

**ERRATA for *An introduction to Chemical Thermodynamics, Second edition 2009***

**Page 15, line 11** ... of one calcium carbide molecule into ...

**Page 25, last line** In the literature, the term exergonic is used for processes that can deliver work and for processes that need work to go the term endergonic is used.

**Page 29, first equation from top** kJ/mol

**Page 42, following eq. 4.5** Footnote added: "Important to note is that the reference temperature for this coefficient is taken to be 0 K which is correct for gasses. For other substances, other reference temperatures may apply."

**Page 41, line 5** ... of one calcium carbide molecule into ...

**Page 44, second line from bottom** ... whereas the enthalpy does not.

**Page 48, line 12** ... the reversible heat plus the lost work, ...

**Page 52, second and third equation from top** 
$$\frac{Q}{T} = \frac{Q_{\text{rev}} + W_{\text{lost}}}{T} = \Delta S + \Delta S_{\text{irr}}$$

which implies that

$$\Delta S_{\text{irr}} \geq 0$$

**Figure 5.4, legend** Horizontal axis velocity in m/s, vertical axis probability density.

**Page 57, eq. 6.5** 
$$\left(\frac{\partial G}{\partial \xi}\right)_{eq} = 0 \quad \text{and} \quad \left(\frac{\partial^2 G}{\partial \xi^2}\right)_{eq} > 0$$

**Page 61, second equation** 
$$K = \frac{a(\text{Ag}^+)a(\text{Cl}^-)}{a(\text{AgCl})} \approx s^2$$

**Page 66, Table 6.2, page 66**  $\Delta_{\text{vap}}H^\ominus = 30.7 \text{ kJ/mol}$

**Page 75, Figure 7.4**  $V_v - V$  and  $V - V_l$  interchanged.

**Page 88, legend figure 8.8** Characteristic shape of a Langmuir isotherm in a semi-logarithmic plot.

**Page 89, paragraph below first equation** ... occurring when the pressure equals  $p = K^{-1}p^\ominus$ .

**Page 97, Label horizontal axis figure 9.5** "mole fraction"

**Page 97, Label horizontal axis figure 9.6** "mole fraction"

**Page 99, legend figure 9.8**  $Tx$ -diagrams ...

**Page 106, eq 10.6** 
$$\Delta_{\text{mix}}H = \frac{1}{2}nx_Ax_B\tilde{h}_{AB}$$

**Page 106, equation at bottom of page** 
$$x_{B,\text{max}} = \exp\left(-\frac{h_{AB}}{RT}\right)$$

**Page 106, bottom line** For the *solubility parameter* defined as  $h_{AB}/RT$  one finds -5.5 for sweet water at room temperature.

**Page 109, top equation** 
$$\Delta_{\text{vap}}S(x_B) = \Delta_{\text{vap}}S + \Delta_{\text{mix}}S_A(x_B) = \Delta_{\text{vap}}S - Rx_B$$

**Page 109, third equation** 
$$\Delta T = \frac{RT_{\text{vap}}}{\Delta_{\text{vap}}S}x_B$$

**Page 113, below eq. 11.2**  $f_j = p_j \exp\left(\frac{(p_j - p^\ominus)B_j}{RT}\right)$

and the last two equations of the intermezzo

$$V_j = \frac{\partial V}{\partial n_j} = \frac{RT}{p_j} + B_j$$

and

$$\mu_j = \mu_j^\ominus + RT \ln \frac{p_j}{p^\ominus} + (p_j - p^\ominus)B_j$$

**Page 115, second equation from top**  $\Delta_{mix}\mu_B = \frac{\partial \Delta_{mix}G}{\partial n_B} = RT \ln x_B + \frac{1}{2}(1 - x_B)^2 h_{AB}$

**Page 117, table 11.1**  $10^{-3}$

**Page 117, equation below table 11.1**  $\kappa^{-1} \approx 0.3 \sqrt{\frac{m^\ominus}{I}} \text{ nm}$

**Page 123, line 5 from bottom** ... between the two parts.

**Page 142, 3rd paragraph from bottom** According to Prigogine, “in the linear regime, the total entropy production in a system subject to flow of energy and matter reaches a minimum value at the non-equilibrium stationary state”.

**Page 154, add to bottom last paragraph** Also, there is a special situation that arises when the second virial coefficient vanishes. Whereas for gases this defines the Boyle temperature, for macro-molecular solutions this defines the theta condition such as the theta temperature.

**Page 157, eq. 15.5** ... be expressed as

$$a = RTK_2V_A$$

where  $V_A$  denotes the molar volume of the solvent.

**Page 159, equation at bottom**  $F_{j,m}^g = -M_j g$

**Page 162, ea. 15.10**  $c(r) \approx c_0 \exp\left\{\frac{(M_p - \rho V_p)\omega^2}{2RT}(r^2 - r_0^2)\right\}$

**Page 163, eq. 15.11**  $c(r) \approx c_0 \exp\left\{\frac{Z_{eff}FE}{RT}(z - z_0)\right\}$  (1)

**Page 165, first equation**  $\frac{\Pi}{RT} = \frac{c}{M} + [B_{HS} - K_2V_m]\left(\frac{c}{M}\right)^2 + \dots$

**Page 165, second equation**  $\frac{\Pi}{RT} = \frac{c}{M} \left\{1 + B_{1s}\frac{c_s}{M_s}\right\} + \left[B_{HS} - K_2V_m \left\{1 + (2B_{1s} - B_{2s})\frac{c_s}{M_s}\right\}\right] \left(\frac{c}{M}\right)^2 + \dots$

**Page 165, third and fourth equation**  $B_{1s} = \frac{2\pi}{3} \mathcal{N}(d + d_s)^3$

and

$$2B_{1s} - B_{2s} = \frac{2\pi}{3} \mathcal{N}d_s^2(3d + 2d_s)$$

**Page 169, last equation of embedded text**  $\theta = \frac{1}{N} \frac{a}{\Xi} \frac{d\Xi}{da} = \frac{1}{N} \frac{d \log \Xi}{d \log a}$

Page 171, Eq. 16.7 
$$\Xi = \sum_{n,m} \binom{N-n+1}{n-m} \binom{n-1}{m} a^n u^m K^n \simeq \left( \frac{1 + aKu + \sqrt{4aK + (1 - aKu)^2}}{2} \right)^N$$

Page 174, Bottom 
$$[B^+](V + \Delta V) = \Delta V c_B$$

Page 180, Embedded text, first paragraph ... the Landau potential or semi grand potential

Page 182, fifth line from bottom ...and hence  $\Xi_2 = a^2(K_t^2 u_t + gK_g^2 u_g)$ .

Page 189, first equation

$$\begin{aligned} \Delta G &\approx n_{s,in} \mu_{s,in} - n_{s,tot} \mu_s^i + n_{s,out} \mu_{s,out} \\ &= RT \left\{ n_{s,in} \ln \frac{n_{s,in}}{n_{o,in}} + n_{s,out} \ln \left( \frac{n_{s,out}}{n_{s,tot}} \frac{n_{o,in}}{n_{o,out}} \right) \right\} < 0 \end{aligned}$$

Page 190, Embedded text

$$\begin{aligned} \frac{d(c_{s,in} - c_{s,out})}{dt} &= \frac{1}{V_{in}} \frac{dn_{s,in}}{dt} - \frac{1}{V_{out}} \frac{dn_{s,out}}{dt} \\ &= AJ_s \left( \frac{1}{V_{out}} + \frac{1}{V_{in}} \right) \\ &= -\frac{AD_s}{L} \left( \frac{1}{V_{out}} + \frac{1}{V_{in}} \right) (c_{s,in} - c_{s,out}) \end{aligned}$$

The above equation is a linear differential equation which yields an equilibration time scale that is equal to

$$\tau = \left\{ \frac{AD_s}{L} \left( \frac{1}{V_{out}} + \frac{1}{V_{in}} \right) \right\}^{-1}$$

Page 191, third equation from top 
$$J_s = -\frac{kT}{f} \frac{dc_s}{dz}$$

Page 192, second equation from bottom 
$$\theta = \frac{\sum_{k=1}^n k[ML_k]}{[M]}$$

Page 192, equation at bottom 
$$[L]_{tot} = [L]_{in} + \frac{V_{in}}{V_{in} + V_{out}} \sum_{k=1}^n k[ML_k]$$

o

Page 193, eq. 17.3 
$$r = \frac{[Cl^-]_{in}}{[Cl^-]_{out}} = \frac{Z[M^{Z+}]}{2[Cl^-]_{out}} + \sqrt{\left( \frac{Z[M^{Z+}]}{2[Cl^-]_{out}} \right)^2 + 1}$$