

# Treatment with *Enterococcus faecalis* CECT7121 Is Not Effective as Therapy in Mice with an Established Allergy Status

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## Abstract

In allergies, an unbalanced immune response towards a T helper (Th) 2 profile with high levels of Immunoglobulin (Ig) E is produced. We have demonstrated that the pre-administration of *Enterococcus faecalis* CECT7121 prevents the development of allergy in ovalbumin-immunized mice. In this work, we evaluated whether this bacterium can also revert an established allergic status. Mice were immunized with ovalbumin (OVA) and after that, were inoculated with an *E. faecalis* CECT7121 suspension. In immunized animals, serum specific immune response, proliferative activity of memory splenocytes, and levels of Th2 cytokines were assessed. The *in vivo* active cutaneous anaphylaxis test was also performed. The treatment with *E. faecalis* CECT7121 only increased anti-OVA IgG2a levels. No differences were observed in other specific immunological parameters. Probiotic-treatment did not prove to have any desensitizing effect on mice. These results, together with those recently published, can be concluded that this bacterium would not be appropriate for the treatment of allergic symptoms.

## Keywords

*Enterococcus faecalis* CECT7121, Ovalbumin-Allergy Murine Model, Immunoglobulin E, Th2 Cytokines, Probiotics

## 1. Introduction

In most individuals suffering from allergic diseases, an imbalance of the immune response towards a Th2 cytokine profile is observed. This Th2 scenario induces

the secretion of high levels of IgE, together with the recruitment of effect or cells to the site where the allergic inflammatory response is taking place [1] [2] [3] [4] [5]. On the other hand, it is known that in healthy individuals, a Th1 response is generated against allergens [6]. According to the hygiene hypothesis [5] [7] [8] [9], the exposition to microorganisms during the early stages of life could revert this Th2-skewed response, and in turn, it could induce the development of a Th1 phenotype and stimulate the activity of Treg cells. This observation has led to the experimental use of microorganisms to prevent or inhibit allergic diseases, being the probiotics, the most promising tool for this purpose [10]. There are prominent studies about probiotic pre-treatment which are successfully employed in allergies in different models. One of the beneficial effects that have been reported is a reduction in the production of serum-specific IgE in mice [11] [12] [13]. Probiotic bacteria can induce immune regulation or immune tolerance in allergic diseases as it had been demonstrated in a  $\beta$ -lactoglobulin-induced intestinal anaphylaxis in a murine model of food allergy where the oral administration of *Clostridium butyricum* CGMCC0313-1 ameliorated intestinal anaphylaxis symptoms and shifted the immune balance towards Th1 and Treg, with significantly increased Foxp3/Ror $\gamma$ t and Foxp3/Gata ratios and a significantly decreased Gata3/Tbet ratio [14]. Furthermore, there are other studies that show that the probiotic administration induces an improvement on allergic symptoms. Yang *et al.* have observed that the treatment for two weeks with *Bifidobacterium infantis* during OVA sensitization attenuated the serum specific IgE and IgG1 secretion as well as reduced Th2-type cytokines in spleen cell supernatants and, after challenge with OVA, probiotic-treated mice showed lower allergic reaction measured as diarrhea than sensitized-mice [15].

*Enterococcus faecalis* strains are frequently isolated from food products, and certain strains have been used as cheese starter and food products in the market worldwide [16]. Moreover, they are also employed as probiotics in many applications for human and animals [17]. We have previously worked with a nonpathogenic strain, *Enterococcus faecalis* CECT7121, isolated from natural corn silage [18]. We have evidenced that this strain implants and remains in the intestinal mucosa of BALB/c mice, stimulating the immune system [19] [20]. We have also demonstrated the adjuvant effect of *E. faecalis* CECT7121 and its broad pro-Th1 immunomodulatory activity, achieving beneficial effects on different experimental models [19] [21] [22] [23].

The immunomodulatory effect of *E. faecalis* CECT7121 has been demonstrated in an allergy model induced by the subcutaneous (s.c.) administration of Ovalbumin (OVA), where this strain was able to diminish the secretion of specific IgE and to induce an increase in the levels of specific IgG2a. This probiotic also decreased the proliferation of splenocytes and the secretion of Th2 cytokines [23]. In the latter model, we have demonstrated that the previous implantation of *E. faecalis* CECT7121 in the intestinal mucosa prevents the development of the allergic status. In order to perform a more insightful assessment of the immunomodulatory capacity of this probiotic, in this work, we evaluated

whether this probiotic can also revert an established allergic status. Considering that probiotic immunomodulatory capacity is broadly assessed as pre-treatment in allergy models, but there is little information about probiotic post-treatment in an established allergy model, in this work we evaluated whether *E. faecalis* CECT7121 can also revert an established allergic status. For that reason, OVA-sensitized mice were treated with the probiotic by the intragastric (i.g.) route, evaluating parameters of the Th-2 biased immune response induced by the allergen.

## 2. Materials and Methods

### 2.1. Mice

Conventional female BALB/c mice were provided by the School of Veterinary Medicine, Universidad de Buenos Aires (Argentina). All animals were housed (n = 5 mice/cage) under specific conditions according to the “Guide for the Care and Use of Laboratory Animals” (National Research Council of the National Academies, USA), with controlled air temperature (20°C - 22°C), humidity, and 12 h light/dark cycles; food and water were provided *ad libitum*. All mice were 6 weeks old at the beginning of the experiment.

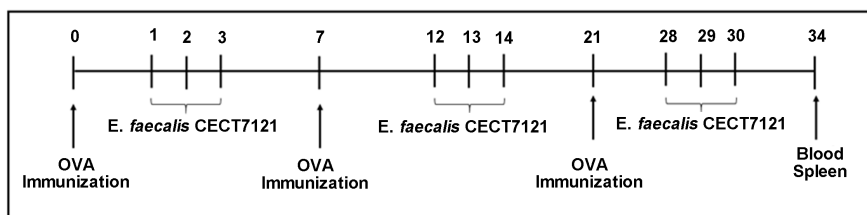
All animal experiments were approved by the Instituto de Estudios de la Inmunidad Humoral “Prof. Dr. Ricardo A. Margni” (IDEHU, CONICET-UBA). Maximum efforts were undertaken to minimize animal suffering in all procedures.

### 2.2. Immunization Protocol with OVA and i.g. Administration of *Ent. faecalis* CECT7121

Mice were divided in two experimental groups: *E. faecalis* CECT7121/ OVA (n = 6), S/OVA (saline) (n = 6). OVA immunization [10 µg OVA and 1 mg Al(OH)<sub>3</sub>] was performed on days 0, 7 and 21 by s.c. injection (0.2 ml) on the back of each mouse (**Figure 1**). An *E. faecalis* CECT7121 suspension ( $3 \times 10^8$  CFU/ml, prepared as previously described [23]) was administered by the i.g. route (0.2 ml/mice/day) with a gavage needle on three doses: days 1, 2 and 3; days 13, 14 and 15 and on days 28, 29 and 30. On day 34, serum samples were obtained and mice were euthanized by CO<sub>2</sub> or cervical dislocation and spleens removed. Another group of mice was subjected to the Active Cutaneous Anaphylaxis Test (ACA). Control non-immunized animals were treated with saline instead OVA or *E. faecalis* CECT7121 (n = 3 each).

### 2.3. Evaluation of Cell Responses

Individual spleen cell suspensions of each mice were prepared in RPMI 1640 medium (Gibco) containing fetal calf serum (Natocor), penicillin (Gibco), streptomycin (Gibco), amphotericin B (Gibco), L -glutamine (Gibco) and pyruvate (J.T. Baker) as previously described [23]. One hundred microlitres of spleen cell suspension ( $4 \times 10^6$  cells/ml) were cultured with 100 µl of culture medium



**Figure 1.** Immunization protocol. Mice were divided in two experimental groups: OVA/*E. faecalis* CECT7121 and OVA/S (saline). OVA immunization [10 µg OVA and 1 mg Al(OH)<sub>3</sub>] was performed on days 0, 7 and 21 with 0.2 ml by s.c. injection (empty arrow) and an *E. faecalis* CECT7121 suspension ( $3 \times 10^8$  CFU/ml) was administrated with a gavage needle by the i.g. route (0.2 ml/mice/day) on days 1, 2 and 3; 13, 14 and 15; and on days 28, 29 and 30. At final day 34, serum samples and splenocyte were taken, and ACA was performed. Control non-immunized animals were treated with saline instead OVA or *E. faecalis* CECT7121.

alone or with 100 µl of OVA (1 mg/ml; Sigma-Aldrich) in quadruplicate at 37°C in presence of 5% CO<sub>2</sub>. Concanavalin A (10 µg/ml; Vector Labs) was used as mitogen. In each experiment, two cell cultures were performed: to assess cell proliferation and to determine the cytokine levels in supernatants. After 72 h, supernatants were collected and stored at -80°C until assayed for cytokine levels. Proliferative responses were assessed after 96 h of culture by [<sup>3</sup>H] thymidine (Perkin- Elmer) uptake and measured by liquid scintillation (Liquid Scintillation Analyzer 1600TR; Packard). Results were expressed as mean counts per minute ± SEM of quadruplicate cultures.

#### 2.4. Levels of Cytokines and Allergen-Specific Antibodies

Commercial kits were employed to determine levels of IL-5, IL-10 (BD Biosciences) and IL-13 (Invitrogen). Specific anti-OVA IgE and IgG antibody levels were measured by indirect ELISA in individual serum samples, employing OVA as coating antigen at 10 µg/ml in phosphate-buffered saline (PBS). A biotinylated anti-mouse IgE antibody (BD Biosciences) followed by an avidine-HRP solution or a HRP-conjugated anti-mouse IgG serum (Cappel) or a HRP-conjugated anti-mouse IgG1 or a HRP-conjugated anti-mouse IgG2a sera (Bethyl) were used as secondary antibodies [23]. To determine the titre of each serum sample, two-fold serial dilutions were performed. All determinations were performed in duplicate. Absorbance values were obtained after spectrophotometric reading (450 - 570 nm) in an ELISA plate reader (Multiskan EX; Thermo Scientific, USA). Antibody titres were calculated as the EC50 (half maximal) value obtained by a four-parameter non-linear regression curve in a log reciprocal-dilution response curve.

#### 2.5. Active Cutaneous Anaphylaxis

Active Cutaneous Anaphylaxis (ACA) was induced by intradermic (i.d.) challenge with the allergen in *E. faecalis* CECT7121/OVA, S/OVA. Briefly, an intravenous injection of 1% w/v Evans Blue (50 µl) was administered by the tail vein,

and ACA was then elicited in the right ear by i.d. inoculation of 50 µl OVA (1 mg/ml). PBS was inoculated in the left ear of each animal as a control reaction. Animals were observed after 30 min.

## 2.6. Statistical Analysis

Two separate immunization schedules were carried out. All values were presented as means with their standard errors. A linear regression of ELISA absorbance values of standards was performed. Student's *t* test was employed to determine significance of specific antibodies results. Cell responses and cytokine levels were analyzed by the Kruskal-Wallis followed by the Dunns multiple comparisons test. In all analysis, GraphPad Prism 5.03 for Windows (GraphPad Software, USA) was used.

## 3. Results and Discussion

The aim of this work was to study the effect *E. faecalis* CECT7121 during the effector phase of the allergic response. We evaluated whether this strain could also revert Th2 parameters of the allergic response in an experimental allergy model induced by s.c. OVA administration. The parameters assessed *ex vivo* were secretion of specific IgE, IgG1, IgG2a antibodies and the Th2 cytokines in splenocyte culture supernatants. In addition, an *in vivo* of the ACA test was performed.

The development of the allergic response comprises two phases. The first one, or sensitization phase, is that CD4<sup>+</sup> T cells become activated and differentiated into Th2 IL-4, IL-5 and IL-13-secreting cells. These cytokines would induce the secretion of IgE and IgG1 by B cells. Both basophils and mast cells have on the cell surface of the high affinity IgE receptor (FcεI), allowing this immunoglobulin to bind to the cell membrane. During a second exposure to the allergen, those IgE molecules bound to the FcεI induce the receptor cross-linking leading to cell activation and the subsequent degranulation and the release of histamine, which is one of the soluble mediators responsible for the appearance of clinical symptoms of allergy. Probiotics as lactic acid bacteria have immunomodulating effects, as a reduction in the production of antigen specific serum IgE production, which is affected by IL-4 and IL-5 produced by Th2 cells [11] [12] [13] [24]. Moreover, Kalliomaki *et al.* have demonstrated, in a double-blind randomized placebo-controlled trial of prevention of atopic disease, that the administration of *Lactobacillus rhamnosus* GG prenatally to mothers and postnatally for 6 months to their infants was effective in prevention of early atopic disease in children [25].

The treatment of allergy employing probiotic microorganisms can be employed to modulate either the sensitization or the effector phase of the response. The immunomodulatory activity of probiotics and the period of time over which they have positive effects are highly dependent on the strain under study, but also with the dose, the timing of supplementation and the method of administration [26] [27] [28]. Previously, our group has demonstrated that the i.g. administration of *Enterococcus faecalis* CECT7121, prior to the sensitization induced

by the administration of OVA, can prevent the establishment of the allergic response [23].

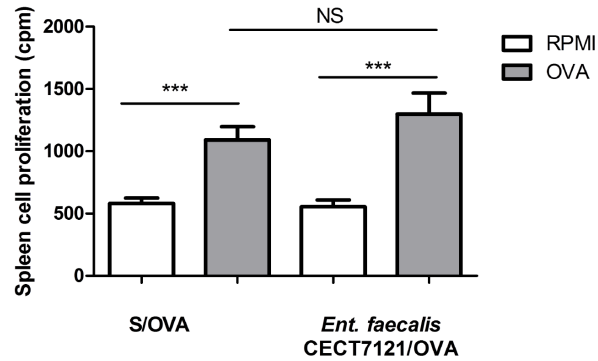
In the present work, on day 34 after the first immunization with OVA and *E. faecalis* CECT7121 treatment protocols, Th1 and Th2 parameters of the immune response were assessed. Levels of specific anti-OVA IgE and IgG1 (Th2-immunoglobulins) and IgG2a (Th1-immunoglobulin) were determined. No detectable specific immunoglobulins were found in non-immunized animals (treated or not with the probiotic; data not shown). Upon analyzing the Th2 immunoglobulin IgE, it was found that the levels of this specific anti-OVA isotype were not affected by the treatment of animals with the probiotic (Table 1). No statistical differences were found in the titres of specific IgG1 between immunized control group and those treated with the probiotic (Table 1). Contrarily, the levels of specific IgG2a were found to be higher in the group of animals immunized with OVA and treated with *E. faecalis* CECT7121 (Table 1). The IgG1/IgG2a ratios, therefore, were found to be significantly different between both groups of animals. The decrease in the anti-OVA IgG1/IgG2a ratios observed in the group of animals treated with the bacterium accounted for a rise in the levels of specific IgG2a. The latter finding is in line with the pro-Th1 activity already reported for *E. faecalis* CECT7121 [19] [21] [22].

After The proliferation rates of splenocytes obtained from both *E. faecalis* CECT7121-treated and untreated control animals did not differ from each other after the *in vitro* stimulation with OVA (Figure 2). Moreover, after stimulation with OVA, these splenocytes were able to secrete the Th2-related cytokines IL-5, IL-13 and the regulatory IL-10; however, no differences were found between animals immunized with OVA and those immunized and treated with *E. faecalis* CECT7121 (Figure 3) which indicates that this bacterium was not able to decrease the levels of the cytokines evaluated (Figure 3). These results are in

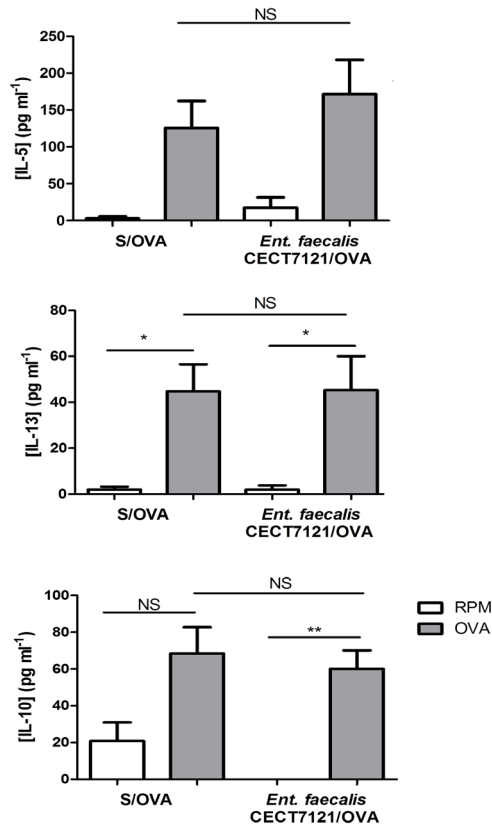
**Table 1.** Determination of specific IgE, IgG, IgG1 and IgG2a levels in serum samples by ELISA after Ova immunization of naïve BALB/c vs. mice treated ig with *E. faecalis* CECT7121.

Antibodies	Mice Group	<i>E. faecalis</i> CECT7121/Ova	S/Ova
anti-Ova IgE (OD450nm)		0.553 ± 0.277	0.629 ± 0.367
anti-Ova IgG (titre)		108.988 ± 15.563	93.561 ± 12.610
anti-Ova IgG1 (titre)		67.057 ± 31.528	59.452 ± 34.763
anti-Ova IgG2a (titre)		5425 ± 2315*	7835 ± 3709
anti-Ova IgG1/anti-Ova IgG2a		12.70 ± 4.88*	8.26 ± 4.08

Antibody titres were defined as the mean inversion of dilution at which the absorbance value is 0.5 of each mouse. Data represent means ± SEM (n:12). \*P < 0.05, Student's t-test.

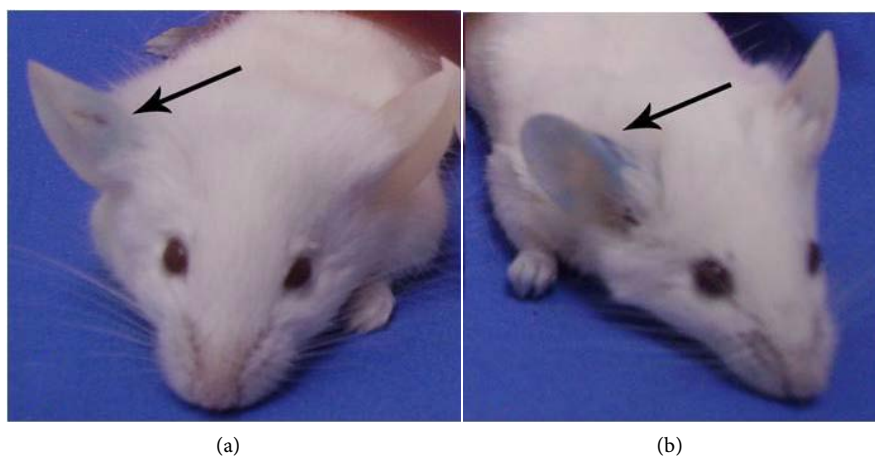


**Figure 2.** Comparative cell responses. On day 34 after first immunization, spleen cells of each immunized mouse were isolated and incubated with culture medium (RPMI) or OVA. Five mice from each group were analyzed in each assay and results expressed as mean count per minute (cpm). Bars represent results of three separate assays. \*\*\*P < 0.001, Kruskal-Wallis followed by the Dunns multiple comparisons test”.



**Figure 3.** Th2 cytokine production. IL-5, IL-13 and IL-10 levels measured by ELISA in splenocytes. IL-5, IL-13 and IL-10 levels measured by ELISA in splenocytes culture supernatants after incubation with culture medium (RPMI) or OVA for 72 h. Five mice from each group were analyzed in each assay and results expressed as pg·ml<sup>-1</sup>. Bars represent results from three separate assays. \*\*P < 0.01, \*P < 0.05, Kruskal-Wallis followed by the Dunns multiple comparisons test.





**Figure 4.** Active cutaneous anaphylaxis. After the intravenous injection of 1% w/v Evans Blue (50  $\mu$ l) by the tail vein, ACA was induced by intradermic challenge with the allergen in OVA-immunized, OVA/*E. faecalis* CECT7121 and control mice (picture not shown) in the right ear by inoculation of 50  $\mu$ l OVA (1 mg/ml). PBS was inoculated in the right ear of each animal as control reaction. Animals were observed after 30 min. a) S/OVA group; b) *E. faecalis* CECT7121/OVA group. Arrows show the inoculation site in immunized animals.

agreement with the titres of specific IgE found in allergic animals, since it is known that IL-4, IL-5 and IL-13 are the cytokines involved in IgE synthesis regulation [2] [29] [30].

Even though the immunological *ex vivo* determinations indicated that *E. faecalis* CECT7121 was incapable of reverting the allergic status, the *in vivo* ACA test was performed, since it has been demonstrated that this test can account for amelioration of the allergic symptoms, as is the case of the treatment with *Lactococcus lactis* NCC2287, even when a decrease in the levels of specific antibodies is not observed [31]. In the ACA test carried out in our model, those animals belonging to the *E. faecalis* CECT7121/OVA group displayed the same degree of dye extravasation as control S/OVA animals (Figure 4), evidencing vasodilatation and a vascular permeability increase caused by mast cells degranulation.

Taking into account the results obtained herein, together with those published demonstrating the preventive effect of *E. faecalis* CECT7121 on the development of allergy [23], it can be concluded that this bacterium as it was administered could decrease the anti-OVA IgG1/IgG2a ratios observed in the group of animals treated with the bacterium due to a rise in the levels of specific IgG2a, but it could not reduce IgE levels nor show differences in the ACA test, so it would not be appropriate for the treatment of allergic symptoms in the OVA allergy induced model. These results reinforce the hypothesis that each probiotic strain can have a differential effect on both phases of the allergic response.

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