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RESEARCH LETTER

Monitoring omalizumab treatment efficacy in chronic urticaria by the basophil activation test

To the Editor,

The mechanism leading to chronic urticaria often remains unclear. Up to half of patients are thought to have functional circulating IgG auto antibodies against the high-affinity IgE receptor (FcεRIα) or, in a few cases, even against IgE.^{1,2} These antibodies are able to release histamine in vitro from basophils and mast cells. The expression of the basophil activation marker CD63 can be induced in healthy subjects by chronic urticaria sera patients with positive autologous serum skin test (ASST).³ A lowering in the CD63 expression after cyclosporine treatment has been assessed.³ The efficacy of omalizumab treatment has been shown in patients with chronic urticaria,⁴⁻⁷ and even in many ASS negative test patients.⁷

We report the case of a 49-year-old female patient, who had suffered from chronic urticaria for 18 months. The severity of the clinical condition progressively increased and soles angio-oedema was added in the last episodes. The disease became uncontrolled despite high doses of antihistamines and low corticosteroid doses every 48 h, so many steroid cycles were required every month. Taking into account that autoimmune aetiology is often related to severe urticaria cases, treatment with omalizumab was indicated. Asthma doses for omalizumab were used. In this case, 300 mg every two weeks was prescribed. Steroid treatment was stopped one week after the first dose, and no new episodes occurred after the beginning of the treatment. Skin prick tests were negative for aliments, latex, anisakis and positive for olea.

Autologous serum skin test was also negative.

Serum complement levels (C2, C4), gave normal results.

Proteinogram, serum immunoglobulins (IgA, IgG, IgM), thyroid function, haemogram and general biochemical serum tests were also normal.

The screening for autoimmune disease, including antithyroid antibodies, was negative.

Basophil activation test was performed before beginning the administration of omalizumab and after several doses of treatment, taking the serum of the patient as stimulus. All the samples were drawn immediately before the next dose administration, thus two weeks after the treatment

with omalizumab. The test was performed with the blood of atopic and non-atopic donors without urticaria, employed to study their basophil response after confrontation with the serum of the patient.

Table 1 Percentage of activated basophils after stimulus with different sera. CU-s, chronic urticaria serum; AID-s, autoimmune disease serum; AU-s, acute urticaria serum; HD-s, healthy donor serum; NA, non-atopic.

Stimulus	% Activated basophils
Donor 1 (NA)	
Negative control	2.50%
Positive control	74.40%
CU-s before treatment	82.3%
CU-s after one dose	2.0%
AI disease serum	3.10%
HD-serum	4.70%
Donor 2 (A)	
Negative control	2.10%
Positive control	13%
CU-s after two doses	2%
AU-s	0.60%
Donor 3 (NA)	
Negative control	2.20%
Positive control	62.5%
CU-s after two doses	13%
AU-s	7.80%
Donor 4 (NA)	
Negative control	5.50%
Positive control	66.2%
CU-s after two doses	11%
AU-s	7.80%
Donor 5 (A)	
Negative control	3.30%
Positive control	18.7%
CU-s after six doses	6%
AU-s	3%
Donor 6 (NA)	
Negative control	4.60%
Positive control	48.20%
CU-s after twelve doses	2.20%

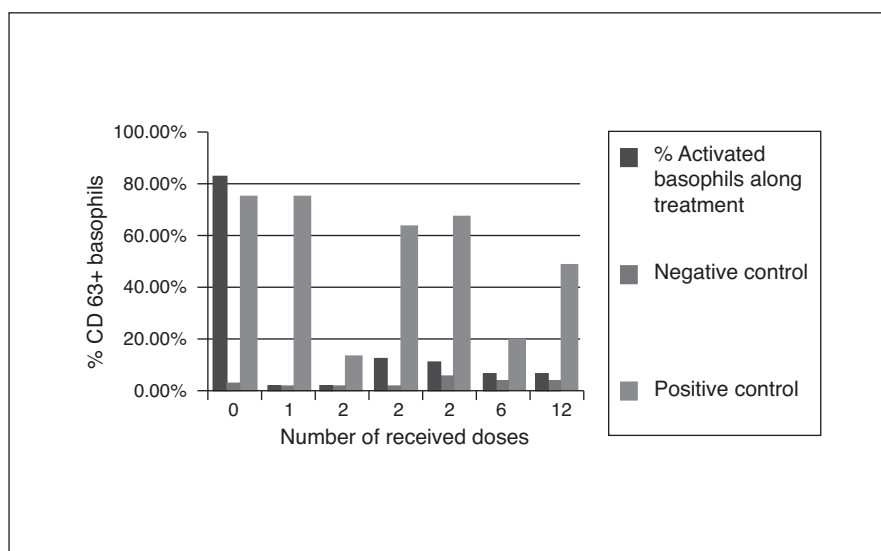


Figure 1 Basophil activation changes throughout the treatment with omalizumab.

Other tests were performed, taking the sera of patients with autoimmune disease as stimulus, to exclude other autoimmune conditions acting as confounding factors; and with the sera of patients with acute urticaria, to evaluate the autoimmune mechanism of chronic urticaria, in contrast with the acute condition. Informed consent was obtained from the patient and the healthy donors.

The assay was performed in accord with the protocol previously described.⁸ The percentage of basophils expressing CD63 with high affinity was the variable used to determine basophil activation. Briefly, a heparinised blood sample was drawn from the donors and aliquotted to test several stimulus. A stimulation buffer (containing IL-3) was first added for ten minutes. Each sample was tested with a negative control (serum saline), a positive control (the chemotactic peptide N-formylmethionyl-leucyl-phenylalanine, that induces basophil activation) and the different sera studied as stimulus. A double blinding was carried out with CD203, to select the basophil population, and CD63, to detect its activation. After a lysing and washing process, the analysis was performed by flow cytometry (approximately 800 basophils per sample). The analysis gate was defined around cells showing high-density CD203c label and low side-scatter, identified as basophils. For inhalants, the test result is considered positive when at least 15% of basophils become activated after the stimulus addition.

Before beginning the treatment, activation even higher than that induced by the positive control was seen in donors (Table 1 and Fig. 1). After one, two, six and twelve doses of treatment, the activation induced by the serum patient decreased to values similar to the negative control or slightly higher than this. In contrast, no evidence of significant activation was seen in donors in whom sera of healthy individuals or patients with autoimmune disease or urticaria were added.

We can conclude that: firstly, in this case, the BAT performed with the serum patient strongly suggested an autoimmune aetiology for the chronic urticaria process and

a possible successful treatment with omalizumab. Secondly, an early decrease in the serum-induced basophil activation coincided with a satisfactory clinical evolution. Moreover, in ASST negative chronic urticaria patients, BAT could be an alternative method to check a possible autoimmune aetiology of the disease. No previous cases of chronic urticaria patients following treatment with omalizumab and monitoring by BAT have been reported.

Conflict of interest

There is not any conflict of interest to disclose. Funding has been completely obtained from Fundesalud (the public foundation for research and education of health care workers in Extremadura) and there has not been any intervention of the Pharmaceutical Industry.

References

1. Hide M, Francis D, Grattan C, Hakimi J, Kochan J, Greaves M. Autoantibodies against the high-affinity IgE Receptor as a cause of histamine release in chronic urticaria. *N Engl J Med*. 1993;328:1599–604.
2. Sabroe RA, Greaves MW. Chronic idiopathic urticaria with functional autoantibodies: 12 years on. *Br J Dermatol*. 2006;154:813–9.
3. Frezzolini A, Provini A, Teofoli P, Pomponi D, De Pità O. Serum-induced basophil CD63 expression by means of a tricolour flow cytometric method for the in vitro diagnosis of chronic urticaria. *Allergy*. 2006;61:1071–7.
4. Kaplan AP, Joseph K, Maykut RJ, Geba GP, Zeldin RK. Treatment of chronic autoimmune urticaria with omalizumab. *J Allergy Clin Immunol*. 2008;122:569–73.
5. Saavedra MC, Sur S. Down regulation of the high-affinity IgE receptor associated with successful treatment of chronic idiopathic urticaria with omalizumab. *Clin Mol Allergy*. 2011; 9:2.
6. Magerl M, Staubach P, Altrichter S, Ardelean E, Krause K, Metz M, et al. Effective treatment of therapy-resistant chronic

spontaneous urticaria with omalizumab. *J Allergy Clin Immunol.* 2010;126:665–6.

7. Spector SL, Tan RA. Effect of omalizumab on patients with chronic urticaria. *Allergy Asthma Immunol.* 2007;99:190–3.
8. Rodríguez-Trabado A, Cámara-Hijón C, Ramos-Cantariño A, Porcel-Carreño SL, Jiménez-Timón S, Pereira-Navarro G, et al. Basophil activation test for the in vitro diagnosis of non-steroidal anti-inflammatory drug hypersensitivity. *Allergy Asthma Proc.* 2008;29:241–9.

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