

SYLLABUS



Marrakesh, Morocco, November 12-17, 2011

XXth WORLD CONGRESS OF NEUROLOGY



World Congress of Neurology



**SOCiete MAROCAINE
DE NEUROLOGIE**

ELECTROMYOGRAPHY (EMG) HANDS ON

Chairperson: **Jun Kimura, USA**

09:00 **MOTOR NERVE CONDUCTION STUDIES**
Jun Kimura, USA

10:30 *Coffee Break*

11:00 **SENSORY NERVE CONDUCTION STUDIES**
Nazha Birouk, Morocco

12:30 *Lunch Break*

14:30 **REPETITIVE NERVE STIMULATION**
William J. Litchy, USA

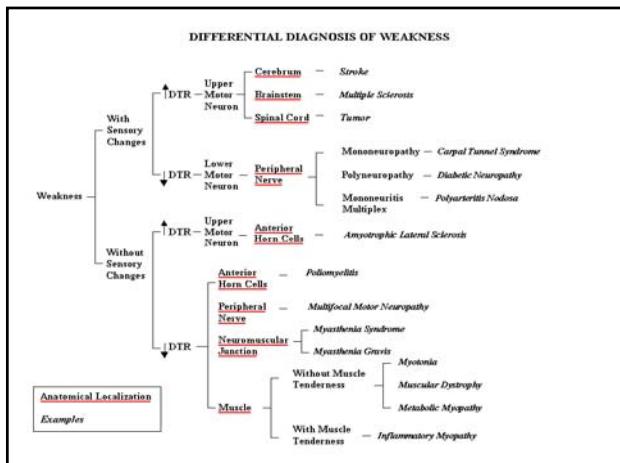
16:00 *Coffee Break*

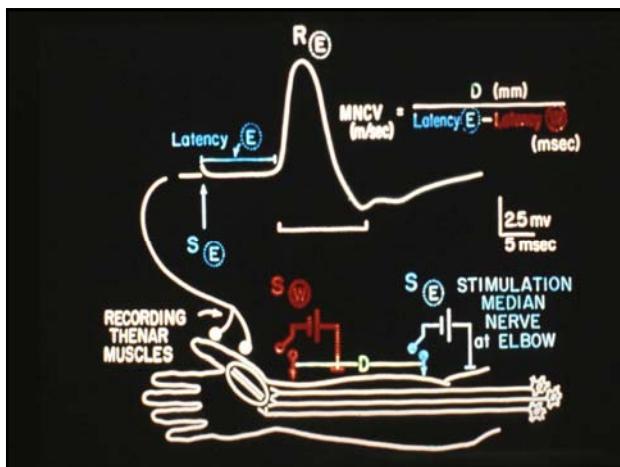
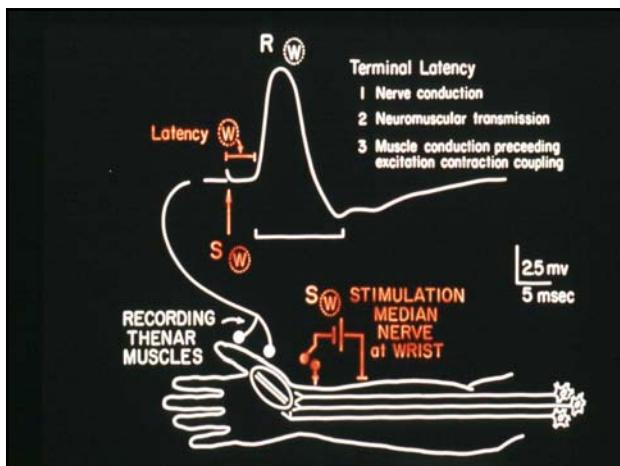
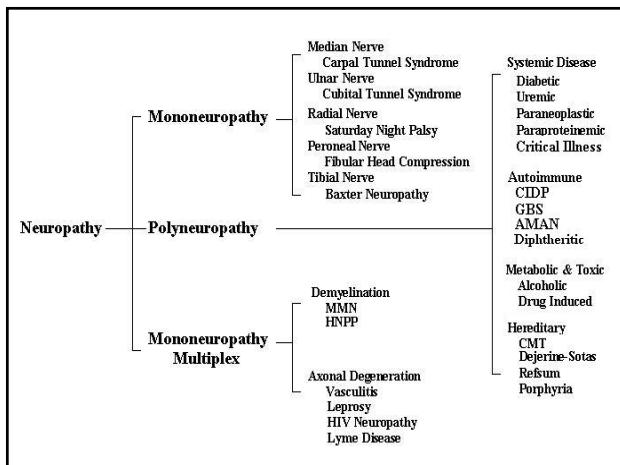
16:30 **REFLEXES AND LATE RESPONSES**
Josep Valls-Sole, Spain



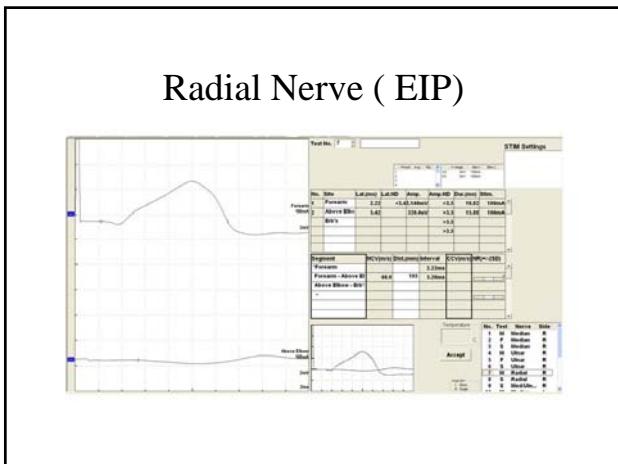
Principles and Practice

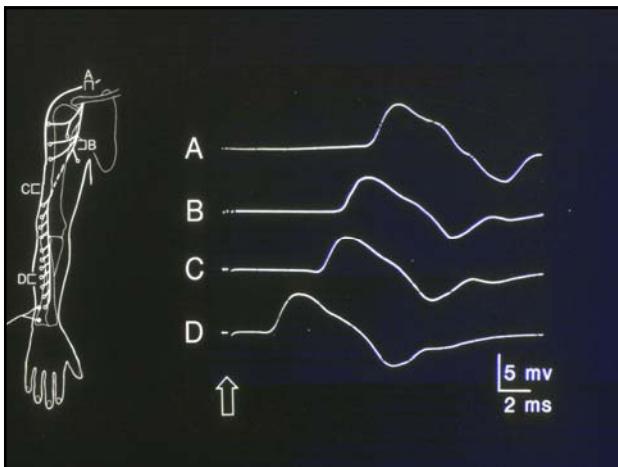
- 1) Hx & Px before EDx
- 2) Watch twitch, then measure
- 3) Distal vs Proximal Shocks
- 4) Linear or Nonlinear Changes
- 5) Short and Long of NCS

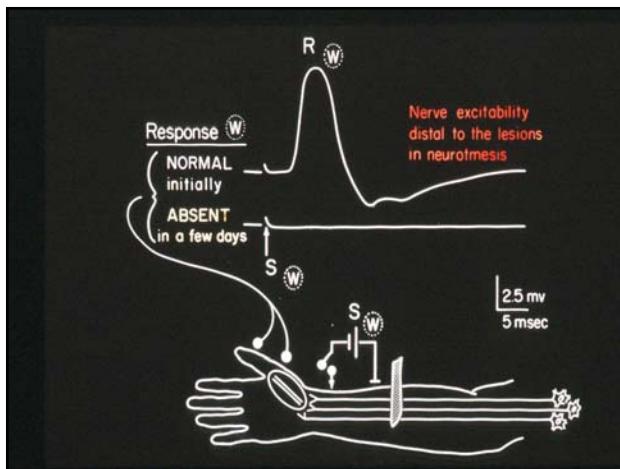


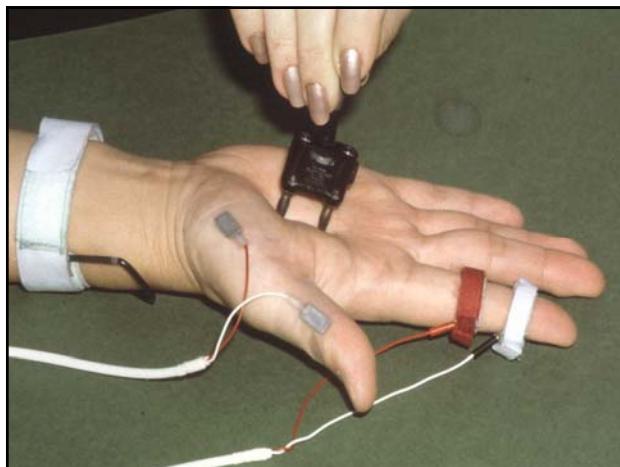


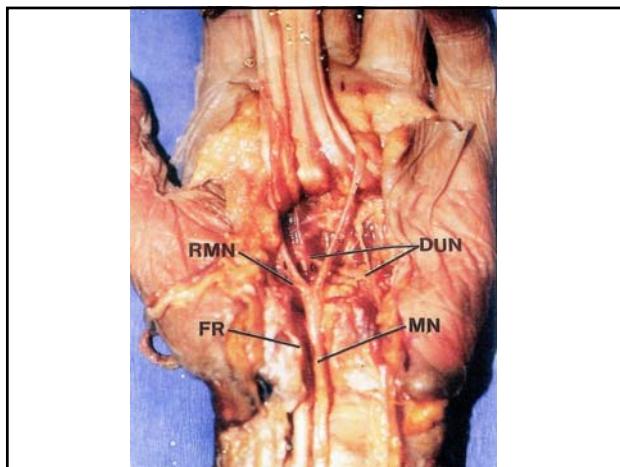






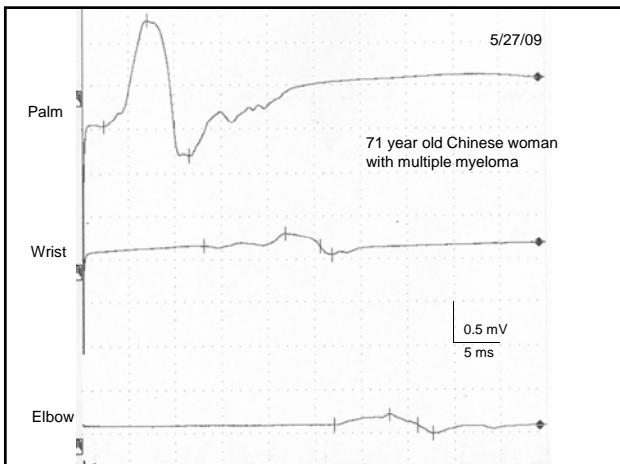


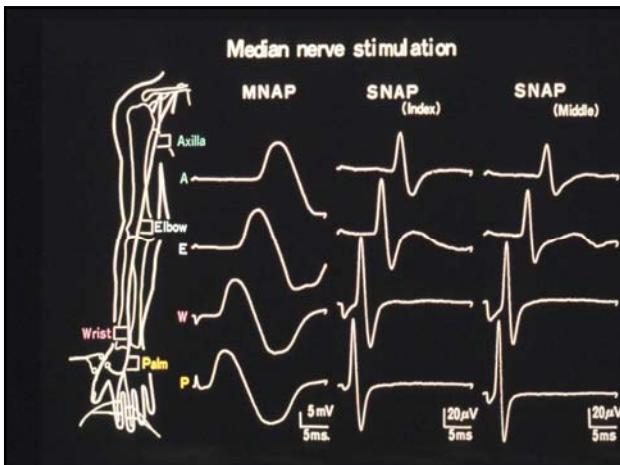


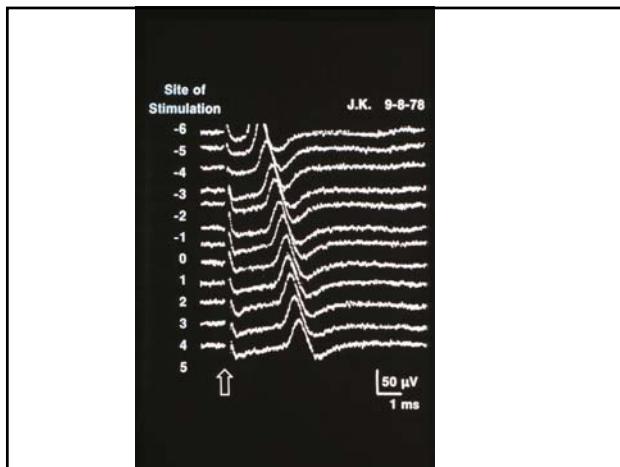
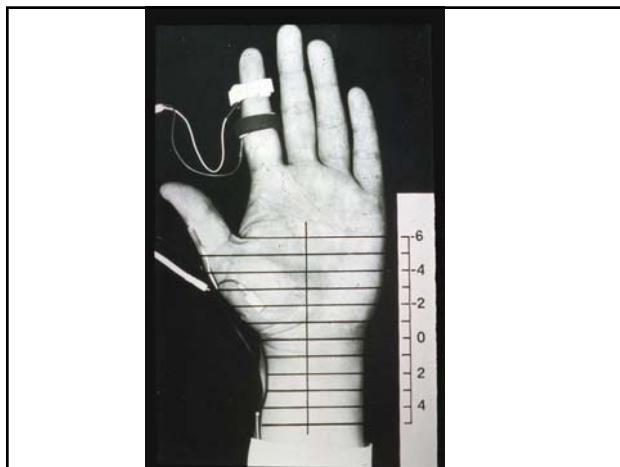
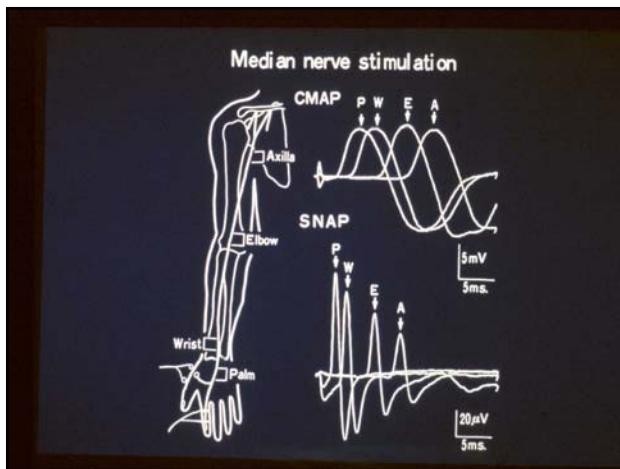


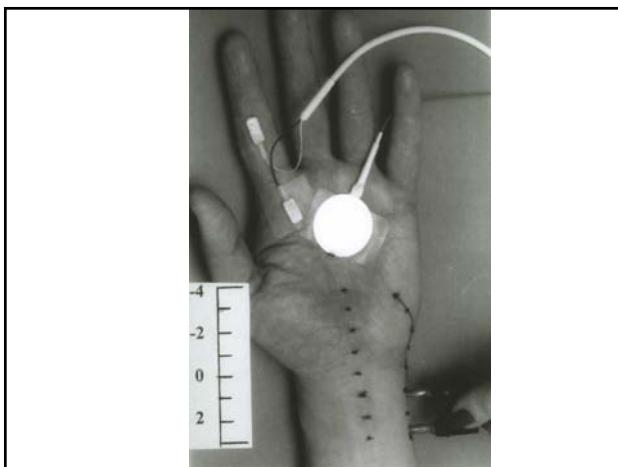
Hx:

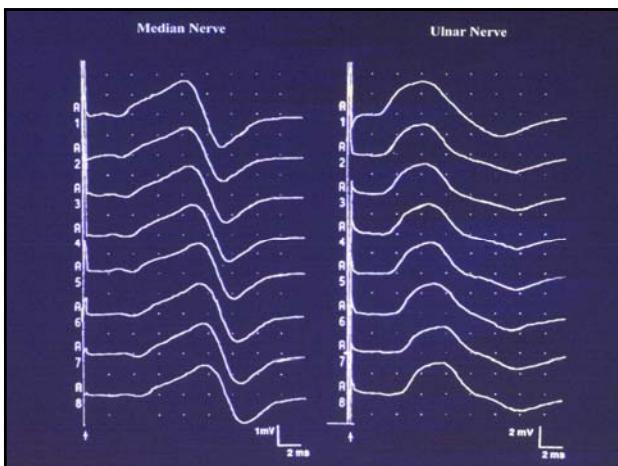
72 year-old Chinese woman
Progressive numbness & weakness involving
Digits 1-3 on left (03/09)
Digits 1-3 on right (07/08)
Feet and toes, bilateral (01/09)



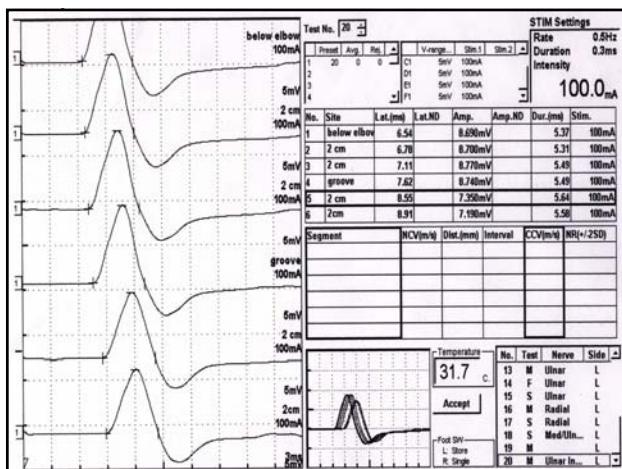
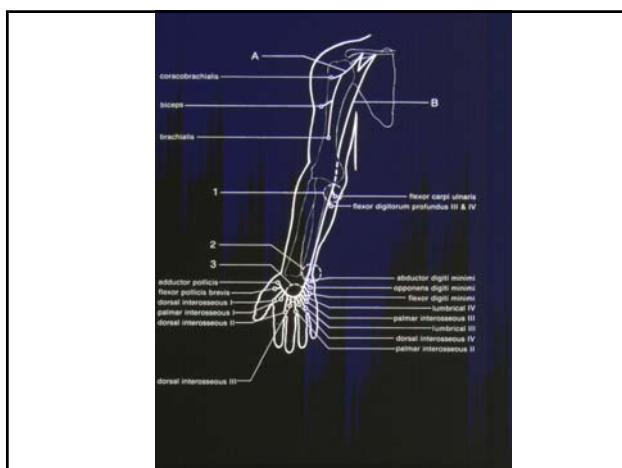
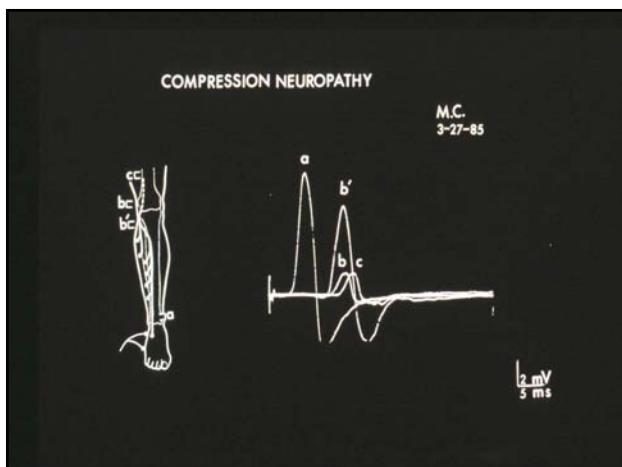


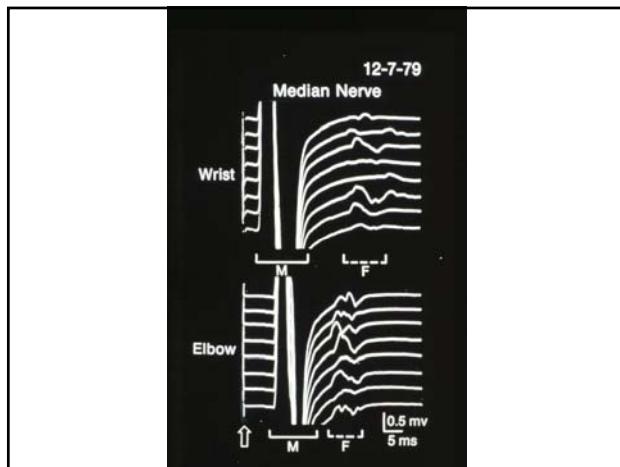
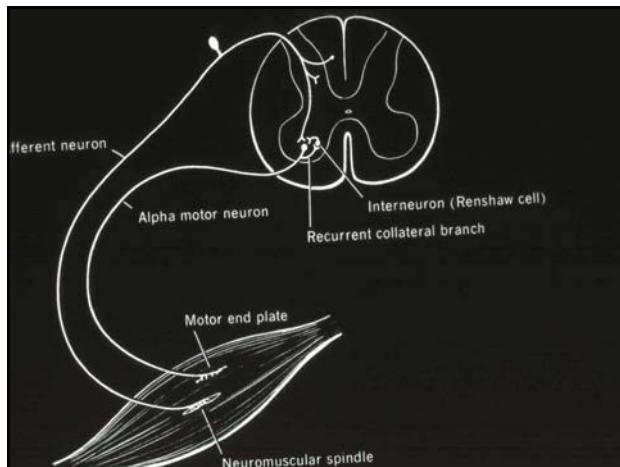
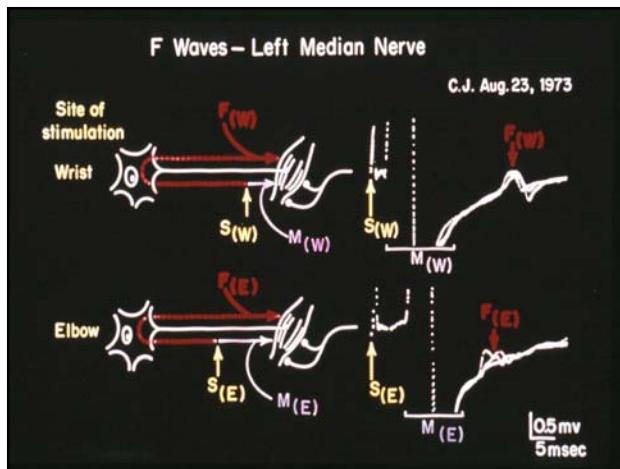


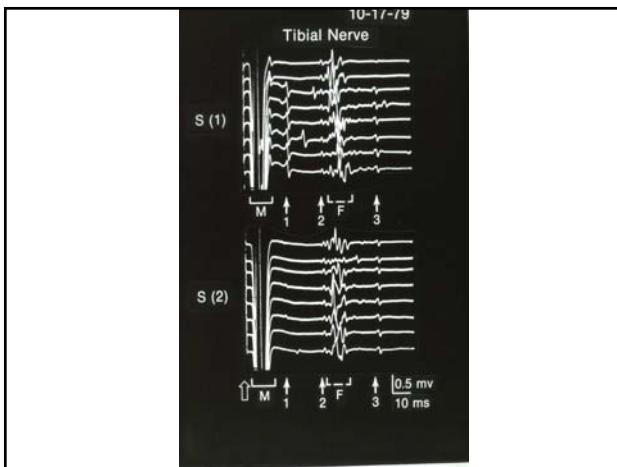


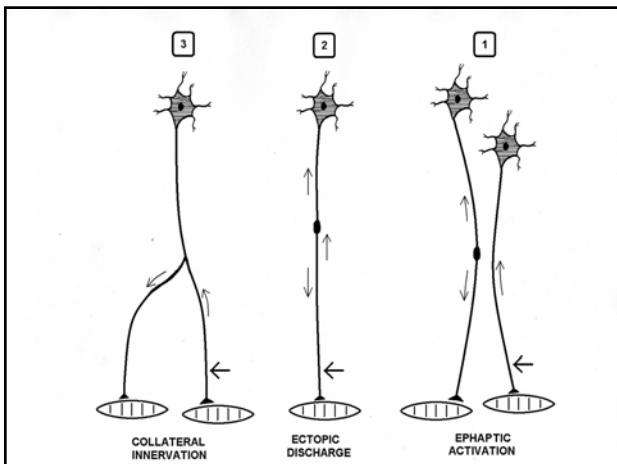






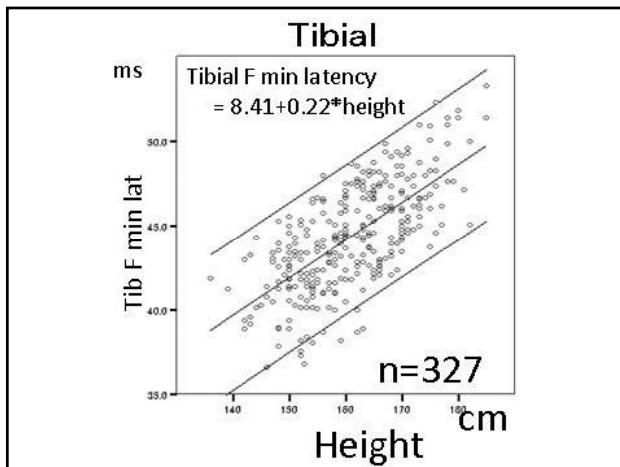
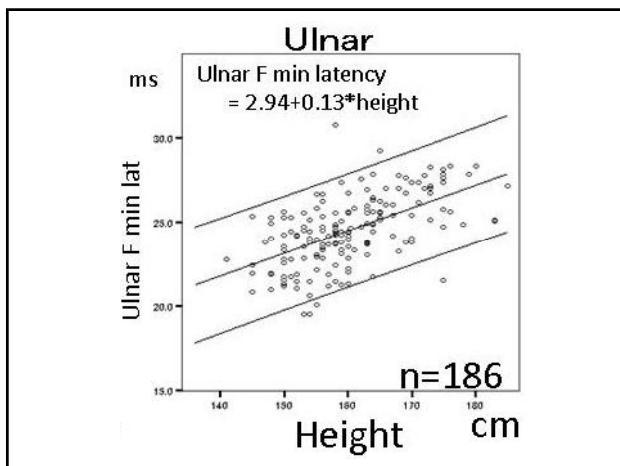
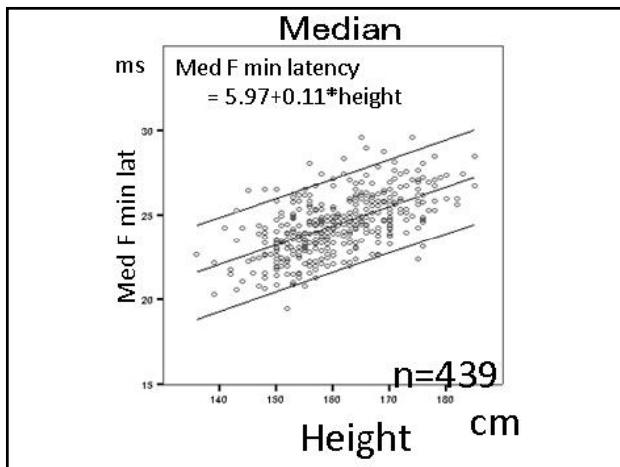


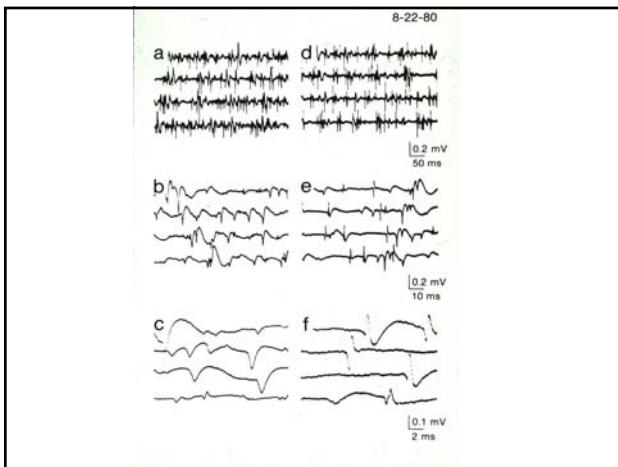
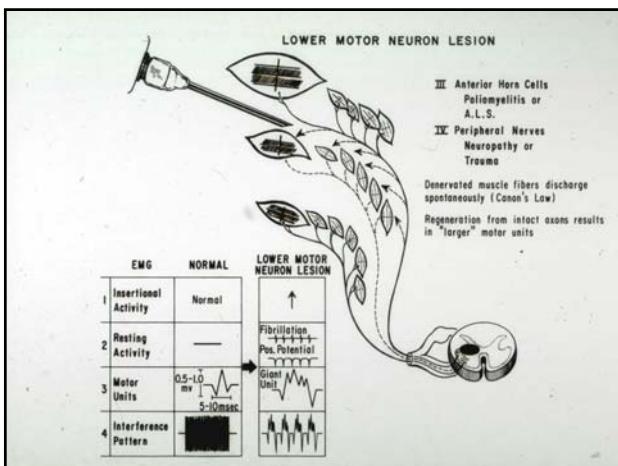
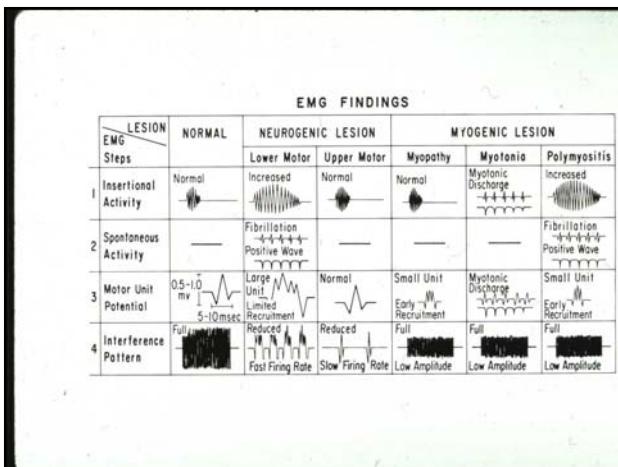


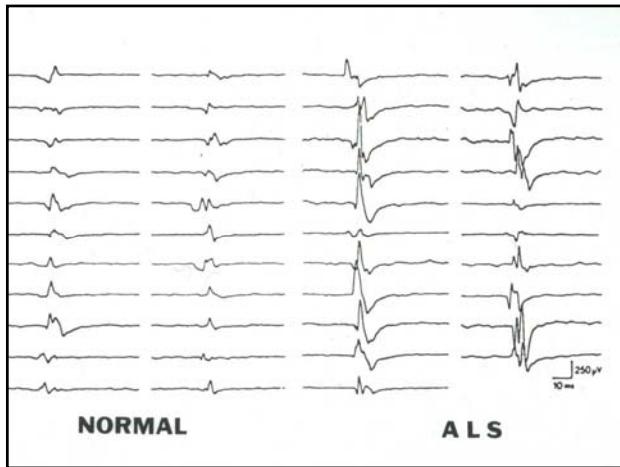
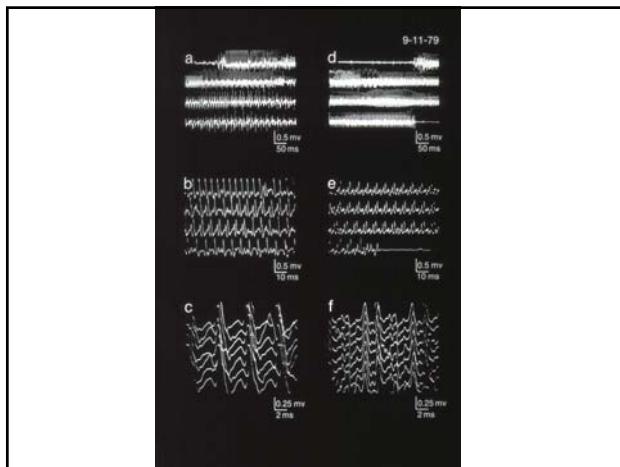
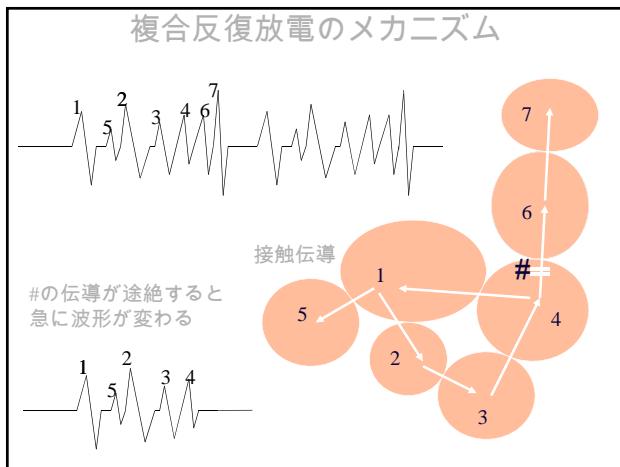


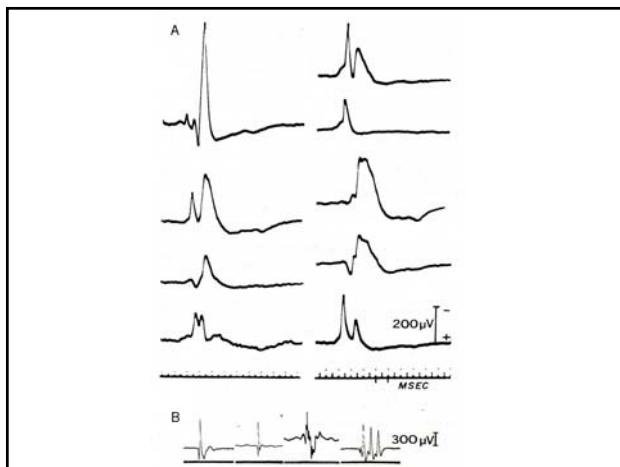
NCS: Long and Short of It

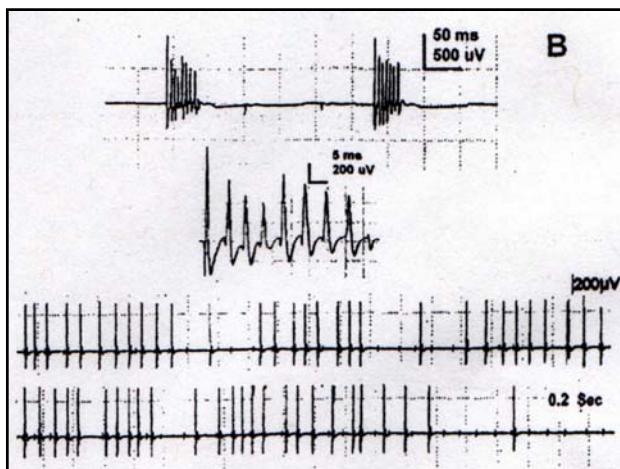
- 1) Short distances magnify focal abnormality despite increased measurement error.
- 2) Long distances, though insensitive to focal lesions, yield better for diffuse process

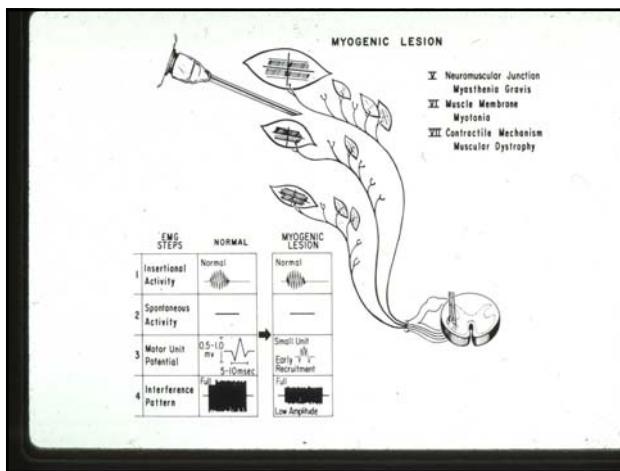


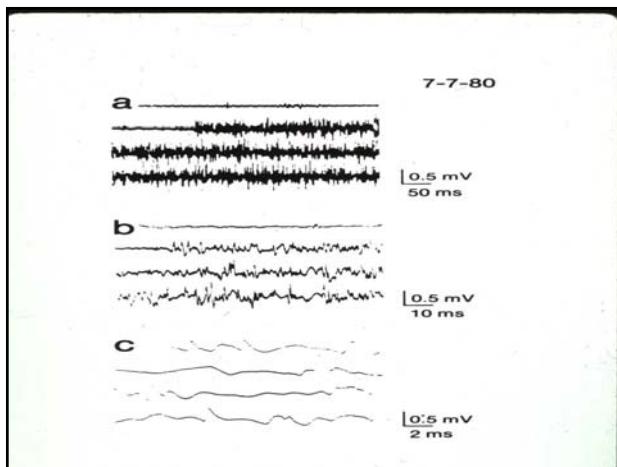












1) Muscle Fiber

- Insertional Positive Waves
- Myotonic Discharge
- Fibrillation Potential
- Positive Sharp Waves
- Complex Repetitive Discharge
- End-plate Noise
- End-plate Spikes

2) Lower Motor Neuron

- Fasciculation Potential
- Myokymic Discharge
- Neuromyotonic Discharge
- Cramp Discharges
- Hemifacial Spasm
- Hemimasticatory Spasm

3) Upper Motor Neuron

- Stiffman Syndrome
- Involuntary Movement

Sensory Nerve Conduction

Prof. Nazha Birouk

Service de Neurophysiologie Clinique
Hôpital des Spécialités, CHU Ibn Sina
Rabat, Morocco

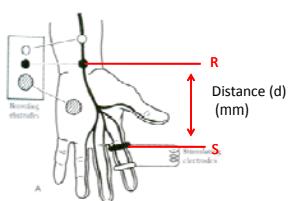
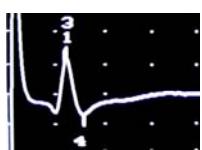


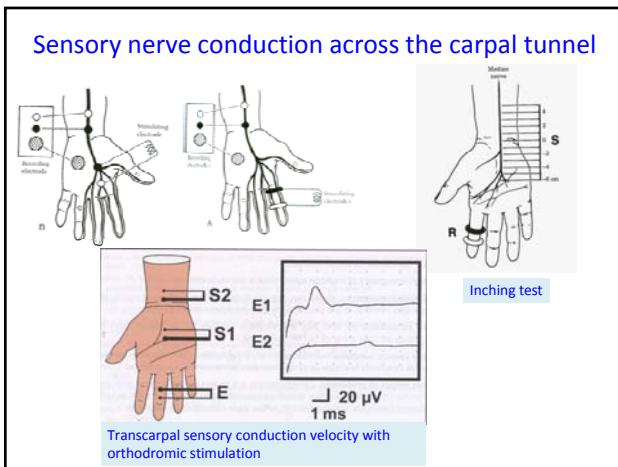
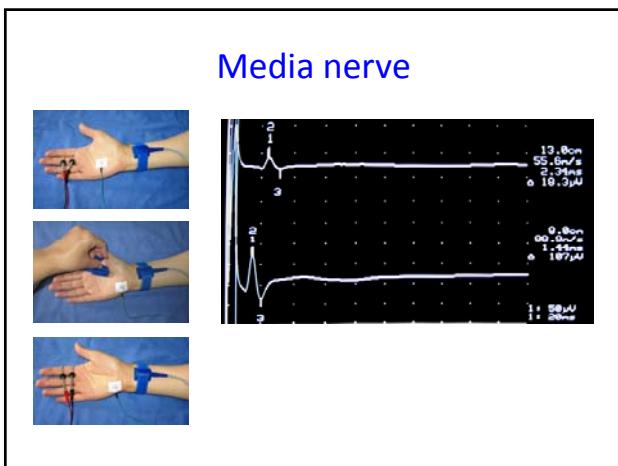
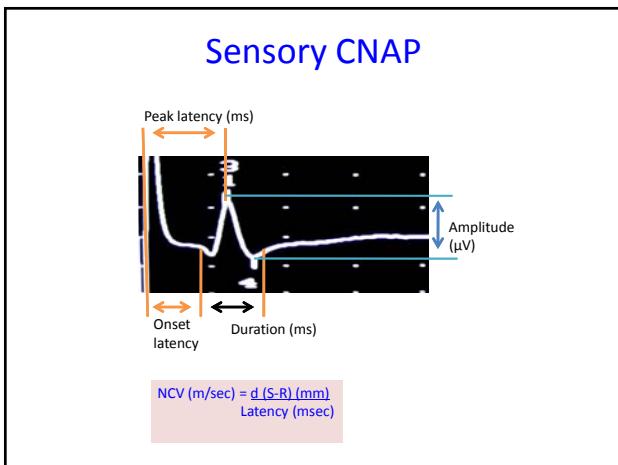
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General considerations

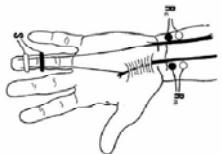
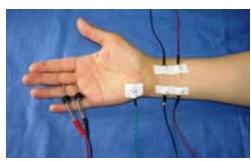
- Large diameter sensory nerve fibers examination
- Stimulation and response on the nerve
- 2 methods :
 - Orthodromic : response proximally to stimulation
 - Antidromic : response distally to stimulation
- Skin preparation
- Skin temperature
- Signal averager : about 20 to 40 responses

Sensory CNAP

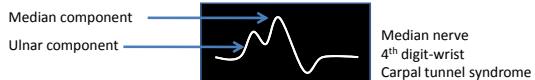




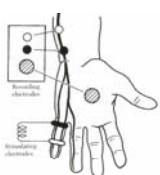
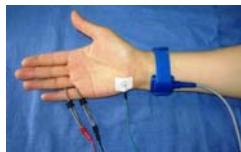
4th digit, Median-Ulnar nerve



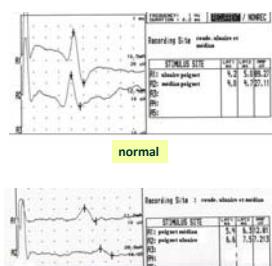
Comparison of median/ulnar latencies with 4th digit ring stimulation and recording at the wrist N ≤ 0.5 msec



Ulnar Nerve



Mixed potentials



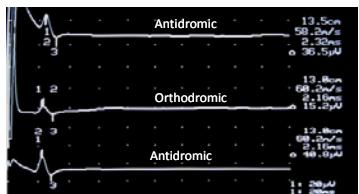
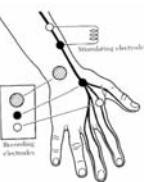
Comparison of median/ulnar latencies above the elbow:
Stimulation at the wrist. Same distances
Usual difference on normal subject ≤ 0.7 msec

Latency difference of 1.3 msec
Ulnar nerve compression at the elbow

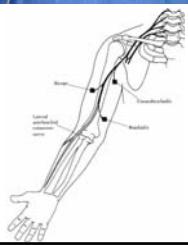
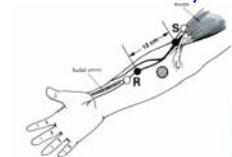
Radial Nerve



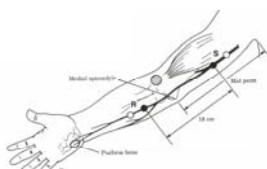
Antidromic method



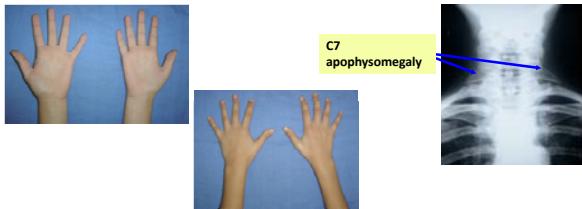
Musculocutaneous nerve (lateral antebrachial cutaneous nerve)



Medial Antebrachial Cutaneous Nerve

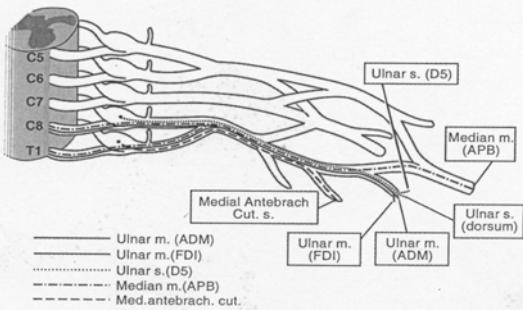


- Girl, 16 years
 - Since 1 year : Neuropathic pain in the right UL + slight motor disability in right hand

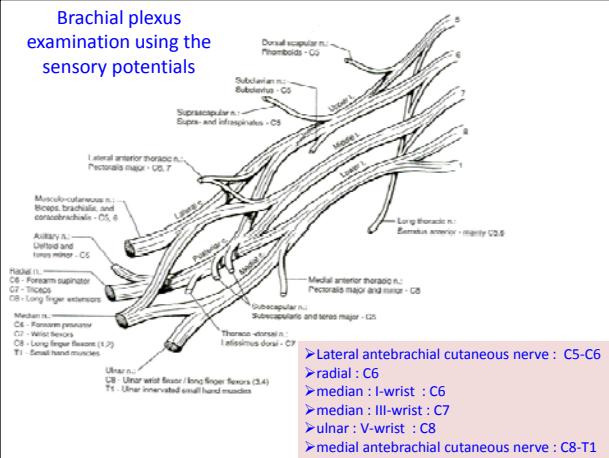


	Right median nerve (II)	Right ulnar nerve	MABCN
Motor response	Low Amplitude	Low Amplitude	-
Sensory response	Normal	Absent	Absent

Brachial plexus



Brachial plexus examination using the sensory potentials



Brachial plexus lesion examination using the sensory potentials

Upper trunk (C5-C6)

Altered potentials:

- LABCN 100% des cas
- median I 100%
- median II 20%
- median III 10%
- radial 60%

Potentials always normal:

- MABCN
- ulnar (V)

Brachial plexus lesion examination using the sensory potentials

middle trunk (C6-C7)

Altered potentials:

- median II 80%
- median III 70%
- radial 40%

Potentials always normal:

- median I
- LABCN
- MABCN
- ulnar (V)

Brachial plexus lesion examination using the sensory potentials

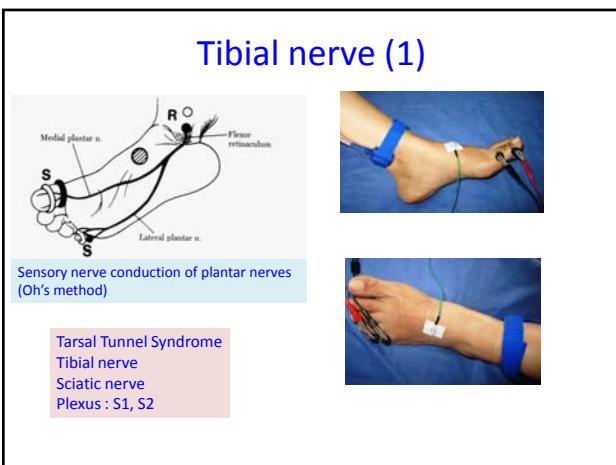
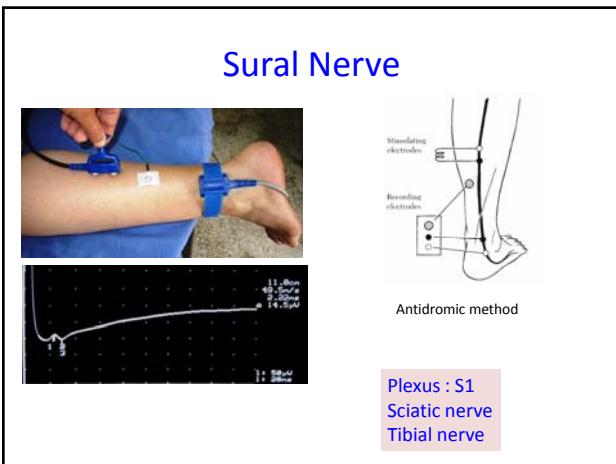
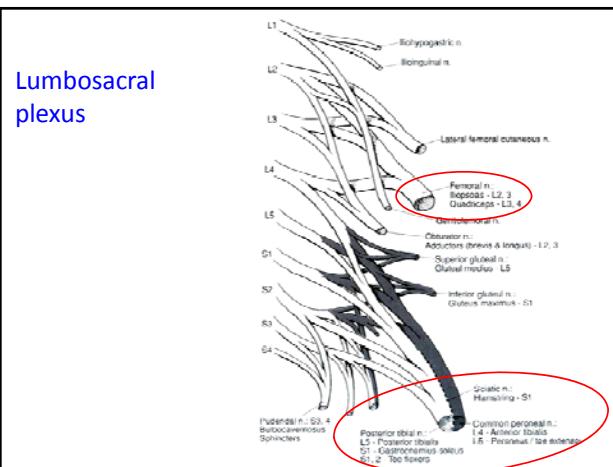
lower trunk (C7-C8-T1)

Altered potentials:

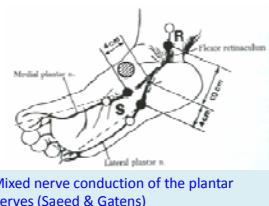
- MABCN 100%
- ulnar 100%
- median III 20%

Potentials always normal:

- median I and II
- LABCN
- radial



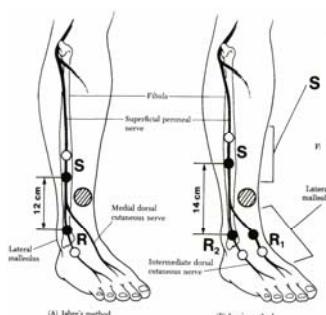
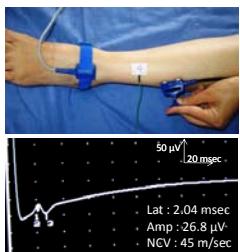
Tibial nerve (2)



Mixed nerve conduction of the plantar nerves (Saeed & Gatens)

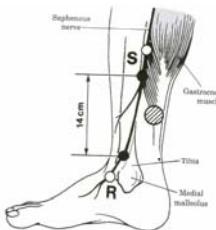
Tarsal Tunnel Syndrome
Tibial nerve
Sciatic nerve
Plexus : S1, S2

Superficial peroneal nerve



Plexus : L5
Sciatic Nerve
Common peroneal nerve

Saphenous Nerve (1)

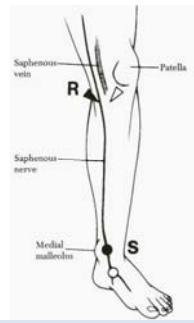


Wainapel's method (antidromic)



Femoral nerve
Plexus : L2, L3, L4

Saphenous Nerve (2)



Senden's method (orthodromic)



Femoral nerve
Plexus : L2, L3, L4

Interpretation of sensory nerve conduction data

Parameter	Axonal degeneration	Segmental demyelination
Amplitude	↓↓ or often absent	N, ↓, or absent
Duration	N	↑ or rarely dispersion
Conduction velocity	N or ↓	↓↓

Polyneuropathy (PN)

- Axonal length dependent PN : sensory responses more altered in the lower limbs : Low amplitudes or absent
- Demyelinating PN : sensory responses often more altered in the upper limbs : reduced NCV, Low amplitude or absent
- Hereditary NP of CMT type : Sensory responses often absent usually contrasting with normal clinical sensory examination (or few clinical sensory signs)
- Small fibers sensory neuropathy : Normal sensory responses.

Case 1

Hajar , 8 years, CMT1A (duplication PMP22)
Onset at 2 years, Typical CMT phenotype, moderate disability, No clinical sensory signs

Motor Nerve Conduction			
	Median (R/L)	Peroneal (R/L)	Tibial (R/L)
DML (ms)	21 / 18.4	0 / 36.3	0 / 30.6
Amplitude D (mV)	0.1 / 0.2	0 / 0.2	0 / 0.16
Amplitude P (mV)	0.07 / 0.2	0 / 0.2	0 / 0.10
NCV (m/s)	5.8 / 6	0 / 6	0 / 6.2

Sensory Nerve Conduction			
	Median (R)	Sural (R / L)	Ulnar (L)
Amplitude (μ V)	0.6	0 / 0	0
NCV (m/s)	19	0 / 0	0

Case 2

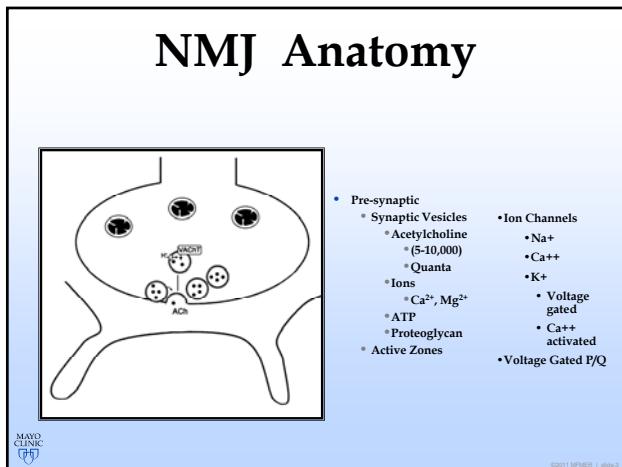
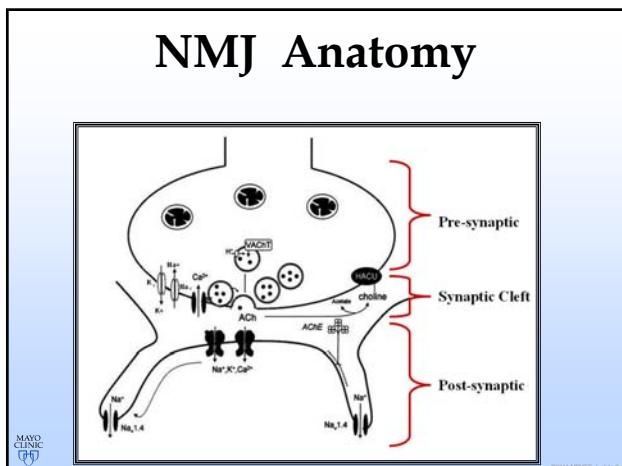
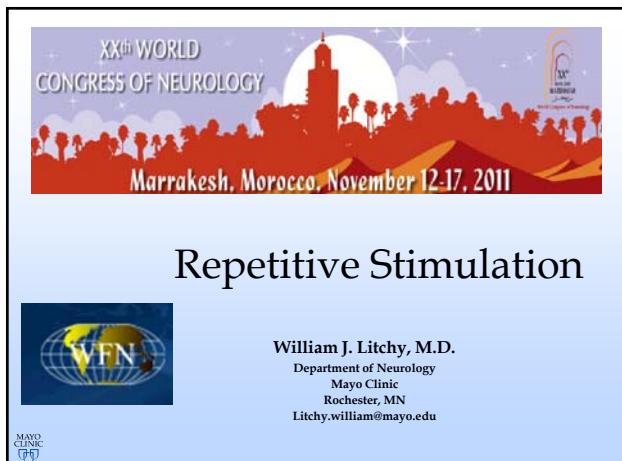
Rayan, 4 years, Recurrent CIDP since 6 months
Complete clinical recovery after steroids therapy

Motor nerve conduction			
	Median R	Peroneal (L)	Tibial (R)
DML (ms)	5.3	7.7	6.1
Amplitude D (mV)	2.8	0.5	0.5
Amplitude P (mV)	2.7 TD	0.5 TD	0.2 TD
NVC (m/s)	16.7	23	20

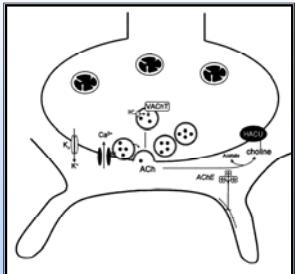
Sensory nerve conduction			
	Median R	Ulnar L	Sural R
Amplitude (μ V)	0.5	0	4
NCV (m/s)	30	0	31

Conclusion

- Important step in ENMG test
- Rigorous technique is necessary :
 - Skin preparation and temperature
 - Electrodes placement
 - Eliminate artifacts sources...
- Nerves examined routinely : Normal values for each laboratory
- Nerves examined occasionally : comparison with other side often useful



NMJ Anatomy

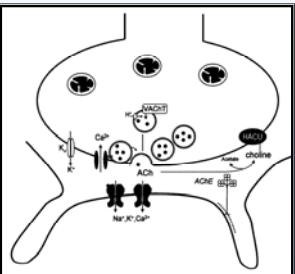


- Synaptic Cleft
 - Small space
 - Acetylcholinesterase (AChE)
 - Two functional sites
 - 5000 molecules/sec
 - Controls ACh content

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NMJ Anatomy

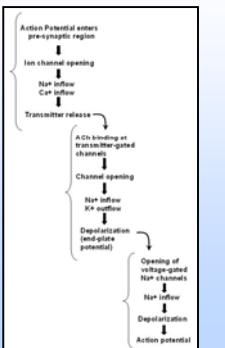
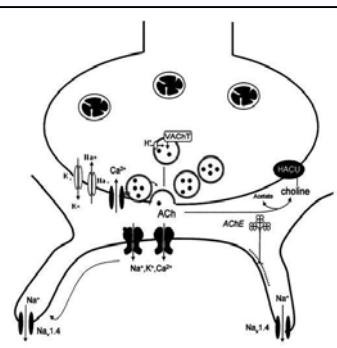


- Post-synaptic
 - Acetylcholine Receptors
 - 5 subunits
 - $\alpha_1, \beta_1, \beta_2, \gamma, \delta$
 - α subunit binds ACh
 - 2 ACh for channel opening
 - Produce non propagating potentials
 - End Plate Potential (EPP)
 - Na^+ gated ion channels
 - EPP activates opening
 - Produces action potential
 - MuSK Receptors

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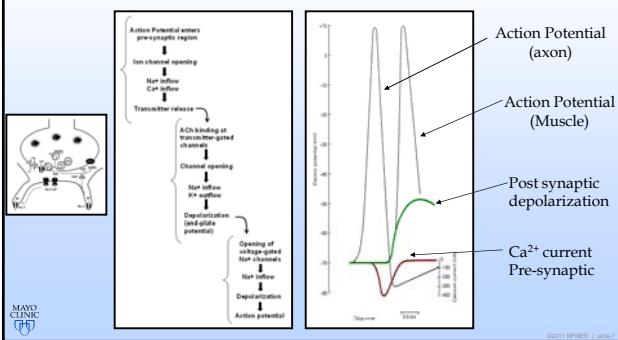
NMJ Physiology



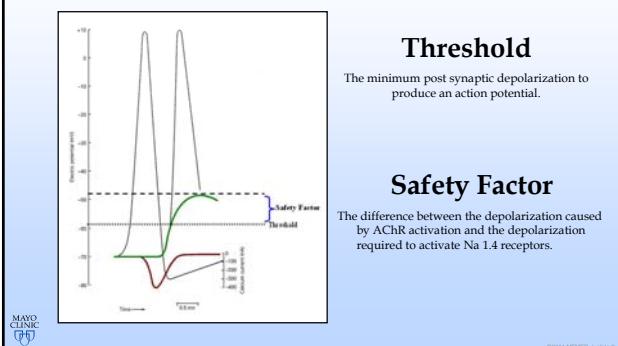
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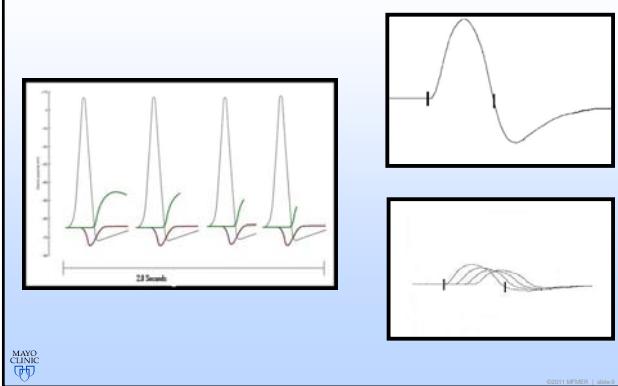
NMJ Physiology



NMJ Physiology



NMJ Physiology



Techniques for Evaluation of NMJ

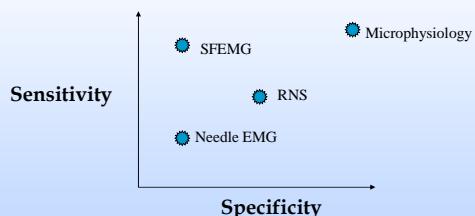
Routine Nerve Conduction Studies

- Repetitive Nerve Stimulation (RNS)
- Needle Electromyography
- Single Fiber EMG
- End-plate potential analysis



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NMJ Evaluation



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Technique for RNS

Patient Preparation

Instruction
Warm

Equipment

Recording Electrodes
Stimulating Electrodes
Instrument
Immobilizing joint

Nerve/Muscle Examined

Clinically Involved
Reliable
Reproducible
Patient Comfort
Physician Comfort



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Nerve/Muscle Choice

Nerve	Muscle	Stimulation Site	Advantages	Disadvantages	Immobilization
Ulnar	Abductor digiti quinti	Wrist	Relatively immobilized, well tolerated	Digital muscle may be spared	Velcro strap, damp
Median	Abductor pollicis brevis	Wrist	Well tolerated	Digital muscle may be spared, difficult to tolerate	Thumb restrained
Musculocutaneous	Biceps	Lower edge axilla	Proximal muscle	Unstable stimulus, difficult to immobilize	Board
Radial	Anconeus	Nerve to anconeus in the forearm	Forearm muscle, comfortable for the patient	Requires a needle for stimulation	None
Radial	Extensor digitorum communis	Elbow	Forearm muscle	Unstable stimulus, muscle artifact	Board
Axillary	Deltoid	Suprascapular	Proximal muscle	Unstable stimulus; difficult to immobilize	Large Velcro strap or sheet
Spinal accessory	Trapezius	Posterior border upper trapeziolaminotub	Proximal muscle	Difficult to immobilize	Large Velcro strap or sheet
Facial	Nasalis	Between mastoid and tragus	Proximal muscle	Unstable stimulus, shock artifact cannot immobilize	None
Peroneal	Anterior tibial	Popliteal fossa	Leg muscle	Digital muscle may be spared	Board
Femoral	Rectus femoris	Femoral triangle	Proximal leg muscle	Difficult to immobilize	Board

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Stimulation Parameters

Duration

Strength

Frequency

Number of Stimuli

Measurements

Decrement

Increment

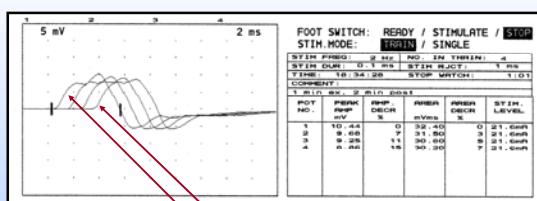
Facilitation

Post activation exhaustion

Decrement Repair



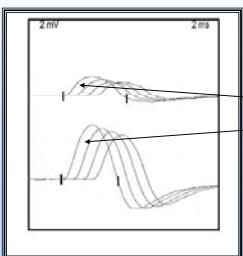
Repetitive Stimulation Decrement



$$\text{Decrement} = (1-4)/1 = \%$$



Repetitive Stimulation Facilitation



Facilitation

The change in the amplitude (area) of the first response in one set compared to the first response in a second set.

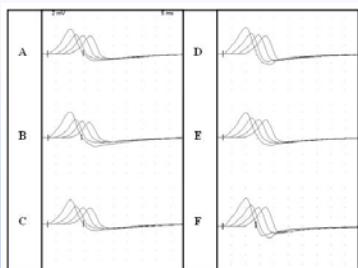


Myasthenia Gravis Protocol

- Warm patient
- Select muscle
- Routine nerve conduction study
- Repetitive stimulation--2Hz
- Exercise--short/long
- Post exercise: immediate, 30,60,120,180



Myasthenia Gravis



Criteria for Abnormality

- Amount of Decrement
 - Amplitude > 10% (1st to 4th response)
- Pattern of Abnormality
 - Largest decrement between 1st and 4th response
 - Decrement repairs with exercise
 - Evidence of post activation exhaustion
- Found with more than one nerve

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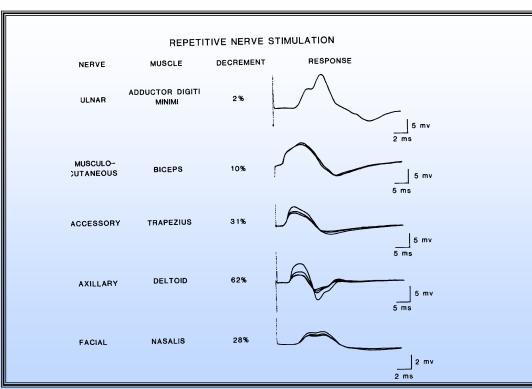
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Results for Myasthenia Gravis

- Overall
 - Class I 15 -17 %
 - Class II- IV 53 – 100%
- Specific Nerve
 - Ulnar < Radial < Accessory < Nasalis < Axillary

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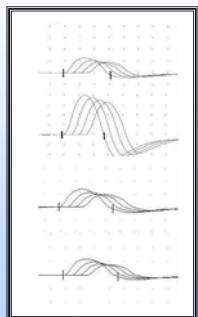
Technical Considerations

Patient
Temperature
Medication

Technical
Over/under stimulation
Choice of nerve



Lambert-Eaton Myasthenic Syndrome



Before Exercise

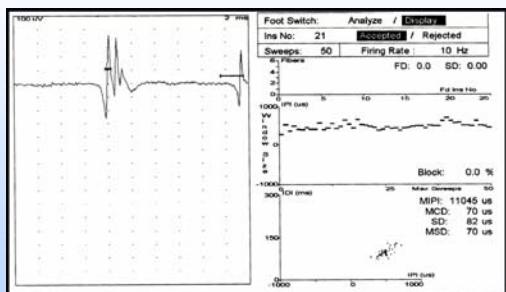
Immediately after exercise

30 seconds after exercise

1 Minute after exercise



Single Fiber EMG



Single Fiber EMG

Concentric Needle



Width Length Area

150 μ 550 μ 0.07 mm²

25 μ 25 μ 0.0005 mm²

Single Fiber Needle



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Concentric Needle Recordings

- Advantages

- Available
- Inexpensive
- Easy

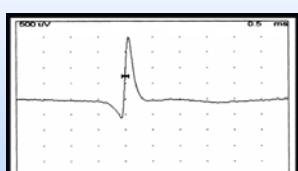
- Disadvantages

- Technically more difficult
- More difficult to record single fiber potentials



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Characteristics of Single Fiber Potentials

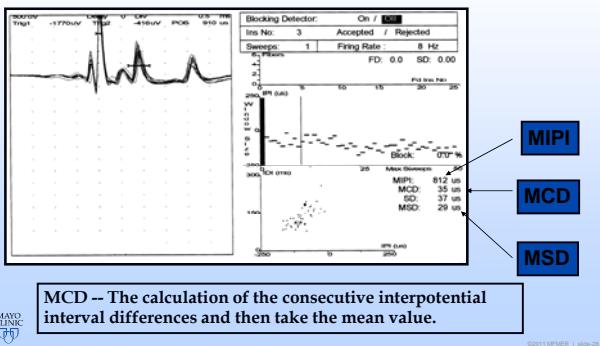


- Amplitude > 100 μ Volt
- Risetime 75-200 μ S
- Duration 1-3 msec

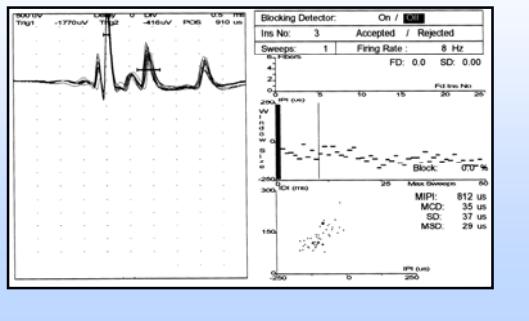


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Characteristics of Single Fiber Pairs



Jitter Measurement



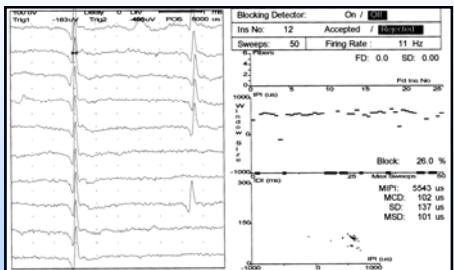
SFEMG - Blocking

A measure of the number of times a motor unit fires and a muscle fiber is not activated

Not present in normal muscles

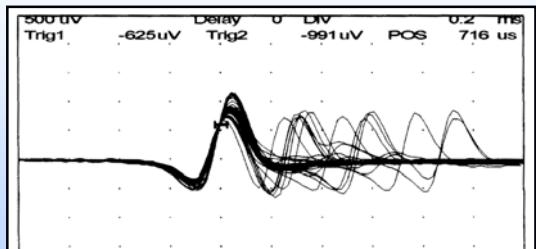
Occurs when jitter is large

SFEMG - Blocking



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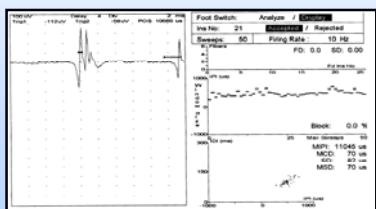
Jitter-Myasthenia Gravis



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SFEMG - Fiber Density

- A measure of the number of muscle fiber potentials recorded on a single recording area.
- Normally between 1.3 and 1.8.
- Increases with age
- Increases in disorders with de/re-innervation



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SFEMG -- Stimulated

- **Technique**

- Nerve branch stimulated
- Record single fiber potential from muscle
- Stimulate at different frequencies
- Record Jitter (MCD)

- **Advantages**

- Any muscle
- Does not require cooperation of patient
- Can be performed with patient (child) sedated
- Can be performed quickly

- **Disadvantages**

- Technically very demanding
- Requires additional skills
- Technical errors
 - Record incorrect potentials
 - Stimulate at incorrect frequencies
 - Requires different reference values



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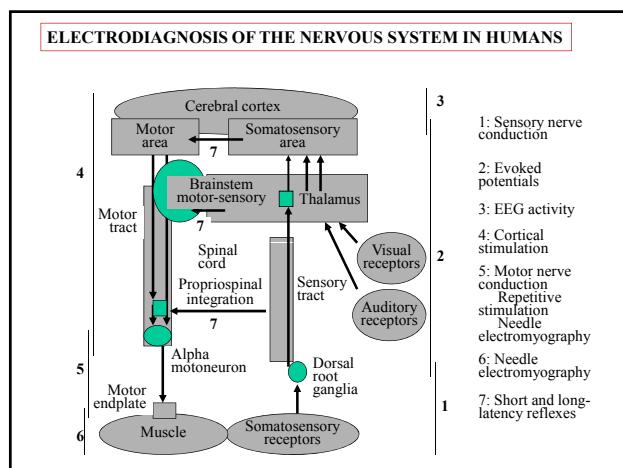
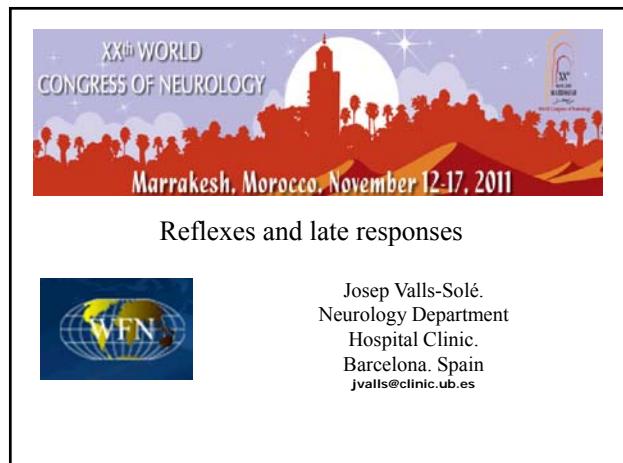
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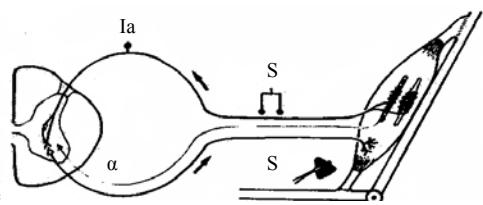
REFLEXES INVOLVING MUSCLE RESPONSES

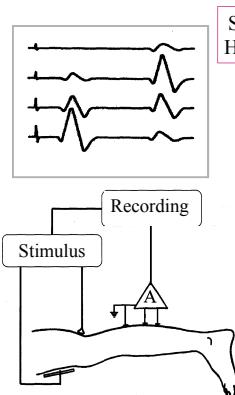
	Stimuli	Resting condition	Sustained muscle contraction
Monosynaptic	Mechanical	Tendon jerk (T wave)	
	Electrical -la afferents	H reflex	-
Polysynaptic	Electrical -Cutaneous	Blink reflex (R1, R2, R3)	Cutaneo-muscular reflex (E) Voluntary potential (V) Masseteric inhibitory reflex (MIR)
	-Cutaneous -Mixed	C reflex (C wave) Nociceptive withdrawal (RIII) Palmomental	Long latency reflexes (LLR)
	Vibration	Tonic vibration reflex (TVR)	
	Muscle stretch		Stretch reflex (M2, M3)
	Muscle shortening	Shortening reaction (SR)	

SHORT LATENCY RESPONSES TO SEGMENTAL AFFERENTS

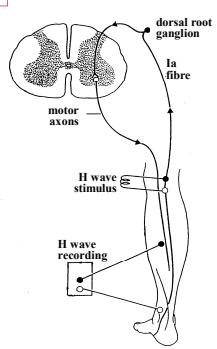
MONOSYNAPTIC REFLEX RESPONSES AND THEIR CIRCUITS AND INTEGRATION SITES

H wave	Ia afferents Alpha motoneurons
T wave	Gamma motoneurons Muscle spindles and Ia afferents Alpha motoneurons

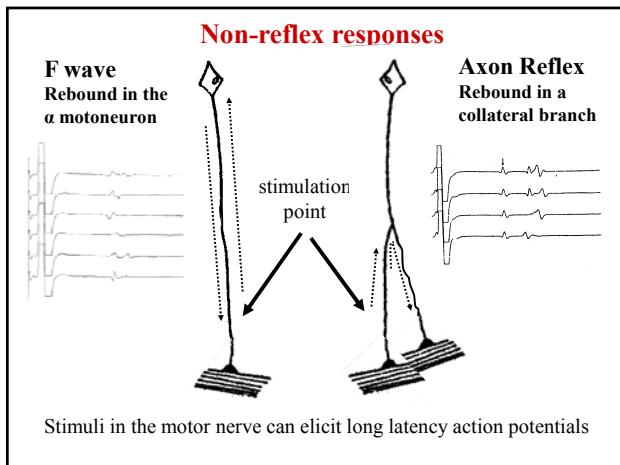
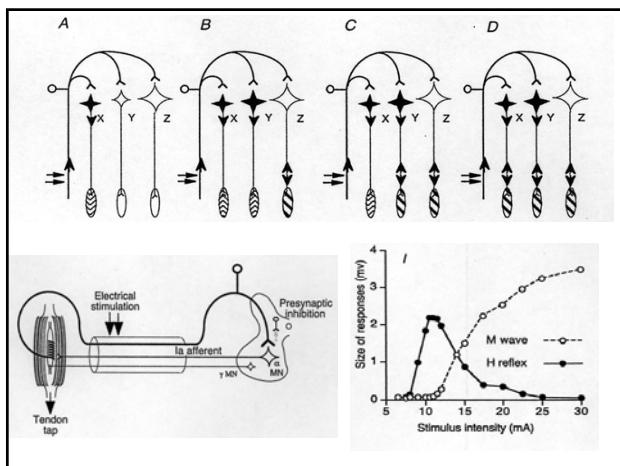
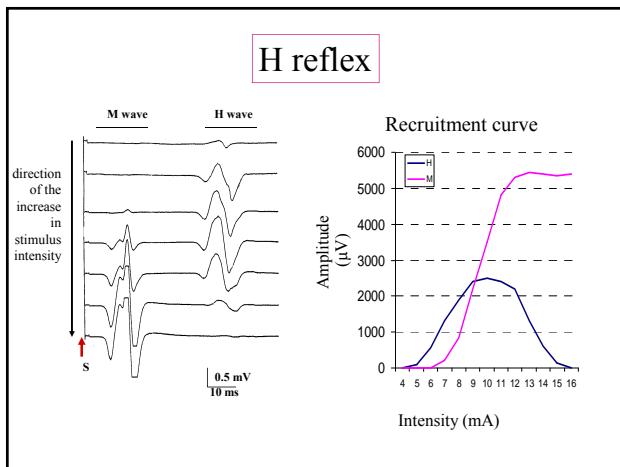


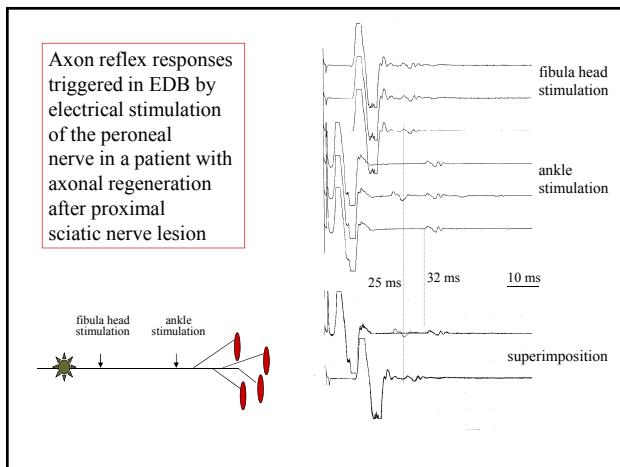


Soleus H wave



Techniques for recording the soleus H reflex





The utility of the H reflex and F waves in clinical practice

Peripheral nervous system

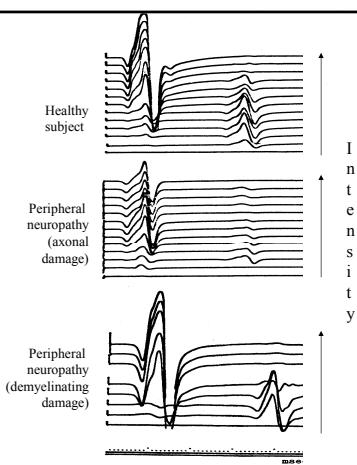
Polineuropathies
Focal lesions
Radiculopathies

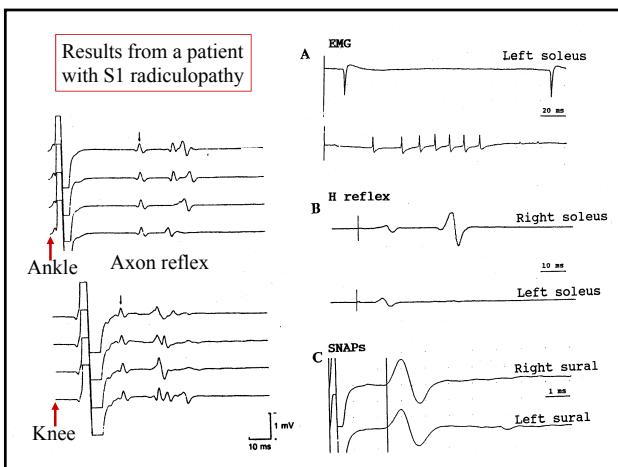
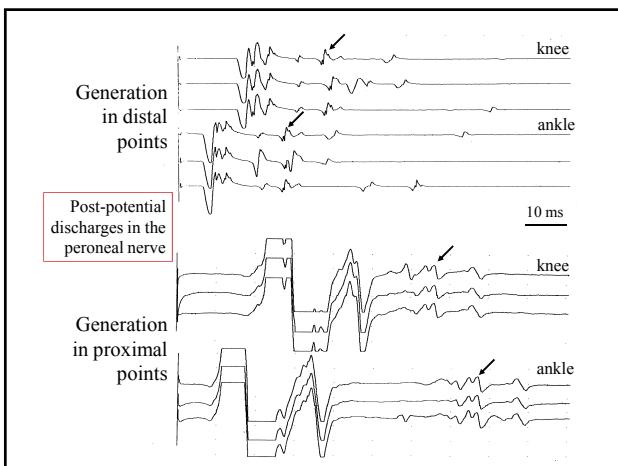
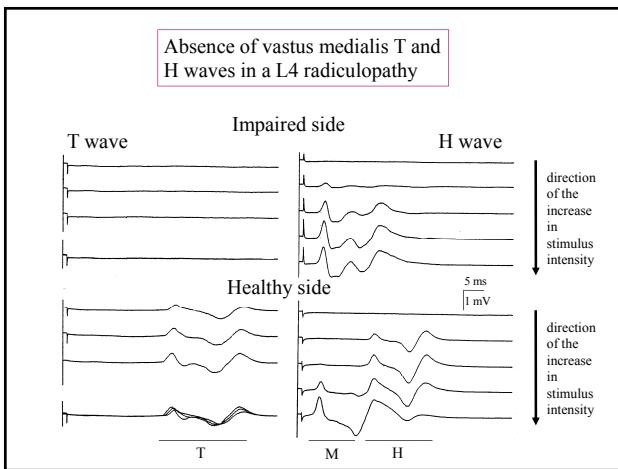
Central nervous system

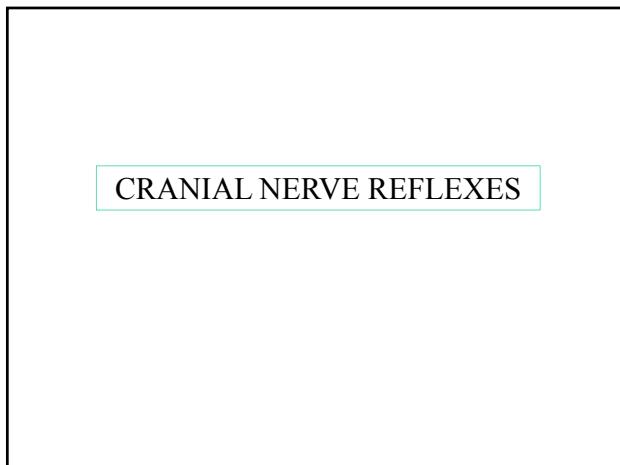
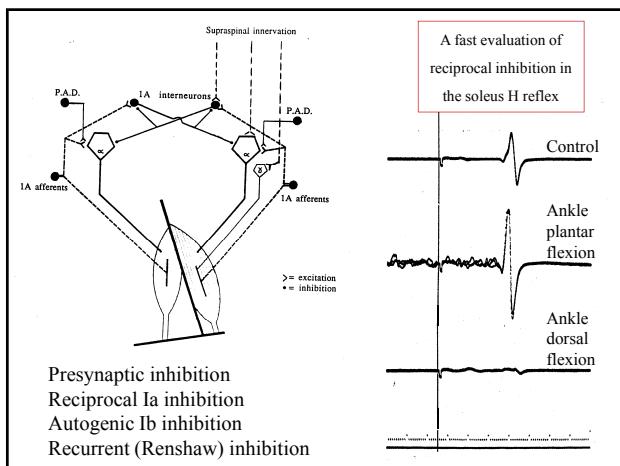
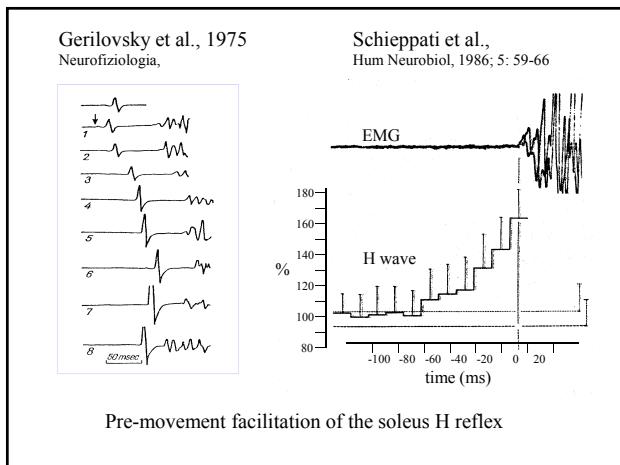
Spasticity
Dystonia
Rigidity and other disorders of propriospinal interneurons

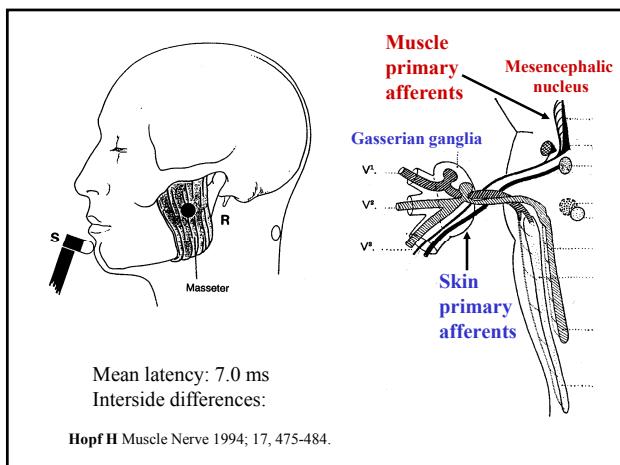
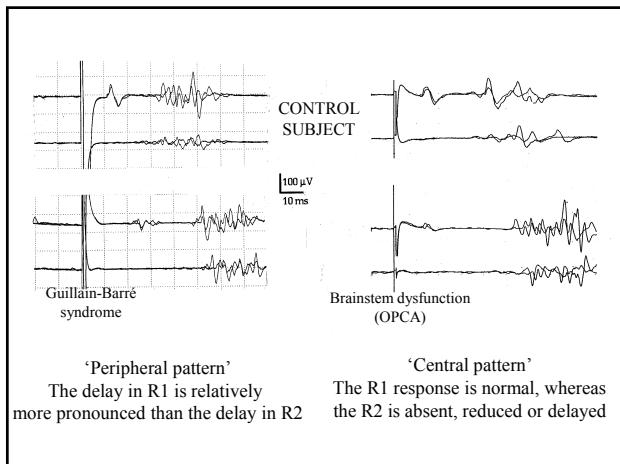
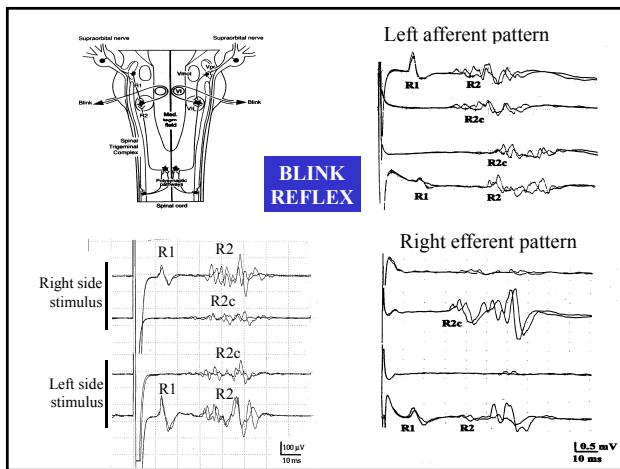
The H reflex in polyneuropathies

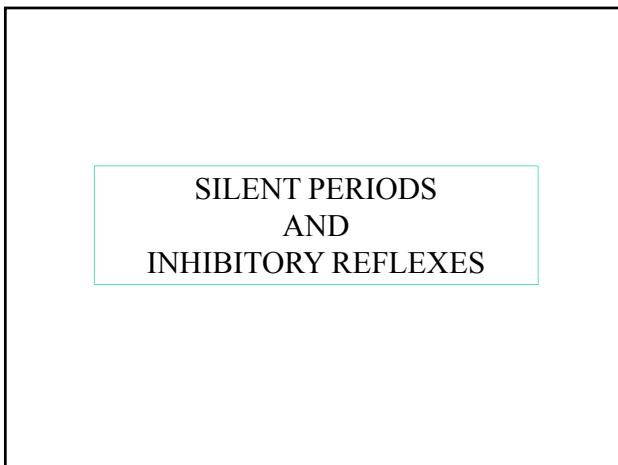
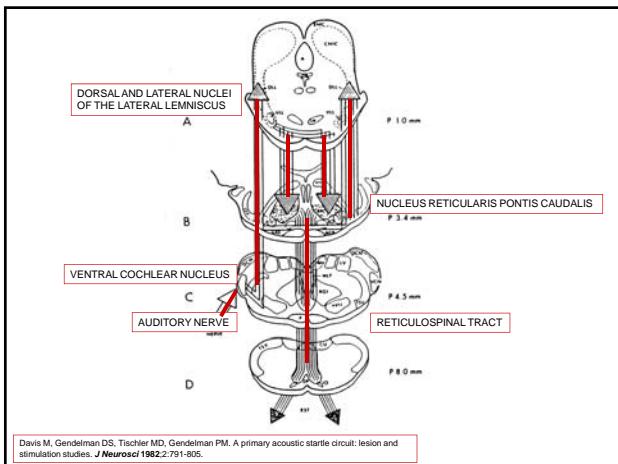
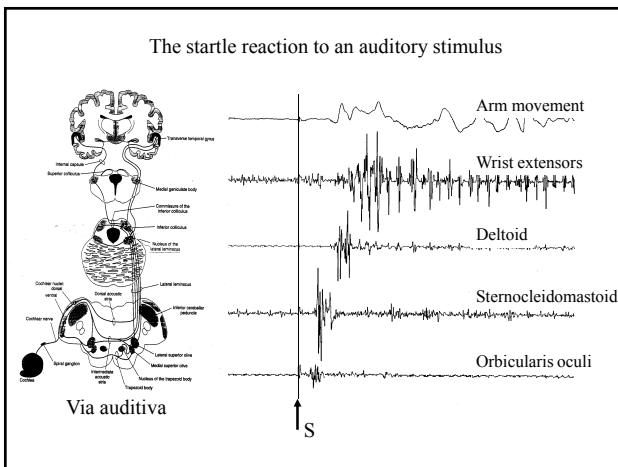
The soleus H reflex in clinical practice

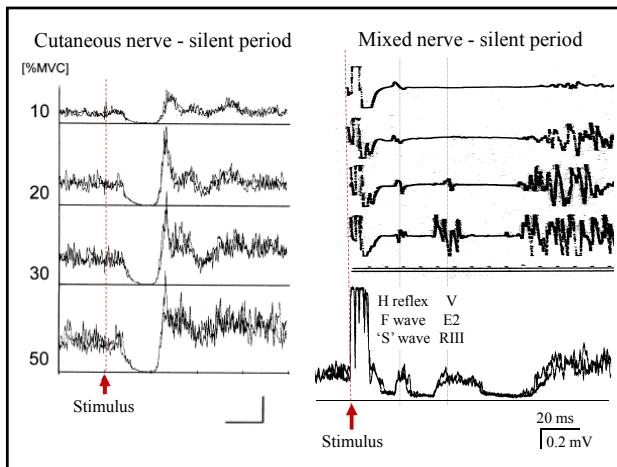
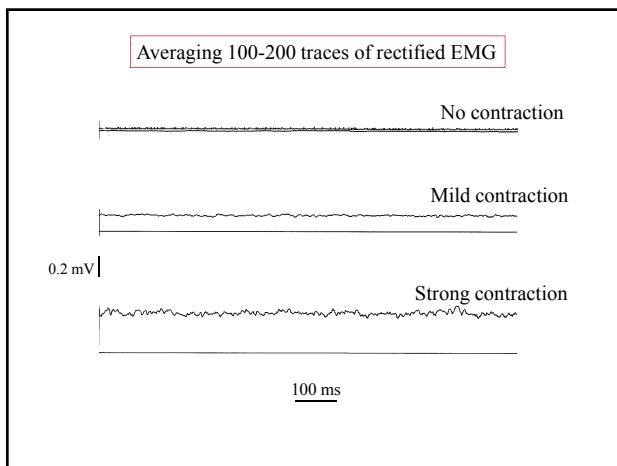
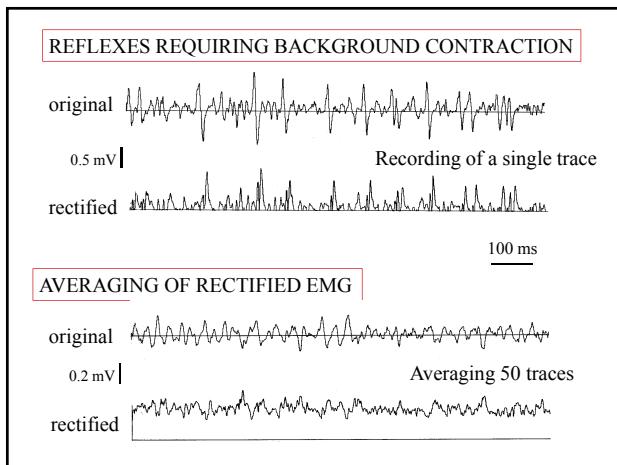


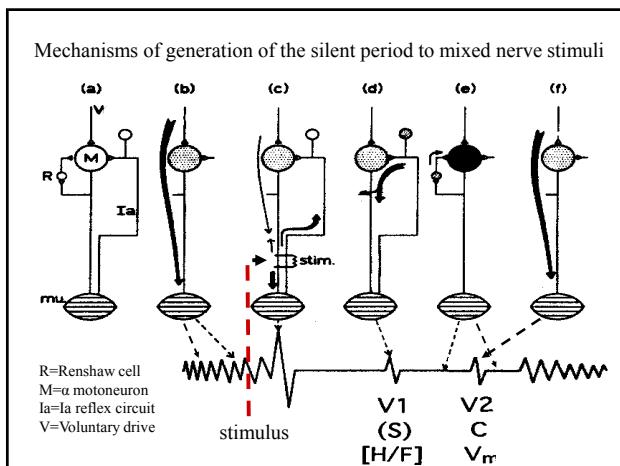
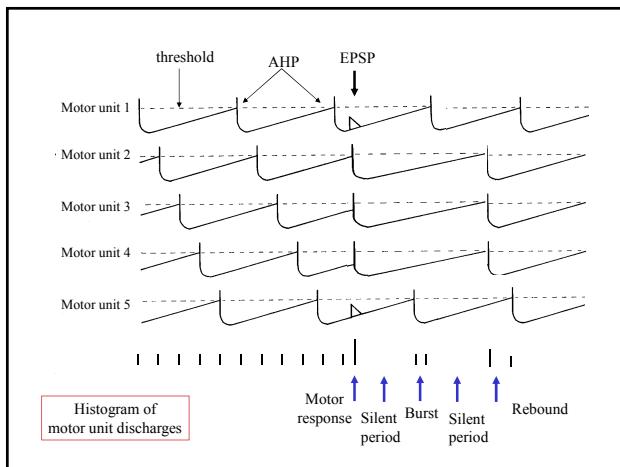












CONCLUSIONS

The H reflex, T wave and F wave should be part of the clinical studies of proximal nerve conduction when suspecting polyneuropathies or radiculopathies

The blink reflex, jaw jerk and masseteric inhibitory reflex are useful in the study of cranial nerves and brainstem lesions

The study of inhibitory reflexes requires modulation of the EMG activity. The silent period to cutaneous afferents differs from that to mixed nerve afferents.

Collision between antidromic impulses and descending volleys explains the presence of bursts in the middle of the silent period to mixed nerve stimuli