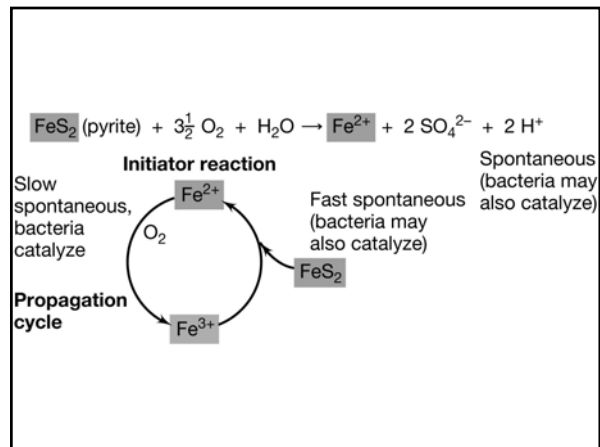
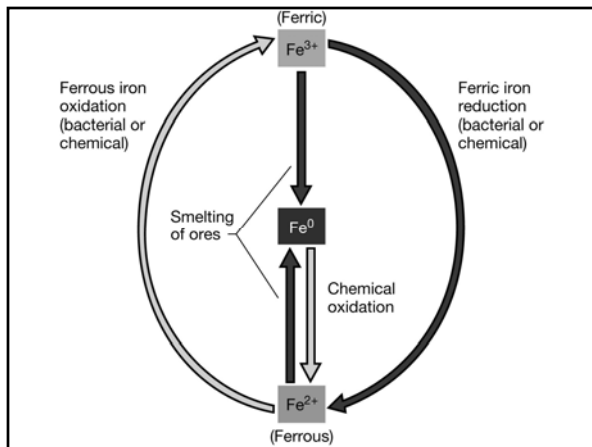
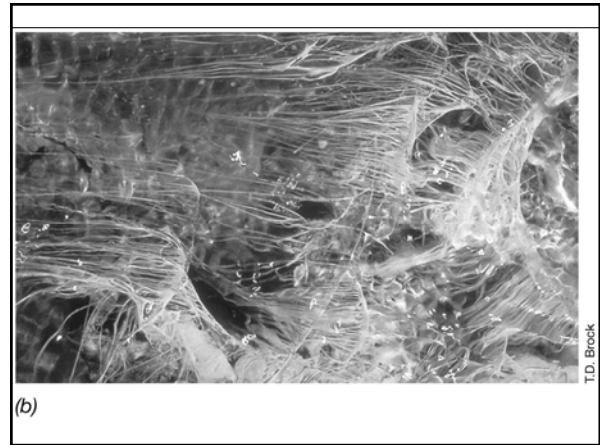
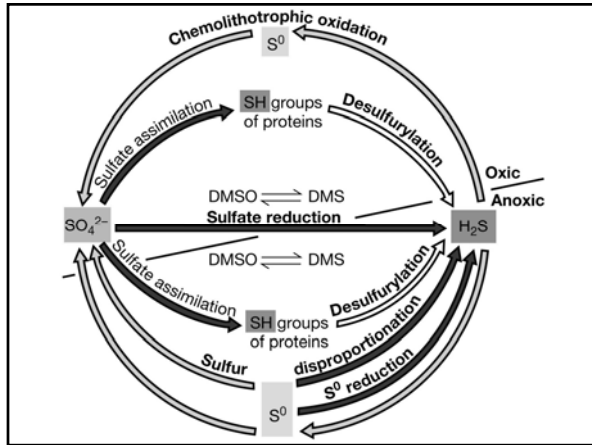


Fig. 22. A comparison between C, S and N oxidation/reductions. The most reduced and the most oxidized compounds of the C, S and N cycles are arranged in pairs, separated by a distance which represents an  $8e^-$  difference between the extremes. Given vertically are the  $G^\circ$  for the oxidation, by  $O_2$ , of the reduced form. There is a decreasing energy yield through the series C, S to N which is represented by the vertical distance between the oxidized and the reduced forms. The location of the lines relative to each other is only approximately correct and is designed to illustrate the decrease in reducing potential through the series  $H_2$ ,  $CH_4$ ,  $H_2S$  to  $NH_3$  and the increase in oxidizing potential through the series  $CO_2$ ,  $SO_4^{2-}$ ,  $NO_3^-$  to  $O_2$ .

### Key Processes and Prokaryotes in the Sulfur Cycle

Process	Organisms
<b>Sulfide/sulfur oxidation</b> ( $H_2S \rightarrow S^0 \rightarrow SO_4^{2-}$ )	
Aerobic	Sulfur chemolithotrophs ( <i>Thiobacillus</i> , <i>Beggiatoa</i> , many others)
Anaerobic	Purple and green phototrophic bacteria, some chemolithotrophs
<b>Sulfate reduction</b> (anaerobic) ( $SO_4^{2-} \rightarrow H_2S$ )	<i>Desulfovibrio</i> , <i>Desulfobacter</i> ,
<b>Sulfur reduction</b> (anaerobic) ( $S^0 \rightarrow H_2S$ )	<i>Desulfuromonas</i> , many hyperthermophilic <i>Archaea</i>
<b>Sulfur disproportionation</b> ( $S_2O_3^{2-} \rightarrow H_2S + SO_4^{2-}$ )	<i>Desulfovibrio</i> , and others
<b>Organic sulfur compound oxidation or reduction</b> ( $CH_3SH \rightarrow CO_2 + H_2S$ ) ( $DMSO \rightarrow DMS$ )	
<b>Desulfurylation</b> (organic-S $\rightarrow H_2S$ )	Many organisms can do this



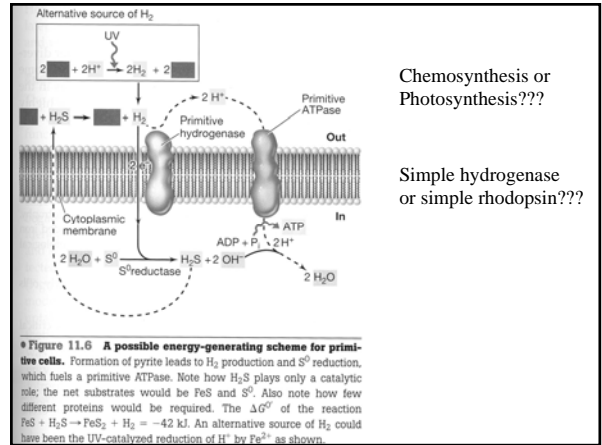
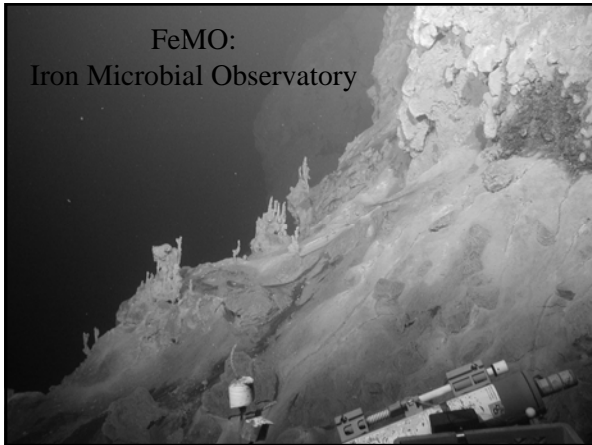
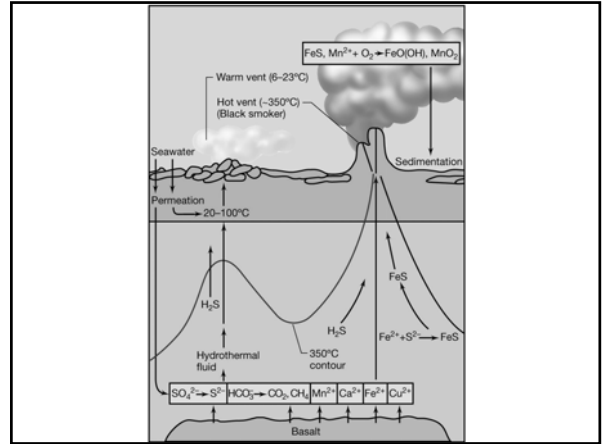
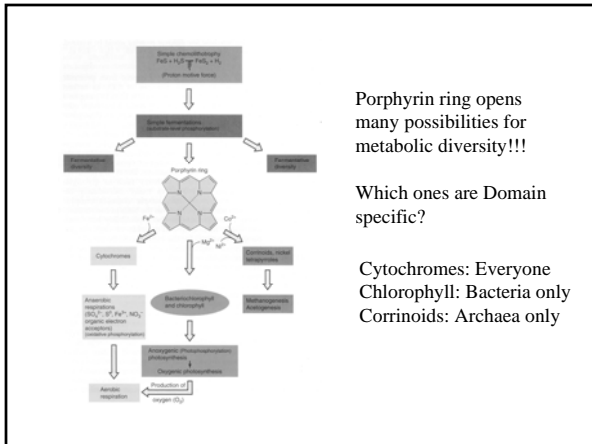


Figure 11.6 A possible energy-generating scheme for primitive cells. Formation of pyrite leads to  $\text{H}_2$  production and  $\text{S}^0$  reduction, which fuels a primitive ATPase. Note how  $\text{H}_2\text{S}$  plays only a catalytic role; the net substrates would be  $\text{FeS}$  and  $\text{S}^0$ . Also note how few different proteins would be required. The  $\Delta G^\circ$  of the reaction  $\text{FeS} + \text{H}_2\text{S} \rightarrow \text{FeS}_2 + \text{H}_2 = -42 \text{ kJ}$ . An alternative source of  $\text{H}_2$  could have been the UV-catalyzed reduction of  $\text{H}^+$  by  $\text{Fe}^{2+}$  as shown.



Porphyrin ring opens many possibilities for metabolic diversity!!!

Which ones are Domain specific?

Cytochromes: Everyone  
 Chlorophyll: Bacteria only  
 Corrinoids: Archaea only