

Anaesthesia recommendations for patients suffering from

Bronchopulmonary dysplasia

Disease name: Bronchopulmonary dysplasia

ICD 10: P27.1

Synonyms: Chronic lung disease of prematurity

Bronchopulmonary dysplasia (BPD) is a chronic lung disease that remains one of the most prevalent long-term sequelae of premature birth. Northway and colleagues first described the condition in 1967 after 32 preterm infants (mean gestational age 34 weeks) with respiratory distress syndrome developed characteristic radiographic changes following the initiation of positive pressure mechanical ventilation [1]. Four stages of lung injury were described in “classic” BPD: exudative (age 1-3 days); necrosis and early repair (age 4-10 days); microcyst formation and pulmonary fibrosis (age 10-12 days); and severe cystic changes and cor pulmonale (after 30 days of age) [2].

The clinical definition of BPD has evolved with time and advancements in neonatal care including surfactant therapy, antenatal steroid administration, and improved ventilator strategies [3,4]. Most infants currently developing BPD are born between 24-28 weeks gestational age, during the time of canalicular and saccular development. “New” BPD is characterized by uniform arrest of lung development, with simplified alveolar structures and dysmorphic capillaries. These infants are more likely to present with a mild respiratory distress syndrome and a continued need for supplemental oxygen [5]. Three levels of BPD severity (mild, moderate, or severe) are determined by the gestational age of the infant, oxygen dependency at 36 weeks post-conceptual age, total duration of oxygen supplementation, and positive pressure requirements [6].

Long-term BPD survivors may experience persistent airway obstruction and pulmonary hypertension, complicating anaesthetic management.

Medicine in progress



Perhaps new knowledge

Every patient is unique

Perhaps the diagnostic is wrong



Find more information on the disease, its centres of reference and patient organisations on Orphanet: www.orpha.net

Typical surgery

Procedures addressing sequelae of prematurity includes: gastrostomy tube placement; inguinal hernia repair; ligation of patent ductus arteriosus; ophthalmologic procedures; ventriculoperitoneal shunt placement.

Type of anaesthesia

There is no definite recommendation for either general or regional anaesthesia.

Long-term BPD survivors experience airway obstruction and hyperreactivity. It is recommended that BPD patients not undergo general anaesthesia for elective procedures during an acute respiratory infection; surgery should ideally be postponed 4-6 weeks. Laryngeal mask airway may decrease bronchospasm risk, but does not protect the airway from aspiration [7]. Regional or local anaesthesia avoids airway manipulation, but may not be feasible for certain sites of surgery or in uncooperative paediatric patients. Avoid spinal anaesthesia in patients with severe pulmonary hypertension and right heart strain, as decreased venous return and bradycardia may precipitate right heart failure.

Necessary additional diagnostic procedures (preoperative)

A thorough medical history must be obtained. For BPD infants requiring surgery while on the neonatal unit, determine the anaesthetic history, current medications, allergies, cough or sputum production, supplemental oxygen requirements, and the extent of positive pressure ventilator support.

Additionally, question parents of BPD patients returning for follow-up procedures after neonatal unit discharge regarding interval hospitalizations (including the need for tracheal intubation), changes in supplemental oxygen requirements, non-invasive positive pressure requirements (ex. CPAP), and exercise tolerance. Poor exercise tolerance may present as diaphoresis or cyanosis during feeding or failure to thrive in patients not yet able to ambulate.

Physical examination should assess vital signs, presence of wheezing or cough, accessory muscle use, cyanosis, and hydration status.

Consider the following testing in patients with persistent disease:

- Electrolyte panel: in patients receiving chronic diuretic therapy
- Arterial blood gas and pulse oximetry: in patients with recent increases in oxygen requirements or in any infant requiring supplementary oxygen
- Chest radiograph

- Echocardiogram: in any infant remaining oxygen dependent or in patients with clinical markers concerning for pulmonary hypertension such as a continued need for positive pressure ventilation, oxygen requirements out of proportion to the degree of lung disease, elevated PaCO₂, recurrent cyanotic episodes, failure to thrive, or recurrent hospitalizations [8]
- Cardiac catheterization: reserved for BPD patients with pulmonary hypertension by echocardiogram who (1) have clinical cardiorespiratory deterioration not related to airways disease; (2) are suspected of having significant pulmonary hypertension despite optimal management of lung disease; (3) are candidates for long-term pulmonary hypertension drug therapy; or (4) have unexplained, recurrent pulmonary edema [9].

Particular preparation for airway management

BPD has no direct correlation with difficult tracheal intubation. However, BPD patients are at risk for subglottic stenosis, airway granulomas, and pseudopolyps if prolonged tracheal intubation occurred during infancy [9-11]. Smaller tracheal tubes may be required.

Particular preparation for transfusion or administration of blood products

Retrospective studies have shown correlations between the receipt of blood transfusions and the development of BPD in very low-birth weight infants [12-14], possibly due to iron-induced oxidative stress [15]. However, there is no clear evidence for a causal relationship.

Patients with known BPD may require diuretics administration in conjunction with transfusion to avoid pulmonary edema and worsening hypoxemia.

Particular preparation for anticoagulation

There is no evidence to support the need of particular anticoagulation.

Particular precautions for positioning, transport or mobilisation

Avoid hypothermia, hypoxia, and hypercarbia as these factors may worsen pulmonary hypertension and precipitate right ventricular failure. Oxygen should be made available for operating theatre transport. Patients requiring positive pressure ventilation should be intubated and ventilated during transport. As ineffective bag ventilation may result in hypercapnea, consider mechanical transport ventilator use to ensure consistent minute ventilation.

Probable interaction between anaesthetic agents and patient's long-term medication

Post-natal systemic or inhaled corticosteroids are administered by some centres to reduce inflammation and improve lung function in infants with evolving or established BPD [16-19]. Long-term BPD survivors may receive oral, intravenous, or inhaled steroids as treatment for severe reactive airway disease in the outpatient setting.

All routes of glucocorticoid administration (oral, inhaled, intranasal, topical, intramuscular, and intravenous) have been associated with suppression of the hypothalamic-pituitary-adrenal axis [20]. Physiologic stress including injury, surgery, or severe infection may precipitate adrenal crisis. Consider adrenal suppression in patients with a history of exogenous steroid use and unexplained hypotension or hypoglycaemia under anaesthesia. Administer stress-dose glucocorticoid supplementation in patients who are on long-standing oral steroid therapy (greater than 8-12 mg/m²/day for greater than 2 weeks), or those who have discontinued steroid use within the last 6 months [21].

Anaesthesiologic procedure

Anaesthetic management goals include avoidance of bronchoconstriction, elevations in pulmonary vascular resistance, or decreases in cardiac contractility.

Propofol is a bronchodilator and is a useful induction agent in the haemodynamically stable patient with reactive airway disease. This agent should be used with caution in patients with pulmonary hypertension, as large boluses can significantly decrease systemic vascular resistance and impair biventricular function [22].

Ketamine maintains arterial pressure and systemic vascular resistance while simultaneously enhancing bronchodilatation. It is useful in the haemodynamically unstable and actively wheezing patient requiring emergency surgery. Ketamine administration in children with pulmonary hypertension remains controversial, as some studies suggest its use may increase pulmonary vascular resistance [23-25].

Volatile agents decrease hypoxic pulmonary vasoconstriction. Sevoflurane is preferable for inhalational inductions due to its bronchodilating effects as well as an association with a decreased incidence of laryngospasm and cardiac arrhythmias when compared to other volatile agents. Carefully titrate volatile agents in patients with pulmonary hypertension, as their use may result in a dose-dependent depression of cardiac contractility and systemic vascular resistance [22].

Tracheal intubation may precipitate bronchospasm. Consider avoiding tracheal intubation by using a mask or supraglottic airway device for appropriate cases [7]. Ensure a deep level of anaesthesia before airway manipulation. Deep extubation reduces the risk of bronchospasm from coughing on the tracheal tube but does not protect the patient from aspiration or laryngospasm [22]. Regional anaesthesia avoids airway manipulation but is not appropriate for all sites of surgery.

Avoiding histamine-releasing neuromuscular blocking agents in patients with BPD is advisable. Acetylcholinesterase inhibitors need to be used with caution during reversal of neuromuscular blockade due to the risk of bronchospasm. Sugammadex reverses neuromuscular blockade without muscarinic side effects but has not been extensively studied in the infant and neonatal populations [26].

Particular or additional monitoring

Monitor pulse oxygen saturation, end tidal carbon dioxide, and body temperature to avoid hypoxia, hypercarbia, and hypothermia. These factors may worsen pulmonary hypertension and lead to right heart failure.

Perform arterial blood gas sampling at regular intervals.

Consider arterial cannulation for invasive blood pressure monitoring and central venous line placement for inotrope administration in patients with pulmonary hypertension for surgical cases of increased length or complexity.

Monitor neuromuscular blockade and ensure its effects are completely reversed after surgery.

Possible complications

Children with BPD frequently experience wheezing, but are less likely to respond to bronchodilators than children with asthma [27,28]. Still, metered dose inhaler administration of bronchodilators is safe and effective for many BPD patients during acute bronchospasm [29]. Refractory bronchospasm may require IV terbutaline or epinephrine administration as a last resort.

In patients with pulmonary hypertension, avoid increases in pulmonary vascular resistance to prevent a pulmonary hypertensive crisis. Moderate hyperventilation with 100% oxygen, correction of acidosis, improved analgesia, and the administration of pulmonary vasodilators are useful in this setting. Inhaled nitric oxide decreases pulmonary vascular resistance without significant systemic effects. Inotropic support is considered for persistent systemic hypotension despite pulmonary vasodilator therapy [30].

Postoperative care

The degree of post-operative monitoring is dependant on the surgical procedure and the physical condition of the patient. Intensive care unit admission may be required for patients experiencing pulmonary hypertensive crisis or requiring mechanical ventilation. Prolonged post-operative ventilation may promote ventilator-associated lung injury and should be avoided.

Information about emergency-like situations / Differential diagnostics

Not reported.

Ambulatory anaesthesia

Typically avoided. Ambulatory anaesthesia is only considered in BPD patients with mild disease (i.e. no baseline wheezing, cyanosis, home oxygen use, or pulmonary hypertension).

Obstetrical anaesthesia

Not reported.

Literature and internet links

1. Northway WH Jr, Rosan RC, Porter DY. Pulmonary disease following respirator therapy of hyaline-membrane disease. Bronchopulmonary dysplasia. *N Engl J Med* 1967;276:357-68
2. Northway WH Jr. Bronchopulmonary dysplasia: then and now. *Arch Dis Child* 1990; 65(10 Spec. No.):1076-81
3. Philip AG. Bronchopulmonary dysplasia: then and now. *Neonatology* 2012;102:1-8
4. Maxwell LG, Goodwin SR, Mancuso TJ, et al. Systemic disorders. In: Davis, PJ, Cladis FP, Motoyama EK, eds. *Smith's Anesthesia for Infants and Children*, 8th Ed. Philadelphia: Elsevier Mosby, 2011:1120
5. Coalson JJ. Pathology of bronchopulmonary dysplasia. *Semin Perinatol* 2006;30:179-84
6. Ehrenkranz RA, Walsh MC, Vohr BR, et al. Validation of the National Institutes of Health consensus definition of bronchopulmonary dysplasia. *Pediatrics* 2005;116:1353-60
7. Ferrari LR and Goudsouzian NG. The use of the laryngeal mask airway in children with bronchopulmonary dysplasia. *Anesth Analg* 1995;81(2):310-3
8. Abman SH, Hansmann G, Archer SL, et al.; on behalf of the American Heart Association Council on Cardiopulmonary, Critical Care, Perioperative and Resuscitation, Council on Clinical Cardiology, Council on Cardiovascular Disease in the Young, Council on Cardiovascular Radiology and Intervention, Council on Cardiovascular Surgery and Anesthesia, and the American Thoracic Society. Pediatric pulmonary hypertension: guidelines from the American Heart Association and American Thoracic Society. *Circulation*. 2015;132:2037-99
9. Doull IJ, Mok Q, Tasker RC. Tracheobronchomalacia in preterm infants with chronic lung disease. *Arch Dis Child Fetal Neonatal Ed* 1997;76:F203-5
10. Duncan S, Eid N. Tracheomalacia and bronchopulmonary dysplasia. *Ann Otol Rhinol Laryngol* 1991;100:856-8
11. Miller RW, Woo P, Kellman RK, Slagle TS. Tracheobronchial abnormalities in infants with bronchopulmonary dysplasia. *J Pediatr* 1987;111:779-82
12. Korhonen P, Tammela O, Koivisto A.-M, et al. Frequency and risk factors in bronchopulmonary dysplasia in a cohort of very low birth weight infants. *Early Hum Dev* 1999;54:245-58
13. Hernandez-Ronquillo L, Tellez-Zenteno JF, Weder-Cisneros, N, et al. Risk factors for the development of bronchopulmonary dysplasia: a case-control study. *Arch Med Res* 2004;35:549-53
14. Zhang H, Fang J, Su H, et al. Risk factors for bronchopulmonary dysplasia in neonates born at ≤ 1500 g (1999-2009). *Pediatr Int* 2011;53:915-20
15. Collard KJ. Transfusion related morbidity in premature babies: possible mechanisms and implications for practice. *World J Clin Pediatr* 2014;3(3):19-29
16. Doyle LW, Ehrenkranz RA, Halliday HL. Early (< 8 days) postnatal corticosteroids for preventing chronic lung disease in preterm infants. *Cochrane Database Syst Rev* 2014; 5:CD001146
17. Doyle LW, Ehrenkranz RA, Halliday HL. Late (> 7 days) postnatal corticosteroids for chronic lung disease in preterm infants. *Cochrane Database Syst Rev* 2014; 5:CD001145
18. Shah V, Ohlsson A, Halliday HL, Dunn MS. Early administration of inhaled corticosteroids for preventing chronic lung disease in ventilated very low birth weight preterm neonates. *Cochrane Database Syst Rev* 2007; :CD001969
19. Onland W, Offringa M, van Kaam A. Late (≥ 7 days) inhalation corticosteroids to reduce bronchopulmonary dysplasia in preterm infants. *Cochrane Database Syst Rev* 2012; 4:CD002311
20. Bancos I, Hahner S, Tomlinson J, et al. Diagnosis and management of adrenal insufficiency. *Lancet Diabetes Endocrinol* 2015;3(3):216-26
21. Ahmet A, Kim H, Spier S. Adrenal suppression: A practical guide to the screening and management of this under-recognized complication of inhaled corticosteroid therapy. *Allergy Asthma Clin Immunol* 2011;7:13
22. Lauer R, Vadi M, Mason L. Anaesthetic management of the child with co-existing pulmonary disease. *Br J Anaesth* 2012;109(S1):i47-i59
23. Morray JP, Lynn AM, Stamm SJ, et al. Hemodynamic effects of ketamine in children with congenital heart disease. *Anesth Analg* 1984;63:895-9

24. Berman W Jr, Fripp RR, Rubler M, et al. Hemodynamic effects of ketamine in children undergoing cardiac catheterization. *Pediatr Cardiol* 1990; 11:72-6
25. Wolfe RR, Loehr JP, Schaffer MS, et al. Hemodynamic effects of ketamine, hypoxia and hyperoxia in children with surgically treated congenital heart disease residing greater than or equal to 1,200 meters above sea level. *Am J Cardiol* 1991;67:84-7
26. Plaud B, Meretoja O, Hofmockel R, et al. Reversal of rocuronium-induced neuromuscular blockade with sugammadex in pediatric and adult surgical patients. *Anesthesiology* 2009;110:284-94
27. Allen JL, Panitch HB. Lung function testing: chronic lung disease of infancy. *Pediatr Pulmonol* 2001; Suppl 23:138.
28. Joshi S, Powell T, Watkins WJ, et al. Exercise-induced bronchoconstriction in school-aged children who had chronic lung disease in infancy. *J Pediatr* 2013; 162:813.
29. Hingston CD, Homes TW, Wise MP. Airway emergency during anaesthesia using a metered-dose inhaler—III. *Anaesthesia* 2011;66(6):531.

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This guideline has been prepared by:

Authors

Marissa Vadi, Anaesthesiologist, Loma Linda University, Loma Linda, California, USA
mvadi@llu.edu

Mathew Malkin, Anaesthesiologist, Loma Linda University, Loma Linda, California, USA
mmalkin@llu.edu

Ryan Lauer, Anaesthesiologist, Loma Linda University, Loma Linda, California, USA
rlauer@llu.edu

Peer revision 1

Anne Greenough, Neonatal intensive care unit, King's College Hospital, London, United Kingdom
anne.greenough@kcl.ac.uk

Peer revision 2

Kathleen M Deakins, Paediatric Respiratory Care, Rainbow Babies and Children's Hospital, Cleveland OH, USA
kathleen.deakins@uhhospitals.org
