

**Change in a cycle** (or “There and back again”)

(a) Isothermal change:  $p_a V_a = p_b V_b = nRT_a$  so

$$\frac{p_b}{p_a} = \frac{V_a}{V_b} = \frac{1}{3.00} = 0.333$$

(b) Adiabatic change:  $p_a V_a^\gamma = p_c V_c^\gamma$  so

$$\frac{p_c}{p_a} = \left(\frac{V_a}{V_c}\right)^\gamma = \frac{1}{(3.00)^{7/5}} = 0.215$$

(c) Adiabatic change:  $T_a V_a^{\gamma-1} = T_c V_c^{\gamma-1}$  so

$$\frac{T_c}{T_a} = \left(\frac{V_a}{V_c}\right)^{\gamma-1} = \frac{1}{(3.00)^{2/5}} = 0.644$$

(d) For the isothermal expansion

$$W_1 = \int_{V_a}^{V_b} p dV = \int_{V_a}^{V_b} \frac{nRT_a}{V} dV = nRT_a \int_{V_a}^{V_b} \frac{dV}{V} = nRT_a \ln\left(\frac{V_b}{V_a}\right)$$

so

$$\frac{W_1}{nRT_a} = \ln\left(\frac{V_b}{V_a}\right) = \ln(3.00) = 1.10.$$

(e) For the isothermal expansion of an ideal gas there is no change in internal energy, so

$$\frac{Q_1}{nRT_a} = \frac{W_1}{nRT_a} = 1.10.$$

(f) For the isothermal expansion of an ideal gas there is no change in internal energy,  $\Delta E_{\text{int},1} = 0$ .

(g) For the isothermal expansion

$$\Delta S_1 = \int_i^f \frac{dQ}{T} = \frac{1}{T_a} \int_i^f dQ = \frac{Q_1}{T_a} \quad \text{so} \quad \frac{\Delta S_1}{nR} = \frac{Q_1}{nRT_a} = 1.10.$$

(h) For the adiabatic compression  $pV^\gamma = p_a V_a^\gamma$  so

$$\begin{aligned} W_3 &= \int_{V_c}^{V_a} p dV = \int_{V_c}^{V_a} \frac{p_a V_a^\gamma}{V^\gamma} dV = p_a V_a^\gamma \int_{V_c}^{V_a} \frac{1}{V^\gamma} dV = p_a V_a^\gamma \left[ -\frac{1}{(\gamma-1)V^{\gamma-1}} \right]_{V_c}^{V_a} \\ &= -\frac{p_a V_a^\gamma}{\gamma-1} \left[ \frac{1}{V_a^{\gamma-1}} - \frac{1}{V_c^{\gamma-1}} \right] = -\frac{p_a V_a}{\gamma-1} \left[ 1 - \left(\frac{V_a}{V_c}\right)^{\gamma-1} \right] \end{aligned}$$

and

$$\frac{W_3}{nRT_a} = -\frac{1}{\gamma-1} \left[ 1 - \left(\frac{V_a}{V_c}\right)^{\gamma-1} \right] = -\frac{1}{2/5} \left[ 1 - \frac{1}{(3.00)^{2/5}} \right] = -0.889.$$

(i) For the adiabatic compression  $Q_3 = 0$ .

(j) For the adiabatic compression

$$\frac{\Delta E_{\text{int},3}}{nRT_a} = \frac{Q_3}{nRT_a} - \frac{W_3}{nRT_a} = 0.889.$$

(k) For the adiabatic compression  $\Delta S_3 = 0$ .

( $\ell$ ) For the cooling at constant volume to reduce pressure,  $W_2 = 0$ .

(m) For the cycle  $\Delta E_{\text{int}} = 0$ , i.e.  $\Delta E_{\text{int},1} + \Delta E_{\text{int},2} + \Delta E_{\text{int},3} = 0$ , but we know  $\Delta E_{\text{int},1}$  from part (f) and  $\Delta E_{\text{int},3}$  from part (j). For the cooling at constant volume to reduce pressure,

$$\frac{\Delta E_{\text{int},2}}{nRT_a} = -0.889.$$

(n) Using  $\Delta E_{\text{int},2} = Q_2 - W_2$  for this cooling,

$$\frac{Q_2}{nRT_a} = -0.889.$$

(o) For the cycle  $\Delta S = 0$ , i.e.  $\Delta S_1 + \Delta S_2 + \Delta S_3 = 0$ , but we know  $\Delta S_1$  from part (g) and  $\Delta S_3$  from part (k). For the cooling at constant volume to reduce pressure,

$$\frac{\Delta S_2}{nR} = -1.10.$$

(p) The efficiency is

$$\frac{W_1 + W_2 + W_3}{Q_1} = 1 + \frac{W_3}{Q_1} = 0.192.$$

(q) The Carnot efficiency (maximum possible) is, from part (c),

$$1 - \frac{T_c}{T_a} = 0.356.$$

As required, the efficiency of this cycle is less than the Carnot efficiency.