Doss Porphyria (δ-Aminolevulinic Acid Dehydratase Porphyria) Presenting with Acute Onset Flaccid Paralysis

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Abstract
δ-Aminolevulinic acid dehydratase porphyria is an autosomal recessive disorder of heme synthesis resulting from deficiency of δ-aminolevulinic acid dehydratase (ALAD). Patients present with fatal neurovisceral manifestations and motor neuropathy. Here we report a patient with rapidly progressive flaccid tetraplegia with respiratory and bulbar paralysis. The importance of early diagnosis, prompt treatment and screening of relatives is stressed.

Keywords: Doss porphyrias, Flaccid tetraplegia, Porphobilinogen, Aminolevulinic acid

Introduction
Porphyrias are rare metabolic diseases resulting from altered activities of specific enzymes of the heme biosynthetic pathway. δ-Aminolevulinic acid dehydratase porphyria (ADP), sometimes termed as Doss porphyria results from a deficiency of δ-aminolevulinic acid dehydratase (ALAD) and is inherited as an autosomal recessive trait. Only six cases have been confirmed by mutation analysis. ALAD is the second enzyme in the heme biosynthetic pathway and catalyzes the condensation of 2 molecules of aminolevulinate acid (ALA) to form a monopyrrole, porphobilinogen (PBG)¹. The prevalence of heterozygous ALAD deficiency was estimated to be <1% in Germany and =2% in Sweden². Clinical presentation of ADP includes autonomic, central, motor and sensory symptoms. We are reporting a case of a 10 years male with ADP presenting as neurovisceral symptoms.

The Case
A 10 years male from Western region of Nepal with normal developmental milestones and no prior neuro-visceral symptoms presented with history of severe abdominal pain for five hours duration associated with vomiting and difficulty in breathing. Eight hours later, he developed decreased level of consciousness. There was no history of fever, headache, seizures, and rashes, intake of canned food or drugs. There was no history of consanguinity or similar illness in family members. Patient was intubated at a district hospital because he was not maintaining saturation and then referred to our hospital. On examination, GCS was 8/15, heart rate was 128 beats/min (reference range 70-110), and respiratory rate was 28 breaths/min, BP: systolic at 70th and diastolic at 50th percentile, SpO₂: 100% on T-piece ventilation. CNS examination revealed 4 mm sluggishly reacting pupils, normal cranial nerves, no signs of meningeal irritation, decreased tone in both limbs with sluggish reflexes and upgoing plantars. Other systemic examination was normal.

Investigation showed WBC: 11,300/cu.mm with neutrophilic predominance, Hb: 13.2 g/dl, platelet: normal counts, ESR: 50 mm in 1st hour. Peripheral smear showed normocytic normochromic RBC, C-reactive protein: negative, normal electrolytes, urine, blood sugar, serum amylase and liver function test (LFT). ABG analysis showed respiratory acidosis, Chest X-ray and CT scan head was normal. CSF analysis was not done as patient was haemodynamically unstable.
During hospital stay, no fever or seizure occurred but there was persistent tachycardia and hypotension requiring inotropic support. On 3rd day of illness he was kept on mechanical ventilator due to repeated desaturation. On 4th day there was marked hypotonia with areflexia and equivocal plantars. On 5th day of illness, colour of urine changed to dark brown on exposure to sunlight (Fig:1). Based on history, examination and urinary findings, a clinical diagnosis of porphyria was made and urine was sent for Porphobilinogen (PBG) and ALA (d-aminolevulinic acid). Urine ALA was 3.18 mg/dl (0.0-0.55 mg/dl); whereas Urine PBG was 0.051 mg/dl (< 0.10mg/dl). Change of colour of urine on exposure to sunlight, increased urine ALA and normal urine PBG suggested a diagnosis of ADP. So 300 grams of 10% dextrose was started. Erythrocyte ALAD could not be done as child developed persistent bradycardia and expired on 9th day of admission. Erythrocyte ALAD activity and urinary ALA were planned in both parents but could not be done due to financial constraints.

**Discussion**

Porphyrias are a group of inherited diseases caused by deficiency of enzymes of the heme synthetic pathway, resulting in accumulation of porphyrins and their precursors. They are classified into hepatic porphyrias with neurological manifestations and erythropoietic porphyrias with no neurological symptoms. ADP, a type of hepatic porphyria is an autosomal recessive condition resulting from deficiency of ALAD. Its prevalence in South East Asia and Nepal is not known; most cases are reported in adolescent males.

ADP presents with severe neurologic manifestations similar to our case\(^5\). The typical neuropathy in ADP is an acute or subacute motor axonopathy predominantly affecting proximal muscles symmetrically and can lead to tetraplegia, with respiratory and bulbar paralysis\(^3\). Initially, tendon reflexes may be hyperactive and become absent later as in this case. Weakness is usually preceded by episodes of abdominal pain secondary to autonomic neuropathy\(^4\). Abdominal pain is the most common manifestation occurring in 85-95% of cases whereas tachycardia is the most common clinical sign\(^5\). Other symptoms of autonomic dysfunction include hyper or hypotension, diaphoresis and hyponatremia. In this case tetraplegia was preceded by abdominal pain, and during PICU stay persistent tachycardia and hypotension developed, which was followed by bradycardia and hypotension.

Acute Encephalitis as a diagnosis was excluded as there was no fever, convulsions, bizarre movements, or hallucinations, whereas weakness in descending pattern and early bulbar involvement made Guillain-Barré syndrome (GBS) less likely.

In our case urinary ALA was raised whereas urinary PBG was normal suggesting the diagnosis. Other causes of increased urine ALA were excluded. Normal LFT excluded acute hepatitis, absence of history of chronic exposure, acute presentation, absence of microcytic hypochromic anaemia excluded lead poisoning, however zinc protoporphyrin was not done for definitive diagnosis. Late presentation, normal odour and absence of hepatic and renal involvement excluded tyrosinemia type-I.

**Confirmation of diagnosis can be done by** erythrocyte ALAD which was not possible in our case as child died before the investigation could be done. Normal Urinary ALA of both parents along with half-normal activity of erythrocyte ALAD would have established the diagnosis but it was not possible due to financial constraints.
Management is symptomatic along with dextrose and hemin. Carbohydrate acts by down-regulation of heme synthesis whereas hemin replenishes the depleted heme pool and provides negative feedback on heme synthesis⁵. However hemin was not available in our part of the world. As this is an autosomal recessive disorder, genetic counselling could have been done and screening should be advised for other siblings as well.

**Conclusion**

Acute Porphyria should always be suspected in patients with gastrointestinal symptoms and neuro-psychiatric manifestations. Families of these individuals should be subjected to suitable enzyme tests, to screen asymptomatic relatives.

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**Reference**