A comparison between two different in vitro basophil activation tests for gluten- and cow’s milk protein sensitivity in irritable bowel syndrome (IBS)-like patients

Abstract

Background: The diagnosis of food hypersensitivity (FH) in adult patients with gastrointestinal symptoms, beyond the immediate IgE-mediated clinical manifestations, is very often difficult. The aims of our study were to: 1) evaluate the frequency of FH in patients with irritable bowel syndrome (IBS)-like clinical presentation; and 2) compare the diagnostic accuracy of two different methods of in vitro basophil activation tests.

Methods: Three hundred and five patients, (235 females, age range 18–66 years) were included and underwent a diagnostic elimination diet and successive double-blind placebo-controlled (DBPC) challenges. Two different methods of in vitro basophil activation tests (BAT) (CD63 expression after in vitro wheat or cow’s milk protein stimulation) were evaluated: one was performed on separated leukocytes, and the other on whole blood.

Results: Ninety patients of the 305 studied (29.5%) were positive to the challenges and were diagnosed as suffering from FH. BAT on separate leukocytes showed a sensitivity of 86% and a specificity of 91% in FH diagnosis. BAT on whole blood showed a sensitivity of 15%–20% and a specificity of 73% in FH diagnosis (p<0.0001 compared to the other method).

Conclusions: About one third of the IBS patients included in the study were suffering from FH and were cured on the elimination diet. The BAT based on CD63 detection on whole blood samples did not work in FH diagnosis and showed a significantly lower sensitivity, specificity and diagnostic accuracy than the assay based on separated leukocytes.

Keywords: basophils activation test; cytofluorimetry; food hypersensitivity; gluten sensitivity; irritable bowel syndrome.

Introduction

The diagnosis of food hypersensitivity (FH) in adult patients with gastrointestinal symptoms, beyond the immediate IgE-mediated clinical manifestations, is very often difficult. The recent US guidelines recommended that the diagnosis of IgE-mediated FH requires both the presence of allergen-specific IgE and the development of specific symptoms on exposure to that food [1]. However, it is well known that most of the gastrointestinal symptoms due to suspect FH lack an IgE-mediated mechanism [2, 3] and in these cases no laboratory assay is considered useful to make the diagnosis [1, 3].

Patients presenting with irritable bowel syndrome (IBS) who are suspected of FH are included in this difficult diagnostic category. There is general agreement that some of these patients are really affected with FH [2] as demonstrated by several studies [4–6], but no immunology assay is helpful in the diagnosis. A double-blind placebo controlled (DBPC) challenge is the gold standard recommended to confirm the suspicion.

Very recently, we showed that a cytological assay based on in vitro basophil activation after food-antigen stimulation had a sensitivity of 86% in the diagnosis of FH in patients with IBS-like symptoms [7].
The present study included patients with IBS diagnosis and suspected FH. The aims of the study were to: 1) evaluate the frequency of FH in patients with IBS-like clinical presentation; and 2) compare the diagnostic accuracy of two different methods of in vitro basophil activation tests (BAT).

Materials and methods

Study design

The study design contained two parts. In the first retrospective part, we included all patients with IBS diagnosis who had undergone DBPC challenges for suspected FH, consecutively referred as outpatients to the Department of Internal Medicine of the University Hospital of Palermo or the Internal Medicine of the Hospital of Sciacca, Italy from January 2007 to June 2011. The inclusion criteria were: 1) age >17 years; 2) diagnosis of IBS; 3) suspected FH diagnosis evaluated by means of DBPC challenges; 4) follow-up duration longer than 12 months after the initial diagnosis; and 5) >1 outpatient visits during the follow-up period.

Exclusion criteria were: 1) patients with a diagnosis of organic gastrointestinal disease; 2) lack of follow-up visits after the initial diagnosis; and 3) lack of DBPC-challenge method in the FH diagnosis.

IBS diagnosis was based on the Rome II criteria for functional gastrointestinal disorders [8]. Furthermore, organic gastrointestinal disorders were excluded by an accurate work-up, as previously described [7].

In the second prospective part of the study we called back 20 patients (15 female, 5 male): 10 of these patients had tested positive at the DBPC challenges and 10 had tested negative; they were chosen at random from those previously included in the first part of the study. All these patients during the first part of the study had undergone the in vitro BAT based on demonstration of altered membrane phenotypes on allergen-activated basophil, with cytofluorometric detection of CD63 protein, after leukocyte separation. During the second part of the study they underwent the in vitro BAT based on the same assay performed on whole blood; furthermore, they underwent open food challenges again.

None of the enrolled patients were on any medication or were undergoing the elimination diet at the time of the first part of the study. They were asked to suspend medications and/or diet at least 3 weeks before the beginning of the study protocol.

The study was conducted in accordance with the principles of the Declaration of Helsinki. The study protocol was approved by the Ethics Committee of the University Hospital of Palermo and all patients gave their informed consent to participate.

Flow cytometric basophil activation test

After enrolment in the first part of the study, the patients underwent flow cytometric BAT. The assay was performed by different operators, unaware of the clinical history of the patients and the results of the other tests.

From January 2007 to March 2010, the BAT was based for the detection of the surface CD63 protein, after centrifugation and leukocyte separation (230 patients: 170 females, median age 33.5 years). From April 2010 to June 2011 the detection of the surface CD63 protein was performed on whole blood samples (75 patients: 65 females, median age 34 years). Both assays were performed with Flow-CAST® (Bühmann Laboratories AG, Schönenbuch, Switzerland), according to the manufacturer’s instructions.

The manufacturer’s allergens used in all patients were α-lactalamin, β-lactoglobulin, casein, and wheat. As a positive control, we used a monoclonal Le27 anti-IgE antibody solution (Bühmann Laboratories AG). As a negative control, only the stimulation buffer containing IL-3 was used.

The blood tubes were centrifuged at 200 g for 5 min at 4°C. The supernatant containing leukocytes was transferred by pipette into polypropylene tubes and centrifuged again at 500 g for 10 min at 4°C. It was then decanted and the cell pellet was re-suspended in 100 μL of stimulation buffer containing interleukin (IL) 3. The 50 μL of reconstituted solutions of allergens were added to 50 μL of cell suspension and the tubes incubated for 40 min at 37°C. The reaction was stopped by adding 100 μL of cold HEPES buffer (pH 7.3) containing EDTA (HEPES 20 mM, NaCl 133 mM, KCl 5 mM, EDTA 0.27 mM) but with no calcium or magnesium (washing buffer) and the tubes were centrifuged at 1000 g for 5 min at 4°C. The basophils from the cell pellet were double-labeled by adding 20 μL of a mixture of anti-CD63 phycoerythrin-labeled antibody and anti-IgE fluorescein isothiocyanate (FITC)-labeled antibody. After the incubation for 30 min at 4°C (protected from light exposure), 4 mL of a pre-warmed (18–28°C) erythrolytic reagent (lyzing reagent) was added to each tube and left at room temperature for 5±1 min. Cell lysis was stopped with 1 mL of washing buffer, the tubes centrifuged for 5 min at 1000 g, the supernatants were decanted and 500 μL of washing buffer added to each tube. The tubes were gently shaken before flow-cytometric analysis that was performed at once. In each assay, at least 500 basophils were counted.

In order to define a result as positive, the percentage of basophils activated after incubation with antigen should be at least threefold the percentage of basophils activated in the background tube. When the percentage of basophils activated spontaneously was below 2.5%, we required an additional condition, namely that the percentage of basophils activated after contact with the antigen should be ≥5%. These cut-offs were chosen on the basis of receiver operating characteristic (ROC) curves plotted in our laboratory, enabling us to achieve the highest possible sensitivity with optimal specificity.

The intra-assay variation of the test was 3% and was calculated from four blood samples taken from two healthy controls and two patients suffering from food allergy stimulated with stimulation buffer and anti-FceRI Ab and consecutively analyzed 10 times by flow cytometry. The inter-assay variation of the test was 6.5% and was calculated from the blood samples of three healthy controls and three patients suffering from food allergy stimulated with stimulation buffer and anti-FceRI Ab, analyzed 10 times by flow cytometry and re-tested after 2 weeks.

Flow cytometric basophil activation test – CD63 detection on separate leukocytes (old test)

The manufacturer’s allergens used in all patients were α-lactalamin, β-lactoglobulin, casein, and wheat. As a positive control, we used a monoclonal Le27 anti-IgE antibody solution (Bühmann Laboratories AG). As a negative control, only the stimulation buffer containing IL-3 was used.

The blood tubes were centrifuged at 200 g for 5 min at 4°C. The supernatant containing leukocytes was transferred by pipette into polypropylene tubes and centrifuged again at 500 g for 10 min at 4°C. It was then decanted and the cell pellet was re-suspended in 100 μL of stimulation buffer containing interleukin (IL) 3. The 50 μL of reconstituted solutions of allergens were added to 50 μL of cell suspension and the tubes incubated for 40 min at 37°C. The reaction was stopped by adding 100 μL of cold HEPES buffer (pH 7.3) containing EDTA (HEPES 20 mM, NaCl 133 mM, KCl 5 mM, EDTA 0.27 mM) but with no calcium or magnesium (washing buffer) and the tubes were centrifuged at 1000 g for 5 min at 4°C. The basophils from the cell pellet were double-labeled by adding 20 μL of a mixture of anti-CD63 phycoerythrin-labeled antibody and anti-IgE fluorescein isothiocyanate (FITC)-labeled antibody. After the incubation for 30 min at 4°C (protected from light exposure), 4 mL of a pre-warmed (18–28°C) erythrolytic reagent (lyzing reagent) was added to each tube and left at room temperature for 5±1 min. Cell lysis was stopped with 1 mL of washing buffer, the tubes centrifuged for 5 min at 1000 g, the supernatants were decanted and 500 μL of washing buffer added to each tube. The tubes were gently shaken before flow-cytometric analysis that was performed at once. In each assay, at least 500 basophils were counted.

In order to define a result as positive, the percentage of basophils activated after incubation with antigen should be at least threefold the percentage of basophils activated in the background tube. When the percentage of basophils activated spontaneously was below 2.5%, we required an additional condition, namely that the percentage of basophils activated after contact with the antigen should be ≥5%. These cut-offs were chosen on the basis of receiver operating characteristic (ROC) curves plotted in our laboratory, enabling us to achieve the highest possible sensitivity with optimal specificity.

The intra-assay variation of the test was 3% and was calculated from four blood samples taken from two healthy controls and two patients suffering from food allergy stimulated with stimulation buffer and anti-FceRI Ab and consecutively analyzed 10 times by flow cytometry. The inter-assay variation of the test was 6.5% and was calculated from the blood samples of three healthy controls and three patients suffering from food allergy stimulated with stimulation buffer and anti-FceRI Ab, analyzed 10 times by flow cytometry and re-tested after 2 weeks.
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Flow cytometric basophil activation test – CD63 detection on whole blood (new test)

The blood collected into tubes with EDTA was not centrifuged. Fifty microliters of patient’s whole blood were added to 3.5 mL polypropylene tubes for each patient and for each allergen tested and controls, then 50 μL of the corresponding stimulus were transferred by pipette, into 100 μL of stimulation buffer and 20 μL staining reagent (anti-CD63 phycoerythrin with anti-CCR3 fluoroscein isothiocyanate). The tubes were incubated for 15 min at 37°C in a water bath. After the incubation 3.5 mL of a pre-warmed (18–28°C) erythrolytic reagent (lysing reagent) was added to each tube and left at room temperature for 5–10 min. The tubes were centrifuged for 5 min at 500 g, the supernatants were decanted, and 500 μL of washing buffer added to each tube. The tubes were gently shaken before flow-cytometric analysis that was performed at once. In each assay, at least 500 basophils were counted. According to the old method of defining a result as positive, the percentage of basophils activated after incubation with antigen should be at least three-fold the percentage of basophils activated in the background tube. When the percentage of basophils activated spontaneously was <2.5%, we required an additional condition, namely that the percentage of basophils activated after contact with the antigen should be ≥5%.

Wheat and cow’s milk proteins hypersensitivity diagnoses

At entry to the study, those patients who had self-reduced wheat and/or cow’s milk consumption were invited to assume a minimum quantity of 30 g of wheat and 150 mL of cow’s milk daily and were observed for 2–4 weeks on a regular diet. All patients then commenced a standard elimination diet, based on our previous experience [7, 9], with the exclusion of wheat, cow’s milk, eggs, tomato and chocolate. Patients self-reporting FH were also asked to avoid ingestion and/or contact with the food(s) causing symptoms.

Furthermore, food diaries were maintained by the patients during the above periods to assess dietary intake and adherence to the diet.

After 4 weeks on the elimination diet, those patients who became asymptomatic underwent DBPC challenges. The challenges were performed with the reintroduction of a single food at a time; details about DBPC challenges have been previously published [7, 10]. In the case of wheat, the DBPC challenge was performed with capsules coded A or B containing wheat or xylose, respectively. Capsules A or B were given for 2 consecutive weeks and then after 1 week of washout the patients received the other capsules for another 2 weeks (cross-over design).

During all the phases of the study, including the challenge period, the severity of symptoms was recorded: the patients completed a 100-mm visual analog scale, with 0 representing no symptoms, which assessed overall symptoms and the specific symptoms they each reported.

DBPC challenge for cow’s milk proteins was performed with an identical method, according to a standard protocol [7, 10], at least 4 weeks before or after wheat challenge and when the VAS score was <10, on the elimination diet.

The challenges were stopped when clinical reactions occurred (increase in VAS score >30) for at least 2 consecutive days (onset of abdominal discomfort or pain, associated with a change in stool frequency and/or appearance). The challenges were considered positive if the same symptoms which had been initially presented reappeared after their disappearance on the elimination diet.

After FH had been excluded or confirmed, all IBS patients were invited to continue the follow-up with regular visits every 6 months for 1 year.

Statistical analysis

We followed the STARD checklist for studies on the diagnostic accuracy of tests [11]. Tables were constructed for frequency and percentage. The sensitivity, the specificity and the diagnostic accuracy of the immunologic assays, along with their 95% confidence intervals, were calculated by standard statistical methods [12]. The Fisher’s test was used to compare the frequencies. To minimize type I errors, p<0.05 was considered statistically significant. All statistical analyses were performed using the SPSS 11.0 statistical package (Systat Software Inc., CA, USA).

Results

First part of the study

Three hundred and five patients, (235 females, 70 males, age range 18–66 years, median age 34 years), fulfilled the inclusion criteria and were studied. Of these, 105 became asymptomatic on the elimination diet (VAS score <10) and underwent DBPC challenges.

Figure 1 Summary of the challenges results. Seventy patients were positive to both wheat and cow’s milk protein challenge (increase in VAS score >30), 12 patients were positive to wheat challenge only and eight patients were positive to cow’s milk protein challenge only. In total, 90 patients of the 305 studied (29.5%) were positive to cow’s milk protein challenge only. In total, 90 patients of the 305 studied (29.5%) were positive to cow’s milk protein challenge only. In total, 90 patients of the 305 studied (29.5%) were positive to cow’s milk protein challenge only. In total, 90 patients of the 305 studied (29.5%) were positive to cow’s milk protein challenge only.

In contrast, a diagnosis of IBS unrelated to FH was made in the remaining 215 patients (IBS group).

Table 2 shows the results of the fluorometric CD63 assays on separate leukocytes and on whole blood in the IBS and in the FH patients. According to these results, Table 3 shows the sensitivity, specificity and diagnostic accuracy of the two assays in the diagnosis of FH in patients with a clinical presentation of IBS. The in vitro BAT based on CD63 detection on separate leukocytes
showed a higher sensitivity than the same assay on whole blood both in cow’s milk protein- and wheat hypersensitivity diagnosis (p<0.0001). In fact, the sensitivity of the assay based on whole blood was very low. The specificity of the CD63 on separate leukocyte assay was also significantly higher than that of the assay based on whole blood (p=0.0004). As a consequence, the diagnostic accuracy in CMP hypersensitivity and in wheat hypersensitivity diagnoses was significantly higher for the CD63 assay on separate leukocytes than for the assay on whole blood (p<0.0001 for both CMP and wheat).

No statistical difference in sensitivity and specificity was observed for the diagnosis of CMP hypersensitivity considering the positive and negative results obtained with each different antigen used as in vitro stimulants – α-lactalbumin, β-lactoglobulin, casein – both for the assay on separate leukocytes and on whole blood (data not shown).

**Second part of the study**

All 10 patients who were diagnosed as suffering from FH by DBPC challenge during the first part of the study and who were included in the prospective second part were still on the elimination diet at this time. In fact, they were well on the elimination diet and continued to suffer from IBS symptoms assuming wheat and/or foods containing cow’s milk. During this part of the study, they underwent open food challenges which confirmed a persistent FH diagnosis.
check and ten as +95%
been rewritten as +95% CI. Please check and amend if necessary.

Table 2: Results of the BAT based on CD63 detection on separate leukocytes and of the same assay on whole blood in the patients with IBS unrelated to FH and in those suffering with cow's milk protein (CMP) hypersensitivity and/or with wheat protein hypersensitivity.

<table>
<thead>
<tr>
<th></th>
<th>CD63 on separate leukocytes</th>
<th>CD63 on whole blood</th>
</tr>
</thead>
<tbody>
<tr>
<td>IBS unrelated to FH</td>
<td>n=160</td>
<td>n=55</td>
</tr>
<tr>
<td>Positive</td>
<td>12</td>
<td>15</td>
</tr>
<tr>
<td>Negative</td>
<td>148</td>
<td>40</td>
</tr>
<tr>
<td>IBS related to cow's milk protein hypersensitivity</td>
<td>n=58</td>
<td>n=20</td>
</tr>
<tr>
<td>Positive</td>
<td>50</td>
<td>3</td>
</tr>
<tr>
<td>Negative</td>
<td>8</td>
<td>17</td>
</tr>
<tr>
<td>IBS related to wheat protein hypersensitivity</td>
<td>n=62</td>
<td>n=20</td>
</tr>
<tr>
<td>Positive</td>
<td>54</td>
<td>4</td>
</tr>
<tr>
<td>Negative</td>
<td>8</td>
<td>16</td>
</tr>
</tbody>
</table>

Table 2: Results of the BAT based on CD63 detection on separate leukocytes and of the same assay on whole blood in the patients with IBS unrelated to FH and in those suffering with cow's milk protein (CMP) hypersensitivity and/or with wheat protein hypersensitivity.

Q5: +95% CI has been rewritten as +95% CI. Please check and amend if necessary.

The other 10 patients with IBS, diagnosed as not suffering from FH during the first part of the study as they did not improve on the elimination diet or did not worsen on DBPC challenges, continued to consume every kind of food without restrictions.

Table 4 summarizes the results of the in vitro BAT performed with both methods. Using the old method based on CD63 detection after leukocyte separation we had found two false negative results in eight patients with wheat hypersensitivity and one false negative result in eight patients with CMP hypersensitivity. When we performed the new method based on CD63 detection on whole blood, we found a much lower concordance between DBPC challenge results and CD63 assay results. The specificity of both methods was high as no patients who were negative to the DBPC challenge were positive to both assays.

Discussion

Patients suffering from gastrointestinal symptoms and suspected FH very often have a difficult clinical history, characterized by many consultations and investigations without reaching any definite diagnosis. This is the clinical history, in particular, of the patients with IBS-like clinical manifestations [2]. In this respect, the possible relationship between IBS and gluten-sensitivity has been underlined very recently [13], although the pathogenesis of this condition still remains to be clarified and probably different subgroups of patients are wrongly included under an unique “gluten-sensitivity category” [10]. The major clinical problem in ascertaining whether patients with IBS really suffer from GS or FH is the lack of accurate diagnostic assays. These patients could have delayed reaction to foods, not IgE-mediated, and obviously none of the assays based on the IgE detection can be helpful in the diagnosis [3]. For this reason, in this kind of patient, the FH diagnosis must be based on the disappearance of symptoms on the elimination diet and a subsequent positive DBPC challenge. This diagnostic methodology has been used in our center for more than 15 years [14], and in the present study we ascertained that about one third of the IBS patients included were suffering from FH and were cured on the elimination diet. This percentage of FH in IBS was almost constant in the different cohorts we have studied in recent years [6, 7].

However, the problem of a simpler and less cumbersome diagnostic approach still exists. Very recently we have showed that in IBS-like patients, the assay based on the cytofluorometric detection of the in vitro basophil

Q6: +95% CI has been rewritten as +95% CI. Please check and amend if necessary.

<table>
<thead>
<tr>
<th></th>
<th>CD63 assay on separate leukocytes</th>
<th>CD63 assay on whole blood</th>
</tr>
</thead>
<tbody>
<tr>
<td>All IBS patients (n=120)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sensitivity in cow's milk hypersensitivity</td>
<td>86% (76%–93%)</td>
<td>15% (6%–41%)</td>
</tr>
<tr>
<td>Sensitivity in wheat protein hypersensitivity</td>
<td>87% (77%–93%)</td>
<td>25% (1%–50%)</td>
</tr>
<tr>
<td>Specificity</td>
<td>92% (87%–95%)</td>
<td>73% (60%–83%)</td>
</tr>
<tr>
<td>Diagnostic accuracy in cow's milk hypersensitivity</td>
<td>91% (86%–94%)</td>
<td>57% (46%–68%)</td>
</tr>
<tr>
<td>Diagnostic accuracy in wheat hypersensitivity</td>
<td>91% (86%–94%)</td>
<td>59% (68%–86%)</td>
</tr>
</tbody>
</table>
activation (CD63 on separate leukocytes) by food antigens diagnosed FH with 86% sensitivity, 88% specificity, and 87% accuracy. This level of sensitivity was significantly higher than that of serum total IgE or food-specific IgE assays [7]. These data have determined an increased interest for the use of the cytofluorometric assay in FH and gluten-sensitive diagnoses and many colleagues searching for methodological details have written to us. However, the commercial assay which we had used in the past changed methods, and since 2010 a new assay on whole blood replaced the old assay performed on separated leukocytes. This inspired us to perform the present study to evaluate the diagnostic accuracy of the two different methods.

Our results clearly and surprisingly demonstrated that the new BAT based CD63 detection on whole blood samples did not work in FH diagnosis. The retrospective analysis performed on over 200 patients, all of whom underwent DBPC challenges, showed that whereas the old test confirmed very good sensitivity (86%–87%) and specificity (91%) in the FH diagnosis, the new test fell to a sensitivity of 15%–20%. This astonishing lack of sensitivity and diagnostic accuracy in FH diagnosis was confirmed by the second part of the study, when we called back 20 patients who had been previously evaluated with the BAT method based on separated leukocytes and tested them with the new method performed on whole blood samples. In this case also, only two of eight patients with CMP hypersensitivity, or with wheat hypersensitivity, were positive to the new test.

It remains to be clarified what the reasons for such an evident fall in diagnostic accuracy could be. It has been pointed out that whole blood is less stable and functional than the isolated leukocytes, and this considerably reduces its range of use [15]. Furthermore, it has been demonstrated that the presence of allergen-specific IgG antibodies in the plasma may influence the results [15]. This could be an important factor in the patients we studied as the presence of anti-gliadin IgG antibodies in 50%–60% of gluten-sensitive, IBS-like patients has been demonstrated [10, 16] and we have demonstrated an evident overlap between patients “classified as gluten-sensitive” and those “classified as suffering from wheat hypersensitivity” [10]. Another factor which could negatively interfere with the accuracy of the new test is the reduced incubation time with the allergens, compared to the old test on separate leukocytes (15 min vs. 60 min) [17].

Table 4 Comparison of BAT results (CD63 detection) performed after leukocyte separation and on whole blood in 10 patients positive to the DBPC challenges to wheat and/or CMP and in 10 patients negative to the DBPC challenges. False negative results (discordant towards the DBPC results) are marked in Bold.

<table>
<thead>
<tr>
<th>Patients/sex/age</th>
<th>DBPC challenge results</th>
<th>CD63 assay on separate leukocytes</th>
<th>CD63 assay on whole blood</th>
</tr>
</thead>
<tbody>
<tr>
<td>n. 1/F/32 years</td>
<td>Wheat positive/CMP positive</td>
<td>Wheat positive/CMP positive</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 2/F/30 years</td>
<td>Wheat positive/CMP positive</td>
<td>Wheat positive/CMP positive</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 3/F/20 years</td>
<td>Wheat positive/CMP positive</td>
<td>Wheat negative/CMP positive</td>
<td>Wheat negative/CMP positive</td>
</tr>
<tr>
<td>n. 4/F/21 years</td>
<td>Wheat positive/CMP negative</td>
<td>Wheat positive/CMP negative</td>
<td>Wheat positive/CMP negative</td>
</tr>
<tr>
<td>n. 5/F/38 years</td>
<td>Wheat positive/CMP positive</td>
<td>Wheat positive/CMP positive</td>
<td>Wheat positive/CMP negative</td>
</tr>
<tr>
<td>n. 6/F/18 years</td>
<td>Wheat positive/CMP negative</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 7/F/21 years</td>
<td>Wheat negative/CMP positive</td>
<td>Wheat negative/CMP positive</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 8/F/38 years</td>
<td>Wheat negative/CMP positive</td>
<td>Wheat negative/CMP positive</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 9/F/21 years</td>
<td>Wheat positive/CMP positive</td>
<td>Wheat positive/CMP positive</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 10/F/42 years</td>
<td>Wheat positive/CMP positive</td>
<td>Wheat positive/CMP positive</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 11/F/19 years</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 12/M/38 years</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
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</tr>
<tr>
<td>n. 13/M/45 years</td>
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<td>Wheat negative/CMP negative</td>
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</tr>
<tr>
<td>n. 14/M/21 years</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
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</tr>
<tr>
<td>n. 15/F/49 years</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 16/F/38 years</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 17/F/18 years</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 18/M/30 years</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 19/M/20 years</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
</tr>
<tr>
<td>n. 20/F/21 years</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
<td>Wheat negative/CMP negative</td>
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</table>
It is evident that the main advantage of using whole blood in vitro basophil activation tests would be speed and simplicity of manipulation. However, our data strongly discouraged its use in FH diagnosis, and induced us to reconsider the use of the assay based on isolated leukocytes. The commercial assay currently available did not seem to be useful in the complicated diagnostic work-up of the suspected FH.

Conflict of interest statement

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