Food allergy affects 11.4% of patients who visit the allergologist for the first time in Spain, and there is increasing evidence of a rise in prevalence. Tree nuts are the second cause of allergy in Spain (28.4% of patients); in contrast, legume allergy is uncommon (3% of patients) [1].

The carob tree (Ceratonia siliqua L.) belongs to the Leguminosae family and is widely cultivated in the Mediterranean area. Locust bean gum (LBG) is a vegetable gum that is extracted from carob tree seed (CTS) and used mainly in food technology as a thickening and gelling agent. Commercial LBG (E410) is a powder composed of galactomannan (80%), proteins (5%-6%), fat (0.5%-0.9%), and fibers (0.8%-1%) that are derived from the endosperm of the CTS [2].

Legume allergens encompass the cupin, prolamin, PR-10, and lipid transfer protein families. Previous studies show that 11S globulins are involved in cross-reactivity between legumes (peanut and soybean), tree nuts (almond, hazelnut, pistachio, and walnut), and seeds (mustard seeds) [3]. We report the case of a 44-year-old woman with a history of rhinoconjunctivitis to pollen who had worked in an ice cream factory for 9 years. She had a 2-year history of ocular and nasal itching, rhinorrhea, nasal congestion, and sneezing every time she handled Cremodan SL29, a powder used as a stabilizer in ice cream. When working, she wore a safety mask and protective eyewear. Her duties included measuring the powder and mixing it with sugar. Her symptoms appeared during the first 5 minutes after handling Cremodan SL29 and disappeared within 24 hours of taking intranasal budesonide and oral loratadine. She reported no bronchial symptoms and was able to ingest the ice cream that she had prepared with Cremodan SL29. She tolerated handling and ingestion of guar gum.

According to the manufacturer, Cremodan SL29 contains LBG (E410), dextrose, milk proteins, gelatin, pectin (E440), and carrageenan (E407).

One year after her initial symptoms presented, she developed pharyngeal itching, eyelid angioedema, and dyspnea when ingesting chickpea. She also experienced cough after oral intake of almonds, pistachio, and sunflower seeds. Since then, she has avoided legumes (except peas and green beans, which she tolerates) and tree nuts.

Pulmonary function tests revealed normal spirometry values; the result of a nonspecific methacholine bronchial challenge was negative.

Prick-prick testing with Cremodan SL29 was positive (13×9 mm), and skin prick tests (SPTs) were positive to CTS extract (8×7 mm) and LBG (4×4 mm). SPTs to dextrose, milk protein, gelatin, pectin, and carrageenan were negative.

SPTs to legume extracts (Roxall) were positive for chickpea (4×4 mm), soybean (6×4 mm), and lentil (4×3 mm) and negative for peanut, pea, and white, red, and green bean. A prick-prick test with soybean was positive (5×5 mm).

The results of SPT with tree nut extract (Roxall) were negative, although the prick-prick test showed positive results for walnut (9×4 mm), almond (7×7 mm), cashew (4×3 mm), pistachio (7×4 mm), pine nut (5×4 mm), and sunflower seeds (6×7 mm). The result of prick-prick testing with hazelnut was negative.

Determination of serum-specific IgE (IMMULITE 2000/ Xpi, Siemens) showed the following values for legume and tree nut extracts: chickpea, 1.70 kU/L; soybean, 0.99 kU/L; lentil, 1.07 kU/L; peanut, 0.28 kU/L; pea, 1.46 kU/L; red bean, 0.58 kU/L; green bean, 0.14 kU/L; walnut, 0.22 kU/L; almond, 1.31 kU/L; cashew, 0.24 kU/L; and pistachio, 0.18 kU/L. Specific IgE to white bean, pine nut, and hazelnut were negative (<0.10 kU/L). Total IgE was 399 IU/mL.

A single-blind oral challenge performed with cooked white bean resulted in intense oral itching and cough 30 minutes after the second dose (3 pieces).

ACTS extract was prepared by homogenization (18% wt/vol) in phosphate-buffered saline (50 mM phosphate buffer, 100 mM NaCl, pH 7.5) for 4 hours at room temperature, followed by dialyzation and lyophilization. The Bradford method was used to assess the protein percentage in the extracts [4]. Sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS-PAGE) immunoblotting was performed to study the molecular mass of the IgE-binding bands in the Cremodan SL29 sample (5% protein). LBG powder (1.6% protein), and CTS extract (15% protein), showing a similar IgE-binding band profile. Bands of 69, 55, 50, 35, 34, 28, 22, and 19.5 kDa were detected (Figure, I).

Cross-reactivity studies were performed with chickpea and almond, as these showed the highest levels of serum-specific IgE, and the patient reported symptoms after ingestion. SDS-PAGE immunoblotting inhibition studies revealed complete inhibition of IgE binding in the LBG sample and CTS extract with Cremodan SL29 and extracts from chickpea (50% protein) and almond (70% protein). Cremodan SL29 completely inhibited IgE binding with chickpea extract and partial inhibition with almond extract (Figure, II). We hypothesized that the patient was sensitized to the remaining CTS proteins present in the LBG sample. The paucity of published data in this area prevents us from speculating about the identity of the IgE-reactive CTS proteins detected. However, the molecular masses of the chickpea and almond proteins (Figure, II) that cross-react with the patient's serum.
with them lead us to suppose that they could belong to 11S globulin type (cupin superfamily) [5,6]. The molecular masses of some of the CTS IgE-reactive proteins detected (67 kDa, 55 kDa, 28 kDa) are similar to some of those observed by Alarcon et al [7].

Of all the cases of LBG sensitization reported to date, few address occupational exposure in the food industry [8-9]. None of the work-related cases involved reactions after the ingestion of legumes or tree nuts.

As LBG is obtained from CTS, a member of the Leguminosae family, some authors studied the cross-reactivity between LBG powder and legume proteins. Fiocchi et al [10] reported sensitization to carob by means of SPT and in vitro testing in peanut-allergic individuals, all of whom tolerated cooked carob. Alarcón et al [7] observed sensitization to legume and tree nut (positive SPT and specific IgE) in a patient with a clinically relevant allergy to LBG (urticaria after ingestion of caramel containing LBG), although this patient tolerated ingestion of legumes and tree nuts.

We present a case of occupational IgE-mediated sensitization to inhaled proteins in an LBG sample with clinical consequences for ingestion of legume. To our knowledge, this is the first report of a clinically relevant allergy to carob gum, legumes, and tree nuts due to cross-reactivity between their proteins.

**Funding**

The authors declare that no funding was received for the present study.

**Conflicts of Interest**

The authors declare that they have no conflicts of interest.

---

**References**


Hyper IgM Syndrome Type 2 Presenting as Intestinal Lymphoid Polyposis Without Recurrent Infection

de la Varga-Martínez R1, Vilches-Moreno M1, Viejo-Almanzor A2, Pérez-Requena J3, Rodríguez C1, Mora-López F1

1Servicio de Inmunología, UGC de Hematología, Inmunología y Genética, Hospital Universitario Puerta del Mar, Cádiz, Spain
2UGC de Aparato Digestivo, Hospital Universitario Puerta del Mar, Cádiz, Spain
3Servicio de Anatomía Patológica, Hospital Universitario Puerta del Mar, Cádiz, Spain

Key words: Hyper IgM syndrome. Immunodeficiency. Benign lymphoid polyposis. Class switch recombination. Activation-induced cytidine deaminase

Palabras clave: Síndrome de hiper-IgM. Inmunodeficiencia. Poliposis linfoide benigna. Recombinación del cambio de clase. Deaminasa de citidina inducida por activación.

Hyper IgM syndrome (HIGMS) comprises a group of primary immunodeficiencies [1] characterized by impaired immunoglobulin class-switch recombination (CSR). Patients with HIGMS have normal or elevated IgM levels, with the other immunoglobulin subtypes absent or strongly decreased. In some HIGMS types, impaired somatic hypermutation (SHM) of immunoglobulin genes and abnormal T-cell function are also present [2].

HIGMS type 1 (HIGMS1) is caused by mutations in the gene encoding CD40 ligand (CD40L), and HIGMS3 is due to mutations in the CD40 gene. HIGMS1 and HIGMS3 are clinically indistinguishable. HIGMS2 is produced by mutations in the activation-induced cytidine deaminase gene (AICDA) [3], and HIGMS5 is caused by mutations in the uracil-DNA glycosylase gene (UNG). These types are clinically similar. Most patients with HIGMS develop clinical symptoms in infancy and early childhood. All the forms of HIGMS are characterized by increased susceptibility to recurrent bacterial infections of the respiratory and digestive tracts [4,5]. Viral infections are more frequent and severe. Patients with defects in the CD40/CD40L pair also have defective cellular immunity and are susceptible to opportunistic infections such as Pneumocystis jiroveci pneumonia. Since cellular immunity is not impaired in HIGMS owing to mutations in AICDA or UNG, patients with HIGMS2 and HIGMS5 do not experience opportunistic infections. In these types, lymphadenopathy due to the presence of expanded germinal centers is frequent, whereas the lymph nodes of patients with mutations in CD40L and CD40 lack germinal centers.

Activation-induced cytidine deaminase (AID) is expressed in B cells in germinal centers, where it plays a key role in both CSR and SHM. Patients with HIGMS2 first experience recurrent infections during early childhood. However, patients with HIGMS2 have no increased susceptibility to opportunistic infections.

Manuscript received October 14, 2019; accepted for publication February 24, 2020.

María Vázquez de la Torre
Allergy Service
Infanta Leonor University Hospital
Av. Gran Vía del Este, 80
28031 Madrid, Spain
E-mail: mvazquezt@salud.madrid.org