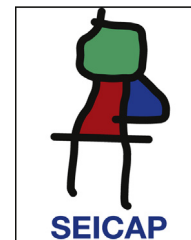


Allergologia et immunopathologia

Sociedad Española de Inmunología Clínica,
Alergología y Asma Pediátrica

www.elsevier.es/ai



ORIGINAL ARTICLE

Overuse of bronchodilators and steroids in bronchiolitis of different severity Bronchiolitis-study of variability, appropriateness, and adequacy

C. Ochoa Sangrador^{a,*}, J. González de Dios^b, Research Group of the aBREVIADo Project[◇]

^a Service of Pediatrics, Hospital Virgen de la Concha, Zamora, Spain

^b Service of Pediatrics, Hospital General Universitario de Alicante, Department of Pediatrics, Universidad Miguel Hernández, Alicante, Spain

Received 6 December 2012; accepted 2 February 2013

Available online 14 June 2013

KEYWORDS

Acute bronchiolitis;
Treatment;
Bronchodilators;
Corticosteroids;
Physician's practice
patterns

Abstract

Background: In the management of acute bronchiolitis there is a generalised use of treatments that have not been shown to be useful or efficacious in clinical studies. The objective of this study was to determine the appropriateness in the treatment of acute bronchiolitis of different severity within different clinical care settings.

Methods: This is a cross-sectional, descriptive study of 5647 cases of acute bronchiolitis in 91 Spanish hospitals and primary care centres. We classified the appropriateness of the treatments according to the recommendations of a consensus conference.

Results: There was an inappropriate use of treatments in 58.3% of the cases during the acute phase and in 45.4% during the maintenance phase. There was a generalised use of inhaled beta 2 agonists, regardless of the severity of the patients (hospitalised patients 69.3%, emergency care 63.2% and ambulatory 64.1%). Adrenaline was used in 30.1% of hospitalised cases and in 80.2% of intensive care patients. Systemic corticosteroids were not only used in one-third of hospitalised patients but also in 25.8% of ambulatory cases.

Conclusions: In acute bronchiolitis in Spain there is a wide use of treatments that are not recommended by the available clinical practice guidelines. Beta 2 agonist bronchodilators and corticosteroids are widely used and maintained, regardless of the severity of the patients.

© 2012 SEICAP. Published by Elsevier España, S.L. All rights reserved.

Introduction

Acute bronchiolitis is the main cause of hospital admissions related to acute lower respiratory airway infections in infants. It has significant repercussions at all health care

* Corresponding author.

E-mail address: cochoas@meditex.es (C. Ochoa Sangrador).

◇ Members of the Research Group specified in Annex 1.

levels. Literature on the management of bronchiolitis is very abundant in diagnostic as well as preventive-therapeutic aspects. The published information has been revised in depth, and different clinical practice guidelines (CPGs) are available.¹⁻⁵ The evidence suggests that in the treatment of bronchiolitis, the use of symptomatic support measures is fundamental for the management of fever, respiratory secretions, hyporexia, respiratory distress and hypoxaemia. Other treatments, in spite of their wide use, have not shown enough efficacy in clinical trials and present unfavourable benefit-risk ratios. A therapy trial with inhaled beta 2 agonists or adrenaline (better with hypertonic saline solution) has been proposed by some CPGs, but only for moderate-to-severe cases. These treatments can only be maintained if there is a documented improvement that compensates their costs and adverse effects.

The objective of our study was to analyse the appropriateness in the treatment of bronchiolitis in a large and representative sample of different health care settings in Spain. This study complements a preliminary one, conducted within the aBREVIADo Project, which describes the global variability in the clinical management. Here, we present an analysis of the appropriateness of the treatments in relation to the severity of the patients.⁶

Materials and methods

Design

This was a cross-sectional, descriptive study of acute bronchiolitis cases in a sample of hospitals, emergency services and primary care centres or offices in Spain. The participating centres belonged to 12 autonomous communities (25 provinces) and corresponded to 31 autonomous communities (25 complete hospitals, 7 hospitalisation services, and 6 emergency services) and 60 primary care centres or offices (Annex 1). The information of this descriptive study is part of the aBREVIADo Project (Bronchiolitis-Study of Variability, Adequacy, and Adherence), in which the recommendations made by the consensus conference of bronchiolitis were used as reference standards.⁷

Study period

From October 2007 to March 2008.

Inclusion criteria

All bronchiolitis cases were diagnosed during the study period according to the McConnochie criteria⁸: first acute episode of respiratory distress with wheezing preceded by a cold-like clinical picture of the upper respiratory airway (rhinitis, cough, with/without fever), which affects children younger than two years of age.

Exclusion criteria

Patients with previous wheezing episodes.

Data gathering

Data gathering included collecting the consecutive records of cases diagnosed by collaborating doctors in the study as well as the periodical review of databases and lists or copies of reports for the records of cases diagnosed by other doctors.

We designed a questionnaire for the collection of the study's variables that included general data, signs-symptoms, risk factors, diagnostic tests, and treatments. A complete description of these items is available in a previous article.⁷ We designed a score of the severity of disease by gathering the variables that have been shown in previous studies to have an adequate interobserver concordance, including the following: respiratory rate (<45; 45-60; >60 per minute), pulmonary ventilation (normal; hypoventilation; silent chest), wheezing (mild expiratory; all expiration; expiratory and inspiratory), retractions (not or mild intercostal; moderate intercostal-suprasternal; severe or nasal flaring), and consciousness (normal; agitated; lethargic); these variables were measured after adequate aspiration of secretions (0-2 for each component; maximum score of 10). The treatments were differentiated according to their use in the acute or maintenance phases of the disease. We considered acute phase treatments in inpatients: those received during admission; in ambulatory patients: treatments administered at the place of diagnosis and those recommended during the following 24 h.

The treatment was classified according to its appropriateness following the recommendations of the consensus conference as: first choice, alternative or inappropriate.^{4,7} Patients admitted to the intensive care unit (ICU) were excluded from this classification. The consensus conference was conducted and published after gathering the cases, so that it did not influence in the management of patients.

Ethical aspects

It was specifically recommended not to modify, in any way, the routine management of patients with bronchiolitis. Data were gathered anonymously without registering the patients' identifying data.

Statistical aspects

Statistical processing was performed with SPSS version 11.5.1 (serial number 9036057). We did not conduct an estimation of the sample size necessary for each setting because in almost all of the centres, all of the patients diagnosed with bronchiolitis were included. However, we had calculated that a subsample of 300 patients would allow the estimation of percentages with a precision of $\pm 5\%$ as well as the ability to discriminate differences of at least 12% (for theoretical most unfavourable percentages of 50%, α value of 5%, and β of 20%).

We estimated confidence intervals (CI) for the main measurements. We compared the variables by health care setting (offices, emergency, hospitalisation, and intensive care) using the χ^2 test or exact tests for the qualitative variables and variance analysis for the quantitative variables.

Results

Between October 2007 and March 2008, we gathered 5647 cases of bronchiolitis from 31 hospitals and 60 primary care centres/offices. The cases were predominantly diagnosed in the emergency departments (2914; 51.6%) and in hospitalisation wards (1576; 27.9%); 1060 (18.8%) were made in primary care offices, and 86 (1.5%) were made in the ICU. The health care setting was not specified in 11 cases.

Of the cases, 1874 (34.7%) children required hospitalisation. Because of the system used for gathering cases, this percentage did not allow for the calculation of overall risk upon admission; however, this was estimated for two of the subsamples: 2655 cases gathered from the emergency wards of ten tertiary hospitals, with 23.5% of admissions (CI 95%: 21.8–25.1%), and 1142 cases from primary care centres (including 181 cases diagnosed at hospital level with 9.1% of admissions; CI 95%: 7.5–11.0%). The mean hospital stay was 5.73 days (CI 95%: 5.54–5.92).

A total of 58.1% of the cases were male, and no differences were observed between different health care settings with regard to gender. The mean age was 0.34 years (CI 95%: 0.32–0.35) with a predominance of children between three months and one year of age (Table 1). Clinical presentation varied by health care setting, and the highest presence of all symptoms was seen among the hospitalised patients (Table 1). Hospitalised and ICU-admitted children had a higher level of respiratory failure, which was reflected in higher scores on the severity score (Table 1).

A total of 11.1% of cases were preterm born (2.6% with ≤ 32 weeks of gestation), and 2.3% of cases had congenital heart disease. Other risk factors were infrequent and included the following: bronchopulmonary dysplasia (0.9%), other chronic lung diseases (0.2%), immunodeficiency (0.1%), and neuromuscular disease (0.11%). Amongst the hospitalised patients, there was a slightly higher frequency of preterm born (hospitalised 13.9%, emergency services 9.9%, offices 9.4%, and ICU 22.4%; $p < 0.001$), congenital heart disease (hospitalised 3.5%, emergency services 1.8%, offices 1.6%, and ICU 2.4%; $p = 0.002$), and bronchopulmonary dysplasia (hospitalised 1.7%, emergency services 0.8%, offices 0.1%, and ICU 1.2%; $p < 0.001$).

Table 2 shows the treatments used during the acute and maintenance phases of bronchiolitis. Fig. 1 shows a comparison of the percentages of use of the main treatments in each health care setting. Table 3 shows the classification of appropriateness of treatments by health care setting, excluding ICU patients.

Discussion

The characteristics of our study, including the number of cases (5647 infants younger than two years of age with a first episode of bronchiolitis from 91 sanitary centres and 25 provinces from 12 autonomous communities) and location of the study (12 autonomous communities) at different health care settings (primary care consults, emergency department, hospitalisation, and ICU), allowed us to obtain representative data of the epidemiological characteristics and diagnostic-therapeutic management of bronchiolitis in

Spain. The researchers prospectively gathered consecutive cases of bronchiolitis during the epidemic period of 2007–2008. They collected the actual management of bronchiolitis in clinical practice, conducted according to their physicians' criteria.

The most frequent types of studies regarding the treatment of bronchiolitis are surveys of physician opinions from different settings and specialities,^{9–18} cross-sectional studies or case series,^{19–24} reviews of medical records,^{25,26} some cohort studies,^{21,27,28} and interventional studies (with previous and posterior analysis to the implementation of a CPG or consensus).^{18,29–33}

In this study, we observed a wide use of bronchodilators, corticosteroids, and other treatments of unclear efficacy (antibiotics, oral bronchodilators, inhaled steroids, ipratropium bromide, etc.). Some treatments had a greater tendency of use associated with greater disease severity; thus, hospitalised patients and ICU patients had a higher use of nebulised adrenaline, antibiotics, parenteral steroids, oxygen therapy, intravenous fluids, and other treatments of selective use (nasal continuous positive airway pressure, heliox, and mechanical ventilation). However, other treatments, including inhaled bronchodilators and inhaled and oral corticosteroids, do not reflect this tendency and were used in a similar manner within the different health care settings.

Most of the hospitalised patients received bronchodilators (69.3% inhaled beta 2 agonists and 30.1% nebulised adrenaline) and approximately one-third systemic steroids. Patients of emergency wards received in similar proportion inhaled beta 2 agonists (63.2%) but less frequently nebulised adrenaline (13.1%) and systemic steroids (13%). In ambulatory patients, we had expected a lower use of inhaled beta 2 agonists and corticosteroids because they were less severe cases. However, they mainly received inhaled beta 2 agonists (64.1%) and frequently systemic steroids (25.8%). As maintenance phase treatment, half of the patients received inhaled or oral bronchodilators, and one-fifth received corticosteroids with small, although significant, differences according to the origin of the cases.

In surveys regarding routine management of patients with bronchiolitis, there is a generalised use of beta 2 agonists (between 44% and 99% of responses), although only a portion of them in a systematic manner (between 2% and 55%).^{9,10,12–15,18} Adrenaline use was indicated by 20–55% of the people surveyed (1–5% for systemic use), inhaled corticosteroids by 24–54%, and antibiotics by 3–69% (0–4% systemically). The wide use of treatments reflected in the surveys contradicts the recommendations of guidelines and protocols available in their respective countries. Only two studies, carried out in Ireland and Australia, show a low use of bronchodilators and corticosteroids.^{16,17}

Additionally, in studies regarding the standard management of patients with bronchiolitis, beta 2 agonists (42–94%), adrenaline (24–69%), corticosteroids (25–85.7% in hospitals; 5–13% in emergencies; and 18.5% in ambulatory), and antibiotics (24–65%) were predominately used.^{19–33} Several studies conducted have shown a different profile with lower use of beta 2 agonists and corticosteroids and generalised use of respiratory physiotherapy.^{23,31,33} With respect to the use of nebulised hypertonic saline, for which recent evidence suggests a certain efficacy,

Table 1 Frequency distributions for the main variables. Distribution according to the place of diagnosis and total.^a

	Hospitalisation 1576		Emergency 2914		Offices 1060		ICU 86		Total 5647		p
	n	%	n	%	n	%	n	%	n	%	
Age											<0.001
Neonates	223	14.2%	106	3.6%	20	1.9%	23	26.7%	372	6.6%	
1–3 months	591	37.5%	539	18.5%	143	13.5%	47	54.7%	1320	23.5%	
>3–11 months	666	42.3%	2008	69.1%	723	68.3%	14	16.3%	3411	60.6%	
≥ 12 months	95	6.0%	253	8.7%	172	16.3%	2	2.3%	522	9.3%	
Clinic											<0.001
Temperature at diagnosis											
<37 °C	569	37.1%	1208	45.1%	573	57.1%	42	49.4%	2392	45.1%	
37–37.9 °C	591	38.6%	845	31.5%	226	22.5%	33	38.8%	1695	32.0%	
>38 °C	372	24.3%	627	23.4%	204	20.3%	10	11.8%	1213	22.9%	
Cough	1497	96.8%	2684	95.9%	1043	99.8%	77	97.5%	5301	96.9%	<0.001
Night cough	1037	92.3%	1414	89.2%	913	96.5%	22	95.7%	3386	92.1%	<0.001
Rhinitis	1283	84.0%	2218	81.5%	824	80.5%	73	97.3%	4398	82.3%	<0.001
Dehydration	27	1.7%	7	0.2%	8	0.8%	2	2.5%	44	0.8%	<0.001
Vomiting	434	28.7%	566	21.2%	208	20.7%	21	30.4%	1229	23.4%	<0.001
Feeding rejection	970	63.6%	1000	37.3%	379	37.8%	59	77.6%	2408	45.6%	<0.001
Apnoea	80	5.2%	38	1.4%	19	1.9%	29	36.3%	166	3.1%	<0.001
Septic picture	26	1.7%	10	0.4%	6	0.6%	9	11.1%	51	0.9%	<0.001
Severity score at diagnosis											<0.001
Mean (SD)	2.4	(1.8)	1.5	(1.5)	1.6	(1.7)	4.0	(1.9)	1.8	(1.7)	
Percentile (25th and 75th)	1.0	4.0	0.0	2.0	0.0	2.0	2.0	5.3	0	3	
Score ≥4	182	13.1%	89	3.8%	56	5.6%	30	45.5%	357	7.4%	
Oxygen saturation < 94%	486	50.7%	438	15.1%	163	15.7%	66	76.7%	1463	26.1%	

SD, standard deviation; ICU, intensive care unit.

^a With some variables, there are cases of unspecified data; thus, the counts do not add up to the total. The place of diagnosis was not specified in 11 cases.

neither our study nor other previously published studies allowed for the description of its implementation in clinical practice.

Due to the extensive information available, the following are well known about the treatment of bronchiolitis^{7,34}: (1) the use of symptomatic support measures is fundamental

for the management of fever, respiratory secretions, hyporexia, respiratory distress and hypoxaemia; (2) the alternative use of a therapeutic trial with inhaled beta 2 agonists or adrenaline (better with hypertonic saline) can be considered in selected moderate-severe cases and maintained only if there is a positive documented response

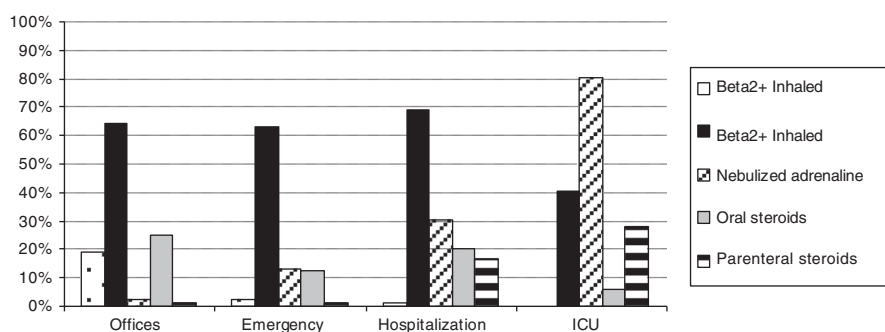


Figure 1 Use of bronchodilators, adrenaline, and corticosteroids in each health care setting (percentages).

Table 2 Main treatments before diagnosis and during the acute and maintenance phases. Distribution according to the place of diagnosis and total.^a

Treatments	Hospitalisation 1576		Emergency 2914		Offices 1060		ICU 86		Total 5647		p
	n	%	n	%	n	%	n	%	n	%	
<i>Acute phase</i>											
Oxygen	719	45.9%	301	10.4%	149	14.4%	61	70.9%	1230	22.0%	<0.001
Intravenous fluids	607	38.9%	125	4.3%	28	2.7%	64	74.4%	824	14.8%	<0.001
Oral Beta 2+	18	1.2%	63	2.2%	197	19.1%	0	0%	278	5.0%	<0.001
Inhaled Beta 2+	1084	69.3%	1828	63.2%	669	64.1%	35	40.7%	3616	64.7%	<0.001
Oral antibiotics	148	9.5%	103	3.6%	127	12.2%	7	8.1%	385	6.9%	<0.001
i.v./i.m. antibiotics	274	17.6%	48	1.7%	6	0.6%	42	48.8%	370	6.6%	<0.001
Respiratory physiotherapy	223	14.4%	81	2.8%	85	8.3%	13	15.3%	402	7.2%	<0.001
Nebulised adrenaline	470	30.1%	379	13.1%	26	2.5%	69	80.2%	944	16.9%	<0.001
Oral corticosteroids	314	20.3%	358	12.4%	259	24.9%	5	5.9%	936	16.8%	<0.001
Inhaled corticosteroids	101	6.5%	91	3.1%	63	6.1%	8	9.4%	263	4.7%	0.001
Parenteral corticosteroids	254	16.4%	36	1.2%	9	0.9%	24	28.2%	323	5.8%	<0.001
Ipratropium bromide	131	8.4%	127	4.4%	17	1.6%	7	8.2%	282	5.1%	<0.001
Nasal CPAP ^b	44	2.8%	16	0.6%	1	0.1%	41	47.7%	102	1.8%	<0.001
Heliox ^b	26	1.7%	5	0.2%	0	0.0%	25	29.1%	56	1.0%	<0.001
Inhaled Ribavirin	1	0.1%	6	0.2%	0	0.0%	0	0.0%	6	0.1%	0.284
Mechanical intubation ^b	17	1.1%	7	0.2%	1	0.1%	14	16.3%	38	0.7%	<0.001
Antipyretic	632	42.2%	566	20.0%	265	26.4%	33	38.4%	1496	27.6%	<0.001
Humidification	190	12.6%	144	5.1%	279	27.9%	2	2.3%	615	11.4%	<0.001
Nasal irrigation	888	60.4%	1089	39.8%	654	66.2%	21	26.3%	2652	50.3%	<0.001
Aspiration of respiratory airway	830	55.0%	787	27.8%	391	39.0%	26	30.2%	2034	37.5%	<0.001
<i>Maintenance</i>											
Antitussives	7	0.5%	13	0.5%	28	2.8%	0	0.0%	48	0.9%	<0.001
Mucolytic decongestants	55	3.6%	71	2.6%	52	5.2%	6	7.1%	184	3.5%	<0.001
Oral Beta 2+	107	6.9%	347	12.9%	76	7.6%	1	1.2%	531	10.0%	<0.001
Inhaled Beta 2+	732	47.2%	1386	51.7%	402	39.9%	19	22.6%	2539	47.7%	<0.001
Antibiotics	213	13.8%	191	7.1%	74	7.3%	12	14.3%	490	9.2%	<0.001
Oral corticosteroids	253	16.4%	445	16.5%	87	8.7%	10	11.9%	795	14.9%	<0.001
Inhaled corticosteroids	113	7.3%	62	2.3%	57	5.8%	7	8.3%	239	4.5%	<0.001
Ipratropium bromide	62	4.0%	20	0.7%	6	0.6%	0	0.0%	88	1.7%	<0.001
Montelukast	13	0.8%	8	0.3%	29	2.9%	0	0.0%	50	0.9%	<0.001

CPAP, continuous positive airway pressure; ICU, intensive care unit; i.m., intramuscular; i.v., intravenous.

^a With some variables, there are cases of unspecified data; thus, the counts do not add up to the total. The place of diagnosis was not specified in 11 cases.

^b Some of the patients diagnosed in a consult or emergency service who required admission received these treatments during admission (CPAP, heliox, assisted ventilation).

Table 3 Appropriateness of the treatments in the acute and maintenance phases. Distribution according to the place of diagnosis (ICU patients excluded) and total.^a

Treatments	Hospitalisation 1576		Emergency 2914		Offices 1060		Total 5550		p
	n	%	n	%	n	%	n	%	
Acute phase									<0.001
<i>First choice</i>	130	8.2%	597	20.5%	158	14.9%	885	15.9%	
<i>Alternative use</i>	788	50.0%	523	17.9%	118	11.1%	1429	25.7%	
<i>Inappropriate use:</i>	658	41.8%	1794	61.6%	784	74.0%	3236	58.3%	
Beta 2+ or Adrenaline in moderate-severe ^b	558	35.4%	446		86	8.1%	1090	19.6%	
Systemic corticosteroids associated with bronchodilators in moderate-severe ^b	230	14.6%	77	2.6%	32	3.0%	339	6.1%	
<i>Inappropriate use:</i>									
Beta 2+ or Adrenaline in mild	20	1.3%	1199	41.1%	301	28.4%	1520	27.4%	
Systemic corticosteroids in mild	9	0.6%	159	5.5%	98	9.2%	266	4.8%	
Other inappropriate treatments ^c	625	39.7%	219	7.5%	193	18.2%	1037	18.7%	
Various inappropriate ^c	4	0.3%	217	7.4%	192	18.1%	413	7.4%	
Maintenance phase^d									<0.001
<i>First choice</i>	634	40.2%	1073	36.8%	496	46.8%	2203	39.7%	
<i>Alternative use</i>	465	29.5%	291	10.0%	69	6.5%	825	14.9%	
<i>Inappropriate</i>	477	30.3%	1550	53.2%	495	46.7%	2522	45.4%	

^a With some variables, there are cases of unspecified data; thus, the counts do not add up to the total.

^b Patients hospitalised or with a severity score ≥ 4 or with an oxygen saturation $\leq 94\%$.

^c Antibiotics, oral salbutamol, inhaled corticosteroids, ipratropium, ribavirin (in severe cases), and physiotherapy. When these treatments were associated with the use of bronchodilators and/or corticosteroids in mild cases they were classified as "various inappropriate".

^d Inappropriate treatments during maintenance phase: corticosteroids (inhaled or systemic), Methylxanthine, Montelukast and bronchodilators (when they were not indicated for use in the acute phase).

(clinical severity score); (3) the use of certain drugs (heliox, surfactant, and/or ribavirin) in well selected severe cases of bronchiolitis can be considered; and (4) the use of the majority of the remaining drugs is considered inappropriate (corticosteroids, oral salbutamol, subcutaneous adrenaline, ipratropium, antibiotics, immunoglobulins, etc.). These recommendations, which were obtained from our consensus conference,^{7,34} are concordant with those of other guidelines previously available.¹⁻³

Upon classifying the appropriateness of our treatments, following the established criteria in the consensus conference^{7,34} and even assuming the optimal or alternative use of certain interventions (trial of bronchodilators with or without corticosteroids in moderate-severe cases), we found that in our study, 58.3% of the treatments in the acute phase (somewhat higher in offices) and 45.4% in the maintenance phase (somewhat higher in emergency services) were inappropriate, which is a clear example of overuse of non-efficient treatments.

The use of bronchodilators and/or corticosteroids, the main cause of inappropriateness in our study, deserves a commentary. Adrenaline is frequently used in inpatients but it has only showed a small effect in reducing the admission risk in ambulatory patients (number needed to treat 17).³⁵ Inhaled beta 2 agonists are also widely used, despite there being enough evidence against their efficacy. With respect to systemic corticosteroids, only some studies have shown a small improvement in clinical scores in inpatients, which has no effect in a reduction of the length of stay. The efficacy of combined nebulised adrenaline plus systemic corticosteroids was discussed in our consensus conference. According to the results of a clinical trial published by Plint et al.³⁶ this combination could slightly reduce the risk of admission on day seven (although the mainly effect is seen in the first hour after a single dose of adrenaline). Nevertheless, this effect was no longer significant after adjustment for multiple comparisons (four treatment groups) and the treated group had a higher atopic risk (non-significant but of the same size as

the observed effect). A recent systematic review has considered this study to support the effect of dexamethasone plus nebulised adrenaline,³⁵ but this was not mentioned in previous reviews.³⁷ Until new studies specifically designed to test this combined treatment are not available we decided to consider it only as an alternative for moderate-severe patients.

If we had accepted the widespread use of inhaled bronchodilators, associated or not with systemic corticosteroids, our estimation of appropriateness had improved. But are we sure that our patients are not harmed by this treatment? In our opinion, physicians caring for these patients try to reduce their respiratory distress or avoid their admissions. To do that, they use available treatments, which have showed limited efficacy, but they are not aware that in mild cases the benefit-risk ratio may be unfavourable. In order to justify a trial of therapy with bronchodilators, we must be aware of the limited efficacy and the associated risks, and inform the family about them. We can resort to bronchodilators if we programme them for a short period of time and if we are willing to reassess their efficacy and tolerance before extending their use.

Ethical disclosures

Protection of human and animal subjects. The authors declare that no experiments were performed on humans or animals for this study.

Confidentiality of data. The authors declare that no patient data appear in this article.

Right to privacy and informed consent. The authors declare that no patient data appear in this article.

Funding

This project was financed by a grant from the Hospital de Torrevieja Foundation between June 2007 and June 2009 (protocol code: BECA0001).

Conflict of interest

No conflicts of interest to report.

Annex 1. Research Group of the aBREVIADO Project: members by regions and centres

Andalucía

Hospital de Torrecárdenas. Almería (MD. Gámez Gómez, J. Battles Garrido, J.E. Cabrera Servilla, I. García Escobar, F. Giménez Sánchez, L. Ruiz Tudela), **C. Salud Candelaria. Sevilla** (A. Fernández Valverde, M.G. Bueno Rodríguez, I. Ramón Faba, M. Praena Crespo).

Aragón

C. Salud Fuentes del Ebro. Zaragoza (J.A. Castillo Laita, R. Macipe Costa), **Hospital Infantil Universitario Miguel**

Servet. Zaragoza (C. Campos Calleja, M.C. García Jiménez, R. Pérez Delgado, Y. Romero Salas).

Asturias

C. Salud Contrueces (M. López Benito), **C. Salud El Llano** (V. Martínez Suárez, M. García Balbuena), **C. Salud Infesto** (I. Mora Gandarillas), **C. Salud La Magdalena** (J.I. Pérez Candás), **C.S. La Felguera** (M. Fernández Pérez, C. Gonzavo Rodríguez), **C. Salud Laviada** (A. Cobo Ruisánchez, B. Yáñez Meana), **C. Salud Natahoyo** (A. Hernandez Encinas), **C. Salud Otero** (B. Domínguez Aurrecochea), **C. Salud Pravia** (M. García Adaro, R. Buznego Sánchez), **C. Salud Tineo** (M. Fernández Francés), **C. Salud Puerta La Villa** (I. Franco, S. Ballesteros), **C. Salud Sama** (M. Benito Martín, A.J. Mira López, M. Fernáñez López), **Hospital Cabueñes. Gijón** (C. Molinos Norniella, C. Pérez Méndez, E. Fernández Fernández, J. Fernández Antuña), **Hospital Central de Asturias. Oviedo** (J. Rodríguez Suárez, S. Jiménez Treviño, F. Álvarez Caro).

Canarias

Hospital Universitario Materno Infantil. Las Palmas de Gran Canarias (S. Todorovic, M.R. García Luzardo).

Cantabria

C. Salud Buelna (A. Bercedo Sanz), **Hospital Marqués de Valdecilla. Santander** (M.J. Cabero Pérez, L. Álvarez Granda, E. Pérez Belmonte).

Castilla y León

Hospital Complejo Asistencial de León. León (S. Lapeña López de Armentia, R. Morales Sánchez, L. Fernández Pérez), **C. Salud Jardinillos. Palencia** (S. Alberola López, I. Pérez García), **C. Salud Pintor Oliva. Palencia** (A.B. Camina Gutiérrez, J. G. Santos García), **C. Salud Venta de Baños. Palencia** (I. Casares Alonso), **C. Salud Villamuriel. Palencia** (A. Cano Garcinuño), **Hospital Río Carrión. Palencia** (C. Urueña Leal, J.M. Andrés de Llano, J.E. Fernández Alonzo, J.M. Bartolomé Porro), **C. Salud Ciudad Rodrigo. Salamanca** (M.C. Sánchez Jiménez, M.J. Estévez Amores), **C. Salud Ledesma. Salamanca** (M. Mendoza Sánchez), **C. Salud Miguel Armijo. Salamanca** (J. López Ávila), **C. Salud San Bernardo. Salamanca** (A. Martín Ruano), **C. Salud Santa Marta. Salamanca** (J. Martín Ruano, B. de Dios Martín), **Hospital Complejo Hospitalario de Salamanca. Salamanca** (S. Fernández de Miguel, J.M. Sánchez Granados, O. Serrano Ayestaran), **Hospital Clínico Universitario. Valladolid** (F. Conde Redondo, A. del Río López, V. Matías del Pozo), **Hospital Río Hortega. Valladolid** (F. Centeno Malfaz, C. Alcalde Martín, B. Bello Martínez, L. Crespo Valderrábano, E. Gutiérrez Abad), **C. Salud Benavente Sur. Zamora** (M.E. Vázquez Fernández), **C. Salud Parada de Molino. Zamora** (A. Cortés Gabaudán), **C. Salud Puerta Nueva. Zamora** (M.M. Miguélez Vara, P. Pérez García), **C. Salud Santa Elena. Zamora** (S. García Vicente), **C. Salud Virgen de la Concha. Zamora** (M. A. Prieto Figuero, M.J.

Piorno Hernández, M^a Jesús Moro Pérez), *Hospital Virgen de la Concha. Zamora* (C. Ochoa Sangrador, A.F. Bajo Delgado, A. Fernández Testa), *Hospital General de Segovia. Segovia* (C. Ortega Casanueva).

Cataluña

ABS Llefià. Badalona. Barcelona (G. Ruiz Aragón), ABS-7 La Salut. Barcelona (P. Aizpurua Galdeano), Hospital Sant Joan de Deu. Barcelona (G. Claret Teruel, S. Fernández Ureña), Hospital Universitari Germans Trias i Pujol. Badalona. Barcelona (M. Méndez Hernández, F. Brossa Guerra, J. Fàbrega Sabaté)- ABS Girona-3. Gerona (R.B. Cortés Marina, E. Fortea Gimeno), ABS Girona-4. Gerona (J.C. Buñuel Álvarez, C. Vila Pablos), Hospital Josep Trueta. Gerona (S. Uriel Prat, Ll. Mayol i Canals).

Comunidad valenciana

C. Salud Acequión. Alicante (C. Buhedo Gordillo, G. Rinero de Campos), *C. Salud El Cabo. Alicante* (M.J. Mateo Moraleda, T. Pérez Martín, A. Redondo, A. Sanguino, B. Sepulcre, B. Serra, A. Tosao), *C. Salud El Campello. Alicante* (J. Galiano Olivares), *C. Salud Guardamar del Segura. Alicante* (C.P. Rico Uriós), *C. Salud Hospital Provincial. Alicante* (M.C. Sirvent Mayor, M.J. Fernández Tarí), *C. Salud La Mata. Alicante* (M.S. Fuggini), *C. Salud Mutxamel. Alicante* (L. Comino Almenara, E. Gutiérrez Roble, A. Melnikova, M. Riva), *C. Salud Rojales i Benijofar. Alicante* (A. Bernabé Gutiérrez, I. Degtyareva), *Hospital de Orihuela. Alicante* (V. Cañadas Olmo, F. Goberna Burguera), *Hospital de San Juan. Alicante* (J.L. Mestre Ricote), *Hospital de Torrevieja. Alicante* (J. González de Dios, C. Rivas Juesas), *C. Salud Gran Vía. Castellón* (E. Fabregat Ferrer, M.J. Palomares Gimeno), *Hospital de La Plana. Villarreal. Castellón* (J. Colomer Pellicer), *C. Salud La Eliana. Valencia* (I. Úbeda Sansano, M. Romero García), *C. Salud de Meliana. Valencia* (A. Plaza Miranda), *Consultorio Auxiliar Albalat de la Ribera. Valencia* (C. Sánchez Medina), *Consultorio Auxiliar Barrio de la Luz. Valencia* (T. Álvarez de Laviada Mulero), *C. Salud Padre Jofré. Valencia* (P. Barona Zamora), *C. Salud Serrería I. Valencia* (M. Asensi Monzó).

Extremadura

C. Salud Talavera la Real. Badajoz (C.M. Gómez Málaga), C. Salud Urbano-I. Badajoz (J.J. Cuervo Valdés), C. Salud Villanueva de la Serena Sur. Badajoz (D. Barroso Espadero).

Galicia

C. Salud Santa Comba. La Coruña (M.E. Amigo Ferreiro), Hospital Arquitecto Marcide. Ferrol. La Coruña (E. García Fernández, A.I. García Villar, R.M. Romaris Santiago, M. Santos Tapia), Hospital Clínico de Santiago. Santiago de Compostela. La Coruña (A. Miras Veiga, F. Martínón Torres, N. Martinón Torres, L. Redondo Collazo), Hospital Virxe da Xunqueira. Cee. La Coruña (M.I. Quintela Fernández), Hospital Monforte. Monforte de Lemos. Lugo (S.A. Fernández Cebrián, M.J. Pita Pérez, F. J. Vadillo González), Hospital

da Costa. Burela. Lugo (A.G. Andrés Andrés, P. Lago Manchado), Hospital Complejo Hospitalario de Ourense. Orense (C. Lorenzo Legerén, M. Berrocal Castañeda, J.M. Iglesias Meleiro), Complejo Hospitalario Universitario de Vigo. Pontevedra (E. González Colmenero, J. Antelo Cortizas, E. García Martínez, A. Ruiz Conde).

Madrid

C. Salud Barrio del Pilar (P. González Rodríguez), *C. Salud Canillejas* (O. Cortés Rico), *C. Salud Entrevías-Área 1* (M. Aparicio Rodrigo), *C. Salud General Ricardos* (G. Orejón de Luna, M.M. Martín Mate), *C. Salud Guayaba* (M. Duelo Marcos, C. Indaberea Iguaran, A. Nuñez Giralda, F. Muñoz Velasco), *C. Salud Juncal* (L. Perdikidis Oliveri), *C. Salud Mar Báltico-Área 4* (J.L. Montón Álvarez, V. Orbe León), *C. Salud Potes. Área 11* (M. Ferrnadero Rodríguez), *Hospital Gregorio Marañón* (M.M. Guerrero, R. Marañón Pardo, A. Peñalba Citores).

País vasco

C. Salud Bidebieta. Guipúzcoa (M. Callén Bleca), *Hospital de Donosita. San Sebastián. Guipúzcoa* (J. Korta Murua, F.J. Mintegui Aramburu, I. Olaciregui Echenique, E. Rezola Arce), *Hospital de Cruces. Baracaldo. Vizcaya* (C. Sánchez Díaz), *Hospital de Cruces. Baracaldo. Vizcaya* (J. Sánchez Echantz).

References

1. American Academy of Pediatrics (AAP). Subcommittee on Diagnosis and Management of Bronchiolitis. Diagnosis and management of bronchiolitis. *Pediatrics*. 2006;118:1774-93.
2. SIGN. Bronchiolitis in children. A national clinical guideline; 2006 <http://www.sign.ac.uk/pdf/sign91.pdf> [accessed 13.11.12].
3. Bronchiolitis Guideline Team, Cincinnati Children's Hospital Medical Center. Evidence-based care guideline for management of bronchiolitis in infants 1 year of age or less with a first time episode, Bronchiolitis Pediatric Evidence-Based Care Guidelines, Cincinnati Children's Hospital Medical Center, Guideline 1; 2010. p. 1-16 <http://www.cincinnatichildrens.org/WorkArea/linkit.aspx?LinkIdentifier=id&ItemID=87885&libID=87573> [accessed 16.11.12].
4. González de Dios J, Ochoa Sangrador C. Grupo de Revisión y Panel de Expertos de la Conferencia de Consenso. Conferencia de Consenso Manejo diagnóstico y terapéutico de la bronquiolitis aguda; 2010 http://www.guiasalud.es/GPC/GPC_463.Bronquiolitis.compl.pdf [accessed 13.11.12].
5. Grupo de Trabajo de la Guía de Práctica Clínica sobre Bronquiolitis Aguda. Fundació Sant Joan de Déu c. Guía de Práctica Clínica sobre Bronquiolitis Aguda. Plan de Calidad para el Sistema Nacional de Salud del Ministerio de Sanidad y Política Social. Guías de Práctica Clínica en el SNS: AATRM. Nº 2007/05.
6. González de Dios J, Ochoa Sangrador C, Grupo Investigador del Proyecto aBREVIADO. Estudio de variabilidad en el manejo de la bronquiolitis aguda en España en relación con la edad de los pacientes. Estudio multicéntrico nacional (Proyecto aBREVIADO). *An Pediatr (Barc)*. 2010;2:4-18.
7. González de Dios J, Ochoa Sangrador C, Grupo de Revisión y Panel de Expertos de la Conferencia de Consenso del Proyecto

- aBREVIADo. Conferencia de Consenso sobre bronquiolitis aguda (I): Metodología y recomendaciones. Revisión de la evidencia científica. *An Pediatr (Barc)*. 2010;72:e1-33.
8. McConnochie KM. Bronchiolitis what's in the name? *Am J Dis Child*. 1983;137:11-3.
 9. Brand PL, Vaessen-Verberne AA. Differences in management of bronchiolitis between hospitals in The Netherlands. *Dutch Pediatric Respiratory Society*. *Eur J Pediatr*. 2000;159:343-7.
 10. Offer I, Ashkenazi S, Livni G, Shalit I. The diagnostic and therapeutic approach to acute bronchiolitis in hospitalized children in Israel: a nationwide survey. *Isr Med Assoc J*. 2000;2:108-10.
 11. Mallory MD, Shay DK, Garrett J, Bordley WC. Bronchiolitis management preferences and the influence of pulse oximetry and respiratory rate on the decision to admit. *Pediatrics*. 2003;111:e45-51.
 12. Martínón Torres F, Contreras Martínón F, Redondo Collazo L, Rodríguez Núñez A, Martínón Sánchez JM. Actitud práctica diagnóstica y terapéutica ante la bronquiolitis aguda del lactante en Galicia (estudio Bronquio-Gal). *Acta Pediatr Esp*. 2007;65:12-20.
 13. Kimpen JL, Schaad UB. Treatment of respiratory syncytial virus bronchiolitis: 1995 poll of members of the European Society for Paediatric Infectious Diseases. *Pediatr Infect Dis J*. 1997;16:479-81.
 14. Barben JU, Robertson CF, Robinson PJ. Implementation of evidence-based management of acute bronchiolitis. *J Paediatr Child Health*. 2000;36:491-7.
 15. Barben J, Hammer J. Current management of acute bronchiolitis in Switzerland. *Swiss Med Wkly*. 2003;133:9-15.
 16. Cahill P, Finan E, Loftus BG. Management of bronchiolitis: current practices in Ireland. *Ir Med J*. 2002;95:167-9.
 17. Babl FE, Sheriff N, Neutze J, Borland M, Oakley E. Bronchiolitis management in pediatric emergency departments in Australia and New Zealand: a PREDICT study. *Pediatr Emerg Care*. 2008;24:656-8.
 18. David M, Luc-Vanuxem C, Loundou A, Bosdure E, Auquier P, Dubus JC. Annullation de la Conférence de Consensus sur la bronchiolite aiguë du nourrisson en médecine générale: évolution entre 2003 et 2008. *Arch Pediatr*. 2010;17:125-31.
 19. Vogel AM, Lennon DR, Harding JE, Pinnock RE, Graham DA, Grimwood K, et al. Variations in bronchiolitis management between five New Zealand hospitals: can we do better. *J Paediatr Child Health*. 2003;39:40-5.
 20. Canalejo González D, García Rodríguez ME, Navas López VM, Sánchez Valderrábanos E, Charlo Molina MT, Alonso Salas MT. Bronquiolitis aguda en pacientes hospitalizados. *Rev Esp Pediatr*. 2004;60:211-6.
 21. Plint AC, Johnson DW, Wiebe N, Bulloch B, Pusic M, Joubert G, et al. Practice variation among pediatric emergency departments in the treatment of bronchiolitis. *Acad Emerg Med*. 2004;11:353-60.
 22. Fernández Díaz M, Fernández EM, Menéndez Arias C, Molinos Norriella C, Viejo de la Guerra G, Solís Sánchez G. Variabilidad del manejo hospitalario de la bronquiolitis por virus respiratorio sincitial en menores de 6 meses en los últimos diez años. *Bol Pediatr*. 2006;46:210-6.
 23. Sebban S, Grimprel E, Bray J. Prise en charge de la bronchiolite aiguë du nourrisson par les medecins liberaux du reseau bronchiolite Ile-de-France pendant l'hiver 2003-2004. *Arch Pediatr*. 2007;14:421-6.
 24. De Raepi F, Pannuti F, Antonelli F, de Seta F, Siani P, de Seta L. Therapeutic approach to bronchiolitis: why pediatricians continue to overprescribe drugs? *Ital J Pediatr*. 2010;36:67.
 25. Christakis DA, Cowan CA, Garrison MM, Molteni R, Marcuse E, Zerr DM. Variation in inpatient diagnostic testing and management of bronchiolitis. *Pediatrics*. 2005;115:878-84.
 26. Mansbach JM, Emond JA, Camargo Jr CA. Bronchiolitis in US emergency departments 1992 to 2000: epidemiology and practice variation. *Pediatr Emerg Care*. 2005;21:242-7.
 27. Luginbuhl LM, Newman TB, Pantell RH, Finch SA, Wasserman RC. Office-based treatment and outcomes for febrile infants with clinically diagnosed bronchiolitis. *Pediatrics*. 2008;122:947-54.
 28. Wang EE, Lahn BJ, Boucher FD, Stephens D, Robinson S, et al. Pediatric Investigators Collaborative Network on Infections in Canada (PICNIC) study of admission and management variation in patients hospitalized with respiratory syncytial viral lower respiratory tract infection. *J Pediatr*. 1996;129:390-5.
 29. Muething S, Schoettker PJ, Gerhardt WE, Atherton HD, Britto MT, Kotagal UR. Decreasing overuse of therapies in the treatment of bronchiolitis by incorporating evidence at the point of care. *J Pediatr*. 2004;144:703-10.
 30. Cheney J, Barber S, Altamirano L, Medico C, Cheney M, Williams C, et al. A clinical pathway for bronchiolitis is effective in reducing readmission rates. *J Pediatr*. 2005;147:622-6.
 31. Halna M, Leblona P, Aissi E, Dumonceaux A, Delépouille F, El Kohen R, et al. Impact de la Conférence de Consensus sur le Traitement Ambulatoire des Bronchiolites du Nourrisson: étude sur 3 années dans le Département du Nord (France). *Presse Medicale*. 2005;34:277-81.
 32. King WJ, Le Paux N, Sampson M, Gaboury I, Norris M, Moher D. Effect of point of care information on inpatient management of bronchiolitis. *BMC Pediatr*. 2007;7:4.
 33. Touzet S, Refabert L, Letrilliart L, Ortolan B, Colin C. Impact of consensus development conference guidelines on primary care of bronchiolitis: are national guidelines being followed. *J Eval Clin Pract*. 2007;13:651-6.
 34. González de Dios J, Ochoa Sangrador C, y Grupo de Revisión del Proyecto aBREVIADo. Conferencia de Consenso sobre bronquiolitis aguda (IV): Tratamiento de la bronquiolitis aguda. Revisión de la evidencia científica. *An Pediatr (Barc)*. 2010;72:e1-42.
 35. Hartling L, Fernandes RM, Bialy L, Milne A, Johnson D, Plint A, et al. Steroids and bronchodilators for acute bronchiolitis: systematic review and meta-analysis. *BMJ*. 2011;342:d1714.
 36. Plint A, Johnson D, Patel H, Wiebe N, Correll R, Brant R, et al. Epinephrine and dexamethasone in children with bronchiolitis. *N Engl J Med*. 2009;360:2079-89.
 37. Patel H, Platt R, Lozano JM, Wang EE. Glucocorticoids for acute viral bronchiolitis in infants and young children. *Cochrane Database Syst Rev*. 2004. CD004878.