OLANZAPINE-INDUCED OCULOGYRIC CRISIS IN A PATIENT WITH SCHIZOPHRENIA

Sir,

Oculogyric crisis is an acute dystonic reaction characterized by sustained upward deviation of eveballs and restlessness. Such reactions generally occur after administration of typical antipsychotics but are considered rare with atypical antipsychotics, particularly olanzapine. Until now, there are only two reports of olanzapine-induced oculogyric crisis from India, one in a patient of bipolar disorder^[1] and another in a child with post-encephalitic syndrome.^[2] Indeed, literature review on June 1, 2008, in ENTREZ PUBMED revealed only another German report in a patient with general anxiety disorder.^[3] Interestingly, although the commonest indication of olanzapine is schizophrenia, olanzapine-induced oculogyric crisis was never reported in schizophrenia. Below, we describe such a case.

A 25-year-old man, suffering from schizophrenia for 6 months, was initially treated with risperidone 8 mg/day and fluoxetine 40 mg/ day (for associated depressive symptoms). But soon, fluoxetine had to be replaced by imipramine 75 mg/day for poor tolerance. The patient improved significantly.

However, once his symptoms resolved, the patient stopped taking medicine and relapsed within 2 weeks. This time, a different consultant placed him on olanzapine 10 mg/day and imipramine 75 mg/day. After 1 month with this regimen, the patient reported partial improvement, along with tremor. As olanzapine is not known to cause tremor commonly,

imipramine was thought to be the reason and was stopped. Indeed, olanzapine was hiked up to 15 mg/day as the patient had partial improvement. But within a few days, the patient started having repeated episodes of sustained upward deviations of eyeballs, along with anxiety; restlessness; and backward flexions of neck, which transiently resolved with injection promethazine 25 mg. However, regular addition of trihexyphenidyl 2 mg/day failed to stop recurrences of the crises, which ultimately required replacement of olanzapine with risperidone. Naranjo's algorithm^[4] indicated a probability score of 8 of olanzapine-induced oculogyric crisis. With risperidone 3 mg/day, the patient is now maintaining remission without having tremor or oculogyric crisis any further.

High dopamine-acetylcholine antagonism or high striatal dopamine inhibition has been suggested to underlie neuroleptic-induced oculogyric crisis.^[1] So olanzapine with an intermediate level of D2-binding affinity^[5] is not expected to cause oculogyric crisis. Probably for this, this patient did not have any problem with olanzapine at lower level except tremor. But in higher doses, olanzapine has been shown to have high D2 affinity, increasing the chance of oculogyric crisis.^[5] Also, high anticholinergic property of imipramine may have prevented the occurrence of oculogyric crisis initially until it was withdrawn. However, this does not explain the failure of trihexyphenidyl to control this side effect. It may imply that probably trihexyphenidyl is not very effective in the treatment of repeated oculogyric crisis with olanzapine. Indeed, we may need to change the antipsychotic in such a case.

Recent Clinical Antipsychotic Trials of

Intervention Effectiveness (CATIE) study^[6] has seriously doubted the prevalent belief of atypical antipsychotics including olanzapine having lesser neurological side effects. Our case report supports this doubt. Although possibility of such a side effect is low, we still need a cautious approach regarding this agent, particularly while prescribing a higher dose.

RUDRAPROSAD CHAKRABORTY, ARUNIMA CHATTERJEE, SUPRAKASH CHAUDHURY

Department of Psychiatry, Ranchi Institute of Neuropsychiatry and Allied Sciences, Ranchi, India

Correspondence:

Dr. Rudraprosad Chakraborty Department of Psychiatry, Ranchi Institute of Neuropsychiatry and Allied Sciences, Kanke, Ranchi, Jharkhand - 834 006, India E-mail: rudrapc@yahoo.com

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