Local anesthesia

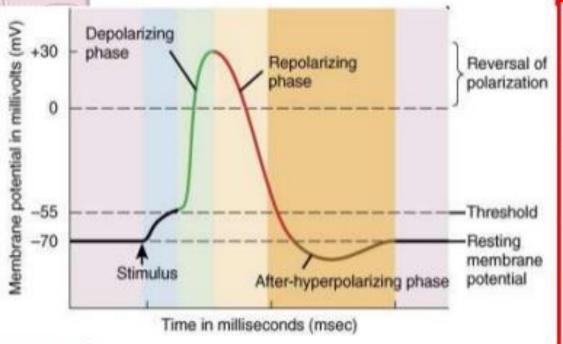
Definition

Tegniques depend on a group of drugs that produces transient loss of sensory, motor, and autonomic function when the drugs are injected or applied in proximity to neural tissue.

Mechanism of action

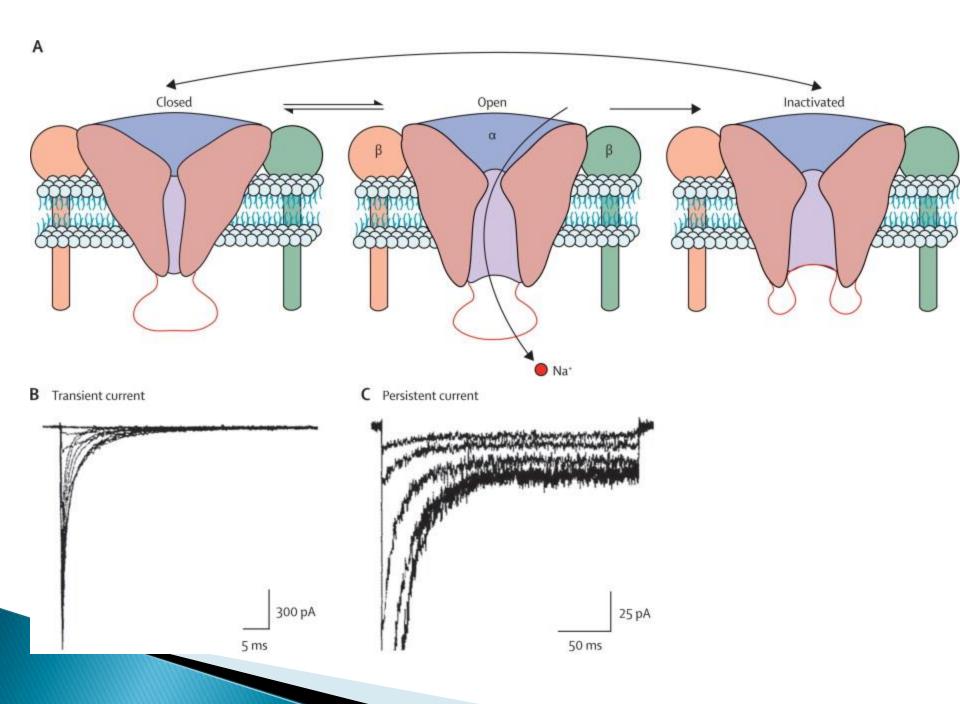
- ▶ An electrogenic Na- K-ATPase pump couples the transport of three Na ions out of the cell for every two K ions moves into the cell, this creats a concentation gradient that favors extracellular diffusion of K and intracellular diffusion of Na. This accounts for the negative resting potential difference (-70 mV polarization).
- If the depolarization exceeds a threshold level (−55mV), sodium channels are activated allowing a sudden influx of Na ions and generating action potential.

Action Potential



- Resting membrane potential is -70mV
- triggered when the membrane potential reaches a threshold usually -55 MV
- if the graded potential change exceeds that of threshold – Action Potential
- Depolarization is the change from -70mV to +30 mV
- Repolarization is the reversal from +30 mV back to -70 mV)

- action potential = nerve impulse
- takes place in two stages: depolarizing phase (more positive) and repolarizing phase (more negative - back toward resting potential)
- followed by a hyperpolarizing phase or refractory period in which no new AP
 can be generated



- Sensitivity to blockade is determined by axonal diameter and degree of myelination
- In spinal nerves, the sensitivity to LA is autonomic> sensory > motor.
- LA consist of benzene ring seperated from tertiary amine by intermediate chain that includes an ester or amide linkage.

Table 2: Nerve Fiber Types and Nerve Blocking

Fiber Type	Function	Diameter (microns)	Mystification	Conduction Velocity (m/s)	Sensitivity to Nerve Block
Type A					
Alpha (α)	Proprioception, motor	12-20	Heavy	70-120	+
Beta (β)	Touch, pressure	5-12	Heavy	30-70	++
Gamma (y)	Muscle spindles	3-6	Heavy	15-30	++
Delta (8)	Pain, temperature	2-5	Heavy	12-30	+++
Туре В	Preganglionic autonomic	<3	Light	3-15	++++
Туре С					
Dorsal root	Pain	0.4-12	None	0.5-2.3	++++
Sympathetic	Postganglionic	0.3-1.3	None	0.7-2.3	++++

- Pain practitioners block the nerves transmitting pain impulses (Type A-8, Type C)
- Lower concentrations of local anesthetic will only block the small unmyelinated and lightly myelinated (Type C and Type A-8) fibers
- Middle-frequency currents (2,000-20,000 Hz) block smaller unmyelinated (Type C) and small myelinated (Type A-8) fibers
- Larger fibers (Type A-α, β, γ) require high-amplitude currents and are usually spared in electrical, low-dose chemical (eg, labor epidural) blocks

Clinical pharmacolgy

- Absorption: systemic absorption of LA depends on blood flow which is determined by factors:
- Site of injection: IV > Tracheal> intercostal > caudal > paracervical > epidural > brachial plexus > sciatic > subcutaneous.
- 2. Presence of vasocontrictors
- 3. Local agent
- Metabolism :
- Ester LA metabolized by pseudocholinesterase
- 2. Amide metabolized by P-450 in the liver .

Agent	Max Dose w/o Epi	Max Dose w/ Epi	Duration of Action	Notes
Lidocaine	5mg/kg	7mg/kg	30 - 90 min	1% = 10mg/mL 2% = 20mg/mL
Bupivacaine	2.5mg/kg	3mg/kg	6 - 8 hrs	0.5% = 5mg/mL
Mepivicaine	7mg/kg	8mg/kg		
Ropivacaine	3mg/kg	Marie Para Marie Para Para Marie Para Para Para		

The systemic effect of LA (toxicity)

- CNS: early symptoms numbness, tongue paresthesia, dizziness, sensory compliants may include tinnitus and blurred vision, excitatory signs may precede CNS depression, muscle twitching and seizures, respiratory arrest often follows
- RESPIRATORY: apnea can result from phrenic and intercostal nerve paralysis or depression of the medullary respiratory center
- CARDIOVASCULAR : the cardiotoxic reaction from accidentally intravascular injection of bupivacaine include hypotension, AV block, arrhythmias

The adjuncts to anesthesia