Neuromyotonia (Isaacs’ Syndrome) In A Patient With Shistosomiasis With Good Response To Plasmapharesis: A Case Report

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CASE REPORT

A 61 years old male patient, farmer from Abo Humos, married with 7 offspring, presented with subacute onset, progressive course of painful spasms of the right upper limb since 1 year (Distal more than proximal) with no precipitating factor that progressed to stiffness of the right upper limb. The condition progressed to involve the left upper limb within 3 months in the form of stiffness and frequent spasms followed by the trunk and the paraspinal muscles, and lastly the lower limbs were affected since 4 month.

The patient developed fasciculations through the course of the illness. It was generalized, spontaneous and involving the eyelids, face, upper and lower limb. Recurrent attacks of muscle cramps were of daily frequency mainly involving the upper limbs. Increase sweating was also reported by the patient. He complained of mild paresthesia involving both upper limbs. No bulbar, sphincteric or cognitive manifestations and no constitutional manifestations or weight loss was seen. No past history of DM or hypertension was found. The patient had a history of Bilharziiasis since 50 years, for which he received tartar emetic. The patient had also a history of oral lesions since 5 years that recurred after surgical intervention. Family History and Drug History were negative.

General examination revealed average body built with stiffness of both hands (Fig 1). Consciousness and MSE were normal. Head and Neck examination was normal except for a submandibular lymph node. Gait was in short steps due to stiffness of the lower limb. Speech and articulation were normal. Cranial nerve examination only showed tongue wasting and fasciculations. Motor examination revealed average muscle status with no wasting. Stiffness all over more in the upper limbs and more distally was found. There was no weakness and the patient showed spontaneous gross fasciculations in both upper and lower limbs and in the face over the masseter muscle and around the eye. Sensory examination revealed short glove and stocking hypothesia. Coordination was normal and plantar reflex was flexor bilaterally.
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Figure 1
Demonstrates muscle contracture and stiff posture more on the right upper limb

Routine laboratory investigations showed pancytopenia with WBC count of 1.900 mm$^3$, RBCs were 3.3 million/mm$^3$ and Platelet count was 20,000/mm$^3$. Other routine laboratory investigations were normal. Collagenic profile and Thyroid Function were Normal. Anti-VGKC Abs were not available in Alexandria.

CT Chest was normal. CT Abdomen showed Liver cirrhosis, mild periportal fibrosis, huge splenomegaly, portal hypertension and a polyp arising from sigmoid colon.

Figure 2
Fig. 2

Nerve conduction study and EMG were done and confirmed the diagnosis showing neuromyotonic discharge associated with denervation and polyphasic units with fibrillation, positive sharp waves and continuous muscle fiber activities (Fig 2).

The patient had many suspicious lesions; the recurrent oral lesion was investigated and was diagnosed as non-neoplastic vascular malformation.

The patient had a submandibular lymph node that was biopsied to exclude lymphoma and it was found to be reactive and non neoplastic. The pancytopenia was investigated thoroughly and together with the history of bilharziasis and blood film that showed Normocytic normochromic anaemia (with anisopoikilocytosis), neutropenia (toxic granules) and thrombocytopenia and the bone marrow aspiration that revealed hypercellular marrow with megakaryopoisis and granulocytic hyperplasia. It was clear that this is a picture of hypersplenism. Colonoscopy and biopsy of the polyp showed adenomatous nature with no granuloma nor malignancy (Bilharzial Polyp) Paraneoplastic syndrome as a cause was ruled out in our patient as well as hereditary causes.

Our patient received 5 sessions of plasma exchange (0.4mg/kg) performed on an every-other-day schedule with marvelous improvement in myokymia, pseudomyokymia, gait difficulties and sweating and he was discharged on Carbamazepine 1000 mg per day.

DISCUSSION
Acquired neuromyotonia (Isaacs’ syndrome) is a rare disorder where hyperexcitability of peripheral motor nerves leads to incapacitating muscle twitching, cramps, myokymia, pseudomyotonia (slow muscle relaxation after forceful contraction) and mild weakness (1). The muscle cramp may be prominent and accompanied by excessive sweating and weight loss (2). This uncommon disorder was first described in 1961 by Isaacs in his paper ‘A syndrome of continuous muscle-fibre activity giving the triad of myokymia, muscular stiffness, and decreased deep tendon reflexes the paper’s name (3).

Isaac’s syndrome has been long recognized by several physicians (4), however its rarity and the variability of its clinical manifestation and ways of presentation is probable the most important reason why its frequently misdiagnosed or wrongly treated (5).

The diagnosis of Isaacs’ syndrome is based on clinical features and electromyographic findings. The cardinal features consist of myokymia, pseudomyotonia and stiffness of trunk and limbs. Stiffness without severe pain is more pronounced in the distal than proximal muscles. This abnormal activity persists during sleep. Dyspnea may occur when respiratory muscle is involved. There have been only a few reports of bulbar and laryngeal involvement in Isaacs’ syndrome. The tongue and jaw become stiff, making swallowing difficult, and the voice turn hoarse. Associating symptoms include weight loss and excessive sweating (1,2). Most patients are sporadic. This is related to the autoimmune mechanism where the autoantibodies are usually detected against the Voltage-Gated Potassium Channels (VGKCs) (6,7).

This syndrome may also be related to other autoimmune diseases such as chronic inflammatory demyelinating...
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polyneuropathy, myasthenia gravis or the presence of antiacetylcholine receptor antibodies(8). The association to hematologic malignancies such as thymoma (9), plasmacytoma(10), Hodgkin’s lymphoma (11) and bronchogenic carcinoma paraneoplastic syndromes, has been documented (12).

Clinical evidence suggesting a possible autoimmune etiology included the presence of oligoclonal bands in the spinal fluid of some patients and clinical improvement following plasma exchange (13).

Classical electrodagnostic studies detect myokymic and neuromyotonic discharges. In addition, fasciculation, doublet, triplet, multiplet and positive sharp waves are also demonstrated in this syndrome. Stimulus-induced repetitive discharges, usually seen after the M wave, are also demonstrated during motor nerve conduction studies (14,15).

Treatment of Isaac’s syndrome with antiepileptic drugs or immunotherapy often improves the clinical and electrophysiologic findings (16). Carbamazepine, phenytoin, lamotrigine and sodium valproate can be used alone or if necessary in combination. Paraneoplastic neuromyotonia usually improves after treatment of the underlying cancer (17). In patients whose symptoms are debilitating or refractory to symptomatic therapy, immunomodulatory therapies should be tried (18,19). Plasma exchange often produces useful clinical improvement lasting from 6 weeks up to 6-17 month accompanied by a reduction in EMG activity and a fall in VGKC antibody titres (7,20). IVIg is useful in combination. Paraneoplastic neuromyotonia is a causal relationship or a mere comorbidity is a matter of debate, as there is lack of publications in this issue, and further research is warranted.

References
5. Foyaca-Sibat H, Ibanez-Valdes LdeF, Awotedu A. Acute renal failure due to herbal medicine intoxication in acquired neuromyotonia.

Only a minority (<5%) of patients will develop CNS symptoms due to schistosomiasis, with cerebral complications being more prevalent than spinal (24).

Neuroschistosomiasis (NS) is the most severe presentation of schistosome infection; cerebral invasion is mostly caused by S. japonicum, with spinal cord involvement due mainly to S. mansoni or S. haematobium (25).

Whether the relation between Shistosomaiasis and Isaac's syndrome is a causal relationship or a mere comorbidity is a matter of debate, as there is lack of publications in this issue, and further research is warranted.
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