CHAPTER 17:
CORROSION AND DEGRADATION

ISSUES TO ADDRESS...

- Why does corrosion occur?
- What metals are most likely to corrode?
- How do temperature and environment affect corrosion rate?
- How do we suppress corrosion?

THE COST OF CORROSION

- Corrosion:
  -- the destructive electrochemical attack of a material.
  -- Al Capone's ship, Sapona, off the coast of Bimini.

Photos courtesy L.M. Maestas, Sandia National Labs. Used with permission.

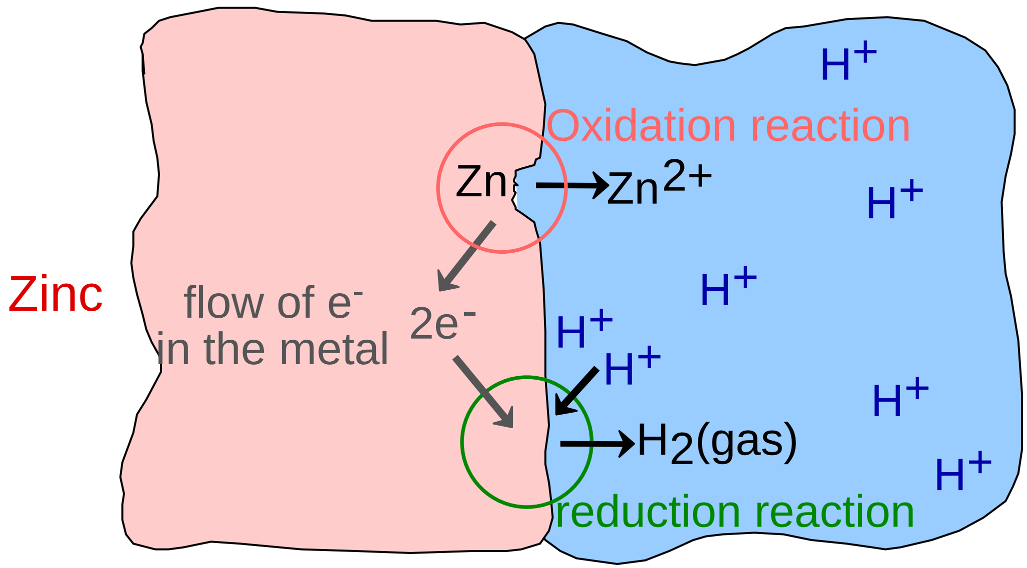
- Cost:
  -- 4 to 5% of the Gross National Product (GNP)*
  -- this amounts to just over $400 billion/yr**

* H.H. Uhlig and W.R. Revie, *Corrosion and Corrosion Control: An Introduction to Corrosion Science and Engineering*, 3rd ed., John Wiley and Sons, Inc., 1985.

**Economic Report of the President (1998).

CORROSION OF ZINC IN ACID

- Two reactions are necessary:
  -- oxidation reaction: $Zn \rightarrow Zn^{2+} + 2e^-$
  -- reduction reaction: $2H^+ + 2e^- \rightarrow H_2(gas)$



Adapted from Fig. 17.1, *Callister 7e*. (Fig. 17.1 is from M.G. Fontana, *Corrosion Engineering*, 3rd ed., McGraw-Hill Book Company, 1986.)

- Other reduction reactions:
  -- in an acid solution

$$O_2 + 4H^+ + 4e^- \rightarrow 2H_2O$$

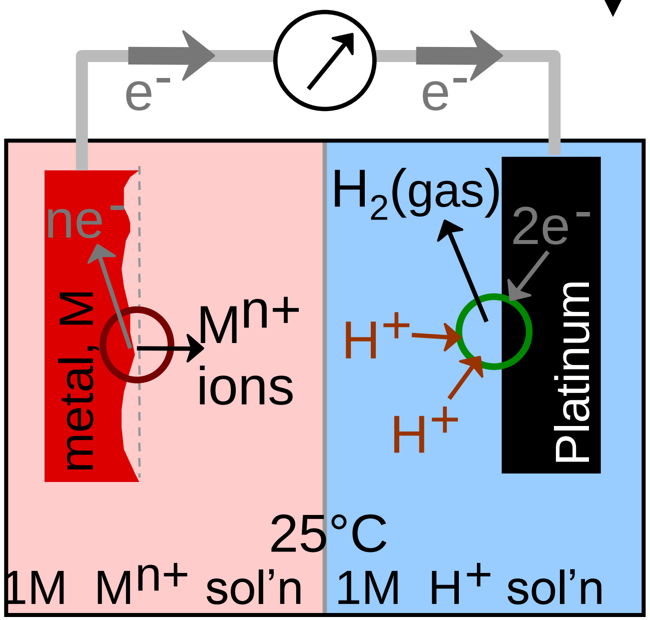
  -- in a neutral or base solution

$$O_2 + 2H_2O + 4e^- \rightarrow 4(OH)^-$$

STANDARD HYDROGEN (EMF) TEST

- Two outcomes:

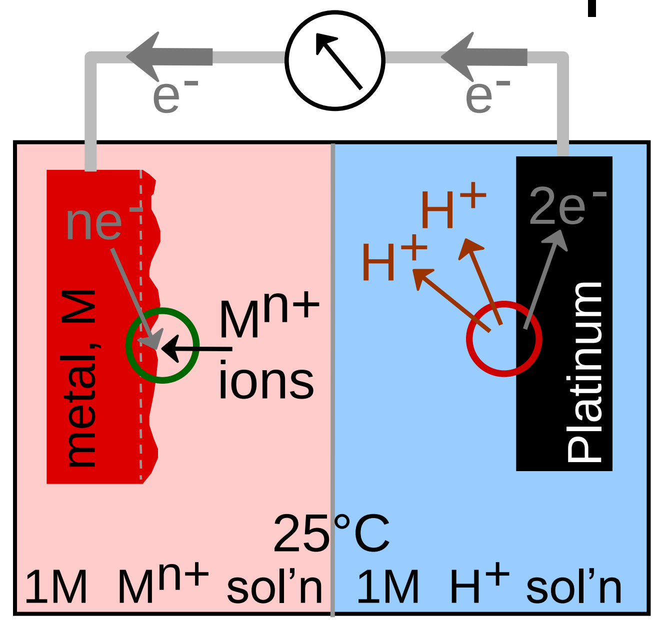
--Metal sample mass ↓



--Metal is the anode (-)

$V^o_{metal} < 0$ (relative to Pt)

--Metal sample mass ↑



--Metal is the cathode (+)

$V^o_{metal} > 0$ (relative to Pt)

Standard Electrode Potential

Adapted from Fig. 17.2, *Callister 7e*.

STANDARD EMF SERIES

- EMF series

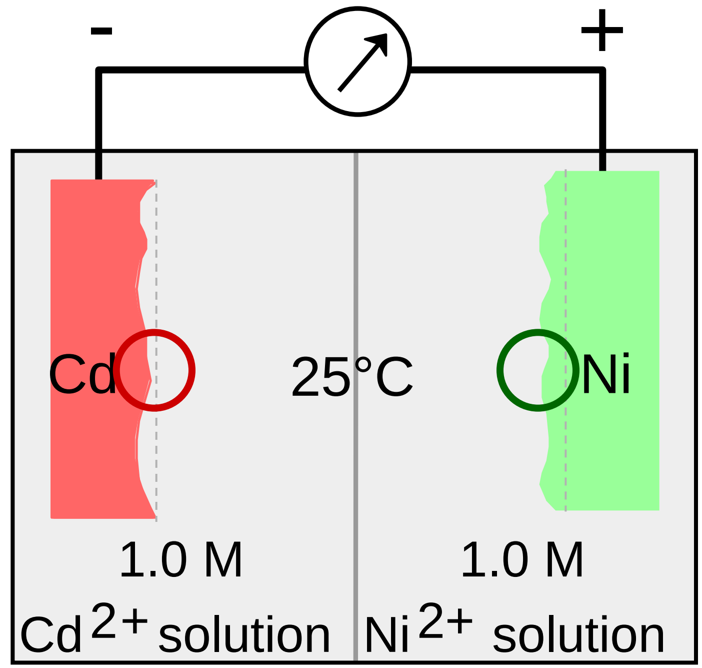
more cathodic ↑ / more anodic ↓

metal	V^o_{metal}
Au	+1.420 V
Cu	+0.340
Pb	- 0.126
Sn	- 0.136
Ni	- 0.250
Co	- 0.277
Cd	- 0.403
Fe	- 0.440
Cr	- 0.744
Zn	- 0.763
Al	- 1.662
Mg	- 2.363
Na	- 2.714
K	- 2.924

$\Delta V^o =$ 0.153V (between Ni and Cd)

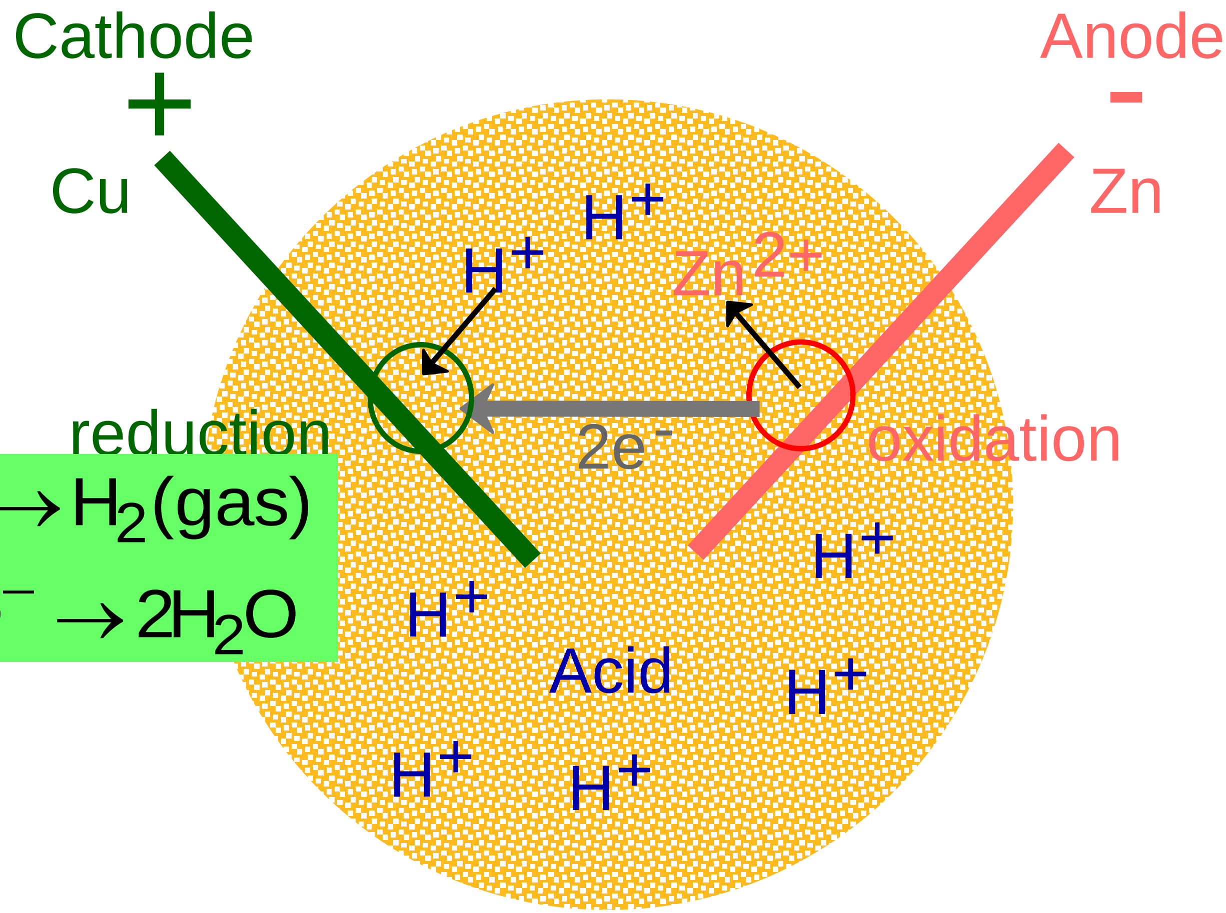
Data based on Table 17.1, *Callister 7e*.

- Metal with smaller V^o_{metal} corrodes.
- Ex: Cd-Ni cell



Adapted from Fig. 17.2, *Callister 7e*.

CORROSION IN A GRAPEFRUIT



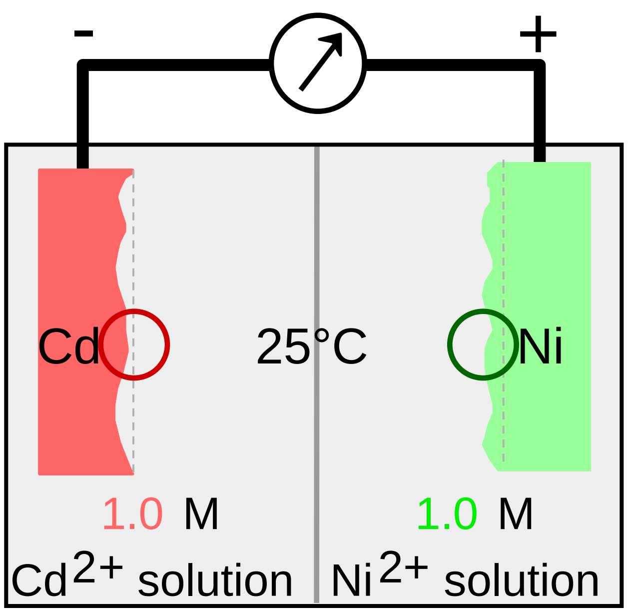
$$2H^+ + 2e^- \rightarrow H_2(\text{gas})$$

$$O_2 + 4H^+ + 4e^- \rightarrow 2H_2O$$

EFFECT OF SOLUTION CONCENTRATION

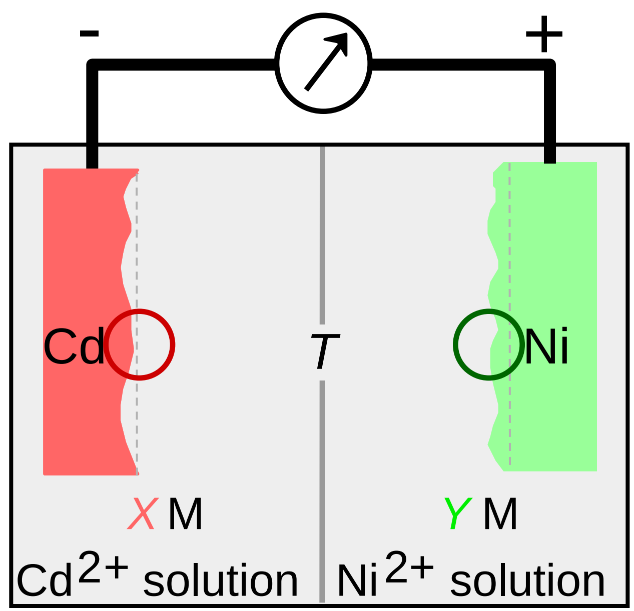
- Ex: Cd-Ni cell with standard 1 M solutions

$$V^{o}_{Ni} - V^{o}_{Cd} = 0.153$$



- Ex: Cd-Ni cell with non-standard solutions

$$V_{Ni} - V_{Cd} = V^{o}_{Ni} - V^{o}_{Cd} - \frac{RT}{nF} ln \frac{X}{Y}$$



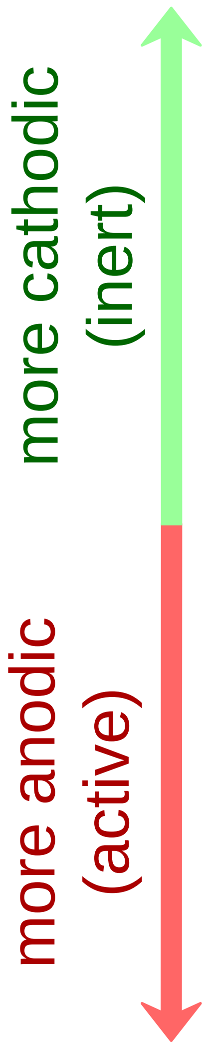
n = #e$^-$ per unit oxid/red reaction (= 2 here)

F = Faraday's constant = 96,500 C/mol.

- Reduce V_{Ni} - V_{Cd} by
  --increasing X
  --decreasing Y

GALVANIC SERIES

- Ranks the reactivity of metals/alloys in seawater



Platinum
Gold
Graphite
Titanium
Silver
316 Stainless Steel
Nickel (passive)
Copper
Nickel (active)
Tin
Lead
316 Stainless Steel
Iron/Steel
Aluminum Alloys
Cadmium
Zinc
Magnesium

Based on Table 17.2, *Callister 7e*. (Source of Table 17.2 is M.G. Fontana, *Corrosion Engineering*, 3rd ed., McGraw-Hill Book Company, 1986.)

FORMS OF CORROSION

- Stress corrosion

Stress & corrosion work together at crack tips.

- Uniform Attack

Oxidation & reduction occur uniformly over surface.

- Erosion-corrosion

Break down of passivating layer by erosion (pipe elbows).

Forms of corrosion

- Selective Leaching

Preferred corrosion of one element/constituent (e.g., Zn from brass (Cu-Zn)).

- Pitting

Downward propagation of small pits & holes.

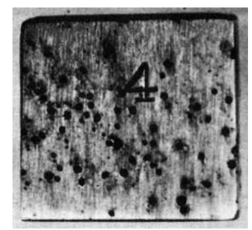


Fig. 17.17, *Callister 7e*. (Fig. 17.17 from M.G. Fontana, *Corrosion Engineering*, 3rd ed., McGraw-Hill Book Company, 1986.)

- Intergranular

Corrosion along grain boundaries, often where special phases exist.

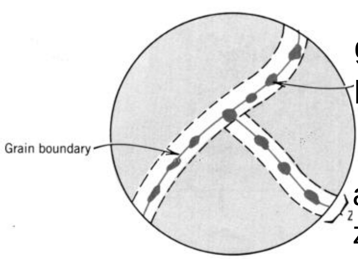


Fig. 17.18, *Callister 7e*.

- Galvanic

Dissimilar metals are physically joined. The more anodic one corrodes.(see Table 17.2) Zn & Mg very anodic.

- Crevice Between two pieces of the same metal.

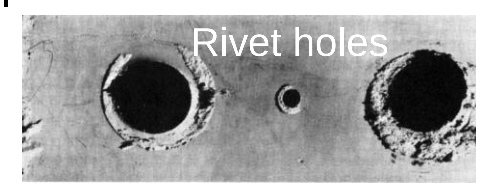
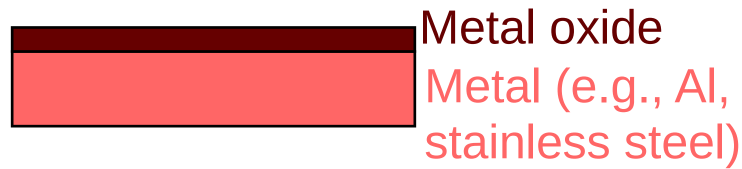


Fig. 17.15, *Callister 7e*. (Fig. 17.15 is courtesy LaQue Center for Corrosion Technology, Inc.)

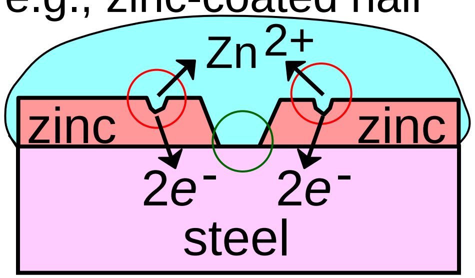
CONTROLLING CORROSION

- Self-protecting metals!
  -- Metal ions combine with O
  to form a thin, adhering oxide layer that slows corrosion.



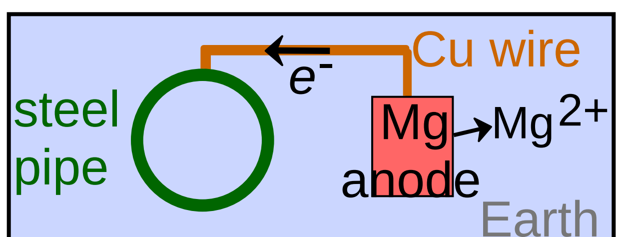
- Reduce *T* (slows kinetics of oxidation and reduction)
- Add inhibitors
  -- Slow oxidation/reduction reactions by removing reactants
  (e.g., remove O_2 gas by reacting it w/an inhibitor).
  -- Slow oxidation reaction by attaching species to
  the surface (e.g., paint it!).
- Cathodic (or sacrificial) protection
  -- Attach a more anodic material to the one to be protected.

e.g., zinc-coated nail



Adapted from Fig. 17.23, *Callister 7e*.

e.g., Mg Anode



Adapted from Fig. 17.22(a), *Callister 7e*. (Fig. 17.22(a) is from M.G. Fontana, *Corrosion Engineering*, 3rd ed., McGraw-Hill Book Co., 1986.)

SUMMARY

- Corrosion occurs due to:
  - -- the natural tendency of metals to give up electrons.
  - -- electrons are given up by an oxidation reaction.
  - -- these electrons then used in a reduction reaction.
- Metals with a more negative Standard Electrode Potential are more likely to corrode relative to other metals.
- The Galvanic Series ranks the reactivity of metals in seawater.
- Increasing *T* speeds up oxidation/reduction reactions.
- Corrosion may be controlled by:
  - -- using metals which form a protective oxide layer
  - -- reducing *T*
  - -- adding inhibitors
  - -- painting
  - -- using cathodic protection.

ANNOUNCEMENTS

Reading:

Core Problems:

Self-help Problems: