Immunological changes in growing mice fed on diets containing casein or peas (Pisum sativum var. Belinda) as the source of protein

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The effects of two different sources of protein: peas (Pisum sativum var. Belinda) and casein on immunocompetence, nutritional utilization and growth performance have been investigated in recently weaned mice. Feeding these animals on a pea diet resulted in an impairment in growth and significant decreases in the weights of liver, muscle, kidneys and femur, while intestine weights increased. No differences in food consumption were observed, but food conversion efficiency (food intake: weight gain) was increased in pea-fed animals compared with those offered the casein diet. Packed cell volume and serum Fe and Zn levels fell significantly after legume-protein intake, and, by contrast, Cu values increased slightly. Serum albumin levels showed a statistically significant reduction in mice fed on the diet containing peas. However, γ-globulins and immunoglobulin G titres were markedly increased. The characterization of spleen-cell subsets using monoclonal antibodies revealed a significantly higher percentage of T-lymphocytes in the pea group compared with casein-fed animals, while no changes were observed in the proportions of B-lymphocytes and macrophages. In vitro mitogenic responses to phytohaemagglutinin, concanavalin A and Escherichia coli lipopolysaccharide S were slightly, but not significantly, lower in the pea-fed animals. Our results describe, apparently for the first time in mice, some immunological disturbances after peak intake. These results may lead to a better understanding of the possible role of antigenic proteins in gastrointestinal disorders and the poor individual performance after legume intake.

Pea (Pisum sativum): Immune response: Nutritional status: Mice

Protein levels and the amino acid composition of the diet play a role in the maintenance of immune response (Bounous et al. 1983; Chandra, 1993; Yamauchi & Suetsuna, 1993). Furthermore, proteins can act as dietary antigens (Perdue & Bienestock, 1991) and the type of protein could affect the immune response through different antigenic or allergic reactions (Le Guen et al. 1991; Hankins et al. 1992; Lallès et al. 1993).

Food-induced allergic responses, especially in young individuals, have been the subject of study in recent years, and alternative protein sources in the treatment of protein allergies have been investigated (Walker, 1992). Moreover, the increasing use of plant proteins in human and animal nutrition has focused great attention on legumes, which are also important sources of other nutrients (Lallès et al. 1993). However, the immunogenicity of certain antinutritional factors such as lectins and some legume proteins seems to indicate that the immune response may be involved in digestive disturbances, impaired availability of different nutrients and reduced growth performance (Barratt et al. 1979; Grant et al. 1983; Lallès et al. 1993). These nutritional imbalances and others produced by the phytate, tannin or fibre content, which diminish mineral bioavailability (Larralde & Martínez, 1989; Cashman & Flynn, 1991; Rubio et al. 1992), may alter immune responses through complex interactions between nutrition and immunocompetence (Sherman, 1992; Chandra, 1993).
The aim of the present study was to evaluate the immunocompetence in mice receiving diets containing peas (*Pisum sativum* var. Belinda) or casein as the source of protein using different indices of the immune response such as the total serum immunoglobulin G (IgG), the mitogenic response of splenic lymphocytes and the percentage of splenic-cell subsets. The nutritional utilization of different nutrients was also assessed. These studies may be of relevance in populations where legumes represent the staple food, but also for nutritional purposes in immunity or allergy-mediated disorders.

**MATERIALS AND METHODS**

*Animals and diets*

Recently weaned male Swiss albino mice (4 weeks old) obtained from Letica S.A. (Barcelona, Spain) and weighing about 21 g were randomly assigned to two dietary groups of eight animals each. The animals were housed in polypropylene cages with wire-meshed bottoms in a room with constant temperature (20–22°C) and with a 12 h light–dark cycle. Feed and water were provided *ad lib.* in feeders specially designed to minimize feed wastage. The diets contained casein (control) or raw pea seeds as the source of protein and were balanced for the remaining nutrients (Table 1). The animals were fed on the same experimental diets for 3 weeks. Mice and feed were weighed daily at the same time (09.00–10.00 hours). Final body and organ weights were assessed at the end of the experimental period as indices of growth performance.

*Packed cell volume and serum mineral measurements*

Mice were ether-anaesthetized and blood samples were obtained by cardiac puncture. Serum Fe was assayed colorimetrically using ferrozine as the chromogen (Thompsen & Motola, 1984), and an Autoanalyser (model Cobas Fara; Roche Diagnostic, Basel, Switzerland). Packed cell volume values, as an index of Fe deficiency, were determined using capillary tubes by centrifugation at 1000 *g* for 10 min.

Serum Zn and Cu determinations were made by atomic absorption spectroscopy (Perkin-Elmer, model 305 B, Norwalk, CT, USA) with an air–acetylene flame at a wavelength of 213.9 nm for Zn and 324.7 nm for Cu (Smith *et al.* 1985).

*Serum protein and IgG determinations*

Serum total proteins, albumin and γ-globulins were measured after agarose-gel electrophoresis (Barta & Porciau, 1984). Samples applied to agarose-gel plates were subjected to 100 V for 20 min, then were scanned in a densitometer (Beckman Instruments, Brea, CA, USA) at 600 nm.

Serum levels of IgG were measured by radial immunodiffusion (Hudson & Hay, 1989) using a kit with anti-mouse IgG antiserum contained in agarose gel (Serotec, Oxford, Oxon). After an appropriate dilution, test samples and standards were applied in 5 *μl* volumes on different wells of the plates. The incubations were carried out at 22°C in moist atmosphere for 72 h. The diameters of the precipitation rings were measured and the concentrations of immunoglobulin in the samples were determined from a calibration curve.

*Isolation of mononuclear cells from spleen*

Spleens were removed aseptically from mice, weighed and briefly stored in NaCl solution (8.5 g/l) at 0°C before being washed and minced in the same solution. After removal of residual tissue, mononuclear cells of the cellular suspension were separated by density gradient centrifugation at 400 *g* for 20 min at room temperature on lymphocyte separation medium (1.083 g/ml; Lymphoprep, Nycomed AS; Böyum, 1983). Mononuclear cells were
Table 1. Composition of experimental diets (g/kg)

<table>
<thead>
<tr>
<th></th>
<th>Casein diet</th>
<th>Pea diet</th>
</tr>
</thead>
<tbody>
<tr>
<td>Casein*</td>
<td>140</td>
<td>—</td>
</tr>
<tr>
<td>Pea (Pisum sativum var. Belinda)</td>
<td>—</td>
<td>600</td>
</tr>
<tr>
<td>Sucrose</td>
<td>350</td>
<td>150</td>
</tr>
<tr>
<td>Wheat starch</td>
<td>350</td>
<td>150</td>
</tr>
<tr>
<td>Olive oil</td>
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<td>50</td>
</tr>
<tr>
<td>Cellulose</td>
<td>50</td>
<td>—</td>
</tr>
<tr>
<td>Mineral mix†</td>
<td>45</td>
<td>40</td>
</tr>
<tr>
<td>Vitamin mix‡</td>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td>Choline chloride</td>
<td>0.6</td>
<td>0.6</td>
</tr>
<tr>
<td>Energy content (KJ/kg)</td>
<td>15758</td>
<td>15591</td>
</tr>
</tbody>
</table>

* Methionine (10 g/kg) was added to the casein diet.
† Harper mixture containing (g/kg): NaCl 130.3, K2HPO4 389.1, MgSO4·7H2O 573.3, CaCO3 381.4, FeSO4·7H2O 27.0, MnSO4·H2O 40, KI 0.79, CuSO4·5H2O 0.5, ZnSO4·7H2O 0.12, CoCl2·6H2O 0.02.
‡ Harper mixture containing (mg/g): retinol 0.5, cholecalciferol 0.37, tocopherol 3.0, menadione 1.5, p-amino benzoic acid 145, γ-aminolevulinic acid 25, nicotinic acid 50, calcium pantothenate 5.7, riboflavin 1.5, thiamine 14, pyridoxine 9.5, pteroylmonoglutamic acid 0.5, cyanocobalamin 6.0, biotin 63.0. Lactose as carrier was added to make up to 1 g.

collected at the interface and washed three times with saline solution. The final cell pellet was suspended in RPMI 1640 medium supplemented with foetal calf serum (100 ml/l), 2 mM-L-glutamine, 10 mM-4-(2-hydroxy-ethyl)-1-piperazine ethanesulphonic acid (HEPES), penicillin 100 units/ml, and 100 μg streptomycin/ml (Gibco, Grand Island, NY, USA). The cells were counted in a haemocytometer and cell viability, as determined by the trypan blue exclusion test, was > 97%.

*Analysis of cell surface markers by flow cytometry*

Percentage of splenic cells was determined by flow cytometry (Hudson & Hay, 1989). The following rat anti-mouse monoclonal antibodies were purchased from Serotec: anti-Thy 1 (T cells), anti H2-1a subregion (B cells) and anti-Ly 40 (macrophage/monocyte). The cell pellets (1 × 10⁶) were incubated with 50 μl of the antibodies for 30 min on ice. After washing twice with phosphate-buffered saline the cell pellets were incubated with 50 μl fluorescein isothiocyanate (FITC)-labelled anti-rat rabbit IgG. After washing twice with phosphate-buffered saline monoclonal antibodies binding to the cells were analysed with a fluorescence-activated cell sorter (FACScan Flow Cytometer; Becton Dickinson, Erembodegem, Belgium). Fluorescence data were collected with logarithmic amplification. For each sample, data from 10000 volume-gated viable cells were collected. Non-mononuclear cells were gated on the basis of the forward and perpendicular light scatter signal.

*Lymphocyte proliferation to mitogen*

Cells were resuspended in RPMI supplemented at a concentration of 1 × 10⁶ cells/ml. Blastogenic response of splenocytes to the mitogens phytohaemagglutinin (PHA; Gibco, Grand Island, NY, USA), concanavalin A (Con A; Sigma Chemical Co., St Louis, MO, USA), and Escherichia coli lipopolysaccharide S (LPS; Difco, Detroit, MI, USA) was assessed by [methyl-³H]thymidine incorporation, as previously described (Martínez et al. 1992). Triplicate cultures were set up in ninety-six-well microplates (Becton Dickinson). Splenocytes (1 × 10⁶ cells in 100 μl culture medium) were incubated with or without...
mitogens at 37°, in an O₂–CO₂ atmosphere (95:5, v/v) for 72 h. Doses of mitogens were 50 µg/ml for PHA, 5 µg/ml for Con A and 100 µg/ml for LPS. At 24 h before cell collection, 3.7 × 10⁴ Bq [methyl-³H]thymidine (Amersham, Bucks) was added to each well. The cultures were then harvested onto fibreglass filter discs and counted in a liquid scintillation counter (LKB 1215). The data are expressed as a stimulation index, defined as a ratio of mean counts/min in stimulated cultures to that in unstimulated cultures of the same cells.

**Statistical analysis**

The results are presented as the arithmetic means with their standard errors for each casein and pea group. Differences among the means of groups were evaluated using the Student’s two-tailed t test and P values < 0.05 were considered to be statistically significant.

**RESULTS**

**Growth performance and feed intake**

Mice fed on pea as the source of protein showed a retardation in growth compared with those fed on casein protein (Fig. 1). Although feed intake in the two groups was similar through the experimental trial, feed efficiency ratios, expressed as g intake/g weight gain, were higher in the legume-fed animals (P < 0.01). Moreover, liver, gastrocnemius muscle, kidneys and femur weights were smaller in mice fed on the pea diet (P < 0.01), while an increase was observed in intestine weights (P < 0.05). No statistical differences were found in lymphoid organ weights between the experimental groups (Table 2).

**Packed cell volume and serum mineral measurements**

Packed cell volume showed a statistically significant decrease in those animals fed on the diet containing peas (P < 0.01). Accordingly, serum Fe levels were found to be markedly reduced (P < 0.01). Serum Zn levels were also decreased in the legume-fed animals (P < 0.05). By contrast, Cu levels were slightly increased, while Fe:Cu and Zn:Cu ratios decreased (P < 0.05 and P < 0.01 respectively), suggesting an interaction in the uptake of these minerals (Table 3).
Table 2. Initial and final body weights and organ weights, daily gain and feed-conversion ratio of mice given diets containing casein or peas (Pisum sativum var. Belinda) as the source of protein‡

(Mean values with their standard errors for eight mice per dietary group)

<table>
<thead>
<tr>
<th></th>
<th>Casein</th>
<th></th>
<th>Pea</th>
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<th>Statistical</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>se</td>
<td>Mean</td>
<td>se</td>
<td>significance</td>
</tr>
<tr>
<td>Initial body wt (g)</td>
<td>21.9</td>
<td>0.3</td>
<td>21.1</td>
<td>0.3</td>
<td>NS</td>
</tr>
<tr>
<td>Final body wt (g)</td>
<td>35.7</td>
<td>0.6</td>
<td>31.6</td>
<td>0.7</td>
<td>**</td>
</tr>
<tr>
<td>Daily gain (g/d)</td>
<td>0.67</td>
<td>0.03</td>
<td>0.50</td>
<td>0.04</td>
<td>*</td>
</tr>
<tr>
<td>Feed conversion ratio (g/g)†</td>
<td>16.1</td>
<td>2.6</td>
<td>32.5</td>
<td>3.6</td>
<td>**</td>
</tr>
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</table>

Organ wt

<table>
<thead>
<tr>
<th></th>
<th>Casein</th>
<th></th>
<th>Pea</th>
<th></th>
<th>Statistical</th>
</tr>
</thead>
<tbody>
<tr>
<td>Liver (g)</td>
<td>1.88</td>
<td>0.07</td>
<td>1.36</td>
<td>0.04</td>
<td>**</td>
</tr>
<tr>
<td>Gastrocnemius (mg)</td>
<td>336</td>
<td>5.5</td>
<td>283</td>
<td>7.2</td>
<td>**</td>
</tr>
<tr>
<td>Kidneys (mg)</td>
<td>532</td>
<td>13.0</td>
<td>449</td>
<td>13.0</td>
<td>**</td>
</tr>
<tr>
<td>Intestine (g)</td>
<td>1.50</td>
<td>0.07</td>
<td>1.86</td>
<td>0.11</td>
<td>*</td>
</tr>
<tr>
<td>Femur (mg)</td>
<td>130</td>
<td>3.4</td>
<td>103</td>
<td>7.4</td>
<td>**</td>
</tr>
<tr>
<td>Thymus (mg)</td>
<td>48</td>
<td>3.6</td>
<td>43</td>
<td>2.7</td>
<td>NS</td>
</tr>
<tr>
<td>Spleen (mg)</td>
<td>177</td>
<td>24</td>
<td>173</td>
<td>30</td>
<td>NS</td>
</tr>
</tbody>
</table>

NS, not significant.
* P < 0.05, ** P < 0.01 (Student’s t test).
† For details of diets and procedures, see Table 1 and p. 88.
‡ Food intake/g body weight gain.

Table 3. Packed cell volume and serum levels of iron, zinc and copper, of mice given diets containing casein or peas (Pisum sativum var. Belinda) as the source of protein†

(Mean values with their standard errors for eight mice per dietary group)

<table>
<thead>
<tr>
<th></th>
<th>Casein</th>
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<th>Pea</th>
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<th>Statistical</th>
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<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>se</td>
<td>Mean</td>
<td>se</td>
<td>significance</td>
</tr>
<tr>
<td>Packed cell volume (%)</td>
<td>46.0</td>
<td>2.1</td>
<td>36.6</td>
<td>2.3</td>
<td>**</td>
</tr>
<tr>
<td>Iron (mg/l)</td>
<td>2.71</td>
<td>0.24</td>
<td>1.20</td>
<td>0.27</td>
<td>**</td>
</tr>
<tr>
<td>Zinc (mg/l)</td>
<td>1.94</td>
<td>0.19</td>
<td>1.46</td>
<td>0.07</td>
<td>*</td>
</tr>
<tr>
<td>Copper (mg/l)</td>
<td>0.78</td>
<td>0.09</td>
<td>1.00</td>
<td>0.06</td>
<td>NS</td>
</tr>
<tr>
<td>Iron:copper</td>
<td>3.80</td>
<td>0.6</td>
<td>1.70</td>
<td>0.4</td>
<td>*</td>
</tr>
<tr>
<td>Zinc:copper</td>
<td>2.53</td>
<td>0.3</td>
<td>1.5</td>
<td>0.1</td>
<td>**</td>
</tr>
</tbody>
</table>

NS, not significant.
* P < 0.05, ** P < 0.01 (Student’s t test).
† For details of diets and procedures, see Table 1 and p. 88.

Serum protein and IgG determinations

Total serum proteins did not decrease significantly in mice fed on pea as the source of protein compared with casein-fed animals. However, serum albumin levels showed a marked decline (P < 0.05), while γ-globulins were increased (P < 0.01) in the legume-fed mice (Table 4).

Total serum IgG levels were increased in those animals fed on pea protein (P < 0.05), suggesting a systemic humoral immune response (Fig. 2).
Table 4. Total serum protein, albumin and γ-globulins of mice given diets containing casein or peas (Pisum sativum var. Belinda) as the source of protein†

(Mean values with their standard errors for eight mice per dietary group)

<table>
<thead>
<tr>
<th></th>
<th>Casein</th>
<th></th>
<th>Pea</th>
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<th>Statistical significance</th>
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<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SE</td>
<td>Mean</td>
<td>SE</td>
<td></td>
</tr>
<tr>
<td>Total protein</td>
<td>59.6</td>
<td>1.1</td>
<td>57.7</td>
<td>1.5</td>
<td>NS</td>
</tr>
<tr>
<td>Albumin</td>
<td>34.8</td>
<td>0.5</td>
<td>32.0</td>
<td>1.0</td>
<td>*</td>
</tr>
<tr>
<td>γ-Globulin</td>
<td>1.40</td>
<td>0.1</td>
<td>2.10</td>
<td>0.2</td>
<td>**</td>
</tr>
</tbody>
</table>

NS, not significant.
* P < 0.05, ** P < 0.01 (Student's t test).
† For details of diets and procedures, see Table 1 and p. 88.

Fig. 2. Serum immunoglobulin G (IgG) levels in mice fed on diets containing casein (□) or pea (Pisum sativum var. Belinda, □) as the source of protein. Values are means for eight mice with their standard errors indicated by vertical bars. The analysis showed that IgG levels were significantly higher in mice fed on pea as the source of protein compared with those fed on casein: * P < 0.05 (Student's t test). For details of diets and procedures, see Table 1 and p. 88.

Table 5. Splenic-cell subsets (%) of mice given diets containing casein or peas (Pisum sativum var. Belinda) as the source of protein†

(Mean values with their standard errors for eight mice per dietary group)

<table>
<thead>
<tr>
<th></th>
<th>Casein</th>
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<th>Pea</th>
<th></th>
<th>Statistical significance</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SE</td>
<td>Mean</td>
<td>SE</td>
<td></td>
</tr>
<tr>
<td>T lymphocytes</td>
<td>46.9</td>
<td>4.5</td>
<td>66.4</td>
<td>7.1</td>
<td>*</td>
</tr>
<tr>
<td>B lymphocytes</td>
<td>34.8</td>
<td>4.0</td>
<td>36.8</td>
<td>7.2</td>
<td>NS</td>
</tr>
<tr>
<td>Macrophages</td>
<td>7.7</td>
<td>1.3</td>
<td>8.5</td>
<td>1.7</td>
<td>NS</td>
</tr>
</tbody>
</table>

NS, not significant.
* P < 0.05 (Student's t test).
† For details of diets and procedures, see Table 1 and pp. 88–89.
Table 6. Mitogenic response (stimulation index, SI) of splenic lymphocytes of mice given diets containing casein or peas (Pisum sativum var. Belinda) as the source of protein†
(Mean values with their standard errors for eight mice per dietary group)

<table>
<thead>
<tr>
<th></th>
<th>Casein</th>
<th></th>
<th>Pea</th>
<th></th>
<th>Statistical significance</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SE</td>
<td>Mean</td>
<td>SE</td>
<td></td>
</tr>
<tr>
<td>PHA (SI)</td>
<td>18·7</td>
<td>6·0</td>
<td>12·2</td>
<td>3·6</td>
<td>NS</td>
</tr>
<tr>
<td>Con A (SI)</td>
<td>12·8</td>
<td>4·0</td>
<td>11·2</td>
<td>3·6</td>
<td>NS</td>
</tr>
<tr>
<td>LPS (SI)</td>
<td>9·9</td>
<td>5·9</td>
<td>6·7</td>
<td>2·1</td>
<td>NS</td>
</tr>
</tbody>
</table>

PHA, phytohaemagglutinin; Con A, concanavalin A; LPS, lipopolysaccharide; NS, not significant.
† For details of diets and procedures, see Table 1 and pp. 88–90.

Spleen-cell subpopulations and mitogenic response of lymphocytes

No statistically significant differences were observed in the percentage of B-lymphocytes and macrophages after the legume-diet intake. However, an increase was found in the splenic proportion of T-lymphocytes ($P < 0·05$), which could be explained by the immunoregulatory function of T-cells in antibody-dependent humoral response (Table 5).

The proliferative responses to T-cell mitogens (PHA and Con A) and B-cell mitogen (LPS) were slightly depressed in the animals fed on the pea-protein diet (Table 6).

DISCUSSION

There are many studies concerning the influence of dietary protein levels on the development of humoral and cellular immunity (Chandra, 1992, 1993), but only a few reports dealing with the influence of the type and quality of dietary protein on the immune system (Bounous et al. 1983; Yamauchi et al. 1993). Thus, although legumes are important sources of protein, little is known about the impact of feeding these seeds on the immune response. In this context we have studied a pea seed (Pisum sativum var. Belinda), which may be used for human nutrition and animal feeding as an important source of protein and other nutrients in different countries (Savage & Deo, 1989).

Mice fed on peas as the source of protein showed a stunting of growth and organ weights at the end of the experimental period, but no changes in feed consumption. These effects could be attributed to a low utilization of different nutrients and the involvement of antinutritional factors (Larralde & Martínez, 1989; Huisman et al. 1990). However, small-intestine weights showed a significant increase which some authors have attributed to the potential mitogenic activity of lectins and their capacity to stimulate crypt hyperplasia (Pusztai et al. 1986).

Serum Fe and packed cell volume decreased significantly in pea-protein-fed mice. These Fe alterations may involve a decrease in Fe storage, Fe transport and haemoglobin formation (Johnson, 1990). In addition, serum Zn levels, as a valid and useful indicator of the size of the exchangeable Zn pool, showed a significant decrease (King, 1990). The availability of these two minerals from legumes may be reduced by phytates, oxalates, tannins and other organic compounds occurring in legumes (Harland, 1989; Cashman & Flynn, 1991; Rubio et al. 1992). This lower bioavailability could have implications for growth, metabolism and immunity (Mengheri et al. 1984; Martínez et al. 1985; Sherman, 1992). However, serum Cu showed a slight increase in the legume group and the Fe:Cu and Zn:Cu ratios were significantly decreased. This could be explained by interactions among these minerals during absorption. Thus, chelating agents such as phytates and tannins
might decrease Fe and Zn absorption and lead to a reduction in competitive interactions, which could increase Cu absorption (Gordon, 1987).

Serum albumin is the usual index of nutritional status in relation to protein intake (Gibson, 1990). In the present experimental trial, serum levels of this protein were significantly lower in legume-fed mice compared with the casein group. A poor content of S-amino acids reported in these seeds and a reduced nutritional utilization of protein affected by e.g. haemagglutinins and protease inhibitors (Savelkoul et al. 1992) could alter plasma protein turnover, which would be linked to a possible situation of subnutrition without external signs, induced by legume intake.

On the other hand, despite the fall in serum albumin levels, serum y-globulins were markedly increased in mice fed on the diet containing the legume protein. Titres for serum IgG, which accounts for the major proportion of the y-globulins, were also increased. It is well known that in normal physiological conditions the gut wall prevents the passage of dietary proteins from the intestinal lumen into the blood circulation. However, the chronic exposure of the small intestine to dietary antigens may cause a lesion to the gut mucosa which increases its permeability, allowing the entry of antigens through the impaired host barrier (Perdue & Bienestock, 1991). Thus, the occurrence of antigenic proteins in peas could explain a loss of nutrients as a result of intestinal damage, the decrease in serum albumin levels and the poorer growth performance observed in mice fed on these seeds. Moreover, the rise of IgG titres is consistent with the passage of these antigens through the intestinal barrier.

Several authors have reported that digestive disturbances associated with feeding soya-bean protein may be attributed to immune mechanisms (Barratt et al. 1978, 1979). Sissons and co-workers found a positive link between systemic anti-soya antibody titres and the severity of gastrointestinal disorders (Kilshaw & Sissons, 1979; Sissons et al. 1989; Sissons & Tolman, 1991). Pre-ruminant calves fed on diets containing high concentrations of soya-bean protein developed long-lasting titres of circulating antibodies specific for soya-bean protein (Barratt et al. 1978, 1979). Similarly, early-weaned pigs fed on a diet containing soya-bean protein generated a systemic humoral immune response, which was specific for soya-bean protein (Li et al. 1991). Furthermore, young piglets and calves given a raw-pea-based diet developed circulatory antibodies against pea legumin and vicilin (Le Guen et al. 1991; Bush et al. 1992).

The expression of immune responses to feed antigens by animals depends on the nature of the antigen, the dose as well as the duration of exposure, age, immune status and genetic background of individuals (Crowe & Perdue, 1992; Pollock et al. 1994). It is known that the newborn’s gastrointestinal tract develops a mucosal barrier against penetration of a great variety and quantity of antigenic substances in food. This transition occurs at a time when systemic and local immunity systems are still immature (Strobel, 1990). Transient humoral and cellular immunodeficiencies have been reported in young piglets or at weaning (Hammerberg et al. 1989), while it has been proved that oral tolerance develops in older pigs (Wilson et al. 1989). In the present study the mice were recently weaned and, therefore, an immaturity in the immune system could explain the observed systemic humoral immune response.

In sensitized animals, immunological reactions to food can involve different types of mechanism. Although knowledge of the effects of immunologically mediated reactions on gut mucosal morphology and function has progressed rapidly in recent years, it is somewhat scanty and many aspects remain unknown (Strobel, 1990). In the present study the identification of cell subsets by monoclonal antibodies made it possible to identify an increase in the percentage of T-lymphocytes in mice fed on the pea diet. These results suggest a specific role for T-cells, which could be involved in a mucosal-cell-mediated...
immune response. The first sign of this reaction is an increase in intra-epithelial lymphocytes followed by an increase in crypt-cell turnover and elongation of the crypt compartment. Depending on the age of the animal, various degrees of villus atrophy have been observed (Strobel, 1990).

On the other hand, the systemic humoral immune response generated by some legume proteins could increase the T-cell subset, which has an immunoregulatory function in antibody-dependent humoral response. Although the percentage of B-lymphocytes did not increase in the pea-fed mice, this may be because a significant increase in B-cells would only occur after polyclonal activation, while the stimulation by some pea antigens would only increase a small percentage of total B-cell clones. No alterations in macrophage:monocyte proportions were found after pea-protein intake, compared with casein protein.

Furthermore, the increase in T-cells could be attributed to a specific effect of some antinutritional factors on the immune system. Thus, lectins when passing from the intestinal lumen into the blood circulation, due to their mitogenic properties, may interact with T-lymphocytes modulating immune responses (Licastro et al. 1993). The role of these molecules as immunoenhancing agents in human diseases is being investigated (Wimer, 1990).

In vitro mitogenic assays are performed to study the functional behaviour of lymphocytes and, thus, a reduction in stimulation index may be an index of lymphocyte hyporeactivity (Hudson & Hay, 1989). An impairment in lymphocyte stimulation responses to mitogens as a result of nutritional deficiencies has been reported (Sherman, 1992; Chandra, 1993). Lectins such as PHA and Con A, extracted from legumes, specifically stimulate most T-cell series and T-helper-cells respectively, while B-cell transformation is specifically achieved by the mitogenic agent LPS. In the present study the proliferative responses of splenocytes to PHA, Con A and LPS were lower in mice fed on the legume diet, although no statistical differences were found. Thus, the decrease in the mitogenic response after legume intake could be attributed to the poor utilization of proteins, Fe and Zn observed in this dietary group. Mice fed on other legumes such as faba bean (Vicia faba) have also shown a statistically significant decrease in the stimulation index (Martinez et al. 1992), which was slightly improved by Zn supplementation (Macarulla et al. 1992). Moreover, previous studies in mice fed on a diet containing peas as the source of protein have shown a significant decrease in interleukin-2 (IL-2) activity, which plays a central role in the generation and regulation of the immune response (Larralde et al. 1983). Zn-deficient diets have led to impairments in IL-2 activity (Dowd et al. 1986) and other immunological variables (Verma et al. 1988).

The present study describes a stunting in growth and a low utilization of different nutrients that could be attributed to the occurrence of some antinutritional factors and the poor content of S-amino acids in pea protein. Moreover, the systemic humoral immune response observed and the increase in T-lymphocytes could help to establish a better understanding of the possible role of antigenic proteins in gastrointestinal disorders and the poor individual performance after legume intake.

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