



# IAEM Clinical Guideline

# Laryngomalacia

Version 1

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#### **DISCLAIMER**

IAEM recognises that patients, their situations, Emergency Departments and staff all vary. These guidelines cannot cover all clinical scenarios. The ultimate responsibility for the interpretation and application of these guidelines, the use of current information and a patient's overall care and wellbeing resides with the treating clinician.

# LARYNGOMALACIA (non-acute stridor in infants)

### INTRODUCTION

Laryngomalacia is defined as collapse of supraglottic structures during inspiration, resulting in intermittent air flow impedance and associated stridor. It is the most common cause of stridor in newborns, affecting 60–75% of all infants with congenital stridor. These patients either self-present or are sometimes referred by their General Practitioner (GP) to the Emergency Department (ED).

Laryngomalacia presents with inspiratory stridor that typically worsens with feeding, crying, supine positioning and agitation. The symptoms begin at birth or within the first few weeks of life, peak at 6 to 8 months, and typically resolve by 12 to 24 months.

### **PARAMETERS**

#### **Target patient population**

This evidence summary applies to well infants presenting with a history of established "noisy breathing". It does not apply to those with acute or severe airway obstruction.

#### Target users

Healthcare professionals engaging in the care of infants in the acute care setting.

### **AIM**

This guideline aims to provide a tool for assessment of infants presenting with laryngomalacia and to guide their management.

### **ASSESSMENT**

## **Assessment of severity**

Along with monitoring vital signs, respiratory and general physical exam, it is practical to observe the infant while taking a feed and notice any difficulty or distress during feeding.

Assess severity based on the feeding and respiratory symptoms

| Severity level | Respiratory symptoms                                                                    | Feeding symptoms                                                   |
|----------------|-----------------------------------------------------------------------------------------|--------------------------------------------------------------------|
| Mild           | Inspiratory stridor with mild increase in work of breathing                             | Occasional cough or regurgitation                                  |
| Moderate       | Recent worsening of inspiratory stridor, significant increase in work of breathing      | Frequent regurgitation  Choking with feeds or other feeding issues |
| Severe         | Inspiratory stridor with cyanosis, apnoea, with associated chest wall signs of distress | Failure to thrive Evidence of aspiration                           |

#### Risk factors for moderate to severe disease

- Gastroesophageal and Laryngopharyngeal Reflux
- Neurological disorders
- Secondary airway disease (SAL)
- Congenital heart disease
- Congenital anomalies, Syndromes, Genetic Disorders

# Differential diagnosis of stridor

| Inspiratory Stridor                       | Biphasic Stridor                                | Expiratory stridor                                | General                               |
|-------------------------------------------|-------------------------------------------------|---------------------------------------------------|---------------------------------------|
| Airway obstruction above the vocal cords. | Airway obstruction at or below the vocal cords. | Airway obstruction at or below the lower trachea. | Airway<br>obstruction at any<br>level |
| Epiglottitis                              | Subglottic stenosis                             | Bronchiolitis                                     | Foreign body                          |
| Croup                                     | Subglottic cyst/                                | Asthma                                            |                                       |
| Vallecular cysts                          | Haemangioma                                     | Tracheomalacia                                    |                                       |
|                                           | Vocal cord paralysis                            | Complete                                          |                                       |
|                                           | Laryngeal web                                   | tracheal ring                                     |                                       |
|                                           | Respiratory papillomatosis                      | Double aortic arch                                |                                       |
|                                           | раршошаючэ                                      | Pulmonary                                         |                                       |
|                                           |                                                 | artery sling                                      |                                       |
|                                           |                                                 | Aberrant innominate artery                        |                                       |
|                                           |                                                 | •                                                 |                                       |

#### Also ask about:

Feeding: regurgitation, emesis, cough, choking, and slow feedings.

Growth: increased metabolic demand of coordinating eating and breathing against the obstruction can lead to weight loss and failure to thrive.

## **Respiratory symptoms:**

- **Common:** tachypnoea, suprasternal and substernal retractions
- **Uncommon:** cyanosis, pectus excavatum, and obstructive sleep apnoea.
- Chronic hypoxia from airway obstruction can lead to pulmonary hypertension.

# **WARNING SIGNS**

| Respiratory                       | Feeding                        |
|-----------------------------------|--------------------------------|
| Stridor with respiratory distress | Choking with feeding           |
| Dyspnoea with retractions         | Episodic cyanosis with feeding |
| Pectus excavatum                  | Recurrent aspiration pneumonia |
| Pulmonary hypertension            | Failure to thrive              |
| Cor pulmonale                     |                                |
| Severe obstructive sleep apnoea   |                                |

#### MANAGEMENT OF LARYNGOMALACIA

#### Mild disease:

- Conservative management with follow up with General Practitioner for weight gain and worsening of respiratory or feeding symptoms.
- Parents should be advised on positional therapy (feeding in upright position),
   feeding interventions (thickening formula or breast feed) and for treatment for reflux where appropriate.
- Use GP proforma letter and give the Parent information leaflet.

#### Moderate to Severe disease:

- Symptomatic management with airway management, supplemental oxygen and nasogastric feed as required.
- These patients need admission under medical paediatric team.
- These patients should be discussed with ENT team for an urgent consultation.

### **COMPANION DOCUMENTS**

- GP proforma letter
- Parent information leaflet
- References / Evidentiary table