

CHEM 116

Redox Reactions and Electrochemistry

November 30, 2006
Prof. Sevian



Note: Start with the lecture slides from Tuesday that we didn't finish

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Agenda

- Review oxidation states (from first semester course)
- Balancing redox reactions
 - Inspection method
 - SOHe method
- Redox reactions that occur spontaneously
 - Predicting whether a reaction will be spontaneous
 - How to separate the halves of a reaction to generate electricity
 - How much electricity?

The **final exam** is scheduled for Monday, December 18, 11:30AM-2:30PM. Location TBA.

Electrochemistry overview

- What you need to know (mostly boils down to vocabulary)
 - How to recognize an electron transfer (redox) reaction
 - Oxidation vs. reduction
 - Loss of electrons is oxidation, gain of electrons is reduction (LEO the lion says GER)
 - Reduction is moving (oxidation number) down on a number line, oxidation is up
 - Agent means "that which causes"
- What you need to be able to do
 - Assign oxidation numbers to elements to figure out which **species** contain elements that are changing
 - Balance redox rxns in neutral, acid, and base solutions
 - Diagram an electrochemical cell and indicate direction of flow of negative charge through the system
 - Read and write electrochemical cell notation
 - Use reduction potentials to predict cell voltage under standard conditions
 - Use the Nernst equation to predict cell voltage at non-standard conditions

Redox Reactions and Electrochemistry

Part I: Foundational

- Assigning oxidation numbers
- Balancing redox equations

Part II: Applied

- Diagramming electrochemical cells
- Facility with cell notation

Part III: Making mathematical predictions of properties

- Calculating cell potential at standard conditions
- Calculating cell potential at non-standard conditions
- Relating the amount of a substance oxidized or reduced to the electrical current and the time during which current is allowed to flow

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Some vocabulary that applies to redox reactions

- **Species** = a reactant that contains an element whose oxidation number changes
 - On the reactant side, there must be one species that gets oxidized (oxidation number of an element in it goes up) **and** one species that gets reduced (oxidation number of an element in it goes down)
- **Oxidation half-reaction**: the reaction showing the species that gets oxidized, its product, and how many electrons get produced
- **Reduction half-reaction**: the reaction showing the species that gets reduced, its product, and how many electrons react in order for this to happen
- Species that gets oxidized is called the **reducing agent** because it causes the reduction half-reaction to occur
- Species that gets reduced is called the **oxidizing agent** because it causes the oxidation half-reaction to occur

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Electrochemical cells

- Electrochemical, voltaic, galvanic = all the same thing
- Divide the half-reactions into separate cells (locations) so that electrons generated at the oxidation half-cell are forced to travel through a wire to get to the reduction half-cell
- <http://www.chem.iastate.edu/group/Greenbowe/sections/projectfolder/flashfiles/electroChem/voltaicCellEMF.html>
- What's the role of the salt bridge?
- Why do reactions occur in only one direction?
- How can you predict the voltage?

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Key ideas so far about electrochem

- You must review ions and ionic charges
- You must be able to assign oxidation numbers to elements in a compound or ion
- Redox reactions are when oxidation numbers change during a reaction
 - One element (part or all of a "species") has its oxidation number increase (this is oxidation)
 - Another element (part or all of a "species") has its oxidation number decrease (this is reduction)
 - Both processes must occur, because redox is electron transfer
- Study the vocabulary

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Parts of an electrochemical cell

- Anode = oxidation half-cell
- Cathode = reduction half-cell
- Wire
- Salt bridge

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Predicting voltage under standard conditions

- Voltage = cell potential (E°_{cell}) = related to ΔH
- Table of reference voltages, all referenced to the hydrogen half-cell
- All reference voltages are reduction potentials, so if you need an oxidation potential you just take the opposite
- See table of standard reduction potentials
- Examples

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Examples

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Non-standard conditions: Nernst equation

- Concentrations of solutions not at standard 1.0 M

$$E_{cell} = E_{cell}^{\circ} - \frac{0.0257}{n} \ln Q$$

$$= E_{cell}^{\circ} - \frac{0.0592}{n} \log Q$$

Reaction quotient

- Temperature not at standard 25 °C

$$E_{cell} = E_{cell}^{\circ} - \frac{RT}{nF} \ln Q$$

Least common multiple of electrons exchanged in the redox reaction

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Electrolysis example

Similar to Sample Exercise 20.14, pp. 885-886

Calculate the mass of O₂ produced in the electrolysis of water, using a current of 0.445 A for a period of 45 minutes.

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Electrolysis: electrochemical cell forced to run backwards (against its will)

- Note: sections 20.7 and 20.8 in the text book are fascinating reading, but they are not required reading
- Electroplating:** The longer you run the cell, the more metal electroplate builds up
- Current means how many electrons pass by per second
- If you know how much metal electroplate you want to make, then stoichiometry tells you how many electrons are required
 - Cu²⁺ (aq) + 2 e⁻ → Cu (s)
 - Au³⁺ (aq) + 3 e⁻ → Au (s)
 - Ag⁺ (aq) + 1 e⁻ → Ag (s)
- If you know what current is being applied (e.g., 0.800 amperes = 0.800 Coulombs/second), then you can figure out how much time you must run the cell for in order to build the amount of electroplated metal that you want