

EMG at full activation Give a few words of argument why each of these TERMS are less than optimal

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Interference pattern Summation pattern Recruitment pattern

50,60Hz not much summation, equally much subtraction Pattern at strong contraction neutral but complicated term leads thoughts to the way MU are orderly recruited

Can a MUP have a longer total duration than the interval between discharges of this MUP



Is it possible to decide whether a recorded EMG signal originates in the nerve or in the muscle Is there any difference in MUP parameters if the recording is obtained 2 cm or 5 cm from the end-plate?

We sometimes record double discharges (extra discharges) in voluntary EMG. Do we require that the two discharges are identical in shape, to separate them from occasional occurrence of discharges from 2 different MUPs

Critical illness: are fibrillations usually a sign of denervation in critical illness?

Critical illness: is the myosin content lower in CIM than in CIP Critical illness: is sural amplitude different in CIM and CIP

Can an A-wave appear after the F-waves

Can an A-wave and an F-response be generated in the same axon by a given stimulus (SFEMG necessary to identify that we record from the same axon) Is there any difference in amplitudes between Awaves and individual F-responses Monopolar recording. Is there any difference in the pattern at voluntary contraction if the distance between the two recording monopolar electrodes ("active" and "reference") is 1 cm or 10 cm?

Can you detect the "size principle" with conventional needle electrodes?

Concentric electrode has an oval recording surface: are the MUP parameters different for transversal or longitudinal insertion of the electrode (in relation to the fiber direction). Which is the concentric needle electrode recording uptake radius (180 or 360 degree) for the duration parameter in a MUP

Which is the concentric needle electrode recording uptake radius (180 or 360 degree) for the spiky part of the MUP Is it possible to make sure that you are stimulating muscle fibers directly and not intramuscular nerves in so called direct muscle stimulation (critical illness tests)

You may stimulate one or very few axons at two different sites (prox and dist) and record a SFEMG response from corresponding muscle and so measure the conduction in a single axon. How do you ascertain that you have stimulated exactly the same axon? SFEMG: how many spikes do you need to record simultaneously to detect neurogenic blocking

SFEMG: how many spikes do you need to record simultaneously to detect neurogenic jitter?

Reinnervation. In the early stage of reinnervation (20 days) after a partial nerve lesion, you start to see MUPs with some jittering spikes. In general is the MUP "small" or "large"?

In monopolar EMG recording you often see a small positive going signal on the end slope of the signal. What is this, and why do you not see that in concentric needle EMG

With increasing force, the EMG amplitude (envelope amplitude) increases. Why?

In concentric needle electrode recordings, one can sometimes obtain low amplitude MUP that looks "upside down". Explanation?

. Neurography: Low ampl CMAP

Normal sensibility, strong muscles: If the CMAP of APB has an amplitude < 4mV and CMAP from ADM is norma and the SNAP from dig III is normal which is a likely reason

CTS
 LEM
 Technical problem
 Polyneuropathy

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LEM is very rare and technical problems common. Answer on these grounds

"Neurogaphy: Low ampl CMAP If the CMAP of ADM and APB have an amplitude < 2mV , sensory signal OK which is a likely reason CTS LEM Technical problem Slight polyneuropathy

Meurography: Low ampl CMAP

If the CMAP $% \left(ABM\right) =0$ of ADM and APB have an amplitude < 2mV , sensory signals OK which is a likely reason

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"Neurography: Root or plexus

Patient 80 yo, weakness and numbness in the right leg. Low sensory amplitude in right fibularis superficialis but normal on the left side. Reduced CMAP ampli ni right ED8 EMG in Tib ant show denervation. EMG in lumbar paraspinals normal

1. L5 root

2. plexopathy
 3. central lesion
 4. polyneuropathy

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Normal sensory and paraspinal EMG favors plexus lesion

"EMG: Myasthenia?

Patient with ptosis but no arm or leg weakness. RNS in nasalis and deltoid normal. SFEMG in orb oculi shows jitter and some blocking. Jitter abnormal in Ext dig.

Ocular MG
 Generalized MG
 Myopathy
 Miller Fisher syndrome

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MG classification is conventionally defined on clinical grounds. Jitter is abnormal in ED in 60% of cases with Ocular MG

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Normal SFEMG findings are strong indications that symptoms are NOT MG. In this case, it may be a myopathy, often with very little of jitter abnormalities

Q 17. In a slight CTS the palm stimulation (wrist recording) may have normal orthodromic latency but abnormal ortho from dig 3 stimulation. Why?

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Pure sensory CTS. Palm stim includes motor nerve fibers, which now give normal latency

Fib and pws

Which alternative(s) is (are) correct?

- generated in the axon
 generated in individual muscle fibers
- usually appear with irregular firing rhythm
 always sign of axonal pathology
- 5. PSW more significant in the EDX interpretation

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EDX in demyelinating neuropathy with conduction block

Which alternative is expected EDX finding in these conditions

- 1. Reduced MUP amplitudes
- 2. Reduced CMAP amplitudes
- 3. Reduced fullness of interference pattern at strong contractions
- 4. Interference pattern at strong contraction most pronounced proximally
- 5. Abundant fibs in distal muscles

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soWhat do you call this phenomenon?



This is CRD, complex repetitive discharges, not myotonia

Abrupt start and stop. No waxing and waning



"EMG: acute weakness

Patients with acute weakness since 2 days. If this is GBS, which combination of findings do we get

- 1. normal CMAP and CV normal MUPs and firing
- 2. normal CMAP , reduced CV early reinnervation
- 3. Reduced CMAP ampl first signs of denervation
- 4. normal CMAP, normal CV, reduced # F waves, presence of A waves
 - abnormal interference pattern

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CMAP and CV changes and denervation signs have not yes developed. Weakness may be due to proximal conduction block; thus reduced F persistence and reduced EMG pattern at effort

Acute weakness

History:

Formerly healthy man with acute weakness from 7 days ago, progressive symptoms.

the patient reports double vision and an unsteady gait.

Clinic:

external ophthalmoplegia, ataxia and areflexia weak facial muscles, ptosis and reduced mobility of the tongue general mild muscle weakness. Sensory diffusely affected

EDX findings con't and QUESTION

Acute weakness, EDX and Questions

- Neurography, MCS e general slightly decreased MCV F latencies extended DLAT extended distal amplitude reduced, ssk in the facial muscles

- Neurography SCS
 abnormal response with low amplitudes, more than
 reduced CVEMG
 reduced interference pattern, 0 fib
 autonomous tests slightly abnormal
 abnormal Bink reflex
 prolonged latency to R1 and R2

Question Likely diagnosis? GBS

GBS Lyme disease brainstem involve Miller Fischer MG

Question What is your next step RNS outpatient CFS analysis test for antibodies against cholinergic receptor SFEMG admission for diagnosis and treatment

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Discrepancies		
70. Good CMAP in weak muscle, no twitch at nerve stimulation.	72. Clear sensory experience with digital nerve stimulation but no SNAP from the wrist	
Normal F response.	Explanation?	
Explanation? conduction block recent axonotmesis tendon rupture pnp MG myopathy	proximal conduction block wrong recording position cold hand bipolar stim pulse has been used the stimulus too short	
71.Good finger SNAP but no sensation at stimulation Why?	73. Normal MUP and low MCV What?	
pnp conduction block myelopathy the wrong digital nerve is stimulated GBS	axonal pnp conduction block demyelination LEM posttraumatic reinnervation	

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Which	is the EDX method of choice for the following diagnostic questions
MG	RNS
	SFEMG
CTS	EMG of APB and ADM
	antidromic sensory neurography
GBS	EMG
	neurography
	thermotest
Root	EMG
	neurography
	evoked potentials
ALS	EMG
	neurography
	Motor unit counting



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	Methods of choice
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Reference List

- Stålberg E and L Karlsson. Simulation of the normal concentric needle electromyogram by using a muscle model. Clin.Neurophysiol.:2001;112(3): 464-71.
- [2] Stålberg E, et al. Single Fiber EMG (ed. 3rd). Uppsala, Edshagen Publishing House. 2010
- [3] Ertas M, et al. Can the size principle be detected in conventional EMG recordings? Muscle Nerve:1995;18: 435-9.
- [4] Nandedkar S D, et al. Selectivity of electromyographic recording techniques; a simulation study. Med Biol Eng Comput:1985;23: 536-40.
- [5] Nandedkar S D and D B Sanders. Recording characteristics of monopolar EMG electrodes. Muscle Nerve: 1991;14: 108-12.
- [6] Stålberg E, et al. Electrical microstimulation with single-fiber electromyography: a useful method to study the physiology of the motor unit. Journal of Clinical Neurophysiology:1992;9: 105-19.
- [7] Gydikov A and D Kosarov. Extraterritorial potential field of impulses from separate motor units in human muscles. Electromyography Clinical Neurophysiology:1972;12: 283-305.