

Case Report

FAMILIAL CHROMHIDROSIS

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ABSTRACT

Chromhidrosis is a rare and benign condition that known producing coloured sweats from eccrine and apocrine sweat glands. However several cases related to drugs and exogen-endogen materials are submitted in the literature; it is usually idiopathic. In this report we present, red-orange sweating case which affected all members of five person family has been determined such a case very rarely reported in literature. Causes, mechanisms and treatment of chromhidrosis affecting life quality of people is clarified in this case.

Key words: Chromhidrosis, benign

ÖZET

Kromhidroz ektrin ve apokrin ter bezlerinden renkli ter üretimiyle bilinen nadir ve benign bir hastalıktır. Bununla birlikte ilaçlar, endojen-ekzojen maddelere ilişkili olgular literatürde bildirilmiştir; genellikle idiyopatiktir. Bu yazıda beş kişilik ailenin tüm bireylerini etkileyen ve kırmızı-portakal rengi terleme gösteren literatürdeki nadir bir olguyu sunuyoruz. İnsanların yaşam kalitesini etkileyen kromhidrozun nedenleri, mekanizmaları ve tedavileri bu olguda tartışılmaktadır.

Anahtar kelimeler: Kromhidroz, benign

INTRODUCTION

Chromhidrosis was first described by Yonge in 1709.¹ It is a rare disease which affects both apocrine and eccrine sweat glands and resulting with different colours of sweat such as yellow, green, blue, blue black and brown.² Familial chromhidrosis

has been reported in the literature in two cases of brothers aged 12 and 9 years.³ We report of five person family with red-orange sweating diagnosed as chromhidrosis.

CASE REPORT

A five person family (35 year-old father, 33 year-old mother and three children aged

11, 9 and 6 years) admitted to our clinics with complaint of red-orange coloured

staining dress. They remarked that red-orange coloured staining of the hands and feet appeared in the morning. Their medical history was unremarkable and they were not on any topical or oral medications. They had no recent history of intake of any colored foods. They stated that they have had such complaints once before.

A complete blood count, urinalysis and homogentisic levels were within normal

limits. Liver function tests were normal. Skin bacterial and fungal cultures were negative.

On physical examination; each of five person had red-orange coloring was observed different part of their bodies and dresses (Figure 1), (Figure 2).

Any treatment was recommended. Patients were followed up. No relaps was observed in 8 month follow up.



Figure 1. Red-orange pigmentation in the plantar region of the child and mother



Figure 2. Red-Orange coloring on the child's sock and mother's athlete

DISCUSSION

Chromhidrosis is defined by Yonge of Plymouth in 1709.¹ It is referred that there are three types of chromhidrosis: i) apocrine chromhidrosis, ii) pseudo-ecrine chromhidrosis, iii) eccrine chromhidrosis. There have been one familial case report in the literature.³

Chromhidrosis, which is characterized producing coloured sweat by eccrine and apocrine sweat glands is very rare condition. Apocrine chromhidrosis is a disease which affects apocrine glands localize only on the face, extremities, and breast areolae. Contrarily, eccrine chromhidrosis is usually generalized on the skin.⁴ Eccrine chromhidrosis occurs releasing chrome or drug which tends to be solved by water from eccrine glands. Furthermore; it is also result in chromogenic bacteria an fungal elements or extrinsic chemicals and nourishments on the skin.^{3,5} Although no relation between chromhidrosis and systemic disease have been documented, it is reported that there is an association between chromhidrosis and hyperbilirubinemia in some case.^{1,5,6} In pseudoecrine chromhidrosis, sweat is actually colorless, discoloration is developing with substances that contact and cause pigmentation from the outside

to the surface of the skin. A large number of chromogenic, porphyrin-producing bacteria and fungi like; *Malassezia furfur*, *Bacillus* spp., *Corynebacterium*, *Piedra*, *Pseudomonas* were reportedly caused to pseudoecrine chromhidrosis.⁷ Our case's skin bacterial and fungal cultures were negative. Therefore pseudoecrin chromohydrosis was not considered. Eccrin chromohydrosis was considered because there was color change in hand and food in our patient.

The differential diagnosis consists of hyperbilirubinemia and alcaptonuria. Routine hemogram, liver and renal profiles, urinalysis and electrolytes should be estimated in case of doubt. Moreover, histopathologic examination and spectrophotometric analysis can be performed.³

Treatment alternatives depends on type and reason of chromhidrosis. Antiperspiration cream which contains aluminum chlorid or capsaicin and botulinium toxin type A (BTX-A) injection can be used for therapy. Besides, it must be withdrewed drugs and foods related to disease.³

CONCLUSION

It should be investigated on underlying causes and planned therapy according to the case.

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