3.01.03 Quantitative Electroencephalography as a Diagnostic Aid for Attention-Deficit/Hyperactivity Disorder

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<th>Original Policy Date:</th>
<th>February 1, 2016</th>
<th>Effective Date:</th>
<th>December 1, 2017</th>
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<tr>
<td>Section:</td>
<td>3.0 Mental Health</td>
<td>Page:</td>
<td>Page 1 of 9</td>
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**Policy Statement**

Quantitative electroencephalographic-based assessment of the theta/beta ratio is considered **investigational** as a diagnostic aid for attention-deficit/hyperactivity disorder.

**Policy Guidelines**

This testing would likely be reported with existing electroencephalography CPT codes. The clinician would report the appropriate code for electroencephalography (e.g., 95812-95813) and the code for digital analysis of electroencephalogram (95957) would be reported for the analysis.

**Description**

Patients with attention-deficit/hyperactivity disorder (ADHD) may have alterations in their brain wave patterns that can be measured by quantitative electroencephalography. A commercially available system, the Neuropsychiatric EEG-based ADHD Assessment Aid (NEBA), measures the resting theta/beta ratio of the electroencephalogram. This technology is being evaluated to aid in the diagnosis of ADHD in adolescents and children for whom there is a clinical suspicion of ADHD.

**Related Policies**

- Neurofeedback

**Benefit Application**

Benefit determinations should be based in all cases on the applicable contract language. To the extent there are any conflicts between these guidelines and the contract language, the contract language will control. Please refer to the member's contract benefits in effect at the time of service to determine coverage or non-coverage of these services as it applies to an individual member.

Some state or federal mandates [e.g., Federal Employee Program (FEP)] prohibits plans from denying Food and Drug Administration (FDA)-approved technologies as investigational. In these instances, plans may have to consider the coverage eligibility of FDA-approved technologies on the basis of medical necessity alone.

**Regulatory Status**

In 2011, the generic device Neuropsychiatric Interpretive Electroencephalograph Assessment Aid was granted a de novo 510(k) classification by the U.S. Food and Drug Administration (FDA; class II, special controls, product code: NCG). According to the FDA documentation, a neuropsychiatric interpretive electroencephalograph assessment aid is a device prescribed by a physician that uses a patient's electroencephalogram to provide an interpretation of the patient's neuropsychiatric condition. In addition to the general controls, approval of these devices is subject to a number of special controls, including the following:

- Clinical performance testing must demonstrate the accuracy, precision, and reproducibility of the EEG-based interpretation, including any specified equivocal ones (cutoffs).
Clinical performance testing must demonstrate the ability of the device to function as an assessment aid for the medical condition for which the device is indicated. Performance measures must demonstrate device performance characteristics per the intended use in the intended use environment. Performance measurements must include sensitivity, specificity, positive predictive value, and negative predictive value per the device intended use. Repeatability of measurement must be demonstrated using interclass correlation coefficients and illustrated by qualitative scatterplots.

The device design must include safeguards to prevent use of the device as a stand-alone diagnostic.

The labeling must bear all information required for the safe and effective use of the device.

In 2013, the Neuropsychiatric EEG-based Assessment Aid (NEBA®; Lexicor Medical Technology, Augusta, GA) for ADHD was granted a de novo 510(k) classification by the FDA. The device is indicated to measure the theta/beta ratio of the electroencephalogram at electrode CZ on patients 6 to 17 years of age, combined with a clinician’s evaluation, to aid in the diagnosis of ADHD (K112711). NEBA® should only be used by a clinician as confirmatory support for a completed clinical evaluation or as support for the clinician’s decision to pursue further testing following a clinical evaluation. The device is not intended to be used as a stand-alone in the evaluation or diagnosis of ADHD.

The Lexicor QEEG system is marketed as a diagnostic aid for ADHD. Lexicor Medical Technology provides an Internet analysis service of the QEEG, producing a DataLex report. FDA product code: NCG.

Rationale

Background

Attention-Deficit/Hyperactivity Disorder

Attention-deficit/hyperactivity disorder (ADHD) is a common disorder in children, adolescents, and adults defined by pervasive symptoms of inattention and/or hyperactivity-impulsivity, which lead to impairment in at least 2 domains of the work, school, or home environments. Stimulant medications reduce symptoms associated with ADHD, although there are concerns about the potential for overdiagnosis and overprescribing of medication.

Diagnosis

Presently, ADHD is diagnosed clinically by assessing behavioral symptoms and impairment via interviews and standard questionnaires. Diagnosis can be challenging because the core symptoms are nonspecific. They may be present in other psychiatric disorders (e.g., learning disabilities, conduct disorders, affective disorders) or result from environmental influences such as a lack of discipline. Also, ADHD is a heterogeneous disorder with multiple subtypes, and frequently coexists with other psychiatric disorders.

There has been a substantial amount of research over the last several decades on whether electroencephalography (EEG) – derived brain wave patterns in patients with ADHD differ from those without ADHD. EEG patterns are typically categorized into 4 frequency ranges: delta (<4 Hz), theta (4-7 Hz), alpha (8-12 Hz), and beta (13-25 Hz). The largest focus of research on brain wave patterns in ADHD has been on whether there is increased theta wave activity and an increased theta/beta ratio in ADHD patients.

The Neuropsychiatric EEG-based ADHD Assessment Aid (NEBA) system is a specific quantitative electroencephalography (QEEG) system that measures the resting theta/beta ratio of the EEG with an electrode located at the central midline position (referred to as position CZ in the international 10-20 EEG system). QEEG uses computer analysis with mathematical transformation from the time domain into the frequency domain (fast-Fourier transform) to determine the total power at each frequency. The relative power of the waveform can then be calculated in

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relation to the total power of the 4 frequency ranges. The NEBA system uses proprietary cutoffs to generate an estimate of the likelihood of ADHD based on the resting theta/beta ratio.

It is proposed that the NEBA system can be used to confirm a clinical diagnosis or support further testing in children and adolescents with ADHD. The system is not intended to evaluate patients in whom the clinician’s diagnosis of ADHD is negative, and the system does not generate an interpretive report in this situation. It is also proposed that the clinician’s diagnostic impression plus the results generated by the NEBA system may reduce the potential for overdiagnosis of ADHD, and thereby reduce the risks of administering unnecessary pharmacologic therapy in the intended use population. Also, as a result of research on EEG brain waves in ADHD, neurofeedback has been developed as a potential treatment for ADHD (see Blue Shield of California Medical Policy: Neurofeedback). This treatment employs principles of biofeedback using EEG brain wave activity and attempts to alter the brain wave patterns in beneficial ways.

Literature Review
Assessment of a diagnostic technology typically focuses on 3 categories of evidence: (1) its technical reliability (test-retest reliability or interrater reliability); (2) clinical validity (sensitivity, specificity, and positive and negative predictive value [NPV]) in relevant populations of patients; and (3) clinical utility demonstrating that the diagnostic information can be used to improve patient outcomes. Subsequent use of a technology outside of the investigational setting may also be evaluated.

Quantitative Electroencephalography
TEC Assessment Overview
A Blue Cross Blue Shield Association Technology Evaluation Center (TEC) Assessment published in 2014 evaluated the evidence related to the use of quantitative electroencephalography (QEEG) with the Neuropsychiatric EEG-based ADHD Assessment Aid (NEBA) system in the diagnosis of attention-deficit/hyperactivity disorder (ADHD).1 This evidence was submitted to the U.S. Food and Drug Administration (FDA) in 2013 and subsequently published by Snyder et al (of NEBA Health) in 2015.2,3 The evidence related to the diagnostic accuracy of NEBA for the diagnosis of ADHD consisted of data submitted to the FDA from 275 children and adolescents (6-18 years) who presented with attention and/or behavioral concerns to one of 13 clinics in the United States. The evidence related to the technical performance of NEBA for the diagnosis of ADHD included test-retest reliability of the NEBA theta/beta ratio for EEG data from 198 patients who had recordings on 2 different days. These studies are described in greater detail in the sections on technical reliability and clinical validity. No studies were identified that assessed whether the reclassification of patients suspected of having ADHD, as reported to the FDA, improves health outcomes.

Technical Reliability
Data submitted to the FDA for the NEBA system included test-retest reliability of the theta/beta ratio for EEG data from 198 patients who had recordings on 2 different days (≈2.5 weeks apart on average).2 EEG data were collected for 10 minutes with patients seated in a chair with eyes open and fixed with attention to a point at eye level on the wall. Epochs (30 seconds in duration) with substantial artifact were screened out by EEG processing technicians at a central facility. Analysis of EEG data required at least 15 epochs of data with little to no artifact. There were 198 patients with 2 sessions of data out of the per protocol set of 275. The theta/beta cutoffs were revised prior to the blind break based on data from a prior study.4 The intraclass correlation coefficient of repeated NEBA theta/beta ratio was 0.83, which is considered to be high.

Clinical Validity
A number of studies have measured theta activity or the theta/beta ratio in children and adolescents with ADHD compared with nonaffected controls. The most commonly reported alteration in EEG is an increase in the theta/beta ratio. However, some studies have reported that other patterns (e.g., increased beta wave activity) are found in some patients, and several
recent studies have found no significant difference in theta activity in a clinical vs nonclinical population.

A 2005 systematic review included 17 studies evaluating theta activity in children and adolescents with ADHD. The meta-analysis found a weighted mean effect size of 0.59 for an absolute increase in theta activity and a mean effect size of 0.91 for theta relative to total EEG activity. A 2006 systematic review by Snyder and Hall included 9 studies (total N=1498 patients) that used DSM-IV (Diagnostic and Statistical Manual of Mental Disorders, 4th Edition) criteria and screening tests in a clinical setting. The meta-analysis identified a mean increase of theta power of 32% and a pooled effect size of 3.08 for the theta/beta ratio in patients who had ADHD compared with unaffected children, adolescents, and adults. It was noted that the included studies often had retrospectively defined limits and that an increase in the theta/beta ratio has also been identified in other conditions. A 2001 study by Monastra et al found a change in QEEG at CZ in a large study (N=469) of children and adolescents who were diagnosed with ADHD vs controls, although this study used an attention index measured across 4 different tasks (eyes fixed, silent reading, listening, drawing).

Other studies found no significant difference in theta activity in a clinical population with ADHD. In 2013, Liechti et al compared the theta/beta ratio in 32 children and 22 adults with ADHD vs healthy controls who were matched for age, sex, and IQ. Resting EEG was measured separately for the 3 midline electrodes (frontal [FZ], central [CZ], parietal [PZ]) and for frontal, central, and parietal regions. The study found a decrease in theta with age, but no consistent increase in theta or theta/beta ratio in patients with ADHD compared with controls. There was no evidence for a maturational lag in patients with ADHD. In 2012, Ogrim et al assessed differences in theta activity measured at CZ in 62 children and adolescents with a tentative diagnosis of ADHD compared with 39 sex- and age-matched controls. The overall accuracy at CZ was 63% for theta and 58% for the theta/beta ratio compared with nonaffected controls. Elevations of theta were found in 25.8% of patients compared with 2.6% of controls. None of the EEG measures were statistically significant for discerning patients from controls. In other studies, subgroups of children with ADHD have been shown to have increased beta activity instead of decreased beta or increased theta/beta ratio. In a study of QEEG in combination with the Connors’ Continuous Performance Test and the Test of Variables of Attention in the diagnosis of ADHD in 157 children, 85 with and 72 without ADHD based on Diagnostic Interview Schedule for Children (Version IV), Kim et al (2015) reported that the ADHD group had significantly higher theta wave values in 13 positions, while for the theta/beta ratio, the ADHD group had higher values for only 1 position (right frontal lobe).

Sensitivity, Specificity, and Predictive Value of the Theta/Beta Ratio in the Diagnosis of ADHD

Data submitted to FDA regarding the diagnostic accuracy of the NEBA system were from the multicenter study of 275 children and adolescents (6-18 years, described above) who presented with attention and/or behavioral concerns to one of 13 clinics in the United States. An additional 89 children and adolescents were recruited but did not complete the study, and, of these, 67 had incomplete EEG recordings. Diagnostic evaluation for ADHD and other disorders was conducted with a clinical interview and rating scales that included behavior rating scales, IQ and achievement testing, and scales of severity and dysfunction. A consensus best-estimated diagnosis was determined by a multidisciplinary clinical team composed of a clinical psychologist, a neurodevelopmental pediatrician, and a child/adolescent psychiatrist. The clinical team had access to de-identified patient files; however, they did not interview patients or have access to the parent rating scales, features considered critical for a criterion standard diagnosis of ADHD. A separate group of investigators who were unaware of the clinical diagnosis collected the EEG data (NEBA system). When compared with the consensus diagnosis, NEBA had a sensitivity of 89%, specificity of 87%, positive predictive value (PPV) of 81% and NPV of 93% for adolescents (12-17 years). For children (ages 6-11 years), NEBA had a sensitivity of 79%, specificity of 97%, PPV of 96% and NPV of 82%. The investigators calculated that the addition of NEBA to the clinician’s ADHD evaluation would have improved the clinician’s diagnostic accuracy from 61% to 88%. This calculation is based on the 275 patients who completed the...
protocol, rather than the intention-to-treat population. The results of this FDA-regulated study suggest that QEEG might be used to decrease the overdiagnosis of ADHD by identifying patients who may not have the disorder. Strengths of this study included its multicenter design and the reclassification analysis of data obtained from a blinded analysis. Limitations were lack of patient interview by the consensus team and lack of intention-to-treat analysis.

Snyder et al also reported on the accuracy of the theta/beta ratio for diagnosis of ADHD in an industry-sponsored, investigator-blinded, multicenter study from 2008. Patients (N=159) ages 6 to 18 who had presented to 1 of 4 psychiatric and pediatric clinics with suspected attention and behavioral symptoms were evaluated in a standardized semi-structured manner according to DSM-IV criteria by a clinical team trained on the study instruments. Rating scales were distributed to parents and teachers and held in sealed envelopes until the blind was broken. EEG was collected separately by investigators, blinded to the clinical diagnosis, using a 19-electrode cap according to the 10-20 system with eyes open and eyes shut. A threshold of 1.5 SD of the theta/beta ratio from normative database values (according to age) at electrode CZ was used to determine ADHD vs non-ADHD. With a prevalence of ADHD of 61% based on clinical diagnosis, the theta/beta ratio had a sensitivity of 87%, specificity of 94%, PPV of 95%, and NPV of 82%. The rating scales provided a sensitivity of 38% to 79% and specificity of 13% to 61%. Results from this study were used to set a new theta/beta threshold for analysis of data from the FDA-regulated study of the NEBA device.

Other studies have reported lower accuracy of QEEG in the diagnosis of ADHD. In the Kim study (2015) previously reported, on receiver operating curve analysis, QEEG theta wave amplitude showed low accuracy for the diagnosis of ADHD (56.4%), and the theta/beta wave amplitude did not significantly predict ADHD diagnosis. Sangal et al (2015) evaluated the discriminatory power of QEEG measurements during auditory and visual tasks requiring selective attention in 28 control children and 58 children with ADHD. Subjects with ADHD had significantly higher average theta/beta ratios (2.6 vs 2.25; p=0.007) and lower average beta-I amplitudes (3.66 vs 4.22, p=0.01). The average theta/beta ratio had a sensitivity and a specificity in diagnosing ADHD of 69% and 50%, respectively, while the theta/beta ratio at the CZ position had sensitivity and specificity of 69% and 43%, respectively.

**Section Summary: Clinical Validity**

Patients who have ADHD may have altered brain wave patterns on QEEG compared with patients who do not. While an increased theta/beta ratio is the most common alteration reported, not all studies have found this association, and some have reported other brain wave patterns in ADHD patients. A few studies have reported on the sensitivity and specificity of quantitative EEG compared with clinical diagnosis. In these studies, sensitivity ranged from 69% to 99% and specificity from 43% to 97%. However, a weakness of these studies is the lack of a true criterion standard for diagnosis of ADHD.

**Clinical Utility**

A proposed benefit of the NEBA system is a reduction in the overdiagnosis of ADHD, thereby lessening the risks of unnecessary pharmacologic therapy in children and adolescents. There were no published studies that directly reported on clinical outcomes, such as measures of disease activity and/or medication use. The pivotal FDA study reported on the reclassification of diagnosis following NEBA; this may be considered an indirect measure that may impact outcomes.

The evidence related to whether QEEG improves the clinical diagnosis of patients with suspected ADHD consists of the material submitted to the FDA as part of the NEBA’s approval process, as previously described. The study included reclassification tables to demonstrate whether NEBA provides additional information beyond the clinician’s initial diagnosis, which are summarized in Table 1. Use of NEBA was consistent with the categorization of patients diagnosed with ADHD by both the initial clinical diagnosis and the consensus diagnosis. For example, 95 (73%) of 130 children and adolescents who were considered to have ADHD by the consensus
diagnosis were classified as ADHD by both the clinician alone and NEBA. Reclassification was observed when using NEBA for patients diagnosed by clinician alone as ADHD and consensus as non-ADHD. For example, 145 children and adolescents had a non-ADHD diagnosis by the consensus. Of the 145, 93 had received an initial clinical diagnosis of ADHD, but 85 (91%) were negative by NEBA.

Table 1. NEBA Reclassification of Patients with Consensus ADHD Diagnosis

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<td>NEBA interpretation</td>
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<tr>
<td></td>
<td>-</td>
<td>21 (18.1)</td>
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<tr>
<td>Total</td>
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</tr>
<tr>
<td>Not ADHD</td>
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<td>NEBA interpretation</td>
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<td>8 (8.6)</td>
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<td></td>
<td>-</td>
<td>85 (91.4)</td>
</tr>
<tr>
<td>Total</td>
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ADHD: attention-deficit hyperactivity disorder; FDA: Food and Drug Administration; NEBA: Neuropsychiatric EEG-Based Assessment Aid; NEBA interpretation: NEBA results plus initial clinical diagnosis.

Section Summary: Clinical Utility

Reclassification results from the pivotal trial suggest that NEBA may support an alternative diagnosis in patients initially suspected of having ADHD but not confirmed by consensus diagnosis. No studies were identified that address whether clinical outcomes are improved for patients with suspected ADHD who are reclassified by NEBA.

Summary of Evidence

For individuals who are suspected of having ADHD who received quantitative electroencephalography, the evidence includes a number of studies on brain wave patterns, particularly the theta/beta ratio. Relevant outcomes are test accuracy, symptoms, functional outcomes, and medication use. Numerous studies have evaluated brain wave patterns with standard electroencephalography equipment, and a pivotal trial, submitted to the U.S. Food and Drug Administration, measured the theta/beta ratio with the NEBA system. In the pivotal trial, both the specificity and positive predictive value of quantitative electroencephalography were high. The reclassification analysis suggests that a negative NEBA might make ADHD less likely, although it is not clear from this study whether the consensus diagnosis was more accurate than the initial clinical diagnosis that included patient interview and parent rating scales. The larger body of evidence also raises questions about the utility of measuring the theta/beta ratio, because it has not been a consistent finding across studies. Given the uncertainty of an increase in the theta/beta ratio in patients with ADHD, additional study is needed to determine whether a low theta/beta ratio can identify children and adolescents who are unlikely to have ADHD. Also, the effect of the test on patient outcomes would allow greater certainty regarding the usefulness of this test. The evidence is insufficient to determine the effects of the technology on health outcomes.
Supplemental Information
Practice Guidelines and Position Statements

American Association of Pediatrics
The American Association of Pediatrics' 2011 practice guidelines on the diagnosis, evaluation, and treatment of attention-deficit/hyperactivity disorder (ADHD) indicated that to make a diagnosis of ADHD, the primary care clinician should determine that Diagnostic and Statistical Manual of Mental Disorders, 4th Edition, Text Revision, criteria have been met (including documentation of impairment in more than 1 major setting), and information should be obtained primarily from reports from parents or guardians, teachers, and other school and mental health clinicians involved in the child’s care.14 The primary care clinician should also rule out any alternative cause (quality of evidence B/strong recommendation). Assessment by quantitative electroencephalography is not mentioned in these guidelines.

American Academy of Neurology
In 2016, the American Academy of Neurology released a technology report on quantitative electroencephalography (EEG) for ADHD.15 The main conclusion of the report was that it remains “unknown whether a combination of standard clinical examination and EEG theta/beta power ratio increases diagnostic certainty of ADHD compared with clinical examination alone.”

U.S. Preventive Services Task Force RECOMMENDATIONS
Not applicable.

Medicare National Coverage
There is no national coverage determination. In the absence of a national coverage determination, coverage decisions are left to the discretion of local Medicare carriers.

Ongoing and Unpublished Clinical Trials
A search of ClinicalTrials.gov in September 2017 did not identify any ongoing or unpublished trials that would likely influence this review.

References

1. Blue Cross and Blue Shield Association Technology Evaluation Center (TEC). Quantitative Electroencephalography as a Diagnostic Aid for Attention-Deficit/Hyperactivity Disorder. TEC Assessments. 2014;Volume 29:Tab 1 PMID

**Documentation for Clinical Review**

- No records required

**Coding**

This Policy relates only to the services or supplies described herein. Benefits may vary according to product design; therefore, contract language should be reviewed before applying the terms of the Policy. Inclusion or exclusion of a procedure, diagnosis or device code(s) does not constitute or imply member coverage or provider reimbursement.

**IE**

The following services may be considered investigational.

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<td>Electroencephalogram (EEG) extended monitoring; 41-60 minutes</td>
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<tr>
<td></td>
<td>95813</td>
<td>Electroencephalogram (EEG) extended monitoring; greater than 1 hour</td>
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<td></td>
<td>95816</td>
<td>Electroencephalogram (EEG); including recording awake and drowsy</td>
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<tr>
<td></td>
<td>95819</td>
<td>Electroencephalogram (EEG); including recording awake and asleep</td>
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<td></td>
<td>95957</td>
<td>Digital analysis of electroencephalogram (EEG) (e.g., for epileptic spike analysis)</td>
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<td>Measurement of Central Nervous Conductivity, External Approach</td>
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<td>Measurement of Central Nervous Electrical Activity, External Approach</td>
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**ICD-10 Diagnosis**

All Diagnoses

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**Policy History**

This section provides a chronological history of the activities, updates and changes that have occurred with this Medical Policy.

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<td>06/01/2017</td>
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**Definitions of Decision Determinations**

**Medically Necessary:** A treatment, procedure, or drug is medically necessary only when it has been established as safe and effective for the particular symptoms or diagnosis, is not investigational or experimental, is not being provided primarily for the convenience of the patient or the provider, and is provided at the most appropriate level to treat the condition.

**Investigational/Experimental:** A treatment, procedure, or drug is investigational when it has not been recognized as safe and effective for use in treating the particular condition in accordance with generally accepted professional medical standards. This includes services where approval by the federal or state governmental is required prior to use, but has not yet been granted.

**Split Evaluation:** Blue Shield of California/Blue Shield of California Life & Health Insurance Company (Blue Shield) policy review can result in a split evaluation, where a treatment, procedure, or drug will be considered to be investigational for certain indications or conditions, but will be deemed safe and effective for other indications or conditions, and therefore potentially medically necessary in those instances.

**Prior Authorization Requirements (as applicable to your plan)**

Within five days before the actual date of service, the provider must confirm with Blue Shield that the member’s health plan coverage is still in effect. Blue Shield reserves the right to revoke an authorization prior to services being rendered based on cancellation of the member’s eligibility. Final determination of benefits will be made after review of the claim for limitations or exclusions.

Questions regarding the applicability of this policy should be directed to the Prior Authorization Department. Please call (800) 541-6652 or visit the provider portal at www.blueshieldca.com/provider.

Disclaimer: This medical policy is a guide in evaluating the medical necessity of a particular service or treatment. Blue Shield of California may consider published peer-reviewed scientific literature, national guidelines, and local standards of practice in developing its medical policy. Federal and state law, as well as contract language, including definitions and specific contract provisions/exclusions, take precedence over medical policy and must be considered first in determining covered services. Member contracts may differ in their benefits. Blue Shield reserves the right to review and update policies as appropriate.