INSTRUCTIONS FOR USE: Company Medical Policies serve as guidance for the administration of plan benefits. Medical policies do not constitute medical advice nor a guarantee of coverage. Company Medical Policies are reviewed annually and are based upon published, peer-reviewed scientific evidence and evidence-based clinical practice guidelines that are available as of the last policy update. The Company reserves the right to determine the application of medical policies and make revisions to medical policies at any time. The scope and availability of all plan benefits are determined in accordance with the applicable coverage agreement. Any conflict or variance between the terms of the coverage agreement and Company Medical Policy will be resolved in favor of the coverage agreement. Coverage decisions are made on the basis of individualized determinations of medical necessity and the experimental or investigational character of the treatment in the individual case. In cases where medical necessity is not established by policy for specific treatment modalities, evidence not previously considered regarding the efficacy of the modality that is presented shall be given consideration to determine if the policy represents current standards of care.

SCOPE: Providence Health Plan, Providence Health Assurance, Providence Plan Partners, and Ayin Health Solutions as applicable (referred to individually as “Company” and collectively as “Companies”).
**PLAN PRODUCT AND BENEFIT APPLICATION**

☐ Commercial  ☒ Medicaid/OHP*  ☐ Medicare**

*Medicaid/OHP Members

Oregon: Services requested for Oregon Health Plan (OHP) members follow the OHP Prioritized List and Oregon Administrative Rules (OARs) as the primary resource for coverage determinations. Medical policy criteria below may be applied when there are no criteria available in the OARs and the OHP Prioritized List.

**Medicare Members

This Company policy may be applied to Medicare Plan members only when directed by a separate Medicare policy. Note that investigational services are considered “not medically necessary” for Medicare members.

**COVERAGE CRITERIA**

Note: Reimbursement for a Speech Generating Device (SGD) must be consistent with what is reasonable and medically necessary to serve the intended purpose. Therefore, payment for an SGD does not include “deluxe” or additional features beyond the function of speech generation and are the least costly alternative.

Devices

I. A non-tablet speech generating device (SGD) (E2500 - E2511) may be considered **medically necessary** when all of the following criteria (A.-G.) are met (for the definition of SGD, see Policy Guidelines):

A. Prior to the delivery of the SGD, the member has had a formal evaluation of their cognitive and communication abilities by a speech-language pathologist (SLP). The formal, written evaluation must include, at a minimum, the following elements (1.-7.):
   1. Current communication impairment, including the type, severity, language skills, cognitive ability, and anticipated course of the impairment;
   2. An assessment of whether the individual's daily communication needs could be met using other natural modes of communication;
   3. A description of the functional communication goals expected to be achieved and treatment options;
   4. Rationale for selection of a specific device and any accessories;
   5. Demonstration that the patient possesses a treatment plan that includes a training schedule for the selected device;
   6. The cognitive and physical abilities to effectively use the selected device and any accessories to communicate;
7. For a subsequent upgrade to a previously issued SGD, information regarding
the functional benefit to the patient of the upgrade compared to the initially
provided SGD; and
B. The member’s medical condition is one resulting in a severe expressive speech
impairment; and
C. The member’s speaking needs cannot be met using natural communication
methods; and
D. Other forms of treatment have been considered and ruled out; and
E. The member’s speech impairment will benefit from the device ordered; and
F. A copy of the SLP’s written evaluation and recommendation have been forwarded to
the member’s treating physician prior to ordering the device; and
G. The SLP performing the member evaluation may not be an employee of or have a
financial relationship with the supplier of the SGD.

II. If criterion I is not met, the SGD is considered **not medically necessary**.

III. A tablet (e.g., iPad) with associated software (E2510 and E2511) when limited to use by
a member with a severe speech impairment and is primarily used for the purpose of
generating speech, may be considered **medically necessary** when general SGD coverage
criteria (see Criterion I) are met.

IV. More than one SGD (tablet or non-tablet) is considered **not medically necessary**.

V. The capability to download updates to the covered features of the device from the
manufacturer or supplier of the device may be considered **medically necessary**.

**Accessories**

VI. Accessories for SGDs (E2599) may be considered **medically necessary** when all of the
following are met (A-C):

   A. General coverage criteria for an SGD are met (See Criterion I); and
   B. The medical need for each accessory is documented within the formal
evaluation by the SLP; and
   C. The accessory is not listed as a non-covered item below.

VII. Accessories are considered **not medically necessary** when coverage criteria for an SGD
are not met.

VIII. Alternative input devices (e.g., specialty keyboards, joysticks, trackballs, trackpads,
buddy buttons, jelly beans, beamers, roller balls, round pads, pal pads) may be
considered **medically necessary** when a member is unable to use standard input
devices.

IX. Alternative input devices for members who are able to use standard input devices is
considered **not medically necessary**.
X. Head control mouse, eye (or ocular) tracking and gaze interaction accessories for speech generating devices may be considered **medically necessary** when furnished to individuals with a demonstrated medical need for such accessories.

XI. The following accessories or features of an SGD are considered **not medically necessary**:

A. A carrying case (with or without a shoulder strap or carrying handle; HCPCS E2599);
B. Features not used to meet functional speaking needs. (See **Policy Guidelines** for examples);
C. Features used for video communications or conferencing;
D. Any computing hardware or software not necessary to allow for generation of speech, email, text or phone messages (e.g., hardware or software used to create documents and spreadsheets or to play games or music).

XII. Internet or phone services or any modification to a member’s home to allow use of the speech generating device is considered **not medically necessary**.

**Replacements**

XIII. A replacement speech generating device may be considered **medically necessary** for when any of the following are met (A.-D.):

A. There is a change in the physical condition of the patient; **or**
B. When replacement is needed due to irreparable *damage* (e.g., fire, flood, etc.) or if the existing equipment is lost or stolen; **or**
C. When replacement is needed due to irreparable *wear* and when the reasonable *useful lifetime* (RUL) of the equipment has been reached (at least 5 years); **or**
D. The cost to repair existing equipment exceeds the purchase price of a replacement.

XIV. Replacement or upgrade of speech generating software loaded onto a covered speech generating device may be considered **medically necessary** when any of the following criteria are met (A.-C.):

A. The replacement software is necessary due to a change in the patient’s condition; **or**
B. The software has been lost, stolen, irreparably damaged; **or**
C. The software has been in continuous use for the reasonable useful lifetime of 5 years.

XV. Replacement or upgrade of speech generating device or device software is considered not medically necessary when criteria XIII or XIV above are not met.

Link to **Evidence Summary**
POLICY CROSS REFERENCES

None

The full Company portfolio of current Medical Policies is available online and can be accessed here.

POLICY GUIDELINES

This policy may be primarily based on the following Center for Medicare and Medicaid Services (CMS) guidance resources:

- Centers for Medicare and Medicaid Services LCD L33739. LCD Title: Speech Generating Devices
- Centers for Medicare and Medicaid Services LCA A52469. LCA Title: Speech Generating Devices
- National Coverage Decision NCD 50.1. Manual Section Title: Speech Generating Devices
- Medicare Claims Processing Manual, Chapter 20 - Durable Medical Equipment, Prosthetics, Orthotics, and Supplies (DMEPOS). Section 90 - Payment for Additional Expenses for Deluxe Features (Rev. 1, 10-01-03) B3-5107, PM AB-02-114.

DOCUMENTATION REQUIREMENTS

While most of the codes in this medical policy are not subject to routine review for medical necessity, the following documentation must be in the medical record to support medically necessity has been established:

- For speech generating devices (SGDs), the following must be available:
  - Description of the item or service
  - Manufacturer name
  - Product name and number
- For accessories, medical necessity for each item must be clearly documented in the formal evaluation by the speech-language pathologist (SLP).
- For alternative input devices, there must be information in the SLP evaluation about why standard input access devices are unable to be used.
- For a multicomponent mounting system, list each component's manufacturer and product name and number.

General

This policy may be primarily based on the following Center for Medicare and Medicaid Services (CMS) guidances:

- Centers for Medicare and Medicaid Services LCD L33739. LCD Title: Speech Generating Devices
- Centers for Medicare and Medicaid Services LCA A52469. LCA Title: Speech Generating Devices
- National Coverage Decision NCD 50.1. Manual Section Title: Speech Generating Devices
- Medicare Claims Processing Manual, Chapter 20 - Durable Medical Equipment, Prosthetics, Orthotics, and Supplies (DMEPOS). Section 90 - Payment for Additional Expenses for Deluxe Features (Rev. 1, 10-01-03) B3-5107, PM AB-02-114.
Speech generation is defined as audible generation of words or phrases and in addition, may include:

1. Communication via written text (i.e., email or text (SMS) messaging); or,
2. Communication via phone messaging.

Speech generating devices are defined by Medicare as durable medical equipment that provides an individual who has severe speech impairment with the ability to meet his or her functional, speaking needs. Speech generating devices are speech aids consisting of devices or software that generate speech (as defined above) and are used solely by the individual who has severe speech impairment. The speech is generated using one of the following methods:

- Digitized audible/verbal speech output, using prerecorded messages;
- Synthesized audible/verbal speech output which requires message formulation by spelling and device access by physical contact with the device-direct selection techniques;
- Synthesized audible/verbal speech output which permits multiple methods of message formulation and multiple methods of device access; or
- Software that allows a computer or other electronic device to generate speech.

- Other covered features of the device include the capability to generate email, text, or phone messages to allow the patient to “speak” or communicate remotely, as well as the capability to download updates to the covered features of the device from the manufacturer or supplier of the device.
- If a speech generating device is limited to use by a patient with a severe speech impairment and is primarily used for the purpose of generating speech, it is not necessary for the device to be dedicated only to audible/verbal speech output to be considered DME. Computers are generally not considered DME because they are useful in the absence of an illness or injury.

The Company medical policy Durable Medical Equipment, Prosthetics, Orthotics, and Supplies (DMEPOS) (All Lines of Business Except Medicare) (MP142) can be reviewed for additional information regarding Medicare guidelines for durable medical equipment, including Medicare’s definition of DME and coverage background for replacement devices.

Desktops, laptops and smartphones are not considered DME, even though they may serve a medical purpose. Medicare will reimburse for speech generating software only (HCPCS code E2511) when installed on one of these general computing devices. While Medicare does not recognize tablets (e.g., iPads) as DME, the Company may choose to allow a tablet as a speech generating device (SGD) when all relevant criteria are met.

Examples of Non-Covered Specific Features

Examples of specific features of a SPG that are not used by the member include any computing hardware or software not necessary to allow for generation of audible/verbal speech, email, text or phone messages, such as hardware or software used to create documents and spreadsheets or play games or music, and any other function a computer can perform that is not directly related to meeting the functional speaking communication needs of the patient, including video communications or
conferencing. These features of a speech generating device do not fall within the scope of § 1861(n) of the Social Security Act and the cost of these features are the responsibility of the patient.

Payment for Additional Expenses for Deluxe Features

The payment amount for a given service or item, whether rented or purchased, must be consistent with what is reasonable and medically necessary to serve the intended purpose. Additional expenses for "deluxe" features, or items that are rented or purchased for aesthetic reasons or added convenience, do not meet the reasonableness test. Thus, where a service or item is medically necessary and covered under the Medicare program, and the patient wishes to obtain such deluxe features, the payment is based upon the payment amount for the kind of service or item normally used to meet the intended purpose (i.e., the standard item.) Usually this is the least costly item.

REGULATORY STATUS

U.S. FOOD AND DRUG ADMINISTRATION (FDA)

Approval or clearance by the Food and Drug Administration (FDA) does not in itself establish medical necessity or serve as a basis for coverage. Therefore, this section is provided for informational purposes only.

BILLING GUIDELINES AND CODING

Codes E2500 - E2511 perform the same essential function - speech generation. Therefore, claims for more than one SGD, regardless of type (tablet or non-tablet) is considered not reasonable and necessary.

Codes E2500, E2502, 22504, E2506, E2508, E2510-E2512 are limited to one every five years.

See the local coverage article (LCA) A52469 for the most up-to-date information regarding coding guidelines for devices, accessories and components.

Different Types of SGDs

Non-tablet SGDs: Codes E2500, E2502-E2506, E2508 and E2510 include all applicable speech generating software programs (whether they are on the device when shipped by the manufacturer or added by the supplier prior to delivery), batteries, battery chargers and AC adapters. These items may not be billed separately. There is also no separate payment if a nonintegrated keyboard is provided with an SGD.

To be coded as E2510, a desktop, smartphone or laptop computer must only be capable of speech generation, as defined above. A desktop, smartphone or laptop computers with additional non-covered features (see Non-Medical Necessity Coverage and Payment Rules above) included at the time of initial issue must be coded A9270. For tablets used as SGDs, see below.
**Tablet SGDs (e.g. iPads)**

While Medicare doesn’t recognize tablets as DME for use as SGD, the Company may allow tablets used as SGDs (when above criteria are met), which can be billed with HCPCS code E2510. If the tablet SGD does not include all applicable speech generating software programs, then E2511 may be billed for the software. Batteries, battery chargers and AC adapters may not be billed separately. There is also no separate payment if a nonintegrated keyboard is provided with a tablet SGD.

**Software Programs**

Code E2511 **must not** be used to code for software programs installed at the time of the initial provision of an SGD accessory or alternative access device. Software for the accessory or alternative access device is included in the reimbursement for the accessory or alternative access device. Claims for code E2511 billed with an accessory or alternative access device will be denied as unbundling.

Code E2511 is also used to code for speech generating software programs that enable a laptop computer, desktop computer, tablet, smartphone or other hand-held general computing device to generate speech. The allowance for code E2511 includes the speech generating software program only. Installation of the program or technical support must not be billed separately. Code E2511 must not be used to code for software programs that are installed at the time of delivery of an SGD (codes E2500, E2502-E2506, E2508 or E2510).

Finally, code E2511 may also be used for upgrade programs for SGDs (codes E2500, E2502-E2506, E2508 or E2510) or when code E2511 is used to bill for software only when installed on a general computing device.

A protective case or cover, any type (E2599) is not separately payable as they are required to make the SGD durable under the definition of DME. Claims for protective cases or covers **when billed with a covered SGD** will be denied as unbundled. However, if a protective case or cover is provided separately, such as when a replacement is required, then it may be reported and reimbursed separately.

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<th>CODES*</th>
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<td><strong>HCPCS</strong></td>
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**E2511** Speech generating software program, for personal computer or personal digital assistant

**E2512** Accessory for speech generating device, mounting system

**E2599** Accessory for speech generating device, not otherwise classified

*Coding Notes:*
- The above code list is provided as a courtesy and may not be all-inclusive. Inclusion or omission of a code from this policy neither implies nor guarantees reimbursement or coverage. Some codes may not require routine review for medical necessity, but they are subject to provider contracts, as well as member benefits, eligibility and potential utilization audit.
- All unlisted codes are reviewed for medical necessity, correct coding, and pricing at the claim level. If an unlisted code is submitted for non-covered services addressed in this policy then it will be denied as not covered. If an unlisted code is submitted for potentially covered services addressed in this policy, to avoid post-service denial, prior authorization is recommended.
- HCPCS/CPT code(s) may be subject to National Correct Coding Initiative (NCCI) procedure-to-procedure (PTP) bundling edits and daily maximum edits known as “medically unlikely edits” (MUEs) published by the Centers for Medicare and Medicaid Services (CMS). This policy does not take precedence over NCCI edits or MUEs. Please refer to the CMS website for coding guidelines and applicable code combinations.

**REFERENCES**


**POLICY REVISION HISTORY**

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<tr>
<td>2/2023</td>
<td>Converted to new policy template.</td>
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| 5/2023  | • Add “Documentation Requirements” and removal of LCA language from Billing Guideline regarding documentation.  
• Add criteria for more than one speech generating device (SGD) (NMN).  
• Criteria changes in the “Accessories” section. |
• Add replacement criteria
• Add language in the “Policy Guidelines” clarifying that tablets may be considered as SGDs.
• Removal/deletion of certain Policy Guideline information to avoid repetition.