ORIGINAL ARTICLE

Non-steroidal anti-inflammatory drug hypersensitivity in children

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Abstract

Introduction: There are rather few publications about hypersensitivity reactions to non-steroidal anti-inflammatory drugs (NSAID) in the paediatric age. In this study, we aimed to assess the frequency of confirmed NSAID hypersensitivity in children with a previous reported reaction to NSAID in order to investigate the role of the drug provocation test (DPT) in the diagnostic workup and to explore the factors associated with confirmed NSAID hypersensitivity. **Methods:** We conducted a retrospective analysis of the clinical files from every patient under 18 years old who attended two Portuguese paediatric allergy outpatient clinics, from January 2009 to August 2014, due to a suspected NSAID hypersensitivity.

Results: We included 119 patients, with a median age of nine years (P_{25} – P_{75} : 5–14). Ibuprofen was the commonest implicated NSAID in the patients' reports ($n=94$ –79%). After DPT, NSAID hypersensitivity was confirmed in nine (7.6%) patients, excluded in 93 (78.2%) and was inconclusive in 17 (14.3%). In the majority ($n=95$ –79.8%), the reaction occurred in the first 24 h after intake. Eighty-four patients (70.6%) reported only cutaneous manifestations and 18 (15.1%) had systemic symptoms. Anaphylaxis represented a relative risk to NSAID hypersensitivity confirmation. No association was found for atopy and the number of previous reactions.

Conclusion: In our study, NSAID hypersensitivity was confirmed in a small proportion of the patients with a previous reported reaction. Ibuprofen was the most implicated drug with urticaria/angio-oedema as the commonest manifestation. Anaphylaxis was associated with confirmed drug hypersensitivity. The drug provocation test was essential to establish the diagnosis.

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Introduction

Non-steroidal anti-inflammatory drugs (NSAID) are a widely used group of medications with antipyretic and analgesic properties. This leads to frequent reports of hypersensitivity reactions to NSAIDs (NSAID-H) by adults as well as children. The reported prevalence of hypersensitivity reactions to NSAIDs in the general population is 0.3% in adults, with similar results in children.¹ In Portugal, the prevalence of self-reported NSAID-H was 0.5% among children attending day-care centres.²

A previous study, which was performed in Portugal among children, showed that, although adverse drug reactions are frequently described in this population, after a thorough investigation, a true drug hypersensitivity will be confirmed in only a few cases.³

In patients evaluated due to a possible NSAID-H, the investigation confirmed this suspicion in a minority of them.^{4,5} The drug provocation test (DPT) with the culprit drug is considered the gold standard to diagnose NSAID hypersensitivity.^{6,7} The prevalence of immunologically mediated reactions to NSAIDs ranges from 0.1% to 3.6%.⁸

There are rather few studies in the paediatric population regarding NSAID hypersensitivity reactions. Atopy, immediate-type reactions or respiratory symptoms as previous manifestations have been identified as risk factors by some authors.⁵

The aim of our study was to assess the frequency of confirmed NSAID hypersensitivity in children with a previous reported reaction to NSAID, to investigate the role of the drug provocation test (DPT) in the diagnostic workup and to explore the factors associated with confirmed NSAID hypersensitivity.

Methods

Study design, setting and participants

We conducted a retrospective analysis of the clinical files from every patient under 18 years old who attended two Portuguese Immunoallergy Departments (one in Lisbon and another in Oporto), from January 2009 to August 2014, due to a suspected NSAID hypersensitivity.

The participating centres were the Immunoallergy Department of Dona Estefânia Hospital, CHLC, in Lisbon and the Immunoallergy Department of Maria Pia Hospital, CHP, in Oporto, which are the two main public paediatric allergy centres in Portugal.

Data collection

Data was retrospectively collected by a standardised questionnaire in order to gather information about NSAID reactions and diagnostic test results. In both allergy centres, the usual diagnostic workup includes a first clinical assessment of the reported reaction followed by a DPT.

The standardised questionnaire comprised questions regarding the chronology of the reaction (acute – if the reaction occurred in the first 24 h after the intake; delayed – if it occurred after 24 h),^{7,8} implicated drugs, clinical

presentation, number of previous reactions, age at the time of the first reaction, drug provocation test result, atopic status (defined as at least one positive skin prick test – SPT – to aero/food allergens) and previous diagnosis of allergic diseases (asthma, atopic eczema, allergic rhinitis, food allergy and chronic urticaria). Anaphylaxis was defined as a severe, life-threatening generalised or systemic hypersensitivity reaction, according to the literature.⁹

Drug provocation test

If there was no contraindication,^{7,8,10} NSAID provocation test was performed to confirm or exclude the presence of hypersensitivity and to classify the reaction.

Open DPTs were conducted by an experienced allergist according to the recommendations of the ENDA (European Academy of Allergology and Clinical Immunology Drug Hypersensitivity Interest Group).¹⁰ The diagnostic workup included single-blind, placebo-controlled DPT if the reported symptoms were subjective or if the open DPT performed ended in an unclear result. Incremental doses of the drug were administered at intervals of 30–60 min, stopping as soon as the first objective symptoms occurred or at the end of the intake of the defined doses of the protocol. The first administration was usually 1/10 of the therapeutic dose, according to the weight and age, and the total cumulative dose was similar to the usual maximal individual intake dose (adjusted to weight and age). Symptoms and vital signs were monitored before, during and at the end of the DPT. Three hours was the minimal period of surveillance after the last drug administration. Written informed consent was obtained from the parents or guardians before the procedure.

The DPT was considered positive if it reproduced the previous reported reaction or elicited objective manifestations.

A NSAID-H confirmation was assumed when the patient had a positive DPT with the implicated or alternative NSAID. A NSAID-H was excluded if the DPT with the implicated drug was negative. The investigation was considered inconclusive when DPT was negative with the alternative NSAID but a DPT with the culprit NSAID was not performed (in this case, a single NSAID-H cannot be excluded), or when a DPT was not performed at all (due to the parent's refusal).

Statistical analysis

An exploratory analysis of the variables of interest (gender, age, presence of atopy and allergic conditions, type of drug, manifestations and chronology, NSAID tolerance) was carried out. The Mann–Whitney *U* test was used for comparisons of differences between medians. A chi-squared test or Fisher Exact Test was used to compare the categorical variables. The level of significance considered was $\alpha = 0.05$. Statistical analysis was performed using IBM SPSS Statistics Version 23® (New York, USA).

Results

We included 119 children, 51.3% boys, with a median age of 9.0 years ($P_{25-75} = 5-14$). Atopy was present in 53 (44.5%) and

Table 1 Demographic characteristics of patients and clinical manifestations of the reported reaction.

	Number of patients
Gender	
Male	61 (51.3%)
Female	58 (48.7%)
Median age (years)	9.0 (P ₂₅₋₇₅ – 5–14)
Atopy	53 (44.5%)
Allergic diseases	
Rhinitis	58 (48.7%)
Asthma	46 (38.7%)
Atopic eczema	8 (6.7%)
Chronic spontaneous urticaria	4 (3.4%)
Food allergy	3 (2.5%)
History of DHR	
With a single NSAID	111 (93.3%)
With multiple NSAID	8 (6.7%)
History of reported reactions	
One reaction	72 (60.5%)
Two or more reactions	47 (39.5%)
Chronology of DHR according to history	
Acute	102 (85.7%)
Delayed	2 (1.7%)
Do not recall	15 (14.7%)
Symptoms	
Mucocutaneous	84 (70.6%)
Urticaria/Urticaria and	49
Angio-oedema	
Angioedema/Facial or periorbital oedema	35
Maculopapular exanthema	27
Others	6
Respiratory	12 (10.1%)
Bronchospasm, cough, wheezing	12
Gastro-intestinal	5 (4.2%)
Vomiting	5
Diarrhoea	1
Systemic	18 (15.1%)
Malaise/pallor/dizziness	1
Anaphylaxis	17

DHR – drug hypersensitivity reaction; NSAID – non-steroidal anti-inflammatory drug.

60.5% had a previous diagnosis of allergic diseases. According to the clinical reports, 111 patients (93.3%) described a reaction to a single NSAID (Table 1).

Prior to the investigation, 70 (58.8%) patients had proven tolerability to at least one alternative NSAID (which was verified after the reported reaction – the patient took an alternative NSAID after the reported reaction but previous to this investigation): 60 (85.7%) referred tolerance to paracetamol, eight (11.4%) to acetylsalicylic acid (ASA), three (4.3%) to metamizole, two (2.9%) to ibuprofen and one (1.4%) to nimesulide.

Table 2 Number of patients and suspected NSAID involved.

Implicated NSAID	Number of patients
Ibuprofen	94
Paracetamol	20
ASA	7
Nimesulide	4
Metamizol	1
Etoricoxib	1
Ketorolac	1

NSAID – non-steroidal anti-inflammatory drugs; ASA – acetylsalicylic acid.

In our sample, 47 (39.5%) patients reported two or more previous reactions (41 with the same drug and six with different NSAID) (Table 1).

Parents reported a previous adverse reaction to other drug class in 24 (20%) of the children, mostly to antibiotics ($n = 17$).

At the time of the first reported reaction, 84 (70.6%) patients had only mucocutaneous manifestations after the intake of the implicated NSAID, 18 (15.1%) had systemic symptoms (17 anaphylaxis), 12 (10.1%) had isolated respiratory manifestations and five (4.2%) referred gastrointestinal symptoms (Table 1).

The most common implicated NSAID in the patients' reaction reports was ibuprofen ($N = 94$; 79%) (Table 2).

The characteristics of the first reaction and the results of the investigation performed in the patients who reported anaphylaxis are detailed in Table 3.

The DPT diagram is presented in Fig. 1. In our sample, 99 (83.2%) patients were challenged with the implicated NSAID (positive in five DPT) and 19 (16%) with an alternative NSAID due to severity of the reactions/patient refusal (positive in four). One patient refused the DPT with the implicated drug as he already had a known tolerance to an alternative NSAID.

The presence of NSAID hypersensitivity was confirmed in nine patients (7.6%) (Table 4), excluded in 93 (78.2%) and inconclusive in 17 (14.3%) patients.

Of the nine patients with proven NSAID-H, seven had a previous diagnosis of an allergic disease (five atopic, sensitised to house dust mites). Of the 93 patients without NSAID-H, 30 had confirmed sensitisation to house dust mites.

Patients with confirmed hypersensitivity were older at the time of the first reaction ($p = 0.008$) than patients in which NSAID-H was excluded. A reported history of anaphylaxis was associated with confirmed NSAID-H ($p = 0.029$) (RR = 5.48, 95% CI 1.69–17.8, $p = 0.0047$). No statistically significant differences were found for asthma, urticaria, atopy and a number of previous reactions. However, patients with asthma or rhinitis presented a higher frequency of confirmed NSAID-H (11.9% versus 6.7% for asthma and 14% versus 3.8% for rhinitis) than those without allergic respiratory disease (Table 5).

Discussion

In the present study, we found a 7.6% frequency of NSAID-H. Ibuprofen was the most frequent implicated drug and

Table 3 Clinical manifestations of the reported reaction, implicated drug and drug provocation tests results in the anaphylaxis cases.

Patient ID	Clinical manifestation	NSAID (implicated)	DPT (implicated drug)	DPT (alternative NSAID)	Diagnosis
A	Angio-oedema/Facial or periorbital oedema; Bronchospasm, cough, wheezing	Paracetamol	Positive	ASA and Nimesulide – Negative	Confirmed
B	Maculopapular exanthema; vomiting	Ibuprofen	Negative	–	Excluded
C	Urticaria; vomiting	Ibuprofen	Negative	–	Excluded
D	Nasal and ocular symptoms; vomiting	Ibuprofen	Negative	–	Excluded
E	Urticaria; Bronchospasm, cough, wheezing; vomiting	Ibuprofen	Negative	–	Excluded
F	Chest tightness, dysphonia, dysphagia; facial angioedema and urticaria	Ibuprofen	Negative	–	Excluded
G	Angio-oedema/Facial or periorbital oedema; angio-oedema of the larynx	Ibuprofen Nimesulide	Not performed	ASA – Positive	Confirmed
H	Angio-oedema/Facial or periorbital oedema; Bronchospasm, cough, wheezing	Ibuprofen	Not performed	Nimesulide – Positive	Confirmed
I	Maculopapular exanthema; Bronchospasm, cough, wheezing	Ibuprofen	Not performed	Nimesulide – Negative	Inconclusive
J	Maculopapular exanthema; Oropharyngeal pruritus	Ibuprofen	Not performed	Meloxicam – Negative	Inconclusive
K	Urticaria; Bronchospasm, cough, wheezing; vomiting	Paracetamol	Negative	–	Excluded
L	Angio-oedema/Facial or periorbital oedema; Bronchospasm, cough, wheezing	ASA	Not performed	Nimesulide – Negative	Inconclusive
M	Urticaria; Bronchospasm, cough, wheezing; vomiting	Ibuprofen	Negative	–	Excluded
N	Urticaria; vomiting and diarrhoea	Ibuprofen	Negative	–	Excluded
O	Urticaria; Bronchospasm, cough, wheezing; Angio-oedema/Facial or periorbital oedema	Ibuprofen	Not performed	Meloxicam and nimesulide – Negative	Inconclusive
P	Angio-oedema/Facial or periorbital oedema; Bronchospasm, cough, wheezing; vomiting	Ibuprofen	Positive	Not performed – tolerance to paracetamol, confirmed after the reaction but before this investigation	Confirmed
Q	Urticaria; Bronchospasm, cough, wheezing; hypotension	Ibuprofen	Negative	–	Excluded

NSAID – non-steroidal anti-inflammatory drug; ASA – acetylsalicylic acid; DPT – drug provocation test.

Total cumulative doses were: ibuprofen 10 mg/kg until 600 mg, paracetamol 15 mg/kg until 1000 mg, nimesulide 100 mg and ASA 20 mg/kg until 1000 mg.

Figure 1 Drug provocation test diagram. DPT – drug provocation test.

Table 4 Characterisation of the patients with confirmed NSAID-H.

Patient ID	Age at the first reaction	Implicated NSAID	Type of reaction	Number of previous reactions	Manifestation	NSAID DPT	DPT manifestation
A 12 yrs Male	5 yrs	Paracetamol	Acute	≥4	Anaphylaxis	Culprit	Bronchospasm and nasal symptoms
G 17 yrs Male	13 yrs	Ibuprofen Nimesulide	Acute	2	Anaphylaxis	Alternative – ASA	Facial angio-oedema
H 11 yrs Male	10 yrs	Ibuprofen	Acute	1	Anaphylaxis	Alternative – Nimesulide	Facial and periorbital angio-oedema
P 14 yrs Male	14 yrs	Ibuprofen	Acute	1	Anaphylaxis	Culprit	Periorbital angio-oedema and ocular pruritus
R 7 yrs Female	6 yrs	Ibuprofen	Acute	1	Urticaria; Angio-oedema/Facial or periorbital oedema	Culprit	Urticaria and periorbital oedema
S 13 yrs Female	11 yrs	Ibuprofen	Acute	1	Urticaria	Alternative – Nimesulide	Hypotension and dizziness
T 8 yrs Male	7 yrs	Nimesulide	Acute	1	Urticaria; Angio-oedema/Facial or periorbital oedema	Alternative – Paracetamol	Urticaria, facial angio-oedema, bronchospasm, cough and wheezing
U 13 yrs Female	13 yrs	Ketorolac	Acute	1	Angio-oedema/Facial or periorbital oedema; Maculopapular exanthema	Culprit	Facial and periorbital angio-oedema
V 8 yrs Male	6 yrs	Ibuprofen Paracetamol	Not clarified	≥4	Angio-oedema/Facial or periorbital oedema	Culprit (paracetamol)	Labial angio-oedema

Yrs – years; NSAID – non-steroidal anti-inflammatory drugs; DPT – drug provocation test.

Table 5 Characterisation of the patients that had confirmed and excluded NSAID-H.

		Confirmed NSAID-H patients (N = 9)	Excluded NSAID-H patients (N = 93)	p-Value
Mean age at the first reaction (years) ^a		9.4	5.5	0.008
Gender	Female (n = 47)	3 (6.4%)	44 (93.6%)	0.657
	Male (n = 55)	6 (10.9%)	49 (89.1%)	
Anaphylaxis ^b	With (n = 13)	4 (30.8%)	9 (69.2%)	0.029
	Without (n = 89)	5 (5.6%)	84 (94.4%)	
Atopy	With (n = 44)	5 (11.4%)	39 (88.6%)	0.430
	Without (n = 58)	4 (6.9%)	54 (93.1%)	
Asthma	With (n = 42)	5 (11.9%)	37 (88.1%)	0.360
	Without (n = 60)	4 (6.7%)	56 (93.3%)	
Rhinitis	With (n = 50)	7 (14%)	43 (86%)	0.140
	Without (n = 52)	2 (3.8%)	50 (96.2%)	
Urticaria	With (n = 4)	0	4 (100%)	0.530
	Without (n = 98)	9 (9.2%)	89 (90.8%)	
Number of previous reactions	1 (n = 61)	6 (9.8%)	55 (90.2%)	0.660
	≥2 (n = 41)	3 (7.3%)	38 (92.7%)	

^a Four patients in whom NSAID-H was excluded did not recall the age at the time of the first reaction.

^b Reported to the implicated NSAID.

urticaria and angio-oedema were the commonest clinical manifestations.

Children often have a history of NSAID reactions, particularly to ibuprofen. According to the literature, in 86% of the single reactors and in 56% of cross-reactors, a NSAID-H is not confirmed.⁶ It is believed that concomitant conditions such as infections, fever, or the use of other drugs might play a role in the pathophysiology of the reaction, eliciting symptoms in these patients.^{6,8}

A well-documented clinical history is essential for the correct diagnosis of NSAID-H and should include a description of the symptoms, time interval between drug administration and the symptoms onset, implicated drugs, drugs tolerated after the reaction, route of administration, number of episodes and underlying diseases.⁸ However, a diagnosis of NSAID-H supported solely by the clinical history may lead to overdiagnosis.⁴

A history of repeated previous reactions was reported to predict a future response to NSAID.⁷ In our study, similarly to the study by Yilmaz et al.⁶ the number of previous reactions was not associated with a positive DPT result.

It was suggested that ibuprofen could be used to detect the cross-reactive type of NSAID-H in children.⁶ According to our results, ibuprofen was the most frequent reported culprit in suspected reactions and the commonest drug involved in challenge-proven NSAID-H. This is in line with Zambonino et al.¹ and Guvenir et al.⁴ This is partially explained because in the paediatric age the prescription of NSAIDs is largely restricted to a few drugs, namely ibuprofen and paracetamol.

During the investigation of a suspected NSAID-H, it is important to check for paracetamol tolerance in all children with cross-intolerance to NSAIDs because there are few medications approved for the treatment of fever or inflammation at younger ages.¹ It has been shown that reactivity to paracetamol among NSAID-H patients ranges from 0% to 25%.^{8,11} In the present study, 2 out of 9 patients (22%) with confirmed NSAID-H reacted to paracetamol.

In this work, as found by other authors,^{1,12,13} urticaria and angio-oedema were the most frequent manifestations reported by the patients, with facial angio-oedema, with or without generalised urticaria, as the most common clinical presentation.^{8,11} In the majority of the patients with cutaneous manifestations (94.0%), NSAID-H was confirmed in only five patients. The low frequency of confirmed diagnosis has been reported by other authors, suggesting that even in patients with a history of urticaria/angio-oedema, the diagnostic procedure could safely start with the offending NSAID.¹⁴

Anaphylaxis is usually considered a contraindication for DPT because of the risk associated. In the work of Zambonino et al.¹ the DPT with the culprit drug was not performed in patients who reported anaphylaxis. However, in the study of Yilmaz et al.⁶ a report of anaphylaxis was not associated with the DPT result. In our work, 17 patients reported anaphylaxis as the clinical manifestation. However, the hypersensitivity was only confirmed in four. These DPT were performed in the majority of anaphylaxis reported cases, as other possible causes, such as viral infections or other drugs, could not be ruled out. Most drug challenges were negative, although this clinical presentation was associated with more positive results than have been described in the literature.⁷

In what concerns atopy, several studies have shown an association between NSAID-H, allergic diseases (asthma, rhinitis) and atopy sensitisation in children and adults, which has been considered a significant risk factor for NSAID-H in young children.^{1,11,15–17} An association particularly with house dust mite sensitisation has been described.¹⁸ The exact mechanism remains unclear.^{15,17} However, this association was not found by other authors.^{4,14}

In our study, patients with confirmed NSAID-H presented a higher frequency of asthma and rhinitis than patients with excluded NSAID-H. Almost half (44.5%) of our patients were atopic. Nevertheless, these differences were not statistically significant possibly due to the low number of positive patients.

In our sample, genders were equally distributed (58 girls and 61 boys). However, in the nine patients with NSAID-H, six were male. These data are consistent with previous reports.¹¹ The male preponderance may be explained considering the association of NSAID-H with atopy and the somewhat increased prevalence of atopic diseases in boys.¹¹

Regarding the median age at the time of the first reaction, patients with confirmed NSAID-H were older than those for which the diagnosis was excluded. This difference, which was previously described,¹⁹ could be related to the fact that younger children are more susceptible to viral infections, causing skin eruptions and leading to a deceptive diagnosis of NSAID-H.

DPT is the gold standard for the NSAID-H diagnosis,^{7,8,10} and a negative result should be reassuring to the patient. Nonetheless, patients frequently continue to avoid the tested NSAID after a negative result,²⁰ emphasising the importance of an effective communication between patients and physicians. There are few published studies about NSAID-H in children. For this reason, we consider that the present study provides additional information about this topic, reinforcing the need to undergo an investigation by a DPT with the implicated NSAID.

Conclusion

We found a low frequency (7.6%) of true NSAID-H in children with a previous NSAID reaction. Ibuprofen was the most frequently implicated drug and urticaria and angio-oedema were the commonest clinical manifestation. Anaphylaxis was a relative risk to confirmed drug hypersensitivity. Despite this, in the face of reported anaphylaxis, the clinical history might not be sufficient to establish a final diagnosis of NSAID-H. The drug provocation test proved to be essential in the NSAID-H diagnosis workup and should be considered, even when systemic symptoms are reported.

Ethical disclosures

Confidentiality of data. The authors declare that they have followed the protocols of their work centre on the publication of patient data and that all the patients included in the study have received sufficient information and have given their informed consent in writing to participate in that study.

Right to privacy and informed consent. The authors have obtained the informed consent of the patients and/or subjects mentioned in the article. The author for correspondence is in possession of this document.

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

Conflicts of interest

None declared.

References

1. Zambonino MA, Torres MJ, Muñoz C, Requena G, Mayorga C, Posadas T, et al. Drug provocation tests in the diagnosis of hypersensitivity reactions to non-steroidal anti-inflammatory drugs in children. *Pediatr Allergy Immunol.* 2013;24:151–9.
2. Martins P, Belo J, Marques J, Papoila AL, Caires I, Araújo-Martins J, et al. Reported drug allergy among children attending day care centers. *Acta Med Port.* 2014;27:444–9.
3. Rebelo Gomes E, Fonseca J, Araújo L, Demoly P. Drug allergy claims in children: from self reporting to confirmed diagnosis. *Clin Exp Allergy.* 2008;38:191–8.
4. Guvenir H, Misirlioglu E, Vezir E, Toyran M, Ginis T, Civelek E, et al. Nonsteroidal anti-inflammatory drug hypersensitivity among children. *Allergy Asthma Proc.* 2015;36:386–93.
5. Cavkaytar O, Yilmaz EA, Karaatmaca B, Buyuktiryaki B, Sackesen C, Sekerel BE, et al. Different phenotypes of non-steroidal anti-inflammatory drug hypersensitivity during childhood. *Int Arch Allergy Immunol.* 2015;167:211–21.
6. Yilmaz O, Ertoylkaragol IH, Bakirtas A, Topal E, Celik GE, Demirsoy MS, et al. Challenge-proven nonsteroidal anti-inflammatory drug hypersensitivity in children. *Allergy.* 2013;68:1555–61.
7. Kowalski ML, Asero R, Bavbek S, Blanca M, Blanca-Lopez N, Bochenek G, et al. Classification and practical approach to the diagnosis and management of hypersensitivity to nonsteroidal anti-inflammatory drugs. *Allergy.* 2013;68:1219–32.
8. Ortega N, Doña I, Moreno E, Audicana MT, Barasona MJ, Berges-Gimero MP, et al. Practical guidelines for diagnosing hypersensitivity reactions to nonsteroidal anti-inflammatory drugs. *J Invest Allergol Clin Immunol.* 2014;24:308–23.
9. Vivolo Aun M, Blanca M, Sabino Garro L, Rosimeire Ribeiro M, Kalil J, Abilio Motta A, et al. Nonsteroidal anti-inflammatory drugs are major causes of drug-induced anaphylaxis. *J Allergy Clin Immunol Pract.* 2014;2:414–20.
10. Aberer W, Bircher A, Romano A, Blanca M, Campi P, Fernandez J, et al. Drug provocation testing in the diagnosis of drug hypersensitivity reactions: general considerations. *Allergy.* 2003;58:854–63.
11. Kidon MI, Kang LW, Chin WC, Hoon LS, Hugo VB. Nonsteroidal anti-inflammatory drug hypersensitivity in preschool children. *Allergy Asthma Clin Immunol.* 2007;3:114–22.
12. Blanca-Lopez N, Torres MJ, Doña I, Campo P, Rondón C, Seoane Reula ME, et al. Value of the clinical history in the diagnosis of urticaria/angioedema induced by NSAIDs with cross-intolerance. *Clin Exp Allergy.* 2013;43:85–91.
13. Kidon MI, Kang LW, Chin CW, Hoon LS, See Y, Goh A, et al. Early presentation with angioedema and urticaria in cross-reactive hypersensitivity to non-steroidal anti-inflammatory drugs among young, Asian, atopic children. *Pediatrics.* 2005;116:e675–80.
14. Zisa G, Riccobono F, Bommarito L, D'Antonio C, Calamari AM, Poppa M, et al. Provocation tests with the offending nonsteroidal anti-inflammatory drugs in patients with urticaria/angioedema reactions. *Allergy Asthma Proc.* 2012;33:421–6.
15. Ayuso P, Blanca-Lopez N, Doña I, Torres MJ, Gueant-Rodriguez RM, Canto G, et al. Advanced phenotyping in hypersensitivity drug reactions to NSAIDs. *Clin Exp Allergy.* 2013;43:1097–109.
16. Asero R. Single NSAID hypersensitivity is associated with atopic status. *Eur Ann Allergy Clin Immunol.* 2015;47:48–53.
17. Sánchez-Borges M, Capriles-Hulett A. Atopy is a risk factor for non-steroidal anti-inflammatory drug sensitivity. *Ann Allergy Asthma Immunol.* 2000;84:101–6.

18. Sánchez-Borges M, Fernández-Caldas E, Capriles-Hulett A, Caballero-Fonseca F. Mite-induced inflammation: more than allergy. *Allergy Rhinol.* 2012;3:e25–9.
19. Rubio M, Bousquet P-J, Gomes E, Romano A, Demoly P. Results of drug hypersensitivity evaluations in a large group of children and adults. *Clin Exp Allergy.* 2012;42:123–30.
20. Bommarito L, Zisa G, Riccobono F, Villa E, D'Antonio C, Calamari AM, et al. Avoidance of nonsteroidal anti-inflammatory drugs after negative provocation test in urticaria/angioedema reactions: real world experience. *Allergy Asthma Proc.* 2014;35:303–6.