MENKES’ SYNDROME: REPORT OF A CASE

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ABSTRACT

An 8 month old boy is presented with clinical and laboratory features of Menkes’ kinky hair syndrome. A brief discussion ensues.

Keywords: Menkes’ syndrome, copper (Cu)


INTRODUCTION

Menkes’ kinky hair disease, also known as trichopoliodystrophy, X-linked copper deficiency, and steely hair disease (with similar hair changes, seen in copper-deficient sheep), was first described in 1962 by Menkes and associates. Due to lack of a certain cellular carrier protein(s), caused by a mutation at a specific gene locus on chromosome X—presently narrowed to X.q.13.3(4)—copper is not delivered to various copper containing enzymes, subserving various systemic and neurological functions, resulting in a progressively fatal neurodegenerative disease. Major manifestations of this disease include abnormal hair, abnormal facies, progressive cerebral degeneration, bone changes, arterial rupture and thrombosis, and hypothermia (Table I). Occipital horn syndrome (also known as X-linked cutis laxa, and X-linked Ehlers-Danlos type IX), is a closely related disease presenting with inguinal hernias, chronic diarrhea, bladder and ureteric diverticula, and skin and joint laxity. Ossified and palpable occipital horns with milder changes in serum copper and ceruloplasmin distinguish this disease from the classical Menkes’ disease. Milder forms of Menkes’ disease, presenting later in life, have also been described.

We report an 8 month old boy fulfilling most of the above clinical stigmata for Menkes’ disease.

Case report

An 8 month old boy from Yasooj was referred to the Pediatric Neurology Ward with refractory upper extremity, eye, and body jerks noted since 5 months of age. He was born by normal vaginal delivery after a full term pregnancy at the hospital. Pregnancy was only complicated by vaginal spotting in the first trimester. He cried somewhat later than normal at birth, and a brief resuscitation was needed. Shortly after birth his serum bilirubin rose up to 11.38 mg/dL with a direct level of 1.64 mg/dL, for which no therapy was required.

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Fig. 1. The patient demonstrating hair and facial features of the disease.
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Fig. 2. The patient’s EEG.

was offered. Postnatally, he was hospitalized for what was
presumed as “sepsis” and was treated accordingly and
discharged. Since 5 months of age he started to have
episodes of brief (several seconds up to 1 min.) increase
in body tone with flexion of the head and body, upward
eye deviation, drooling, and lip cyanosis. Upon referral to a
pediatrician in Yasooj he received subsequent courses of
phenobarbital and phenytoin with no effect. He was noted
to be hypotonic and inattentive to his parents and
surroundings.

Nonconsanguineous parents gave a history of 2
miscarriages at the second trimester of each of the two
pregnancies, and no other sibling or close relative was
known to have the same problem as the patient. He received
his vaccinations as routine. Before 8 months of age he
could not hold his neck or roll over, and did not pay any
attention to his parents or any other visual or auditory
stimuli.

On physical exam he was found to be hypotonic with a
certain facies (Fig. 1.). Head circumference: 41 cm (<2 SD
for age); chest circumference: 41 cm; length: 64 cm
(about 5% for age); weight: 7 kg (about 5% for age); body
temperature ranged from 36°C in the mornings to 37.5°C in
the afternoons. Anterior fontanelle: 1.5 x 1 cm, not bulged;
posterior fontanelle, closed. Positive findings were global
hypotonia with preserved deep tendon reflexes, a bony
prominence at the right parietal area of the head, total
inattention to visual or auditory stimuli, crying only for
hunger, kinked, blond hair, bilaterally positive Babinski
sign, and no organomegaly. Paraclinical work-up revealed
the following abnormalities:

1. EEG: slow background, periods of relatively higher
voltage sharp waves followed by slowing (pseudoperiodic
pattern?) (Fig. 2).

2. Brain CT: bilateral temporo-occipital, and to a lesser
degree, frontal cortical and white matter atrophy (Fig. 3).

3. Skull series: increase in right parietal bone outer table
thickness—possible previous subperiosteal hematoma (Fig.
4).

4. Serum ceruloplasmin level: 0.052 g/dL (normal:
0.233-0.402 g/L).

5. Serum copper (Cu) level: 12.4 µg/dL (normal: 70-
150 µg/dL).

6. Pili torti: the patient’s hair was fragile, several shafts
always found shed on his bed mattress on daily hospital
rounds, and under low power light microscopy was twisted
at several millimeter intervals. Due to excess shedding of
hair the patient had a sparse hair population on his head.
The patient did not show hyperelasticity, joint
hypermobility, easy bruisability, or paper-thin scars on his
skin.

7. Long bone X-ray (Fig. 5): curving and decreased
density of the shaft.

8. A near 10 cm diverticulum at the postero-superior
aspect of the bladder. Other biochemical tests were
Table I. Clinical and paraclinical features of Menkes’ disease

- Hx of premature delivery, neonatal hypothermia and hyperbilirubinemia frequently elicited
- Hypothermia, poor feeding, poor weight gain, seizures, hypotonia, progressive deterioration of all neurologic functions
- Cherubic facies, depressed nasal bridge, pale optic discs, retinal degeneration, emphysema, iris cysts, gastric hemorrhage, subdural hematoma, colorless, friable, twisted hair (pili torti)
- Trichorrhexis nodosa (fracture of hair shaft at regular intervals)
- Very low serum copper and ceruloplasmin levels
- Decreased liver and brain Cu/increased intestinal mucosal copper
- Reduced activity of cytochrome oxidase, superoxide dismutase, tyrosinase, dopamine beta-hydroxylase, lysyl oxidase
- Poor Cu absorption/L.V. and subcutaneous Cu → serum Cu and ceruloplasmin increase
- Urinary tract dilation/bladder diverticula → rupture or U.T.I.
- X-ray: progressive osteoporosis, flared anterior ribs, periosteal reaction, increased density of provisional zone of calcification, small mandible, scalloping of posterior surface of the vertebral bodies; differential Dx. of bone changes, child abuse and scurvy
- Brain CT: macroscopic changes of brain degeneration
- Angiography: elongated, tortuous arteries in the cerebral and systemic vascular system with variable narrowing and dilatation and increased branching points
- Death: usually between 3 months-3 years (most often at 12 months).

unremarkable. More elaborate biochemical tests were not feasible. Angiography was not performed.

In his 2 hospital admissions and outpatient follow-up,
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the patient had not responded to subsequent trials of phenobarbital, phenytoin, prednisolone, clonazepam, vitamin B₆, vigabatrin, and carbamazepine, almost all the anti-convulsants available in our region. No parenteral copper was available in Iran for a trial of therapy and his seizures have remained intractable, although his general condition is fairly stable, except for some irritability and excessive crying.

DISCUSSION

This infant represents a typical case of Menkes’ kinky hair disease. He manifested hyperbilirubinemia shortly postnatally, a sepsis-like episode at early infancy, and neural, urinary, hair, bone, and biochemical changes peculiar to this disease. His seizures have resisted any available anti-convulsant therapy. Various parenteral copper compounds have been tried in animal models⁶ and affected humans²,³ with resultant enzymatic and some clinical recovery. Thus far efforts to find a parenteral form of copper have failed, as have all drug trials to bring the patient’s seizures under control. Our case demonstrates the importance of looking for a systemic disease whenever an infant or child presents primarily with neurological signs and symptoms.

REFERENCES