

Universitair Medisch Centrum Groningen Afdeling Anesthesiologie

Anesthesia for Cardiovascular Surgery





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Basic monitoring:

•ECG

- \geq 2 leads (II and V₅)
- Baseline print of all leads
- ST-segment analysis
- •Pulse oxymetry
- •Backup manual or automatic blood pressure cuff



Venous Access:

- Two large-bore (16-gauge or larger) peripheral IV catheters
- •One central venous line, usually IJV
- •Measurement of central venous pressure (?)
- •PA-catheters: on indication
 - Low EF, pulm. Hypertension, complex procedures

Arterial cannulation

- Insert before induction of anesthesia
- Non-dominant hand (caveat: radial art. harvesting)
- Direct and continuous measurement of arterial blood pressure



Other:

- Indwelling urinary catheter
 - Urine output, bladder temperature
- •Temperature probes
 - Esophageal, nasopharyngeal, skin, bladder, tympanic, blood
- Cross-matched blood available
 - Especially if patient has already had a midline sternotomy
- •Consider thoracic epidural anesthesia (only Europe)
 - Problem: epidural hematoma formation following heparinization



Induction of Anesthesia:

- •Goal: hemodynamic stability
- •Selection of induction agents:
 - High-dose opiates
 - ± Benzodiazepine
 - Modest dose propofol
 - Muscle relaxant, endotracheal intubation
 - Vasopressor if blood pressure falls >20%

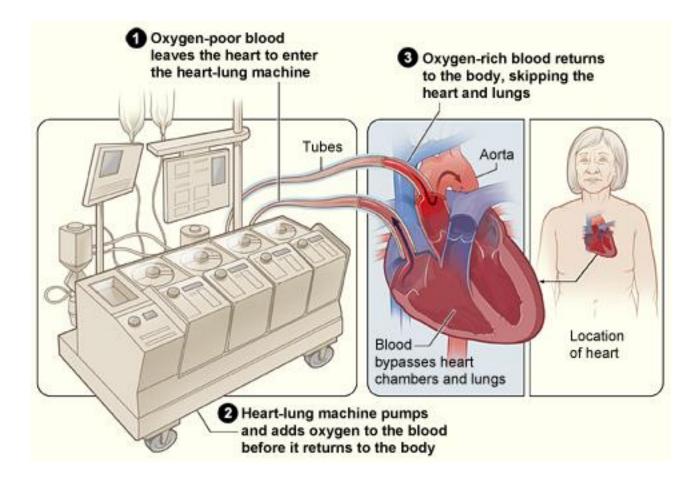


Maintenance of anesthesia:

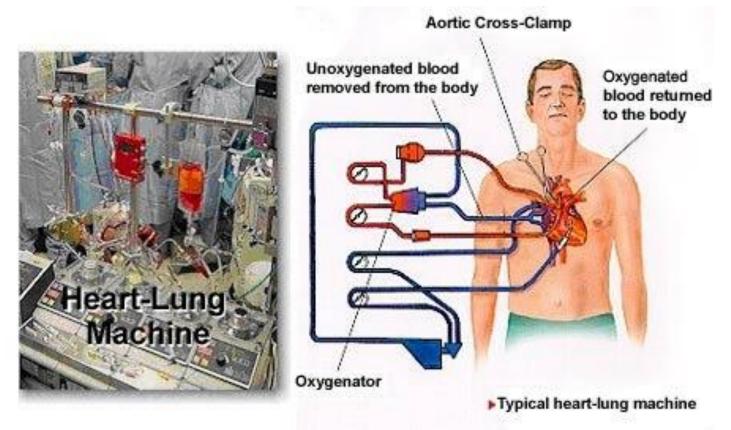
•Selection of anesthetic technique/agents:

- TIVA with short acting agents, covers whole procedure
- Volatile anesthetic agents
 - Cardioprotection
 - Difficult to use during CPB
- Avoid N₂O! (expansion of intravascular air bubbles, pneumothorax)
- High-dose opiates
- Muscle relaxants
- Vasopressor if blood pressure falls >20%
- Goal: early extubation (1–6 h postop.), fast track?





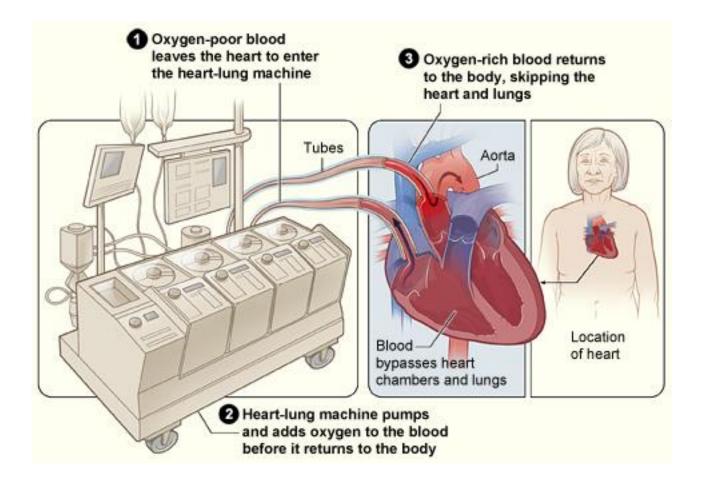




The CPB machine has five basic components:

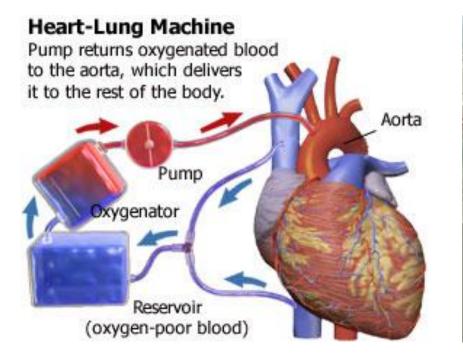
- 1. venous reservoir,
- 2. oxygenator,
- 3. heat exchanger,
- 4. main pump (roller pumps of centrifugal pumps), and
- 5. arterial filter (air, thrombi, fat globules, calcium, tissue debris)

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CPB diverts venous blood away from the heart, adds oxygen, removes CO_2 , and returns the blood to a large artery. As a result, most blood flow through the heart and most of the flow through the lungs cease.







Flow = non-pulsatile Organ protection: Perfusionist

- Hypothermia
- Cardioplegia
 - (cold, K⁺-rich, blood or crystalloids, ante- and retrograde, repeat every 30 min)



PHYSIOLOGICAL EFFECTS OF CARDIOPULMONARY BYPASS

Initiation of CPB is associated with:

- •a marked increase in stress hormones
- •a variable systemic inflammatory response (sepsis-like)
 - Generation of oxygen-derived free radicals
- •Activation of multiple humoral systems, including complement, coagulation,

fibrinolysis, and the kallikrein system

- •Mechanical trauma alters platelets and activates leukocytes
 - Depletion of glycoprotein receptors on the surface of platelets
 - Increased perioperative bleeding



PHYSIOLOGICAL EFFECTS OF CARDIOPULMONARY BYPASS

Potential methods to prevent complications (experimental):

- •Glucocorticoids?
- •Cyclokapron acid?
- •Leukocyte depletion?
- Intraoperative hemofiltration?
- •Antioxidants (Vit. C, E, Mannitol)?
- cyclooxygenase-2 inhibitors?
- •Pentoxifylline?



Going on-pump

- Confirm adequate anticoagulation (ACT) before cannulation
- Venous cannulation
- Aortic cannulation
 - Reduce systemic arterial pressure (to 90–100 mm Hg systolic)
 - Complications: Aortic dissection, Cerebral embolism (plaque, air)
- Start CPB, pump flow gradually increased to 2–2.5 L/min/m²
- Cold blood cardioplegia
- Continue ventilation until heart stops ejecting blood
- Maintain MAP 50 80 mmHg (organ perfusion)
- Hemodilution due to CPB priming
- umcg Keep hematocrit between 20% and 25%

Intraoperative laboratory monitoring:

- •Blood gas analysis (point of care):
 - Hematocrit
 - Serum potassium
 - Ionized calcium
 - Glucose
- •Activated clotting time (ACT)
 - Activators: celite, kaolin, glass
 - Reference value ranges between 70-180 sec
 - During CPB the desired range is >400-500 sec
 - During OPCAB procedures usually >300-400 sec
- •Thrombelastography (TEG)?



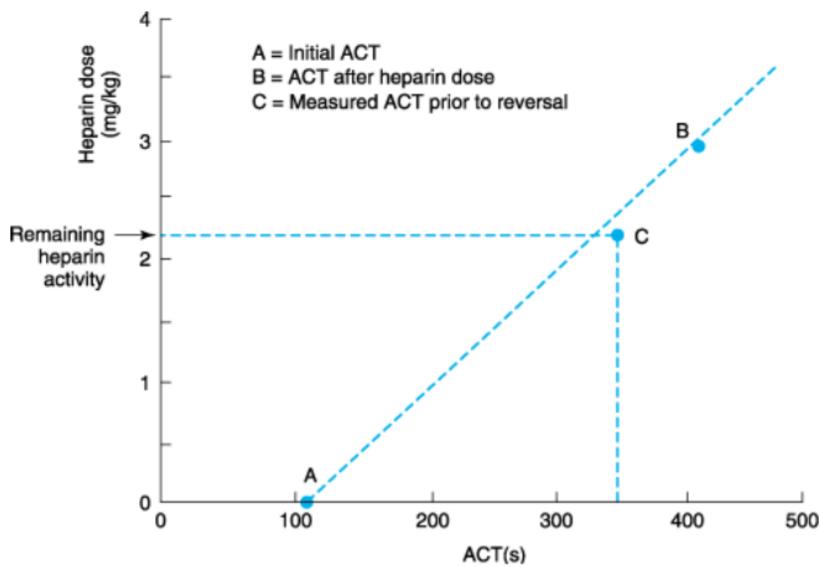
Anticoagulation:

- •must be established prior to CPB
- •must be confirmed with determination of the ACT
- •Heparin: 3 mg/kg (OPCAB: 2 mg/kg)
 - Problem: resistance to heparin (AT III deficiency)
 - Solution: infuse FFP (2 units) or AT III
 - Problem: heparin-induced thrombocytopenia (HIT)
 - Solution: consider alternative anticoagulants (hirudin, bivalirudin, argatroban)
 - Problem: previous administration of glycoprotein IIb/IIIa inhibitors (abciximab [RheoPro] or tirofiban [Aggrastat]) or ADP receptor antagonist clopidogrel (Plavix)
 - Solution: aminocaproic acid (5–10 g followed by 1 g/h) or tranexamic acid (10 mg/kg followed by 1 mg/kg/h)



Anticoagulation:

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Management of Respiratory Gases

- Problem: the solubility of a gas increases with hypothermia
 - As a result, the partial pressure of the gas will decrease
 - most significant for PaCO₂ (effect on cerebral blood flow)
- Example: Blood with PaCO₂ of 5 kPa and a pH of 7.40 at 37°C, when cooled to 25°C, will have a PaCO₂ of about 2.6 kPa and a pH of 7.60



Management of Respiratory Gases

- Problem: blood samples are heated to 37°C in blood gas analyzers
- Solution 1: pH-stat management
 - Temperature correcting gas tensions and maintaining a "normal" PaCO₂ of 40 mm Hg and a pH of 7.40 during hypothermia
 - May require adding CO₂ to the oxygenator
 - Increases total blood CO₂-content
 - Impairs cerebral blood flow autoregulation
- Solution 2: α-stat management (more common)
 - Use of uncorrected gas tensions during hypothermia
 - Preserves cerebral autoregulation of blood flow



Troubleshooting during CPB

- Problem: after start of CPB, venous reservoir empties (air enters pump circuit)
- Solution: check venous return:
 - forgotten clamps?
 - Cannula malposition?
 - Kinking?



Troubleshooting during CPB

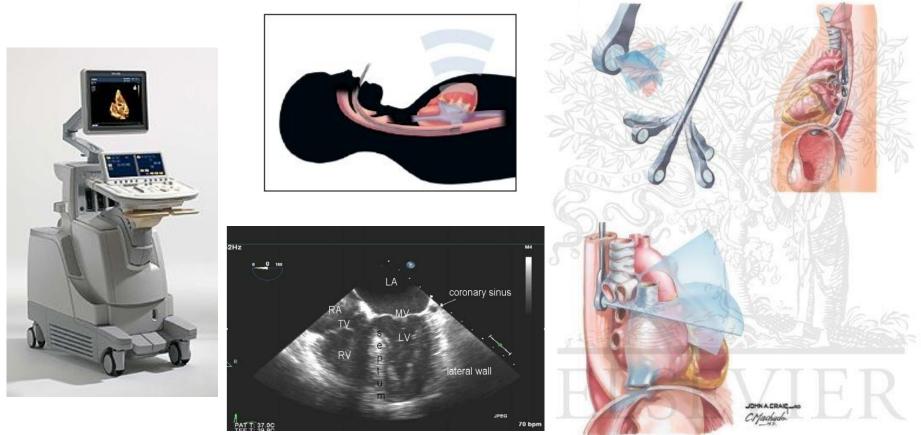
- Heart does not empty
 - Solution: aortic regurgitation? Malpositioning of the venous cannula?
- MAP decreases <30 mmHg
 - Solution: search for unrecognized aortic dissection, recannulate aorta (distal)



Monitoring: Transesophageal Echocardiography (TEE)

TEE provides extremely valuable information about

- Cardiac anatomy and
- Cardiac function during surgery (valvular function, pump function, ischemia, hypovolemia)
- Surgical result, need for re-intervention





Monitoring: Transesophageal Echocardiography (TEE)

Future: 3D echo

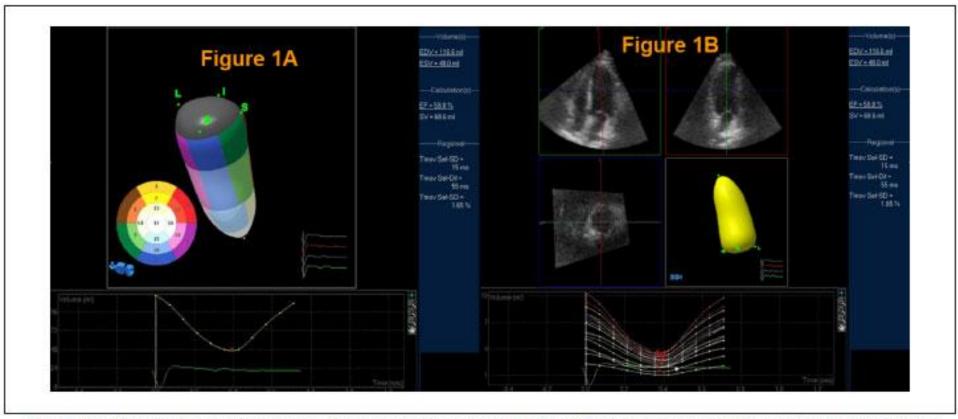


Figure 1 - Three-dimensional echocardiographic image of the left ventricle (17-segment model, Figure 1A), with measurements of ventricular ejection fraction and volumes. Volume and ejection fraction measurements were derived from the two-dimensional echocardiography (Figure 1B) as analyzed in multiple observation planes.



Weaning from CPB (preparation):

- •A "hot shot" or warm blood cardioplegia can be administered
 - To wash out byproducts
 - Replenish metabolic substrates
- Optimise physiological conditions
 - Acidosis and hypoxia should be corrected
 - Lung ventilation must be resumed
 - Normothermia (≥36°C) should be achieved
 - Normovolemia should be achieved
 - Hb should be kept $\geq 5 \text{ mmol/L}$



Weaning from CPB (preparation):

- •Surgeon unclamps the aorta
 - Washes out cardioplegia (heart re-starts)
- •Re-start pulmonary ventilation
- •Continue CPB additional 5 10 mins after heart re-start
 - Keep heart in empty and beating state
 - Stabilizes heart, minimal metabolic requirements
- •A stable rhythm (preferably sinus) must be present
 - Atrioventricular pacing may be necessary (80–100 bpm)
- •TEE monitoring (chamber vols, contractility, valvular function)



Weaning problems:

•Problem: residual air (TEE, ECG)?

•Solution:

- Evacuate air from the heart and any bypass grafts
- increase perfusion pressure (norepinephrine)



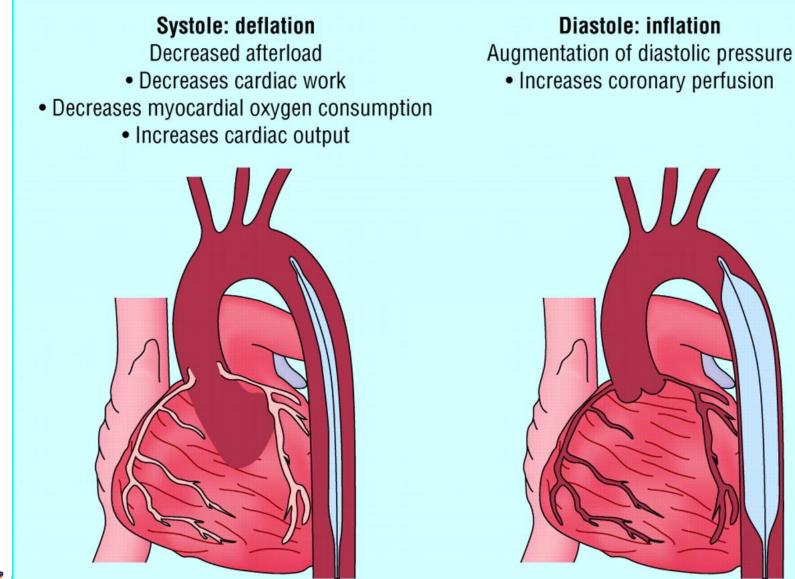
Weaning problems:

Poor cardiac function

- Consider inotropic support:
 - dobutamine (1st choice)
 - milrinone (esp. right ventricular failure)
 - Norepinephrine
 - (epinephrine, dopamine)
 - Levosimendan (not approved in NL)
- Consider reperfusion
- Consider afterload reduction (nitroprusside, milrinone)
- Consider intraaortic balloon pump (IABP)
- Consider left or right ventricular assist device

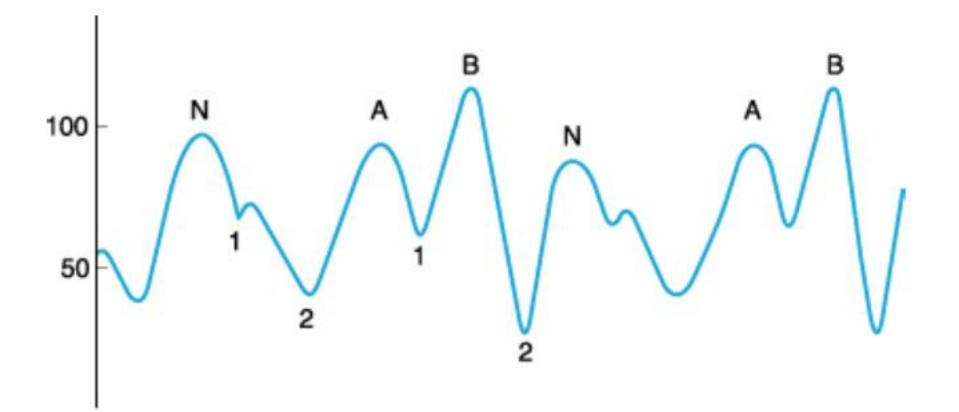


Intraaortic balloon pump





Intraaortic balloon pump





Reversal of anticoagulation:

- •Protamine binds and effectively inactivates heparin
- •Dose: 1 mg of protamine per mg of (initial) heparin
- Infuse slowly
 - Hemodynamic side-effects
 - Hypotension (vasodilation)
 - myocardial depression
 - pulmonary hypertension
 - Allergic reactions
- Check effect with ACT

•Consider supplemental protamine (50–100 mg) after administration of CPB blood



PATHOLOGICAL EFFECTS OF CARDIOPULMONARY BYPASS

• Underlying problems

- Age
- Comorbidities
- Procedural complexity
- Equipment issues

Postoperative problems

- Stroke, incidence 1-3%, higher in aortic surgery
- Delirium, incidence 10-60%
- POCD, incidence 24-53%
- Longer hospital LOS
- Higher costs



PATHOLOGICAL EFFECTS OF CARDIOPULMONARY BYPASS

Intra- and Postoperative Predictors of Stroke After Coronary Artery Bypass Grafting

Donald S. Likosky, PhD, Bruce J. Leavitt, MD, Charles A. S. Marrin, MB, BS, David J. Malenka, MD, Alexander G. Reeves, MD, Ronald M. Weintraub, MD, Louis R. Caplan, MD, Yvon R. Baribeau, MD, David C. Charlesworth, MD, Cathy S. Ross, MS, John H. Braxton, MD, Felix Hernandez, Jr, MD, and Gerald T. O'Connor, DSc, PhD, for the Northern New England Cardiovascular Disease Study Group (Ann Thorac Surg 2003;76:428–35)

- 11,825 CABG Patients, 1.5% Incidence of Stroke
- 75% of strokes occurred among low or medium preoperative risk patients
 - Many of these strokes may be preventable
 - Traditional pre-op risk assessment is unreliable
 - Additional intraoperative monitoring (e.g. NIRS) may be of value



Cerebral Oxymetry Changes in INVOS values are influenced by the critical balance between arterial oxygen delivery and cerebral consumption.

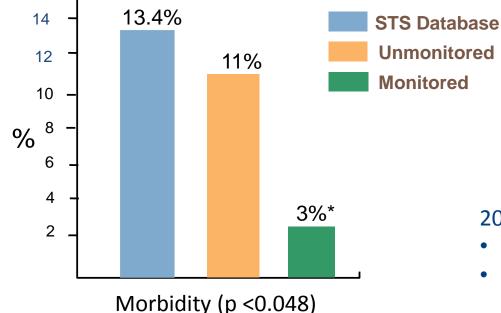
Imbalances are identified by changes in rSO₂.



Cerebral Oxymetry: Results

Monitoring Brain Oxygen Saturation During Coronary Bypass Surgery: A Randomized, Prospective Study

Murkin JM, et al. Anesth Analg. 2007 Jan;104(1):51-8.



200 CABG patients;

- 100 blinded rSO₂ monitoring
- 100 intervention protocol



PATHOLOGICAL EFFECTS OF CARDIOPULMONARY BYPASS

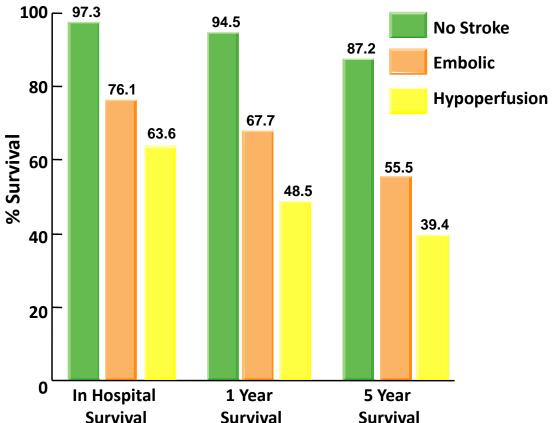
Perioperative Stroke and Long-Term Survival After Coronary Bypass Graft Surgery (Ann Thorac Surg 2005;79:532–7)

Lawrence J. Dacey, MD, Donald S. Likosky, PhD, Bruce J. Leavitt, MD, Stephen J. Lahey, MD, Reed D. Quinn, MD, Felix Hernandez, Jr, MD, Hebe B. Quinton, MS, Joseph P. Desimone, MD, Cathy S. Ross, MS and Gerald T. O'Connor, DSc, PhD, for the Northern New England Cardiovascular Disease Study Group

35,733 CABG patients

Stroke incidence 1.61%

"Patients who had perioperative stroke were at a significantly increased risk for death...Survival at each time point was lowest among patients who had hypoperfusion strokes."



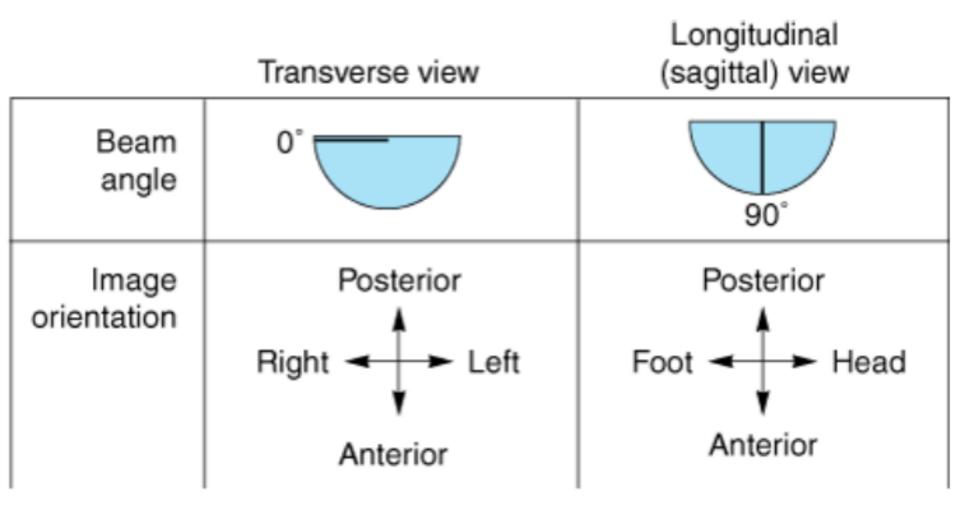




Any questions?

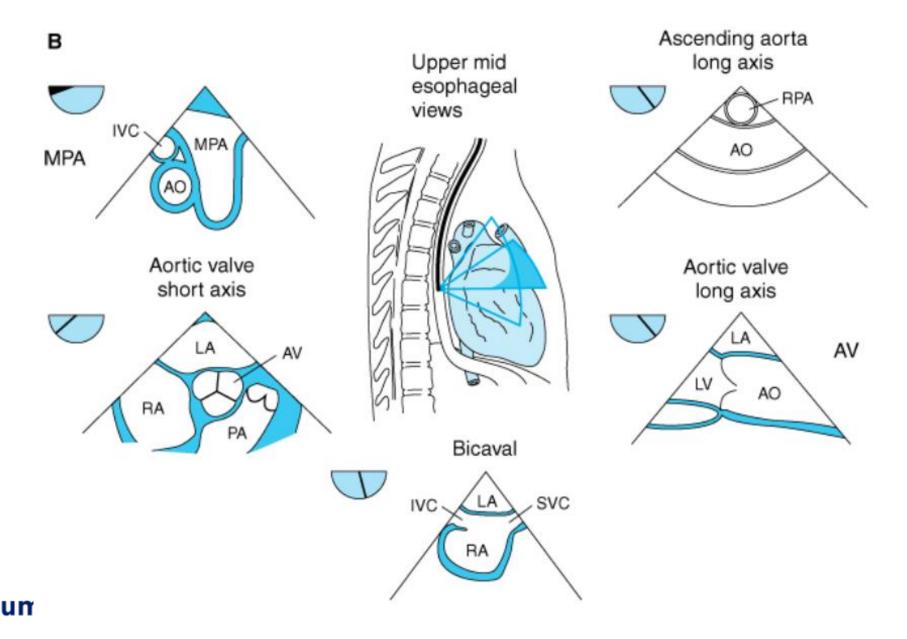


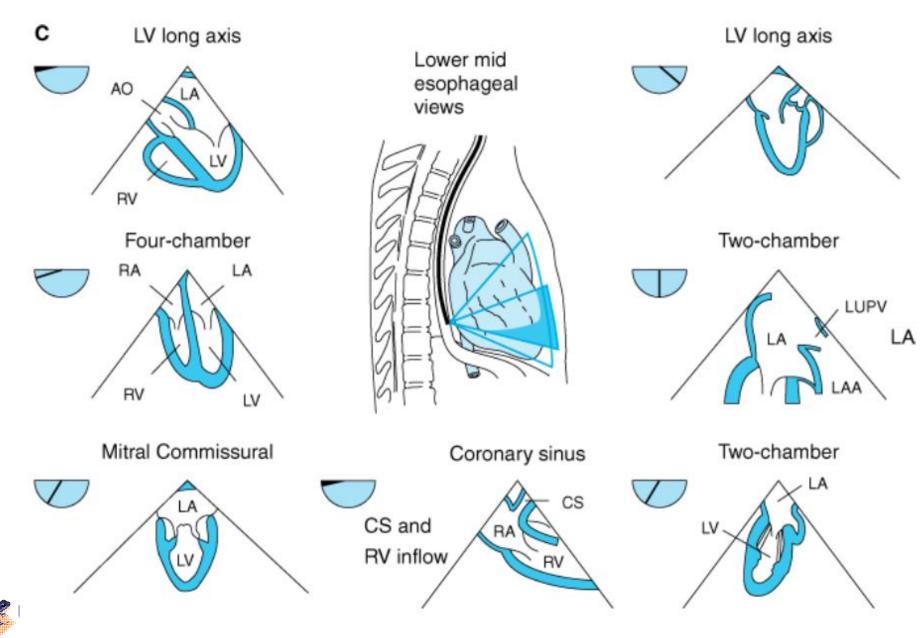
Monitoring: Transesophageal Echocardiography (TEE)

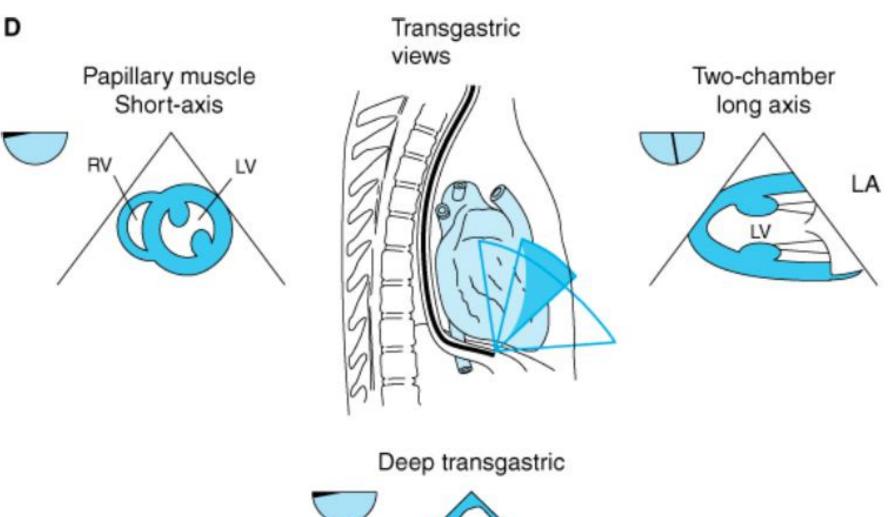


Twodimensional, multiplane TEE: nomenclature



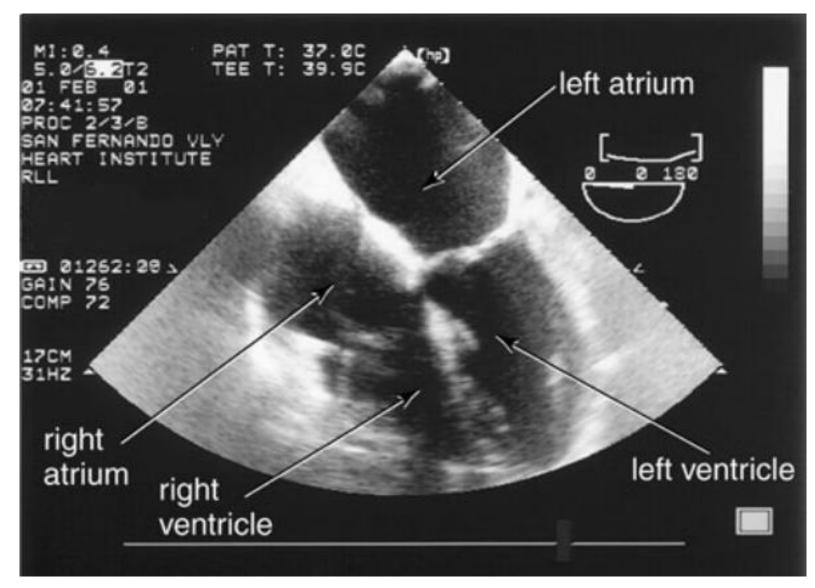




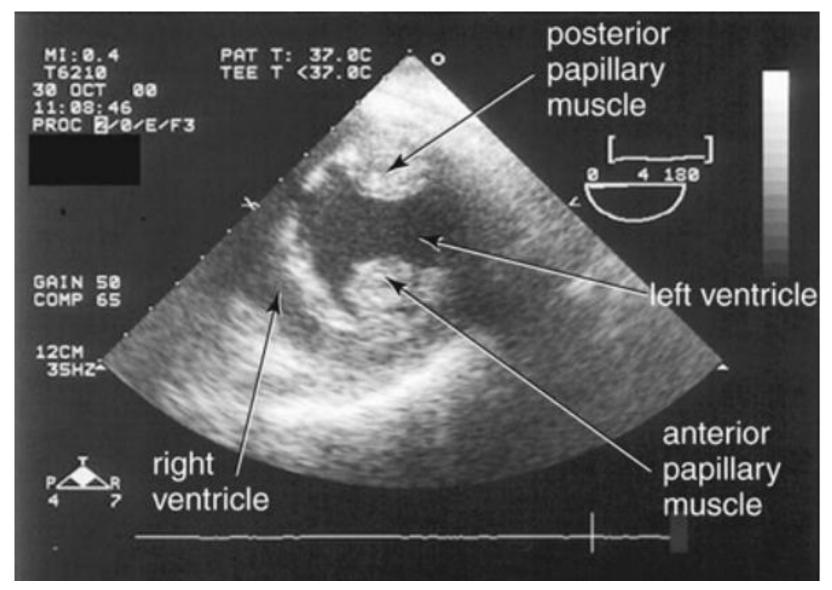


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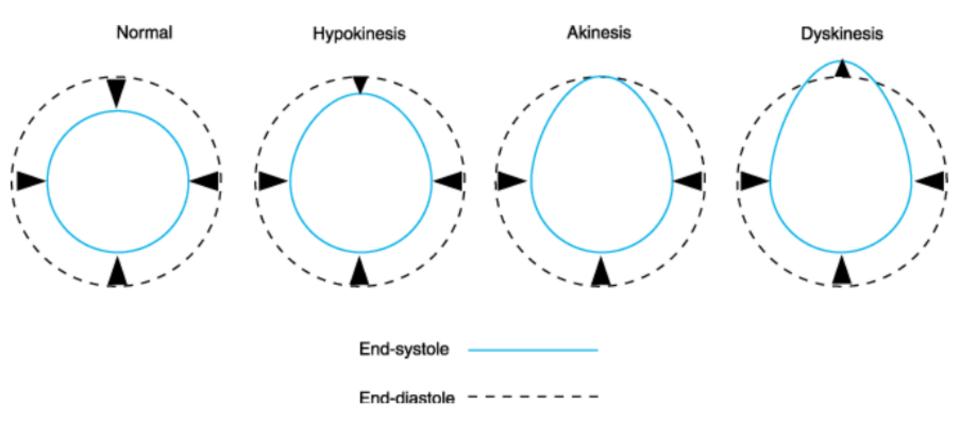






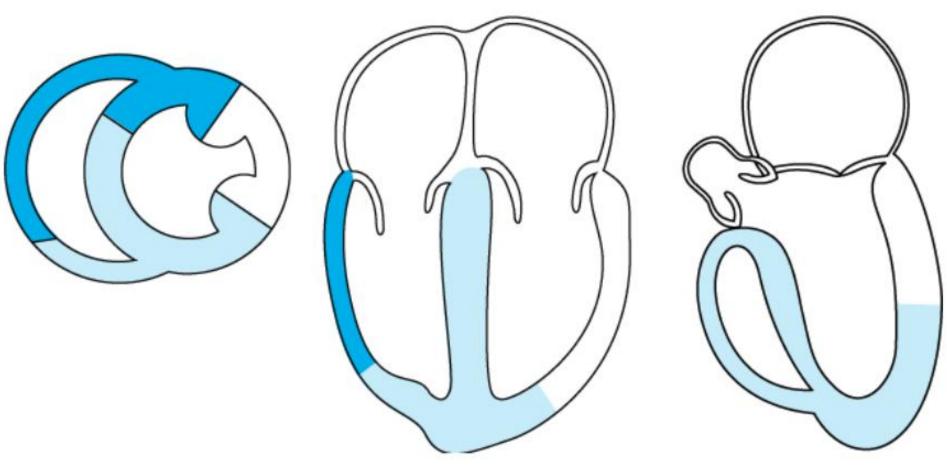


Assessing Ventricular Function





Coronary artery supply of the left and right ventricles

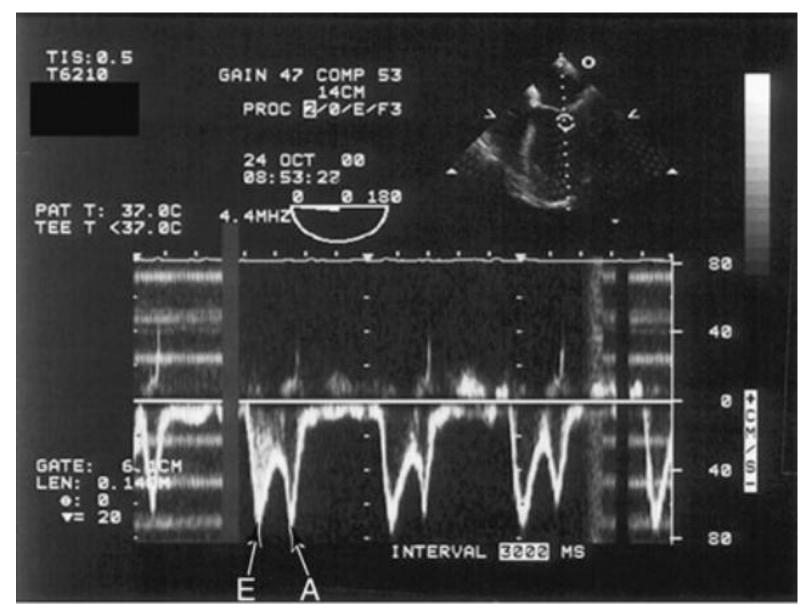


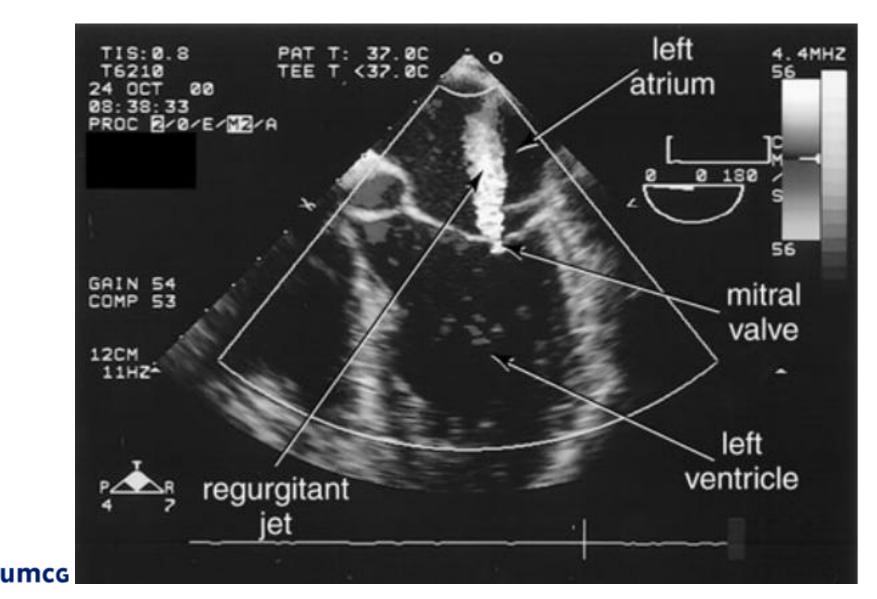
Dark blue, RCA; light blue, LAD; white, CX.

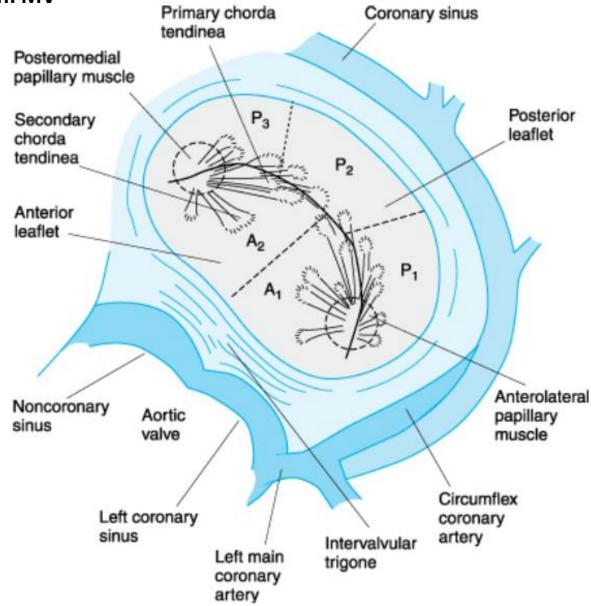


Assessing Valvular Function: MV

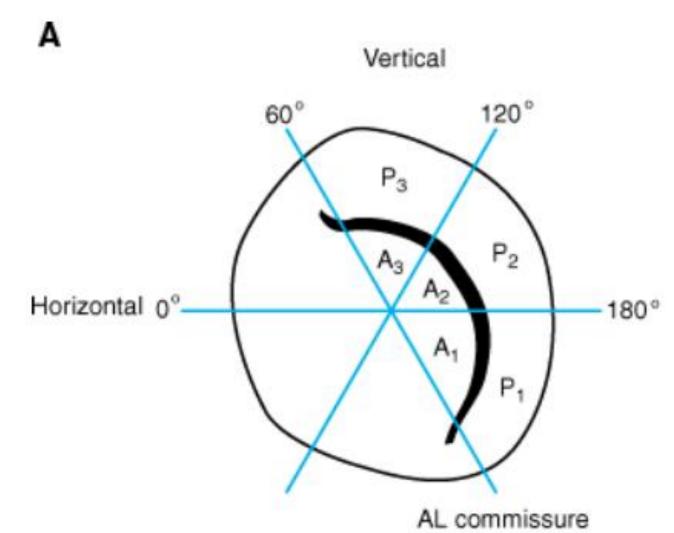
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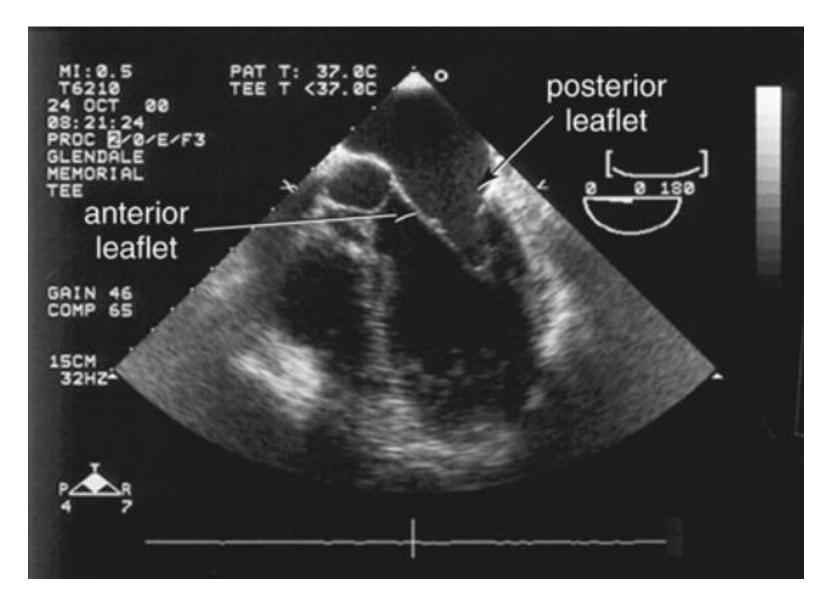




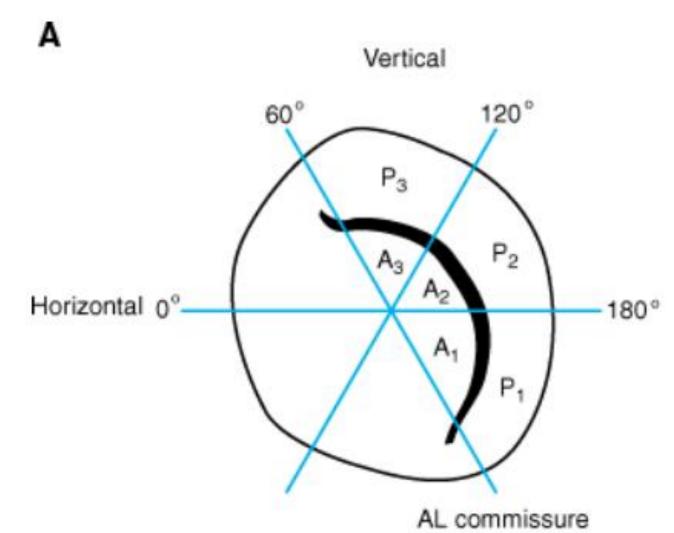




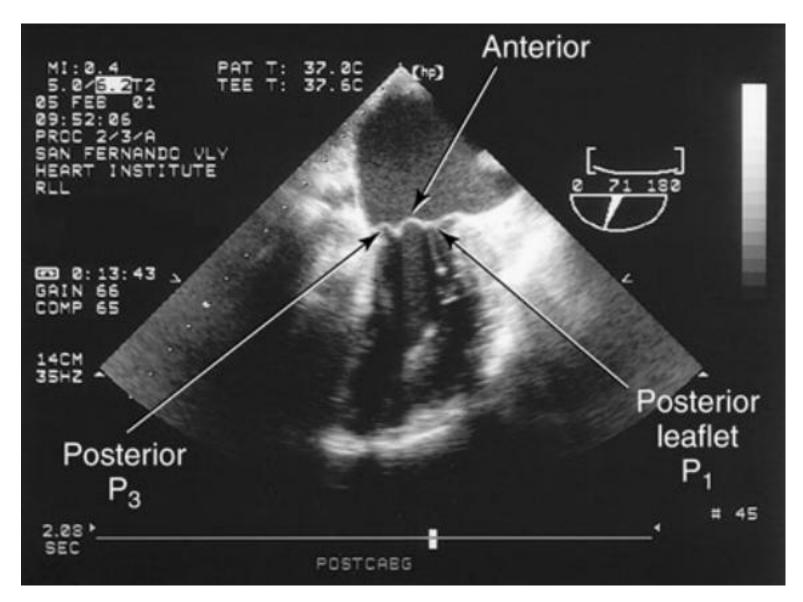




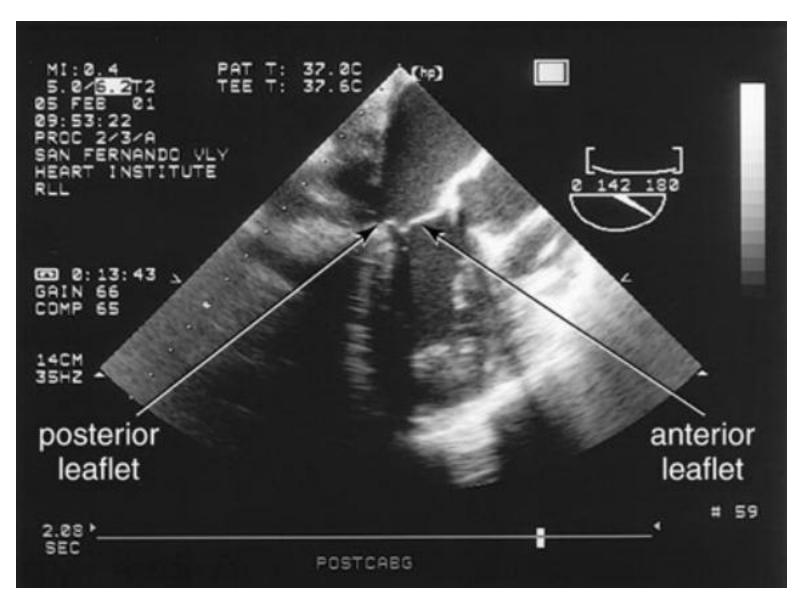




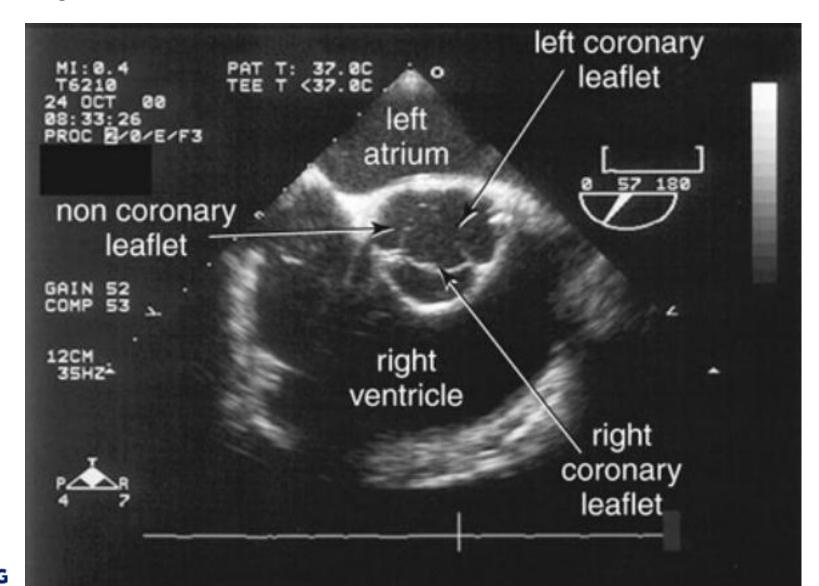




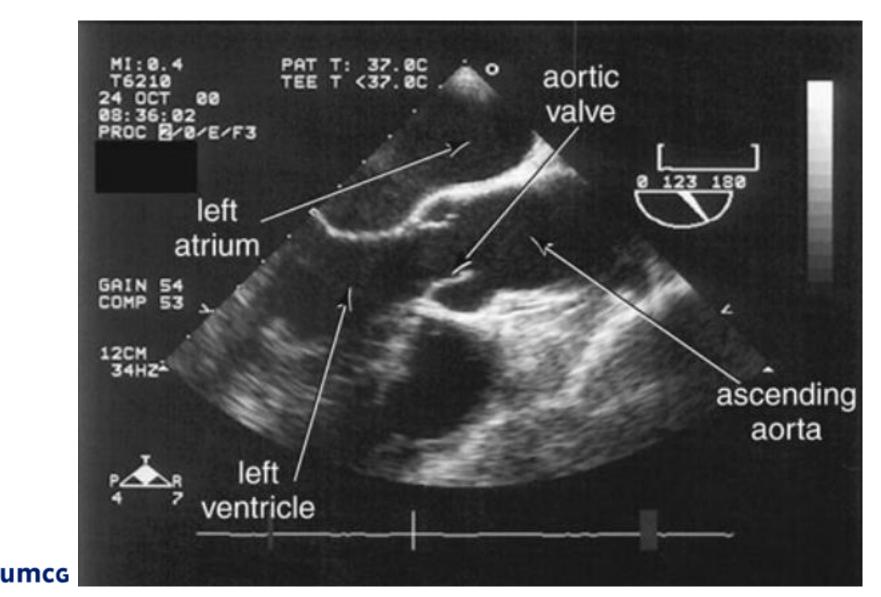


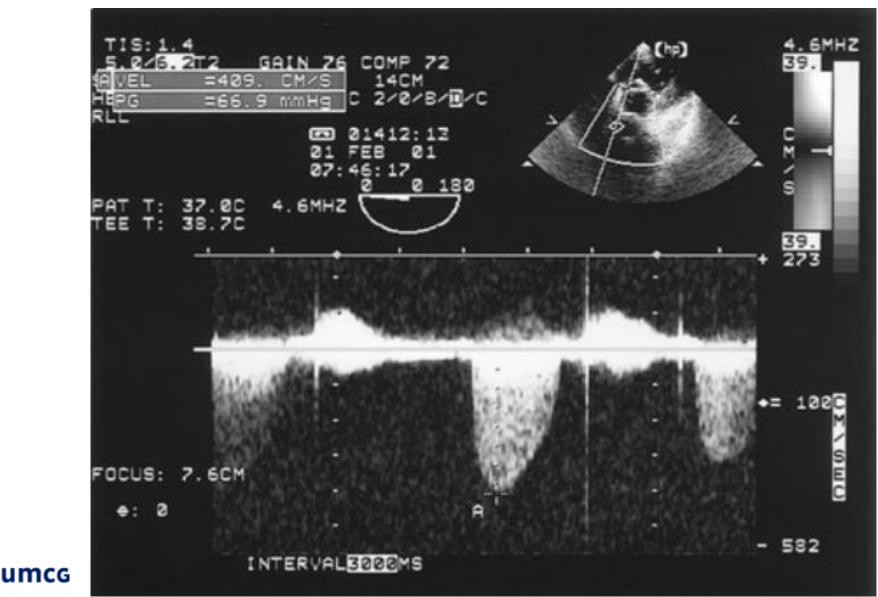




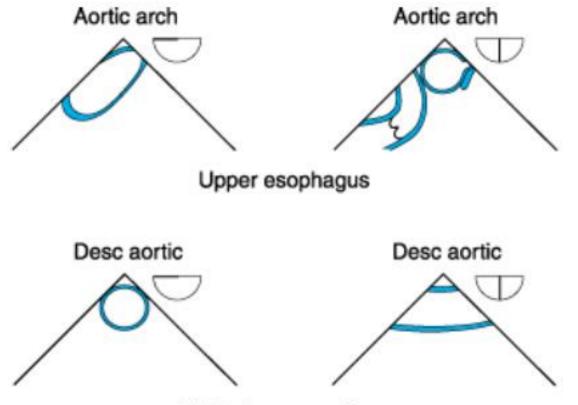








Views of the aortic arch and descending aorta



Mid to lower esophagus

