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Risk factors for wheezing in a subtropical environment: Role of respiratory viruses and allergen sensitization

Ataide A. Camara, MD,a,b Jogete M. Silva, MD,a Virginia P. L. Ferriani, MD,a Kátia R. C. Tobias, MSc,a Izolète S. Macedo, MSc,b Márcio A. Padovani, MD,c Charlotte M. Harsi, PhD,d M. Regina A. Cardoso, PhD,e Martin D. Chapman, PhD,f Eurico Arruda, MD,e Thomas A. E. Platts-Mills, MD, PhD,g and L. Karla Arruda, MDa Ribeirão Preto and São Paulo, Brazil, and Charlottesville, Va

Background: Risk factors for acute wheezing among children in subtropical areas are largely unknown.

Objective: To investigate the role of viral infections, allergen sensitization, and exposure to indoor allergens as risk factors for acute wheezing in children 0 to 12 years old.

Methods: One hundred thirty-two children 0 to 12 years of age who sought emergency department care for wheezing and 65 children with no history of wheezing were enrolled in this case-control study. Detection of respiratory syncytial virus antigen, rhinovirus and coronavirus RNA, adenovirus, influenza, and parainfluenza antigens was performed in nasal washes. Total IgE and specific IgE to mites, cockroach, cat, and dog were measured with the CAP system. Major allergens from mites, cockroach, cat, and dog were quantified in dust samples by ELISA. Univariate and multivariate analyses were performed by logistic regression.

Results: In children under 2 years of age, infection with respiratory viruses and family history of allergy were independently associated with wheezing (odds ratio, 15.5 and 4.2; P = .0001 and P = .008, respectively). Among children 2 to 12 years old, sensitization to inhalant allergens was the major risk factor for wheezing (odds ratio, 2.7; P = .03). High-level allergen exposure, exposure to tobacco smoke, and lack of breast-feeding showed no association with wheezing.

Conclusions: Some risk factors for wheezing previously identified in temperate climates were present in a subtropical area, including respiratory syncytial virus infection in infants and allergy in children older than 2 years. Rhinovirus was not associated with wheezing and did not appear to be a trigger for asthma exacerbations. (J Allergy Clin Immunol 2004;113:551-7.)

Key words: Wheezing, asthma, allergy, mites, cockroach, respiratory viruses, respiratory syncytial virus, rhinovirus, allergens

Asthma is the most common chronic disease of childhood. Evidence indicates that more than half of the cases of persistent asthma start before the age of 3 years, and it is now clear that airway inflammation and irreversible remodeling may be initiated early in life.1,2 Definitive diagnosis of asthma and subsequent decision to initiate long-term anti-inflammatory treatment at an early age has been a challenge in pediatric practice.3 Although asthma in young children appears to represent a heterogeneous disease with distinct phenotypes, sensitization to common inhalant allergens including those derived from dust mites, cockroach, cat, and Alternaria remains a major risk factor for persistent asthma associated with airway hyperreactivity and decreased lung function.4,6

Respiratory viruses have been recognized as major triggers of wheezing episodes and asthma exacerbation.7 In the northern hemisphere, respiratory syncytial virus (RSV) accounts for the majority of wheezing episodes in children younger than 2 years of age, being detected in 60% to 80% of infants.7 Results of the Tucson Children’s Respiratory Study8 revealed that lower respiratory tract infection with RSV early in life was associated with persistent wheezing; however, by age 13 years, lung function abnormalities were no longer present and wheezing resolved in most children. In one study, severe RSV bronchiolitis in the first year of life has been linked to asthma and allergic sensitization at age 7 years;9 however, studies to determine whether RSV infections enhance allergy have provided conflicting results.7 It has been definitely established that specific IgE response to RSV is not an important element in the pathogenesis of RSV-induced wheezing;10 however, the virus has been shown
to induce expression of cytokines and chemokines and activation of monocytes, macrophages, and lymphocytes, which may influence the mechanisms involved in the development of asthma in children.7 In older children, rhinovirus becomes the dominant viral pathogen linked to wheezing exacerbations. This has been demonstrated in community-based and emergency department studies.11,12 In the emergency department setting, rhinovirus infections act synergistically with allergen sensitization and/or allergen exposure to induce severe asthma symptoms in both children and adults.12,13 In keeping with this, lower respiratory tract symptoms and markers of inflammation in young adults with mild asthma experimentally infected with rhinovirus are more pronounced in those with high total IgE.14 It is thought that a deficient Th1-type response with decreased IFN-γ production underlies the observation that rhinovirus infections in atopic individuals are more severe and longer-lasting and induce more lower respiratory symptoms as compared with nonatopic subjects.15

The purpose of the current study was to investigate the role of viral infections, allergen sensitization, and exposure to indoor allergens as risk factors for acute wheezing in children 0 to 12 years old living in a subtropical environment. The results of the current study may contribute to promote better care of children with wheezing in these areas.

**METHODS**

Subjects

One hundred thirty-two children 0 to 12 years of age (80 boys) who sought emergency department (ED) care for wheezing either at the Clinical Hospital of the University of São Paulo School of Medicine of Ribeirão Preto or Santa Lydia community Hospital were enrolled in this case-control study. Patients were selected for the study if they presented with wheezing that required therapy with inhaled β2-agonists as judged by the attending physician. The study protocol was approved by the ethics committee of both hospitals and the children’s parents and/or guardians gave written informed consent to participate in the study.

Parents or guardians completed a questionnaire that included questions on personal and family history of allergies and asthma, passive exposure to tobacco smoke, breast-feeding, and housing conditions. A positive personal history of allergy was defined as a report of physician-diagnosed asthma, rhinitis, and/or atopic dermatitis. A positive family history for allergy was characterized by the report of asthma, rhinitis, and/or atopic dermatitis diagnosed by a physician in the mother, father, and/or sibling(s). History of prior wheezing illnesses was reported for 48% of the wheezing children younger than 2 years and for 93% of the older wheezing children. However, only 16% of the wheezing infants and 51% of the wheezing children older than 2 years reported asthma diagnosed by a physician. The study protocol was approved by the ethics committee of both hospitals and the children’s parents and/or guardians gave written informed consent to participate in the study.

Detection of respiratory viruses

Nasal washings were collected and processed for detection of respiratory viruses as previously described.15 RSV antigen was detected by using both a rapid enzyme immunoassay (Test-Pack, Abbott, Chicago, Ill) and an indirect immunofluorescence (Chemicon, Temecula, Calif). Human rhinovirus and coronavirus RNA was detected by RT-PCR.12,16 Influenza and parainfluenza viruses were detected by indirect immunofluorescence.17,18 In a subset consisting of 90 of 132 wheezing children and 48 of 65 control subjects, from whom remaining material was available, detection of adenovirus was investigated by culture of frozen nasal aspirates on A549 cells, with cytopathic effect confirmed by immunofluorescence. Typing of adenovirus isolated from cultures to the subgenus level was performed by amplification of the VA-RNA region of the viral genome by PCR, according to previously described method.19 PCR products of 520 to 522 bp and 505 bp in length corresponded to adenovirus subgenera B and C, respectively. Certain adenovirus serotypes including those of subgenus C can be identified in nasal aspirates up to several weeks after acute infection.20 To minimize the interference of latent adenovirus, only subgenus B was included in the analysis.

Total IgE, specific IgE antibodies, and peripheral blood eosinophils

Serum levels of total IgE and specific IgE to mites (Dermatophagoides pteronyssinus, D farinae, and Blomia tropicalis), cockroach (Blattella germanica), cat, dog, foods (cow’s milk, egg, wheat, soy, peanut, and fish), and to Ascaris lumbricoides were measured with the use of the UniCAP system (Pharmacia, Uppsala, Sweden). Specific IgE levels ≥0.7 kU/L (CAP score ≥2) were considered positive. Automated white blood cell counts were carried out with the use of a Colter T540 system, and differential counts, including eosinophil quantification, were performed manually.

Allergen levels in house dust samples

Dust samples were collected from four sites in each subject’s home: bedroom, living room, TV room, and kitchen, within 3 weeks of the ED visit. Dust collection and preparation of dust extracts were performed according to standard procedures.21 Measurements of major allergens from mites (group 1 and group 2), cockroach (Bla g 1 and Bla g 2), cat, (Fel d 1), and dog (Can f 1) in dust extracts were carried out with the use of monoclonal antibody–based ELISA.22 Group 1 mite allergens comprised the sum of levels of Der p 1 and Der f 1. Group 2 allergens (Der p 2 and Der f 2) were simultaneously detected in the monoclonal antibody ELISA with a cross-reactive capture antibody (1D8). High-level exposure was defined as the presence of at least one dust sample in the house containing concentrations of group 1 mite allergens (≥10 µg/g, Bla g 1 ≥8 U/g, and/or Bla g 2 ≥0.32 µg/g, Fel d 1 ≥8 µg/g, and Can f 1 ≥10 µg/g).22
**Statistical analysis**

Data from wheezing and control children were compared in two age groups: children younger than 2 years of age (n = 74 and n = 30, respectively) and children 2 to 12 years of age (n = 58 and n = 35, respectively). Data on total IgE levels were log-transformed and analyzed by Student t test. Peripheral blood eosinophils and allergen levels in house dust were analyzed by Mann-Whitney test. Chi-square analysis was used to compare the prevalence of elevated specific IgE, positive viral tests, frequency of passive smoke exposure, and high-level allergen exposure. Univariate and multivariate analysis were carried out by logistic regression, using the software STATA version 6.0.

**RESULTS**

**Detection of respiratory viruses**

In the group of children under 2 years of age, respiratory viruses were detected in 60.8% (45 of 74) wheezing infants and in 13.3% (4 of 30) control infants (P < .001). RSV was detected in 39% wheezing children and none of the control infants (P < .001). Rhinovirus RNA was found in 20.2% and 10% of the wheezing and control children, respectively (P = .21). Adenovirus was cultured from 29.4% (15/51) of wheezing children and 20.8% (5/24) of control children; 4 and 1 isolates were identified as subgenus B (7.8% and 4.1%), respectively (P = .55). Coronavirus RNA was found in 3 of 73 wheezing subjects and none of 30 control subjects. Influenza A and B and parainfluenza viruses were not detected (Fig 1). Five wheezing children were positive for more than one virus.

In the group of children 2 to 12 years of age, respiratory viruses were not significantly associated with wheezing. Rhinovirus RNA was detected in 19.3% wheezing children as compared with 17.1% of the non-wheezing control children (P = .80); RSV antigen was detected in 10.3% and 8.5%, respectively (P = .70), and adenovirus was isolated from 23% (9/39) and 17.8% (5/28), respectively; 4 and 0 (10.5% and 0%) of the adenovirus isolates were identified as subgenus B (P = .07). Coronavirus, influenza A and B, and parainfluenza viruses were not found in children older than 2 years (Fig 1).

**Relation of wheezing, IgE response, and eosinophilia**

In children under 2 years of age, geometric mean (GM) total IgE levels were 19.7 IU/mL (range < 2 IU/mL to 1709 IU/mL) in wheezing infants and 9.4 IU/mL (range < 2 IU/mL to 620 IU/mL) in control infants (P = .09). Only four wheezing infants, all over 1 year old, presented IgE antibodies to inhalant allergens: three to mites and one to mites, dog, and egg; three of these had a reported history of allergy in both mother and father. Both wheezing and control infants presented IgE to foods: 8.9% (6 of 67) and 11.1% (3 of 27) were sensitized to at least one food allergen, respectively (P = .74). The most common allergens were egg and cow's milk, followed by wheat and soy. No sensitization to peanut or fish was found. IgE to *Ascaris lumbricoides* was not detectable in children under 2 years of age. No signifi-
cant differences in peripheral blood eosinophil counts were observed in the wheezing and control groups (GM = 218/mm³ and 208/mm³, respectively) (P = .21).

In children older than 2 years, GM value of total IgE was 278 IU/mL (range, 7.9 to 2627 IU/mL) in wheezing children and 97.4 IU/mL (range, 2.0 to 1067 IU/mL) in control children (P = .002). Sensitization to inhalant allergens was present in 72.4% of wheezing children and 42.8% of control children (P = .005). Mites and cockroach were the sensitizing allergens associated with wheezing. Specific IgE to A. lumbricoides was more frequently found in wheezing children, 33.9% (19 of 56) as compared with 14.2% (5 of 35) of the control children (P = .04). Peripheral blood eosinophil counts were higher but did not reach significance (GM = 350/mm³ and 199/mm³ in wheezing and control children, respectively) (P = .08).

Role of allergen exposure

Concentrations of mite, cockroach, cat, and dog allergens in dust samples of homes of wheezing children and control children were compared (Table I). Levels of group 2 mite allergens were significantly higher in homes of wheezing children under 2 years of age. Levels of group 1 mite allergens, Bla g 1 and Bla g 2, Fel d 1, and Can f 1, in the homes were not different between wheezing and control children for either age group (Table I). The highest levels of mite allergens were found in bedding samples.

Analysis of risk factors for wheezing

Infection with respiratory viruses, particularly with RSV, was the major risk factor for wheezing among children younger than 2 years of age, as revealed by univariate analysis. Personal and family history of allergy were both significantly associated with wheezing (Table II). In the multivariate model, viral infection and family history of allergy were each independently associated with wheezing, and sensitization to inhalant allergens played no significant role in wheezing in infants (Table III).

Among children 2 to 12 years old, allergy was the most important risk factor for wheezing, indicated by the higher frequency of positive tests for specific IgE to inhalant allergens and elevated total IgE levels (Table II). IgE to A. lumbricoides and personal history of allergy were both associated with wheezing (Table II). On the other hand, viral infection was not a significant risk factor for wheezing in this age group. Two models of multivariate analysis were generated by using levels of specific IgE antibodies ≥ 0.7 kUA/L or ≥ 3.5 kUA/L (considered as high level). In both models, sensitization to inhalant allergens was the major independent risk factor for wheezing (Table IV).

Current exposure to high levels of indoor allergens was not significantly associated with wheezing (Table I). Levels of group 1 mite allergens ≥10 µg/g of dust were found in approximately 70% of the homes of children older than 2 years and in 36% to 46% of the homes of children under 2 years, with no differences between wheezing and control groups. Approximately 25% of homes presented high levels of cockroach allergens in dust samples. In 13 of 172 (7.5%) homes, cats were present in the house. Levels of cat allergen were undetectable or very low in the majority of samples; however, Fel d 1 levels in homes with cats were higher than in those without a cat (GM = 1.06 µg/g and 0.19 µg/g, P < .0001). The presence of dogs was found in 87 of 172 (50.2%) homes visited, and Can f 1 levels were also higher in homes with dogs as compared with homes without dogs (GM = 6.4 µg/g and 0.5 µg/g, P < .0001). No synergistic effect was observed when the combination of sensitization and exposure to specific allergens was analyzed.
DISCUSSION

In the current study, we have identified for the first time risk factors associated with acute wheezing among children living in a subtropical area of South America. We have found that RSV infection was strongly associated with wheezing among children under 2 years of age.7,12 The frequency of RSV was lower than that reported in temperate regions (39% versus 60% to 80%).7 However, if we consider the subgroup of infants 0 to 6 months old, 61% tested positive for RSV antigen. RSV infections were predominantly found in the months of February to May, corresponding to late summer and early to mid-fall, indicating that the virus occurs in a different seasonal pattern as compared with that of the northern hemisphere. The available data indicate that the main age group affected by RSV in developing countries is children under 6 months of age and that in tropical or subtropical climates RSV outbreaks are frequently associated with the rainy season.23 Household and day care–based prospective studies in coastal northeast Brazil have identified RSV at an unexpectedly low frequency, 0% to 4%.24,25 A previous study in Ribeirão Preto, analyzing samples of 829 children under 5 years of age with nonspecific respiratory tract symptoms seen in the ED revealed that RSV was present in 21.6% of nasal washes. However, the vast majority of RSV infections (157/188 episodes, 83.5%) were found in children under 1 year of age.26

### TABLE II. Univariate analysis of risk factors for acute wheezing among children 0 to 12 years old

| Risk factor                              | Under 2 y (n = 74) | Control (n = 30) | Odds ratio (95% CI) | P value | 2 to 12 y (n = 58) | Control (n = 35) | Odds ratio (95% CI) | P value |
|------------------------------------------|--------------------|------------------|---------------------|---------|--------------------|------------------|---------------------|---------|
| Age in mo (median)                       | 6                  | 4                | 1.0 (0.9 to 1.0)    | .6      | 48                 | 60               | 0.9 (0.9 to 1.0)    | .2      |
| Male sex (%)                             | 62.1               | 56.6             | 0.8 (0.3 to 1.9)    | .6      | 58.6               | 62.8             | 1.1 (0.5 to 2.8)    | .6      |
| Personal history of allergy (%)          | 33.7               | 13.3             | 3.3 (1.04 to 10.5)  | .04     | 75.8               | 11.4             | 24 (7.3 to 81.0)    | <.001   |
| Family history of allergy (%)            | 72.9               | 46.6             | 3.0 (1.2 to 7.4)    | .01     | 65.5               | 62.8             | 1.1 (0.4 to 2.6)    | .79     |
| Total serum IgE ≥400 IU/mL (%)           | 4.1                | 3.4              | 1.2 (0.1 to 12.0)   | .8      | 44.8               | 22.8             | 2.7 (1.06 to 7.0)   | .03     |
| Sensitization to inhalant allergens (%)  | 5.4                | 0                | —                   | —       | 72.4               | 42.8             | 3.5 (1.4 to 8.4)    | .005    |
| Mite (%)                                 | 5.4                | 0                | —                   | —       | 68.9               | 40               | 3.3 (1.3 to 8.0)    | .007    |
| Cockroach (%)                            | 0                  | 0                | —                   | —       | 26.3               | 2.8              | 12.1 (1.5 to 96.6)  | .018    |
| Cat (%)                                  | 0                  | 0                | —                   | —       | 5.2                | 5.7              | 0.9 (0.1 to 5.7)    | .9      |
| Dog (%)                                  | 1.4                | 0                | —                   | —       | 7.0                | 0               | —                   | —       |
| Sensitization to food allergens (%)      | 8.9                | 11.1             | 0.7 (0.1 to 3.4)    | .7      | n.d.               | n.d.             | n.d.                | n.d.    |
| Specific IgE to Ascaris lumbricoides (%) | 0                  | 0                | —                   | —       | 33.9               | 14.2             | 3.0 (1.02 to 9.2)   | .04     |

*Levels of specific IgE antibodies ≥0.7 kUA/L (CAP class ≥2) to at least one inhalant allergen (mite, cockroach, cat, or dog) or food allergen (egg, milk, soy, wheat, fish, or peanut).

†Levels of IgE antibodies to A. lumbricoides ≥0.7 kUA/L (CAP class ≥2).

‡Detection of RSV antigen, rhinovirus RNA, adenoivirus B, and/or coronavirus RNA in nasal washings.

§High-level exposure to mites, cockroach, cat, and dog were defined as group 1 mite allergen levels ≥10 µg/g of dust; cockroach allergens Bla g 1 ≥8 U/g or Bla g 2 ≥32 µg/g of dust; cat allergen Fel d 1 ≥10 µg/g of dust; and dog allergen Can f 1 ≥10 µg/g of dust, respectively, at least in one site of the home.

n.d., Not done. IgE to food allergens was not assayed on children ages 2 to 12 years; —, odds ratios for variables with a zero value could not be calculated with STATA 6.0.

### TABLE III. Multivariate analysis of risk factors for wheezing in children under 2 years of age

| Risk factor                          | Wheeze (%) (n = 74) | Control (%) (n = 30) | Odds ratio (95% CI) | P value |
|--------------------------------------|---------------------|----------------------|---------------------|---------|
| Viral infection*                     | 45/74 (60.8)        | 4/30 (13.3)          | 15.5 (4.0 to 60.5)  | .0001   |
| Family history of allergy*           | 54/74 (72.9)        | 14/30 (46.6)         | 4.2 (1.4 to 12.4)   | .008    |
| Sensitization to inhalant and/or food allergens* | 9/70 (12.8) | 3/29 (10.3) | 1.3 (0.2 to 6.5) | .7      |
| Sex, male                            | 46/74 (62.1)        | 17/30 (56.6)         | 0.9 (0.3 to 2.7)    | .9      |

*Detection of rhinovirus RNA, coronavirus RNA, adenoivirus B, and/or RSV antigen in nasal washings.

†History of asthma, rhinitis, and/or atopic dermatitis in parent(s) and/or sibling(s).

‡Sensitization defined as IgE antibody levels ≥0.7 kUA/L (CAP class ≥2) to at least one inhalant allergen (mites, cockroach, cat, dog) or food allergen (egg, milk, soy, wheat, fish, or peanut).
The lack of evidence for association of rhinovirus and wheezing in the current study is striking. It is possible that genetic factors may play a role in determining immunologic responses to rhinovirus in different populations. In addition, rhinovirus serotypes circulating in tropical areas may be different from those in temperate climates, particularly regarding their ability to replicate in the lower respiratory airways. Temperatures in the lower airways may be critical for the intensity of replication, at least for some rhinovirus serotypes. It has been demonstrated that inhalation of cold air results in substantial temperature decrease in the lower airways that may favor rhinovirus replication. In the northern hemisphere, rhinovirus peak prevalence occurs in early fall and late spring and has been associated with new school terms. In subtropical climates, it is possible that higher temperatures year-round may be less conducive to productive growth of rhinovirus. Reduced rhinovirus spread in these areas could, at least in part, be due to the styles of housing peculiar to warmer climates, with homes and schools being ventilated by keeping windows open.

The role of adenovirus in causing wheezing and persistent asthma in children is largely unknown. It has been suggested that latency of adenovirus in infected lung epithelial cells is associated with amplification of cigarette smoke–induced inflammation and may cause allergen-induced eosinophilic inflammation to become steroid resistant. Adenoviruses play an important role as a cause of lower respiratory infections among children in Argentina and Chile, being secondary only to RSV.

Similar to previous studies, sensitization to inhalant allergens was the major independent risk factor for wheezing among children 2 to 12 years of age, and home exposure to high levels of inhalant allergens was not associated with current wheezing. We have previously reported that mites and cockroach were major causes of sensitization among children with asthma and/or allergic rhinitis, seen at university-based allergy clinics in Ribeirão Preto and São Paulo, Brazil. In the current study, the frequency of allergic sensitization was very high, notably among control children (43% of the children). Part of the reason for finding a higher-than-expected frequency of sensitization among control subjects may be due to the fact that we have not aimed to exclude subjects with allergic rhinitis from the control group. It would be difficult to have a definitive diagnosis of allergic rhinitis among children with respiratory distress at the ED setting. It is possible that some subjects with allergic rhinitis may have been included as control subjects, particularly in the group of children older than 2 years of age.

The relation of asthma and allergy with parasitic infection is controversial. Our studies in children living in poverty in northeast Brazil have shown that current infection with *A. lumbricoides* was an independent risk factor for wheezing, in addition to positive skin tests to inhalant allergens. A study of 2164 children 8 to 18 years of age from China revealed that infection with *A. lumbricoides* was associated with increased risk of asthma and with sensitization to common inhalant allergens. On the other hand, studies in rural Africa point to an inverse correlation between helminthic infections and atopy and asthma. Medeiros et al. have recently shown that subjects from a *Schistosoma mansoni* endemic area in Brazil had lower frequency of skin test positivity to indoor allergens and a milder course of asthma than individuals with asthma not living in an area endemic for *S. mansoni*. In the current study, we have found that IgE antibodies to *A. lumbricoides* were more frequently found in wheezing children than in control children; however, this association was no longer significant after multivariate analysis.

In conclusion, results of the current study have shed light on risk factors associated with acute wheezing in children living in a subtropical environment, an area where the prevalence of asthma is comparable with that in the United States and other developed countries. Wheezing in the prior 12 months, which has been shown to correlate better with diagnosis of asthma in Brazil, was reported in 22.5% and 16.7% of children 6 to 7 years old and 13 to 14 years old, respectively, evaluated by the ISAAC questionnaire. Some of the risk factors for acute wheezing identified in the current study, including RSV infections in the first 2 years of life, are similar to those previously reported in studies carried out in temperate developed countries. Inhalant allergens appear to be a major cause of asthma.
attacks in this environment, and rhinovirus does not appear to play a remarkable role in triggering asthma symptoms, as previously described in temperate regions.

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