



# 4- MUSCLE PHYSIOLOGY



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2024

# The simple muscle twitch

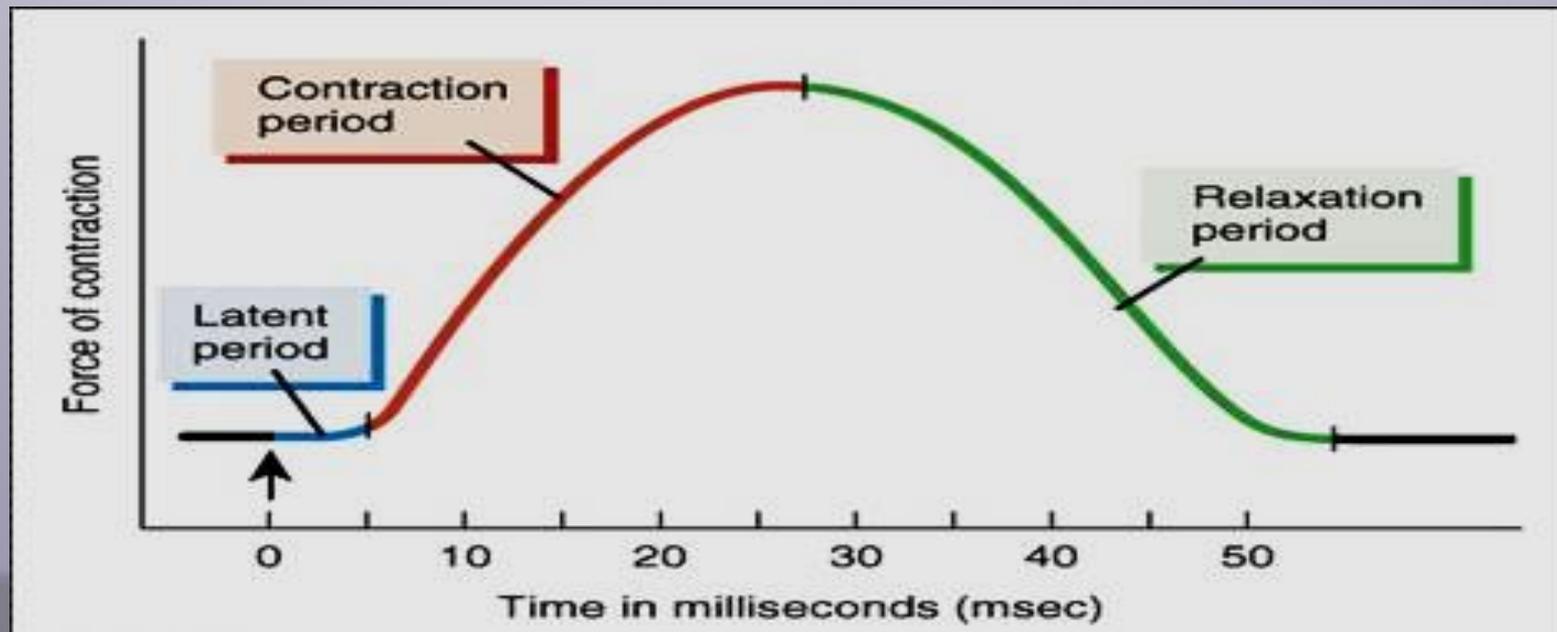
Definition: It is the response of the muscle to a single maximal stimulus and consists of:

**1) Latent period:** -It is the time between time of stimulus & response.

-About 0.01 second duration. - Due to: 1- conduction of impulse in nerve 2- production of MEP potential. 3-conduction of impulse in the muscle. 4- contraction and 5- the time of recording.

**2) Contraction period:** during it the muscle contracts either isometrically or isotonicly.(0.04 sec.)

**3) Relaxation period:** the muscle relaxed (= 0.05 sec. In isotonic relaxation).



## Factors affecting the simple muscle twitch:

**1-Type of muscle:** there are 2 types of muscle fibers:

<b>Red muscle fibers</b>	<b>White (pale) muscle fibers</b>
1- Of type I & slow fibers. 2- Rich in myoglobin (red) 3-fibres are small in size 4-supplied by small, slow nerve 5- More blood supply 6-Contain large number of mitochondria and depend on aerobic metabolism 7-Respond slowly but with long duration 8- Not early fatigued 9-Adapted for prolonged muscle activity (Static function) 10- e.g antigravity muscles to maintain body posture.	- of type II fibers & fast fiber - poor in myoglobin (pale) - the fibers are large in size - supplied by large rapid nerve - less blood supply -contains few number of mitochondria and depend on anaerobic metabolism  -it responds rapidly but with short duration - it early fatigued -Adapted for rapid, fine, skilled Movement (Phasic function)  -e.g. extraocular muscle

**N.B.:** most muscle contain both types but one is predominant.

## 2- Temperature:

Warming of the muscle as in muscular exercise leads to stronger and rapid contraction by acceleration of the chemical reactions and decrease the muscle viscosity.

But overheating ( $> 45^{\circ}\text{C}$ )  $\rightarrow$  heat rigor (stiffness).

## 3- Initial length:

The strength of contraction (in isotonic contraction) and the developed tension (in isometric contraction) are directly proportional to the initial length of the muscle fibre up to limit (**Starling's law**).

## 4- Fatigue:

• **Definition:-** It is the gradual decrease in the muscle contraction and prolonged duration of all phases of the SMT, especially relaxation due to repeated and strong stimulation of the muscle. • **The effect:** decrease strength and prolonged duration of contraction and incomplete or absent relaxation

• **The cause of fatigue:** - In case of **indirect** stimulation (via stimulation of its motor nerve) is the gradual exhaustion of Ach at the MEP.

- Also **direct** stimulation of the muscle may lead to fatigue due to exhaustion of energy sources (ATP) or accumulation of metabolites.

- In living muscle (after exercise), **fatigue** is caused by:

1- Decrease blood supply to the muscle.

2- Decrease energy sources.

3- Accumulation of metabolites which depress the brain and spinal cord (central effect).

- **Contracture** may occur with fatigue due to decrease in ATP required for separation between the thin and thick filaments and muscle relaxation.

## 5- Stair-case (Treppe) phenomenon:

-It occurs in the skeletal and cardiac muscle.

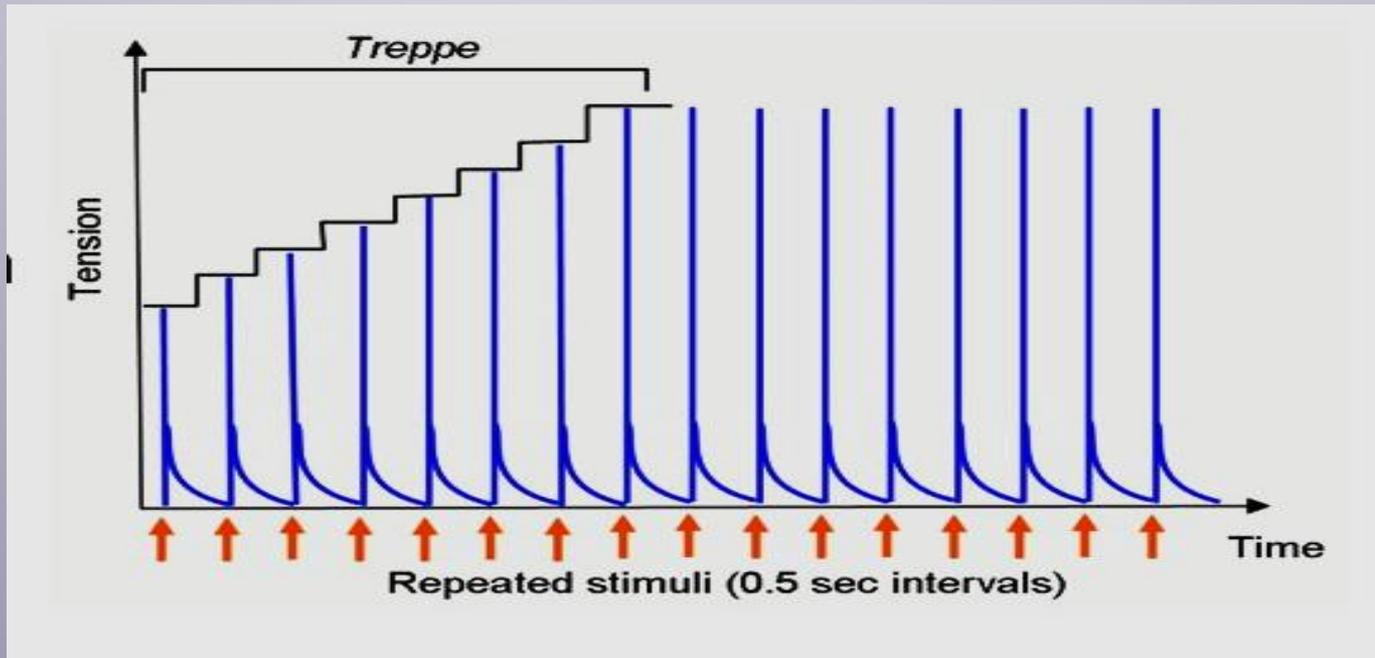
-It is a gradual increase in muscle contraction until plateau.

-This occurs by application of series of maximal stimuli just after relaxation period of each muscle twitch.

-This is due to: 1- accumulation of  $\text{Ca}^{++}$  intracellular.

2-  $\uparrow$  temperature of the muscle.

3-  $\downarrow \text{K}^+$  &  $\uparrow \text{Na}^+$  intra-cellular  $\rightarrow \uparrow \text{Ca}^{+2}$  release from sarcoplasmic reticulum  $\rightarrow \uparrow$  contraction.



# Summation of muscle contractions

Since the contraction phase in the skeletal muscle starts with the relative refractory period, the muscle respond to another stimulus during either cont. or relaxation → summation of contraction.

## (a) Effect of two successive stimuli:

According to frequency of stimulation:

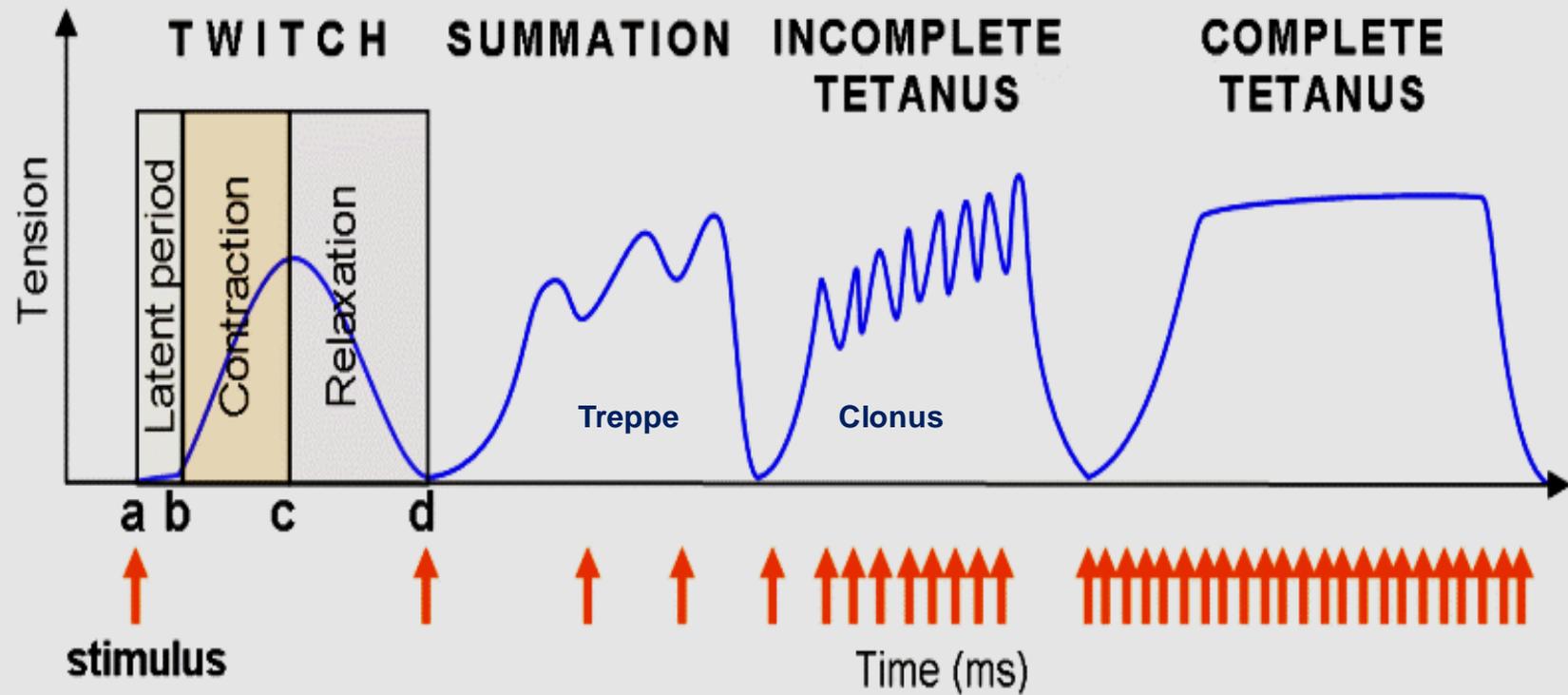
If the 2<sup>nd</sup> **stimulus** falls in relation to preceding one:

- 1- During the **latent period** → no response (during ARP).
- 2- During the **contraction period** → more strong contract.
- 3- During the **relaxation period** → 2 peaks contraction.
- 4- Just **after the relaxation period** → stair-case phenomenon.
- 5- **After relaxation** → normal second contraction.

## (b) Effect of multiple successive stimuli:

- 1- If the frequency is low → separate twitches with **Stair – case** phenomenon.
- 2- If the frequency increases and stimuli falls during relaxation phase of preceding twitch → **Clonus** (incomplete tetanus).
- 3- If the frequency increases more and stimuli falls during contraction phase → sustained contraction (**complete tetanus**).

N.B.: Cooling, fatigue & anticholinesterase (Eserine) change **clonus into complete tetanus**. However, warmth and rest cause the reverse.



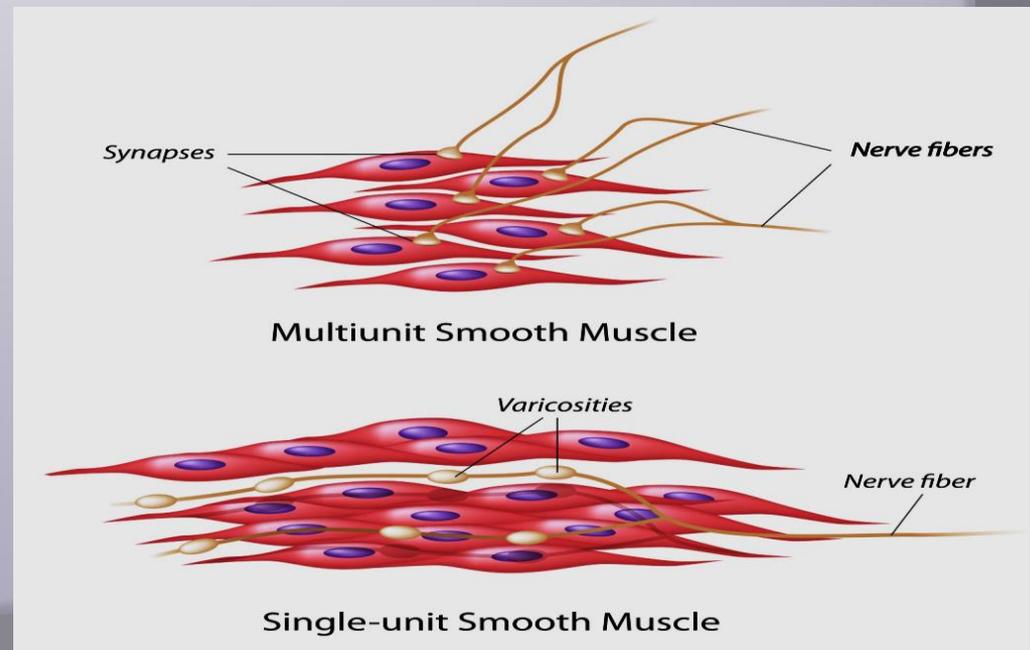
# Smooth Muscle

•Site: it presents in wall of most viscera, bl. vessels, some glands, intraocular muscles and erector pilae ms. So, it controls the involuntary activities.

## •Structure:

- Smooth muscle fibers are spindle-shaped, non striated (plain) cells with central long nuclei.
- Fiber's length is 20-500 microns and diameter 2-5 microns.
- S.M. contains **more** actin filaments which attached to each other and to dense bodies .
- S.M contain **calmodulin** instead of troponin-tropomyosin.
- S.M contain **less** mitochondria and endoplasmic reticulum.
- S.M innervated by autonomic nervous system.

## Types of smooth muscles



•**S.M of two types:**

-S.M has no motor end plate but at nerve endings (sympathetic or parasympathetic) there are special nodes (varicosities) via it neuromuscular transmission occurs → depolarization followed by contraction or hyperpolarization followed by relaxation according to the type of chemical transmitter.

<b>Multi-unit S.M.</b>	<b>Single unit S.M (unitary)</b>
<ul style="list-style-type: none"> <li>-Separate fibers without connection. (Except via the chemical transmitters).</li> <li>- One nerve for each fiber</li> <li>- Not obey all or none law</li> <li>-Sensitive to chemical transmitter</li> <li>-Rare spontaneous cont. but controlled by nerve impulses.</li> <li>-Not respond to stretch</li> <li>-e.g iris, wall of bl. vs., pilomotor muscle&amp; ciliary muscle</li> </ul>	<ul style="list-style-type: none"> <li>-Aggregated fibers attached by gap junctions facilitate conduction of action potential (functional syncytium).</li> <li>-one nerve for many fibers.</li> <li>-Obey all or non law</li> <li>- less sensitive.</li> <li>-Contract spontaneously</li> <li>- Respond to stretch</li> <li>- e.g wall of viscera as uterus, GIT, ureter,.....</li> </ul>

# Excitability of S.M

- RMP: is **unstable** and about  $-40$  to  $-60$  mvolt. with slow sine waves.

- Action potential of **four** types:

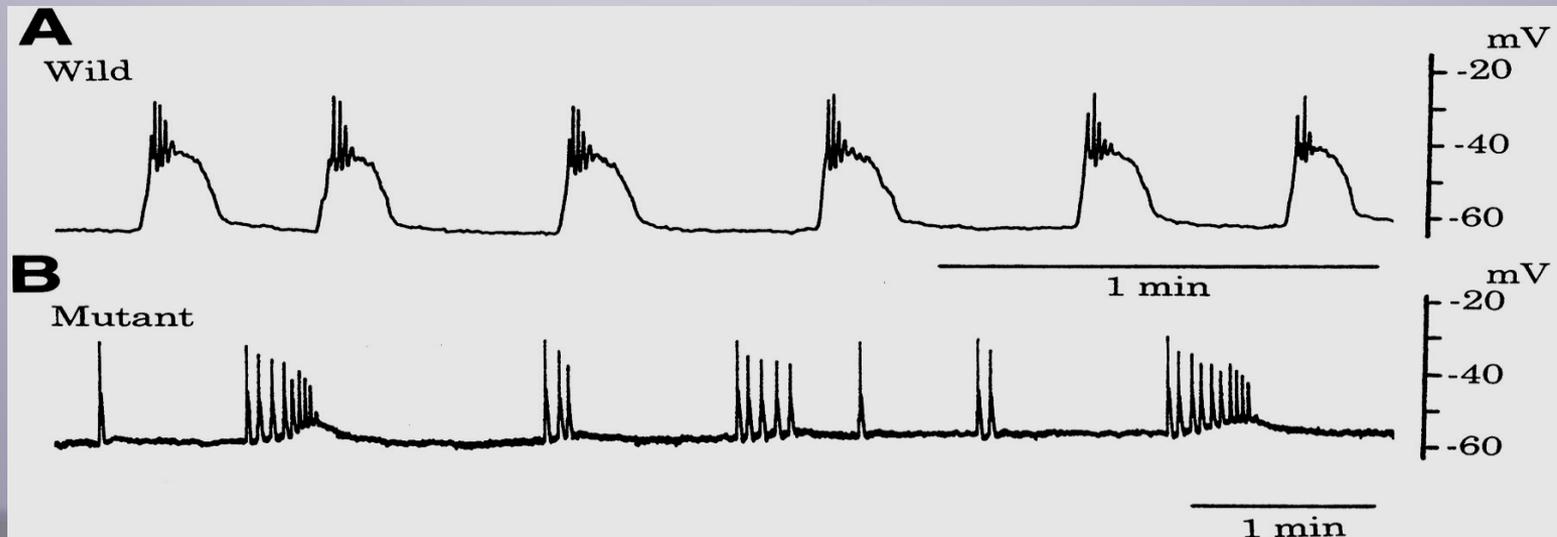
- a-Spike potential** as in sk. ms with duration of 50 msec. Present in the single unit S.M fibers.

- b-Action potential with prolonged plateau** (hundreds of m.sec) due to delayed repolarization as in uterus (similar to cardiac potential).

- c-Spike pot. with serrations** as in small intestine.

- d-Pacemaker potential** (slow – wave potential): It occurs due to rhythmical activity of  $\text{Na}^+$ - $\text{K}^+$  pump. When the wave reach the firing level ( $-35$  mv)  $\rightarrow$  action potentials which spread over the muscle. This type of potential initiates rhythmical contraction of GIT (as in the automatic cardiac fibers).

- Ionic base of action potential:** may due to  $\text{Na}^+$  **influx** or  $\text{Ca}^{++}$  **influx** or **both**.



# Contractility of S.M.

## •Excitation contraction coupling:

-**Contraction:** Extracellular  $Ca^{++}$  influx or intracellular  $Ca^{++}$  release from the sarcoplasmic reticulum  $\rightarrow$   $\uparrow$  intracellular  $Ca^{++}$  which combine with calmodulin  $\rightarrow$  activate myosin light chain kinase enzyme  $\rightarrow$  phosphorylation of the light chain of myosin  $\rightarrow$  binding of actin & myosin  $\rightarrow$  shortening (contraction).

-**Relaxation:**  $\downarrow$  intra-celular  $Ca^{++}$  (by  $Ca^{++}$  pump)  $\rightarrow$  stimulate myosin phosphatase enzyme  $\rightarrow$  removal of phosphate from light chain of myosin  $\rightarrow$  stop contraction  $\rightarrow$  relaxation.

## -Characters of S.M. contractility:

1-Spontaneous contraction but under nervous regulation.

2-**Slow** cycling of cross bridges

3-**Slow** onset of contraction and relaxation.

4-Energy and  $O_2$  consumption is **low** and depends mainly on anaerobic glycolysis. So it is not easily fatigued.

5-SM has great ability to shorten as far greater percentage of its length.

6-Its contraction is **sluggish** and excitation / contraction coupling is **very slow** also  $Ca^{++}$  pump is slow so contraction is maintained than in skeletal muscle.

7-**Latch mechanism**, as prolonged tonic contraction needs less energy, less nervous or chemical stimulation than initial activity. So this **delays fatigue**.

8- **Stress relaxation (plasticity)** in which if SM is slowly stretched  $\rightarrow$  increased tension at first then the tension gradually decreases inspite of continuous stretch (e.g., the urinary bladder can receive large volumes of urine without marked increase in wall tension).

9-Visceral SM shows: **Tone** = continuous mild contraction

**Rhythm** = irregular cont. due to repetitive discharge of spike potential.

## Factors affecting excitability & contractility

	↑Excitability → contraction	↓Excitability → relaxation.
-Motor neurons -Temperature -Stretch -pH <sup>+</sup> -Osm. Pressure -Ions -Autonomic drugs -Hormones	Parasympathetic Cooling Rapid moderate stretch Alkalinity Low ↓ Ca <sup>++</sup> & ↑ K <sup>+</sup> Parasympathomimetics Vasopressin, Oxytocin and Estrogen.	Sympathetic Warmth Severe stretch Acidity High ↑ Ca <sup>++</sup> & ↓ K <sup>+</sup> Sympathomimetics Catecholamines, Progesterone

# Thank You

