

Botanical Extract Coacillium for Management of Paediatric Atopic Dermatitis: A Case Report

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Accepted Oct 18, 2023; Published Nov 21, 2023

Acta Derm Venereol 2023; 103: adv13376. DOI: 10.2340/actadv.v103.13376

Atopic dermatitis (AD) is associated with the activation of type 2 inflammation (1). Environmental and psychological factors complete the aetiological characteristics and aggravate the clinical picture (2). In addition to recently highlighted therapeutic approaches (anti-IL13, anti-PD4, anti-JAK) (3), anti-inflammatory herbal mixtures have been studied (4). Transient receptor potential cation channel subfamily M member 8 (TRPM8) and cannabinoid agonists represent novel pathways, but also Coacillium (Coa) acting as an immunomodulator in the type 2 inflammatory pathway but also through pro-apoptotic B Cell Lymphoma 2-6 gene expression (BCL2-6) (5). Coa is a topical hydroalcoholic solution composed of 4 plants extracts: Onion (*Allium cepa*), Lemon (*Citrus limon*), Guarana (*Paullinia cupana*) and Theobroma cacao. Coa was initially commercialized as an anti-hair loss lotion and is now being studied as a drug for treatment of alopecia areata in children (6). Coacillium is the only name used for the product. It has not yet been approved for any indication in Switzerland but has been filed for approval with the European Medical Agency (EMA) for the treatment of Alopecia Areata as a « Herbal Medicine ». We report here the experimental use of Coa to manage symptoms in a paediatric patient with AD.

MATERIALS AND METHODS

The product used in this case report is an alcohol-free Coa formulation (with reliable test data related to safety)(7). The constituent plant extracts listed in the introduction are all classed as Generally Regarded as Safe (GRAS) or equivalent by the American Food and Drug Administration (FDA), who placed the development and regulation of Coa under the category of “Botanical Drugs” (also called “herbal medicine” by EMA). Unlike chemical entities, botanical drugs do not have International Nonproprietary Names (INN). The alcohol-free product batch 18 3 21 5, manufactured

by Delpharm Tours, Chambray-Les-Tours, France, contains the following excipients regularly used in cutaneous solutions: water, glycerin, sodium chloride, potassium sorbate, sodium benzoate, maltodextrin, citric acid. Coa has no allergenic potential. In this case, Coa was used as an adjunctive therapy to decrease the sensation of pruritus and to control skin lesions in a patient with AD. An 8-year-old girl presented with persistent cheilitis with intermittent acute flares. Dry skin, erythema, desquamation, and the Dennie-Morgan sign associated with nocturnal pruritus complete the clinical picture of diagnosed AD in early childhood (Fig. 1A). Previous conventional approaches such as topical corticosteroids led to a partially favourable situation.

RESULTS

The patient experienced immediate improvement with a significant decrease in itch, dryness, rhagades, and inflammation with 1 application locally (fingertip-unit amount) daily for 5 days, which allowed us to taper the treatment (1-month duration interval therapy twice a week). Usual prior to Coa moisturizing treatment (products containing a combination of humectants, emollients and occlusives for general skincare) and, especially for the lips, a protective barrier function with ingredients such as shea butter and cocoa butter with very low allergenic potential allowed long-term control of her disease (Fig. 1B). To date, in the 12 months after treatment, she has not experienced major worsening of her symptoms.

DISCUSSION

The patient in this single case report experienced subjective improvement of her AD symptoms: an immediate anti-inflammatory response with long-term maintenance of symptom control. Long-term use of corticosteroids, even at regular intervals, can often pose problems with



Fig. 1. Atopic cheilitis (A) before treatment and (B) after 1 month of treatment.

compliance and side-effects (vasoconstriction, fragility of the skin barrier). Calcineurin inhibitors are often poorly tolerated due to the associated mast cell stimulation, especially on the lips. Many patients with AD or their parents seek alternatives to standard treatment. This single preliminary case study indicates the potential of an herbal medicine in the management of AD. No contact sensitization due to Coa has yet been shown. There are a few reports of allergic reactions following onion (*Allium cepa*) consumption (8). The CoA path mechanism, mainly around the B Cell Lymphoma gene 2 (BCL-2) expression affinity, suggests the involvement of apoptosis in AD (1,5). This intrinsic process (mitochondrial driven pathway), where apoptosis is driven by cytotoxic drugs or DNA-damage, is regulated by BCL-2 family proteins and the apoptosis-inducing factor (9). This single case report found Coa to be a possible therapeutic modality. However, further research is needed to determine its efficacy.

ACKNOWLEDGEMENTS

Funding sources: Legacy Healthcare Ltd, Route de la Corniche 3B, 1066 Epalinges, Vaud, Switzerland.

Conflicts of interest: The author works as a consultant for Legacy Healthcare Ltd. He receives honoraria related to the academic activity of the company.

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