

**Table 17.2.** Normal Oxidation-Reduction Potentials of Some Biologically Important Systems at pH 7.0

SYSTEM	$E'_0$	T IN °C.
Ketoglutarate $\rightleftharpoons$ succinate + CO <sub>2</sub> + 2H <sup>+</sup> + 2e	-0.68	-‡
Ferredoxin	-0.432	-§
Formate $\rightleftharpoons$ CO <sub>2</sub> + H <sub>2</sub>	-0.420	38
H <sub>2</sub> $\rightleftharpoons$ 2H <sup>+</sup> + 2e	-0.414	25
NADH + H <sup>+</sup> $\rightleftharpoons$ NAD <sup>+</sup> + 2H <sup>+</sup> + 2e	-0.317	30†
NADPH + H <sup>+</sup> $\rightleftharpoons$ NADP <sup>+</sup> + 2H <sup>+</sup> + 2e	-0.316	30†
Horseradish oxidase	-0.27	-†
FADH <sub>2</sub> $\rightleftharpoons$ FAD + 2H <sup>+</sup> + 2e	-0.219	30†
FMNH <sub>2</sub> $\rightleftharpoons$ FMM + 2H <sup>+</sup> + 2e	-0.219	30†
Lactate $\rightleftharpoons$ pyruvate + 2H <sup>+</sup> + 2e	-0.180	35
Malate $\rightleftharpoons$ oxaloacetate + 2H <sup>+</sup> + 2e	-0.102	37
Reduced flavin enzyme $\rightleftharpoons$ flavin enzyme + 2H <sup>+</sup> + 2e	-0.063	38
Luciferin* $\rightleftharpoons$ oxyluciferin + 2H <sup>+</sup> + 2e	-0.050	?*
Ferrocyanochrome B $\rightleftharpoons$ ferricyanochrome B + e	-0.04	25
Succinate $\rightleftharpoons$ fumarate + 2H <sup>+</sup> + 2e	-0.015	30
Decarboxylase	+0.19	-†
Ferrocyanochrome C $\rightleftharpoons$ ferricyanochrome C + e	+0.26	25
Ferrocyanochrome A $\rightleftharpoons$ ferricyanochrome A + e	+0.29	25
Ferrocyanochrome A <sub>3</sub> $\rightleftharpoons$ ferricyanochrome A <sub>3</sub> + e	?	-‡
H <sub>2</sub> O $\rightleftharpoons$ ½O <sub>2</sub> + 2H <sup>+</sup> + 2e	+0.815	25

Data from Goddard, 1945. Potentials in all cases are at or near neutrality.

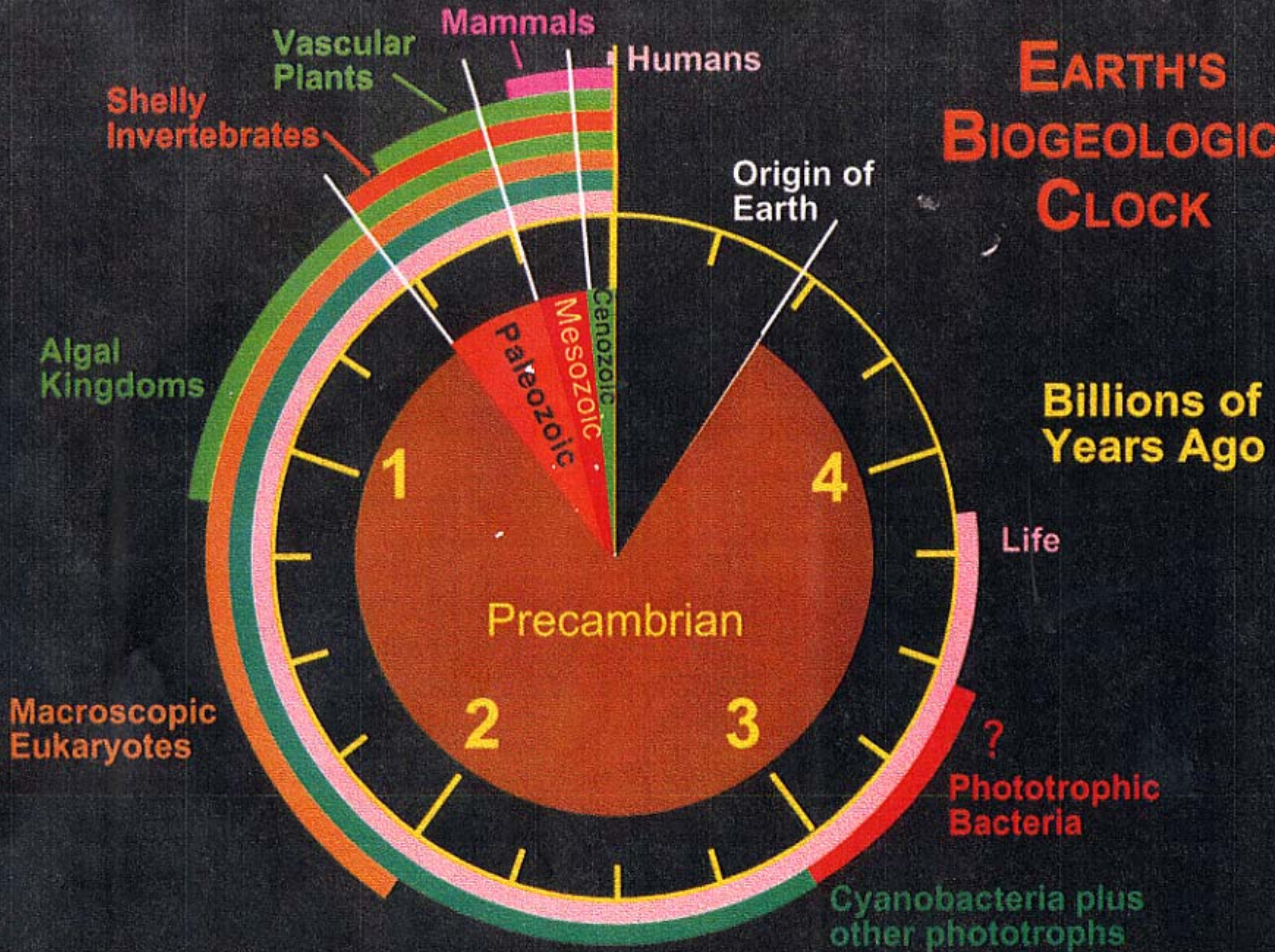
\* From McElroy and Strehler, 1954: *Bact. Rev.* 18.

† From Clark, 1960.

‡ From Goddard and Bonner, 1960: *In Plant Physiology, a Treatise*. Steward, ed. Academic Press, New York. Goddard and Bonner give the NADPH/NADP<sup>+</sup> system as -0.324, and NADH/NAD<sup>+</sup> as -0.320.

§ From Tagawa and Arnon, 1962: *Nature* 195:537-543. The value cited is for spinach ferredoxin.

# EARTH'S BIOGEOLOGIC CLOCK



D. J. Des Marais (2000)  
*Science* 289: 1703-1705.

# Carbon Pools in the Major Reservoirs on Earth

**Table 5.1** Carbon pools in the major reservoirs on Earth

Pools	Quantity ( $\times 10^{15}$ g)
Atmosphere	720
Oceans	38,400
Total inorganic	37,400
Surface layer	670
Deep layer	36,730
Total organic	1,000
Lithosphere	
Sedimentary carbonates	>60,000,000
Kerogens	15,000,000
Terrestrial biosphere (total)	2,000
Living biomass	600–1,000
Dead biomass	1,200
Aquatic biosphere	1–2
Fossil fuels	4,130
Coal	3,510
Oil	230
Gas	140
Other (peat)	250

From: Falkowski & Raven. Aquatic Photosynthesis. p. 130 (1997)

# REDOX REACTIONS ARE COUPLED ON

MICROSCOPIC SCALES

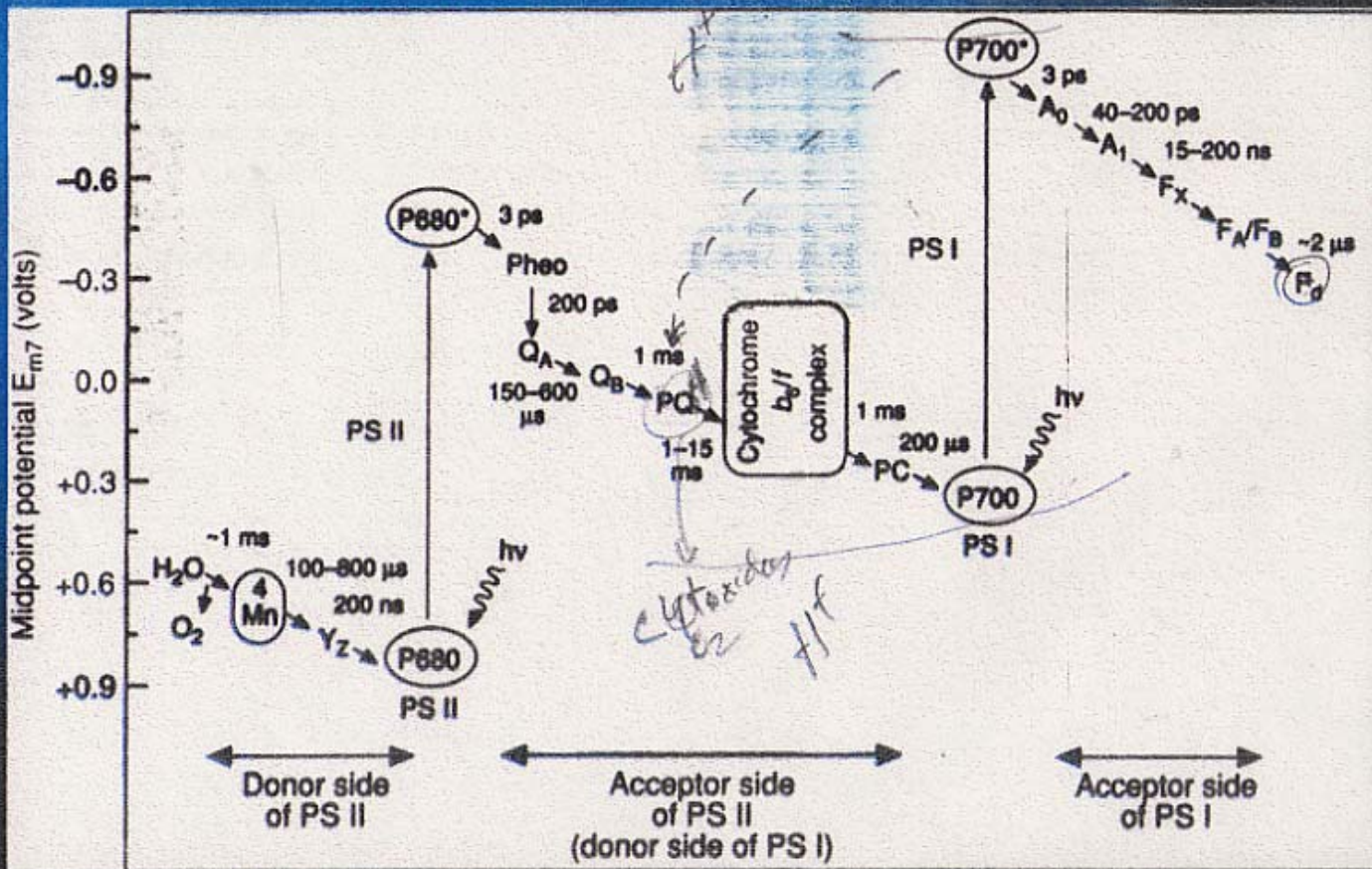
General Reaction  $\longrightarrow$



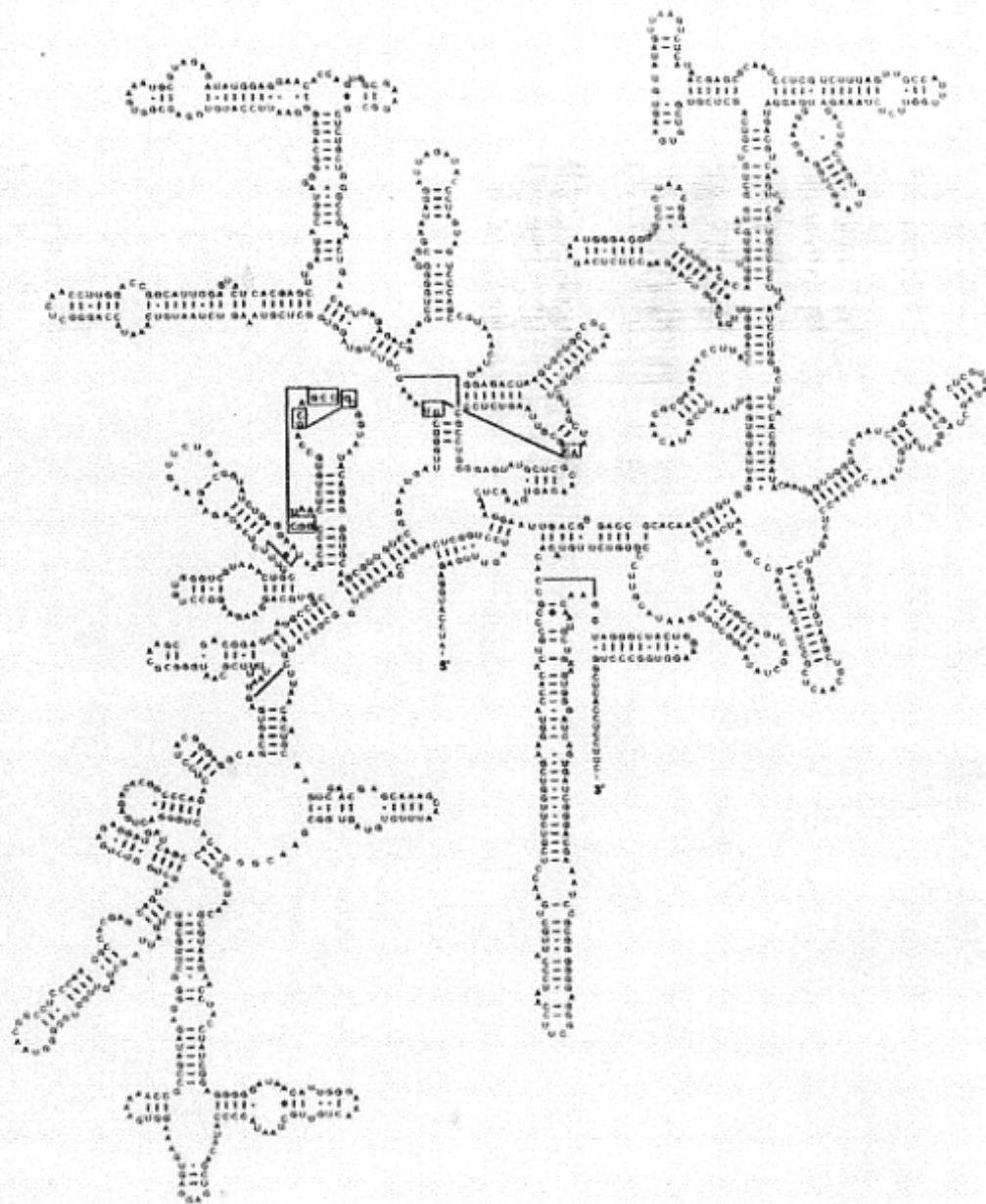
Photosynthesis



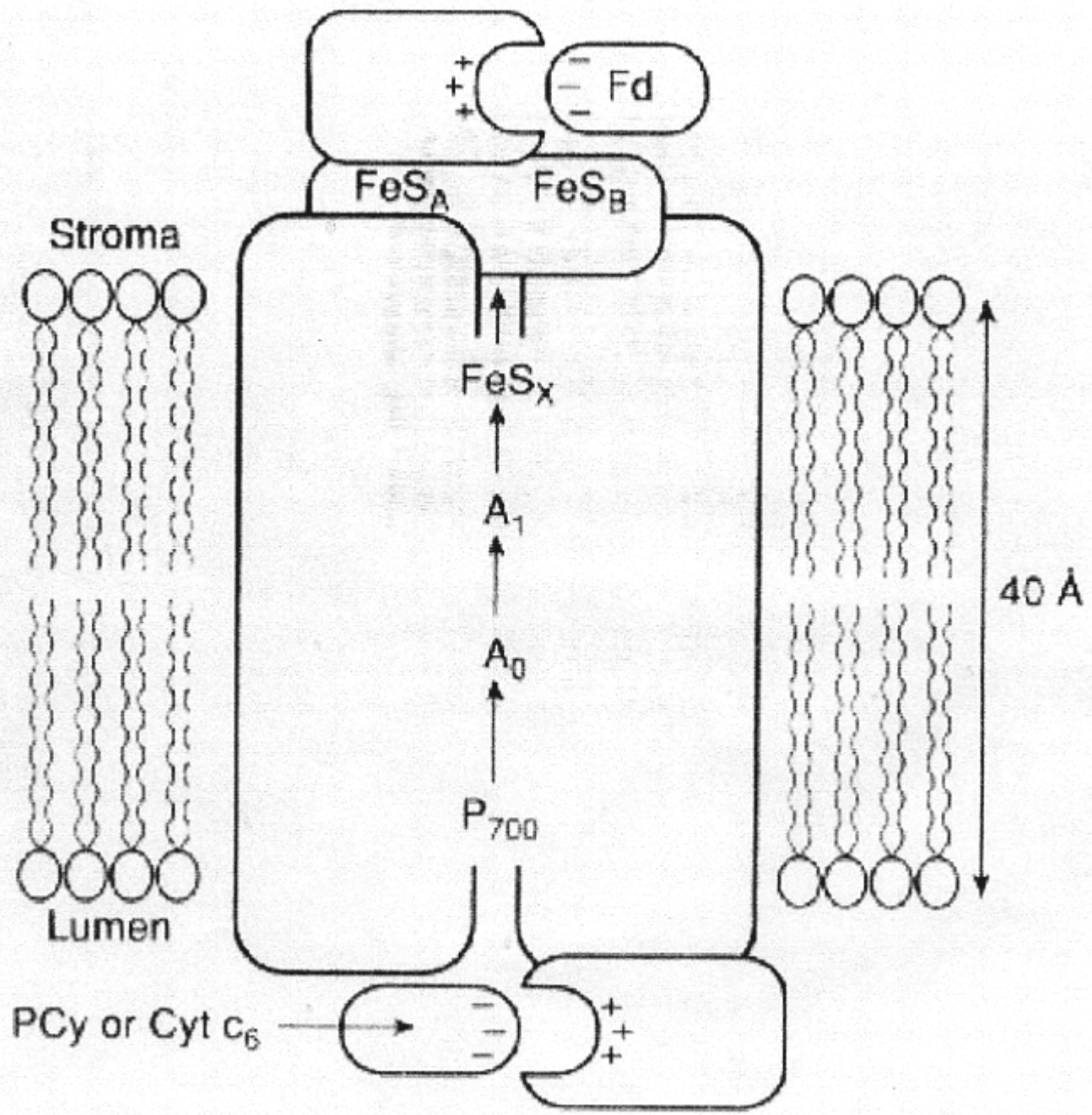
# Oxygenic Photosynthetic Electron Transport



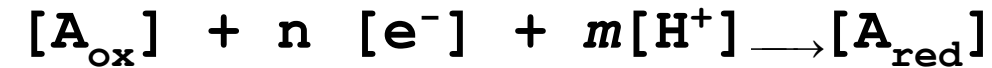
From: Falkowski & Raven. Aquatic Photosynthesis. (1997)



**Figure 1.7** The secondary structure of the small subunit (16S) rRNA from *Chlamydomonas reinhardtii*. The structure is inferred from homology with known structures in yeast and prokaryotes. Hollow circles and unpaired regions represent areas of generally higher variability between organisms.



## The Nernst Equation



where  $m$  is the number of protons involved in the reduction of  $\text{A}_{\text{ox}}$ .

The redox potential for this reaction can be calculated by:

$$E = E_{\text{m7}} + 59/n \log [\text{A}_{\text{red}}] / [\text{A}_{\text{ox}}] [\text{H}^+]^m$$

which can be rewritten as:

$$E = E_{\text{m7}} + 59/n \log ([\text{A}_{\text{red}}] / [\text{A}_{\text{ox}}]) + 59 (m/n) \text{pH}$$