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Prevalence of asthma, allergic rhinitis and eczema in 6–7-year-old schoolchildren from Luanda, Angola



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Risk factors

Abstract

Background: Epidemiological data have shown that the prevalence of asthma, rhinoconjunctivitis and eczema in children is still increasing, namely in Africa. However, there are no epidemiological studies on asthma or allergic diseases in Angolan children.

Objective: To study the prevalence of asthma and other allergic diseases in Angolan children.

Methods: Descriptive, observational, cross-sectional study, using the ISAAC study methodology, in the province of Luanda, Angola in 6–7-year-old children. Forty-six (8.3%) public schools were randomly selected. Data were analysed using the SPSS Statistics version 24.0 software.

Results: A total of 3080 children were studied. Results showed that the prevalence of asthma (wheezing in the previous 12 months) was 15.8%, that of rhinitis (sneezing, runny or blocked nose in the previous 12 months) was 19%, and that of eczema (itchy skin lesions in the previous 12 months) was 22%, without differences between sexes. Rhinitis was associated with a higher number of episodes of wheezing episodes, disturbed sleep and night cough, in children with asthma. Rhinitis, eczema, Split-type air conditioning system, antibiotic intake in the child's first year of life, frequent intake (more than once per month) of paracetamol and active maternal smoking were associated with a higher risk of having asthma, whereas electrical cooking was associated with a protective effect.

Conclusion: Asthma and allergic diseases are highly prevalent in children from Luanda. A strategy for preventive and control measures should be implemented.

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Introduction

Asthma is associated with a relevant burden of disease worldwide, which may still be increasing.^{1,2} The International Study of Asthma and Allergies in Childhood (ISAAC), part of which included two phases (Phases I and III) separated by a 5–10-year interval, was implemented in multiple centres worldwide. Although there were differences in prevalence values of asthma, rhinitis and eczema among participating countries, prevalence was increasing in children, particularly in countries with a lower prevalence in Phase I, and which mostly included developing countries.^{3,4}

Epidemiological data regarding children from Africa are scarce but global analysis of the ISAAC study and other reports showed that the prevalence of asthma averaged around 10% for 6–7-year olds.^{2,5} Furthermore, prevalence values for asthma varied significantly throughout Africa: 16% in Botswana,⁶ 13.3% in Mozambique,⁷ 11.1% in South Africa,⁸ 9% in Senegal,⁹ and 4.8% in Nigeria.³ In addition, increases in prevalence were detected between Phases I and III, as reported in Nigeria, with values increasing from 4.8% to 5.6%.^{3,10} Furthermore, African countries had a high proportion of children reporting symptoms of severe asthma.^{2,11} In Angola, asthma is one of the main causes for visits to emergency units in children. However, although we have previously studied the prevalence of asthma and allergic diseases in Angolan adolescents,¹² no studies were performed in children. We therefore decided to study the prevalence of asthma and allergic diseases in 6–7-year-old children from Luanda.

Methods

Population sample

Cross-sectional, observational study performed in the province of Luanda, Angola, between August and October 2014, and between March and May 2015, in 6–7-year-old schoolchildren. Luanda is the capital and includes seven boroughs in which 97.5% of the population is urban. In Luanda, 46 (8.3%) primary public schools were randomly selected out of a total of 552, to meet the ISAAC criterium of analysing at least 3000 children.¹³ Children's parents/guardians were classified, in sociodemographic terms, as low, middle or upper class, in accordance with criteria of the Angolan 2015–2016 IIMS Inquiry on Multiple and Health Indicators.

Written questionnaires

We used the Portuguese version of the ISAAC questionnaire,^{13,14} which has questions on symptoms of asthma, allergic rhinitis and eczema and which was filled out by children's parents/guardians. The ISAAC Phase III Environmental exposure and risk factor questionnaire was also used.¹⁵ All questions and explanations about the questionnaire were supplied in a standardised manner, in Portuguese.

Current asthma was defined as positive replies to the question "Has your child had wheezing or whistling in the chest in the past 12 months?".¹³ Parents also answered

questions on the number of wheezing episodes, interference of wheezing with sleep or speech, relation to physical exercise and episodes of nocturnal cough, in the previous 12 months.

Current rhinitis was defined as sneezing bouts, rhinorrhoea or nasal obstruction, in the absence of flu, in the previous 12 months, and rhinoconjunctivitis involved the presence of rhinitis and conjunctivitis symptoms.¹³ Parents were also asked whether nasal symptoms interfered with their child's daily activities and whether their child had ever had "hay fever".

Eczema was considered if parents/guardians reported cutaneous lesions with pruritus, which waxed and waned, in the previous 12 months.¹³ Additional questions were asked regarding specific location and age of appearance of the lesions and whether the latter interfered with sleep.

The questionnaire on environmental exposure included questions on the type of fuel used for cooking, type of indoor home-cooling device, frequency of passage of trucks in front of their homes, presence of cats and dogs at home, child's passive exposure to tobacco smoke, use of antibiotics in their child's first year of life, frequency of use of paracetamol, breastfeeding and the number of siblings living in the home.

Measurement of lung function by peak flow metre

Peak flow metre recordings (Mini-Wright Peak Flow Metre, Clement-Clarke, Harlow, UK) were performed in all children with reported current asthma. Children with symptoms of infectious acute respiratory illness were excluded. Readings were taken in triplicate with the child standing and the highest value was recorded only if the coefficient of variation was below 5%. Since there are no reference values for Peak Expiratory Flow (PEF) for Angolan children, we used values from Nigerian schoolchildren¹⁶ for the calculation of percentage predicted values and comparison with ranges of reference values (above 80%, 50–80% or below 50%). We defined confirmed current asthma as current asthma symptoms associated with PEF values below 80% predicted.

Height and weight measurements

The height of each child was measured using a portable 200 cm stadiometer, accurate to 0.1 cm (SECA 123, Hamburg, Germany) and recorded in centimetres. Children had their backs turned to the stadiometer and their heads were positioned in the Frankfurt horizontal plane. Weight was measured using a portable, calibrated scales, with a 150 kg capacity and a 0.1 kg precision (SECA 780 digital scale, Hamburg, Germany) and recorded in kilograms. For both measurements, children were standing upright, and barefoot.

Calculation of body mass index (BMI)

Body mass index (BMI) was calculated with the standard formula: weight (in kg)/height (in m²). Since there are no BMI reference values for Angola, children were classified as "underweight", "normal weight", "overweight", and

‘‘obese’’, in accordance with World Health Organisation definitions regarding BMI values.¹⁷

Ethical considerations

This study was approved by the Ethics Committees of the Angolan Ministry of Health and the Faculty of Health Sciences, University of Beira Interior, Portugal, by the Provincial Board of Education, Luanda, Angola, and by the Directors of the selected schools. All parents/guardians were informed about the study in a face-to-face session as well as via a leaflet, and those who agreed to participate signed a written consent form.

Statistical analysis

Data were analysed using the Software Package for Social Sciences (SPSS) version 24.0[®]. Descriptive analysis was used for frequencies, percentages, means and standard deviations. Prevalence values were estimated by dividing the number of positive responses to the questions selected for diagnosis by the number of completed questionnaires. Comparison of proportions was performed using Chi-Square Test or Fisher’s Exact Test, as appropriate. Odds ratios (OR) were calculated for characterisation of possible risk factors for asthma. A logistic regression model was developed using the *logit* function. For categorical variables, the ‘‘normal’’ situation was defined as the reference category and odds were estimated for the other categories against the reference one. Quality and assumptions of the model were tested using the Omnibus and Hosmer–Lemeshow tests, as well as by analysis of residuals and outliers. A Receiver Operating Characteristic (ROC) analysis of the model was also carried out. A *p*-value of less than 0.05 was regarded as significant with all two-tailed statistical tests.

Results

Demographics

All directors of the 46 randomly selected schools agreed to participate in the study. The final sample included 4505 children whose parents received information and the questionnaire. From these, 83 parents did not return the questionnaire (98% reply rate), and 1342 questionnaires were excluded because they were incomplete or incorrectly filled in. Thus, we obtained 3080 valid questionnaires (68.3% reply rate). There was no concentration of non-responders or responders with invalidated questionnaires in any specific school. Of the validated 3080 questionnaires, 1608 (52.2%) were female and 1472 were male (47.7%) (Table 1). Gender and age distributions were similar to those in the non-responders or responders with invalid questionnaires. All children who participated in the study lived in an urban area. The borough of Kissama was excluded because most parents/guardians were illiterate. In socio-demographic terms, just over 40% of the children (1241) belonged to a low social class, whereas most belonged to middle or upper classes. Although only around 23% of the

Table 1 Sociodemographic data of study sample of 6–7-year-old children from Luanda.

Parameter	Total (n)
<i>Boys: Girls (n; %)</i>	1472 (47.8%): 1608 (52.2%)
<i>Urban: Rural (%)</i>	100: 0
<i>Boroughs of dwelling (n; %)</i>	
Luanda	2036 (66.0%)
Belas	350 (11.4%)
Cacuaco	104 (3.4%)
Viana	153 (5.0%)
Icolo e Bengo	65 (2.1%)
Cazenga	372 (12.1%)
<i>Social status and income (n; %)</i>	
High	692 (22.5%)
Medium	1147 (37.2%)
Low	1241 (40.3%)
<i>Parental schooling (n; %)</i>	
Primary school (up to 4 years)	1143 (37.1%)
Intermediate school (up to 10 years)	1220 (39.6%)
High school (up to 14 years)	717 (23.3%)

mothers had high school/university level schooling, significant proportions had lower secondary or primary schooling.

Prevalence of asthma-like symptoms

Of the 3080 children included in the study, almost 24% reported that they had already had wheezing episodes in their lives (Table 2). However, 485 children had had wheezing in the last 12 months, indicating a prevalence of current asthma of 15.8% (95% CI: 14.5–17.1%), without significant differences between boys and girls. Only around 7% of the children reported having wheezing during or after physical exercise, but 26.4% reported episodes of nocturnal dry cough, not associated with respiratory infections in the previous 12 months (Table 2).

The prevalence of ‘‘Wheezing ever’’, ‘‘Wheezing with physical exercise in the last 12 months’’ and ‘‘Nocturnal cough in the last 12 months’’ was significantly higher in girls than in boys.

Of the 485 children with current asthma, only 37 (8.6%) were regularly seen by a paediatrician due to their asthma symptoms, and 268 (55.2%) had already been seen more than once at an emergency department and occasionally prescribed a β 2-agonist.

Prevalence of rhinitis

The prevalence of current rhinitis was 19% (95% CI: 17.7–20.5%; *n* = 586), and that of current rhinoconjunctivitis was 10% (95% CI: 9.0–11.1%; *n* = 309) (Table 2). Symptoms of rhinitis interfered with the daily activities in only 4.5% of the children. Around 15% of the children had had ‘‘Hay fever ever’’. No significant differences in the prevalence of rhinitis or rhinoconjunctivitis symptoms were seen between sexes.

Table 2 Prevalence rates for asthma, rhinitis and eczema.

	Total (n)	% (n)	F	%	M (n)	%	p Value
<i>Bronchial asthma</i>							
Wheezing ever	724	23.5	354	22.0	370	25.1	0.041
Wheezing last 12 months	485	15.7	238	14.8	247	16.8	0.132
Asthma ever	558	18.1	271	16.9	287	19.5	0.057
Exercise-induced wheezing last 12 months	227	7.4	99	6.2	128	8.7	0.007
Nocturnal cough last 12 months	812	26.4	388	24.1	424	28.8	0.003
<i>Rhinitis</i>							
Sneezing, runny or blocked nose ever	692	22.5	352	21.9	340	23.1	0.423
Sneezing, runny or blocked nose last 12 months	586	19.0	294	18.3	292	19.8	0.273
Rhinoconjunctivitis last 12 months	309	10.0	149	9.3	160	10.9	0.139
<i>Interference with activities last 12 months</i>							
None or few	2941	95.5	1540	95.8	1401	95.2	0.427
More or less or very	139	4.5	68	4.2	71	4.8	
Hay fever ever	464	15.1	230	14.3	234	15.9	0.217
<i>Eczema</i>							
Itchy rash ever	671	21.8	347	21.6	324	22.0	0.772
Itchy rash last 12 months	568	18.4	296	18.4	272	18.5	0.960
Itchy flexural areas	351	11.4	178	11.1	173	11.8	0.547
Itchy before 2 years old	184	6.0	88	5.5	96	6.5	0.221
Itchy between 2 and 4 years old	188	6.1	95	5.9	93	6.3	0.635
Itchy with 5 or more years old	196	6.4	116	7.2	80	5.4	0.043
Clearance of rash last 12 months	337	10.9	181	11.3	156	10.6	0.559
Interference with sleep last 12 months	107	3.5	59	3.7	48	3.3	0.532
Eczema ever	450	14.6	239	14.9	211	14.3	0.679

Prevalence of eczema

Itchy rash or eczema “ever” were reported in almost 22% of the children (Table 2), and 18.4% of the children (95% CI: 17.0–19.8%; $n = 568$) had had such lesions in the previous 12 months. The lesions affected specific areas of the body in 11.4% of the children and disappeared at least temporarily, in around 11% of the cases. Cutaneous symptoms only affected sleep in 3.5% of the children. No significant differences in the prevalence of eczema symptoms were seen between sexes.

Respiratory symptoms and function in adolescents with current asthma

Of the 485 children with reported current asthma, most (74.2%) had only had one to three wheezing episodes (Table 3). However, almost 11% reported more than 12 episodes and 22.1% of the children woke up during the night, more than once weekly, because of wheezing episodes. In addition, almost 27% of the children had had episodes of wheezing that interfered with speech, 34% had had wheezing episodes during or after physical exercise, and 71% reported dry cough during the night. Finally, PEF recordings showed that a high proportion (47.3%; 229) of the children had a moderate degree of obstruction and around 3% (16 children) had severe obstruction (Table 3), confirming the presence of asthma in 50% of the children reporting

symptoms in the previous 12 months, and suggesting a prevalence of confirmed current asthma of 8.0% (95% CI: 7.0–9.0%).

Influence of rhinitis symptoms upon symptoms of asthma

In the 485 children with current asthma, the presence of rhinitis in the last 12 months was significantly associated with a higher number of episodes of nocturnal cough ($p < 0.001$; Table 4). In fact, having current rhinitis symptoms increased the risk of having a high number of wheezing episodes, disturbed sleep, and nocturnal episodes of dry cough about four-fold (*Odds ratio*).

Risk factors for asthma

Current rhinitis or eczema, AC-split home cooling system, excessive intake of paracetamol, intake of antibiotics in the first year of life, frequent passage of trucks, the presence of animals at home during pregnancy or the child's first year of life, and active maternal smoking during the child's infancy as well as the number of smokers at home were significantly associated with asthma, using univariate analysis, whereas having a fan as a cooling system, having a higher number of siblings at home, and breastfeeding significantly reduced the risk of asthma, and using electricity for

Table 3 Clinical features of asthma in children with asthma symptoms ("Wheezing in the last 12 months"; $n = 485$).

	Total (n)	%	F (n)	%	M (n)	%	p Value
<i>Wheezing episodes last 12 months</i>							
1–3	360	74.2	180	75.6	180	72.9	0.488
4–12	72	14.8	36	15.1	36	14.6	0.864
>12	53	10.9	22	9.2	31	12.6	0.243
<i>Sleep disturbance episodes last 12 months</i>							
<1/week	244	50.3	126	52.9	118	47.8	0.255
≥1/week	107	22.1	53	22.3	54	21.9	0.914
Speech disturbance last 12 months	130	26.8	65	27.3	65	26.3	0.805
Asthma ever	261	53.8	122	51.3	139	56.3	0.268
Exercise-induced wheezing last 12 months	167	34.4	71	29.8	96	38.9	0.036
Nocturnal cough last 12 months	345	71.1	163	68.5	182	73.7	0.207
<i>Peak-flow recordings (% predicted)</i>							
>80%	239	49.4	99	41.8	140	56.7	0.003
50–80%	229	47.3	127	53.6	102	41.3	
<50%	16	3.3	11	4.6	5	2.0	

Table 4 Associations between the presence of rhinitis in the last 12 months and clinical asthma parameters in children with asthma symptoms (wheezing episodes in the last 12 months; $n = 485$).

	Rhinitis last 12 months		Odds ratio (95% CI)	p Value
	Yes	No		
<i>Wheezing episodes last 12 months</i>				
<4	189	171	1	0.017
4–12	49	23	1.93 (1.13; 3.30)	
>12	29	24	1.09 (0.61; 1.95)	
<i>Sleep disturbance episodes last 12 months</i>				
<1/week	143	101	1	0.125
≥1/week	72	35	1.45 (0.90; 2.34)	
<i>Nocturnal cough last 12 months</i>				
No	42	98	1	<0.001
Yes	225	120	4.38 (2.86; 6.69)	
<i>Peak-flow recordings (% predicted)</i>				
Above 80%	137	102	1	0.432
50–80%	123	106	0.86 (0.60; 1.24)	
Below 50%	6	10	0.45 (0.16; 1.27)	

For each categorical variable, the "normal" situation was defined as the reference category and odds were estimated for the other categories against the reference one.

cooking was almost significantly protective (Table 5). However, in the logistic regression model, only rhinitis, eczema, AC-split type of cooling system, high intake of paracetamol, antibiotic intake and active maternal smoking during the child's first year of life were confirmed as significant risk factors, whereas electricity as cooking system was a protective factor (Table 6).

Discussion

This is the first study of asthma prevalence in Angolan children, one of few studies in 6–7-year-olds in Africa,

and showed a value of 15.8%, without significant differences between boys and girls, and that 8% of asthmatics had confirmed bronchial obstruction. The prevalence of rhinitis was 19%, that of eczema was 22%, again without differences between genders. Rhinitis was associated with clearly more symptomatic asthma. Rhinitis and eczema, use of AC-split home cooling system, frequent intake of paracetamol, antibiotic use and active maternal smoking in the child's first year of life were significantly associated with an increased risk of asthma, whereas cooking with electricity was protective.

We followed the ISAAC methodology in a random sample of more than 3000 children, had a high reply rate,

Table 5 Risk factors for probable asthma (Wheezing last 12 months).

Risk factors	Total	%	Positive wheezing 12 M	%	Negative wheezing 12 M	%	Odds ratio (95% CI); Logistic regression	p Value ^a
<i>Rhinitis last 12 months</i>								
No	2494	81.0	218	44.9	2276	87.7	1	
Yes	586	19.0	267	55.1	319	12.3	8.74 (7.06; 10.82)	<0.001
<i>Itchy rash last 12 months</i>								
No	2512	81.6	282	58.1	2230	85.9	1	<0.001
Yes	568	18.4	203	41.9	365	14.1	4.40 (3.56; 5.44)	
Cooking system used at home								
<i>Electricity</i>								
No	2940	95.5	471	97.1	2469	95.1	1	0.059
Yes	140	4.5	14	2.9	126	4.9	0.58 (0.33; 1.02)	
<i>Gas</i>								
No	5	0.2	0	0	5	0.2		–
Yes	3075	99.8	485	100	2590	99.8	–	
<i>Coal</i>								
No	2701	87.7	430	88.7	2271	85.5	1	0.481
Yes	379	12.3	55	11.3	324	12.5	0.90 (0.66; 1.22)	
<i>Other</i>								
No	3080	100	485	100	2595	100		–
Yes	0	0	0	0	0	0	–	
Indoor home cooling system								
<i>AC-split</i>								
No	1943	63.1	207	42.7	1736	66.9	1	<0.001
Yes	1137	36.9	278	57.3	859	33.1	2.71 (2.23; 3.31)	
<i>Window AC</i>								
No	2513	81.6	409	84.3	2104	81.1	1	
Yes	567	18.4	76	15.7	491	18.9	0.80 (0.61; 1.04)	0.091
<i>Fan</i>								
No	1511	49.1	295	60.8	1216	46.9	1	<0.001
Yes	1569	50.9	190	39.2	1379	53.1	0.57 (0.47; 0.69)	
<i>Other</i>								
No	3080	100	485	100	2595	100		–
Yes	0	0	0	0	0	0	–	
<i>None</i>								
No	2885	93.7	463	95.5	2422	93.3	1	0.079
Yes	195	6.3	22	4.5	173	6.7	0.67 (0.42; 1.05)	
<i>Frequency of paracetamol intake</i>								
Never	406	13.2	15	3.1	391	15.1	1	
>Once/year	1050	34.1	138	28.5	912	35.2	3.94 (2.29; 6.81)	<0.001
>Once/month	1621	52.7	332	68.5	1289	49.7	6.71 (3.93; 11.40)	
<i>Antibiotic intake</i>								
No	909	29.5	76	15.7	833	32.1	1	
Yes	2171	70.5	409	84.3	1762	67.9	2.54 (1.97; 3.29)	<0.001
<i>Breast-feeding</i>								
No	150	4.9	34	7.0	116	4.5	1	
Yes	2930	95.1	451	93.0	2479	95.5	0.62 (0.42; 0.92)	<0.001

Table 5 (Continued)

Risk factors	Total	%	Positive wheezing 12 M	%	Negative wheezing 12 M	%	Odds ratio (95% CI); Logistic regression	p Value ^a
<i>Number of siblings</i>								
Mean±SD	2.6±2.2	–	2.3±1.9	–	2.7±2.3	–		0.018
Median (range)	2 (0–18)		2 (0–10)		2 (0–18)		0.93 (0.89; 0.97)	
<i>Frequency of passage of trucks in front of home</i>								
Never	420	13.6	55	11.3	365	14.1	1	0.002
Seldom	1617	52.5	229	47.2	1388	53.5	1.10 (0.80; 1.50)	
Frequently in the day	708	23.0	140	28.9	568	21.9	1.64 (1.17; 2.30)	0.574
Almost the whole day	335	10.9	61	12.6	274	10.6	1.48 (0.99; 2.20)	0.004
Pets at home								
<i>Cat (first year of life)</i>								
No	2899	94.1	456	94.0	2443	94.1	1	
Yes	181	5.9	29	6.0	152	5.9	1.02 (0.68; 1.54)	0.054
<i>Cat (last 12 months)</i>								
No	2894	94.0	449	92.6	2445	94.2	1	
Yes	186	6.0	36	7.4	150	5.8	1.31 (0.90; 1.91)	0.917
<i>Dog (first year of life)</i>								
No	2196	71.3	339	69.9	1857	71.6	1	
Yes	884	28.7	146	30.1	738	28.4	1.08 (0.88; 1.34)	0.165
<i>Dog (last 12 months)</i>								
No	2037	66.1	313	64.5	1724	66.4	1	
Yes	1043	33.9	172	35.5	871	33.6	1.09 (0.89; 1.33)	0.457
<i>Animals (first year of life)</i>								
No	2834	92.0	429	88.5	2405	92.7	1	
Yes	246	8.0	56	11.5	190	7.3	1.65 (1.21; 2.27)	0.417
<i>Animals (during pregnancy)</i>								
No	2816	91.4	425	87.6	2391	92.1	1	
Yes	264	8.6	60	12.4	204	7.9	1.66 (1.22; 2.25)	0.002
Smoking at home								
<i>Mother</i>								
No	3034	98.5	473	97.5	2561	98.7	1	
Yes	46	1.5	12	2.5	34	1.3	1.91 (0.98; 3.72)	0.001
<i>Mother: n. cigarettes/day</i>								
Mean±SD	7.4±5.8	95.6	7.4±4.9	93.8	7.4±6.2			
Median (range)	5 (1–24)	4.4	6.5 (2–16)	6.2	5 (1–24)	–	–	0.056
<i>Father</i>								
No	2945	–	455	–	2490	96.0	1	
Yes	135		30		105	4.0	1.56 (1.03; 2.38)	0.819 ^b
<i>Father: n. cigarettes/day</i>								
Mean±SD	8.0±5.7	98.4	9.3±6.0	96.5	7.7±5.6			
Median (range)	6 (1–30)	1.6	9 (2–20)	3.5	5 (1–30)	–	–	0.036
<i>Mother (first year of life)</i>								
No	3030	92.4	468	89.3	2562	98.7	1	
Yes	50	7.6	17	10.7	33	1.3	2.82 (1.56; 5.11)	0.171 ^b
<i>n. smokers ≥1</i>								
No	2846	92.4	433	89.3	2413	93.0	1	0.001
Yes	234	7.6	52	10.7	182	7.0	1.59 (1.15; 2.20)	

Table 5 (Continued)

Risk factors	Total	%	Positive wheezing 12 M	%	Negative wheezing 12 M	%	Odds ratio (95% CI); Logistic regression	p Value ^a
BMI								
<i>Normal</i>								
Weight – mean; SD (kg)	46 20.2 ± 1.6	4.0	22 20.1 ± 1.6	4.5	24 20.3 ± 1.7	3.6	1	0.005
<i>Underweight</i>								
Weight – mean; SD (kg)	110514.7 ± 1.7	95.7	46114.7 ± 1.6	95.1	64414.7 ± 1.7	96.1	0.78 (0.43; 1.41)	0.412
<i>Overweight</i>								
Weight – mean; SD (kg)	4 26.6 ± 1.0	0.3	2 26.9 ± 0.6	0.4	2 26.3 ± 1.6	0.3	1.09 (0.14; 8.42)	0.933

^a Wald's test.^b Mann-Whitney test.

For each categorical variable, the "normal" situation was defined as the reference category and odds were estimated for the other categories against the reference one.

and used "Wheezing episodes in the last 12 months" for the diagnosis of current asthma, since it has high sensitivity for this purpose.^{18,19} The prevalence of asthma in our study (15.8%) places Angola as the country with the 11th highest prevalence, when compared with countries that participated in ISAAC Phases I and III, and which showed Phase III values ranging between 37.6% (Costa Rica) and 4.1% (Indonesia).³ Furthermore, the prevalence value we found is higher than the mean for 6–7-year-old children in Africa (10%).¹¹ The highest value was reported in a 2014/2015 study in Botswana (15.9%),⁶ which is similar to our report, although only 385 schoolchildren were included in the former study. In Mozambique, data from 2004 showed a relatively similar prevalence of asthma (13.3%).⁷ Prevalence was lower both in South Africa (11.1%),^{2,8} and in Nigeria (5.5%), in 2001/2002.^{3,10} Finally, a study using the ISAAC protocol, performed in rural Senegal showed a prevalence of 9.0% in 5–8-year-old schoolchildren.⁹ Since these studies used the same questionnaire, discrepant values may be due to time, genetic, environmental or lifestyle differences, as was seen in Mozambique, with prevalence of cough being higher in suburban and semi-rural children.⁷ However, our study as well as others may have underestimated the prevalence of asthma, since some of the parents/guardians did not know the concept of "wheezing", and symptom recognition may be poor or not well accepted.⁷ The prevalence of nocturnal cough in our study (26.4%) was high, but similar to that in Mozambique (27.5%),⁷ slightly above that seen in Botswana⁶ and Senegal,⁹ and clearly above that reported in Nigeria (6.5%).³ However, it is possible that cough was not always associated with asthma.

Although the prevalence of current rhinitis (22.5%) was high, that of current rhinoconjunctivitis was lower (10%), and places Angola at the bottom of the top third of ISAAC participating countries worldwide, above the global mean of 8.5%.^{3,20} In Africa, it was similar to prevalence in South Africa (10.6%)²⁰ and Mozambique (8.9%)⁷ and much higher than in Nigeria (3.6%).^{3,20} In our study, only 15% of the

parents/guardians reported that their children had "Hay fever ever", which, again was similar to Mozambique (12%),⁷ and much lower than the value in Angolan (33%)¹² or Mozambican adolescents.⁷ This suggests that either the prevalence of rhinoconjunctivitis tends to increase with age, or that adolescents more frequently overestimate the situation. Nevertheless, in non-English speaking countries, as well as in countries without a clear pollen season, as happens in Luanda, "hay fever" is a concept that is not easy to interpret.

The prevalence of current eczema in our study (18.4%) is the second highest in all ISAAC reporting countries, significantly higher than the mean world prevalence (9.3%).^{3,22} In Africa, it is much higher than the values reported in Mozambique (12.8%),⁷ South Africa (12.3%)²¹ and Nigeria (5%).^{3,21} In contrast with results in 13–14-year-old adolescents,^{3,10,12} the highest prevalences of eczema in the ISAAC study of 6–7-year olds came essentially from scattered centres, including the UK, Australia, New Zealand, Panama and Chile which also reported the highest asthma prevalences,¹⁰ and Angola has a similar situation. Although eczema is a significant public health problem in developing countries,²¹ non-eczema-related manifestations may have been reported in our study, and in others.⁷ Nevertheless, comparison of ISAAC Phases I and III showed that the prevalence of eczema increased in most countries, independently of their socioeconomic status.^{2,3,21}

Since our focus was asthma, we further analysed clinical features in the 485 children who reported symptoms in the previous 12 months. Almost 11% reported more than 12 episodes of wheezing in that period and about a quarter had frequent sleep disturbance episodes. Furthermore, a high percentage (27%) had episodes of wheezing that interfered with speech, as seen in other countries,^{5–7} about one third had exercise-induced wheezing and a high proportion (71%) reported episodes of nocturnal cough. In addition, almost 50% had a moderate degree of bronchial obstruction. Although these findings may have been biased by manifestations misinterpreted as wheezing, by reporting of cough

Table 6 Adjusted odds ratios of risk factors for probable asthma (wheezing last 12 months).

Risk factors	Adjusted odds ratio (95% CI); Logistic regression	p Value ^a
<i>Rhinitis last 12 months</i>		
No	1	
Yes	6.48 (5.14; 8.17)	<0.001
<i>Itchy rash last 12 months</i>		
No	1	
Yes	2.15 (1.66; 2.80)	<0.001
Cooking system used at home		
<i>Electricity</i>		
No	1	
Yes	0.38 (0.20; 0.74)	0.004
Indoor home cooling system		
<i>AC-split</i>		
No	1	
Yes	2.66 (1.95; 3.62)	<0.001
<i>Frequency of paracetamol intake</i>		
Never	1	
≥Once/year	2.34 (1.31; 4.18)	0.004
≥Once/month	3.24 (1.84; 5.70)	<0.001
<i>Antibiotic intake</i>		
No	1	
Yes	1.75 (1.31; 2.34)	<0.001
<i>Breast-feeding</i>		
No	1	
Yes	0.69 (0.43; 1.09)	0.115
<i>Number of siblings</i>	0.95 (0.90; 1.01)	0.095
Pets at home		
<i>Cat (first year of life)</i>		
No	1	
Yes	0.69 (0.40; 1.17)	0.168
Smoking at home		
<i>Mother (first year of life)</i>		
No	1	
Yes	2.82 (1.13; 7.00)	0.026

^a Wald's test; OR's adjusted for all factors in Table 5, except BMI and "n. of cigarettes/day father and mother"; Only the results are shown when $p < 0.2$; Omnibus test: $p < 0.001$; Hosmer–Lemeshow test: $p = 0.330$; Nagelkerke pseudo- $R^2 = 0.303$; ROC analysis: area under curve = 0.809 (95%CI: (0.787; 0.831)); sensitivity = 72.8%, specificity = 76.1%, overall = 75.6% (probability cut-off = 0.148). For each categorical variable, the "normal" situation was defined as the reference category and odds were estimated for the other categories against the reference one.

due to causes other than asthma or by suboptimal technical performance of peak flow by some children, it should be stressed that, in a high proportion of cases symptoms clustered together in the same children. Thus, our results show that a high percentage of children in Luanda are asthmatic and frequently have uncontrolled symptoms. This is in line with ISAAC study findings of the highest prevalence of symptoms of severe asthma among children with current wheeze being observed in low and middle-income countries.^{1,11} Globally, asthma should be regarded as a

priority in terms of non-communicable diseases, as stated in the 2018 GAN - Global Asthma Network Report.²

We also identified risk factors for asthma. In the total sample of children, rhinitis increased almost nine-fold the risk of having asthma, as seen in other countries^{10,22–24} and in Angolan adolescents.¹² Rhinitis is a known risk factor for asthma and may worsen asthma symptoms.^{10,25} In our study, in children with asthma, rhinitis was associated with significantly more wheezing episodes and nocturnal cough. We also identified eczema as another risk factor, since current

eczema increased the risk of having asthma four-fold, as also seen in adolescents,¹² and in reports showing a relationship between early onset of atopic eczema and subsequent respiratory disease in schoolchildren.^{26,27}

Using AC-split air conditioning system was also a significant risk factor, as seen in other studies.^{12,28} This system may constitute a risk because cleaning it is difficult, which may allow accumulation of allergens,²⁹ microorganisms and irritating substances.

We also detected drugs as risk factors for asthma. Antibiotics given to children during their first year of life increased the risk of asthma, as seen in ISAAC studies.³⁰ A high frequency of paracetamol intake was another risk factor, which is also in agreement with ISAAC findings,^{30,31} was also reported in Angolan adolescents,¹⁰ and is also a risk factor for rhinoconjunctivitis and eczema.^{30,31} Maternal smoking during the child's first year of life also increased the risk of asthma, as reported in the ISAAC study, which also showed an increased risk of rhinoconjunctivitis, and eczema.³² Furthermore, multinational, longitudinal studies performed in Europe showed that maternal smoking during pregnancy and the child's first year of life is a significant risk factor for subsequent development of wheezing in early childhood or adolescence.^{33,34} A small study performed in Mozambique, in asthmatic and non-asthmatic children, aged between 18 months and eight years, also showed that having at least one parent who smoked was a significant risk factor for asthma.²³ Further studies are needed in African and other developing countries.

In contrast, using electricity as cooking system was a protective factor against asthma, which may be explained by the fact that children in homes that use this form of energy are less exposed to toxic fumes than those from homes where coal (open fire) is used for cooking. In fact, coal-based cooking has been shown to be a risk factor for asthma, in many studies.^{2,35}

Our study had several limitations. It is based on self-reports by parents/guardians of the children and may, therefore, be influenced by various types of bias, although the ISAAC questionnaire makes it likely that reported symptoms significantly reflect the clinical situation.³⁶ Some parents/guardians did not know some of the terms used in the questionnaire, as seen in other ISAAC studies. In addition, all children were from urban areas and relatively well-off families and results may not be fully extrapolated to children from poorer, rural areas. Some other potentially relevant risk factors, such as family history of asthma, were not included in our analysis, which partially impaired full comparisons with other studies. The ISAAC questionnaire on environmental exposure is validated but its level of detail may not be sufficient for some of the risk factors. Lastly, the cross-sectional design of the study does not allow analysis of interrelationships between different diseases, in patterns of multimorbidity or risk factors.

Conclusion

Asthma and related allergic diseases are a public health problem in children from Luanda, and a high proportion of children with asthma are frequently symptomatic and this

may also apply to other developing countries. Thus, preventive and control measures should be implemented to deal with this problem.

Author contributions

MA participated in the study design, data collection and analysis, as well as in writing the manuscript; OL and FQ participated in data collection and analysis; JRP participated in the study design and writing the manuscript; JMRG carried out the statistical analyses and participated in writing the manuscript; LTB supervised the whole project and participated in study design, data analysis and writing the manuscript.

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Conflict of interest

OL, FQ, and JMRG have no conflicts of interest and have not been funded.

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LTB has received support to attend EAACI congresses from Victoria Laboratories and Menarini and has been paid lecture fees by Novartis, AstraZeneca, Merck Sharp & Dohme and Menarini.

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