

**Colorado State University
Department of Chemistry**

**CHEM 431
Instrumental Analysis Laboratory**

**Notes for
Electrochemistry - Voltammetry**

The following is a set of short notes to outline the experiment in question and to provide helpful guidance to those executing the experiment.

- 1. Voltammetry is the method of following reduction-oxidation reactions through the imposition of an electrical potential (a voltage) across an electrochemical cell and measuring the current (an electron flow) passing through the cell.**
- 2. Study up on voltammetry and cyclic voltammetry from references. At each major step in the experimental work collect a voltammogram for inclusion in a laboratory report.**
- 3. Configure the potentiostat using a dummy resistor set to simulate an electrochemical cell. Learn how to collect a voltammogram (including the exportation of experimental data and import into Igor). Use +1.0 and -1.0 V for the endpoints and 200 mV s⁻¹ for the sweep rate.**
- 4. Prepare your own silver-silver chloride reference electrode with an agar-salt (KCl) bridge. Use a Luggin-style construction.**
- 5. Configure an electrochemical cell with the platinum working, platinum counter and the reference electrodes using approximately (but known exactly) 10 mM FeCl₃ in approximately 1 M H₂SO₄ as the electrolyte. Learn how to select appropriate potentials for the endpoints.**
- 6. See if you can successfully and convincingly observe voltammograms using lower concentrations of FeCl₃. It might be necessary to sparge the cell with nitrogen to reduce the background reactions.**
- 7. Collect a comprehensive set of experimental data in support of the Nernst equation using voltammetry measurements on the quinone-hydroquinone redox couple in buffered aqueous solution according to reference J. Chem. Ed. 1997, 74 1195-1197. Use your platinum working electrode instead of the gold one described in the reference (adjust the endpoint potentials accordingly). Be clever in preparing the necessary solutions.**