The 2nd and 3rd law of thermodynamics

- P21. Calculate the change of entropy for the surroundings when 1.00 mol of $H_2O(l)$ is formed from its elements under standard conditions at 298.15 K. $\Delta_{\text{form}}H^{\bullet}(H_2O, l) = -285.8 \text{ kJ mol}^{-1}$. Provide a general statement on how the entropy of the surroundings changes in exothermic and endothermic processes. [958.6 J/K. In exothermic processes (like this one), the entropy for the surroundings increases, in endothermic processes, it decreases.]
- P22. Calculate the change of entropy for 1.00 mol perfect gas (ΔS), its surroundings (ΔS '), and ΔS_{tot} , when the volume of the gas is doubled in an
 - a) isotherm reversible, or $\triangle S = 5.76 \text{ J/K}$, $\triangle S' = -5.76 \text{ J/K}$, $\triangle S_{\text{tot}} = 0 \text{ J/K}$
 - b) adiabatic reversible expansion. $[\Delta S = \Delta S' = \Delta S_{tot} = 0 J/K]$
- P24. 2.00 mol of CO₂ is heated at atmospheric pressure from 0 °C to 1000 °C. Calculate the change in entropy for CO₂. The molar heat capacity of CO₂ at constant pressure is $C_{p,m} = (28.8 + 3.59 \times 10^{-2} T 1.04 \times 10^{-5} T^2) \text{ J mol}^{-1} \text{ K}^{-1}$. [$\Delta S = 144.4 \text{ J/K}$]
- P25. Calculate ΔS_{tot} when 100 g of 300 °C iron is placed into 1000 g of 15 °C water. The ironwater system is isolated from its surroundings. Specific heat values for iron and water are as follows: 0.460 J g⁻¹ K⁻¹ and 4.184 J g⁻¹ K⁻¹. [$\Delta S = 13.7 \text{ J/K}$]
- P26. Calculate the Carnot efficiency for a ruddy steam engine working with 100 °C steam and 60 °C condensate. Do the same calculation for a modern steam turbine operating with 300 °C steam and 80 °C condensate. [10.7%, 38.4%]