

**MAKENA
(HYDROXYPROGESTERONE CAPROATE)**



Keystone First

PERFORMRxSM
Next Generation Pharmacy Benefits

PRIOR AUTHORIZATION FORM

(form effective 1/1/20)

Fax to PerformRxSM at **1-215-937-5018**, or to speak to a representative call **1-800-588-6767**.

PRIOR AUTHORIZATION REQUEST INFORMATION

<input type="checkbox"/> New request	<input type="checkbox"/> Renewal request	total # pages:	Name of office contact:
Contact's phone number:		LTC facility contact/phone:	

PATIENT INFORMATION

Patient name:	Patient ID #:	DOB:
Street address:	Apt. #:	City/state/zip:

PRESCRIBER INFORMATION

Prescriber name:	Specialty:	
State license #:	NPI:	MA Provider ID #
Street address:	Suite #:	City/state/zip:
Phone:	Fax:	

PHARMACY INFORMATION (PRESCRIBER TO IDENTIFY THE PHARMACY THAT IS TO DISPENSE THE MEDICATION):

Deliver to: Patient's Home Physician's Office Patient's Preferred Pharmacy Name:

Pharmacy Phone #: _____ Pharmacy Fax #: _____

I acknowledge that the patient agrees with the pharmacy chosen for delivery of this medication.

CLINICAL INFORMATION

Medication requested:

<input type="checkbox"/> hydroxyprogesterone caproate injection (non-preferred)	<input type="checkbox"/> Makena 250 mg/ml (1 ml) single-dose vial*
<input type="checkbox"/> Makena 275 mg/1.1 ml autoinjector*	<input type="checkbox"/> Makena 250 mg/ml (5 ml) multi-dose vial*
<input type="checkbox"/> _____	

*preferred with clinical prior authorization required

Dose/directions:	Quantity:	Refills:
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Diagnosis (submit documentation): pregnancy with history of pre-term labor other: _____

DX codes (required): _____ Start date of therapy: _____ / _____ / 20_____

1. Is the patient currently pregnant with a single fetus? Yes No *Submit supporting documentation.*

2. What is the current gestational age? Weeks: _____ Days: _____

3. Does the patient have a documented history of a prior spontaneous pre-term singleton birth (defined as prior to 37 weeks' gestation)? Yes No *Submit supporting documentation.*

4. Does the patient have any of the following contraindications to the use of Makena? **Check all that apply.**

<input type="checkbox"/> current or history of thrombosis or thromboembolic disorders	<input type="checkbox"/> Yes <input type="checkbox"/> No <i>If yes, submit supporting documentation.</i>
<input type="checkbox"/> history of or current known or suspected breast cancer or other hormone-sensitive cancer	
<input type="checkbox"/> undiagnosed abnormal vaginal bleeding unrelated to pregnancy	
<input type="checkbox"/> cholestatic jaundice of pregnancy	
<input type="checkbox"/> benign or malignant liver tumors or active liver disease	
<input type="checkbox"/> uncontrolled hypertension	

5. Does the patient have any of the following conditions?. **Check all that apply.**

<input type="checkbox"/> history of, or plans for, a cervical cerclage	<input type="checkbox"/> Yes <input type="checkbox"/> No <i>If yes, submit supporting documentation.</i>
<input type="checkbox"/> known fetal anomaly	
<input type="checkbox"/> history of seizure disorder	

6. For non-preferred hydroxyprogesterone caproate (generic Makena): Does the patient have a history of trial and failure, contraindication, or intolerance of the preferred agent – Makena? Yes No *Submit documentation.*

PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION

Prescriber signature:	Date:
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