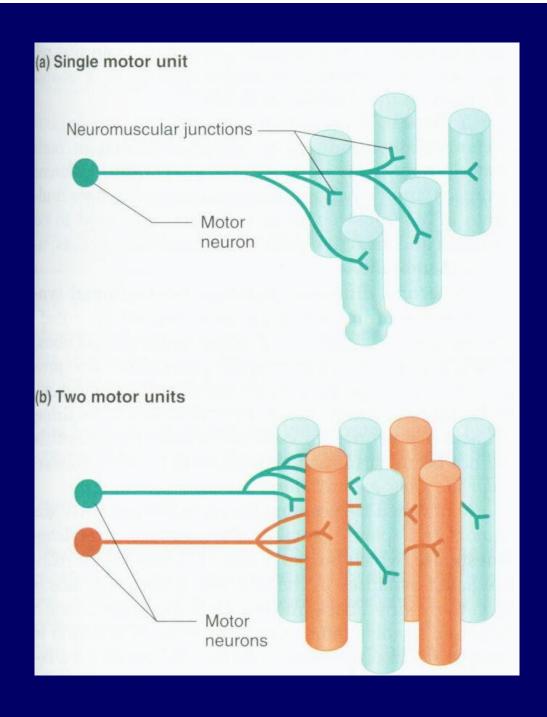


### **Motor Unit**

Consists of a motor neuron and all the muscle fibers it innervates

When an action potential occurs in a motor neuron, all the Msl fibers in its MU are stimulated to contract



**EMG** is the recording of electrical activity of a Msl at rest & during contraction: (to evaluate the electrophysiology of a MU)

Activity is amplified and displayed on an oscilloscope.

Instrument: Electromyograph

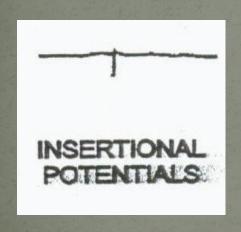
**Record:** Electromyogram

## MA concentric needle Ede inserted into the belly of the Msl.



Needle EMG does not introduce any electrical stimulation instead it records the intrinsic electrical activity of skeletal muscle fibers.

Normally a muscle is **silent at rest** after insertional activity has ceased.





 Then the patient is asked to contract the Msl smoothly.

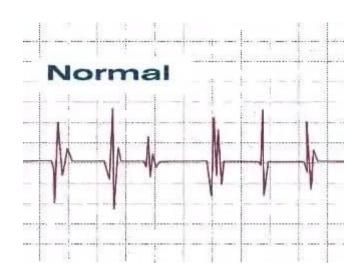
With muscle contraction, MUs are activated and MUAPs appear on the screen:



Motor unit potential: represents the summation of the potentials generated by µsl fibers belonging to the MU

### Normal MUPs

- Bi Triphasic
- Duration 3 16 mSec.
- Amplitude 300μV 5 mV

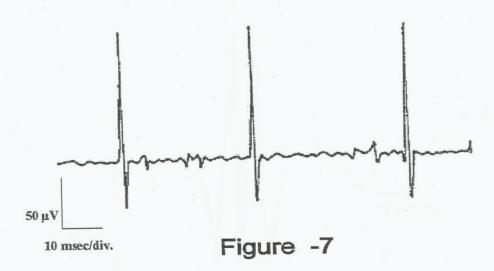


With increasing strength of contract<sup>o</sup>

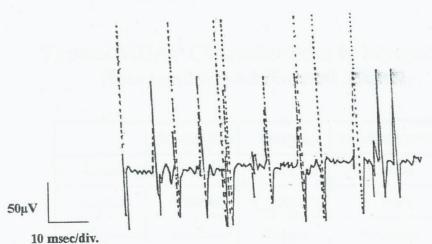
→recruitment of MUs →↑number & size of MUAPs

At full contraction separate MUAP will be indistinguishable resulting in a complete recruitment = interference pattern

## MOTOR UNIT POTENTIAL DURING MILD EFFORT



## MOTOR UNIT POTENTIAL DURING MODERATE EFFORT



### MOTOR UNIT POTENTIAL AT FULL VOLUNTARY EFFORT



## **Analysis**

The EMG is used to investigate both neuropathic and myopathic disorders (weakness, numbness, pain )

 The size, duration, frequency of the electrical signals generated by Msl cells help determine if there is damage to the Msl or to the nerve leading to that Msl.  Myopathy: progressive degeneration of skleletal muscle fibers

Eg: Duchenne Muscular dystrophy

 Neuropathy: Damage to the distal part of the nerve.
 peripheral neuropathy mainly affects feet & legs

#### Most common etiologies:

- Guillain Barré syndrome
- Diabetes mellitus
- Alcohol abuse

LMN lesions: interrupt the spinal reflex arc (a motor N) → Partial or complete loss of voluntary contraction, muscle wasting, treflexes, fasciculation

**Example:** Polyomyelitis

# In neurogenic lesion or in active myositis, the following spontaneous activity is noted:

- Positive sharp waves
- Fibrillations
- Giant motor unit potentials

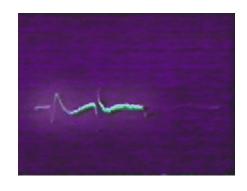
## Fibrillation potentials:

Low amplitude, short duration potentials, correspond to the spontaneous discharge of a denervated single muscle fiber due to denervato hypersensitivity to acetylcholine.

Fine invisible, irregular contractions of individual muscle fibers.

### Positive sharp waves

Small fibrillation APs (50 to 100  $\mu$ V, 5 to 10 msec duration) whose propagation is blocked at the level of the recording Ede



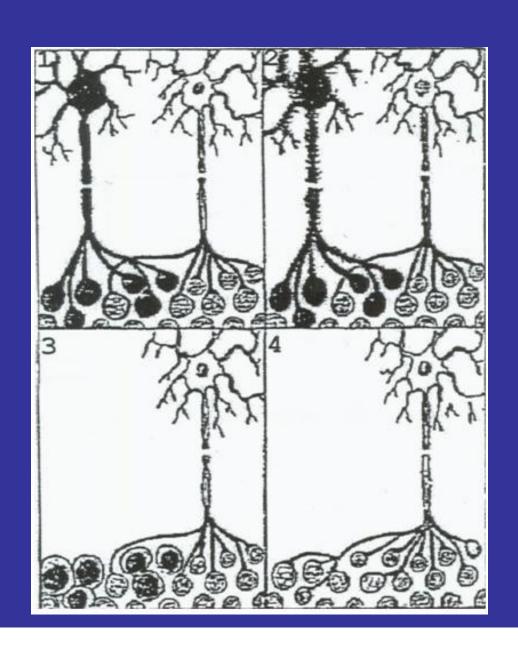


## Fasciculation potentials

spontaneous discharge of a MU at rest, can be seen and felt by the patients

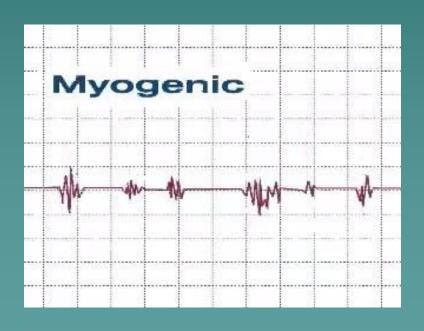
• Partial re-innervation of denervated muscle, by sprouting of the remaining nerve terminals, produces abnormally large, long polyphasic potentials (giant potential)

#### REINNERVATION BY COLLATERAL SROUTING



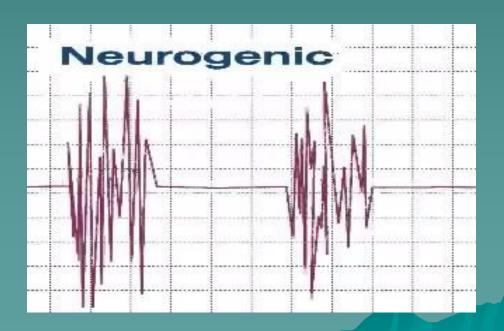
### Myopathic alteration of the EMG:

## Polyphasia ,short duration ,reduced voltage of MUPs



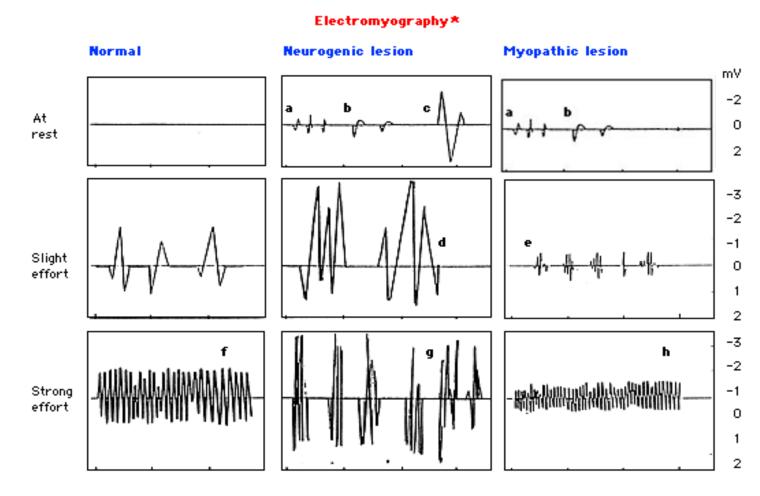
## Neuropathic alteration of the EMG:

◆ Polyphasia ,long duration ,high voltage of MUPs



## **Analysis of MUP**

MUP	NORMAL	NEUROGENIC	MYOPATHIC
Duration msec.	3 – 16 msec	> 16 msec	< 3 msec
Amplitude	300 – 5000 μV	> 5 mV	< 300 μV
Phases	Biphasic / triphasic	Polyphasic	May be polyphasic
Resting Activity	Absent	Present	Present
Interference pattern	full	partial	full



- 1. At rest (spontaneous activity): a. fibrilations, b. positive sharp waves, c. fasiculation.
- 2. Slight effort (motor unit potentials): d. giant polyphasic, e. BSAPS (brief-small-abundant polyphasic).
- 3. Strong effort (interference pattern); f. full, g. reduced units, h. reduced amplitude.
- \* (helpful in selecting denervated muscles [in radiculopathies (myotomal), mononeuropathies (distal to lesion), generalized neuropathies (distal muscles)] and myopathies)

### **Nerve Conduction studies**

A nerve conduction study (NCS) is a test commonly used to evaluate the function, especially the ability of electrical conduction, of the motor and sensory nerves of the human body.

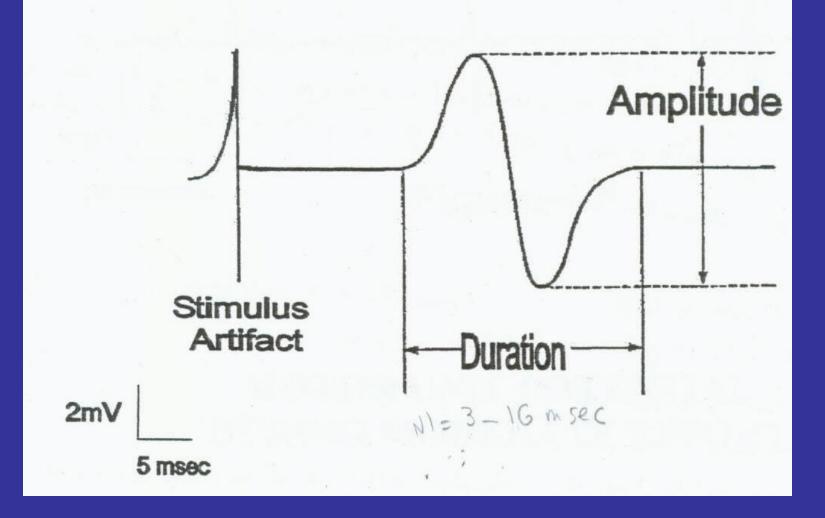
## Motor Nerve Conduction Study

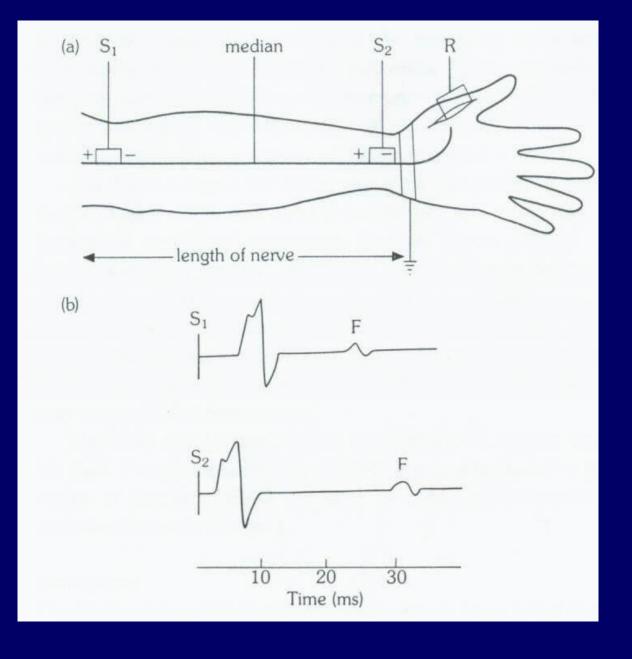
 Stimulato of median nerve at two points until visible muscle contracto is seen and a reproducible Compound Muscle A P is recorded



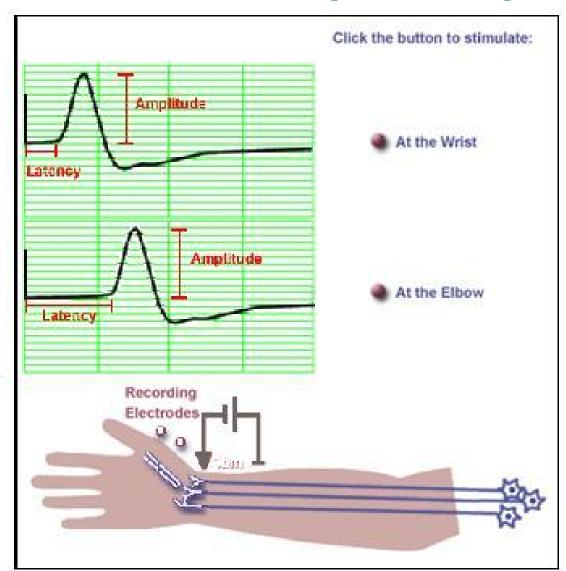
CMAP: summated potentials from all Motor Units in a muscle

#### COMPONENTS OF THE CMAP





## MOTOR NERVE CONDUCTION VELOCITY (MNCV)



$$\begin{array}{c} + \text{MNCV} = & \frac{\text{distance}}{l_1 - l_2} \end{array}$$
 (m/sec)

 $I_1$  = latency at elbow.  $I_2$  = latency at wrist

Distance between the two stimulating electrodes

abNI if < 40 m/sec</pre>

## Normal values for conduction velocity

- In arm
  - 50 to 70 m / sec.
- In leg
  - 40 to 60 m / sec.

Conduction is faster in myelinated fibres.

♣ Diseases which produce demyelinated peripheral nerves (diabetes, Gillain Barré)slow the conducto greatly(20-30 m/s).

