

fuel equivalence ratio.

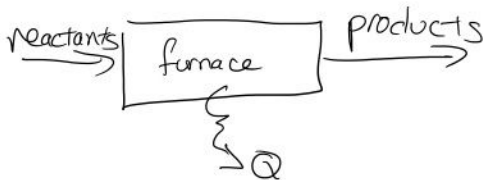
$$\phi = \frac{(A/F)_{\text{stoc.}}}{(A/F)_{\text{act.}}} = \frac{(F/A)_{\text{act.}}}{(F/A)_{\text{stoc.}}}$$

- $\phi = 1 \rightarrow$ stoch.
- $\phi < 1 \rightarrow$ fuel lean
- $\phi > 1 \rightarrow$ fuel rich

$$\% \text{ stoichiometric air} = \frac{100\%}{\phi}$$

$$\% \text{ excess air} = \frac{(1 - \phi)}{\phi} 100\%$$

heat (if exothermic)



$$\dot{Q} - \dot{W} = \Delta E$$

$$-\dot{Q} = \Delta h = \underline{h_{\text{prod}}} - h_{\text{react}}$$

Standard enthalpy: the sum of + the energy associated w/ chemical bonds, the enthalpy of formation h_f , and a temperature dependent enthalpy called the sensible enthalpy change Δh_s

$$\bar{h}_i(T) = \bar{h}_{f,i}^0(T_{\text{ref}}) + \Delta \bar{h}_{s,i}(T)$$

Standard enthalpy @ T for i

enthalpy of formation at STP (T_{ref}, P^0) for i

sensible enthalpy for $T_{\text{ref}} - T$

$\bar{h}_i(T) = \bar{h}_{f,i}^0(T_{\text{ref}}) + \Delta \bar{h}_{s,i}(T)$

$$\Delta h_{s,i} \equiv h_i(T) - h_{f,i}(T_{ref})$$

$$\bar{h}_f^{\circ} \rightarrow \text{STP (298K, 1atm)}$$

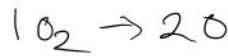
O: O₂

H: H₂

C: graphite

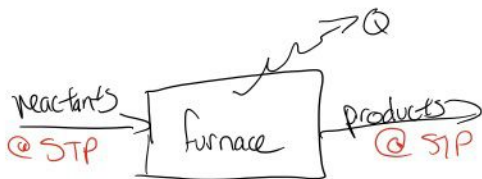
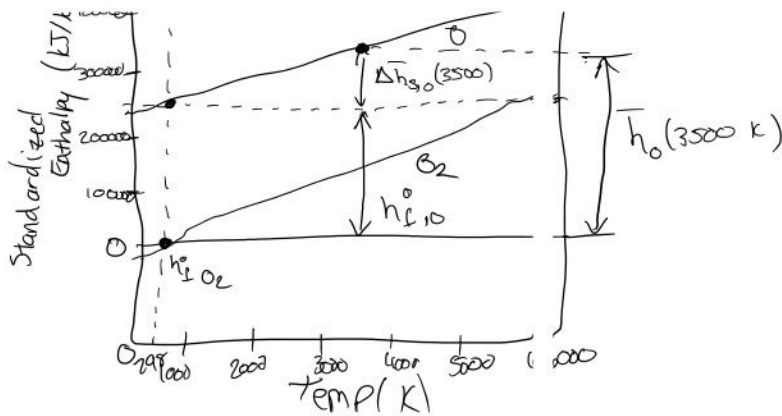
$$\left(\bar{h}_{f,i}^{\circ}\right)_{298} = 0$$

⊙: Bond Dissociation Energy O₂ @ 298K
= 498,340 kJ/kmol O₂



$$\hookrightarrow \left(\bar{h}_{f,i}^{\circ}\right) = 2 \cdot 19,195 \text{ kJ/kmol O}$$

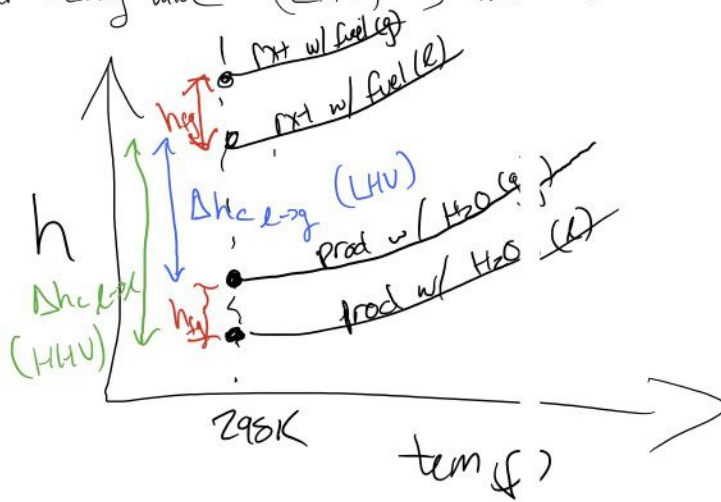
Standard Enthalpy



$$h_{prod} - h_{react} = -q \equiv \Delta h_{rxn}$$

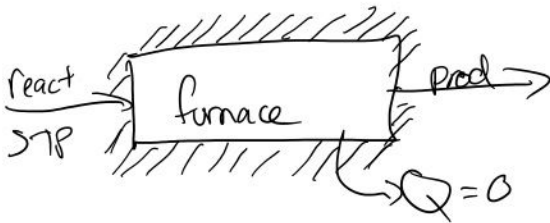
$$\Delta h_{rxn} = -\Delta h_{comb.}$$

- ↳ heating value
- upper heating value (HHV) → all water is liq.
 - lower heating value (LHV) → all water is vapor.



$$HHV > LHV$$

vaporized fuel has ↑ Δh_c



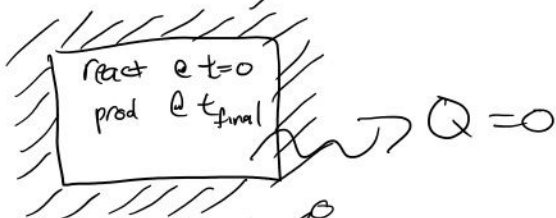
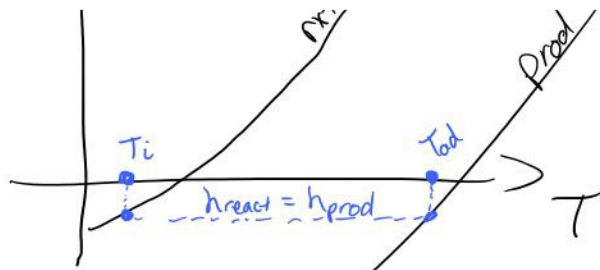
- assume adiabatic
- rxn. @ STP
- const P.

$$Q - W = \Delta h \Rightarrow h_{prod} = h_{rxn}$$

$$h_{rxn}(T_i, P) = h_{prod}(T_{ad}, P)$$

$h \uparrow$

x / /



$$\Delta E = U_{\text{prod}} - U_{\text{react}}$$

$$\rightarrow U_{\text{react}}(T_i, P_i) = U_{\text{prod}}(T_{\text{ad}}, P_f)$$

$$\rightarrow H_{\text{react}} = U_{\text{react}} + P_i V$$

$$H_{\text{react}} - U_{\text{react}} - P_i V + P_i V = 0$$

$$H_{\text{react}} - H_{\text{prod}} - R_u (T_i n_{\text{react}} - T_{\text{ad}} n_{\text{prod}}) = 0$$

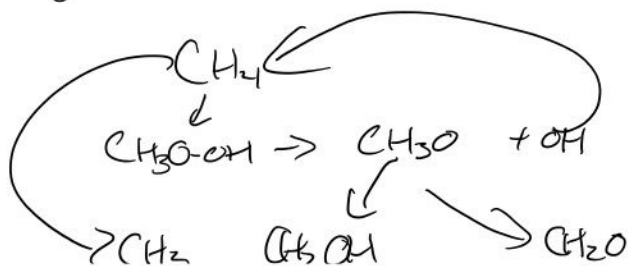
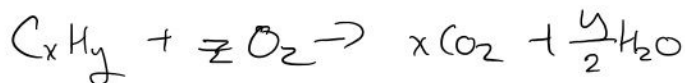
$$P_i V = n_{\text{react}} R_u T_i$$

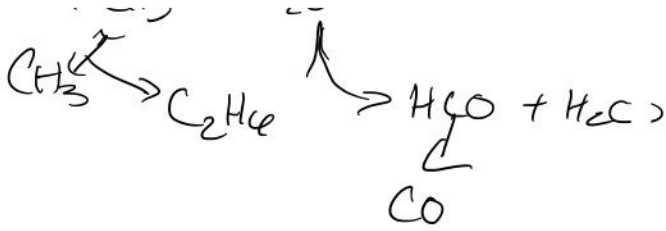
$$H_{\text{react}} - H_{\text{prod}} - R_u (T_i n_{\text{react}} - T_{\text{ad}} n_{\text{prod}}) = 0$$

$$n_{\text{react}} = n_{\text{prod}} = n$$

$$n_{\text{react}} M_{w_{\text{react}}} = n_{\text{prod}} M_{w_{\text{prod}}}$$

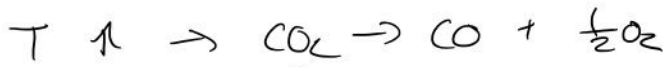
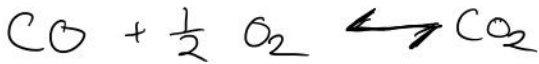
$$h_{\text{react}} - h_{\text{prod}} - R_u \left(\frac{T_i}{M_{w_{\text{react}}}} - \frac{T_{\text{ad}}}{M_{w_{\text{prod}}}} \right) = 0$$



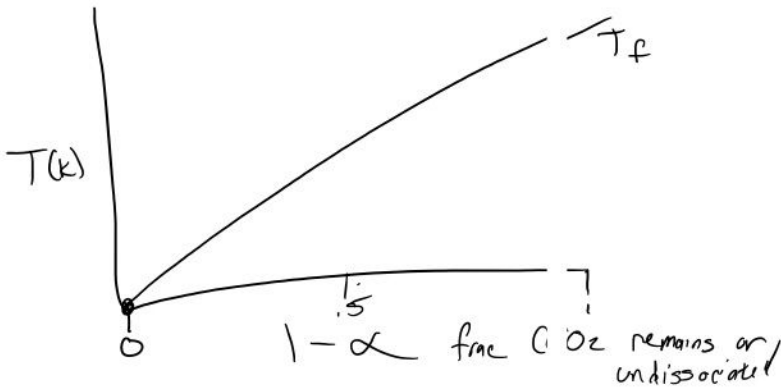
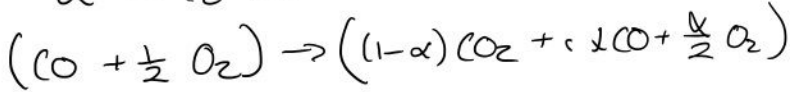


what is true?

- mass is conserved
- elements are conserved
- 1st 2nd laws apply



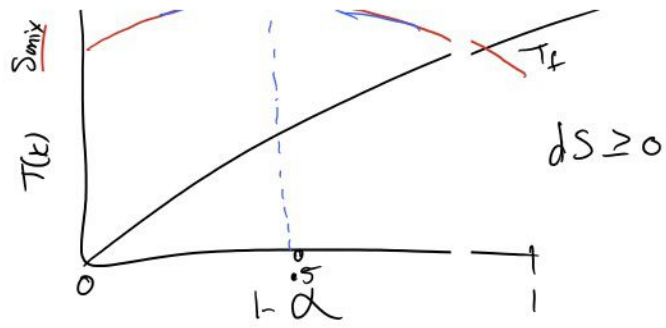
$$\alpha = \frac{x}{1-x}$$



$$\begin{aligned} S_{\text{final}} &= S_{\text{mix}}(T_f, P) = \sum_{\text{prod}} N_i \bar{s}_i(T_f, P) \\ &= (1-\alpha) \bar{s}_{\text{CO}_2} + \alpha \left(\bar{s}_{\text{CO}} + \frac{x}{2} \bar{s}_{\text{O}_2} \right) \end{aligned}$$

$$\bar{s}_i = \bar{s}_i(T_{\text{ref}}) + \int_{T_{\text{ref}}}^{T_f} \frac{c_{p,i}}{T} dT - R_u \ln\left(\frac{P_i}{P}\right)$$

$\xrightarrow{\text{Smax}}$



$$(dS)_{u,v,m} = 0 \text{ @ eq.}$$

NASA Gas Eq \leftarrow