**Evoked Potentials**

**Evoked Potential (EP)** – electrical response recorded from CNS, elicited by external stimuli; synonyms: EVENT-RELATED POTENTIAL (ERP) or EVENT-RELATED RESPONSE (ERR)

**CLINICAL USES**

1. Assessing functional integrity (and detecting lesions) in different pathways under study.
   - most useful when identify subclinical abnormalities (esp. in multiple sclerosis) or confirm abnormalities corresponding to vague or equivocal symptoms.
   - may reveal abnormalities missed by MRI, and vice versa.
   - precise localization on basis of electrophysiological findings may not be possible (because generators of many components of EP are unknown).
   - changes produced by disease states:
     1. delayed responses - reflect conduction delays in responsible pathways.
     2. attenuation or loss of component waveforms – reflect conduction block or dysfunction of responsible generator.


3. Evaluating patients in coma, suspected brain death for BAER role - see p. E280 >>

ıt is most accurate in assessment of neurologic outcome: patients with absent cortical SEPs bilaterally are unlikely to recover cognition (esp. bilateral loss of N20 response after median stimulation is associated with fatal outcome or development of persistent vegetative state).
  - presence of normal SEPs does not predict useful recovery.

4. Determining completeness of lesion in spinal cord injuries.
   - absence of any cortical response in acute stage doesn’t mean that lesion is complete;
   - preserved responses (or their early return) indicate better prognosis.

5. Determining auditory acuity in patients whose age / mental state precludes their cooperation for behavioral testing. see p. E280 >>

6. Intracranial monitoring see below >>

**SOMATOSENSORY EVOKED POTENTIALS (SSSEP)**

Stimulation of sensory systems leads to generation of CORTICAL EVOKED POTENTIALS - can be recorded with exploring electrode (connected to another electrode at indifferent point some distance away):
- a) over scalp (surface electrode)
- b) over parietal cortex (samples activity to depth of only 0.3-0.6 mm)
- c) microelectrode (inserted in layers 2-6 of underlying cortex)
  - best seen in animals under barbiturate anesthesia (eliminates background electrical activity).
  - in unanesthetized animals / humans, evoked potential is obscured by spontaneous brain activity (i.e. not apparent in ordinary EEG); evoked potential can be demonstrated by superimposing multiple traces - signal averaging technique (signals that are time locked to stimuli are enhanced, whereas background EEG activity is averaged out).

**PRIMARY EVOKED POTENTIAL**

- latency 5-12 ms; latency and morphology depends on eliciting stimulus.
- highly specific in location (can be observed only over primary receiving area for particular sense).
- primary response is negative- positive when it is recorded with microelectrode (indicates depolarization on dendrites and somas in cortex, followed by hyperpolarization).

2. Record positive-negative wave sequence is DIFFUSE SECONDARY RESPONSE

- larger, more prolonged; latency 20-80 ms.
- not highly localized - appears at same time over most of cortex - due to activity in projections from medulla and related thalamic nuclei (not due to lateral spread of primary potential).

- 7.5-Hz ELECTRICAL STIMULATION of peripheral nerve:
  - a) sufficient to produce slight muscle twitch (when mixed nerve is stimulated)
  - b) sufficient to generate sensory nerve action potential that is ≥ 50% of maximum (when sensory nerve is stimulated).
  - best recorded with SURFACE ELECTRODES:
    - a) bipolar derivation (both recording electrodes placed on scalp - over posterior and lateral regions).
    - b) referential derivation involving noncephalic reference electrode:

**BRAIN STEM AUDITORY EVOKED RESPONSE (BAER)** see p. E280 >>

**VISUAL EVOKED POTENTIALS (VEP)** see p. E280 >>

**OLFACTORY EVOKED POTENTIALS** see p. CNI >>

**Electrical brain activity is either spontaneous or event-related (i.e. elicited by stimulus).**

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Clinical Uses

1. SOMATOSENSORY EVOKED POTENTIALS (SSSEP) ... Clinical Uses

2. Trigeminal Evoked Responses...

3. MAGNETIC STIMULATION, MOTOR EVOKED POTENTIALS (MEP)...

4. COGNITIVE EVOKED POTENTIALS...

5. INTRAOPERATIVE MONITORING (IOM)...

6. Indications...

7. BIPOLAR DERIVATION...

8. SSEP...

9. Sensitivity, specificity...

10. Stagnara test...

11. CN5-7, 10 monitoring...

12. Brain Stem Auditory Evoked Response (BAER)...

13. Visual Evoked Potentials (VEP)...

14. Olfactory Evoked Potentials...

15. Electrical brain activity is either spontaneous or event-related (i.e. elicited by stimulus).
- over cervical spine, Erb point (for median nerve stimulation at wrist);
- over lumbar spine, popliteal fossa (for peroneal or posterior tibial nerve stimulation at ankle).

- response is small - necessary to average 2000 responses in arm or 4000 responses in leg.

N.B. physiological transmission must be distinguished from electrical conduction!

SEP components are defined by polarity (P/N) and latency (number); obligate components:
in arm nerve stimulation:
- P9 - activity at or just beyond brachial plexus.
- P13-P14 - activity in medial lemniscus (P13 in cervical cord, P14 in lower brain stem).
- N18 - rostral brain stem.
- N20 - primary somatosensory cortex.
in tibial nerve stimulation at ankle:
- P38 - primary somatosensory cortex.

Most important features:
1. presence or absence of obligate components; amplitude size is not so important.
2. absolute and interpeak latencies of components (N.B. absolute latency of individual components, but not interpeak latency, varies with limb length!).

CLINICAL USES

1. Intraoperative monitoring see below >>

2. Detecting lesions of somatosensory pathways within CNS (esp. dorsal column-mediallemniscal system):
   - SEP is abnormal in multiple sclerosis (≈ 80%) - loss (or marked attenuation) of cervical response after median stimulation, increase in central conduction time.
   - abnormally large amplitude SEP (enhanced cortical excitability) are seen in progressive myoclonus epilepsy, photosensitive epilepsy, late infantile ceroid lipofuscinosis.

3. Little value in evaluating peripheral nervous system (except functional integrity of nerves that are not easily accessible for conventional nerve conduction studies); e.g. SEPs to evaluate radicalopathies;
   - stimulation of polysegmental nerve trunk would not cause abnormal SEP in isolated root lesions;
   - cutaneous nerve or dermatomal stimulation gives conflicting results.

Median-elicited SEP
(EP - Erb's point; CV 5 - 5th cervical spine; i - ipsilateral; c - contralateral):
A. Normal subject.
B. Multiple sclerosis (Erb point potential is present but responses over cervical spine and scalp are absent).

Tibial-elicited SEP
(PP - popliteal fossa, L1 - 1st lumbar spine; L3 - 3rd lumbar spine; d - distal; p - proximal):
A. Normal subject.
B. Multiple sclerosis (no response over scalp).
TRIGEMINAL EVOKED RESPONSES

- infraorbital nerve is stimulated with electrode inserted into infraorbital foramen.
- recording electrodes are placed at Cz with reference to C7 vertebral spinous process (Cv7).
- bilateral studies provide opportunity for evaluation of control and determination of symmetry.
- waves 1 (entrance of maxillary division into gasserian ganglion), 2 (root entry zone into pons), and 3 (trigeminal tract within pons) have latencies of 0.88, 1.80, and 2.44 ms, respectively.
- interwave latencies 1-2 and 2-3 are 0.90 and 1.55 ms, respectively; increase in wave 1 latency > 0.32 ms (when compared to normal side) is considered abnormal.
- absence of waves 2 and 3 is after successful surgery for trigeminal neuralgia.
- evoked responses may also be obtained from stimulation of supraorbital nerve (harder to obtain, need to anesthetize scalp).

MAGNETIC STIMULATION, MOTOR EVOKED POTENTIALS (MEP)

- magnetic stimulation of brain / spine elicits motor evoked potentials (i.e. compound muscle action potential over appropriate target muscle).
- assesses descending motor pathways!
- magnetic stimulation of peripheral nerves elicits somatosensory evoked potentials.
- assesses ascending sensory pathways!

- magnetic stimulation effects are similar to electrical stimulation.
- magnetic impulses travel through tissues painlessly and without attenuation* (vs. electrical impulses).
- *but magnetic impulses decrease in relation to inverse square of distance from stimulator coil
- procedure is noninvasive (!), painless and apparently safe.
- latency of motor responses can be measured.
- central conduction time can be estimated by comparing latency of cerebral and spinal stimulation.
- motor latencies↑↑↑ in MS, cerebral myelopathy, cervical spondylosis, spinal cord trauma, hemiplegia, hereditary spastic paraparesis, etc.
- clinical utility is investigational.
- with development of accurate focal stimulation, cortical mapping could be done noninvasively!

Electrical stimulation (painful in alert patients) may be preferable for intraoperative monitoring where patient is anesthetized and paralyzed, since equipment is less complicated to organize in operating room environment;
- response is best recorded from peripheral nerves, using needle electrodes.

COGNITIVE EVOKED POTENTIALS

- evoked potential components depending upon mental attention of subject and setting in which stimulus occurs (rather than on physical characteristics of stimulus), i.e. such endogenous "event-related" potentials (ERP) are related to cognitive aspects of distinguishing infrequently occurring target stimulus from other stimuli occurring more frequently (usually randomly alternating low and high pitch auditory stimuli).
- most important is P3 component (s. P300 component - because of 300 ms latency after auditory target stimulus).
- P3 latency is prolonged in dementia.
- P3 is normal in depression or other psychiatric disorders (that might be mistaken for dementia).
INTRAOPERATIVE MONITORING (IOM)

IOMs can reliably detect and predict neurological damage but there are 2 major problems:

a) it is too late (damage is done)

b) IOM is not useful if corrective action is not available

- when evoked potential abnormality occurs during surgical procedure, it is hoped that alteration / reversal of procedure will minimize damage; examples:
  - monitoring CN2 (visual evoked potentials) during transsphenoidal removal of pituitary tumor.
  - monitoring CN7 and CN8 (brain stem auditory evoked potentials) during posterior fossa surgery.
  - monitoring spinal cord (somatosensory evoked potentials, motor evoked potentials) during scoliosis / myelomeningocele / intramedullary tumor / degenerative * cervical spine surgery; repair of coartation of aorta.
- *reduces complications rate 10-fold (because effective corrective action exists if IOM signal changes – popping rod)
- **most likely no benefit at all (studies show, IOM does not prevent complications)

- for kids < 4 years old, white matter long tracts are immature – motor evoked potentials, SSEP are unreliable.

INDICATIONS

Spine surgery:
1) severe spinal cord compression
2) deformity correction
3) intradural tumor removal

Avoid use for simpler surgeries (e.g. ACDF without myelopathy, lumbar microdiscectomy).

MEP

N.B. MEP (motor evoked potentials) is gold standard but are highly affected by anesthesia and muscle relaxation in particular.

- need to pause surgery to run stimulation – gives warning too late!
- run baseline after patient positioning

SSEP

- use dorsal column pathway to assess somatosensory cortex noninvasively.

- frequent* stimulation of bilateral median or posterior tibial nerves --> response measurement via contralateral cortical electrodes.
- *averages signal over several minutes – gives warning too late
- changes in latency or amplitude of SSEP waveforms indicate disruption of somatosensory pathway.

What changes in SSEP should trigger concern:

a) increased signal LATENCY (typically > 10% prolongation)

b) decreased signal AMPLITUDE (typically > 50% reduction)

- change is called irreversible when it fails to return to baseline before end of procedure.

False-positive SSEP signal change may be caused by:

1) blood pressure (mean and diastolic)
amplitudes of SSEPs are very sensitive to changes in mean arterial pressure, making them useful for detecting ischemia.

2) heart rate
3) temperature
4) partial pressure of alveolar carbon dioxide
5) anesthetic drugs

Anesthesia considerations:
- all volatile anesthetics produce dose-dependent reduction in SSEP peak amplitude and increase in peak latency; adding nitrous oxide increases this sensitivity to anesthetic agents.
- helpful anesthesia measures:
  - minimize pentothal dose during induction (produces 30 minutes of suppression of EPs), or use ETOMIDATE (which increases both SSEP amplitude and latency)
  - total intravenous anesthesia is ideal; use < 1 MAC (maximal allowable concentration), ideally < 0.5 MAC
  - avoid older agents such as Halothane
- nondepolarizing muscle relaxants have little effect on EP (in monkeys)
- PROPOFOL has a mild effect on EP; total anesthesia with propofol causes less EP depression than inhalational agents at the same depth of anesthesia
- continuous infusion of anesthetic drugs is preferred over intermittent boluses
- hypocapnia (down to end tidal CO\textsubscript{2} = 21) causes minimal reduction in peak latencies
- antiepileptic drugs (phenytoin, carbamazepine, phenobarbital) do not affect SSEP.

Sensitivity, Specificity
- SSEP sensitivity 99%, specificity only 27%.
- MEP sensitivity 90-100%, specificity 90-100%.
- combined SSEP+MEP sensitivity 81% and specificity 99%

STACUS-7 TEST
- awakening patient during surgery (e.g. under remifentanil balanced anesthesia) and performing neuro exam.
4. Connect electrodes to Patient Interface. Avoid entwining anode return (white) with other electrodes. Attach Patient Interface to the NIM-Response 2.0.

5. Plug in power cord and attach to the NIM-Response 2.0. Turn unit on. Confirm the NIM-Response 2.0 test results. Pass, Pass, Pass

6. Select physician setting from Quick Start menu. Verify the Stimulus setting is what the surgeon has requested.

7. Check baseline. Peak values = 5–15uV.

8. Press "ELECTRODES" tab to check Impedance and Difference. Reposition if necessary. Press "EMG" tab to return to monitoring.

9. Clip Muting Detector to ESU. For bipolar ESU, route single conductor cable through the clamp. Do not include grounding pad. Plug into Muting Probe Input jack #3 on rear panel of unit.

10. Connect the Prass Monopolar Stimulating Probe to the black (-) Stim 1 jack of the Patient Interface.

Please refer to the NIM-Response 2.0 User’s Guide (825061) for complete operating instructions. For technical questions or guidance through OR protocol with the NIM-Response 2.0, call the NIM+ Helpline or Medtronic ENT Customer Service at 1-800-874-5797 or 904-296-9600. For general information, you may also visit our website at www.MedtronicENT.com.