Acroqueratodermia Aquagénica e Mutações no Gene da Fibrose Quística: Uma Pista Diagnóstica

Aquagenic Wrinkling of the Palms and Mutations in the Cystic Fibrosis Gene: A Diagnostic Clue

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Resumo:
Descrita pela primeira vez em 1974, a acroqueratodermia aquagénica (AA) é uma entidade caracterizada por um enrugamento pruriginoso das palmas das mãos com edema associado após breve contacto com a água. Foi inicialmente reportado em crianças com diagnóstico de fibrose quística e, nos últimos anos, uma associação entre estas duas condições tem sido estabelecida. Descreve-se o caso de uma mulher caucasiana de 21 anos com história de rinossinusite e diagnóstico clínico de AA. O estudo realizado revelou uma mutação F508del num dos alelos do gene da fibrose quística (CFTR). A AA pode constituir uma manifestação de mutações ao nível do gene CFTR e a sua identificação deve conduzir a um aconselhamento genético.

Palavras-chave: Queratodermia Palmar e Plantar; Fibrose Quística; Mutação; Regulador de Condutância Transmembrana em Fibrose Quística.

Abstract:
Described for the first time in 1974, aquagenic wrinkling of the palms (AWP) is a dermatological condition characterized by skin wrinkling of the palms with oedema after brief contact with water. It was first noticed in children diagnosed with cystic fibrosis (CF), the most frequent severe autosomal inherited abnormality, caused by mutations in the cystic fibrosis transmembrane conductance regulator (CFTR) gene. Since then, the association has been confirmed by several smaller studies and case reports and also in individuals where a single mutation of the CFTR gene has been reported, as well as in patients with non-classic CF.

Case Report
A 21-year-old Caucasian woman presented with a 12 months history of pruritic wrinkling of the palmar skin after brief contact with water. Symptoms were temporary and spontaneously regressed within a few minutes (15 minutes) after hands drying. On clinical examination, she developed substantial oedema and a whitish papillomatous appearance of the palms after immersion of the hands in water for about 5 to 7 minutes (Fig. 1). She had no history of hyperhidrosis or other skin condition, as well as no other concomitant diseases, except for chronic rhinosinusitis. She was not on any medication and had no family history of CF or upper airway pathology. A clinical diagnosis of AWP was made and a full medical routine was ordered, namely chest X-ray, chest computed tomography (CT) scan, sinus X-ray, sweat test and spirometry with no abnormality found. A complete CFTR gene sequencing was performed, leading to the identification of a heterozygous mutation for p.Phe508del. No specific treatment was applied; the patient and her family were referred to a genetic counselling consultation where global genetic counselling was made without family testing.

Discussion
CF is an autosomal recessive disease in the CFTR gene leading to abnormal transport of chloride over epithelial membranes that results in reduced electrolyte reabsorption and raised concentrations of chloride in sweat and organ damage in
the respiratory, pancreatic, biliary, reproductive and gastrointestinal systems.\textsuperscript{2,4} Currently, nearly 2,000 mutations have been identified and in recent years, with help from the field of genetics, the diagnosis of CF has been extended to include milder phenotypes regarded as non-classic CF or CFTR related disorders.\textsuperscript{2,5} Some of these patients have mild or no lung disease with unusual clinical features such as chronic sinusitis or male infertility, and AWP could be another feature that may fall in this category.\textsuperscript{4}

The link between CF and AWP was first established in 1974 but the exact physiological mechanism remains unknown. In the past decades, studies documented a prevalence of AWP between 78\% and 84\% in patients with CF and of 25\% in CF carriers.\textsuperscript{3,5,6} Nonetheless, in recent years, reports of AWP related to a carrier state of CF like the one in this article have exponentially increase.\textsuperscript{5}

In conclusion, AWP is yet an unknown condition with a strong established connection with classic CF, non-classic CF and CF carriers. The identification of this phenomenon with a directed clinical history, physical examination and appropriate complementary study is mandatory and can be of great impact in individuals of childbearing age, knowing that the search for a mutation in the \textit{CFTR} gene must be extended to the full gene sequencing and not only to the most common mutations associated with this disease. \textsuperscript{[1]}

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