Ischemic Stroke and Epilepsy in a Patient with Tourette’s Syndrome: Association with the Antiphospholipid Syndrome and Good Response to Levetiracetam

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Abstract: The role played by different humoral factors, including antiphospholipid antibodies, in the pathogenesis of Tourette syndrome (TS) is still presently unclear. We present a patient with chronic and severe TS who, at the age of 16 years, presented an ischemic stroke in the left posterior cerebral artery and/or postero-inferior temporal branch of the left medial cerebral artery. A complete study was negative with the exception of a positive lupus anticoagulant. The stroke was related with the primary antiphospholipid syndrome (APS). The stroke manifested visual abnormalities and thereafter by secondary generalized complex partial seizures. The epileptic syndrome was initially difficult to control but responded dramatically to levetiracetam. With this therapy, the manifestations of TS, especially the tics, improved. We conclude that some TS cases may present APS. In addition, levetiracetam may be useful in the management of TS. Further investigations should pursue both these facts.

Keywords: Tourette syndrome, antiphospholipid syndrome, antiphospholipid antibodies, levetiracetam.

INTRODUCTION

The pathogenesis of the Tourette syndrome (TS), the main component of the primary tic-syndrome [1], is presently unclear. Both genetic and non-genetic factors are important considerations [2-8]. The antiphospholipid antibodies (aPLAs) are a group of heterogeneous antibodies which bind with negatively charged phospholipids. When present in association with vascular thrombosis or recurrent fetal loss, the diagnostic criteria of the antiphospholipid syndrome (APS) are fulfilled [6]. The neurological complications of APS are varied and include movement disorders and chorea. However, the mechanism underlying non-thrombotic movement disorders in APS remains presently unknown. A direct immune interaction of aPLAs with diverse brain structures has been hypothesized. Presently, there is conflicting evidence of an association of aPLA with TS [7-8].

Herein, we present a young patient with severe TS who presented an ischemic stroke in association with APS and secondarily presented epileptic syndrome. We reappraise the hypothesis that aPLAs may play a role in TS. In addition, we highlight the fact that levetiracetam therapy had a significant beneficial effect in the control of the tics and the epileptic syndrome.

CASE REPORT

The patient is presently a 25 year-old male with a family history for a transient tic-disorder present in the infancies of both his father and paternal grandfather. He was born after an uneventful pregnancy and birth. From the age of 8 years he presented a progressive and severe complex motor and vocal tic disorder including echolalia, palilalia and coprolalia in association with obsessive-compulsive behavior. He was diagnosed of TS. The severity of the syndrome required multiple evaluations over the years in various neurological centers and by the psychiatry department. He was treated with multiple combination therapies with neuroleptics, anticholinergics and tricyclics but the clinical response was basically poor. The tic disorder had a severe and negative impact in his family, social relations and on his academic results. In this period, an extensive and complete study including neuroimaging studies with CT and MRI were normal. At the age of 16 years, he presented sudden-onset blurring in his left hemifield followed by a tonic-clonic seizure of minutes’s duration. No motor, sensory or language abnormalities were present. The left-sided visual abnormalities improved over a few weeks. However, the seizure disorder persisted. The seizures were fairly stereotyped and consisted in paroxistic onset of
an oppressive epigastric sensation, tachycardia, diaphoresis and visual alterations. These visual phenomena were rich and varied in type: metamorphopsia, positive phenomena of complex nature in the left hemifield, and a vague sensation described as “far vision”. Some of these episodes were followed by a head turning to the left and impaired consciousness and confusion of variable but, generally brief, duration. He was diagnosed of secondary generalized complex partial (autonomic, visual and versive) epilepsy and the patient was managed with antiepileptic drugs (AED) in various combinations (valproate, carbamacepine, phenobarbital and phenytoin). These agents were mainly effective on the generalized convulsive fits, but the other phenomena persisted, were frequent (2–4 per week) and motivated many consultations in the emergency room.

At the age of 21 years levetiracetam 1000mg bid was initiated as add-on therapy to phenytoin 300 mg/day and the seizure disorder came under complete and total control. Simultaneously, with the initiation of levetiracetam, the tic-disorder (both the motor and vocal tics) improved in parallel with the control of the seizure disorder. The patient maintains treatment with platelet inhibitors and levetiracetam; the dose of phenytoin is being progressively lowered (presently at 100 mg/day). He is seizure-free since the initiation of levetiracetam therapy.

The patient still presents occasional vocal and motor tics but is able to lead a satisfactory personal and social life. In addition, he has earned an employment as a public servant.

DISCUSSION

This patient with severe TS presented an ischemic stroke in the posterior cerebral artery and/or postero-inferior temporal branch of the right medial cerebral artery. A stroke work-up was negative with the exception of a positive lupus anticoagulant repeated on various occasions. Systemic diseases were excluded and the patient satisfied the diagnostic criteria for primary APS [6]. The stroke manifested visual abnormalities and secondary generalized complex partial seizures. Upon reaching a good control of the seizures the manifestations of the TS improved.

The pathogenesis and pathophysiology of TS are still presently unknown. Abnormalities in both the dopaminergic and, to a lesser extent, serotonergic neurotransmission within the basal ganglia and the cortico-ganglionic-thalamocortical circuitry have been suggested [2].

Both genetic and non-genetic factors may play a role in the pathogenesis of TS. The presence of certain cytogenetic abnormalities, including chromosomal translocations and inversions, and epigenetic risk factors suggest that certain genetic or chromosomal irregularities may be common to some cases of TS [2]. There is also evidence for an immunological basis in TS. Anti-basal ganglia antibodies appear to play a role in certain TS-spectrum disorders including the pediatric autoimmune neuropsychiatric disorders associated with streptococcal infection (PANDAS). These antibodies have been found in 65% of patients with atypical movement disorders [3]. Anti-basal ganglia antibodies are often found in children with Sydenham’s chorea. In this disease, antibodies seem to attack the basal ganglia by a mechanism of molecular mimicry triggered by a previous streptococcal infection [8]. In TS there are higher concentrations of antiputaminal antibodies compared to normal controls [5] and immunomodulatory treatment is effective in some patients. There are therapies that may act on a putative immune-mediated attack directed against the nigrostriatal or prefrontal dopaminergic pathways [3,9].

Some studies have approached the possible association between aPLAs and TS. Antiphospholipid antibodies have been identified in children with TS and have led to the speculation that these antibodies might play a pathophysiological and possibly pathogenic role in this condition [7]. If confirmed, the presence of aPLAs has diagnostic and therapeutic implications in TS. However, aPLAs in TS may represent an epiphenomenon rather than a pathophysiological mechanism [8]. Unfortunately, the number studies on this aspect are scarce and have not included patients with APS.

To our knowledge, this is the first report of a TS patient presenting a stroke in association with APS. Acknowledging a possible chance association, in some TS cases immune phenomena may be involved in the movement disorders. The presence of a particular genetic load or predisposition may increase susceptibility in developing specific antibodies and thrombotic phenomena.

Also of interest in this case is the favorable response of both the epileptic syndrome and the tic-disorder to
levetiracetam. It is possible that the seizure disorder may have aggravated the frequency and severity of the tics but we do not exclude a direct effect of levetiracetam on the favorable outcome of the tic-disorder [10].

In conclusion, this report indicates that aPLAs may be present in some patients with TS. We believe that the mechanisms that trigger the generation of these antibodies leading to vascular events and possibly movement disorders should be further investigated.

REFERENCES

[1] The Tourette Syndrome Classification Study Group. Definitions and classification of tic disorders. Arch Neurol 1993; 50: 1013-6.
[2] Albin RL, Mink JW. Recent advances in Tourette syndrome research. Trends Neurosci 2006; 29: 175-82.
[3] Edwards MJ, Trikouli E, Martino D, et al. Anti-basal ganglia antibodies in patients with atypical dystonia and tics. Neurology 2004; 63: 156-8.
[4] Church AJ, Cardoso F, Dale RC, Lees AJ, Thompson EJ, Giovannoni G. Anti-basal ganglia antibodies in acute and persistent Sydenham’s chorea. Neurology 2002; 59: 227-31.
[5] Singer HS, Giuliano JD, Hansen BH, et al. Antibodies against human putamen in children with Tourette syndrome. Neurology 1998; 50: 1618-24.
[6] Wilson WA, Gharavi AE, Koike T, et al. International consensus statement on preliminary classification criteria for definite antiphospholipid syndrome: report of an international workshop. Arthritis Rheum 1999; 42: 1309-11.
[7] Toren P, Toren A, Weizman A, et al. Tourette’s Disorder: Is there an association with the antiphospholipid syndrome? Biol Psychiatry 1994; 35: 495-8.
[8] Singer HS, Krumholz A, Giuliano J, Kiessling LS. Antiphospholipid antibodies: an epiphenomenon in Tourette syndrome. Mov Disord 1997; 12: 738-42.
[9] Perlmutter SJ, Leitman SF, Garvey MA, et al. Therapeutic plasma exchange and intravenous immunoglobulin for obsessive-compulsive disorder and tic disorders in childhood. Lancet 1999; 354: 1153-8.
[10] Awaad Y, Michon AM, Minarik S. Use of levetiracetam to treat tics in children and adolescents with Tourette syndrome. Mov Disord 2005; 20: 714-8.