Wilson’s Disease with Neurological Presentation, without Hepatic Involvement in Two Siblings

MANI KANT KUMAR, VIJAY KUMAR, PRAPUL KUMAR SINGH

ABSTRACT
Wilson’s Disease (WD) is a rare, autosomal, recessive, inborn error of the copper metabolism, which is caused by a mutation in the copper-transporting gene, ATP7B. The presentation is usually neurologic or hepatic, which is seen in 40% of the patients. The diagnosis depends primarily on the clinical features, the biochemical parameters and the presence of the Kayser – Fleischer ring. Here, we are reporting two siblings who were affected by Wilson’s disease, with only neurological manifestations, without any hepatic involvement.

Key words: ATP7B, Dystonia, Kayser – Fleischer ring, Wilson’s Disease

INTRODUCTION
Wilson’s Disease (WD), which is also known as hepatolenticular degeneration, was first defined by Dr Samuel Alexander Kinnier Wilson in 1912, [1]. Wilson’s disease (WD) is a rare, autosomal, recessive inborn error of the copper metabolism, which is caused by a mutation in the copper-transporting gene, ATP7B. The WD gene, ATP7B, is located on chromosome 13q14.3, which encodes a metal-transporting P-type adenosine triphosphatase (ATPase), which is expressed mainly in hepatocytes. The incidence of WD is estimated to be 1 in 30,000 individuals and the carrier frequency is approximately 1 in 90 [2]. An absent or a reduced function of the ATP7B protein leads to a decreased hepatocellular excretion of copper into bile. This results in hepatic copper accumulation and injury. Eventually, copper is released into the bloodstream and it is deposited in other organs, notably in the brain, kidneys, and the cornea. The hepatic production and the secretion of the ceruloplasmin protein without copper and apoceruloplasmin, result in the decreased blood level of ceruloplasmin which is found in most of the patients with WD, due to the reduced half-life of apoceruloplasmin [3]. The manifestations are more likely to be hepatic in early childhood and to be neurological in adolescents [4]. The neurological manifestations of Wilson’s disease can vary extremely and they are often diagnosed after long delays. Here, we are reporting Wilson’s disease with only neurological presentations in two siblings.

CASE REPORT
Case 1: A 14 years old Hindu boy, who was the youngest of 4 siblings, who was a product of a non – consanguineous marriage, presented with pain in right knee joint and difficulty in walking for 4 months, 10 months back, which was treated by many physicians and neurologists for these problems but the diagnosis was missed. There was no improvement, and he was then brought to our institute. This abnormal movements subsided during sleep. The patient also had progressive dysarthria. His elder brother also had a history of very slowly progressive abnormal movements and dysarthria. There was no history of measles during early childhood and convulsion. His developmental milestones were normal. On examination, his vital signs were found to be normal. There was no pallor, icterus or significant lymphadenopathy. His nervous system examination revealed dystonia, exaggerated deep tendon reflexes, ankle clonus and a positive Babinski’s sign. His muscle power was >4/5 in all the limbs. The Kayser – Fleischer ring (K – F ring) was visible on both sides by the naked eye [Table/Fig-1], which was confirmed on an ophthalmoscopic examination which was done by an ophthalmologist by slit lamp examination. His other systemic examinations did not reveal any abnormality.

His complete blood count revealed haemoglobin 10 gm/dl, a total leukocyte count of 5600/cumm (neutrophil 62%, lymphocytes 33%, monocytes 2% and eosinophils 3%) and platelets ~340,000/ cumm. The serum electrolytes and the renal functions were normal. The total serum bilirubin was 0.6 mg/dl (direct bilirubin 0.2 mg/dl), the total serum protein was 6.8 gm/dl (Albumin 4.2 gm/dl) and the serum transaminases (AST, ALT) and Alkaline Phosphatise (ALP) were 38, 24, and 168 IU/L respectively. His Prothrombin Time (PT) and his activated Partial Thromboplastin Time (aPTT) were within normal limits. His serum ceruloplasmin was 95 mg/L (normal 180–350 mg/L) and his 24 hours urine copper excretion was increased to 160 µg (normal 24 hours urine excretion 20 – 50 µg). Liver biopsy was not done because his parents had not given their consents for that invasive procedure. Genetic testing for mutation analysis was not done due to financial constraints. His urine routine and microscopic examinations were normal. Ultrasonography of the abdomen was normal. Magnetic resonance imaging (MRI) of the brain, on the T2 weighted axial sequence through the pons revealed the “Face of the miniature Panda” [Table/Fig-2]. The T2 weighted image also revealed hyperintensities in the thalami and the pontine tegmentum.
(normal 180-350 mg/L) and his 24-hours urine copper excretion was increased (175 µg). His urine routine and microscopic examinations were normal.

The diagnosis of Wilson’s disease with a neurological manifestation was made on the basis of the dystonia, the K–F ring in both eyes, the low serum ceruloplasmin, and the 24-hours urinary copper excretion. His parents were explained regarding the prognosis of the disease and the available treatment in our country. Both the patients were started on oral zinc (as zinc acetate) at a dose of 1mg/kg/dose of elemental zinc 8 hourly and on Trihexiphenidyl to control the dystonia. He was advised to avoid food with a high copper content, such as chocolates, nuts, legumes, mushrooms, shellfish and liver. His parents were also advised to avoid the use of copper utensils in the household for the storage of water and for cooking food. After two weeks of treatment, his dystonia reduced in intensity.

DISCUSSION

Wilson’s Disease (WD) is a rare, autosomal, recessive, inborn error of the copper metabolism, which is caused by a mutation in the copper-transporting gene, ATP7B [2]. Copper first accumulates in the liver; after the liver storage capacity for copper gets saturated, copper gets redistributed, with accumulation in the nervous system, the cornea, the kidneys and other organs [5]. Most of the patients present in the second decade of life with a primary hepatic involvement, with the remainder of the patients presenting during the third and fourth decades, with a primarily neurologic or a psychiatric presentation [6]. However, in our study, both the cases presented with neurological manifestations in the 2nd decade of life (adolescent), which was similar to the findings of Kalra et al., as was seen in their study which was done at the All India Institute of Medical Sciences (AIIMS) [4]. In most of the Indian studies, the disease was found to manifest at a younger age in Indian children. This could be most likely due to higher average intake of copper in India, which ranges from 5.7 – 7.1 mg/day and it is higher than the reported 0.34 – 1.1 mg/day in the western countries. The practice of cooking food and storing drinking water in copper/copper alloy pots might be contributory [5,7].

In Wilson’s disease with neurological presentations, the sympotomatology is predominantly extra pyramidal, like dystonia, tremors, dysphasia, dysarthria, and ataxia. The neurological symptoms are usually secondary to the cerebral copper deposition, which is sufficient to destroy the nerve cells. Both our cases had neurological manifestations, predominantly dystonia, dysarthria and some cognitive impairment. None of our patients had either a clinical nor a biochemical evidence of a hepatic involvement.

The serum ceruloplasmin levels should not be considered for making a definitive diagnosis, because they are normal in up to 10% of the affected patients and are reduced in 20% of the carriers. The Kayser – Fleischer rings can only be diagnosed definitively by an ophthalmologist by using a slit lamp.

Urine copper is an important diagnostic tool but it must be collected carefully to avoid contamination. An estimation of the 24-hour urinary copper excretion is another reliable test which can be done for the confirmation of WD. The normal excretion of copper is between 20 and 50µg per day; in the cases of WD, the excretion is increased to in excess of 100 Normal excretion is between 20 and 50 µg per day; in cases of WD, excretion is increased to in excess of 100µg per day. The “Gold standard” for the diagnosis remains a liver biopsy with quantitative copper assays. The affected patients

Case 2: A 18 years old Hindu boy, the 2nd youngest of 4 siblings (the elder brother of the above mentioned patient), a product of a non-consanguineous marriage, presented with a history of progressive dystonia for the past 5 years and a progressive, purposeless, repetitive, abnormal movement in the form of dystonia for the past 2 years. This abnormal movement subsided during sleep. He had been treated by many physicians and neurologists with antiepileptic drugs, without any improvement. There was no history of measles during early childhood and convulsion. On examination, his vital signs were found to be normal. His nervous system examination revealed dystonia, dystonia, exaggerated deep tendon reflexes, ankle clonus and a positive Babinski’s sign. His muscle power was normal. On ophthalmoscopic examination, the Kayser – Fleischer ring (K – F ring) was found to be present on both the sides. His other systemic examinations were normal. His haemogram, liver function test and renal function test and ultrasonography of the abdomen were normal. His serum ceruloplasmin was 110 mg/L.
have values of > 200μg/gm dry weight of the liver. The copper
stains are not reliable [8].

In both the cases, we had made the diagnoses on the basis of the clinical
displays, the serum ceruloplasmin levels, the 24 hours
urinary copper excretion and neuroimaging (MRI). Although a liver
biopsy with quantitative copper assay is the “Gold standard”,
due to the parents’ refusal, liver biopsies were not done in our
patients.

MRI is a very sensitive method for revealing the abnormalities
in WD. On the T1 – weighted sequence, hypointensities in the
basal ganglia are seen in two-thirds of the cases while in the T2 –
weighted images, hyperintensities in the basal ganglia, the white
matter, the thalamus or the brainstem are seen. These abnormalities
are caused by a neuronal loss, gliosis, degeneration of the fibres,
and vacuolisation, which are associated with the increased water
content in the brain. The typical ‘face of the giant panda’ can be
seen in the midbrain on the T2 – weighted axial MRI sequence of
the brain and the ‘face of the miniature panda’, can be seen in the
tegmentum region of the pons in the T2 – weighted sequence
[9,10]. In our case, the T2 – weighted axial sequence through
pons revealed the “face of the miniature panda”, which was also
suggestive of Wilson’s disease.

Penicillamine was previously the primary anticonvulsant
therapy, but now, it plays only a minor role because of its toxicity and
because it often worsens the existing neurologic disease if it is
used as the initial therapy. If penicillamine is given, it should always
be accompanied by 25 mg/d of pyridoxine. Trientine is a less
toxic chelator and it supplants penicillamine when a chelator is
used as the initial therapy. If penicillamine is given, it should always
be accompanied by 25 mg/d of pyridoxine. Trientine is a less

CONCLUSION
The present cases gave us valuable information that Wilson’s
disease is an uncommon, autosomal, recessive, metabolic
disorder, which is often missed but easily treatable. A high index
of suspicion is required while dealing with adolescents and
young adults with abnormal movements and neurobehavioural
abnormalities. A high degree of suspicion and an early detection
of WD is critical, because an early initiation of the treatment can
prevent a catastrophic outcome.

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Author(s):
1. Dr Mani Kant Kumar
2. Dr Vijay Kumar
3. Dr Praphul Kumar Singh

Particulars of Contributors:
1. Assistant Professor, Department of Pediatrics,
2. Assistant Professor, Department of Biochemistry,
3. MBBS, DCH, Clinical Tutor, Department of Pediatrics,
Narayan Medical College and Hospital, Jamuhr,
Sasaram, Dist – Rohtas, Bihar – 821305, India.

Name, Address, E-Mail Id of The Corresponding Author:
Dr Mani Kant Kumar,
Assistant Professor, Department of Pediatrics,
Narayan Medical College and Hospital, Jamuhr,
Sasaram, Dist – Rohtas, Bihar – 821305, India.
Phone: +91 9162095353
E-mail: manikant7@yahoo.com

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