Evaluation of the usefulness of a standard multivitamin supplement designed for patients with cystic fibrosis*,**

Valoración de la utilidad de un suplemento multivitamínico estándar diseñado para pacientes con fibrosis quística

Dear Editor:

Fat and fat-soluble vitamin (FSV) malabsorption is frequent in individuals with cystic fibrosis (CF) and exocrine pancreatic insufficiency (EPI). Consensus guidelines recommend FSV supplementation for patients that receive this diagnosis. Traditional supplementation may call for the ingestion of four or more units of medication a day, which poses a barrier to adherence. A new formulation (AquADEKs®) has been available in recent years that contains FSVs, water-soluble vitamins and antioxidants in doses specifically conceived for patients with CF (Table 1). All its preparations (liquid suspension, chewable tablets and soft gel capsules) have a micellar structure with a hydrophilic outer layer and a lipophilic inner layer that houses the FSVs, facilitating their absorption. Thus, our expectation was that post-intervention blood levels would remain within the normal range.

The aim of this study was to assess the efficacy of AquADEKs® in maintaining the serum levels of FSVs of children and adults with CF and EPI within the normal range, analysing the serum levels of FSVs at the time the administration of the new formulation was initiated, when the patients were receiving FSV supplementation with traditional formulations (pre-intervention), and 3–6 months after starting supplementation with AquADEKs® (post-intervention).

We conducted a prospective quasi-experimental study. Between September 2012 and July 2013, 31 patients with CF and EPI that were receiving FSV supplementation were included in the study; 21 were male, the mean age was 10.7 ± 9.8 years, 12 were homozygous for Phe.508del with a mean force expiratory volume in the first second (FEV1, % over the vital capacity) of 82.8 ± 22.2. Comparing the pre- and post-intervention periods, the mean ingestion of medication units was 3.2 ± 1.4 versus 2 ± 1 (P = 0.045); the serum levels of vitamin A (retinol), 34.04 ± 11.7 μg/dL versus 34.2 ± 10.1 μg/dL, remained stable (P = 0.90); the vitamin E (alpha-tocopherol) levels, 838.5 ± 272.6 ng/dL versus 1005.1 ± 289.9 ng/dL (P = 0.02), were higher post intervention, and the vitamin D (25-OH-D3) levels, 29.6 ± 11.4 ng/mL versus 22.6 ± 8.5 ng/mL (P = 0.002), were lower post intervention. Prothrombin time was normal in every case, 11.69 ± 0.53s pre intervention compared to 11.66 ± 0.63s post intervention, a difference that was not statistically significant.

We found two scientific studies published in the literature up to date in which this product was used in patients with CF and EPI.

When it came to vitamin A, our findings were consistent with those of the study in 11 patients by Sagel et al., in which the serum concentrations of retinol remained stable and within the normal range after supplementation with AquADEKs® (pre, 39 [32–48] μg/dL vs post, 39 [35–53] μg/dL). Another study conducted by Moen et al. on 30 children found a significant increase in serum retinol after using AquADEKs® (pre, 31.46 ± 8.5 μg/dL vs post, 37.18 ± 8.5 μg/dL; P = 0.04), with concentrations within the normal range.

As for the increase in serum vitamin E concentrations, our results were similar to those of the studies by Sagel et al. and Moen et al., which found post-intervention increases in vitamin E serum levels that did not reach statistical significance.

As for vitamin D, most patients in the study by Sagel et al. had insufficient levels of this vitamin (<30 ng/dL) before and after the treatment. In the study by Moen et al., the percentage of insufficiency was slightly higher than the one in our study (50%), but deficiency was lower in theirs (25%). Since optimum levels of vitamin D were not achieved...


** Previous presentation: this study was presented as an oral communication under the title “Simplificación del tratamiento multivitamínico en la fibrosis quística: ¿una solución para nuestros pacientes?” at the XII Congreso Nacional de Fibrosis Quística; November 14–16, 2013; Palma de Mallorca, Spain.
(Fig. 1) patients needed supplementation with an additional medication unit, but the number of medication units per day was still lower than under traditional supplementation (2 vs 5). To meet the current recommendations for vitamin D supplementation in patients with CF, a new formulation of AquADEKs® with a higher dose of vitamin D has been introduced recently that aims at achieving adequate concentrations of this vitamin.

One limitation of our study is that the sample size was determined by convenience sampling, with the study including the first 31 patients that met the inclusion criteria. The power was high enough for the observed decrease in vitamin D levels and the increase in vitamin E levels, and was low for the vitamin A results, although the values that we obtained lead us to believe that the clinical differences are not important or relevant.

Adherence to treatment was good, with good tolerance, and we did not observe any adverse reactions to the new formulation.

Supplementation with AquADEKs® makes multivitamin treatment easier, helps improve quality of life and facilitates adherence to treatment in patients with CF. While using this product, some patients required additional vitamin D supplements and others had high levels of vitamin E.

**Funding**

This study received 12,000€ of funding from Praxis Pharmaceutical, S.A.

**Acknowledgments**

We want to thank nurses Diana San Miguel and Maria Concepción Rodríguez and nurse assistant Carmen Espallargas for their collaboration in this study; as well as Javier Zamora and Alfonso Muriel, from the Biostatistics Department, Itziar de Pablo and María Ángeles Galvez, from the Ethical Committee for Clinical Research (Comité Ético de Investigación Clínica [CEIC]), and Carmen Santistebe, Óscar Pastor, Diana Cuesta and Carmen Bayón from the Biochemistry Laboratory of the Hospital Universitario Ramón y Cajal.

**References**


M. Garriga a, A. Horrisberger b, A. Lamas c,d, M. Ruiz de Valbuena e, L. Suárez e,f

a Unidad de Fibrosis Quística, Servicio de Endocrinología y Nutrición, Hospital Universitario Ramón y Cajal, Madrid, Spain
b Unidad de Fibrosis Quística, Hospital Universitario Ramón y Cajal, Madrid, Spain
c Sección de Neumología Pediátrica, Unidad de Fibrosis Quística, Servicio de Pediatría, Hospital Universitario Ramón y Cajal, Madrid, Spain
d Instituto Ramón y Cajal de Investigación Sanitaria (IRYCS), Hospital Universitario Ramón y Cajal, Madrid, Spain
e Servicio de Gastroenterología Pediátrica, Unidad de Fibrosis Quística, Hospital Universitario Ramón y Cajal, Madrid, Spain
f Facultad de Medicina, Universidad de Alcalá de Henares, Madrid, Spain
g Corresponding author.
E-mail address: maria.garriga@salud.madrid.org
(M. Garriga).

Gastrointestinal involvement in Langerhans cell histiocytosis

Histiocitosis de células de Langerhans con afectación gastrointestinal

Dear Editor:

Langerhans cell histiocytosis (LCH) is characterised by the accumulation of dendritic cells in granulomatous lesions that mostly involve bone and skin.1 Gastrointestinal involvement is rare in LCH (1–5%), although its incidence is probably underestimated due to the nonspecific nature of its presenting symptoms and the association of the latter to the chronic course of disease or the treatments for it.2 We present two cases of LCH with gastrointestinal involvement.

Figure 1 Colorectal mucosa showing a histiocytic infiltrate in the lamina propria (a). Positive CD1a staining in the previously described cellular tissue (b).

Case 1

Girl, 4 months of age presenting with a papular rash in the torso and scalp lasting 4 months associated with anorexia, delayed growth (3rd percentile) and haematocrit. The skin biopsy was consistent with LCH and displayed intense reactivity to CD1a and S100. After ruling out a cow’s milk allergy, we performed a gastrointestinal endoscopy that showed nonspecific signs of duodenitis and colitis. The histological examination revealed a histiocytic infiltrate accompanied by eosinophils in the colorectal mucosa with immunohistochemical findings consistent with LCH (Fig. 1). The remaining staging studies ruled out disease at other locations. As a case of multisystem LCH without involvement of risk organs it was treated according to the LCH III group B protocol, consisting of a 6-week induction

* Please cite this article as: Andión Catalán M, Ruano Domínguez D, Azorín Cuadrillero D, de Rojas de Pablos T, Madero López L. Histiocitosis de células de Langerhans con afectación gastrointestinal. An Pediatr (Barc). 2015;83:279–280.