<u>5160-10-01</u> <u>Durable medical equipment, prostheses, orthoses, and supplies</u> (DMEPOS): general provisions.

- (A) This rule sets forth general coverage and payment policies for durable medical equipment (DME), prostheses, orthotic devices, medical/surgical supplies, and supplier services.
 - (1) Additional conditions specific to a particular DMEPOS item or service may be set forth in other rules in this chapter of the Administrative Code.
 - (2) Policies set forth in other rules in this chapter supersede any provisions in this rule with which they conflict.
- (B) <u>Definitions that apply to rules in this chapter of the Administrative Code.</u>
 - (1) "Certificate of medical necessity (CMN)" is a written statement by a practitioner attesting that a particular item or service is medically necessary for an individual.
 - (a) If no other form or format is specified, the CMN form specified in the relevant rule in this chapter of the Administrative Code is the default that is to be used.
 - (b) A CMN is not invalidated by a change in an individual's status from one medicaid eligibility category to another (e.g., from fee-for-service medicaid to medicaid managed care).
 - (c) Renewal of lifetime certification is not necessary.
 - (d) An illegible CMN will not be accepted.
 - (2) "Coverage" is the principle that medicaid payment is routinely made for a particular medically necessary item or service.
 - (a) The department maintains several payment schedules of covered items and services, which are posted on the department's web site. These schedules are neither all-inclusive nor exclusive. Neither the appearance of an item or service on a payment schedule nor its absence determines, in and of itself, coverage or non-coverage.
 - (b) For most covered items and services, medical necessity has already been established and is simply confirmed on a case-by-case basis through the completion of a CMN (when applicable). For certain items and services, medical necessity and coverage are established through a prior authorization (PA) process.

(c) Only the department can determine coverage. Providers cannot decide on their own that an item or service is not covered or would not be covered with PA. Providers should submit a PA request to obtain an official decision.

- (3) "Department" is the Ohio department of medicaid or, when applicable, its designee. The address of the department's web site is http://medicaid.ohio.gov.
- (4) "DMEPOS item" is a collective term for a covered durable medical equipment (DME) item, prosthetic device, orthotic device, or medical supply item furnished by an eligible provider to an eligible recipient.
- (5) "DMEPOS provider" is a collective term for the following eligible providers:
 - (a) A basic DME supplier, which holds licensure or certification in accordance with Chapter 4752. of the Revised Code and furnishes items other than life-sustaining or technologically sophisticated equipment;
 - (b) A specialized DME supplier, which holds licensure or certification in accordance with Chapter 4752. of the Revised Code and furnishes lifesustaining or technologically sophisticated equipment; and
 - (c) An orthotics and prosthetics (O&P) supplier, which holds licensure or certification in accordance with section 4779.02 of the Revised Code and furnishes orthotic and prosthetic devices.
- (6) "DMEPOS service" is a covered service, such as labor for repair or replacement, that is furnished by an eligible provider and is related directly to a DMEPOS item.
- (7) "Frequency limit" is the average expected useful life of a DMEPOS item. A frequency limit is not an absolute restriction but a general guideline and therefore may be exceeded with medical justification. For certain DMEPOS items that can be dispensed in multiple units (such as fasteners or items with left/right orientation), a frequency limit applies to each unit that is requested.
- (8) "Long-term care facility (LTCF)" is a collective term for a nursing facility (NF), a skilled nursing facility (SNF), and an intermediate care facility for individuals with intellectual disabilities (ICFIID).
- (9) "Need verification" is a process by which the department determines whether to make payment for a DMEPOS item or service that exceeds the established cost threshold or frequency guideline. Because need verification is applied only to items or services for which medical necessity has been established or presumed.

no extensive or in-depth clinical assessment is necessary (as it is with prior authorization). One purpose of need verification is to enable the department to consider whether the purchase of a new piece of equipment might be more costeffective than continued repair.

- (10) "Private residence" is a recipient's place of residence other than a long-term care facility (LTCF).
- (11) "Provider cost" is the amount paid for an item by a DMEPOS provider to a supplier or manufacturer, exclusive of discounts, rebates, and situation-specific adjustments. Documentation of provider cost is subject to approval by the department; a figure that has been entered, superimposed, modified, obscured, or obliterated by the provider will not be accepted. Suitable documents for substantiating provider cost include but are not limited to the following examples:
 - (a) An invoice submitted by the supplier or manufacturer to the provider;
 - (b) A bona fide quotation (quote) submitted by the supplier or manufacturer to the provider; or
 - (c) A standard supplier or manufacturer price list that can be independently verified by the department.

(C) Coverage.

- (1) The provision of or payment for a medically necessary DME item or medical supply for a resident of a LTCF is the responsibility of the LTCF, in accordance with Chapter 5160-3 of the Administrative Code. In turn, the LTCF receives medicaid per diem payment on the basis of its cost report. Therefore, claims submitted for such items or supplies furnished to LTCF residents will be denied. Any exceptions are set forth in other rules in this chapter of the Administrative Code.
- (2) Separate payment may be made for a prosthesis or orthotic device supplied to a resident of a LTCF.
- (3) A medically necessary DMEPOS item can be dispensed only by prescription. The following provisions apply:
 - (a) Eligible medicaid providers of the following types having prescriptive authority under Ohio law may certify the medical necessity of a DMEPOS item:

- (i) A physician;
- (ii) A podiatrist;
- (iii) An advanced practice registered nurse with a relevant specialty; or
- (iv) A physician assistant.
- (b) Before writing a prescription for certain DMEPOS items, a practitioner conducts a face-to-face encounter with the medicaid recipient and documents it in the recipient's medical record. Items for which an encounter is a prerequisite are listed on the website of the centers for medicare and medicaid services (CMS) at http://www.cms.gov.
- (c) A prescription cannot be written before an encounter.
- (d) Unless a different length of time is specified, a prescription for a particular DMEPOS item is valid for sixty days, regardless of whether it is based on a face-to-face encounter.
- (e) A single encounter can serve for twelve months as the basis for a single prescription or for more than one prescription addressing the same medical condition for which a DMEPOS item is being prescribed.
- (f) The medical practitioner acting as prescriber needs to be actively involved in managing the recipient's healthcare. The department may disallow a prescription written by a practitioner who has no professional relationship with the recipient.
- (g) There needs to be a direct relationship between the prescribed DMEPOS item and a medical condition of the recipient that the practitioner evaluates, assesses, or actively treats during the encounter.
- (h) Each prescription should specify a quantity (e.g., "TID," "thirty per month").

 An unstated quantity is assumed to be one unit.
- (4) A prescription serves as an order to dispense, and a DMEPOS provider may dispense an item on receipt of a valid prescription. Payment, however, depends on the establishment of medical necessity, which is separate from the prescription process. For most DMEPOS items, a provider has the applicable CMN completed and signed by a prescribing practitioner after the prescription is written and before it expires. If no CMN is specified for an item, then the prescription itself establishes medical necessity.

(5) Certain DMEPOS items are subject to prior authorization (PA). A list of such items is posted on the department's web site.

- (a) The following DMEPOS items are always subject to PA:
 - (i) A custom or a specialized DMEPOS item;
 - (ii) A "not otherwise specified," "miscellaneous," or "unlisted" item or service; and
 - (iii) Used DME.
- (b) When PA is given, it may specify a quantity, manufacturer, model, part number, or other information identifying a particular item. When such identifying information is present, a provider may supply and subsequently submit claims for the specified items only. No changes or substitutions are allowed without explicit authorization by the department.
- (c) The department, on the basis of clinical indications, may grant PA for an item other than one that has been requested.
- (d) For items subject to PA, the provider submits the following documentation within sixty days after the date on which the CMN was signed (or, if there is no applicable CMN, within the validity period of the prescription):
 - (i) The fully completed and signed CMN (or, if there is no applicable CMN, the prescription):
 - (ii) Related information, such as a full description of any similar item currently in possession of the recipient or an explanation of a change in the recipient's condition that warrants a change in equipment;
 - (iii) For a "not otherwise specified," "miscellaneous," or "unlisted" item, a complete description of the item (including, as applicable, the manufacturer, model or style, and size), a list of all bundled components, and an itemization of all charges; and
 - (iv) Any other information requested by the department, as detailed in this chapter of the Administrative Code.
- (e) A request for PA of a preparatory prosthesis includes the reason for the amputation, the date of the amputation, and an explanation of the benefit

- to be derived from having the recipient use a preparatory prosthesis before a definitive prosthesis is designed.
- (f) A claim for an item or service that exceeds the specified maximum quantity or frequency but is not otherwise subject to PA may be subject to need verification before payment will be considered.
- (g) A request for PA or need verification may be denied in cases involving malicious damage, neglect, culpable irresponsibility, or wrongful disposition.
- (6) For items not subject to PA, the provider keeps on file the prescription and, if applicable, the fully completed and signed CMN. The provider cannot submit a claim until these documents have been obtained.
- (7) For an item that is shipped directly to a recipient, the shipping date is the dispensing date.
- (8) For an item that needs multiple fittings and special construction, the first date of service is the dispensing date.
- (9) If a recipient dies after measurements for a prescribed custom item have been taken but before the item has been dispensed, then payment for the item may be made under the following conditions:
 - (a) The code set description for the item indicates that it is designed or intended for a specific individual;
 - (b) The item is substantially complete and cannot be modified for use by another individual;
 - (c) No information available to the provider indicated that the death of the recipient was imminent;
 - (d) The provider can document the date of measurement; and
 - (e) On the claim, the provider reports the date of measurement as the date of service.
- (10) Any request for a DMEPOS item or service needs to originate with an individual recipient, the recipient's authorized representative, or a medical practitioner acting as the prescriber with the recipient's full knowledge and consent.

(11) A request that is determined by the department to have resulted from a mass screening or examination will be denied.

- (12) When instruction in the safe and appropriate use of a particular DMEPOS item is indicated, it is the responsibility of the provider to ensure that the recipient or someone authorized to assist the recipient has received such instruction.
- (13) Payment for repair of a DME item, prosthetic device, or orthotic device or for purchase of a related medical supply item or service can be made only if the medical necessity of the DME item, prosthetic device, or orthotic device itself has been established. The medical necessity of an item purchased by the department is established during the purchasing process. For an item not purchased by the department, medical necessity may be documented on an appropriate medicaid certificate of medical necessity, on a prescription that addresses all specified criteria, or on any other form that is acceptable to the department. No additional documentation of medical necessity is necessary for subsequent repairs made to an item. The determination that an item not purchased by the department is medically necessary does not indicate that the item would be authorized for purchase.
- (14) Payment may be made for covered repair, maintenance, parts, accessories, or supplies for a DME item that is owned by an individual but has not been purchased by the department. Payment for the initial service or delivery is subject to PA; payment for subsequent service or deliveries is not subject to PA.
- (15) Unless otherwise specified elsewhere in this chapter of the Administrative Code, for each claim submitted for payment, a provider keeps the following supporting documents on file:
 - (a) A completed and signed CMN, if needed;
 - (b) If no CMN is needed, a legible prescription that specifies a diagnosis;
 - (c) <u>Information such as practitioner orders or chart notes, used to establish the medical necessity of the DMEPOS item;</u>
 - (d) Any record indicating a change in an individual's needs or plan of care;
 - (e) Proof of delivery;
 - (f) Confirmation that the recipient or the recipient's authorized representative has been instructed in the safe use of the DMEPOS item, if applicable;
 - (g) A copy of the manufacturer's or dealer's warranty, if applicable; and

(h) A record of any repair or service that has been performed on equipment not paid for by medicaid, if applicable.

- (16) The default CMN form for general DME items and supplies is the ODM 01913, "Certificate of Medical Necessity / Request for Need Verification: General Medical Supplies and Equipment" (rev. 7/2021).
- (17) Proof is needed to show that a DMEPOS item has been delivered to the intended recipient.
 - (a) Providers, their employees, and anyone else having a financial interest in the delivery of DMEPOS items are not permitted to accept delivery of an item on behalf of a medicaid recipient.
 - (b) If a provider delivers directly to a recipient, then acceptable proof of delivery includes the signature of the recipient or the recipient's authorized representative. For a DMEPOS item delivered to a resident of a LTCF, the LTCF is responsible for furnishing proof of delivery.
 - (c) If a provider uses a third-party shipper, then acceptable proof of delivery includes the shipper's tracking slip or a returned postage-paid delivery invoice.
 - (d) If a signature obtained physically at the time of delivery is not legible, then the provider or shipper records the name of the person accepting delivery and the relationship of the person to the recipient. If the provider or shipper records such information for a particular person and maintains it in a readily accessible format, then on subsequent deliveries only the signature is needed.
- (18) If more than one DMEPOS item or service will meet a recipient's needs, then the maximum payment amount cannot exceed the least costly alternative, in accordance with rule 5160-1-01 of the Administrative Code.
- (19) No separate payment will be made under this chapter of the Administrative Code for the following items or services:
 - (a) Items presumed to be nonmedical in nature and for which no medical necessity can therefore be demonstrated, including but not limited to the following examples:
 - (i) Environmental control devices;

(ii) Items that have no medical benefit but are intended solely for the comfort or convenience of the user;

- (iii) Physical fitness equipment;
- (iv) Precautionary items (e.g., emergency alert systems);
- (v) Training equipment (e.g., speech-teaching machines);
- (vi) Communication aids, except as specified elsewhere in this chapter of the Administrative Code;
- (vii) Educational aids; and
- (viii) Hygiene equipment (e.g., bidets);
- (b) Routine over-the-counter treatment supplies (e.g., adhesive bandages, antiseptic solutions, antibiotic ointments) and personal hygiene items (e.g., soap, diapers for children younger than three years of age);
- (c) Medical supplies or DME items that are used during a visit with a medical practitioner (i.e., that are incidental to a professional service) in the practitioner's office, in a clinic, or in the recipient's private residence;
- (d) Items or services that are covered under manufacturer or dealer warranty;
- (e) Items or services for which full remuneration is made through other payment mechanisms;
- (f) Costs of delivery (including postage), setup and assembly, pickup, and routine cleaning and maintenance associated with a covered DME item;
- (g) <u>Labor, measuring, casting, fitting, travel by the supplier, and shipping or mailing associated with a covered orthotic device or prosthesis;</u>
- (h) Maintenance and repair of equipment during a rental period;
- (i) Supporting wires, power supplies, cables, or attachment kits;
- (j) Related supplies and accessories that are furnished either during a rental period or with the dispensing or delivery of a purchased equipment item and for which no payment amount exists for separate purchase or rental;
- (k) A service call in addition to materials and labor;

(1) Repairs, adjustments, or modifications that are made within ninety days after delivery or during the total rental period, unless necessitated by major changes in the recipient's condition;

- (m) <u>Instruction of the recipient or the recipient's authorized representative in</u> the safe use of an item; and
- (n) Education, training, instruction, counseling, or monitoring conducted in support of an individual's ordered treatment plan.
- (20) Payment is not available for DMEPOS items that duplicate or conflict with another item currently in the recipient's possession, regardless of payment or supply source. Providers are responsible for ascertaining whether duplication or conflict exists.
- (21) Certain DMEPOS items may be dispensed on a recurring basis. A provider is to confirm a recipient's current need before the next delivery. If DMEPOS items are routinely delivered without necessary confirmation of need, then any payment for excess quantities is subject to recovery.
- (22) No prescription for disposable items dispensed on a recurring basis (e.g., incontinence garments, wound dressings) can be renewed earlier than ninety days before the expiration of the current prescription.
- (23) Most covered DME items are purchased and become the property of the recipient. Some covered DME items that need ongoing servicing are rented exclusively. Some covered DME items may be rented on a short-term basis, purchased, or rented and then purchased.
 - (a) The short-term rental of a covered DME item other than a wheelchair is subject to PA, which may be given if rental is determined to be more costeffective than purchase.
 - (b) Unless a different length of time is specified elsewhere in this chapter of the Administrative Code, the initial rental period does not exceed six months.
 - (c) PA may be given for additional rental periods.
 - (d) Regardless of its authorized length, a rental period ends when the rented item is no longer medically necessary.
 - (e) A monthly rental payment secures the rented item for the entire calendar month.

(f) During a rental period and for ninety days afterward, all rental amounts paid apply toward purchase.

- (g) The department reserves the right to determine whether an item will be rented or purchased.
- (h) The provider is to notify the recipient when an item in effect has been purchased through rental.
- (24) Medical supply items such as gauze pads and wound fillers/packing are dispensed in bulk. No payment amount per unit has been established for such items; instead, an overall payment limit per period is specified. The charge submitted by the provider cannot exceed one hundred forty-seven per cent of the provider cost for the quantity of the item.
- (25) The purchase of torsion cables may be authorized only for the treatment of children with neuromuscular diseases and related conditions. Requests for torsion cables to treat positional deformities will be denied because of anticipated resolution that occurs with maturation.
- (26) No provider can submit a claim for a DMEPOS item or service before the item or service has been supplied.

(D) Claim payment.

- (1) The payment amount specified in another rule in this chapter of the Administrative Code supersedes any payment amount established by provisions in this rule.
- (2) For a covered DMEPOS item or service represented by a new or newly adopted healthcare common procedure coding system (HCPCS) procedure code, the initial maximum payment amount may be established in accordance with rule 5160-1-60 of the Administrative Code. New or newly adopted HCPCS codes are published in a separate table on the department's web site and remain there until the appropriate DMEPOS payment schedules can be updated.
- (3) For any covered DMEPOS item or service not represented by a new or newly adopted HCPCS procedure code, the payment amount is the lesser of the submitted charge (which is to reflect any discounts or rebates available to the provider at the time of claim submission but need not reflect subsequent discounts or rebates) or the first applicable medicaid maximum from the following ordered list:
 - (a) The amount listed in the appendix to this rule;

(b) For a "by report" DMEPOS item or service, an amount determined on a case-by-case basis;

- (c) For a supply item for which payment is determined by PA, one hundred forty-seven per cent of the provider cost (minus discounts or rebates);
- (d) For a non-supply DMEPOS item or service for which payment is determined by PA, an amount determined on a case-by-case basis;
- (e) For a bulk item having an overall payment limit per period, the submitted charge;
- (f) For the authorized purchase of a DMEPOS item in used condition, eighty per cent of the payment amount for the item in new condition;
- (g) For monthly payment for a "rental/purchase" DME item, ten per cent of the medicaid maximum specified for purchase; or
- (h) For a professional service for which separate payment is made (such as an evaluation), the applicable amount listed in appendix DD to rule 5160-1-60 of the Administrative Code.
- (4) In accordance with the principle stated in rule 5160-1-60 of the Administrative Code concerning correct coding, a "not otherwise specified," "miscellaneous," or "unlisted" procedure code of the appropriate DMEPOS type may be reported on a claim only if no other code listed on a payment schedule indicates coverage of the item or service. The department may deny a claim that omits necessary information or that includes a "not otherwise specified," "miscellaneous," or "unlisted" procedure code when an appropriate procedure-specific code is available.

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