## 3.3 Change of entropy

Here we provide some fundamental equations regarding the entropy for the expansion of a perfect gas under different boundary conditions resulting in reversible changes between an initial state i and a final state f. As stated above we use the first law to calculate  $dq_{rev}$  and finally apply Eq. (3.1) to calculation  $\Delta S$ :

• Isothermal expansion, i.e. U = 3/2 n RT = const. or  $0 = dU = \delta q + \delta w$  (cf. Eq. (2.16)):

$$\Delta S = \int_{i}^{f} \frac{dq_{rev}}{T} = \frac{1}{T} \int_{i}^{f} dq_{rev} = -\frac{1}{T} \int_{i}^{f} dw = n R \ln \frac{V_f}{V_i} = n R \ln \frac{p_i}{p_f}$$
(3.3)

• Isochoric expansion, i.e.  $\delta w = 0$ , or  $C_V dT = \delta q = dU$ :

$$\Delta S = \int_{i}^{f} \frac{dq_{rev}}{T} = C_V \int_{i}^{f} \frac{dT}{T} = C_V \ln \frac{T_f}{T_i}$$
(3.4)

• Adiabatic expansion:

$$\Delta S = 0 \quad \text{since} \quad \delta q = 0 \tag{3.5}$$

Fig. 3.1 visualizes the logarithmic behavior described by Eq. (3.3) and Eq. (3.4)

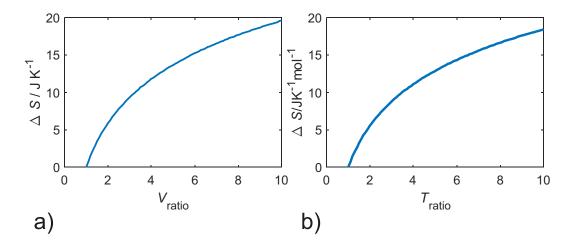


Figure 3.1: Entropy change for a) isothermal expansion; b) isochoric change.