Introduction

Infantile tremor syndrome (ITS) is a syndrome characterized by anemia, skin pigmentation, tremors, physical, and mental regression. It has been reported in children between 5 months and 3 years of age with a male predominance. The exact incidence of ITS is not available. However, in various studies 0.87–1.55% hospital admissions were due to ITS. The exact etiopathogenesis of this condition remains obscure. The purpose of this case series was to determine factors responsible for its continuing prevalence despite better knowledge and weaning practices.

Case Report

We have studied 9 patients presenting to us with typical features of ITS and pre-ITS in a tertiary care hospital of hilly terrain [Table 1]. A detailed history and examination were carried out in each patient followed by investigations. Investigations included complete hemogram, peripheral smear to type the anemia and malaria parasite, serum electrolytes, renal and liver function tests along with sepsis screen, and urine routine examination. Blood and urine cultures, serum B12 levels, chest X-ray, and tuberculosis screen were done when indicated. Written informed consent was taken from all. Resources available at the hospital were utilized for the same.

Total 9 patients, 3 males and 6 females presented over the period of 1 year. The mean age of presentation was 13.5 months, ranging from 7 months to 18 months. All (100%) patients admitted with the complaint of lethargy and weakness with a mean duration of 21 days. Neuroregression was present in all (100%) with a mean delay of 7 months ranging from 4 to 11 months in all spheres of development. Tremors were present in 5 (55%) and hypotonia in 2 (22%) reflecting neurological changes. All (100%) were healthy term babies born by normal vaginal delivery. History of faulty feeding was present in all. Seven (78%) were exclusively breastfed and 2 (22%) improperly weaned with diluted milk. Protein energy malnutrition was present in 5 (55%) out of which 2 (40%) had Grade I and 3 had Grade II (60%) protein energy malnutrition as per IAP classification. Three (33%) had severe wasting according to WHO criteria. There was no stunting in any patient. OFC records of all patients were within normal range for their age [Table 2].
All patients were hemodynamically stable at admission. All had severe pallor (100%) with knuckle hyperpigmentation and sparse hair in most. Three (33%) also had signs of rickets. Hepatomegaly was present in 4 (44%), splenomegaly in 2 (22%), and pneumonia in 1 (11%).

Anemia (4–7.5 g/dl) was present in all. Peripheral blood film examination revealed the macrocytic picture in 4 (44%), dimorphic in 4 (44%), and microcytic in 1 (11%). Chest X-rays had infiltrated in 3 (33%) but workup for tubercular screening was negative. Sepsis screen was positive in 6 (66%). Urine cultures were positive in 2 (22%) with *Escherichia coli* and *Pseudomonas* in 1 each. Dyselectrolemia was present in 2 patients with hyponatremia in one and hypokalemia in other. Serum B12 levels were low in 4 out of 5 patients but could not be performed in all due to financial constraints. Renal and liver functions were normal in all the patients.

The mean duration of hospital stay was 9 days ranging from 4 to 16 days. Total 6 (66%) with positive sepsis screen or cultures received antibiotics for the average duration of 9 days. Two also required blood transfusions. All signs and symptoms improved after treatment with supplements such as Vitamins A, D, K, B-complex, zinc, calcium, and magnesium along with dietary management. Propranolol was given to patients with ITS. On monthly follow-up, those with pre-ITS had better catch up in growth and development in all spheres. Some degree of developmental lag persisted in those with ITS.

**Discussion**

Infantile and pre-ITS is a well-known clinical entity reported primarily from the developing nations.[4-6] Despite its decreasing incidence, the finding of additional clinical features made us present this series. The etiology of ITS is still elusive. Among various theories, the nutritional theory is the most accepted. Vitamin B12 deficiency has been found to be associated with ITS in many studies.[1] It is usually seen in children who are exclusively breastfed. Iron, magnesium, and zinc deficiency have also been postulated to cause ITS. Associated nutritional deficiencies such as protein, Vitamin A, D, C, and B-complex may be present.[7,8] Other speculations for its etiology include viral encephalitis and degenerative processes.[1,4,5]

Our series also found that all children with this syndrome had anemia, lethargy, and neuroregression and presented throughout the year with no seasonal clustering as previously stated.[4,6,9] Furthermore, the type of anemia in this disease is not specific. It reflects the pattern of prevalence of the hematinic deficiencies in the region. However, unlike previous reports, there were more female patients, which may be due to bias in nutritional status.

The study adds the need for rational antibiotic use in patients of ITS, unlike many previous reports where all patients have been administered antibiotics. It reinforces the concept of sepsis screens and avoiding antibiotic abuse. The disease process may be superadded by urinary infections probably due to lower immunity state as a result of malnutrition, the leading cause of acquired immunodeficiency. It is also a reflection of changing epidemiology with no seasonal
variation, showing the role of non-viral etiologies although previous studies show a seasonal pattern of disease.[10,11] It highlights the need for better education for weaning practices and need for better community support.

References


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