Polisomnographic findings on children with laryngopathies

Michele Themis Moraes Gonçalves, Juliana Sato, Melissa A. G. Avelino, Gilberto U. Pizarro, Gustavo A. Moreira, Márcia Pradella Hallinan, Reginaldo R. Fujita, Luc Louis Maurice Weckx

Summary

Polysomnography is the gold standard exam for child OSAS. When possible, polysomnography clearly distinguishes between those with isolated primary snoring and patients with sleep apnea (obstructive, central and mixed). The most common cause of OSAS in childhood is adenotonsillar hypertrophy. Laryngomalacia is the most common cause of stridor in childhood, though its physiopathology remains unknown. Among the most prominent theories are immaturity of the cartilaginous framework of the larynx and/or neuromuscular immaturity. Objective: Our proposal was to describe polysomnographic findings in children with laryngomalacia or other isolated laryngeal alterations, that is, without other alterations in the upper airways. Methods: The sample included 29 children with exclusively laryngeal alterations. All of them underwent an otorhinolaryngological exam, nasofibrolaryngoscopy and polysomnography. Information was recorded concerning age, nasofibrolaryngoscopy and polysomnography. For analysis, the children were divided into two groups: those with laryngomalacia and those with other laryngeal diseases. Results: Among the 18 children with a diagnosis of laryngomalacia, 18 had central breathing events, knowing that the majority had showed desaturation of oxihemoglobin and bradicardia. In this same group, 3 children had obstructive events. On the other hand, 11 children with other laryngeal alterations showed no predominance of one type or another of apnea. Of these, 4 had central type breathing events and 2 obstructive type. Conclusion: The majority of patients with laryngomalacia showed a central type apnea. Patients with various laryngeal diseases did not present a predominant type of apnea.

Key words: Polysomnography, Larynx, Apnea, Children.
INTRODUCTION

Snoring and mouth breathing are common complaints in childhood, with incidence rates varying from 3 to 26%. In the pediatric population, we have 1 to 3% of patients with Sleep Obstruction Apnea Syndrome (SOAS), and snoring is the most common symptom.

Snoring is what we call the vibratory sound produced in the nasopharynx during inspiration. Primary snoring is not related to apnea, hypoventilation, hypoxia, hypercarbia, restless sleep or excessive daily sleepiness.

SOAS in children is characterized by partial or complete obstruction of the upper airways during sleep, usually followed by a drop in oxyhemoglobin saturation and/or hypercarbia. The first series of SOAS cases in children was published in 1976 by Guilleminault et al. SOAS in children is different from its adult counterpart. In children the disease comprises episodes of isolated obstructive hypoventilation, apnea associated to obstructive hypoventilation, and plain obstructive apnea. In other words, the sleep obstructive respiratory disorders are called SOAS even when there are no apnea episodes.

By definition, apnea is the lack of airflow by either the nose or the mouth. It may be caused by an interruption in respiratory movements (central-type apnea) or by a collapse of the upper airways (obstructive-type apnea). When we have central and obstructive apnea occurring simultaneously in children, in order to be considered relevant, they should last more than 2 respiratory cycles and are not always associated to hypoxemia. Short duration central apnea (less than 15 seconds) is a common finding in the sleep of normal newborn children.

Insofar there is no criterion that defines an acceptable number of central apnea episodes. Notwithstanding, if associated to bradycardia or cyanosis, regardless of its duration, it is considered significant. Heart-related, blood-related, infectious, metabolic, neurological, pulmonary, duration, it is considered significant. Heart-related, blood-related, infectious, metabolic, neurological, pulmonary, gastrointestinal and neuromuscular causes are mentioned for central apnea, although most of the times they are not identified.

Hypoventilation means a drop in pulmonary ventilation below the minimum necessary to sustain normal oxygen saturation. Just like apnea, it may also happen from non-obstructive causes (lower central respiratory stimulus, neuromuscular abnormalities or pulmonary restrictive diseases) or obstructive (partial obstruction of upper airways causing inadequate pulmonary ventilation) causes.

Contrary to what happens in adults, in whom the major SOAS symptom is daily sleepiness, children are most often taken to the doctor, because of snoring or respiratory difficulty during sleep. Night time awakenings, chronic mouth breathing, excessive daily sleepiness, psychomotor restlessness, excessive movements during sleep are also frequent. Children with SOAS may have low school performance, disciplinary problems, attention deficits, mood alterations (irritability, aggressiveness), swallowing disorders and voice changes. Besides all of these, there may be important growth deficits because of alterations in GH secretion during sleep. Other hypothesis, still not confirmed, is that weight and height deficit may be secondary to greater energy expenditure during a more laborious breathing or due to less calorie ingestion caused by poor appetite or dysphagia. Severe cases may be associated to cor pulmonale and death.

Pediatric SOAS is not fully understood. We know that there is no single mechanism responsible, and that it is very likely that pediatric SOAS with adenotonsillar hypertrophy be different from pediatric SOAS with craniofacial or neurological syndromes.

The otorhinolaryngological exam is very important and starts by facial inspection, looking for retrognatism or hypognatism. After that we assess tongue size and its relationship with the dental arch, presence of tonsil hypertrophy, high palate and cross bite. In the nose, we look for inferior turbinates hypertrophy or cyanosis, septal deviations, secretions, polyps or tumors. Nasofibroscopy is necessary and very important because it may define, in children, the sites of upper airway narrowing, adenoid-chouma ratio, tongue base and oropharynx with the pharyngeal posterior wall, side to side tonsil collapse, signs of gastroesophageal reflux or glottic and subglottic alterations.

As it happens with adult patients, the gold standard tests for childhood SOAS is the polysomnography, in order to assess both the severity and the therapy mode to be followed. It is a test that requires skilled professionals in dealing with children, and very important is active parent participation. When carried out, polysomnography clearly separates those bearers of primary snoring from sleep apnea patients (obstructive, central and mixed).

Children SOAS diagnostic criteria were only recently defined. In children, one or more obstructive-type apnea episodes per hour and/or obstructive-type hypoventilation, characterized by peak CO2 exhaled (ETCO2) = 53 mmHg, ETCO2 > 50 mmHg in more than 10% of total sleep time or ETCO2 > 45 mmHg in more than 60% of total sleep time, are considered to be abnormal. HAI greater than 10 is considered very severe. The polysomnography test of primary snorers have hypopnea/apnea index (HAI) lower than 1 episode per hour, oxygen saturation average during sleep greater than 90% and ETCO2 peak < 53 mmHg or ETCO2 peak = 50 mmHg in less than 10% of total sleep time. As to the assessment of sleep efficacy, it is considered abnormal when it is lower or equal to 85%.

The most common cause of SOAS in children is adenotonsillar hypertrophy, and chronic hypertrophic allergic rhinitis makes it even worse. It is also very important to check other anatomical causes (nasal septum...
deviation, nasal polyps, choanal stenosis, tongue hypertrophy, micrognatia, laryngomalacia, laryngeal diaphragm), congenital and syndrome diseases (Apert, Cri du Chat, Crouzon, Down, Pierre Robin, Treacher-Collins syndromes; mucopolysaccharidosis, cystic fibrosis, Arnold-Chiari, brain palsy) among others (myopathy, neuromuscular disease, neoplasia, family issues, gastroesophageal reflux).

Laryngomalacia is the most common cause of children stridor\textsuperscript{15}, although its prevalence is unknown. Literature data report incidence rates varying from 19.4 to 75\%\textsuperscript{16,17}. Stridor usually appears after the first weeks of life and may remain until 18-24 months. Stridor may increase in restlessness, feeding or supine position. It happens because of supraglottic collapse secondary to excess mucosal tissue in the larynx posterior wall, shortening of the aryepiglottic fold or epiglottis cartilage falling over the larynx lumen. Its cause is still debated. There are many theories proposed to explain its physiopathology, such as disorders in the cartilaginous framework of the larynx and trachea, causing a greater laxity in supraglottic structures, anatomical alterations and neuromuscular immaturity\textsuperscript{18-21}.

Back in 1897, Sutherland & Lack\textsuperscript{22} proposed an anatomical theory after studying 18 cases of congenital laryngeal obstruction, in which they concluded that the disorder was associated to the immaturity of the children cartilaginous tissue. Prescott\textsuperscript{23} added to this hypothesis by studying 40 patients with laryngomalacia and noticed that all of them had short aryepiglottic folds and 30\% had neuromuscular disorders.

Thompson and Turner\textsuperscript{24} demonstrated that a passive medial supraglottic prolapse could be induced from the denervation of children larynxes. This study, together with others such as the Perón et al.\textsuperscript{25} and Wiggs and Dinardo\textsuperscript{26}, established the neurological hypothesis. The neurological theory explains the association between laryngomalacia and other neurological disorders\textsuperscript{22,27}. Belmont and Grundfast\textsuperscript{28} found 80\% of LFR, 13\% of hypotonia and 10\% of central apnea in 30 children with laryngomalacia.

The diagnosis of laryngomalacia is achieved through flexible laryngoscopy during spontaneous breathing. It is usually of spontaneous resolution. Literature reports indicate that severe laryngomalacia is uncommon to the point of causing cyanosis, feeding difficulties, cor pulmonale, apnea or growth impairment. When it happens, patients with severe laryngomalacia are usually treated by tracheostomy, notwithstanding there are a number of other surgical treatments (aryepiglottic fold resection, epiglottoplasty and supraglottoplasty).

**OBJECTIVE**

Our goal was to describe polysomnographic findings in children with laryngomalacia and other isolated larynx alterations, in other words, without alterations in their upper airways.

**MATERIALS AND METHODS**

This work was carried out in the Mouth Breather Center - Otorhinolaryngology Department - Pediatric Otorhinolaryngology - Federal University of São Paulo, and the Sleep Institute from the Psychobiology Department of the Federal University of São Paulo. From 1999 to 2004, 29 children with laryngomalacia or other laryngeal alterations were selected. Their ages varied from 11 days of life to 8 years and 10 months. All of them underwent otorhinolaryngological exam, nasofibroscopy and polysomnography. We included only children with snoring and laryngeal stridor or vocal alteration in whom the fibroscopy exam showed laryngomalacia or other isolate laryngeal alteration. Laryngomalacia was considered when there was an excess of mucosal tissue in the posterior larynx wall, shortening of the aryepiglottic fold or collapse of epiglottis cartilage over the laryngeal lumen.

Children with adenotonsillar hypertrophy, craniofacial malformations or genetic syndromes were excluded.

Patients underwent day time polysomnography at the Sleep Institute - UNIFESP-EMF, where the exam was carried out with the patient sleeping in a comfortable, in a dark and quiet room. Electrophysiological and cardio-respiratory parameters were recorded in a computerized system (Alice®): electroencephalogram (4 channels), submentonian and tibial electromyogram, right and left electro-oculography, oronasal air flow, thorax and abdominal movements, microphone, oxyhemoglobin saturation (\textit{SaO2}), and sleeping position. The patient was studied under regular room air. We plotted data related to age, nasofibroscopy and polysomnography. In order to analyze them, the children were divided in two groups: those with laryngomalacia and those with other laryngeal diseases (subglottic stenosis, hemangioma, synchiae, vocal fold paralysis, vocal gap, arytenoid fixation and/or vocal fold thickening).

In order to carry out a data statistical analysis, we used the Fisher Test, which did not show significance among the results.

All the patients handed in a duly signed informed consent form, and this research was approved by the Ethics Committee. (Protocol # 0122/02).

**RESULTS**

Tables 1 and 2 depict the polysomnographic results found among the laryngomalacia patients and other laryngeal lesions, respectively.

Among the 18 children diagnosed with laryngomalacia, all of them had central-type respiratory events, and most of the episodes were related to oxygen desaturation and some to bradycardia. In this same group, 3 children had obstructive-type apnea. As to the 11 children with other laryngeal alterations, 4 had central-type respiratory
Oxygen saturation during sleep for these children varied from 76 to 98%.

Patient INJ, of three months of age has severe laryngomalacia, for he presented baseline oxygen saturation (awaken and during sleep) of 89%. Although the patient had 2 central-type respiratory events, there was an improvement in oxygen saturation during sleep.

DISCUSSION

From 3 to 26% of the pediatric population seek specialized medical help because of snoring and mouth breathing, and only 1 to 3% of those have Sleep Obstructive Apnea Syndrome (SOAS). This disease is suspected through history taking; however this alone is unable to separate primary snorers from SOAS patients. Polysomnography is indicated in these cases, because besides defining diagnosis, it also characterizes the type of event that happened, obstructive, central or mixed, and also assessing disease severity through the Hypopnea/apnea index (HAI) and oxygen saturation level. It is important to remember that, as far as physiopathology and polysomnographic diagnostic criteria are concerned, SOAS in children is considered a different disease from the one that affects adults.

Otorhinolaryngological exam, together with nasofibroscopy is able to find most of the apnea causes. Although most of the times it is caused by adenotonsillar hypertrophy, we must bear other causes in mind, laryngeal diseases amongst them.

Laryngomalacia physiopathology is still controversial. Among the most accepted theories we have the immaturity of the nervous system. This finding is in agreement with what has been seen in the paper by Milczuk & Johnson29, in which children with laryngomalacia presented comorbidities such as LFR, prematurity and DNPM delays, amongst others, all of them related to some degree of neurological immaturity.

Table 1. Polysomnography data observed in patients with laryngomalacia.

| Patient | Age | naso | sat vig | SC | SA | NED | NEC | IAC | NEO | nadir | FC V | FC A | FC Q |
|---------|-----|------|--------|----|----|-----|-----|-----|-----|-------|------|------|------|
| TMS     | 6m  | LM   | 98%    | 96%| 93%| 1    | 2   | 0.9 | 0   | 89%   | 167.5| 135.1| 135.6|
| LAB     | 5m  | LM   | 95%    | 92%| 92%| 1    | 1   | 0.4 | 0   | 89%   | 145.0| 123.5| 126.0|
| ECM     | 5m  | LM   | 90%    | 92%| 93%| 1    | 1   | 0.7 | 0   | 85%   | 173.0| 151.4| 158.8|
| GSS     | 6m  | LM   | 96%    | 96%| 96%| 20   | 20  | 7.7 | 0   | 88%   | 163.2| 139.1| 134.2|
| DASS    | 9m  | LM   | 98%    | 97%| 98%| 7    | 7   | 1.0 | 0   | 92%   | 138.2| 126.9| 119.8|
| SNR     | 5m  | LM   | 95%    | 95%| 95%| -    | 1   | 0.5 | 0   | 95%   | indisp| indisp| indisp|
| TNB     | 2m  | LM   | 96%    | 94%| 96%| 7    | 7   | 2.6 | 1   | 90%   | 174.9| 136.9| 141.0|
| SMS     | 11m | LM   | 92%    | 90%| 89%| 3    | 12  | 7.5 | 0   | 76%   | 158.7| 141.9| 138.4|
| PJS     | 3m  | LM   | 98%    | 96%| 95%| 3    | 5   | 2.0 | 0   | 91%   | 160.2| 130.1| 134.0|
| MGF     | 6m  | LM   | 97%    | 95%| 93%| 5    | 6   | 2.8 | 0   | 87%   | indisp| indisp| indisp|
| LFSS    | 21d | LM   | 96%    | 96%| 92%| 6    | 8   | 4.2 | 0   | 91%   | 164.3| 151.7| 138.4|
| KMS     | 7m  | LM   | 97%    | 97%| 97%| 10   | 12  | 4.6 | 0   | 88%   | 177.9| 151.8| 152.1|
| KSS     | 3m  | LM   | 96%    | 96%| 95%| 4    | 18  | 6.9 | 0   | 92%   | 154.6| 137.8| 134.0|
| ENR     | 2m  | LM   | 99%    | 98%| 98%| 3    | 18  | 8.9 | 0   | 93%   | indisp| indisp| indisp|
| INJ     | 3m  | LM   | 89%    | 93%| 93%| -    | 2   | 1.0 | 0   | 90%   | 182.4| 143.6| indisp|
| EGS     | 17m | LM   | 98%    | 95%| 96%| -    | 24  | 2.6 | 7   | 92%   | 129.4| 129.0| 124.3|
| BNM     | 4m  | LM   | 96%    | 96%| 96%| 1    | 2   | 1.5 | 0   | 92%   | 152.5| 121.8| 119.1|
| ASO     | 4m  | LM   | 96%    | 94%| 94%| 13   | 30  | 25.0| 1   | 86%   | 136.8| 125.0| 122.4|
Table 2. Polysomnography data observed in patients with other laryngeal diseases.

| Patient | Age | Diagnosis                        | Sat vig | SC | SA | NED | NEC | IAC | IAH O | Nadir |
|---------|-----|----------------------------------|---------|----|----|-----|-----|-----|-------|-------|
| CRM     | 6a  | bilateral vocal nodules          | 99%     | 98%| 98%| indisp | 0  | 0.4  |       | 92%   |
| JCS     | 2a8m| Post muc fall glottis            | 99%     | 99%| 99%| indisp | 0  | indisp |       | 89%   |
| PRC     | 8a  | post arythenoid fix *            | 99%     | 97%| 97%| indisp | 0  | indisp | 0.8   | 95%   |
| RAS     | 7a  | Bilat VF fixation*              | 98%     | 98%| 98%| indisp | 3   | indisp |       | 94%   |
| SFC     | 7a  | bilateral vocal nodules          | 98%     | 96%| 96%| indisp | 0  | indisp | 0.9   | 89%   |
| KMF     | 13d | LVF idiopathic paralysis         | 98%     | 98%| 97%| indisp | 162 | indisp |       | 87%   |
| MHS     | 1a  | Obstant subglottic stenosis 70%  | 96%     | 96%| 97%| indisp | 2   | indisp |       | 94%   |
| OGST    | 1a  | aryepiglottic fold shorten       | 93%     | 92%| 92%| indisp | 16  | indisp |       | 87%   |
| AAA     | 6a  | papilomatose obst larynx         | 98%     | 96%| 95%| indisp | 0  | indisp |       | 92%   |
| EPC     | 2a  | arythenoid hemangioma            | 93%     | 92%| 91%| indisp | 0  | indisp |       | 72%   |
| BGN     | 8a10m| subglottic stenosis             | 95%     | 95%| 93%| indisp | 0  | indisp |       | 84%   |

Legend:
- naso: nasofibroscopic diagnosis
- IAC: Central apnea index
- Sat vig: saturation when awake
- NEO: number of obstructive events
- SC: calm sleep
- SA: active sleep
- NED: desaturation events
- IAH O: Heart rate at calm sleep
- NEC: number of central events
- FC Q: Heart rate during active sleep
- indisp: not available

the majority: we may mention ENR, that despite having 18 respiratory events, none of them were associated to oxygen desaturation below 93%, while OGST had practically the same number of events, however with relevant oxygen desaturation (87%).

CONCLUSION

Most of the children with laryngomalacia or other laryngeal diseases did not have significant respiratory alterations during sleep. However the group of infants with laryngomalacia presented more central-type apneas. In this group, although most of the respiratory events may be considered benign, their greater frequency suggests partial obstruction of the upper airways.

Having seen the findings of our study, we believe the evaluation of respiratory disorders during sleep in patients with laryngeal disorders, using modern methods provided with pressure sensors to detect nasal air flow, may bring new data for the understanding of the physiopathology of these diseases, and this is what we intend to do in a continuation of the present investigation.

REFERENCES

1. Ali NJ, Pitson DJ, Stradling JR. Snoring, sleep disturbance and behaviour in 4-5 year olds. Arch Dis Child 1993;68:360-6.
2. Gislason T, Benediktsdottir B. Snoring, apnic episodes, and nocturnal hypoxemia among children 6 months to 6 years old. Chest 1995;107:563-6.
3. Guilleminault C, Eldrige FL, Simmons FB, Dement WC. Sleep apnea in eight children. Pediatrics 1976;58(1):23-30.
4. Carroll JL, Loughlin GM. In Ferber R, Kryger M. Principles and Practice of sleep medicine in the child. WB Saunders Company; 1995:163-91.
5. Guilleminault C. In: Guilleminault C. Sleep and its disorders in children. New York: Raven Press; 1987:195.
6. Bar A, Tarasiuk A, Segev Y, Phillip M, Tal A. The effect of adenotonsillectomy on serum insulin-like growth factor-I and growth in children with obstructive sleep apnea syndrome. J Pediatr 1999;135(1):76-80.
7. Grunstein RR, Handelsman DJ, Lawrence SJ, Blackwell C, Caterson ID, Sullivan CE. Neuroendocrine dysfunction in sleep apnea: reversal by continuous positive airways pressure therapy. J Clin Endocrinol Metab 1989;68:352-8.
8. Niiminen P, Lopponen T, Tolenon U, Lanning P, Knip M, Lopponen H. Growth and biochemical markers of growth in children with snoring and obstructive sleep apnea. Pediatrics 2002;109(4):55.
9. Williams EF, Woo P, Miller R, Kellman RM. The effects of adenotonsillectomy on growth in young children. Otolaryngol Head Neck Surg 1991;104:509-16.
10. Akhyozi-Rastad J, Hultcrantz E, Melander H, Svahnholm H. Body growth in relation to tonsillar enlargement and tonsillectomy. Int J Pediatr Otorhinolaryngol 1992;24:55-61.
11. Stradling JR, Thomas G, Warley HRA, Williams P, Freeland A. Effect
of adenotonsillectomy on nocturnal hypoaemia, sleep disturbance, and symptoms in snoring children. Lancet 1990;335:249-53.
12. Marcus CL, Carroll JL, Koerner CB, Hamer A, Lutz J, Loughlin GM. Determinants of growth in children with obstructive sleep apnea syndrome. J Pediatr 1994;125:556-62.
13. Marcus CL, Omlin KJ, Basinski DJ, Bailey SL, Rachal AB, Von Pechmann WS, Keens TG, Ward SL. Normal polysomnographic values for children and adolescents. Am Rev Respir Dis 1992;146 (5 Pt1):1235-9.
14. Marcus CL, Hamer A, Loughlin GM. Natural history of primary snoring in children. Pediat Pulmonol 1998;26 (1):6-11.
15. Richardson MA, Cotton RT. Anatomic abnormalities of pediatric airway. Pediatr Clin North Am 1984;31(4):821-34.
16. Rupa V, Raman R. Aetiological profile of paediatric laryngeal stridor in an Indian hospital. Ann Trop Paediatr 1991;11(2):137-41.
17. Lane RW, Wieder DJ, Steinem C, Marin-Padilla M. Laryngomalacia: a review and case report of surgical treatment with resolution of pectus excavatum. Arch Otolaryngol 1984;110(8):546-51.
18. Zalzal GH, Anon JB, Cotton RT. Epiglottoplasty for the treatment of laryngomalacia. Ann Otol Rhinol Laryngol 1987;96:72-6.
19. Belmont JR, Grundfast K. Congenital laryngeal stridor (laryngomalacia): etiologic factors and associated disorders. Ann Otol Rhinol Laryngol 1984;93:430-7.
20. Schulman JB, Hollister DW, Thibeault DW, Krugman ME. Familial laryngomalacia: a case report. Laryngoscope 1976;86(1):84-91.
21. McSwiney PF, Cavanagh NP, Langoth P. Outcome in congenital stridor (laryngomalacia). Arch Dis Child 1977;52:215-8.
22. Sutherland GA, Lack HL. Congenital laryngeal obstruction. Lancet 1897;2:653-5.
23. Prescott GA. The current status of corrective surgery for laryngomalacia. Am J Otolaryngol 1991;12(4):230-5.
24. Thompson J, Turner A. On the causation of congenital laryngeal stridor of infants. Br Med J 1900;2:1561-3.
25. Peron DL, Graffino DB, Zenker DO. The redundant aryepiglottic fold: report of a new cause of stridor. Laryngoscope 1988;98(6 Pt1):659-63.
26. Wiggs WJ, DiNardo LJ. Acquired laryngomalacia: resolution after neurologic recovery. Otolaryngol Head Neck Surg 1995;112(6):773-6.
27. Apley J. The infant with stridor: a follow-up survey of 80 cases. Arch Dis Child 1953;28:423-35.
28. Belmont JR, Grundfast K. Congenital laryngeal stridor (laryngomalacia): etiologic factors and associated disorders. Ann Otol Rhinol Laryngol 1984;93(5 Pt1):430-7.
29. Milczuk HA, Johnson SM. Effect on families and caregivers of caring for a child with laryngomalacia. Ann Otol Rhinol Laryngol 2000;109(4):546-54.
30. Kaemingk KL, Pasvogel AE, Goodwin JL, Mulvaney SA, Martinez F, Enright PL, Rosen GM, Morgan WJ, Fregosi RF, Quan SF. Learning in children and sleep disordered breathing: findings of the Tucson Children’s Assessment of Sleep Apnea (tUCASA) prospective cohort study. J Int Neuropsychol Soc 2003;9(7):1016-26.
31. Archbold KH, Giordani B, Ruzicka DL, Chervin RD. Cognitive executive dysfunction in children with mild sleep-disordered breathing. Biol Res Nurs 2004;5(3):168-76.