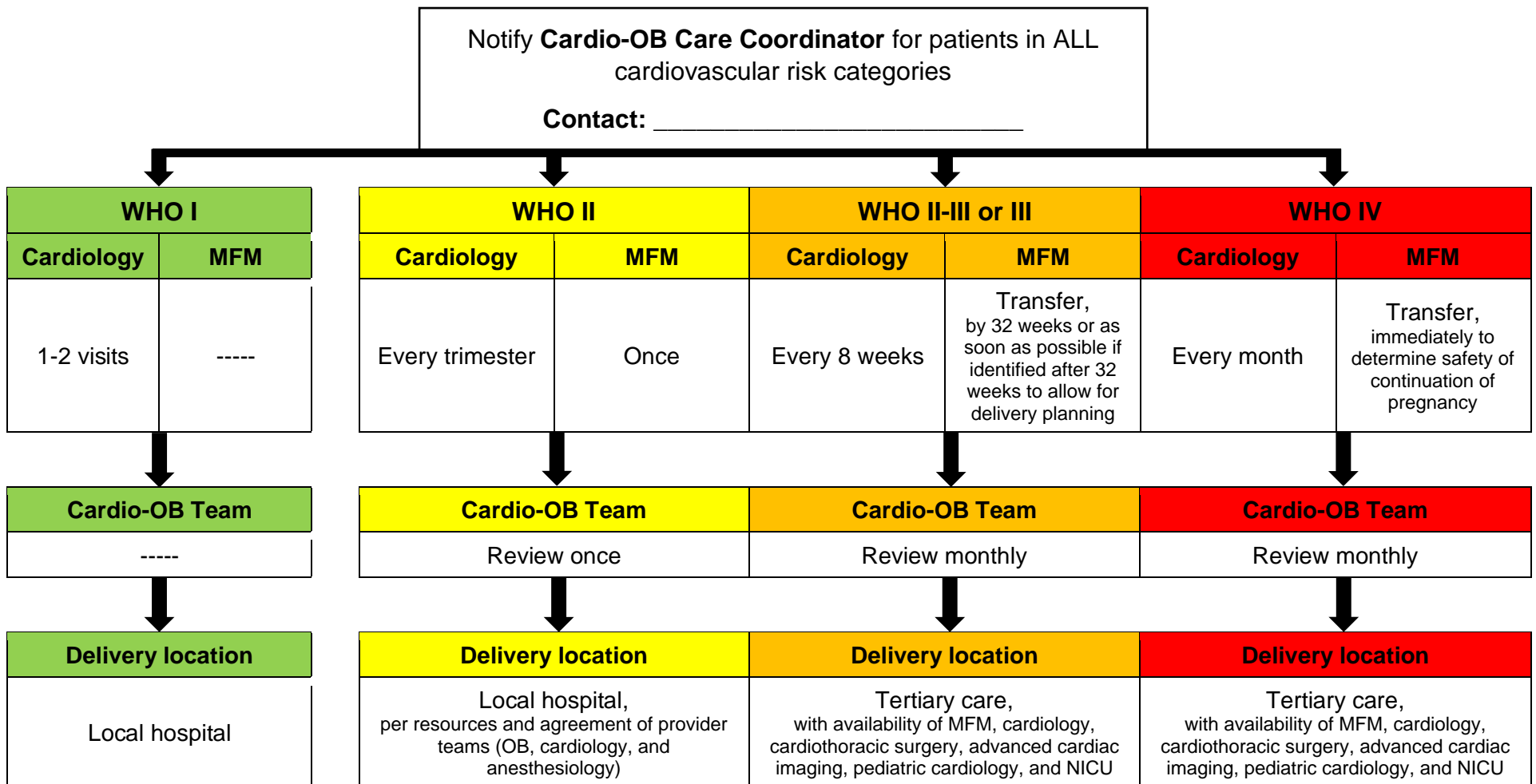




Cardiac Conditions in Obstetric Care

Referral Protocol for Patients with Known Cardiovascular Diagnoses



Cardio-OB Team should include the Cardio-OB Care Coordinator and representatives from cardiology, MFM, obstetrics, anesthesiology, nursing, and pharmacy. Additional team members may include social work, case managers, primary care providers, CT, surgery, neonatology, etc.

Modified World Health Organization (WHO) Classification of Maternal Cardiovascular Risk¹

WHO I	WHO II	WHO II-III	WHO III	WHO IV
<ul style="list-style-type: none"> • Small or mild: <ul style="list-style-type: none"> ○ Pulmonary stenosis ○ Patent 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 	<ul style="list-style-type: none"> • Unoperated atrial or ventricular septal defect • Repaired tetralogy of Fallot • Most arrhythmias (supraventricular arrhythmias) • Turner syndrome without aortic dilatation 	<ul style="list-style-type: none"> • Mild left ventricular impairment (EF >45%) • Hypertrophic cardiomyopathy • Native or tissue valve disease not considered WHO I or IV (mild mitral stenosis, moderate aortic stenosis) • Marfan or other HTAD syndrome without aortic dilatation • Aorta <45 mm in bicuspid aortic valve pathology • Repaired coarctation • Atrioventricular septal defect 	<ul style="list-style-type: none"> • Moderate left ventricular impairment (EF 30–45%) • Previous peripartum cardiomyopathy without residual left ventricular impairment • Mechanical valve • Systemic right ventricle with good or mildly decreased ventricular function • Fontan circulation, if otherwise well and cardiac condition uncomplicated • Unrepaired cyanotic heart disease • Other complex heart disease • Moderate mitral stenosis • Severe asymptomatic aortic stenosis • Moderate aortic dilatation (40–45 mm in Marfan syndrome or other HTAD; 45–50 mm in bicuspid aortic valve, Turner syndrome ASI 20–25 mm/m, tetralogy of Fallot <50 mm) • Ventricular tachycardia 	<ul style="list-style-type: none"> • Pulmonary arterial hypertension • Severe systemic ventricular dysfunction (EF <30%) • Previous peripartum cardiomyopathy with residual left ventricular impairment • Severe mitral stenosis • Severe symptomatic aortic stenosis • Systemic right ventricle with moderate or severely decreased ventricular function • Severe aortic dilatation (>45 mm in Marfan syndrome or other HTAD, >50 mm in bicuspid aortic valve, Turner syndrome ASI >25 mm/m, tetralogy of Fallot >50 mm) • Vascular Ehlers–Danlos • Severe (re)coarctation • Fontan with any complication

¹ Regitz-Zagrosek V, Blomstrom Lundqvist C, Borghi C, et al. 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. doi:10.1093/EURHEARTJ/EHR218