The Otolaryngologist's Approach to the Patient with Down Syndrome

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KEYWORDS

- Otolaryngology Down syndrome Quality of life Morphology Otitis media
- Hearing loss Obstructive sleep apnea Tonsillectomy Sinusitis Intubation
- · Acquired subglottic stenosis

KEY POINTS

- The child with Down syndrome should undergo behavioral audiologic testing every 6
 months, or every 3 months if the patient has very stenotic ear canals. Treatment of hearing
 loss caused by recurrent otitis media and otitis media with effusion (OME) should be
 aggressive, with close follow-up. Parents should be prepared for multiple sets of pressure
 equalization (PE) tubes throughout the child's life.
- Up to 80% of patients with Down syndrome have obstructive sleep apnea. All children with Down syndrome should get an overnight polysomnography study between the ages of 3 and 4 years. Parents are not reliable in assessing sleep apnea.
- Primary treatment of obstructive sleep apnea syndrome (OSAS) is tonsillectomy and adenoidectomy, but parents should be prepared that this is curative in only 25% to 50% of children with Down syndrome, and that their child may require further surgery to alleviate obstruction, or the child may continue to require continuous positive airway pressure (CPAP).
- Airway anomalies are more common in the Down syndrome population, including subglottic stenosis. Prevention of subglottic stenosis can be achieved in part by using an endotracheal tube 2 sizes smaller than predicted for the patient's age, and ensuring that an audible air leak is present around the tube.
- During surgery in patients with Down syndrome, the surgeon should always be aware of the possibility of atlantoaxial instability, and no dramatic head movement should be made.

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