

Recommendations on the Drug Treatment of Psychosis in Parkinson's Disease

To the Editor:

In an otherwise excellent and balanced summary of the problem of psychosis in parkinsonian patients, Hasnain et al¹ make the unfortunate recommendation of equating quetiapine and risperidone for the treatment of psychotic symptoms in non-demented patients with Parkinson's disease. They also fail to note that in the trials of clozapine for treating Parkinson's disease psychosis, demented patients were included, and in one subgroup analysis there were no differences in efficacy between demented and non-demented subjects with Parkinson's disease.²

Although the authors are clearly correct in noting that the only double-blind, placebo-controlled trials of quetiapine for the treatment of Parkinson's disease psychosis showed no benefit,³⁻⁵ these trials also concluded that the drug caused no worsening of motor symptoms. All open-label trials of quetiapine have been positive,⁶ whereas risperidone's motor side effects have often been extreme.^{7,8} It is true that the published reports on risperidone have been mixed in terms of the motor side effects,⁶ but The American Academy of Neurology's task force on practice parameters in the treatment of Parkinson's disease published its recommendations in 2006⁹ recommending quetiapine as the drug of first choice for treating psychosis in Parkinson's disease, although noting that there are no level 1 data to support its

Funding: Over the past 12 months the author has received money for research, lectures or consulting from: Teva, Ingelheim-Boehringer, Glaxo-SmithKline, Cephalon, EMD Serono, Acadia, Schering Plough, and Novartis. Since 2001 the author has received money from: Astra Zeneca, Eli Lilly, Janssen, Bristol Myers Squibb, and Pfizer (all are manufacturers of the atypical antipsychotics).

Conflict of Interest: I have received funds for lectures, clinical research, or consultations over the last 12 months from Astra Zeneca, Ingelheim Boehringer, GlaxoSmithKline, Acadia, EMD Serono, Cephalon, Teva, Valeant, and Pfizer.

use, with clozapine as the second-line choice. I think it highly unlikely that any Parkinson's disease specialist would recommend risperidone until quetiapine, clozapine, aripiprazole, and cholinesterase inhibitors had failed.

We suggest that risperidone, which causes all the extrapyramidal side effects of the first generation of antipsychotics, be removed as a recommended treatment for patients with Parkinson's disease psychosis.

Joseph H. Friedman, MD

*Parkinson's Disease and Movement Disorders Center
 Warwick, RI*

doi:10.1016/j.amjmed.2009.09.027

References

1. Hasnain M, Vieweg WVR, Baron MS, et al. Pharmacological management of psychosis in elderly patients with Parkinsonism. *Am J Med.* 2009;122:614-622.
2. Parkinson Study Group. Low dose clozapine for the treatment of drug-induced psychosis in Parkinson's disease. *N Engl J Med.* 1999;340:757-763.
3. Ondo WG, Tintner R, Young KD, et al. Double blind placebo controlled unforced titration parallel trial of quetiapine for dopamine-induced hallucinations in Parkinson's disease. *Mov Disord.* 2005;20:958-963.
4. Rabey JM, Prokhorov T, Minovitz A, et al. Effect of quetiapine in psychotic Parkinson disease patients: a double blind labeled study of 3 months duration. *Mov Disord.* 2007;22:313-318.
5. Shotbolt P, Samuel M, Fox C, David AS. A randomized controlled trial of quetiapine for psychosis in Parkinson's disease. *Neuropsychiatr Dis Treatment.* 2009;5:327-332.
6. Chou KL, Borek LL, Friedman JH. Management of psychosis in movement disorder patients. *Exp Opin Pharmacother.* 2007;8:935-943.
7. Ford B, Lynch T, Greene P. Risperidone in Parkinson's disease. *Lancet.* 1994;344:681.
8. Rich SS, Friedman JH, Ott BR. Risperidone versus clozapine in the treatment of psychosis in six patients with Parkinson's disease and other akinetic-rigid syndromes. *J Clin Psychiatry.* 1995;56:556-559.
9. Miyasaki J, Shannon K, Voon V, et al. Practice Parameter: evaluation and treatment of depression, psychosis and dementia in Parkinson disease (an evidenced-based review): report of the Quality Standards Subcommittee of the American Academy of Neurology. *Neurology.* 2006;66:996-1002.