Editors' Note: In WriteClick this week, the authors of "Evidence-based guideline: Treatment of tardive syndromes," Bhidayasiri et al., address the concerns of Lerner et al. regarding inclusion of articles in evidence-based guideline analysis and further clarify the American Academy of Neurology's classification scheme for rating therapeutic articles. Brenner and author DiMauro discuss mitochondrial DNA dysfunction in neurodegenerative disorders without typical mendelian inheritance patterns.

—Megan Alcauskas, MD, and Robert C. Griggs, MD

EVIDENCE-BASED GUIDELINE: TREATMENT OF TARDIVE SYNDROMES: REPORT OF THE GUIDELINE DEVELOPMENT SUBCOMMITTEE OF THE AMERICAN ACADEMY OF NEUROLOGY

Vladimir Lerner, Chanoch Miodownik, Be'er Sheva, Israel: We read the article by Bhidayasiri et al.¹ with interest. We found some inaccuracies. The authors cited only one of our articles regarding treatment of tardive dyskinesia (TD).² Even though the authors performed a search from 1966 to 2011, they did not include 2 studies published in 2007 that could broaden the knowledge about new options of TD management.^{3,4} The first deals with vitamin B₆ and the other with piracetam. Both studies include greater samples (50 and 40 subjects, respectively) and could be considered Class I evidence according to the authors' classification. Inclusion of this information could positively influence the weight and emphasize the significance of these medications. Our experience shows that different types of TD react uniquely to different types of medications.

Author Response: Roongroj Bhidayasiri, Bangkok, Thailand; Stanley Fahn, New York; Gary S. Gronseth, Kansas City, KS; Kelly L. Sullivan, Theresa A. Zesiewicz, Tampa, FL: We thank Lerner et al. for their comments and interest in our article.¹ We agree that various forms of tardive syndromes (TDS) can respond differently to medications or interventions. Well-designed randomized controlled trials with specific TDS inclusion criteria/subtypes are necessary to determine the most effective interventions for TDS symptoms. As Lerner et al. noted, 2 studies on vitamin B₆ and piracetam were not included in our original analysis.^{3,4} The first study, a double-blind, placebo-controlled

trial on vitamin B₆ treatment in 50 inpatients with schizophrenia/schizoaffective disorders, was rated Class III for no allocation concealment and a >20% (14/50) dropout rate.³ The second study, involving 40 patients with schizophrenic/schizoaffective disorders who received piracetam or placebo over 4 weeks, was rated Class III for dropout rate.⁴ After applying the AAN's classification scheme for rating therapeutic articles,⁵ the data are still insufficient to support or refute use of vitamin B₆ and piracetam for TDS. Although the AAN endeavors to find all pertinent literature,⁵ no literature search is 100% effective, and we thank the authors for alerting us to these 2 studies. We corrected errors for an article cited in table e-1 and this table is updated online: <http://www.neurology.org/content/81/5/463/suppl/DC1>.¹

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1. Bhidayasiri R, Fahn S, Weiner WJ, Gronseth GS, Sullivan KL, Zesiewicz TA. Evidence-based guideline: treatment of tardive syndromes: report of the Guideline Development Subcommittee of the American Academy of Neurology. *Neurology* 2013;81:463–469.
2. Lerner V, Miodownik C, Kaptsan A, et al. Vitamin B6 in the treatment of tardive dyskinesia: a double-blind, placebo-controlled, crossover study. *Am J Psychiatry* 2001;158:1511–1514.
3. Lerner V, Miodownik C, Kaptsan A, et al. Vitamin B6 treatment for tardive dyskinesia: a randomized, double-blind, placebo-controlled, crossover study. *J Clin Psychiatry* 2007;68:1648–1654.
4. Libov I, Miodownik C, Bersudsky Y, Dwolatzky T, Lerner V. Efficacy of piracetam in the treatment of tardive dyskinesia in schizophrenic patients: a randomized, double-blind, placebo-controlled crossover study. *J Clin Psychiatry* 2007;68:1031–1037.
5. Gronseth GS, Woodroffe LM, Getchius TSD. Clinical Practice Guideline Process Manual. St. Paul, MN: The American Academy of Neurology; 2011.

MITOCHONDRIAL ENCEPHALOMYOPATHIES—FIFTY YEARS ON: THE ROBERT WARTENBERG LECTURE

Steven R. Brenner, St. Louis: I read the article on mitochondrial diseases with interest.¹ Multiple sclerosis,² Alzheimer disease,³ and Parkinson disease⁴ have an epidemiologic maternal effect. However, the cause is still unclear except in a minority of familial cases with mendelian inheritance. Perhaps the

maternal bias in inheritance is an indication of abnormal mitochondrial DNA (mtDNA). It is possible that the variations in age at onset are related to the degree of heteroplasmy of the mitochondrial genome, as has been seen in mitochondrial inheritance in diabetes and deafness.⁵

Author Response: Salvatore DiMauro, New York: I thank Dr. Brenner for his comments. The possible involvement of mtDNA in neurodegenerative diseases without typical mendelian inheritance has attracted the interest of neuroscientists since the discovery of pathogenic mtDNA mutations. Mitochondrial dysfunction clearly plays a role in the pathogenesis of these disorders,^{6,7} yet there is no evidence of any underlying mtDNA mutations. Even epidemiologic evidence of maternal inheritance is weak and often anecdotal, as in the article cited by Dr. Brenner.⁴ Dr. Brenner also cited an article on the syndrome of diabetes and deafness.⁵ This syndrome is a bona fide mtDNA-related disease and the effect of heteroplasmy on the phenotype is to be expected. To be fair, there is increasing evidence that mtDNA mutations may be related to neurodegenerative diseases, aging, and even cancer, but

these are somatic, de novo mutations not related to maternal inheritance.⁸

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1. DiMauro S. Mitochondrial encephalomyopathies—Fifty years on: the Robert Wartenberg Lecture. *Neurology* 2013;81:281–291.
2. Hoppenbrouwers IA, Loi F, Aulchenko YS, et al. Maternal transmission of multiple sclerosis in a Dutch population. *Arch Neurol* 2008;65:345–348.
3. Honea RA, Vidoni ED, Swerdlow RH, Burns JM; Alzheimer Disease Neuroimaging Initiative. Maternal family history is associated with Alzheimer's disease biomarkers. *J Alzheimers Dis* 2012;31:659–668.
4. Wooten GF, Currie LJ, Bennett JP, Harrison MB, Trugman JM, Parker WD Jr. Maternal inheritance in Parkinson's disease. *Ann Neurol* 1997; 41; 265–268.
5. Olsson C, Zethelius B, Lagerstrom-Fermer M, Asplund J, Berne C, Landegren U. Level of heteroplasmy for the mitochondrial mutation A3243G correlates with age at onset of diabetes and deafness. *Hum Mutat* 1998;12:52–58.
6. DiMauro S, Schon EA. Mitochondrial disorders in the nervous system. *Annu Rev Neurosci* 2008;31:91–123.
7. Schon EA, Przedborski S. Mitochondria in the next (neuro)generation. *Neuron* 2011;70:1033–1053.
8. Schon EA, DiMauro S, Hirano K-I. Human mitochondrial DNA: roles of inherited and somatic mutations. *Nat Rev Genet* 2012;13:878–890.

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Mitochondrial encephalomyopathies—Fifty years on: The Robert Wartenberg Lecture

Steven R. Brenner and Salvatore DiMauro

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