

499.2

THE CONTRIBUTION OF AFTERHYPERPOLARIZATION CURRENTS TO CHOLINERGIC ENHANCEMENT OF EXCITABILITY AND PRESERVATION OF SPIKE TIMING IN A NEOCORTICAL NEURON MODEL.
A.M. Bartels, M.J. McKeown, A.C. Tang, T.J. Sejnowski*

Computational Neurobiology Lab., Salk Institute for Biological Studies, La Jolla, CA 92037-1099

Recent physiological studies have shown that the effects of cholinergic modulation can be different when physiologically more realistic fluctuating rather than constant depolarizing currents are injected into a neocortical neuron (see companion poster, Tang-Sejnowski, 1996). In a modified version of a 10 compartment neocortical neuron model (Bush-Sejnowski, 93), we show that the effect of cholinergic modulation is a stimulus-dependent simultaneous preservation of spike timing and enhancement of neuronal excitability. We focus on the hypothesis that although playing a dominant role in cholinergic enhancement of excitability under constant depolarizing input I_{AHP} is less important in the modulation of the firing rate under the physiologically more realistic fluctuating inputs. We first constrained the model to reproduce the phenomenon of spike frequency adaptation (SFA) and a reduction in SFA due to cholinergic modulation (modeled as a decrease in maximum conductance, g_{KCa} .) Next, by varying the fluctuation amplitude and mean intensity of the current injection, we compute a spike count-based index, D , for the contribution of I_{AHP} to enhanced excitability. We found that D can decrease as the amplitude of current fluctuation increases. This suggests that the contribution of I_{AHP} to cholinergic enhancement of neuronal excitability will be more prominent for physiologically unrealistic stimuli than that observed under *in vivo* conditions.

Research supported by the Howard Hughes Medical Institute.