may be quite useful.

When motor restlessness or akathisia is prominent, propranolol 10 to 20 mg three times daily is helpful.

Use of antiepileptics. Finally, if the neuroleptic or atypical neuroleptic agents are ineffective or are accompanied by intolerable side effects, antiepileptics such as carba-mazepine or valproic acid may be very useful for agitation and psychosis. Carbamazepine is started at small doses of 100 mg twice daily and is gradually titrated to 200 mg three times daily, while valproic acid is begun at doses of 125 mg twice daily and is gradually and is gradually titrated to 500 mg three times daily as tolerated, or higher if necessary.

Sleep disturbances in Alzheimer disease are exceedingly common and may contribute to the other behavioral disorders. Traditional sedating medications such as benzodiazepines may increase confusion and daytime drowsiness. A better alternative would be trazodone 25 to 100 mg at bedtime or thioridazine 25 to 75 mg at night.

## REFERENCES

- Mulnard RA, Cotman CW, Kawas C, et al. Estrogen replacement therapy for treatment of mild to moderate Alzheimer disease: a randomized controlled trial. JAMA 2000; 283:1007–1015.
- Knopman D, Schneider L, Davis K, et al. Long-term tacrine treatment: Effects on nursing home placement and mortality. The Tacrine Study Group. Neurology 1996; 47:166–177.
- Rogers SC, Friedhoff LT. The efficacy and safety of donepezil in patients with Alheimer's disease: Results of a US multicenter randomized, double-blind, controlled trial. The Donepezil Study Group. Dementia 1996; 7:293–303.
- Rogers SC, Farlow MR, Doody RS, Molis R, Friedhoff LT. Donepezil improves cognition and global function in Alzheimer disease: A 15-week, double-blind, placebo trial study. Donepezil Study Group. Arch Intern Med 1998; 158:1021–1030.
- Samuels SC, Davis DL. Dementia and delirium. In: Enna ST, Coyle JT, editors. Pharmacological management of neurological and psychiatric disorders. New York: McGraw-Hill, 1998: 274–279.
- Tierney MC, Szalai JP, Snow WG, et al. Prediction of probable Alzheimer's disease in memory-impaired patients. Neurology 1996; 46:661–665.
- Corey-Bloom J, Arnand R, Veach J, et al. A randomized trial evaluating the efficacy of ENA-713 (rivastigmine tartrate), a new acetylcholinesterase inhibitor, in patients with mild to moderate severe Alzheimer's disease. International Journal of Geriatric Psychopharmacology 1998; 1:55–65.
- 8. Mulnard RA, Cotman CW, Kawas C, et al. Estrogen

replacement therapy for treatment of mild to moderate Alzheimer disease. JAMA 2000; 283:1007-1015.

- McGreer PL, Schulzer M, McGreer EG. Arthritis and anti-inflammatory agents as possible protective factors for Alzheimer's disease: A review of 17 epidemiological studies. Neurology 1996; 47:425–432.
- Rogers J, Kirby LC, Hempelman SR, et al. Clinical trial of indomethacin in Alzheimer's disease. Neurology 1993; 43;1609–1611.
- Yan SD, Chen X, Fu J, et al. RAGE and amyloid—peptide neurotoxicity in Alzheimer's disease. Nature 1996; 382:685–691.
- Sano M, Ernesto C, Thomas RG, et al. A controlled trial of selegiline and alpha-tocopherol or both as treatment for Alzheimer's disease. The Alzheimer's Disease Cooperative Study. N Engl J Med 1997; 278:1327–1332.
- Tariot PN, Blazina L. The psychopathology of dementia. In: Morris J, editor. Handbook of dementing illnesses. New York: Marcel Decker, 1993: 461–475.
- Tariot PN. General approaches to behavioral disturbances. In: Reichman ME, Katz P, editors. Psychiatric care in the nursing home. New York: Oxford University Press, 1996: 10–22.
- Tariot PN. Treatment for agitation and psychosis in dementia. J Clin Psychiatr 1996; 57 Suppl 14:21–29.

**ADDRESS:** Maurice R. Hanson, MD, Department of Neurology, Cleveland Clinic Florida, 3000 West Cyprus Creek Road, Fort Lauderdale, FL 33309.

## CORRECTION

## New therapies for allergic rhinitis

In the article "New therapies for allergic rhinitis" by David F. Graft, MD in the March 2000 issue (*Cleve Clin J Med* 2000; 67:165–168), the first paragraph in the section on newer second-generation antihistamines (page 166) did not list all of the available agents, and also indicated that all of the newer agents have no sedating effect. The corrected paragraph should read as follows:

Newer second-generation antihistamines have fewer side effects and are safe. Acrivastine (the antihistamine ingredient in Semprex-D), azelastine, (Astelin), cetirizine (Zyrtec), fexofenadine (Allegra), and loratadine (Claritin) are as effective as older antihistamines. Acrivastine, azelastine and cetirizine have some potential for causing sedation, but much less than with older agents, and fexofenadine and loratadine do not appear to cause sedation at all. All of the newer agents have very little anticholinergic activity, and thus have very low rates of the other side effects.