Injection laryngoplasty for laryngeal cleft type I in an 8-week-old infant

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SUMMARY
A laryngeal cleft is a rare anatomical deformity which is increasingly treated with injection laryngoplasty. Since diagnosis of laryngeal cleft type I is often made between 2 and 5 years of age, this treatment is rarely performed on very young children. In this case, we describe how injection laryngoplasty is performed safely on an 8-week-old child, and we illustrate its added value for the diagnostic process and for temporary symptom relief.

BACKGROUND
A laryngeal cleft (LC) is an anatomical defect, consisting of a deeper notch in the interarytenoid region. This can result in overflow of food and liquids towards the trachea, and cause symptoms of aspiration, cough, recurring respiratory tract infections and respiratory distress. This can lead to diminished quality of life (QoL). A type I laryngeal cleft (T1LC) only extends to the supraglottic interarytenoid region, no further than the true vocal folds. A subclassification of T1LC is the deep interarytenoid notch (DIN). A DIN has normal musculature and a shorter interarytenoid notch height compared to T1LC. Symptoms of T1LC are often non-specific. Because of this less distinct presentation of symptoms, diagnosis of T1LC is often delayed. According to previous research, type I and II LC are usually diagnosed between the age of 2 and 5 years. Gold standard for diagnosis is through palpation of the interarytenoid region using a probe, during suspension laryngoscopy under general anaesthetic. In approximately 58%–68% of patients with an LC, other congenital malformations are found. Associated anomalies are most commonly found in the gastrointestinal tract, including oesophageal atresia, tracheoesophageal fistula (TEF) and intestinal malrotation. Furthermore, an LC can lead to diminished QoL, by causing emotional and financial stress. Injection laryngoplasty (IL) using a dermal filler is increasingly used in the management of LC. This treatment is not often performed on newborn children, as in this case. Literature on the efficacy and safety of IL in younger children is therefore limited.

INVESTIGATIONS
At 7 weeks of age, Videofluoroscopic Swallowing Study (VFSS) was carried out to assess the suspected LC. The patient was assessed as a level 2 on the Penetration Aspiration Scale. During swallowing of slightly thick liquid (International Dysphagia Diet Standardization Initiative (IDDSI) level 1), penetration was seen several times, but no aspiration. There was no penetration or aspiration seen while swallowing moderately thick liquids (IDDSI level 2). To evaluate possible TEF recurrence, the patient underwent a dLTB at 8 weeks of age. Due to respiratory distress during this investigation, the presence of a recurred TEF could not be assessed, but severe obstructing tracheobronchomalacia with CO₂ accumulation was observed in addition to an oesophageal stenosis. Also, palpation of the interarytenoid area indicated either a T1LC or a DIN.

DIFFERENTIAL DIAGNOSIS
Since TEF recurrence could not be determined during dLTB, presenting symptoms could be attributed to either one or several of the following diagnoses; a fistula recurrence, the observed obstructing tracheobronchomalacia or an LC.

TREATMENT
Considering the symptoms of aspiration, the associated aspiration pneumonia, and the uncertainty which diagnosis was causative for presented symptoms, treating physicians decided to perform IL using a hyaluronic acid (HA) agent (figure 1). This is a way to evaluate if the LC is significant in causing the symptoms. The HA agent used in our institution is Juvederm Ultra 3 (Allergan, France). During suspension laryngoscopy, the HA derivative is injected into the interarytenoid region. This

CASE PRESENTATION
A 31+5-week premature boy, with a birth weight of 1275 g, presented with respiratory distress and persistent continuous positive airway pressure (CPAP) dependency since birth. Nasogastric feeding tube insertion failed at 10 cm. Further diagnostics indicated a type C oesophageal atresia with a TEF and a ventricular septal defect. At 5 days of age, thoracoscopic correction of the oesophageal atresia was performed. Per-operative laryngeal inspection revealed a possible mild T1LC, without signs of laryngomalacia. Postoperatively, oxygen dependency persisted, without the need for CPAP Oral feeding was started under supervision of a speech and language therapist, and bottle feeding using only thickened liquids improved gradually. Additional portions could be administered through a nasoduodenal tube when the child (partially) refused bottle feeding. Nonetheless, incidents of aspiration occurred, and the child developed aspiration pneumonia. This clinical presentation and the possibility of an LC gave reason to perform a diagnostic laryngotracheobronchoscopy (dLTB).
is done in order to add bulk and height to the interarytenoid region, and to thus decrease overflow of liquids towards the trachea. The injected volume of the HA agent is not measured. The treating physician injects an amount of filler that gives sufficient bulking, but causes no airway obstruction. Expected is that, as a result of less overflow into the trachea, symptoms of aspiration and symptoms caused by the aspiration, such as recurrent respiratory tract infections, will decrease. After some time, the HA derivative dissolves, and the deeper notch in the interarytenoid region reappears. If the LC was indeed causative, symptoms should subsequently return.

Mean duration of symptom improvement after IL differs between previous studies, partly because of different agents injected. Cohen et al found that the mean duration of improvement of symptoms was 3.3 months. In another study using HA agent in IL for vocal cord medialisation, a median duration of 10.6 months was found.

OUTCOME AND FOLLOW-UP
Initially, treating physicians saw an indication for posterior tracheopexy because of the oxygen dependency, CO2 accumulation and bronchial collapse. Per contra, the patient recovered postoperatively: oxygen therapy was unnecessary, CO2 levels and symptoms caused by the aspiration, such as recurrent respiratory tract infections, will decrease. After some time, the HA derivative dissolves, and the deeper notch in the interarytenoid region reappears. If the LC was indeed causative, symptoms should subsequently return.

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Currently, IL is mostly performed on older children. The mean age at performance of IL was 11.8 months and 29 months in the research of Cohen et al and Ramazani et al respectively. In this case, IL was performed on a child of 8 weeks old and again at 15 weeks. Known complications of IL include stridor, croup-like cough, and respiratory distress as a result of postoperative swelling. According to a previous study, there was no difference in efficacy and safety of IL in children between 0–3 months of age and older children. However, one can presume that a smaller airway lumen is more easily affected by little swelling, according to Poiseuille’s law. Also, a smaller airway lumen can make the procedure more intricate. Volume of filler injected should be assessed by the treating physician during the procedure. This depends on the extent of the cleft, the lumen of the airway treated and the way the filler automatically distributes inside the cleft, which differs per procedure. The physician cannot control the distribution of the filler and it can also create bulking into the airway lumen, causing obstruction and thereby stridor. It is therefore important to be cautious and precise with the amount of filler injected.

In our experience, IL is not often performed in smaller children and literature concerning IL in those cases is sparse. In our case, IL was performed safely in an 8-week-old child, and revised at 15 weeks of age. Diagnosis and intervention at an earlier age is associated with better outcomes.

In this case, the infant did receive a posterior tracheopexy eventually, yet there was a clear indication for IL during the dLTB. As in this patient, children with an LC are often known to have other congenital malformations or diseases. These cloud the diagnostic process, as symptoms could derive from more than one condition, but these comorbidities could also enhance risks associated with surgery and anaesthesia. IL can bridge a few

DISCUSSION
IL is used to determine whether presenting symptoms can be attributed to this small anatomical defect. Therefore, it serves both as a diagnostic tool and a temporary treatment. In this case, there were two other possible causes for these symptoms, each with a different treatment approach. For an LC, standard treatment is surgical repair of the cleft by approximating cleft sides using sutures. According to Chew et al, complications occur in 10%–65% of paediatric airway procedures. The paediatric airway anatomically differs to that of adults, particularly in infants younger than 1 year old. Laryngoscopy is consequently more difficult in this age group and the view is suboptimal. Therefore, there should be a confident indication before surgical intervention, especially in a child of this age, which can be provided by IL.


Figure 1  Interarytenoid injection laryngoplasty. Palpation of the laryngeal cleft using a probe (A), injection of hyaluronic acid filler (B), after injection laryngoplasty (C).
months in which the child’s anatomy can mature, which could possibly lead to both the resolving of symptoms attributed to a newborn’s anatomy, and a more developed infant which can be easier to perform anaesthesia and operate on. This way, IL can prevent unnecessary operative procedures in infants when certain diagnoses are confirmed or disproved. Also, as in this case, IL can be revised multiple times in one child, to extend the period of HA effect if necessary.

Learning points

- Injection laryngoplasty is rarely performed in newborns, and literature regarding safety in these children is sparse. In some cases, however, it can be a good way to relieve symptoms and improve the overall health of a child.
- Injection laryngoplasty has a temporary effect. It can, however, be performed more than once.
- For children with comorbidities in need of surgery, in addition to a laryngeal cleft, injection laryngoplasty at a young age can provide a period of symptom improvement, to bridge the time that is needed for the child to get in better condition for surgery.
- Injection laryngoplasty can prevent unnecessary operative procedures in infants, by confirming or eliminating a laryngeal cleft as the cause of the symptoms.

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Case reports provide a valuable learning resource for the scientific community and can indicate areas of interest for future research. They should not be used in isolation to guide treatment choices or public health policy.

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