

section, the last screening barium enema was performed.

(4) In the case of an individual who is not at high risk for colorectal cancer as described in paragraph (a)(3) of this section but who has had a screening flexible sigmoidoscopy performed, payment may be made for a screening colonoscopy only after at least 47 months have passed following the month in which the last screening flexible sigmoidoscopy was performed.

(h) *Conditions for coverage of screening barium enemas.* Medicare Part B pays for a screening barium enema if it is ordered in writing by the beneficiary's attending physician.

(i) *Limitations on coverage of screening barium enemas.* (1) In the case of an individual age 50 or over who is not at high risk of colorectal cancer, payment may be made for a screening barium enema examination performed after at least 47 months have passed following the month in which the last screening barium enema or screening flexible sigmoidoscopy was performed.

(2) In the case of an individual who is at high risk for colorectal cancer, payment may be made for a screening barium enema examination performed after at least 23 months have passed following the month in which the last screening barium enema or the last screening colonoscopy was performed.

[62 FR 59100, Oct. 31, 1997, as amended at 66 FR 55329, Nov. 1, 2001; 67 FR 80040, Dec. 31, 2002]

**§ 410.38 Durable medical equipment: Scope and conditions.**

(a) Medicare Part B pays for the rental or purchase of durable medical equipment, including iron lungs, oxygen tents, hospital beds, and wheelchairs, if the equipment is used in the patient's home or in an institution that is used as a home.

(b) An institution that is used as a home may not be a hospital or a CAH or a SNF as defined in sections 1861(e)(1), 1861(mm)(1) and 1819(a)(1) of the Act, respectively.

(c) Wheelchairs may include a power-operated vehicle that may be appropriately used as a wheelchair, but only if the vehicle—

(1) Is determined to be necessary on the basis of the individual's medical and physical condition;

(2) Meets any safety requirements specified by CMS; and

(3) Except as provided in paragraph (c)(2) of this section, is ordered in writing by a specialist in physical medicine, orthopedic surgery, neurology, or rheumatology, the written order is furnished to the supplier before the delivery of the vehicle to the beneficiary, and the beneficiary requires the vehicle and is capable of using it.

(4) A written prescription from the beneficiary's physician is acceptable for ordering a power-operated vehicle if a specialist in physical medicine, orthopedic surgery, neurology, or rheumatology is not reasonably accessible. For example, if travel to the specialist would be more than one day's trip from the beneficiary's home or if the beneficiary's medical condition precluded travel to the nearest available specialist, these circumstances would satisfy the "not reasonably accessible" requirement.

(d) Medicare Part B pays for medically necessary equipment that is used for treatment of decubitus ulcers if—

(1) The equipment is ordered in writing by the beneficiary's attending physician, or by a specialty physician on referral from the beneficiary's attending physician, and the written order is furnished to the supplier before the delivery of the equipment; and

(2) The prescribing physician has specified in the prescription that he or she will be supervising the use of the equipment in connection with the course of treatment.

(e) Medicare Part B pays for a medically necessary seat-lift if it—

(1) Is ordered in writing by the beneficiary's attending physician, or by a specialty physician on referral from the beneficiary's attending physician, and the written order is furnished to the supplier before the delivery of the seat-lift;

(2) Is for a beneficiary who has a diagnosis designated by CMS as requiring a seat-lift; and

(3) Meets safety requirements specified by CMS.

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(f) Medicare Part B pays for transcutaneous electrical nerve stimulator units that are—

(1) Determined to be medically necessary; and

(2) Ordered in writing by the beneficiary's attending physician, or by a specialty physician on referral from the beneficiary's attending physician, and the written order is furnished to the supplier before the delivery of the unit to the beneficiary.

(g) As a requirement for payment, CMS may determine through carrier instructions, or carriers may determine that an item of durable medical equipment requires a written physician order before delivery of the item.

[51 FR 41339, Nov. 14, 1986, as amended at 57 FR 57688, Dec. 7, 1992; 58 FR 30668, May 26, 1993]

EFFECTIVE DATE NOTE: At 70 FR 50946, Aug. 26, 2005, § 410.38 was amended by revising paragraph (c), effective October 25, 2005. For the convenience of the user, the revised text is set forth as follows:

§ 410.38 Durable medical equipment: Scope and conditions.

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(c) *Power mobility devices (PMDs)*. (1) *Definitions*. For the purposes of this paragraph (c), the following definitions apply:

*Physician* has the same meaning as in section 1861(r)(1) of the Act.

*Power mobility device* means a covered item of durable medical equipment that is in a class of wheelchairs that includes a power wheelchair (a four-wheeled motorized vehicle whose steering is operated by an electronic device or a joystick to control direction and turning) or a power-operated vehicle (a three or four-wheeled motorized scooter that is operated by a tiller) that a beneficiary uses in the home.

*Prescription* means a written order completed by the physician or treating practitioner who performed the face-to-face examination and that includes, the beneficiary's name, the date of the face-to-face examination, the diagnoses and conditions that the PMD is expected to modify, a description of the item (for example, a narrative description of the specific type of PMD), the length of need, and the physician or treating practitioner's signature and the date the prescription was written.

*Treating practitioner* means a physician assistant, nurse practitioner, or clinical nurse specialist as those terms are defined in section 1861(aa)(5) of the Act, who has con-

ducted a face-to-face examination of the beneficiary.

*Supplier* means a durable medical equipment (DME) supplier.

(2) *Conditions of payment*. Medicare Part B pays for a power mobility device if the physician or treating practitioner, as defined in paragraph (c)(1) of this section:

(i) Conducts a face-to-face examination of the beneficiary for the purpose of evaluating and treating the beneficiary for his or her medical condition and determining the medical necessity for the PMD as part of an appropriate overall treatment plan;

(ii) Writes a prescription, as defined in paragraph (c)(1) of this section, which is provided to the beneficiary or supplier, and is received by the supplier within 30 days of the face-to-face examination.

(iii) Provides supporting documentation, including pertinent parts of the beneficiary's medical record (e.g., history, physical examination, diagnostic tests, summary of findings, diagnoses, treatment plans and/or other information as may be appropriate) that supports the medical necessity for the power mobility device, which is received by the supplier within 30 days after the face-to-face examination.

(3) *Exceptions*. (i) Beneficiaries discharged from a hospital do not need to receive a separate face-to-face examination as long as the physician or treating practitioner who performed the face-to-face examination of the beneficiary in the hospital issues a PMD prescription and supporting documentation that is received by the supplier within 30 days after the date of discharge.

(ii) Accessories for PMDs may be ordered by the physician or treating practitioner without conducting a face-to-face examination of the beneficiary.

(4) *Dispensing a power mobility device*. Suppliers may not dispense a PMD to a beneficiary until the PMD prescription and the supporting documentation have been received from the physician or treating practitioner who performed the face-to-face examination of the beneficiary. Such documents must be received within 30 days after the date of the face-to-face examination.

(5) *Documentation*. (i) A supplier must maintain the prescription and the supporting documentation provided by the physician or treating practitioner and make them available to CMS and its agents upon request.

(ii) Upon request by CMS or its agents, a supplier must submit additional documentation to CMS or its agents to support and/or substantiate the medical necessity for the power mobility device.

(6) *Safety requirements*. The PMD must meet any safety requirements specified by CMS.

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