

Perioperative Implications

Preoperative Preparation

- PFTs for BOS staging and resp status and bronchoscopy for biopsy and culture.
- Treat active infections aggressively.
- Evaluate renal functions and adjust periop medications where appropriate.
- Continue anti-infective and immunosuppressive therapy during the periop period and adjust dosing to keep within the indicated therapeutic range.
- Strict aseptic techniques due to immunosuppression.
- Premedication useful due to excessive secretions, but avoid excessive resp depression.
- Reflux prophylaxis.
- Corticosteroids supplementation especially for long, invasive, and stressful procedures.
- Watch for increased sensitivity to opioids, hypercarbia, resp acidosis, bronchial hyperresponsiveness (bronchoconstriction), V/Q mismatch, GE reflux, hyperkalemia, and hypomagnesemia.
- Most common side effect of immunosuppressive drugs: Cyclosporine and tacrolimus (Htn, diabetes, neurotoxicity, and renal failure), glucocorticoids (hyperglycemia, weight gain, osteoporosis, and adrenal insufficiency), and azathioprine (anemia and thrombocytopenia).

Monitoring

- Routine.
- Consider arterial line placement if hypoxic, acidotic, or O₂ saturation is inadequate: invasive monitoring must be carefully weighed against the possibility of infection from intravascular catheters.
- TEE or other continuous CO monitoring systems may be helpful in assessing cardiac function in post

heart-lung transplant pts, and when there is evidence of pulm edema and pulm Htn.

- CVP insertion recommended (when necessary) on side of native lung (one-lung transplant).

Airway

- ETT cuff placement should avoid tracheal anastomosis.
- Oral intubation is preferred over nasal intubation (due to infection and thrombocytopenia).
- Anticipate difficult intubation if on chronic corticosteroids due to Cushingoid (moon face) features and limited atlanto-occipital joint mobility.
- Use aseptic tracheal suction technique.

Induction

- Short-acting agents preferred; adjust doses to pt status and to avoid prolonged CV depression.

Maintenance

- Avoid fluid overload; renal dysfunction due to immunosuppressants and disruption of lymphatic drainage in posttransplant pts can lead to pulm edema with fluid overload.
- Significant reductions of cyclosporine or tacrolimus blood levels can be caused by dilution with IV fluids.
- Adjust neuromuscular blocking dosage due to interactions with immunosuppressive agents and adjust dosage if renal impairment. (Cyclosporine enhances the effect of muscle relaxants producing a prolonged block.)
- NSAIDs can cause further renal toxicity in addition to immunosuppressants.
- Prevent additional mechanical obstruction (ventilator-induced disease and excessive tidal volumes) and employ ventilator with capability for variable inspiratory and expiratory ratios.
- Lateral decubitus position may aggravate V/Q mismatch.

- Hyperventilation during mechanical ventilation should be avoided because seizure threshold in pts taking immunosuppressive agents may be lowered.
- Use shorter-acting agents to avoid prolonged CNS, CV, and resp depression to facilitate a swift recovery of functions and timely extubation.

Extubation

- Delay until adequate ventilation is assured (sustained tetanus on monitoring).
- The lack of cough reflex below the tracheal anastomosis makes pts unable to clear secretions, unless they are awake, increasing the risk of silent aspiration.

Adjuvants

- Consider regional technique because it allows opioid sparing, but dense intercostal blockade can delay extubation in pts with poor respiratory reserve.

Postoperative Period

- Monitor for and aggressively treat resp depression, infection, and fluid overload.

Anticipated Problems/Concerns

- Many pts with resting hypoxia and marginal compensated lung functions come to OR for diagnostic lung biopsy. A thoracoscopic technique may be impossible owing to adhesions post heart/lung transplantation or pt's inability to tolerate one-lung ventilation.
- Anticipate further perioperative resp decompensation after open-lung biopsy.
- Arrange postop disposition (monitored bed and ventilator support) depending on preop functional status and the potential for periop complications.

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Bronchitis, Chronic

C. William Hanson III

Risk

- Incidence in USA: 14 million
- Race with highest prevalence: Caucasian
- M:F ratio 1:2
- Smoking, second-hand smoke, occupational exposure to pulm toxic substances (radon, coal, silicates, and asbestos)

Perioperative Risks

- Bronchospasm

Worry About

- Airway stimulation at light levels of anesthesia
- Laryngospasm (due to secretions and hyperreactivity)
- Hypoxia
- Hypercarbia

Overview

- Chronic productive cough with periodic exacerbations (most d for at least 3 mo and for at least 2 consecutive y)
- Enlargement of the mucus-secreting glands in the airways with excessive sputum production
- Obstruction of expiratory airways
- Derangement in V/Q relationships
- Chronic hypoxia with right heart failure
- Exacerbations with intercurrent bacterial or viral infections

Etiology

- Acquired, usually due to smoking
- May also be due to asthma or frequent childhood resp infections

Usual Treatment

- Avoidance of environmental irritants such as cigarette smoke (preferably >8–10 wk before elective surgery)
- Antibiotics for acute exacerbations; inefficacious for prophylactic treatment
- Oral glucocorticoids: appropriate for acute exacerbations but not for maintenance therapy
- Periop stress dose glucocorticoid (methylprednisolone, dexamethasone, and hydrocortisone) administration: may be appropriate in pts on prolonged (>3 wk) high dose (≥20 mg prednisone per day) oral steroids
- Short-acting bronchodilators, such as beta agonists or anticholinergics, for acute exacerbations and long-acting beta agonist bronchodilators plus inhaled steroids for long-term maintenance therapy; pts on inhalers may be treated with preintubation inhalation of a beta agonist

Assessment Points

System	Effect	Assessment by Hx	PE	Test
HEENT	Short, fat neck			
CV	Right heart failure	Exercise tolerance	RV heave Dependent edema	ECG ECHO
	Pulm Htn			PA catheter
RESP	Airways obstruction	Smoking Hx (current, recent, emote) Number and severity of recent exacerbations	Cyanosis	PFT, DLCO, ABGs
MS			Clubbing of fingers	

Key References: Kim V, Criner GJ: Chronic bronchitis and chronic obstructive pulmonary disease, *Am J Respir Crit Care Med* 187: 228–237, 2013; Yamakage M, Iwasaki S, Namiki A: Guideline-oriented perioperative management of patients with bronchial asthma and chronic obstructive pulmonary disease, *J Anesth* 22:412–428, 2008.

Perioperative Implications**Preoperative Preparation**

- Smoking cessation
- Antibiotics to decrease sputum production
- Resp conditioning

Monitoring

- Consider arterial line to monitor blood gases
- Consider pulm artery catheter for large fluid-shift operations

Airway

- Often, truncal obesity (especially with corticosteroids); may have redundant soft tissue in airway or a short, fat neck

Preinduction/Induction

- Avoid stimulating the airway while pt is in light levels of anesthesia because it may precipitate bronchospasm (although less likely than with asthma).
- Regional anesthesia may be preferable.

Maintenance

- Frequent suctioning of ETT
- Limit narcotic administration (danger of periop CO₂ retention)
- Adjuvant regional anesthesia for postop pain management in procedures that affect resp mechanics (e.g., intercostal nerve blocks, epidural analgesia)

Extubation

- Administer intratracheal bronchodilator in responsive pts before extubation.
- Consider IV lidocaine before extubation.

Anticipated Problems/Concerns

- Postop resp complications (secretions, mucus plugging, atelectasis, pneumonia, and prolonged requirement for mechanical ventilation)

Bronchopulmonary Dysplasia

Marissa G. Vadi | Ryan E. Lauer

Risk

- Incidence in USA: 10,000-15,000 infants annually
- Risk increases with decreasing gestational age and birth weight
- Affects at least one-quarter of infants with birth weights <1500 g
- No race or gender predilection

Perioperative Risks

- Bronchospasm
- Pulm Htn
- Cor pulmonale

Worry About

- Airway obstruction and hyperreactivity
- Pulm Htn and cor pulmonale
- “BPD spells”: Acute cyanotic events caused by increases in central airway resistance
- Tracheomalacia and/or bronchomalacia
- Recurrent pulm infections

Overview

- Chronic lung disease associated with premature birth and positive pressure mechanical ventilation, the clinical definition of which has evolved over time
 - “Classic BPD”: Associated with characteristic radiographic changes and four stages of lung injury: exudative → necrosis → pulm fibrosis → severe cystic changes, and cor pulmonale
 - “New BPD”: Seen after introduction of surfactant therapy, antenatal steroid administration, and improved neonatal ventilator strategies; mild respiratory distress syndrome and continued need for supplemental oxygen; and lung development is uniformly arrested, with simplified alveolar structures and dysmorphic capillaries
- Disease severity (mild, moderate, or severe) determined by the gestational age of the infant, oxygen dependency at 36 wk postconceptional age, total duration of oxygen supplementation, and positive pressure requirements

- Chronic airway obstruction and hyperreactivity present in long-term survivors
- High risk of periop morbidity if pulm Htn present

Etiology

- Multifactorial; arrest of pulm development ± inflammation
- Major risk factors: premature birth, respiratory failure, oxygen supplementation, and mechanical ventilation
- Impaired angiogenesis, which reduces alveolar-capillary gas exchange, leading to hypoxemia and increased PVR

Usual Treatment

- Supplemental oxygen
- Inhaled bronchodilators (e.g., β-agonists)
- Pulm vasodilator therapies (e.g., sildenafil, calcium channel blockers, bosentan) if pulm Htn present

Assessment Points

System	Effect	Assessment by Hx	PE	Test
HEENT	Possible tracheomalacia	SOB Stridor	Retractions Audible stridor	Bronchoscopy for severe cases
CV	PulmHtn Cor pulmonale	Exertional dyspnea Syncope Cyanosis, esp. during feeds Supplemental O ₂ dependence	Hypoxia Cyanosis ± clubbing Rales Peripheral edema Elevated JVP	ECG ECHO Cardiac cath
RESP	Possible bronchomalacia Decreased tidal volumes Increased airway resistance Decreased dynamic lung compliance Hypoxia Hypercapnia	SOB Cyanotic spells Supplemental O ₂ dependence Asthmalike symptoms Recurrent respiratory infections	Tachypnea Retractions Cyanosis ± clubbing Expiratory wheezing Rales	CXR ABG
GI	Failure to thrive	Poor feeding	Low BMI	Generally not needed

Key References: Jensen EA, Schmidt B: Epidemiology of bronchopulmonary dysplasia, *Birth Defects Res A Clin Mol Teratol* 100(3):145–157, 2014; Lauer R, Vadi M, Mason L: Anaesthetic management of the child with co-existing pulmonary disease, *Br J Anaesth* 109(21):i47–i59, 2012.

Perioperative Implications**Preoperative Preparation**

- Determine room-air oxygen saturation and baseline supplemental oxygen requirements.
- Obtain electrolytes (if pt receiving chronic diuretic therapy); ABG (if oxygen requirements recently increased); and ECHO (if clinical markers concerning for pulm Htn).

- Avoid general anesthesia for elective procedures during acute respiratory infection.
- Avoid spinal anesthesia in patients with severe pulm Htn; decreased venous return and bradycardia may precipitate right heart failure.
- Consider preoperative nebulized β₂ agonist and/or steroid administration.
- Administer premedication cautiously in pts with pulm Htn.

Monitoring

- Standard ASA monitors.
 - Monitor pulse oxygen saturation, end-tidal carbon dioxide, and body temperature closely. Abnormalities may worsen pulm Htn.
- Consider arterial cannulation for invasive blood pressure monitoring and central venous line placement for inotrope administration in pts with pulm Htn.