## DIVISION OF HEALTH CARE FINANCING AND POLICY NEVADA MEDICAID DRUG USE REVIEW (DUR) BOARD PROPOSED CRITERIA

## Makena<sup>TM</sup> (hydroxyprogesterone caproate) Injection

Makena<sup>TM</sup> is a covered benefit of Nevada Medicaid for recipients who meet the criteria for coverage.

1. Coverage and Limitations:

Authorization will be given if all of the following criteria are met and documented:

- a. Treatment with Makena<sup>TM</sup> is ordered by or recommended by a physician specializing in Obstetrics/Gynecology, Perinatology or Maternal/Fetal Medicine; AND
- b. The recipient is female, 16 years of age or older, and pregnant with a singleton pregnancy; AND
- c. The recipient is between 16 weeks 0 days and 20 weeks 6 days of gestation when therapy begins AND
- d. The recipient has a history of singleton spontaneous preterm birth (prior to 37 weeks gestation); AND
- e. The recipient does not have other risk factors for preterm birth; AND
- f. There is no known major fetal anomaly or fetal demise; AND
- g. The recipient has not been treated with heparin therapy during the current pregnancy; AND
- h. The recipient has no history of thromboembolic disease; AND
- i. The recipient has no maternal/obstetrical complication (e.g., current or planned cerclage, hypertension requiring medication or seizure disorder)

## 2. Quantity Limitations:

Quantity limit: 1 vial per rolling 30 days (5 doses)

**3.** Length of Approval: Makena will be approved for use until the recipient is 36 weeks 6 days of gestation or delivery, whichever occurs first.