Children with Upper Airway Dysfunction: At Risk of Obstructive Sleep Apnea

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Abstract

Obstructive sleep apnea is characterized by prolonged partial upper airway obstruction or intermittent complete obstruction that disrupts normal ventilation during sleep and alters normal sleep patterns. Patients with obstructive sleep apnea tend to develop neurocognitive, cardiovascular, behavioral, attention issues, and poor academic performance. Therefore, it is essential to diagnose and treat obstructive sleep apnea early and avoid significant and long-lasting adverse outcomes. Most commonly, upper airway obstruction is caused by enlarged lymphoid tissues within the upper airway, and therefore adenotonsillectomy is considered as the first-line treatment of obstructive sleep apnea in children. Fifty to 70% of patients who have obstructive sleep apnea and treated by surgery are not entirely cured on follow-up polysomnography. In light of this, it is recommended that patients with suspected obstructive sleep apnea undergo a thorough evaluation, and all potential risk factors are identified and treated. The purpose of this review is to familiarize pediatricians with developmental, anatomical, and physiological risk factors involved in the development of obstructive sleep apnea. Additionally, we will present an array of evaluation techniques that can offer adequate assessment of the patient’s upper airway anatomy and physiology.

Keywords
► upper airway
► sleep apnea
► sleep breathing
► tongue base
► snoring
► pathophysiology

Introduction

Obstructive sleep apnea (OSA) is a subtype of a larger class of sleep-related breathing disorders and is characterized by prolonged partial upper airway obstruction or intermittent complete obstruction that disrupts normal ventilation during sleep and alters normal sleep patterns.1 The prevalence of OSA is 2 to 4% among the general pediatric population. However, this prevalence changes with age and peaks between 1.5 and 5 years of age, coinciding with the peak growth of adenotonsillar tissues in the upper airway. In patients with Down's syndrome (DS), the prevalence can be as high as 66%.2 OSA is characterized by nighttime symptoms such as frequent snoring (typically ≥ 3 nights per week), labored breathing during sleep, mouth breathing, witnessed apneas, choking- or gasping-induced arousals, sleep enuresis, and restless sleep. Morning symptoms may include headaches on awakening, excessive daytime sleepiness, attention deficit/hyperactive disorder, and behavioral and learning problems. Also, these patients tend to have neurocognitive, cardiovascular, behavioral, attention issues, and poor academic performance.3–6 Risk factors encountered in children with OSA include adenoid and tonsil hypertrophy, turbinate hypertrophy, obesity, and other craniofacial abnormalities such as micrognathia/retrognathia and high-arched palate.7 Therefore, it is essential to diagnose and treat
OSA syndrome early and to avoid significant and long-lasting adverse outcomes.

OSA is caused by a complex interaction of structural, functional, and behavioral elements influencing the airways. The most common anatomical cause of upper airway obstruction is enlarged lymphoid tissues within the upper airway, and therefore adenotonsillectomy (AT) is considered as the first-line treatment of OSA in children. Yet, it has been shown in several studies and subsequently by a meta-analysis that AT is not very effective in curing OSA (i.e., normalizing the sleep study and all respiratory disturbances during sleep) in most children. For example, in a prospective study involving 56 children (age: 16 months to 12.5 years), Guillenlmault et al showed that in patients who had undergone AT or orthodontic intervention, 29 (51.7%) children still had an apnea–hypopnea index (AHI) of >1 event per hour of sleep and 12 (41.3%) patients had an AHI of >5 events per hour of sleep. Additionally, these researchers found that when the recommended multidisciplinary approach was not followed, rates of failures (i.e., AHI > 1 event per hour of sleep) were much higher (88.9 vs. 10%).

Two years later, Tauman et al showed in a prospective series of 110 children that obesity and OSA severity were important factors in the resolution of OSA after AT but also confirmed that the “cure rate” after AT was exceedingly low and averaged approximately 30%. Later in 2010, Bhattacharjee et al in a multicenter retrospective study looked at 578 children who underwent AT. They found that only 157 (27.2%) had complete resolution of their OSA by PSG (i.e., AHI < one event per hour of sleep) after AT. Additionally, they found that age > 7 years, obesity, and a history of asthma contributed to the persistence of OSA after AT.

These results reflect highly complex interactions of various factors involved in the pathogenesis of OSA. Hence, a comprehensive multidisciplinary evaluation for the management of OSA is critical for an accurate and personalized evaluation. In this article, we will review the various factors involved in the development of upper airway dysfunction and discuss some diagnostic strategies to evaluate these children.

**Airway Development**

To better understand the development of airway dysfunction, it is critical to acquire a good understanding of upper airway development. There are critical periods during airway development that can lead to dysfunction, which, if unrecognized or untreated, can lead to significant dysmorphism, necessitating surgical intervention. The upper airway extends from the nasal opening and mouth to the vocal cords in the larynx. The upper airway is a complex structure composed of various muscles from different embryological origins and is involved in several functions such as breathing, swallowing, and speech. Craniofacial development begins during the fourth week of embryonic life with the migration of neural crest cells, which progressively develop into many facial bones known as the maxillary and mandibular prominences through the activations of several genes such as Hox, FgF, and Pax-6. Following anatomical development, functional development of the swallowing and sucking reflex begins. These suck/swallow reflexes along with postnatal nasal breathing are crucial for the optimal development of craniofacial and upper airway anatomy and function.

**Prematurity**

Studies have shown that premature infants are at an increased risk of upper airway dysfunction due to decreased upper airway muscle tone and high nasal airway resistance. In addition, premature infants tend to have smaller airways. In one study, 8.1% of premature patients were reported to manifest snoring for three or more nights per week, and in another study, children with a history of prematurity were more likely to snore and undergo AT. Additionally, patients with a history of prematurity had a significantly higher AHI (AHI > 5 [0.9% in full-term vs. 4.3% in premature infants]) and significantly lower oxygen saturation during sleep. Preliminary evidence also shows that premature infants with high and narrow palates, with low muscle tone, and who are mouth-breathers create an inflammatory environment, leading to an increased risk of allergies and enlarged adenoids and tonsils. Infants are obligatory nasal breathers, and therefore premature infants who are at a high risk of increased nasal resistance due to underdeveloped nasomaxillary complex are also at a much higher risk of OSA. In addition, low muscle tone seen in these infants predisposes to habitual mouth breathing, leading to further craniomaxillary complex maldevelopment. Although anatomical problems tend to play a major role in causing OSA in premature infants, some studies suggest that blunted ventilatory responses to hypoxia during sleep in premature infants may also worsen gas exchange which can cause central respiratory pauses following desaturation.

**Anatomical Consideration of Upper Airway Dysfunction**

**Nose**

Previously stated embryological structures are necessary for the development of the base of the nasal cavity and the base and roof of the oral cavity. After birth, facial structures continue to develop. By age six, the face is 60% the size of the adult face dimensions, reaching maximal growth by the age of 12 years. During this time, facial growth is governed by the growth of the cranial base through endochondral ossification, which is under the influence of many physiological conditions. Breathing through the nasal cavity, the primary route of breathing in infants, provides stimulus for the proper growth of the sinuses, nasal, and oral cavity. In nasal airflow, two areas of major resistance are encountered: the nasal valve and the turbinate mucosa. These factors help in the proper development of the palate and dental arches. Cyclic changes in the blood supply to the mucosa alter the space for airflow through each nasal chamber; a phenomenon referred to as the nasal cycle. The nasal cycle can be altered in children with nasal allergies or nasal blockage. More detailed description about nasal airflow can be found elsewhere.

Obstructions at the level of the nasal cavity can cause a significant degree of obstructive sleep-disordered breathing.
resulting in sleep fragmentation. In adult studies, nasal obstruction has been shown to reduce time in deep nonrapid eye movement (NREM; stage N3) sleep and increase the incidence of apneic episodes. Based on the Starling law, upstream obstruction at the nasal level increases the turbulence of airflow and leads to increased propensity for airway collapse at the pharyngeal level. In children with allergic rhinitis, studies have shown increased incidence of obstructive SDB, which can be reduced by the administration of intranasal steroids and by other measures that can help increase nasal patency and reduce upper airway resistance. During childhood, altering normal nasal breathing aerodynamics can lead to craniofacial skeletal abnormalities. In these children, the floor of the nasal cavity (palate) moves up (high arch palate), and the maxilla is more constricted and retruded. Additionally, there is lowering of the inferior portion of the posterior nasal fossa, causing impingement on the pharyngeal region, thus increasing the chances of velopharyngeal obstruction. As a consequence of these changes, children can be easily distinguished in the clinic by the presence of “adenoid facies,” which include maxillary growth restriction, incomplete lip seal, narrow upper dental arch (posterior crossbite), crowded teeth, increased lower anterior face height, open mouth posture, a steep mandibular plane angle, and a retrognathic mandible.  

Oropharynx

Mastication and swallowing are primary functions of the oral cavity, and the oral cavity is not specifically programmed for normal breathing. As noted earlier, oral breathing is a default route to breathe in children with nasal obstruction. The oropharynx begins to develop in the second month of pregnancy. During this time, tongue placement on the roof of palate allows for closure of the mouth. This developmental process is a critical part in upper airway development as it sets up a physiological environment for breathing, sucking, swallowing, and phonation. The location of the tongue on the roof of the mouth in resting position acts like a natural expander to the hard palate and helps in normal maxillary and dental arch development. Integrated into the development of the oropharynx is positioning of the hard and soft palate relative to the tongue and adenoid tissues, which are positioned posteriorly to the nasopharynx. As discussed earlier, children with nasal obstruction become mouth-breathers and are predisposed to low tongue posture, which may contribute to a narrow palate and a reduction of the space for the tongue in the oropharynx (small mouth size or relative macroglossia). Hence, during the development of the oropharynx, if physiological homeostasis in breathing is altered, obstruction to breathing can happen at various locations such as tonsils, soft palate, and tongue base, and then at the laryngeal level.

To summarize the site and type of obstruction, Kezirian et al developed the VOTE (velum, oropharynx lateral walls, tongue base, epiglottis) classification. Based on observations from drug-induced sleep endoscopies (DISEs), the VOTE classification focuses on the most common sites of obstruction and their configuration. The structures included in the classification are the velum, oropharynx and lateral walls, tongue base, and epiglottis. The degree of obstruction can be classified into no obstruction, partial obstruction, or complete obstruction. Finally, the configuration of airway collapse can be stated as anteroposterior (AP), lateral, or concentric type.  

**Table 1** The VOTE classification (reproduced with permission from authors)

<table>
<thead>
<tr>
<th>Structure</th>
<th>Degree of obstruction</th>
<th>Configuration</th>
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<tbody>
<tr>
<td>Velum</td>
<td></td>
<td></td>
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<tr>
<td>Oropharynx lateral walls</td>
<td></td>
<td></td>
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<tr>
<td>Tongue base</td>
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<tr>
<td>Epiglottis</td>
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Abbreviation: AP, anteroposterior.  
Note: for each structure, there should be a classification of the degree of obstruction and configuration of obstruction. Open boxes reflect the potential configuration that can be visualized related to a specific structure. Black boxes reflect the fact that a specific structure configuration cannot be seen.

Degree of obstruction has one number for each structure: 0, no obstruction (no vibration); 1, partial obstruction (vibration); 2, complete obstruction (collapse); X, not visualized.

Configuration noted for structures with degree of obstruction greater than 0.

Oropharynx obstruction can be distinguished as related solely to the tonsils or including the lateral walls, with or without tonsil component.
positive airway pressure (CPAP), and accordingly, other therapies could be offered to these patients.

**Tongue**

The tongue plays an important role in the development of upper airway obstruction. As part of the inspiratory phasic upper airway muscle dilator (along with influencing the hyoid position and the muscles of the palate), the genioglossal muscle is tensed during inspiration, increasing negative pressure (i.e., decreasing $P_{crit}$). As previously stated, as the patient goes into NREM sleep, glossal and pharyngeal muscle tonic activity is reduced, causing an increase in the capacitance and collapsibility of the upper airway.\(^{25,26}\) This mechanism for collapsibility has been shown to be augmented in patients with OSA. Additionally, the reflex that blunts this response in normal patients also affects the patients with OSA.\(^{27}\) For example, the genioglossus muscle has been shown to function at 40% of its capacity in patients with OSA.\(^{28}\) Therefore, tongue base obstruction is a major site of obstruction in children with OSA. Previous research has shown that upper airway muscle tone, mandible size, and tongue volume are all variables that influence the development of obstruction.\(^{29}\) As seen in many of the studies of OSA in patients with DS, the decreased ratio of mandibular area to tongue area can cause airway overcrowding and posterior displacement of tongue, consequently increasing the risk of obstruction at the level of the tongue base.\(^{30}\) In one study, the researchers found that the previously stated ratio by magnetic resonance imaging (MRI) caused airway obstruction by glossoptosis and hypopharyngeal collapse in nearly two-thirds of children with OSA and DS after AT.\(^{31}\) Soft tissue surrounding the upper airway also plays an essential role in upper airway collapse. It has been clearly shown that fatty deposition in the tongue and around the airway is a risk factor for OSA.\(^{32,33}\)

**Epiglottis**

At the level of the epiglottis, laxity has also been shown to obstruct the upper airway. The epiglottis might fold posteriorly or laterally and cause airway obstruction.\(^{23}\) In adults, this can happen in 12% or more of patients with OSA. Sleep-state laryngomalacia is a distinct condition where the epiglottis tends to become floppy and completely collapses, blocking airflow. However, in awake nasal endoscopy, the larynx appears normal. This is readily seen in pediatric patients diagnosed with laryngomalacia.\(^{34}\) As the upper airway becomes obstructed, changes in the oropharynx occur, which are often missed by the clinicians. Specifically, with prolonged obstructive breathing, children develop cephalometric parameter changes that have been associated with OSA.

**Obstructive Breathing and Changes in Skeletal and Dental Structures**

Luzzi et al measured several parameters in patients with primary snoring using lateral cephalograms.\(^{35}\) In this study, researchers measured sagittal measurements such as the sella–nasion–A (SNA) point angle, the sella–nasion–B (SNB) point angle, and the A point–nasion–B point angle (ANB) (Fig. 1A). These measurements defined the AP relationship between the maxilla and the mandible. In the vertical axis, the Frankfurt mandibular angle (FMA) was measured. The FMA angle is the angle between the Frankfurt horizontal plane and the mandibular plane, a well-established measurement of total facial divergence (Fig. 1B). The FMA angle is considered normal between 20 and 30 degrees. These investigators

![Fig. 1 Cephalometric parameters. (A) Frankfurt mandibular angle (FMA) measures the angle between the Frankfurt horizontal plane and the mandibular plane. (B) SNA, “sella–nasion–A point” angle, measures the anteroposterior relationship of maxillary basal arch on anterior cranial base. SNB, “sella–nasion–B point” angle, measures the anterior limit of the mandibular basal arch in relation to the anterior cranial base. Go, gonion or mandibular angle; Me, mental protuberance; PM, plane of mandible; PFH: plane of Frankfurt. Reproduced with permission from Luzzi et al.\(^{35}\)](image)
showed that the FMA angle was the only orthodontic parameter that was statistically associated with the level of airway obstruction. When the FMA angle is increased, the degree of airway obstruction is also increased.\textsuperscript{38} Likewise, other studies have shown that patients who are mouth-breathers develop specific phenotypic characteristics that should alert physicians to evaluate these children for upper airway dysfunction. Pacheco et al studied the prevalence of morphological changes in 520 healthy children, ranging in age from 7 to 12 years old. Within their population, 167 (24.3\%) patients were classified as mouth-breathers. Within this group, 26.1\% had excessive overjet and 17.7\% had an anterior open bite.\textsuperscript{36} Additionally, within the mouth-breather group, 53.9\% had atretic palates. Finally, this group also had a significantly higher prevalence of dolicho facial pattern as well as a convex facial profile.\textsuperscript{36} In adults, Lee et al studied craniofacial morphology with objective facial measurements. As seen in other studies, patients with OSA tended to have shorter and retruded mandible, a smaller area within the mandible, and more fat deposition on the anterior neck. These changes cause patients with OSA to have a brachy cephalic head and euryprosopic facial characteristics (more extensive laterally, shorter AP dimensions, and short vertically) when corrected for BMI.\textsuperscript{37}

Dental morphology has also been shown to be drastically altered in patients with upper airway obstruction and therefore should be inspected. Pirlä-Parkkinen et al analyzed dental arch morphology in children with SDB and evaluated children who snored, those diagnosed with OSA, and healthy controls. The mean age of the children in the study was 7.2 years (range: 4.3–11.4). Results revealed again that children with OSA had a narrower upper dental arch, increased overjet, reduced overbite, and shorter length of the lower dental arch when compared with nonobstructed controls.\textsuperscript{38} Additionally, this study showed that patients who snore, but are do not fulfill criteria for OSA, still have alteration in their dental arch morphology. Interestingly, many of the patients with OSA did not have enlarged tonsils.\textsuperscript{38}

**Functional Characteristics of Upper Airway Dysfunction**

**Control of Breathing during Sleep**

Breathing is controlled by both peripheral and central inputs. Peripheral chemoreceptors in the carotid and aortic bodies and central chemoreceptors in the ventral area of the medulla oblongata and other areas of the central nervous system sense physiological metabolites (i.e., hydrogen ions and carbon dioxide), which activate the central pattern generator (CPG) area of the brainstem.\textsuperscript{16} Activation of the CPG transmits signals to the respiratory muscles to maintain a normal breathing pattern and gas exchange. During sleep, there is a decrease in the ventilatory drive, shown as a decrease in the slope of the ventilation/CO\textsubscript{2} relationship in NREM sleep and rapid eye movement (REM) sleep. Studies have also shown a decrease in hypoxic ventilatory sensitivity during NREM and REM (39\% and 52\%, respectively).\textsuperscript{39–41} Finally, upper airway muscle tone can also be affected during sleep. Because these muscles are under the influence of respiratory chemoreceptors, a decreased or blunted response to hypoxia and hypercapnia allows for increasing resistance of the upper airway during sleep and the emergence of prolonged obstructive events.\textsuperscript{27}

**Upper Airway Neuromuscular Tone and Critical Closing Pressure (P\textsubscript{crit})**

The pharynx constitutes the last portion of the upper airway, made up of mainly skeletal musculature. The patency of the muscular tube is dependent on muscle tone and neurologic activity during sleep. During sleep, this highly compliant muscle tube is under the influence of negative pressures created by the expansion and compression of the diaphragm. Therefore, a small change in pressure can drastically alter its diameter. Critical closing pressure (P\textsubscript{crit}) is the pressure at which the airway collapses and closes completely. Usually, P\textsubscript{crit} in the awake state is between –7.4 cm H\textsubscript{2}O, and –25 cm H\textsubscript{2}O during sleep. The more negative P\textsubscript{crit} during sleep signifies a decrease in compliance of the pharynx and an increase in muscle tonicity during sleep, and hence more stable and less collapsible airways. Conversely, the closer the P\textsubscript{crit} gets to zero or becomes positive, the more collapsible the airway becomes. Infants and children tend to have more negative P\textsubscript{crit} than adults, and hence their airways are less collapsible compared with adults.\textsuperscript{42} Therefore, in children, OSA is caused mainly by anatomical factors and less by neuromuscular factors. However, patients with OSA tend to have more positive (higher) P\textsubscript{crit} during sleep, which predisposes their airway to collapse easily. Huang et al analyzed this effect by measuring several parameters during sleep in patients with OSA and controls.\textsuperscript{43} Their study revealed that patients with OSA had pressure drops during REM sleep, which significantly affected airflow when compared with controls. For example, the mean drop in airflow in the OSA group was –44.33 ± 14.09 mL/seconds. Yet, many of these patients were able to compensate and maintain minute ventilation by increasing their respiratory rate. Unfortunately, other studies have revealed that many patients with OSA have decreased compensatory mechanisms, such as the one seen in the previous study.\textsuperscript{27} Therefore, it is essential that patients with OSA maintain lower P\textsubscript{crit} pressures to avoid airway collapse during sleep, which is known to cause apneas and hypopneas. Besides the inherent characteristics of the pharynx, many extrinsic characteristics may mitigate or enhance the collapsibility of the pharynx. Adipose tissue deposit in the surrounding connective tissue and around the pharynx in the cervical region, and the neck circumference has been shown to be a risk factor for OSA.\textsuperscript{44} Finally, flexion of the neck has shown to worsen upper airway collapsibility, and neck extension may reduce upper airway collapsibility in patients with OSA.\textsuperscript{35} For this reason, many children with severe OSA will adopt a hyperextended neck posture over a pillow during their sleep.

**Loop Gain and Arousal Threshold**

Besides airway pressures, neurologic responses to hypoxia can be measured and have been shown to be related to the
development of OSA. Loop gain, originally an engineering term, is a method used to measure the stability of a system’s feedback loops. In medicine, loop gain has been used to measure the stability of the upper airway during sleep. This method specifically measures the response of the central nervous system to a disturbance in the upper airway. For example, an excessive ventilatory response (hyper-ventilation) to hypoxia (disturbance) will decrease PaCO₂ due to ventilatory overshooting more than necessary (increased loop gain) and also generate upper airway hypotonia. As a result of this excessive response to hypoxia, upper airway obstruction can be recurring and perpetuating. Loop gain has been shown to have a causal relationship with OSA, complementing the theory that OSA has both structural and functional components of the upper airways that need to be considered during the clinical evaluation and treatment. Finally, another pathological disturbance that decreases the nervous system from compensating for hypoxia or upper airway obstruction is reduced arousal threshold, leading to premature termination of the obstructive events, which impedes the central ventilatory drive from acting on the upper airway, reducing stable breathing during sleep.

**Evaluation of Upper Airway Obstruction**

**Imaging Studies**

Imaging of upper airways has become an essential part of the evaluation of patients with OSA. As more physicians recognize that patients with OSA might have several sites of obstruction, identifying the anatomy of these patients is essential. Both structural and dynamic evaluation of the airways is essential to assess both size and stability of the airways. Cine magnetic resonance imaging (Cine MRI) has become an integral part of evaluating the upper airway anatomy of patients with DS and OSA. Cine and static MRI allow for proper assessment of the airway and its surrounding tissues. Several studies have shown that MRI is a useful tool for evaluating upper airway anatomy and collapsibility and assessing different sites of obstruction simultaneously in children with DS with OSA. One meta-analysis revealed that patients with OSA had a significantly reduced total volume of the upper airways when compared with controls (1.4 ± 0.7 vs. 1.6 ± 0.7 cm³). A slight reduction in the diameter of airway has a significant impact on the airway resistance. MRI could be used to evaluate the size of lymphoid tissue as well as the cross-sectional area regions surrounding hypertrophied lymphoid tissue. This meta-analysis revealed that MRI could evaluate differences in upper airway volume in pediatric patients with OSA compared with healthy controls (mean difference: −0.53 cm³; 95% confidence interval: −1.07 to −0.07), as well as soft tissue and skeletal muscle surrounding the airways.

In another study, researchers evaluated the effect of dynamic (cine) MRI to localize and identify the sites of obstruction in the upper airways. Using either 1.5- or 3.0-T MRI equipment, sagittal three-dimensional respiratory-triggered fast spin echo, or proton-density sequence, researchers captured sequential videos of patients with OSA. Slices of 1.6 mm thickness were used to captures images from the tip of the nose anteriorly to the occiput posteriorly, and from the aspect of the nasal airway superiorly to the subglottic trachea inferiorly. The researchers used dexmedetomidine.
for sedation, which does not cause significant respiratory depression.\textsuperscript{52}

Currently, the expert opinion states that a 50\% collapse of the airway is abnormal and that 80\% is very abnormal and should prompt further evaluation. Chen et al also showed that upper airway loop gain (UALG) could identify anatomical risk factors for OSA.\textsuperscript{53} In this study, UALG was determined by measuring the ratio of the change in area drop to the change in area overshoot before and after airway collapse. Although patients with OSA did not have significantly different UALG ($n = 4$) from controls ($n = 3$), patients with higher AHI tended to have higher UALG values. Additionally,
functional upper airway area did differ significantly between OSA patients and controls, suggesting simultaneous multislice real-time MRI (SMS RT-MRI) as a valuable tool to assess anatomy and physiology of patients with OSA. Therefore, UALG is both a measurement of airway collapse and a response to airway collapse that can be used to evaluate anatomy and pathophysiology of OSA patients.

Drug-Induced Sleep Endoscopy

Finally, DISE has evolved to become a valuable tool for the evaluation of upper airway obstruction. Specifically, this non-invasive procedure allows clinicians to precisely locate points of obstruction during medication-induced sleep. Controversy exists regarding which anesthetic drugs are best mimickers of natural sleep, and hence there is no standardized protocol. When anesthetized, the patient’s sleep stage can be approximated using the bispectral index score. Finally, the endoscope should be introduced through the nasal cavity, allowing visualization of the retrolarval, oropharyngeal, retrobasillar, retroepiglottic, and glottic regions. If done correctly under optimal anesthesia to mimic natural sleep, DISE procedure can provide a comprehensive evaluation of the multisite obstruction in the upper airway. The site and the severity of the upper airway obstruction can be assessed based on the VOTE classification mentioned previously.

Conclusion

In conclusion, airway development starting in utero through adolescence has a significant impact on the normal development of upper airway anatomy and physiology. Patients who present with minor signs of upper airway obstruction (i.e., snoring and adenoid facies) should be evaluated closely in the clinic, and studies assessing the degree of obstruction (polysomnography), upper airway anatomy (imaging studies), and function (DISE, UALG) should be performed to identify the site and severity of upper airway obstruction. These complementary and informative approaches will allow physicians to accurately phenotype upper airways, leading to a targeted and personalized treatment strategy and to treat, cure, or potentially prevent OSA before it can have its lasting effects on the child’s overall health and quality of life.

Conflict of Interest

None declared.

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