

Recurrent Acute gastroenteritis with septic shock and acute kidney injury detected as Addison's disease after 5 years

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Abstract- Introduction: Addison's disease is a rare endocrinal disorder, with several oral and systemic manifestations. A variety of pathological processes may cause Addison's disease. Classically, hyperpigmentation is associated with the disease, and intraoral pigmentation is perceived as the initial sign and develops earlier than the dermatological pigmentation. The symptoms of the disease usually progress slowly and an event of illness or accident can make the condition worse and may lead to a life-threatening crisis. In this case, several oral as well as systemic manifestation of the Addison's disease was encountered.

A 45 year old male became acutely unwell and had a background of recurrent admissions in past for AGE like illness requiring pressor support and in between had generalized weakness, nausea, weight loss and fatigue. He had an acute diarrhoeal illness which didn't respond completely to intravenous fluids and antibiotics. In view of patient having such episodes in past 5 years and features of hyperpigmentation and generalized weakness, nausea and weight loss. With this background, patient was evaluated for adrenal insufficiency which was confirmed by basal cortisol levels and Short synacthen test(SST).

Conclusion: Addison's disease can remain unrecognized until precipitated by acute stress.

Index Terms- Addison's disease, hyperpigmentation, prednisolone.

I. INTRODUCTION

Addison's disease is rare endocrinal disorder that affects 1 in 100,000 people. It is seen in all age groups and affects male and female equally. This disease is named after Thomas Addison, who first described patients affected by this disorder in 1855, in the book titled "On the constitutional and local effects of the disease of supra renal capsule.[1,2] Addison's disease can present as a life-threatening crisis, because it is frequently unrecognized in its early stages. The basis of Addison's disease has dramatically changed from an infectious cause to autoimmune pathology since its initial description. However, tuberculosis is still the predominant cause of Addison's disease in developing countries.[3]

The symptom of Addison's disease begins gradually, chronic worsening fatigue, loss of appetite, generalized weakness, hypotension, and weight loss. The clinical features of hypoadrenocorticism do not actually begin to appear until at least 90% of the glandular tissue has been destroyed. Generalized hyperpigmentation of skin is seen, which is classically described as "bronzing" the hyperpigmentation is generally more prominent on sun-exposed skin and over pressure points, such as the elbows and knees which are caused by increased levels of beta-lipotropin or Adrenocorticotrophic hormone, each of which can stimulate melanocyte production. Hyperpigmentation of the mucous membrane and skin usually proceeds over other symptoms by month to year. Vitiligo may also be seen in association with hyperpigmentation in idiopathic Addison's disease due to autoimmune destruction of melanocytes.[4-6] Hyponatremia and hyperkalemia are commonly associated with Addison's disease, while hypoglycaemia uncommon. The patient usually complains of gastrointestinal upset with anorexia, nausea, vomiting, diarrhea, and a peculiar craving for salt. Adrenal calcification and enlargement are commonly seen in Addison's disease associated with tuberculosis.[7-9]

The symptoms of Addison' disease progress slowly and are usually ignored, an event of illness or accident can make the condition worsen and lead to Addisonian crisis. Sudden penetrating pains in the lower back region, abdomen or legs are symptoms of Addisonian crisis with severe vomiting and diarrhea, which is followed by dehydration, low blood pressure and loss of consciousness.[6,9]. The current case not only presents with the typical features of the disorder but also with the extra and particularly unique extensive intraoral pigmentation which is often considered as the initial presentation of the Addison's disease.

Case : A 45 year old man presented with complaints of multiple episodes of vomiting and loose stools. He was evidently fatigued and weak. He had abdominal pain and in past had occasional nausea ,vomiting and abdominal pain episodes for which he had multiple admissions in past. On clinical examination he had thin and brittle nails, scanty body hair, hyperpigmentation of skin in the neck , palmar crease (fig 1) and feet (fig 2) was present. There was a pulse of 106 bpm, and blood pressure 90/65 mmHg supine (after fluids and noradrenaline). Extraoral examination showed brown pigmentation affecting the perioral region. Intraoral examination demonstrated pigmentation with bilateral involvement of buccal mucosa, gingival, mucosal surface of lower lip, alveolar mucosa, and hard palate.(fig 3) Tongue has pigmentation on the anterior and lateral surfaces . (fig4)

Laboratory investigations revealed hemoglobin level 10.8 g/dl, with normal red blood cell morphology, erythrocyte sedimentation rate (ESR) 59 mm/h, fasting blood sugar 70 mg/dl. The metabolic profile - serum urea 96mg/dl, creatinine 2mg/dl and electrolytes sodium 126meq/l, potassium 5.7meq/l . Early morning 8a.m cortisol level was well below normal level 1.6 µg/dl. A SST was done which revealed flat cortisol level at 30 minutes. Anti-HIV, anti hepatitis C virus hepatitis B surface antigen (HCV ,HBs Ag) factors were negative. Mantoux tuberculin skin test was negative and chest radiograph also ruled out tuberculosis. Based on patient history, clinical findings and laboratory investigations, we reached to the diagnosis of Addison's disease. Initially, scaling was done and instruction was given to maintain oral hygiene. Cortisol replacement in from of prednisolone 20 mg in morning and 10 mg in evening was started, as her condition improved after being on hydrocortisone 10 mg/hr infusion. He was managed with intravenous fluids, parental supplement glucose, hydrocortisone. He had full recovery after 20 days of hospitalization and is maintained on prednisolone.

Discussion:

A variety of pathological processes may cause Addison's disease, which was first described by Thomas Addison.[1–3] The commonest causes of Addison's disease are autoimmune and tuberculosis. Several autoimmune processes can lead to adrenal insufficiency affecting exclusively the adrenal glands or be part of a more complex inherited autoimmune polyglandular syndrome.[2,3,6,10] Tuberculosis is the most common cause of Addison's disease in developing countries.

Fungal infection, hemochromatosis, metastatic neoplasm, and X-linked adrenoleukodystrophy are other causes of Addison's disease.[2,11] Several investigators have found dysphagia, fatigue, weight loss and hypotension, abdominal pain, amenorrhea, nausea, and vomiting, thin and brittle nail, scanty body hair in Addison's disease, which is also present in this study.[2–5,7–9,11,12] Psychiatric symptoms such as mood disturbances, decreased motivation, and behavior changes are frequently associated with Addison's disease. According to Anglin et al., the etiology of neuropsychiatric symptoms associated with Addison's disease is unknown, but may be related with the disturbances in the electrophysiological, electrolyte, and metabolic activity. In this case, positive history of mood disturbances and behavior changes is also present.[13] Sleep disturbances on periodic exacerbation is present in this case, which is according to the study by Løvås et al.[14] One of the hallmark signs of Addison's disease is cutaneous and mucosal hyperpigmentations related to ACTH melanogenesis action.[3–5] Soule reported that the presenting features among 50 patients seen over a 17-year period, as including hyperpigmentation (86%), weight loss (67%), abdominal pain (20%), and diarrhea (16%).[7] Pigmentation can be homogeneous or blotchy. The pigmentation may involve skin, oral cavity, conjunctiva, and genitalia.[3,5] Brown patches of gingival, vermilion border of the lips, buccal mucosa, palate, and tongue may represent the first signs of Addison's disease.[5] In this study, pigmentation was present in the feet and with bilateral involvement of buccal mucosa, gingival, mucosal surface of lower lip, and alveolar mucosa. Tongue has pigmentation on the posterior surface . This case is managed by hydrocortisone in hospital and after fully recovered the patient is now maintained with prednisolone.

Conclusion:

Addison disease is rare endocrinal disorder, in the developed nations it usually related to auto-immune disorder but in the developing nations it is widely associated with tuberculosis. Addison disease is usually associated with dysphagia, fatigue, weight loss, hypotension, abdominal pain, amenorrhea, nausea, vomiting, thin and brittle nail, scanty and body hair. Addison disease is classically seen with hyperpigmentation due to ACTH melanogenesis. Intraorally pigmentation over the gingival, vermilion border of lip, buccal mucosa, palate tongue, is evident and perceived as first sign. As the oral manifestation of the Addison's disease, particularly oral pigmentation may develop earlier than their dermatological counterpart, dental surgeon may be the first medical professional to encounter disease and early diagnosis of the disease essential for proper medical management.

FIGURES



FIGURE 1. HYPERPIGMENTATION OF SKIN
AND PALMAR CREASE.



FIGURE 2. HYPERPIGMENTATION OF FEET.



FIGURE 3. PIGMENTATION WITH BILATERAL
INVOLVEMENT OF BUCCAL MUCOSA, GINGIVAL,
MUCOSAL SURFACE OF LOWER LIP,
ALVEOLAR MUCOSA, AND HARD PALATE

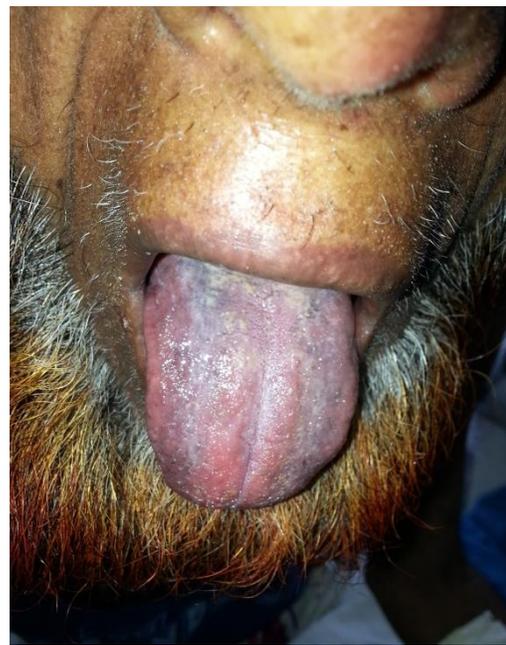


FIGURE 4. TONGUE HAS PIGMENTATION ON THE ANTERIOR
AND LATERAL SURFACES

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