PRE-OP A&P IN PT WITH CARDIAC DISEASES AND



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Topics

Introduction

- GA and Heart performance
- special cardiac risk factors
- Pre-op cardiac assessment & preparation
- **1.** Assessment of overall status
- 2. Cardiac surgery
- 3. Non-cardiac surgery in Pt with cardiac diseases
 - congenital heart diseases (CHD)
 - Heart failure (CHF)
- 4. SBE prophylaxis
- Post-operative care



Introduction

- All anesthetic agents can affect the normal cardiovascular system profoundly and adversely.
- 1- The sinus node, conduction system ,myocardial contractility all can be depressed by general anesthetics.
- 2- These drugs alter both **preload** and **after load** by relaxing vascular smooth-muscle tone.
- 3- General anesthetics also attenuate hypoxic pulmonary vasoconstriction and thereby impair ventilation perfusion matching.

Special cardiac risks for op /GA

- Functional shortage especially heart failure , conductive abnormalities, HTn,
- Structural defects (depends of its functional implication and its interaction to GA agent and operation stress)
- Cardiac surgery or catheterization
- Risk of **SBE** and need for SBE prophylaxis.
- Variable effects of some syndromes that may involve <u>difficult ETI</u>, or presence of <u>autism</u> and other behavioral-communicative disorders.

Inadequate pro-op preparation <u>e.g1</u>:Prolonged pre-op fasting without IVF support [] hypovolemia
 e.g.2:The alkalotic, hypokalemic, hypercalcemic, hypotensive, dilated, digoxin-bound myocardium fibrillates with ease.

Pre-operative Assessment

- History and Examination :
 1- general Hx for the Pt. complaints
 2- relevant Hx :
 - S/S if cardiac performance (preload/congestion , pump action , postload/circulatory shortage)
 - co-morbidities and complications
 - Type of surgery :1- Emergency 2- Elective (curative, pallitive)
- Ped. cardiologist consultation with direct communication with op. team
- Lab, investigations :
 - routine tests including CXR ,
 - relevant test : ECG, ECHO, cath!! etcaccordingly

The Golden rule

Complete and accurate pre-operative judgment of cardiac disease and its patho-physiological implication will lead to predictable intraoperative and postoperative course and thus appropriate management plan

Adequacy of resuscitation, rather than severity of illness at presentation, appears to influence postoperative outcome

ambulatory surgical ?facilities

- Current use of cardiac medications,
- prolonged QT syndrome and
- residual cardiac disease

all <u>disqualify</u> a child from having procedures performed at freestanding ambulatory surgical facilities.



CARDIAC SURGERY

Preoperative Care of the Pediatric Cardiac Surgical Patient

- The Crucial point in this process is the continued communication among medical, surgical, and nursing disciplines.
- Optimal preoperative care involves

 (a) initial stabilization, airway
 management, and establishment of
 vascular access;

Preoperative Care of the Pediatric Cardiac Surgical Patient

(b) A complete and thorough non invasive delineation of the anatomic defect(s);
(c) Resuscitation with evaluation and treatment of secondary organ dysfunction, particularly the brain, kidneys, and liver...
(d) Preparation for cardiac catheterization if necessary.

(e) surgical management when cardiac, pulmonary, renal, and central nervous systems are optimized.

HEART FAILURE

Table 70A.1. Signs of heart failure or low cardiac output states

Signs

Cool extremities/poor perfusion Oliguria and other end-organ failure Tachycardia Hypotension Acidosis Cardiomegaly Pleural effusions

Monitor and measure

Heart rate, blood pressure, intracardiac pressure Extremity temperature, central temperature Urine output Mixed venous oxygen saturation Arterial blood gas pH and lactate Laboratory measures of end-organ function Echocardiography

Ten intensive care strategies to diagnose and support low cardiac output states

- 1) Know the cardiac **anatomy** in detail and its **physiologic consequences**.
- 2) Understand the specialized considerations of the newborn and implications of reparative rather than palliative surgery.
- 3) Diversify personnel to include expertise in neonatal and adult congenital heart disease.
- 4) <u>Monitor(invasive)</u>, <u>measure</u>, and <u>image</u> the heart to rule out <u>residual</u> disease as a cause of postoperative hemodynamic instability or low cardiac output.

Ten intensive care strategies to diagnose and support low cardiac output states

- 5) Maintain aortic **perfusion** and improve the contractile state.
- 6) Optimize preload (including a trial shunting).
- 7) Reduce after load.
- 8) Control heart rate, rhythm, and synchrony.
- 9) Optimize heart-lung interactions.
- 10) Provide mechanical support when needed.

Blalock-Taussig Bidirectional Glenn shunt shunt Modified Blalock-Taussig shunt Waterston-Cooley Potts shunt shunt

CONGENITAL HEART DISEASE

Review CHD headlines

- General assessment and preparations
- Recognizing innocent murmurs
- Evaluation of pathological murmurs
- Full orientation about anatomical defects and its implication on cardiac performance with subsequent complications

Heart Murmur – Innocent

1- Innocent murmurs through childhood(75% to 90%) most of these children will not require a cardiology consultation. A).Still's murmur. **B)**.pulmonary murmur. **C).** supraclavicular bruit..

Heart Murmur – pathological Congenital heart disease

- routine evaluation : CHD (whether repaired or not) should have a cardiac evaluation within 1 year before surgery, even if asymptomatic.
- Current use of cardiac medication.
- co-morbidities:e.g. prolonged QT syndrome and residual cardiac disease
- may require **SBE** prophylaxis

Some implications of CHD

- R L shunt: risk of hypoxemia , polycythemia and embolic phenomenon
- Long-term L R shunt : pul. HTn, shunt reversal (E.S = in-operable)
- Ebstein anomaly : risk of arrhythmia
- Pulmonary hypertension : risk of some anesthetic drugs
- Duct-dependent CHD : sudden collapse, emergency use of PGE2 infusion
- Etc

SPECIAL CONDITIONS

Prolonged QT Syndrome



- increased risk for V.tac (Torsades de pointes) either <u>congenital</u> genetic mutations or the effects of <u>drugs</u> or metabolic abnormalities on the ion channels responsible for repolarization.
- 2. so be vigilant and prepare suitable antiarrhythmic drugs for intra-operative and post=operative period .
- 3. ensure that serum levels of potassium, calcium, and magnesium are normal.
- 4. avoid drugs that prolog QT interval

Drugs that prolong QT interval

Dopamine, Droperidol, Ephedrine, Epinephrine Erythromycin, Felbamate, Flecainide, Foscarnet Fosphenytoin, Gemifloxacin, Granisetron, Halofantrine Haloperidol, Isoproterenol, Isradipine, Levalbuterol Levofloxacin, Lithium, Metaproterenol, Methadone Methylphenidate, Milodrine, Moxifloxacin, Nicardipine Norepinephrine, Octreotide, Ofloxacin, Ondansetron Pentamidine, Phenylephrine, Pimozide, Procainamide Pseudoephedrine, Quetiapine, Quinidine, Risperidone Salmeterol, Sotalol, Tacrolimus, Telithromycin Terbutaline, Thioridazine, Tizanidine, Venlafaxine Ziprasidone



Williams Syndrome

Williams syndrome is a constellation of dysmorphic facies, intellectual disability, growth deficiency, neonatal hypercalcemia, genitourinary anomalies, and congenital heart disease.

- The principal cardiac manifestation is supravalvular aortic stenosis, which is often progressive Pulmonary arterial obstruction is seen in 80% of cases.
- Other end-organ manifestations include epilepsy, persistent hypercalcemia, hypothyroidism, renovascular hypertension, and proteinuria.
- Sudden death is estimated to occur in at least 3% of patients with Williams syndrome; many deaths occur during the perioperative period (myocardial ischemia, decreased cardiac output, and ventricular dysrhythmias (Holter monitor).

Williams Syndrome - preparations

- Full evaluation for heart function (including outflow tract) and comorbidities in other systems .
- High risk consent.
- Anesthetic induction must be performed with great care, **avoiding** extremes in heart rate and blood pressure.



DOWN SYNDROME

40-50% associated CHD, often requiring surgical correction. As well as the need for **SBE** prophylaxis. increased risk of OSAS and Pul. Hypertension and Pulmonary vascular disease results from these lesions, \rightarrow **Difficult ETI** due to : mid-face hypoplasia, smaller trachea ETT must 1-2smaller size than peers, OSAS, and risk of cervical **spine subluxation** (neck-protection strategies)



Figure 62–9. Relationship of the arch of C1 and odontoid process of C2. (Hata T, Todd MM. Cervical spine considerations when anesthetizing patients with Down syndrome. Anesthesiology. 2005;102:680–685. Reprinted by permission of Lippincott Williams & Wilkins.)



Figure 62–10. Increased distance (subluxation) between C1 and odontoid in Down syndrome. (Hata T, Todd MM. Cervical spine considerations when anesthetizing patients with Down syndrome. Anesthesiology. 2005;102:680–685. Reprinted by permission of Lippincott Williams & Wilkins.)

atlantodens interval of greater than 5 mm (usually maximal in the flexion view), then the child should be referred for orthopedic or neurosurgical consultation before elective surgery



SUBACUTE BACTERIAL ENDOCARDITIS PROPHYLAXIS

Antibiotic prophylaxis to prevent bacterial endocarditis has long been recommended For children who have congenital heart disease and are undergoing any procedure in which the patient is at risk for transient bacteremia.

SUBACUTE BACTERIAL ENDOCARDITIS PROPHYLAXIS

Oral endotracheal intubation by itself is not an indication for SBE prophylaxis, but nasotracheal intubation requires it. **Patients with hemodynamically** insignificant lesions such as bicuspid aortic valve, mitral valve prolapse no longer require prophylaxis for any procedure.

SUBACUTE BACTERIAL ENDOCARDITIS PROPHYLAXIS

Patients with congenital heart disease repaired with prosthetic material require prophylaxis only for the first 6 months after repair, after which endothelialization has occurred. Such is the case for VSD as well as ASD repairs as long as no residual defect is present.
Patients with prosthetic valves or those palliated with shunts or conduits require prophylaxis.

BOX 7-4

CARDIAC CONDITIONS FOR WHICH ANTIBIOTIC PROPHYLAXIS IS RECOMMENDED FOR DENTAL, RESPIRATORY TRACT, INFECTED SKIN, SKIN STRUCTURES, OR MUSCULOSKELETAL TISSUE PROCEDURES

- Prosthetic cardiac valve
- Previous bacterial endocarditis
- Congenital heart disease (CHD)—Limited to the following conditions*
 - Unrepaired cyanotic defect, including palliative shunts and conduits
 - Completely repaired CHD with prosthetic material/device (placed by surgery or catheterization), during first 6 months after procedure[†]
 - Repaired CHD with residual defects at or adjacent to the site of prosthetic patch or device (which inhibit endothelialization)
 - Cardiac transplantation patients who develop cardiac valvulopathy

*Conditions associated with the highest risk of adverse outcome from endocarditis. [†]Endothelialization process of prosthetic material occurs within 6 months after the procedure.

Are rheumatic heart diseases still an indication for SBE prophylaxis ??
 Are Pts on LABP for RHD , still need EBE prophylaxis (if involved in previous indications)

Key Points in Preoperative Care

- Pediatric cardiac intensive care has emerged as an important and necessary subspecialty.
- The complexity of heart disease and the expertise necessary to treat these
 patients require multidisciplinary training and collaborative, integrated care.
- Expertise is required for care of premature and full-term newborns, infants, and children, as well as rapidly growing numbers of adults with long-term survival and continuing need for care of their CHD.
- Reparative surgery in the newborn is the objective of advanced cardiovascular programs whenever feasible.
- Diagnosis is usually based in echocardiography and physical examination, with catheterization reserved usually for complex cases or interventional procedures.
- Adequate resuscitation of preoperative patients is essential to good outcomes.
- Balancing single-ventricle physiology and maintaining adequate cardiac output is important to achieving preoperative stabilization.
- Risk stratification and identification of biologic markers are maturing as useful tool sets to guide therapy and benchmarks for outcomes.
- Therapies for the future will target genetic factors and tailor treatments to the polymorphisms and individual inherited profiles of patients.
- Fetal diagnosis will increasingly continue to provide advanced knowledge, counseling, and therapeutic planning and eventually eliminate unanticipated postnatal circulatory collapse.

POST-OPERATIVE CARE

Assessment and supports

- Ass essment:verify the accuracy of the preoperative diagnosis depending on intra-op findings and the adequacy of the surgical repair. And bulk of supportive therapy needed in post-op state.
- **Standard post-op care:** stabilization, supportive & symptomatic
- Cardiac monitoring & strategies to improve C.O.P.
- Search for special anticipated complications ,(e.g. arrhythmias, hypoperfusion....) and deal according previously constructed plan .
- Effects of cardiopulmonary bypass (CPB) in cardiac surgery which may include ; hypothermia, alkalosis, generalized SIR, platelets dysfunction and deficiency ,

Key Points post cardiac operation

- Know the anatomy and surgical procedure in detail.
- Search systematically for residual disease in the postoperative patient.
- Anticipate low cardiac output.
- Preserve right-to-left shunts for transient posto perative benefit in select patients.
- Optimize afterload reduction and avoid high doses of catecholamines.
- Appreciate heart-lung interactions and effects of positive-pressure breathing.
- Understand limitations and opportunities to manipulate single-ventricle physiology.
- Estimate the limit of cardiac reserve at the nadir of postoperative cardiac output.
- Embrace mechanical support of the circulation as a vital tool for bridge to recovery, bridge to transplantation as well as destination therapy.
- The management of the postoperative pediatric cardiac surgical patient requires a comprehensive understanding of the basic principles of oxygen delivery, cardiovascular physiology, and the anatomy and physiology of congenital heart disease.
- Signs and symptoms of low cardiac output syndrome should be treated aggressively, and diagnostic and therapeutic strategies should address both universal and lesion-specific problems.

Postoperative complications **1- VENOUS THROMBOEMBOLISM** Although not as common as in adults, venous thrombo-embolism (VTE) occurs in children in the postoperative period, with the incidence increasing in adolescence.

Postoperative complications

- **1- VENOUS THROMBOEMBOLISM**
- The primary care physician should be alert to symptoms of VTE, such as extremity pain, swelling, and discoloration, which may indicate deep-vein thrombosis and should be referred for immediate evaluation.
- Patients at highest risk are those who are immobilized after surgery and have at least 1 other risk factor. Risk factors for VTE are listed.

Box 63-2 Risk Factors for Venous Thromboembolism (VTE) in the Postoperative Period Immobility Major lower extremity orthopedic surgery Spinal cord injury Major trauma or trauma to the lower extremities Previous history of deep-vein thrombosis or VTE or pulmonary embolism Pregnancy Oral contraceptive use Inflammatory bowel disease Nephrotic syndrome Burns Obesity Central venous catheter in the lower extremity Known acquired or inherited thrombophilia Acute infection

Postoperative complications **1- VENOUS THROMBOEMBOLISM** Patients who develop VTE are at risk for pulmonary embolism, which has a mortality rate as high as 20%. **Symptoms** of pulmonary embolism include dyspnea, chest pain, cough, hemoptysis, and fever. Patients at risk for VTE should receive prophylactic measures, which may include compression stockings or pneumatic sequential compression devices (or both) until ambulatory.

Postoperative complications **1- VENOUS THROMBOEMBOLISM** • Patients with 3 or more risk factors may be treated with pharmacologic prophylaxis: subcutaneous heparin or low-molecular-weight heparin.



