Case Report

Chronic Upper Airway Obstruction as a Main Cause of Pulmonary Hypertension

Samah Alasrawi1* and Lemis Yavus1

1Specialist Pediatric Cardiologist, Al Jalila Children Heart Center, Dubai UAE

Abstract

Introduction
The obstruction of the superior airways in childhood turns from several causes, among them the commonest is the adenotonsillar hypertrophy [1,2]. This chronic obstruction frequently is associated by syndrome of the apnea of the sleep and, in the most serious cases, they evolve with pulmonary hypertension [3]. The pulmonary hypertension is an indisputable indication for realization of adenotonsillectomy [4,5]. Because that we describe a case of severe pulmonary hypertension secondary of the adenotonsillar hypertrophy in the cardiology department of a university hospital.

Objective
To present a case of pulmonary hypertension secondary of the adenotonsillar hypertrophy in Cardiology department of a university hospital.

Case Report
3.5 years old girl admitted with respiratory difficulty, coughing, nocturnal snores and mouth breath. Cardiomegaly was found in chest XR. The Doppler echocardiogram study showed pulmonar arterial hypertension. The lateral X ray films showed important adenoid hypertrophy. Adenotonsillectomy was carried without problems. After six months, the patient did a new Doppler echocardiogram that showed normal, showing clear reversion of all alterations shown in the first echocardiogram.

Conclusion
Patients with adenotonsillar hypertrophy and chronic upper airway obstruction, may present with evaluated a pulmonary hypertension. The Doppler echocardiogram is safest exam and little invasive for determination of the diagnosis [5,6] and the adenotonsillectomy is the definitive treatment

Keywords: Adenoid Hypertrophy; Adenotonsillar Hypertrophy; Pulmonary Hypertension; Upper Airway Obstruction

Introduction
The obstruction of the superior airways in childhood turns from several causes, among them the commonest is the adenotonsillar hypertrophy [1,2]. This chronic obstruction frequently is associated by syndrome of the apnea of the sleep and, in the most serious cases; they evolve with pulmonary hypertension [3]. The pulmonary hypertension is an indisputable indication for realization of adenotonsillectomy [4,5]. Because that we describe a case of severe pulmonary hypertension secondary of the adenotonsillar hypertrophy in the cardiology department of a university hospital.

Case Report
3.5 years old girl admitted to the hospital with cough, history of recurrent chest infections from 2 years (4 to 5 times a year) treated by oral antibiotics without hospitalization, breathing difficulty with any exercise

No history of cyanosis or syncope, Failure to thrive (WT=12kg <3%, HT=90cm <3%), History of snoring and mouth breathing

On physical examination: Loud S2, soft systolic murmur II/VI, chest: Harsh crackles bilateral

Hypertrophied tonsils, other systemic examination was normal

Labs: WBC: 6200 (N 33%, L 66%), Hgb 11g/dl, CRP negative

Electrolytes, liver function, renal function and Immune globulins were normal.

Blood gases: During the day: PH=7.41, PCO2=34, PO2=72, Hco3=25, So2=96%

During sleep: PH = 7.35, PCO2=42, PO2 =60, Hco3=26, So2=94%

*Corresponding Author: Samah Alasrawi, Specialist Pediatric Cardiologist, Al Jalila Children Heart center, Dubai, UAE, Tel: 00971561387382; E-mail: samahisrawi@gmail.com

Sub Date: January 29th 2018, Acc Date: February 5th 2018, Pub Date: February 6th 2018,

Citation: Samah Alasrawi and Lemis Yavus (2018) Chronic Upper Airway Obstruction as a Main Cause of Pulmonary Hypertension. BAOJ Pediat 4: 52.

Copyright: © 2018 Samah Alasrawi , et al. This is an open access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.
Citation: Samah Alasrawi and Lemis Yavus (2018) Chronic Upper Airway Obstruction as a Main Cause of Pulmonary Hypertension. BAOJ Pediat 4: 52.

Chest XR: Cardiomegaly, increased pulmonary vascularity bilateral
(figure 1)

![Chest XR Image](image1.png)

Figure 1: chest XR: Cardiomegaly, increased pulmonary vascularity bilateral

ECG: signs of dilated right atrium, dilated right ventricle (Figure 2)

![ECG Image](image2.png)

Figure 2: ECG: signs of dilated right heart.

ECHO cardiogram: dilated right atrium, dilated right ventricle, moderate to severe tricuspid regurgitation, RVSP=80mmHg, no structure abnormality was seen.

Conclusion

Pulmonary hypertension, normal cardiac anatomy, (figure3)

So we started to search for extra cardiac reason for the pulmonary hypertension, we prescribed Sildenafil but the parents didn't start it.

![ECHO Image](image3.png)

Figure 3: Echo: pulmonary hypertension.
Chest CT scan: cardiomegaly, dilated main pulmonary artery and its branches, lungs about normal (figure 4)

Lateral skull XR showed Large adenoids in the nasal pharynx and narrow air way (figure 5)

After surgery the girl became better, no more snoring or breathing difficulty and began to gain weight

Scull XR after surgery (wide upper air way)

Chest XR after 3 month of surgery: mild cardiomegaly (figure 7)
Discussion

The symptoms caused by adenotonsillar hypertrophy can be nasal obstruction; mouth breathing, snoring, sleep apnea, irritability, and low development at school [5,6]. Complications of such obstruction cover from alterations on craniofacial development to pulmonary hypertension and Cor Pulmonale [1,3,4]. The current case already presented pulmonary involvement seen in the clinical history, assured by initial complaints.

Although adenotonsillar hypertrophy is very common, secondary pulmonary hypertension to it has been hardly diagnosed due to great amount of adenotonsillectomy approach performed in the past [4]. Doppler echocardiogram is highly useful when diagnosing pulmonary hypertension in infants with adenotonsillar hypertrophy, as it is considered a very safe, practical and non-invasive exam [5,6]. That exam was very important when diagnosing right heart involvement of the current patient. Chronic obstruction of the upper airways followed by pulmonary hypertension and Cor Pulmonale is an unquestionable sign of adenotonsillectomy, once it is already known that clinical and echocardiographic findings are fully reversible by surgery [3,6]. The current child was clinically benefited by adenotonsillectomy and few months later there was Doppler echocardiography in normal condition.

Final Comments

Adenotonsillar hypertrophy is a significant cause of pulmonary hypertension in children’s.

Echocardiogram is the best exam to diagnose pulmonary hypertension for being safe and non-invasive. We suggest this exam becomes part of the investigations on children with adenotonsillar hypertrophy associated to sleep apnea [5,6].

Children with symptomatic adenotonsillar hypertrophy adenotonsillectomy required, to prevent pulmonary hypertension as complication of chronic obstruction by an enlargement of adenotonsil tissue [7,8].

References


