**Coverage Summary**

**Speech Generating Devices**

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<th>Policy Number: S-011</th>
<th>Products: UnitedHealthcare Medicare Advantage Plans</th>
<th>Original Approval Date: 02/18/2009</th>
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<tr>
<td>Approved by: UnitedHealthcare Medicare Benefit Interpretation Committee</td>
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**Related Medicare Advantage Policy Guideline:** Speech Generating Devices (NCD 50.1)

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The benefit information in this Coverage Summary is based on existing national coverage policy, however, Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable.

There are instances where this document may direct readers to a UnitedHealthcare Commercial Medical Policy, Medical Benefit Drug Policy, and/or Coverage Determination Guideline (CDG). In the absence of a Medicare National Coverage Determination (NCD), Local Coverage Determination (LCD), or other Medicare coverage guidance, CMS allows a Medicare Advantage Organization (MAO) to create its own coverage determinations, using objective evidence-based rationale relying on authoritative evidence (Medicare IOM Pub. No. 100-16, Ch. 4, §90.5).

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**I. COVERAGE**

**Coverage Statement:** Speech generating devices are covered when the treating physician determines that the patient suffers from severe speech impairment and that the medical condition warrants the use of a device based on the Medicare definitions outlined below.

**DME Face to Face Requirement:** Effective July 1, 2013, Section 6407 of the Affordable Care Act (ACA) established a face-to-face encounter requirement for certain items of DME (including speech generating devices). For DME Face to Face Requirement information, refer to the Coverage Summary of Durable Medical Equipment (DME), Prosthetics, Corrective Appliances/Orthotics (Non-Foot Orthotics) and Medical Supplies Grid.
Guidelines/Notes

1. Speech generating devices are considered to fall within the durable medical equipment benefit category established by § 1861(n) of the Social Security Act. They are covered for patients who suffer from a severe speech impairment and have a medical condition that warrants the use of a device based on the following definitions.

Speech generating devices are defined as durable medical equipment that provides an individual who has a 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 and are used solely by the individual who has a severe speech impairment.

The speech is generated using one of the following methods:

a. Digitized audible/verbal speech output, using pre-recorded messages;

b. Synthesized audible/verbal speech output, which requires message formulation by spelling and device access by physical contact with the device-direct selection techniques;

c. Synthesized audible/verbal speech output, which permits multiple methods of message formulation and multiple methods of device access; or

d. Software that allows a computer or other electronic device to generate audible/verbal speech.

2. 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.

3. The following devices, modifications and services do not meet the definition of a speech generating device and are not covered:

a. Computers and tablets in general are not considered DME because they are useful in the absence of an illness or injury.

b. Internet or phone services or any modification to a patient’s home to allow use of the speech generating device are not covered by Medicare because such services or modifications could be used for non-medical equipment such as standard phones or personal computers.

c. Specific features of a speech generating device that are not used by the individual who has a severe speech impairment to meet his or her functional speaking needs are not covered.

d. 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 is not covered.

See the NCD for Speech Generating Devices (50.1), (Accessed January 30, 2020)

4. A speech generating device (SGD) is covered when all of the following criteria are met. If one
or more of the above coverage criteria is not met, the SGD will be denied as not medically necessary.

a. Prior to the delivery of the SGD, the patient 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) 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's evaluation may not be an employee of or have a financial relationship with the supplier of the SGD.

5. Accessories

a. Accessories are covered if the basic coverage criteria above for the base device are met and reasonable and necessary criteria for each accessory is clearly documented in the formal evaluation by the SLP.

b. Alternative input devices are covered when a member is unable to use standard input devices. Claims for alternative input devices for member’s who are able to use standard input devices will be denied as not reasonable and necessary.

c. Eye tracking and gaze interaction accessories for speech generating devices are covered when furnished to individuals with a demonstrated medical need for such accessories.

For additional coverage and coding information, see the DME MAC LCD for Speech Generation Devices (L33739). Compliance with these policies is required where appropriate. (Accessed January 30, 2020)
II. DEFINITIONS

III. REFERENCES

See above

IV. REVISION HISTORY

02/18/2020  Guideline 3 (Not Covered Devices, Modifications and Services)
- Replaced language indicating “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 a speech-generating device to be dedicated only to speech generation to be considered DME” with “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”