Clinical Policy: Ambulatory EEG
Reference Number: CP.MP.96
Effective Date: 09/2015
Last Review Date: 09/2015

IMPORTANT REMINDER
This Clinical Policy has been developed by appropriately experienced and licensed health care professionals based on a thorough review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by the policy; and other indicia of medical necessity. Centene Corporation makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this policy.

The purpose of this Clinical Policy is to provide a guide to medical necessity. Benefit determinations should be based on the applicable contract provisions governing plan benefits (“Benefit Plan Contract”) and applicable state and federal requirements, as well as applicable plan-level administrative policies and procedures. To the extent there are any conflicts between this Clinical Policy and the Benefit Plan Contract provisions, the Benefit Plan Contract provisions will control.

Clinical policies are intended to be reflective of current scientific research and clinical thinking. This policy is current at the time of approval, may be updated and therefore is subject to change. This Clinical Policy is not intended to dictate to providers how to practice medicine, nor does it constitute a contract or guarantee regarding payment or results. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members.

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Note: For Medicare members, to ensure consistency with the Medicare National Coverage Determinations (NCD) and Local Coverage Determinations (LCD), all applicable NCDs and LCDs should be reviewed prior to applying the criteria set forth in this Policy. Refer to the CMS website at http://www.cms.gov for additional information.

Description
Ambulatory electroencephalogram (EEG) testing in the outpatient setting (e.g., at home) is a diagnostic test used to evaluate an individual in whom a seizure disorder is suspected but not conclusively confirmed by the person's medical history, physical examination, and a previous routine or standard (awake and asleep) EEG.

Policy/Criteria
I. It is the policy of health plans affiliated with Centene Corporation® that ambulatory EEG is medically necessary following an inconclusive or nondiagnostic standard (awake and asleep) EEG for any of the following indications:
   A. To investigate episodic events where epilepsy is suspected but the history, examination, and routine EEG do not resolve the diagnostic uncertainties;
B. To confirm epilepsy in those individuals experiencing suspected nonepileptic events or for classification of seizure type;
C. To differentiate between neurological and cardiac related episodes;
D. To adjust antiepileptic medication levels;
E. To localize seizure focus for enhanced patient management;
F. To identify and medicate absence seizures;
G. To differentiate between epileptic and sleep disorder related episodes; and/or
H. To evaluate seizures precipitated by naturally occurring cyclic events or environmental stimuli that are not reproducible in the hospital or clinic setting.

II. It is the policy of health plans affiliated with Centene Corporation® that ambulatory EEG is considered not medically necessary for the following indications:
A. Studies of neonates;
B. Studies of unattended, non-cooperative patients; and/or
C. Studies to localize seizure focus/foci when the seizure symptoms and/or other EEG recordings indicate the presence of bilateral foci or rapid generalization.

Ambulatory EEG (CPT code 95950 or 95953) should always be preceded by an awake and drowsy/sleep EEG (CPT code 95816, 95819, 95822 or 95827).

Background
In most instances, a standard EEG performed at a clinic or outpatient epilepsy facility can identify brain activity specific to seizures; however, when routine EEG is inconclusive and the clinical history strongly suggests seizure activity, an ambulatory EEG may be indicated. An ambulatory EEG may increase the chance of detecting an epileptiform abnormality in these individuals and significantly impact clinical management. An estimated 12% to 25% of individuals who previously had a normal or non-diagnostic routine EEG have epileptiform activity on ambulatory EEG. ³

Clinical events known as psychogenic nonepileptic spells (PNES) (previously referred to as pseudoseizures) are nonepileptic seizures where the person perceives altered movement, emotion, sensation, or an experience similar to those involved with epilepsy. These events are without an EEG-documented ictal association. PNES occur in as many as 20% of persons evaluated at inpatient epilepsy monitoring centers and in 5% to 20% of outpatient populations. Both PNES and epileptic seizures are concurrent in an estimated 10% to 60% of individuals with epilepsy. ³

Ambulatory EEG recordings can be utilized in the evaluation and differential diagnosis of other conditions, including cardiac arrhythmias, sleep disorders, syncope, and transient ischemic attacks, if these episodes are not diagnosed by conventional studies. It may also allow an estimate of seizure frequency, which may at times help to evaluate the effectiveness of a drug and determine its appropriate dosage.

Ambulatory EEG testing provides a continuous recording of the brain's electrical activity that can range from several hours to several days (typically 48 hours to 72 hours). In the outpatient setting (physician office, clinic), a set of electrodes with leads is secured to the person's scalp and
a digital recording unit is attached to the waist or a shoulder harness. Currently, portable recordings of up to 32 channels can record computer-assisted spike and seizure detection rates over several days. Event detection computer software is designed to increase the chance of recording an ictal event during a seizure or interictal epileptiform discharges occurring between seizures, during the person's routine daily activities and sleep. The person being tested and observers (family members, caregiver) have the opportunity to "tag" portions of the recording during clinical events using a push button device to signal when an observable event occurs.

The gold standard for evaluating the large amount of data collected by a computer-assisted system is visual analysis at the end of the testing period by a highly trained individual. Digital analysis of an EEG can be used to diagnose neurological conditions when routine EEG outcomes and neurological imaging are inconclusive to confirm suspicious but nondiagnostic symptoms. Digital analysis of an EEG requires the analysis of an EEG using quantitative analytical techniques such as data selection, quantitative software processing, and dipole source analysis.

**Coding Implications**

The following is a list of procedures codes for which coverage may be provided when billed with a diagnosis code(s) that supports medical necessity criteria (see list of ICD-10-CM codes supporting medical necessity further below). They are current at time of review of this policy. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

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<th>CPT®* Codes</th>
<th>Description</th>
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<tr>
<td>95950</td>
<td>Monitoring for identification and lateralization of cerebral seizure focus, electroencephalographic (eg, 8 channel EEG) recording and interpretation, each 24 hours</td>
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<tr>
<td>95953</td>
<td>Monitoring for localization of cerebral seizure focus by computerized portable 16 or more channel EEG, electroencephalographic (EEG) recording and interpretation, each 24 hours, unattended</td>
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**ICD-10-CM Diagnosis Codes that Support Coverage Criteria**

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<th>ICD-10-CM Code</th>
<th>Description</th>
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<tr>
<td>F44.5</td>
<td>Conversion disorder with seizures or convulsions</td>
</tr>
<tr>
<td>G40.00 – G40.B19</td>
<td>Epilepsy and recurrent seizures</td>
</tr>
<tr>
<td>G40.811 – G40.814</td>
<td>Lennox-Gastaut syndrome</td>
</tr>
<tr>
<td>R25.0 – R25.8</td>
<td>Abnormal involuntary movements</td>
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<tr>
<td>R56.1</td>
<td>Post-traumatic seizures</td>
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<tr>
<td>R56.9</td>
<td>Unspecified convulsions</td>
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**Reviews, Revisions, and Approvals**

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References