

Principles of Amplitude Integrated EEG (aEEG) & Basic Trace Pattern Qualifications

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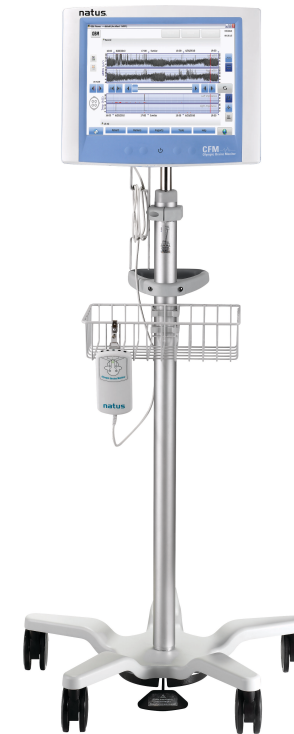
Objectives

Upon completion of this program, the clinician will be able to:

- Describe aEEG trending components including filtering, compression, rectification
- Describe aEEG trace nomenclature and qualifications as they pertain to term babies and premature babies at various gestational ages
- Assess aEEG trace patterns and apply trace nomenclature/qualifications to define aEEG background patterns
- Assess aEEG trace patterns and apply trace qualifications for areas of suspicion/suspected seizures
- Describe how artifact and/or impedance may impact trace purity

What is aEEG?

- aEEG is:
 - 1, 2, or 3 channel bedside brain *monitor*
 - Basic neurologic function trending tool
 - Long-term monitoring capability
 - Used to measure global electro-cortical activity or specific site brain activity
 - Developed by Neonatologists, for Neonatologists
 - Complimentary tool to quickly obtain information regarding the baby's neurological status



Monitoring Tools in the NICU



What Do We Want to Know When We Monitor the Brain with aEEG?

- **What is the neurological status of the patient?**
 - Is there cerebral injury?
 - What is the severity of the injury?
 - What changes are occurring over time?
 - Is there improvement or worsening of the neurological status
 - What is the impact of NICU treatments to the patient's brain function?
- **Is the patient having seizures?**
 - What is causing the seizures?
 - Are the seizures occurring more frequently, or for longer/shorter duration?
 - Are the seizures responding to medical therapy?
 - Is there electromechanical disassociation after medication?

Who Should Be Monitored? | Clinical Applications

- **Infants that have experienced a sentinel event during delivery and are at risk for hypoxic ischemic encephalopathy (HIE):**
 - Low Apgar
 - Low pH
 - Required resuscitation or artificial ventilation at birth
 - Poor tone/poor reflexes
- **Infants receiving hypothermia treatment for HIE**
- **Infants with definite or questionable seizures (clinical or subclinical):**
- **Infants with unexplained neurological symptoms (i.e. severe apnea)**

*Thoresen M, Hellstrom-Westas L, Liu X, de Vries L.
“Effect of Hypothermia on Amplitude-Integrated
Electroencephalogram in Infants With Asphyxia”.
Pediatrics published online June 21, 2010;
DOI: 10.1542/peds.2009-2938



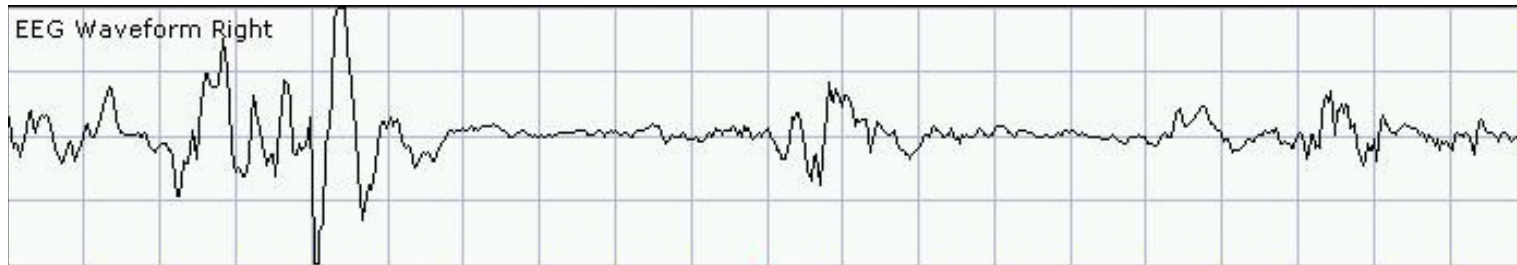
- **Infants who are at higher risk for cerebral complications due to circulatory instability**
 - Sepsis
 - Hypoxia
 - Persistent pulmonary hypertension
 - Meconium aspiration
 - Cardiac malformations
 - Diaphragmatic hernia
- **Additional clinical applications**
 - Muscle relaxed/neuromuscular blockade
 - Grade 3 or 4 IVH
 - ELBW infants
 - Inborn errors of metabolism (e.g. urea cycle disorders, hypoglycemia, hypocalcemia)
 - Neonatal abstinence syndrome (e.g. alcohol/opiate withdrawal)
 - Post surgical
 - Post cardiac arrest
 - Infants requiring ECMO or surgery for CHD

NeoReviews Vol 7 No. 2
February 2006
Hellstrom-Westas, Rosen, deVries, Greisen

Breakdown - How Does aEEG Work?

- **aEEG** (“a”=amplitude integrated / EEG = electroencephalography):
 - One, two, or three channels of EEG that go through a number of modifications:
 - special filtering
 - rectification
 - compression
 - very slow, trend display
 - aEEG is a process of taking a raw EEG, modifying it, and producing a trending pattern that allows clinicians to measure and view the microvoltage of the brain over time

Background Information - Channel



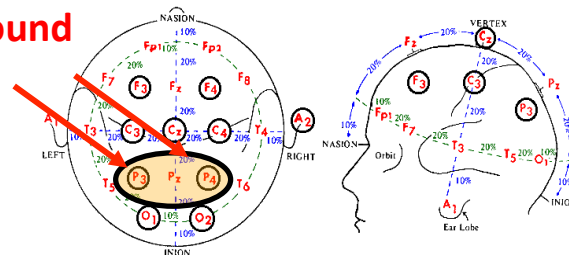
- Two electrodes are needed to create a single channel
- EEG waves reflect electrical voltage differences between these two electrodes sites
 - Measured in microvolts (μV)

aEEG Channels & the 10-20 System

- The Olympic Brainz Monitor may be used to monitor and record aEEG patterns through either:

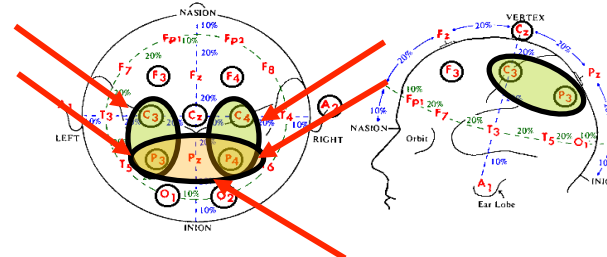
- **Cross-Cerebral (default mode)**

- **3 electrodes - 2 active & 1 hydrogel ground**
- 1 aEEG channel (P3/P4)
- 1 EEG channel (P3/P4)



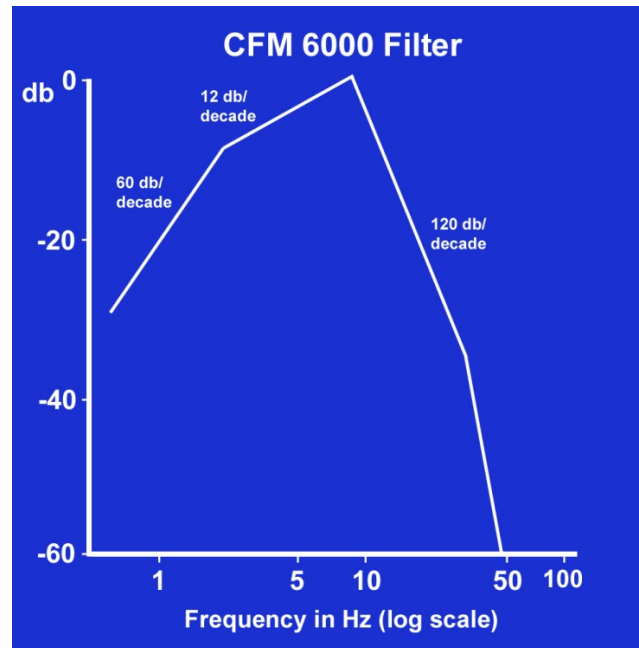
- **Bilateral**

- **5 electrodes – 4 active & 1 hydrogel ground**
- 3 aEEG channels (C3/P3, C4/P4, P3/P4)
- 3 EEG channels (C3/P3, C4/P4, P3/P4)

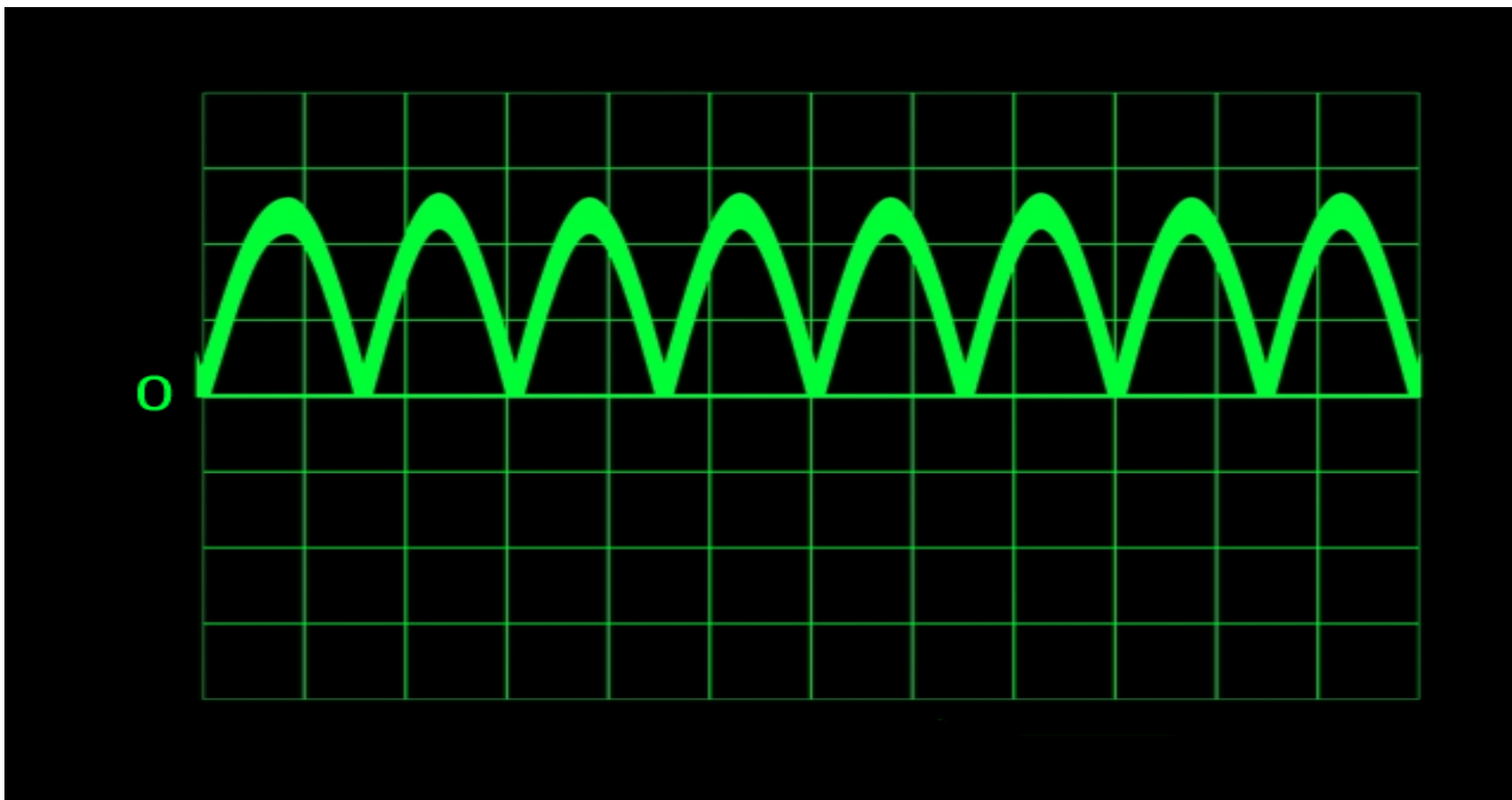


Filtering

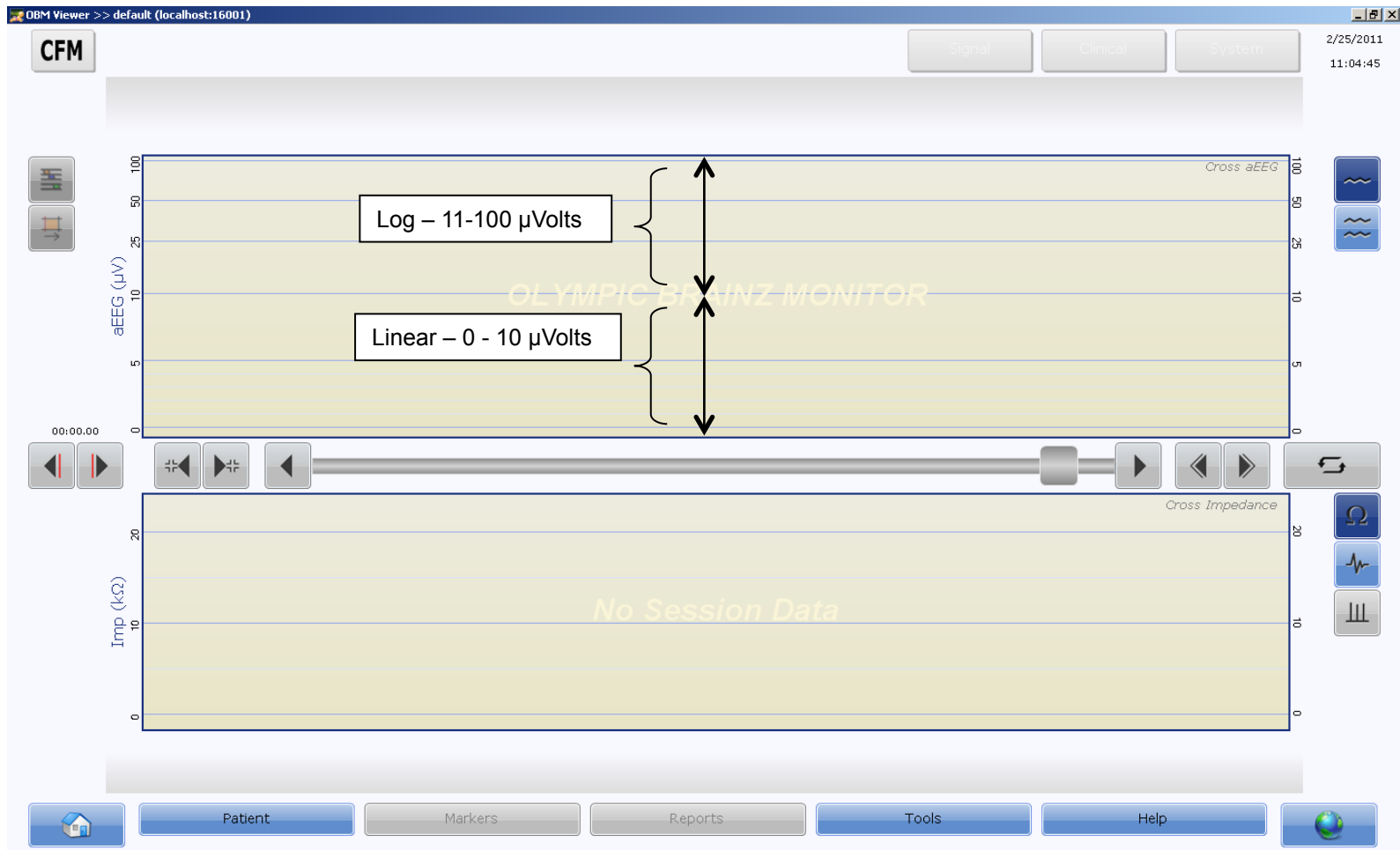
- The EEG signal is filtered 2–15 Hz
- Specially shaped filter
- Reduces muscle and other artifacts



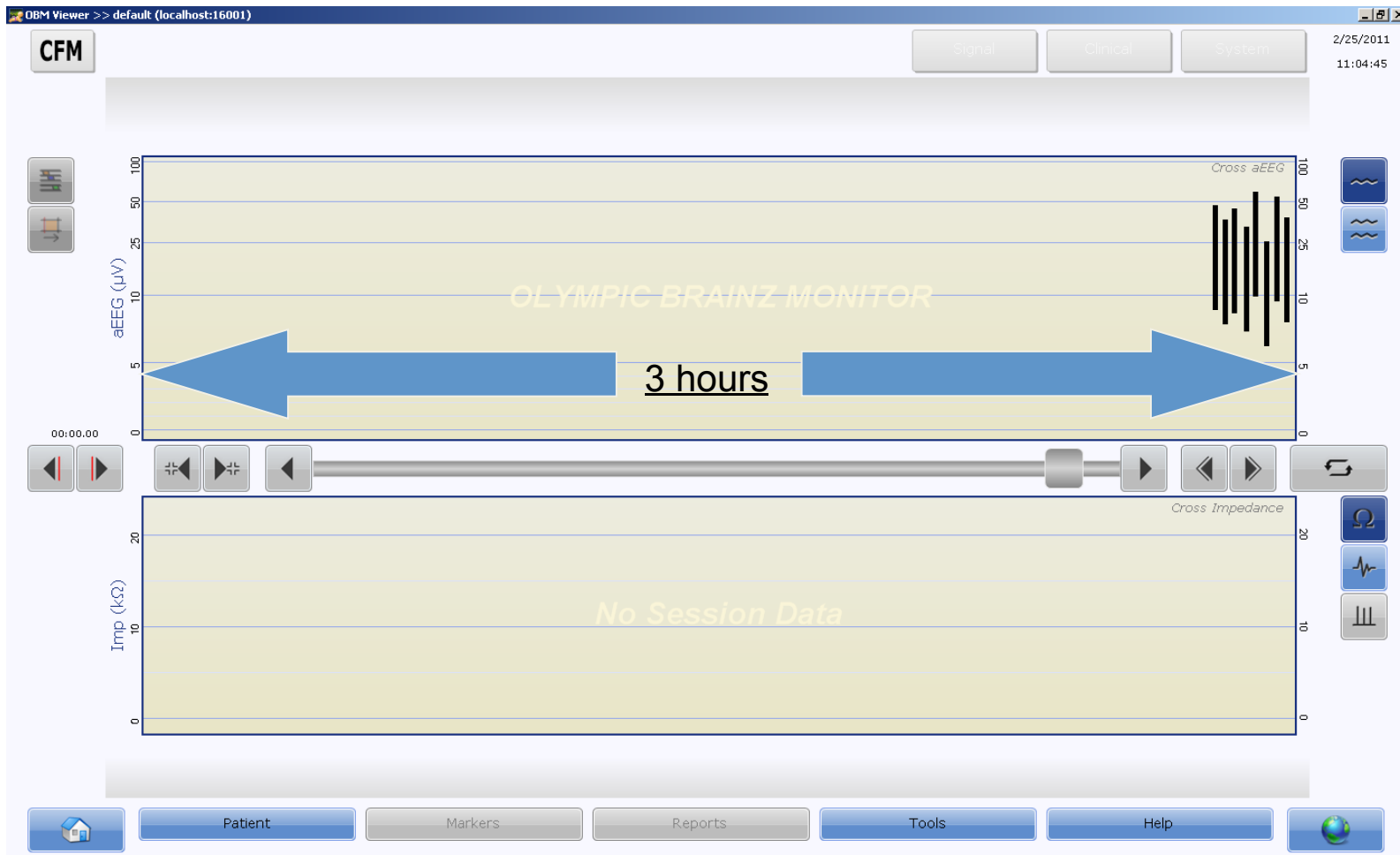
Rectification



Compression



Very Slow, Trend Display

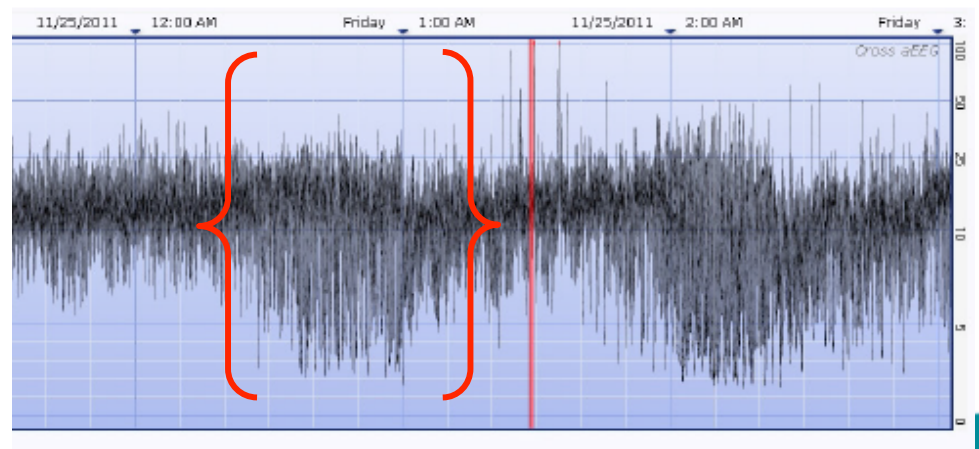


Background Information - Margins



Background Information – Sleep Wake Cycling (SWC)

- **SWC characterized by:**
 - Smooth sinusoidal variations, mostly in the lower margin
 - Broader bandwidth represents discontinuous background activity during quiet sleep
 - More narrow bandwidth corresponds to more continuous activity during wakefulness and active sleep
 - Quiet Sleep Cycle duration ≥ 20 minutes
 - Total SWC $\sim 60-90$ minutes



aEEG Classification Framework

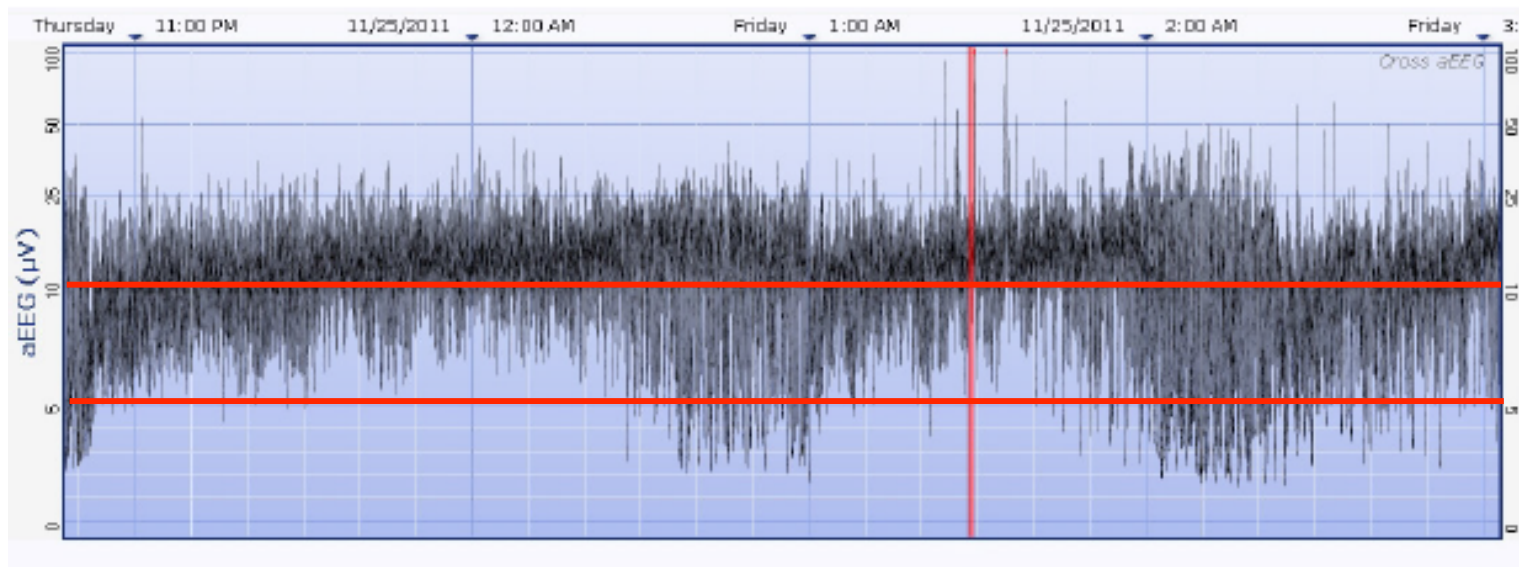
Feb 2006 – NeoReviews – Hellstrom Westas

<i>Pattern Definition (Hellstrom-Westas & Toet)</i>	<i>Lower Margin (in μV)</i>	<i>Upper Margin (in μV)</i>
Continuous Normal Voltage	> 5	>10
Discontinuous Normal Voltage	<5	>10
Burst Suppression	<5	>10 due to high voltage bursts
Continuous Low Voltage	<5	<10
Isoelectric/Flat	< 5	<5

aEEG and TERM Babies

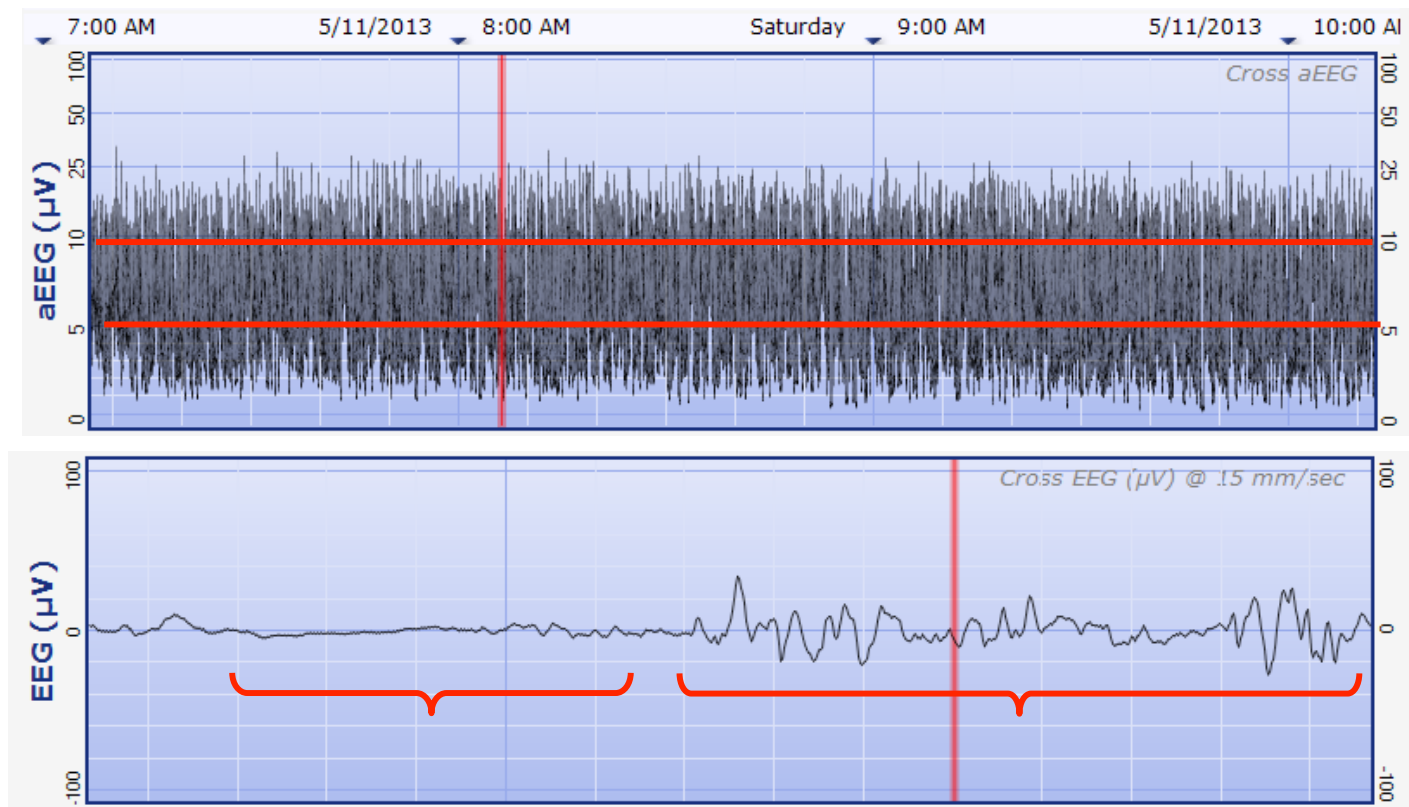
Continuous Normal Voltage

- Sleep/Wake Cycling
- Upper Margin $> 10 \mu\text{Volts}$
- Lower Margin $> 5 \mu\text{Volts}$
- Limited Bandwidth Variability (between upper and lower margin)
 - $\sim 5\text{-}10 \mu\text{Volts}$

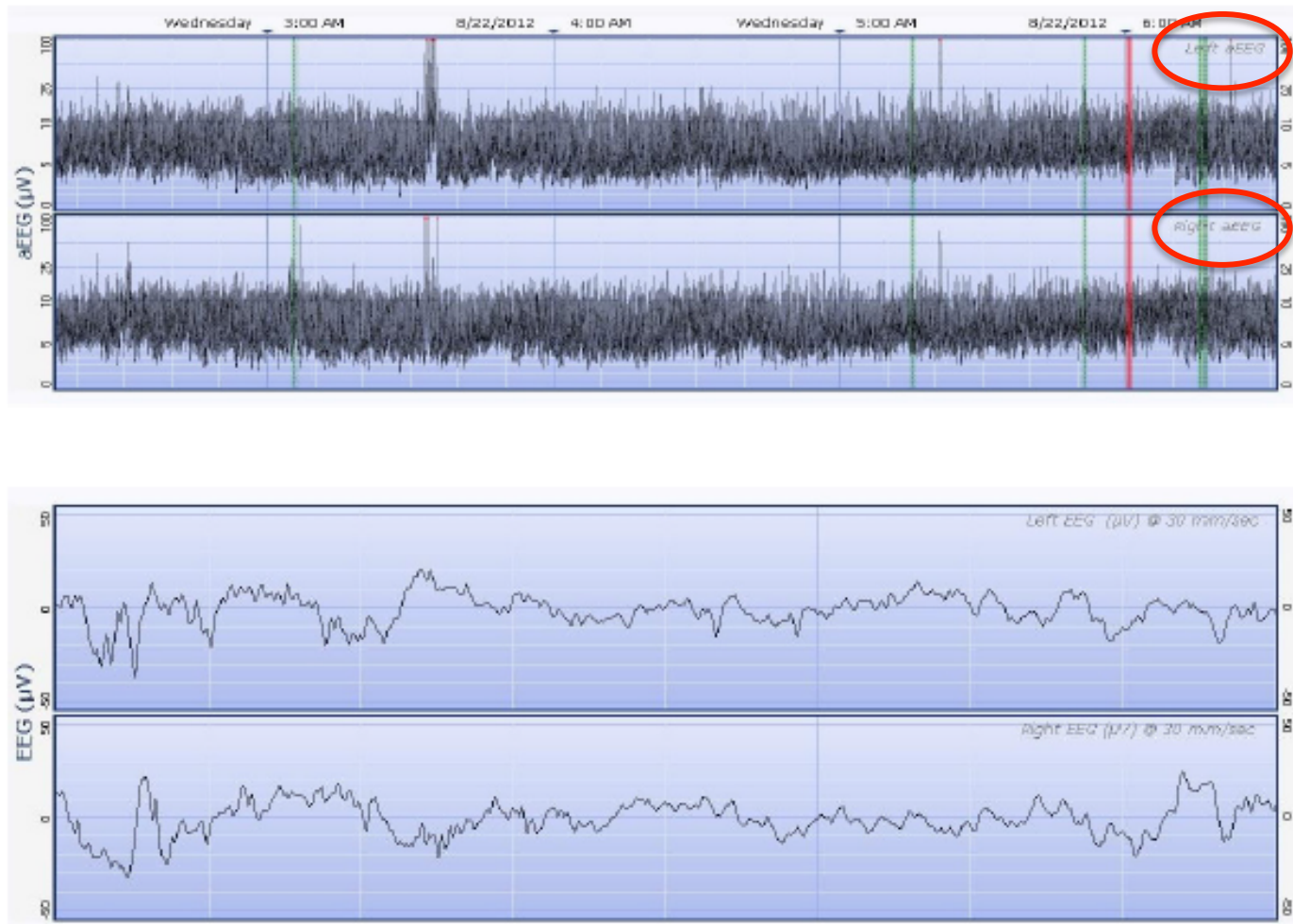


Discontinuous Normal Voltage

- No Sleep/Wake
- Upper Margin $> 10 \mu\text{Volts}$
- Lower Margin $< 5 \mu\text{Volts}$
- Increased Bandwidth Variability
 - $\sim 30 - 40 \mu\text{Volts}$

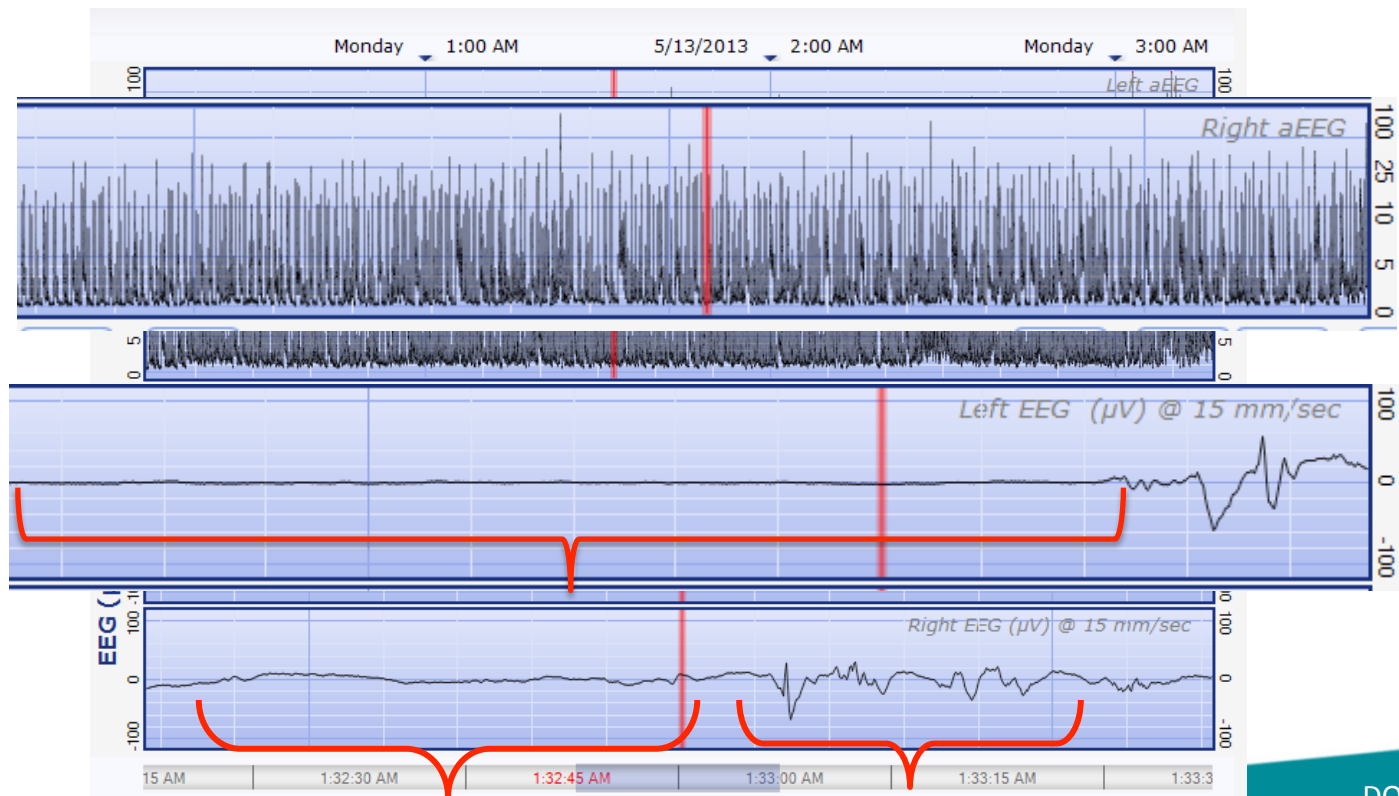


Discontinuous Normal Voltage



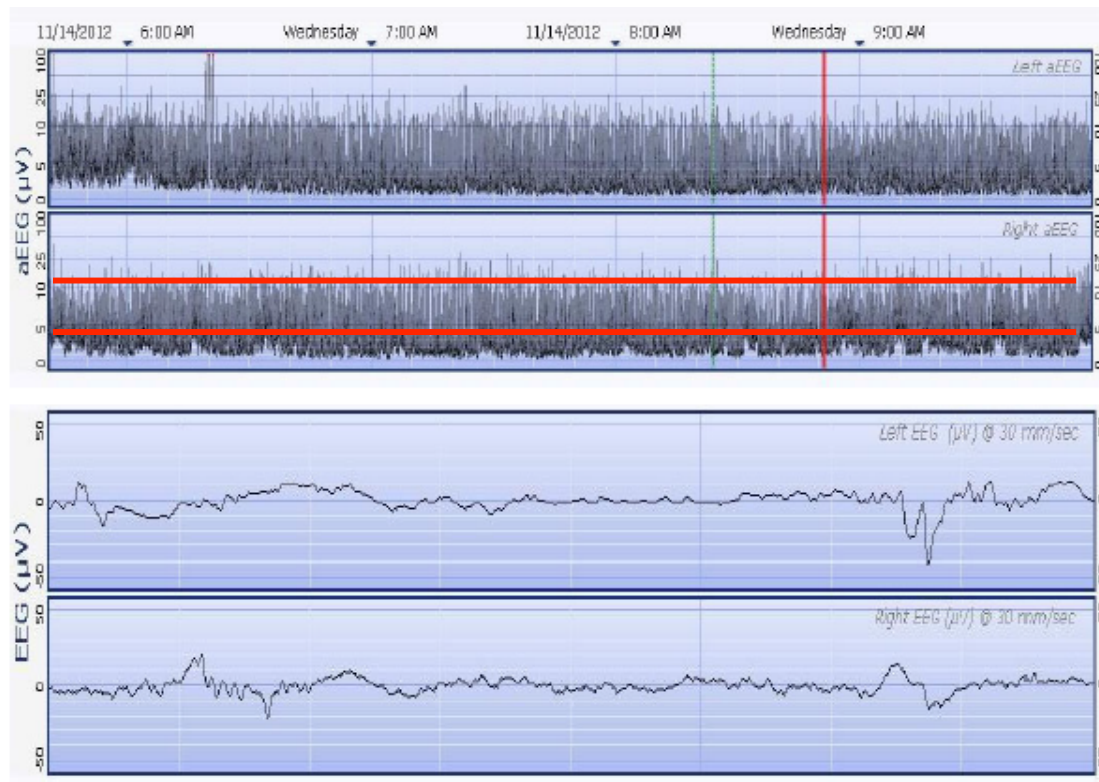
Burst Suppression

- No Sleep Wake Cycling
- Upper margin $>10\mu\text{V}$ (due to high voltage bursts)
- Lower margin $<5\mu\text{V}$
- Limited variability of lower margin



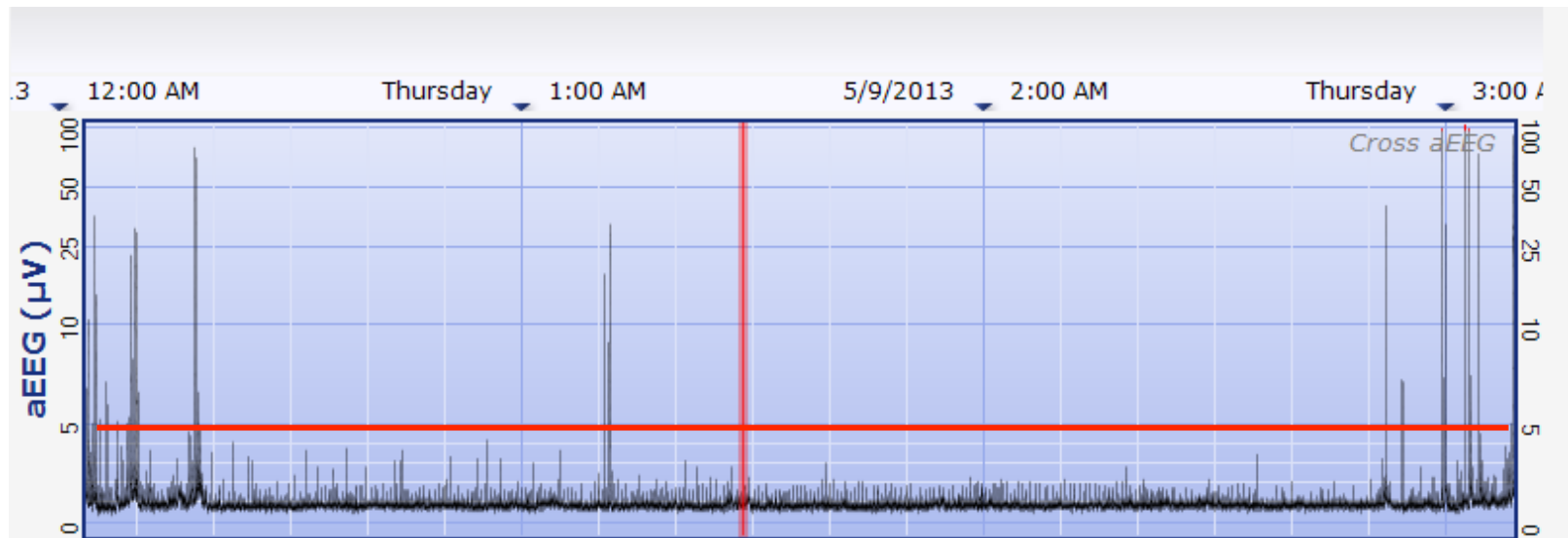
Continuous Low Voltage

- No Sleep/Wake Cycling
- Upper margin <10uV
- Lower margin <5uV

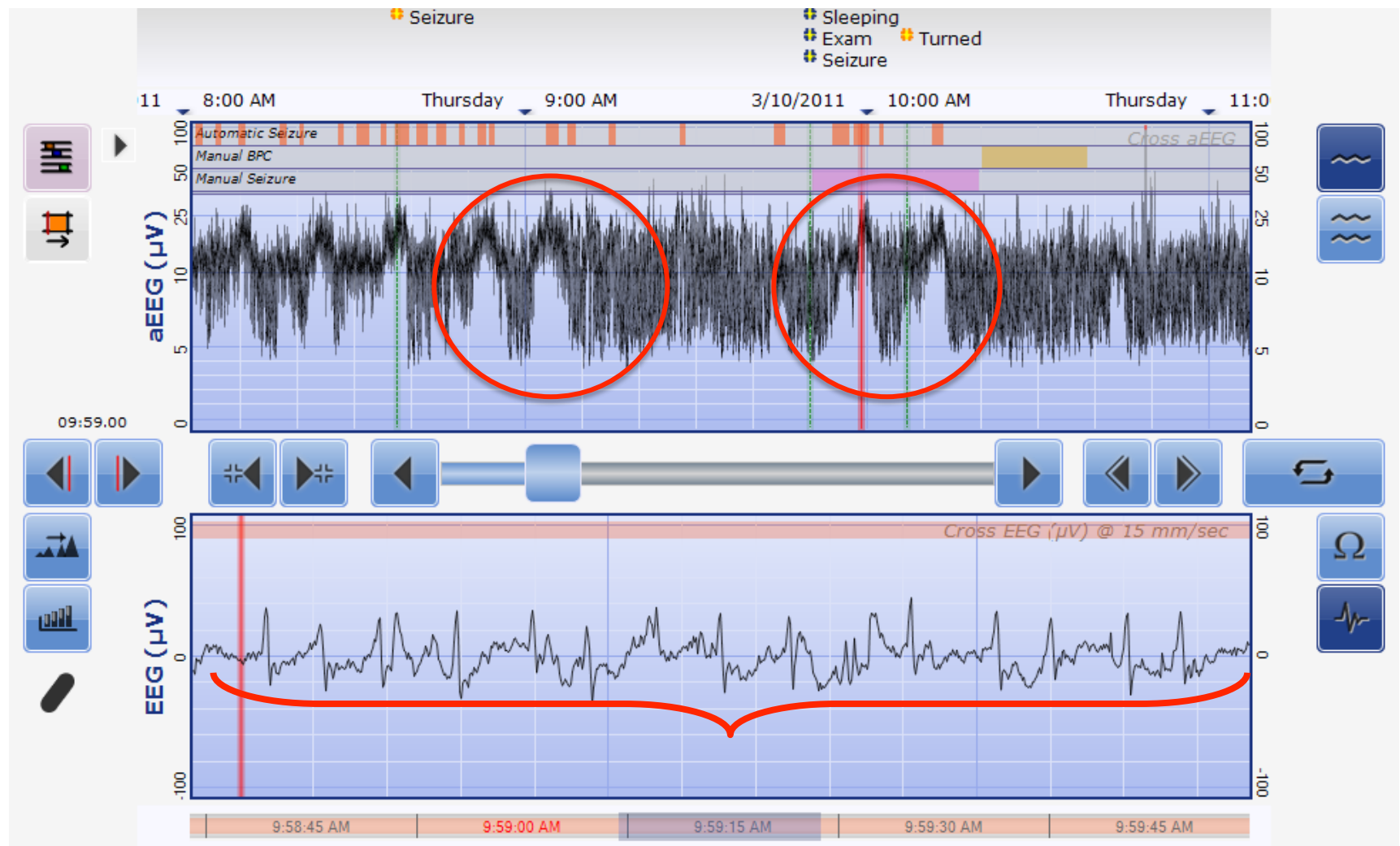


Isoelectric or Flat

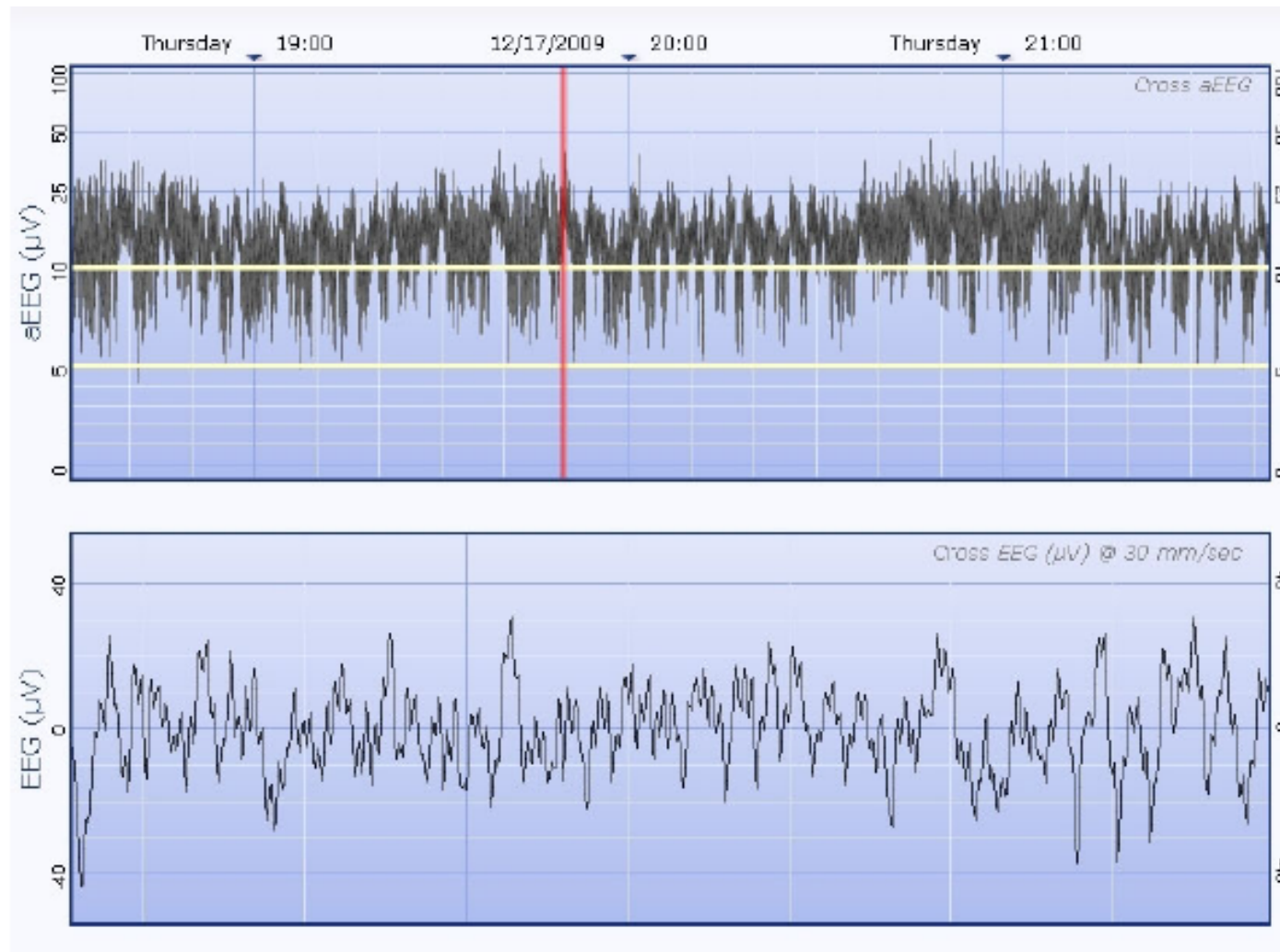
- No Sleep/Wake
- Upper Margin $< 5 \mu\text{Volts}$
- Greatly reduced bandwidth variability
 - $\sim 1 \mu\text{Volt}$



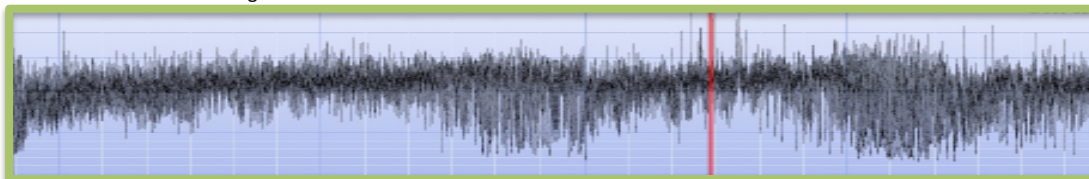
Seizure EEG



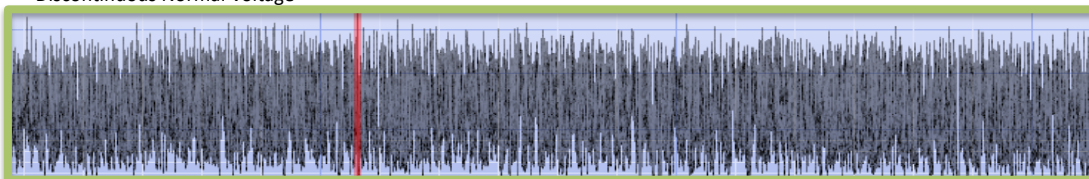
Status Epilepticus



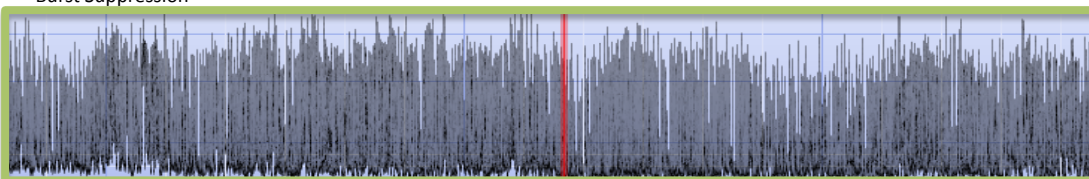
Continuous Normal Voltage



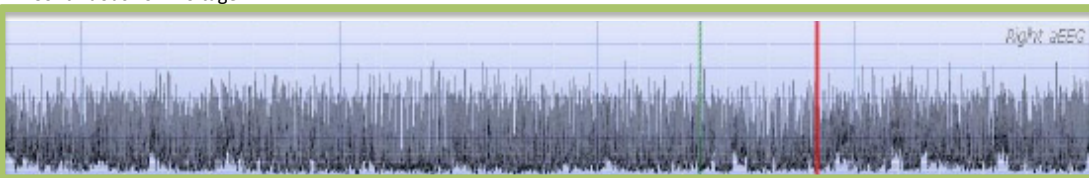
Discontinuous Normal Voltage



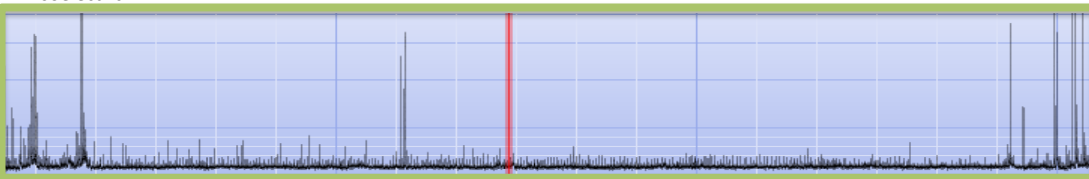
Burst Suppression



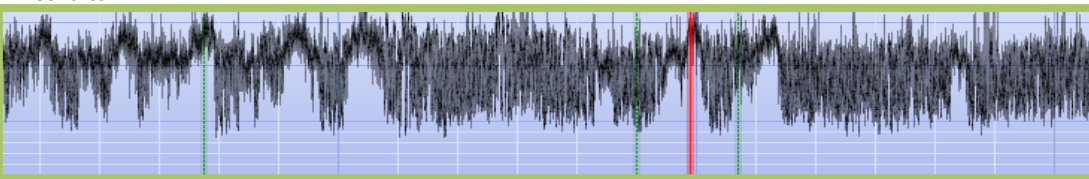
Continuous Low Voltage



Isoelectric



Seizures



Impedance and Artifact

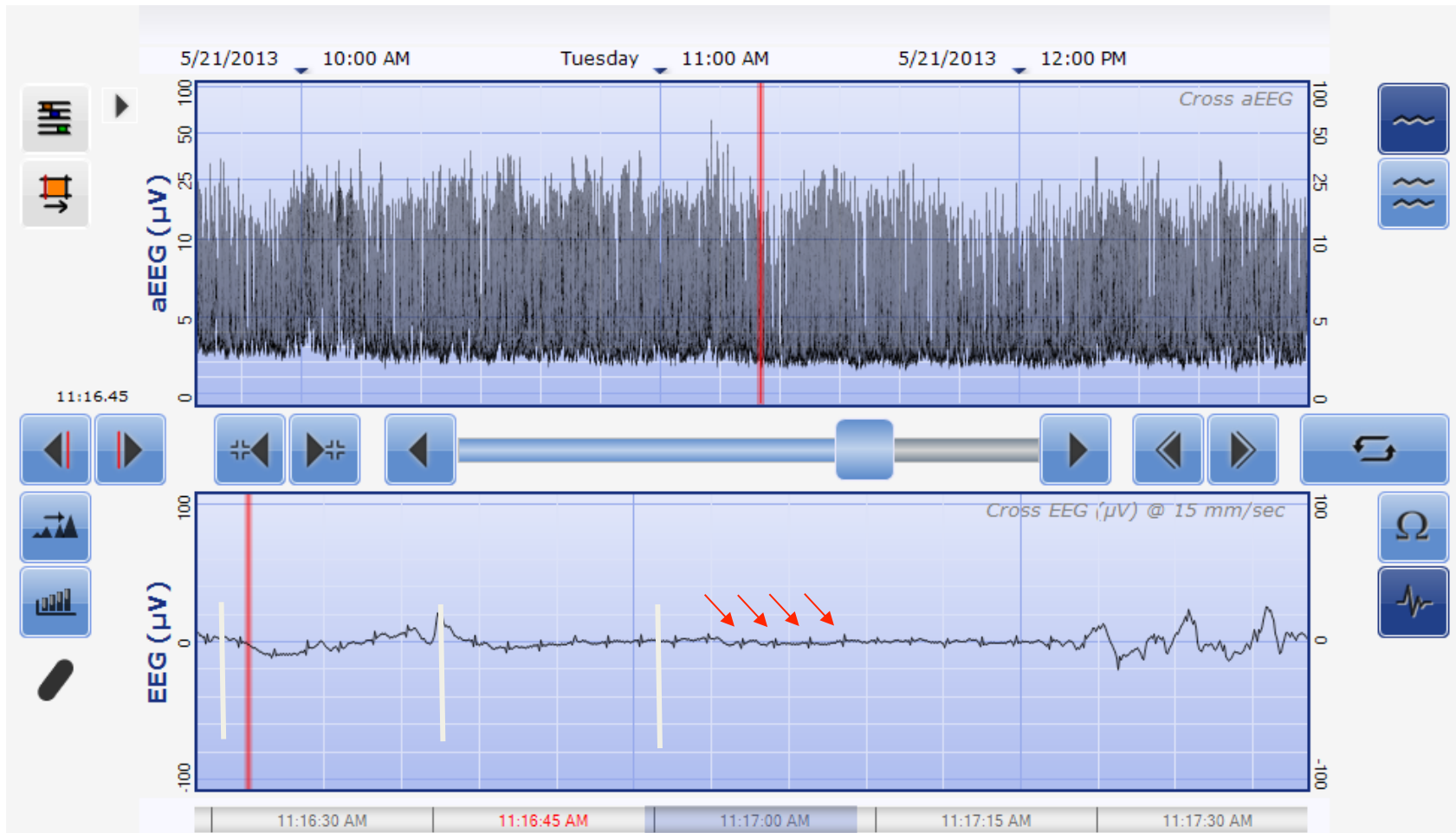
- **Impedance**

- A measure of the quality of electrode contact
- Anything that gets between the sensor (hydrogel or low impedance needles) and “impedes” or interferes with the devices ability to read the brain signal (hair, dry skin, vernix)

- **Artifact**

- Any electrical activity other than the brain’s electrical activity (monitors, IV pumps, ventilators, etc.)
- Live EEG signal is used as a point of reference to confirm suspected brain activity OR to distinguish artifact from the real signal

ECG Artifact



aEEG and Premature Babies

Pre-Term Infants

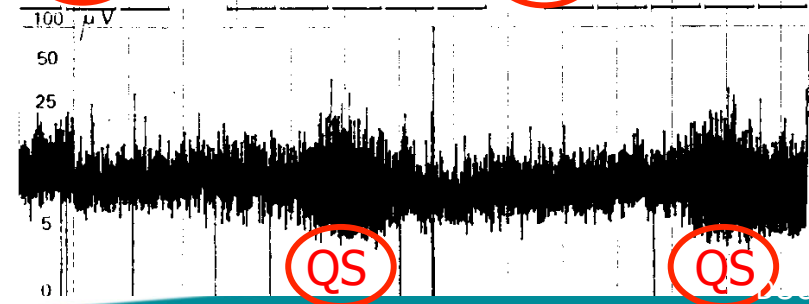
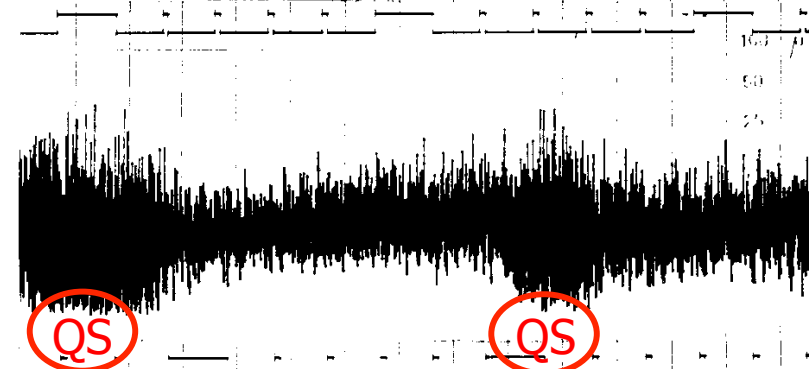
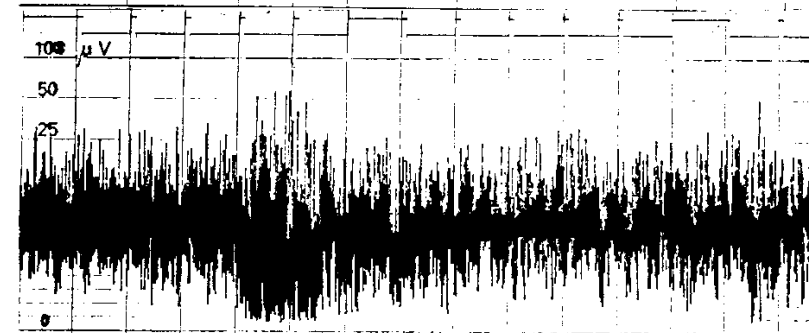
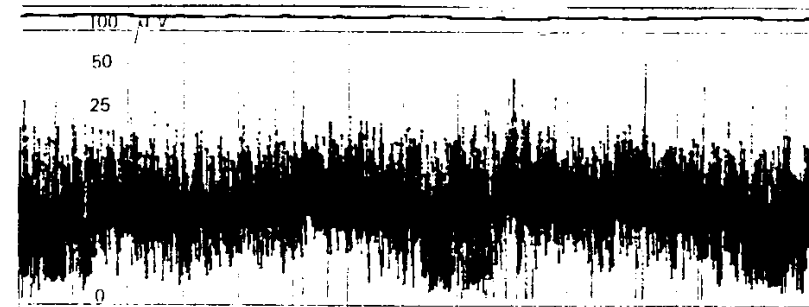
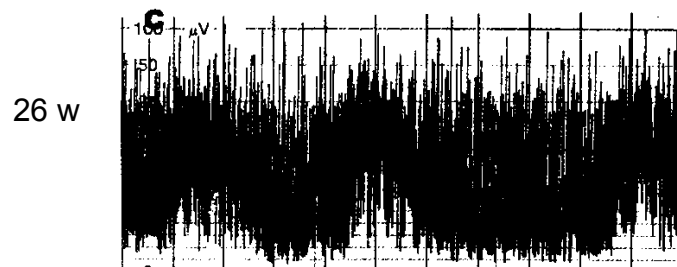
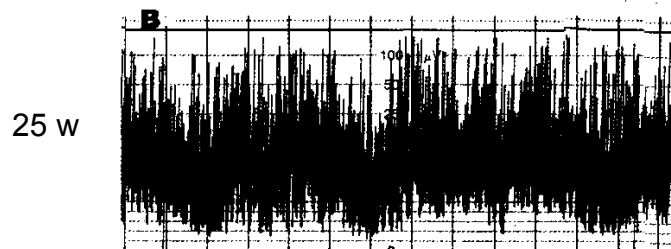
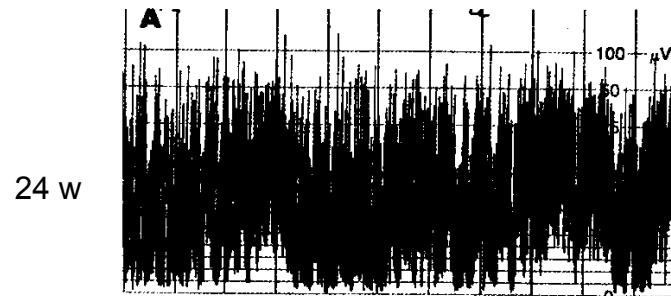
Gestational or Postconceptual Age (wk)	Dominating Background Pattern	SWC	Minimum Amplitude (mcV)	Maximum Amplitude (mcV)	Burst/h
24 through 25	DC	(+)	2 to 5	25 to 50 (to 100)	>100
26 through 27	DC	(+)	2 to 5	25 to 50 (to 100)	>100
28 through 29	DC/(C)	(+)/+	2 to 5	25 to 30	>100
30 through 31	C/(DC)	+	2 to 6	20 to 30	> 100
32 through 33	C/DC in QS	+	2 to 6	20 to 30	> 100
34 through 35	C/DC in QS	+	3 to 7	15 to 25	>100
36 through 37	C/DC in QS	+	4 to 8	17 to 35	> 100
38+	C/DC in QS	+	7 to 8	15 to 25	> 100

SWC: (+) = imminent/immature; SWC: + = developed; SWC: QS = quiet/deep sleep;
DC = discontinuous background pattern; (C) = continuous

NeoReviews. Vol 7 No. 2 February 2006
Hellstrom-Westas, Rosen, deVries, Greisen

Normal aEEG's at Various Gestational Ages

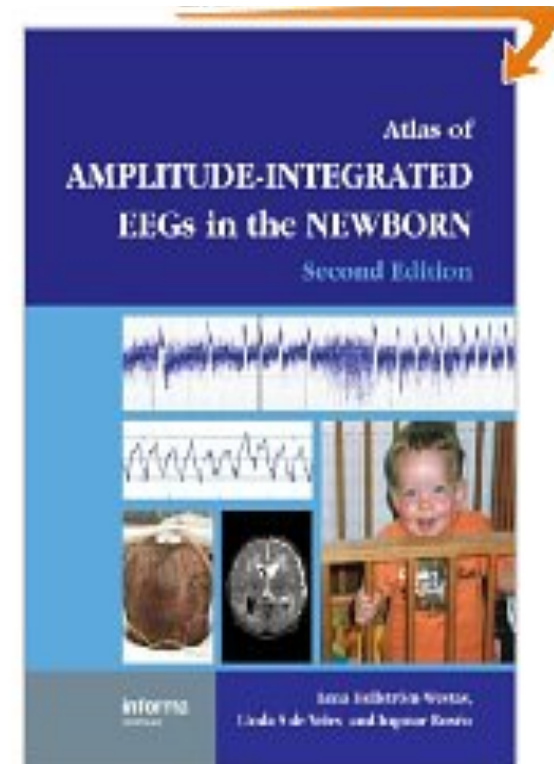
*Thornberg & Thiringer 1990,
Kuhle et al 1999)*



aEEG Reference Literature

Reference:

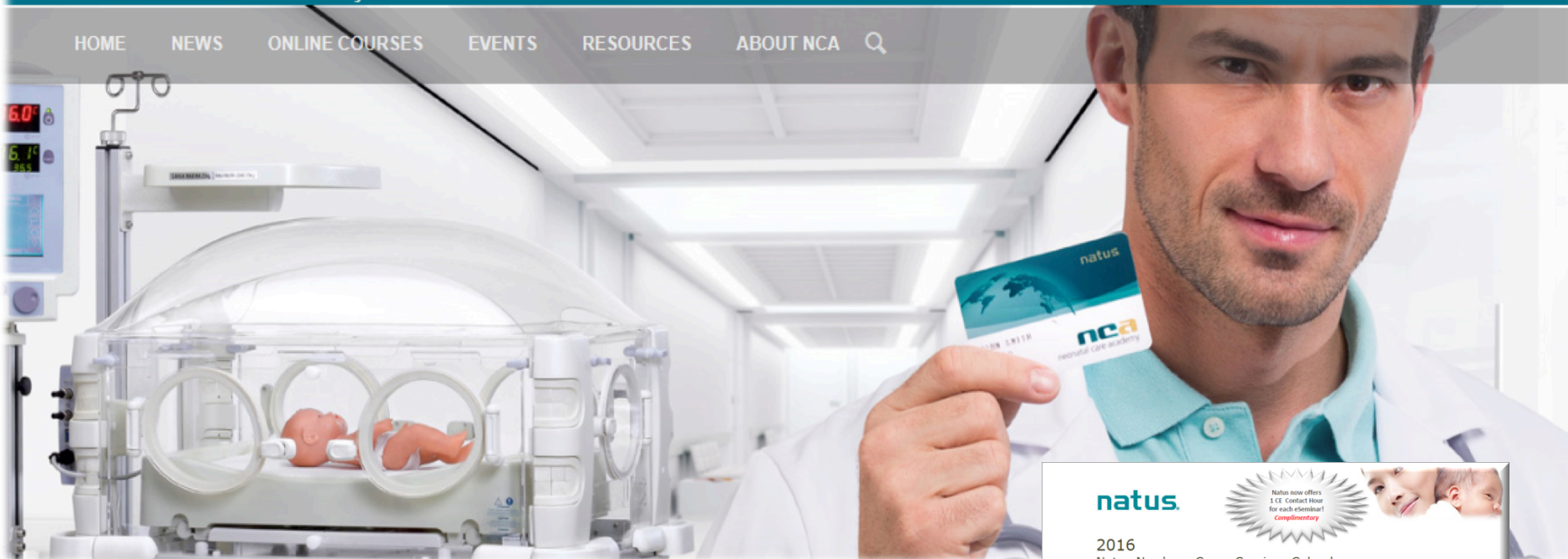
- Atlas of Amplitude-Integrated EEGs in the Newborn: Second Edition (*Hellstrom-Westas, De Vries, Rosen; 2008*)



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Natus Newborn Care eSeminar Calendar**

Month	Topic	Date
FEBRUARY	Thermoregulation 101	February 2, 2016 eSeminar
MARCH	Basics of aEEG	March 15, 2016 eSeminar
APRIL	Assessing Jaundice in the Newborn Infant	April 26, 2016 eSeminar
MAY	Humidifying the Microenvironment	May 3, 2016 eSeminar
JUNE	Family Centered Care <small>By Personal Invitation to Neonatal Care Academy Members</small>	June 21, 2016 eSeminar
JULY	Optimizing the Thermal Environment - Creating a Comfort Zone	July 26, 2016 eSeminar
SEPTEMBER	Basics of aEEG	September 20, 2016 eSeminar
OCTOBER	Creating a Healing Environment	October 4, 2016 eSeminar
NOVEMBER	Science of Hyperbilirubinemia and Clinical Evidence Review	November 1, 2016 eSeminar
World Prematurity Day - Topic TBD <small>By Personal Invitation to Neonatal Care Academy Members</small>		
		November 17, 2016 eSeminar

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