



Entropy change for an ideal gas

(Visit <https://www.nasa.gov/centers/armstrong/multimedia/imagegallery/Schlieren/index.html> for more shock wave photos)

From the T - ds relations

$$Tds = du + pdv \quad \text{and} \quad Tds = dh - vdp$$

For an ideal gas

$$pv = RT \Rightarrow vdp + pdv = RdT \quad \text{and} \quad du = c_v(T)dT \quad \text{and} \quad dh = c_p(T)dT$$

| Temp. [K] | h [kJ/kg] | u [kJ/kg] | s° [kJ/kg/K] | pr | vr |
|-----------|-----------|-----------|--------------|--------|--------|
| 200 | 200.0 | 142.5 | 1.309 | 0.3363 | 1707.0 |
| 210 | 210.0 | 149.7 | 1.352 | 0.3987 | 1512.0 |
| 220 | 220.0 | 156.8 | 1.395 | 0.4690 | 1346.0 |
| 230 | 230.0 | 164.0 | 1.437 | 0.5477 | 1205.0 |
| 240 | 240.0 | 171.1 | 1.479 | 0.6355 | 1084.0 |
| 250 | 250.0 | 178.3 | 1.520 | 0.7329 | 979.0 |
| 260 | 260.0 | 185.4 | 1.559 | 0.8405 | 887.8 |
| 270 | 270.0 | 192.6 | 1.597 | 0.9590 | 808.0 |
| 280 | 280.1 | 199.8 | 1.633 | 1.0889 | 738.0 |

For the Ideal Gas model

- $p v = R T$ where $R = \bar{R}_u / M$ and $\bar{R}_u = 8.314 \text{ kJ}/(\text{kmol}\cdot\text{K})$ and M is the molecular weight. For example, $M_{\text{air}} = 28.98 \text{ kg}/\text{kmol} \Rightarrow R_{\text{air}} = 0.287 \text{ kJ}/(\text{kg}\cdot\text{K})$
- $c_p = c_v + R$ and $k \equiv \frac{c_p}{c_v}$
- $c_v = c_v(T)$, $c_p = c_p(T)$, $u = u(T)$, $h = h(T)$, $s = s(T, p)$
- $u(T_2) - u(T_1) = \int_{T_1}^{T_2} c_v(T) dT$ (find $u(T)$ in Ideal Gas Tables (IGTs))
- $h(T_2) - h(T_1) = \int_{T_1}^{T_2} c_p(T) dT$ (find $h(T)$ in IGTs)
- $s(T_2, p_2) - s(T_1, p_1) = s^0(T_2) - s^0(T_1) - R \ln \left(\frac{p_2}{p_1} \right)$ (find $s^0(T)$ in IGTs)
 - If the process is isentropic, i.e., $s_2 = s_1$, then,
 - $s^0(T_2) - s^0(T_1) = R \ln \left(\frac{p_2}{p_1} \right)$ (find $s^0(T)$ in IGTs)
 - $\frac{p_2}{p_1} = \frac{p_r(T_2)}{p_r(T_1)}$ (find $p_r(T)$ in IGTs)
 - $\frac{v_2}{v_1} = \frac{v_r(T_2)}{v_r(T_1)}$ (find $v_r(T)$ in IGTs)
- If the temperature range isn't too large, then it's reasonable to assume $c_v, c_p \approx \text{constants}$ (i.e., a perfect gas assumption) and,
 - $u(T_2) - u(T_1) \approx c_v(T_2 - T_1)$,
 - $h(T_2) - h(T_1) \approx c_p(T_2 - T_1)$
 - $s(T_2, v_2) - s(T_1, v_1) \approx c_v \ln \left(\frac{T_2}{T_1} \right) + R \ln \left(\frac{v_2}{v_1} \right)$
 - $s(T_2, p_2) - s(T_1, p_1) \approx c_p \ln \left(\frac{T_2}{T_1} \right) - R \ln \left(\frac{p_2}{p_1} \right)$
 - If the process is isentropic, i.e., $s_2 = s_1$, then,
 - $\frac{v_2}{v_1} = \left(\frac{T_2}{T_1} \right)^{\frac{1}{1-k}}$ $\frac{p_2}{p_1} = \left(\frac{T_2}{T_1} \right)^{\frac{k}{k-1}}$ $\frac{p_2}{p_1} = \left(\frac{v_2}{v_1} \right)^{-k}$