

EMG signs of denervation and reinnervation

Markus Weber, MD

Scandinavia goes St.Gallen

March 29- April 1, 2023

St.Gallen

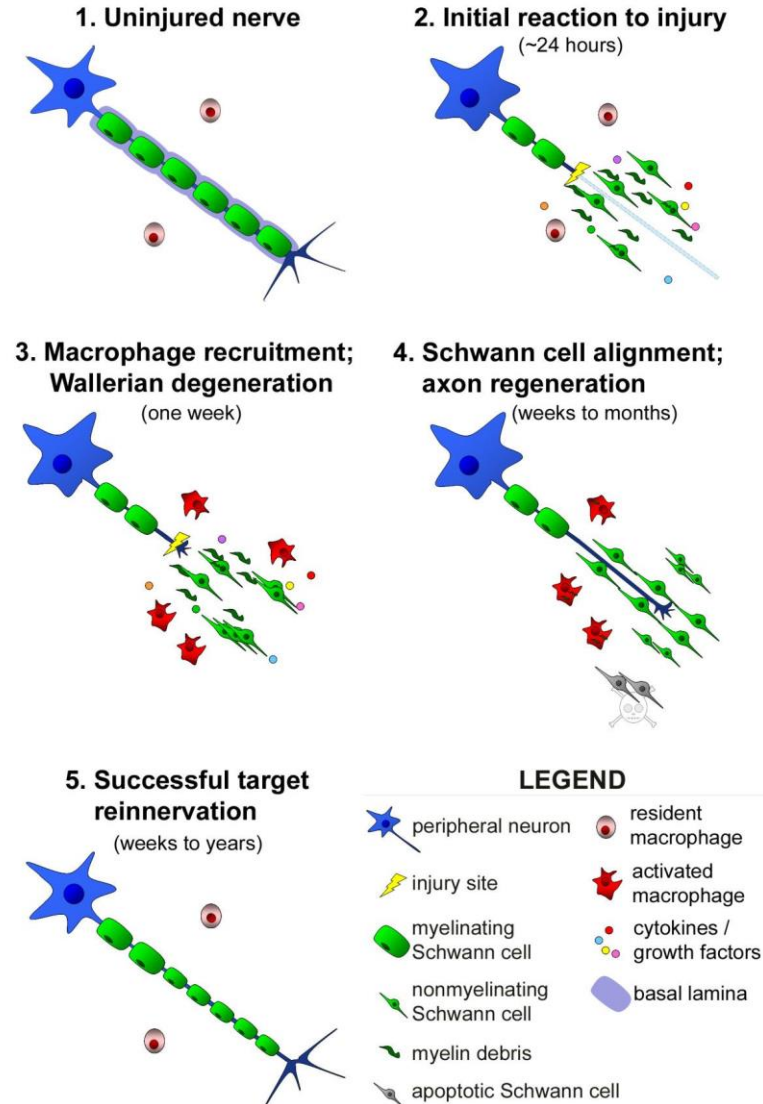
The motor endplate



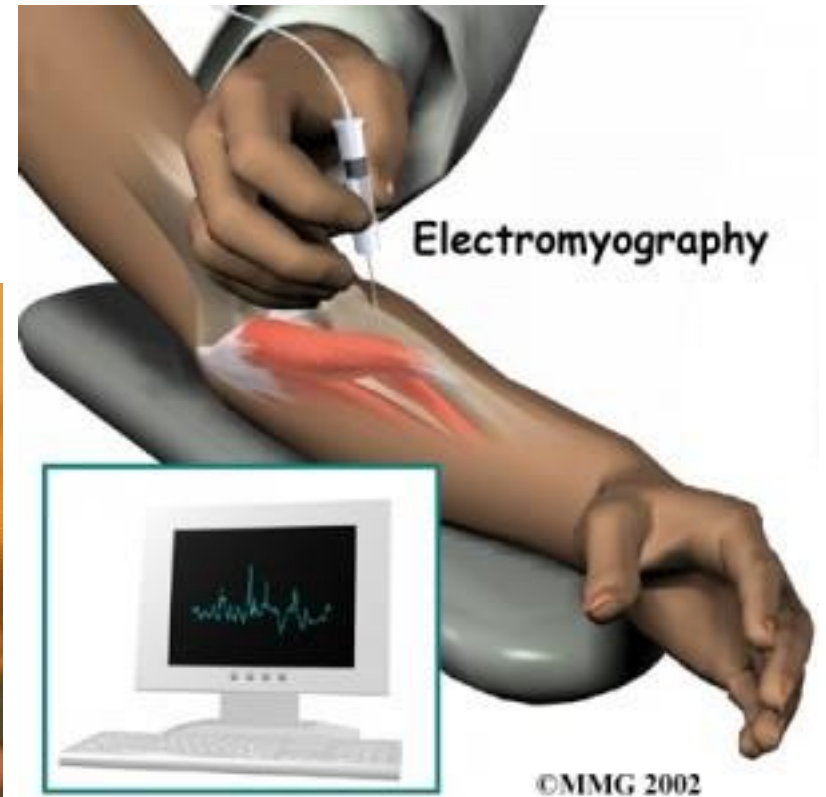
Etiology of axonal damage and denervation

- Acute (injury, trauma)
 - Complete axonal damage
 - Incomplete (partial) axonal damage
- Chronic
 - Compression (carpal tunnel syndrome, herniated disc, etc)
 - Neuropathies (diabetes, alcohol abuse, autoimmune)
 - Axonal
 - Inherited/acquired
 - Demyelinating neuropathies
 - Inherited/acquired
 - Neurodegeneration/ amyotrophic lateral sclerosis

Pathophysiology of denervation and reinnervation

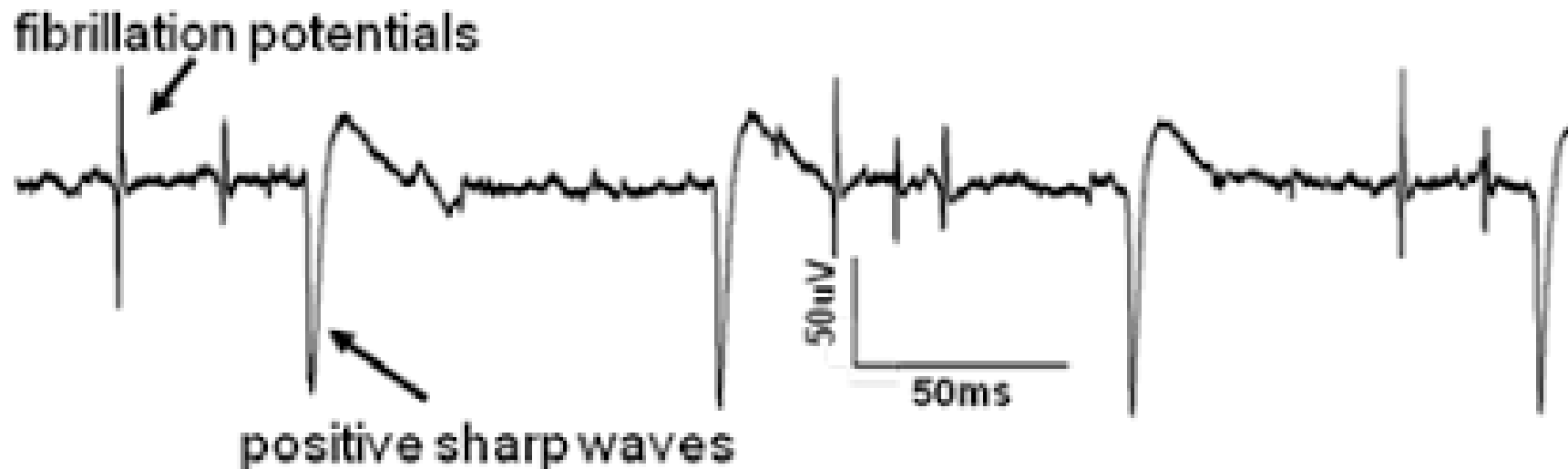


Clinical Neurophysiology



EMG signs of denervation

- Abnormal spontaneous activity
 - positive sharp waves
 - fibrillation potentials



SINGLE MUSCLE FIBER DISCHARGES (INSERTIONAL ACTIVITY, END-PLATE POTENTIALS, POSITIVE SHARP WAVES, AND FIBRILLATION POTENTIALS): A UNIFYING PROPOSAL

DANIEL DUMITRU, MD

MUSCLE & NERVE 19:216–220 1996

ARE FIBRILLATION POTENTIALS AND POSITIVE SHARP WAVES THE SAME? NO

GEORGE H. KRAFT, MD

tion of PSWs conform to that of a blocked fibrillation potential.

Muscle Nerve **36**: 349–356, 2007

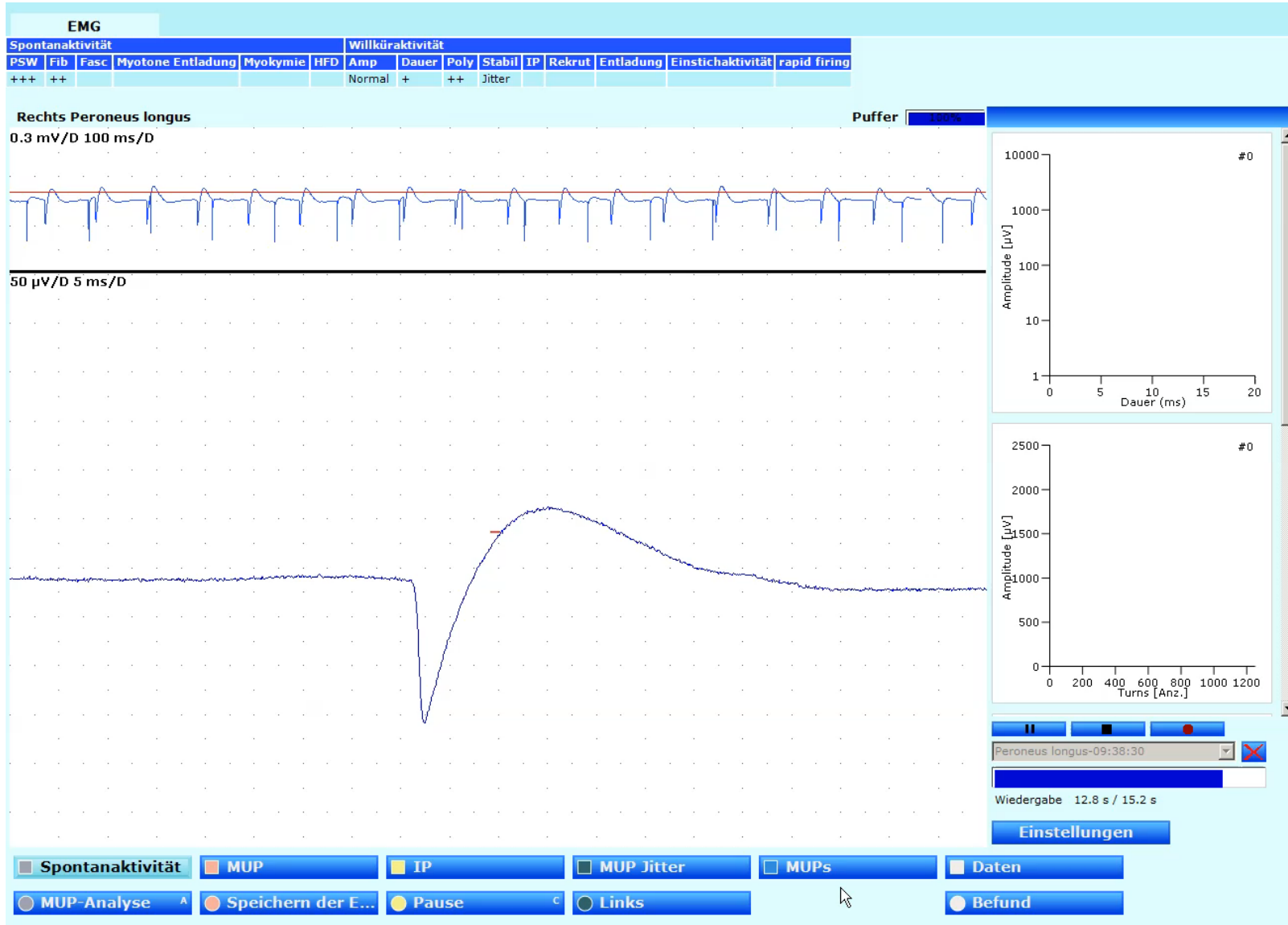
POSITIVE SHARP WAVE ORIGIN: EVIDENCE SUPPORTING THE ELECTRODE INITIATION HYPOTHESIS

DANIEL DUMITRU, MD, PhD, and DANIEL L. SANTA MARIA, MD

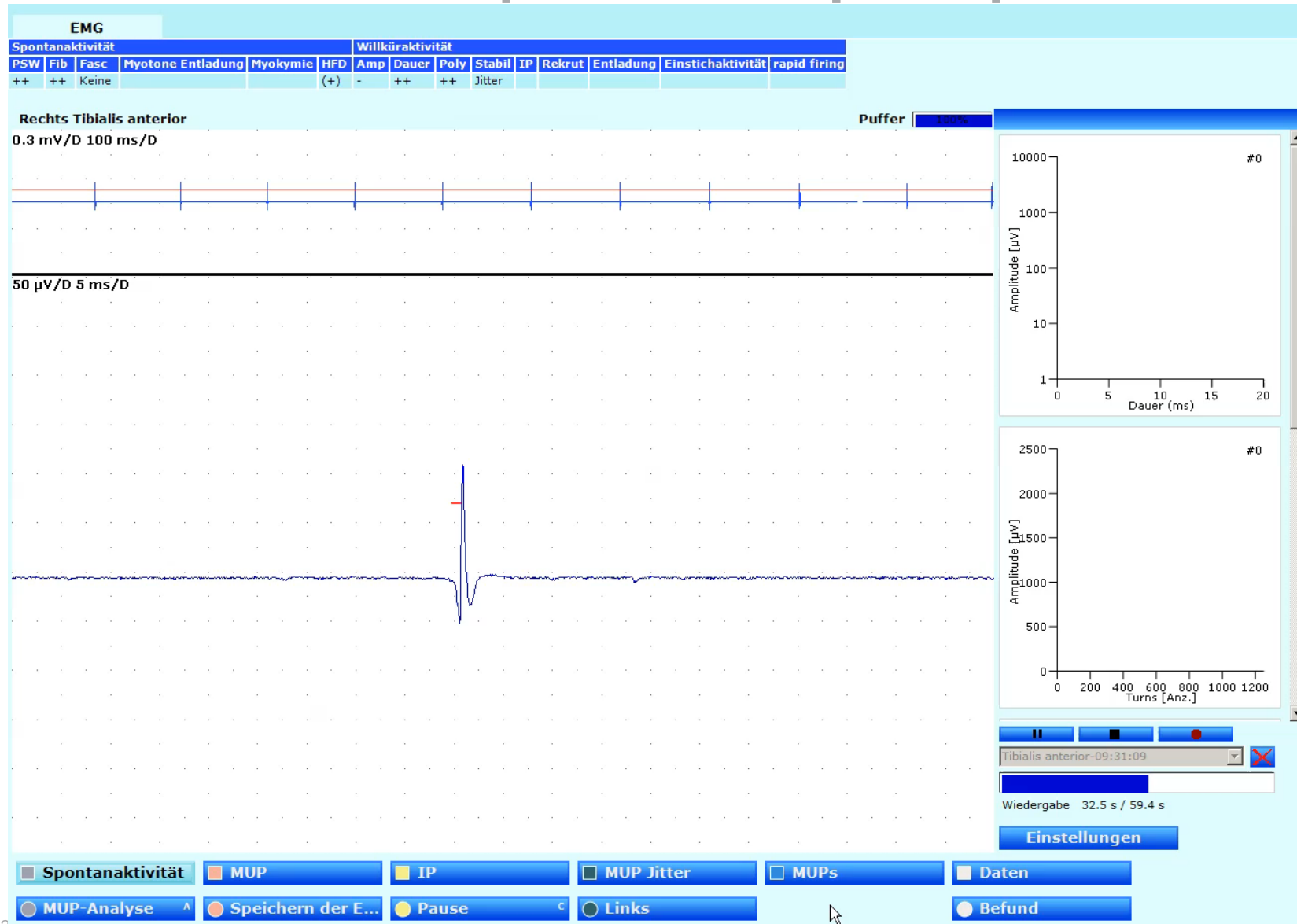
Department of Rehabilitation Medicine, University of Texas Health Science Center,
7703 Floyd Curl Drive, San Antonio, Texas 78229-3900, USA

Accepted 28 March 2007

Denervation after complete nerve injury



Fibrillation potential superimposed



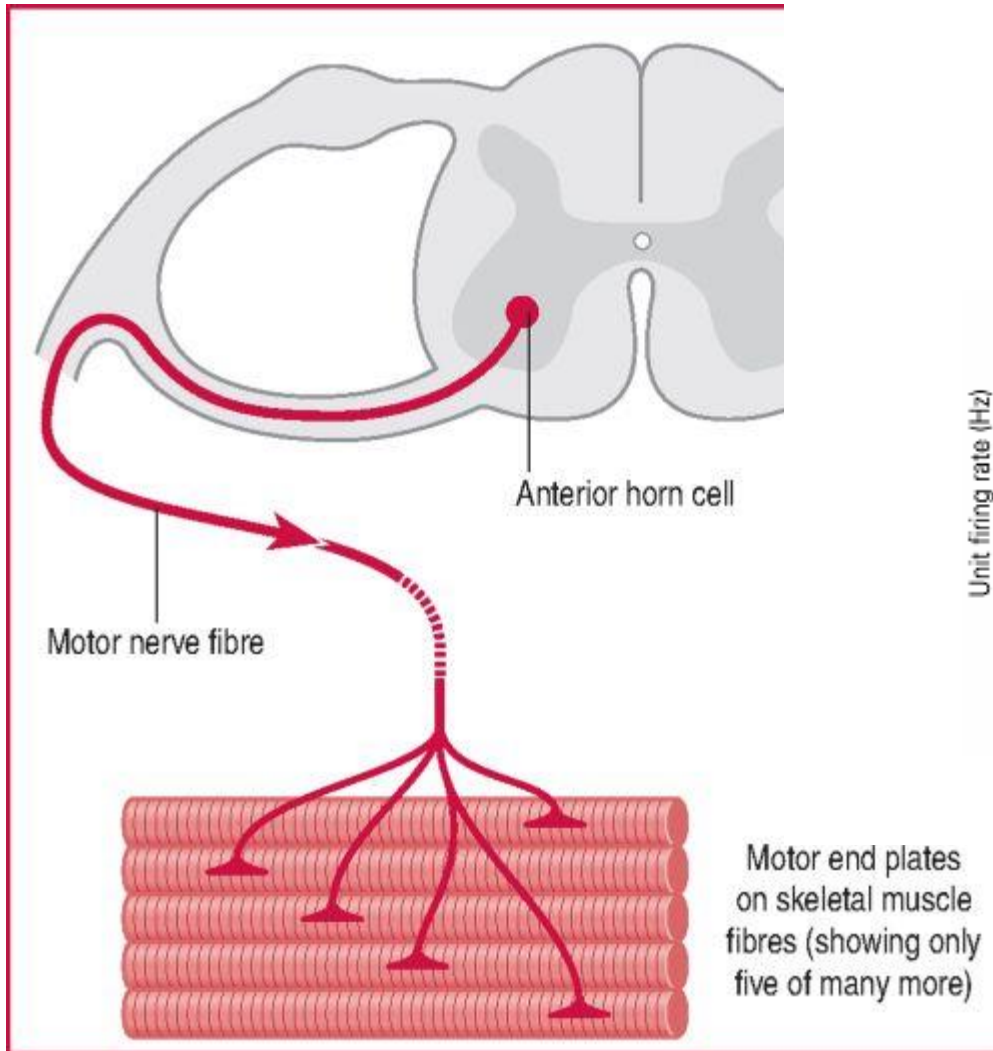
Nerve lesion and voluntary EMG

- Complete axonal
 - Spontaneous activity (positive waves and fibs)
 - No voluntary activity
- Partial axonal
 - Spontaneous activity
 - Voluntary activity
 - Time dependent

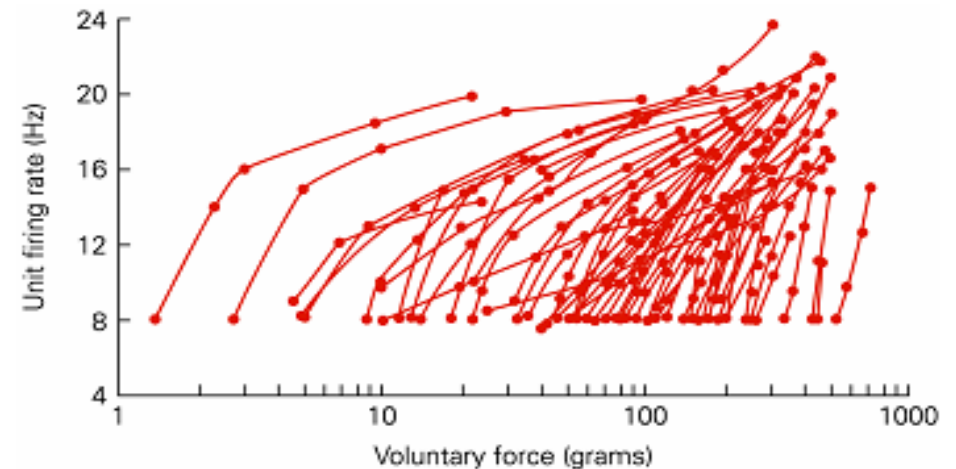
Motor unit firing



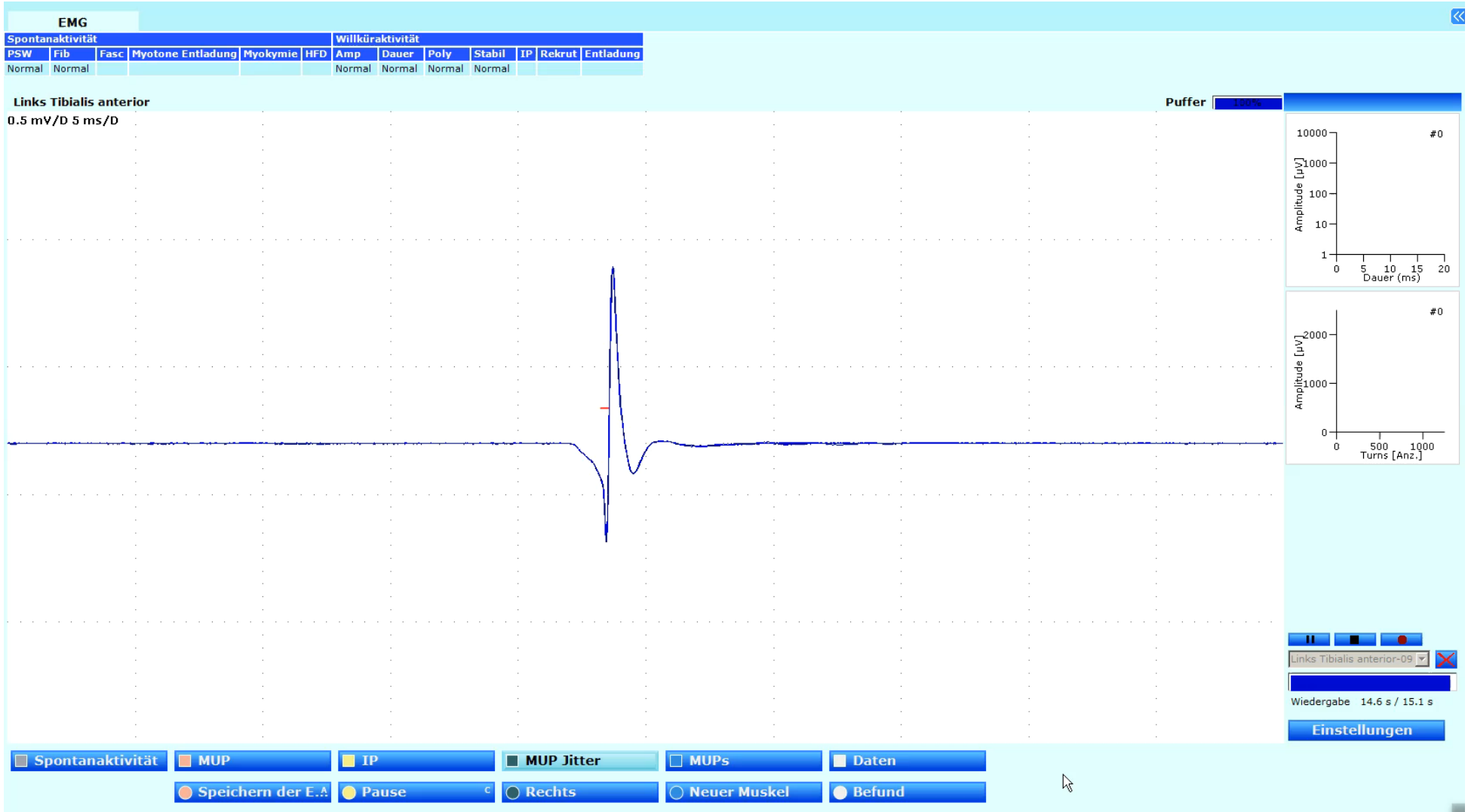
The motor unit and voluntary recruitment



Recruitment and Rate Coding

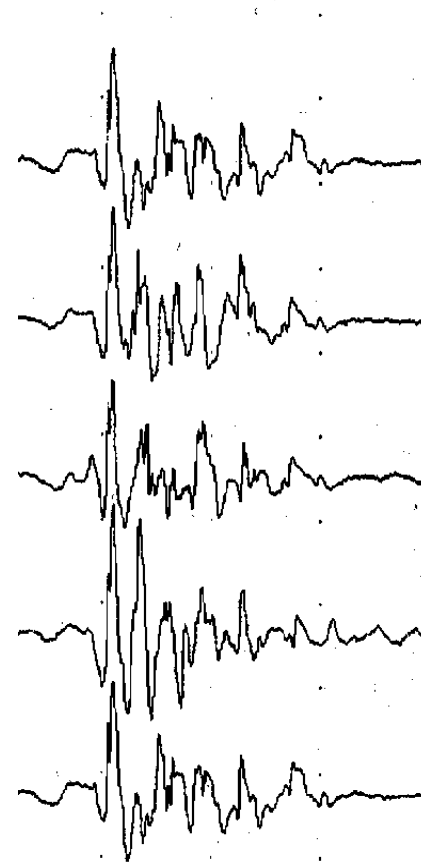
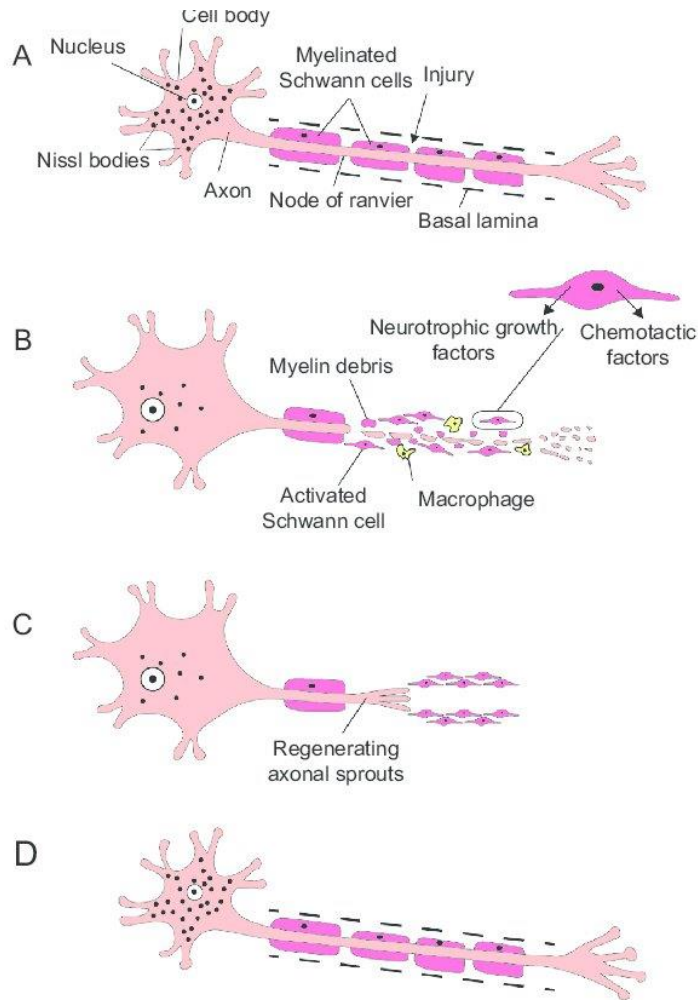


Normal MU stable superimpose



Nascent reinnervation after complete axonal damage and Wallerian degeneration

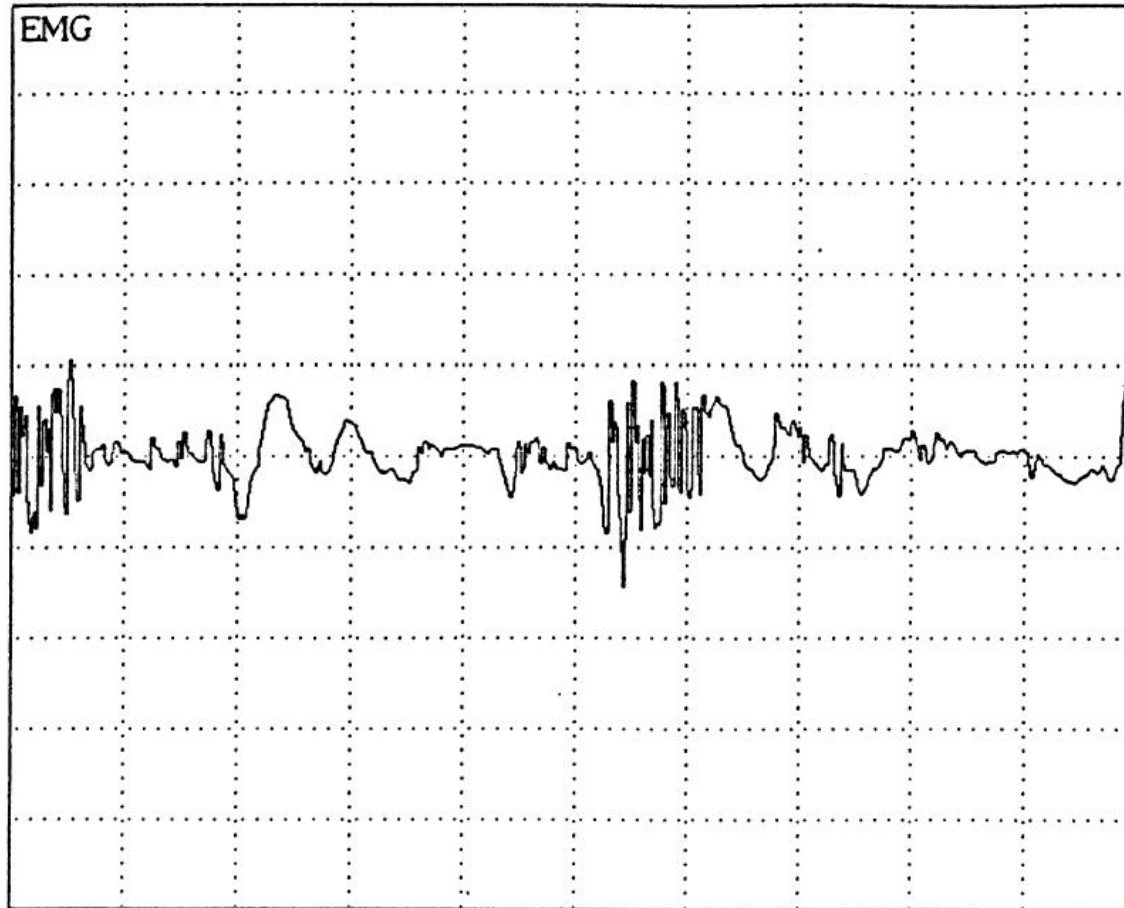
EMG



<https://www.researchgate.net/publication/289525755>

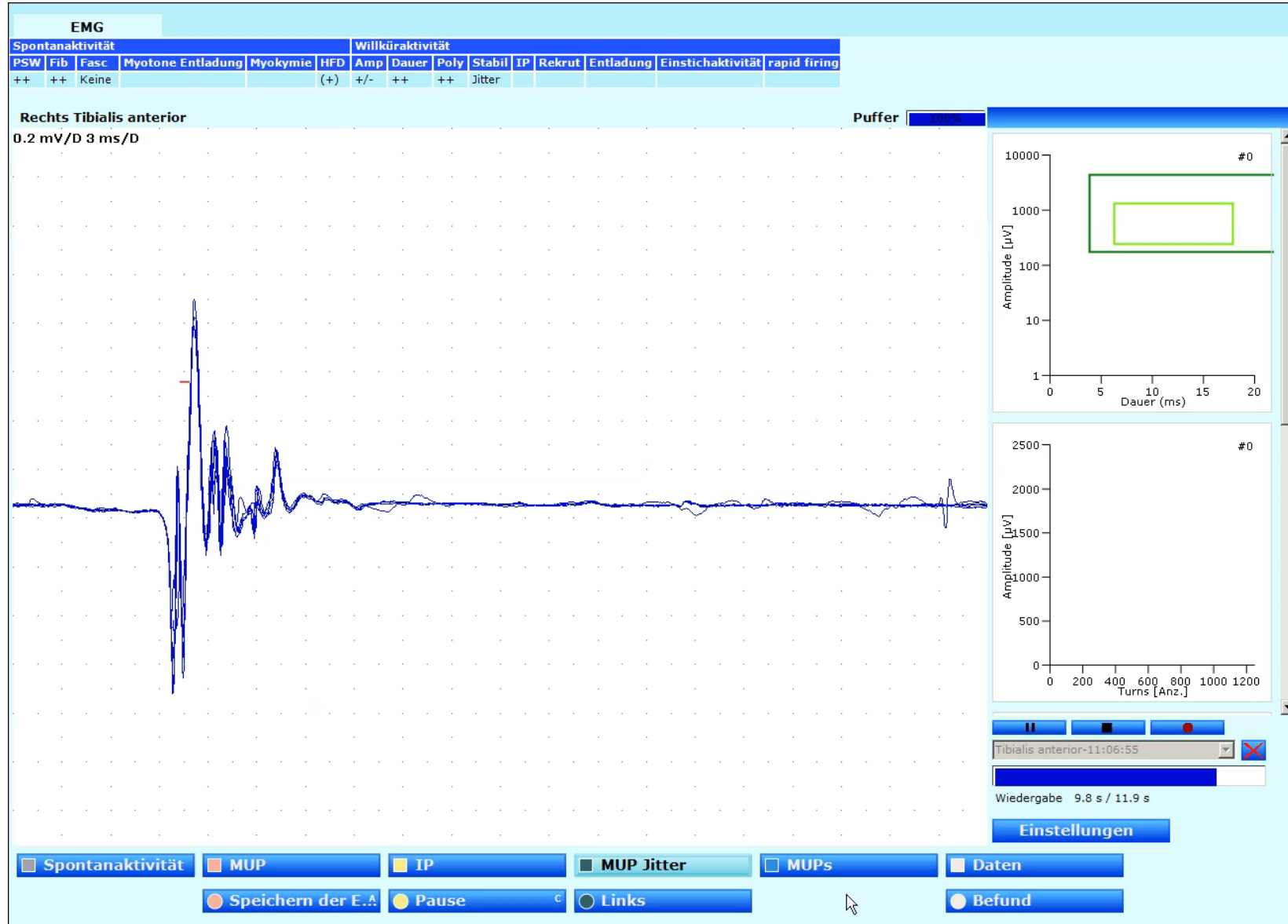
Nascent reinnervation

a



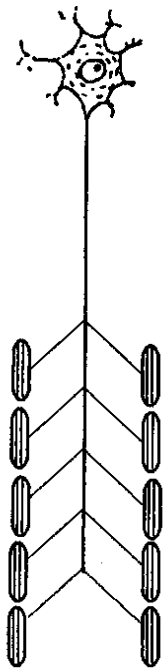
Ch	Hicut	Locut	Gain ($\mu\text{V}/\text{div}$)	Sweep (ms/div)
1	10000	10.00	200.0	20.0

Nascent reinnervation after complete nerve injury



EMG changes with partial nerve lesion: terminal sprouting

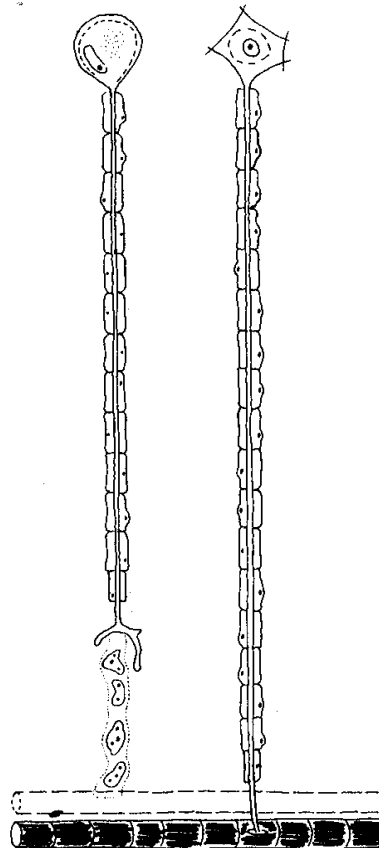
Normal



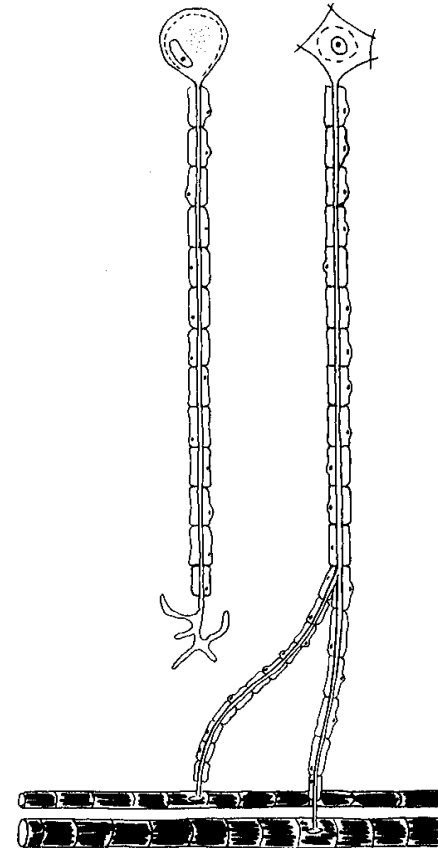
MUAP

Histology

- partial



- with collateral reinnervation



Jiggle

EMG **EMG Befunde** F8 **Einzel-faser-EMG** F9 **Stim SF EMG** F10

Spontanaktivität			Willküraktivität											
PSW	Fib	Fasc	Myotone Entladung	Myokymie	HFD	Amp	Dauer	Poly	Stabil	IP	Rekrut	Entladung	Einstichaktivität	rapid firing
+	+	Normal				+	++	++	Jitter					

Rechts Tibialis anterior (AMP1) Puffer 100%

0.5 mV/D 3 ms/D

Amplitude [µV]

Dauer (ms)

Amplitude [µV]

Turns [Anz.]

#0

#0

Tibialis anterior-09:56:53

Wiedergabe 0.0 s / 25.8 s

Einstellungen

Spontanaktivität

MUP

IP

MUP Jitter

MUPs

Daten

Speichern der E.A.

Pause

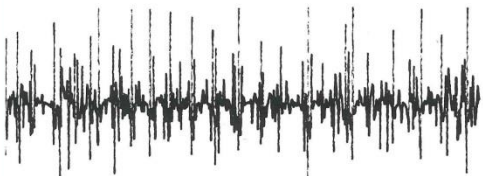
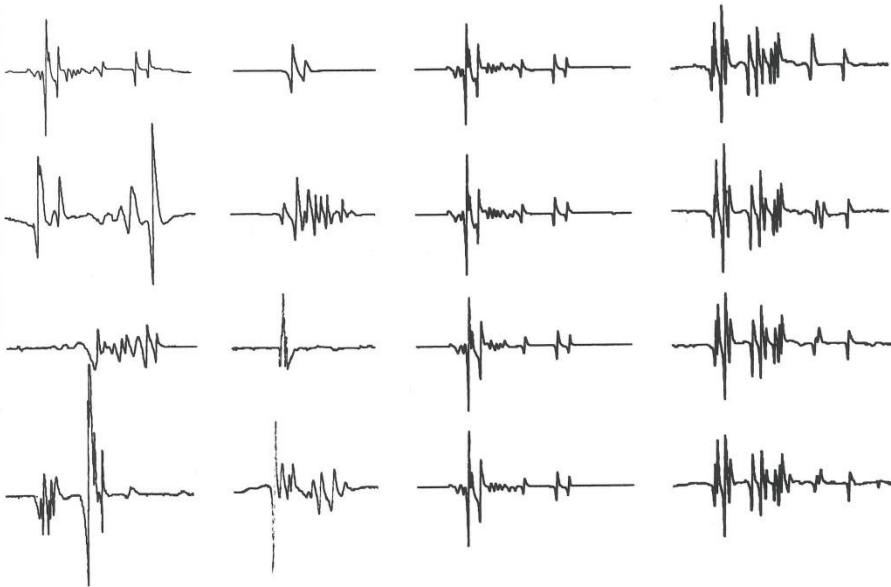
Links

Neuem Muskel

Befund

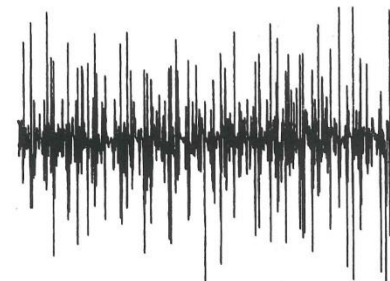
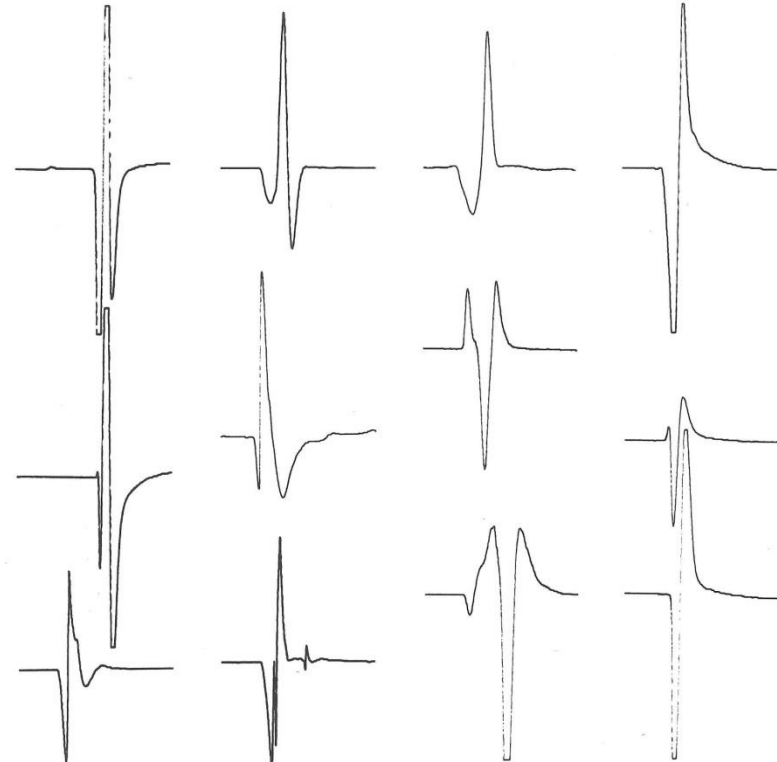
EMG changes over time

2.1. EMG-Befunde



0.1 mV
10 ms
100 ms
1 mV

Abb. 118: Frische Reinnervation (M. abductor pollicis brevis)



0.5 mV
10 ms
100 ms
1 mV

Abb. 120: Spätes Reinnervationsstadium

Comparison of EMG changes

complete

- Spontaneous activity +++
- Nascent reinnervation
 - Months-years
 - MUPS:
 - small
 - complex
 - unstable
 - fatigues easily

partial

- Spontaneous activity +, ++
- Terminal sprouting
 - Weeks-years
 - MUPS
 - Normal to large, «mother» unit
 - complex (satellites)
 - unstable (Jiggle)

Clinical examples: Amotrophic lateral sclerosis

- Signs of active denervation
- signs of chronic denervation («chronic neurogenic»)
- Fasciculation potentials

EMG changes in ALS: denervation

EMG										Willküraktivität				
Spontanaktivität														
PSW	Fib	Fasc	Myotone Entladung	Myokymie	HFD	Amp	Dauer	Poly	Stabil	IP	Rekrut	Entladung	Einstichaktivität	rapid firing
++	+				Keine	+	+	+	Jitter	Normal				

Links Tibialis anterior (AMP1)

0.3 mV/D 100 ms/D

50 µV/D 8 ms/D

Puffer 100%

#1

#42

Tibialis anterior-11:19:51

Wiedergabe 23.4 s / 29.9 s

Einstellungen

Spontanaktivität

MUP

IP

MUP Jitter

MUPs

Daten

MUP-Analyse

Speichern der E...

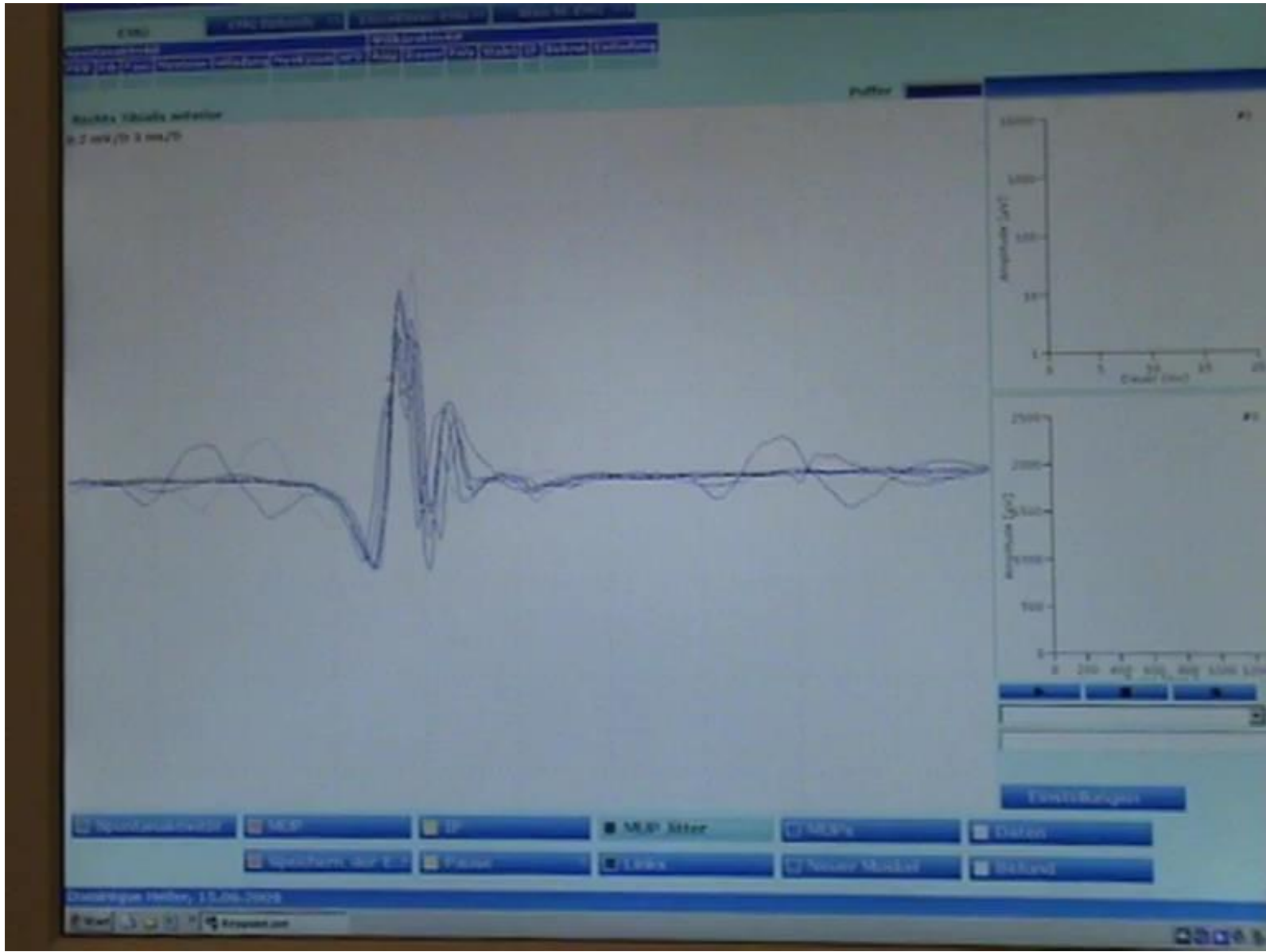
Pause

Rechts

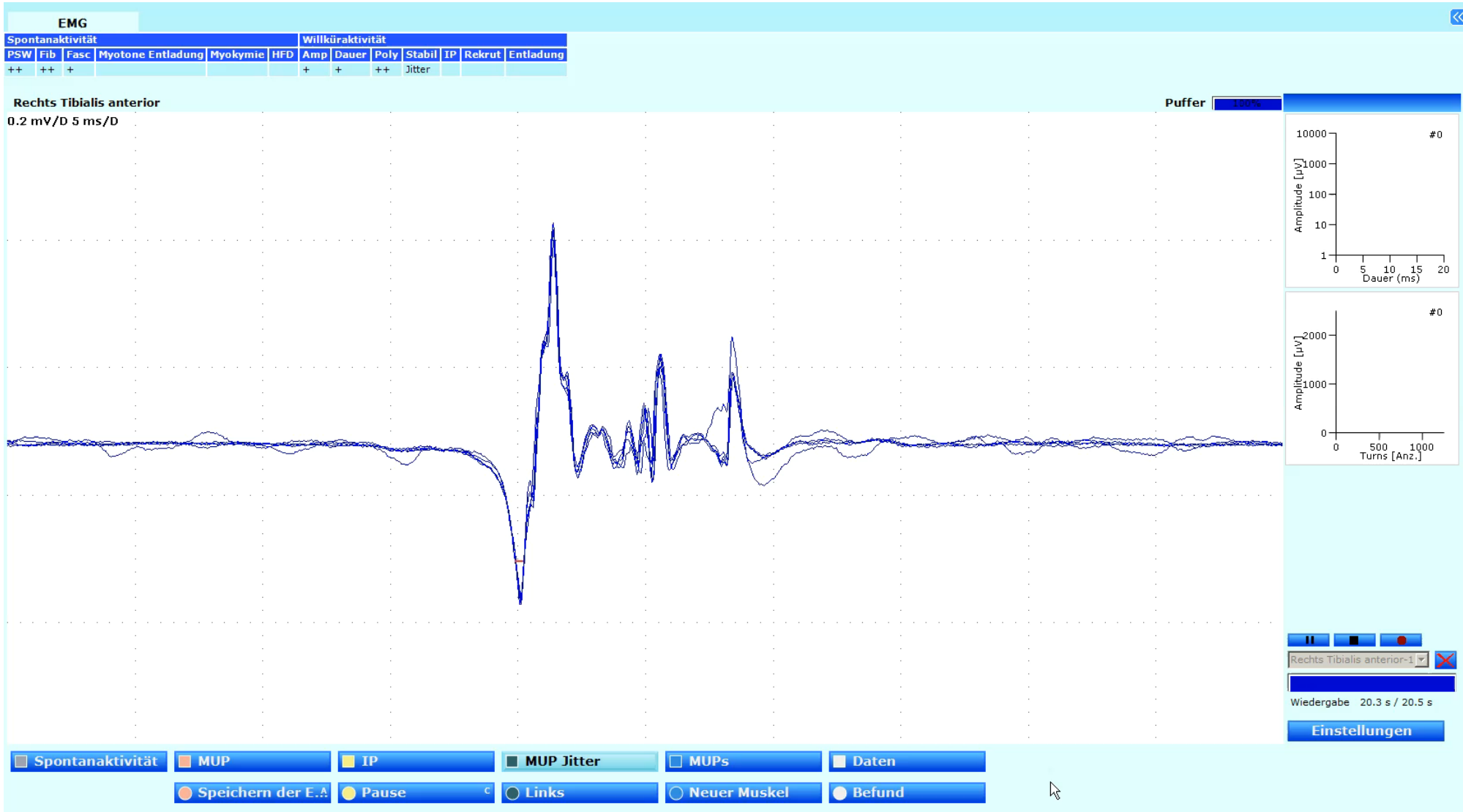
Neu Muskel

Befund

Chronic neurogenic... unstable MUPs (Jiggle)



EMG changes in ALS: chronic neurogenic



Electrophysiological Features of LMN Dysfunction (revised El Escorial criteria 2000)

- **Conventional EMG studies**
The features of LMN dysfunction ...are defined by electromyographicevidence of
 - **active *and***
 - **chronic denervation**
 - **fasciculations.**

EMG findings

- Active denervation:
 - spontaneous activity (fibrillation potentials, positive sharp waves)
- Chronic denervation:
 - impaired MUP recruitment (rapid firing)
 - unstable MUPs (Jiggle)
 - abnormal MUP size and shape (polyphasic potentials)



Clinical Neurophysiology 119 (2008) 497–503



www.elsevier.com/locate/clinph

Review

Electrodiagnostic criteria for diagnosis of ALS [☆]

Mamede de Carvalho ^a, Reinhard Dengler ^b, Andrew Eisen ^c, John D. England ^d,
Ryuji Kaji ^e, Jun Kimura ^f, Kerry Mills ^g, Hiroshi Mitsumoto ^h,
Hiroyuki Nodera ⁱ, Jeremy Shefner ^j, Michael Swash ^{k,*}

^a Department of Neurology, Hospital de Santa Maria, University of Lisbon, Lisbon, Portugal

^b Department of Neurology, Medizinische Hochschule Hannover, Germany

^c Department of Neurology, University of British Columbia, Vancouver, Canada

^d Department of Neurology, Billings Clinic, Billings, MT, USA

^e Department of Neurology, Tokushima University Graduate School of Medicine, Tokushima-city, Japan

^f Department of Neurology, University of Iowa, Iowa City, USA

^g Department of Neurology, Kings College Hospital, Guys Kings and St. Thomas's School of Medicine, London, UK

^h Eleanor and Lou Gehrig ALS Center, Neurological Institute, Columbia University, NY, USA

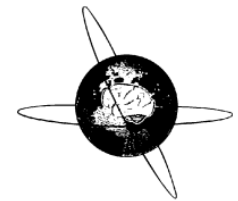
ⁱ Department of Neurology, Tokushima University, Tokushima-city, Japan

^j Department of Neurology, Upstate Medical University, Syracuse, NY, USA

^k Department of Neurology, Royal London Hospital, Queen Mary University of London, London, UK

Awaji consensus

1. *Edx and clinical data* are of equal and interchangeable value in diagnosing ALS
2. *In the presence of signs of partial denervation*, Fasciculation potentials (preferably of complex morphology) are equivalent to fibs-psw, indicating ongoing denervation
3. *Fibs and psws* are usually recorded in **strong, non-wasted** muscles
4. *Unstable MUPs & FPs* are especially relevant



Opinion Paper

A proposal for new diagnostic criteria for ALS



Jeremy M. Shefner^{a,*}, Ammar Al-Chalabi^b, Mark R. Baker^c, Li-Ying Cui^d,

- EMG abnormalities that must include:
- Both evidence of chronic neurogenic change, defined by large motor unit potentials of increased duration and/or increased amplitude, **with polyphasia and motor unit instability regarded as supportive but not obligatory evidence.**
- And evidence of ongoing denervation including Fibrillation potentials or positive sharp waves, or fasciculation potentials

RESEARCH PAPER

Fasciculation potentials and earliest changes in motor unit physiology in ALS

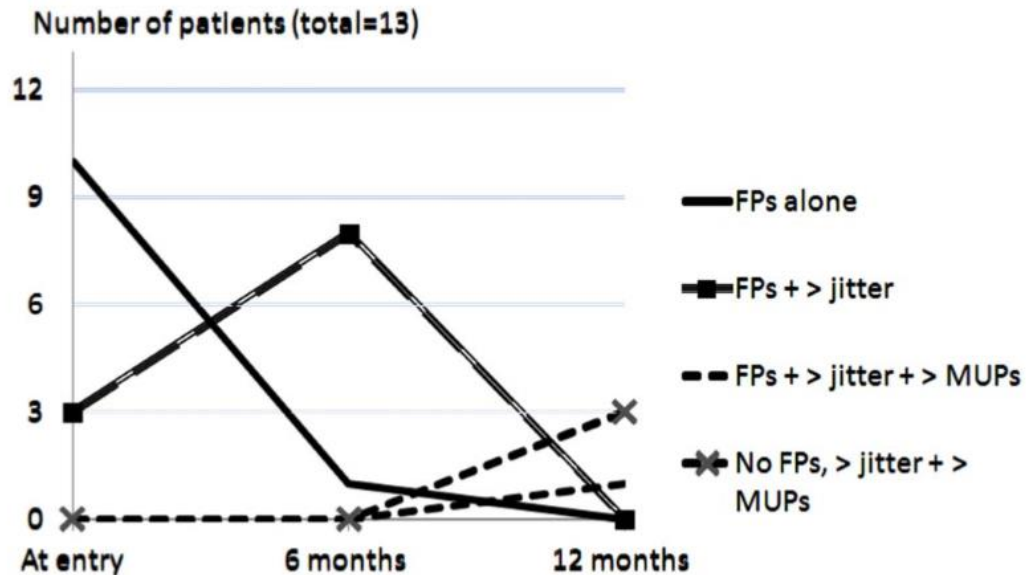
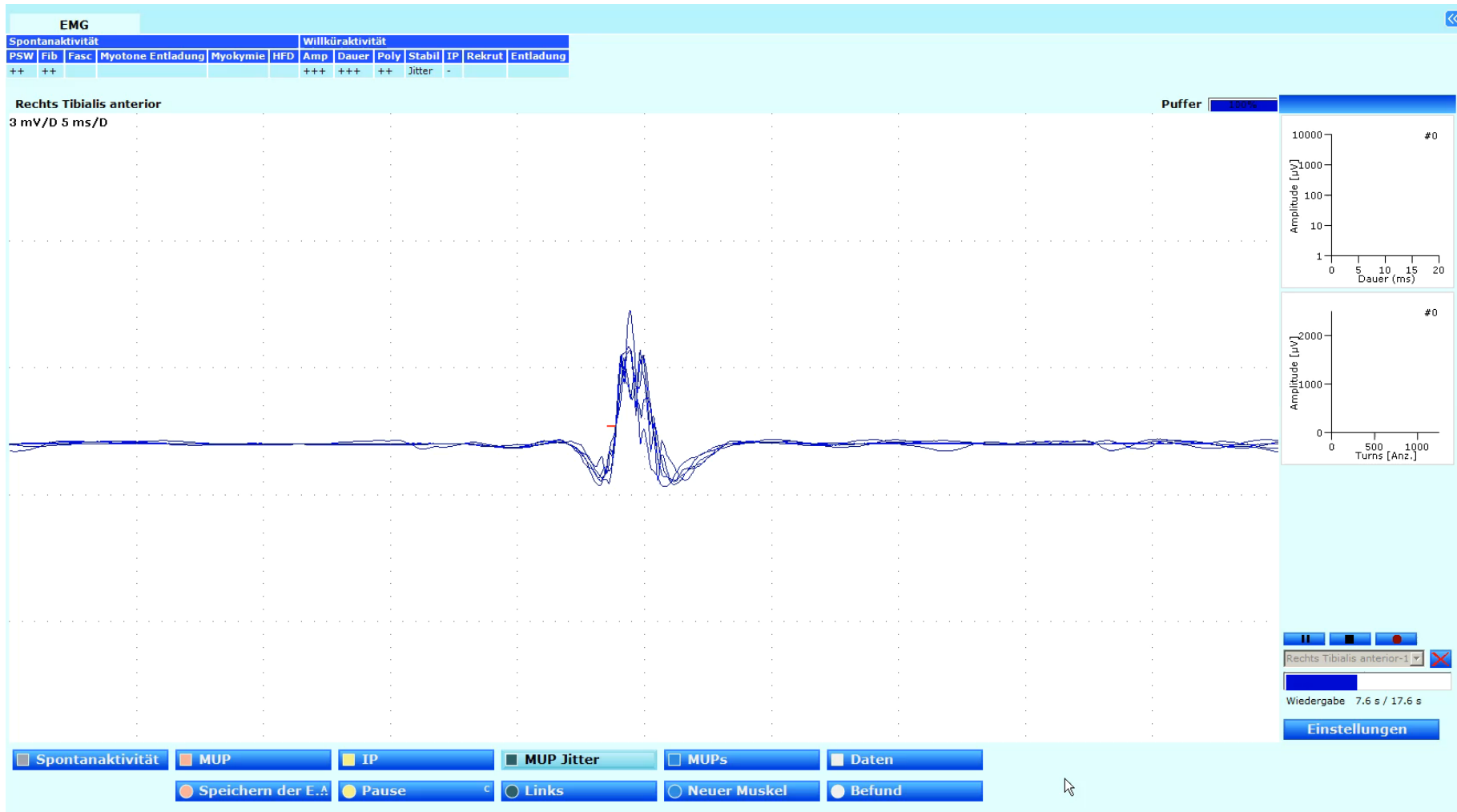
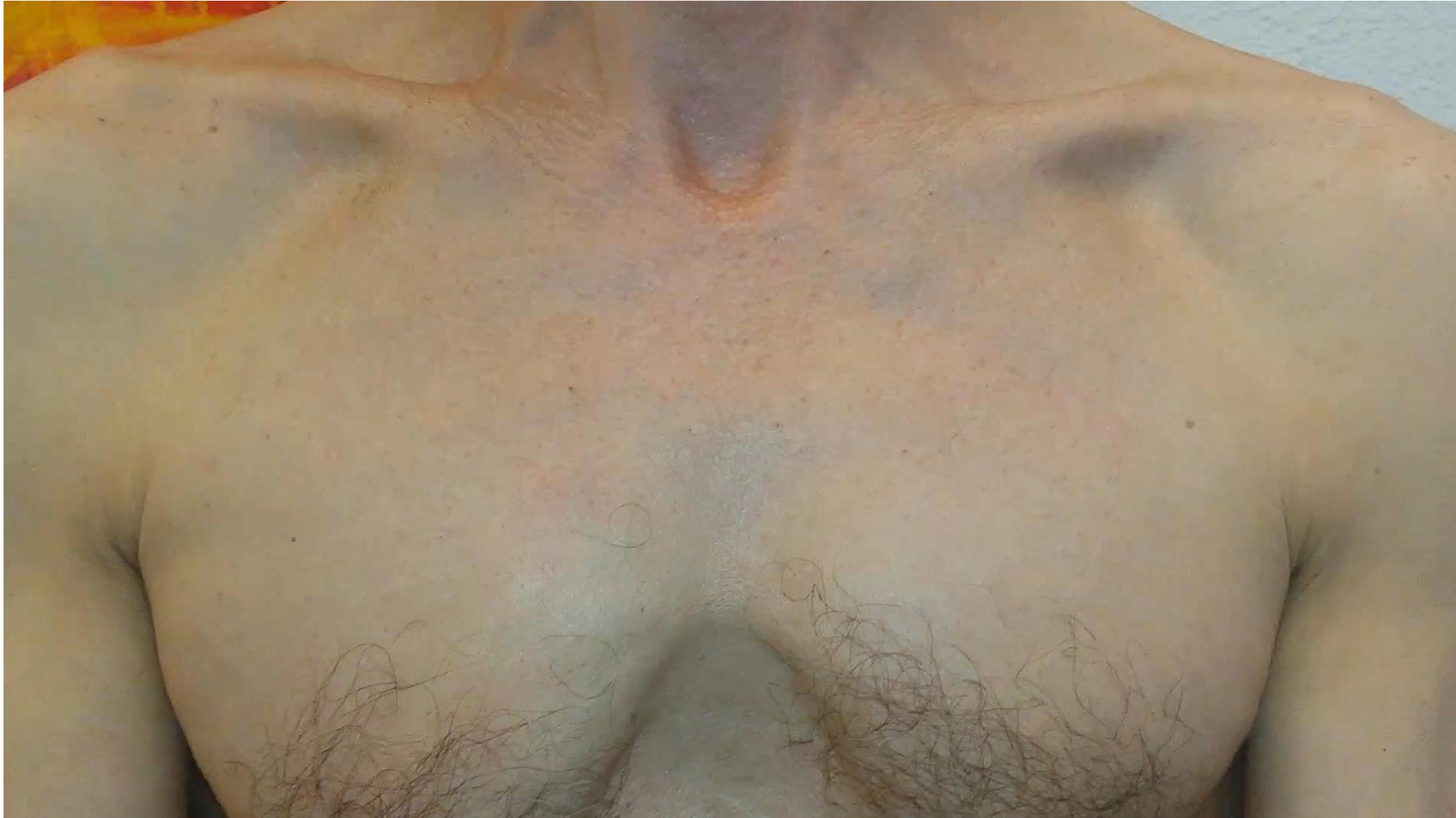
Mamede de Carvalho,^{1,2} Michael Swash^{1,2,3}Carvalho M, et al. *J Neurol Neurosurg Psychiatry* 2013;84:963–968

Figure 2 Progression of 13 amyotrophic lateral sclerosis patients with isolated fasciculation potentials (FPs) (normal motor unit potential (MUP) and no fibrillation/sharp-waves (fibs-sw)). All patients were evaluate 6 months later, but only four had preserved normal tibialis anterior strength 12 months after study entry. The Y-axis represents the number of patients.

Jiggle CMT II



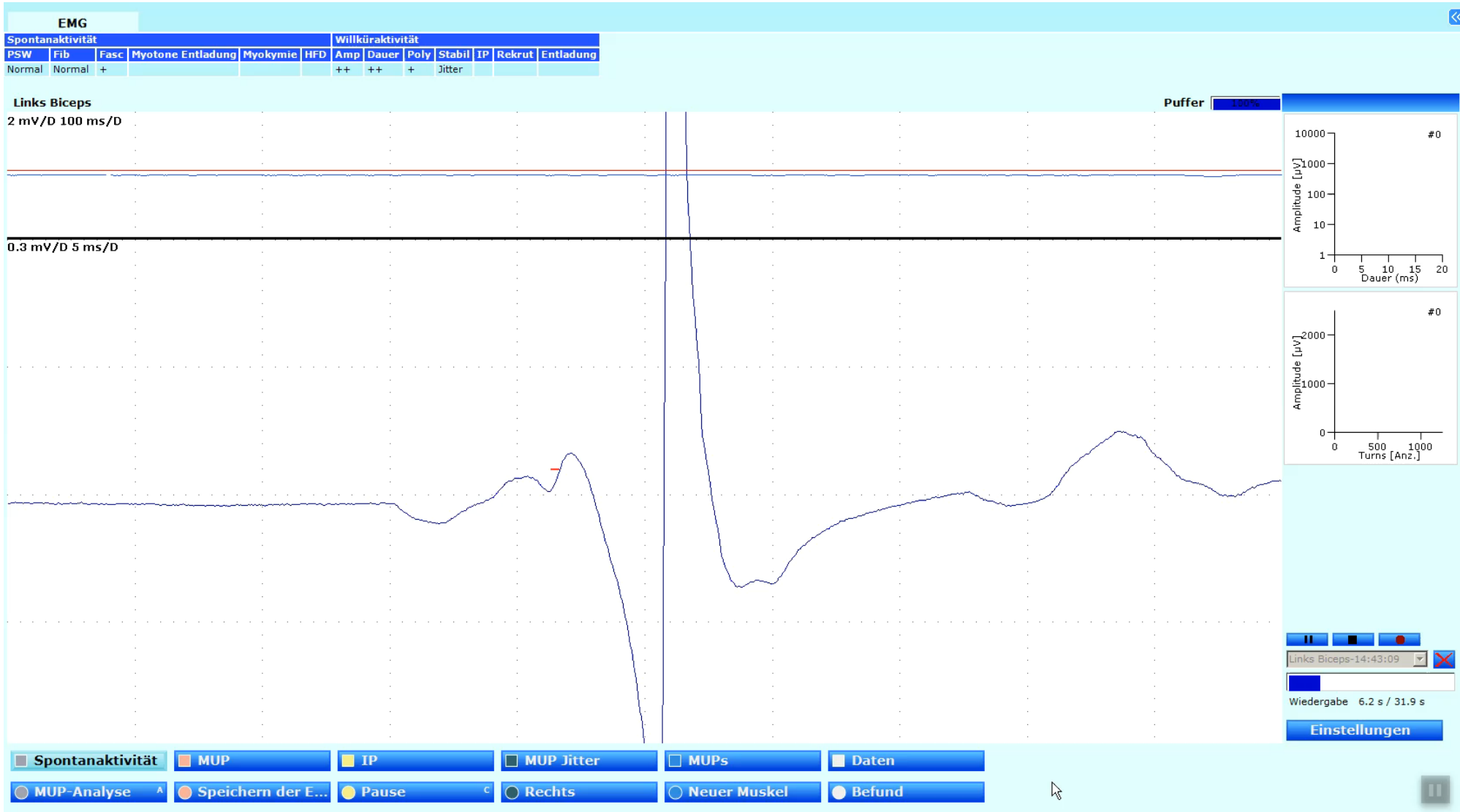
Fasciculations



Fasciculations



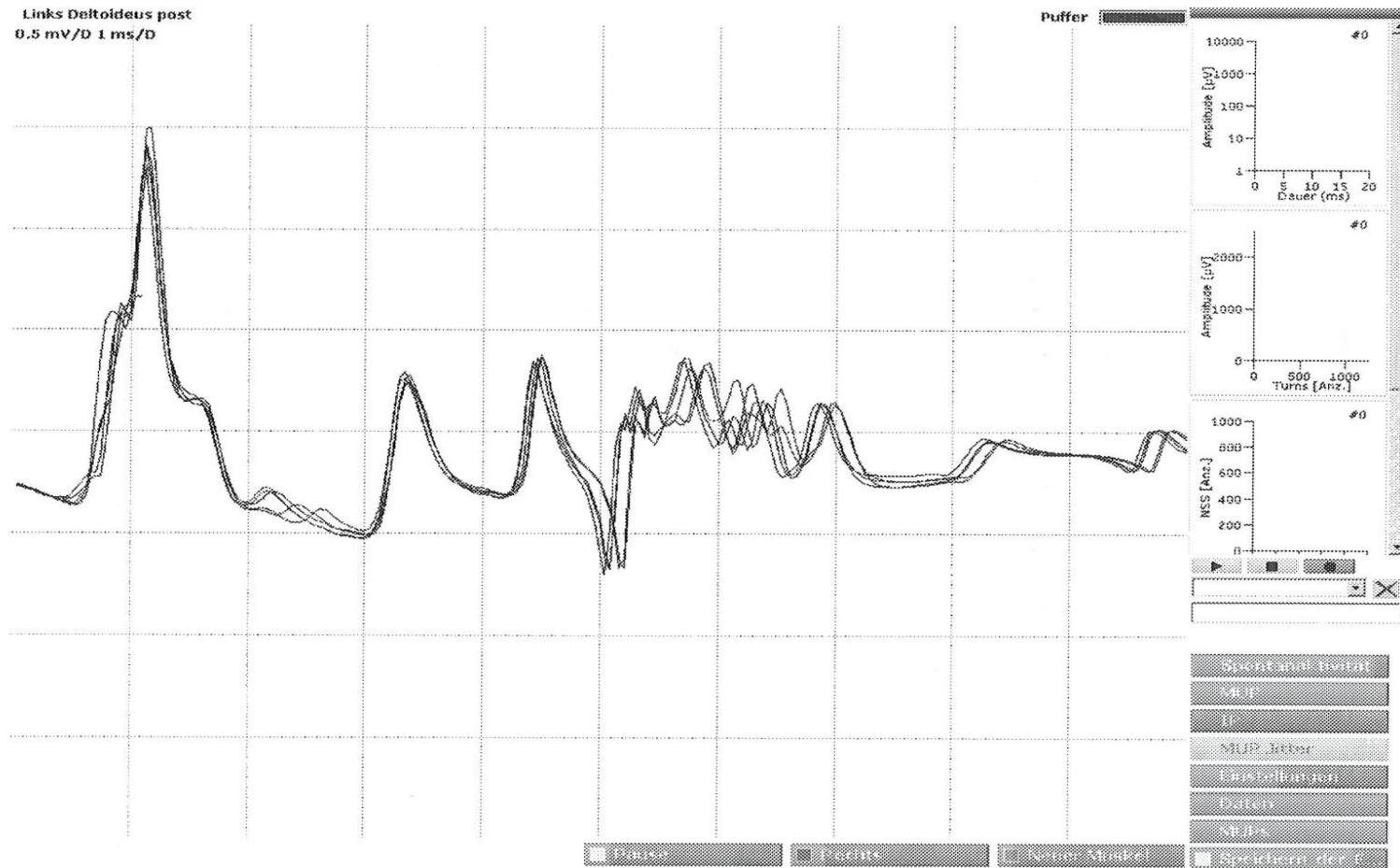
EMG changes in ALS: Fasciculations



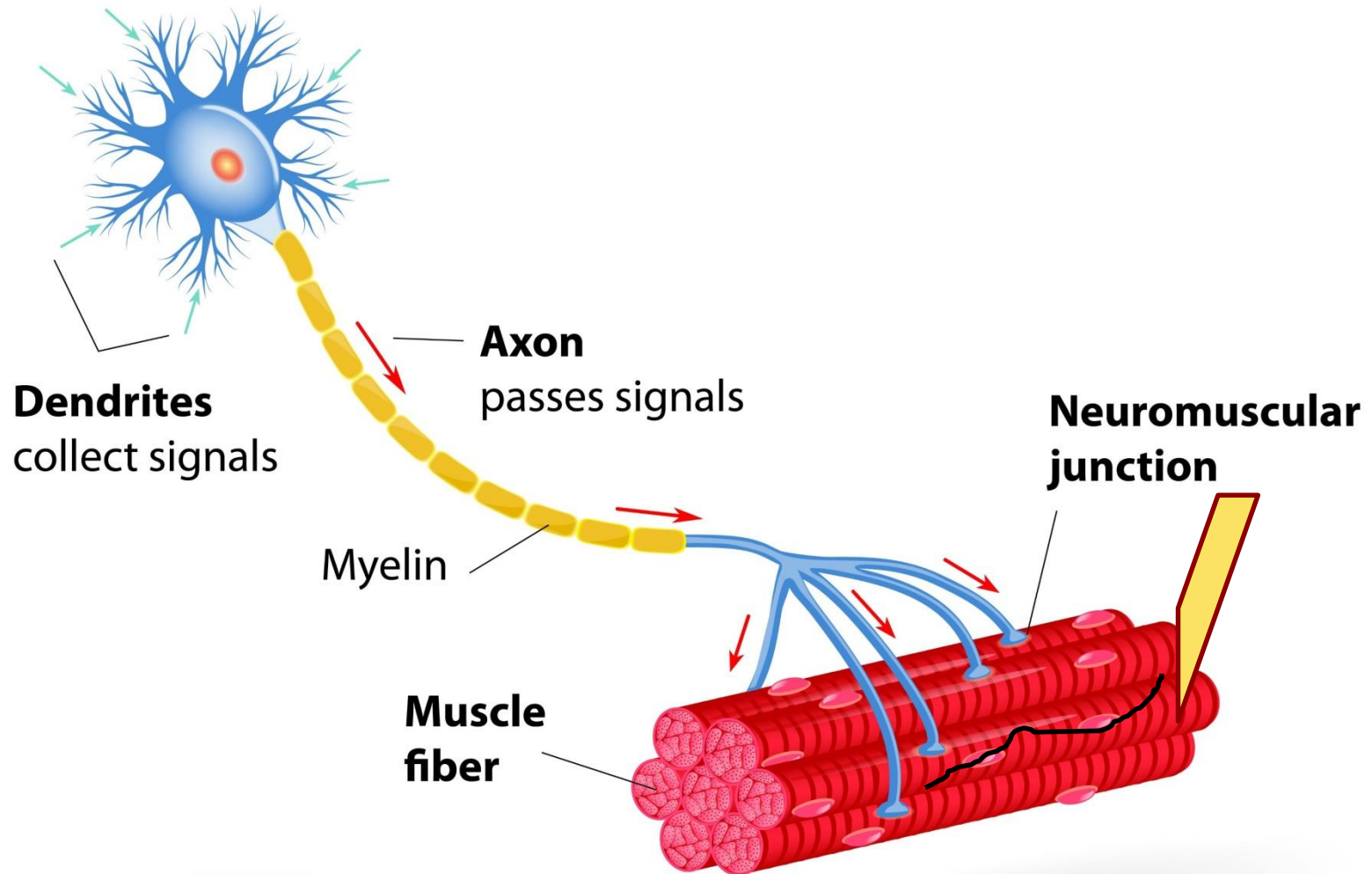
Clinical examples: Chronic inflammatory demyelinating neuropathy (CIDP)

- Demyelinating disease
- Autoimmune
- Secondary axonal loss
 - Very slowly
 - Lots of time to reinnervate
 - Very complex unstable MUPs

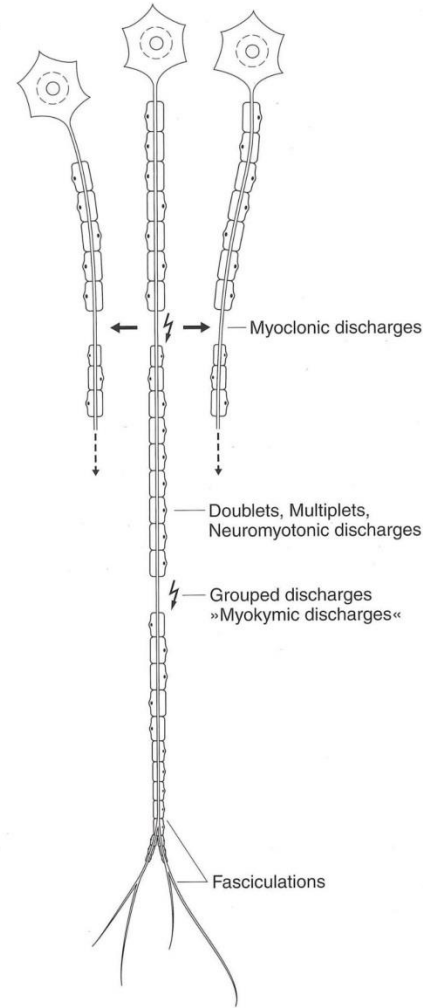
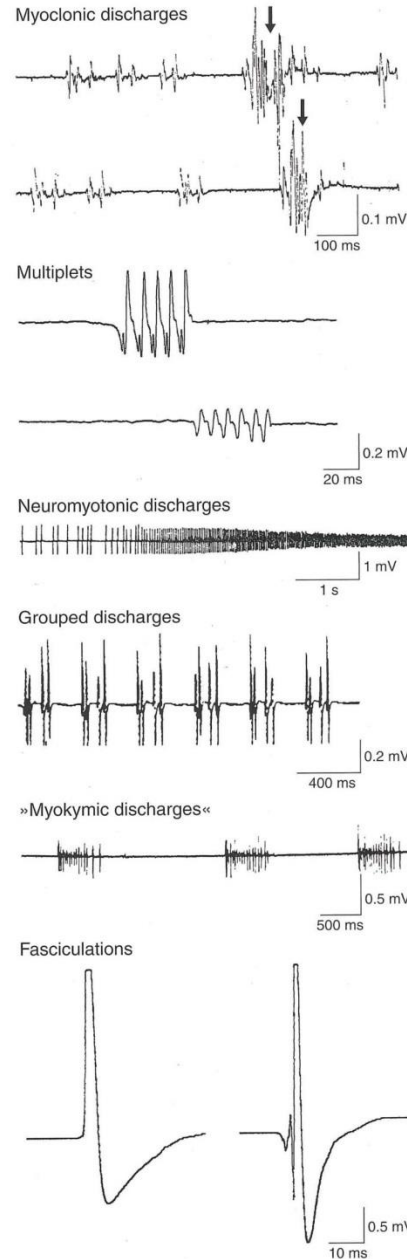
CIDP



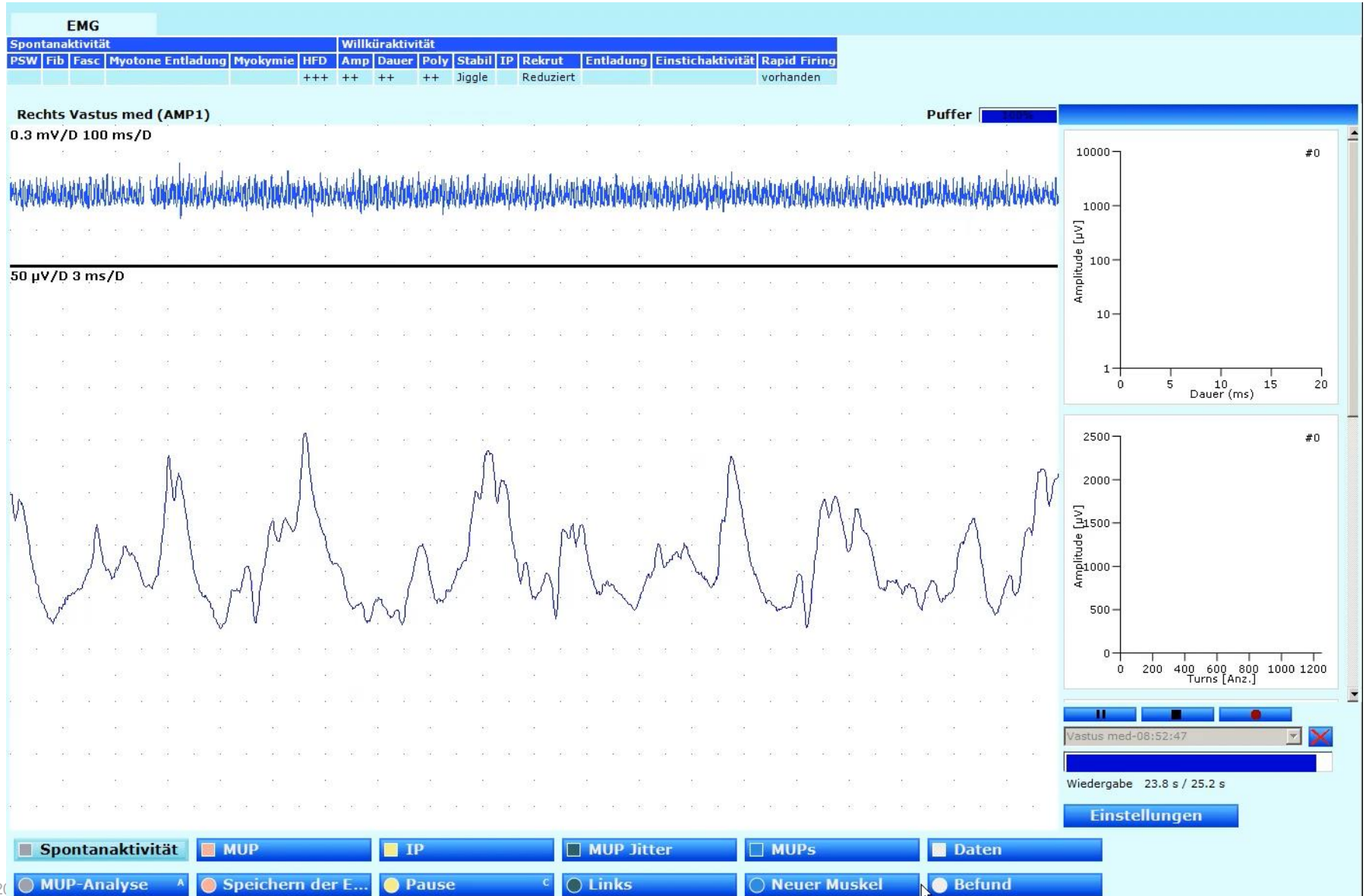
Signs of denervation in myopathy



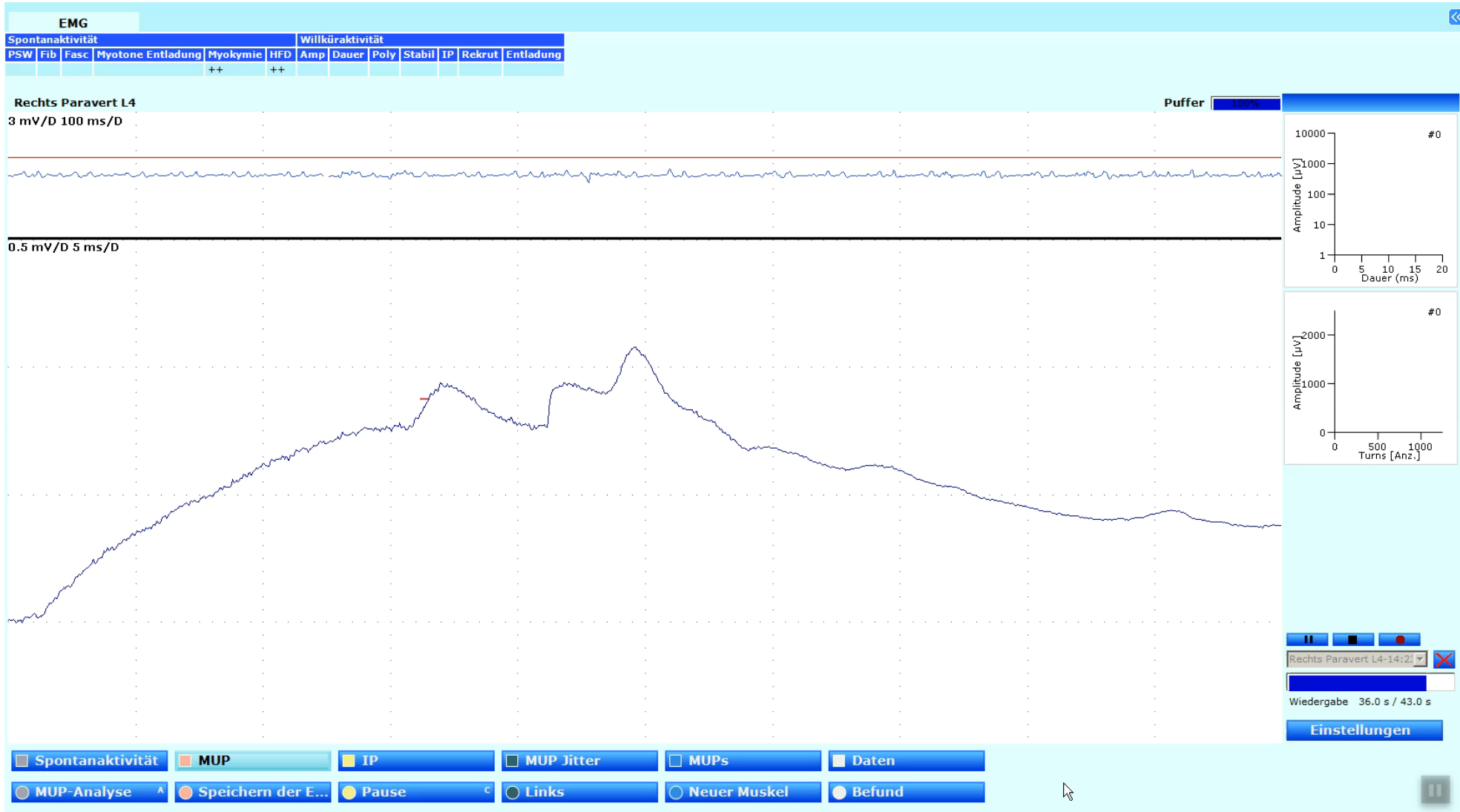
Other signs of hyperexcitability



Hyperexcitability



Hyperexcitability





Question 1

- Nascent reinnervation occurs within

A. Hours

B. Days

C. Weeks

D. Months

E. Years

Question 2

- Terminal sprouting occurs within

A. Hours

B. Days

C. Weeks

D. Months

E. Years

Question 3

- Which statement is **wrong**
- Signs of active denervation include:
 - A. Fibrillation potentials
 - B. Positive sharp waves
 - C. Fasciculation potentials
 - D. All of the above

Question 4

- Which statement is **wrong**
- With nascent reinnervation MUPs are
 - A. Complex
 - B. Unstable
 - C. Large
 - D. Fatigue easily

Question 5

- Which statement is **wrong**
- With terminal sprouting MUPs are
 - A. Complex
 - B. Stable
 - C. Normal to large
 - D. Show Jiggle
 - E. Long duration

Question 6

- The following statements relate to the triggered MU
 - What do you see ?
- A. A complex unstable MU
- B. A normal MU
- C. Satellite potential
- D. All 3

Question 7

- The following statements relate to the recorded potentials.
 - Which statement is **wrong** ?
- A. All potentials are fasciculation potentials
- B. They might occur in demyelinating diseases (e.g. CIDP)
- C. They are typical in ALS patients
- D. Fasciculation potentials are always abnormal

Question 8

- The following MUP suggests
 - A. An acute complete nerve lesion
 - B. An acute partial nerve lesion
 - C. An intermediate stage of reinnervation
 - D. A very chronic disease

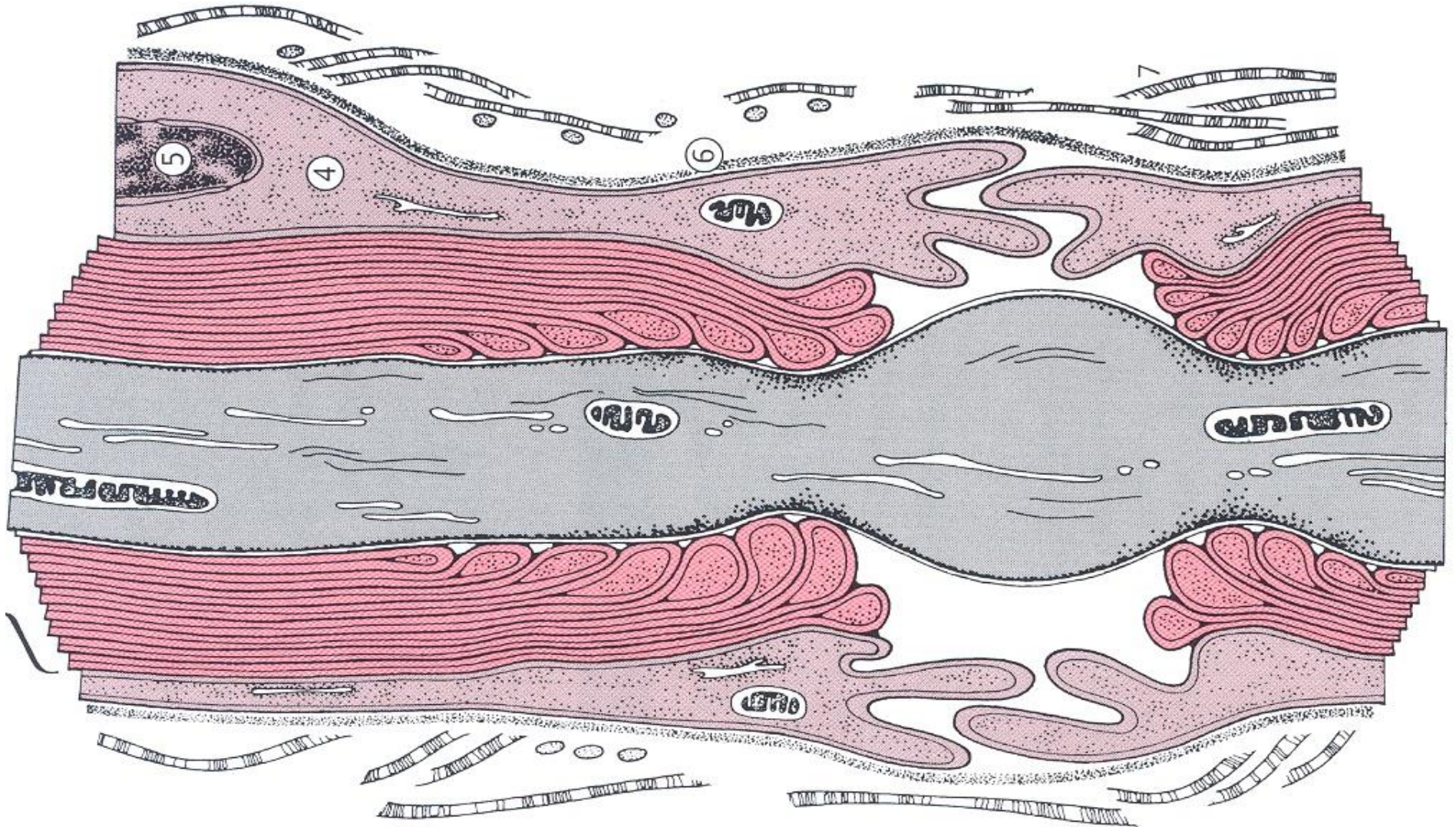
Question 9

- The following recording suggests which etiology
 - A. Complete nerve cut
 - B. Partial nerve lesion
 - C. Radiation
 - D. Chronic nerve compression

Question 10

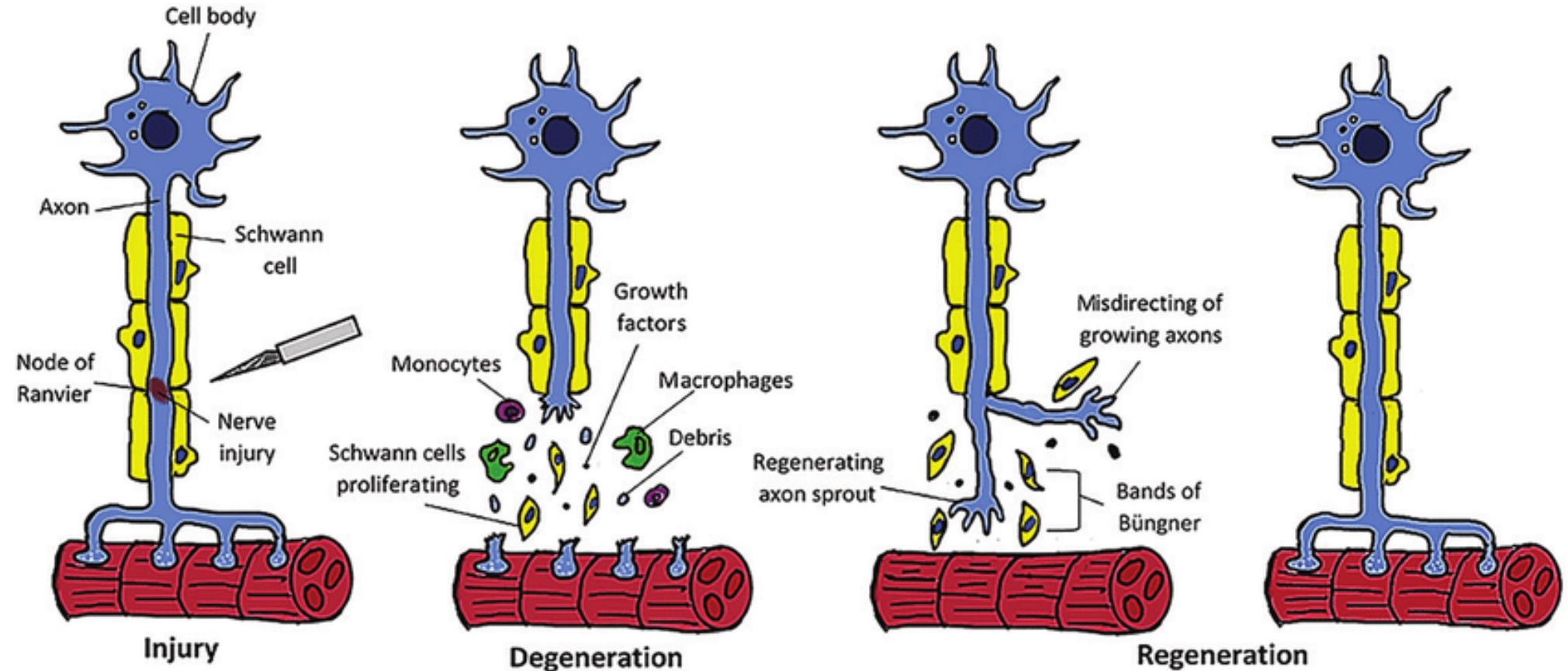
- The duration of the recorded MUP is (sweep speed 8 ms/Div)
 - A. 8 ms
 - B. 16 ms
 - C. 32 ms
 - D. More than 32 ms

Nerve longitudinal



Pathophysiology of denervation and reinnervation

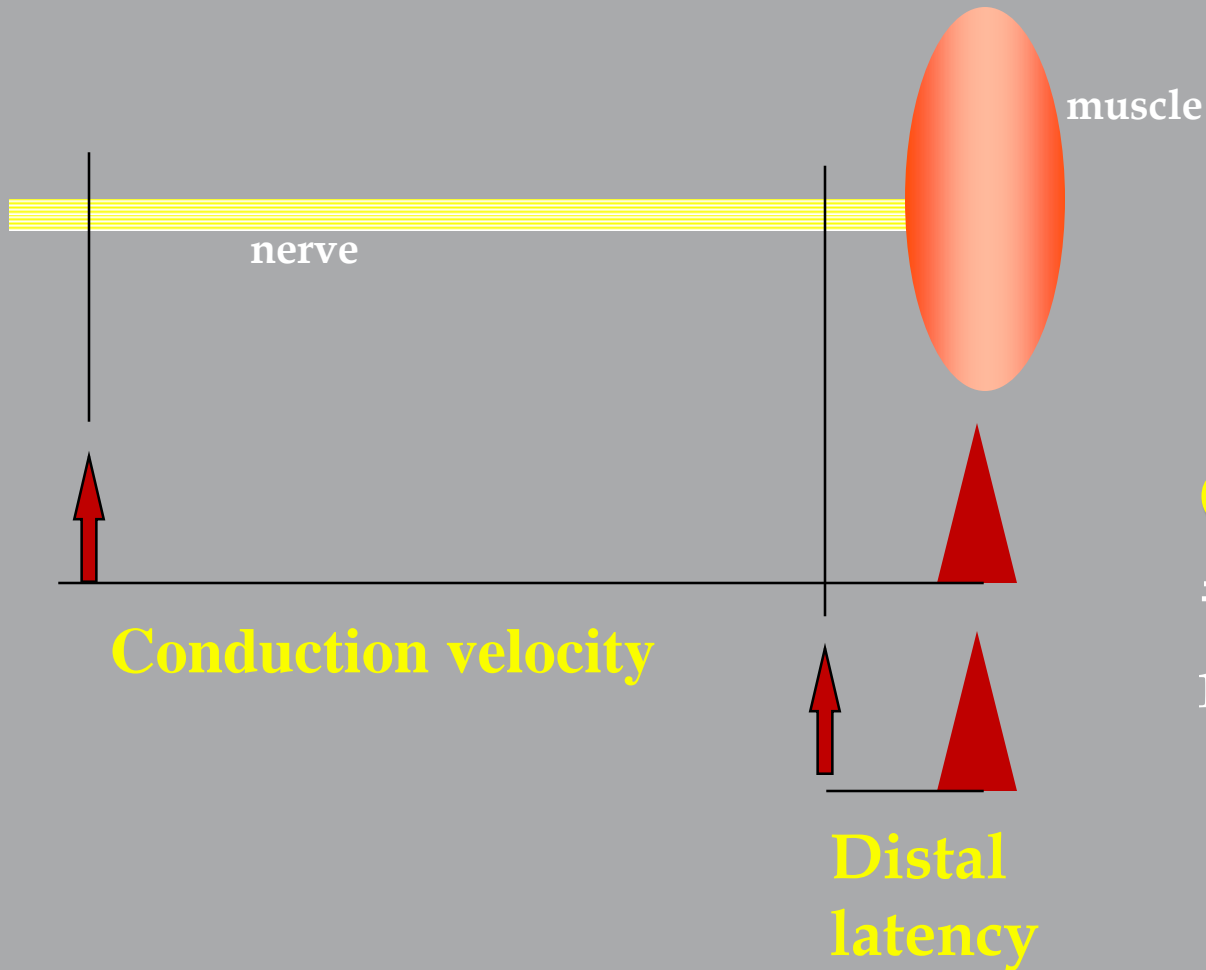
Wallerian degeneration



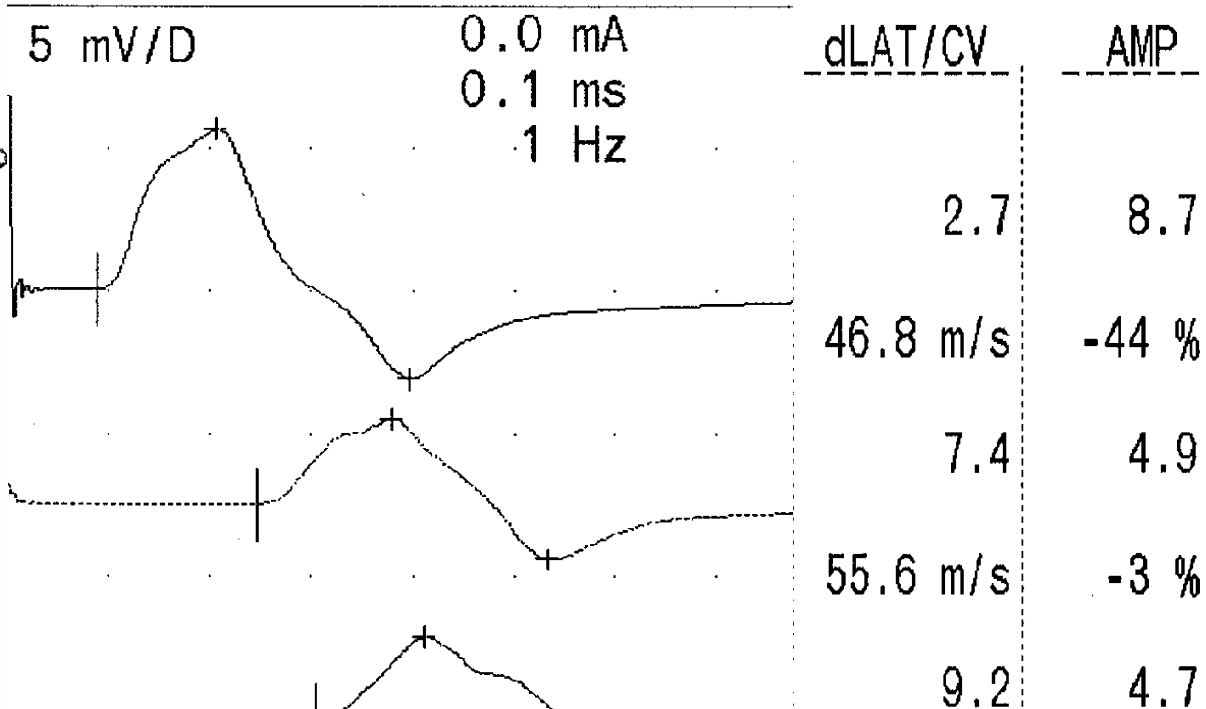
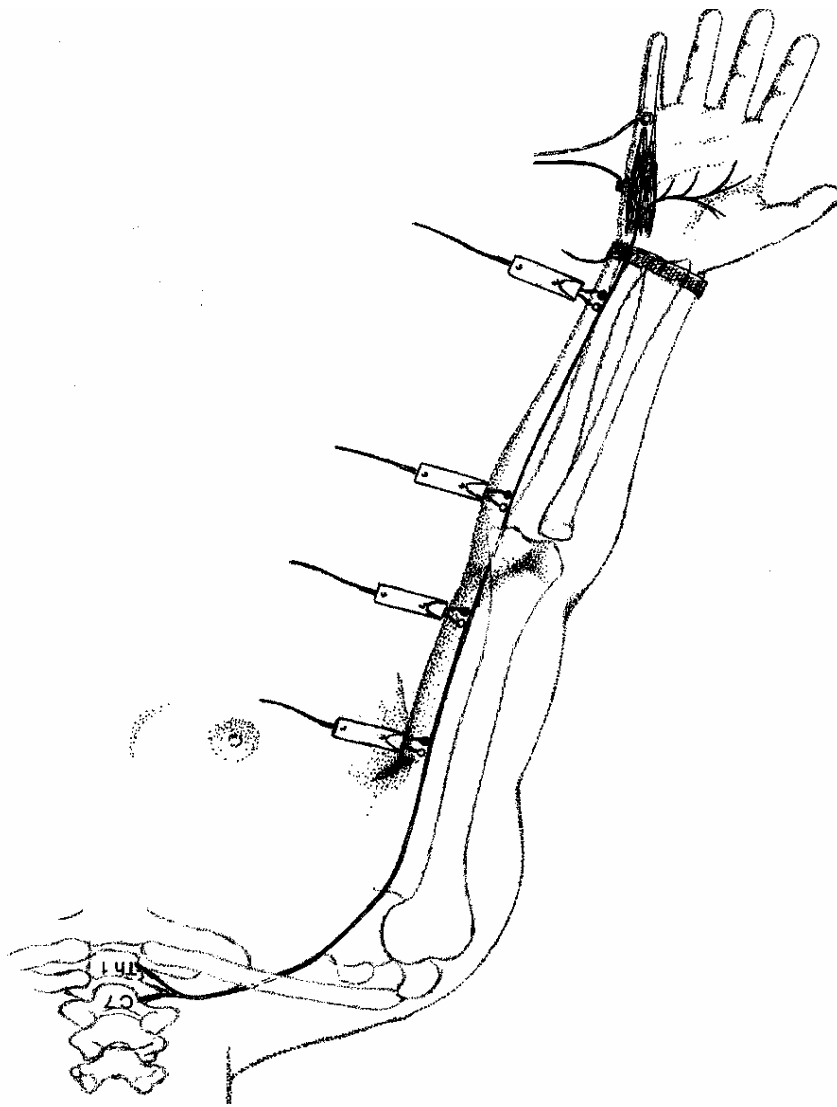
- DOI:10.5772/intechopen.68174 Alvites et al.
- In book: Mesenchymal Stem Cells - Isolation, Characterization and Applications



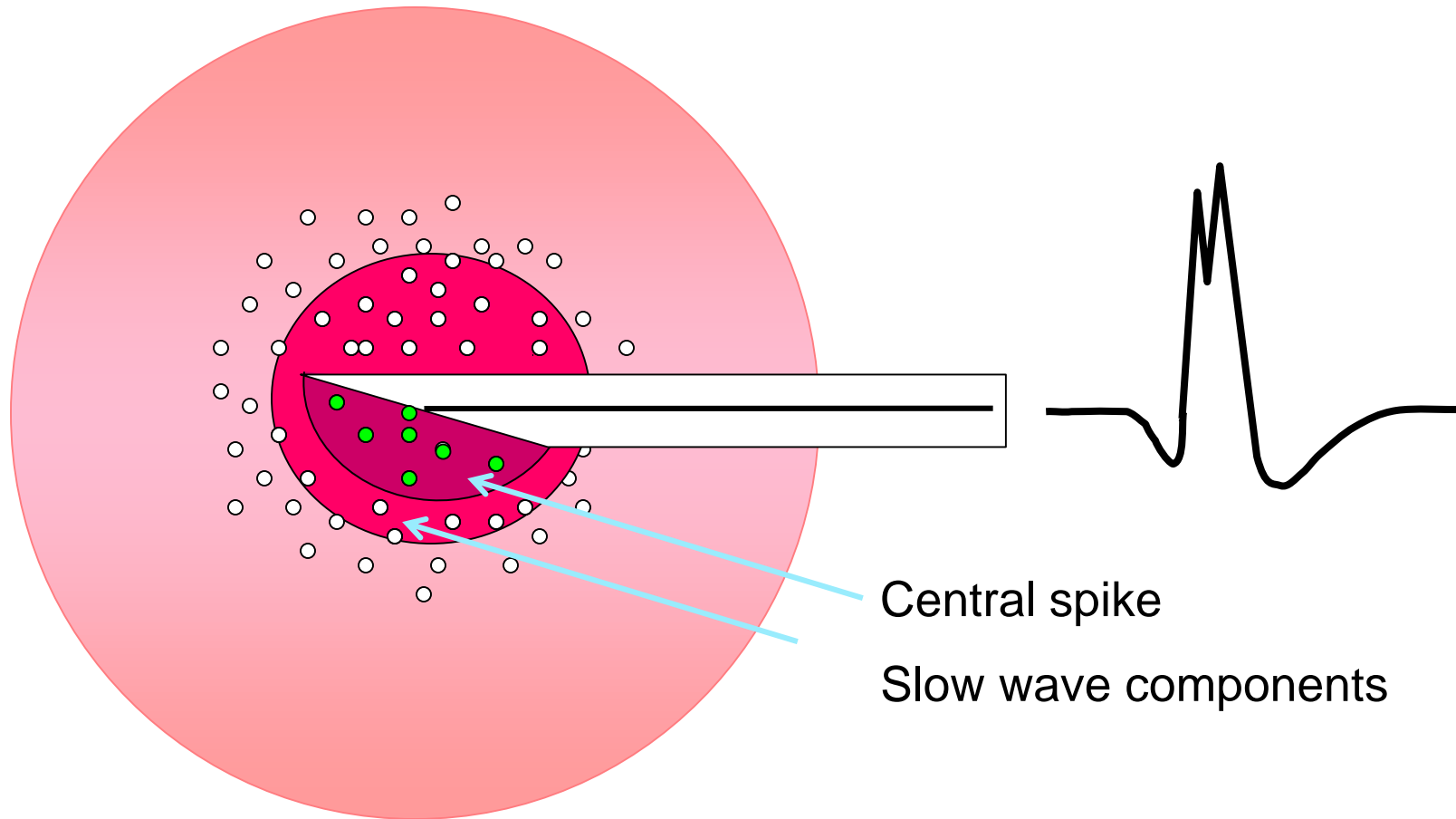
Motor nerve conduction study



Motor nerve conduction study and compound muscle action potential (CMAP)



Electromyography with concentric needle: signals from 2-15 muscle fibres



Spontaneous activity in normal

- insertional activity
- end-plate noise
- "nerve spikes"
- positive wave at end-plate zone