# **Medical Coverage Policy** | Home Uterine Activity Monitoring (HUAM)



**EFFECTIVE DATE:** 10 | 01 | 2015 **POLICY LAST UPDATED:** 09 | 18 | 2018

### **OVERVIEW**

The home uterine activity monitor (HUAM) is a device intended to provide early detection of preterm labor in women at high risk of preterm labor and preterm birth. A monitoring device worn by the patient collects data on uterine activity. After using the device, the patient transmits data recordings to a provider who assesses risk of preterm labor-onset, based on frequency of uterine contractions and responses to interview questions.

## **MEDICAL CRITERIA**

Not applicable

## **PRIOR AUTHORIZATION**

Not applicable

#### **POLICY STATEMENT**

#### BlueCHiP for Medicare

Home uterine activity monitoring, with or without nursing contact, is not covered, including use with tocolytic therapy (medications used to slow contractions). Despite numerous scientific studies, there is insufficient evidence to determine the effects of the technology on health outcomes. There is also no evidence that the use of home uterine monitoring improves health outcomes for mother or baby. It has not been shown to improve outcomes for twins, triplets, high-risk situations, or women who have already experienced preterm labor. According to U.S. Preventive Services Task Force, use of these monitors has not been shown to improve gestational age at birth, the baby's weight at birth, or neonatal morbidity.

# **Commercial Products**

Home uterine activity monitoring, with or without nursing contact, is considered not medically necessary, including use with tocolytic therapy (medications used to slow contractions). Despite numerous scientific studies, there is insufficient evidence to determine the effects of the technology on health outcomes. There is also no evidence that the use of home uterine monitoring improves health outcomes for mother or baby. It has not been shown to improve outcomes for twins, triplets, high-risk situations, or women who have already experienced preterm labor. According to U.S. Preventive Services Task Force, use of these monitors has not been shown to improve gestational age at birth, the baby's weight at birth, or neonatal morbidity.

## **COVERAGE**

Benefits may vary between groups/contracts. Please refer to the appropriate Benefit Booklet, Evidence of Coverage or Subscriber Agreement for applicable not medically necessary/not covered benefits/coverage.

#### **BACKGROUND**

The home uterine activity monitor (HUAM) is a monitoring device that consists of a guard-ring tocodynamometer (worn as a belt around the abdomen), a data recorder, and a data transmitter. Typically, the patient is instructed to use the device daily for two 1-hour periods. After monitoring, the patient transmits the recordings by telephone modem link to a remote base station. Base station nurses not only facilitate transmission and analysis of the monitor tracings, they also maintain daily telephone contact with the patient to assess signs and symptoms and to provide advice and counseling.

Nurses employed in HUAM services look for evidence of the onset of preterm labor, either on the basis of uterine activity exceeding a threshold level or from the findings of a telephone interview with the patient. Signs and symptoms of preterm labor include back pain, increased vaginal discharge, menstrual-like cramps, and pelvic pressure or heaviness. The threshold number of uterine contractions signaling the possible onset of preterm labor is usually 4 to 6 per hour. If signs and symptoms are present or the uterine activity exceeds a certain threshold, patients are instructed to perform the following: empty the bladder, hydrate orally, and assume the left lateral recumbent position. The patient is also instructed to re-monitor for 1 additional hour. If uterine activity still exceeds threshold or signs and symptoms persist, the patient is instructed to see her physician immediately for a cervical examination. The cervical examination would then play a pivotal role in diagnosing whether preterm labor is occurring and whether to initiate tocolytic therapy.

The HUAM is described as an electronic system for at-home antepartum measurement of uterine contractions, data transmission by telephone to a clinical setting, and for receipt and display of the uterine contraction data at the clinic. The HUAM system comprises a tocotransducer, an at-home recorder, a modem, and a computer and monitor that receive, process and display data. The FDA indicates that the device is intended for use in women at least 24 weeks' gestation with a previous preterm delivery to aid in the detection of preterm labor.

The available evidence suggests that HUAM does not improve health outcomes, and HUAM is not recommended by national organizations.

#### **CODING**

The following codes are not covered for BlueCHiP for Medicare and not medically necessary for Commercial Products:

99500 Home visit for prenatal monitoring and assessment to include fetal heart rate, non-stress test, uterine monitoring, and gestational diabetes monitoring

**S9001** Home uterine monitor with or without associated nursing services

## **RELATED POLICIES**

Not applicable

## **PUBLISHED**

Provider Update, November/December 2018 Provider Update, November 2017 Provider Update, January 2017 Provider Update, June 2015 Provider Update, June 2014

## **REFERENCES**

- 1. Blue Cross and Blue Shield Association Technology Evaluation Center (TEC). Home uterine activity monitoring for secondary prevention of preterm birth. TEC Assessments 1996; Volume 11, Tab 15.
- 2. Urquhart C, Currell R, Harlow F et al. Home uterine monitoring for detecting preterm labour. Cochrane Database Syst Rev 2012; 5:CD006172.
- 3. Honest H, Forbes CA, Duree KH et al. Screening to prevent spontaneous preterm birth: systematic reviews of accuracy and effectiveness literature with economic modeling. Health Technol Assess 2009; 13(43):1-627.
- A multicenter randomized controlled trial of home uterine monitoring: active versus sham device. The Collaborative Home Uterine Monitoring Study (CHUMS) Group. Am J Obstet Gynecol 1995; 173(4):1120-7.
- 5. Dyson DC, Danbe KH, Bamber JA et al. Monitoring women at risk for preterm labor. N Engl J Med 1998; 338(1):15-9.

- 6. Iams JD, Johnson FF, O'Shaughnessy RW. Ambulatory uterine activity monitoring in the post-hospital care of patients with preterm labor. Am J Perinatol 1990; 7(2):170-3.
- 7. Blondel B, Breart G, Berthoux Y et al. Home uterine activity monitoring in France: a randomized, controlled trial. Am J Obstet Gynecol 1992; 167(2):424-9.
- 8. Nagey DA, Bailey-Jones C, Herman AA. Randomized comparison of home uterine activity monitoring and routine care in patients discharged after treatment for preterm labor. Obstet Gynecol 1993; 82(3):319-23.
- 9. Brown HL, Britton KA, Brizendine EJ et al. A randomized comparison of home uterine activity monitoring in the outpatient management of women treated for preterm labor. Am J Obstet Gynecol 1999; 180(4):798-805.
- 10. U.S. Preventive Services Task Force. Screening Home Uterine Activity Monitoring. 1996. Available online at: http://www.uspreventiveservicestaskforce.org/uspstf/uspshuam.htm. Last accessed October, 2013.
- 11. National Institute of Child Health and Human Development. Home uterine monitors not useful for predicting premature birth. Available online at: http://www.nih.gov/news/pr/jan2002/nichda23.htm. Last accessed October, 2013.

# ----- CLICK THE ENVELOPE ICON BELOW TO SUBMIT COMMENTS

This medical policy is made available to you for informational purposes only. It is not a guarantee of payment or a substitute for your medical judgment in the treatment of your patients. Benefits and eligibility are determined by the member's subscriber agreement or member certificate and/or the employer agreement, and those documents will supersede the provisions of this medical policy. For information on member-specific benefits, call the provider call center. If you provide services to a member which are determined to not be medically necessary (or in some cases medically necessary services which are non-covered benefits), you may not charge the member for the services unless you have informed the member and they have agreed in writing in advance to continue with the treatment at their own expense. Please refer to your participation agreement(s) for the applicable provisions. This policy is current at the time of publication; however, medical practices, technology, and knowledge are constantly changing. BCBSRI reserves the right to review and revise this policy for any reason and at any time, with or without notice. Blue Cross & Blue Shield of Rhode Island is an independent licensee of the Blue Cross and Blue Shield Association.

