

Medical prior authorization form Fax completed form to: 877.974.4411 toll free, or 616.942.8206 This form applies to: Commercial (Traditional) Commercial (Individual/Optimized) Medicaid This request is: Urgent (life threatening) Non-Urgent (standard review) Urgent means the standard review time may seriously jeopardize the life or health of the patient or the patient's ability to regain maximum function. Hydroxyprogesterone caproate (non-Makena J1729) Member

Member			
Last Name:		First Name:	
		DOB:	Gender:
Requesting Provider:		Prov. Phone:	Prov. Fax:
Provider Address:			
Provider NPI:		Contact Name:	
Provider Signature:		Date:	
Product and Billing	g Information		
☐ New Request ☐ Co	ontinuation Request		
Drug product: ☐ Hydroxyprogesterone caproate 250 mg/mL (J1729)		Start date (or date of next dose):	
		Date of last dose (if applicable):	
		Dosing frequency:	
Dlace of administration	Dhysician's office	Current gestational age:	_ weeks, days
Place of administration:	<u> </u>		
	Outpatient infusion		_
	Facility:	NPI:	Fax:
	☐ Home infusion		
	Facility:	NPI:	Fax:
Billing:	☐ Physician to buy and bill		
	☐ Facility to buy and bill		
	☐ Specialty Pharmacy		
	Pharmacy:	NPI:	Fax:
ICD code(s):			

Precertification Requirements

Before this drug is covered, the patient must meet all of the following requirements:

- 1. Must be used for a FDA-approved diagnosis or to reduce the risk of preterm birth
- 2. If using to reduce risk of preterm birth all of the following must be met:
 - a. Must be a singleton pregnancy
 - b. Woman must have a history of a prior spontaneous preterm birth of singleton pregnancy
 - c. The first weekly injection of hydroxyprogesterone caproate must be started on or after 16 weeks gestation, but before 27 weeks gestation
 - d. Hydroxyprogesterone caproate must be stopped at 36 weeks, 6 days gestation or delivery, whichever comes first



Additional information

Priority Health supports the prescriber's discretion to use either hydroxyprogesterone caproate (non-Makena J1729) or compounded preservative-free 17-P based on the facts and circumstances of the individual patient. The compounded formulation will continue to be a covered alternative and does not require prior approval.

NOTE: Hydroxyprogesterone caproate (Makena J1726) is not covered.

Note: Authorization for indications, dosing, or a route of administration not approved by the Food and Drug Administration (FDA) or recognized in CMS-accepted compendia (e.g. DrugDex, AHFS, U.S. Pharmacopeia, and also Clinical Pharmacology for oncology indications only) require supporting evidence for coverage. Please provide two published peer-reviewed literature articles supporting the appropriateness of the drug, the dosing of the drug, or the route of administration to be used for the identified indication.

Pr	iorit	y Health Precertification Documentation
Α.	Wh	at condition is this drug being requested for? Advanced adenocarcinoma of the uterine corpus (Stage III or IV) Management of amenorrhea (primary and secondary) Abnormal uterine bleeding due to hormonal imbalance in the absence of organic pathology Test for endogenous estrogen production Production of secretory endometrium and desquamation Reduce risk of preterm birth Other – the patient's condition is:
В.	ls t	he patient pregnant? ☐ Yes ☐ No
lf h	ıydr	oxyprogesterone is being requested to reduce risk of preterm birth, answer the following:
	A.	Will hydroxyprogesterone caproate be used for a singleton pregnancy? ☐ Yes ☐ No – explain why hydroxyprogesterone caproate is required:
	В.	Does the patient have a history of a spontaneous preterm birth of singleton pregnancy? ☐ Yes ☐ No – explain why hydroxyprogesterone caproate is required: