Pregnancy Notification (First Prenatal Visit)

Please fax **both pages** to 702-691-5620 (Must be faxed within 15 days of first visit)



type of Referral:		
☐ Pregnancy Notification	☐ High Risk Pregnancy	☐ Miscarriage/Termination Notification
Member ID#:		
Patient Name:		
Street Address:		
City/State:		
Phone:		
Date of Birth:		
LMP:	EDC:	GESTATIONAL AGE:
PARA	GRAVIDA	
Physician:		
Street Address:		
City/State/Zip:		
Phone:		
Tax ID#:		

The information contained in this facsimile is confidential and includes protected patient health information. The information is intended only for the use of CHF and its designees.

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For Office Use Only

Background Summary for Questionnaires.

These are evidence based guidelines which treating providers are encouraged to use, along with their own clinical judgment:

- 17-H Progesterone Use in Pregnancy Guidelines
 - 17-OH progesterone has been shown to decrease the risk of preterm delivery.
- Aspirin Use in Pregnancy Guidelines

Low dose aspirin has been shown to decrease the risk of preeclampsia in pregnancy and the associated complications.

Patient Name
Culinary ID#

High Risk Factor Questions		No
1. Is there a history of preeclampsia in a prior pregnancy?		
2. Is the current pregnancy multiple gestation?		
3. Is chronic hypertension present?		
4. Is pre-gestation diabetes present?		
5. Is renal disease present?		
6. Is autoimmune disease or are antiphospholipid antibodies present? (antiphospholipid syndrome, lupus, etc.)		

If the answer to ANY ONE (1) High Risk Factor Question is YES, the patient may be a candidate for Aspirin treatment, if no contraindications exist. Move to section B - Exclusion Criteria.

B. Exclusion Criteria

Do not use Aspirin if any of the following contraindications exist.

Contraindications to the use of Aspirin	Yes	;	No
1. Allergy to Aspirin			
2. Bleeding disorder			
3. Platelet disorder			
4. Recent vaginal bleeding			
5. Reye's Syndrome			
6. Inadequate vitamin K			
7. G6PD deficiency			
If ANY contraindication is present, do not use	e Aspirin		

17 - OH Progesterone

A. Inclusion Criteria

Inclusion Criteria		No
1. Is there a history of previous singleton preterm delivery from 20-36 6/7 weeks gestation?		
2. Was the previous preterm delivery from spontaneous labor or PPROM?		

If the answer to BOTH Inclusion Criteria questions are YES, the patient may be a candidate for weekly 17-OH progesterone treatment, if no contraindications exist. Move to section **B** - **Exclusion Criteria**

B. Exclusion Criteria

Do not use 17-OH progesterone if any of the following contraindications exist.

Contraindications to the use of 17-OH Progesterone		No
Current pregnancy is multiple (2 or more) gestation		
2. Current pregnancy is more than 20 6/7 weeks gestation		
3. Current or history of thrombosis or thromboembolic disorders		
4. Known or suspected breast cancer, other hormone sensitive cancer or a history of these conditions		
5. Undiagnosed abnormal vaginal bleeding unrelated to pregnancy		
6. Cholestatic jaundice of pregnancy		
7. Liver tumors, benign or malignant or active liver disease		
8. Uncontrolled hypertension		
9. Allergy to hydroxyprogesterone caproate, castor oil or any of the other ingredients in 17-OH Progesterone (Makena)		
10. Any other contraindications to 17-OH progesterone: (list here)		

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