Electrochemistry of Cytochrome *c* Mutants at Functionalized Self-Assembled Monolayers.

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Electrochemistry of cytochrome c (cyt. c) bound to carboxylate-terminated alkanethiol SAMs provides a simple molecular assembly for investigating both long-range electron transfer (ET) and intermolecular protein interactions [1-3]. ET gating has been observed at short alkanethiol-modified surfaces, and has been attributed to dynamic interactions between the charged protein and the carboxylate terminus of the SAMs [3]. We have developed a theoretical model for ET in these systems, and have proposed that electrostatic interactions between lysine-13 and the negatively charged monolayer provide an efficient pathway for electron transfer [3]. Our present study involves determining the dynamics of ET between functionalized alkanethiol SAMs and a series of cyt. c mutants in which lysine-13 has been replaced by alanine; these data should provide a more detailed picture of the protein/SAM interface, as well as its effect on the ET pathway.

References:

- 1) Tarlov, M.J.; Bowden, E.F. J. Am. Chem. Soc., **1991**, 113, 1847
- Feng, Z-Q.; Imabayashi, S.,;Kakiuchi, T.; Niki, K. J. Chem. Soc., Faraday Trans., 1997, 93, 1367.
- Avila, A.; Gregory, B.W.; Niki, K.; Cotton, T.M. J. Phys. Chem. B, 2000, 104, 2759.