



## Cardiomyopathy in Pregnancy

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### Course Description:

- In all of the United States' public health, pregnancy-related mortality has the widest and most persistent racial and ethnic disparity (inequality). Alarming is the fact that African-American women have a 3 to 4-fold greater risk of maternal mortality than women of other ethnic or racial groups. The disparities stem from social, medical, clinical care and health system factors. Provider and institutional biases are factors impacting these health disparities as well.
- Cardiomyopathy has both intrigued and terrified obstetricians for more than 150 years. It is the leading cause of maternal mortality with cardiovascular disease accounting for one-third of all pregnancy-related deaths! Terrorizing is the fact that only a small fraction of these women had a known diagnosis of heart disease prior to death. However, the majority of women who died had presented with symptoms either during pregnancy or postpartum.

**Approximate Time to Complete:** 60 minutes



Introduction





**Hi! My name is Chantel. I will be guiding you through an introduction to cardiomyopathy in pregnancy as well as a case study.**



**My name is Sachi and I will guide you through peripartum cardiomyopathy in more depth.**

Meet the Presenters





**Here's how to use this module:  
The buttons above from left to  
right - Table of Contents,  
Home, Help, References and  
the X to close.**





**This course will:**

- Help participants develop sound critical judgement regarding the possibility of cardiomyopathy.
- Expand participant's knowledge base on learning theories and instructional implications regarding symptoms of cardiomyopathy.
- Enable participants to develop, implement and evaluate healthcare delivery in practice setting prior to an actual event.
- Enhance participant's ability to put knowledge into active health care delivery.
- Prepare participants to address issues and implement changes in the health care setting as necessary to ensure a safe environment.



- Introduction
- Cardiovascular Disease (CVD)
- Intro to Case Example
- Case Example
- Racial Disparities
- Racial Disparities - Maternal and Pregnancy-Related Mortality
- Racial Disparities - CVD Incidence and Comorbidities
- Racial Disparities - Peripartum Cardiomyopathy (PPCM) Prevalence
- Racial Disparities - Cont'd
- Racial Disparities- Provider and Institutional Bias
- Racial Disparities - Clinical Implications
- Algorithm 1 – Red Flags Not Present, Move on to Next Algorithm
- Algorithm 2 Assessment
- To Impact Change for the Presented Case, Other Orders Would Be 1
- B-type Natriuretic Peptide (BNP)
- Presentation of Women with CVD
- American College of Obstetricians & Gynecologists (ACOG) table
- Contributing Factors to Maternal Morbidity and Mortality with CVD – I
- Incidence of Peripartum Cardiomyopathy
- Defining Peripartum Cardiomyopathy
- Risk Factors
- Review of Cardiac Physiology Related to Pregnancy
- PPCM Introduction
- Conditions Associated with PPCM
- Theory on Pathophysiology
- Pathophysiology, Other Theories
- Intro to Pearls
- Needs for Providers



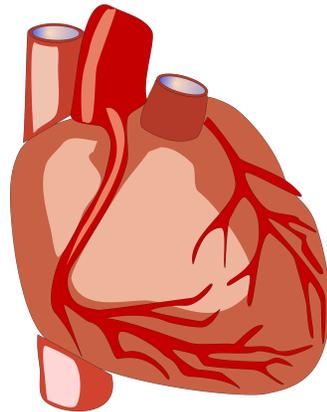


-  Incidence of Peripartum Cardiomyopathy
-  Defining Peripartum Cardiomyopathy
-  Risk Factors
-  Review of Cardiac Physiology Related to Pregnancy
-  PPCM Introduction
-  Conditions Associated with PPCM
-  Theory on Pathophysiology
-  Pathophysiology, Other Theories
-  Intro to Pearls
-  Pearls for Providers
-  Algorithm 1 – Red flags not present, move on to next algorithm.
-  Algorithm 2 Assessment
-  B-type Natriuretic Peptide (BNP) a tool
-  Diagnosis
-  Studies
-  Summary
-  Summary Cont'd
-  Summary Cont'd
-  Summary Cont'd
-  Summary Cont'd
-  Summary Cont'd
-  Summary Cont'd
-  Summary Cont'd
-  Summary Cont'd
-  Pregnancy Complications Infographic
-  Patient Education



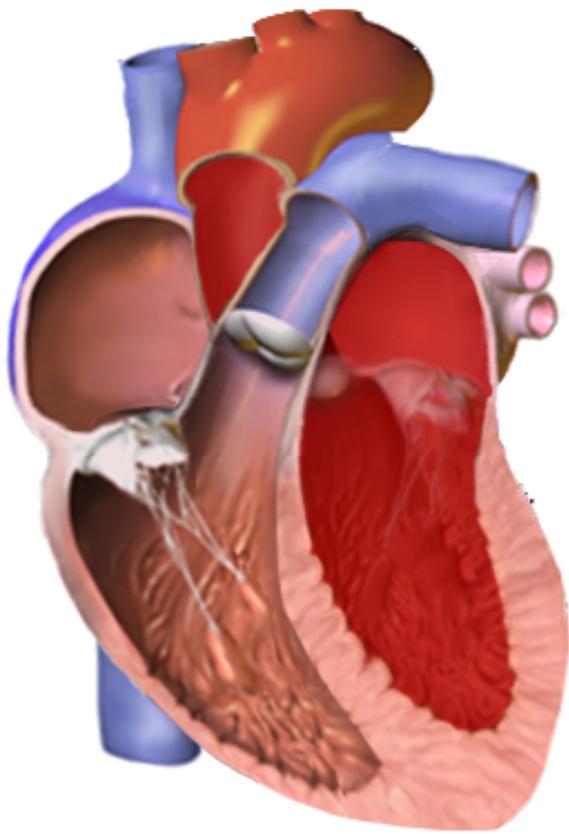
## Introduction [\[1.2\]](#)

**Peripartum heart failure has both intrigued and terrified obstetricians for more than 150 years.**

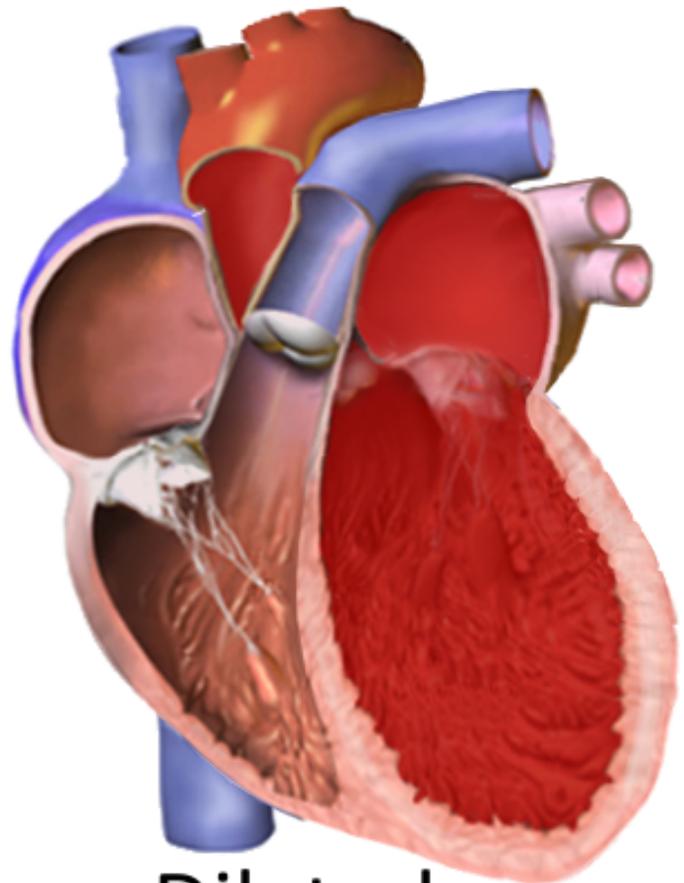


Click the heart for some pictures of common cardiomyopathies.

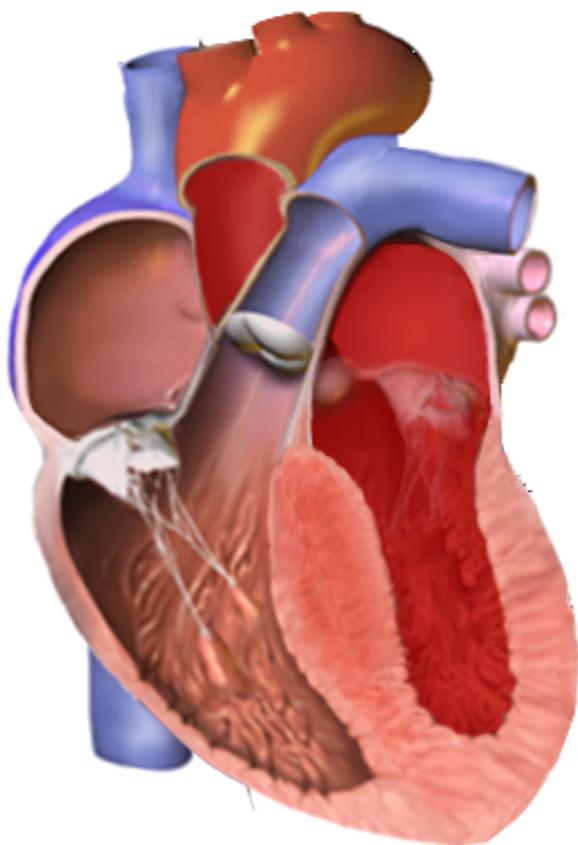
- Cardiomyopathy is a chronic disease of the heart muscle.
- The group of disorders where the heart muscle is structurally and functionally abnormal in the absence of other disease is cardiomyopathy.
- The most common cardiomyopathies are:
  - Hypertrophic
  - Dilated
- Rare types are:
  - Arrhythmogenic right ventricular
  - Restrictive
  - Takotsubo and left ventricular non-compaction cardiomyopathies.



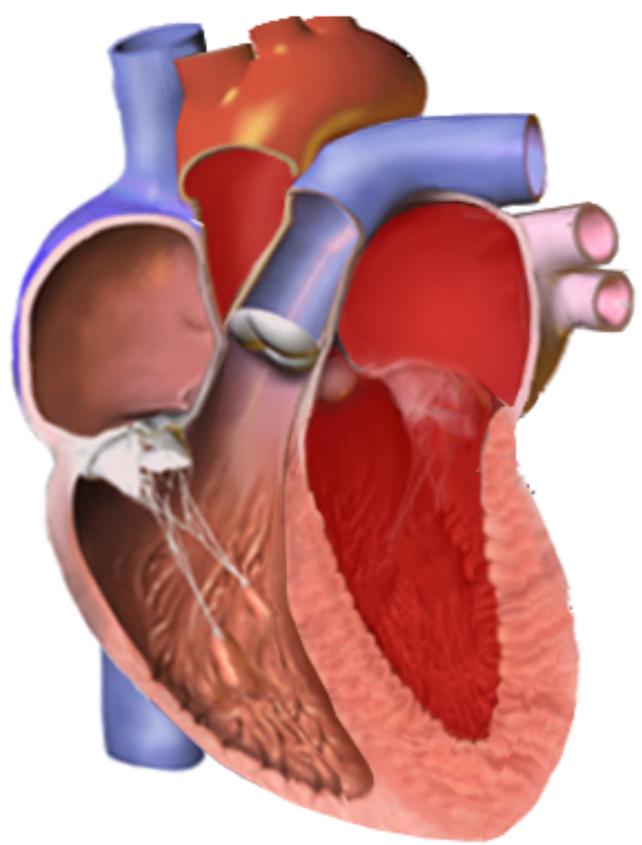
Normal



Dilated



Hypertrophic



Restrictive

## Cardiovascular Disease (CVD) [3]



**The leading cause of maternal mortality lies in cardiovascular disease in the United States (U.S.) and accounts for one-third of all pregnancy-related deaths!**

1 of 3 



Click the next arrows to see what Chantel has to say about cardiovascular disease

Cardiovascular Disease (CVD)



# Cardiovascular Disease (CVD) <sup>[3]</sup>



**Terrorizing is the fact that only a small fraction of these women had a known diagnosis of heart disease prior to death. However, the majority of women who died presented with symptoms either during pregnancy or postpartum.**

◀ 2 of 3 ▶



Click the next arrows to see what Chantel has to say about cardiovascular disease

Cardiovascular Disease (CVD)



## Cardiovascular Disease (CVD) <sup>[3]</sup>



**Despite numerous attempts to uncover the etiology of peripartum cardiomyopathy (PPCM), the cause remains unknown and may be multifactorial.**

◀ 3 of 3 ▶

Cardiovascular Disease (CVD)



Click the next arrow to continue





**Let's start  
learning more by  
taking a look at a  
case example.**





Case Example

“Sample Case Presentation A 25-year-old obese (Body Mass Index (BMI) 38) African-American G2P2 underwent an uncomplicated vaginal delivery 10 days ago. She presents to the urgent care clinic with complaints of fatigue and persistent cough since delivery. She is afebrile with blood pressure of 110/80 mmHg, heart rate 110 bpm and respiratory rate of 28 per minute. Chest X-ray reveals bilateral infiltrates. Oxygen saturation is 94% on room air.”

1 of 3 



Click the next arrows to  
read the case study





"The patient is diagnosed with a respiratory infection. Fatigue is attributed to the lack of sleep due to care of the newborn. She is prescribed an antibiotic and sent home. One week later, she presents again with continued symptoms. Antibiotics are switched at this time, and beta agonists are added due to presumptive diagnosis of "new-onset asthma" as evidenced by physical examination findings. Two days later, the patient experiences cardiac arrest at home. Resuscitation attempts are unsuccessful. Autopsy findings were indicative of cardiomyopathy."

◀ 2 of 3 ▶



Click the next arrows to  
read the case study

Case Example





"This case is representative of similar deaths attributed to cardiovascular disease reviewed by California Pregnancy-Associated Mortality Review (CA-PAMR). Maternal mortality due to cardiac disease primarily revolved around the lack of awareness of CVD at both patient and provider levels, coupled with delays in diagnosis. In most cases, diagnosis was made in the perimortem period or at the time of autopsy." [4]

◀ 3 of 3 ▶



Click the next arrow to continue



Case Example



## Racial Disparities [\[4\]](#)

- Clinical implications for African-American women with cardiovascular disease.
  - Retaining a high index of suspicion when CVD risk factors are present is needed to help impact the known racial disparities.

# Racial Disparities - Maternal and Pregnancy-Related Mortality



**Pregnancy-related mortality has the widest and most persistent racial and ethnic disparity (inequality) in all of U.S. public health [5].**

- The fact that African-American women have a 3-4 fold greater risk of maternal mortality than women of other ethnic or racial groups is alarming. [5].
- In California, pregnancy-related CVD is more than 8 times higher than that for white women [6].



## Racial Disparities - CVD Incidence and Comorbidities [5,6]

- Higher rates of pre-existing CVD occurs among African-American women [5].
- Comorbid conditions, such as hypertensive disorders of pregnancy, are more pervasive among African-American women.
  - Hypertensive disorders correlate to all types of CVD.
- It has been reported that African-American women have a lower number of visits for prenatal care and seek care later in pregnancy when compared to women of other racial and ethnic groups [7].



## Racial Disparities - Peripartum Cardiomyopathy (PPCM) Prevalence [8-11]



**When compared to white women, African-American women have an increased incidence of CVD in general and PPCM.**

- All forms of hypertensive disease in pregnancy highly correlate to PPCM
  - Women with gestational hypertension, chronic hypertension, and mild preeclampsia had 2- to 5-fold increases in odds of PPCM.
  - Women with severe preeclampsia had a 17-fold increase in odds of PPCM and eclampsia had a 25-fold increase.
- Reviewing the population with PPCM, African-American women were typically younger, more frequently presented with severe symptoms, and were more likely to be diagnosed postpartum than white women.



## Racial Disparities

Factors Affecting Racial Disparities in CVD-Related Diagnoses [[12-19](#)]:

Maternal mortality and morbidity are impacted by racial and ethnic disparities that stem from social, medical, clinical care, and health system factors.

1 of 5 



Click the next arrows to read more information.



## Racial Disparities

Factors Affecting Racial Disparities in CVD-Related Diagnoses [[12-19](#)]:

The incidence of hypertension and preeclampsia is higher with African-American women, yet these women are less likely to be hospitalized for treatment.

◀ 2 of 5 ▶



Click the next arrows to read more information.



## Racial Disparities

Factors Affecting Racial Disparities in CVD-Related Diagnoses [[12-19](#)]:

Maternal health outcomes are likely impacted by the chronic stress and experiences of racism over the course of life contributing to the higher rates of low birth weight infants and infant mortality.

◀ 3 of 5 ▶



Click the next arrows to read more information.



## Racial Disparities

Factors Affecting Racial Disparities in CVD-Related Diagnoses [[12-19](#)]:

Birth outcomes have known factors where high stress levels (the allostatic load) both before and after pregnancy impact the outcomes. African-American women are subject to higher loads which is multi-factorial with 'embodied inequality' arising from environmental, social, and genetic factors.

◀ 4 of 5 ▶



Click the next arrows to read more information.



# Racial Disparities

Factors Affecting Racial Disparities in CVD-Related Diagnoses [[12-19](#)]:

These are some of the socially patterned factors impacting maternal health, such as: malnutrition, toxic substance exposure, intimate partner violence, smoking, infections, racial discrimination, inadequate medical, and dental care.

- These factors highly correlate to low socioeconomic status.
- African-American women living in underserved areas often lack resources to maintain a healthy lifestyle, further impacting the negative cycle of poor reproductive and maternal health.
- What is even more frightening is the fact that African-American women with higher socioeconomic status and education does not necessarily resolve nor provide a protective effect against poor maternal outcomes.

◀ 5 of 5



# Racial Disparities - Provider and Institutional Bias [19-23]



The doctor-patient interaction is influenced by racial bias (conscious or not), gender bias (conscious or not), social ideas about race and class. These issues can shape and influence treatment disparities.



Click the picture to learn more about provider and institutional bias.

Research has examined physicians' implicit racial attitudes and found:

- [African-American Physicians](#)
- [Woman Physicians](#)
- [White Male Physicians](#)



Click the links to see implicit racial attitudes of each physician type.



African-American patients are less likely to have cardiovascular therapy of proven benefit and worse outcomes after the procedures. This has been shown in multiple studies.

- Unfortunately, multiple studies show, surgical and medical care often occurs at lower quality health care facilities and experience higher mortality regarding African-American patients.





**African-American doctors show no preference toward care for White or African-American patients.**

picture credit: [www.audio-luci-store.it](http://www.audio-luci-store.it)



Women physicians showed less implicit bias than their male counterparts.



White male physicians demonstrated significant implicit preference for White compared to non-White patients.



## Racial Disparities - Clinical Implications [24-26]

Complaints from patients need to be taken seriously, and clinicians need to maintain a high suspicion for CVD, especially when seeing African-American women who are pregnant or postpartum.



Click the next arrows to read more information.

1 of 3 





## Racial Disparities - Clinical Implications [24-26]

- Careful, thorough evaluations need to occur when African-American women having chronic or gestational hypertension with high pre-pregnancy BMI ( $\geq 35$ ) presenting with:
  - Extreme shortness of breath, especially when lying down
  - Persistent cough unrelieved with treatment
  - Significant fatigue
  - Palpitations
  - Swelling
  - Chest pain
  - Abnormal vital signs



Click the next arrows to read more information.

◀ 2 of 3 ▶



## Racial Disparities - Clinical Implications [24-26]

- We each have an opportunity to reduce disparities when diagnosing and treating CVD by implementing standardized protocols:
  - Assessment algorithms (an option follows)
  - Patient-level interventions with signs and symptoms
  - Examining the *process*
  - Content of educational interventions



Click the next arrow to continue

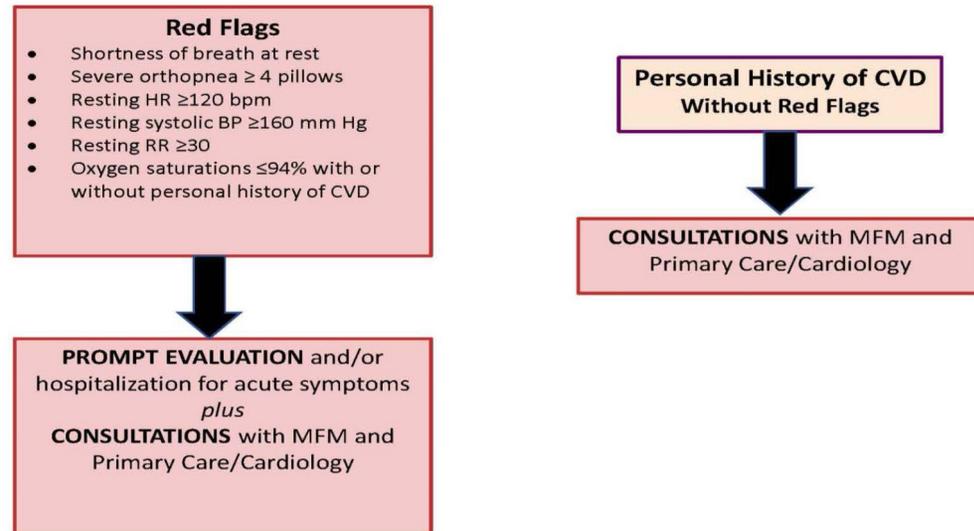
3 of 3



## Algorithm 1 – Red Flags Not Present, Move on to Next Algorithm [4]



Check for red flags first



Click the picture for a larger view

### Red Flags

- Shortness of breath at rest
- Severe orthopnea  $\geq 4$  pillows
- Resting HR  $\geq 120$  bpm
- Resting systolic BP  $\geq 160$  mm Hg
- Resting RR  $\geq 30$
- Oxygen saturations  $\leq 94\%$  with or without personal history of CVD



**PROMPT EVALUATION** and/or hospitalization for acute symptoms  
*plus*  
**CONSULTATIONS** with MFM and Primary Care/Cardiology

**Personal History of CVD  
Without Red Flags**



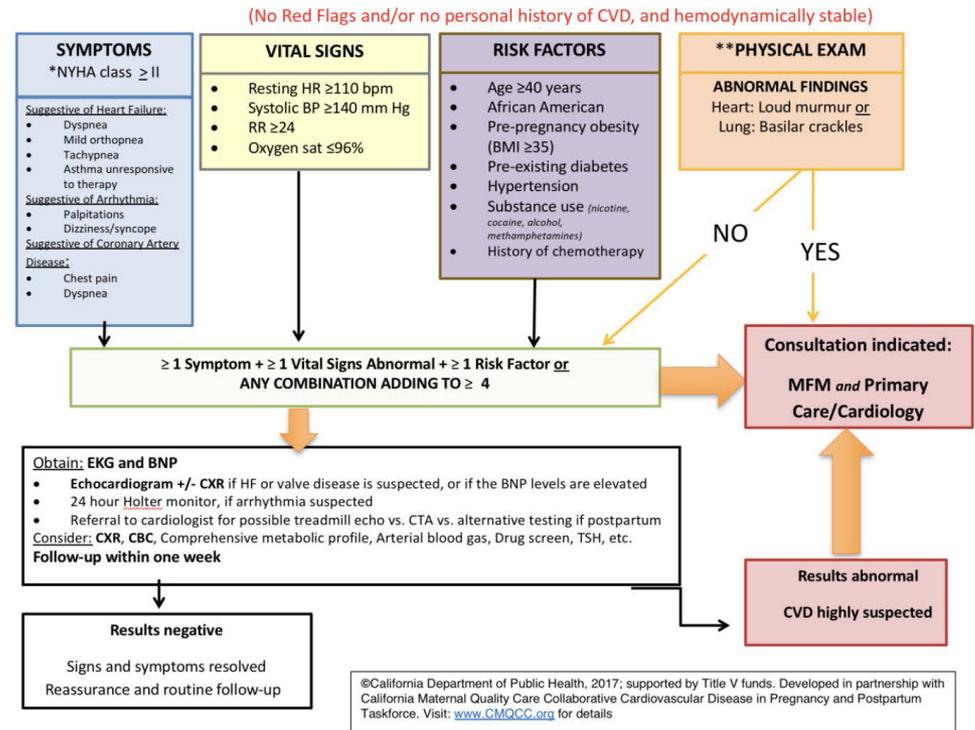
**CONSULTATIONS** with MFM and Primary Care/Cardiology

[4]

# Algorithm 2 Assessment



When there are no Red Flags nor personal history of CVD and the patient is hemodynamically stable this algorithm can be utilized:

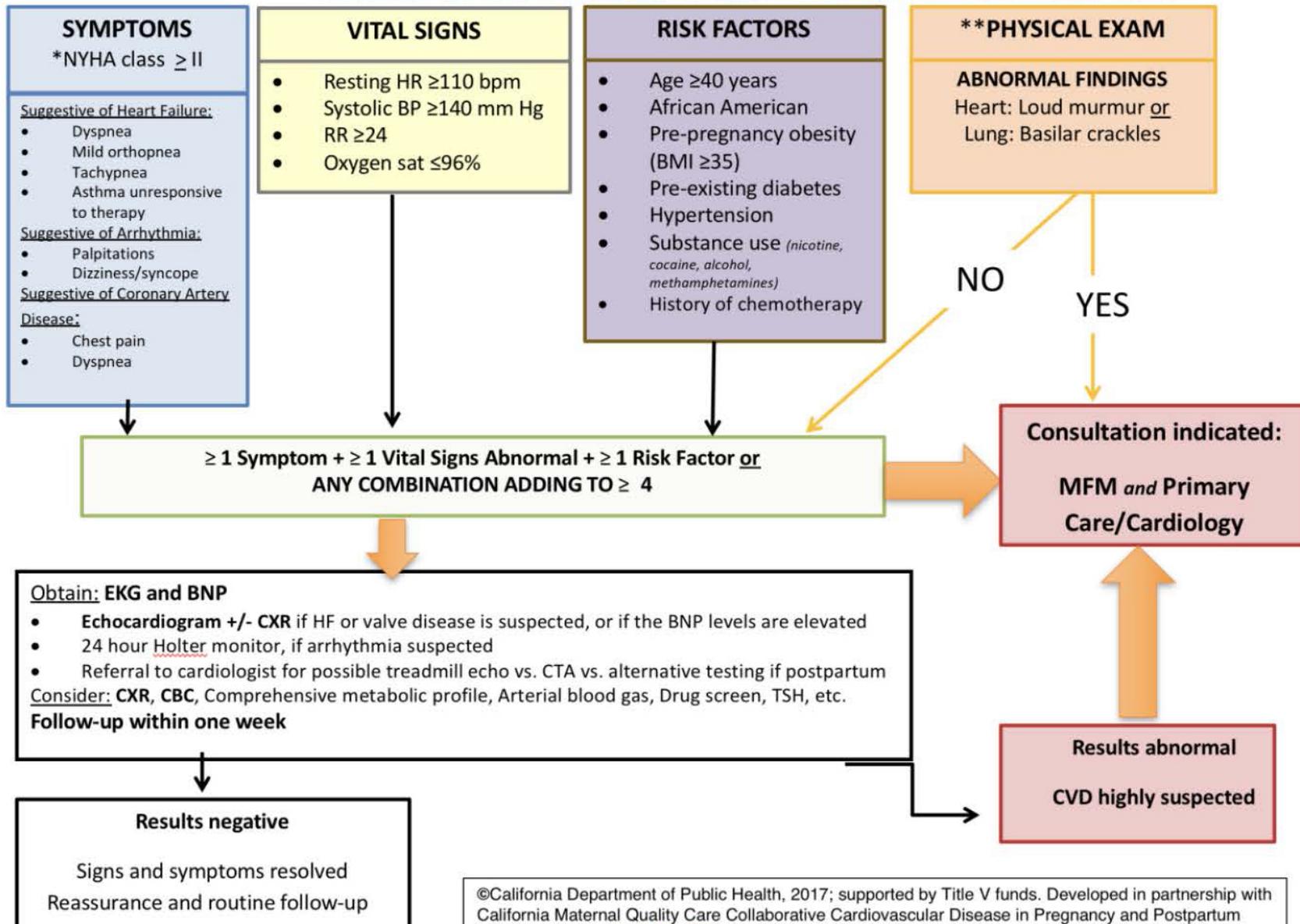


Click the picture for a larger view

Algorithm 2 Assessment



(No Red Flags and/or no personal history of CVD, and hemodynamically stable)



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To impact change for the presented case, other orders would be needed [4].

Click here to review the case study.



- Electrocardiogram (EKG)
- B-type natriuretic peptide (BNP)
- Other tests to consider:
  - Arrhythmia monitor
  - Echocardiogram
  - Chest X-ray
  - Complete blood count (CBC)
  - Comprehensive metabolic panel (CMP)
  - Arterial blood gas (ABG)
  - Assessment of thyroid function
  - A drug screen may also be considered



Click the next arrows on the clipboard to read more information.



To impact change for the presented case, other orders would be needed [4].

Click here to review the case study.



- The following Red Flags MUST signify further evaluation and possibly involve hospitalization or Maternal Fetal Medicine (MFM) physicians:
  - Shortness of breath at rest
  - Severe orthopnea  $\geq 4$  pillows
  - Resting HR  $\geq 120$  bpm
  - Resting systolic BP  $\geq 160$  mmHg
  - Resting RR  $\geq 30$
  - Oxygen saturations  $\leq 94\%$  with or without personal history of cardiovascular disease



Click the next arrow to continue

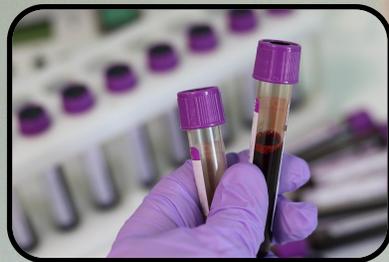


“Sample Case Presentation A 25-year-old obese (Body Mass Index (BMI) 38) African-American G2P2 underwent an uncomplicated vaginal delivery 10 days ago. She presents to the urgent care clinic with complaints of fatigue and persistent cough since delivery. She is afebrile with blood pressure of 110/80 mmHg, heart rate 110 bpm and respiratory rate of 28 per minute. Chest X-ray reveals bilateral infiltrates. Oxygen saturation is 94% on room air. The patient is diagnosed with a respiratory infection. Fatigue is attributed to the lack of sleep due to care of the newborn. She is prescribed an antibiotic and sent home. One week later, she presents again with continued symptoms. Antibiotics are switched at this time, and beta agonists are added due to presumptive diagnosis of “new-onset asthma” as evidenced by physical examination findings. Two days later, the patient experiences cardiac arrest at home. Resuscitation attempts are unsuccessful. Autopsy findings were indicative of cardiomyopathy.

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BNP is a readily available test. It may help identify asymptomatic women with left ventricular dysfunction and assist in triaging pregnant or postpartum women who present with symptoms.



Click the image above to read more information.

## B-type Natriuretic Peptide (BNP) [[27-29](#)]

- BNP is a neurohormone. It is secreted predominantly by the cardiac ventricles in response to volume or pressure overload.
- A BNP level of  $<100$  pg/mL is considered normal. BNP's half-life is 20 minutes.
- BNP's use has been validated in the diagnosis of systolic and diastolic heart failure.

B-type Natriuretic Peptide (BNP)





**BNP is a readily available test. It may help identify asymptomatic women with left ventricular dysfunction and assist in triaging pregnant or postpartum women who present with symptoms.**



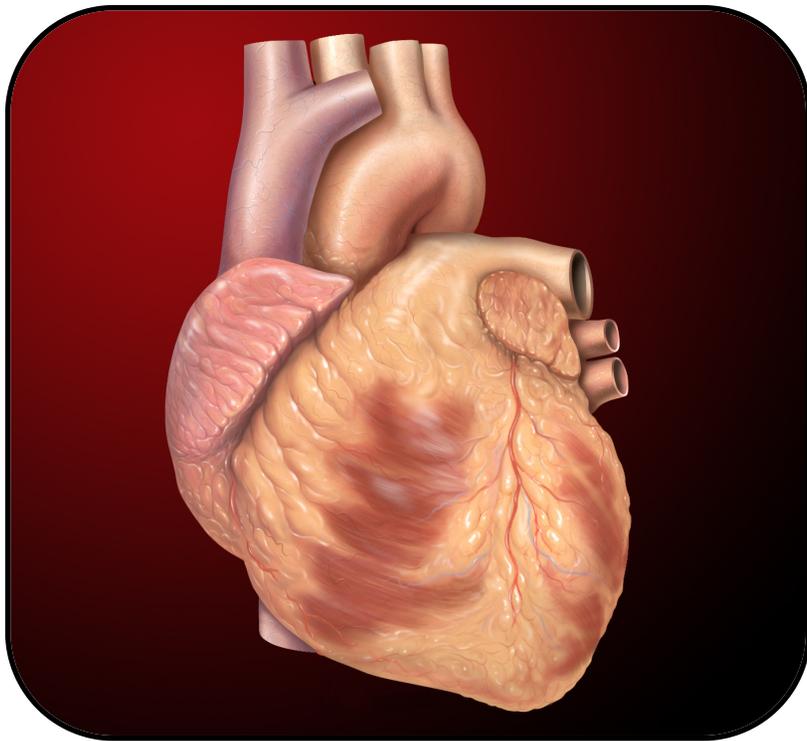
**Click the image above to read more information.**

## **B-type Natriuretic Peptide (BNP) [[27-29](#)]**

- BNP levels in pregnancy remain within normal range despite significant volume overload in pregnancy, and the levels are higher in pathologic conditions.
- An elevated BNP level should trigger an echocardiogram to evaluate cardiac function.
- Serial measurements of BNP in pregnant women with dilated cardiomyopathy are shown to be predictive of adverse cardiovascular outcomes.

**B-type Natriuretic Peptide (BNP)**





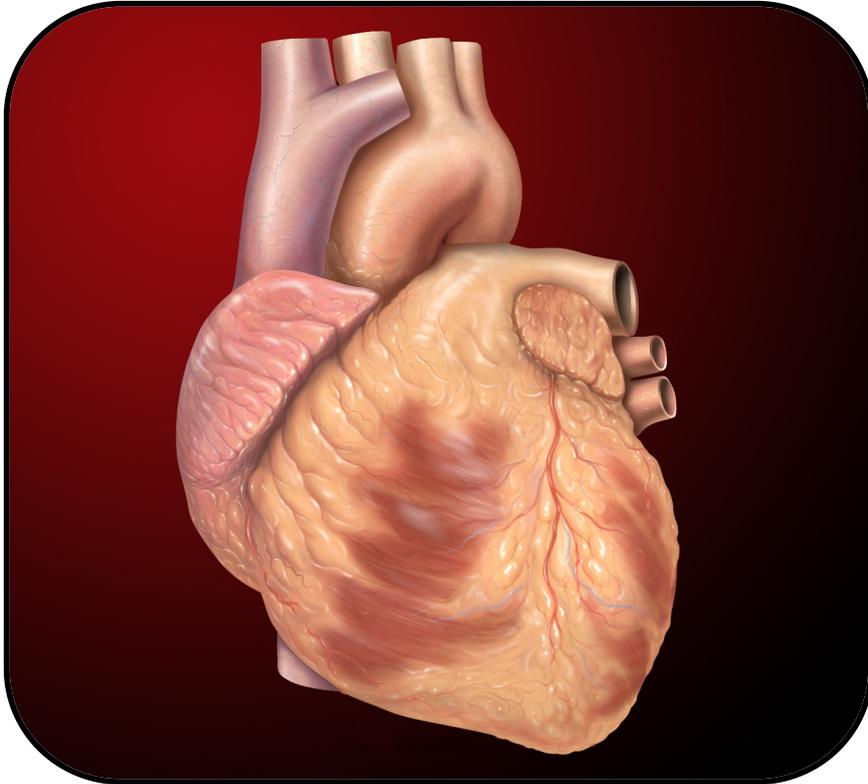
Click the heart graphic above to read more information.

Presentation of Women with CVD

### Presentation of Women with CVD:

- In the California Collaborative study, only 2 women entered pregnancy with known CVD [6].
- However, the prevalence of CVD symptoms (SOB, wheezing, palpitations, edema, chest pain, dizziness, or extreme fatigue) follow:
  - Prenatal period 43%
  - Labor and delivery 51%
  - Postpartum 80%





Click the heart graphic above to read more information.

## Presentation of Women with CVD:

- It can be tricky to determine pathology from normal pregnancy variance since many young healthy pregnant women without cardiac disease have signs and symptoms of CVD such as shortness of breath, fatigue, limitation of exercise capacity, and swelling [4].
- The people providing care to a pregnant population must become familiar with risk factors, warning signs and certain physical exam findings that suggest underlying cardiac conditions [4].
- Global cardiovascular risk assessment should be obtained in all pregnant women at their first encounter with an obstetric provider [4].



## How to differentiate common signs and symptoms of normal pregnancy versus those that are abnormal and indicative of underlying cardiac disease\*

	<b>ROUTINE CARE</b> Reassurance	<b>CAUTION</b> Nonemergent Evaluation	<b>STOP</b> Prompt Evaluation Pregnancy Heart Team
<b>HISTORY OF CVD</b>	None	None	Yes
<b>SELF-REPORTED SYMPTOMS</b>	None or mild	Yes	Yes
Shortness of Breath	No interference with activities of daily living; with heavy exertion only	With moderate exertion, new-onset asthma, persistent cough, or moderate or severe OSA	At rest; paroxysmal nocturnal dyspnea or orthopnea; bilateral chest infiltrates on CXR or refractory pneumonia
Fatigue	Mild	Mild or moderate	Extreme
<b>VITAL SIGNS</b>			
Heart Rate	<90	90-119	≥120
<b>PHYSICAL EXAMINATION</b>			
Edema	Mild	Moderate	Marked

\*This is an abbreviated list—see Practice Bulletin *Pregnancy and Heart Disease* for complete table of symptoms.



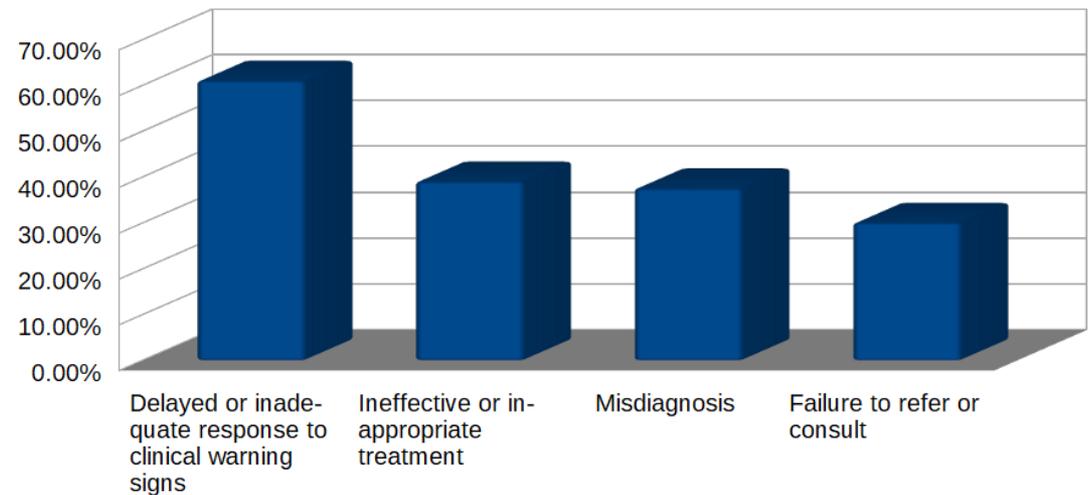
Eliminate preventable maternal mortality  
#EveryMomEveryTime



**Another contributing factor to maternal morbidity and mortality with CVD from the health care provider perspective:  
Insufficient use of hypertensive medications and misdiagnosis of “new onset asthma” or other respiratory illness [4].**

[Click here for a list of improvement opportunities.](#)

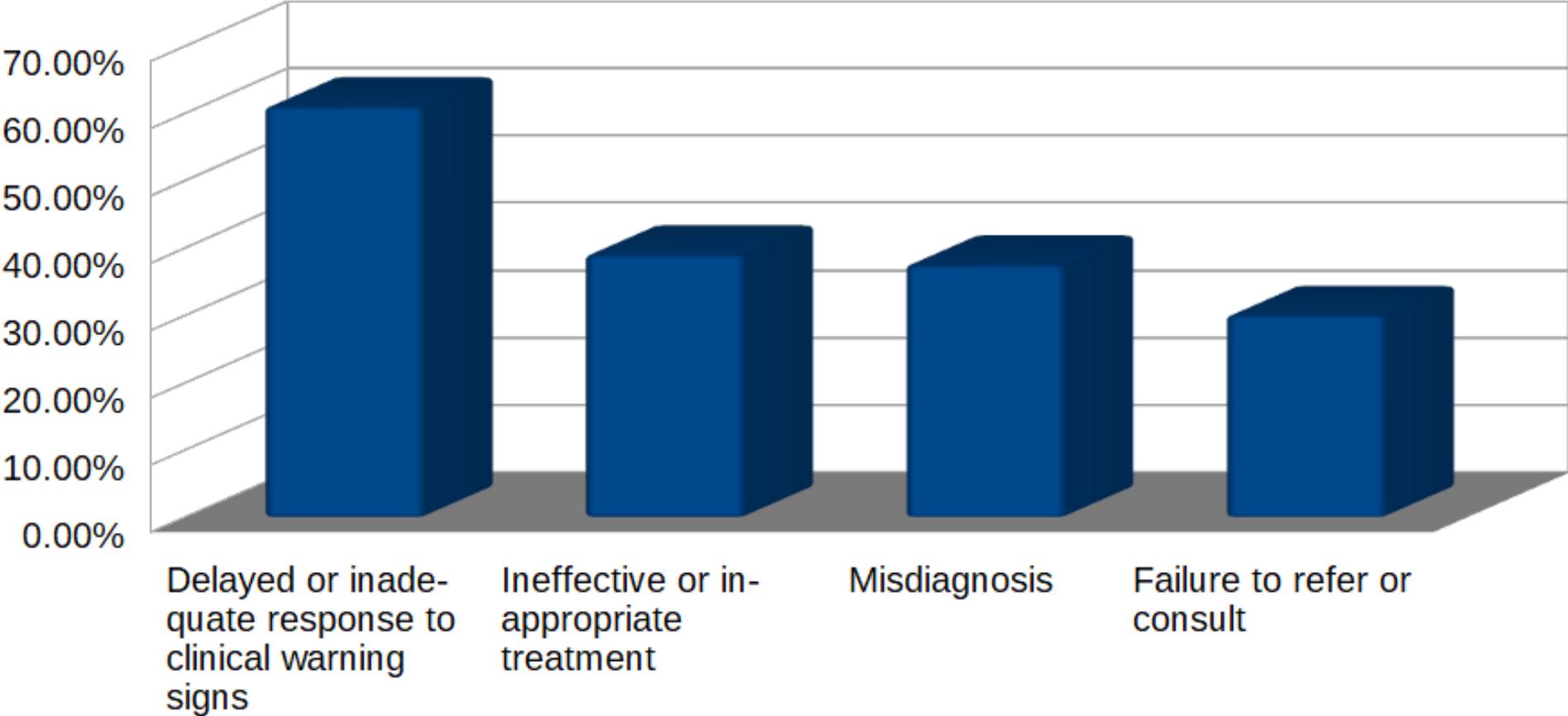
Contributing Factors (69% of all cases)



**Click the graph above for a larger**



### Contributing Factors (69% of all cases)



## Improvement opportunities [4]:

- Better recognition of signs and symptoms of CVD in pregnancy:
  - Shortness of breath, fatigue
  - Tachycardia, BP change, and low oxygen saturation
  - Improved hypertensive management



Incidence is important for careful exclusion of alternative etiologies of heart failure and paramount in both defining prevalence in the population and in caring for individual patients [31].

## Incidence of Peripartum Cardiomyopathy [30]:

- Incidence varies widely owing to numerous factors [32]:
  - Geographic difficult area
  - Differing definitions
  - Evolving diagnostic criteria
  - Lacking of accurate data



Africa and Haiti



United States



Mouse over each map above for more on cardiomyopathy incidence in each location



The incidence of peripartum cardiomyopathy in the U.S. has increased from 1 in 4,350 births in the early 1990s to 1 in 2,230 births in the mid-2000's [[31](#)].

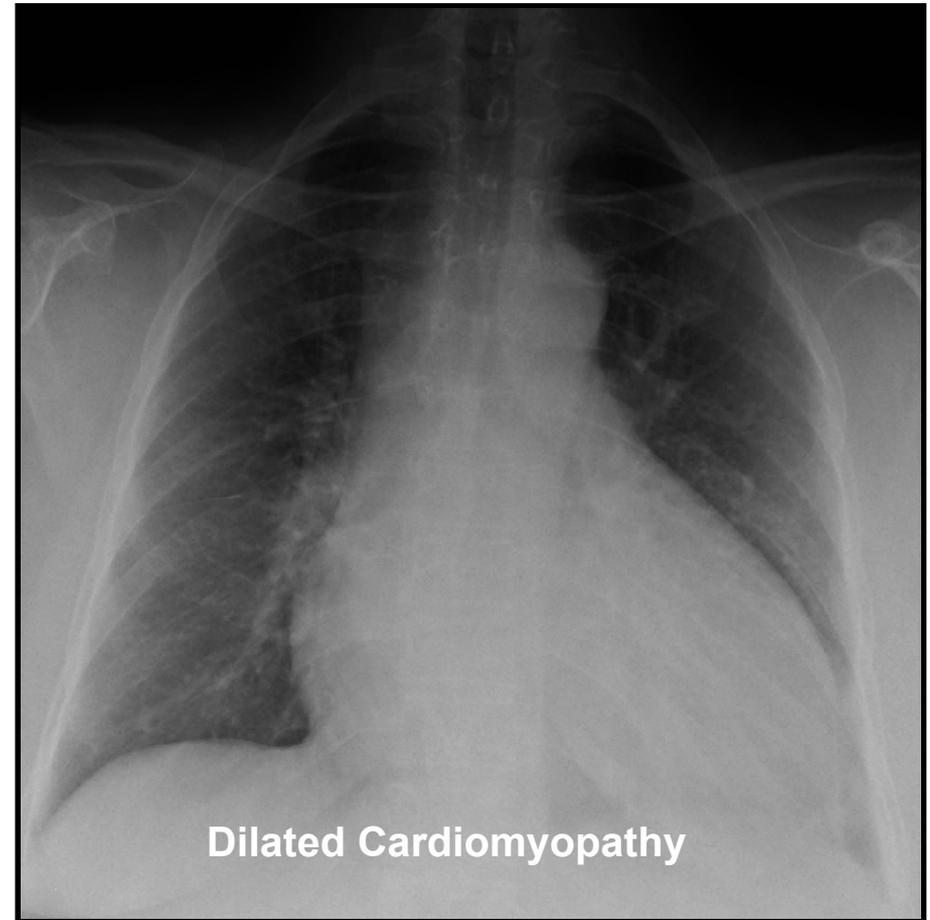
Worldwide, peripartum cardiomyopathy affects approximately 1 per 1,000 pregnancies with geographic hot spots found in Africa--up to 1 per 100 pregnancies-- and Haiti--1 in 300 births [\[30\]](#). Genetic predisposition may contribute to geographical variation.



Peripartum cardiomyopathy is defined by left ventricular dysfunction and development of cardiac failure without a known cause and occurring in the final month of pregnancy and up to 5 months postpartum [1].



Mouse over the xray for more information



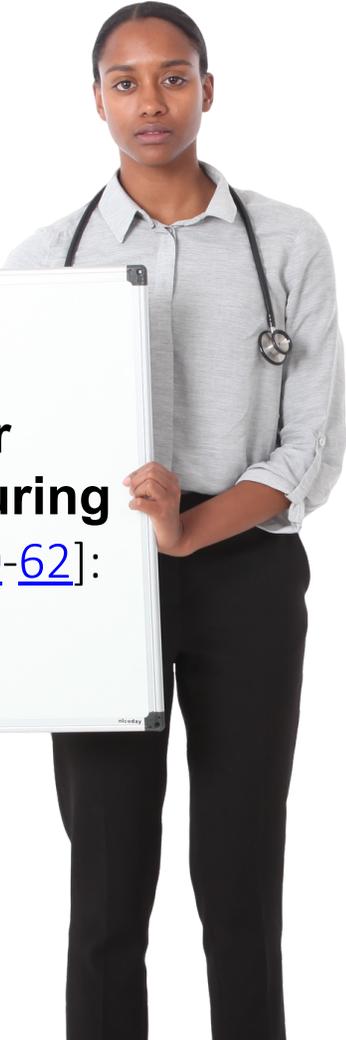
Dilated Cardiomyopathy

Defining Peripartum Cardiomyopathy



Absence of another identifiable cause for the heart failure [\[55\]](#):

- Left ventricular (LV) systolic dysfunction with an LV ejection fraction (LVEF) of less than 45% [\[55\]](#).
  - The LV may or may not be dilated [\[55\]](#).



**Risk factors for  
cardiomyopathy during  
pregnancy [[1,34](#), [60-62](#)]:**

- Multiparity
- Advanced maternal age
- Multifetal pregnancy
- Preeclampsia
- Gestational HTN
- African-American race
  - African-American's may also have more severe disease
- History of preeclampsia, eclampsia, or postpartum hypertension
- Maternal cocaine abuse
- >4 weeks of oral tocolytic therapy with beta adrenergic agonists such as terbutaline



## Review of Cardiac Physiology Related to Pregnancy

A 30-50% increase of cardiac output occurs to meet the needs of a colossal uterine blood flow and fetal perfusion while allowing the mother to function without impairment [[1,35](#)].

- Hormones secreted by the corpus luteum likely orchestrate these early cardiovascular changes [[1,35](#)].
- These hormones induce profound changes in systemic vascular resistance and result in significantly decreased systolic and diastolic blood pressures as early as 5 weeks [[1,36](#)].



Click the next arrow to review information on cardiac physiology in pregnancy.

1 of 3





## Review of Cardiac Physiology Related to Pregnancy

At least half of the total rise in cardiac output during pregnancy has manifested as early as 8 weeks gestation [37,38].

On the other hand, mean BP nadirs at 16-29 weeks gestation [37,38].

- These changes persist until the 3<sup>rd</sup> trimester [37,38].
- "From early pregnancy until about 16 weeks of gestation a reduction in peripheral vascular resistance occurs causing stroke volume to be augmented and increase after this time in gestation, typically stroke volume plateaus along with cardiac output [1]."



Click the next arrow to review information on cardiac physiology in pregnancy.



2 of 3





## Review of Cardiac Physiology Related to Pregnancy

Intrinsic left ventricular contractility does not appear to change appreciably, even with all the dramatic hemodynamic alterations that occur while pregnant.

It is uncharacteristic in a high output cardiac state with hyperdynamic function to maintain normal left ventricular function as it is in pregnancy [35].

The pregnancy induced hypervolemia involves an increase in end-systolic and end-diastolic dimensions is accommodated by cardiac atria and ventricles.

- Due to these facts, the left ventricular mass is increased proportional to maternal size.
- The cardiac atria have a similar increase in mass.

Most, but not all, agree the cardiac remodeling is a normal physiologic response resolving by 3 months postpartum.



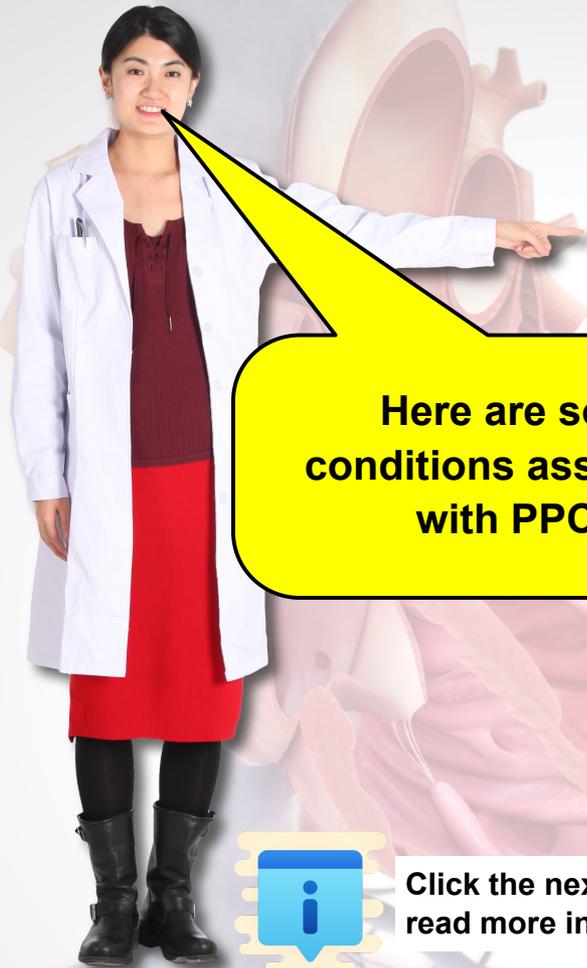
Click the next arrow to continue

◀ 3 of 3





**After reviewing normal physiologic cardiac changes, we will now discuss peripartum cardiomyopathy (PPCM) in more depth.**



Here are some conditions associated with PPCM [1].



Click the next arrows to read more information.

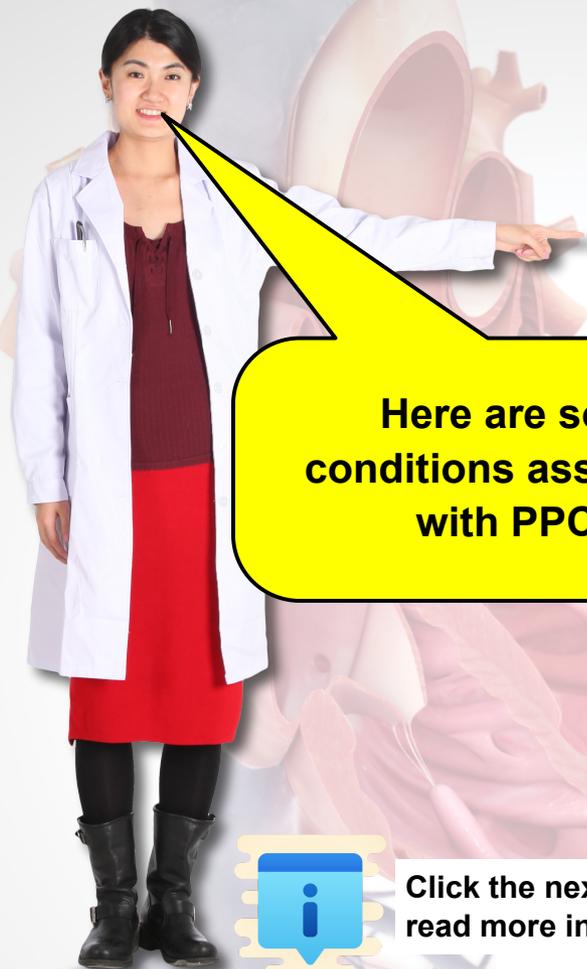
Conditions Associated with PPCM

## Advanced maternal age [39]:

- More than half of PPCM occurs in women >30 years old.
- Interestingly, the incidence of PPCM is 10-fold higher in women >40 years old compared with those <20 years old.

1 of 4 >>





Here are some conditions associated with PPCM [1].



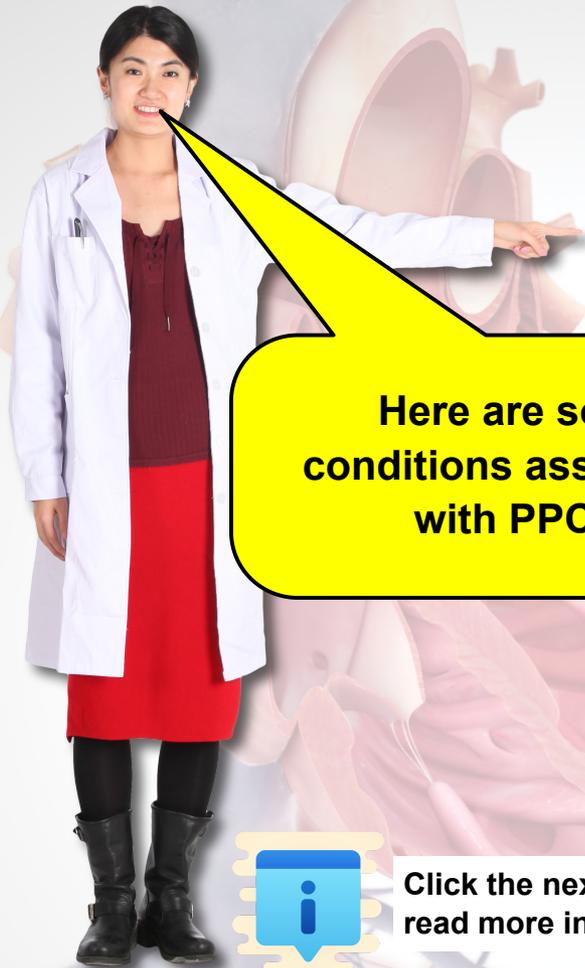
Click the next arrows to read more information.

Conditions Associated with PPCM

### Black race is strongly associated risk factor for PPCM [10]:

- This translates to the geographic propensity with incidences as high as 1% in some populations.
- In the U.S. a 5- to 15-fold increased risk for black women compared with other races has been reported.





Here are some conditions associated with PPCM [1].



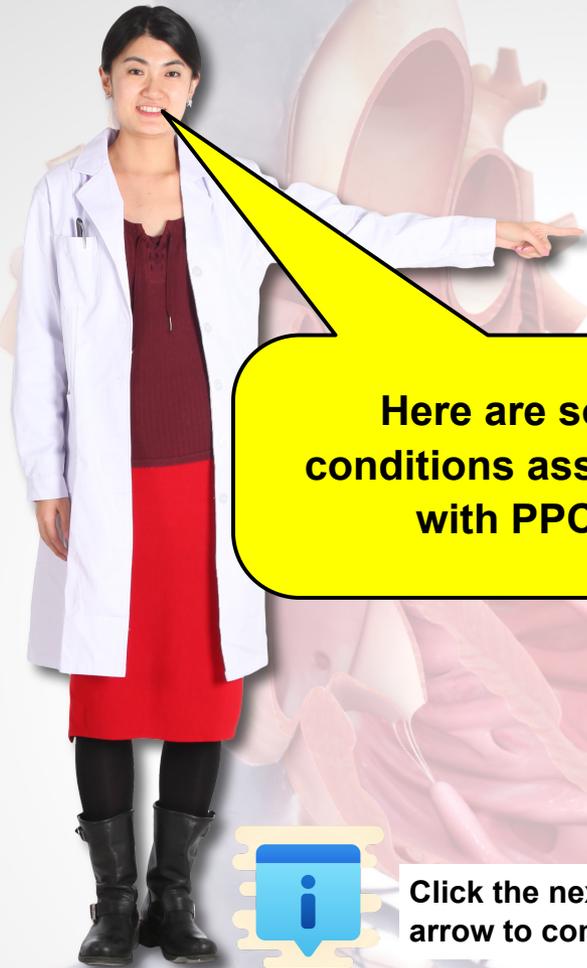
Click the next arrows to read more information.

Conditions Associated with PPCM

## Pregnancy induced hypertension (PIH)

- Particularly preeclampsia, strongly disposes patients to the development of peripartum cardiomyopathy.
- The incidence associated with hypertensive disorders increases from 5- to 30-fold.
  - Permeability pulmonary edema caused by preeclampsia can mimic cardiogenic edema caused by heart failure from peripartum cardiomyopathy.
  - Distinguishing the 2 is imperative to her outcome.





Here are some conditions associated with PPCM [1].



Click the next arrow to continue

Conditions Associated with PPCM

## Multifetal pregnancy increases her risk:

- 9% of cases of peripartum cardiomyopathy were in women with a multifetal pregnancy.

## Common with peripartum cardiomyopathy are common obstetric comorbidities:

- Obesity
- Anemia
- Infection

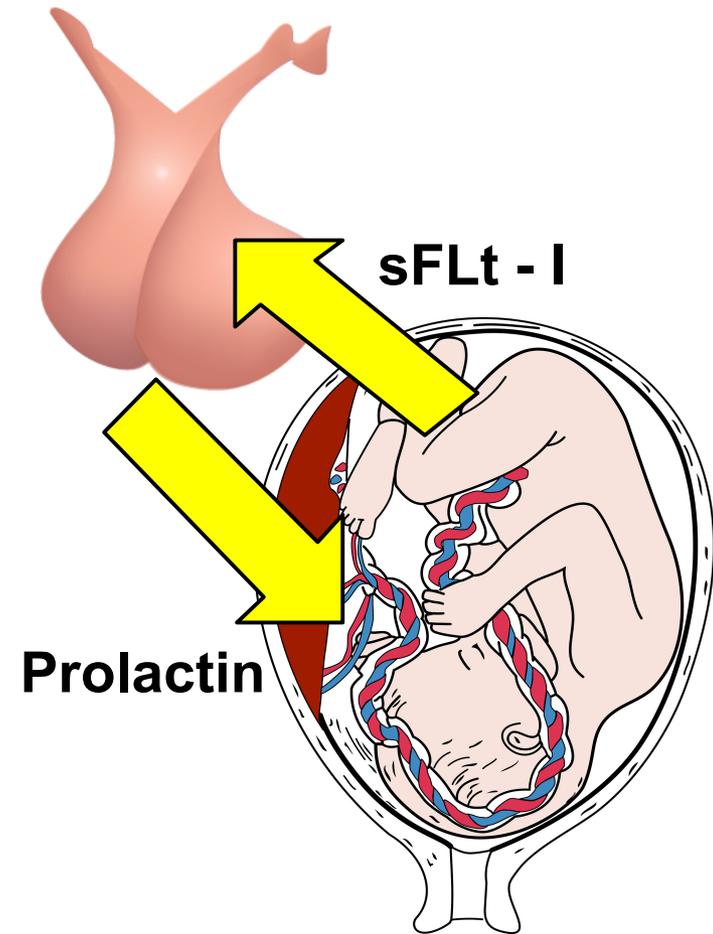


**“Two hit Hypothesis”**  
Peripartum cardiomyopathy affects genetically susceptible women who carry one of the following gene mutations [1]:

- TTNC1
- TTN
- STAT3



Click the picture to read more information.





**“Two hit Hypothesis”  
Peripartum cardiomyopathy  
affects genetically  
susceptible women who  
carry one of the following  
gene mutations [1]:**

- **TTNC1**
- **TTN**
- **STAT3**



Click the picture to  
read more information.

Term pregnancy is also characterized by prodigious secretions of prolactin by the maternal pituitary and at the same time the placenta secretes high levels of the antiangiogenic molecule, sFlt-1:

- There is a fragment of the prolactin molecule that is hypothesized to trigger events of vasoinhibin-acts which is thought to cause myocardial damage with ventricular dysfunction.
- This is made worse by secretions of high levels of vascular endothelial growth factor inhibitory molecule, sFlt-1 which is superabundant in women with preeclampsia, multifetal pregnancy, or both.
- So altered prolactin processing is believed to be involved in the pathogenesis of PPCM.



## Angiogenic Imbalance [56]:

- PPCM may be caused by systemic angiogenic imbalance which may explain why preeclampsia and multiple gestations are risk factors for PPCM.
- Angiogenic imbalances damage the vasculature leading to PPCM issues.

1 of 6 >>





## **Prolactin** [\[57\]](#):

- Altered prolactin processing is believed to be involved in the pathogenesis of PPCM.
- Alterations in prolactin processing may contribute to angiogenic imbalances.

◀◀ 2 of 6 ▶▶



## Myocarditis [58]:

- Viral genomes have been noted in research where the endomyocardium was biopsied.
- However, myocarditis is not always present with PPCM.

◀◀ 3 of 6 ▶▶





## Abnormal Immune Response [\[59\]](#):

- The maternal immunologic response to a fetal antigen has been suggested as an etiology leading to PPCM.
- If these cells lodge in the cardiac tissue, they can trigger an autoimmune response.

◀◀ 4 of 6 ▶▶



## Genetic Predisposition:

- Several studies support the hypothesis that PPCM may develop as a result of interaction between pregnancy-related factors, such as late pregnancy oxidative stress, and a susceptible genetic background.
- African genomic ancestry may be a risk factor for the development of PPCM and explain the high prevalence of PPCM in Haitian, African, and black women in the U.S.

◀◀ 5 of 6 ▶▶



## Hemodynamic Factors:

- As discussed, in pregnancy there is a 40 to 50% increase in blood volume and cardiac output resulting in left ventricular remodeling and hypertrophy. It is possible this remodeling is an exaggerated response with a decrease in left ventricular systolic function in women who develop PPCM.
- The hemodynamic stress of gestational hypertension may contribute to the development of heart failure with gestational hypertension being more common in women with PPCM.

◀◀ 6 of 6





**Next I will share with you some pearls of knowledge for providers on cardiomyopathy in pregnancy.**





## Pearls for Providers

The highest risk period for a preexisting cardiac condition to manifest is generally in the late 2nd trimester — 24-28 weeks — or in the postpartum period [4].

1 of 13 



Click the next arrows to read more information.



## Pearls for Providers

The first presentation of cardiovascular disease may occur during pregnancy or in the early postpartum time frame [\[4\]](#).

◀ 2 of 13 ▶



Click the next arrows to read more information.



## Pearls for Providers

Cardiovascular risk assessment needs to occur in all pregnant women, with or without symptoms [\[4\]](#).

◀ 3 of 13 ▶



Click the next arrows to read more information.



## Pearls for Providers

Pregnant or postpartum women presenting with symptoms of shortness of breath, cough, or excessive fatigue should be evaluated in the context of risk factors, vital sign abnormalities, and abnormal physical examination findings [[4](#)].

◀ 4 of 13 ▶



Click the next arrows to read more information.





## Pearls for Providers

A symptom of heart failure may present as persistent respiratory symptoms and 'new-onset' asthma [\[4\]](#).



5 of 13



Click the next arrows to read more information.



## Pearls for Providers

When an chest x-ray shows bilateral infiltrates, this may be due to heart failure rather than pneumonia, correlate these findings clinically [\[4\]](#).

◀ 6 of 13 ▶



Click the next arrows to read more information.



## Pearls for Providers

Counsel women currently pregnant or postpartum with significant cardiovascular risk factors regarding their future cardiovascular risk [\[4\]](#).

Involving the woman's primary care provider (PCP) early is crucial to ensure a smooth transition postpartum [\[4\]](#).

◀◀ 7 of 13 ▶▶



Click the next arrows to read more information.



## Pearls for Providers

Emphasizing awareness of risk factors, sign and symptoms of cardiac disease, and compliance with follow-up are key patient education points [\[4\]](#).

◀ 8 of 13 ▶



Click the next arrows to read more information.



## Pearls for Providers

Optimally, when a woman has known CVD, she should receive preconception and interconception care by perinatology and cardiology with a high risk pregnancy in a center with access to cardiovascular care [4].

◀ 9 of 13 ▶



Click the next arrows to read more information.



## Pearls for Providers

Education for the patient and the provider is essential [4].

◀◀ 10 of 13 ▶▶



Click the next arrows to read more information.





## Pearls for Providers

Tailoring contraception choices is a must for the type of CVD present [\[4\]](#).



11 of 13



Click the next arrows to read more information.





## Pearls for Providers

Key elements to a successful outcome includes [\[4\]](#):

- High index of suspicion
- Early diagnosis
- Appropriate referrals
- Follow up

◀ 12 of 13 ▶



Click the next arrows to read more information.



## Pearls for Providers

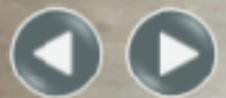
When Red Flags are present then immediate evaluation and/or hospitalization [4]:

- Shortness of breath at rest
- Severe orthopnea  $\geq 4$  pillows
- Resting HR  $\geq 120$  bpm
- Resting systolic BP  $\geq 160$  mmHg
- Resting respiratory rate  $\geq 30$
- Oxygen saturation  $\leq 94\%$  with or without personal history of CVD

◀ 13 of 13



Click the next arrow to continue



# Algorithm 1 – Red Flags Not Present, Move on to Next Algorithm



Check for red flags first

- Red Flags**
- Shortness of breath at rest
  - Severe orthopnea  $\geq 4$  pillows
  - Resting HR  $\geq 120$  bpm
  - Resting systolic BP  $\geq 160$  mm Hg
  - Resting RR  $\geq 30$
  - Oxygen saturations  $\leq 94\%$  with or without personal history of CVD

**PROMPT EVALUATION** and/or hospitalization for acute symptoms *plus* **CONSULTATIONS** with MFM and Primary Care/Cardiology

**Personal History of CVD Without Red Flags**

**CONSULTATIONS** with MFM and Primary Care/Cardiology



Click the picture for a larger view

Algorithm 1 – Red flags not present, move on to next algorithm.



### Red Flags

- Shortness of breath at rest
- Severe orthopnea  $\geq 4$  pillows
- Resting HR  $\geq 120$  bpm
- Resting systolic BP  $\geq 160$  mm Hg
- Resting RR  $\geq 30$
- Oxygen saturations  $\leq 94\%$  with or without personal history of CVD



**PROMPT EVALUATION** and/or hospitalization for acute symptoms  
*plus*  
**CONSULTATIONS** with MFM and Primary Care/Cardiology

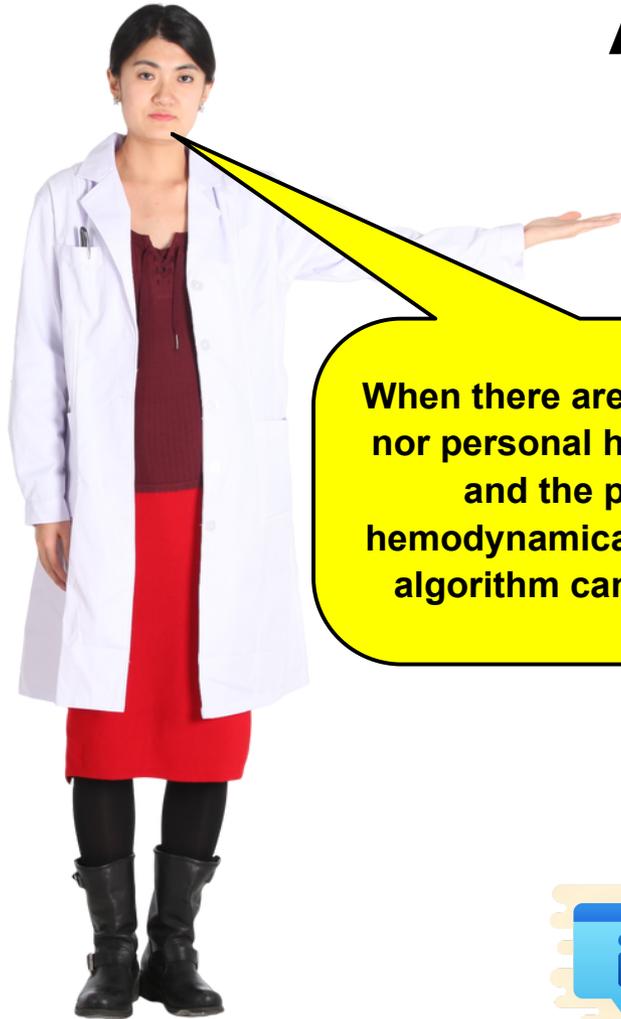
**Personal History of CVD Without Red Flags**



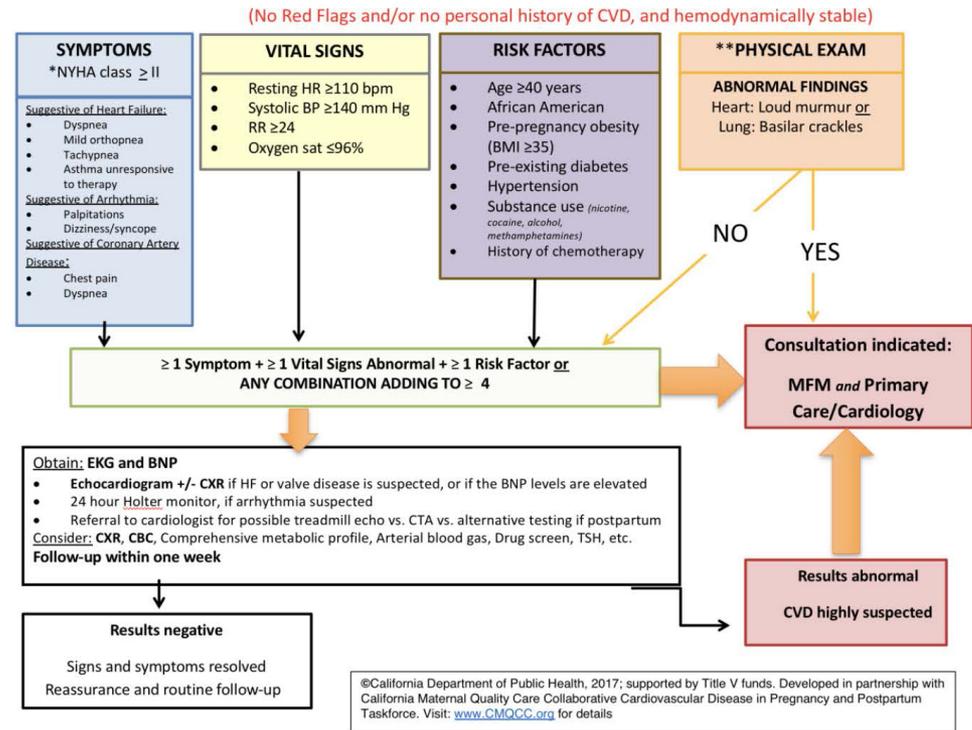
**CONSULTATIONS** with MFM and Primary Care/Cardiology

[4]

# Algorithm 2 Assessment



**When there are no Red Flags nor personal history of CVD and the patient is hemodynamically stable this algorithm can be utilized:**

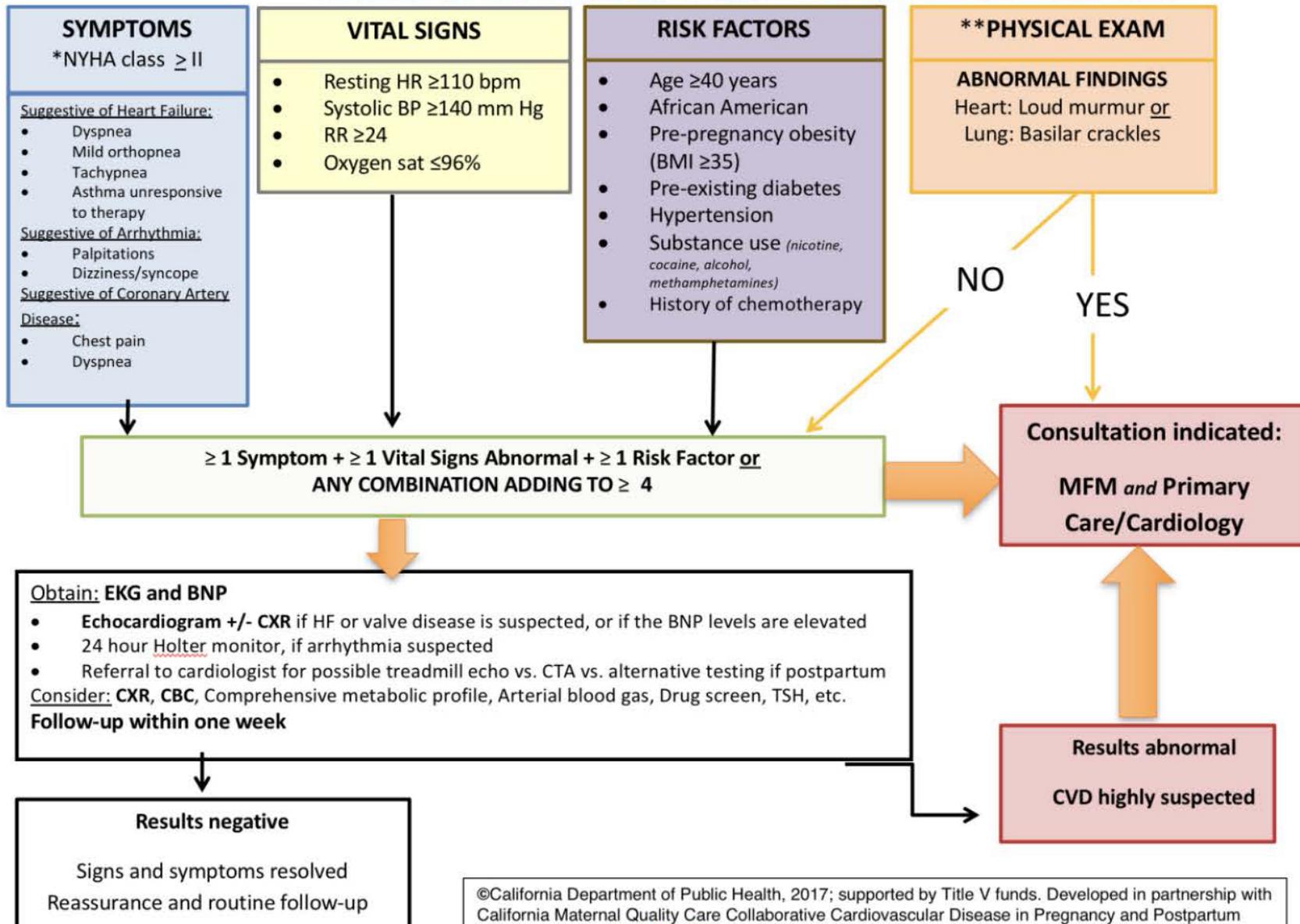


Click the picture for a larger view

Algorithm 2 Assessment



(No Red Flags and/or no personal history of CVD, and hemodynamically stable)



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**B-type Natriuretic Peptide (BNP) is a tool to help identify asymptomatic women with left ventricular dysfunction [4].**

BNP is a B-type Natriuretic Peptide:

- BNP can also be utilized to triage patients presenting with symptoms in need of further diagnostic test [4].

1 of 9 >>



Click the next arrows to read more information.





**B-type Natriuretic Peptide (BNP) is a tool to help identify asymptomatic women with left ventricular dysfunction [4].**

◀ 2 of 9 ▶



Click the next arrows to read more information.

- BNP is a neurohormone secreted predominantly by the cardiac ventricles in response to volume expansion or pressure overload.
- BNP acts as the body's defense against volume overload by virtue of its vasodilatory and renin-angiotensin-aldosterone system inhibitory properties that lead to natriuresis and diuresis.





**B-type Natriuretic Peptide (BNP) is a tool to help identify asymptomatic women with left ventricular dysfunction [4].**

◀ 3 of 9 ▶



Click the next arrows to read more information.

### **BNP Facts [4,48]:**

- Half-life is 20 minutes
- BNP level <100pg/mL is considered normal
- Women tend to have higher levels than men
- Known to be elevated in patients with renal insufficiency or renal failure
- Obesity is associated with lower plasma BNP compared to non-obese
- BNP level under 50pg/mL has a negative predictive value of 96% in excluding heart failure





**B-type Natriuretic Peptide (BNP) is a tool to help identify asymptomatic women with left ventricular dysfunction [4].**

◀ 4 of 9 ▶



Click the next arrows to read more information.

## **Asymptomatic Left Ventricular Dysfunction [4, 49]:**

- When a BNP level of 50pg/mL is used as a cutoff, it has been shown to detect asymptomatic left ventricular dysfunction with sensitivity of 88% and specificity of 67%.
- In this situation, BNP may be used as an initial low-cost modality to identify asymptomatic high risk individuals who would need further testing.





**B-type Natriuretic Peptide (BNP) is a tool to help identify asymptomatic women with left ventricular dysfunction [4].**

◀ 5 of 9 ▶



Click the next arrows to read more information.

### **Predictor of Adverse Cardiovascular Outcomes in an Older Population [50,51]:**

- The strongest predictor of serious *adverse* cardiovascular outcomes is a BNP level >50 pg/mL in the elderly.
- BNP level correlates directly with left ventricular hypertrophy and is increased in diastolic left ventricular dysfunction.





**B-type Natriuretic Peptide (BNP) is a tool to help identify asymptomatic women with left ventricular dysfunction [4].**

◀ 6 of 9 ▶



Click the next arrows to read more information.

## BNP and Pregnancy:

- BNP levels remain stable throughout the pregnancy and postpartum period, despite an increase in left ventricular wall mass and end-diastolic dimensions during normal pregnancy [52].

[Click here for more information on BNP and pregnancy.](#)





**B-type Natriuretic Peptide (BNP) is a tool to help identify asymptomatic women with left ventricular dysfunction [4].**

◀ 6 of 9 ▶



Click the next arrows to read more information.

### **BNP Facts in Pregnancy [52]:**

- The median level of BNP was noted to be 19pg/mL during pregnancy versus 10pg/mL in the non-pregnant state from a longitudinal study group of plasma levels.
- BNP levels stay well within the normal range during uncomplicated pregnancy.
- Significant elevations are seen in patients with hypertensive disorders including preeclampsia.





**B-type Natriuretic Peptide (BNP) is a tool to help identify asymptomatic women with left ventricular dysfunction [4].**

◀◀ 7 of 9 ▶▶



Click the next arrows to read more information.

### Preexisting Heart Disease:

- Serial measurements of N-terminal pro-BNP (NT-proBNP) are shown to be predictive of adverse cardiovascular outcomes in women with *preexisting* dilated cardiomyopathy [29].
- Another study, with 66 women having cardiac symptoms, all remained event-free during pregnancy and had BNP < 100 pg/mL [53].





**B-type Natriuretic Peptide (BNP) is a tool to help identify asymptomatic women with left ventricular dysfunction [4].**

◀ 8 of 9 ▶



Click the next arrows to read more information.

## **Pregnant Women with Symptoms [54]:**

- BNP plays an important role for pregnant women presenting with cardiac symptoms such as shortness of breath:
  - BNP levels correlate with elevated left ventricular filling pressures in symptomatic pregnant women.
  - Helping to determine both systolic and diastolic left ventricular dysfunction.





**B-type Natriuretic Peptide (BNP) is a tool to help identify asymptomatic women with left ventricular dysfunction [4].**

### **In Summary [4]:**

- BNP may assist clinicians in triaging patients who present with symptoms for further diagnostic testing:
  - Simple test
  - Relatively inexpensive
  - Readily available

◀ 9 of 9

[Click here for more information](#)



Click the next arrow to continue





**B-type Natriuretic Peptide (BNP) is a tool to help identify asymptomatic women with left ventricular dysfunction [4].**



9 of 9

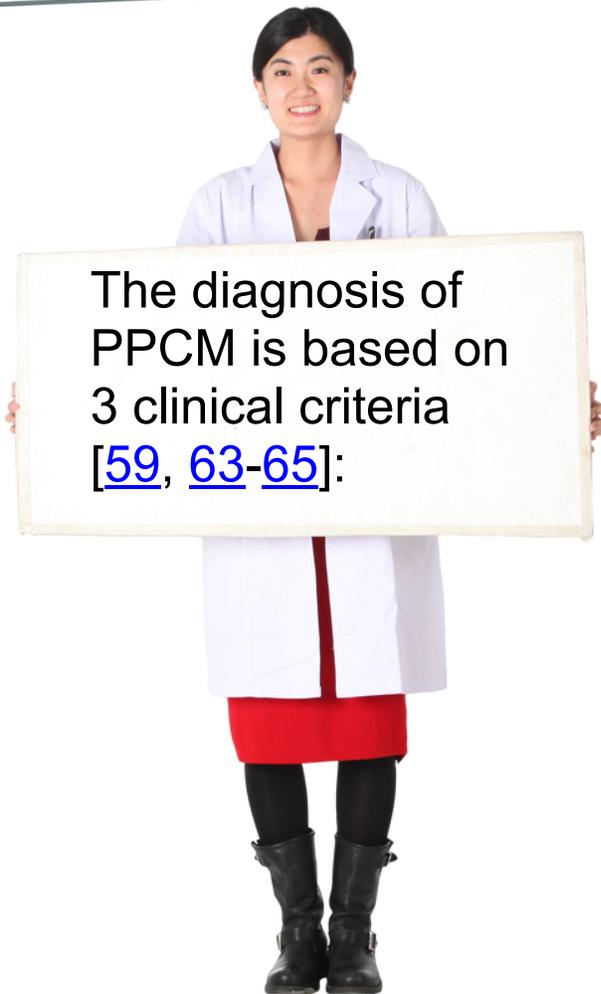
## Summary Continued

- Many pregnant women have some degree of fatigue, shortness of breath, palpitations, and/or edema during pregnancy. BNP added to routine evaluation in cases with symptoms out of proportion to pregnancy will be a useful tool in triaging.
- Certainly, the women presenting with symptoms suggestive of cardiac disease, evaluating the BNP may reduce potential morbidity.



Click the next arrow to continue





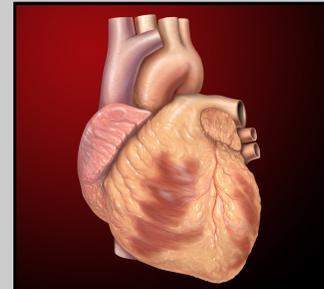
The diagnosis of PPCM is based on 3 clinical criteria [[59](#), [63-65](#)]:

1. Development of heart failure toward the end of pregnancy or in the months following delivery.
2. Absence of another identifiable cause of heart failure.
3. Left ventricular systolic dysfunction with a left ventricular ejection fraction generally  $<45\%$ .

Directing investigations to a timely diagnosis and treatment is necessary.

## Studies

- There are no pathognomonic findings in PPCM and is a diagnosis of exclusion.
- A variable portion of patients have evidence of myocarditis.
- Ordering an electrocardiogram (ECG) and echocardiogram are prudent actions in those suspected of having PPCM.
  - A normal ECG does not rule out PPCM [66].
  - An ECG may be helpful in differentiating between other diagnosis such as a myocardial infarction (MI) or pulmonary embolism (PE).
  - An echocardiogram generally reveals global reduction in left ventricular function with LVEF nearly always  $<45\%$ ; the left ventricle is frequently dilated, but not always.



Click the heart for more information.



## Studies

- BNP, chest X-ray (with fetal shielding when pregnant), cardiac magnetic resonance (CMR), cardiac catheterization and even endomyocardial biopsy (EMB) may be helpful in certain cases.
- BNP can help evaluate patients suspected of having heart failure when the diagnosis is uncertain with higher levels noted than their healthy female counterparts [[67](#)].





## Summary

Preexisting cardiovascular disease and/or new-onset peripartum cardiomyopathy may initially present during pregnancy or in the postpartum period [4].

- By 2 weeks postpartum, the physiologic changes associated with pregnancy gradually return to baseline [55].
- Mostly, women with peripartum cardiomyopathy present in the 1st week postpartum with 75% of women with peripartum cardiomyopathy present in the 1st month [56].
- New-onset cough or shortness of breath are the most frequent complaints of women presenting with pregnant or postpartum women with CVD [6].
- A high index of suspicion for underlying CVD needs to occur by all providers (emergency, primary care and obstetrical providers) when a woman presents with symptoms, signs and risk factors concerning for heart disease for up to 5 months postpartum.





# Summary

Be certain to ask the following questions when a woman presents in the postpartum period with complaints of shortness of breath [4]:

- Intolerance of exercise
- Activities of daily living are difficult to complete
- Deteriorating symptoms
- Dizziness
- Palpitations
- Chest pain
- New-onset cough
- Wheezing
- Lower extremity edema or pedal edema
- Fatigue that is not expected (i.e. frequent stops while out walking)
- Orthopnea (if present, ask how many pillows are used)
- Weight gain or inability to lose weight
- History of cardiac or pulmonary conditions
- History of tobacco and/or substance abuse
- Has received care by other providers or Emergency Departments (ED) during the postpartum period





## Summary

Common symptoms of pregnancy may be falsely attributed, failing to recognize cardiac disease [\[4\]](#):

- Shortness of breath
- Fatigue
- Edema





## Summary

### Differential diagnosis for postpartum dyspnea [4]:

- Congestive heart failure (CHF)
- Myocarditis
- Endocarditis
- PE
- Pulmonary hypertension
- Asthma
- Infection





## Summary

### Key Points [4]:

- The postpartum period should provide improving symptoms related to physiologic changes of pregnancy.
- ED visits for dyspnea need to raise suspicion for CVD.
  - The women with these symptoms, of child bearing age, should be questioned about recent pregnancies and when her last menstrual period (LMP) occurred.
- Concerns for CVD should be raised when postpartum dyspnea or new-onset cough is present.





## Summary

Physical examination should pay particular attention to [\[4\]](#):

- The vital signs (and investigate for underlying causes needing further tests):
  - HR  $\geq 120$  bpm
  - BP  $\geq 160$  mm Hg
  - RR  $\geq 30$
  - Oxygen saturation ( $O_2$  Sat)  $\leq 94\%$
- Lung exam:
  - Crackles
  - Wheezing
- Cardiac exam:
  - Loud murmur
  - Jugular venous distention (JVD)
- Extremities:
  - Edema
  - Taunt shiny skin





## Summary

Workup considerations for postpartum dyspnea [\[4\]](#):

- Chest Radiograph
  - Frequently normal in asthma
- EKG
  - May be normal in cardiomyopathy, except for sinus tachycardia
  - Should be obtained on an emergency basis if the patient has abnormal vital signs or is very symptomatic
    - Normal LV ejection fraction does not exclude heart failure
    - Normal RV function does not exclude pulmonary embolism





## Summary

Workup considerations for postpartum dyspnea [4]:

- Labs
  - CBC
  - Basic Metabolic Panel (BMP)
  - Thyroid Function Test
  - BNP - An elevated BNP should raise suspicion for CHF
  - D-dimer - May normally be elevated in pregnancy, however, may be considered for negative predictive value
  - Toxicology screen - Substance use (e.g., meth or cocaine) is a strong risk factor for pregnancy-related CVD
- Venous doppler ultrasound and/or computed tomography (CT) pulmonary angiogram for PE
- Cardiology consultation as needed





## Summary

Key points [4]:

- New-onset asthma is rare in adults.
- Bilateral crackles on lung examination are most likely associated with CHF.
  - Improvement of dyspnea with bronchodilators does not confirm the diagnosis of asthma, as CHF may also improve with bronchodilators.
  - Response to bronchodilators should prompt the consideration of a diagnosis other than asthma.





## Summary

### Disposition:

- If considering discharge:
  - Repeat vital signs to ensure they are persistently normal, the symptoms have improved, and the patient is stable for discharge.
  - Arrange for early follow-up with PCP or cardiologist as indicated.
- Admission and cardiology consultation may be indicated for:
  - Persistent symptoms or abnormal vital signs, in particular, HR >120 bpm, BP >160 mmHg, RR >30, and O<sub>2</sub> Sat <94%.
  - Lack of response to treatment.
  - Newly-diagnosed cardiomyopathy or pulmonary hypertension.



## DID YOU HAVE COMPLICATIONS DURING PREGNANCY?

You may be at a higher risk for heart disease over your lifetime

Which pregnancy complications can increase your risk for heart disease as you age?



### HIGH BLOOD PRESSURE

5-10% of all pregnant women



### GESTATIONAL DIABETES

7-14% of all pregnancies



### PRETERM BIRTH

11.5% of babies were born preterm in 2012.

**Can include:**

- ♥ Gestational hypertension
- ♥ Preeclampsia once known as Pregnancy Induced Hypertension (PIH) and Toxemia
- ♥ Eclampsia
- ♥ HELLP syndrome

**!** Mothers who had gestational diabetes are more likely to have the condition again in a future pregnancy.

**i** Babies born before 37 completed weeks of pregnancy are preterm, or premature.

If you had **PREECLAMPSIA**, you have **2x** the risk of **stroke, heart muscle damage, or blood clot** and **4x** the risk of developing **high blood pressure** for the rest of your life!

If you had **GESTATIONAL DIABETES**, you are **50%** more likely to develop **Type II diabetes** within 5 years, putting you at **higher risk** for heart disease.

Women with **PRETERM BIRTH AND PREECLAMPSIA** have an **8-10x** higher chance of **death** from heart disease.

If you had complications in pregnancy, you can lower your risk:

#### New Mothers

- See your health care provider 3-6 months after birth to check your overall physical health. Discuss your pregnancy and any complications you experienced.
- Get a copy of your pregnancy and post-delivery medical records to share with your providers for the rest of your life. Don't wait – records may be destroyed.
- Breastfeed as long as possible. Women whose total lifetime breastfeeding is 6-12 months were 10% less likely to develop heart disease (and it's good for baby too).

If you had one of these complications, speak with your provider when planning your next pregnancy to optimize your health.



It's a **MYTH** that **ALL** pregnancy related high blood pressure and gestational diabetes complications go away after the baby is born!

Get more information and stay heart healthy.  
[www.cmqcc.org](http://www.cmqcc.org)

#### Mothers With Kids Over One Year

- Get annual checkups and be screened for heart disease. At this visit, your provider should check your overall physical condition.
- Ask your provider what your test results mean and how you can lower your heart disease risk.
 

These screening numbers show desirable results:	Blood Pressure < 120/80 mm hg	Fasting Blood Glucose < 100 mg/dl
	Total Cholesterol < 200 mg/dl	Body Mass Index < 25 kg/m <sup>2</sup>
- Try a mobile app to automatically retrieve and store your medical records, so you always have them handy.
- Eat healthy! A diet low in salt, fat, cholesterol and sugar can help you lower your risk for obesity, diabetes and heart disease.
- Maintain a healthy weight. Body Mass Index (BMI) is an estimate of body fat based on height and weight. Less than 25 is healthy.
- Get active for 30 minutes a day, or as recommended by your provider.
- If you smoke, make a plan to quit. Your provider may have resources to support you.
- Take medications as directed. Sometimes a healthy diet and exercise is not enough to lower your risk for heart disease, so your provider may prescribe medications to help.



Click the picture to download this infographic



Click the picture to download this infographic

## Signs & Symptoms of Heart Disease

Heart disease is the leading cause of death among women in the U.S. who are pregnant or gave birth in the last 5 months (postpartum).

*During Pregnancy and Postpartum*

**Symptoms to watch for in late pregnancy and up to five months postpartum:**



**NOTE:** While some of these symptoms are common in late pregnancy, they may be a sign of heart disease especially if they are severe and do not go away after treatment.

**If you have any of these symptoms and they don't go away:**

- ♥ Contact your OB, midwife, family medicine doctor, or your primary care provider
- ♥ Describe your symptoms clearly and explain how sick you feel
- ♥ If your symptoms arise postpartum, be sure to tell the provider that you recently had a baby
- ♥ If your provider says your symptoms are normal, ask what symptoms should cause you to call or come back

**Go to the Emergency Department**

If you have persistent chest pain or severe shortness of breath, or otherwise feel extremely sick. If possible, take someone with you.

**Any woman can develop heart disease in pregnancy or postpartum, but you are at higher risk if you:**

- ♥ Have prior heart disease
- ♥ Are over 40 years old
- ♥ Have preeclampsia or high blood pressure (hypertension)
- ♥ Are African-American (4X greater risk and 8-10X more likely to die of heart disease)
- ♥ Are obese



### Bottom line

- \* Trust your instincts when you feel something is wrong
- \* When you see a healthcare provider, bring your partner, friend or family member who can support you and help explain these symptoms are not normal for you
- \* Seek a second opinion if you don't feel listened to or your symptoms are not taken seriously

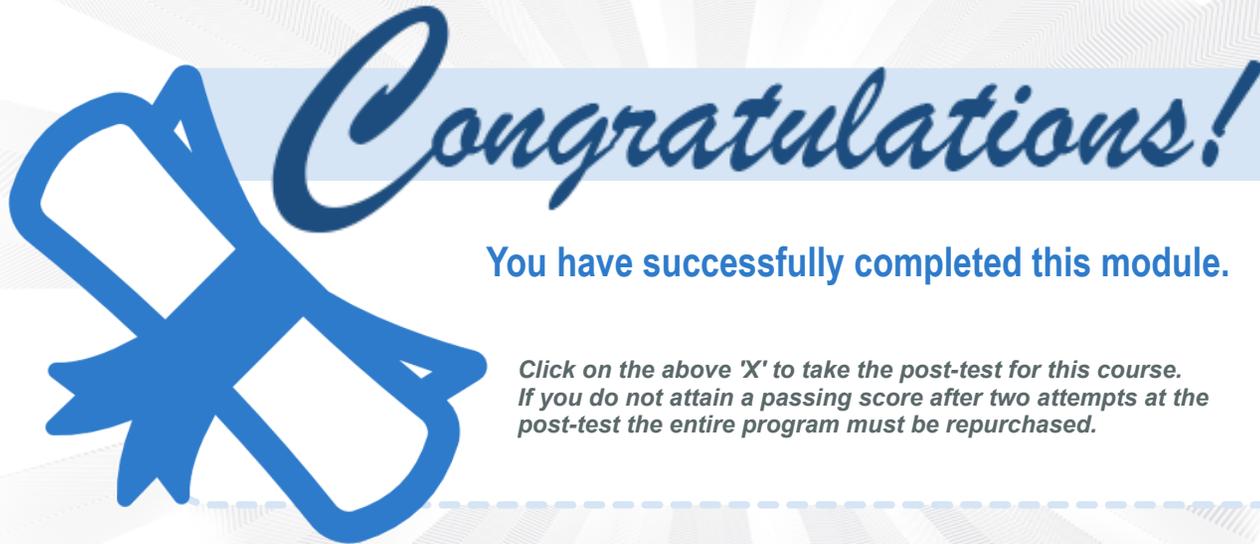
Get online support and information: [www.myheartsisters.com](http://www.myheartsisters.com) | [www.womenheart.org](http://www.womenheart.org)

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QUALITY CARE COLLABORATIVE  
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Funding for the development of this Infographic was provided by Federal Title V MCH block grant funding from the California Department of Public Health, Maternal Child Adolescent Health Division, and Stanford University.





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*Click on the above 'X' to take the post-test for this course.  
If you do not attain a passing score after two attempts at the  
post-test the entire program must be repurchased.*

## References

1. Cunningham, F.G., Byrne, J. Nelson, D. Peripartum Cardiomyopathy. *Maternal Morbidity and Mortality: Clinical Expert Series. Obstet Gynecol* Volume 133 No 1 January 2019.
2. Schaufelberger M. *Heart* 2019;105:1543–1551. doi:10.1136/heartjnl-2018-313476.
3. Creanga AA, Berg CJ, Syverson C, Seed K, Bruce FC, Callaghan WM. Pregnancy-related mortality in the United States, 2006-2010. *Obstet Gynecol.* 2015;125(1):5-12.
4. Afshan B. Hameed, Christine H. Morton, and Allana Moore. Improving Health Care Response to Cardiovascular Disease in Pregnancy and Postpartum Developed under contract #11-10006 with the California Department of Public Health, Maternal, Child and Adolescent Health Division. Published by the California Department of Public Health, 2017.
5. Creanga AA, Berg CJ, Syverson C, Seed K, Bruce FC, Callaghan WM. Pregnancy-related mortality in the United States, 2006-2010. *Obstet Gynecol.* 2015;125(1):5-12.
6. Hameed A, Lawton E, McCain C, et al. Pregnancy-related cardiovascular deaths in California: Beyond peripartum cardiomyopathy. *American Journal of Obstetrics and Gynecology.* 2015.
7. Harper M, Dugan E, Espeland M, Martinez-Borges A, McQuellon C. Why African-American women are at greater risk for pregnancy-related death. *Annals of Epidemiology.* 2007;17(3):180-185.
8. Williams RA. Cardiovascular disease in African American women: a health care disparities issue. *J Natl Med Assoc.* 2009;101(6):536-540.
9. Fong A, Lovell S, Gabby L, Pan D, Ogunyemi D, Hameed A. Peripartum cardiomyopathy: demographics, antenatal factors, and a strong association with hypertensive disorders. *American Journal of Obstetrics and Gynecology.* 2014;210(1):S136.
10. Gentry MB, Dias JK, Luis A, Patel R, Thornton J, Reed GL. African-American women have a higher risk for developing peripartum cardiomyopathy. *J Am Coll Cardiol.* 2010;55(7):654-659.
11. Goland S, Modi K, Hatamizadeh P, Elkayam U. Differences in clinical profile of African-American women with peripartum cardiomyopathy in the United States. *J Card Fail.* 2013;19(4):214-218.
12. Creanga AA, Bateman BT, Kuklina EV, Callaghan WM. Racial and ethnic disparities in severe maternal morbidity: a multistate analysis, 2008-2010. *Am J Obstet Gynecol.* 2014;210(5):435 e431-438.
13. Tanaka M, Jaamaa, G., Kaiser, M., Hills, E., Soim, A., Zhu, M., Shcherbatykh, I., Samelson, R., Bell, E., Zdeb, M., McNutt, L. Racial Disparity in Hypertensive Disorders of Pregnancy in New York State: A 10-Year longitudinal Population- Based Study. *American Journal of Public Health.* 2007;97(1):163-170.
14. Dominguez TP. Race, racism, and racial disparities in adverse birth outcomes. *Clinical Obstetrics and Gynecology.* 2008;51(2):360-370.
15. Dominguez TP. Adverse birth outcomes in African American women: the social context of persistent reproductive disadvantage. *Soc Work Public Health.* 2011;26(1):3-16.
16. Francis LE, Berger CS, Giardini M, Steinman C, Kim K. Pregnant and poor in the suburb: The experiences of economically disadvantaged women of color with prenatal services in a wealthy suburban county. *Journal of Sociology and Social Welfare.* 2009;36(2):133-157.
17. Nuru-Jeter A, Dominguez TP, Hammond WP, et al. "It's the skin you're in": African-American women talk about their experiences of racism: An exploratory study to develop measures of racism for birth outcome studies. *Maternal and Child Health Journal.* 2009;13(1):29-39.

18. Wyatt SB, Williams, D.R., Calvin, R., Henderson F.C., Walker E.R., Winters, K. Racism and Cardiovascular Disease in African Americans. *The American Journal of the Medical Sciences*. 2003;325(6):315-331.
19. Cox KJ. Midwifery and Health Disparities: Theories and Intersections. *Journal of Midwifery & Womens Health*. 2009;54(1):57-64.
20. Malat J. Expanding research on the racial disparity in medical treatment with ideas from sociology. *Health (London)*. 2006;10(3):303-321.
21. Wheatley RR, Kelley MA, Peacock N, Delgado J. Women's narratives on quality in prenatal care: a multicultural perspective. *Qual Health Res*. 2008;18(11):1586-1598.
22. Sabin J, Nosek BA, Greenwald A, Rivara FP. Physicians' implicit and explicit attitudes about race by MD race, ethnicity, and gender. *Journal of health care for the poor and underserved*. 2009;20(3):896-913.
23. Lewey J, Choudhry NK. The current state of ethnic and racial disparities in cardiovascular care: lessons from the past and opportunities for the future. *Curr Cardiol Rep*. 2014;16(10):530.
24. Hameed A. How not to miss pregnancy related cardiomyopathy? *International Journal of Cardiovascular Research* 2013;2(5).
25. Reiner T, Sonicki Z, Tedeschi-Reiner E. Physicians' perception, knowledge and awareness of cardiovascular risk factors and adherence to prevention guidelines: The PERCRO-DOC survey. *Atherosclerosis*. 2010;213(2):598-603.
26. Clarke AR, Goddu AP, Nocon RS, et al. Thirty years of disparities intervention research: what are we doing to close racial and ethnic gaps in health care? *Med Care*. 2013;51(11):1020-1026.
27. Wei T, Zeng C, Chen L, et al. Systolic and diastolic heart failure are associated with different plasma levels of B-type natriuretic peptide. *Int J Clin Pract*. 2005;59(8):891-894.
28. Grewal J, McKelvie R, Lonn E, et al. BNP and NT-proBNP predict echocardiographic severity of diastolic dysfunction. *Eur J Heart Fail*. 2008;10(3):252-259.3.
29. Blatt A, Svirski R, Morawsky G, et al. Short and long-term outcome of pregnant women with preexisting dilated cardiomyopathy: an NTproBNP and echocardiography-guided study. *Isr Med Assoc J*. 2010;12(10):613-616.
30. Pfeffer TJ, Hilfiker-Kleiner D. Pregnancy and heart disease: pregnancy-associated hypertension and peripartum cardiomyopathy. *Curr Probl Cardiol* 2018;43:364–88.
31. Mielniczuk LM, Williams L, Davis DR, Tang AS, Lemery R, Green MS, et al. Frequency of peripartum cardiomyopathy. *Am J Cardiol* 2006;97:1765–8.
32. Maternal Morbidity and Mortality: Clinical Expert Series Peripartum Cardiomyopathy F. Gary Cunningham, MD, John J. Byrne, MD, MPH, and David B. Nelson, M (Obstet Gynecol 2019;133:167–79).
33. <https://twitter.com/acog/status/1154455817690779650>
34. Demakis JG, Rahimtoola SH, Sutton GC. et al. Natural course of peripartum cardiomyopathy. *Circulation*.1971;44:1053-1061.
35. Hibbard JU, Shroff SG, Cunningham FG. Cardiovascular alterations in normal and preeclamptic pregnancies. In: Taylor RN, Roberts JM, Cunningham FG, editors. *Chesley's hypertensive disorders in pregnancy*. 4th ed. Amsterdam (the Netherlands): Academic Press; 2014.
36. ACOG Practice Bulletin No. 212. American College of Obstetricians and Gynecologists. *Obstet Gynecol* 2019;133:e320–56.

37. Simpson LL. Maternal cardiac disease: update for the clinician. *Obstet Gynecol* 2012;119:345–59.
38. Clapp JF 3rd, Capeless E. Cardiovascular function before, during, and after the first and subsequent pregnancies. *Am J Cardiol* 1997;80:1469–73.
39. Arany Z, Elkayam U. Peripartum cardiomyopathy. *Circulation* 2016;133:1397–409.
40. Martin RB, Nelson DB, Stewart R, Matulevicius S, McIntire DD, Cunningham FG. Impact of pregnancy on maternal cardiac atria. *Am J Perinatol* 2017;34:1097–101.
41. Ducas RA, Elliott JE, Melnyk SF, Premecz S, DaSilva M, Cleverly K, et al. Cardiovascular magnetic resonance in pregnancy: insights from the cardiac hemodynamic imaging and remodeling in pregnancy (CHIRP) study. *J Cardiovasc Magn Reson* 2014;16:1.
42. Stewart RD, Nelson DB, Matulevicius SA, Morgan JL, McIntire DD, Drazner MH, et al. Cardiac magnetic resonance imaging to assess the impact of maternal habitus on cardiac remodeling during pregnancy. *Am J Obstet Gynecol* 2016;214:640.e1–6.
43. Savu O, Jurcut, R, Guisca S, van Mieghem T, Gussi I, Popescu BA, et al. Morphological and functional adaptation of the maternal heart during pregnancy. *Circ Cardiovasc Imaging* 2012;5:289–97.
44. Clark SL, Cotton DB, Lee W, Bishop C, Hill T, Southwick J, et al. Central hemodynamic assessment of normal term pregnancy. *Am J Obstet Gynecol* 1989;161:1439–42.
45. Lindley KJ, Conner SN, Cahill AG, Novak E, Mann DL. Impact of preeclampsia on clinical and functional outcomes in women with peripartum cardiomyopathy. *Circ Heart Fail* 2017;10. pii: e003797.
46. Kolte D, Khera S, Aronow WS, Palaniswamy C, Mujib M, Ahn C, et al. Temporal trends in incidence and outcomes of peripartum cardiomyopathy in the United States: a nationwide population-based study. *J Am Heart Assoc* 2014;3:e001056.
47. Cunningham FG, Pritchard JA, Hankins GD, Anderson PL, Lucas MJ, Armstrong KF. Peripartum heart failure: idiopathic cardiomyopathy or compounding cardiovascular events? *Obstet Gynecol* 1986;67:157–68.
48. Maisel A, Mueller C, Adams K, Jr., et al. State of the art: using natriuretic peptide levels in clinical practice. *Eur J Heart Fail*. 2008;10(9):824-839.
49. Macabasco-O'Connell A, Meymandi S, Bryg R. B-type Natriuretic Peptide (BNP) is useful in detecting asymptomatic left ventricular dysfunction in low-income, uninsured patients. *Biol Res Nurs*. 2010;11(3):280-287.
50. Karuppiyah S, Graham F, Ledwidge M, et al. Elevated BNP with normal systolic function in asymptomatic individuals at-risk for heart failure: a marker of diastolic dysfunction and clinical risk. *Ir J Med Sci*. 2006;175(4):5-13.
51. Lukowicz TV, Fischer M, Hense HW, et al. BNP as a marker of diastolic dysfunction in the general population: Importance of left ventricular hypertrophy. *Eur J Heart Fail*. 2005;7(4):525-531.
52. Resnik JL, Hong C, Resnik R, et al. Evaluation of B-type natriuretic peptide (BNP) levels in normal and preeclamptic women. *Am J Obstet Gynecol*. 2005;193(2):450-454.
53. Tanous D, Siu SC, Mason J, et al. B-type natriuretic peptide in pregnant women with heart disease. *J Am Coll Cardiol*. 2010;56(15):1247-1253.
54. Kansal M, Hibbard JU, Briller J. Diastolic function in pregnant patients with cardiac symptoms. *Hypertens Pregnancy*. 2012;31(3):367-374.
55. Bauersachs J, König T, van der Meer P, et al. Pathophysiology, diagnosis and management of peripartum cardiomyopathy: a position statement from the Heart Failure Association of the

- European Society of Cardiology Study Group on peripartum cardiomyopathy. *Eur J Heart Fail* 2019; 21:827.
56. Patten IS, Rana S, Shahul S, et al. Cardiac angiogenic imbalance leads to peripartum cardiomyopathy. *Nature* 2012; 485:333.
  57. O'Connell JB, Costanzo-Nordin MR, Subramanian R, et al. Peripartum cardiomyopathy: clinical, hemodynamic, histologic and prognostic characteristics. *J Am Coll Cardiol* 1986; 8:52.
  58. Hilfiker-Kleiner D, Kaminski K, Podewski E, et al. A cathepsin D-cleaved 16 kDa form of prolactin mediates postpartum cardiomyopathy. *Cell* 2007; 128:589.
  59. Pearson GD, Veille JC, Rahimtoola S, et al. Peripartum cardiomyopathy: National Heart, Lung, and Blood Institute and Office of Rare Diseases (National Institutes of Health) workshop recommendations and review. *JAMA* 2000; 283:1183.
  60. Bello N, Rendon IS, Arany Z. The relationship between pre-eclampsia and peripartum cardiomyopathy: a systematic review and meta-analysis. *J Am Coll Cardiol* 2013; 62:1715.
  61. Mendelson MA, Chandler J. Postpartum cardiomyopathy associated with maternal cocaine abuse. *Am J Cardiol* 1992; 70:1092.
  62. Lampert MB, Hibbard J, Weinert L, et al. Peripartum heart failure associated with prolonged tocolytic therapy. *Am J Obstet Gynecol* 1993; 168:493.
  63. Sliwa K, Hilfiker-Kleiner D, Petrie MC, et al. Current state of knowledge on aetiology, diagnosis, management, and therapy of peripartum cardiomyopathy: a position statement from the Heart Failure Association of the European Society of Cardiology Working Group on peripartum cardiomyopathy. *Eur J Heart Fail* 2010; 12:767.
  64. Demakis JG, Rahimtoola SH, Sutton GC, et al. Natural course of peripartum cardiomyopathy. *Circulation* 1971; 44:1053.
  65. Hibbard JU, Lindheimer M, Lang RM. A modified definition for peripartum cardiomyopathy and prognosis based on echocardiography. *Obstet Gynecol* 1999; 94:311.
  66. Honigberg MC, Elkayam U, Rajagopalan N, et al. Electrocardiographic findings in peripartum cardiomyopathy. *Clin Cardiol* 2019; 42:524.
  67. Forster O, Hilfiker-Kleiner D, Ansari AA, et al. Reversal of IFN-gamma, oxLDL and prolactin serum levels correlate with clinical improvement in patients with peripartum cardiomyopathy. *Eur J Heart Fail* 2008; 10:861.