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## A Randomized Controlled Trial of the Effect of Hostility Reduction on Cardiac Autonomic Regulation

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**Objective:** To test whether reduction in hostility increases autonomic regulation of the heart.

**Methods:** In this randomized controlled trial, participants were 158 healthy adults, 20 to 45 years of age, who were one standard deviation (SD) above national norms on the Cook-Medley Hostility Scale and the Spielberger Trait Anger Index. Participants also were interviewed using the Interpersonal Hostility Assessment Technique (IHAT). They were randomly assigned to a 12-week cognitive behavior therapy (CBT) program for hostility reduction or a wait-list control condition. The main outcome measure was cardiac autonomic modulation, measured as

RR interval variability (RRV) derived from 24-ECG recordings.

**Results:** In a MANOVA assessing psychological outcomes of hostility, anger, and IHAT scores, there was a significant treatment effect with an average reduction across the three outcomes that was approximately 0.7 SD (ES = 0.685, se = 0.184, P < 0.001) greater for the intervention group than for the control group.

In contrast, the change in HR was -0.14 bpm (95% CI, -2.43 to 2.14) in treatment participants and -1.36 bpm (95% CI, -3.28 to 0.61) in wait-list participants. HF RRV, an index of cardiac parasympathetic modulation, increased by 0.07 ln msec<sup>2</sup> (95% CI, -0.10 to 0.24) for participants in the treatment condition and decreased by 0.04 ln msec<sup>2</sup> (95% CI, -0.18 to 0.10) for participants in the wait-list condition. These differences were not significant. The findings for other indices of RRV were similar.

**Conclusions:** Reduction of hostility and anger was not accompanied by increases in cardiac autonomic modulation. These findings raise questions about the status of disordered ANS regulation of the heart as a pathophysiological mechanism underlying the hostility–heart disease relationship and about whether hostility itself is a mechanism or merely a marker of elevated risk of heart disease.