

HHS Public Access

Author manuscript

Am J Psychiatry. Author manuscript; available in PMC 2020 March 15.

Published in final edited form as:

Am J Psychiatry. 2017 April 01; 174(4): 397–398. doi:10.1176/appi.ajp.2017.16111322r.

A Step Toward Optimizing Treatment Schedules for Continuation ECT: Response to Rasmussen

Charles H. Kellner, M.D., Rebecca G. Knapp, Ph.D., Georgios Petrides, M.D., W. Vaughn McCall, M.D., M.S., Robert C. Young, M.D., Mustafa M. Husain, M.D., Sarah H. Lisanby, M.D. CORE/PRIDE Work Group

Department of Psychiatry, Icahn School of Medicine at Mount Sinai, New York; the Department of Psychiatry, Columbia University and New York State Psychiatric Institute, New York; the Department of Public Health Sciences, College of Medicine, Medical University of South Carolina, Charleston; the Department of Psychiatry, Zucker Hillside Hospital/Northwell Health System, New York; the Department of Psychiatry, University of Texas Southwestern Medical Center, Dallas; the Department of Psychiatry, New York Presbyterian/Weill Cornell Medical Center, New York and White Plains; the Department of Psychiatry and Health Behavior, Augusta University/Medical College of Georgia, Augusta; and the Division of Services and Intervention Research, NIMH, Bethesda, Md.

TO THE EDITOR: We thank Dr. Rasmussen for his careful reading of our study and his speculations as to the reasons the data turned out as they did. Our a priori hypothesis was that additional ECT after the acute course would be beneficial in the ensuing 6-month period; our preplanned data analysis was a comparison of symptom severity at the primary study endpoint of 24 weeks, with power to control for a false positive rate of 5% or less. Our data showed significant difference at the primary time point, as well as a consistent benefit at all other time points. We believe that we were cautious in interpreting these results and that various biological explanations could be invoked to explain them. One-third of the patients in the Symptom-Titrated, Algorithm-Based Longitudinal ECT (STABLE) treatment arm did, in fact, receive additional ECT past the 4-week time point, and even for those who did not, there is no a priori reason to dismiss the possibility of ongoing biological benefit from tapering, rather than abruptly stopping, the acute course of ECT for the treatment of an episode of depression. We did not advocate that STABLE be considered "the standard of care for continuation ECT" at this point because considerable additional research is needed to establish optimal treatment schedules. Our data stand, however, as the accurately reported and conservatively interpreted results of a randomized clinical trial, and we disagree with characterizing our results as "inexplicable" or "fortuitous."