

NHS MEDICAL POLICY

Continuous Passive Motion Devices DME 2014-006

Continuous passive motion devices may be indicated when **ONE** of the following is present:

- The member has undergone (or will undergo) any knee surgery and one of the following applies:
 The device is ordered post-operatively for up to 21 days of use as part of a comprehensive
 - rehabilitation program.
 - The device is ordered post-operatively for up to 21 days of use because the patient is not able to participate in physical or occupational therapy.
 - The device is ordered post-operatively for up to 21 days of use because physical or occupational therapy has been inadequate.

2 The member has undergone (or will undergo) surgical release of arthrofibrosis/adhesive capsulitis or manipulation under anesthesia of any joint (except in the spine) and one of the following applies:

- The device is ordered post-operatively for up to 21 days of use as part of a comprehensive rehabilitation program.
- The device is ordered post-operatively for up to 21 days of use because the patient is not able to participate in physical or occupational therapy.
- The device is ordered post-operatively for up to 21 days of use because physical or occupational therapy has been inadequate.

SOURCES

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- 1. UpToDate.com was accessed August 21, 2014 and January 5, 2016.
- 2. NIH.gov was accessed January 5, 2016.
- 3. Harvey LA, Brosseau L, Herbert RD. Continuous passive motion following total knee arthroplasty in people with arthritis. Cochrane Database Syst Rev 2010; CD004260.

- 4. Ritter MA, Stringer EA. Predictive range of motion after total knee replacement. Clin Orthop Relat Res 1979; 115.
- 5. O'Driscoll SW, Giori NJ. Continuous passive motion (CPM): Theory and principles of clinical application. J Rehabil Res Dev. 2000;37(2):179-188.
- 6. Postel JM, Thoumie P, Missaoui B, et al; French Physical Medicine and Rehabilitation Society. Continuous passive motion compared with intermittent mobilization after total knee arthroplasty. Elaboration of French clinical practice guidelines. Ann Readapt Med Phys. 2007;50(4):244-257.
- 7. Thien TB, Becker JH, Theis J-C. Rehabilitation after surgery for flexor tendon injuries in the hand. Cochrane Database Syst Rev. 2004;(4):CD003979

CODE REFERENCE (This may not be a comprehensive list of codes to apply to this policy.)

HCPCS: E0935, E0936

POLICY HISTORY/REVISION INFORMATION

Date	Action/Description
09/25/15	Annual review and approval by UM Committee
2/26/2016	Additions made to #1 and #2 and approved by UM Committee
12/14/2016	Annual review and approval by UM Committee
12/13/2017	Annual review and approval by UM Committee
12/13/2018	Annual review and approval by UM Committee
12/12/2019	Annual review and approval by UM Committee
12/10/2020	Annual review and approval by UM Committee
12/10/2021	Annual review and approval by UM Committee
12/21/2022	Annual review and approval by UM Committee
12/20/2023	Annual review and approval by UM/QM Committee