



Standards and Clinical Applications of Visual, Auditory and Somatosensory Evoked Potentials

- § Technical basics
 - Components of the system
 - Averager
 - Nomenclature and normal values
- § Neurophysics of the generation of evoked potentials
 - Intra-extra cellular currents
 - Electrical fields
 - Generation of far-field potentials
- § Visual potentials / Auditory potentials / Somatosensory potentials
 - Anatomy and physiology of the system
 - Stimulus
 - Recommended standard protocols
 - Clinical applications and artifacts

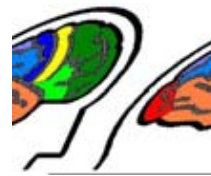
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www.richardjungkolleg.de

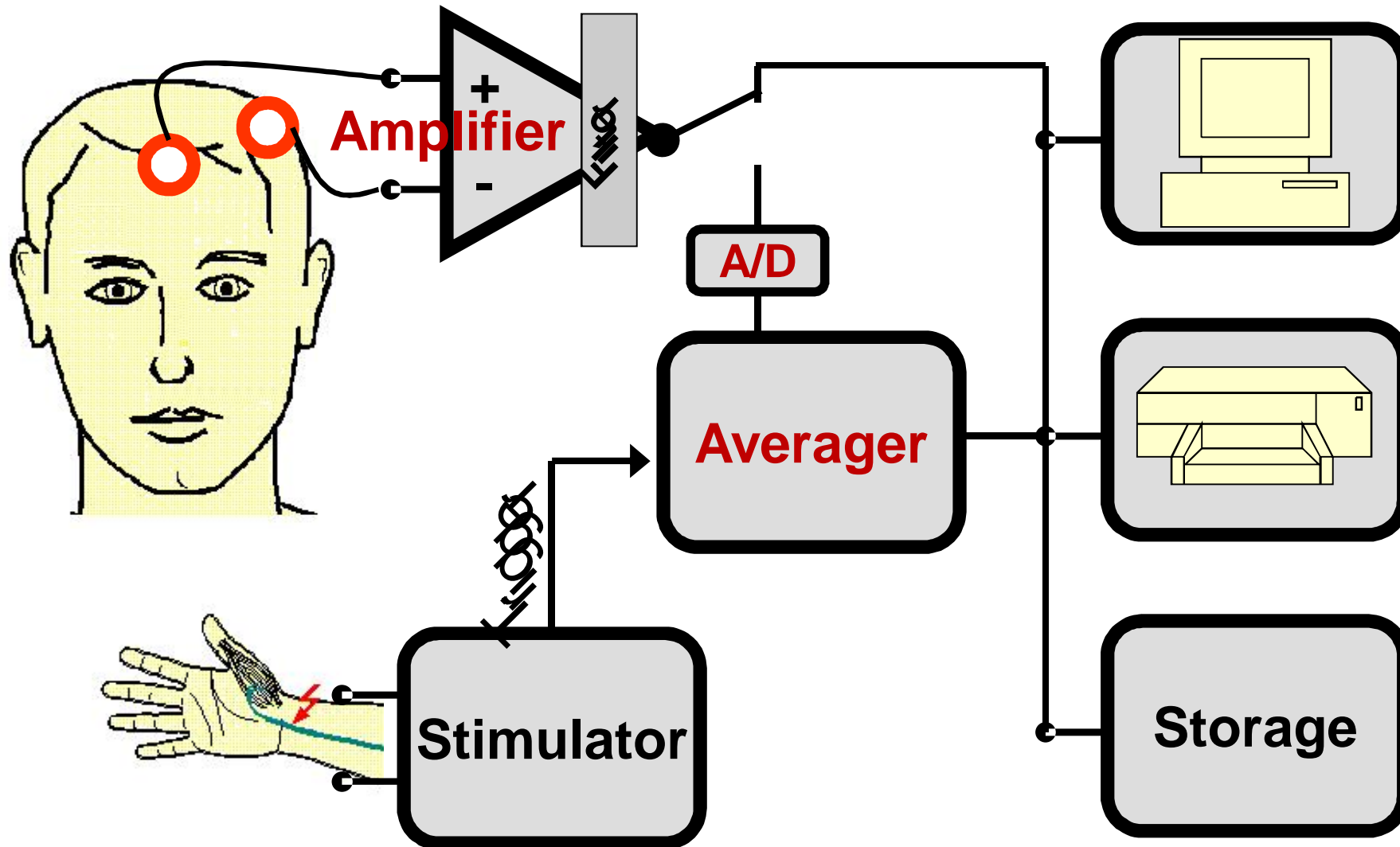


Evoked Potentials are electric potentials occurring in the peripheral and the central nervous system time locked to a natural or artificial sensory stimulus.

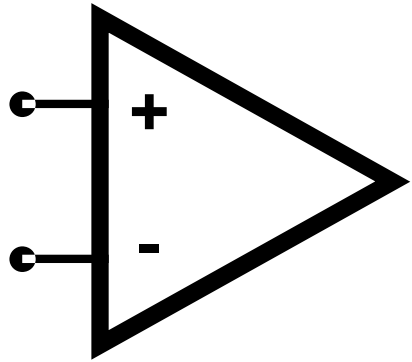
I have nothing to disclose



§ Components of the system



§ Averager



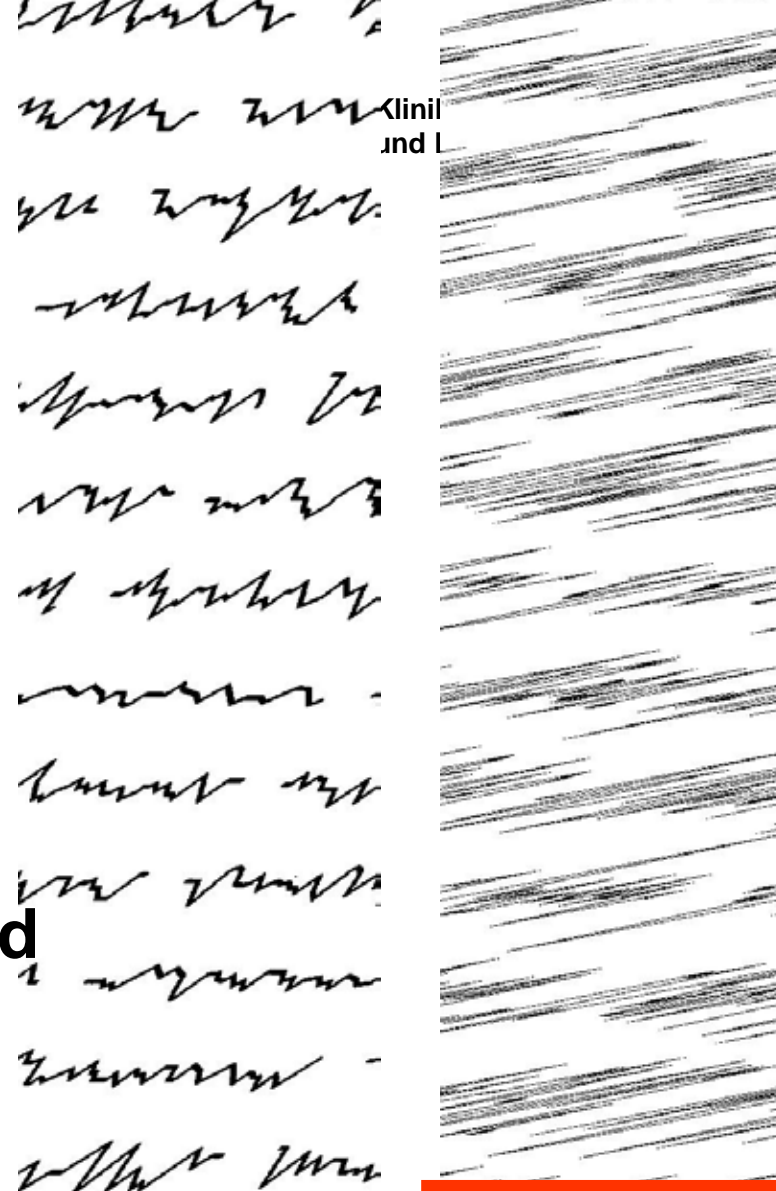
Prerequisites:

- Signal Stimulus time-locked
- Noise random

$$\frac{1+2+3+\dots+16}{16}$$

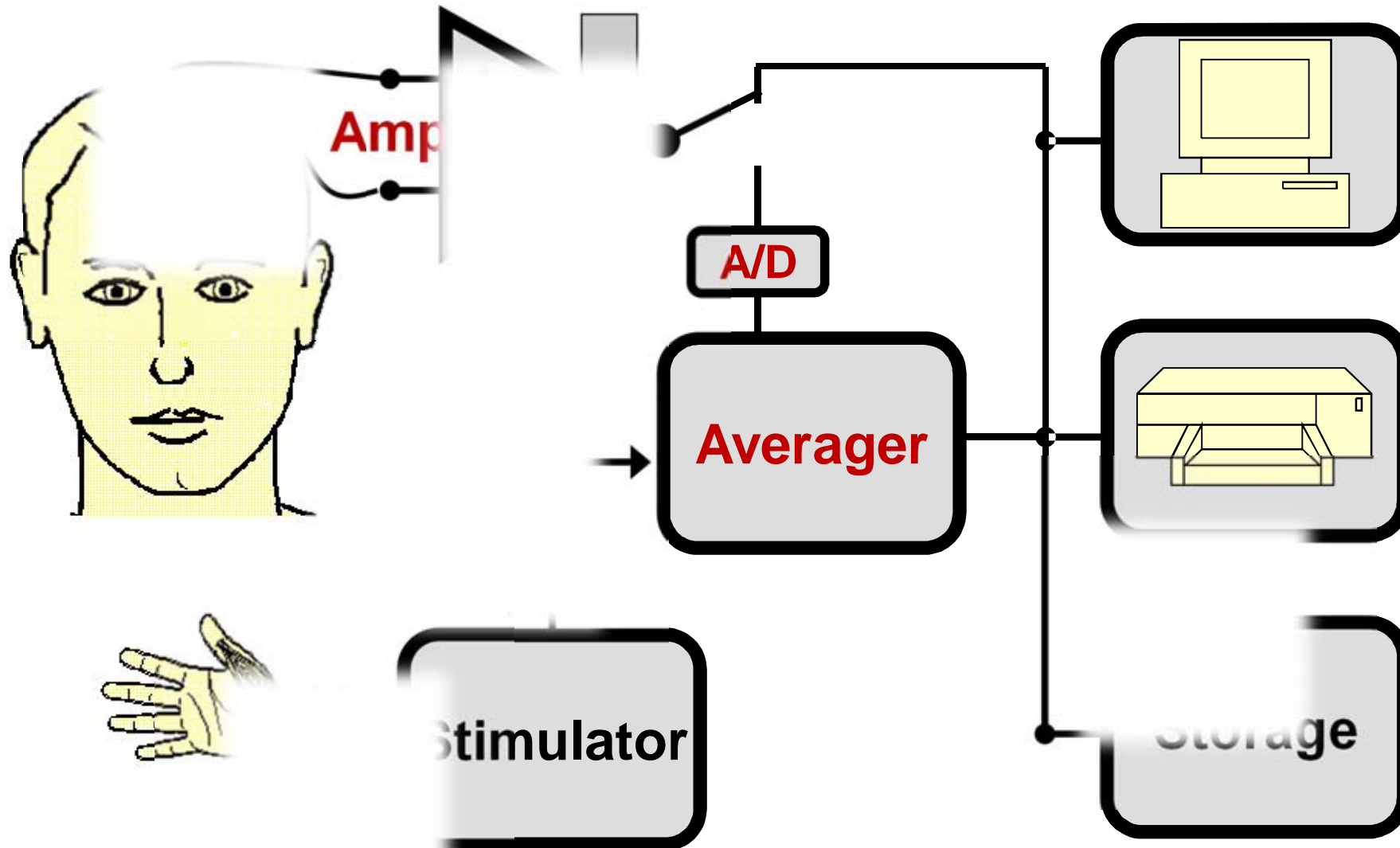
16

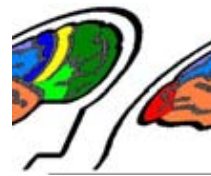
$$S/N \sim \frac{1}{\sqrt{16}}$$



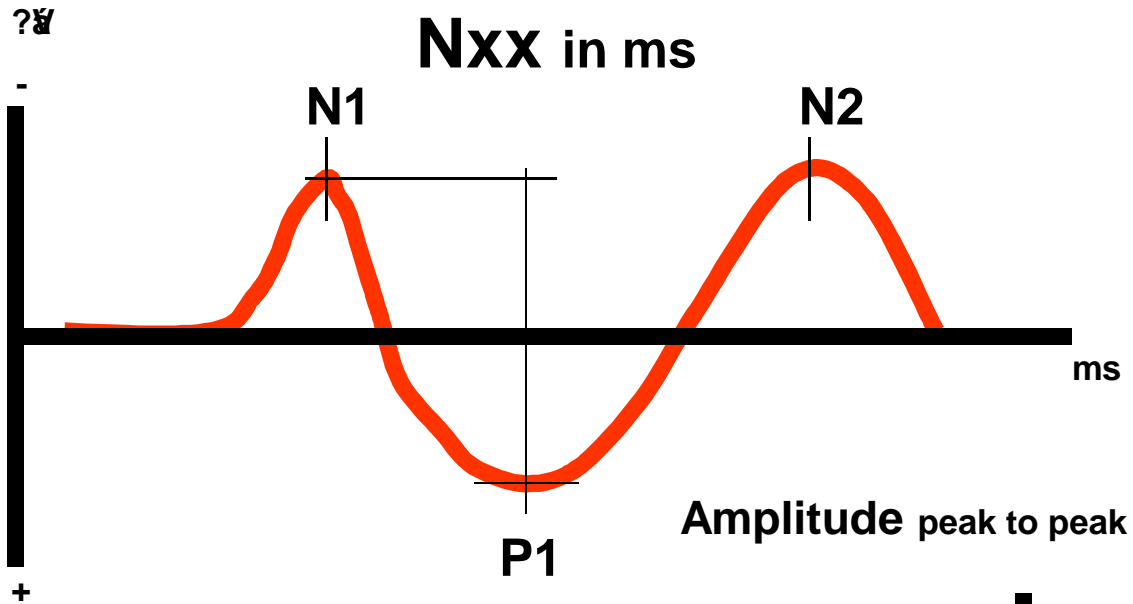


§ Check of the system

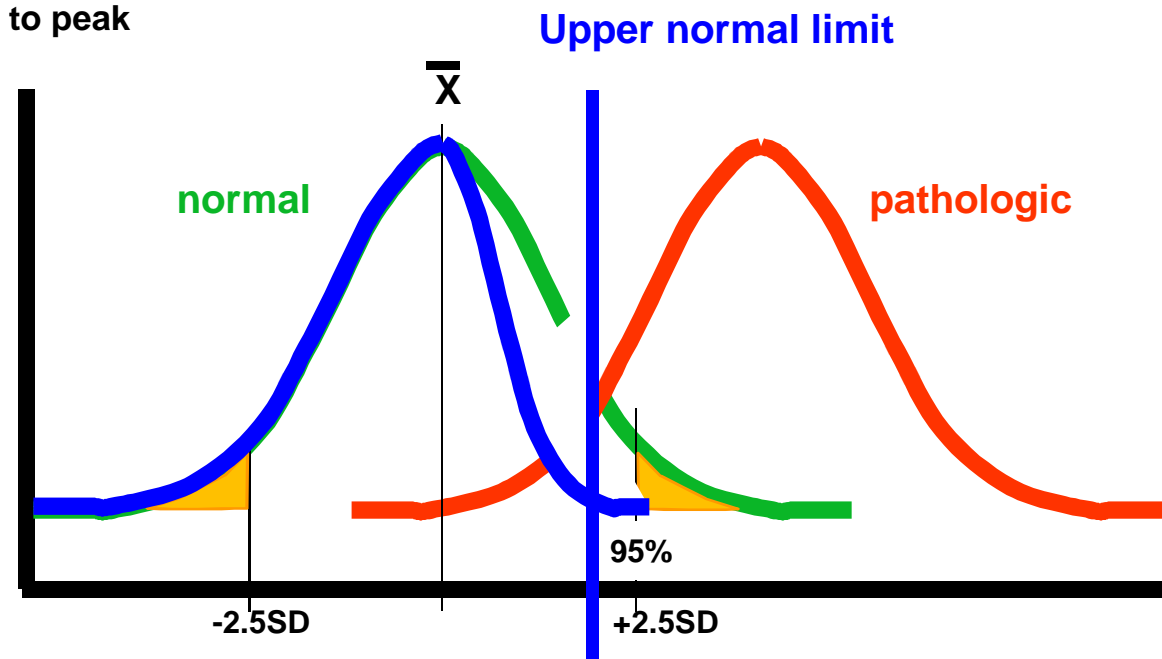




§ Nomenclature



§ Normal values

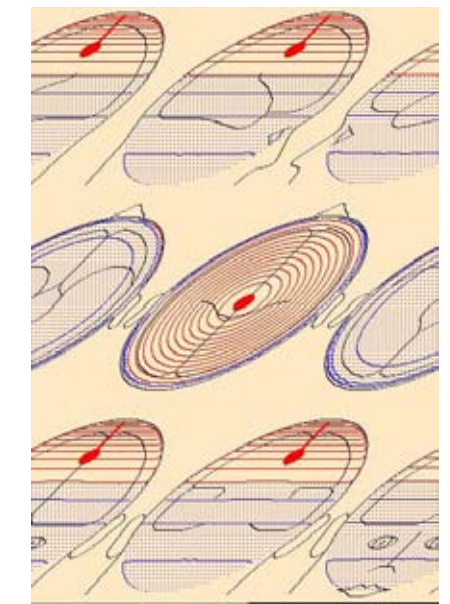
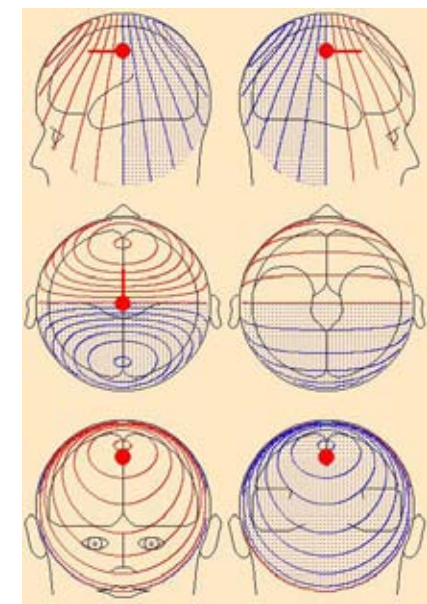
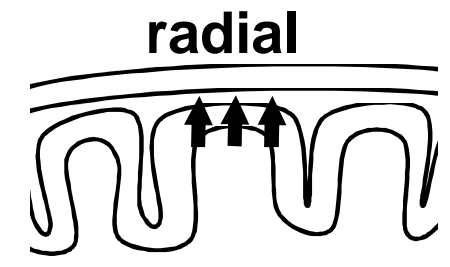
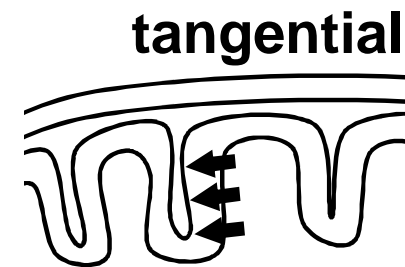
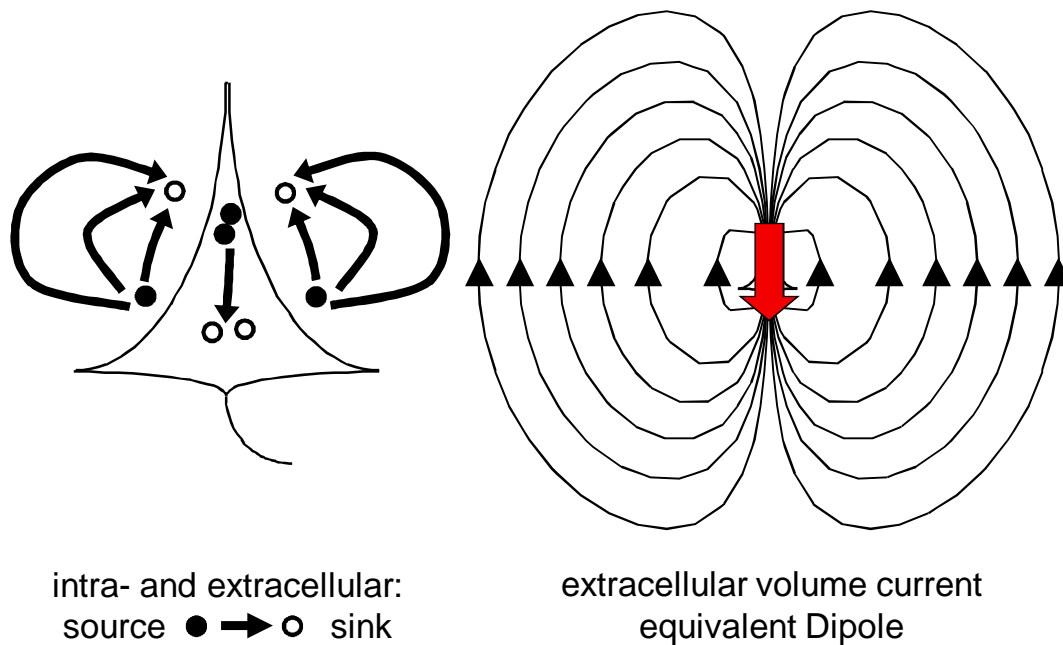


§ Generation of evoked potentials

für Neurologie
nische Neurophysiologie

§ Intra-extra cellular currents

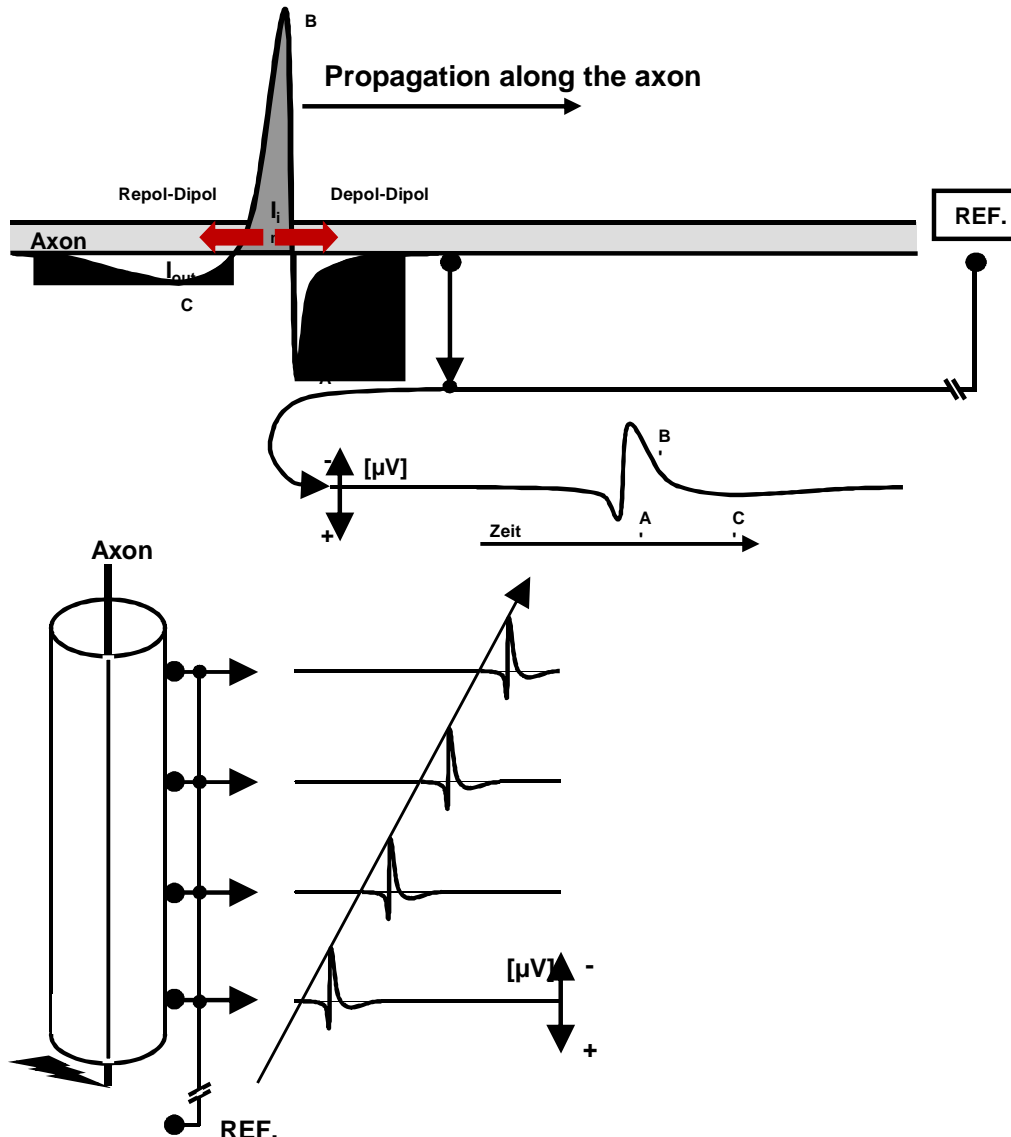
§ near-fields



DipoleSimulator © P. Berg (2002)

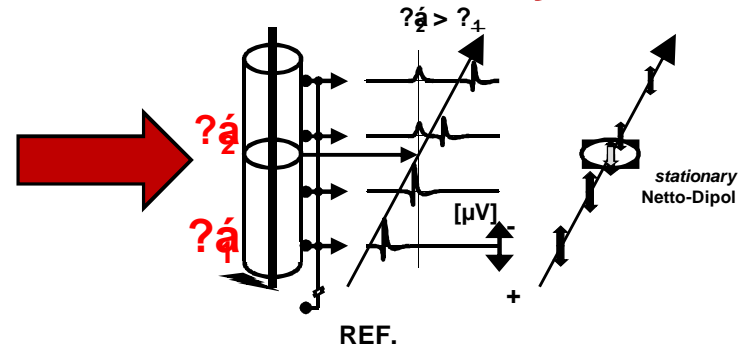
§ Generation of evoked potentials

§ Generation of far-field potentials

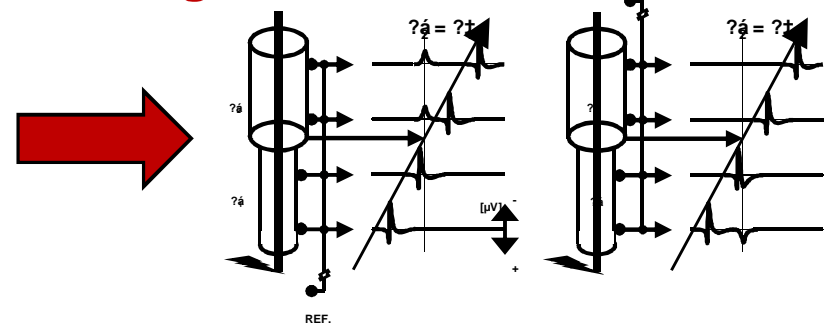


Curio G. Praxisbuch Evozierte Potenziale Thieme 2014

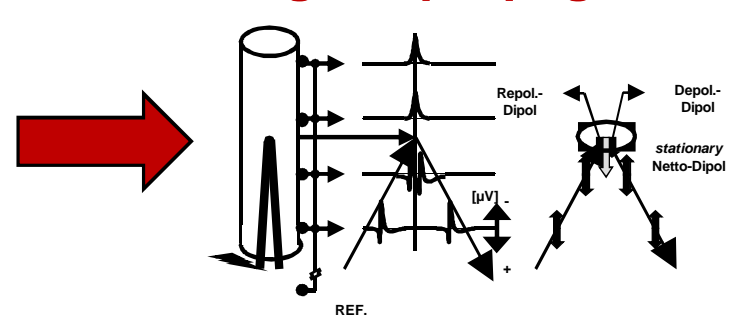
conductivity



geometrie of volume conductor



change in propagation direction



§ Visual evoked potentials

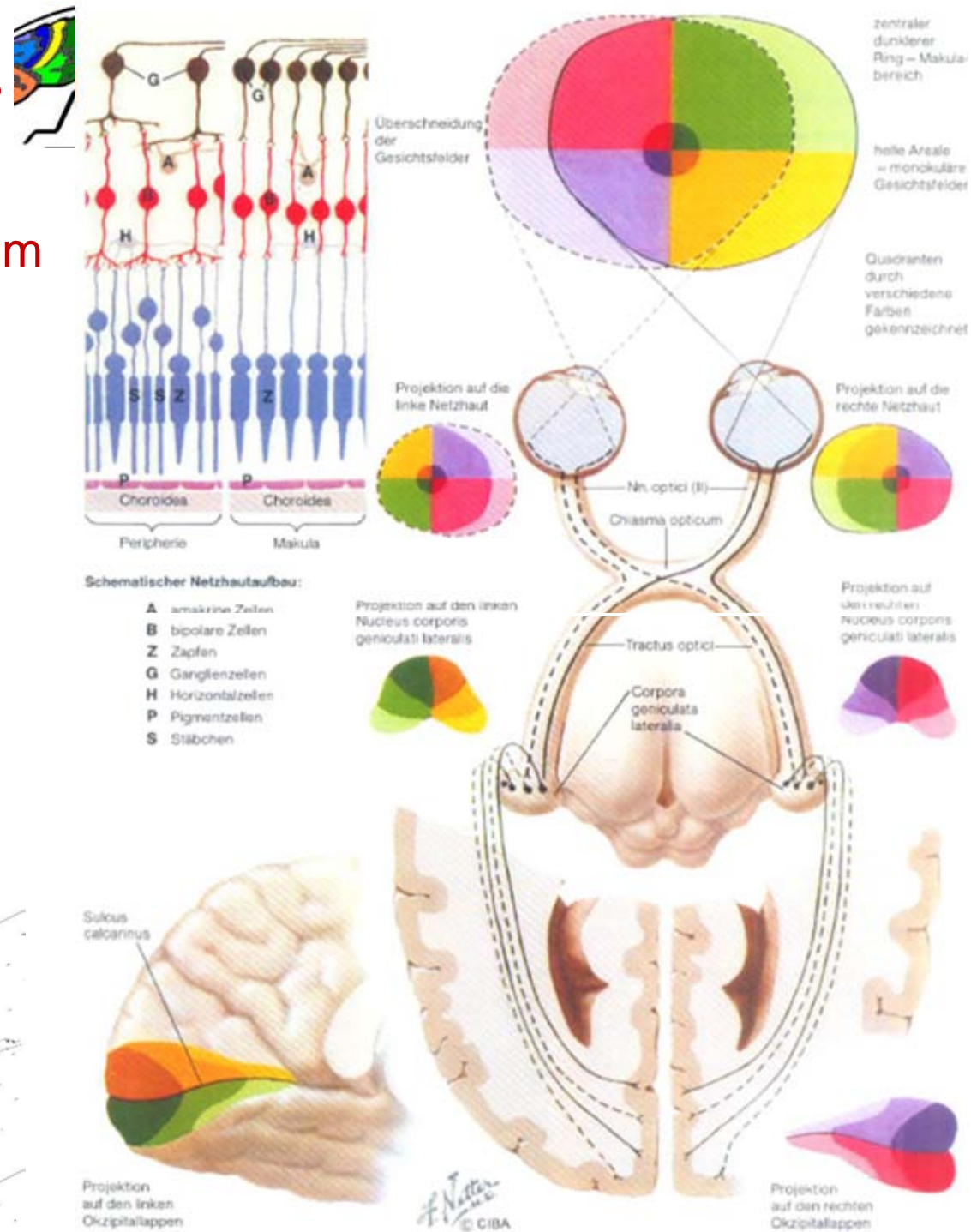
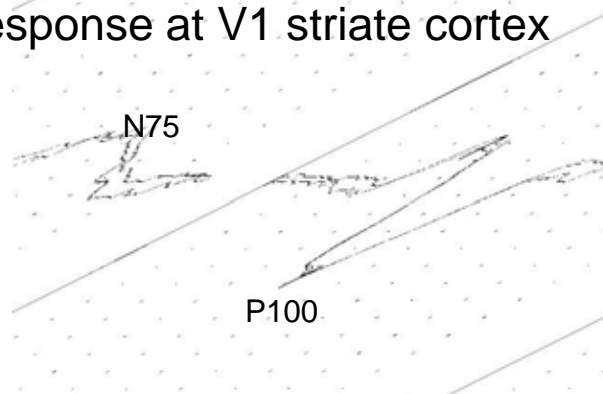
§ Anatomy and physiology of the system

- Stimulus
- Recommended standard protocols and sources of errors
- Clinical applications and advanced use

- Cones – color sensitive, fovea
- Rods – light sensitive, outside fovea
- Retinal ganglion cells
 - magnocellular – direction of contrast and motion
 - parvocellular - color

N75 – input from LGN to striate cortex

P100 – secondary response at V1 striate cortex



§ Visual evoked potentials

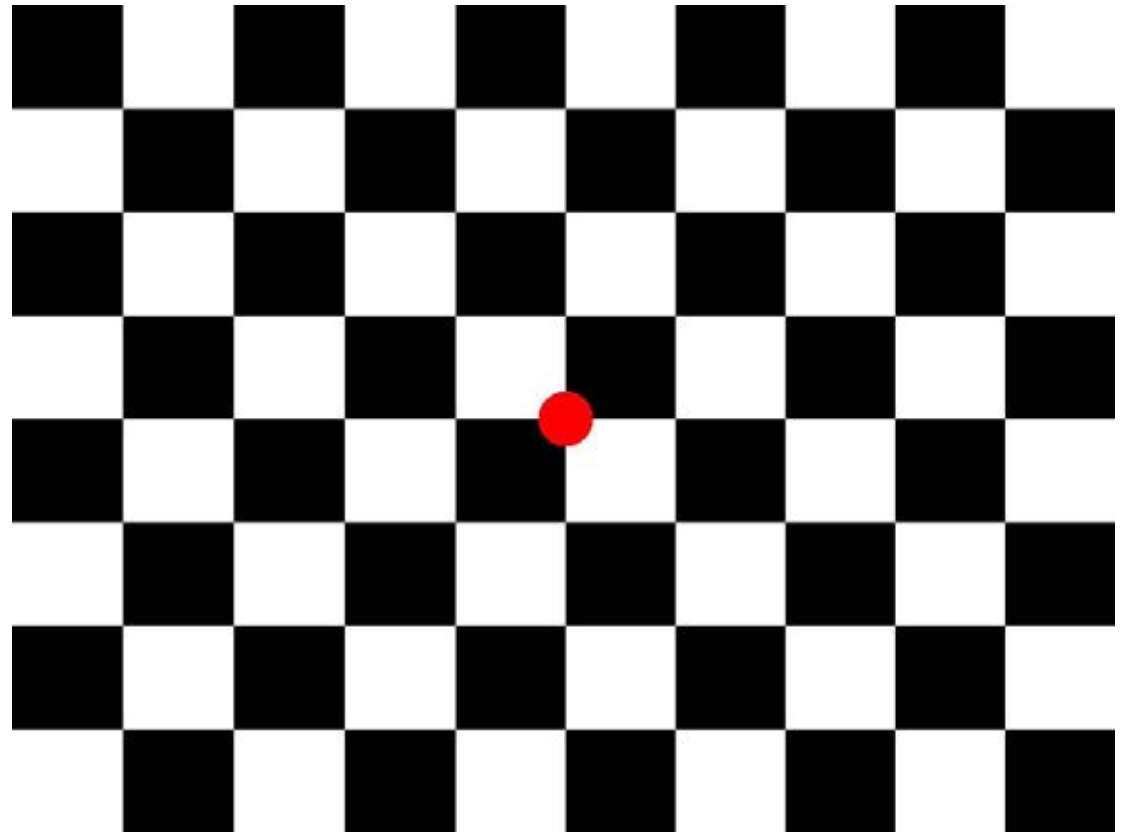


∅ Pattern reversal

§ Stimulus

- Recommended standard protocols and sources of errors
- Clinical applications and advanced use

- One eye full-field stimulation
- Stimulus field $\geq 12-15^\circ$
- Check size (15'/50-60')
- **Contrast black/white >80%**
- Reversal rate 1-2 Hz
- Fixation: central
- app. 80% of VEP from central 8 dg. of visual field

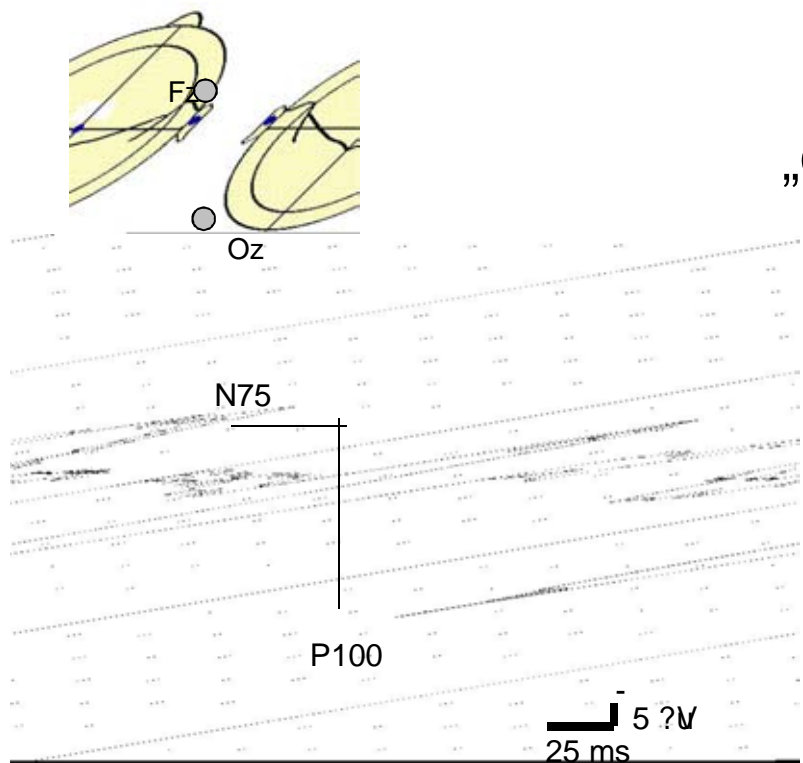


§ Visual evoked potentials

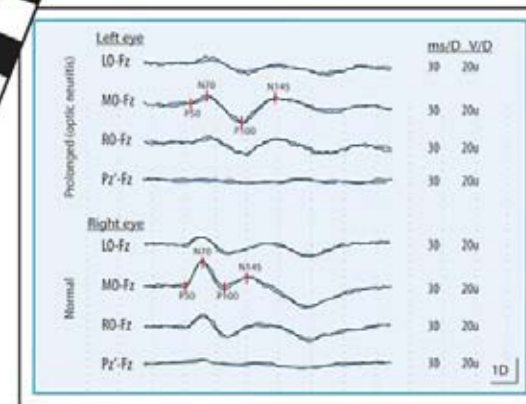
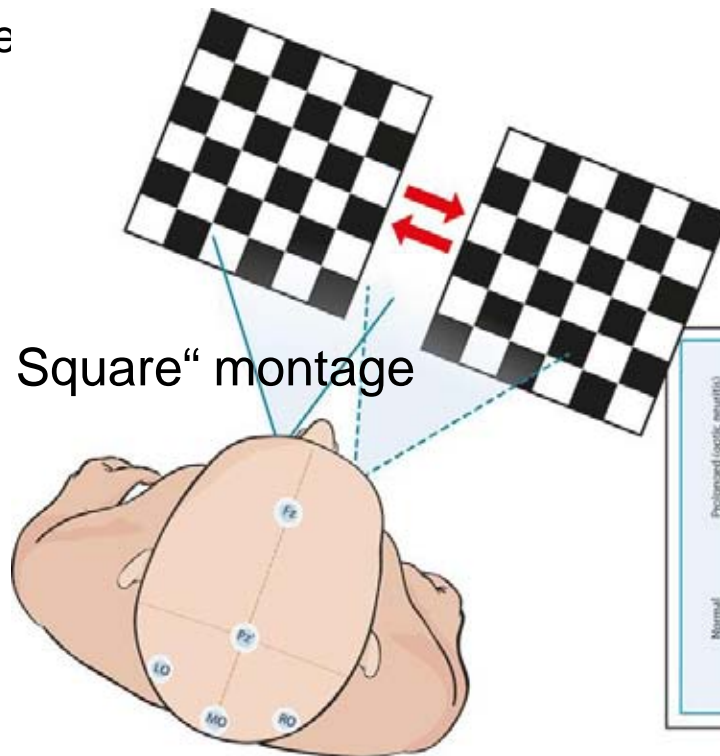


§ Recommended standard protocols and sources of errors

- Clinical applications and advanced use



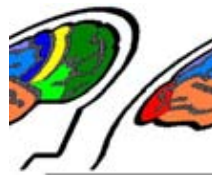
„Queen Square“ montage



quality control

- § two measurements
- § reproduction with:
- § Latencies below 1 ms
- § Amplitudes of + / - 20%

§ Visual evoked potentials



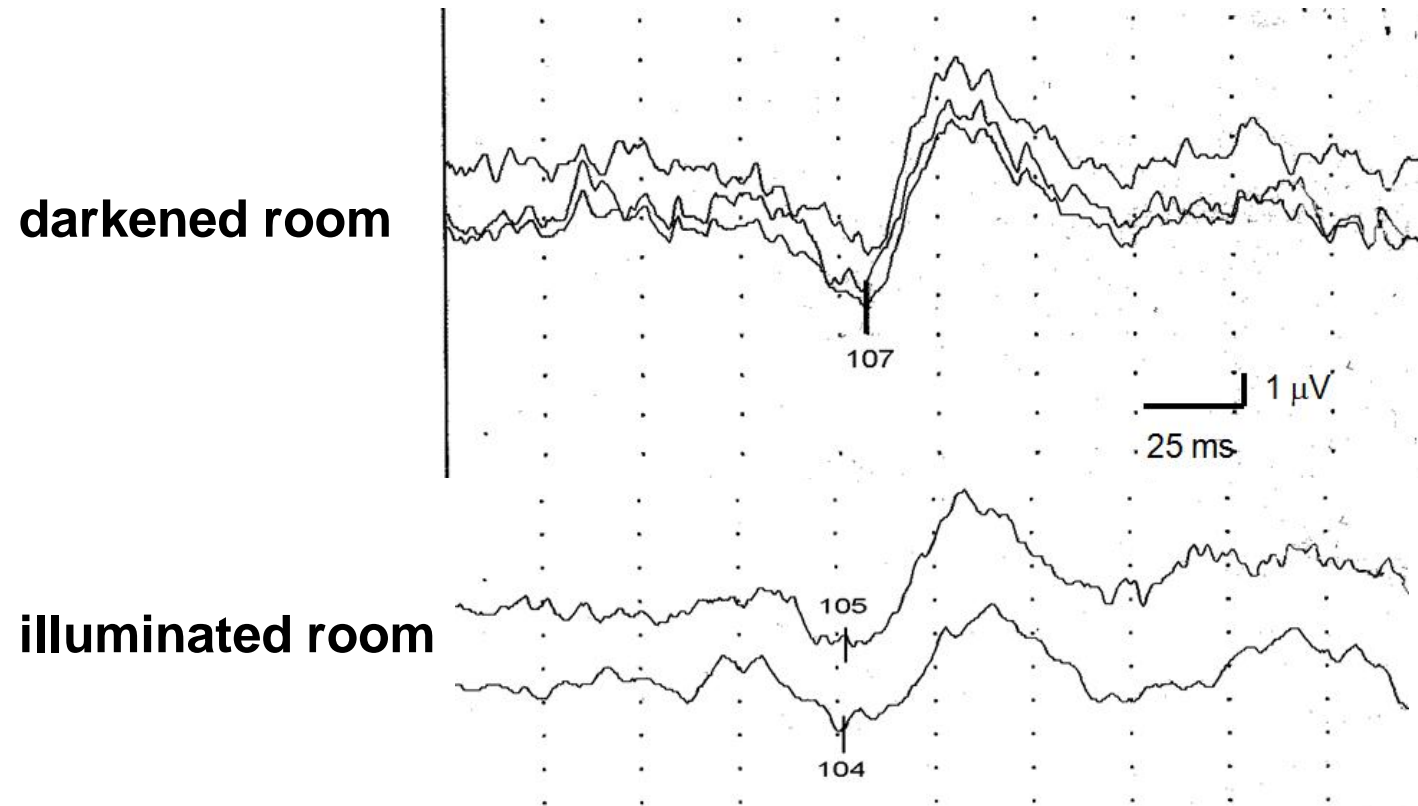
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und klinische Neurophysiologie



§ Sources of errors

- Stimulus parameters

- Contrast black/white > 80%



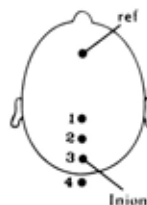
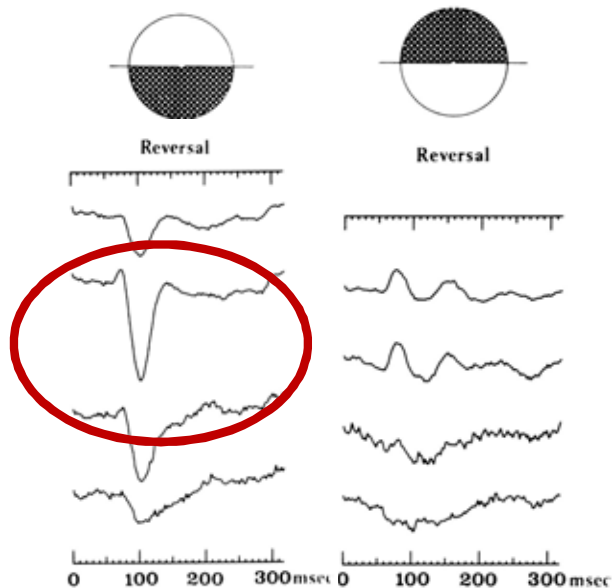
§ Visual evoked potentials



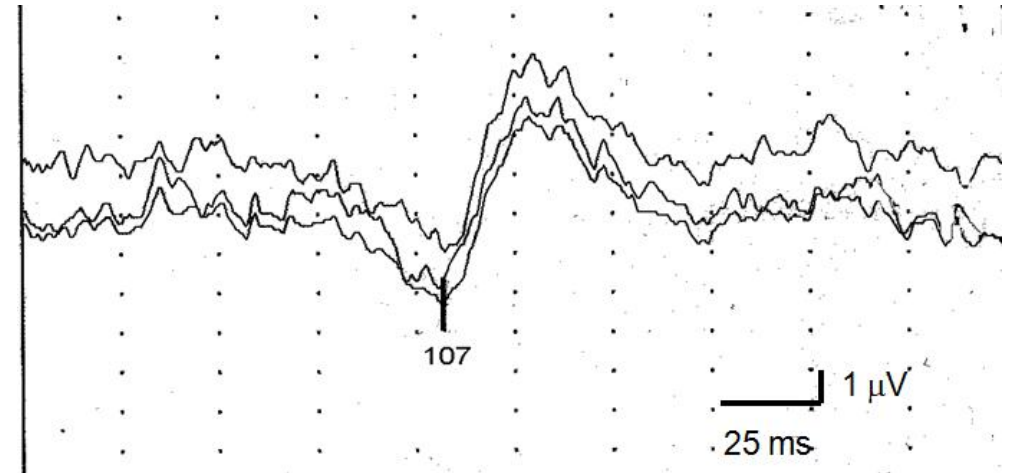
§ Sources of errors

– Fixation of stimulus

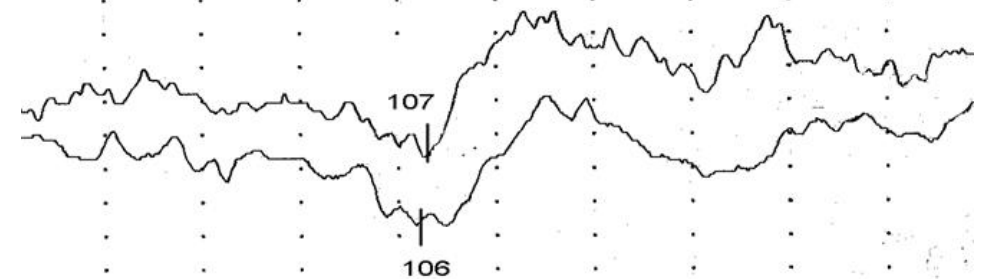
• **Fixation: central**



attending



inattentive



• **Minor influence**

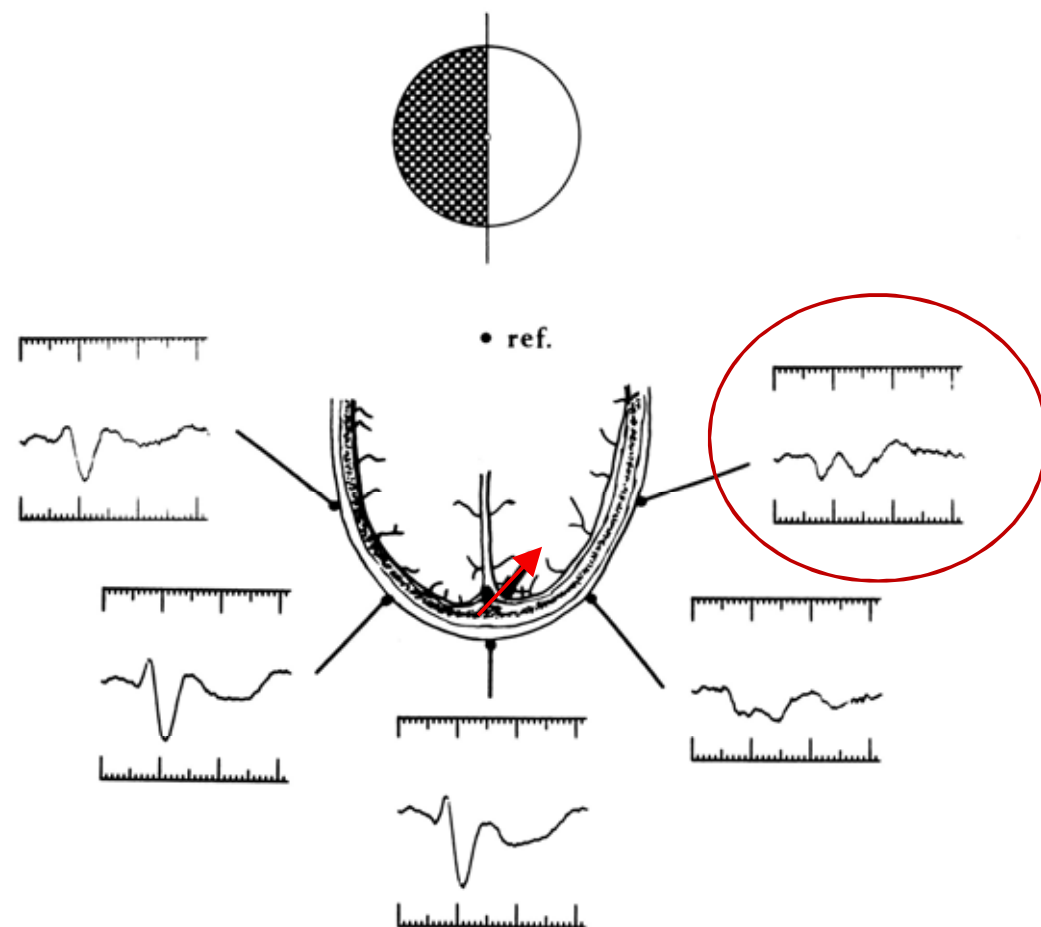
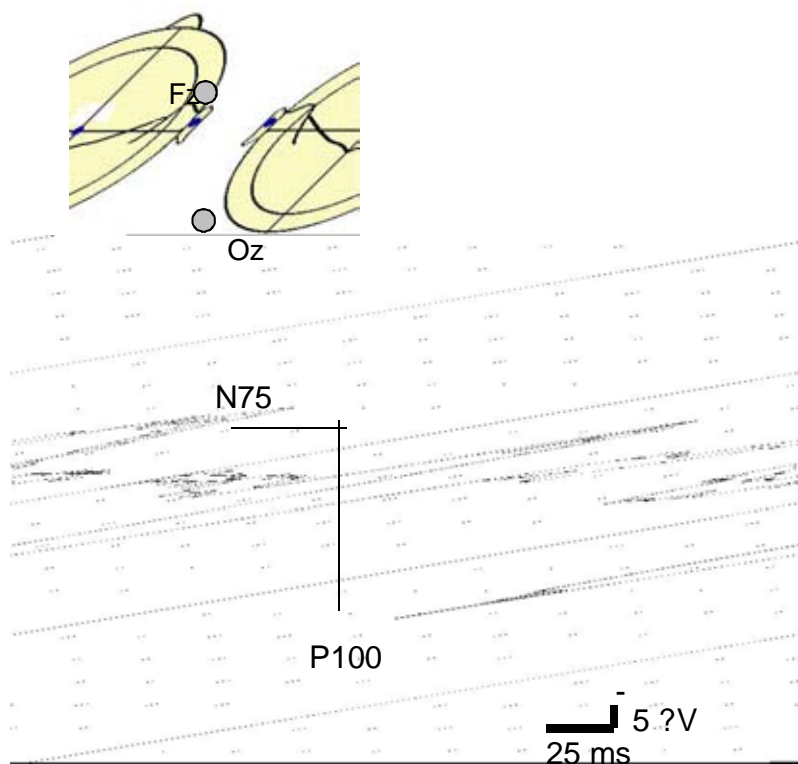
- Age / Gender
- Head Size
- Pupil Size
- Temperature

§ Visual evoked potentials



§ Normal and pathological VEPs

- P100 identification
- P100 latency
- P100 amplitude



Barrett et al., 1976

§ Visual evoked potentials

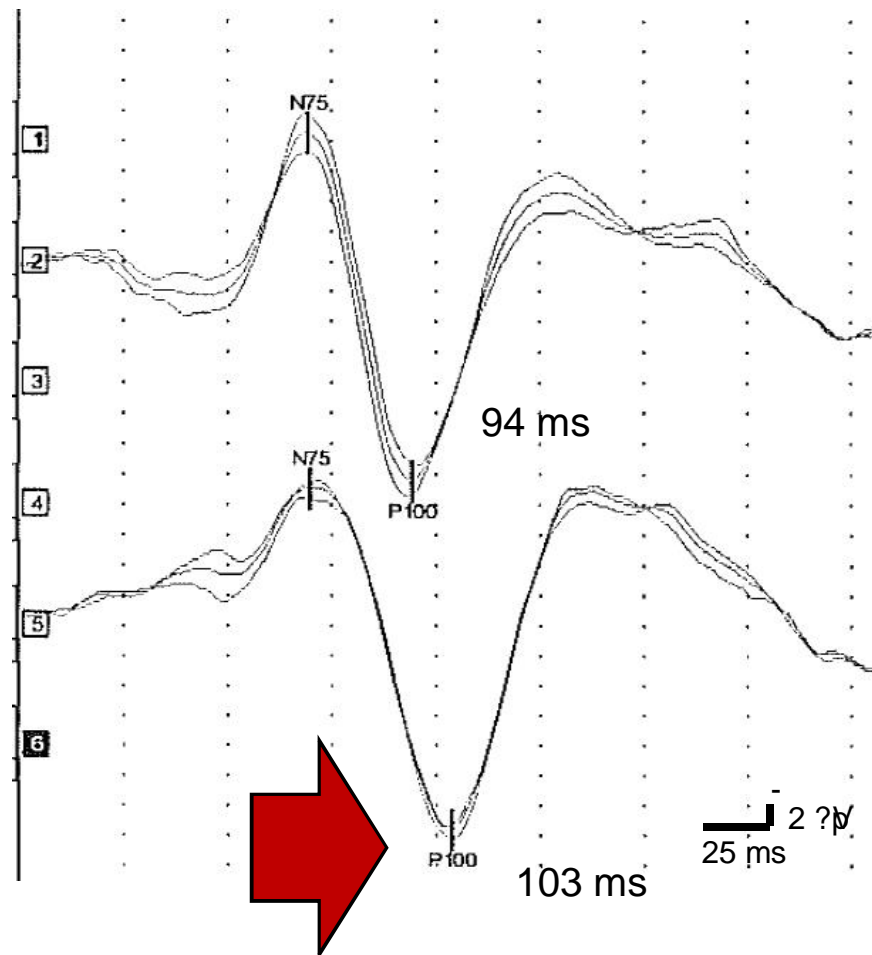


§ Normal and pathological VEPs

- P100 identification
- P100 latency
- P100 amplitude

P100

- Upper normal limit **111 ms**
- Upper normal limit of side difference **5 ms**



§ Visual evoked potentials



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§ Clinical applications and advanced use

∅ Hard rocks

MS – diagnosis

optic neuritis

neuromyelitis optica

anterior ischemic optic neuropathy

compressive optic neuropathy

∅ Soft rocks

MS – history

Degenerative diseases

Friedreich ataxia, familial spastic paraplegia, hereditary neuropathies ...

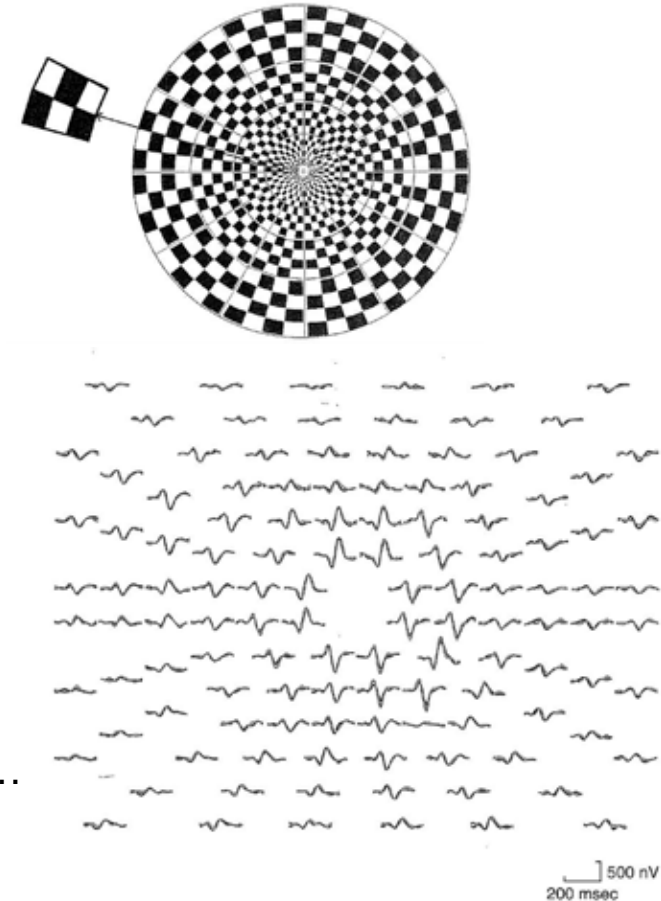
Toxic neuropathies

∅ The beach

Paraneoplastic disorders

Visual field defects

Cortical blindness



multifocal stimulated VEP

With compliments to D. J. Jewett Berlin conference 1986

subjective opinion and incomplete list

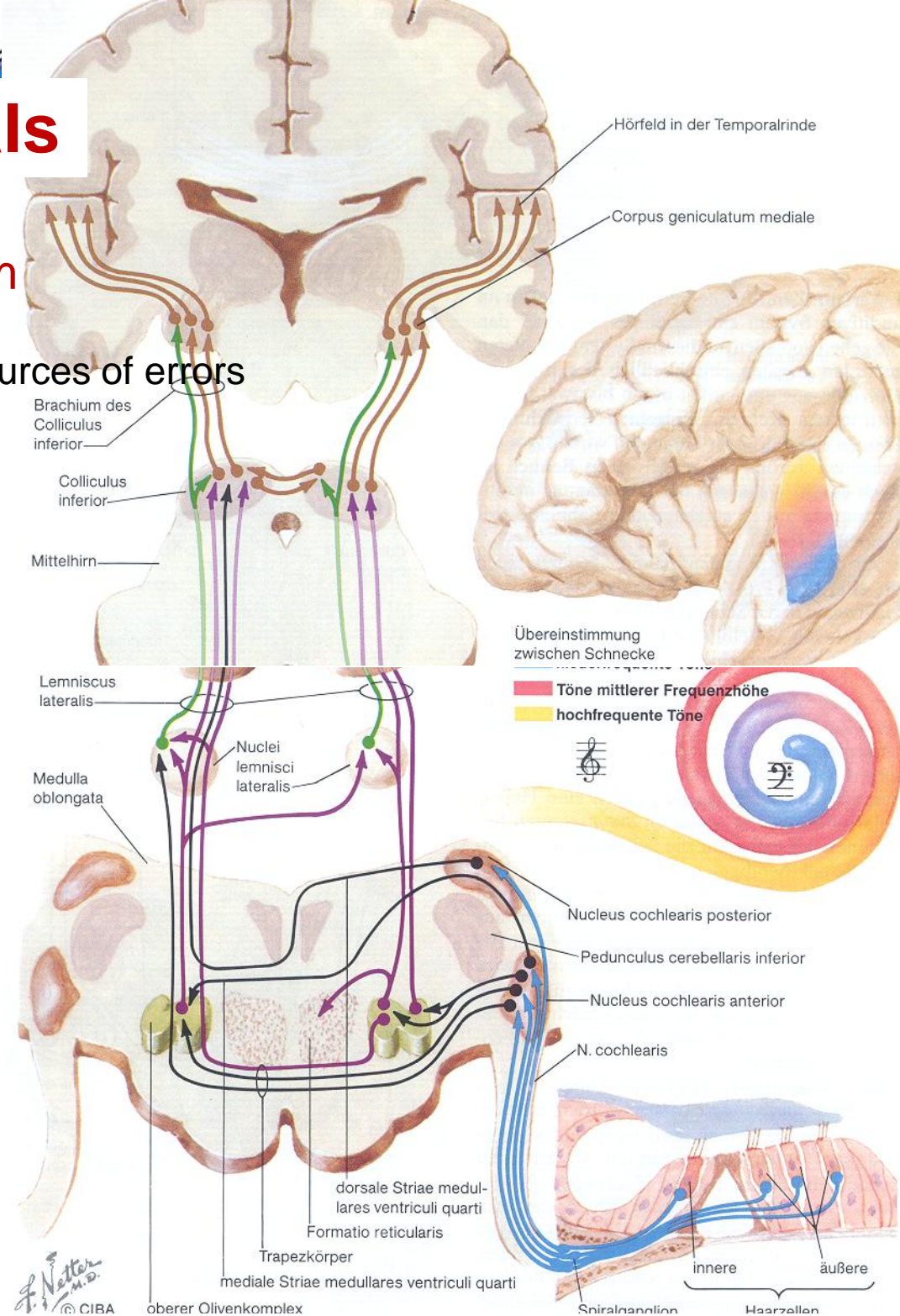
§ Auditory evoked potentials

§ Anatomy and physiology of the system

- Stimulus
- Recommended standard protocols and sources of errors
- Clinical applications and advanced use

Anatomy and Physiology

- § ear - sound management
- § compressor – cavity
- § inner ear - Sound-mechanical conversion
- § gain 1:22
- § cochlea - mechanical-electrical conversion
- § "Traveling wave" hypothesis
- § cochlear nerve
- § compound action potential
- § brainstem



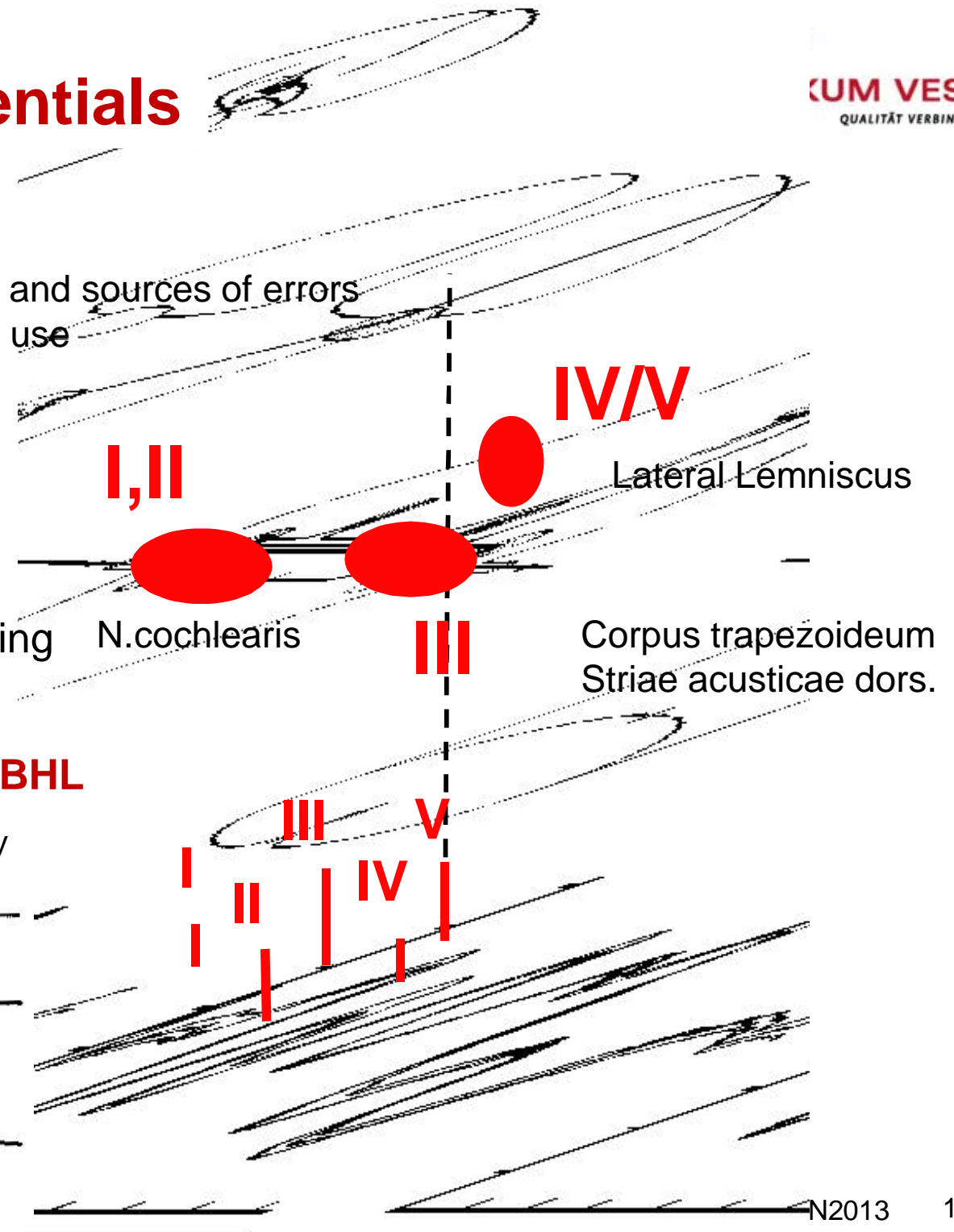
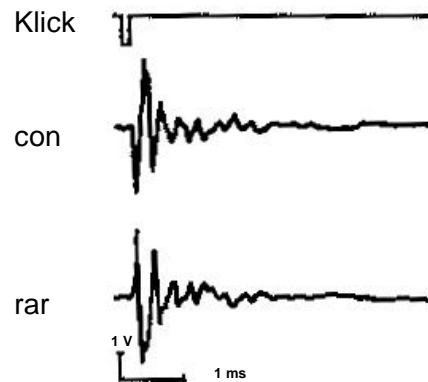
§ Auditory evoked potentials

§ Stimulus

- Recommended standard protocols and sources of errors
- Clinical applications and advanced use

Stimulus parameter

- Klick duration 100 us
- Polarity
- condensation - rarefaction = alternating
- Frequency 10 - 20 Hz (14,7)
- **Intensity 70 dBSL max. 90 dBHL**
- Contralateral - 40 dB of stimulus intensity



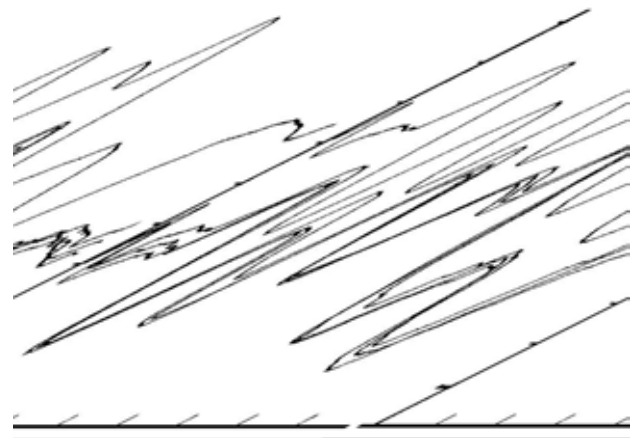
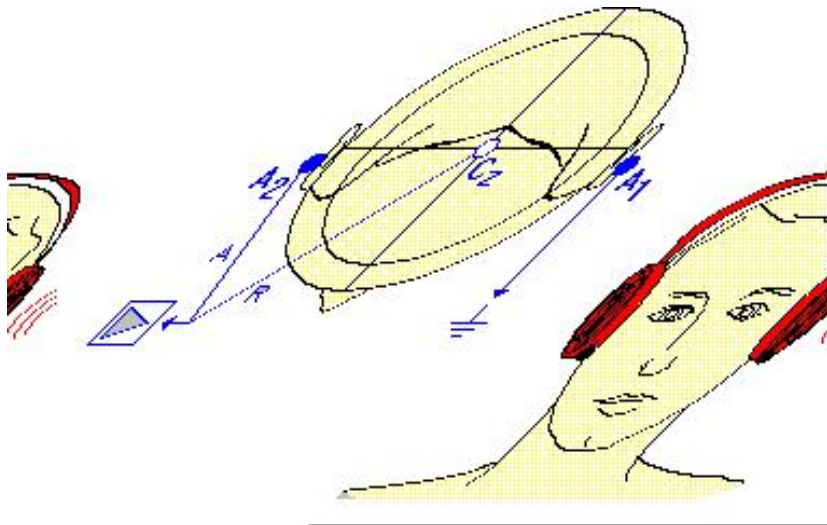


§ Auditory evoked potentials

§ Recommended standard protocols and sources of errors

- Clinical applications and advanced use

§ Acquisition



quality control

- § two measurements
- § Reproduction with:
- § Latencies below 0.1 ms
- § Amplitudes of + / - 20%

measurement parameters

- § Channel 1 Ai – Cz
- § Channel 2 Ac – Cz
- § Filters 100 - 3000 Hz
- § Gain about 100,000 times (about 1-5 uV / dev)
- § Measuring time 10 ms

Upper normal limits

- I-III 2.5 ms – side difference 0.5 ms
- III-V 2.4 ms – side difference 0.5 ms
- I-V 4.5 ms – side difference 0.5 ms

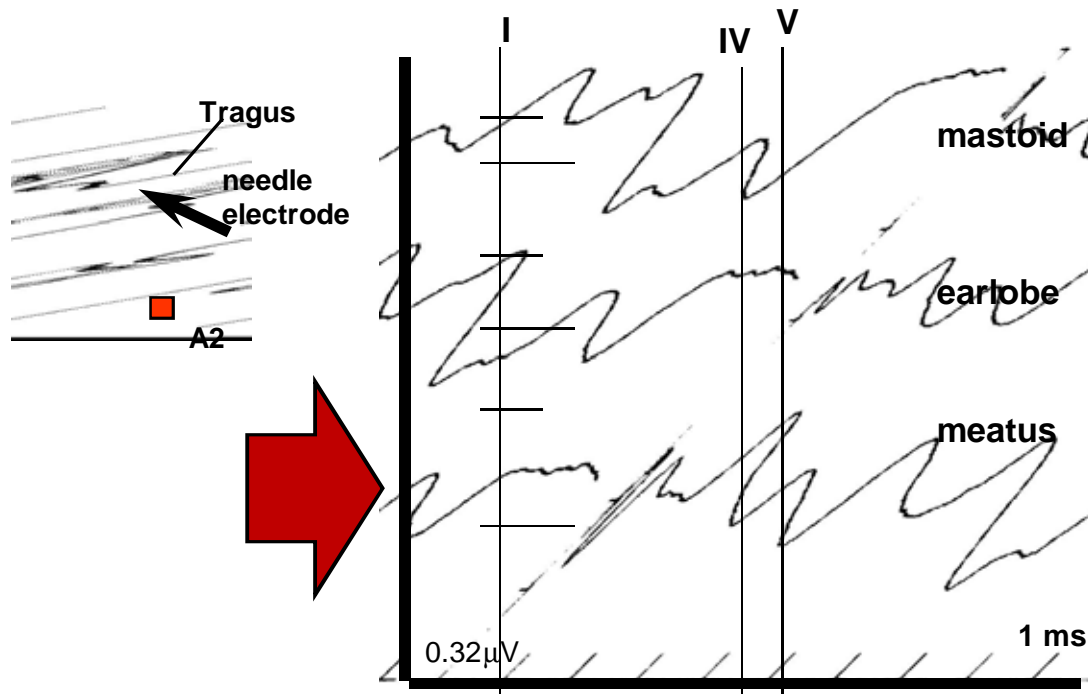
§ Auditory evoked potentials



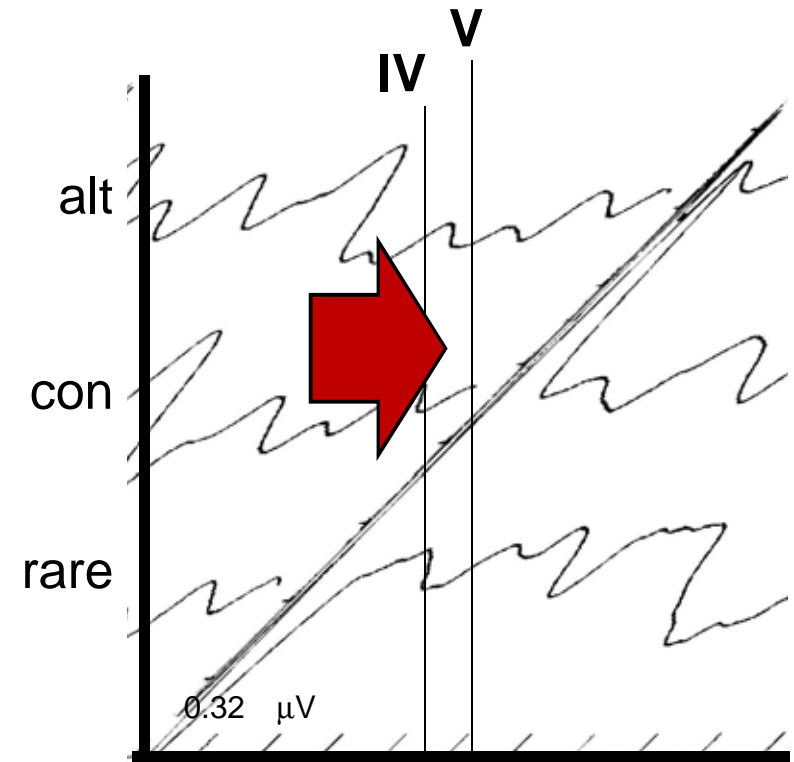
§ Recommended standard protocols and sources of errors

- Clinical applications and advanced use

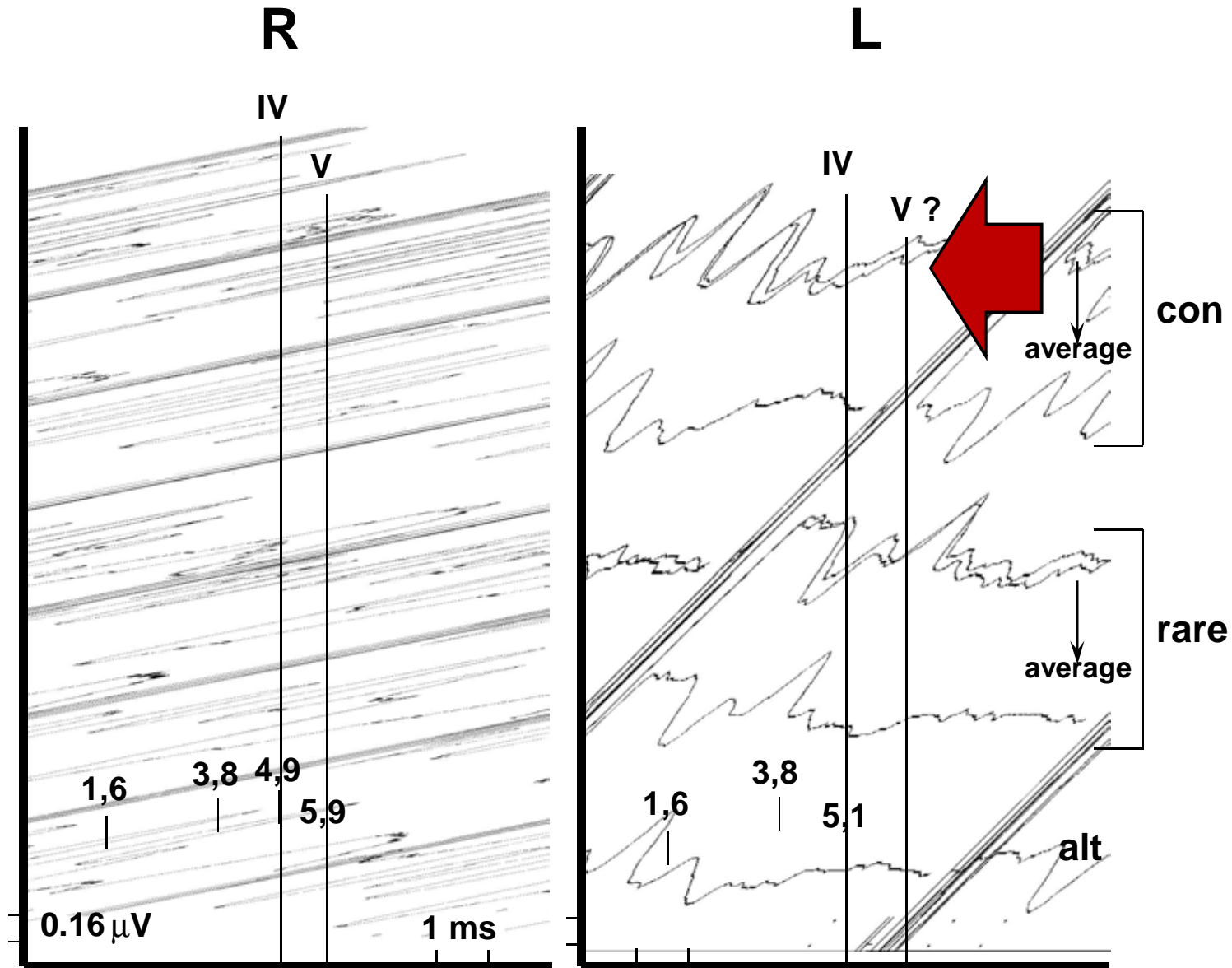
§ Acquisition



Identification of wave V



§ Auditory evoked potentials

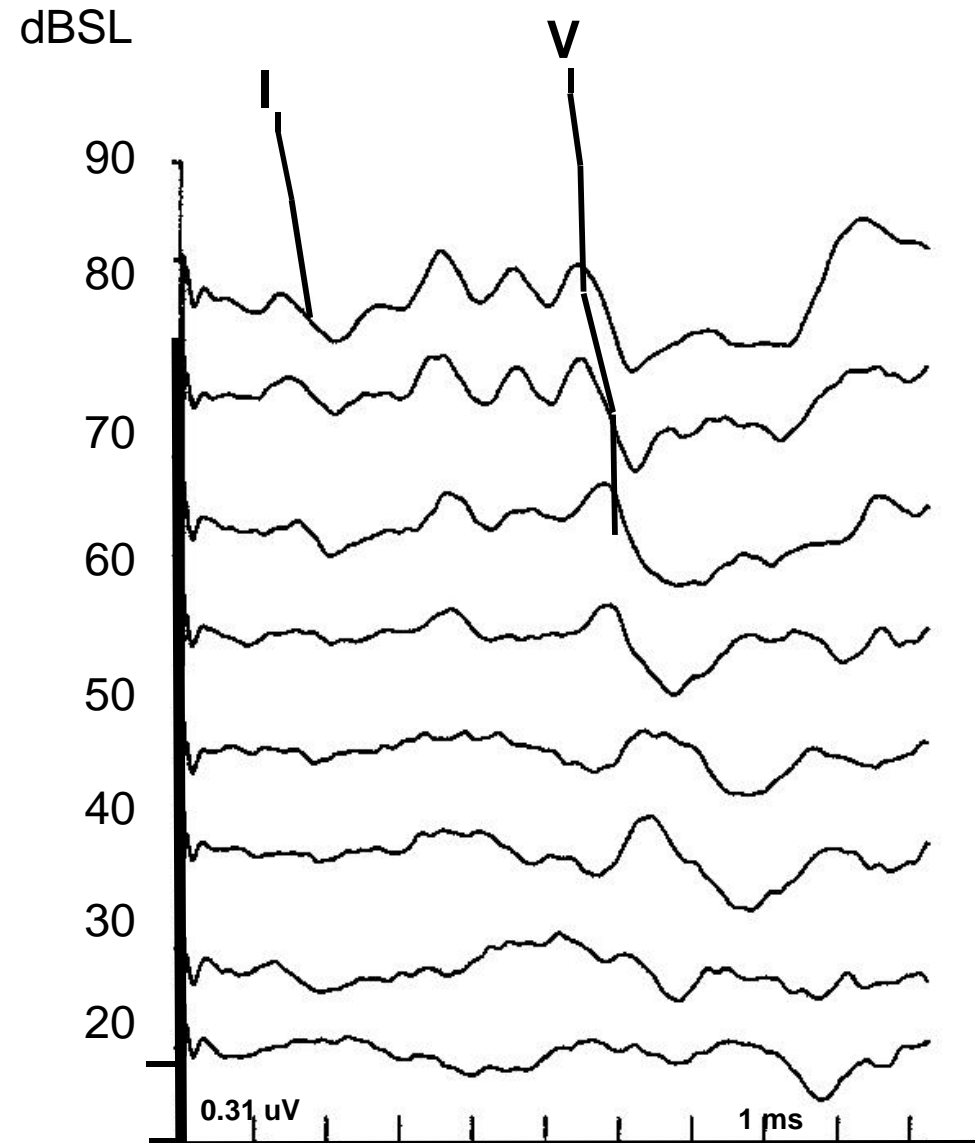
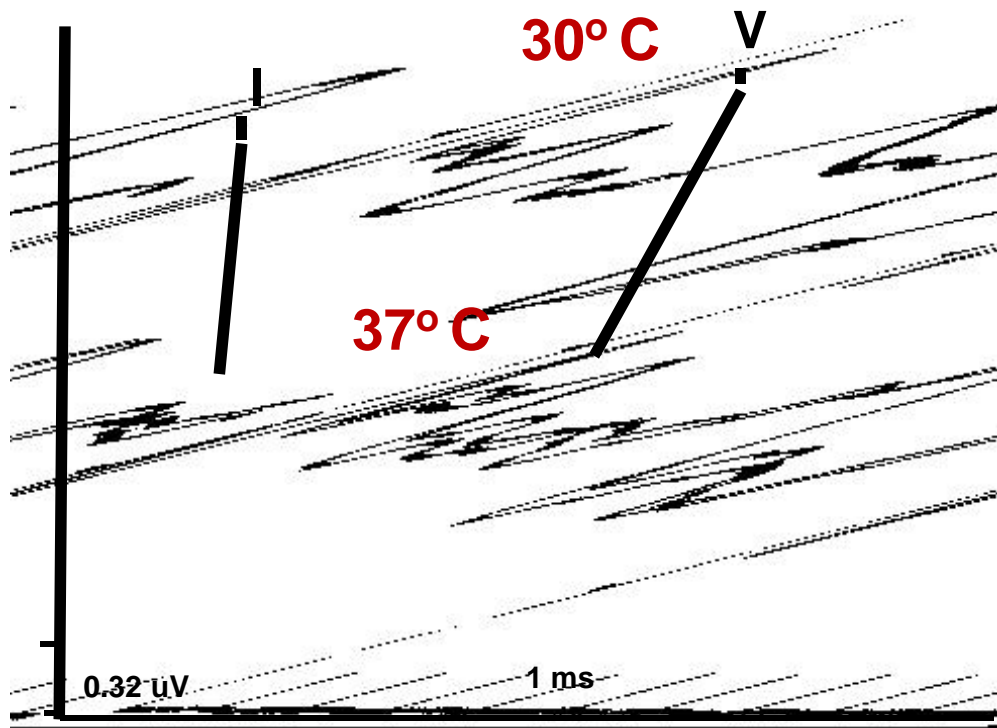


§ Auditory evoked potentials

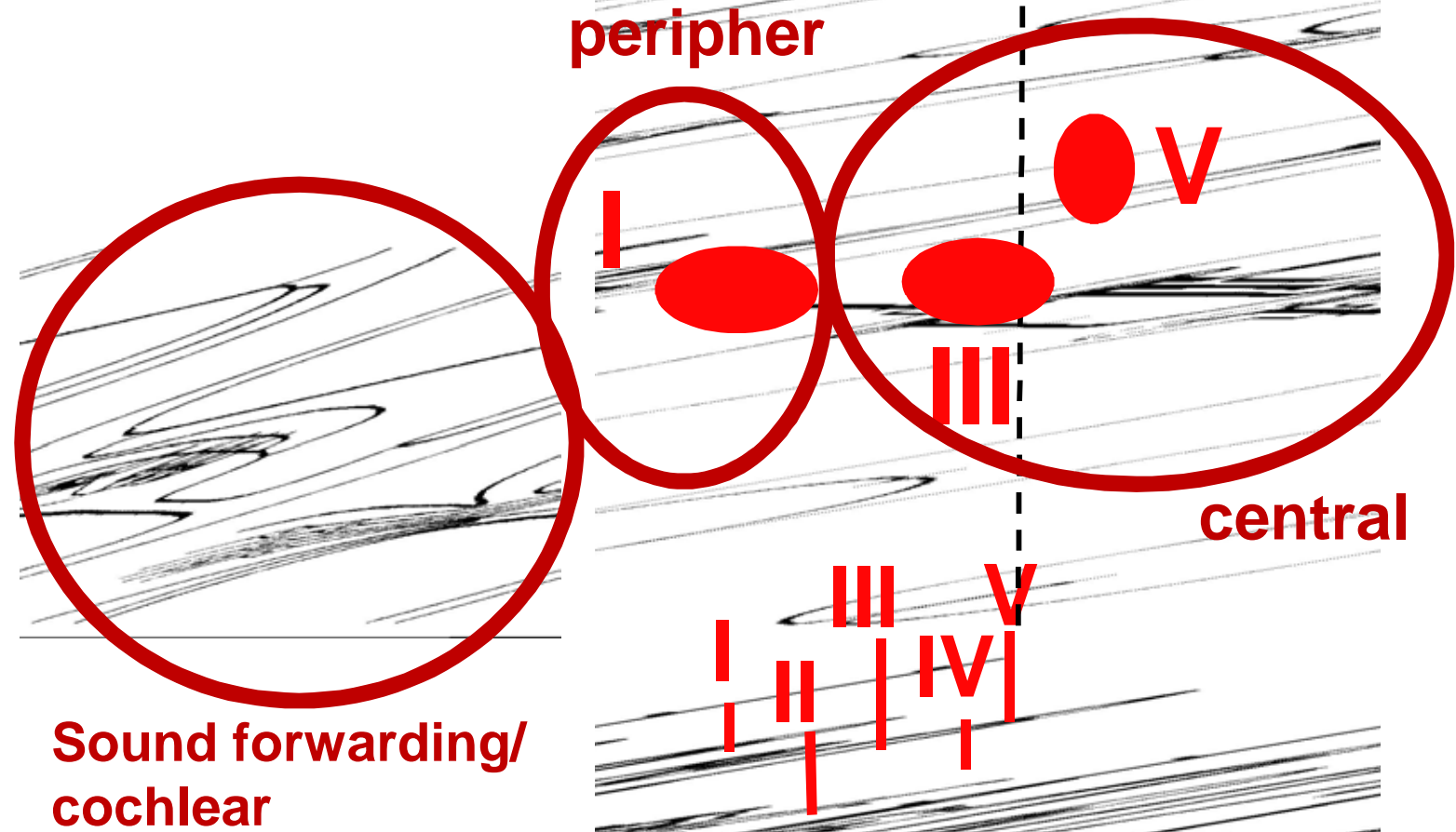


§ sources of errors

- body temperature
- peripheral vs. central hearing



§ Auditory evoked potentials

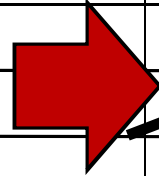


normal

0.5 uV

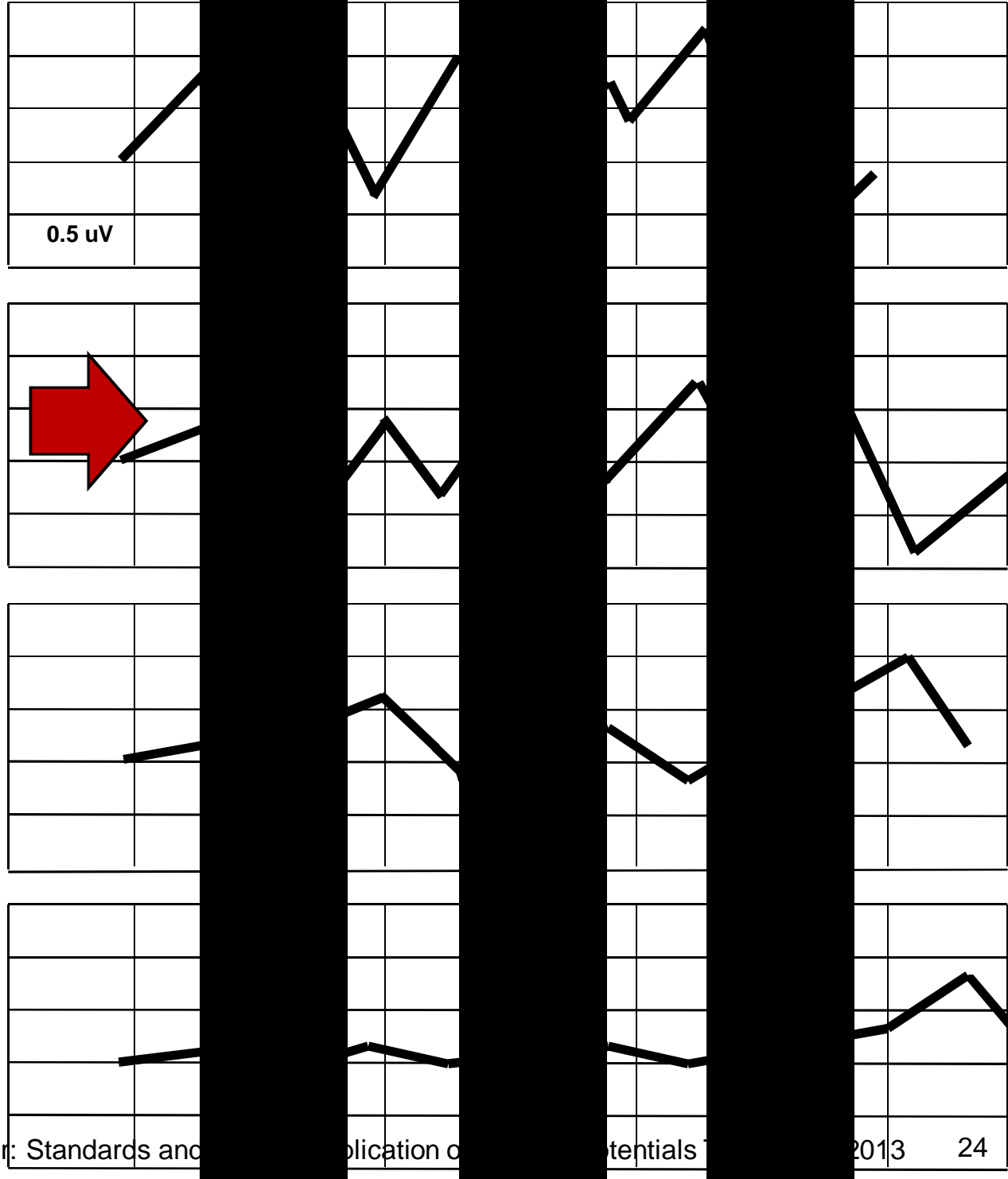
1 ms

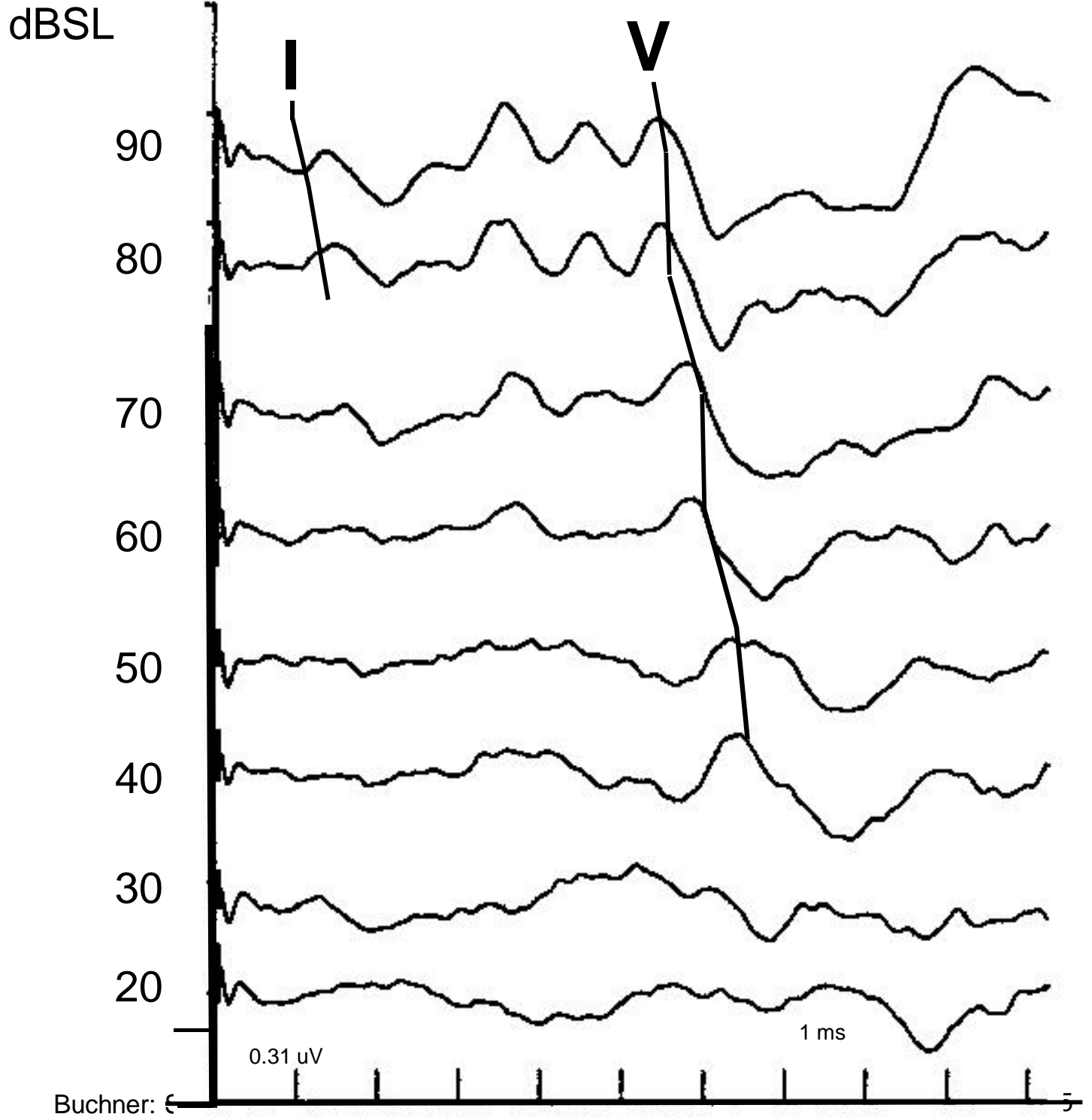
delayed I to V



delayed I to V
loss of II to IV

delayed V
loss of I to IV

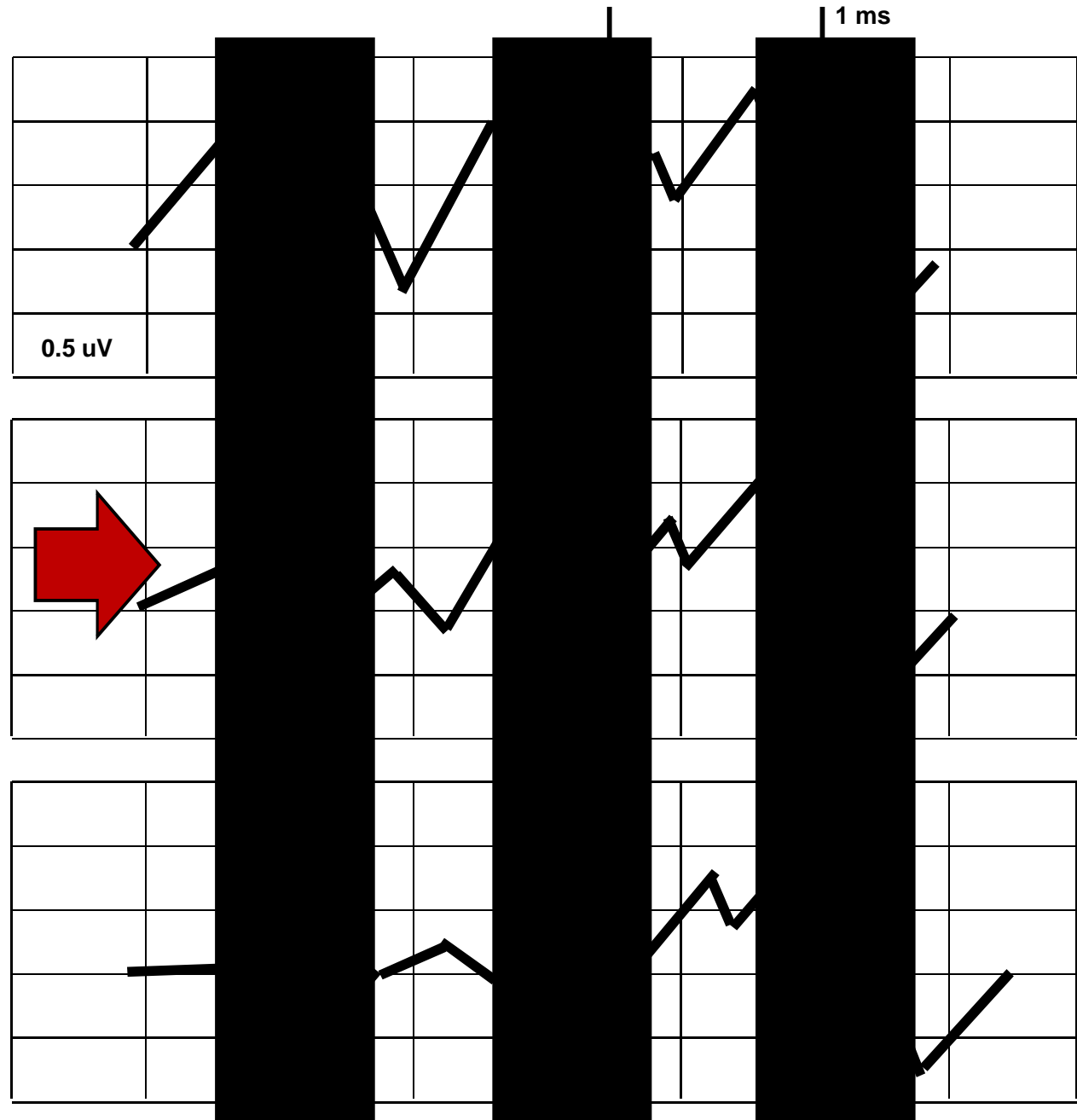




normal

delayed I
normal V
shortened I-V

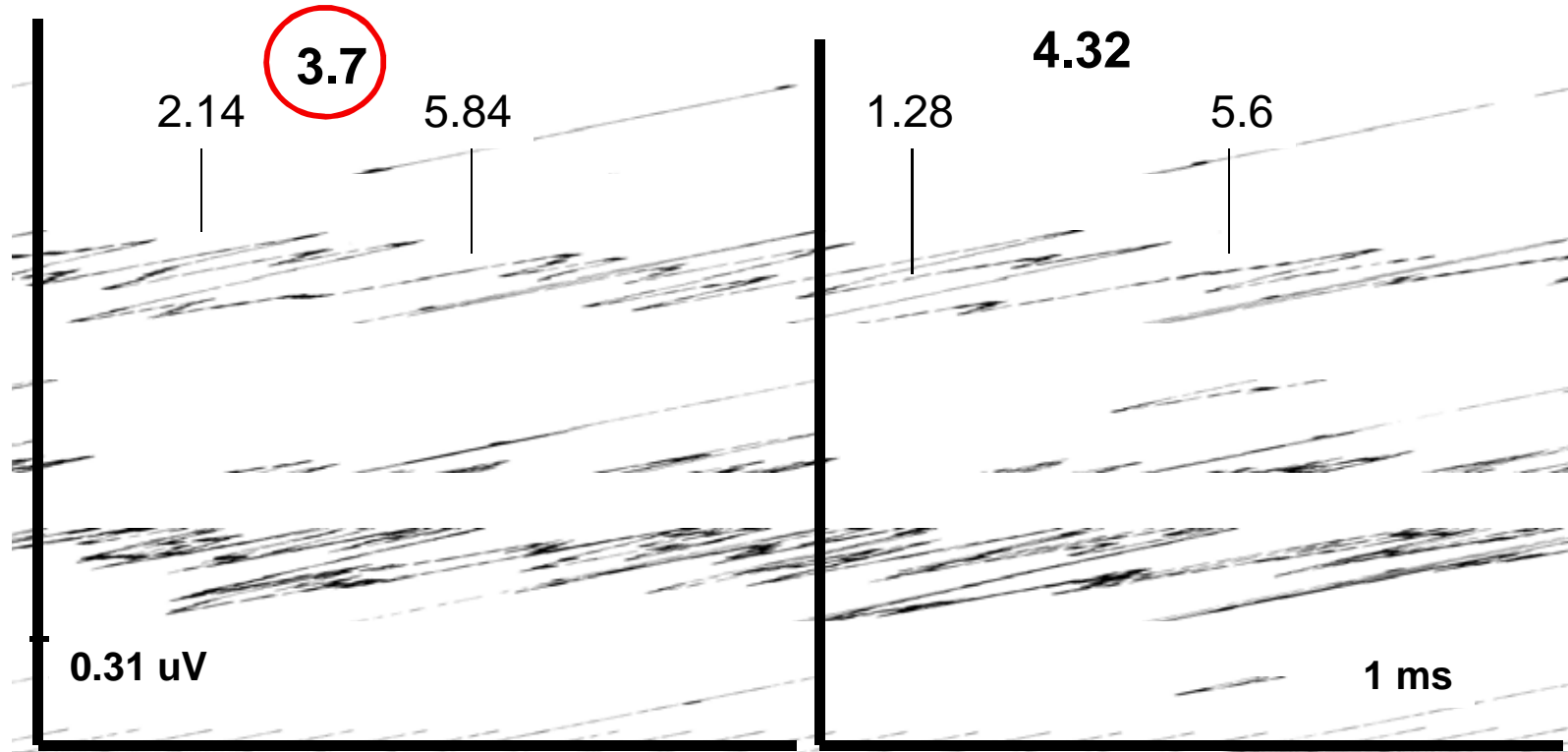
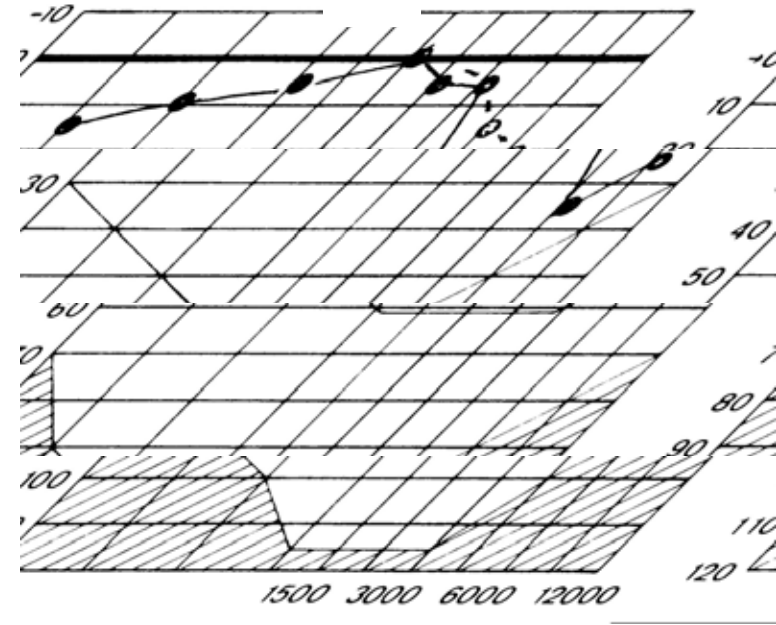
delayed I
delayed V
shortened I-V



R



L



Upper limits

- I-III 2.5 ms
- III-V 2.4 ms
- I-V 4.5 ms

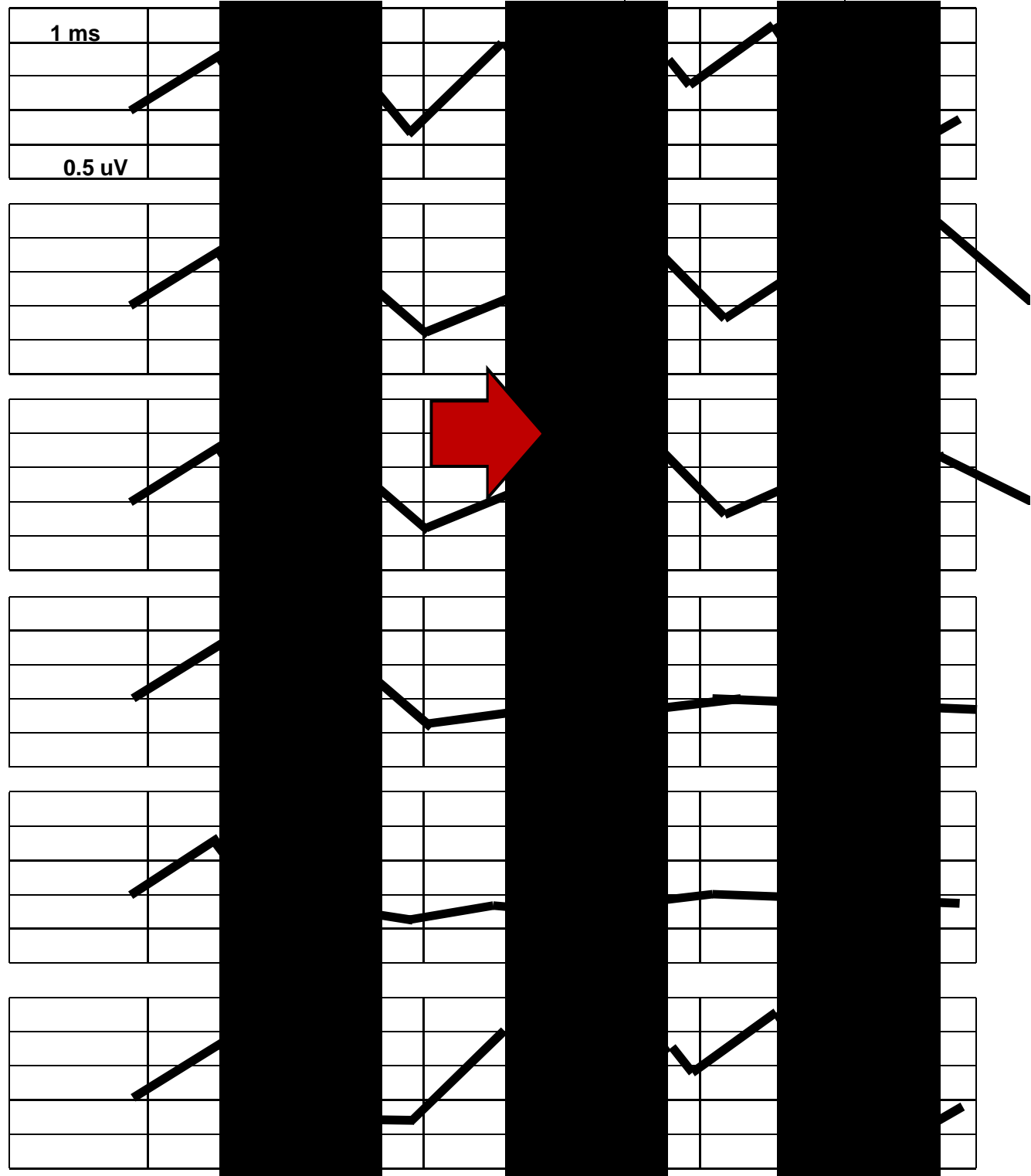
normal

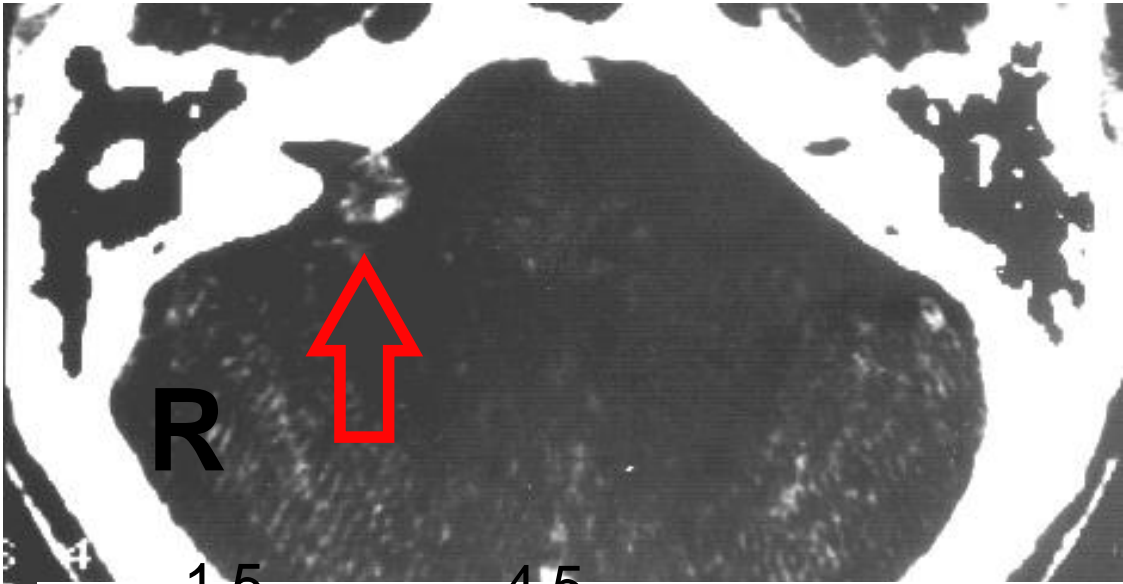
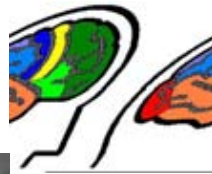
delayed III
prolonged I-III

delayed III
prolonged I-III
reduced amplitude V

reduced amplitude III
reduced amplitude II

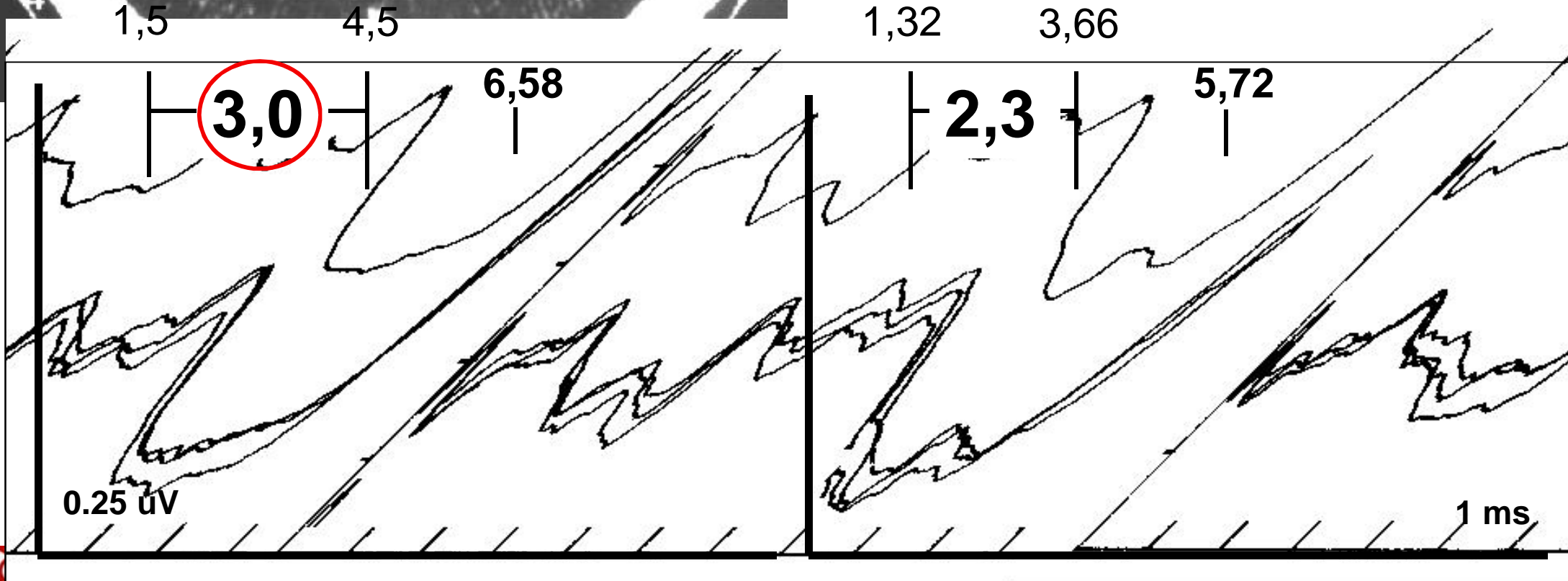
Isolated
loss of II





Upper limits

- I-III 2.5 ms
- III-V 2.4 ms
- I-V 4.5 ms



normal

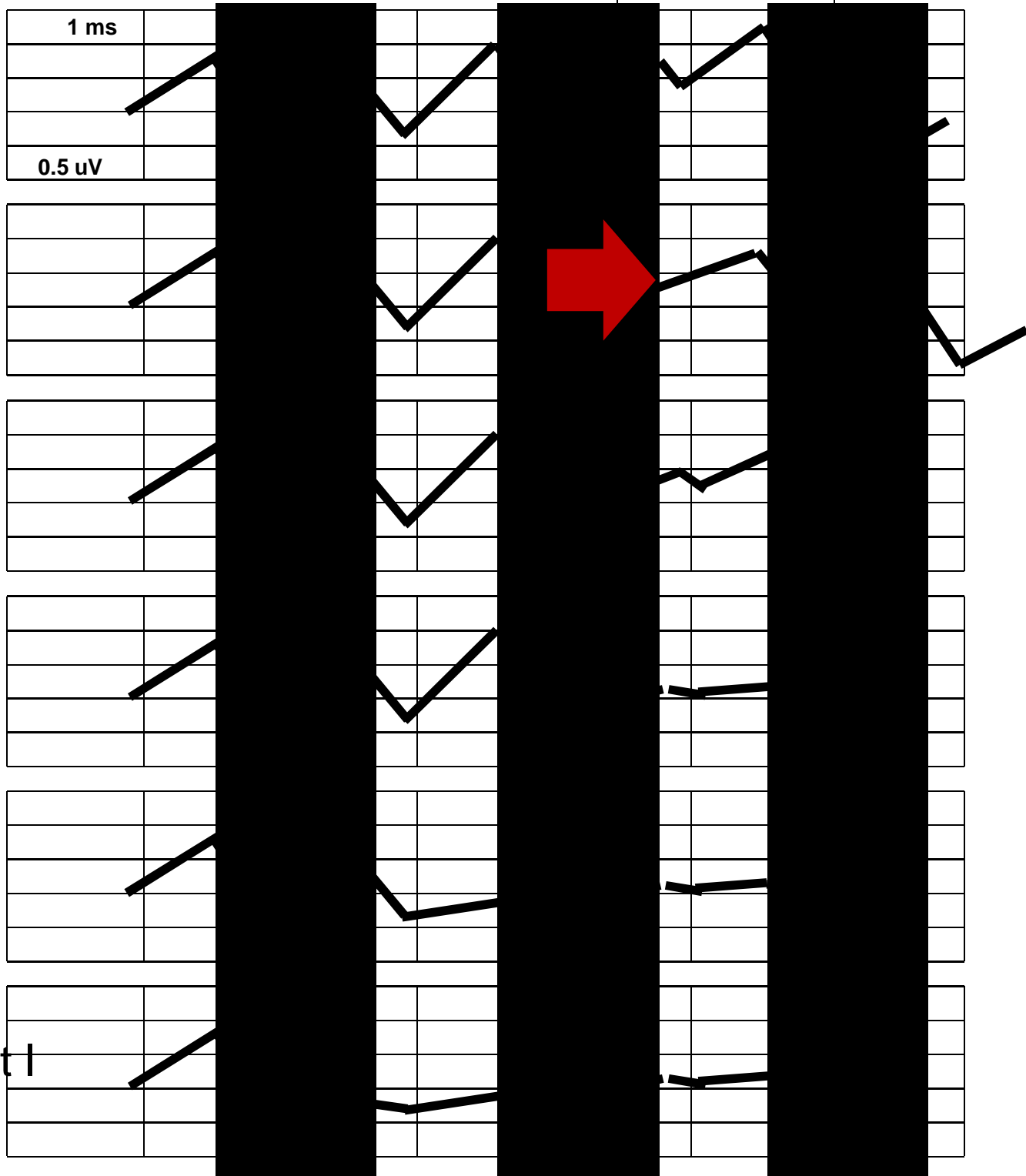
delayed V

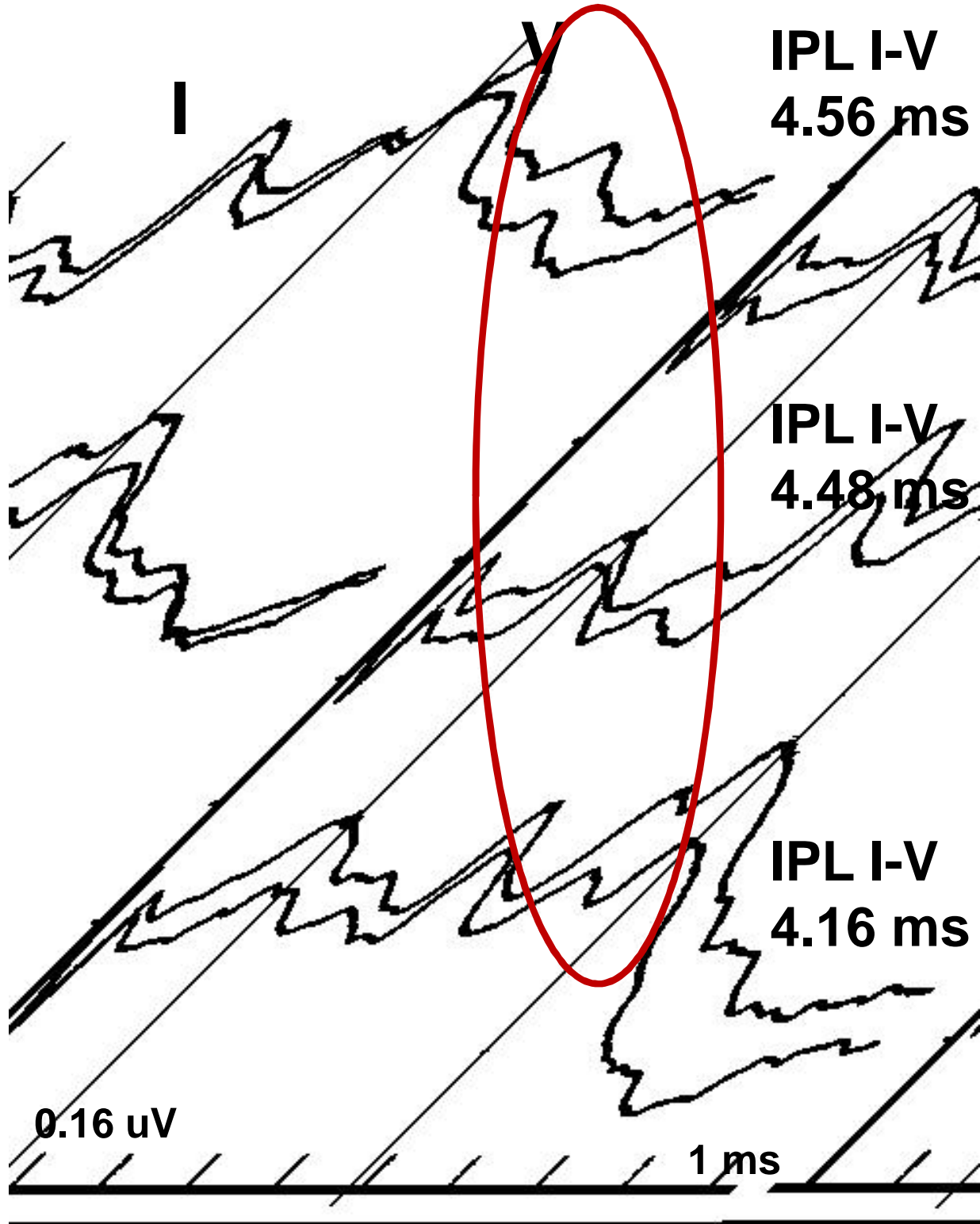
delayed V
reduced amplitude V

loss of IV/V

loss of III and IV/V

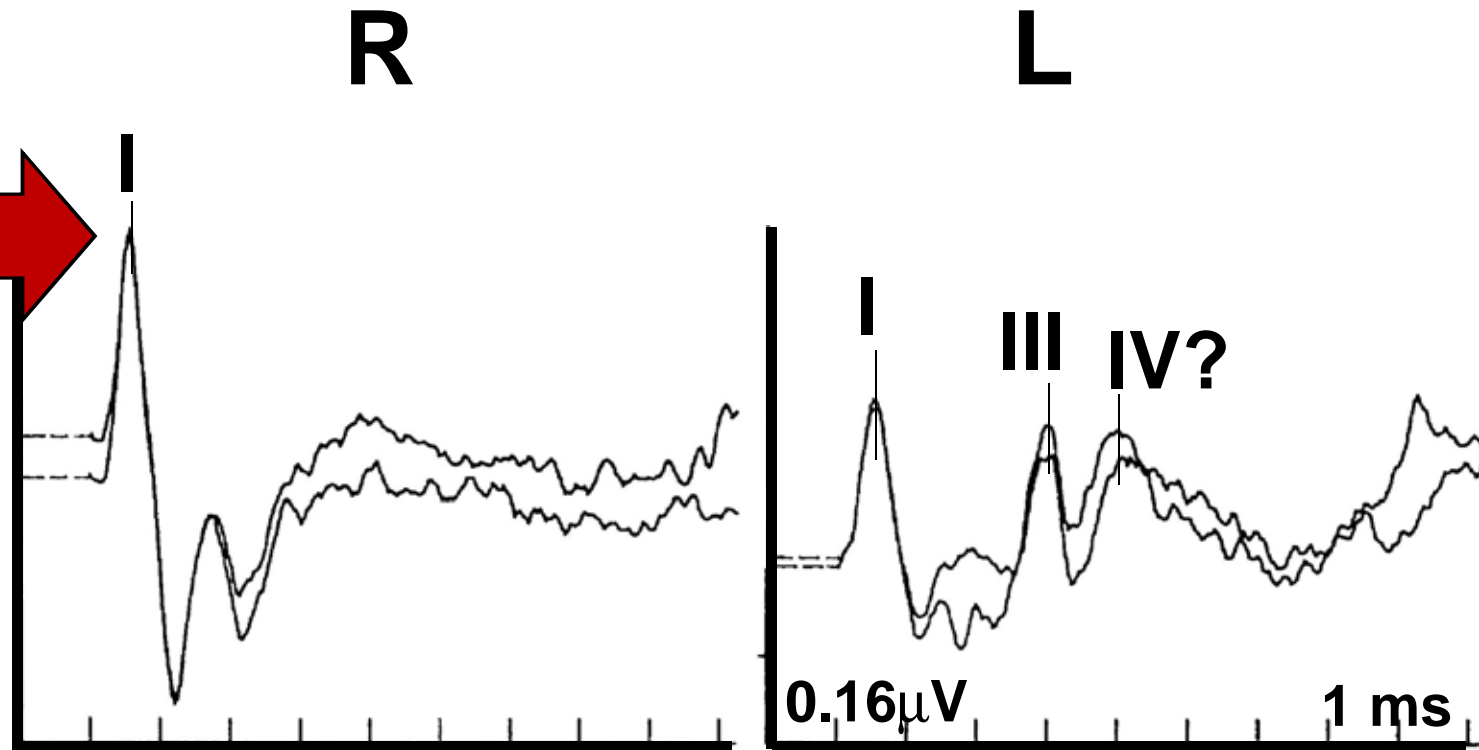
loss of all waves except I





upper normal
limit 4,5 ms

§ Auditory evoked potentials



§ Auditory evoked potentials



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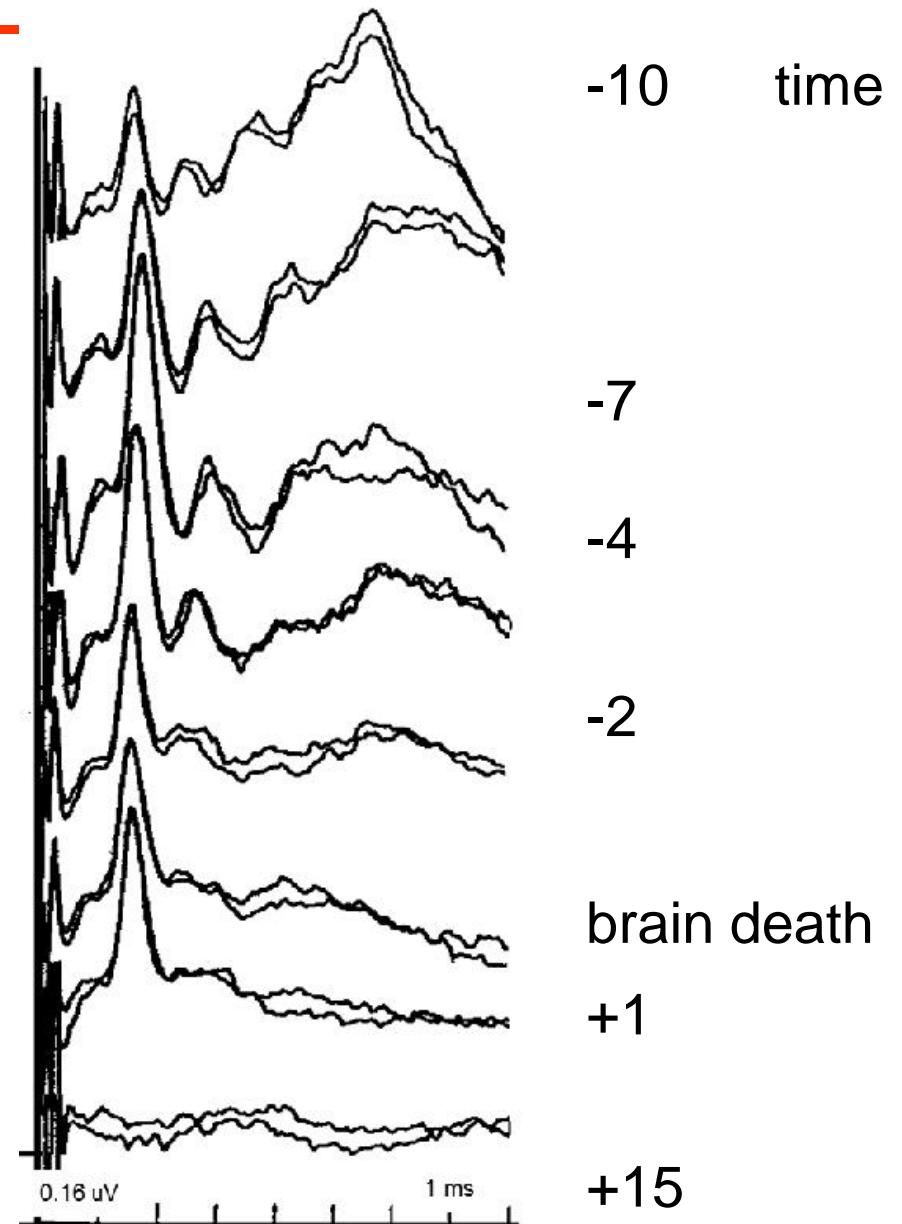


brain death

validity

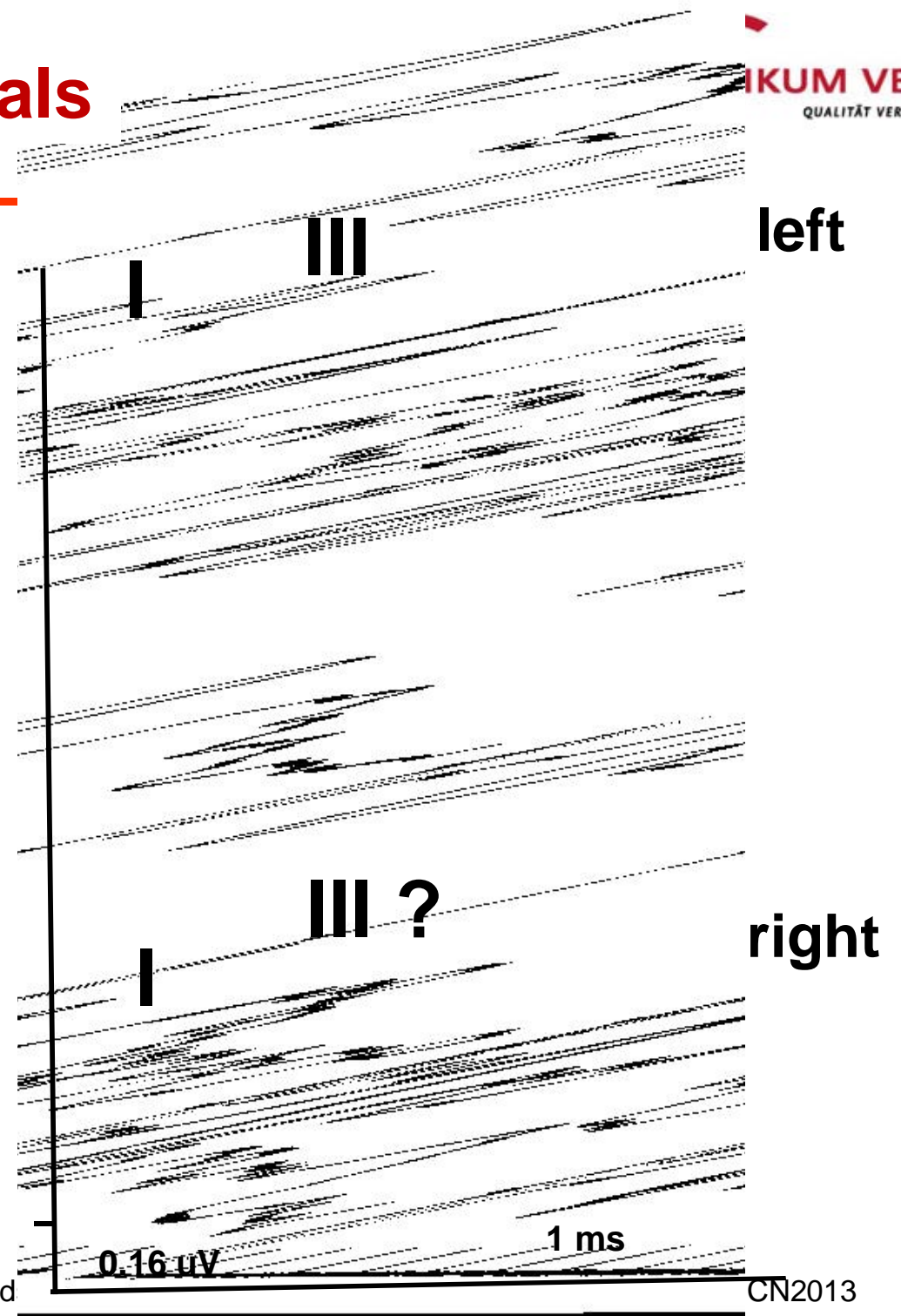
130 HT from examination
6 (4,6%) with preserved waves III or IV/V

Facco et al. 2002 Clin Neurophysiol



§ Auditory evoked potentials

Multiple sclerosis



§ Auditory evoked potentials



§ Clinical applications and advanced use

∅ **Hard rocks**

retrocochlear hearing impairment
posterior fossa tumors
MS - diagnosis
Prognosis in coma

∅ **Soft rocks**

brain death
intoxication
peripheral hearing loss

∅ **The beach**

dizziness
MS - history
diagnosis of ischemic brainstem lesions

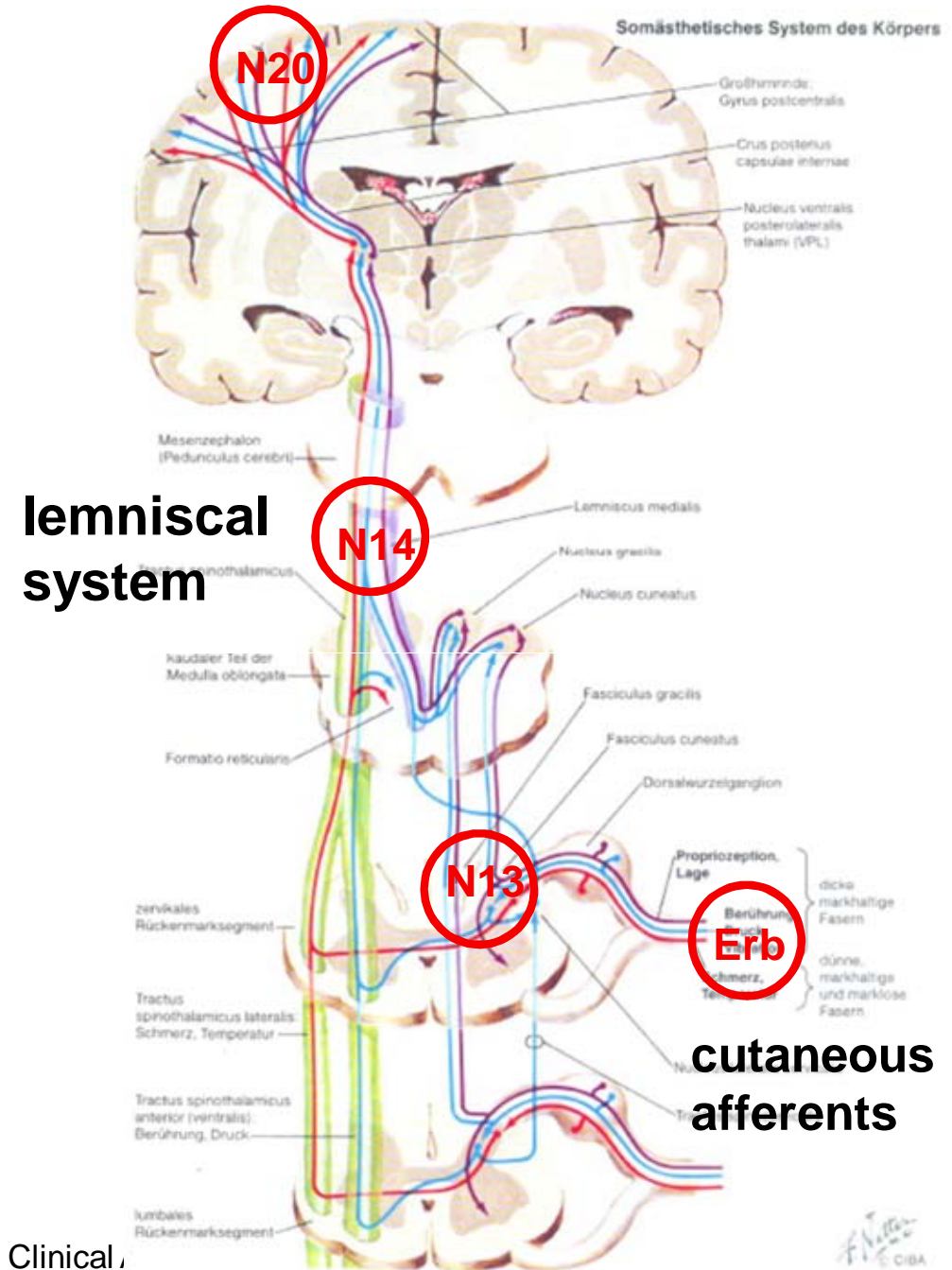
subjective opinion and incomplete list

With compliments to D. J. Jewett Berlin conference 1986

§ Somatosensory evoked potentials

§ Anatomy and physiology of the system

- Stimulus
- Recommended standard protocols and sources of errors
- Clinical applications and advanced use





§ Somatosensory evoked potentials

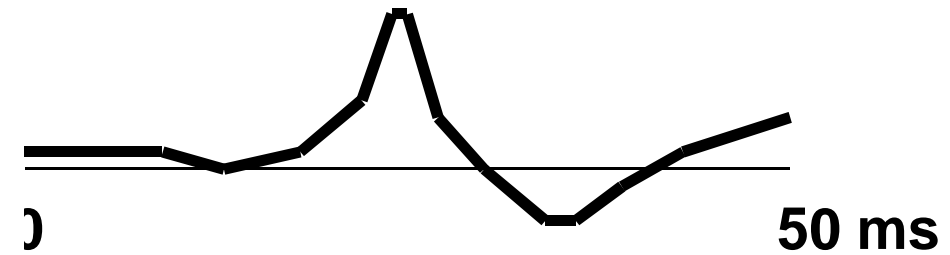
§ Stimulus

- Recommended standard protocols and sources of errors
- Clinical applications an advanced use

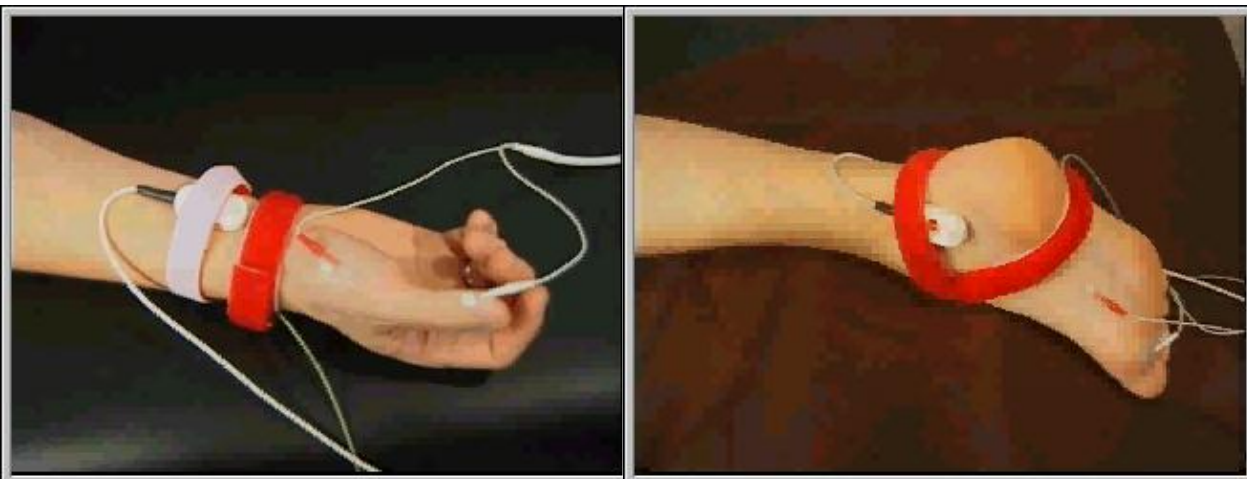
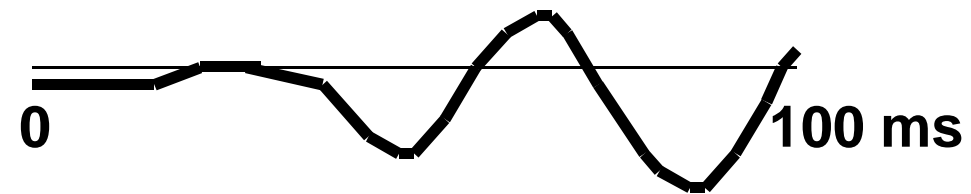
Stimulus parameter

- constant current; duration 0,2 ms
- cathode proximal
- 1 - 10 /sec; 4,3 /sec
- **intensity above motor threshold**

median N.



tibial N.





§ Somatosensory evoked potentials

§ quality control

§ two measurements

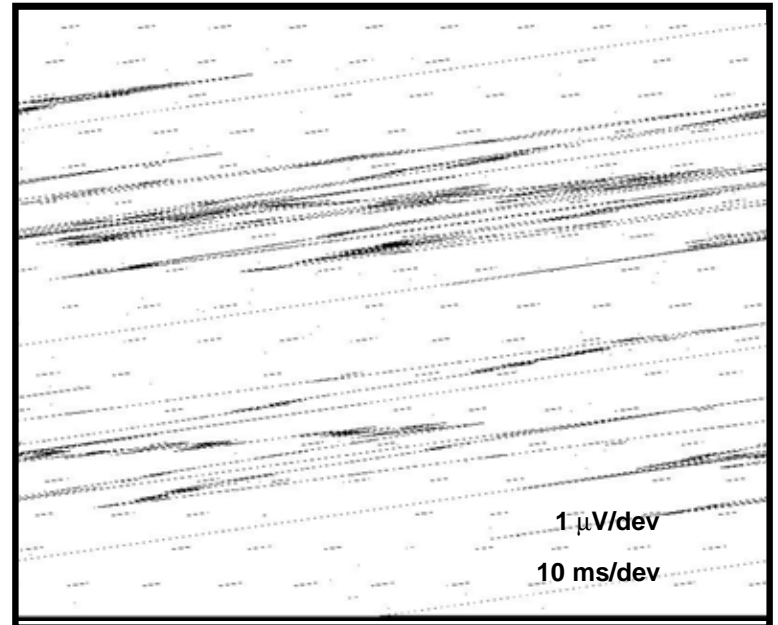
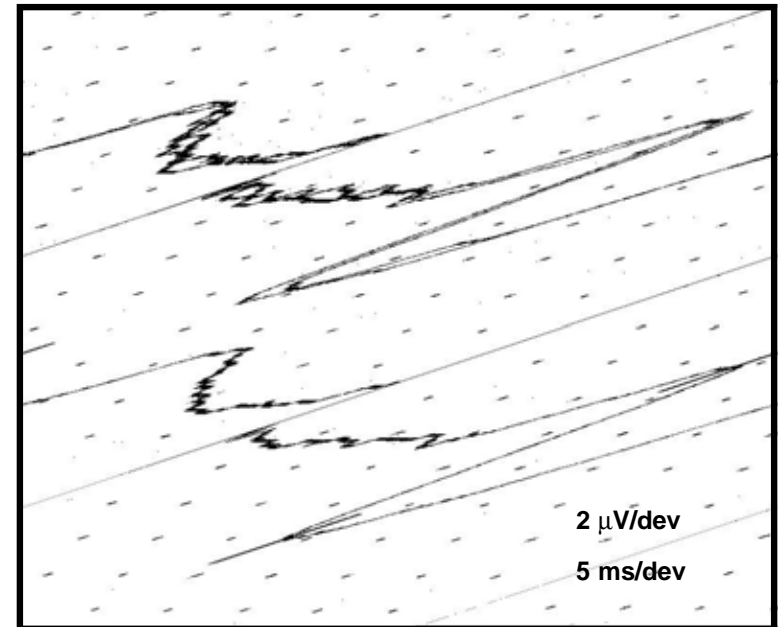
§ Reproduction with:

§ Latencies

§ Arm nerves below 0.25 ms

§ Leg nerves below 0.5 ms

§ Amplitudes of + / - 20%



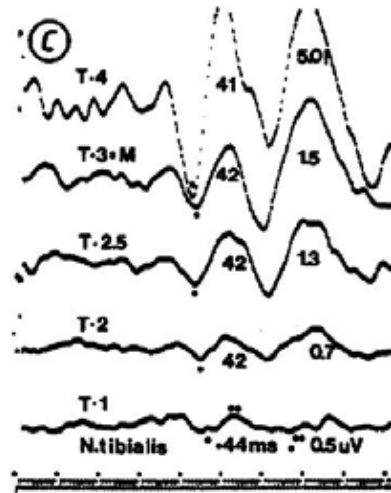


§ Somatosensory evoked potentials

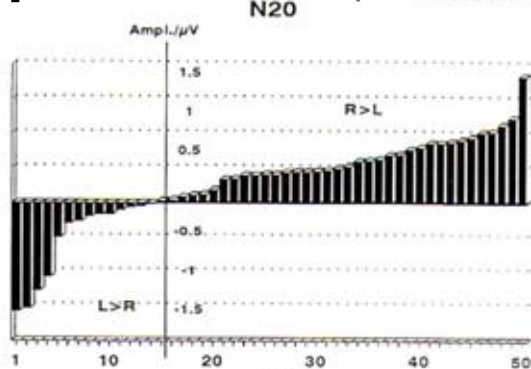
§ sources of errors

- body temperature
- body size
- muscle artifacts

Stimulus intensity



Electrode placement



Vogel et. al. J Neurol 1982, 228,97-111

Buchner et al. Electromyogr. Clin. Neurophysiol. 1995, 35, 207-215

Factors

§ Height

parallel with body size - but only very small effects on
N. median N9-N20 and N. tibial N22-P40
1.50 to 1.90 cm

§ Temperature

in parallel with the conduction time of the peripheral nerves

§ Gender

Women shorter latencies

§ Drugs

little to no effect - only when intoxication prolonged N13-N20

§ Sleep – attention

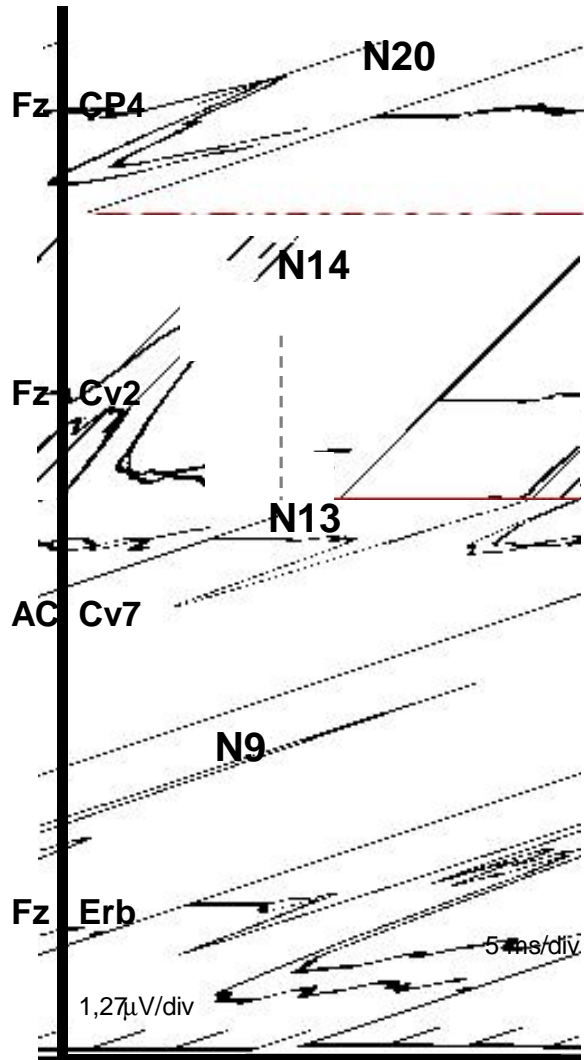
no effects

§ Age

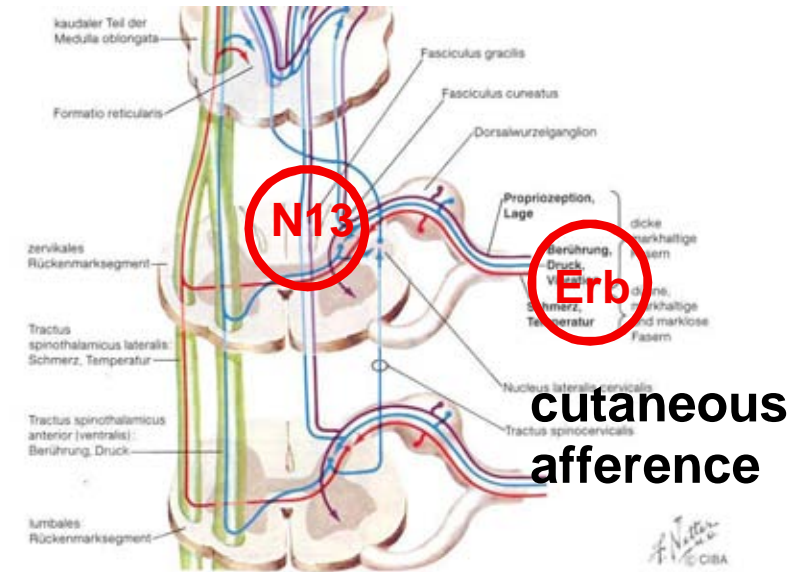
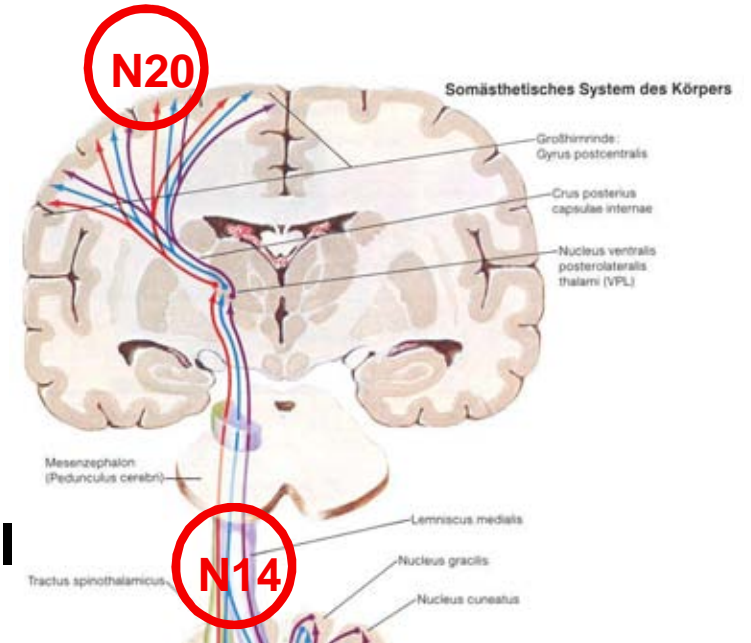
Tendency to higher amplitudes and longer latencies

§ Somatosensory evoked potentials

§ Median nerve



lemniscal system





§ Normal values N. medianus

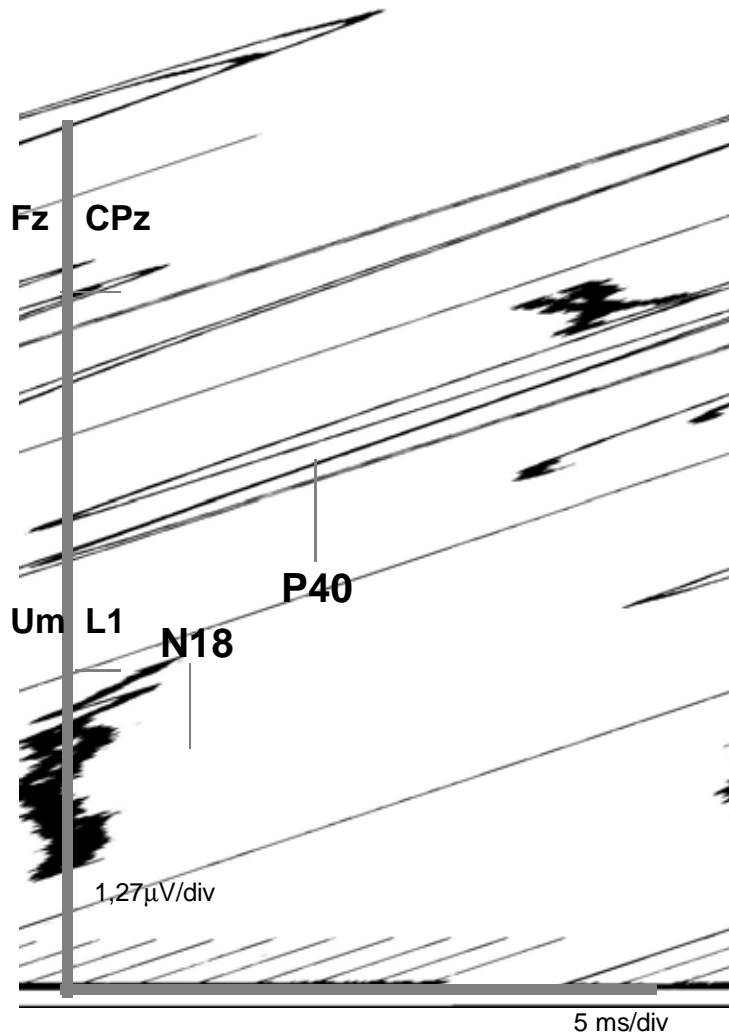
Latency			
	mean	+ 3 SD	re - li
N9	9.8	11.5	-
N13	10.1	14.5	-
N14	14.3	16.7	0.8
N20	19.8	23.0	1.4
N9-N13	3.5	4.5	1.3
N13-N20	5.7	7.2	1.0
N14-N20	4.6	6.0	1.1

Amplitude			
	mean	lower limit	re-li
N9 (baseline to peak)	4.8	1.0	50%
N13 (baseline to peak)	2.3	0.5	-
N20 (baseline to peak)	2.2	0.6	47%
N20-P25 (peak to peak)	3.2	0.8	-

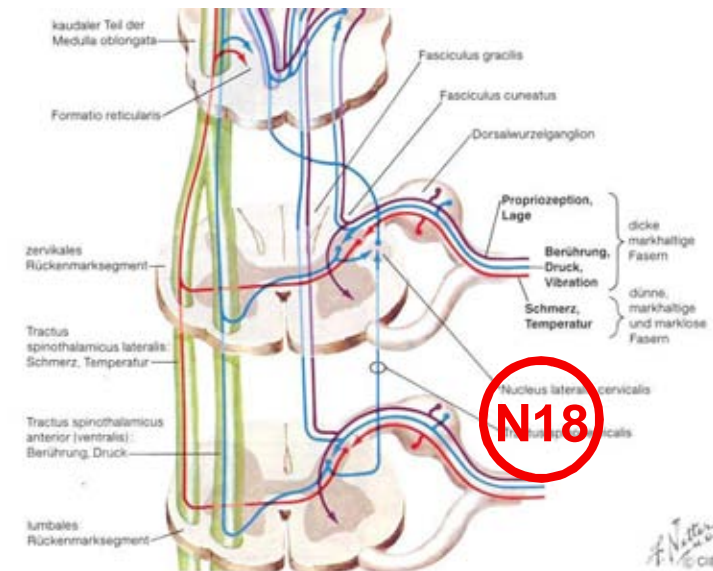
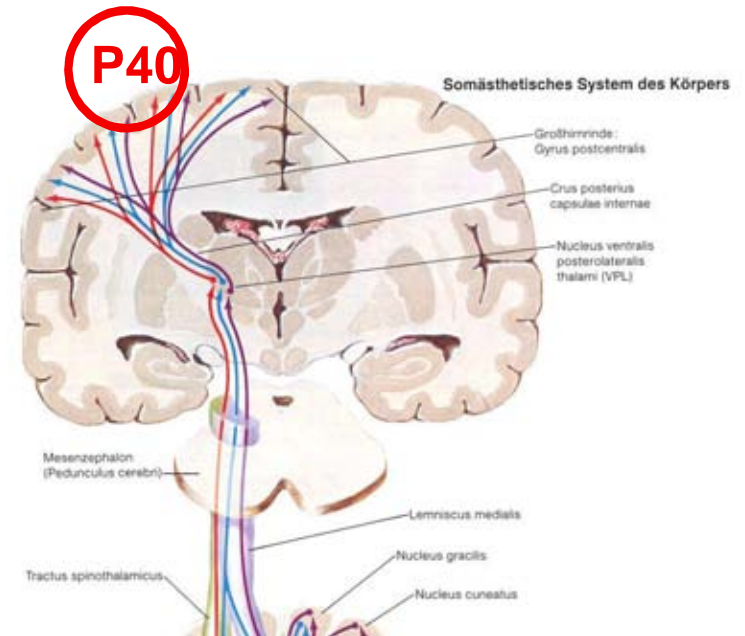
Mauguière et al. (1999) Electroenceph. clin. Neurophysiol. supp. 52 79-90

§ Somatosensory evoked potentials

§ Tibial nerve



**Lemniscal
plus
extra-
lemniscal**





§ Normal values N. tibialis

Latency			
	mean	+ 3 SD	re - li
N22	21.8	25.2	1.1
P40	38.0	43.9	2.1
N22-P40	16.0	20.6	2.1

Amplitude			
	mean	lower limit	re-li
N22 (baseline to peak)	1.1	0.3	-
P40-N50 (peak to peak)	1.8	0.5	-

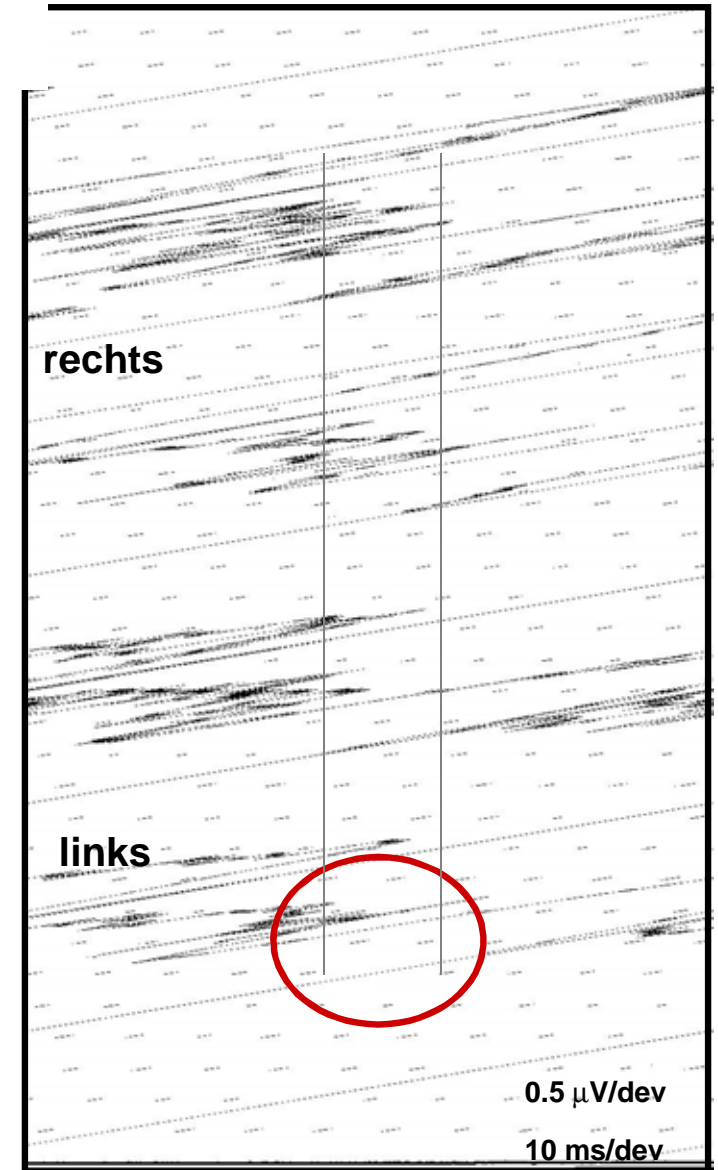
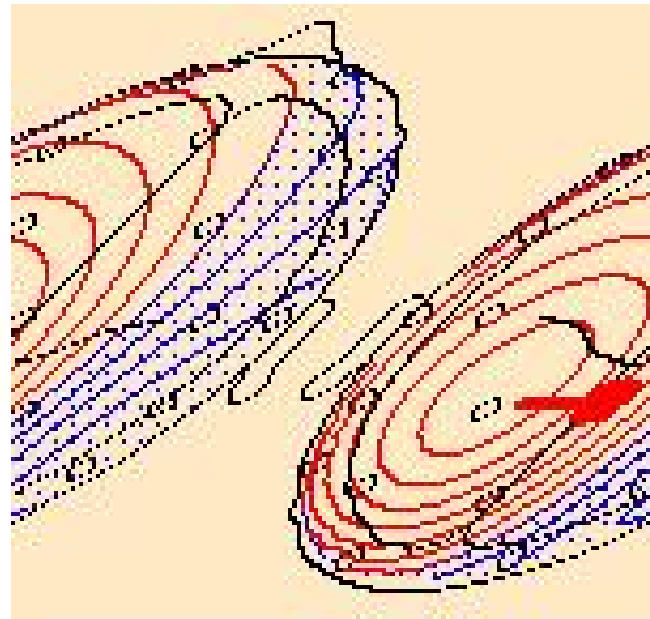
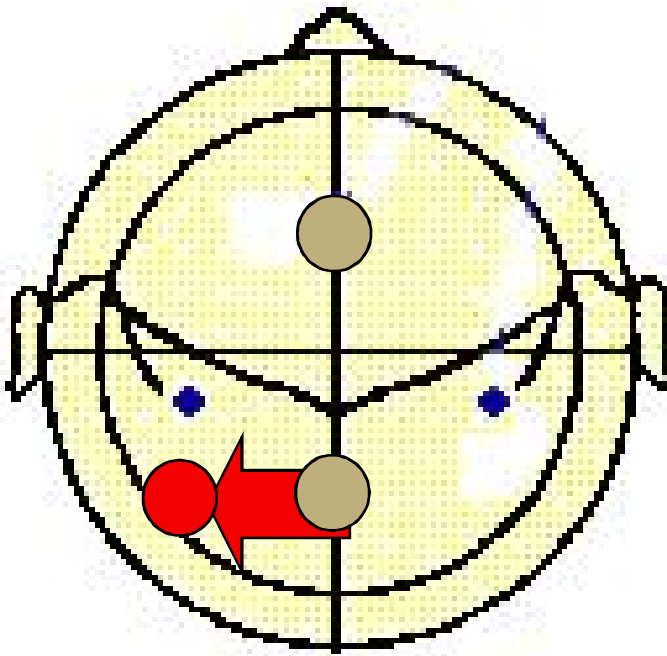
Mauguière et al. (1999) Electroenceph. clin. Neurophysiol. supp. 52 79-90



§ Somatosensory evoked potentials

§ Tibial nerve

– paradoxical lateralization of P40

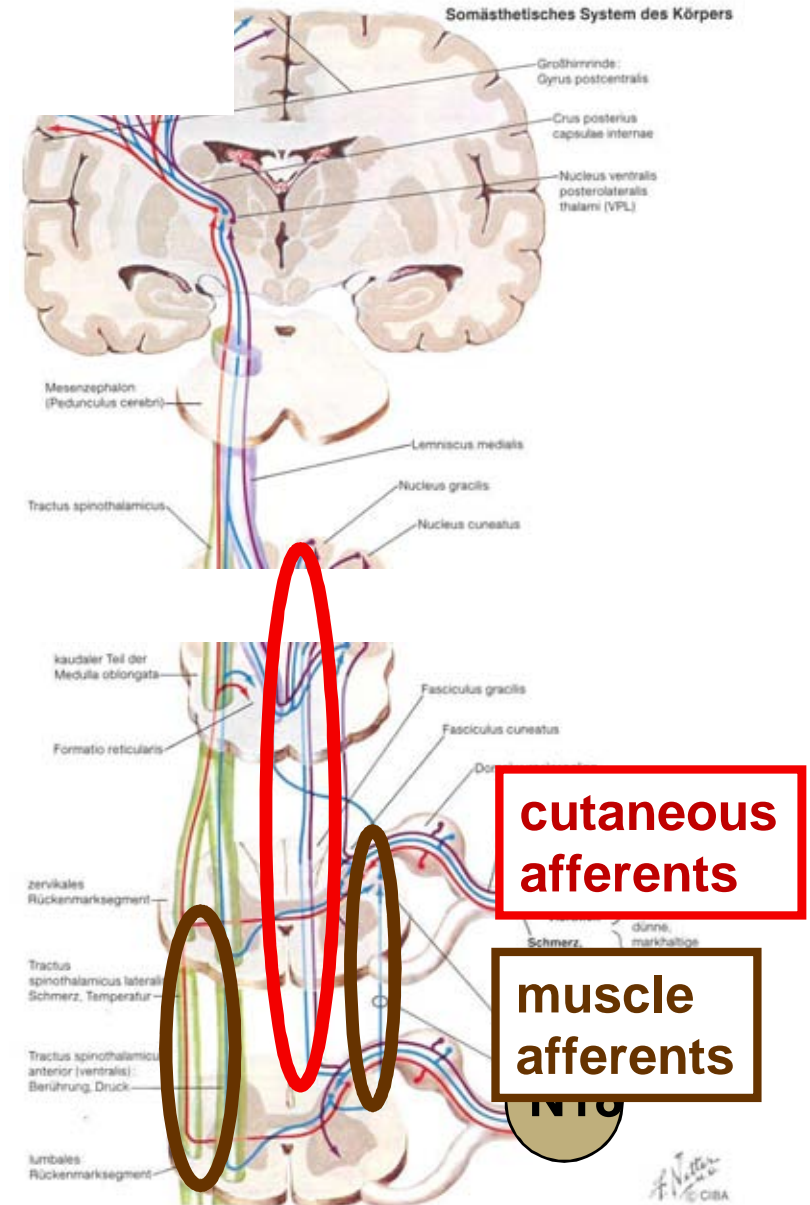




§ Somatosensory evoked potentials

§ Tibial vs. Sural nerve

- Spinal pathway
 - **Tibial nerve**
muscle afferents and tractus spino-cervicalis and / or frontal column
 - **Sural nerve**
Cutaneous afferents and dorsal column

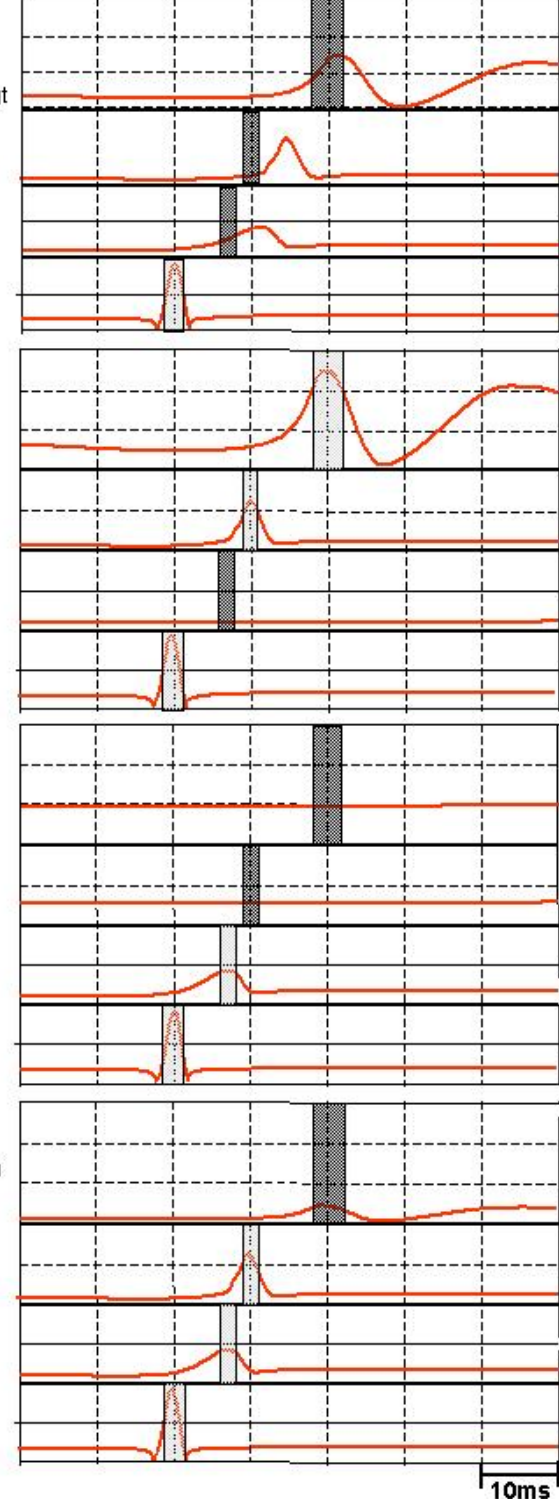
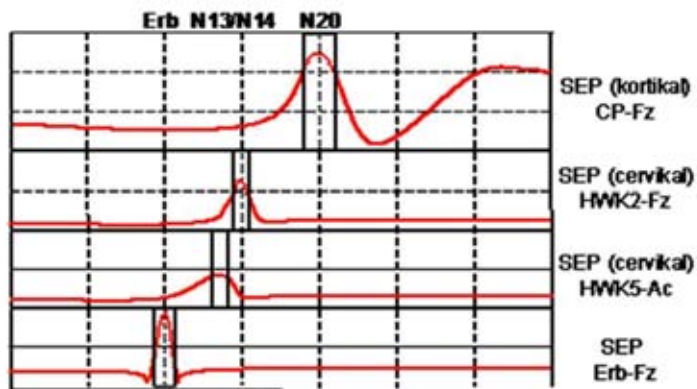


Zervikale
extramedulläre Läsion
N9-N13/N14 verzögert
N20 normal oder verzögert/erniedrigt

Zervikale
intratramedulläre Läsion
N13 ausgefallen

Medulla oblongata Läsion
Ausfall ab N14

Pontine Läsion
Normal bis N14
amplitudengeminderte / Ausfall N20



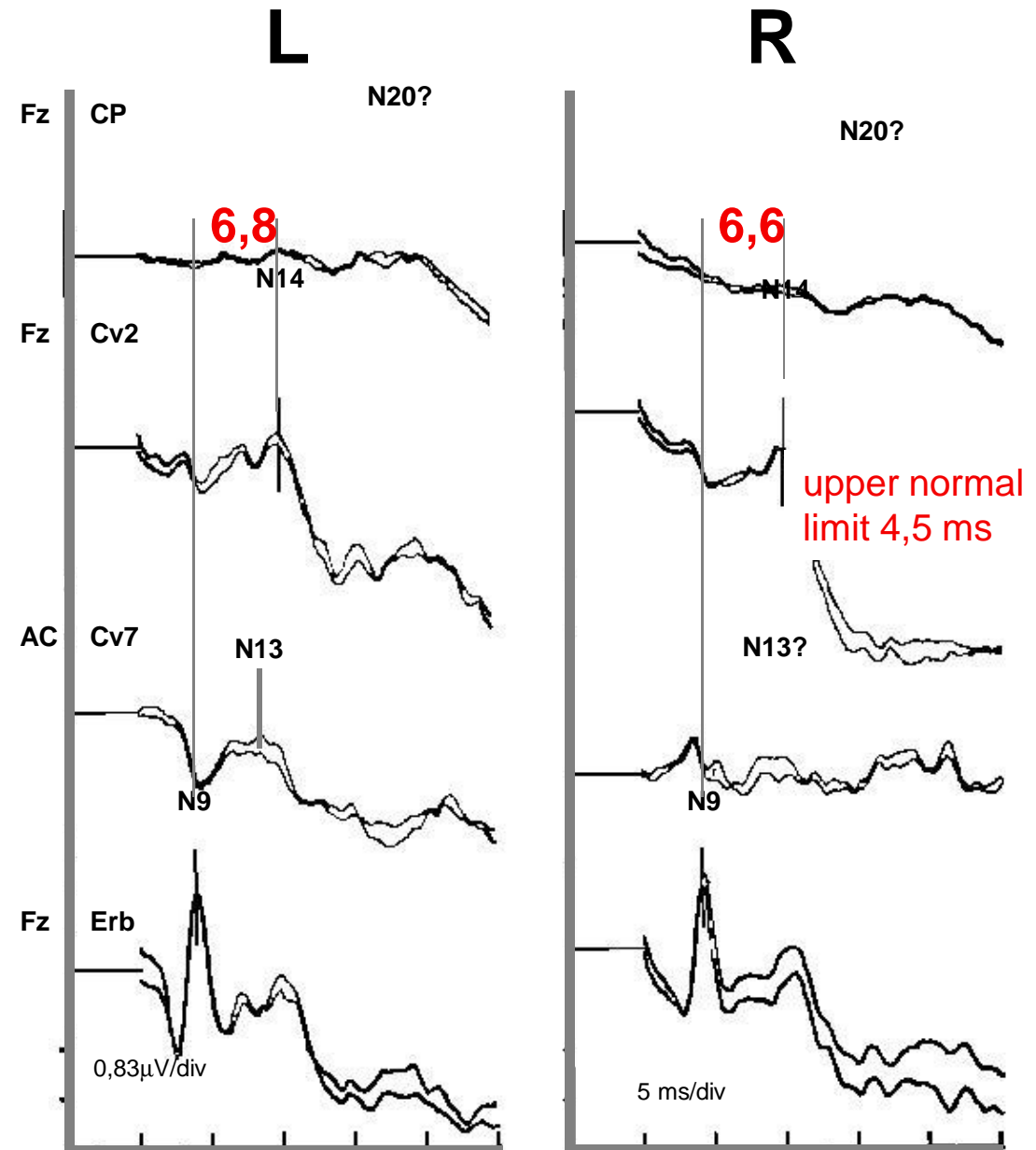
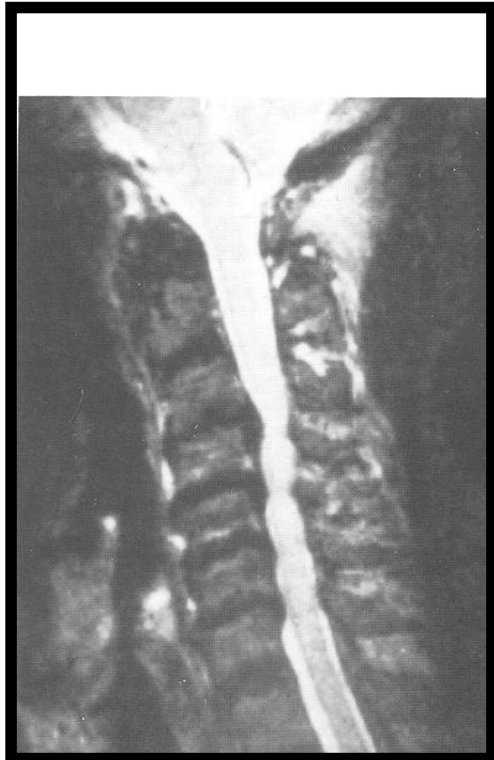
§ Somatosensory evoked potentials



cervical

extra medullary Lesion

- prolonged N9-N14



§ Somatosensory evoked potentials



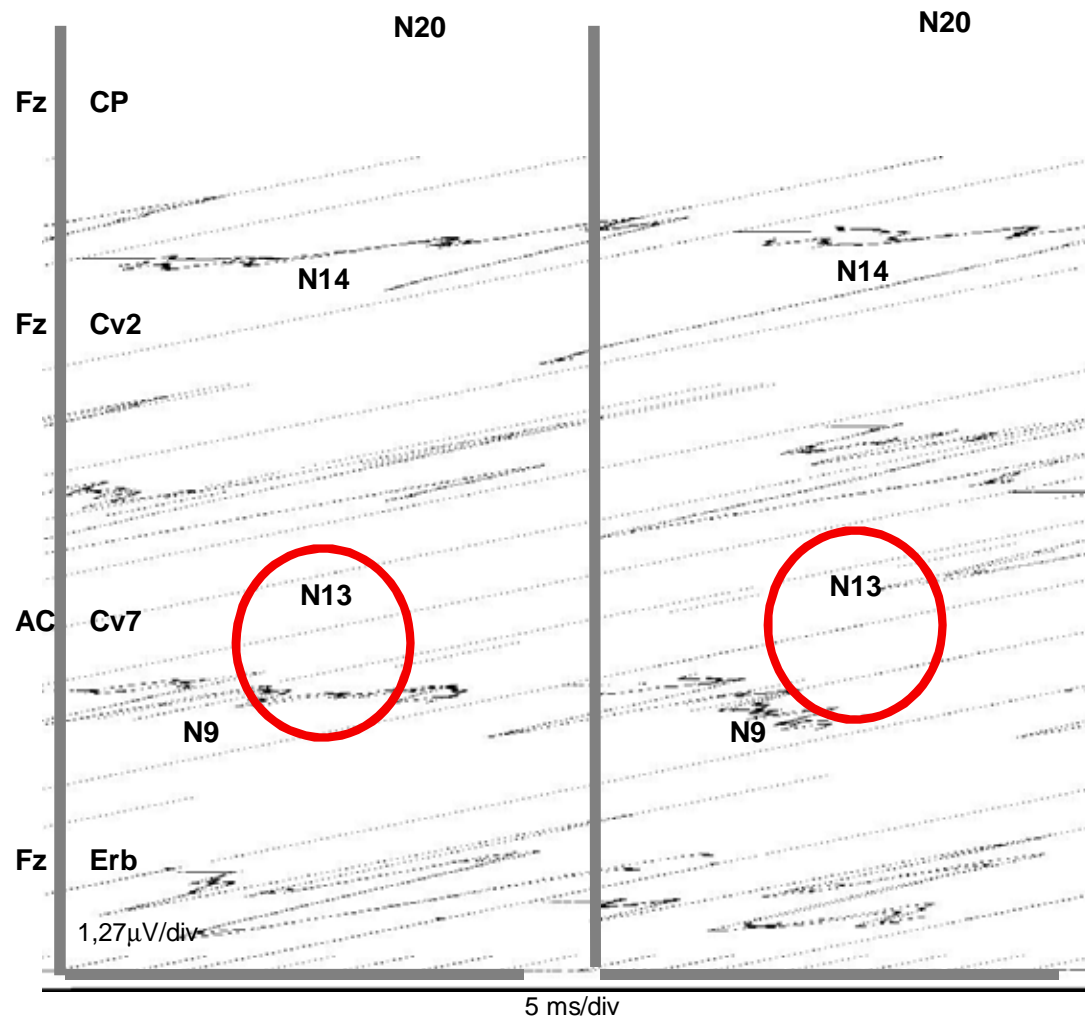
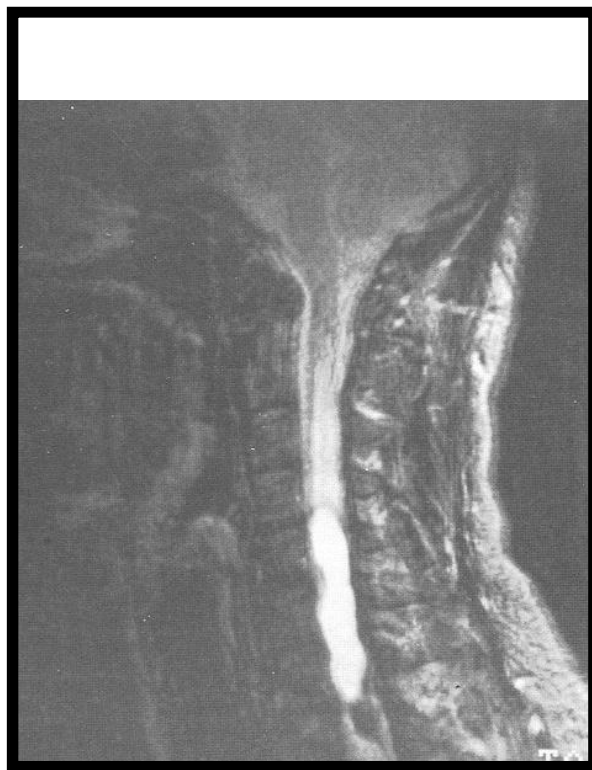
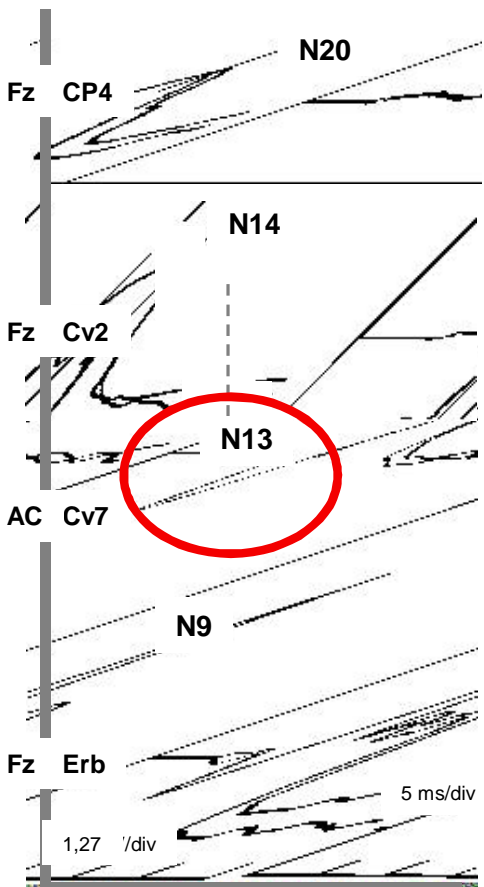
cervical

Intra medullary Lesion

- **loss of N13**

L

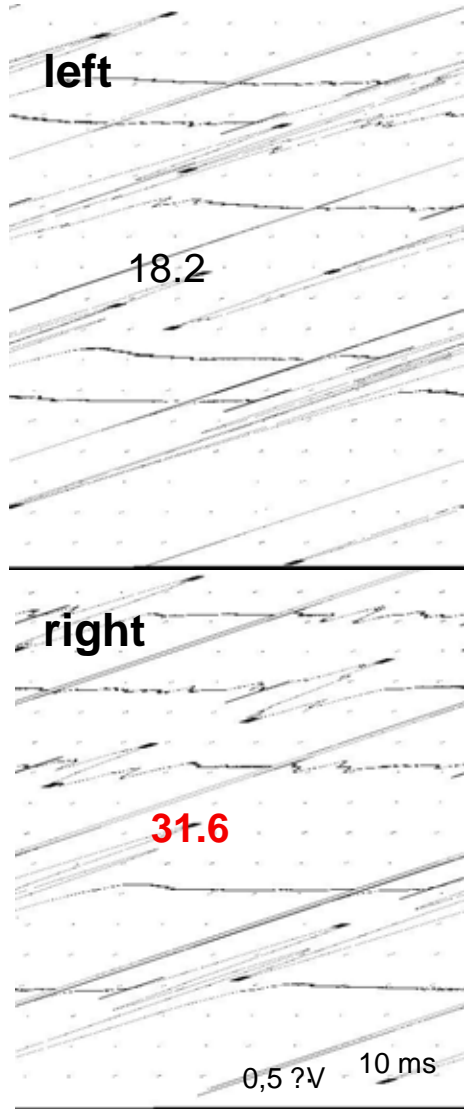
R



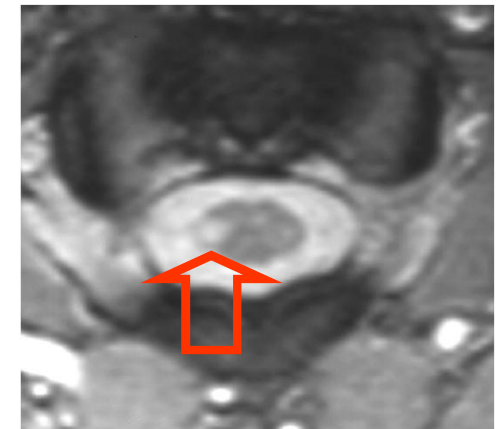
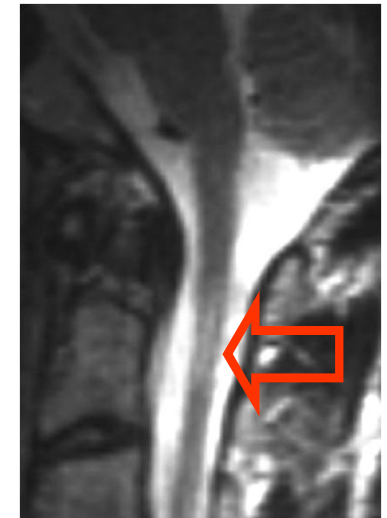
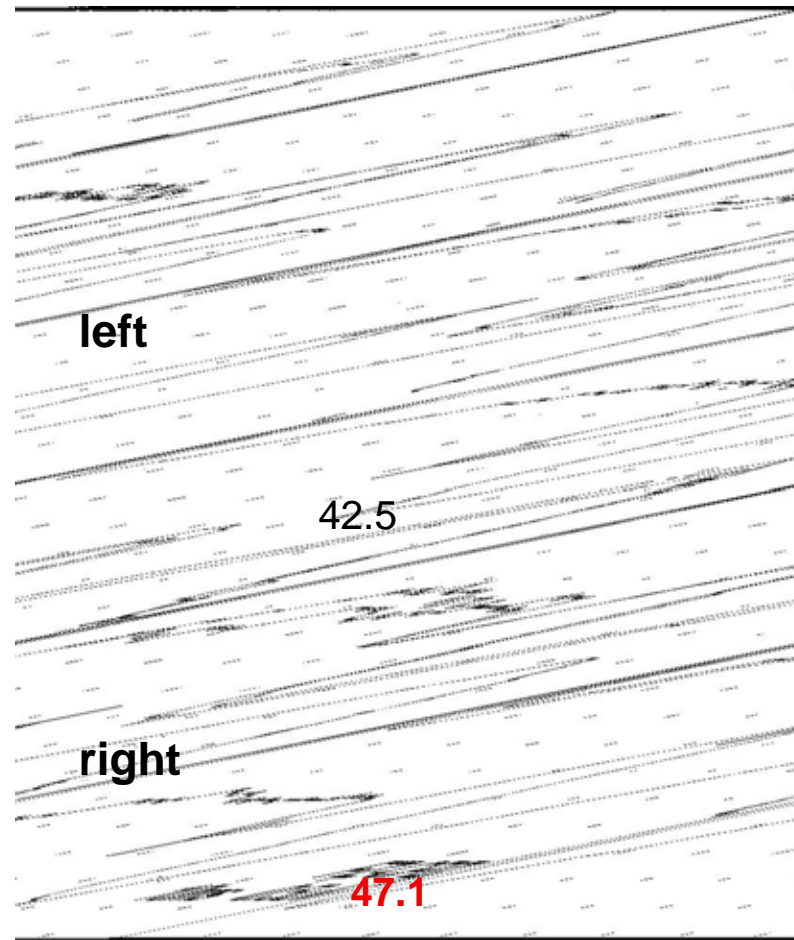
§ Somatosensory evoked potentials



MEP



SEP



§ Somatosensory evoked potentials

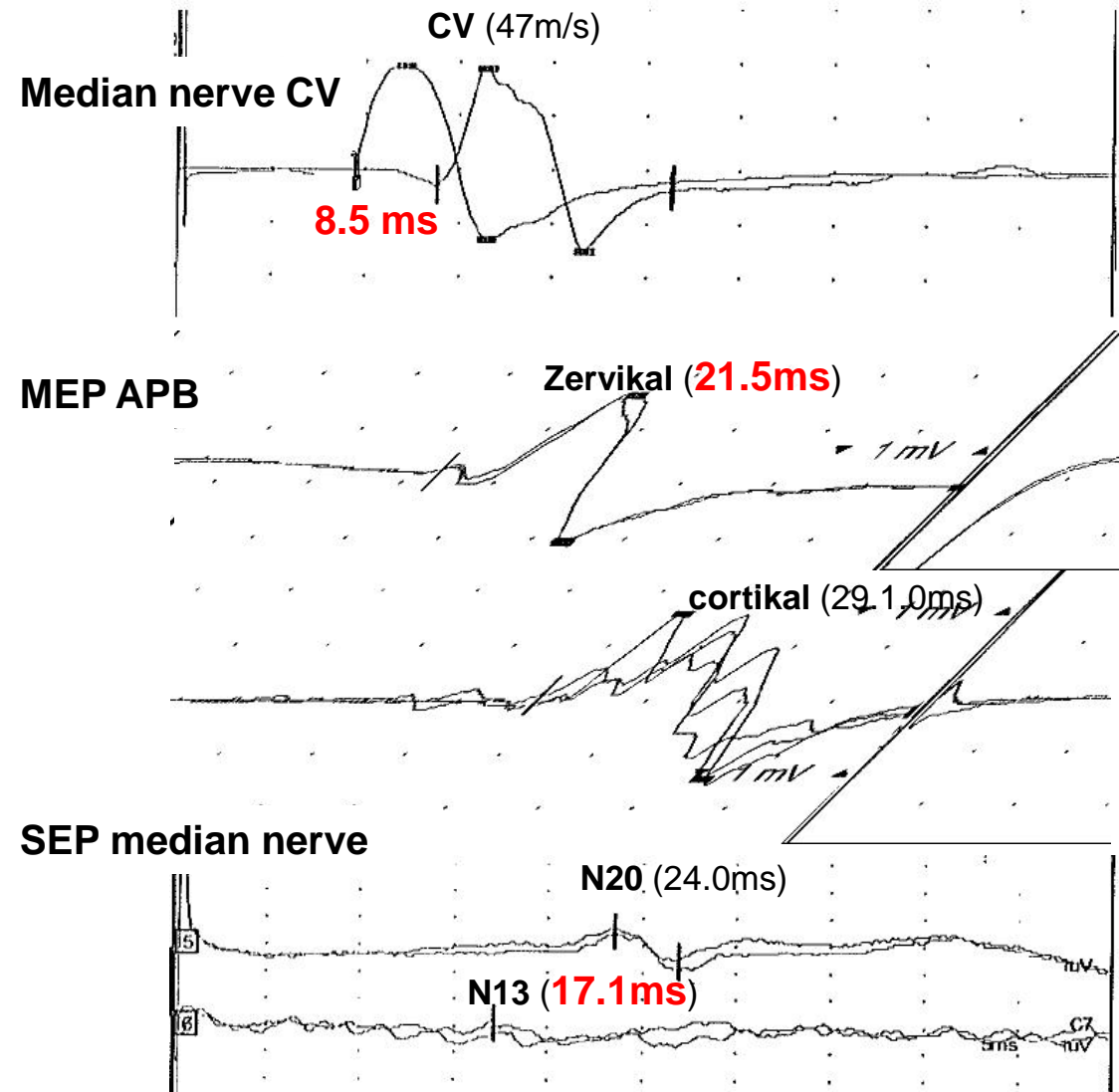


- peripheral nerve lesions
proximal neuropathies

HNPP

Hereditary neuropathy with liability to pressure palsies
17p11.2-p12 deletion

Prominent slowing of
distal and proximal and motor
conduction



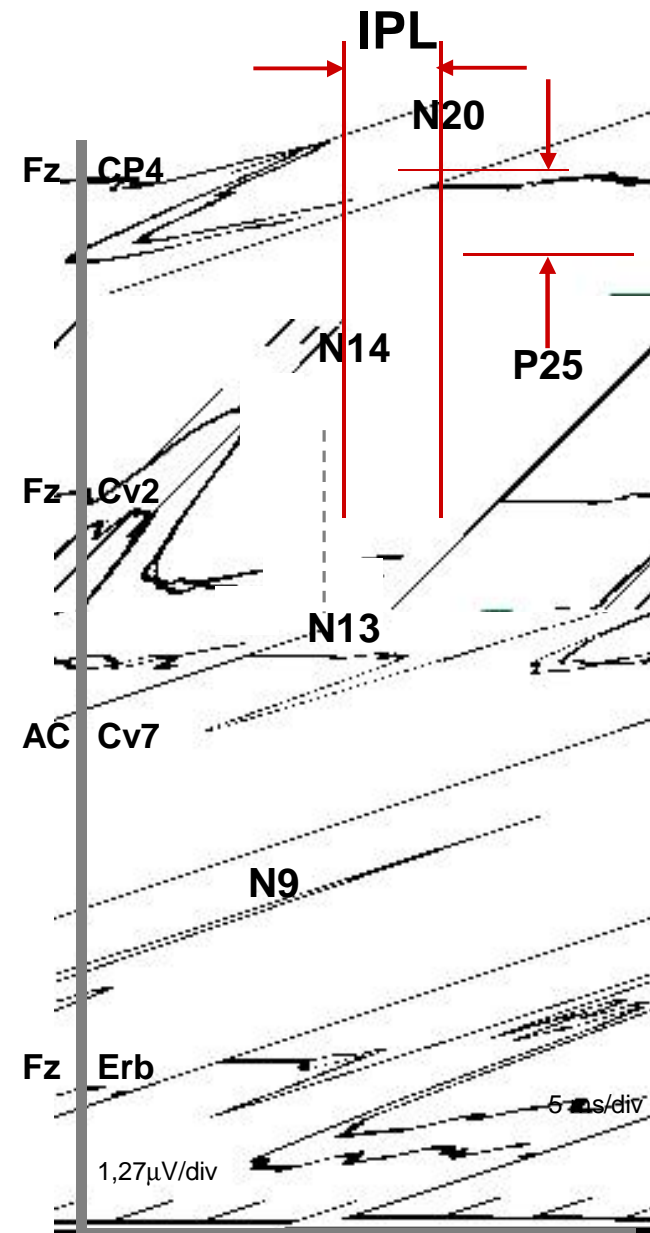


§ Somatosensory evoked potentials

§ Prognosis in coma

- Prolonged N14-N20
- Reduced N20/P25 amplitude
- loss of N20

∅ **bilateral loss N20 =
poor prognosis**





§ Somatosensory evoked potentials

§ Prognosis in coma

Prognoses – outcome – sever **SHT**

N20 – Bilateral loss – Prognosis: wake up <1%

Guérit 2005 Prog Brain Res

Morgalla et al. 2006 Anaesthesist

Amantini et al. 2005 Clin Neurophysiol

Prognoses – outcome – cerebral **Hypoxia**

N20 – Bilateral loss – Prognosis: persistent coma

Tiainen et al. 2005 Crit Care Med

Zandbergen et al. 2006 Neurology

Robinson et al. 2003 Crit Care Med

60 Pat. Hypothermia within 24-28h after Hypoxia –

Prognosis: persistent coma

Tiainen et al. 2005 Crit Care Med

∅ **Recovery from bilateral loss of N20 in rare cases after cardiac arrest and hypothermia**

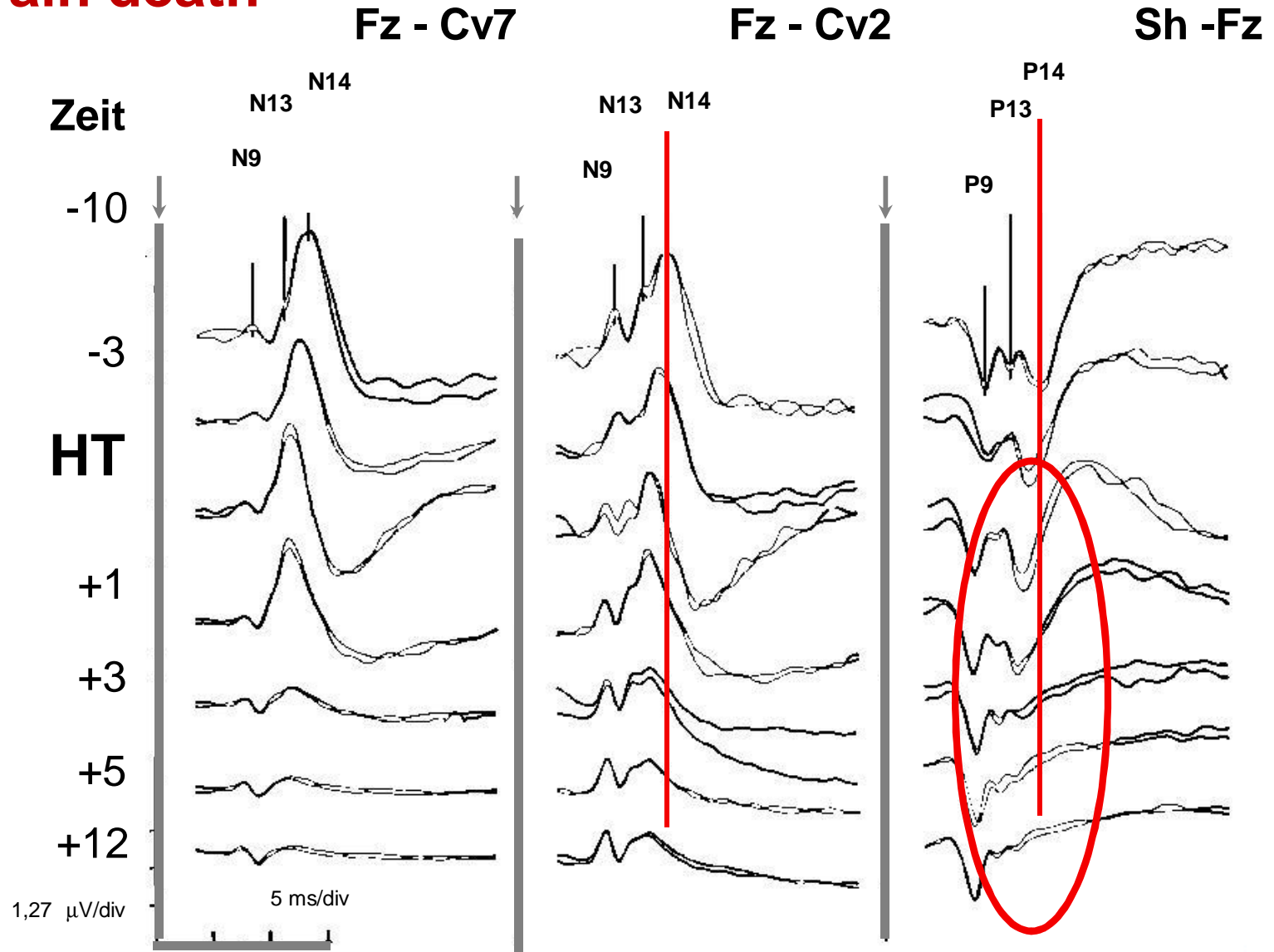
Leithner et. al. Neurology 2010; 74:965-969



§ Somatosensory evoked potentials

§ Diagnosis of brain death

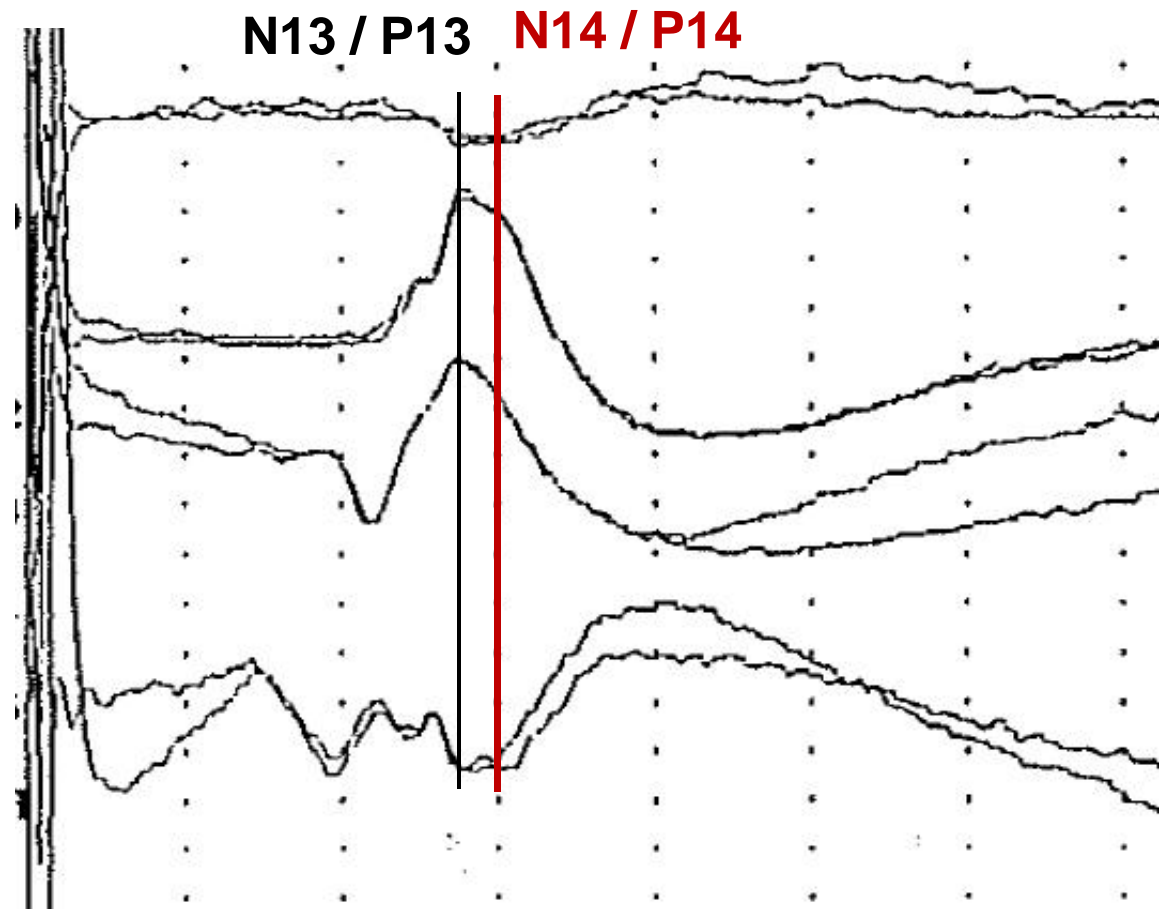
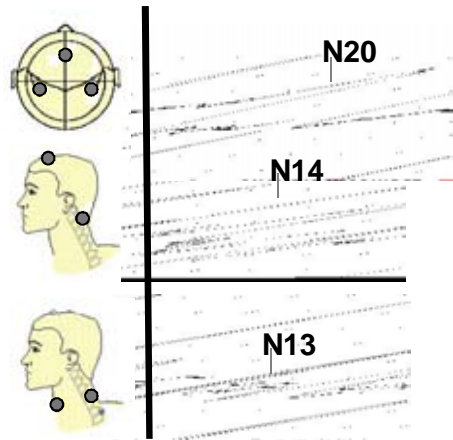
∅ bilateral loss
N14 / P14





§ Somatosensory evoked potentials

§ Diagnosis of brain death

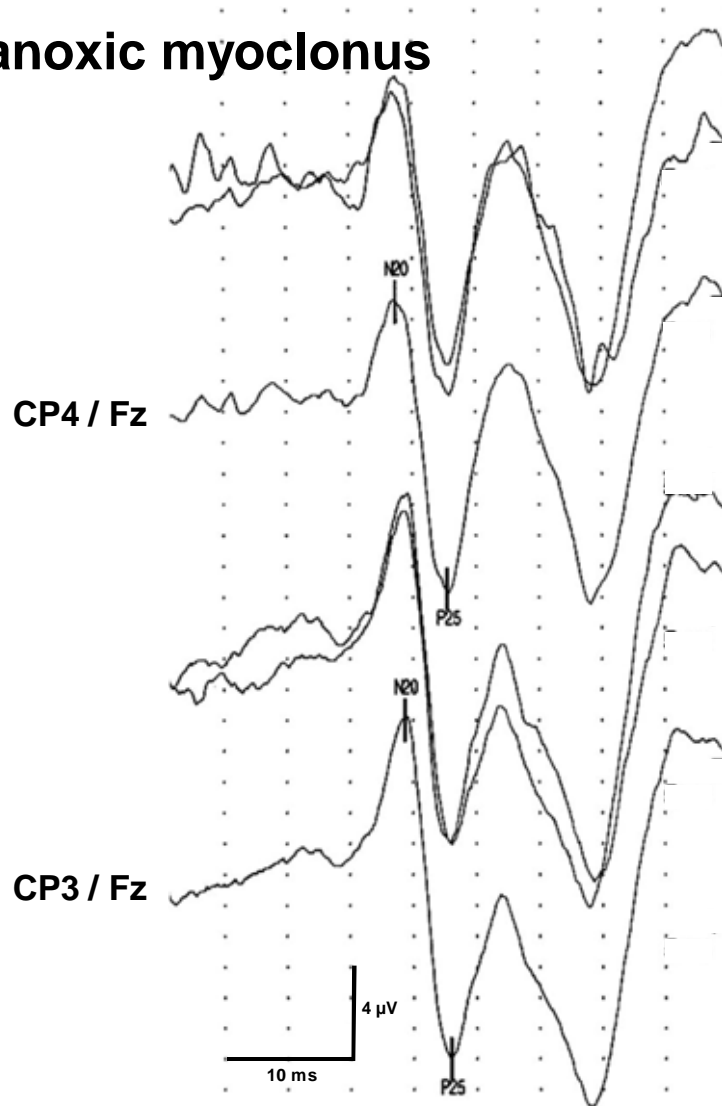




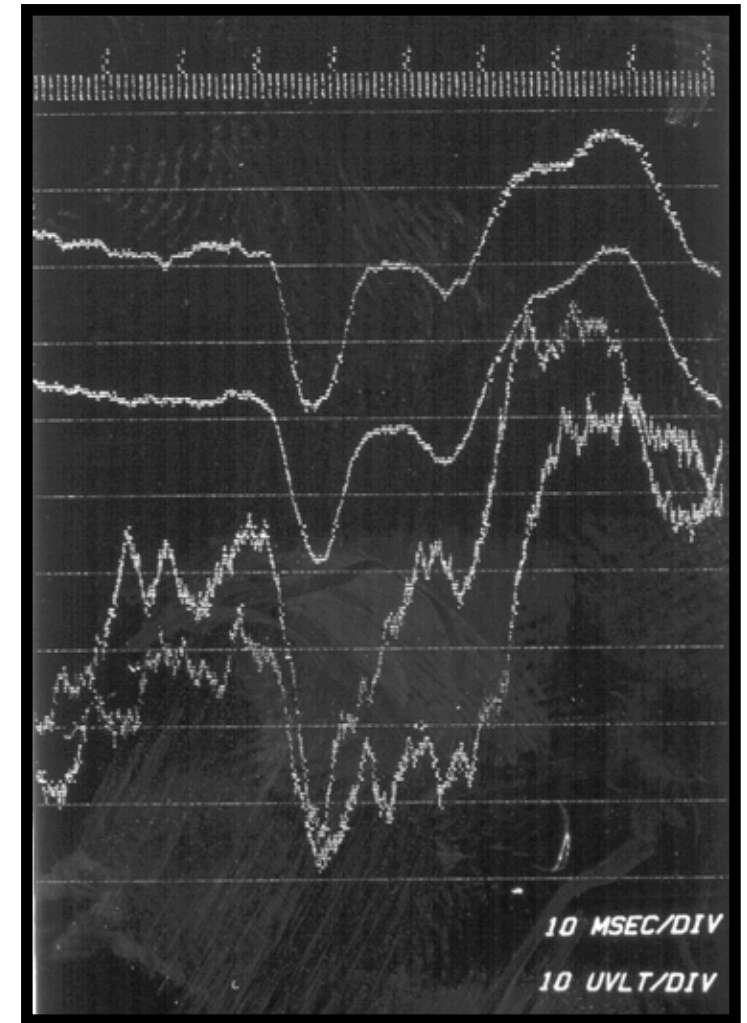
§ Somatosensory evoked potentials

§ Giant SEP

post anoxic myoclonus



myoclonus epilepsy





§ Somatosensory evoked potentials

urologie
Neurophysiologie



§ further SEPs

- **sural - SEP**
 - Additional to tibial nerve SEP in some cases
- **ulnar - SEP**
 - Only in rare questions concerning the brachial plexus
- **dermatomal - SEP**
 - Not reliable
- **trigeminal - SEP**
 - Not reliable



§ Somatosensory evoked potentials

§ Clinical applications and advanced use

∅ Hard rocks

central demyelinating lesions

spinal lesions

MS - Diagnosis

Prognosis in coma

peripheral nerve lesions

proximal neuropathies

intraoperative monitoring / location of central sulcus

cortical myoclonus

∅ Soft rocks

brain death

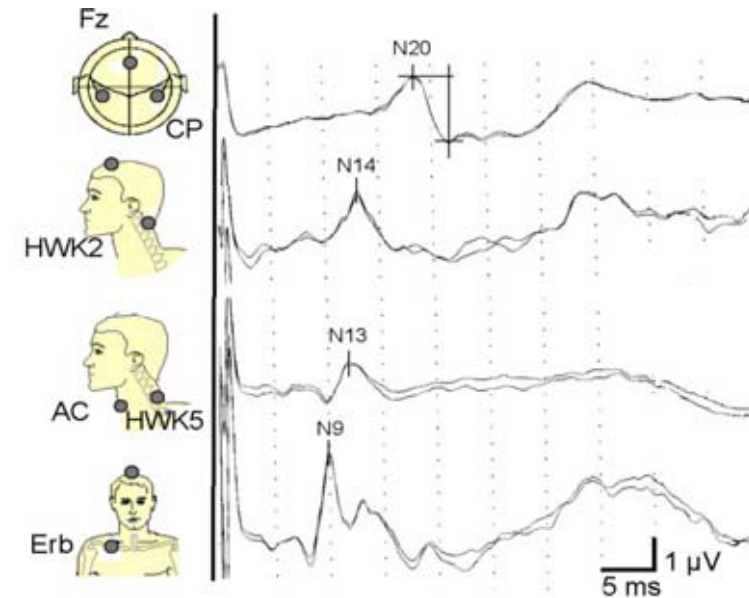
intoxication

MS - History

∅ The beach

stimulation of cutaneous nerves / dermatomal stimulation

pudendal / trigeminal stimulation



multichannel SEP



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