



OPEN ACCESS

Review article

Cardiovascular adverse events in pregnancy: A global perspective

Susy Kotit*, Magdi Yacoub

ABSTRACT

Pregnant women with heart disease are vulnerable to many adverse cardiovascular events (AE). AEs during and after pregnancy continue to be important causes of maternal mortality and morbidity worldwide, with huge variations in burden in different countries and regions. These AEs are classified as having direct or indirect causes, depending on whether they are directly caused by pregnancy or due to some pre-existing disease and/or non-obstetric cause, respectively.

The risks continue throughout pregnancy and even after childbirth. Apart from immediate complications during pregnancy, there is increasing evidence of a significant link between several events and the risk of cardiovascular disease (CVD) later in life.

A significant number of pregnancy-related deaths caused by cardiovascular disease are preventable. This prevention can be realized through increasing awareness of cardiovascular AE in pregnancy, coupled with the application of strategies for prevention and treatment. Knowledge of the risks associated with CVD and pregnancy is of extreme importance in that regard.

We discuss the global distribution of cardiovascular maternal mortality, adverse events during and after pregnancy, their predictors and risk stratification. In addition, we enumerate possible solutions, particularly the role of cardio-obstetric clinics.

Aswan Heart Centre, Aswan, Egypt
*Email: susykotit@hotmail.com

[https://doi.org/
10.21542/gcsp.2021.5](https://doi.org/10.21542/gcsp.2021.5)

Received: 13 February 2021
Accepted: 21 March 2021
© 2021 The Author(s), licensee
Magdi Yacoub Institute. This is an
open access article distributed un-
der the terms of the Creative Com-
mons Attribution license CC BY-4.0,
which permits unrestricted use, dis-
tribution and reproduction in any
medium, provided the original work
is properly cited.

Cite this article as: Kotit S, Yacoub M. Cardiovascular adverse events in pregnancy: A global perspective, *Global Cardiology Science and Practice* 2021;5
<https://doi.org/10.21542/gcsp.2021.5>

INTRODUCTION

Pregnant women with heart disease have pronounced vulnerability to adverse cardiovascular events. Diagnosis and treatment of heart disease in pregnancy is difficult due to similarities between disease manifestations and normal physiological changes.

Cardiovascular adverse events (AEs), during and after pregnancy continue to be important causes of maternal mortality and morbidity worldwide, with huge variations in burden in different countries and regions.^{1,2} These AEs are classified as having direct or indirect causes, depending on whether they are directly caused by pregnancy or due to some pre-existing disease and/or non-obstetric cause, respectively.³⁻⁵

Cardiovascular disease is the single largest cause of indirect maternal mortality,^{3,6} accounting for over 33% of pregnancy-related maternal deaths.⁶⁻¹¹ Additionally, maternal heart disease complicates up to 4% of pregnancies¹²⁻¹⁴ and up to 16% of pregnancies in women with previous cardiac conditions,¹³ with risk depending on the underlying cardiac condition.^{13,15-17}

Over 50% of maternal deaths occur post-partum.¹⁸ Late maternal mortality is defined as death more than 42 days (and up to one year) after child birth.¹⁹ Importantly, the currently cited figures almost certainly constitute an underestimate.¹⁹

It is estimated that up to 68% pregnancy-related deaths caused by cardiovascular conditions are preventable.^{6,20} This can be achieved, through increasing awareness, coupled with applying strategies for prevention and treatment. Knowledge of the risks associated with CVD and pregnancy is of extreme importance in that regard.

We discuss the global distribution of cardiovascular maternal mortality, the adverse events during and after pregnancy, their predictors and risk stratification. In addition, we enumerate possible solutions, particularly the role of cardio-obstetric clinics.

Incidence and global distribution of maternal mortality

According to the World Health Organization (WHO)²¹ and Global Burden of Disease (GBD),^{1,2} in 2017 there were up to 295,000 maternal deaths globally, leading to an estimated global maternal mortality rate (MMR) of 211 per 100,000 live births.

In 2017, every day approximately 810 women died from preventable causes related to pregnancy and childbirth, with 94% of all maternal deaths occurring in low and middle-income countries (LMIC's).²² Sub-Saharan Africa alone accounted for roughly two-thirds of the estimated global maternal deaths.²¹ In Egypt the number of maternal deaths was 1,316 in 2017 with a MMR of 62 per 100,000 livebirths.^{8,23} The regional distribution and pattern of maternal death reported by the GBD varies considerably around the world.²³⁻²⁵ (Figure 1).

Causes of maternal mortality and burden of disease

Up to 1990, direct causes accounted for over two-thirds of maternal mortality worldwide. There has been a significant decrease in overall maternal mortality over the years due to progressive drops in direct causes, however, the contribution of indirect and late maternal deaths, as well as maternal hypertensive disorders, has remained unchanged (Figure 2).^{1,2}

Regional and national variation in incidence and causes of maternal mortality

According to the GBD, maternal mortality rate (MMR) varied from 1 to 496 per 100,000 live births (Figure 3) in 2017.^{1,2} The alarmingly high incidence in MMR in some countries continues, with very few exceptions.

The regional incidence and cause of maternal mortality varies considerably from one country to another (Table 1). In addition, the representation of maternal hypertensive

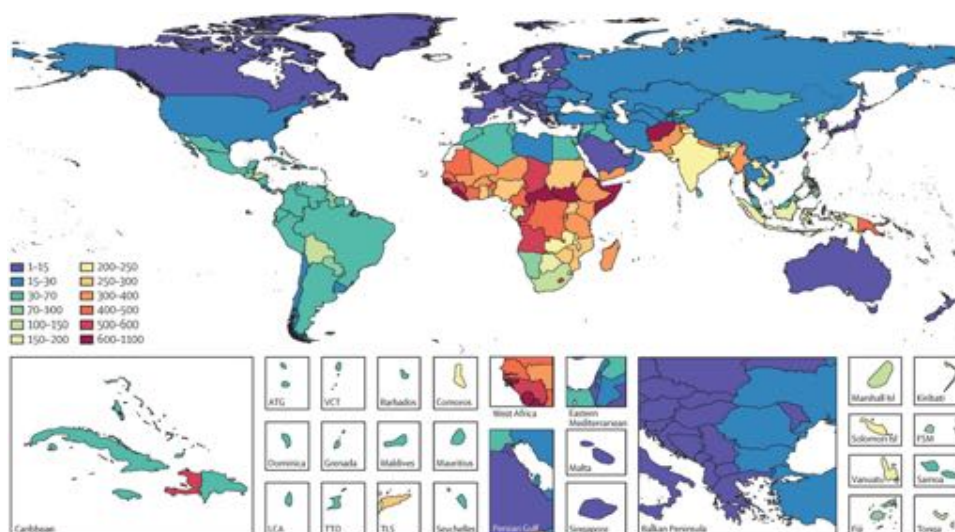


Figure 1. Maternal mortality ratio (MMR; number of deaths per 100,000 livebirths) for countries and territories, GBD 2015.²³

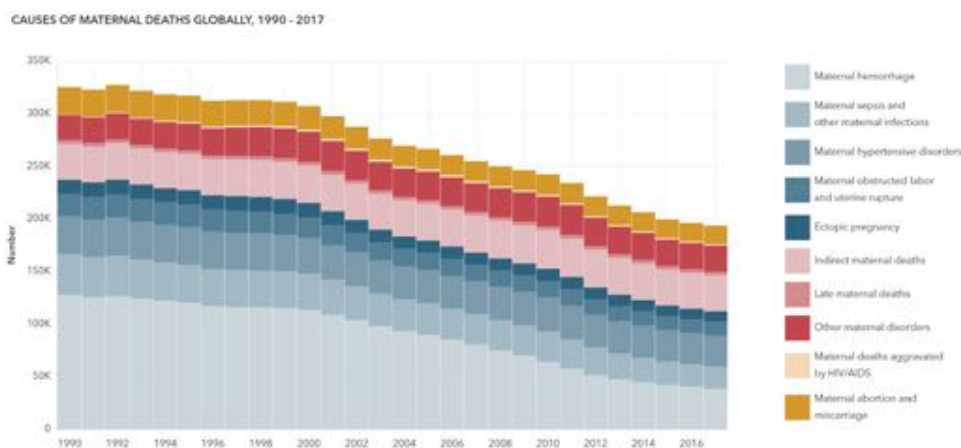


Figure 2. Causes of maternal deaths globally, 1990–2017.^{1,2} (A) maternal hypertensive disorders: High blood pressure during pregnancy in women who did not already have hypertension, or preeclampsia in women with preexisting hypertension. (B) Indirect maternal deaths: Deaths due to preexisting conditions made worse by physiologic effects of pregnancy. (C) Late maternal deaths: Deaths due to any cause that occurs six weeks to 12 months after pregnancy. (D) Other maternal disorders: All other direct maternal disorders, including anemia in pregnancy, gestational diabetes and embolism.

disorders, indirect maternal deaths as well as the reported late maternal mortality also differs across countries.

Details of adverse cardiovascular events

Direct maternal adverse cardiovascular events

Hypertensive disorders of pregnancy. Chronic hypertension, gestational hypertension and preeclampsia²⁶ are important causes of maternal and perinatal morbidity and mortality,^{9,27–31} particularly toward the end of pregnancy (Figure 4). Novel diagnostic methods and therapies have recently been reviewed.³²

Gestational hypertension is defined as new-onset hypertension arising after 20 weeks of gestation^{33,34} and occurs in 10% of women.³⁵ It is associated with acute and chronic

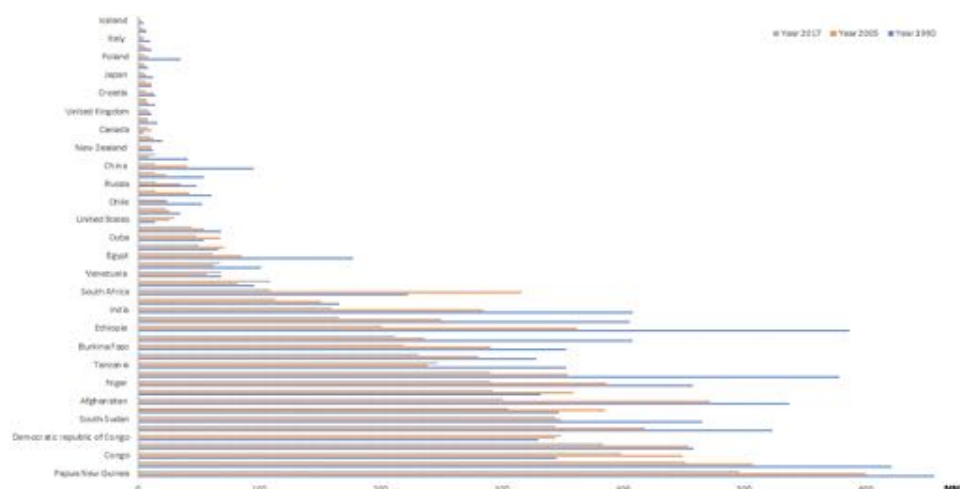


Figure 3. MMR per country across 1990, 2005 and 2017.^{1,2}

cardiovascular changes leading to an increased risk for hypertension throughout life,^{36,37} and a 4-fold increased risk of future maternal cardiovascular events.³⁷⁻⁴⁰

Indirect maternal adverse cardiovascular events

Cardiovascular disease is the single largest cause of indirect maternal mortality.^{3,6} Patients who experience complications during pregnancy may also be at higher risk of cardiac events later in life.⁴⁴

Arrhythmias

The altered cardiac anatomy during pregnancy can elicit new onset arrhythmia or prompt the recurrence of preexisting arrhythmias.^{45,46} An increased incidence of cardiac arrhythmias has been reported during pregnancy⁴⁷ in patients with and without identifiable heart disease. Arrhythmias are responsible for complications in 67 per 100,000 pregnancies,⁴⁹ mainly in the form of atrial fibrillation (27 per 100,000 pregnancies) and supraventricular tachycardia (22 per 100,000 pregnancies).⁴⁸⁻⁵⁰

Heart failure

Heart failure (HF) remains the most common complication during pregnancy among all women with heart disease, regardless of the cardiac pathology.^{42,51} Heart failure can occur during, or immediately after pregnancy (Figure 5). Despite the poor prognosis associated with the diagnosis of HF during pregnancy, and the fact that prevalence of HF among pregnant women has increased over the years - particularly in the post-partum period,⁵² data in the literature are scarce.

Peri-partum cardiomyopathy

Peri-partum cardiomyopathy (PPCM) is an idiopathic cardiomyopathy presenting with heart failure secondary to left ventricle systolic dysfunction towards the end of pregnancy or in the months following child birth.⁵³⁻⁵⁶ At diagnosis, the majority of patients have severe symptoms (NYHA III/IV) and LVEF < 35%, with regional variations in presentation and outcomes⁵⁷⁻⁵⁹ (Figure 6).

The incidence of PPCM varies markedly from 1-100 per 10,000 live births depending on the region.^{59,61-63} PPCM has the highest mortality rate in pregnancy, with a worldwide mortality of 2.4%,⁵⁷ although it may be underdiagnosed. PPCM leads to substantial maternal and neonatal morbidity and mortality, with less than half of all cases recovering

Table 1 Global distribution in incidence, causes and timing of maternal mortality by MMR (2017).^{1,2} The reported late maternal deaths represented 21.9% of total maternal mortality in Chile, compared to 0.08% in Iceland.

Country	Number of maternal deaths (2017)	MMR (2017)	Top mortality disorder	Maternal hypertensive disorders (% total maternal deaths)	Indirect maternal deaths (% total maternal deaths)	Late maternal deaths (% total maternal deaths)
Globally	193,639	140	Other maternal disorders	15.17	17.61	1.74
Iceland	0	1	Other maternal disorders	0.08	0.03	0.08
Sweden	4	3	Other maternal disorders	7.25	2.50	7.50
Italy	17	4	Other maternal disorders	7.82	1.29	1.65
Spain	20	5	Other maternal disorders	11.40	1.20	1.50
Poland	16	5	Other maternal disorders	6.31	0.88	6.50
Australia	15	5	Other maternal disorders	5.07	1.53	9.07
Japan	46	5	Other maternal disorders	7.87	9.93	3.50
Netherlands	11	6	Other maternal disorders	11.09	2.36	2.91
Croatia	2	6	Other maternal disorders	5.50	7.00	8.00
France	52	7	Other maternal disorders	11.27	9.54	2.54
United Kingdom	62	8	Other maternal disorders	9.11	14.56	6.27
Portugal	7	8	Other maternal disorders	11.14	4.29	0.86
Canada	31	8	Other maternal disorders	6.10	12.26	16.94
South Korea	42	10	Other maternal disorders	10.24	6.98	2.29
New Zealand	6	11	Other maternal disorders	1.33	37.00	12.17
Latvia	3	14	Other maternal disorders	1.67	1.67	5.33
China	2241	14	Other maternal disorders	14.80	6.80	0.98
Vietnam	213	14	Maternal hemorrhage	14.63	13.50	2.77
Russia	250	15	Other maternal disorders	11.33	12.35	2.08
Kazakhstan	53	15	Other maternal disorders	8.32	6.53	7.66
Chile	56	23	Late maternal death	17.27	19.84	21.91
Iran	298	23	Indirect maternal disease	10.58	33.20	9.95
United States	1171	30	Other maternal disorders	10.12	15.65	17.12
Mexico	1120	44	Maternal hemorrhage	23.45	19.17	7.22
Cuba	53	48	Indirect maternal disease	7.09	21.43	17.06
Colombia	427	50	Indirect maternal disease	23.16	24.17	11.71
Egypt	1316	62	Other maternal disorders	15.76	19.68	3.67
Brazil	2,054	68	Maternal hypertensive disorders	20.93	18.23	6.23
Venezuela	395	69	Maternal hemorrhage	26.26	22.81	2.83
Dominican Republic	237	109	Hypertensive disorders	26.21	17.34	9.92
South Africa	1191	109	Indirect maternal disease	15.44	35.03	1.89
Uganda	1753	113	Indirect maternal disease	15.22	22.27	0.72
India	39,428	160	Other maternal disorders	11.50	13.16	1.78
Indonesia	6627	165	Maternal hemorrhage	26.69	2.19	1.41
Ethiopia	7451	201	Maternal hemorrhage	15.56	17.38	0.73
Mozambique	2090	212	Indirect maternal disease	15.07	25.13	5.29
Burkina Faso	1863	219	Indirect maternal disease	12.40	28.46	0.74
Nigeria	17982	231	Maternal hemorrhage	5.22	16.79	1.68
Tanzania	4916	247	Maternal hypertensive disorders	24.61	21.65	0.79
Mali	2559	291	Indirect maternal disease	6.63	31.18	1.23

(continued on next page)

Table 1 (continued)

Country	Number of maternal deaths (2017)	MMR (2017)	Top mortality disorder	Maternal hypertensive disorders (% total maternal deaths)	Indirect maternal deaths (% total maternal deaths)	Late maternal deaths (% total maternal deaths)
Niger	2930	291	Maternal sepsis and other maternal infections	2.00	26.25	0.87
Kenya	3990	292	Indirect maternal disease	15.70	23.88	1.19
Afghanistan	4095	301	Indirect maternal disease	9.11	38.79	0.66
Cote d'Ivoire	2631	305	Indirect maternal disease	9.47	28.27	0.67
South Sudan	1410	344	Maternal hemorrhage	5.95	8.82	0.62
Somalia	2363	344	Maternal hemorrhage	9.10	16.98	0.56
Democratic republic of Congo	10166	349	Indirect maternal disease	10.99	26.59	0.68
Chad	2745	383	Maternal hemorrhage	6.29	14.41	0.78
Congo	522	398	Indirect maternal disease	11.60	26.63	0.80
Guinea	1916	451	Maternal hemorrhage	8.35	25.25	0.82
Papua New Guinea	1532	496	Indirect maternal disease	8.86	34.92	0.59

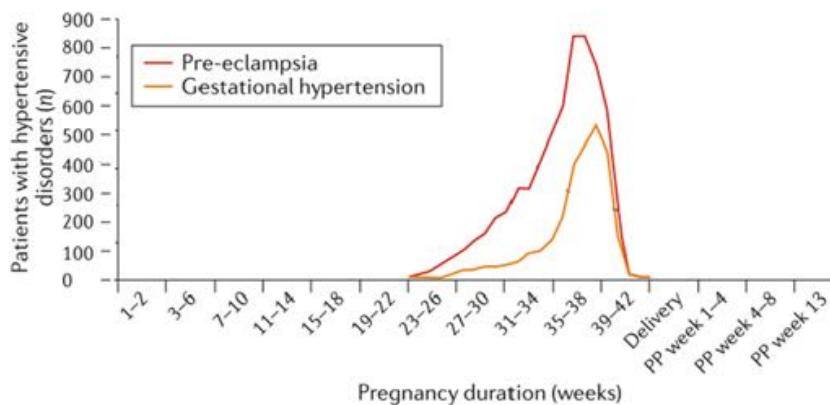


Figure 4. Onset of hypertensive disorders at different stages of pregnancy and postpartum (PP) among women without chronic hypertension (adapted from Ramlakhan et al).^{41,43}



Figure 5. Timing of heart failure in women with structural heart disease at different stages of pregnancy and postpartum (PP) (adapted from Ramlakhan et al).^{41,42}

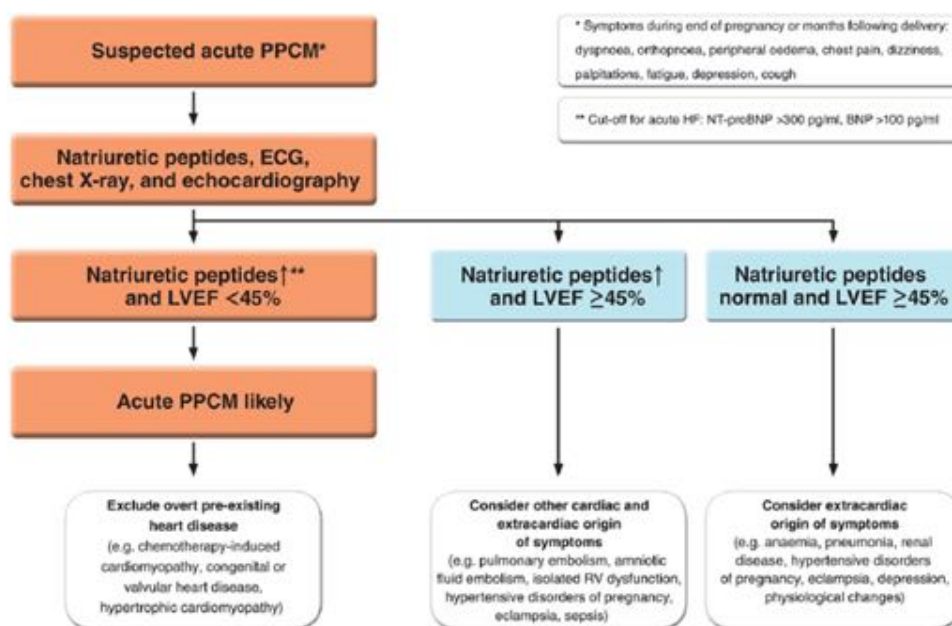


Figure 6. Diagnostic pathway in patients with suspected peripartum cardiomyopathy (PPCM). BNP, B-type natriuretic peptide; ECG, electrocardiogram; HF, heart failure; LVEF, left ventricular ejection fraction; NT-proBNP, N-terminal pro-B-type natriuretic peptide; RV, right ventricular.⁶⁰

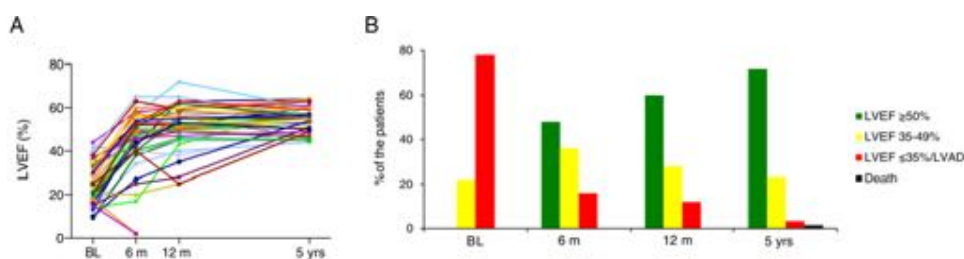


Figure 7. Time course of left ventricular function (adapted from.⁶⁸) (A) Changes in left ventricular ejection fraction (LVEF) from baseline (BL) to 5-year follow-up. Remarkably, LVEF further improves even after 1 year. (B) Proportion of patients with full cardiac recovery constantly increases. At 5-year follow-up, 72% had recovered completely and 23% partially. No recovery was observed in 5%. Death occurs up to 5 years.

full cardiac function^{58,64-67} (Figure 7). Six-month mortality is around 6%, mainly due to heart failure.⁵⁸ Neonatal death is around 5%, although with marked regional variation.⁶³

Management follows the guidelines of HF. In non-responsive patients, other pharmacological agents - such as bromocriptine and prolactin-blocking therapy with dopamine D₂ receptor agonists - have been tried with variable results.⁶⁸

Contributing factors to PPCM include genetic predisposition⁶⁹ as well as auto-immune responses.⁷⁰⁻⁷² Using such factors, all be it indirectly, to define phenocusters⁷³ - it could be possible to identify novel therapeutic targets to guide personalized medicine in PPCM.

For a small proportion of patients with rapidly-progressive PPCM resistant to conventional therapy, the use of the current generation of left ventricular assist devices can give long-lasting “cures” in some patients (personal experience in Harefield and Aswan series).^{74,75}

Registries on the condition will provide fundamental data on predisposing factors, potential aetiologia and regional variations.^{57,63,76,77}

Mechanical valve thrombosis

Even with the right care, the incidence of thromboembolism during pregnancy is estimated from 7–23% with half of these cases being mechanical valve thrombosis (MVT),⁷⁸ which is associated with 20% mortality.⁷⁹ Thrombosis is the most life-threatening complication for women with prosthetic heart valve, during pregnancy.^{78,80} The chance of a successful uncomplicated pregnancy, which depends on the balance between the thrombotic and bleeding risks, is around 57%.

Acute coronary syndromes

The incidence of coronary artery disease (CAD) in women of childbearing age is unclear and varies between countries.^{81,82} CAD, is a major cause of maternal death and accounts for over 20% of maternal cardiac deaths,⁸³ especially in the form of acute coronary syndromes (ACSs).⁸⁴ The estimated incidence of 6.2 per 100,000 deliveries^{85–88} nearly 4 times higher than in non-pregnant women⁸⁹ and reflects the growing prevalence of cardiovascular risk factors in the pregnant population.⁹⁰

Risk stratification

Several tools have been developed to estimate morbidity and mortality risk in pregnant women with cardiac disease, such as CARPREG¹³ and ZAHARA.¹² However, the best estimate of the risk of cardiovascular events during pregnancy in pre-existing heart disease is the WHO's⁹¹ (Figure 8), as it integrates congenital and acquired heart disease. The risk of cardiac adverse events during pregnancy are significantly increased in mWHO IV compared to mWHO I (Figure 9).

Based on the modified WHO risk classification for milder conditions, where the risk of pregnancy is very low to moderate, the needed care might be limited to a few visits during pregnancy, while in case of high risk of complications, a more frequent follow-up schedule is recommended. Women in the highest risk group (WHO IV) should be advised against pregnancy. In case of pregnancy, strict monitoring is required.⁹² More comparative studies should be performed in order to define the most accurate risk index for pregnant women with heart disease.^{91,95}

The role of comprehensive cardio-obstetric clinics

The concept of multidisciplinary cardio-obstetric clinics (Figure 10) has evolved and been applied in several countries, with extremely encouraging results^{96–114} and excellent survival rates of mothers even with complex diseases, and their offspring.²⁴

However, the need for cardio-obstetric clinics remains un-met, with most programs to be found in developed countries and with only a few centers continuing follow-up long term.^{24,101,102}

Although these initiatives might be feasible in large metropolitan areas, smaller towns, rural communities and remote regions are completely neglected. There is an urgent need for action and worldwide implementation of cardio-obstetric clinics, strategically placed, in order to reach the majority of those in need.

Integrated, tailored and dynamic healthcare services responding to current state of disease burden and initiatives are essential. Cardio-obstetric care clinics (COcare) have been initiated in the metropolitan Aswan Heart Center as well as its rural branch in Ballana.¹¹⁶

CONCLUSIONS

Preventable maternal cardiovascular adverse events continue to be a global problem with an unacceptably high burden of disease. This requires urgent concerted efforts

WHO Pregnancy Risk Classification (Risk of pregnancy by medical condition)	Cardiovascular Conditions by WHO Risk Class
WHO Risk Class I No detectable increased risk of maternal mortality and no or mild increase in morbidity.	<ul style="list-style-type: none"> Uncomplicated, small or mild <ul style="list-style-type: none"> Pulmonary stenosis Patient ductus arteriosus Mitral valve prolapse Successfully repaired simple lesions (atrial or ventricular septal defect, patent ductus arteriosus, anomalous pulmonary venous drainage). Atrial or ventricular ectopic beats, isolated
WHO Risk Class II (If otherwise well and uncomplicated) Small increased risk of maternal mortality or moderate increase in morbidity.	<ul style="list-style-type: none"> Unoperated atrial or ventricular septal defect Repaired tetralogy of Fallot Most arrhythmias
WHO Risk Class II or III (Depending on individual) Risk as indicated in Class II (above) or Class III (below).	<ul style="list-style-type: none"> Mild left ventricular impairment Hypertrophic cardiomyopathy Native or tissue valvular heart disease not considered WHO I or IV Marfan syndrome without aortic dilatation Aorta <45 mm in aortic disease associated with bicuspid aortic valve Repaired Coarctation
WHO Risk Class III 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.	<ul style="list-style-type: none"> Mechanical valve Systemic right ventricle Fontan circulation Cyanotic heart disease (unrepaired) Other complex congenital heart disease Aortic dilatation 40-45 mm in Marfan Syndrome Aortic dilatation 45-50 mm in aortic disease associated with bicuspid aortic valve
WHO Risk Class IV (Pregnancy contraindicated) 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.	<ul style="list-style-type: none"> Pulmonary arterial hypertension of any cause Severe systemic ventricular dysfunction (LVEF <30%, NYHA III-IV) Previous peripartum cardiomyopathy with any residual impairment of left ventricular function Severe symptomatic mitral or aortic stenosis Marfan syndrome with aorta dilated >45 mm Aortic dilation >50 mm in aortic disease associated with bicuspid aortic valve Native severe Coarctation

Figure 8. Modified WHO classification of maternal cardiovascular risk: application.⁹²

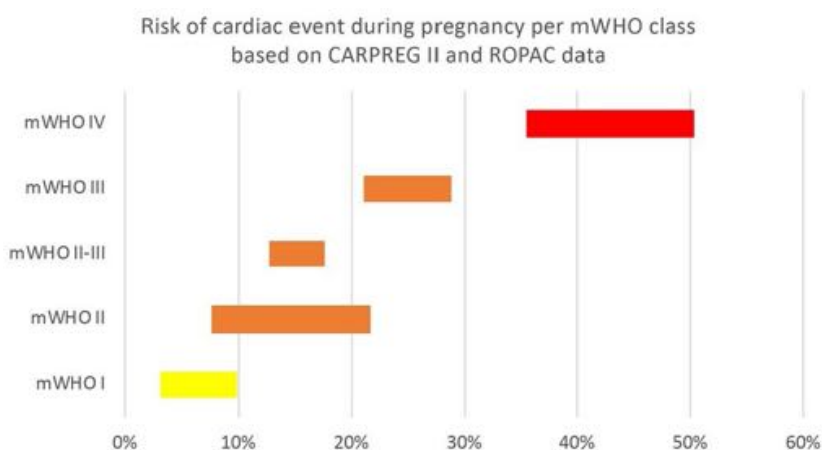


Figure 9. Risk of cardiac event during pregnancy per mWHO class based on CARPREG II and ROPAC data. 12,13,93 adapted from.⁹⁴

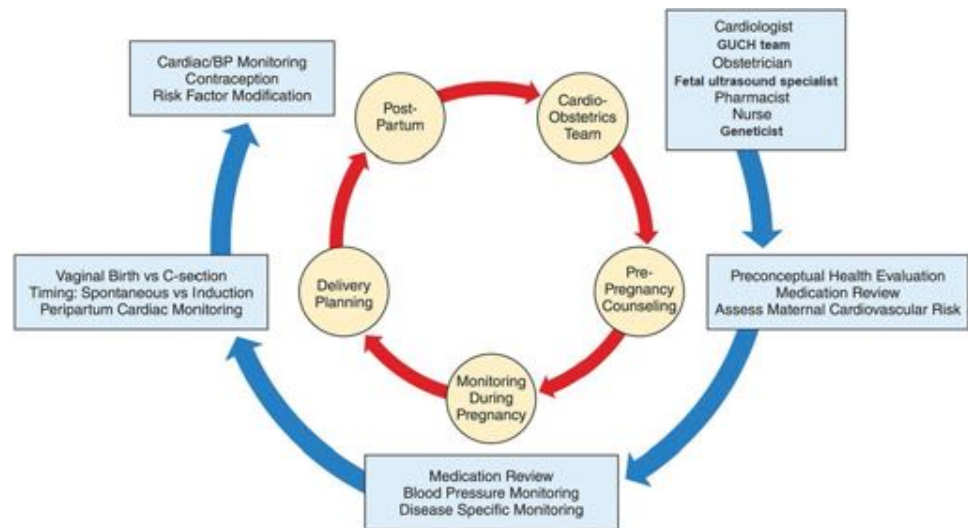


Figure 10. Cardio-obstetrics team in the management of women before pregnancy, during pregnancy, and postpartum (GUCH: grown-up congenital heart disease) Adapted from.¹¹⁵

from governments, individuals and professionals, who have first-hand experience of the magnitude of the problem. Multidisciplinary cardio-obstetric care should also play an important role.

REFERENCES

- [1] IHME. Findings from the Global Burden of Disease Study 2017 (GBD 2017). In: *Inst Heal Metrics Eval (IHME) Seattle*. WA IHME; 2018.
- [2] Maternal Health Atlas. <https://maternalhealthatlas.org/> (Accessed 4 December 2020).
- [3] Nair M, Nelson-Piercy C, Knight M. Indirect maternal deaths: UK and global perspectives. *Obstet Med*. 2017;10(1):10–15 doi: 10.1177/1753495X16689444.
- [4] Storm F, Agampodi S, Eddleston M, Sørensen JB, Konradsen F, Rheinländer T. Indirect causes of maternal death. *Lancet Glob Heal*. 2014;2(10):e566 doi: 10.1016/S2214-109X(14)70297-9.
- [5] 2012. The WHO Application of ICD-10 to Deaths during Pregnancy, Childbirth and the Puerperium: ICD-MM. <http://www.who.int/about/licensing/> (Accessed 13 January 2021).
- [6] Knight M, Kenyon S, Brocklehurst P, Neilson J, Shakepeare J, Kurinczuk JJ. Saving Lives, Improving Mothers' Care Lessons learned to inform future maternity care from the UK and Ireland confidential enquiries into maternal deaths and morbidity 2009-2012. *Published online*. 2012;2012:1–120.
- [7] Regitz-Zagrosek V, Blomstrom Lundqvist C, Borghi C, et al. ESC Guidelines on the management of cardiovascular diseases during pregnancy. *Eur Heart J*. 2011;32(24):3147–3197 doi: 10.1093/eurheartj/ehr218.
- [8] Alkema L, Chou D, Hogan D, et al. Global, regional, and national levels and trends in maternal mortality between 1990 and 2015, with scenario-based projections to 2030: a systematic analysis by the un maternal mortality estimation inter-agency group. *Lancet*. 2016;387(10017):462–474 doi: 10.1016/S0140-6736(15)00838-7.
- [9] Khan KS, Wojdyla D, Say L, Gülmezoglu AM, Van Look PF. WHO analysis of causes of maternal death: a systematic review. *Lancet*. 2006;367(9516):1066–1074 doi: 10.1016/S0140-6736(06)68397-9.
- [10] Macdorman MF, Declercq E, Cabral H, Morton C. Recent increases in the U.S. maternal mortality rate: disentangling trends from measurement issues. *Obstet Gynecol*. 2016;128(3):447–455 doi: 10.1097/AOG.0000000000001556.
- [11] Ferranti EP, Jones EJ, Hernandez TL. Pregnancy reveals evolving risk for cardiometabolic disease in women. *JOGNN - J Obstet Gynecol Neonatal Nurs*. 2016;45(3):413–425 doi: 10.1016/j.jogn.2016.02.004.
- [12] Drenthen W, Boersma E, Balci A, et al. Predictors of pregnancy complications in women with congenital heart disease. *Eur Heart J*. 2010;31(17):2124–2132 doi: 10.1093/eurheartj/ehq200.
- [13] Silversides CK, Grewal J, Mason J, et al. Pregnancy outcomes in women with heart disease: the CARPREG II study. *J Am Coll Cardiol*. 2018;71(21):2419–2430 doi: 10.1016/j.jacc.2018.02.076.
- [14] Roos-Hesselink JW, Ruys TPE, Stein JI, et al. Outcome of pregnancy in patients with structural or ischaemic heart disease: results of a registry of the european society of cardiology. *Eur Heart J*. 2013;34(9):657–665 doi: 10.1093/eurheartj/ehs270.
- [15] Khursheed R, Tabasum AZB. Maternal and fetal outcome in pregnancies [9]complicated with maternal cardiac diseases: experience at tertiary care hospital. *Internet J Gynecol Obstet*. 2015;19(1):01–06.

- [16] McFaul PB, Dornan JC, Lamki HBD. Pregnancy complicated by maternal [10]heart disease. A review of 519 women. *BJOG An Int J Obstet Gynaecol*. 1988;95(9):861–67.
- [17] Regitz-Zagrosek V, Roos-Hesselink JW, Bauersachs J, et al. ESC Guidelines for the management of cardiovascular diseases during pregnancy. *Eur Heart J*. 2018;39(34):3165–3241 doi: 10.1093/eurheartj/ehy340.
- [18] Lameijer H, Schutte JM, Pieper PG, Van Roosmalen JJM, Schuitemaker NWE. Maternal mortality due to cardiovascular disease in the Netherlands: a 21-year experience for Dutch Maternal Mortality and Morbidity Committee. *Netherlands Hear J*. 2020;28:27–36 doi: 10.1007/s12471-019-01340-w.
- [19] Sliwa K, Anthony J. Late maternal deaths: a neglected responsibility. *Published online 2016*. 2016; doi: 10.1016/S0140-6736(16)00651-6.
- [20] Building U.S.. Capacity to review and prevent maternal deaths: report from nine maternal mortality review committees. https://reviewtoaction.org/Report_fro.
- [21] WHO. *Trends in maternal mortality: 2000 to 2017: estimates by WHO, UNICEF, UNFPA, World Bank Group and the United Nations Population Division*. Geneva: World Health Organization; WHO 2019; 2019.
- [22] WHO. 2021. Maternal mortality. <https://www.who.int/news-room/fact-sheets/detail/maternal-mortality> (Accessed 21 January 2021).
- [23] Kassebaum NJ, Barber RM, Dandona L, et al. Global, regional, and national levels of maternal mortality, 1990–2015: a systematic analysis for the Global Burden of Disease Study 2015. *Lancet*. 2016;388(10053):1775–1812 doi: 10.1016/S0140-6736(16)31470-2.
- [24] Sliwa K, Libhaber E, Elliott C, et al. Spectrum of cardiac disease in maternity in a low-resource cohort in South Africa. *Heart*. 2014;100(24):1967–1974 doi: 10.1136/heartjnl-2014-306199.
- [25] Roth GA, Mensah GA, Johnson CO, et al. Global burden of cardiovascular diseases and risk factors, 1990–2019. *J Am Coll Cardiol*. 2020;76(25):2982–3021 doi: 10.1016/j.jacc.2020.11.010.
- [26] Rana S, Lemoine E, Granger J, Karumanchi SA. Preeclampsia: pathophysiology, challenges, and perspectives. *Circ Res*. 2019;124(7):1094–1112 doi: 10.1161/CIRCRESAHA.118.313276.
- [27] Duley L. The global impact of pre-eclampsia and eclampsia. *Semin Perinatol*. 2009;33(3):130–137 doi: 10.1053/j.semperi.2009.02.010.
- [28] Steegers EAP, Von Dadelszen P, Duvekot JJ, Pijnenborg R. Pre-eclampsia. *The Lancet*. 2010;376:631–644 doi: 10.1016/S0140-6736(10)60279-6.
- [29] Sammour MB, El-Kabarity H, Fawzy MM, Schindler AE. 2011. WHO recommendations for prevention and treatment of Pre-Eclampsia and Eclampsia. Vol 97. World Health Organization; 2011. <https://www.ncbi.nlm.nih.gov/books/NBK140561/> (Accessed 6 November 2011).
- [30] Peters RM, Flack JM. Hypertensive disorders of pregnancy. *JOGNN - J Obstet Gynecol Neonatal Nurs*. 2004;33(2):209–220 doi: 10.1177/0884217504262970.
- [31] Creanga AA, Syverson C, Seed K, Callaghan WM. Pregnancy-related mortality in the United States, 2011–2013. *Obstet Gynecol*. 2017;130(2):366–373 doi: 10.1097/AOG.0000000000002114.
- [32] Phipps EA, Thadhani R, Benzing T, Karumanchi SA. Pre-eclampsia: pathogenesis, novel diagnostics and therapies. *Nat Rev Nephrol*. 2019;15(5):275–289 doi: 10.1038/s41581-019-0119-6.
- [33] O'Brien TE, Ray JG, Chan WS. Maternal body mass index and the risk of preeclampsia: A systematic overview. *Epidemiology*. 2003;14(3):368–374 doi: 10.1097/00001648-200305000-00020.
- [34] Robertson RM. Women and cardiovascular disease. *Circulation*. 2001;103(19):2318–2320 doi: 10.1161/01.CIR.103.19.2318.
- [35] Ros HS, Cnattingius S, Lipworth L. Comparison of risk factors preeclampsia and gestational hypertension in a population-based cohort study. *Am J Epidemiol*. 1998;147(11):1062–1070 doi: 10.1093/oxfordjournals.aje.a009400.
- [36] Wilson BJ, Watson MS, Prescott GJ, et al. Hypertensive diseases of pregnancy and risk of hypertension and stroke in later life: Results from cohort study. *Br Med J*. 2003;326(7394):845–849 doi: 10.1136/bmj.326.7394.845.
- [37] Garovic VD, Bailey KR, Boerwinkle E, et al. Hypertension in pregnancy as a risk factor for cardiovascular disease later in life. *J Hypertens*. 2010;28(4):826–833 doi: 10.1097/HJH.0b013e328335c29a.
- [38] Ray JG, Vermeulen MJ, Schull MJ, Redelmeier DA. Cardiovascular health after maternal placental syndromes (CHAMPS): Population-based retrospective cohort study. *Lancet*. 2005;366(9499):1797–1803 doi: 10.1016/S0140-6736(05)67726-4.
- [39] Bellamy L, Casas JP, Hingorani AD, Williams DJ. Pre-eclampsia and risk of cardiovascular disease and cancer in later life: Systematic review and meta-analysis. *Br Med J*. 2007;335(7627):974–977 doi: 10.1136/bmj.39335.385301.BE.
- [40] Roccella EJ. Report of the national high blood pressure education program working group on high blood pressure in pregnancy. *Am J Obstet Gynecol*. 2000;183(1):s1–s22 doi: 10.1067/mob.2000.107928.
- [41] Ramlakhan KP, Johnson MR, Roos-Hesselink JW. Pregnancy and cardiovascular disease. *Nat Rev Cardiol*. 2020;17(11):718–731 doi: 10.1038/s41569-020-0390-z.
- [42] Ruys TEP, Roos-Hesselink JW, Hall R, et al. Heart failure in pregnant women with cardiac disease: Data from the ROPAC. *Heart*. 2014;100(3):231–238 doi: 10.1136/heartjnl-2013-304888.
- [43] Shiozaki A, Matsuda Y, Satoh S, Saito S. Comparison of risk factors for gestational hypertension and preeclampsia in Japanese singleton pregnancies. *J Obstet Gynaecol Res*. 2013;39(2):492–499 doi: 10.1111/j.1447-0756.2012.01990.x.
- [44] Balint OH, Siu SC, Mason J, et al. Cardiac outcomes after pregnancy in women with congenital heart disease. *Heart*. 2010;96(20):1656–1661 doi: 10.1136/hrt.2010.202838.

- [45] Brodsky M, Doria R, Allen B, Sato D, Thomas G, Sada M. New-onset ventricular tachycardia during pregnancy. *Am Heart J*. 1992;123(4 PART 1):933–941 doi: [10.1016/0002-8703\(92\)90699-V](https://doi.org/10.1016/0002-8703(92)90699-V).
- [46] Meah VL, Cockcroft JR, Backx K, Shave R, Stöhr EJ. Cardiac output and related haemodynamics during pregnancy: a series of meta-analyses. *Heart*. 2016;102(7):518–526 doi: [10.1136/heartjnl-2015-308476](https://doi.org/10.1136/heartjnl-2015-308476).
- [47] CK S. Physiological changes in pregnancy. In: Warn CA, Oakley C, eds. *Hear Dis pregnancy Massachusetts Blackwell Publ 2007..*
- [48] Shotan A, Ostrzega E, Mehra A, Johnson JV, Elkayam U. Incidence of arrhythmias in normal pregnancy and relation to palpitations, dizziness, and syncope. *Am J Cardiol*. 1997;79(8):1061–1064 doi: [10.1016/S0002-9149\(97\)00047-7](https://doi.org/10.1016/S0002-9149(97)00047-7).
- [49] Vaidya VR, Arora S, Patel N, et al. Burden of Arrhythmia in Pregnancy. *Circulation*. 2017;135(6):619–621 doi: [10.1161/CIRCULATIONAHA.116.026681](https://doi.org/10.1161/CIRCULATIONAHA.116.026681).
- [50] Li JM, Nguyen C, Joglar JA, Hamdan MH, Page RL. Frequency and outcome of arrhythmias complicating admission during pregnancy: experience from a high-volume and ethnically-diverse obstetric service. *Clin Cardiol*. 2008;31(11):538–541 doi: [10.1002/clc.20326](https://doi.org/10.1002/clc.20326).
- [51] Sliwa K, Anthony J. Decompensated heart failure in pregnancy. *Card Fail Rev*. 2016;2(1):20 doi: [10.15420/cfr.2015.24.2](https://doi.org/10.15420/cfr.2015.24.2).
- [52] Mogos MF, Piano MR, McFarlin BL, Salemi JL, Liese KL, Briller JE. Heart failure in pregnant women: a concern across the pregnancy continuum. *Circ Heart Fail*. 2018;11(1): doi: [10.1161/CIRCHEARTFAILURE.117.004005](https://doi.org/10.1161/CIRCHEARTFAILURE.117.004005).
- [53] Hilfiker-Kleiner D, Sliwa K. Pathophysiology and epidemiology of peripartum cardiomyopathy. *Nat Rev Cardiol*. 2014;11(6):364–370 doi: [10.1038/nrcardio.2014.37](https://doi.org/10.1038/nrcardio.2014.37).
- [54] Azibani F, Sliwa K. Peripartum cardiomyopathy: an update. *Curr Heart Fail Rep*. 2018;15(5):297–306 doi: [10.1007/s11897-018-0404-x](https://doi.org/10.1007/s11897-018-0404-x).
- [55] Ricke-Hoch M, Pfeffer TJ, Hilfiker-Kleiner D. Peripartum cardiomyopathy: Basic mechanisms and hope for new therapies. *Cardiovasc Res*. 2020;116(3):520–531 doi: [10.1093/cvr/cvz252](https://doi.org/10.1093/cvr/cvz252).
- [56] Jackson AM, Dalzell JR, Walker NL, Coats CJ, Jhund PS, Petrie MC. Peripartum cardiomyopathy: diagnosis and management. *Heart*. 2018;104(9):779–786 doi: [10.1136/heartjnl-2016-310599](https://doi.org/10.1136/heartjnl-2016-310599).
- [57] Sliwa K, Mebazaa A, Hilfiker-Kleiner D, et al. Clinical characteristics of patients from the worldwide registry on peripartum cardiomyopathy (PPCM): EURObservational Research Programme in conjunction with the Heart Failure Association of the European Society of Cardiology Study Group on PPCM. *Eur J Heart Fail*. 2017;19(9):1131–1141 doi: [10.1002/ejhf.780](https://doi.org/10.1002/ejhf.780).
- [58] Sliwa K, Petrie MC, Meer P, VanDer, et al. Clinical presentation, management, and 6-month outcomes in women with peripartum cardiomyopathy: An ESC EORP registry. *Eur Heart J*. 2020;41(39):3787–3797 doi: [10.1093/eurheartj/ehaa455](https://doi.org/10.1093/eurheartj/ehaa455).
- [59] 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(7):827–843 doi: [10.1002/ejhf.1493](https://doi.org/10.1002/ejhf.1493).
- [60] Bauersachs J, König T, 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(7):827–843 doi: [10.1002/ejhf.1493](https://doi.org/10.1002/ejhf.1493).
- [61] 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(8):767–778 doi: [10.1093/eurjhf/hfq120](https://doi.org/10.1093/eurjhf/hfq120).
- [62] Demakis JG, Rahimtoola SH. Peripartum cardiomyopathy. *Circulation*. 1971;44(5):964–968 doi: [10.1161/01.CIR.44.5.964](https://doi.org/10.1161/01.CIR.44.5.964).
- [63] Sliwa K, Petrie MC, van der Meer P, et al. Clinical presentation, management, and 6-month outcomes in women with peripartum cardiomyopathy: an ESC EORP registry. *Eur Heart J*. 2020;41(39):3787–3797 doi: [10.1093/eurheartj/ehaa455](https://doi.org/10.1093/eurheartj/ehaa455).
- [64] Blauwet LA, Libhaber E, Forster O, Tibazarwa K, Mebazaa A, Hilfiker-Kleiner DKS. Predictors of outcome in 176 South African patients with peripartum cardiomyopathy. *Heart*. 2013;99:308–313.
- [65] Haghikia A, Podewski E, Libhaber E, Labidi S, Fischer D, Roentgen PTD, Jordan J, Lichthagen R, Kaisenberg CS, Struman I, Bovy N, Sliwa KB, H-KD J. Phenotyping and outcome on contemporary management in a German cohort of patients with peripartum cardiomyopathy. *Basic Res Cardiol*. 2013;2013:108366.
- [66] Irizarry OC, Levine LD, Lewey J, Boyer T, Riis V, Elovitz MAAZ. Comparison of clinical characteristics and outcomes of peripartum cardiomyopathy between African American and non-African American women. *JAMA Cardiol*. 2017;21256.
- [67] McNamara DM, Elkayam U, Alharethi R, et al. Clinical outcomes for peripartum cardiomyopathy in North America: Results of the IPAC study (Investigations of Pregnancy-Associated Cardiomyopathy). *J Am Coll Cardiol*. 2015;66(8):905–914 doi: [10.1016/j.jacc.2015.06.1309](https://doi.org/10.1016/j.jacc.2015.06.1309).
- [68] Moulig V, Pfeffer TJ, Ricke-Hoch M, et al. Long-term follow-up in peripartum cardiomyopathy patients with contemporary treatment: low mortality, high cardiac recovery, but significant cardiovascular co-morbidities. *Eur J Heart Fail*. 2019;21(12):1534–1542 doi: [10.1002/ejhf.1624](https://doi.org/10.1002/ejhf.1624).
- [69] Ware JS, Li J, Mazaika E, et al. Shared genetic predisposition in peripartum and dilated cardiomyopathies. *N Engl J Med*. 2016;374(3):233–241 doi: [10.1056/nejmoa1505517](https://doi.org/10.1056/nejmoa1505517).

- [70] Caforio ALP, Adler Y, Agostini C, Allanore Y, Anastasakis A, Arad MBM, Charron P, Elliott PM, Eriksson U, Felix SB, Garcia-Pavia P, Hachulla E, Heymans S, Imazio M, Klingel K, Marcolongo R, Cerinic MM, Pantazis A, Plein S, Poli V, Rigopoulos A, Seferovic P, Shoenfeld Y, Zamorano JLLA. Diagnosis and management of myocardial involvement in systemic immune-mediated diseases: a position statement of the European Society of Cardiology Working Group on Myocardial and Pericardial Disease. *Eur Heart J*. 2017;38:2649–2662.
- [71] Warraich RS, Sliwa K, Damasceno A, et al. Impact of pregnancy-related heart failure on humoral immunity: Clinical relevance of G3-subclass immunoglobulins in peripartum cardiomyopathy. *Am Heart J*. 2005;150(2):263–269 doi: 10.1016/j.ahj.2004.09.008.
- [72] Ansari AA, Fett JD, Carraway RE, Mayne AE, Onlamoon N, Sundstrom JB. Autoimmune mechanisms as the basis for human peripartum cardiomyopathy. *Clin Rev Allergy Immunol*. 2002;23(3):301–324 doi: 10.1385/CRIAL:23:3:301.
- [73] Verdonschot JAJ, Merlo M, Dominguez F, et al. Phenotypic clustering of dilated cardiomyopathy patients highlights important pathophysiological differences. *Eur Heart J*. 2021;42(2):162–174 doi: 10.1093/eurheartj/ehaa841.
- [74] Oosterom L, Jonge NDe, Kirkels JH, Klöpping C, Lahpor JR. Left ventricular assist device as a bridge to recovery in a young woman admitted with peripartum cardiomyopathy. *Netherlands Heart J*. 2008;16(12):426–428 doi: 10.1007/BF03086192.
- [75] Hayworth R. From labor and delivery to left ventricular assist device: A peripartum cardiomyopathy case report. *Hear Lung*. 2020;49(1):54–57 doi: 10.1016/j.hrtlng.2019.06.005.
- [76] Sliwa K, Hilfiker-Kleiner D, Mebazaa A, et al. EURObservational research programme: A worldwide registry on peripartum cardiomyopathy (PPCM) in conjunction with the heart failure association of the European Society of Cardiology working group on PPCM. *Eur J Heart Fail*. 2014;16(5):583–591 doi: 10.1002/ejhf.68.
- [77] PeriPartum CardioMyopathy (PPCM) Registry. <https://www.escardio.org/Research/Registries-&-surveys/Observational-researchprogramme/PeriPartum-CardioMyopathy-PPCM-Registry> (Accessed 26 January 2021).
- [78] Elkayam U, Bitar F. Valvular heart disease and pregnancy - Part II: Prosthetic valves. *J Am Coll Cardiol*. 2005;46(3):403–410 doi: 10.1016/j.jacc.2005.02.087.
- [79] Van Hagen IM, Roos-Hesselink JW, Ruys TPE, et al. Pregnancy in women with a mechanical heart valve. *Circulation*. 2015;132(2):132–142 doi: 10.1161/CIRCULATIONAHA.115.015242.
- [80] Mahgoub A, Kotit S, Bakry K, Magdy A, Hosny H, Yacoub M. Thrombosis of mechanical mitral valve prosthesis during pregnancy: An ongoing saga in need of comprehensive solutions. *Glob Cardiol Sci Pract*. 2020;2020(3): doi: 10.21542/gcsp.2020.32.
- [81] Petitti DB, Sidney S, Quesenberry CP, Bernstein A. Incidence of stroke and myocardial infarction in women of reproductive age. *Stroke*. 1997;28(2):280–283 doi: 10.1161/01.STR.28.2.280.
- [82] Etchill E, Clarke N, Giuliano K, Lawton J, Choi CW. Post-partum spontaneous coronary artery dissection refractory to conservative management. *Glob Cardiol Sci Pract*. 2020;2020(3): doi: 10.21542/gcsp.2020.34.
- [83] Knight M, Nair M, Tuffnell D, et al. *Saving Lives, Improving Mothers' Care Maternal, Newborn and Infant Clinical Outcome Review Programme*. 2009.
- [84] Cantwell R, Clutton-Brock T, Cooper G, et al. Saving Mothers' Lives: Reviewing maternal deaths to make motherhood safer: 2006–2008. *BJOG An Int J Obstet Gynaecol*. 2011;118:1–203 doi: 10.1111/j.1471-0528.2010.02847.x.
- [85] Ginz B. MYOCARDIAL INFARCTION IN PREGNANCY. *BJOG An Int J Obstet Gynaecol*. 1970;77(7):610–615 doi: 10.1111/j.1471-0528.1970.tb03578.x.
- [86] Ladner HE, Danielsen B, Gilbert WM. Acute myocardial infarction in pregnancy and the puerperium: A population-based study. *Obstet Gynecol*. 2005;105(3):480–484 doi: 10.1097/01.AOG.0000151998.50852.31.
- [87] James AH, Jamison MG, Biswas MS, Branciazio LR, Swamy GK, Myers ER. Acute myocardial infarction in pregnancy: A United States population-based study. *Circulation*. 2006;113(12):1564–1571 doi: 10.1161/CIRCULATIONAHA.105.576751.
- [88] Bush N, Nelson-Piercy C, Spark P, Kurinczuk JJ, Brocklehurst P, Knight M. Myocardial infarction in pregnancy and postpartum in the UK. *Eur J Prev Cardiol*. 2013;20(1):12–20 doi: 10.1177/1741826711432117.
- [89] Caspi J, Pettitt TW, Sperrazza C, Mulder TSA. Reimplantation of anomalous left coronary artery from the pulmonary artery without mitral valve repair. *Ann Thorac Surg*. 2007;84:619–23.
- [90] James AH, Jamison MG, Biswas MS, Branciazio LR, Swamy GK, Myers ER. Acute myocardial infarction in pregnancy: A United States population-based study. *Circulation*. 2006;113(12):1564–1571 doi: 10.1161/CIRCULATIONAHA.105.576751.
- [91] Balci A, Sollie-Szarynska KM, Van Der Bijl AGL, et al. Prospective validation and assessment of cardiovascular and offspring risk models for pregnant women with congenital heart disease. *Heart*. 2014;100(17):1373–1381 doi: 10.1136/heartjnl-2014-305597.
- [92] European ES of G (ESG); A for, Medicine PC (AEPC); GS for G, (DGesGM), Regitz-Zagrosek V, Blomstrom Lundqvist C, Al Borghi CE. ESC Guidelines on the management of cardiovascular diseases during pregnancy: the task force on the management of cardiovascular diseases during pregnancy of the European Society of Cardiology (ESC). *Eur Heart J*. 2011;32(24):3147–3197.
- [93] van Hagen IM, Boersma E, Johnson MR, et al. Global cardiac risk assessment in the registry of pregnancy and cardiac disease: results of a registry from the European Society of Cardiology. *Eur J Heart Fail*. 2016;18(5):523–533 doi: 10.1002/ejhf.501.

- [94] Van Hagen IM, Roos-Hesselink JW. Pregnancy in congenital heart disease: Risk prediction and counselling. *Heart*. 2020;106(23):1853–1861 doi: [10.1136/heartjnl-2019-314702](https://doi.org/10.1136/heartjnl-2019-314702).
- [95] Lu CW, Shih JC, Chen SY, et al. Comparison of 3 risk estimation methods for predicting cardiac outcomes in pregnant women with congenital heart disease. *Circ J*. 2015;79(7):1609–1617 doi: [10.1253/circj.CJ-14-1368](https://doi.org/10.1253/circj.CJ-14-1368).
- [96] Davis MB, Walsh MN. Cardio-obstetrics: Team-based care to improve maternal outcomes. *Circ Cardiovasc Qual Outcomes*. 2019;1(12): doi: [10.1161/CIRCOUTCOMES.118.005417](https://doi.org/10.1161/CIRCOUTCOMES.118.005417).
- [97] Magun E, DeFilippis EM, Noble S, et al. Cardiovascular Care for Pregnant Women With Cardiovascular Disease. *J Am Coll Cardiol*. 2020;76(18):2102–2113 doi: [10.1016/j.jacc.2020.08.071](https://doi.org/10.1016/j.jacc.2020.08.071).
- [98] UCSF Pregnancy and Cardiac Treatment (PACT) Program | Obstetrics, Gynecology & Reproductive Sciences. <https://obgyn.ucsf.edu/maternal-fetal-medicine/ucsf-pregnancy-and-cardiac-treatment-pact-program> (Accessed 17 January 2021).
- [99] Penn Pregnancy and Heart Disease Program—Penn Medicine. <https://www.pennmedicine.org/for-patients-and-visitors/find-a-program-or-service/obstetrics/pregnancy-and-heart-disease> (Accessed 17 January 2021).
- [100] Heart Disease in Pregnancy Program | Cardiovascular Medicine | Medical College of Wisconsin. <https://www.mcw.edu/departments/medicine/divisions/cardiovascularmedicine/patient-care/heart-disease-in-pregnancy-program> (Accessed 17 January 2021).
- [101] Perinatology Maternal Cardiac Program | Center for Women’s Health | OHSU. <https://www.ohsu.edu/womens-health/perinatology-maternalcardiac-program> (Accessed 17 January 2021).
- [102] Pregnancy and Cardiac Treatment (PACT) Program | Clinics | UCSF Health. <https://www.ucsfhealth.org/clinics/pregnancy-and-cardiac-treatment-pact-program> (Accessed 17 January 2021).
- [103] Cardiac Obstetrics | Heart Centre. <https://heartcentre.ca/services/cardiac-obstetrics> (Accessed 17 January 2021).
- [104] Pregnancy and heart disease centre | Royal Brompton & Harefield NHS Foundation Trust. <https://www.rbht.nhs.uk/our-services/heart/inherited-cardiovascular-conditions/pregnancy-and-heart-disease-centre> (Accessed 17 January 2021).
- [105] Cardiac antenatal clinic | University Hospitals Bristol NHS Foundation Trust. <http://www.uhbristol.nhs.uk/patients-and-visitors/yourhospitals/bristol-heart-institute-clinical-services/adult-congenital-heartdisease/cardiac-antenatal-clinic/> (Accessed 17 January 2021).
- [106] Sliwa K, Libhaber E, Elliott C, et al. Spectrum of cardiac disease in maternity in a low-resource cohort in South Africa. *Hear*. 2014;1001967–1974.
- [107] Sharma G, Zakaria S, Michos ED, et al. Improving cardiovascular workforce competencies in cardio-obstetrics: current challenges and future directions. *J Am Heart Assoc*. 2020;9(12): doi: [10.1161/JAHA.119.015569](https://doi.org/10.1161/JAHA.119.015569).
- [108] Easter SR, Valente AM, Economy KE. Creating a multidisciplinary pregnancy heart team. *Curr Treat Options Cardiovasc Med*. 2020;22(1): doi: [10.1007/s11936-020-0800-x](https://doi.org/10.1007/s11936-020-0800-x).
- [109] Ouyang P, Sharma G. The potential for pregnancy heart teams to reduce maternal mortality in women with cardiovascular disease. *J Am Coll Cardiol*. 2020;76(18):2114–2116 doi: [10.1016/j.jacc.2020.09.007](https://doi.org/10.1016/j.jacc.2020.09.007).
- [110] Brown HL, Warner JJ, Gianos E, et al. Promoting Risk Identification and Reduction of Cardiovascular Disease in Women Through Collaboration with Obstetricians and Gynecologists: A Presidential Advisory from the American Heart Association and the American College of Obstetricians and Gynecologists. *Obstet Gynecol Surv*. 2018;73(10):574–576 doi: [10.1097/OGX.0000000000000600](https://doi.org/10.1097/OGX.0000000000000600).
- [111] Khairy P, Ouyang DW, Fernandes SM, Lee-Parritz A, Economy KE, Landzberg MJ. Pregnancy outcomes in women with congenital heart disease. *Circulation*. 2006;113(4):517–524 doi: [10.1161/CIRCULATION-AHA.105.589655](https://doi.org/10.1161/CIRCULATION-AHA.105.589655).
- [112] Cardio-Obstetrics Program | Von Voigtlander Women’s Hospital | Michigan Medicine. <https://www.umwomenshealth.org/conditions-treatments/cardio-obstetrics-program> (Accessed 17 January 2021).
- [113] Cardio-Obstetrics Program | NorthShore. <https://www.northshore.org/cardiovascular-institute/our-programs/cardio-obstetrics-program/> (Accessed 17 January 2021).
- [114] Center for Maternal Cardiac Care | TriHealth. <https://www.trihealth.com/institutes-and-services/womens-services/high-risk-pregnancy/center-for-maternal-cardiac-care> (Accessed 17 January 2021).
- [115] Mehta LS, Warnes CA, Bradley E, et al. Cardiovascular considerations in caring for pregnant patients: a scientific statement from the American Heart Association. *Circulation*. 2020;141(23):e884–e903 doi: [10.1161/CIR.0000000000000772](https://doi.org/10.1161/CIR.0000000000000772).
- [116] Ballana Heart Study. <https://myglobalheart.org/wp-content/uploads/2019/03/Ballana-Heart-Study-022819.pdf>.