

Sri Lakshmi Narayana Institute of Medical Sciences

Date 02/09/2018

From Dr.R.Venkataramanan, Professor and Head, otorhinolaryngology, SLIMS Bharath Institute of Higher Education and Research, Puducherry.

To The Dean, SLIMS Bharath Institute of Higher Education and Research, Puducherry.

Sub: Permission to conduct value-added course: : Hands on training and an Extensive Discussion on BERA reg.

Dear Sir,

With reference to the subject mentioned above, the department proposes to conduct a value-added course titled: Simulation Based Training In Audiology on Sep 2018 to Dec 2018. We solicit your kind permission for the same.

Kind Regards

Dr.R.Venkataramanan

FOR THE USE OF DEANS OFFICE

Names of Committee members for evaluating the course: The Dean: The HOD: The Expert:

The committee has discussed about the course and is approved.

Dean

(Sign&Seal) DEAN Prof.K.BALAGURUNATHAN,M.S (General surgeon) SRI LAKSHMI NARAYANA INSTITUTE OF MEDICAL SCIENCES OSUDU PONDICHERRY



Subject expert (Sign &Seal)

Seal & Signature of the HOD PROFESSOR & HOD DEPARTMENT OF ENT Britdshain Nersynna Institute Of Medical Sciences PONDICHERRY - 605 502



OFFICE OF THE DEAN

Sri Lakshmi Narayana Institute of Medical Sciences

OSUDU, AGARAM VILLAGE, VILLIANUR COMMUNE, KUDAPAKKAM POST,

PUDUCHERRY - 605 502.

[Recognised by Medical Council of India, Ministry of Health letter No. U/12012/249/2005-ME (P -II) dt. 11/07/2011] [Affliated to Bharath University, Chennai - TN]

Ref. No. SLIMS/Dean Off/VAC/024

Date:03/09/18

From The Dean Sri Lakshmi Narayana Institute of Medical sciences, Pondicherry – 605502

То

The Registrar, Bharath Institute of Higher Education and Research, Chennai - 600073.

Respected Sir

Sub: Request for permission and approval of Syllabus for certificate course (Value Added course) for the academic year 2018-19 - Reg
 Ref: Requesting letter received from Departments

With reference to the above, herewith forwarding the proposed list of Value-added

courses for necessary permission and approval of syllabus to conduct the same.

This is for your kind information and needful action.

Thankingyou

Yours faithfully



DEAN Prof.K.BALAGURUNATHAN,M.S (General surgeon) SRI LAKSHMI NARAYANA INSTITUTE OF MEDICAL SCIENCES OSUDU PONDICHERRY

Encl's:

- 1. Requesting letter received from department
- 2. Syllabus of thecourse
- 3. Details of faculty handlingcourse

Sri Lakshmi Narayana Institute of Medical Sciences, Puducherry

VALUE ADDED COURSE : Hands on training and an Extensive Discussion on BERA

COURSE CO-ORDINATOR DETAILS

Faculty Name: Dr. K.Venkataramanan

Email ID:entslims@gmail.com



Ref. No. BHIER/ VAC/B-02

Date:05.09.2018

From

The Registrar, Bharath Institute of Higher Education and Research, Chennai - 600073.

То

The Dean Sri Lakshmi Narayana Institute of Medical sciences, Pondicherry – 605502

Sir / Madam,

- Sub: Approval of Syllabus to conduct certificate course (Value Added course) for the academic year 2018-2019 Reg.
- Ref: Ref. No. SLIMS/Dean Off/VAC /024 Dated: 03.09.2018

With reference to the above, it is to inform that the proposal submitted to conduct Value Added Course has been accepted and approved by BIHER, council meeting. List of the VAC are mentioned below for the academic year 2018–2019. The abstract of the VAC course completion detail should be submitted to the Registrar office.

Thanking you

Yours faithfully

REGISTRAR



Sri Lakshmi Narayana Institute of Medical Sciences

OSUDU, AGARAM VILLAGE, VILLIANUR COMMUNE, KUDAPAKKAM POST, PUDUCHERRY - 605 502.

[Recognised by Medical Council of India, Ministry of Health letter No. U/12012/249/2005-ME (P -II) dt. 11/07/2011] [Affliated to Bharath University, Chennai - TN]

Circular

07/09/2018

Sub: Organising Value-added Course: Hands on training and an Extensive Discussion on BERA reg.

With reference to the above mentioned subject, it is to bring to your notice that SLIMS, **Bharath Institute of Higher Education and Research**, is organising **"Hands on training and an Extensive Discussion on BERA"**. The course content and registration form is enclosed below.

The application must reach the institution along with all the necessary documents as mentioned. The hard copy of the application should be sent to the institution by registered/ speed post only so as to reach on or before 15/09/2018. Applications received after the mentioned date shall not be entertained under any circumstances.

DEAN

DEAN Prof.K.BALAGURUNATHAN,M.S (General surgeon) SRI LAKSHMI NARAYANA INSTITUTE OF MEDICAL SCIENCES OSUDU PONDICHERRY

Encl: Copy of Course content

VALUE ADDED COURSE

1. Name of the programme &Code

Hands on training and an Extensive Discussion on BERA - A value added course for the medical students.

& ENT 08

2. Duration & Period

30 hrs & Sep 2018-Dec 2018

3. Information Brochure and Course Content of Value Added Courses

Enclosed as Annexure- I

4. List of students enrolled

Enclosed as Annexure- II

5. Assessment procedures:

Pre test and post test which includes 10 mcqs - Enclosed as Annexure- III

6. Certificate model

Enclosed as Annexure- IV

7. No. of times offered during the same year:

1 time Sep 2018- Dec 2018

8. Year of discontinuation:2019

9. Summary report of each program year-wise

Value Added Course- Sep 2018- Dec 2018										
Sl. No	Course Code	Course Name	Resource Persons	Target Students	Strength & Year					
1	ENT 08	Hands on training and an Extensive Discussion on BERA	1.Dr.Venkataramanm 2. Dr. Sreedhar.B	3 rd year MBBS students	14 students & 2018					

10. Course FeedBack

Enclosed as Annexure- V

Tr.R.Jum

RESOURCEPERSON 1. Dr.K.R.Jothikumar 2. Dr. Sreedhar.B

COORDINATOR Dr.R.Venkataramanan

COURSE PROPOSAL

1. NAME OF THE PROGRAMME

Hands on training and an ExtensiveDiscussion on BERA- A value added course for the medical students.

2. AIM

Training the students with hands on experience in BERA as an objective audiological test

3. OBJECTIVES

- a) To teach the students to connect the electrodes of BERA
- b) To teach the students to interpret the BERA waves

4. METHODOLOGY

- Students who are interested in participating in value added course are enrolled and the course is conducted for them during the non college hours for a period of 30 hours from Sep 2018 Dec 2018 . This course is conducted every 6 months.
- Course Audience: 3rd year MBBS students
- Course Coordinator: Dr.R.Venkataramanan
- **Course Faculties with Qualification and Designation:**
- 1.Dr.K.R. jothikumar
- 2.Dr. Sreedhar.B

Schedule followed during the course

No	Topic	Title	Duration	Date and time
1	Hands on training and an	Introduction of Brain Stem Evoked Response	4 hrs	4pm-6pm(19/9/18),4pm-6pm(25/9/18)
	Extensive Discussion on BERA	Audiometry (BERA)		
		Principle and method of recording BERA	4 hrs	4pm-6pm(29/9/18),4pm-6pm(4/10/18)
		Latency and amplitude studies of BERA	4 hrs	4pm-6pm(10/10/18),4pm-
				6pm(15/10/18)
		Clinical uses of BERA	4 hrs	4pm-6pm(19/10/18),4pm-
				6pm(23/10/18)
		Neurophysiologic basis of BERA	4 hrs	4pm-6pm(29/10/18),4pm-6pm(4/11/18)
		Demonstration of BERA and interpretation of	5 hrs	4pm-6pm(10/11/18),4pm-
		results		7pm(15/11/18)
		Hands on training on BERA lead connection and	5 hrs	4pm-6pm(23/11/18),4pm-
		interpretation of results and DOPS		6pm(3/12/18),4pm-5pm(8/12/18)
		TOTAL	30HRS	

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REFERENCE BOOKS	(1) SCOTT BROWN 7th edition	
KEFERENCE DOORS		
	2) ANIRBAN BISWAS 3rd edition	
		\square

Brainstem Auditory Evoked Potentials



Sohmer and Feinmesser

Signal-averaged ECochG studies

Jewett and co-workers

- Identified the short-latency scalp-recorded AEPs as far- field potentials volume-conducted from the brainstem, described the components and their properties
- Established the Roman numeral labeling of the peaks

Brainstem Auditory Evoked Potentials

- Following a transient acoustic stimulus, ear and parts of the nervous system generate a series of electrical signals with latencies ranging from milliseconds to hundreds of milliseconds
- Recorded from electrodes placed on the skin
- To evaluate noninvasively the function of the ear and portions
 of the nervous system activated by the acoustic stimulation

BAEPs

□ Generated by an anatomically distinguishable neuronal

subsystem for sound localization within the brainstem

BAEPs can be used only to assess the status of the ear,
 auditory nerve, and brainstem auditory pathways up through
 the level of the mesencephalon

Auditory pathway



5

BAEPs

- Ascending projections from the cochlear nucleus are bilateral but are more extensive contralaterally than ipsilaterally
- Despite this anatomic asymmetry, the BAEPs appear to reflect predominantly activity in the ipsilateral ascending pathways

BAEPs

Short-latency components, with
 latencies of under 10 msec in adults

Long-latency AEPs, with latencies
 exceeding 50 msec

 Middle-latency AEPs, with intermediate latencies



Long-latency AEPs

Affected profoundly by the degree to which the subject is

attending to the stimuli and analyzing stimulus features

- Used as probes of cognitive processes
- Their variability, as well as uncertainty about the precise identity of their cortical generators, limits their utility for neurologic diagnosis

Middle-latency AEPs

Small

- Subject to contamination by myogenic signals
- □ Variable from subject to subject

Middle- and Long-latency AEP

generated predominantly by postsynaptic potentials

within areas of cerebral cortex that are activated by the acoustic stimulus

 affected increasingly by the state of the subject and by anesthesia as their latency increases

Greatest clinical utility because

- □ Relatively easy to record
- Waveforms and latencies are highly consistent across normal subjects
- Unaffected by the subject's degree of attention to the stimuli and are almost identical in the waking and sleeping states
- □ Minor differences related to changes in body temperature

Although short-latency AEPs commonly are called brainstem auditory evoked potentials, this term is not completely accurate because the roster of generators clearly includes the distal (with respect to the brainstem) cochlear nerve and may also include the thalamocortical auditory radiations, neither of which is within the brainstem

- Brief acoustic click stimuli that are produced by delivering monophasic square pulses of 100-µsec duration to headphones or other electromechanical transducers at a rate of about 10 hz
- A rate of exactly 10 hz or another submultiple of the power line frequency should be avoided because of line frequency artifact
- □ Stimulus intensity of 60 to 65 db HL is a typical level

If hearing loss is present, stimulus intensity may be

- □ Stimuli are delivered monaurally
- To prevent contralateral ear stimulation it is masked with continuous white noise at an intensity 30 to 40 dB below that of the BAEP stimulus
- Activate region of the cochlea (base) responding to 2,000to 4,000-Hz sounds

Stimulation at Several Intensities Differentiate peripheral from neural abnormalities, especially when wave I is not clear

- In Conductive hearing loss, if the stimulus intensity is increased and no coexisting sensorineural hearing loss is present, a normal BAEP will be recorded
- In contrast, BAEPs that are delayed as a result of abnormally slowed neural
- conduction do not normalize



Rapid Stimulation

- Approximately 10 Hz is used for routine clinical testing
- As the stimulus rate is increased above approximately 10 per sec, component amplitudes decrease and the peaks tend to become less well defined
- □ Wave V is most resistant to these effects
- More rapid rates may be used to facilitate recordings to measure the wave V threshold

Stimulus clicks according to polarity

Compression click (condensation click)

□ If the electrical square pulse causes the diaphragm of the acoustic transducer to move toward the patient's ear

Rarefaction click

- Reversing the polarity of the electrical square pulse
- Generally preferable because BAEP peaks tend to be clearer

Alternative Auditory Stimuli

Stimulation with brief tone pips

□ To probe specific parts of the cochlea

Acoustic masking

 Used to obtain frequency-specific information from BAEPs

Relatively poor signal-to-noise ratios

BAEPs by bone-conducted stimuli

 Most useful in assessing patients who may have conductive hearing losses, such as neonates in whom BAEPs performed with air-conducted stimuli are suggestive of a hearing loss

19

BAEPs to Electrical Stimuli

- Electrical stimulation of eighth nerve fibers through the electrodes of a cochlear prosthesis
- Used to assess the proximity of these electrodes to the spiral ganglion during implantation and the adequacy of eighth nerve stimulation during programming of the processor
- Correlate well with auditory outcomes, and may prove to be useful in guiding therapy in young children with questionable auditory nerve integrity

Recording electrodes

- Typically are placed at the vertex
 (location Cz of the International 10–
 20 System) and at both ear lobes
 (Ai and Ac)
- Electrodes at the mastoids (Mi and Mc) may be substituted, although wave I tends to be smaller because of muscle noise





- Patients usually are tested while lying comfortably so that their neck musculature is relaxed
- Patients should be requested to let their mouth hang open if the muscles of mastication are tensed
- encouraged to sleep during testing
- If the patient cannot relax sufficiently, sedation can be induced with agents such as chloral hydrate (little or no effect on BAEPs in the usual sedative doses)

Data analysis

- Amplifier filters out all of the delta, theta, alpha, and beta bands of the EEG
- Biologically derived noise in the recordings is derived predominantly from muscle activity
- Therefore, patient relaxation during the recording session is essential to obtain "clean" waveforms with a good signal-tonoise ratio

- Data typically are digitized over an epoch duration or analysis time of approximately 10 msec
- Longer analysis time of 15 msec may be required for recording pathologically delayed BAEPs, BAEPs to lowered stimulus intensities (as when recording a latency–intensity study), BAEPs in children, and BAEPs during intraoperative monitoring
 - Signal averaging is required for improvement in the
 - signal-to-noise ratio

- Cz–Ai BAEP typically is displayed so that positivity at the vertex relative to the stimulated ear is displayed as an upward deflection
- Upward-pointing peaks are labeled with Roman numerals
- Downward-pointing peaks are labeled with the suffix N according to the peak that they follow

BAEP waveform

□ Typically begins with an electrical stimulus artifact that is

synchronous with stimulus production at the transducer


Reducing the Stimulus Artifact

May overlap with wave I and impair the identification and

measurement

 Using shielded headphones and headphones with piezoelectric transducers instead of voice coil transducers

□ Transducers that are connected to an ear insert by

flexible plastic tubing several centimeters in length

Wave I

- □ first major upgoing peak of the Cz–Ai BAEP
- upgoing peak of similar amplitude in the Ac–Ai waveform
- markedly attenuated

or absent in the Cz-

Ac waveform



- Arises from at the most distal
 (i.E., Closest to the cochlea)
 portion of auditory nerve
- Circumscribed negativity
 around the stimulated ear
- Appears in Cz–Ai and Ac– Ai recordings but is minimal or absent in Cz–Ac



Cochlear microphonic Vs Wave I

Visible as a separate peak preceding wave I, especially if

the stimulus artifact is small

 Distinguished by reversing the stimulus polarity, which will reverse the polarity of the cochlear microphonic; Wave I may show a latency shift, but will not reverse polarity

- Represents contributions to wave I from different portions
 of the cochlea
- Earlier of the two peaks, which reflects activation of the base of the cochlea, corresponds to the single wave I that is typically present in the Cz–Ai waveform
- Reversal of stimulus polarity can be used to distinguish a bifid wave I from a cochlear microphonic followed by (a single) wave I

Techniques to obtain a clearer wave I

- □ Electrode within the external auditory canal
- Ac-Ai recording channel can yield a somewhat larger and clearer wave I than that in the standard Cz-Ai recording
- Reduction in the stimulus repetition rate
- □ Increasing the stimulus intensity

- □ Present at substantial amplitude in the Cz–Ac channel
- □ Usually the earliest BAEP component in that waveform



35

- From auditory nerve as it passes the internal auditory meatus and moves from a nerve encased in bone to one surrounded by cerebrospinal fluid
- Field includes positivity at the mastoid and far-field negativity around the vertex



In contrast to wave I,

Wave II

- □ First major upward deflection in the Cz–Ac waveform
- □ Similar amplitude in the Cz–Ai and Cz–Ac channels
- may be small and difficult to identify in some normal subjects



- Wave II
 - Arises from two loci
 - a) distal auditory nerve
 - b) brainstem, specifically the cochlear nucleus or its outflow and proximal end of the auditory nerve

V, VI?,

- Earliest component affected by pontomedullary CVAs involving the cochlear nucleus
- Usually predominant over the dorsal part of the head and a clear wave II in the Cz–Ac waveform

Arises from
 auditory nerve as it
 passes the internal
 auditory meatus



usually present in
both the Cz–Ai and
Ac–Ai channels
substantially
smaller in the Cz–
Ac



- Arises from caudal pontine
 tegmentum in the superior
 olivary complexes or their
 outflow within the lateral
 - lemniscus
- Abnormal either ipsilateral or contralateral to the major pathology in patients with asymmetric lesions



Wave III variants

Bifid wave III

normal variant

Poorly formed or absent wave II

 normal variant in a patient with a clear wave V and a normal I–V interpeak interval



Role of Descending Pathways in BAEPs

- Waves I and II may be quite large or waves II and III are delayed in latency in patients with rostral brainstem pathology
- probably reflects loss of activity in descending inhibitory
 pathways originating in or traversing the region of the
 inferior colliculus

□ often fused into a IV/V

complex

- most prominent
 component in the BAEP
 waveform
- morphology varies from
 one subject to another, and
 may differ between the
 - two ears in the same



Waves IV and V

- Earliest components that are absent and usually are the earliest that are abnormal in patients with lesions of the midpons, rostral pons, or mesencephalon
- Usually is followed by a large negative deflection that lasts several milliseconds and brings the waveform to a point below the prestimulus b.



Various IV/V complex morphologie s in Cz–Ai waveforms recorded in normal subjects



45_

Totally fused IV/V complex Vs single wave IV or V

Complex has a "base" that is greater than 1.5 msec in duration, whereas the width of a single wave is less than 1.5 msec



- Reflect activity
 predominantly in ascending
 auditory fibers within the
 dorsal and rostral pons,
 caudal to the inferior
 colliculus
- Affected by tumors or
 cerebrovascular accidents



or rostral pons



48

- □ Arises at the level of the mesencephalon, either from the inferior colliculus itself or, from the fibers in the rostral portion of the lateral lemniscus as they terminate in the inferior colliculus Intracranial data suggest C/L
 - mesencephalon but clinically
 - associated most often with

ipsilateral pathology



Wave IV vs V

- Wave V most resistant to
 the effects of decreasing
 stimulus intensity or
 increasing stimulus rate
- If either of these stimulus modifications is performed progressively until only one component remains, that peak can be identified as



wave V

- Occasionally, wave V may
 be present following
 stimulation with one click
 polarity but not the other
- Therefore, recording a
 BAEP with the opposite
 stimulus polarity may be
 useful if wave V is not
 identifiable with the
 - standard laboratory



Differential affection of waves IV and V

- Multilevel demyelination
- Brainstem infarct

51

□ Small brainstem hemorrhage in the lateral lemniscus

Wave VN

Downward deflection following

wave V(slow negativity (SN)

- Typically wider than the positive components and the earlier negative peaks
- Reflects postsynaptic potentials
 within brainstem auditory nuclei,
 primarily the inferior colliculus



53

- Generation within the medial geniculate nuclei or their outflow tracts
- Absent in Cz–Ai and Cz–Ac recordings in some normal individuals
- Abnormalities in patients with tumors of the rostral midbrain and caudal thalamus at the level of the medical geniculate nucleus and the brachium of the inferior colliculus

BAEPs cannot be used to assess the status of the



Often absent in conventionally

recorded normal BAEPs

 Generation near the auditory cortex, predominantly contralaterally

Does not provide clinically useful
 information about the status of the
 auditory pathways



Wave V latency measurement Should be taken from the

55

second subcomponent of the IV/V complex, even if this is not the highest peak (in contrast to the amplitude measurement, which is taken from the highest point in the complex)



Wave V latency measurement

□ Measurement in a Cz−

Ac

- Overlapping peaks are
 separated more clearly
 because the latency of
 wave IV is typically
 earlier, and that of wave
 V is later, than in the Cz–
 - Ai waveform



IV/V:I amplitude ratio

□ With respect to the most negative point that follows it in

the waveform (I to IN and IV/V to VN), and their ratio is



27-year-old woman with probable multiple sclerosis

IV/V:I amplitude ratio is 0.28; all absolute latencies and interpeak intervals are normal

¢linical interpretation of BAEPs

□ Waves II, IV, VI, and VII are sometimes not identifiable in

normal individuals, and their peak latencies display more interindividual variability

- Amplitude measurements of the individual components are also highly variable
- Ratio between the amplitude of the IV/V complex and that
 of wave I has proved to be a clinically useful measure

¢linical interpretation of BAEP

□ Identification of waveforms - Presence or absence of

waves I, III, and V

- □ Latencies of waves I, III, and V
- □ I–III, III–V, and I–V interpeak interv
- Right-left differences of these values
- □ IV/V:I amplitude ratios



Measurement of right-left differences

□ Increases test sensitivity because the intersubject

variability of these measures is less than that of the

absolute component latencies and interpeak intervals



¢linical interpretation of BAEPs

Peripheral transmission time (PTT)

□ Latency of wave I

Central transmission time (CTT)

□ I–V interpeak interval

- Control data should have been acquired under the same conditions used to test the patient, including the polarity, rate, and intensity of the stimulus and the filter settings used for data recording
- Limits of the normal range are typically set at 2.5 or 3
 standard deviations from the mean of normally distributed
 data
- I–V and III–V interpeak intervals are, on average, shorter in women than in men

Normative Data



□ Wave I=1.50

□ Wave III=3.57

 \Box Wave V=5.53
Delay Versus Absence of Components

Evoked potentials represent the summated activity of

large populations of neurons firing in synchrony

- Delay If delayed uniformly, a delayed evoked potential component will result
- Absence If the delay is nonuniform due to temporal dispersion
- Either delay or absence of a BAEP peak indicate dysfunction,
 but not necessarily complete loss of activity, in a part of the
- infratentorial auditory pathways

Criteria for retrocochlear dysfunction Absence of all BAEP waves I through V unexplained by extreme hearing loss determined by formal audiometric testing.

- Absence of all waves following waves I, II, or III.
- Abnormal prolongation of I-III, III-V. and I-V interpeak intervals
- Abnormal diminution of the IV-V/I amplitude ratio, especially when accompanied by other abnormalities.
- Abnormally increased differences between the two ears (interaural differences) when not explained by unilateral or asymmetric middle and/or ear dysfunction determined by appropriate audiometric tests.

65

Obtaining formal audiometric testing in patients undergoing

- Reflect peripheral auditory dysfunction, either conductive or cochlear, or pathology involving the most distal portion of the eighth nerve
- Poorly formed or absent wave I but a clear wave V may reflect high-frequency hearing loss.
- May reflect intracranial pathology because the cochlea receives its blood supply from the intracranial circulation via the internal auditory artery

Abnormalities of the I–III Interpeak Interval

- Prolongation reflects an abnormality within the neural auditory pathways between the distal eighth nerve on the stimulated side and the lower pons
- □ Seen in acoustic neuromas, demyelinating disease,
 - brainstem tumors, or vascular lesions of the brainstem



- Abnormalities of the III-V Interpeak Interval
 Reflects an abnormality between the lower pons and the mesencephalon most often, although not always, ipsilateral²to/ the lesion
- Prolongation not an abnormality if the I–V interpeak interval is normal.
- □ Seen in a variety of disease processes involving the

brainstem, including demyelination, tumor, and vascular disease

Abnormalities of the IV/V:I Amplitude Ratio

- Reflects dysfunction within the auditory pathways between the distal eighth nerve and the mesencephalon
- □ False increase in ratio in
- Decreasing the stimulus intensity
- Suboptimal placement of the Ai recording electrode (may decrease the amplitude of wave I)

BAEPs and Hearing Loss

- can detect subtle neuronal dysfunction that is not
 clinically apparent on the neurologic and audiologic
 examination
- Relatively insensitive to isolated low-frequency hearing losses

BAEPs and Hearing Loss

71

Central pattern

□ CTT (I–V interpeak interval) is prolonged

Peripheral pattern

□ Wave I is delayed

A single waveform may contain both abnormalities

Classification of Hearing

LOSS Clinical Classification	Location of Pathology	BAEP Classification
Conductive hearing	External or middle	
loss	ear	Peripheral hearing
Sensorineural hearing loss	Inner ear (cochlea)	loss
	Eighth nerve or	"Central" hearing loss
	(retrocochlear)	



Latency-intensity curves

□ May increase the sensitivity of BAEPs for detecting small



Latency-intensity curves stimulation recorded before and after surgery in left-sided intracanalicular acoustic neuroma

acoustic neuromas

BAEPs abnormal but normal hearing

Unilateral brainstem lesions because the ascending

projections from each ear are bilateral

Lesions of subsystem involved in sound localization
 sparing other portions of the brainstem auditory
 pathways

Absence of a component may reflect temporal dispersion
 rather than conduction block, so hearing may even be present
 when there is no identifiable wave V

BAEPs and functional hearing loss

□ Abnormal BAEP study demonstrates the existence of

pathology within the auditory system

- Normal study does not prove that the symptoms are psychogenic
- If they maintain a degree of tension in their cranial and neck muscles, the EMG activity picked up by the recording electrodes may be sufficient to prevent recording of an interpretable BAEP study

Neuroma

- □ Abnormal BAEPs in more than 95 % with acoustic neuromas
- Abnormal BAEPs is less in patients with small (less than 1 cm) tumors
- Small, intracanalicular tumors in whom BAEPs to standard high-intensity stimuli are normal, latency– intensity studies may reveal abnormal cochlear function resulting from compression of the internal auditory artery

BAEP in Acoustic

Neuroma

Typically originate from the distal vestibular nerve at the vestibular ganglion, and the auditory portion of the nerve may be unaffected early in the course of the disease.

↓ As it enlarges, compress the auditory nerve Prolongation of the I–III interpeak interval ↓

Complete eradication of wave III and subsequent BAEP components

BAEP in Acoustic

- Wave II may be relatively spared, a reflection of the contribution to that component originating in the distal eighth nerve
- Wave I may become delayed as the degree of cochlear ischemia increases

BAEP in Large Acoustic Neuroma

- Infarction of the cochlea may cause elimination of all
 BAEPs
- Prolongation of the III–V interpeak interval in response to stimulation of the ear contralateral to the tumor due to compression of brainstem



BAEP in Other Posterior Fossa Tumors

82

□ Almost always abnormal in brainstem gliomas and other

intrinsic brainstem tumors except that within the medulla

- Abnormalities in the I–III or III–V interpeak interval, or a combination of both
- Serial recordings may show deterioration of the BAEPs
 because of tumor growth
- Response to treatment can be demonstrated as an improvement in conduction within the brainstem auditory pathways

BAEP in Cerebrovascular Disease

- Usually normal in medullary infarcts or lesions confined to the basis pontis, cerebral peduncles, and cerebellar hemispheres
- Presence of BAEP abnormalities is associated with an adverse clinical outcome
- □ Abnormalities in the I–III or III–V interpeak interval, or a combination of both, may be seen
- If a cochlear stroke accompanies the brainstem stroke, all
 BAEP components will be absent following stimulation

BAEP in vertebrobasilar TIA

- Abnormalities present in most cases
- □ Findings tend to resolve over time
- □ Yield lower if recorded after more than a week
- Persistent BAEP changes may represent small infarcts that are clinically silent

BAEP in Demyelinating Disease

85

Can demonstrate a residual abnormality related to a prior

symptom that has cleared

- Abnormality rate higher in definite MS and with brainstem lesion.
- Abnormalities in the I–III and/or III–V interpeak interval
- □ Abnormally small IV/V:I amplitude ratio in the presence of

normal component latencies and interpeak intervals

Prolongations of the PTT (wave I latency) without any otologic cause

BAEP in MS

VIIIn fibers are ensheathed along most of their lengths by

central-type myelin, produced by oligodendrocytes unlike other CNs which have Schwann cells

In MS vulnerable to demyelination along most of its length

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Abnormalities of the I–III interpeak interval

- Markedly abnormal BAEPs are likely to have poor neurologic outcomes attributable to the brainstem damage
- Typically normal in patients with coma caused entirely by supratentorial disease
- May deteriorate subsequently because of transtentorial herniation
- Abnormal BAEPs in patients with supratentorialinfarctions or hemorrhages are correlated with poor

- Normal-appearing BAEPs in a patient whose examination shows widespread brainstem dysfunction should prompt suspicion of a metabolic etiology such as a drug overdose
- BAEPs are highly resistant to central nervous system depressant drugs



BAEP in Locked-in syndrome

BAEPs may be either normal or abnormal, depending on

the extent to which the lesion extends outside the ventral pons and involves the auditory pathways

BAEP in brain death

- Contains no identifiable components, or consists of wave I alone, or contains only a wave I followed by a wave IN
- Rarely, waves II and IIN may also be present and reflect the contribution to these components from the auditory nerve
- Although consistent with brain death, negative BAEP
 cannot be used as evidence that the brainstem is
 nonfunctional

Intra operative BAEP monitoring

esp during surgery in the cerebellopontine angle with the

goal of preserving auditory nerve function

- □ bulky earphone replaced by a small insertable earphone
- faster stimulation rate typically about 30 Hz compared with
 10 Hz which allows more rapid signal acquisition
- □ Stable, robust BAEPs are recorded readily in the

presence of general anesthetic agents

Intra operative BAEP



nerve, a loss of wave V amplitude of 50 percent or more or an increase in wave V latency by 0.5 msec generally is recognized as a potentially important alteration, particularly when it occurs suddenly



BAEPs in infants and children

To detect and measure hearing loss in children who

cannot be tested behaviorally

- To evaluate the auditory brainstem pathways in children who may have neurologic problems
- Requires close cooperation between audiologists and neurologists because it is impossible to interpret these responses correctly without paying careful attention to both the ear and the brain.

ANNEXURE 2 BHARATH INSTITUTE OF HIGHER EDUCATION AND RESEARCH SLIMS

1	U15MB287	GAYATHRI .M
2	U15MB288	GOGUL SUGAN. K
3	U15MB289	GOKULA KRISHNAN. E
4	U15MB290	GOWTHAM. M .R
5	U15MB291	GOWTHAM.S
6	U15MB292	HARIHARAN.S
7	U15MB293	HARINI .L
8	U15MB294	ILAMATHI.S
9	U15MB295	ILAYARAJA .B.U
10	U15MB296	JAMZER. J
11	U15MB297	JANARTHANAM. M
12	U15MB298	JANISHA MARAGATHA J P
13	U15MB299	JAWATH. S
14	U15MB300	JAYA AKSHAIY. J

ANNEXURE 3

SRI LAKSHMI NARAYANA INSTITUTE OF MEDICAL SCIENCES

PUDUCHERRY

TOPIC: Brainstem evoked response audiometry (ENT08) STUDENT NAME: UNIVERSITY NO:

1.Earliest age for doing BERAa. In uterob. At birthc. 3 monthsd. 6months

2.BERA is most accurate at;a.28wkb.30 wkc.32wkd.34wk

3.True about BERA findings in acoustic neuroma a.Latency of waveV increased in the affected ear b.Latency of wave I increased in the affected ear c.No waves on BERA d. No change in BERA

4.Brainstrem evoked response audiometry true is:a.Can differentiate cochlear and retrocochlear lesionb.cannot differentiate the site of lesion in sensorineural hearing lossc.. Can differentiate barbiturate poisionong from other causes of comad.As a screening procedure for infants

5.In normal adult, waveV is generated from b. superior olivary complex a. cochlear nucleus d. Inferior colliculus c. Lateral lemniscus 6. To distinguish between cochlear and post cochlear damage The test is done a.BERA b.Impedance audiometry d.All of the above c.Pure tone audiometry 7. which is the investigation of choice in assessing hearing loss in infants a) Impedance audiometry b)BERA c) Free field audiometry d) behavior audiometry

8) A 6month old child developed labrinthintis following meningitis

a)BERA b)Impedance audiometry c)Pure tone audiometry d) OAE

9) screening test for neonate in ICU
a)Distortion product OAE
b) OAE
c) BERA
d) Impedance audiometry

10. objective method of hearing testa) tuning fork testb) Pure tone audiometryc) Speech audiometryd) BERA

DR.GIRISH.S

PRE TEST


SRI LAKSHMI NARAYANA INSTITUTE OF MEDICAL SCIENCES

PUDUCHERRY

TOPIC: Brainstern evoked response audiometry (ENTO8)

STUDENT NAME: Jawath-S UNIVERSITY NO: U15MB299.

1.Earliest age for doing BERA a. In utero b. At birth c. 3 months d. 6months

2.BERA is most accurate at; a.28wk b.30 wk X c.32Wk d.34wk

> 3.True about BERA findings in acoustic neuroma a.Latency of waveV increased in the affected ear b.tatency of wave I increased in the affected ear c.No waves on BERA d. No change in BERA

4.Brainstrem evoked response audiometry true is:
a.Can differentiate cochlear and retrocochlear lesion
b.caphot differentiate the site of lesion in sensorineural hearing loss
c.. Can differentiate barbiturate polsionong from other causes of coma
d.As a screening procedure for infants

5.In normal adult, waveV is generated from a. cochlear nucleus b-superior olivary complex

c. Lateral lemniscus d. Inferior colliculus

X

X

6. To distinguish between cochlear and post cochlear damage The test is done a.BERA b.Impedance audiometry

c.Pure tone audiometry d.All of the above

7. which is the investigation of choice in assessing hearing loss in infants
a) Impedance audiometry
b) BERA
c) Free field audiometry
d) behavior audiometry

8) A 6month old child developed labrinthintis following meningitis

a)BERA b)Impedance audiometry c)Pure tone audiometry d) OAE

9) screening test for neonate in ICU a)Distortion product OAE b) OAE c) BERA d) Impedance audiometry

10. objective method of hearing test

a) tuning fork test b) Pure tone audiometry c) Speech audiometry d) BERA 8

SRI LAKSHMI NARAYANA INSTITUTE OF MEDICAL SCIENCES

PUDUCHERRY

TOPIC: Brainstem evoked response audiometry (ENTOB) STUDENT NAME: Hariharan. S UNIVERSITY NO: U15 MB292.

1.Earliest age for doing BERA a. In utero b.At birth c. 3 months d. 6months

X

2.BERA is most accurate at; a.28wk b.30 wk C.32Wk d.34wk

> 3. True about BERA findings in acoustic neuroma a.Latency of waveV increased in the affected ear b.tatency of wave I increased in the affected ear c.No waves on BERA d. No change in BERA

4.Brainstrem evoked response audiometry true is:
a.Can differentiate cochlear and retrocochlear lesion
b.cannøt differentiate the site of lesion in sensorineural hearing loss
c.. Can differentiate barbiturate poisionong from other causes of coma
d.As a screening procedure for infants

POST TEST

 S.In normal adult, waveV is generated from

 a. cochlear nucleus
 b. superior olivary complex

 c. tateral lemniscus
 d. Inferior colliculus

6. To distinguish between cochlear and post cochlear damage The test is done **BERA** b.Impedance audiometry c.Pure tone audiometry d.All of the above

7. which is the investigation of choice in assessing hearing loss in infants
a) Impedance audiometry
b) BERA
c) Free field audiometry
d) behavior audiometry

8) A 6month old child developed labrinthintis following meningitis a)BERA b)Impedance audiometry c)Pure tone audiometry d) OAE

9) screening test for neonate in ICU a)Distortion product OAE b) OAE CHEERA d) Impedance audiometry

 10. objective method of hearing test

 a) tuning fork test
 b) Pure tone audiometry

 c) Speech audiometry
 d) BERA

8

SRI LAKSHMI NARAYANA INSTITUTE OF MEDICAL SCIENCES

PUDUCHERRY

TOPIC: Brainstem evoked response audiometry (ENTO8) STUDENT NAME: Harcharan .S UNIVERSITY NO: V) 5 M B 2 9 2.

1.Earliest age for doing BERA

c. 3 months d. 6months

2.BERA is most accurate at; a.28wk b.30 wk

c.32wk d.34wk

3.True about BERA findings in acoustic neuroma -a.tatency of waveV increased in the affected ear b.Latency of wave I increased in the affected ear c.No waves on BERA d. No change in BERA

4.Brainstrem evoked response audiometry true is:
a.Can differentiate cochlear and retrocochlear lesion
b.cannot differentiate the site of lesion in sensorineural hearing loss
c.. Can differentiate barbiturate poisionong from other causes of coma
d.As a screening procedure for infants

 5.In normal adult, waveV is generated from

 a. cochlear nucleus
 b. superior olivary complex

 c-tateral lemniscus
 d. Inferior colliculus

6. To distinguish between cochlear and post cochlear damage The test is done A-BERA b.Impedance audiometry c.Pure tone audiometry d.All of the above

7. which is the investigation of choice in assessing hearing loss in infants
a) Impedance audiometry
b)BERA
c) Free field audiometry
d) behavior audiometry

8) A 6month old child developed labrinthintis following meningitis
 a)BERA
 b)Impedance audiometry
 c)Pure tone audiometry
 d) OAE

9) screening test for neonate in ICU a)Distortion product OAE b) OAE -e) BERA d) Impedance audiometry

10. objective method of hearing test

a) tuning fork test b) Pure tone audiometry c) Speech audiometry 8

SRI LAKSHMI NARAYANA INSTITUTE OF MEDICAL SCIENCES

PUDUCHERRY

TOPIC: Brainstem evoked response audiometry (ENTOB) STUDENT NAME: Jawath - S UNIVERSITY NO: U(SIV) 8299.

1.Earliest age for doing BERA a. In utero b. At birth c. 3 months d. 6months

2.BERA is most accurate at; a.28wk b.30 wk c.32wk d.34wk

3. True about BERA findings in acoustic neuroma actatency of waveV increased in the affected ear b.Latency of wave I increased in the affected ear c.No waves on BERA d. No change in BERA

4. Brainstrem evoked response audiometry true is:
a. Can differentiate cochlear and retrocochlear lesion
b. cannot differentiate the site of lesion in sensorineural hearing loss
c.. Can differentiate barbiturate poisionong from other causes of coma
d. As a screening procedure for infants



Annexure 5 **Course/Training Feedback Form Student Feedback Form**

Course Name: Hands on training and an ExtensiveDiscussion on BERA

Subject Code: ENT08

Name of Student: _____ Roll No.: _____

We are constantly looking to improve our classes and deliver the best training to you. Your

evaluations, comments and suggestions will help us to improve our performance

SI. NO	Particulars	1	2	3	4	5
1	Objective of the course is clear					
2	Course contents met with your expectations					
3	Lecturer sequence was well planned					
4	Lectures were clear and easy to understand					
5	Teaching aids were effective					
6	Instructors encourage interaction and were helpful					
7	The level of the course					
8	Overall rating of the course	1	2	3	4	5

* Rating: 5 – Outstanding; 4 - Excellent; 3 – Good; 2 – Satisfactory; 1 - Not-Satisfactory

Suggestions if any:

Annexure 5

Course/Training Feedback Form

Student Feedback Form

Course Name: Hands on training and an ExtensiveDiscussion on BERA

Subject Code: ENT08

Name of Student: HARIHARAN, 5

Roll No .: U15 MB292

We are constantly looking to improve our classes and deliver the best training to you. Your

evaluations, comments and suggestions will help us to improve our performance

SI.	Particulars	1	2	3	4	5
1	Objective of the course is clear	<				
2	Course contents met with your expectations		-			
3	Lecturer sequence was well planned		1			1914
4	Lectures were clear and easy to understand	1	-			
5	Teaching aids were effective		/	T		
6	Instructors encourage interaction and were helpful			1		1
7	The level of the course	1			/	T
8	Overall rating of the course	1	2	3	4	5

* Rating: 5 - Outstanding; 4 - Excellent; 3 - Good; 2- Satisfactory; 1 - Not-Satisfactory

Excellent.

Suggestions if any:

Annexure 5

Course/Training Feedback Form

Student Feedback Form

Course Name: Hands on training and an ExtensiveDiscussion on BERA

Subject Code: ENTOS

Name of Student: L-HARINI Roll No.: MSM B 293

We are constantly looking to improve our classes and deliver the best training to you. Your evaluations, comments and suggestions will help us to improve our performance

SL NO	Particulars	1	2	3	4	
1	Objective of the course is clear	-	1			-
2	Course contents met with your expectations		-	0		
3	Lecturer sequence was well planned	-	-	- 1		
4	Lectures were clear and easy to understand			0	1	
5	Teaching aids were effective			- /		
6	Instructors encourage interaction and were helpful			0	1	
7	The level of the course					
8	Overall rating of the course	1	2		1	

- Excellent; 3 - Good; 2- Satisfactory; 1 - Not-Satisfactory naing;

Suggestions if any:

Jansfactory	
P A	

From Dr.Venkataramanan.R, Dept of Otorhinolaryngology, SLIMS Bharath Institute of Higher Education and Research, Puducherry.

Through Proper Channel

To The Dean, SLIMS, Bharath Institute of Higher Education and Research, Puducherry.

Sub: Completion of value-added course: Hands on training and an ExtensiveDiscussion on BERA reg.

Dear Sir,

With reference to the subject mentioned above, the department has conducted thevalue-added course titled:**Hands on training and an Extensive Discussion on BERA** on Sep 2018 to Dec2018. We solicit your kind action to send certificates for the participants, that is attached with this letter. Also, I am attaching the photographs captured during the conduct of the course. Kind Regards

Dr.K.R.Jothikumar <HOD Sign and Seal>

Tr.R.Jymm Seal & Signature of the HOD PROFESSOR & HOD DEPARTMENT OF E.N.T Sri Lakshmi Narayana Institute Of Medical Sciences PONDICHERRY - 605 502

