Solutions to exercises week 38

FYS2160

Kristian Bjørke, Knut Oddvar Høie Vadla

November 7, 2017

Schroeder 2.30

a) $\Omega_{\text{total}} = \frac{2^{4N}}{\sqrt{8\pi N}}$ $\frac{S}{k} = \ln \Omega_{\text{total}} = \ln \left[\frac{2^{4N}}{\sqrt{8\pi N}} \right] = 4N \cdot \ln 2 - \ln \sqrt{8\pi N}$ $N = 10^{23} \Rightarrow \frac{S}{k} = 2.77 \cdot 10^{23} - 28.1$ **b)** $\Omega_{\rm most~likely} = \frac{2^{4N}}{4\pi N}$ 4*πN* $\frac{S}{k} = \ln \Omega_{\rm most~likely} = \ln \left[\frac{2^{4N}}{4\pi N} \right] = 4N \cdot \ln 2 - \ln 4\pi N$ $N = 10^{23} \Rightarrow \frac{S}{k} = 2.77 \cdot 10^{23} - 55.5$ **c)**

Small difference in entropy, $\Delta S/k = 55.5 - 28.1 = 27.4$, time scale irrelevant. **d)**

Entropy decrease by 27 units out of 2.8×10^{23} , insignificant.

Schroeder 2.37

 $N_A = (1 - x)N$, $N_B = xN$

 $V_i^A = (1 - x)V_f, V_i^B = xV_f$ Use $\Delta S_X = -N_X k \ln \frac{V_i^X}{V_f}$ to get:

$$
\Delta S_{\text{mixing}} = \Delta S_A + \Delta S_B
$$

= -(1 - x)Nk ln(1 + x) - xNk ln x
= -Nk [x ln x + (1 - x) ln(1 - x)]

For $x=\frac{1}{2}$ $\frac{1}{2}$: $\Delta S_\text{mixing} = -Nk \left[\ln \frac{1}{2} \right]$ $\big] = Nk \ln 2$ (*N* is total number of molecules, which is 2*N* in eq. 2.54)

Schroeder 2.38

When we allow the system to mix, assuming an ideal mixture, the only changes will be that the different molecules can change places with each other. This gives:

 $\Delta\Omega = \begin{pmatrix} N \\ N \end{pmatrix}$ *N^A* $= \frac{N!}{N+1}$ *NA*!*NB*! $\Delta S_{\text{mixing}} = k \ln \left(\frac{N}{N} \right)$ *N^A* $\left(\int \frac{N!}{N_A!N_B!} \right)$ Use Stirling's approximation (ln $N! \approx N \ln N - N$) to get: $\Delta S_{\text{mixing}} = -Nk [x \ln x + (1 - x) \ln(1 - x)]$

Schroeder 3.3

Initially: $\frac{\delta S_A}{\delta U_A} > \frac{\delta S_B}{\delta U_B}$ δU_B U_A will increase, U_B will decrease until: *δS^A* $\frac{\delta S_{A}}{\delta U_{A}}=\frac{\delta S_{B}}{\delta U_{B}}$ δU_B

Schroeder 3.38

From ideal gas law:

 $P_i = x_i P = x_i N kT/V = N_i kT/V$

, so if P_i is fixed the gas component i is unchanged when adding another component to the mix.

For a mix of two ideal gases total entropy given by:

$$
S_{\text{total}} = S_A(U_A, V, N_A) + S_B(U_B, V, N_B)
$$

Chemical potential for gas A is:

 $\mu_A = -T\left(\frac{\delta S}{\delta N}\right)$ *δN^A* \setminus $U_{,V,N_B} = -T\left(\frac{\delta S_A}{\delta N_A}\right)$ *δN^A* \setminus *U,V* , same as if gas *B* was not present.

Schroeder 4.3

a)
\n
$$
Q_c = W\left(\frac{1}{e} - 1\right) = 1.5GW
$$
\nb)

Every second 1.5×10^9 J dumped in 10^5 kg water.

Heat capacity of water: $C = 4186$ J/[∘]C.

$$
\Delta T = \frac{Q}{C} = \frac{15 \text{kJ}}{4.2 \text{kJ}/^{\circ}\text{C}} = 3.6 \text{ °C}
$$

c)

Latent heat at room temp: $L = 2.4$ kJ/g Evaporation rate: $\frac{Q}{L} = 600 \text{ kg/s} = 0.6 \text{ m}^2/\text{s}$, 0.6% of river

Schroeder 4.7

Need to dump heat into "cold" reservoir separate from room to be effective, otherwise it would raise temperature not lower it.

Schroeder 4.8

Q^h always bigger than *Qc*, so temperature in room will increase.

Schroeder 4.14

a)

COP is defined as benefit divided by cost, heat pump aims to heat up so:

$$
COP = \frac{\text{benefit}}{\text{cost}} = \frac{Q_h}{W}
$$

b)

$$
Q_h = Q_c + W
$$

$$
COP = \frac{1}{1 - Q_c/Q_h} > 1
$$

c)

$$
Q_h = \frac{1}{2Q_c/Q_h} > 1
$$

2. law of thermodynamics:

$$
\frac{Q_h}{T_h} \ge \frac{Q_c}{T_c}
$$

$$
COP \ge \frac{T_h}{T_h - T_c}
$$

d)

Electric heater: $Q_h = W$, $COP = 1$

Heat pump: *COP >* 1

For $T_h = 25$ °C and $T_c = 0$ °C *COP* can in principle be as high as 12 for heat pump (usually much lower but still larger than 1).

Compendium 6.1

