ABSTRACT

Background: Asthma is an important childhood disease. Recent surveys of the International Study of Asthma and Allergies in Childhood (ISAAC) suggest that the prevalence of asthma is increasing but these surveys do not include any pulmonary tests to confirm the possible diagnosis of asthma.

Objective: To compare bronchodilator reversibility with the albuterol test in symptomatic and asymptomatic 6-7-year-old children with asthma participating in the ISAAC survey and living in Mexico City.

Patients and methods: We performed an observational, descriptive, comparative, cross sectional study in children participating in phase 3b of the ISAAC study. According to the ISAAC questionnaire children were classified as asthma symptomatic or asymptomatic. Both groups had bronchodilator reversibility with the albuterol test, using the guidelines of the American Thoracic Society to confirm or rule out the diagnosis of asthma.

Results: The asymptomatic group had a baseline FEV1 of 1.70 ± 0.34 l/sec (mean ± SD) and an endpoint FEV1 of 1.76 ± 0.42 l/sec; in the symptomatic group the respective values were 1.51 ± 0.41 l/sec and 1.57 ± 0.44 l/sec (p < 0.05). A positive reversibility test was found in 13/136 (9.6%) children in the asymptomatic group and in 22/112 (19.6%) children in the symptomatic group (p < 0.05).

Conclusion: Because of its low sensitivity, bronchodilator reversibility cannot be considered a diagnostic tool to confirm diagnosis of asthma.

Key words: Bronchodilator reversibility. Asthma. ISAAC. Survey.

RESUMEN

Introducción: El asma es una enfermedad infantil importante. Las encuestas recientes del Estudio Internacional sobre Asma y Alergias en la Infancia (ISAAC, por sus siglas en inglés) sugieren un aumento de la prevalencia del asma, si bien este instrumento de estudio no incluye ninguna prueba pulmonar para confirmar el posible diagnóstico de la enfermedad.

Objetivo: Comparar la reversibilidad con tratamiento broncodilatador a base de albuterol entre niños asintomáticos sintomáticos y asintomáticos según el estudio ISAAC, de edades comprendidas entre los 6 y 7 años y residentes en Ciudad de México.

Pacientes y métodos: Se trató de un ensayo comparativo transversal, de carácter descriptivo y basta-
INTRODUCTION

Asthma is one of the most frequent chronic diseases in childhood and it is an important cause of functional limitation and school absenteeism. The worldwide information generated from 1970, suggests an increasing trend, not only in frequency, but also in severity of the disease. There was a rise of the number of hospitalizations and mortality, between 1979 and 1997, the absolute number of dead increased dramatically from 2603 to 5434. In the last decade it has been observed an increase on asthma prevalence in diverse countries, in spite of a better understanding of its pathogenesis and a better treatment for asma. Ambos grupos fueron sometidos a la prueba de reversibilidad con tratamiento broncodilatador a base de albuterol, siguiendo las directrices de la Sociedad Torácica Americana, para confirmar o descartar el diagnóstico de asma.

Results: The group symptomatic presented an FEV1 inicial de 1,70 ± 0,34 l/seg (media ± DE) y un FEV1 final de 1,76 ± 0,42 l/seg; en el grupo sintomático, los valores respectivos fueron 1,51 ± 0,41 l/seg y 1,57 ± 0,44 l/seg (p < 0,05). En el grupo asintomático, 13 de 196 niños (el 9,6 %) dieron positivo en la prueba de reversibilidad; en el grupo sintomático dieron positivo 22 de 112 niños (el 19,6 %) (p < 0,05).

Conclusión: La reversibilidad con tratamiento broncodilatador no puede considerarse una herramienta para confirmar el diagnóstico del asma debido a su bajo nivel de precisión.

Palabras clave: Asma. Encuesta ISAAC. Reversibilidad con tratamiento broncodilatador.
A clinical history with vital signs and anthropometric measures of weight and height was completed. The weight (kg) was obtained by a Health o Meter inc platform scale (Bridgeview, Illinois, USA), and the height (cm) was obtained using a Holtain Limited Crymych, Difec stadiometer (made in Britain) registering the results in centimeters (cm). The platform scale was calibrated daily to zero. Height and weight were assessed without shoes and with the children standing with the legs together, touching the wall, their legs, buttocks and scapulas aligned. Their head was positioned maintaining their seeing from the front, their eyes in horizontal position, related with ears. The age was determined according the born date and it was expressing in decimal fractions.

All procedures were realized in the Allergy department, in a room 5 × 5 meters, with a room temperature of 20 °C. Patients performed three spirometries and the best one was selected as the baseline, then patients inhaled 100 mcg of albuterol twice with metered dose inhalator with an aero chamber spacer (made in London Ontario Canada), afterward patients performed three spirometries and the best one was selected.

For this study Easy ware, 2001, spirometers (made in Zurich, Switzerland) were used; they complied with the quality requirements of American Thoracic Society (ATS). The daily calibration was realized with a syringe of 3 L flow-volume, with daily review of the quality control. The temperature of the spirometer before each study and the barometric pressure of Mexico City (585 mmHg) were entered to the spirometer. The physicians were trained specifically for this project, using the guidelines of American Thoracic Society (ATS).

Patients followed the standard procedures suggested by ATS, adding some specifications as not inhaling since the spirometers, only register the expiratory phase of the forced expiratory procedure, using disposable mouthpieces, and bacterial filters. Before patients perform spirometry investigators explained the purpose of the test and showed the correct procedure. In general, patients needed less than 6 maneuvers to get 3 useful curves. When children made a double inspiration, they repeated the procedure. FEV1 and FVC were considered reproducible according guidelines of ATS when the best of 3 spirometric curves varied no more than 200 ml or 5 %.

The size of the sample was 100 children, based in a study of Laprise and the ISAAC requirements. We calculated mean and standard deviation for parametric variables, and frequencies for reversibility rates. One way ANOVA was used for comparison between the groups considering baseline and endpoint spirometric values.

<table>
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<th>Table I</th>
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<td>ISAAC Questionnaire for 6/7 years children</td>
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1. Has your child ever had wheezing or whistling in the chest at any time in the past?
2. Has your child had wheezing or whistling in the chest in the past 12 months?
3. How many attacks of wheezing has your child had in the past 12 months?
4. In the past 12 months, how often, on average, has your child sleep been disturbed due to wheezing?
5. In the past 12 months, has wheezing ever been severe enough to limit your child speech to only one or two words at a time between breaths?
6. Have your child ever had asthma?
7. In the past 12 months, has your child's chest sounded wheezy during or after exercise?
8. In the past 12 months, has your child had a dry cough at night, apart from a cough associated with a cold or chest infection?

Children with positive answer to questions 1 and 2 of the ISAAC questionnaire were considered as the symptomatic group and those with negative answers were regarded as the asymptomatic group.

Participants should not have respiratory tract infection in the last four weeks and they should not use beta agonist, bronchodilators, asthma preventive medication (sodium chromoglycate, inhaled steroids or antileukotriene drugs), or antihistaminics in the last 45 days. Patients suffering from other pulmonary disease, cardiovascular disorders, endocrine diseases or somatodysmorphic syndromes were not included in the study. Patient parents must sign written informed consent before any study procedures. Children not able to perform spirometric or peak flow procedures were excluded.

The study protocol was approved by the Investigation and Ethics Committee of the Hospital Infantil de Mexico “Federico Gómez” and the Public Education Department of Mexico City.
RESULTS

In this study 248 children between 6 and 7 years were included, 112 children had a questionnaire suggestive of asthma (symptomatic group) and 136 had negative answers to the questionnaire questions (asymptomatic group). 248 spirometries sets were performed from September 2002 to January 2003. Both groups were similar regarding sex distribution, weight and height (table II).

In the asymptomatic group the baseline FEV1 was $1.70 \pm 0.34$ l/seg (mean ± SD) and the endpoint FEV1 was $1.76 \pm 0.42$ l/seg; in the symptomatic group the respective values were $1.51 \pm 0.41$ l/seg and $1.57 \pm 0.44$ l/seg. One way ANOVA showed a significant difference between the groups ($p < 0.05$). In the asymptomatic group 13 out of 136 (9.6 %) children had a positive reversibility test, while in the symptomatic group 22 out of 112 (19.6 %) children had a positive test ($p < 0.019$) (table III). There were no difference in the reversibility rates between boys and girls in the symptomatic group ($p > 0.05$) (table IV).

The sensibility and specificity of the reversibility test were 19 % and 90 % respectively, with a positive predictive value of 62 % and negative predictive value of 57 %.

DISCUSSION

Currently, there is great interest about prevalence of asthma worldwide. Because this, ISAAC established 3 phases of study since 1970. The first phase investigate the actual and accumulated prevalence by means of a questionnaire, second phase represents the search of the etiology by means of diagnosis testing, and the third phase is as the first, 5 years latter. When a specific center realizes the third phase without the previous ones, it is considered as a 3b phase. In phase 1 and 3 standardized and validated questionnaires with the official language of the center are used to determine the prevalence on age specific groups. (Children 6 to 7 years)\(^4\).

In the population studied in the North area of Mexico City, we observed an accumulated prevalence of asthma (considered as the presence of wheezing) of 19 %, asthma diagnosis prevalence of 4.5 % and current prevalence of asthma (wheezing in the last 12 months) of 6.8 % (unpublished data).

Although we could not consider the present study as a second phase of ISAAC, we pretended to evaluate an easy and quickly method of bronchial reversibility in children with questionnaire suggestive of asthma.

### Table II

<table>
<thead>
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<th>Characteristics of asthma symptomatic and asymptomatic children according to ISAAC questionnaire</th>
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<tr>
<td>Symptomatic</td>
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<tr>
<td>N</td>
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<tr>
<td>Weight (kg)</td>
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<td>Height (cm)</td>
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### Table III

<table>
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<th>Bronchodilator reversibility in asthma symptomatic and asymptomatic children according to ISAAC questionnaire</th>
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<tr>
<td>Reversibility test</td>
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<tr>
<td>Negative n (%)</td>
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<td>Positive n (%)</td>
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$p < 0.02$ by Fisher exact test.

### Table IV

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<th>Comparison of the bronchodilator reversibility according a gender in the symptomatic group according to ISAAC questionnaire</th>
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<tbody>
<tr>
<td>Reversibility test</td>
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<tr>
<td>Negative n (%)</td>
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<tr>
<td>Positive n (%)</td>
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$p = 0.05$ by Fisher exact test.

The validation of the asthma questionnaires has demonstrated that the written questionnaire has a negative predictive value and positive predictive value of 61 % and 94 % respectively, and a sensibility of 85 % and specificity of 81 %. In contrast, in the video questionnaire sensibility is close to 90 % and specificity to 68 %,\(^2\)-\(^4\).

Eventhough, surveys based on questionnaires have some disadvantages, they provide guide to elaborate hypothesis. Bronchial hyperresponsiveness is a mechanism implied in the asthma physiopathology and the bronchial reversibility test is the method to corroborate the presence of hyperresponsiveness and therefore of asthma.
Other two test are used to confirm asthma diagnosis: the provocation choline challenge has a specificity and sensitivity of 89 % with a negative predictive value of 94 % and a positive predictive value of 81 % and it is used to discard asthma; the hypertonic saline challenge has a sensitivity of 72 % and specificity close to 100 % and it is used to assess big populations as it is easy and inexpensive.

There are some studies with ISAAC’s methodology which have assessed bronchial hyperresponsiveness. The most of them have used exercise challenge, some provochoine and few hypertonic saline. These differences may be ascribed to the distinct sensibility of the tests themselves or to the absence of inflammatory process in these subjects.

In Chinese studies using provochoine challenge test in 10 year children, the subjects with asthma diagnosis in ISAAC questionnaire had a rate of 8.3 % of positive exercise challenge while the control group had a positive rate of 5.3 %.

In the present study the reversibility test had a lower positive test rate than the challenge test with provochoine and hypertonic saline. These differences may be ascribed to the distinct sensibility of the tests themselves or to the absence of inflammatory process in our subjects at the moment of the reversibility test.

In the studies using hypertonic saline challenge the subjects with a positive questionnaire had a positive rate of 40 % versus 9 % in the subjects with negative diagnosis.

In the studies using hypertonic saline challenge the subjects with a positive questionnaire had a positive test rate from 50 to 60 % while the subjects with negative questionnaire had a positive test rate of 12.8 %.

In the present study the reversibility test had a lower positive test rate than the challenge test with provochoine and hypertonic saline. These differences may be ascribed to the distinct sensibility of the tests themselves or to the absence of inflammatory process in our subjects at the moment of the reversibility test.

It is very important to stress that BHR can be an intermediate state between asthma and normality. Bronchial hyperresponsiveness is a characteristic finding in patients with asthma, however, when a patient is under control it is difficult to probe. The use of beta agonist is the easiest form to demonstrate BHR, but because its low sensitivity and specificity, it is not recommended in big populations as a detection procedure, neither to complement the ISAAC questionnaire.

CONCLUSIONS

In our conditions bronchial reversibility test may not be considered as a valuable diagnostic tool to complement the ISAAC questionnaire and to determine the asthma prevalence because of its low sensitivity and relative specificity.

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