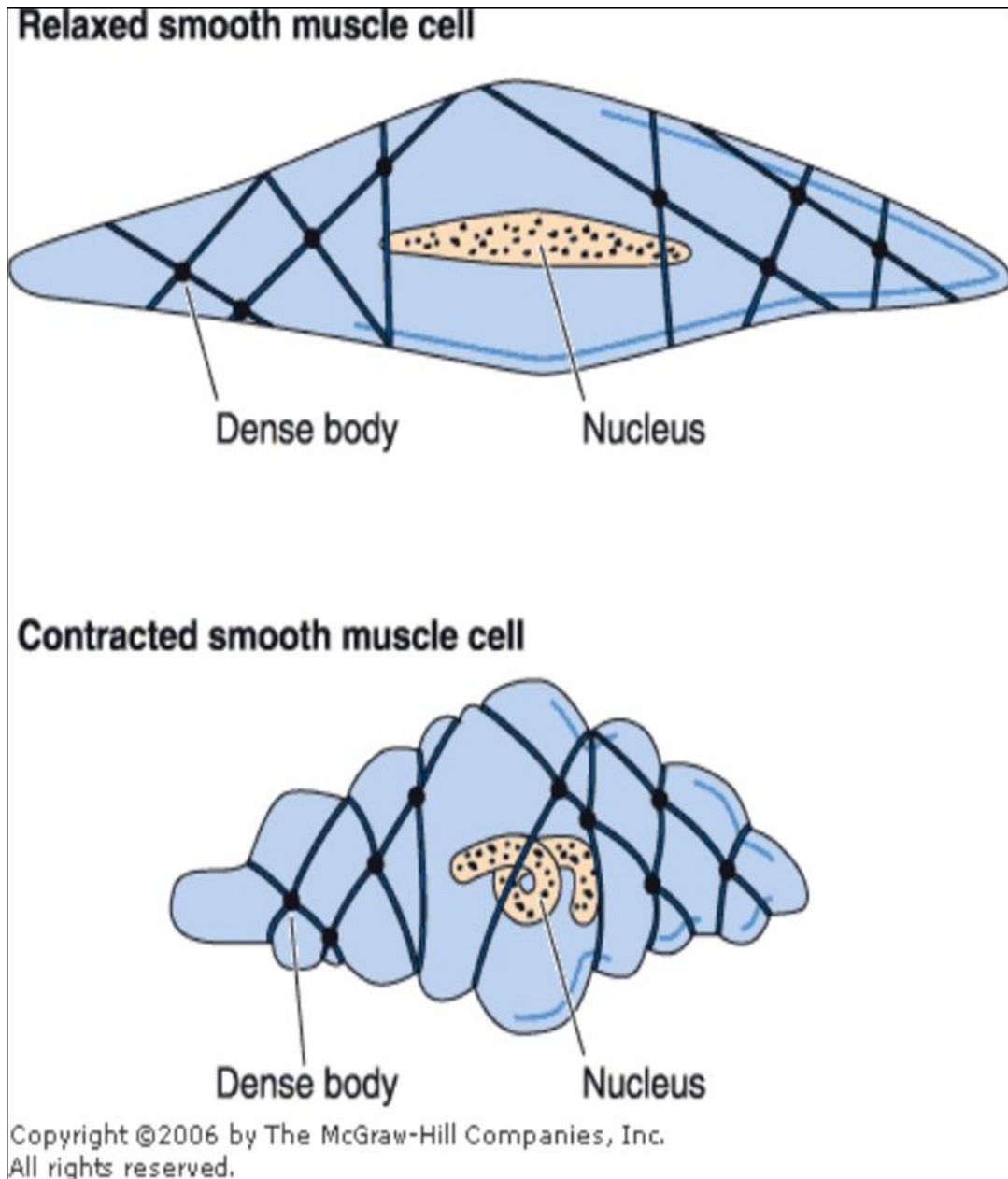
Structure of smooth muscle fibers



Smooth muscle sheath

- **Sheath (proteoglycan, reticular, collagen & elastic fibers)**
- **Myofilaments:**
 - 1. thin myofilaments (actin)** which are the most common type
 - 2. thick myofilaments (myosin)** which are less common
 - 3. intermediate filaments (desmin)** These may be grouped as "dense bodies" and are also found in contact with the sarcolemma (attachment plaques of thin and intermediate filaments that are functionally similar to Z disc of skeletal and cardiac muscles).

No T-tubule
sacculles
caveolae.



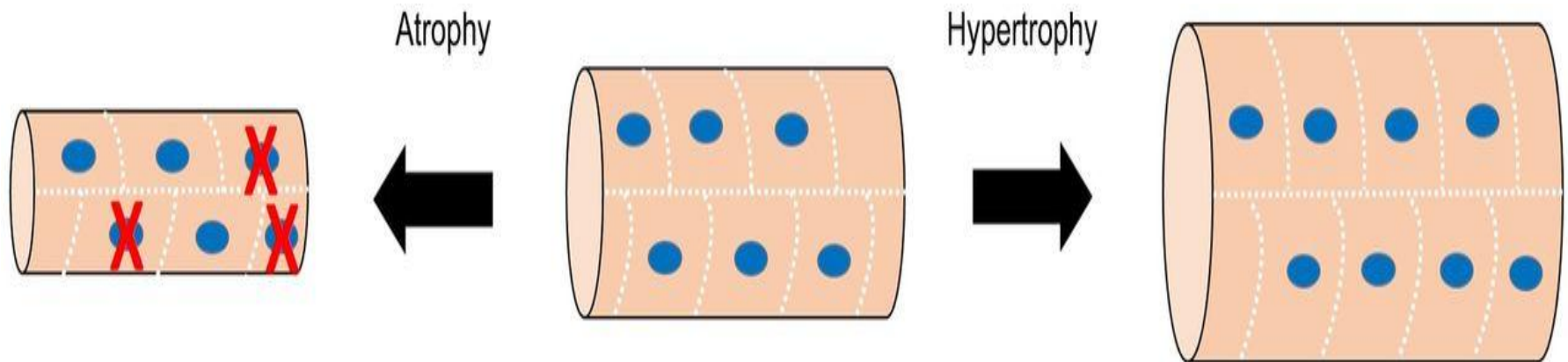
Origin of smooth muscle

- **Mesoderm**
- **From mesenchyme as connective tissue cells**
- **myoepithelial cells**
- **part of the esophagus, anal sphincter, tarsi of eyelids**

Repair and regeneration after injury

Skeletal muscle

- hypertrophy of use
- disuse myopathy or atrophy

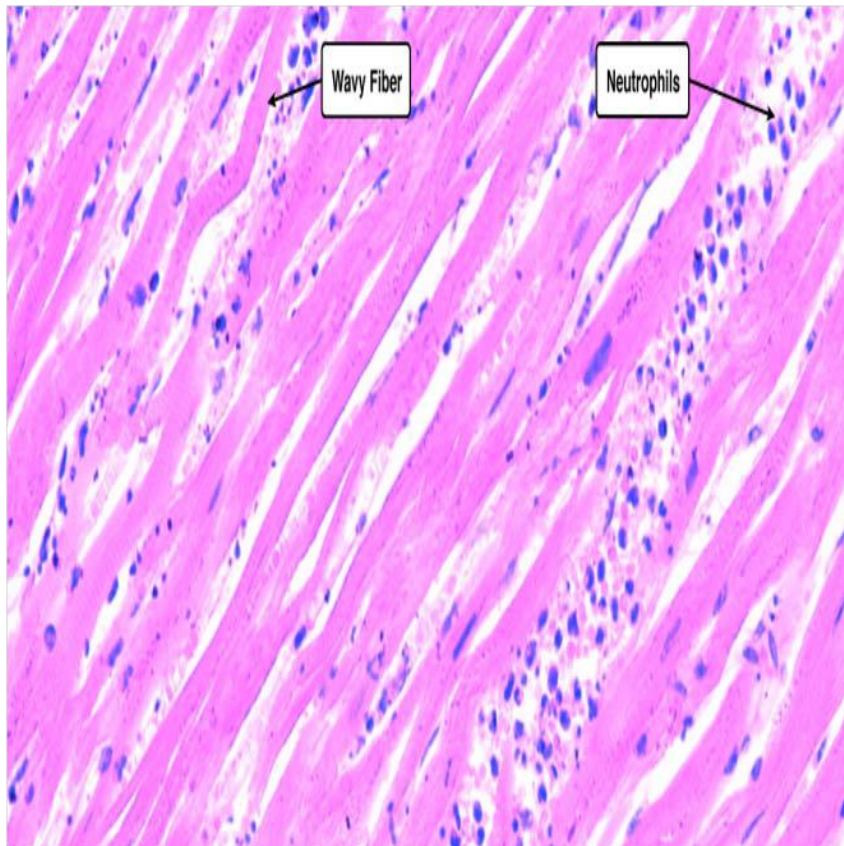


- Loss of myonuclei (apoptosis)
- Myofibre CSA ↓
- Myofibrillar proteins ↓

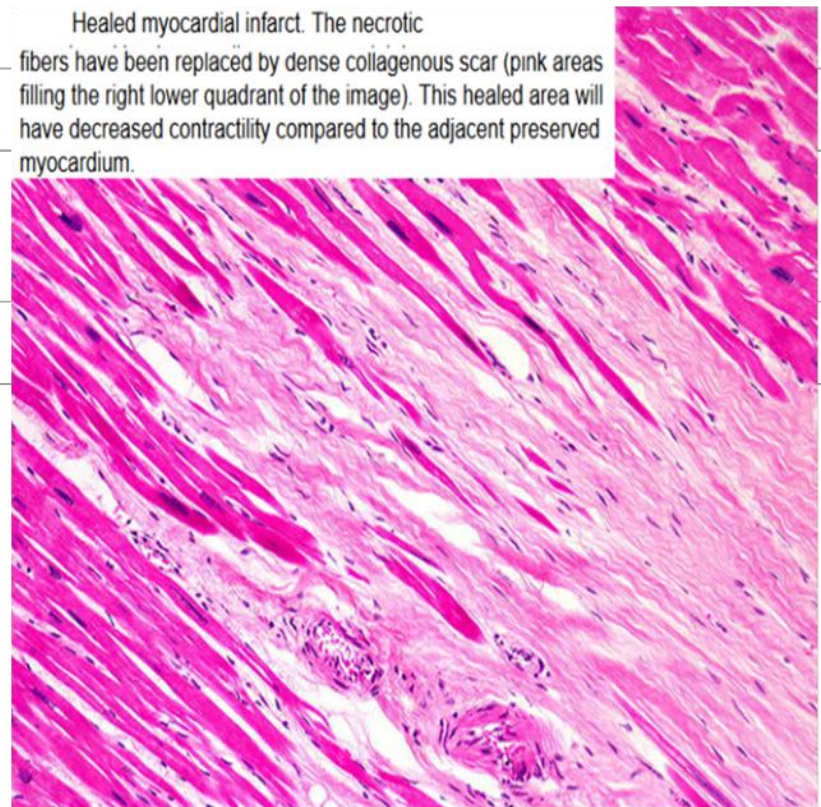
- Addition of myonuclei
- Myofibre CSA ↑
- Myofibrillar proteins ↑

Regeneration of cardiac muscle

Acute Myocardial Infarction



Healed myocardial infarction





Regeneration of Smooth muscle

hyperplasia and hypertrophy

Muscular dystrophy

Duchenne muscular dystrophy

- Typical feature – the child uses his hands to climb up, while getting up from the floor.
- Wheelchair by age 12
- Fatal by age 30

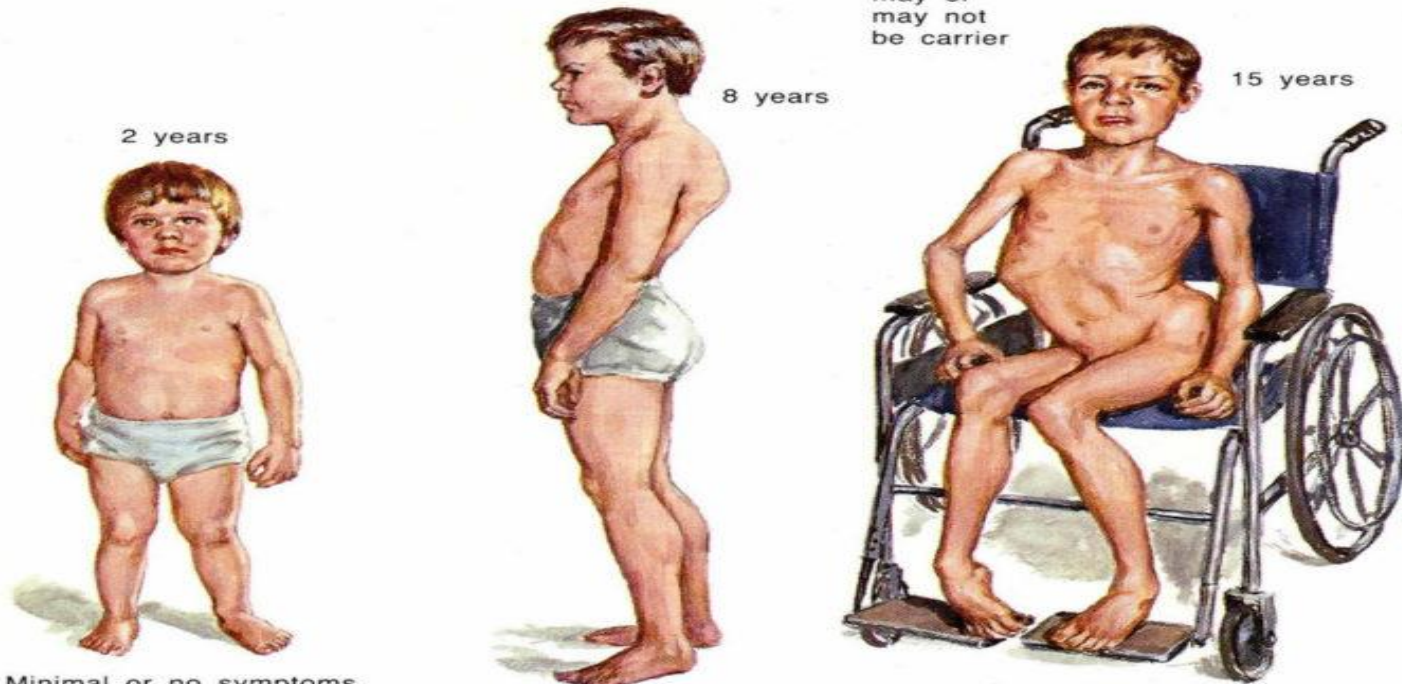
Duchenne's Muscular Dystrophy

Sex-linked recessive inheritance

Mother normal, carrier

Father normal

Only males affected, but females may be carriers



Minimal or no symptoms

Severe crippling deformities and contractures

Progression with age { Weakness, especially of pelvic girdle muscles; marked lordosis, enlarged calves

Calf muscles usually but not always enlarged



Lordosis disappears when child sits

Elevidys (delandistrogene moxeparvovec-rokl) is a one-time [gene therapy treatment](#) for [Duchenne muscular dystrophy \(DMD\)](#), made by the pharmaceutical company Sarepta. Approved by the [U.S. Food and Drug Administration \(FDA\)](#) in 2023, Elevidys is designed to strengthen muscles by delivering a shortened version of the dystrophin gene, known as micro-dystrophin. This compensates for the missing or broken dystrophin gene that is the cause of DMD.

To deliver the micro-dystrophin gene, Elevidys uses a “vector” made from a harmless virus called *AAVrh74* that is good at getting inside of cells. Once infused into the bloodstream, Elevidys gets into muscle cells, prompting them to produce the micro-dystrophin protein. This protein helps protect muscle cells and prevent them from degenerating.



Thank you &
Good luck