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Upper airway obstruction due to acute infection is not uncommon in children, and many parents have experienced an anxious night with a “croupy” child. Although infants and young children are most commonly affected because they have relatively narrow upper airways, older children and adults can also have significant symptoms. Fortunately, these are mostly due to self-limiting viral laryngotracheobronchitis (LTB), but there is also a group of bacterial infections (e.g., epiglottitis, bacterial tracheitis, diphtheria, retropharyngeal abscess, and peritonsillar abscess) that can occasionally cause significant obstruction. It is the job of the emergency physician, pediatrician, pediatric pulmonologist, or otolaryngologist to diagnose more serious infections promptly so that treatment can be instituted early and disastrous obstruction avoided. It is also important to recognize when a simple viral LTB is causing significant problems, so that appropriate treatment can be given immediately. This chapter is clinically oriented and outlines the principal infective causes of upper airway obstruction, with an emphasis on diagnosis and treatment. Confusion exists regarding the nomenclature for these disorders, with some using the term croup to refer to any inflammatory disorder of the upper airway, whereas others restrict its use to subglottic disease (i.e., LTB, which is usually of viral origin). Therefore, for the sake of clarity, the term croup will be largely avoided in this chapter.

The consequence of these upper airway infections is usually stridor, which is a clinical sign and should not be considered a definitive diagnosis. This section briefly outlines the principles behind what causes stridor, which should clarify why this condition mostly affects infants and young children. The appendix in Holinger and colleagues’ Pediatric Laryngology and Bronchoesophagology discusses the physics of air flow and fluid dynamics. The laws of fluid dynamics are based on flow through fixed tubes and may not always apply to dynamic airways in vivo. Normally, air flow through the upper airways is laminar, and the moving column of air produces slight negative pressure on the airway walls. Inflammation resulting from infection causes a degree of airway narrowing, which increases the flow rate through the narrowed segment (the Venturi effect). This, in turn, causes a reduction in the pressure exerted on the airway wall. This is the Bernoulli principle. In other words, negative intraluminal pressure increases. This enhances the tendency of the airway to collapse inward, further narrowing the airway and causing turbulent air flow. The respiratory phase (inspiration or expiration) has a differential effect on air flow, depending on whether the obstruction is intrathoracic or extrathoracic (Fig. 23.1). Stridor is the sound made by rapid, turbulent flow of air through a narrowed segment of a large airway. It is most often loud, with medium or low pitch, and inspiratory. It usually originates from the larynx, upper trachea, or hypopharynx. Progression of the disease process may make stridor softer, higher-pitched, and biphasic (inspiratory and expiratory). With the onset of complete obstruction, stridor may become barely audible as minimal air moves through the critically narrowed airway.

The laryngeal anatomy of children makes them particularly susceptible to narrowing of the upper airways. The larynx of a neonate is situated high in the neck, and the epiglottis is narrow, omega-shaped (ω), and vertically positioned. The narrowest segment of the pediatric airway is the subglottic region (in adults, it is at the glottic level), which is encircled by the rigid cricoid cartilage ring. There is nonlibrinos, loosely attached mucosa in this region that is easily obstructed in the presence of subglottic edema. In addition, the cartilaginous support of the infant airway is soft and compliant, easily allowing dynamic collapse of the airways during inspiration. Young children have proportionally large heads and relatively lax neck support; this combination increases the likelihood of airway obstruction when supine. Also, their tongues are relatively large for the size of the oropharynx. Simple mathematics shows why a small amount of edema has such a profound effect on the cross-sectional area and hence air flow. The diameter of the subglottis in a normal newborn is approximately 5 mm, and 0.5-mm edema in this region reduces the cross-sectional area to 64% of normal (area = πr²). Air flow is directly proportional to the airway radius to the fourth power (Poiseuille’s law), so a small reduction in caliber has a major effect on flow rate. The same 5-mm airway with 0.5-mm edema will have a flow rate of only 41% of baseline, assuming that pressure remains unchanged—a situation that is not necessarily the case if the Bernoulli principle is in play. Because the caliber of the airway is almost inevitably reduced further in accord with the Bernoulli principle, and Poiseuille flow is not established, the flow rate is much further reduced, and the work of breathing is greatly increased to maintain ventilation.

**Viral Laryngotracheobronchitis**

**EPIDEMIOLOGY**

Viral LTB is the most common cause of infective upper airway obstruction in the pediatric age group. Affected children are usually of preschool age, with a peak incidence between 18 and 24 months of age. Although viral LTB episodes become uncommon beyond 6 years of age, cases have been reported during later childhood and adolescence, and rarely described in adults. Reported annual incidence rates in preschool children vary from 1.5% to 6%, but less than 5% of these require...
ABSTRACT
This chapter discusses the common and potentially serious infective causes of acute upper airway obstruction in children. The laryngeal anatomy of young children makes them particularly susceptible to upper airway obstruction, and during acute infections this is exacerbated by inflammation and edema of the airway mucosa. The most common cause of infective upper airway obstruction in children is viral laryngotracheobronchitis, or croup, which is usually a mild and self-limiting illness, but management with corticosteroids may still be necessary. Bacterial causes of upper airway obstruction have fortunately become rare since the introduction of the Haemophilus influenzae B (HiB) immunization, but a few cases of epiglottitis do still occur due to nonimmunization, vaccine failure, and infection with non-HiB organisms. These cases constitute a medical emergency due to the risk of rapid progression to complete airway obstruction. Other rare conditions are discussed including bacterial tracheitis, diphtheria, retropharyngeal, and peritonsillar abscesses. Key learning points in this chapter include the main discriminating factors of the various causes of infective upper airway obstruction, the importance of a calm and minimally distressing approach to the child presenting with stridor, and the need for early anesthetic team involvement in cases with a suspected bacterial etiology or signs of impending airway obstruction.

KEYWORDS
upper respiratory tract infection
upper airway obstruction
croup
viral laryngotracheobronchitis
epiglottitis
bacterial tracheitis
Cases may occur in epidemics, with those caused by parainfluenza virus (PIV) type 1 typically presenting in fall and winter months. Infections caused by other common organisms, including other PIV subtypes, occur more commonly as isolated infections. Infection is via droplet spread or direct inoculation from the hands. Viruses can survive for long periods on dry surfaces, such as clothes and toys, emphasizing the importance of infection-control practices.

**ETIOLOGY**

The most common etiologic agents are the PIVs, of which PIV 1 is found most frequently and leads to epidemics. PIV 2 may account for many sporadic cases, and PIV 3 is a less common cause of viral LTB, usually targeting the epithelium of the smaller airways and leading to bronchiolitic illness. Less is known about the clinical presentation of PIV 4, but it is reported to result in milder clinical illness and is infrequently associated with croup symptoms. The PIVs belong to the *Paramyxoviridae* family, along with respiratory syncytial virus, measles, mumps, and the recently identified human metapneumovirus. Together the PIVs account for more than 75% of viral LTB cases, although other respiratory viruses (e.g., respiratory syncytial virus, rhinovirus, adenovirus, coronavirus, human bocavirus, and enteroviruses) can produce a similar clinical syndrome. Herpes viruses and influenza viruses tend to cause a more severe and protracted form of the disease. LTB can also occur with some systemic infections, such as measles and, less commonly, mycoplasma. In general, however, it is not usually possible to identify the cause of infection from the child’s symptoms because severity does not correlate with any particular etiologic agent.

**PATHOLOGY**

Infection affects the larynx, trachea, and bronchi, although swelling and inflammation in the subglottic area leads to the characteristic clinical features of viral LTB. In addition to relative differences in airway size (discussed earlier in the chapter), it is suggested that poor cell-mediated immunity in younger age groups also accounts for differences observed between adults and children. The epithelium of the subglottis possesses abundant mucous glands, secretions from which can further narrow the airway lumen in response to infection. The PIVs are trophic for the respiratory epithelium, binding in particular to ciliated cells via an interaction between the viral hemagglutinin-neuraminidase protein and its receptor, sialic acid. Other viral proteins (the F protein in particular) are important in membrane fusion and the passage of viral particles between cells. Many strains of PIV are cytopathic, with infection leading to the formation of giant cells and cell death. As with many infective processes, the ensuing inflammatory response is involved in the evolution of symptoms. Both polymorphonuclear and mononuclear leukocytes infiltrate the subepithelium, which leads to vascular congestion and airway wall edema. In addition, the symptoms of viral LTB are believed to be caused by the release of spasmodic mediators, leading to decreased airway diameter. This may result from a type I hypersensitivity response to PIV, and some authors have postulated a role for anti-PIV-specific immunoglobulin E (IgE) in the development of airway narrowing. These factors may play a relatively greater role.
in patients with recurrent (spasmodic) croup, and these patients may have hyperreactivity of the extrathoracic and intrathoracic airways.\textsuperscript{16}

The etiology of recurrent, spasmodic croup remains unclear, with authors expressing differing views on whether it is usually virus-related\textsuperscript{17} or is a separate disease entity; suggested triggers include gastroesophageal reflux, eosinophilic esophagitis,\textsuperscript{18} and anatomical abnormalities, in addition to an allergic predisposition.\textsuperscript{19}

**CLINICAL FEATURES**

**Mild**

Most children are affected mildly by viruses that cause LTB.\textsuperscript{7} The exact incidence remains unknown, because many of them do not receive medical attention, but are managed by parents at home. Children have a barking cough and a hoarse cry or voice; these symptoms are worse in the evening and at night. They may also have inspiratory stridor on exertion, but stridor at rest is usually absent, as are other signs of respiratory distress. There is most commonly a coryzal prodrome accompanied by a low-grade fever, but children are not particularly unwell or toxic. They remain interested in their surroundings, are playful, and still eat and drink.

**Moderate**

Features of moderate viral LTB include those discussed earlier, but with inspiratory stridor present at rest, as well as a degree of respiratory distress manifest by chest wall recession, tachypnea, and the use of accessory muscles of respiration. There is usually accompanying tachycardia, but children remain interactive and are able to take at least liquids orally.

**Severe**

Progression from moderate to severe infection can occur rapidly and may be precipitated by the distress caused by clinical examination. Worrisome signs include those of increasing respiratory distress, with the child appearing anxious or preoccupied and tired. Drooling may occur (but not as commonly as in epiglottitis), and the child will often refuse liquids or be unable to coordinate swallowing and breathing. However, the child with viral LTB will not appear toxic, with high fever and flushed face, as do those with the classic signs of bacterial epiglottitis (Table 23.1). Another difference is in the nature of the cough: a harsh, barking cough is not commonly associated with epiglottitis (in which there is often a muffled cough and cry). Restlessness and agitation are late signs of airway obstruction of any cause, as is cyanosis, pallor, or decreased level of consciousness.

### Table 23.1 Differentiation of Principal Infective Causes of Upper Airway Obstruction

| Viral Laryngotracheobronchitis | Epiglottitis | Bacterial Tracheitis | Diphtheria | Retropharyngeal Abscess |
|-------------------------------|-------------|---------------------|-----------|------------------------|
| **Principal organisms**       | Parainfluenza 1–3 Adenovirus respiratory syncytial virus | Haemophilus influenzae, Moraxella catarrhalis, H. influenzae | Corynebacterium diphtheria | Mixed flora, including S. aureus, Streptococcus, H. influenzae, anaerobes |
| **Age range**                 | 6 months–4 years (peak, 1–2 years) | 2–7 years | 6 months–8 years | All ages |
| **Incidence**                 | Common | Rare | Rare if vaccinated | Uncommon |
| **Onset**                     | Insidious usually follows upper respiratory tract infection | Rapid | Slow, with sudden deterioration | Gradual |
| **Site**                      | Below the vocal cords | Supraglottis | Trachea | Retropharyngeal space |
| **Clinical manifestations**   | Low-grade fever Nontoxic barking (seallike) cough Stridor hoarseness Restlessness | High fever Severe sore throat Minimal nonbarking cough Toxic stridor Drioling Dysphagia Muffled voice | High fever Toxic, brassy cough Stridor Hoarse voice Neck pain Choking | Fever Sore throat Neck pain and stiffness (especially on extension) Dysphagia Stridor (less common) Drioling Retropharyngeal bulge |
| **Endoscopic findings**       | Deep red mucosa Subglottic edema | Deep red mucosa Ulcerations Copious, thick tracheal secretions Subglottic edema, with normal epiglottis and arytenoids | Deep red mucosa Ulcerations Copious, thick tracheal secretions Subglottic edema, with normal epiglottis and arytenoids | Gray, adherent membrane on the pharynx |
| **Intubation Therapy**        | Occasional Corticosteroids Nebulized epinephrine | Usual Intubation (1–3 days) IV antibiotics | Usual Intubation (3–7 days) IV antibiotics Tracheal suction | Occasional Diphtheria antitoxin IV antibiotics Immunization during convalescence |

IV, Intravenous.
Pulse oximetry should be performed, but limitations must be recognized. Oxygen saturation may be well preserved until the late stages of severe viral LTB, and it can lead to significant underestimation of respiratory compromise in a patient who is receiving supplementary oxygen. Conversely, desaturation may be seen in children with relatively mild airway obstruction (presumably reflecting lower airway involvement and ventilation-perfusion mismatch). Pulsus paradoxus is present in the group with severe disease, but in clinical practice, it is difficult to assess, and attempts to do so could worsen symptoms by causing distress.

**Diagnosis and Differential Diagnosis**

The diagnosis of croup is made clinically, based on the features described earlier; there is no role for laboratory tests or radiography in the assessment of acute airways obstruction. In skilled hands, plain lateral neck radiographs may demonstrate sites of obstruction, but this rarely influences management; it also wastes time and can be dangerous. The neck extension that is required could precipitate sudden worsening of airway obstruction, which can be fatal in severe cases. Investigations during the acute illness should be reserved for children with an atypical presentation or who fail to respond to conventional treatment. In such cases, alternative infective (e.g., epiglottitis) and noninfective (e.g., inhaled foreign body) causes of acute airway obstruction require careful exclusion (see Box 23.1 and Table 23.1). Advice from a senior clinician is vital when croup is severe.

**RECURRENT OR SPASMODIC CROUP**

Symptoms are similar to those of the more typical forms of viral LTB, but children are often older, do not have the same coryzal prodrome, and may be afebrile during the episode. There may be links with atopy, often with a positive family history. Episodes are often similar to acute asthma except the child has stridor rather than wheeze. Attacks often occur suddenly, at night, and may resolve equally quickly. Treatment must be guided by the degree of severity, and is similar to that for viral LTB. Some practitioners prescribe oral or inhaled corticosteroids (via a nebulizer) to be kept at home and administered by the parents in case of an episode, although there is a paucity of evidence for or against this practice in spasmodic croup.

**Box 23.1 Infectious and Noninfectious Causes of Acute Upper Airway Obstruction**

| Infectious                        | Noninfectious               |
|-----------------------------------|-----------------------------|
| Viral laryngotracheobronchitis     | Foreign body                |
| Epiglottitis                      | Trauma                      |
| Bacterial tracheitis              | Caustic burns               |
| Diphtheria                        | Spasmodic croup             |
| Retropharyngeal abscess           | Angioneurotic edema         |
| Peritonsillar abscess             | Hypocalcemia (e.g.,         |
| Infectious mononucleosis          | hypoparathyroidism)        |

**NONINFECTIVE CAUSES OF ACUTE AIRWAY OBSTRUCTION**

There are a number of noninfective causes of upper airway obstruction, and these must be considered in the differential diagnosis of infective causes (see Box 23.1). Foreign body inhalation is the most common noninfective cause in children. Symptoms may partly mimic those of viral LTB and will depend on the location of the foreign body, the degree of resultant airway obstruction, and (to a lesser extent) the nature of the foreign body. Onset of symptoms may be either acute or insidious; a large foreign body may cause severe obstruction, whereas a smaller one may simply lead to laryngeal and tracheal irritation and airway edema. In cases of severe airway obstruction, the voice may be lost and breath sounds quiet. This condition is an emergency, and requires immediate visualization of the larynx and trachea and removal of the foreign body by a physician or surgeon experienced in this procedure. Occasionally an unrecognized inhaled foreign body leads to chronic stridor. Acute upper airway obstruction may also result from the ingestion of caustic substances, with resulting pharyngeal burns, edema, and inflammation of the epiglottis, aryepiglottic folds, larynx, and trachea. This diagnosis is usually clear from the history. Rarely, angioneurotic edema may cause acute laryngeal swelling and airway obstruction. Patients appear nontoxic and may exhibit other signs of allergic disease, such as urticaria and abdominal pain. In hereditary angioneurotic edema due to C1 esterase inhibitor deficiency, the family history may be positive, although the first presentation is more common in adults than in children. Hypocalcemia, for example, due to hypoparathyroidism, can lead to laryngospasm (a form of tetany), which can cause stridor with significant airway obstruction.

There are numerous causes of chronic airway obstruction that are discussed elsewhere in this book. Confusion may arise when an upper respiratory tract infection unmasks a previously asymptomatic congenital abnormality. For example, mild subglottic stenosis may cause symptoms only with the additional burden of airway edema due to a simple viral upper respiratory infection. It is important to ensure that there is no history of intubation (which may have been brief, as in resuscitation of a newborn in the maternity unit) or of any coexisting signs (e.g., a cutaneous hemangioma) that may increase the index of suspicion for a congenital airway abnormality. One catch with subglottic hemangioma is that they may initially respond to systemic steroids, and thus their diagnosis may be masked with the assumption that stridor was due to viral croup.

**MANAGEMENT OF VIRAL LARYNGOTRACHEOBRONCHITIS**

Management of viral LTB must be based on clinical assessment of severity. Several scoring systems have been devised, and the most commonly applied system (the 17-point Westley scale, which assesses degree of stridor, chest retractions, air entry, cyanosis, and level of consciousness) has been well validated. However, these are mainly used in the context of clinical trials and are not a substitute for experienced clinical assessment.
Supportive Care

Children with mild croup can be managed at home. They should be treated with plenty of fluids and antipyretics as required. Because the vast majority of cases are of viral etiology, there is no role for the routine use of antibiotics in the absence of other features suggestive of bacterial infection. Parents should be warned that symptoms are usually worse at night and may recur after apparently disappearing during the day.

Humidification

Both at home and in the hospital setting, humidified air (either steam or cool mist) has been used for more than a century to produce symptomatic relief from croup. Despite this, there is very little supportive evidence; most early studies, some of which may have been underpowered, generally suggested no benefit.24–26 A larger study of 140 moderately affected children showed no differences in signs or requirement for additional treatments with optimally delivered 100% humidity,27 and the most recent Cochrane Systematic Review has also concluded there is no evidence of benefit.28 Case reports have described severe burns caused by spilling of boiling water and facial scalds from the use of steam, so this type of treatment is not without the potential for harm.29

Corticosteroids

The use of corticosteroids has received much attention for more than a decade, and their therapeutic role is well established. Their mechanism of action, however, remains unclear, although is believed to relate to rapid-onset antiinflammatory properties. The cumulative evidence strongly supports their use in children with moderate to severe symptoms, although there are still outstanding questions, including the optimal route of administration, the most appropriate dosing regimen, and the best oral agent.

The role of corticosteroids in the management of croup in children has been the subject of several Cochrane reviews, with the most recent update in January 2011.30 In this review, the authors identified 38 studies that fulfilled their criteria for inclusion—namely, randomized controlled trials in children measuring the effectiveness of corticosteroids (any route of administration) against either a placebo or another treatment. A total of 4299 children were included, the majority from the emergency department or the hospital was also significantly decreased, as was the requirement for nebulized epinephrine. Importantly, and in contrast to the first version of this Cochrane review, the authors concluded with funnel plots and other statistical methods that these results were not influenced by publication bias. In conclusion, there is convincing evidence that corticosteroids provide effective and sustained treatment of croup symptoms, leading to clinical improvement within 6 hours of administration. In severe disease, rates of intubation are significantly decreased and the duration of intubation is reduced, and in moderate disease admission, the need for additional treatment and return visits are reduced.8 More recent studies have focused attention on the optimal formulation, dose, and treatment regimen.

Optimal Route of Administration, Formulation, and Dosing Regimen

Studies included in the Cochrane review (discussed earlier in the chapter) and others conducted since then have used the intramuscular, oral, or nebulized route to administer different corticosteroid preparations. This area has been well reviewed recently.9 From the studies that have attempted to address the route of administration, nebulized, oral, and intramuscular routes appear, in general, to be roughly equivalent. Nebulization could potentially increase distress of the child and worsen upper airway obstruction, although it may be preferable in a child who is vomiting or having difficulty swallowing.

Similarly, studies using oral agents have used either dexamethasone or prednisolone, and both in varying doses. Many primary care physicians who visit homes do not routinely carry dexamethasone but do carry oral prednisolone. There is no strong evidence in support of one preparation over the other, although one recent study favored dexamethasone, which led to a reduced frequency of re-presentation.31 In contrast, a recent Australian trial compared 1 mg/kg prednisolone with a single dose of oral dexamethasone using two different dosing regimens (0.15 and 0.6 mg/kg) and found no difference in croup score, requirement for further treatment, or re-presentation.32 Similarly, a community-based double-blind trial comparing a single dose of dexamethasone 0.6 mg/kg with three doses of 2 mg/kg prednisolone on consecutive days found no difference in additional health care attendances, duration of symptoms, parental stress, or sleep disturbance.33 With regard to dexamethasone, 0.6 mg/kg has been the dose most widely used, but several studies have demonstrated that this dose may be higher than required and that 0.15 mg/kg is just as effective.34–36 A practical approach might be to use dexamethasone, if available, at a dose of 0.15 mg/kg. If this preparation were not available at a home visit, prednisolone (at an equivalent dose of 1 mg/kg) could provide a useful substitute. Due to the short duration of symptoms in a typically episode of croup, one dose of steroid is usually sufficient treatment. However, a second dose should be considered if residual symptoms are still present the following day, and can be given to the parents to administer at their discretion.

Nebulized Epinephrine (Adrenaline). The most recent update of the Cochrane Review of nebulized epinephrine use in croup was published in 2013.37 It included eight randomized controlled studies, with a total of 225 subjects comparing the efficacy of nebulized epinephrine to placebo.
(six studies), or comparing different epinephrine formulations and delivery methods (two studies).

All studies involved the management of moderate to severe croup in either an emergency department or hospital setting. Outcome measures used were croup score, intubation rates, side effect profile, and total health care utilization. Nebulized epinephrine was associated with improved clinical severity score 30 minutes posttreatment, but this effect was not sustained at 2 or 6 hours posttreatment. Magnitude of benefit was consistent between studies, and similar, irrespective of whether epinephrine was administered in an inpatient or outpatient setting.

Duration of hospitalization was also significantly reduced with epinephrine (mean difference of 32 hours), although this outcome was only assessed in one study. No studies provided a comparison of intubation rate or side effect profile. In summary, nebulized epinephrine has been shown to improve clinical severity in the short term and reduce hospital admission in cases of moderate to severe croup. It should be used in any child who has severe signs and symptoms, and it should be considered for those with moderate signs and symptoms, depending on the signs of respiratory distress and possible response to corticosteroid administration. It can be administered in the home setting while awaiting an ambulance, but clearly any child requiring this treatment at home must be transferred promptly to the hospital for monitoring. Multiple doses may be administered, although the requirement for this must lead to consideration of the need for intensive care management. Although rebound worsening of symptoms after administration of nebulized epinephrine is often alluded to, in practice, this phenomenon does not appear to be a real risk. Traditionally, children treated with epinephrine have been admitted to the hospital, but recent studies have confirmed that discharge home is safe after 3 to 4 hours of observation if the child has made significant improvement.38

Most clinical trials have used the racemic form of this drug,39 although there is evidence that the L-isomer used alone (which is the only available formulation in some units) may be equally effective and has a longer duration of action.40 The mechanisms of action are believed to be a combination of rapid reduction in airway wall edema and bronchodilation. The recommended dose is 0.4 to 0.5 mL/kg (to a maximum of 5 mL) of the 1:1000 preparation that is put undiluted into the nebulizer cup. Intramuscular epinephrine is not used in severe stridor, so it is important to ensure the stridor is not due to acute anaphylaxis, in which case it should be given.

**Other Treatments for Severe Cases.** Oxygen should be administered to any child with severe airway obstruction, even in the absence of severe hypoxia, because it will aid respiratory muscle function. As mentioned earlier, a child with severe respiratory distress and obstruction may have normal pulse oximetry readings when breathing oxygen, which can be dangerous if misinterpreted by staff who are unaware of this limitation. Heliox (70% to 80% helium with 20% to 30% oxygen) has been used in both upper airway obstruction41 and severe asthma, and it is the focus of a recent Cochrane review.42

Three randomized controlled trials were identified totaling 91 participants with croup of varying severity. Heliox was compared with either 30% humidified oxygen, 100% oxygen plus epinephrine, or against no treatment. No pooled analysis was possible, as each study used a different comparator. Collectively, these studies provided evidence of a short-term benefit of Heliox inhalation, with improved croup scores at 60 to 90 minutes following administration. However, this clinical improvement was not sustained over subsequent hours. The authors summarized that no additional benefit appeared to have been provided by Heliox beyond that delivered by administration of 30% oxygen in cases of mild croup, or 100% oxygen plus nebulized epinephrine in moderate to severe croup. Adequately powered studies are required to further evaluate the role of Heliox in the management of moderate to severe croup.

**Endotracheal Intubation**

Some children with severe croup either do not respond to the usual therapies or are too severely compromised at presentation to permit their use. These children require urgent endotracheal intubation and mechanical ventilation to avoid potentially catastrophic complete airway obstruction and the serious sequelae of hypoxia and hypercapnia (e.g., hypoxic ischemic encephalopathy). Intubation should be performed by the most experienced person available, and it should be attempted with an uncuffed endotracheal tube one size smaller than the usual size for the child.43 Facilities for immediate tracheostomy must be available at the time of intubation. Children may have coexisting lower airway and parenchymal involvement that impairs gas exchange and may lead to slower than expected clinical improvement after intubation. Rarely, pulmonary edema may develop after relief of airway obstruction, particularly if the disease course has been prolonged. Most children without severe parenchymal involvement require respiratory support for 3 to 5 days.43 This is one context in which multiple rather than single doses of corticosteroids are often administered. The timing of extubation will depend on the development of an air leak around the endotracheal tube, indicating resolution of airway narrowing.44 Reintubation rates of approximately 10% have been reported.44

**PREVENTION**

Because of the substantial health care costs associated with management of PIV infections (estimated annual costs of $43, $58, and $158 million for hospitalizations in the United States for PIV-associated bronchiolitis, croup, and pneumonia, respectively), there is much interest in development of an effective PIV vaccine.45 The most promising candidate vaccine developed to date is an intranasally administered cold-adapted PIV 3 vaccine that appears to be well-tolerated and immunogenic in infants as young as 1 month of age.46 Also under evaluation is a PIV3-vectored ing candidate vaccine developed to date is an intranasally administered PIV 3 vaccine that appears to be well-tolerated and immunogenic in infants as young as 1 month of age.46 Also under evaluation is a PIV3-vectored

**PROGNOSIS AND FURTHER EVALUATION**

Most children with viral LTB were previously well, have short, self-limiting symptoms, and make a full recovery. The lack
of complete immunity and the variety of agents that can cause viral LTB mean that more than one episode is not uncommon, particularly in separate seasons.

Recurrent (≥3) episodes have been reported in more than 60% of affected children, with family history of croup identified as the most significant risk factor for recurrence in one case-control study. Although further evaluation is not necessary in every case of recurrent croup, it should be considered in cases that are particularly severe or frequent, if symptoms are particularly slow to resolve, or if symptoms occur between or in the absence of obvious infections. Evaluation of patients in this group is aimed at identifying an underlying airway abnormality that would predispose the child to more severe airway narrowing with viral infections, or that could cause problems independently of such infection. Investigation is usually centered on airway endoscopy. This must be performed in a unit and by an operator who is experienced in the technique because there is a risk of exacerbating the airway obstruction. Spontaneous breathing is necessary to identify vocal cord problems or airway malacia, and anesthetic techniques must be carefully considered. If an inhaled foreign body is considered likely, rigid bronchoscopy is the study of choice. Additional studies that might be considered once the acute episode has resolved include plain lateral neck and chest radiographs, computed tomography or magnetic resonance imaging, contrast assessment of the upper airway (e.g., videofluoroscopy, barium swallow), and a pH probe study. Polysomnography may help determine the severity of chronic symptoms. Rarer causes of recurrent stridor (e.g., hypocalcaemia or angioneurotic edema) are diagnosed by blood testing.

**Epiglottitis**

**EPIDEMIOLOGY**

*Haemophilus influenzae* type B (HiB) vaccines were first licensed in the United States in 1988, with widespread immunization programs in place by the early 1990s. Since then, reported cases of invasive HiB disease (including epiglottitis) in children younger than 5 years of age have declined by 99%. By 2008, the overall incidence in the United States had fallen to 1.55 per 100,000, having been 4.39 per 100,000 in 1989. The same pattern has been repeated in Europe, with significant reductions in the United Kingdom. Immunization was introduced in the United Kingdom in 1992, and with immunization coverage exceeding 90%, the decline in incidence was more than 95%. In 1998, the incidence in those younger than 5 years of age was 0.6 per 100,000, compared with 31 to 36 per 100,000 in England and Wales before the introduction of the vaccine. However, in 2003 there was a resurgence of HiB infections in the United Kingdom, which led to the launch of a booster program. By 2012, in England and Wales, the incidence had fallen further to 0.06 per 100,000 aged less than 5 years (0.02 per 100,000 for all ages). Ethnicity plays a part in vaccine efficacy. Data from 1996 to 1997 in the United States show that the average annual incidence of HiB invasive disease per 100,000 children younger than 5 years of age was 0.5 among non-Hispanic whites, 0.6 among Asians and Pacific Islanders, 0.7 among non-Hispanic blacks, 0.7 among Hispanic Americans, and 12.4 among Native Americans and Alaskan natives. Nevertheless, cases of epiglottitis due to HiB continue to be reported, as do cases due to other organisms.

Cases of invasive HiB occur principally in nonimmunized children, but also rarely in some true vaccine failures. Clinical risk factors for vaccine failure include prematurity, Down syndrome, malignancy, developmental delay, and congenital or acquired immunodeficiency, principally with reduced immunoglobulin concentrations (IgG2 subclass, IgA, IgM) or neutropenia. However, these factors explain fewer than 50% of cases of vaccine failures. An HiB IgG antibody titer of ≥0.15 µg/mL confers protection from disease, but given the natural waning in antibody levels, it is estimated that a titer of ≥1.0 µg/mL should provide long-term protection. However, sometimes there are qualitative functional problems with antibody responses that are not yet fully elucidated.

Among the pediatric cohort, epiglottitis tends to occur in children 2 to 7 years of age, but cases have been reported in those younger than 1 year of age. Since the introduction of HiB vaccine, the peak age distribution has increased slightly. A review of a national US dataset from 1998 to 2006 has shown that the mean age of a patient admitted with epiglottitis is 45 years, and the national mortality rate is 0.89%; there is a decrease in admissions for those under 18 years of age with greatest risk at <1 year and an increase in the 46- to 64-year-old group. This is similar in England and Wales, where 68% of culture-positive cases of epiglottitis occurred in patients aged 45 or over. Most cases occur in the fall and winter.

**ETIOLOGY**

Historically, HiB was responsible for almost all (approximately 99%) cases of epiglottitis in otherwise healthy children. Since the introduction of HiB immunization, other organisms have been implicated, including groups A, B, C, and G β-hemolytic *Streptococcus*. Other responsible organisms include *H. parainfluenzae*, *Staphylococcus aureus*, *Moraxella catarrhalis*, *Streptococcus pneumoniae*, Klebsiella, Pseudomonas, Candida, and viruses (e.g., herpes simplex, varicella, PIH, influenza, and Epstein-Barr virus).

**PATHOLOGY**

Although HiB has a low point-prevalence of nasopharyngeal carriage (1% to 5%), most young children become colonized with HiB in the first 2 to 5 years. The relationship between asymptomatic carriage, immunity, and the development of invasive disease is not clearly understood. Viral coinfection may have a role in the transition from colonization to invasion. Colonies of HiB organisms reside in the nasal mucosal epithelium and submucosa. Invasive disease occurs when organisms disseminate from the mucosa of the upper respiratory tract via the bloodstream; bacteremia increases over a period of hours, and metastatic seeding can occur. Thus, although situated in close proximity to the nose, the supraglottic area is likely to be affected via the bloodstream; direct spread along mucosal surfaces may also play a part. This may account for the relatively high yield of positive blood cultures in epiglottitis and the relatively low incidence of epiglottitis among carriers of HiB.
Acute Infections That Produce Upper Airway Obstruction

However, a 13-year case series demonstrated that cardiac arrest occurred in 3 of 40 cases (7.5%), although there were no long-term sequelae. Secondary sites of HiB infection may be present in approximately half of cases, and include meningitis, otitis media, pneumonia, and cellulitis; descending necrotizing mediastinitis has been reported. Therefore repeated physical examination during the admission is critical. The pneumonia may contribute to poor gas exchange.

The presentation and clinical course of epiglottitis caused by the various types of β-hemolytic streptococcal pathogens are similar to each other, but they differ from those associated with HiB. The onset of disease is more gradual, but the resolution of tissue damage and the time to recovery are longer, with a mean intubation time of 6 days.

**MANAGEMENT**

The first priority and key response to the diagnosis must be to secure the airway in a controlled environment. Physical

**CLINICAL FEATURES**

Classic epiglottitis caused by HiB is a fulminant disease in an otherwise healthy child, who can be near death in a few hours. It is a medical emergency that can be alarming for the medical staff and devastating for the family. Epiglottitis clearly has not been eliminated, but due to its rarity there are concerns about a potential lack of familiarity with its management among emergency physicians, pediatricians, anesthesiologists, and otolaryngologists. Up to 20% of infants with epiglottitis are misdiagnosed initially, usually with viral LTB. Typically there is a short history of fever, severe throat pain, stridor, and respiratory distress, but the symptoms progress rapidly. Children become toxic and tend to sit anxiously in the classic tripod position (sitting upright, with the chin up, mouth open, bracing themselves on their hands) as air hunger develops. They often drool because they cannot swallow their secretions, and the voice is muffled due to pain and soft tissue swelling. Stridor may progress, and when marked (or if it disappears completely), signals almost complete obstruction of the airways. Complete, fatal airway obstruction may occur suddenly and without warning. The most serious complication of this disease process (and any infective upper airway obstruction) is hypoxic ischemic encephalopathy resulting from respiratory arrest. This tragic complication is almost always preventable with clinical suspicion, prompt diagnosis, and correct management.

However, a 13-year case series demonstrated that cardiac arrest occurred in 3 of 40 cases (7.5%), although there were no long-term sequelae. Secondary sites of HiB infection may be present in approximately half of cases, and include meningitis, otitis media, pneumonia, and cellulitis; descending necrotizing mediastinitis has been reported. Therefore repeated physical examination during the admission is critical. The pneumonia may contribute to poor gas exchange.

In general, distinction from standard viral LTB is based on the older age of the child, the lack of history of upper respiratory tract infection, the speed of progression, the degree of toxicity, the extent of drooling, the use of the tripod position, and minimal cough (see Table 23.1). However, it is important to remember that most of these symptoms can be present in acute severe upper airway obstruction from other causes.

The presentation and clinical course of epiglottitis caused by the various types of β-hemolytic streptococcal pathogens are similar to each other, but they differ from those associated with HiB. The onset of disease is more gradual, but the resolution of tissue damage and the time to recovery are longer, with a mean intubation time of 6 days.

**MANAGEMENT**

The first priority and key response to the diagnosis must be to secure the airway in a controlled environment. Physical
of intravenous ceftriaxone was as efficacious as 5 days of have traditionally been given for 7 to 10 days; however, a changed once antibiotic sensitivities are available. Antibiotics ceftriaxone or cefotaxime) is usually given and may be prepared to perform this immediately if intubation is unsuc-

tomy tray. If complete airway obstruction develops suddenly, performance of a Heimlich maneuver may relieve the obstruction temporarily; alternatively, forward traction may be applied to the mandible.

Inhalational induction of anesthesia is preferred. Laryn-
goscopy should then be performed and the diagnosis con-
firmed, based on the appearance of the epiglottic region, as described earlier in the chapter (erythema and edema of the supraglottis). Endotracheal intubation is then achieved using an orotracheal tube, which is later changed to a nasotracheal tube, because this is less likely to be displaced leading to a potentially disastrous extubation. Intubation may be done using the Seldinger technique over a bronchoscope. Although tracheostomy is rarely necessary, a surgical team should be prepared to perform this immediately if intubation is unsuccess-

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Some authors have advocated the use of a lateral neck radiograph if the child is stable before intubation, claiming it to be the “single most useful study.” We strongly disagree, and this is not our recommendation. Because it can precip-

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CLINICAL FEATURES

The clinical picture is initially similar to that of viral LTB, with mild fever, cough, and stridor for several days. However, the patient’s condition deteriorates rapidly, with a high fever and often a toxic appearance, with respiratory distress and airway obstruction. Other symptoms include choking episodes, orthopnea, dysphagia, and neck pain. The clinical picture differs from that of epiglottitis in that its onset tends to be more insidious. Patients have a substantial brassy cough, are more able to lie flat, and tend not to drool (see Table 23.1). Children are more ill than with simple viral LTB and do not respond to expected therapies (e.g., corticosteroids or nebulized epinephrine). There may be other coinfections, particularly pneumonia. Other reported complications include cardiopulmonary arrest, with subsequent hypoxic encephalopathy and seizures, pneumothorax, subglottic stenosis, septicemia, toxic shock syndrome, pulmonary edema, and adult respiratory distress syndrome.

The white blood cell count shows polymorphonuclear leukocytosis, often with a left shift. A lateral neck radiograph may show a hazy tracheal air column, with multiple luminal soft tissue irregularities due to pseudomembrane detachment from the soft tissue, but radiographs should be taken only after the patient is stabilized and safe. There are, however, no clinical or radiographic features capable of confirming the diagnosis. This must be confirmed by upper airway endoscopy and a positive bacterial culture.

MANAGEMENT

Diagnostic rigid endoscopy, which should be done under general anesthesia, is also therapeutic because it enables removal of secretions and sloughed tissue from the airway lumen; sometimes the procedure must be repeated. Many patients (especially younger ones) require endotracheal intubation and mechanical ventilation to overcome airway obstruction (reports of 50% to 100% intubation rates), usually for 3 to 7 days. Frequent tracheal suction is necessary. In a 1998 case series, 57% of patients required intubation, which is lower than the rate previously reported. The decision to extubate is based on clinical improvement, with reduction of fever, decreased airway secretions, and development of an air leak around the endotracheal tube. Corticosteroids may be given before extubation. Tracheostomy lumen; sometimes the procedure must be repeated. Many

ETIOLOGY

Diphtheria is caused by toxigenic strains of the bacterium Corynebacterium diphtheriae and, less frequently, C. ulcerans. The organism may be isolated on bacterial culture of nasal and pharyngeal swabs, and serologic studies may detect antibodies to diphtheria toxin. Polymerase chain reaction can confirm C. diphtheriae tox genes. There has been a case reported where nontoxigenic Corynebacterium diphtheriae caused necrotizing epiglottitis in an immunocompromised child.

PATHOLOGY

Diphtheria is an acute disease, primarily involving the tonsils, pharynx, larynx, nose, skin, and occasionally other mucous membranes. In the milder, catarrhal form, there is no membrane formation, but in the more severe, membranous form, there is a characteristic lesion of one or more patches of an adherent grayish-white membrane, surrounded by inflammation. The toxin causes local tissue destruction at the site of membrane formation, which promotes multiplication and transmission of the bacteria.

CLINICAL FEATURES

The incubation period is 1 to 6 days. Classic respiratory diphtheria is characterized by an insidious onset, and patients typically present with a 3- to 4-day history of upper respiratory infection. They have a fever, membranous pharyngitis with a sore throat, characteristic fetor oris, cervical lymphadenopathy, and sometimes edema of the surrounding soft
vascular coagulation and renal failure). Myocarditis/cardiomyopathy, or sepsis (disseminated intra-
break. Deaths are usually a result of airway obstruction, myocarditis/cardiomyopathy, neuritis or paralysis, and adrenal failure.

**ETIOLOGY**

Retropharyngeal abscesses generally result from lymphatic spread of infection, although direct spread from adjacent areas, penetrating pharyngeal trauma (e.g., a fall with a pencil in the mouth), or foreign bodies can also play a role. It has also been reported as a complication of adenoidectomy and adenotonsillectomy. Infection is usually due to mixed flora, including *Staphylococcus aureus* (methicillin-sensitive and resistant), various streptococcal species (in particular group A beta-hemolytic *Streptococcus*), *Hib*, and anaerobes. A case has been reported due to tuberculosis (cervical Pott’s disease).

**PATHOLOGY**

The retropharyngeal space (between the posterior pharyngeal wall and the prevertebral layer of deep cervical fascia) contains tissues (bull-neck appearance). They may have a serosanguineous nasal discharge. Although not always present, the membrane is typically gray, thick, fibrinous, and firmly adherent, so it may bleed on attempted removal (Fig. 23.4). Laryngeal diphtheria most commonly occurs as an extension of pharyngeal involvement in children, leading to increasing hoarseness and stridor. The patient appears toxic, with symptoms of LTB, and signs of severe airway obstruction may develop quickly if the pharyngeal membrane dislodges and obstructs the airway. Complications include secondary pneumonia and toxin-mediated disease including myocarditis or cardiomyopathy, neuritis or paralysis, and adrenal failure with hypotension. Cardiomyopathy can be predicted from a combination of a pseudomembrane score greater than 2 (range is up to 4) and a bull-neck appearance. Case fatality rates vary and can reach 20%, but the mortality rate was only 3% in a series of 676 patients (30% younger than 15 years of age) reported from the 1995 Kyrgyz Republic outbreak. Deaths are usually a result of airway obstruction, myocarditis/cardiomyopathy, or sepsis (disseminated intravascular coagulation and renal failure).

**MANAGEMENT**

Because diphtheria is now rare, a high index of suspicion is important to make the diagnosis and institute prompt treatment. Patients should be strictly isolated. Diphtheria antitoxin, which is a hyperimmune equine antiserum (available from the Centers for Disease Control and Prevention), should be administered without waiting for laboratory confirmation because it neutralizes circulating toxin, but not toxin bound to tissues. Antibiotics are not a substitute for antitoxin, but they are given to eradicate the organism, stop toxin production, and reduce the likelihood of transmission. Intravenous penicillin or erythromycin is used, and once the child can swallow comfortably, treatment can be given orally, for a total of 14 days. Mechanical ventilation and tracheostomy may be required. Intravenous dexamethasone has been given to children with laryngeal diphtheria and airway obstruction, and a small case series suggested that it was beneficial. Because the disease may not confer immunity, patients should be given a diphtheria toxoid-containing vaccine during convalescence. Antibiotic prophylaxis (penicillin or erythromycin) is recommended for close contacts after nasal and pharyngeal specimens are taken, and immunization should be given to those who have not been vaccinated in the preceding 5 years.
contains loose connective tissue and lymph nodes that drain the nasopharynx, paranasal sinuses, middle ear, teeth, and adjacent bones. The space extends from the base of the skull down to vertebra C7 or T1. Acute bacterial infection in this region may start as retropharyngeal cellulitis with localized edema of the tissues, which can progress to purulent inflammation of the tissues and retropharyngeal adenitis. However, when this process is caused by lymphatic spread, it starts as adenitis. If liquefaction of one of the nodes occurs, an abscess can form and is usually contained within the inflammatory rind of the infected node.

CLINICAL FEATURES

Presentation is often nonspecific, and there may be overlap with the presentation of croup, epiglottitis, tracheitis, and peritonsillar abscess (see Table 23.1). Children with acute epiglottitis tend to appear more toxic and progress to respiratory distress more rapidly. Excluding causes secondary to foreign bodies or trauma, patients usually have a history of a viral upper respiratory infection that lasts several days and then worsens. Children then have high fever, sore throat, dysphagia, poor feeding, neck pain, and stiffness. Limitation of neck extension and torticollis are more common than limited neck flexion. Although the occurrence of neck signs with fever may suggest meningitis, affected children tend not to be as toxic as those with meningitis. Further deterioration may lead to extrathoracic airway compromise, with drooling, stridor, and respiratory distress. The classical picture of stridor and airway obstruction seems to be less common now, and in a series of cases reported by pediatricians from 1993 to 1998, only 3% of 64 children in Salt Lake City had stridor or wheezing. However, in a series of cases from 2002 to 2007 from the same center, but published by the otolaryngologists, 14/130 (11%) presented with airway obstruction, and 50% required intubation, with an average age of 1.4 years. Older series have reported a higher incidence of stridor. Reported rates include 71% in patients younger than 1 year of age, 43% in patients older than 1 year of age, and no stridor in those older than 3 years of age in a series of 31 children in Sydney, Australia; a rate of 56% in 17 patients in Denver; and a rate of 23% in 65 children in Los Angeles. It is possible that the spectrum of disease is changing, but it is more likely that the diagnosis is being made earlier, before the airways are compromised. Sometimes a retropharyngeal mass is visible in the mouth, seen as an asymmetrical bulge of the posterior pharyngeal wall (Fig. 23.5), or a neck mass (marked lymphadenopathy or parapharyngeal abscess) is visible and palpable. Complications include rupture of the abscess with aspiration, asphyxiation or pneumonia, extension to a mediastinal abscess with or without mediastinitis, and Lemierre syndrome with vascular complications (e.g., anaerobic septic thrombophlebitis of the internal jugular vein with multiple septic pulmonary emboli, and potential erosion through the carotid artery sheath).

Once the child is stable and safe, a lateral neck radiograph with the neck in full extension may confirm the diagnosis. Widened prevertebral soft-tissue shadow and air-fluid levels in the retropharyngeal space are all indicators. The radiograph must be taken in the correct position (true lateral orientation, with the neck in extension, and if possible, during full inspiration) to ensure that the retropharyngeal space is not falsely thickened. A contrast-enhanced computed tomography scan is useful because it differentiates a fully developed abscess from cellulitis and delineates the full extent of the abscess. Blood cultures findings are usually negative, but the white blood cell count will be increased.

MANAGEMENT

Traditionally, management involves surgical drainage of the abscess (plus antibiotics). However, more cases are being managed by intravenous antibiotics alone, although surgery must be considered early if there is a compromised airway. Percutaneous CT-guided aspiration has also been described. In one older series, 25% of patients required no surgery, but in the Salt Lake City series that covered 1993–1998, 58% of 64 patients had antibiotics alone, with no treatment failures. Data from KID 2003 (discussed earlier in the chapter) showed that 43% of cases admitted required surgical drainage. Update of the KID database shows that surgical management has continued to fall to 38% in 2009. Many children start treatment with antibiotics before the diagnosis is made, and certainly retropharyngeal cellulitis can be treated with antibiotics alone. When necessary, surgical drainage is performed through the intraoral route. Care must be taken to avoid aspiration of the infected material. Occasionally, if there is extension lateral to the great vessels, external drainage through the neck is necessary. Mortality is rare now, with no deaths in recent reports.

Peritonsillar Abscess (Quinsy)

EPIDEMIOLOGY

Peritonsillar abscess is the most common deep-space head and neck infection in both adults and children. However, it is more common in young adults than in children. It tends to affect older children and adolescents, and in one large,
10-year series, the mean age was 12 years of age, with two-thirds of those affected older than 10 years of age. In this series, 62% of cases occurred on the left side versus 38% on the right (with no obvious explanation). There is no seasonal predilection. The reduction of antibiotic prescribing to children by general practitioners in the United Kingdom from 1993 to 2003 has not been accompanied by an increase in hospital admissions for peritonsillar abscess. The KID database in the United States found no significant change in incidence of peritonsillar abscess between 2000 and 2009 (0.82 to 0.94 per 10,000 children). In England, there was a 31% rise in peritonsillar abscess admission rate for children and adults between 1991 and 2011, that was thought to be associated with the 41% fall in overall tonsillectomy rate in that same period.

ETIOLOGY

A peritonsillar abscess is usually a complication of acute tonsillitis, but it may follow pharyngitis or a previous peritonsillar abscess. The infection usually involves mixed bacterial flora, with Streptococcus pyogenes as the predominant organism. In a small series of children younger than 5 years of age, Streptococcus viridans was the most common organism detected. In a large series of 457 children and adults in Israel, while S. pyogenes was the most common organism isolated, there was a sharp rise in anaerobes cultured, particularly Prevotella and Peptostreptococcus.

PATHOLOGY

Peritonsillar abscess is believed to arise from the spread of infection from the tonsil or the mucous glands of Weber, located in the superior tonsillar pole. There is a spectrum of peritonsillar cellulitis that may then result in a collection of pus located between the tonsillar capsule (pharyngobasilar fascia), the superior constrictor, and the palatopharyngeus muscle. There is a risk of spread through the muscle into the parapharyngeal space or other deep neck spaces. The abscess pushes the adjacent tonsil downward and medially, and the uvula may be so edematous, as to resemble a white grape.

CLINICAL FEATURES

The child who is already affected by acute tonsillitis becomes more ill with a high fever and has a severe sore throat or neck pain, as well as marked dysphagia with referred earache. Absent or decreased oral intake can lead to dehydration, particularly in younger children.

Cervical lymphadenopathy is almost always present. The uvula is edematous and deviated to one side, and there is fetor oris. A striking feature may be trismus, with limited mouth opening. Examination may be difficult in a young, uncooperative child who refuses to open the mouth. The white blood cell count will be elevated.

The relevance of this condition to pediatric pulmonologists is that acute enlargement of the tonsils can cause airway compromise, and a ruptured abscess can lead to aspiration of infected material and subsequent pneumonia. In one large series of 169 children under 18 years of age, 8% presented with airway compromise.

MANAGEMENT

Children may need intravenous fluids. Antibiotics are necessary, and intravenous penicillin is as effective as broad-spectrum antibiotics, although additional anaerobic coverage should be considered. Analgesia is important. Corticosteroids are not uncommonly used but are of no obvious benefit (or harm). Treatment often involves needle aspiration or incision and drainage; a meta-analysis of 10 studies with 496 patients revealed an average 94% success rate with simple needle aspiration. The KID database revealed that about a third of cases required incision and drainage in 2009, with an increase over the decade (26% to 34%). Nevertheless, an initial period of medical treatment is appropriate in the absence of airway compromise and systemic toxicity, especially in younger children. One case series of 88 children from Korea suggested that younger age (<7.5 years), fewer episodes of acute tonsillitis, and smaller abscess size predicted successful nonsurgical treatment. There is some debate among otolaryngologists about the role and timing of tonsillectomy, and peritonsillar abscess is no longer considered an absolute indication for tonsillectomy, although a history of recurrent tonsillitis prior to developing the peritonsillar abscess leads to a higher recurrence rate. In the United States, the KID database found a significant reduction in tonsillectomy carried out between 2000 and 2009 (13% to 8% cases). In a large nationwide study from Taiwan of almost 29,000 cases of peritonsillar abscess, the recurrence rate was 5% and was associated with the incidence of prior episodes of tonsillitis in all ages, and management by needle aspiration (vs. incision and drainage) in children.

Infectious Mononucleosis

Infectious mononucleosis is caused by Epstein-Barr virus (EBV) and is common in adolescents and young adults. The clinical syndrome is characterized by fever, fatigue, malaise, lymphadenopathy, and sore throat. Diagnosis is confirmed by positive EBV serology, and positive heterophile antibodies (Monospot), although the latter are only positive in 50% of children and 90% of adults. The illness is usually self-limiting, but malaise and exhaustion can persist.

Some degree of airway obstruction is not uncommon (reported in 25% to 60% cases), but significant airway compromise is rare, occurring in an estimated 1% to 3.5% cases. Nevertheless, given the high frequency of EBV infection, this small proportion still represents many patients. Acute upper airway obstruction may occur, but the cardinal signs of acute obstruction (stridor and respiratory distress with recession and tachypnea) can be absent until late in the process. Obstruction arises from a combination of inflammation and hypertrophy of the palatal and nasopharyngeal tonsils, edema of the pharynx and epiglottis, and pseudomembrane formation in the large airways. Acute onset of obstructive sleep apnea has also been described in a child who required continuous positive airway pressure when asleep, until recovery after 3 days. Peritonsillar abscess formation is a rare complication that can further compromise the airway, and it is now believed that this is not significantly associated with the use of corticosteroids, which contradicts earlier reports.
Management is with systemic corticosteroids (in the presence of obstruction) and supportive care, which may include ventilation. A tracheostomy is sometimes necessary, as the severity of the inflammation in the oropharynx can make intubation difficult. If corticosteroids do not help, the role of acute tonsillectomy has been advocated, but it is controversial due to the high risk of perioperative bleeding.

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Access the reference list online at ExpertConsult.com.

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