

# Obstetric Anesthesia Considerations in Heart Failure

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# Sarah

25 y/o G2P1

Dx PPCM during 2<sup>nd</sup> pregnancy

1<sup>st</sup> misdx as PNA

PP decompensation to LVEF 5%

LVAD → TIA

Heart transplant

*“A lot of times, when you hear the words ‘heart failure,’ you think of the elderly.*

*As a healthy 25-year-old, never did I expect to hear those two words. And never did I ever expect to hear those two words while pregnant with my second.”*



<https://transplantliving.org/stories/i-only-had-a-2-chance/>  
<https://expectinghearts.com/sarahs-story>



# Objectives:

**Current Data on Maternal Mortality**

**Peripartum Hemodynamics Review**

**Maternal Cardiac Risk Scoring Systems  
& Heart Failure Identification Tools**

**Heart Failure from  
Preeclampsia & Peripartum Cardiomyopathy**

**Delivery Management Considerations in:**

- **Heart Failure**
- **Pulmonary Edema**

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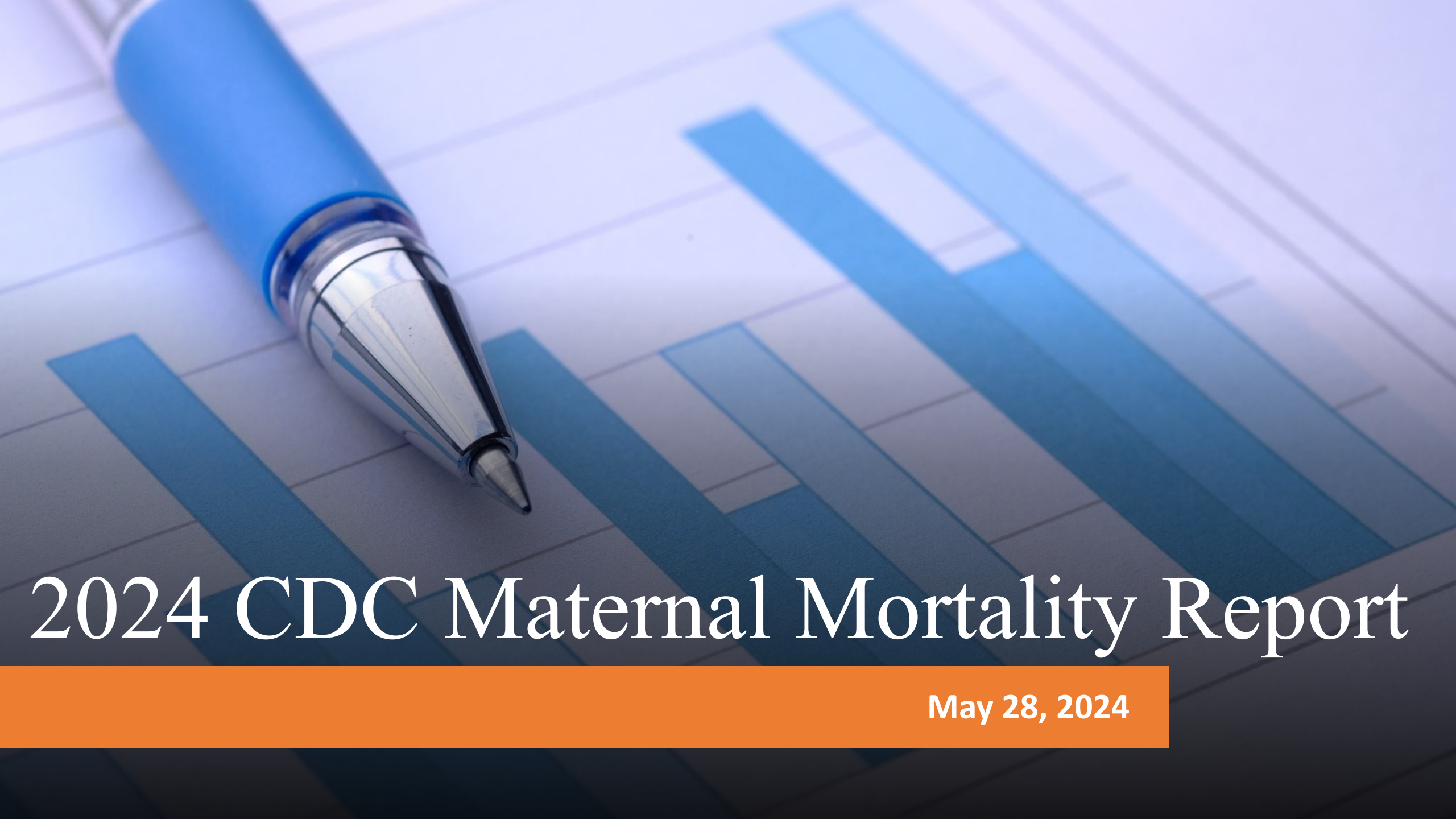
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# 2024 CDC Maternal Mortality Report

May 28, 2024



**Among pregnancy-related deaths in 2020, MMRCs  
determined **84% to be preventable.** ( $^{430}_{515}$ )<sup>1</sup>**

**Cardiac disease is 2<sup>nd</sup> leading cause of Pregnancy-Related  
Death (PRD) (16.2%)<sup>1</sup>**



# CDC 2024 Maternal Mortality Report

- **CV disease 2<sup>nd</sup> leading cause of PRD** (16.2% total mortality) after mental health (21.9% total mortality)
  - **Cardiomyopathy 41.2 % (6.7% total mortality)** *other CV conditions 58.8% (9.5% total mortality)*
    - *Other CV conditions: CAD, pulm HTN, acquired & congenital valvular heart dx, vascular aneurysm, HTN, Marfan dx, conduction defects, vascular malformations, and other CV dx; \*excluding cardiomyopathy & HTN disorders of pregnancy.*
    - **HTN d/o of pregnancy was 6<sup>th</sup> leading cause of PRD, accounting for 6%,** behind: infection (16%), hemorrhage (10.9%), and embolism (8.4%)<sup>1</sup>

## Slide Notes:

For comparison: Cardiomyopathy kills as many women as HTN disorders of pregnancy (PreE/Eclampsia/+/-HELLP).

- Medical community very vigilant with routine screening of ante- & postpartum women for development of HTN disorders, yet no standard assessments are in place to routinely assess for heart failure, despite ~ equal mortality %.

# CDC 2024 Maternal Mortality Report

	Number of pregnancy-related deaths	%
During pregnancy	135	25.7
Day of delivery	58	11.1
1–6 days postpartum	85	16.2
7–42 days postpartum	106	20.2
43–365 days postpartum	141	26.9
<sup>a</sup> Percentages might not sum to 100 because of rounding.		

Among pregnancy-related deaths in 2020:  
63% occurred postpartum, and **47% occurred 7–365 days postpartum (PP).**<sup>1</sup>

Slide Notes:

Over half of maternal deaths (63%) occurred PP, with 47% occurring 1wk - 1 yr PP1, after most women have left the hospital. Women many require PP (within ~6mos of end of pregnancy) anesthesia care for various reasons: Delayed urgent or elective surgeries/procedures until PP, PP tubals, PPH D&C which can occur weeks after delivery, emergency surgery

WE MUST BE AWARE OF POSSIBLTY OF HEART FAILURE IF S/S PRESENT PERIOP IN BOTH PREGNANT &

1. Trost SL, Busacker A, Leonard M, et al., CDC, 2024

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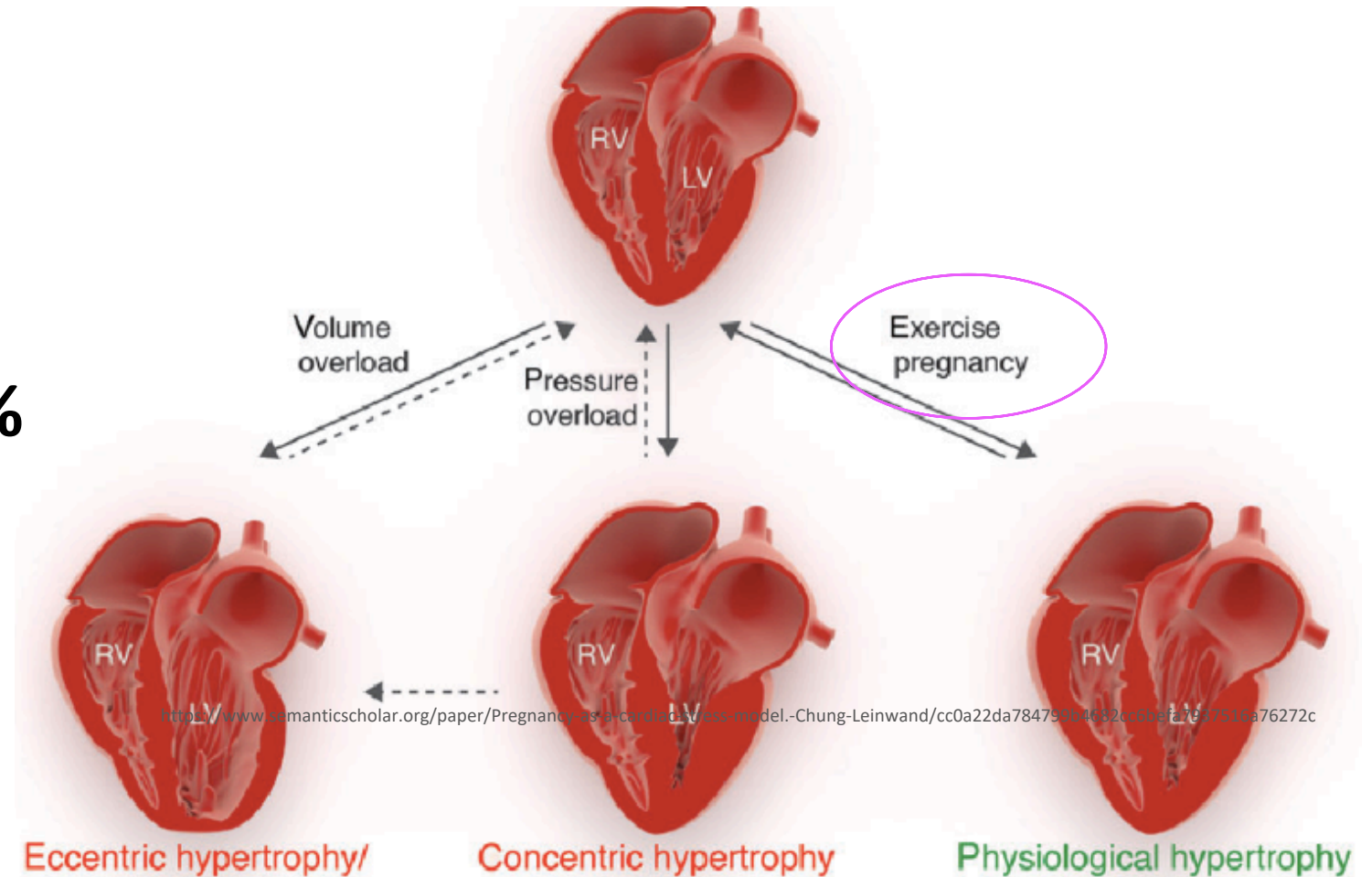
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# Peripartum Myocardial Remodeling

- LV wall thickness ↑ 28%
- LV mass ↑ 52%
- RV mass ↑ 40%



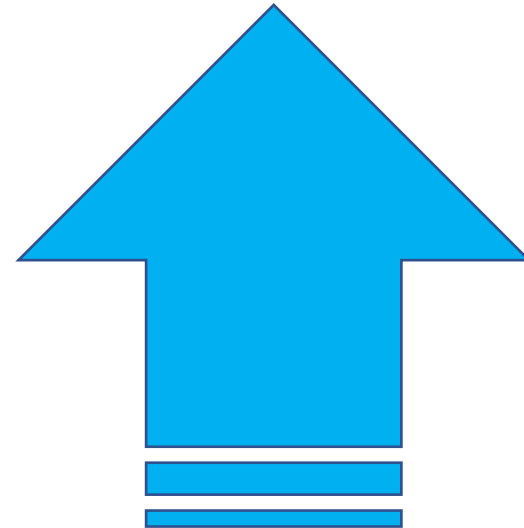


Peripartum  
fluid shifts..

can hit like a  
truck.

Cardiac output is highest **during labor and immediately after delivery**

- Increases 60 - 80% above pre-labor levels.





# Peripartum fluid shifts..

# can hit like a truck.

## Immediately on Delivery:

- **Increased preload:**
  - 1) Gravid uterus offloaded from great vessels
  - 2) Auto-Transfusion
    - [ 300-500cc with each labor cxn ]
    - Occurs during labor & immediately after delivery
- CO ~150% above non-pregnant levels.
- CO, HR, SV remain significantly elevated for first few days PP to accommodate IV fluid shifts.<sub>3</sub>
- Hemodynamics take ~ 2 wks to return to normal in patients with *normal cardiac function* as extravascular fluid shifts back into intravascular space.<sub>2</sub>

2. Sanghavi & Rutherford, *Circulation*, 2014

3. Meng ML, Arendt KW, *Anesthesiology*, 2021



The sudden ↑ in preload and C.O. on delivery can overwhelm a failing ventricle



## Jessica

30 y/o G2P1 HTN at 32w → PO Anti-HTN, IOL 37wk → Breech c/s.

**Coded in PACU.**

10 min CPR , Impella, then ECMO for 2 wks.

Dx: paraganglioma: rare adrenal tumor likely contributed to HF.

*“This pregnancy was different. I was more tired, I got winded easily, and I had terrible headaches when I would lay down at night.”*

<https://www.stroke.org/en/news/2020/11/19/mother-brought-back-to-life-the-day-she-gave-birth>  
<https://www.today.com/health/womens-health/peripartum-cardiomyopathy-mom-30-shares-symptoms-rcna50204>  
<https://thebirthtraumamama.com/jessicas-story-surviving-peripartum-cardiomyopathy/>



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# Current Maternal Cardiac Risk Scoring Systems

**mWHO**

**CARPREG II**

**ZAHARA**

# mWHO

## Modified World Health Organization Classification of Maternal Cardiovascular Risk

Slide Notes:

Ex: EF 45% is class II-III, but EF 45% with hx PPCM = residual impairment and is class IV, which is a significantly more dangerous pregnancy.

**Table 2: Modified World Health Organization (WHO) Classification of Maternal Cardiovascular Risk: Application**

WHO Pregnancy Risk Classification (Risk of pregnancy by medical condition)	Cardiovascular Conditions by WHO Risk Class
<b>WHO Risk Class I</b> <i>No detectable increased risk of maternal mortality and no or mild increase in morbidity.</i>	<ul style="list-style-type: none"> <li>Uncomplicated, small or mild               <ul style="list-style-type: none"> <li>Pulmonary stenosis</li> <li>Patent ductus arteriosus</li> <li>Mitral valve prolapse</li> </ul> </li> <li>Successfully repaired simple lesions (atrial or ventricular septal defect, patent ductus arteriosus, anomalous pulmonary venous drainage).</li> <li>Atrial or ventricular ectopic beats, isolated</li> </ul>
<b>WHO Risk Class II</b> (If otherwise well and uncomplicated) <i>Small increased risk of maternal mortality or moderate increase in morbidity.</i>	<ul style="list-style-type: none"> <li>Unoperated atrial or ventricular septal defect</li> <li>Repaired tetralogy of Fallot</li> <li>Most arrhythmias</li> </ul>
<b>WHO Risk Class II or III</b> (Depending on individual) <i>Risk as indicated in Class II (above) or Class III (below).</i>	<ul style="list-style-type: none"> <li>Mild left ventricular impairment</li> <li>Hypertrophic cardiomyopathy</li> <li>Native or tissue valvular heart disease not considered WHO I or IV</li> <li>Marfan syndrome without aortic dilatation</li> <li>Aorta &lt;45 mm in aortic disease associated with bicuspid aortic valve</li> <li>Repaired Coarctation</li> </ul>
<b>WHO Risk Class III</b> <i>Significantly increased risk of maternal mortality or severe morbidity. Expert counseling required. If pregnancy is decided upon, intensive specialist cardiac and obstetric monitoring needed throughout pregnancy, childbirth and the puerperium.</i>	<ul style="list-style-type: none"> <li>Mechanical valve</li> <li>Systemic right ventricle</li> <li>Fontan circulation</li> <li>Cyanotic heart disease (unrepaired)</li> <li>Other complex congenital heart disease</li> <li>Aortic dilatation 40-45 mm in Marfan Syndrome</li> <li>Aortic dilatation 45-50 mm in aortic disease associated with bicuspid aortic valve</li> </ul>
<b>WHO Risk Class IV</b> (Pregnancy contraindicated) <i>Extremely high risk of maternal mortality or severe morbidity; pregnancy contraindicated. If pregnancy occurs termination should be discussed. If pregnancy continues, care as for class III.</i>	<ul style="list-style-type: none"> <li>Pulmonary arterial hypertension of any cause</li> <li>Severe systemic ventricular dysfunction (LVEF &lt;30%, NYHA III-IV)*</li> <li>Previous peripartum cardiomyopathy with any residual impairment of left ventricular function</li> <li>Severe symptomatic mitral or aortic stenosis</li> <li>Marfan syndrome with aorta dilated &gt;45 mm</li> <li>Aortic dilation &gt;50 mm in aortic disease associated with bicuspid aortic valve</li> <li>Native severe Coarctation</li> </ul>

# Cardiac Event Rate Based on mWHO classification

**I: 2.5%-5%**

No detectable increased risk of maternal mortality  
No or mild increased risk in morbidity

**II: 5.7-10.5%**

Small increased risk of maternal mortality  
Moderate increase in morbidity

**II-III: 10%-19%**

Intermediate increased risk of maternal mortality  
Moderate to severe increase in morbidity

**III: 19%-27%**

Significantly increased risk of maternal mortality  
Significant increase in severe morbidity

**IV: 40%-100%**

Extremely high risk of maternal mortality  
Extremely high risk of severe morbidity

# ZAHARA & CARPREG II

## Slide Notes:

- CARPREG system replaced by more comprehensive CARPREG II\*
- CARPREG = CARDiac dx in PREGnancy.
- ZAHARA = same concept but in Dutch
- CARPREG II & mWHO most commonly used together.

**Table 1: Clinical Factors Identified as Risks in the Current Scoring Tools**

Current Scoring Tools	Points per Factor	Maternal Risk
<b>CARPREG</b>		
<ul style="list-style-type: none"> <li>• Prior cardiac event (including arrhythmia)</li> <li>• NYHA III/IV or cyanosis</li> <li>• Systemic ventricular dysfunction (EF &lt;40)</li> <li>• Left heart obstruction</li> </ul>	1 for each factor	0 points – 5% 1 point – 27% >1 point – 75%
<b>ZAHARA</b>		
<ul style="list-style-type: none"> <li>• Prior arrhythmia</li> <li>• NYHA III/IV</li> <li>• Left heart obstruction</li> <li>• Mechanical valve prosthesis (strongest weighed)</li> <li>• Cyanotic</li> <li>• Cardiac medication before pregnancy</li> <li>• Moderate/severe AVV regurgitation (systemic)</li> <li>• Moderate/severe AVV regurgitation (sub-pulmonary)</li> </ul>	Arrhythmia – 1.5 Cardiac medication – 1.5 NYHA class – 0.75 Left heart obstruction – 2.5 Systemic AVVR – 0.75 Sub-pulmonary AVVR – 0.75 Mechanical valve – 4.25 Cyanosis – 1	>0.5 points – 2.9% 0.51–1.5 points – 7.5% 1.51–2.5 points – 17.5% 2.51–3.5 points – 43% >3.51 – 70%
<b>CARPREG II</b>		
<ul style="list-style-type: none"> <li>• Prior cardiac event (including arrhythmia)</li> <li>• NYHA III/IV or cyanosis</li> <li>• Systemic ventricular dysfunction (EF &lt;40)</li> <li>• Left heart obstruction</li> <li>• Mechanical valve prosthesis (strongest weighed)</li> <li>• Pulmonary hypertension</li> </ul>	Prior event/arrhythmia – 3 NYHA III/IV/cyanosis – 3 Systemic EF <40 – 2 Left heart obstruction – 2 Mechanical valve – 2 Pulmonary hypertension – 2 Coronary artery disease – 2 High risk aortopathy – 2 No cardiac intervention – 1 Late presentation – 1	0–1 point – 5% 2 points – 10% 3 points – 15% 4 points – 22% >4 points – 41%



# Cardiac Event Risk

## Highest risk patients are:

- Older
- Identify as Black or African American
- Acquire heart disease during pregnancy (eg: PPCM)
- Have unrecognized cardiovascular disease and become pregnant <sup>5</sup>

# IMPLEMENTING CARDIAC RISK SCORING SYSTEMS

Each risk scoring system has individual limitations.

Careful interpretation and combining scoring systems can help estimate individual risk

mWHO classification is the 1<sup>st</sup> step in guiding management and follow-up during pregnancy.

# IDENTIFYING HEART FAILURE IN PREGNANCY



# IDENTIFYING HEART FAILURE IN PREGNANCY

Many symptoms of HF are  
vague and mimic normal  
pregnancy symptoms  
(eg: fatigue, SOB,  
peripheral edema.)



Failure to recognize HF  
contributes significantly to  
maternal deaths in the US.

# IDENTIFYING HEART FAILURE IN PREGNANCY

ACOG Practice Bulletin No. 212: *Pregnancy and Heart Disease* contains a table of signs and symptoms which, if reported, require cardiac evaluation ASAP.

If (+): Low threshold to obtain further imaging: TTE, CXR, LUS, ECG



Vital Signs	Physical Exam Signs	History and Symptoms
Heart rate $\geq$ 120 beats/min	Jugular venous pressure visible 2 cm above clavicle at 45°	History of cardiovascular disease
Systolic blood pressure $\geq$ 160 mm Hg	Loud systolic murmur or S4	Shortness of breath at rest, paroxysmal nocturnal dyspnea, orthopnea, refractory pneumonia, or bilateral chest infiltrates on chest radiography
Symptomatic low blood pressure	Wheezing	Chest pain at rest or minimal exertion
Respiratory rate $\geq$ 25 breaths/min	Lung crackles	Exertional or unprovoked syncope or palpitations associated with near syncope or syncope
Oxygen saturation $<$ 95%	Marked peripheral edema	Extreme fatigue

Modified from American College of Obstetricians and Gynecologists.<sup>14</sup>

National Partnership for  
Maternal Safety

## Maternal Early Warning Criteria (MEWC)

- To aid in faster recognition, evaluation, diagnosis, and treatment of signs of developing critical illness, to reduce preventable maternal death.

If (+): Low threshold  
to obtain further  
imaging: TTE, CXR,  
LUS, ECG

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Systolic BP (mm Hg)	<90 or >160
Diastolic BP (mm Hg)	>100
Heart rate (beats per min)	<50 or >120
Respiratory rate (breaths per min)	<10 or >30
Oxygen saturation on room air, at sea level, %	<95
Oliguria, mL/hr for $\geq 2$ hours	<35
Maternal agitation, confusion, or unresponsiveness; Patient with preeclampsia reporting a non-remitting headache or shortness of breath	

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# “Fett Self Test” for Early Identification of Heart Failure

> 4 pts → further workup (BNP & TTE)<sup>27</sup>

Table 1. Self-test for early diagnosis of peripartum cardiomyopathy.

Symptoms	0 points	1 point	2 points
Orthopnea	None	Need to elevate head	Need to elevate 45 degrees or more
Dyspnea	None	Climbing 8 or more steps	Walking on level
Unexplained cough	None	At night	Day and night
Lower extremity swelling	None	Below knee	Above and below knee
Excessive weight gain during last month of pregnancy	Under 2 pounds per week	2–4 pounds per week	Over 4 pounds per week
Palpitations	None	When lying down at night	Day and night, any position

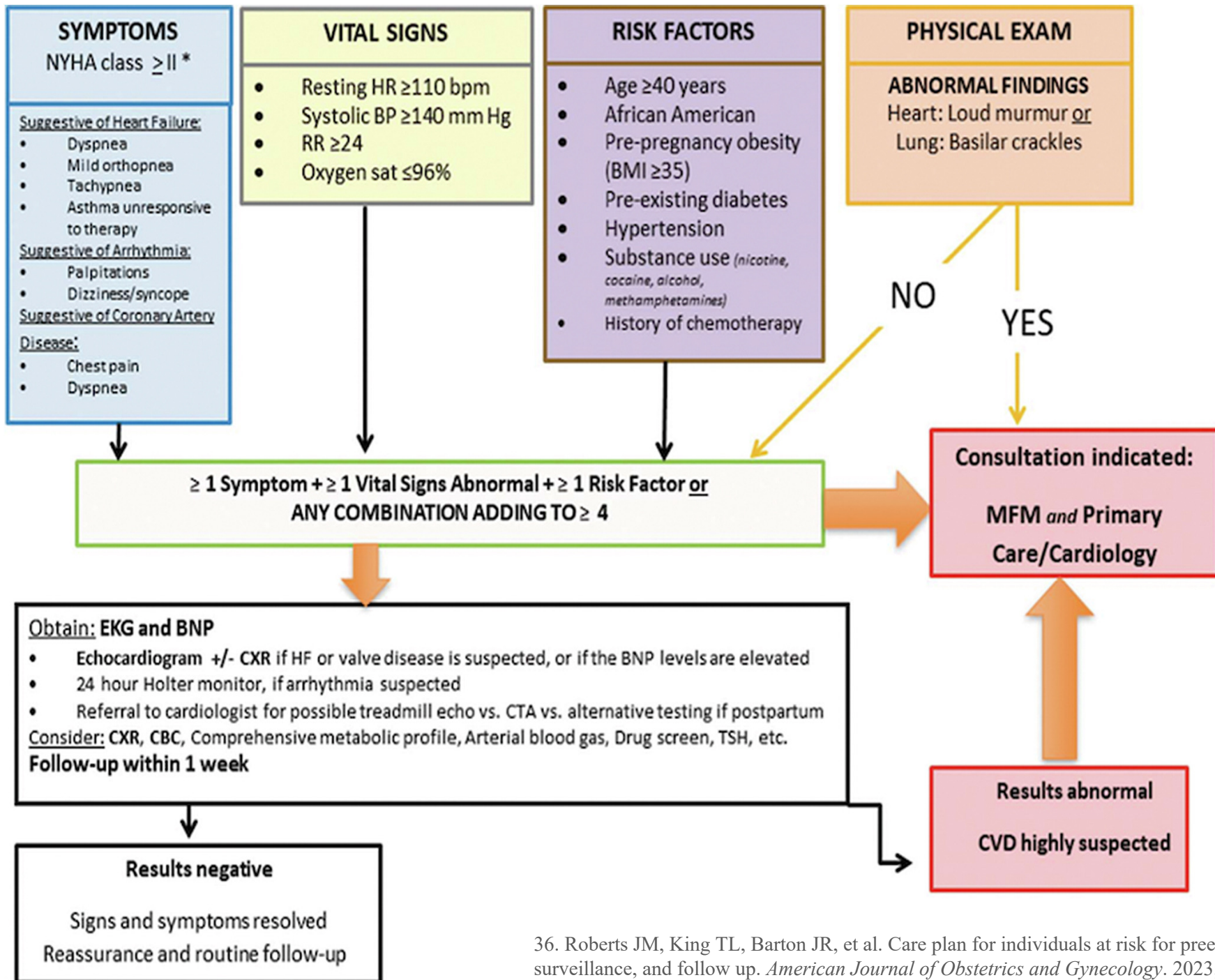
The presence of 4 or more points should prompt additional investigation.  
Data taken from [30].

Davis M, Duvernoy C. *Women's Health*. 2015

In validation study: **100% of women who presented with > 4pts had LV systolic dysfunction.**<sup>27</sup>

26. Davis M, Duvernoy C. Peripartum Cardiomyopathy: Current Knowledge and Future Directions. *Women's Health*. 2015.

27. Fett JD. Validation of a self-test for early diagnosis of heart failure in peripartum cardiomyopathy. *Critical Pathways in Cardiology*. 2011;10(1):44-45. doi:<https://doi.org/10.1097/HPC.0b013e31820b887b>



Pts with risk factors:  
AMA, obesity, CHTN, DM,  
or other risk factors for  
PreE (and therefore CV  
disease) will benefit from  
baseline cardiac eval  
including ECG & ECHO.

# BNP

## Reference Intervals in Pregnancy:

- BNP: **> 50** pg/ml
- NT-proBNP:
  - TM 1&2: **> 200** pg/ml
  - TM 3: **> 150** pg/ml<sub>33</sub>

- BNP shown to improve diagnostic accuracy in women presenting with acute dyspnea.<sup>12</sup>
- BNP =/↓ in pregnancy, despite ↑ plasma volume; as normal adaptations of cardiac remodeling and ↓ SVR leave central filling pressures unchanged,<sup>13</sup> and plasma dilution and ↑ GFR will ↓ circulating levels.<sup>33</sup>
- BNP ↑ in PreE... *as a marker of diastolic dysfunction.*<sup>11</sup>
- **BNP LOWER in obesity and diabetes, with additive lowering effect**<sup>34, 35</sup>
- Traditionally, BNP < 100 rules out HF in *non-pregnant* patients.

11. Borges VTM, Zanati SG, Peraçoli MTS, et al. *Ultrasound in Obstetrics & Gynecology*. 2018

12. Mueller C, Scholer A, Laule-Kilian K, et al. *N Engl J Med*. 2004

13. Estensen ME, Beitnes JO, Grindheim G, et al. *Ultrasound in Obstetrics & Gynecology*. 2013

33. Stocktree, et al. *J Endocr Soc*. 2021

34. Wang TJ, et al. *Circulation*. 2004

35. Nishikimi T, Nakagawa Y. *Journal of Cardiology*. 2021.

# IDENTIFYING HEART FAILURE IN PREGNANCY

- Chances of full recovery higher when LVEF > 35%;
  - With early detection, LVEF likely to be higher at time of diagnosis and initiation of treatment.
- Most serious complications of PPCM arise when diagnostic or baseline LVEF < 30-35%
  - Ventricular tachyarrhythmias
  - Thromboembolic events
  - Chronic cardiomyopathy
- *These complications can be either avoided or decreased with earlier diagnosis...*







*...But women's symptoms of heart failure are frequently misdiagnosed, or dismissed as pregnancy symptoms, which delays treatment and leads to further decompensation.*

## Rachel

s/s 3<sup>rd</sup> TM: excessive swelling – dismissed “bc it’s twins”

Leg swelling after delivery – told “it’s nothing”

Cough developed - told “it’s bronchitis”

Workup 1 month PP

LVEF 15%

Ext defib & cardiac rehab - Recovered

*“And I kept saying something was wrong, and they told me, 'No, you're just carrying multiples. This is what happens with multiples. There's nothing wrong with you'...  
...But come to find out, I was dying.”*



# Claire

Unable to lie flat & severe cough 6d postpartum after 1<sup>st</sup> delivery.

She went to 8 emergency appts, saw 5 different doctors and was given incorrect dxs of a PE and gastric asthma.

***Even despite mentioning she thought her s/s were cardiac and could be PPCM***

***“I was told I was just a hormonal, overanxious new mother.”***

S/S worsened: “SOB, breathless, gasping for air”

→ ED Echo → **EF < 18%**

Managed medically

**Over 7 years later, EF 48-50%, still medication dependent.**



Peripartum Cardiomyopathy:  
Claire's story ISSN 2516-5852 (Online)  
AIMS Journal, 2024, Vol 36, No 2

# Marian

35y/o

6 months PP: fatigue, bloat, ankle swelling, brief vision loss

1<sup>st</sup> and 2<sup>nd</sup> GP dismissed s/s as viral:

*“At the second appointment, the physician emphasized that I was young and healthy”*

3<sup>rd</sup>: ED doc: misdx “severe congestion” as PNA

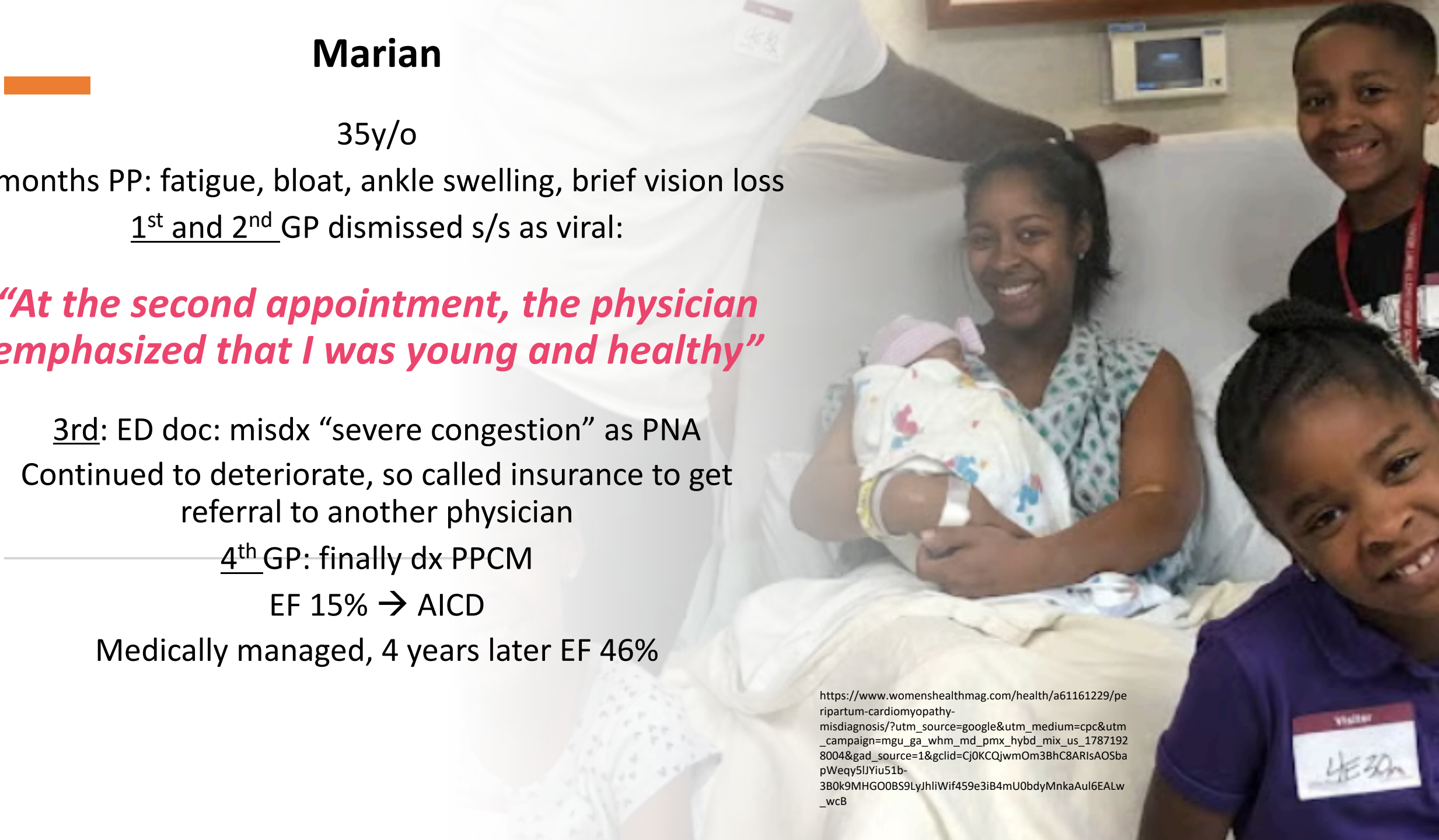
Continued to deteriorate, so called insurance to get referral to another physician

4<sup>th</sup> GP: finally dx PPCM

EF 15% → AICD

Medically managed, 4 years later EF 46%

[https://www.womenshealthmag.com/health/a61161229/peri-partum-cardiomyopathy-misdiagnosis/?utm\\_source=google&utm\\_medium=cpc&utm\\_campaign=mgu\\_ga\\_whm\\_md\\_pmx\\_hybd\\_mix\\_us\\_17871928004&gad\\_source=1&gclid=Cj0KCQjwmOm3BhC8ARIsAOSbaPWeqy5IJYiu51b-3B0k9MHGO0BS9LyJhliWif459e3iB4mU0bdyMnkaAul6EALw\\_wcB](https://www.womenshealthmag.com/health/a61161229/peri-partum-cardiomyopathy-misdiagnosis/?utm_source=google&utm_medium=cpc&utm_campaign=mgu_ga_whm_md_pmx_hybd_mix_us_17871928004&gad_source=1&gclid=Cj0KCQjwmOm3BhC8ARIsAOSbaPWeqy5IJYiu51b-3B0k9MHGO0BS9LyJhliWif459e3iB4mU0bdyMnkaAul6EALw_wcB)





# Lacresha

31 y/o 2<sup>nd</sup> child

Gained 10 lbs in 1 wk + SOB

***"I couldn't breathe"***

dismissed as "normal pregnancy s/s"

Collapsed at home shortly PP

→ Cardiogenic shock & MODS

EF 5%

ECMO, Impella, & CRRT

Now LVAD-dependent

***"Though we don't have the medical degree, we know our bodies... And no one knows our bodies better than we do"***





## Brittany

39-y/o noticed s/s as avid runner during 5<sup>th</sup> pregnancy - more SOB than usual.

**5<sup>th</sup> child: 3<sup>rd</sup> TM SOB: dismissed twice by providers as “normal pregnancy s/s.”**

**6<sup>th</sup> child, significant SOB though pregnancy,** diagnosed PP PPCM, and began PO meds for HF  
**5 months PP she experienced rapid decompensation.**

**EF 10%**

Heart transplant recipient

*"I've had five other pregnancies...from the very beginning, I struggled to breathe..."*  
*I just wish ...that this would have been caught a little earlier, and maybe things could have been different.."*



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# Hypertension & Heart Failure

HTN in pregnancy is a risk factor for heart failure development

Severe PreE causes diastolic dysfunction, and some of these women decompensate to systolic dysfunction heart failure.



# PreE and Heart Failure

- Compared with normal cohorts, PreE has a 71% increased risk of CVD mortality
  - 2.5-fold increased risk of CAD
  - 4-fold increased risk of heart failure.<sup>8</sup>
- PreE with PPCM has been associated with a higher incidence of adverse CV outcomes.<sup>9</sup>



**Madi**

17y/o

**Emergent c/s 25w5d for severe PreE**

Decompensated HF: dx PPCM  
IABP → ECMO → RVAD & LVAD

Remains with LVAD

8. Mehta LS, Warnes CA, Bradley E, et al. *Circulation*. 2020

9. Zoltan. *N Engl J Med*, 2024.

# PreE and Heart Failure with Preserved EF

- Prior to decompensation (reduced EF), echos on preE women often look...  
...*normal*... with normal to increased CO, EF & Contractility<sub>32</sub>
- Further investigation often shows:
  - Diastolic dysfunction\*\*
  - Increased pericardial effusions
  - Increased LV wall dimensions compared to healthy parturients<sub>32</sub>
- So a quick cardiac POCUS will probably look normal unless obvious LV hypertrophy (PSA) or LA dilation visible (4C, PLA)
- Formal ECHO with more in-depth assessment needed, esp if symptomatic (SOB, cough, orthopnea, etc r/o pulm edema)

# PreE and Heart Failure with Preserved EF

- Difficult to diagnose and easy to miss.
- Seemingly normal EF on systolic assessment WITHOUT assessing diastolic function *can miss diagnosis of HFpEF*
- ***However 9.5 % of PreE pts w severe features found to have HFpEF! Almost 1 in 10!***
  - SOB & pulm edema are common presenting signs of HFpEF
  - Often precipitated by excess IV fluids → pulmonary edema

# PreE and Heart Failure

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- Well-established that diastolic dysfunction can be seen in PreE
  - Especially early-onset PreE < 34wks
  - Increased incidence of concentric hypertrophy, LV diastolic dysfunction, and higher BNP levels in early –onset PreE, compared with late-onset PreE. <sup>10,11</sup>
- **Despite brief time period of HTN, maternal heart can still adapt to acute and transient ↑ SVR with remodeling and concentric hypertrophy.** <sup>11</sup>
  - In fact, *most women with PreE undergo adaptive myocardial remodeling from ↑SVR, but a small subgroup demonstrate signs of overt decompensation.*
    - Pre-term PreE can lead to severe LVH assc. with advanced cardiac dysfunction. <sup>10, 11</sup>

10. Melchiorre K, Sutherland GR, Watt-Coote I, Liberati M, Thilaganathan B., *Hypertension in Pregnancy*. 2012

11. Borges VTM, Zanati SG, Peraçoli MTS, et al. *Ultrasound in Obstetrics & Gynecology*. 2018

## Cynthia

G5P4

*“Two weeks after birth, I was back in the hospital with my BP at 215/127. ...I couldn't breathe. Every time I lay down, I felt as though I was drowning.”*

EF 25%

Ext defib

Medically managed

Full recovery







# PERIPARTUM CARDIOMYOPATHY



<https://www.health.harvard.edu/womens-health/research-were-watching-womens-hearts-age-differently-than-mens-do>

<https://www.nursingtimes.net/clinical-archive/womens-health/pregnancy-1-effects-on-haematological-and-cardiovascular-systems-29-08-2023/>

# PPCM

## Defined as:

- LVEF  $\leq$  45% with no prior hx cardiac dysfunction
  - LV systolic dysfunction<sup>14</sup>
- Dx by exclusion
- No serum or biopsy markers:
  - dx by TTE and usually  $\uparrow$ BNP
- Usually LV dilation, but not always
  - most-similar to non-ischemic DCM.<sup>9</sup>
- Can have RV dysfunction as well, which is a poor prognostic indicator.<sup>15</sup>



<https://medium.com/@nelore/whos-taylor-swift-anyway-ew-0b698037cd92>

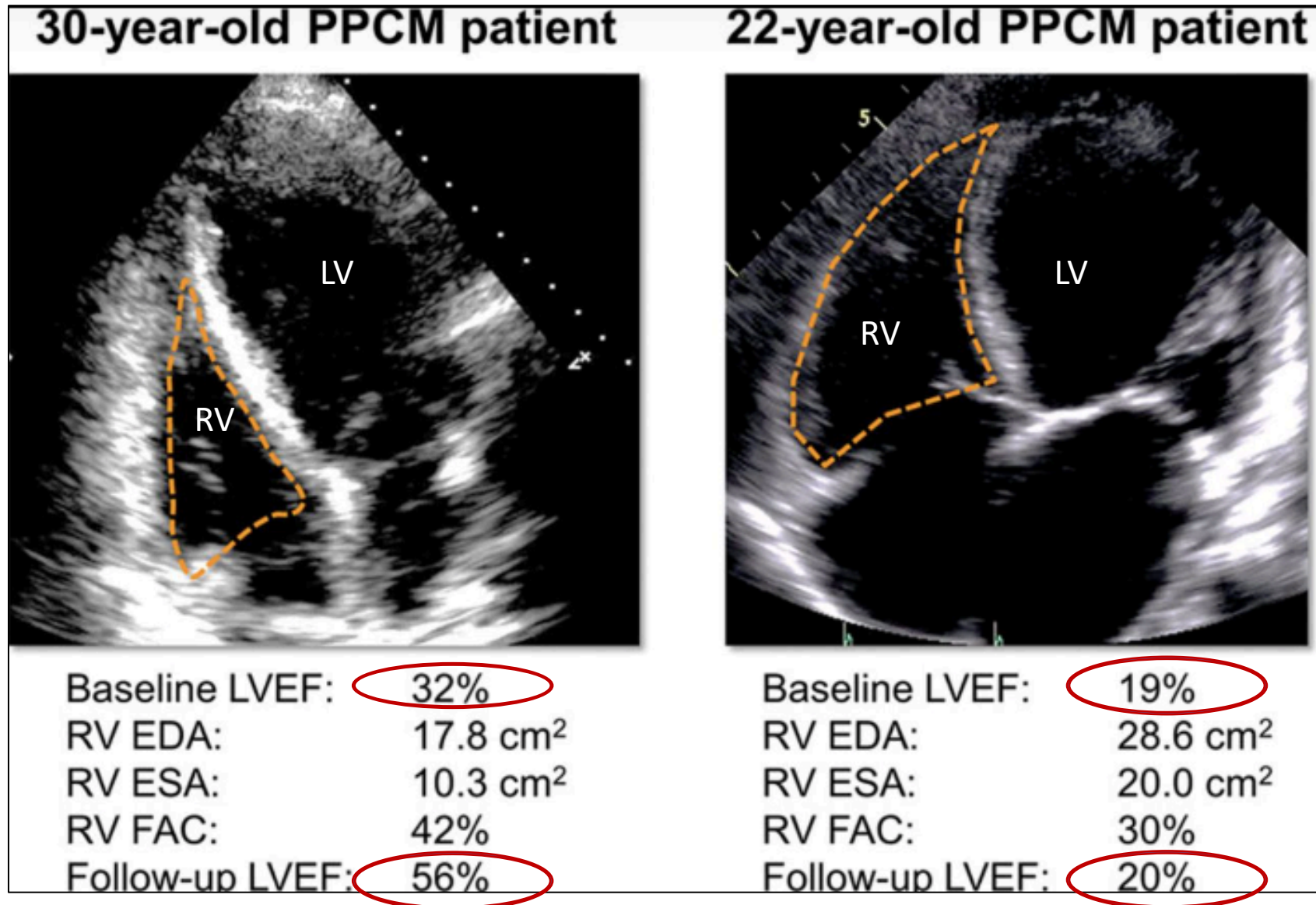
9. Zoltan. *N Engl J Med*, 2024.

14. Bauersachs J, König T, Meer P, et al. *European Journal of Heart Failure*. 2019

15. Blauwet LA, Delgado-Montero A, Ryo K, et al. *Circulation-heart Failure*. 2016



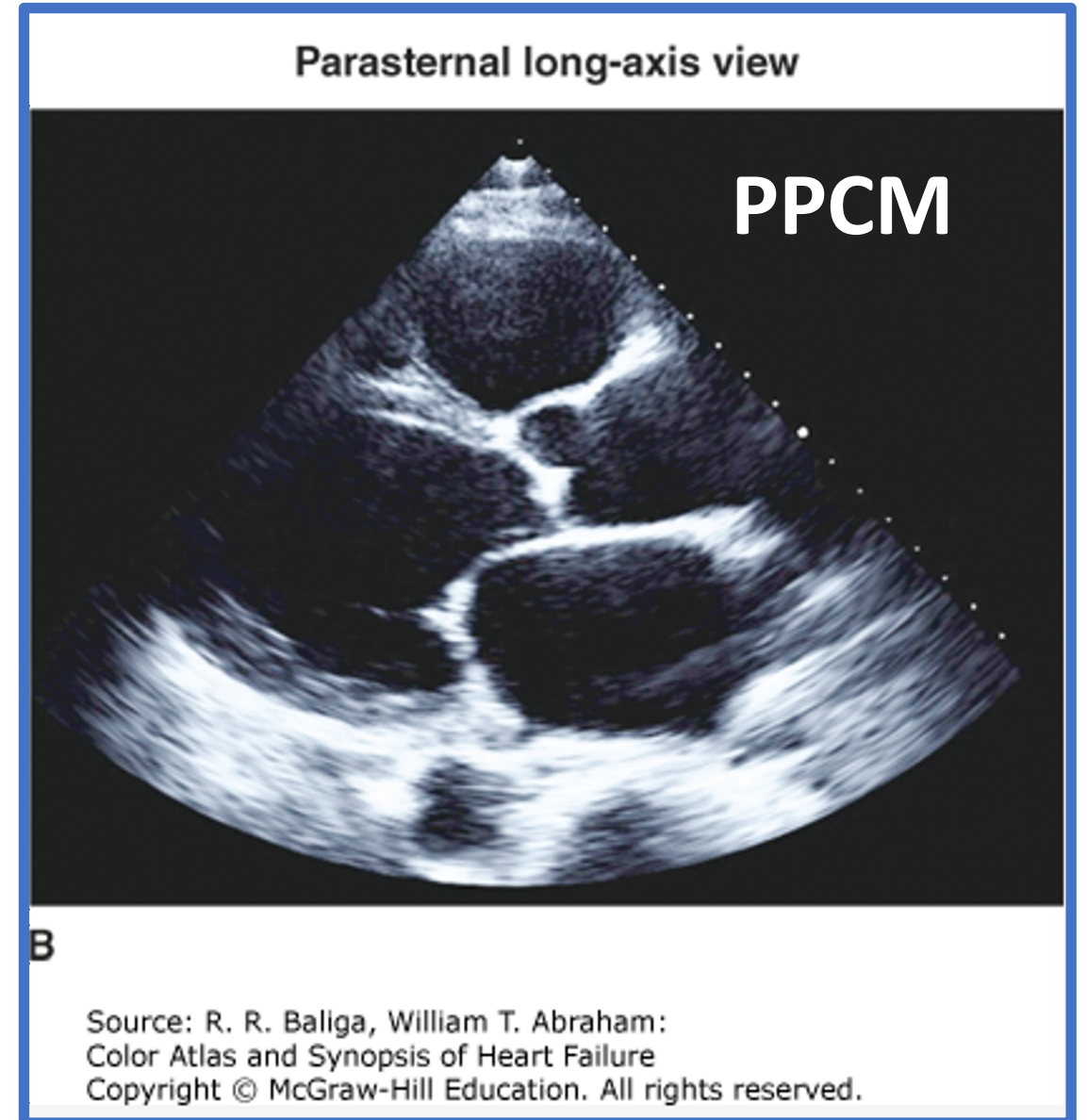
## RV dysfunction in PPCM has poorer prognosis



# PPCM

---

- Current pathogenesis theory:
  - Imbalances in peripartum hormones secreted by pituitary and placenta affect cardiac vasculature
    - Causes CV dysfunction and consequent HF in susceptible women.
- Possible genetic susceptibility.



# PPCM: Onset and Recovery

## **ONSET:**

- Can develop **RAPIDLY over the course of days: LVEF<45%.<sup>16</sup>**
  - Often presents as clinical congestion 2/2 elevated cardiac filling pressures:
    - orthopnea, bendopnea, JVD, leg edema <sup>17</sup>
  - Use acute heart failure treatment guidelines <sup>18</sup>
- Usually either during last month of pregnancy or most commonly, postpartum.
  - **60-90% of cases occur within 1<sup>st</sup> week postpartum**
  - **Can occur up to 5 months postpartum, or later...<sup>9</sup>**

16. Karaye et al. *BMC Cardiovasc Disord.* 2016

17. Heidenreich PA, Bozkurt B, Aguilar D, et al. *Circulation.* 2022

18. Regitz-Zagrosek V, Roos-Hesselink JW, Bauersachs J, et al. *European Heart Journal.* 2018

9. Zoltan. *N Engl J Med*, 2024



# PPCM: Onset and Recovery

## **RECOVERY:**

- CV function recovers in >50% of patients
  - However, some never recover and need LVAD +/- heart transplant.
- Incomplete recovery:
  - Cardiac cellular and molecular recovery may lag behind LVEF.
  - Despite recovered EF, exercise or dobutamine stress test may reveal persistent cardiac dysfunction.
- In the US, Black women have twice the risk of persistent cardiac impairment, and if recovery is achieved, it can take twice as long.
- Thorough cardiac workup if any hx PPCM – can take years to recover..

# MANY PPCM PATIENTS NEVER RECOVER LV FUNCTION



Danecia

24 y/o, 1<sup>st</sup> baby

**1 wk PP** s/s: chest pain & SOB

EF 25%

AICD

**SIX YEARS LATER** suffered stroke

**Workup: LVEF had declined to 10-15%**

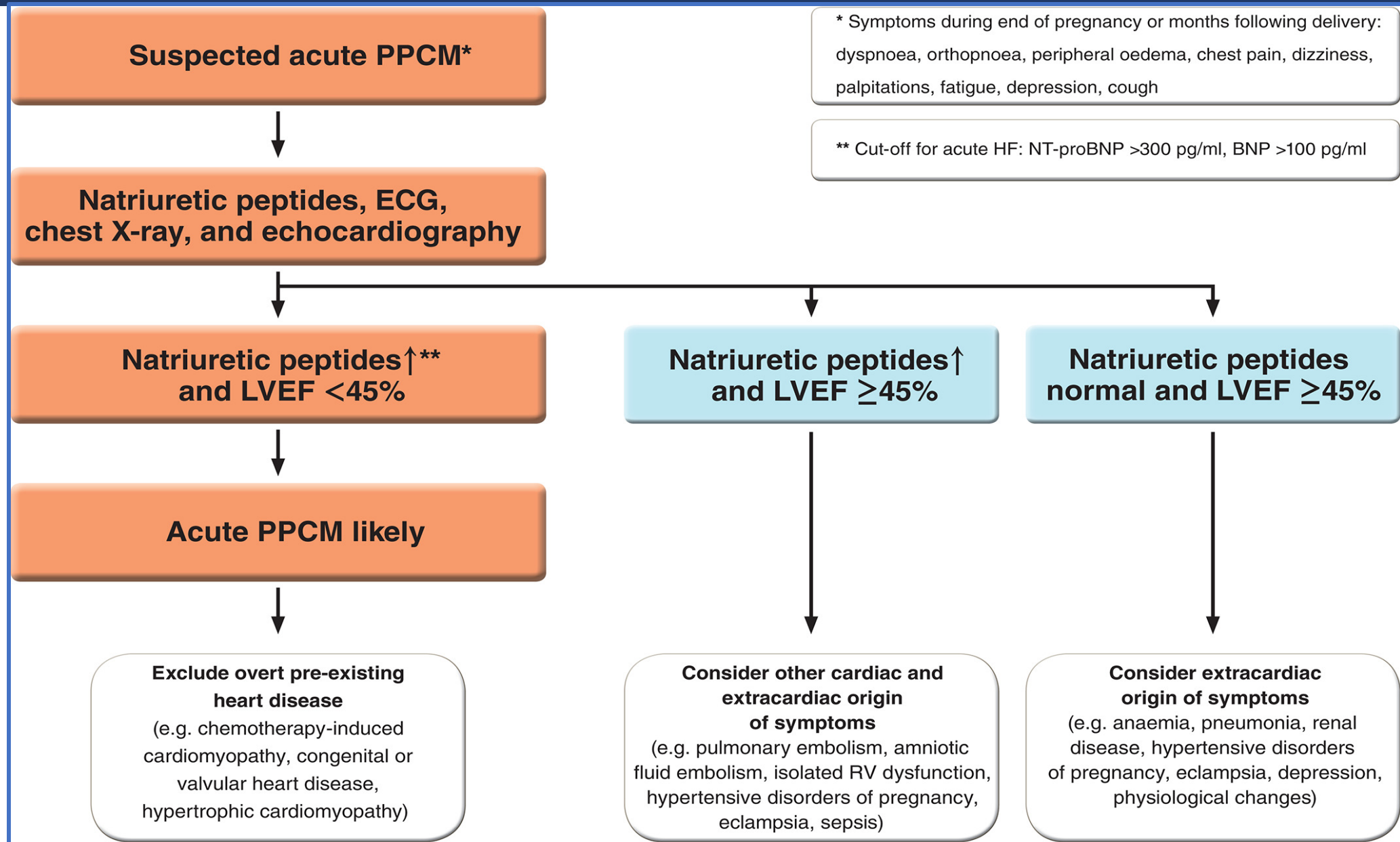
Waited >1yr for heart transplant

LVEF further declined

LVAD while she waited

Then received heart transplant.

# PPCM Diagnostic Pathway



# Objectives:

Current Data on Maternal Mortality

Peripartum Hemodynamics Review

Maternal Cardiac Risk Scoring Systems  
& Heart Failure Identification Tools

Heart Failure from  
Preeclampsia & Peripartum Cardiomyopathy

Delivery Management Considerations in:

- **Heart Failure**
- Pulmonary Edema



# Delivery Mode: Vaginal with Neuraxial

## Vaginal delivery best with good neuraxial analgesia.

- Pain control = ↓ catecholamines = ↓ CV stress
- Vaginal delivery has less severe hemodynamic swings
  - ✓ lower EBL & less neuraxial sympathectomy<sup>5</sup>
- Vaginal delivery has lower risk of infection and thrombotic complications than C/S<sup>8</sup>
- ACOG (No. 212): IOL 39-40 wks for women w/ heart dx if no spontaneous labor<sup>8,19</sup>
  - IOL at 39 wks may reduce rates of C/S and improve maternal outcomes by decreasing risk of HTN disorders of pregnancy<sup>9</sup> (ARRIVE & HYPITAT)<sup>20,21</sup>

### Slide Notes:

ARRIVE & HYPITAT are both RCTs evaluating elective IOL vs expectant management.

- ARRIVE showed eIOL at 39wks in healthy primips reduced C/S & reduced HTN d/o.
- HYPITAT showed women with gHTN or mild PreE had lower composite poor maternal outcomes w/ eIOL > 37wks.

5. Girnius, A, Meng ML. *Journal of Cardiothoracic and Vascular Anesthesia*, 2021

8. Mehta LS, Warnes CA, Bradley E, et al. *Circulation*. 2020

9. Zoltan. *N Engl J Med*, 2024

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20. Grobman WA, Rice MM, Reddy UM, et al. *New England Journal of Medicine*. 2018

21. Koopmans CM, Bijlenga D, Groen H, et al. *Lancet*. 2009

# Delivery Mode: Vaginal with Neuraxial

## Monitors:

- Continuous pulse oximetry & ECG
- NIBP Q 2-5 min after neuraxial analgesia initiated
  - May require A-line, especially before neuraxial to monitor hemodynamic response
    - AS, MS, HOCM, PPCM or PreE w/ severely reduced EF, RV dysfunction, Pulmonary HTN.
- Consider staffing if L&D RNs are not trained for invasive hemodynamic monitoring or continuous vasoactive gtts, etc.
- **Plan for → ICU if inotropes/vasoactive meds needed peripartum, as they should be maintained through postpartum period to assist ventricles with postpartum fluid shifts.**

# Delivery Mode: Cesarean with Neuraxial

- **Cesarean** reserved for obstetric or fetal indications (malpositioned/breech, FTP, eRPT c/s, FHR)
  - *Or* high-risk maternal lesions:
    - severe aortopathy, severe AS or MS, or any maternal decompensation.<sup>3,5</sup>
- **Neuraxial anesthesia preferred**
  - Spinal may be well-tolerated in mWHO class 1 & 2 lesions
  - Slowly-dosed epidural best for mWHO class 3 & 4 lesions<sup>3</sup>
- **Plan for → ICU if inotropes/vasoactive meds needed peripartum, as they should be maintained through postpartum period to assist ventricles w/ postpartum fluid shifts.**

# Delivery Mode: Cesarean with GA

- **General Anesthesia** reserved for emergency requiring ETT for cardiopulmonary indication, or contraindication to neuraxial in necessary c/s.
  - If complex requirements: (eg: TEE, ECMO) use cardiac OR <sup>3</sup>
- PPCM possibly 1<sup>st</sup> sign of rare myopathy:
  - Danon Disease (LAMP2 variants) or Duchenne Muscular Dystrophy variants <sup>9, 22</sup>
    - Succinylcholine use in RSI ?

3. Meng ML, Arendt KW. *Anesthesiology*. 2021

9. Zoltan. *N Engl J Med*, 2024

22. Ware JS, Li J, Mazaika E, et al. *N Engl J Med*. 2016



# Neuraxial Anesthesia/Analgesia Technique

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- Slow-dosed epidural offers slower sympathectomy
- C/S: IT narcotics via CSE +/- low-dose bupi ~2.5-5mg to improve surgical anesthetic while minimizing hemodynamic swings, then slowly dose epidural.
- Can avoid routine pre-procedure IVF bolus and manage sympathectomy with pressors, as HF patients at higher risk of pulmonary edema
  - or use smaller ~200cc bolus if necessary

# Thromboembolic Risk

- Consider neuraxial timing:
  - LVEF < 35% likely on anticoagulation
  - 5-20% cases develop LV thrombus<sup>9</sup>

<https://www.everydayhealth.com/heart-failure/i-wanted-to-make-it-to-her-first-birthday-new-mom-diagnosed-with-heart-failure-days-after-giving-birth-to-daughter/>



## Eloise

32 y/o, 1<sup>st</sup> child

“health nut - 5am gym & green smoothie every day”

s/s HF immediately after delivery

C/o persistent cough & SOB.

PPD#6 Dx PPCM.

4 days in ICU, stabilized and sent home.

The day after d/c, **suffered stroke at home**

Lost function of R arm and hand.

PT/OT and has made full recovery.

# Delivery Planning: Maternal Optimization

- **Cardiopulmonary workup**

- Labs: BNP (in addition to standard CBC/CMP/troponin if cp, etc..)
- ECG, TTE, CXR, LUS

- **Diuresis/Fluid restriction**

- To accommodate postpartum fluid volume
  - Targeted tx tailored to conditions (eg,  $\beta$ -blockers for LVOTO or arrhythmias)
  - Cautious diuresis in:
    - Volume overload if preload-dependent condition (eg, LVOTO) <sup>3</sup>
    - Pre-E, intravascularly dry 2/2 endothelial damage and capillary leak
  - Careful in early postpartum period, where dramatic fluid shifts and SVR changes may create hemodynamic instability. <sup>8</sup>

3. Meng ML, Arendt KW. *Anesthesiology*. 2021

8. Mehta LS, Warnes CA, Bradley E, et al. *Circulation*. 2020

# Delivery Planning: Maternal Optimization

- Assess ability to **tolerate C/S positioning**:
  - If patient unable to lie flat for c/s (ie: increased SOB, O<sub>2</sub> requirement)
    - May require ETT, as persistent PP fluid shifts can/will acutely worsen condition.
    - Controlled intubation preferred before emergency occurs.<sup>3</sup>
      - Parturients have higher incidence difficult airway, and swelling may be exacerbated by PreE and labor/pushing.<sup>30</sup>
      - *Appropriate airway positioning before start of case\**

3. Meng ML, Arendt KW. *Anesthesiology*. 2021

30. Kodali et al., *Anesthesiology*. 2008



# Delivery Planning: Maternal Optimization

- **Inotropic gtts** available in OR, depending on cardiac eval, to assist ventricle with PP fluid load
  - ✓ Norepinephrine, Dobutamine, Milrinone, Epinephrine, Dopamine
- Invasive lines: A-line, CVC if needed.
  - CCO monitor option if A-line in place.
  - **A-line during labor recommended in; RV dysfunction, severe LVOTO, severe MS, CM with severely decreased LVEF, pulm HTN, or PreE with HF.**
- Cardiac OR if ECMO, IABP LVAD considered

# INOTROPES

## Milrinone

- 0.125 – 0.375 mcg/kg/min
- Phosphodiesterase-3 inhibitor
- Loading dose can decrease SVR
  - Careful in Pre-E or preload – dependent lesion.
  - Slow titration of loading dose preferred

## Dobutamine

- 5-10 mcg/kg/min
- Beta adrenergic agonist
  - At low doses: Primarily B<sub>1</sub> agonism, some B<sub>2</sub> agonism
- Rapid onset (< 2min)
- Minimal side effects

## Dopamine

- Dose - dependent receptor response
  - D1: 1-2mcg/kg/min
  - B2: 2-10 mcg/kg/min
  - A1: >10mcg/kg/min

## Epinephrine

- Dose –dependent response
- Predominantly B1 & B2 agonism at lower doses
- 0.02-0.1 mcg/kg/min

## Extra points:

- ✓ With contractile dysfunction, maternal heart cannot increase contractility to accommodate increased preload, especially after delivery. May be advantageous to begin inotropes before signs of ventricular failure present (tachycardia, hypoxemia, hypotension). By then, myocytes may be hypoxic and less responsive.  $\Delta O_2$  demand on an already stressed ventricle..
- ✓ CCO monitoring may be beneficial in guiding and balancing vasoactive/inotropic therapy
- ✓ If parturient started on inotropes at any time peripartum (eg, optimization for pulmonary edema, systolic dysfunction, peri-delivery in anticipation of autotransfusion, etc.. **DO NOT WEAN OFF** after delivery/before case ends. **Maintain gtts and transfer to ICU.** Pt will need continued inotropic support to manage the continued increased preload, CO, and fluid volume shifts PP (~ 24-48h) to prevent cardiac decompensation <sub>5</sub>

The sudden ↑ in preload and C.O. on delivery can overwhelm a failing ventricle

*\*If inotropic/vasoactive gtts are needed during labor/delivery to optimize cardiac function, do not d/c in stable patient. Maintain gtts → transfer to ICU*

### Jessica

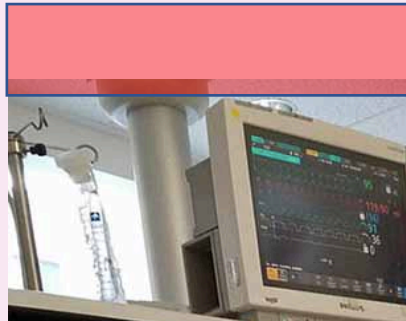
30 y/o G2P1 HTN at 32w → PO Anti-HTN. IOL 37wk → Breech → c/s.

**Coded in PACU**

10 min CPR, Impella, then ECMO for 2 wks.

Dx: paraganglioma: rare adrenal tumor likely contributed to HF.

*“This pregnancy was different. I was more tired, I got winded easily, and I had terrible headaches when I would lay down at night.”*



<https://www.stroke.org/en/news/2020/11/19/mother-brought-back-to-life-the-day-she-gave-birth>  
<https://www.today.com/health/womens-health/peripartum-cardiomyopathy-mom-30-shares-symptoms-rcna50204>  
<https://thebirthtraumamama.com/jessicas-story-surviving-peripartum-cardiomyopathy/>



# Significant Cardiac Effects of OB Meds

## Oxytocin (Pitocin)

- Usually well tolerated if titrated slowly, but rapid IV bolus ↓ SVR, → tachycardia and myocardial ischemia
  - Can give smaller, slower doses to effect. Treat ↓ SVR with pressors.<sup>3,5</sup>
  - Ex: **Rule of 3s**: Dilute 10u in 10cc NS0.9% and give 3u slowly(3cc) Q~3 min to effect.<sup>23</sup>
    - Can prevent over-dosing oxytocin. Reduced dose = reduced side effects<sup>3,23</sup>

## Methylergonovine (Methergine)

- 5-HT, Dopamine, and α-agonist – increases smooth muscle contraction
  - ↑ SVR and **PVR**
  - Can cause HTN, coronary vasospasm, and myocardial ischemia
  - *Relatively contraindicated in HTN conditions, aneurysms, CAD.*
    - If life-threatening bleeding, can give as small DILUTE IV pushes to effect
      - Ex: dilute 0.2mg into 20cc – 10mcg/cc and give 1-2cc at a time.
  - \***Avoid in Pulm HTN, intracardiac shunts, Ischemic heart dx, aortopathy, right heart failure.**<sup>3,5</sup>

3. Meng ML, Arendt KW. *Anesthesiology*. 2021

5. Girnius A, Meng ML. *Journal of Cardiothoracic and Vascular Anesthesia*. 2021

23. Kovacheva VP, Soens MA, Tsen LC. *Anesthesiology*. 2015



# Significant Cardiac Effects of OB Meds

## Carboprost (Hemabate)

- Prostaglandin F2 $\alpha$ -agonist
  - Induces significant pulmonary vasoconstriction if bronchospasm - can double PVR
  - Induces bronchoconstriction - can cause bronchospasm → V/Q mismatch, shunting, ↑ PVR
    - **Avoid** in asthma, Pulm HTN, intracardiac shunts, right heart failure

## Misoprostol (Cytotec)

- not for emergency use, no significant CV effects, used prophylactically

# Significant Cardiac Effects of OB Meds

## **Terbutaline:**

- Uterine relaxant
- Selective B<sub>2</sub> agonist
- ↑HR ↑ contractility ↓SVR
  - **Avoid in HOCM and hx tachyarrhythmias**

## • **Nitroglycerin:**

- Rapid uterine relaxation via N.O. mediated smooth muscle relaxation
- Rapid ↓SVR, resultant tachycardia
  - **Caution in HOCM, AS, R→L shunt, etc**



# Additional Considerations

## Epidural IV Test Dose & Tachycardia:

- Epi 15 mcg
- Risk of tachycardia may be problematic in:
  - hx arrhythmias, HCM, AS, MS, or severe aortopathy
- Could instead use fentanyl 50-100 mcg and ask pt to report any s/s of IV opioids.

## Uterine Eversion & RV Strain

- Uterus lifted up and out of abdomen for repair after C/S entrains micro air-emboli, which upon entering pulmonary circulation will ↑ PVR.
  - Theoretical risk of consequent increased RV strain.
- Can repair uterus in situ to mitigate this risk in pts with RV strain, pulm HTN, or shunt lesions.

Education | July 2021

# Obstetric Anesthesia and Heart Disease: Practical Clinical Considerations **FREE**

Marie-Louise Meng, M.D.; Katherine W. Arendt, M.D.

+ Author and Article Information

*Anesthesiology* July 2021, Vol. 135, 164–183.

<https://doi.org/10.1097/ALN.0000000000003833>

Slide Notes:

Recommend saving this article to your phone. Table 4 (next slide) has concise delivery management key points for specific cardiac conditions as a quick reference.



**Table 4** has concise delivery management key points for specific cardiac conditions as a quick reference.

3.Meng ML, Arendt KW. *Anesthesiology*. 2021

# Objectives:

Current Data on Maternal Mortality

Peripartum Hemodynamics Review

Maternal Cardiac Risk Scoring Systems  
& Heart Failure Identification Tools

Heart Failure from  
Preeclampsia & Peripartum Cardiomyopathy

Delivery Management Considerations in:

- Heart Failure
- Pulmonary Edema

A large, dark, triangular-shaped ultrasound image of a lung, showing a granular texture, serves as the background for the title. The title text is white and positioned on the left side of the image.

# **Delivery Management Considerations For Pulmonary Edema**

# Pulmonary Edema in Pregnancy

## Parturients with heart disease are at high risk of pulmonary edema.

- Cardiogenic: heart failure, or fluid overload (can be iatrogenic)
- Non-cardiogenic: increased vascular permeability 2/2 damaged endothelium.
- Mixed: Parturient with PPCM and PreE/Eclampsia

## Heart failure can increase pulmonary capillary hydrostatic pressure

- further increased risk of transudation of fluid into pulmonary interstitium.

## Preparation for potential emergent delivery in acute pulmonary edema 2/2 heart failure includes:

- Rapid management of hypoxemia to prevent maternal or fetal compromise
- diuresis, +/- inotropic support, O<sub>2</sub> supplementation and possible tracheal intubation.



# DIAGNOSTIC TESTS

## PULMONARY EDEMA

BNP

TTE, POCUS: PLA,  
PSA, 4-chamber  
view

POCUS lung US

CXR

ECG

High BNP suggests  
HF and warrants  
further workup →  
TTE, POCUS

BNP > 100

NT- proBNP >300

# LUNG US vs CXR in HF

Original Investigation | Emergency Medicine

## Diagnostic Accuracy of Point-of-Care Lung Ultrasonography and Chest Radiography in Adults With Symptoms Suggestive of Acute Decompensated Heart Failure A Systematic Review and Meta-analysis

Anna M. Maw, MD, MS; Ahmed Hassanin, MD; P. Michael Ho, MD, PhD; Matthew D. F. McInnes, MD, PhD; Angela Moss, MS; Elizabeth Juarez-Colunga, PhD; Nilam J. Soni, MD, MS; Marcelo H. Miglioranza, MD, MHSC, PhD; Elke Platz, MD, MS; Kristen DeSanto, MSLS, MS, RD; Anthony P. Sertich, MD; Gerald Salame, MD; Stacie L. Daugherty, MD, MSPH

### 2019 SRMA: 6 prospective cohort studies, 1827 patients

- Lung US more sensitive than CXR for detection of cardiogenic pulmonary edema, with similar specificity:
  - 15% absolute difference in sensitivity found between LUS and CXR (0.88 vs 0.73) ( $P < .001$ )
    - In other words: Q 100 patients c/c dyspnea 2/2 cardiogenic pulmonary edema, LUS can dx 15 more cases than CXR without increase in false (+).
- Additional Benefits: LUS easier to see real-time edema resolution, as confirmatory CXR images can lag behind both development and resolution. There is no radiation exposure, and learning curve for use is quick.

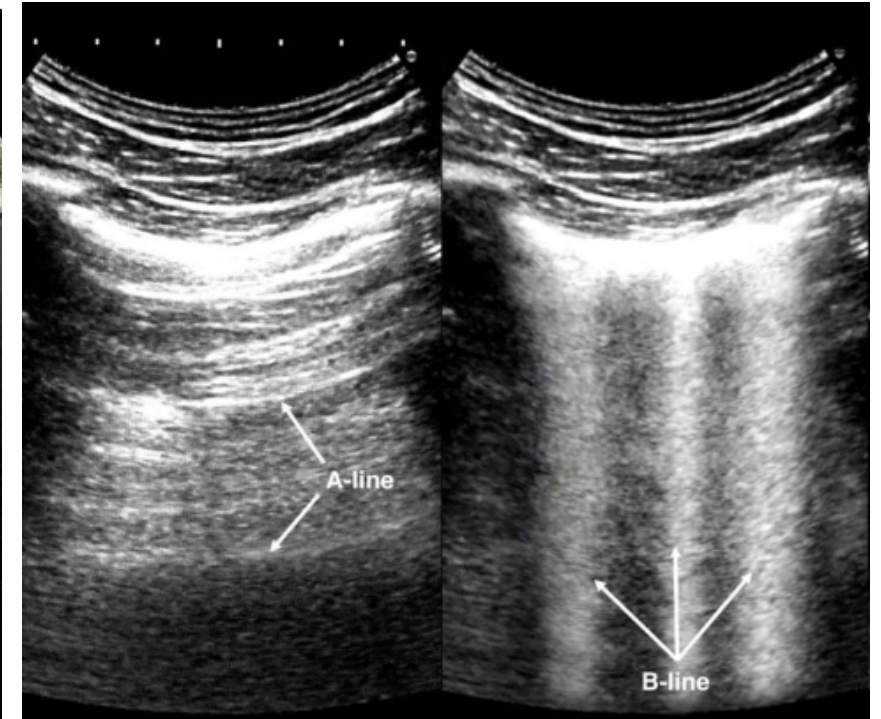
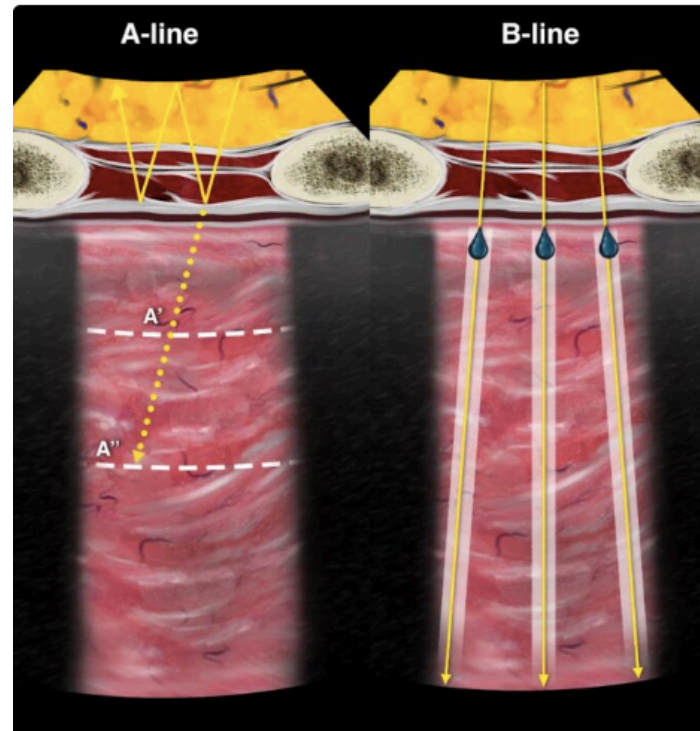
# POCUS LUNG ULTRASOUND

## **A-LINE:**

horizontal evenly-spaced  
hyperechoic reverberation air  
artifacts of the pleural line.<sup>25</sup>

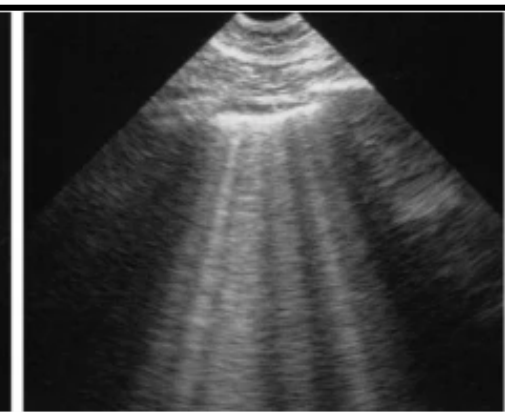
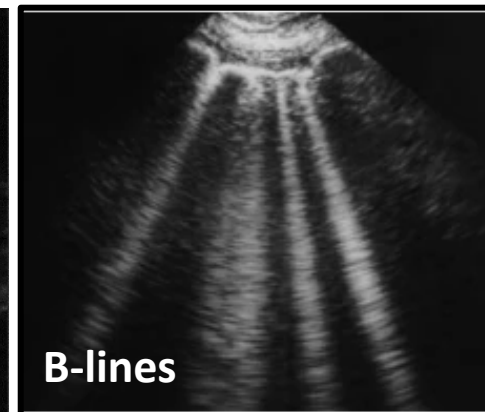
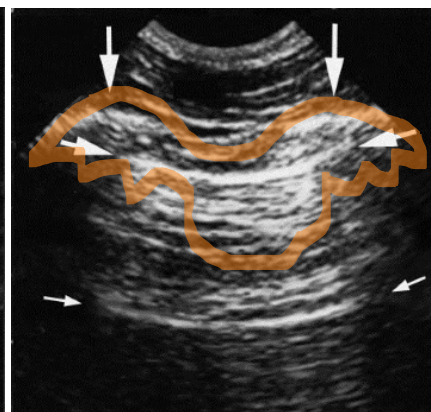
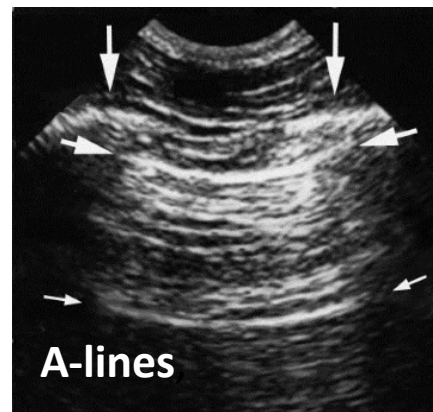
## **B-LINE:**

well-defined, hyperechoic water  
artifacts extending from the  
pleural line down to bottom of  
screen without fading, obliterating  
A-lines.<sup>25</sup>



25. NYSORA. Acute respiratory failure: The BLUE protocol. *NYSORA*. 2023.

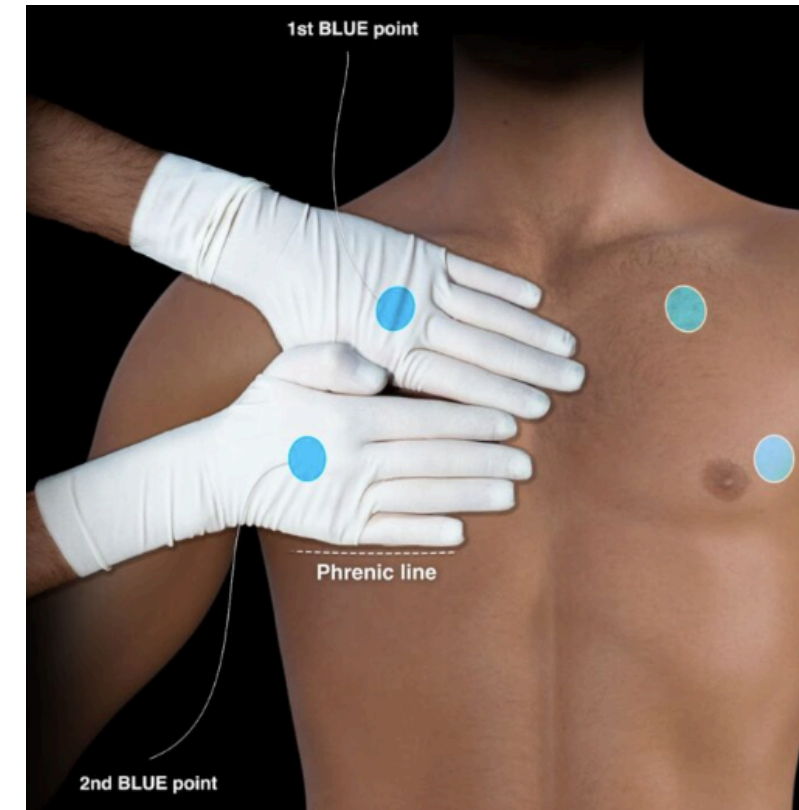
“bat sign” of rib shadows  
and pleural line



# POCUS LUNG ULTRASOUND

Indicator cephalad with linear or curvilinear probe

- 3+ B-lines in one field = “wet” lung.
  - 1-3 B-lines in bases can be normal finding in pregnancy.
- Visualized lung sliding\*, with:
  - **3+ B-lines at all 4 points** → likely pulmonary edema
    - *lack of lung sliding suggests pneumonia*
    - ❖ [BLUE protocol – Bedside Lung Ultrasound in Emergency]





# Pulmonary Edema: *Maternal Optimization*

## Diuresis

- Careful with high doses in PreE – intravascularly dry patient 2/2 capillary leak.

## Supplemental O2

- HFNC, CPAP, BiPAP: +PEEP

## Fluid restriction during delivery (C/S or VD)

- Careful IVF titration
- Smaller reconstitution doses for antibiotics (eg: 20cc syringe slow IVP vs 250cc NS bag for azithromycin)
- **Pitocin:** (usually 10u over 10min ~ 166cc in 30u/500cc bag)
  - Instead:
    - “Rule of 3s”: 10u Pitocin in 10cc NS: Administer 3u Q3min to effect.
    - OBGYN can give IU Pitocin in C/S

# Pulmonary Edema: *Maternal Optimization*

## HYPERTENSION MANAGEMENT

- In PreE: IV BBs, hydralazine, or oral nifedipine often 1<sup>st</sup> line anti-HTN agents.
- However, **when PreE is associated with pulmonary edema, IV nitroglycerin is preferred**.<sup>18</sup>
- ***Especially*** in Pre-E/HTN with LV systolic dysfunction/PPCM:
  - Titrated NTG gtt can offload ventricles without affecting contractility, while inotropes (eg: dobutamine) can augment contractility and improve forward flow.
    - NTG benefit: tighter HD control with rapid onset and short  $E_t^{1/2}$
    - Dose: **IV gtt: 5 mcg/min, titrated Q 3–5 min to max dose of 100 mcg/min**.<sup>18</sup>
      - Initiation of high-dose BBs may be detrimental in setting of poor LV contractile function/HR-dependent CO.

**IMPORTANCE OF BNP → +/- TTE to assess which agents would be most beneficial if myocardial dysfunction present.**

# PPCM: Where to Look

**WE MUST BE AWARE OF POSSIBILITY OF HEART FAILURE IF S/S PRESENT PERIOD IN PREGNANT OR POSTPARTUM WOMEN**

- ✓ S/S on labor floor :
  - ↓SpO2, tachycardia, positioning, pt c/o not feeling right:
    - ✓ Maternal Early Warning Criteria/ACOG criteria/Fett Self Test.
    - ✓ Pulmonary edema: Cardiac workup, BNP, +/-TTE.
- ✓ Postpartum tubals, D&C for PPH – can occur in days, weeks PP
- ✓ Postpartum non-obstetric elective surgery/procedure
- ✓ Ante-/Postpartum urgent or emergent surgery

# PPCM: Where to Look

- In women of childbearing age, asking pre-op if any recent pregnancies:
  - If within 6mos, prudent to ask pointed cardiac questions:
    - ✓ METs, SOB, palpitations, chest pain, abnormal fatigue, s/s at rest vs active, peripheral edema, able to lay flat at night? Etc..

After 6wk PP checkup with OBGYN,  
we may be the *only* other providers women see during PP  
period who know to look for this.



# “Fett Self Test” for Early Identification of Heart Failure

> 4 pts → further workup (BNP & TTE)<sup>27</sup>

Table 1. Self-test for early diagnosis of peripartum cardiomyopathy.

Symptoms	0 points	1 point	2 points
Orthopnea	None	Need to elevate head	Need to elevate 45 degrees or more
Dyspnea	None	Climbing 8 or more steps	Walking on level
Unexplained cough	None	At night	Day and night
Lower extremity swelling	None	Below knee	Above and below knee
Excessive weight gain during last month of pregnancy	Under 2 pounds per week	2–4 pounds per week	Over 4 pounds per week
Palpitations	None	When lying down at night	Day and night, any position

The presence of 4 or more points should prompt additional investigation.  
Data taken from [30].

26: Davis M, Duvernoy C. *Women's Health*. 2015

In validation study: **100% of women who presented with > 4pts had LV systolic dysfunction.**<sup>27</sup>

26. Davis M, Duvernoy C. Peripartum Cardiomyopathy: Current Knowledge and Future Directions. *Women's Health*. 2015.

27. Fett JD. Validation of a self-test for early diagnosis of heart failure in peripartum cardiomyopathy. *Critical Pathways in Cardiology*. 2011;10(1):44-45. doi:<https://doi.org/10.1097/HPC.0b013e31820b887b>

“Although it is possible that a fulminant myocarditis/cardiomyopathy can suddenly appear without prior warning and awareness, *almost all of these women, upon reflection, can recognize that they experienced signs and symptoms earlier by days and weeks.*”

“My incessant theme is this: Physicians, nurses and patients must be alert to the possibility that *a young woman, despite the lack of any type of heart problem in her medical history, may develop a serious cardiomyopathy with acute onset of heart failure in the setting of pregnancy*”

- James D Fett, MD, MPH

## TLDR: Key Points Summary

- Cardiomyopathy is a significant cause of maternal death in the United States.
- During pregnancy and postpartum, heart failure from either HTN or PPCM can develop rapidly and acutely over just days, in otherwise low-risk women. The condition is both under-screened and under-diagnosed.
- Pulmonary edema may be presenting sign of heart failure, and pulmonary ultrasound is a fast and reliable diagnostic tool.
- The significantly increased cardiac demands of labor, delivery, and **especially the postpartum period**, can push an impaired ventricle into failure.
- Uterotonic and tocolytic agents given peripartum can have **significant and deleterious cardiac effects in patients with heart failure**.
- Pregnancy-induced heart failure develops most commonly within 6 months postpartum, so it is prudent to include directed cardiac questions in pre-op eval for women who have recently been pregnant.
  - Fett Test is a useful, validated tool.

## References

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