

# Clinical Guidance

## Paediatric Critical Care: Severe Bronchiolitis

### Summary

This guideline is for those patients with severe bronchiolitis who are being considered for critical care, thus interventions like chest x-ray, bloods, fluid restriction and antibiotics are appropriate. For other patients please see the NICE guideline.

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<p>This clinical guideline has been produced by the South Thames Retrieval Service (STRS) at Evelina London for nurses, doctors and ambulance staff to refer to in the emergency care of critically ill children.</p> <p>This guideline represents the views of STRS and was produced after careful consideration of available evidence in conjunction with clinical expertise and experience. The guidance does not override the individual responsibility of healthcare professionals to make decisions appropriate to the circumstances of the individual patient.</p>	

Change History		
Date	Change details, since approval	Approved by
Aug 18	Reviewed, minor clarifications made. Milesi reference added Use of PCT (Procalcitonin) added as per ELCH PICU policy	ELGG

Bronchiolitis is a common viral respiratory condition associated with lower airway obstruction, air trapping and atelectasis. This guideline is for those patients with severe bronchiolitis who are being considered for critical care, thus interventions like chest x-ray (CXR), bloods, fluid restriction and antibiotics are appropriate. For other patients please see the [NICE guideline](#).

**Clinical presentation**

- Fever, rhinitis and cough
- increased work of breathing, tachypnoea or wheeze
- Apnoeas (esp. <2 months)
- Cyanosis
- Poor feeding

**Criteria for severe disease**

- Saturations < 92% despite oxygen
  - Respiratory rate > 70 breaths / minute
  - Signs of severe respiratory distress
  - Apnoea
  - Decreased level of consciousness
- Lower threshold in high risk group (see right)

**High risk groups**

- Prematurity and neonates
- Pre-existing respiratory condition
- Congenital heart disease (CHD)
- Neuromuscular conditions (may not have signs of distress)
- Immune deficiency

**Baseline investigations**

- Nasopharyngeal aspirate for respiratory viruses (RSV)
- Baseline CXR
- Full blood count with differential (pertussis-lymphocytosis)
- Electrolytes (including plasma sodium)
- Blood gases unhelpful in directing need for resp support

**Management principles on ward / HDU**

- Maintain oxygen saturations  $\geq 90\%$  unless congenital heart disease/CHD (seek advice)
  - Apnoea monitoring if required (history, <2 months)
  - Minimal handling
  - Suction nasal secretions if obstructed with mucus
  - Non-invasive respiratory support by humidified high flow nasal cannula at 2 L/kg/min or CPAP 5-6cm H<sub>2</sub>O (CPAP offers better resp support). Reassess frequently.
  - Small volume, frequent nasogastric feeds if possible
  - Reduce enteric fluid intake to 50ml/kg/day (risk of fluid overload/hyponatraemia/seizures)
  - If IV fluid required-must be isotonic (e.g. 0.9% sodium chloride with 5% glucose). Run at 50ml/kg/day.
  - Antibiotics: Co-amoxiclav IV: treat empirically if referred for PICU (30-40% have bacterial isolates). Consider cefotaxime <1month or ceftriaxone >1month if CNS concerns (+/- aciclovir) or penicillin allergy\*.
- No role for nebulised salbutamol, ipratropium bromide, adrenaline, hypertonic sodium chloride or steroids
  - Respiratory stimulants (e.g. caffeine) have no role

**Assessment for ventilatory support & PICU transfer**

- Severe respiratory distress or risk of respiratory arrest
- Lack of clinical improvement or deterioration on non-invasive respiratory support
- O<sub>2</sub> requirement  $\geq 60\%$  to maintain saturations  $\geq 90\%$
- Persistent or recurrent apnoeas
- Lack of improvement in level of consciousness
- Neuromuscular patients may not be able increase their work of breathing- consider gas in these patients
- Seizures

**Intubation and ventilation**

- **Optimise pre-oxygenation**
- Decompress stomach by nasogastric tube aspiration
- Consider volume bolus (10ml/kg) prior to anaesthesia
- Anticipate non-compliant chest once anaesthetised and muscle relaxed. Mask ventilate with slow respiratory rate (20-30) to achieve good chest movement
- Choose appropriate endotracheal tube(ETT) to minimise leak
- Ensure end tidal (ET)CO<sub>2</sub> monitoring available
- **Initial ventilation: I time 0.8s, RR 20-30, PEEP 5cm H<sub>2</sub>O, Enough PIP to move chest (ideally <30cm H<sub>2</sub>O)**
- Secure ETT (see guideline) and CXR
- Review chest and ventilation settings regularly
- Target oxygen saturations >92% unless CHD
- Target ETCO<sub>2</sub> 5-10KPa
- Sedate with morphine & muscle relax as required
- Suctioning of ETT may be helpful
- Arterial line not usually required for management

**Troubleshooting difficulties on ventilator- DOPES**

- **D**isplaced ETT-check ETCO<sub>2</sub> and exact length of tube
  - **O**bstruction- suction ETT and check passes to end of ETT
  - **P**neumothorax-clinical examination- can be difficult to exclude if chest hyper-expanded due to air trapping
  - **E**quipment- check ventilator settings including O<sub>2</sub>.
  - **S**tomach- Ensure decompressed with nasogastric tube
- Assess DOPES first, CXR if problem not resolved**

**Further management on PICU**

- Bronchoalveolar lavage specimen for viral+bacterial culture
- White cell count, haemoglobin, electrolytes, CRP & PCT
- Continue antibiotics (see left) for 48hrs then review stopping or continue for 5 days. Enteral co-amoxiclav if no sepsis concerns. Consider gentamicin if high fever/WCC/CRP/PCT
- Consider further tests if recurrent resp. presentations

**Outcome and discharge from PICU to ward**

- Mortality <1%. Underlying co-morbid group has highest risk
- Length of ventilation, usually 3-7 days
- Oxygen requirement <2litres/min nasal cannula
  - No apnoeas for 12 hours

\*Consult microbiology if severe allergy (anaphylaxis: angioedema, airway swelling, hypotension)