

### Kaiser Foundation Health Plan of Washington

# *Clinical Review Criteria* Speech Generating Devices

Augmented and Alternative Communication Devices or Communicators

**NOTICE:** Kaiser Foundation Health Plan of Washington and Kaiser Foundation Health Plan of Washington Options, Inc. (Kaiser Permanente) provide these Clinical Review Criteria for internal use by their members and health care providers. The Clinical Review Criteria only apply to Kaiser Foundation Health Plan of Washington and Kaiser Foundation Health Plan of Washington Options, Inc. Use of the Clinical Review Criteria or any Kaiser Permanente entity name, logo, trade name, trademark, or service mark for marketing or publicity purposes, including on any website, or in any press release or promotional material, is strictly prohibited.

Kaiser Permanente Clinical Review Criteria are developed to assist in administering plan benefits. These criteria neither offer medical advice nor guarantee coverage. Kaiser Permanente reserves the exclusive right to modify, revoke, suspend or change any or all of these Clinical Review Criteria, at Kaiser Permanente's sole discretion, at any time, with or without notice. **Member contracts differ in health plan benefits. Always consult the patient's Evidence of Coverage or call Kaiser Permanente Member Services at 1-888-901-4636 (TTY 711), Monday through Friday, 8 a.m. to 5 p.m. to determine coverage for a specific medical service.** 

## Criteria

#### **For Medicare Members**

Source	Policy
CMS Coverage Manuals	None
National Coverage Determinations (NCD)	Speech Generating Device (50.1)
Local Coverage Determinations (LCD)	Speech Generating Device (L33739)
Local Coverage Article	None

#### **For Non-Medicare Members**

Kaiser Permanente has elected to use the Augmentative Communication Devices, Electronic (KP-0516) MCG\* for medical necessity determinations. For access to the MCG Clinical Guidelines criteria, please see the MCG Guideline Index through the provider portal under Quick Access.

**MCG\*** are proprietary and cannot be published and/or distributed. However, on an individual member basis, Kaiser Permanente can share a copy of the specific criteria document used to make a utilization management decision. If one of your patients is being reviewed using these criteria, you may request a copy of the criteria by calling the Kaiser Permanente Clinical Review staff at 1-800-289-1363 or access the MCG Guideline Index using the link provided above.

#### If requesting this service, please send the following documentation to support medical necessity:

- Last 6 months of clinical notes from requesting provider and/or specialist (neurology)
- Speech therapy notes

The following information was used in the development of this document and is provided as background only. It is provided for historical purposes and does not necessarily reflect the most current published literature. When significant new articles are published that impact treatment option, Kaiser Permanente will review as needed. This information is not to be used as coverage criteria. Please only refer to the criteria listed above for coverage determinations.

## Background

Augmentative and alternative communication (AAC) is an area of clinical practice that attempts to temporarily or to permanently compensate for the impairment and disability patterns of children with severe oral and written expressive communication disorders. Interventions that use AAC should incorporate the individual's full communication abilities e.g. any existing speech or vocalization, gestures, manual signs, communication boards, and speech output communication devices. Abilities may change over time and the AAC may need to be modified as a child grows and develops.

AAC has four components: symbols, aids, techniques, and strategies. Aids are the physical objects or devices used to transmit or receive messages. These include books, communication boards, charts, mechanical or electronic devices, and computers. The AAC devices have variable capabilities, durability, and cost. The delivery of AAC services to children with severe spoken language disorders requires the collaboration and competence of

<u>Criteria | Codes | Revision History</u> families, professionals, and paraprofessionals. Effective, co-coordinated multidisciplinary and an integrated service is crucial in achieving optimal outcome for the children.

The role an AAC system plays in a particular child's life varies with the type and severity of the language disorder. Children with congenital language disorders who may benefit from AAC include those with cerebral palsy, dual sensory impairments, developmental apraxia, oro-motor dyspraxia, language learning disabilities, mental retardation, autism, and pervasive developmental disorders. Acquired language disorders include: traumatic brain injury, aphasia, spinal cord injuries, and other physical disabilities. Not all these indications are covered by health insurance companies.

## Medical Technology Assessment Committee (MTAC)

# Augmentative Communication Devices

02/13/2002: MTAC REVIEW

**Evidence Conclusion:** The study reviewed had several limitations; it had a small sample size, lacked a control group, used only subjective measures, and was subject to selection and observation biases. In conclusion the literature available does not provide enough evidence to determine the effect of the augmentative communication devices on the communication skills of children with speech impairments.

<u>Articles</u>: The search yielded 43 articles. Most were reviews, tutorials, notes, and discussions. The search did not reveal any randomized controlled trials, or meta-analyses, only four case reports and two studies that only measured young patients' or parents' satisfactions and /or utilization of the communication systems. The study with the larger sample size was selected for critical appraisal. *An evidence table was created for the following study:* Ko MLB, et al. Outcome of recommendations for augmentative communication in children. *Child Care, Health and Development* 1998; 24(3): 195-205. See Evidence Table.

The use of augmentative communication devices on the communication skills of children with speech impairments not voted using the *Kaiser Permanente Medical Technology Assessment Criteria*.

# **Applicable Codes**

CPT<sup>®</sup> or **HCPC** Codes E1902 Communication board, nonelectronic augmentative or alternative communication device E2500 Speech generating device, digitized speech, using prerecorded messages, less than or equal to eight minutes recording time E2502 Speech generating device, digitized speech, using prerecorded messages, greater than eight minutes but less than or equal to 20 minutes recording time E2504 Speech generating device, digitized speech, using prerecorded messages, greater than 20 minutes but less than or equal to 40 minutes recording time E2506 Speech generating device, digitized speech, using prerecorded messages, greater than 40 minutes recording time E2508 Speech generating device, synthesized speech, requiring message formulation by spelling and access by physical contact with the device E2510 Speech generating device, synthesized speech, permitting multiple methods of message formulation and multiple methods of device access 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

Considered Medically Necessary when criteria in the applicable policy statements listed above are met:

\*Note: Codes may not be all-inclusive. Deleted codes and codes not in effect at the time of service may not be covered.

\*\*To verify authorization requirements for a specific code by plan type, please use the Pre-authorization Code Check.

CPT codes, descriptions and materials are copyrighted by the American Medical Association (AMA). HCPCS codes, descriptions and materials are copyrighted by Centers for Medicare Services (CMS).

Creation Date	Review Date	Date Last Revised
06/18/2001	03/02/2010 <sup>MDCRPC</sup> , 02/10/2011 <sup>MDCRPC</sup> , 12/06/2011 <sup>MDCRPC</sup> , 10/02/2012 <sup>MDCRPC</sup> , 08/06/2013 <sup>MPC</sup> , 11/05/2013 <sup>MPC</sup> , 09/02/2014 <sup>MPC</sup> , 07/07/2015 <sup>MPC</sup> , 05/03/2016 <sup>MPC</sup> , 03/07/2017 <sup>MPC</sup> , 01/09/2018 <sup>MPC</sup> , 12/04/2018 <sup>MPC</sup> , 12/03/2019 <sup>MPC</sup> , 12/03/2019 <sup>MPC</sup> , 12/01/2020 <sup>MPC</sup> , 12/07/2021 <sup>MPC</sup> , 12/06/2022 <sup>MPC</sup> , 12/09/2023 <sup>MPC</sup> , 02/13/2024 <sup>MPC</sup>	08/31/2015

MDCRPC Medical Director Clinical Review and Policy Committee

MPC Medical Policy Committee

Revision	Description
History	
08/31/2015	Added Update to Pub. 100-03 NCD Manual
02/26/2024	Removed CPT 92609 from criteria page as this code is for the service and not the device.