One man's magic is another man's engineering.

Lazarus Long

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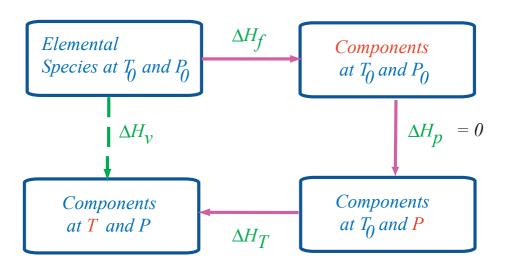
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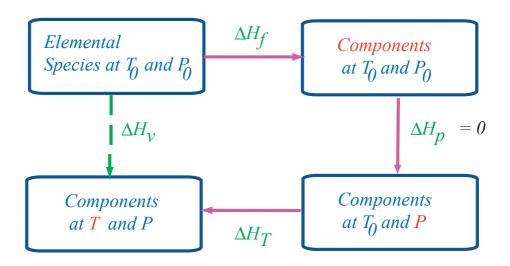
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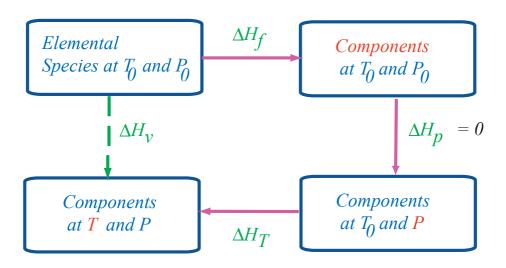
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 - 3. Standard enthalpy reference ($\Delta H = 0$) is at $P_0 = 1 \ atm, T_0 = 298 \ K$ and elemental species.

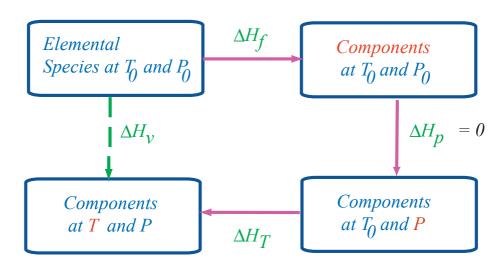




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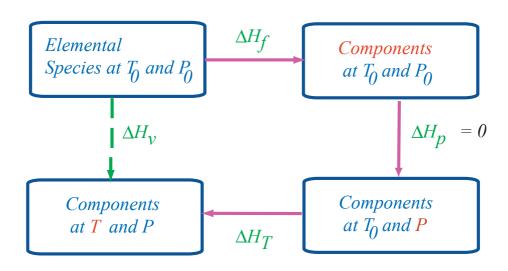


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$$\Delta H_f = y_1 H_{f,1}(T_0) + \dots + y_n H_{f,n}(T_0)$$

where $H_{f,k}(T_0)$ is the heat of formation of component k at temperature T_0 .

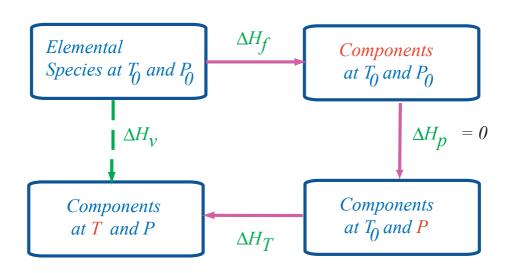


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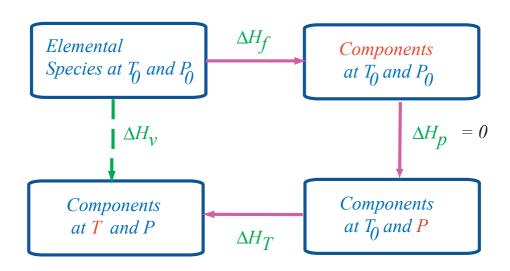
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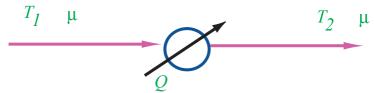
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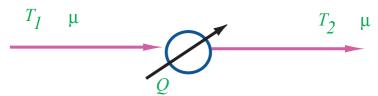
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Consider a heat exchanger with no phase change.



There is temperature change but no composition change.

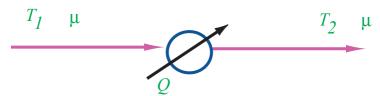
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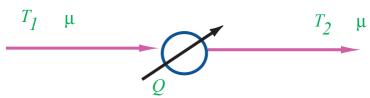
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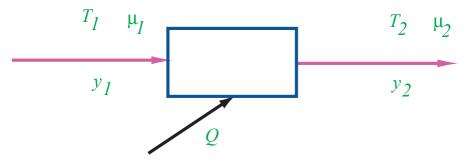
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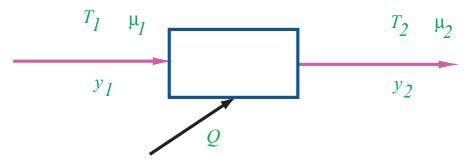
$$Q = \mu \left(y_1 \int_{T_1}^{T_2} c_{p,1}^0 dT + \dots + y_n \int_{T_1}^{T_2} c_{p,n}^0 dT \right)$$

Consider a gas phase reactor.



There is temperature change as well as composition change.

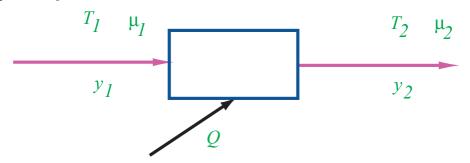
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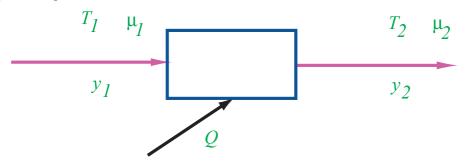
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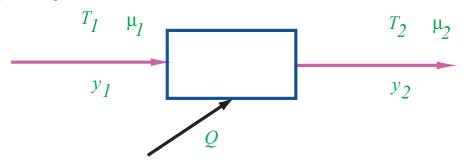
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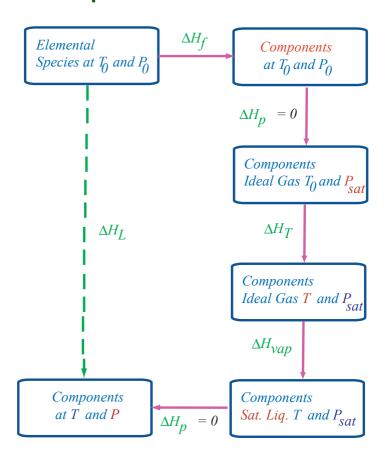
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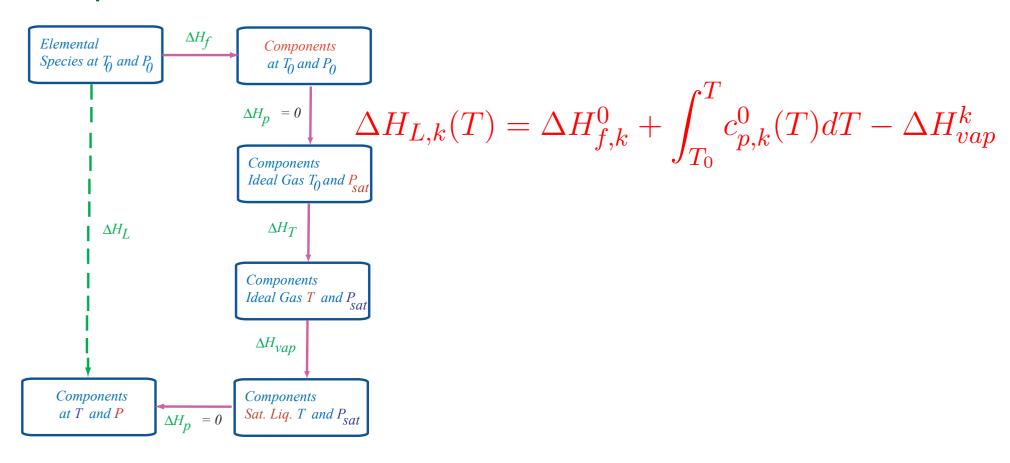
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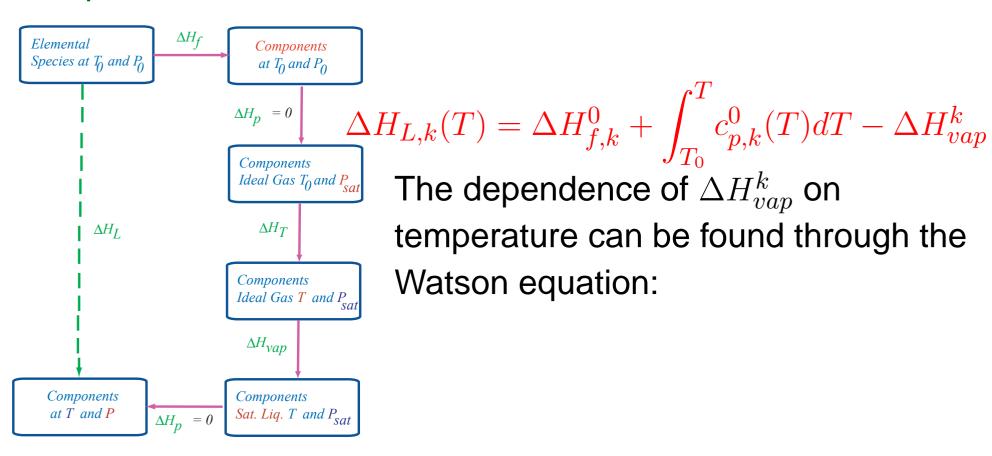
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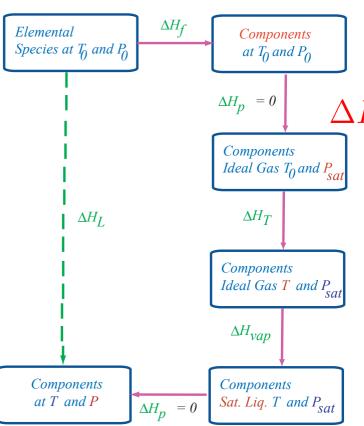
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Enthalpies of liquid mixtures are evaluated directly from the ideal vapor enthalpy and subtracting the heat of vaporization at saturation conditions.



$$\Delta H_{L,k}(T) = \Delta H_{f,k}^0 + \int_{T_0}^T c_{p,k}^0(T) dT - \Delta H_{vap}^k$$
 The dependence of $\Delta H_{p,k}^k$ on

The dependence of ΔH_{vap}^k on temperature can be found through the Watson equation:

$$\Delta H_{vap}^{k}(T) = \Delta H_{vap}^{k}(T_b) \left[\frac{(T_c^k - T)}{(T_c^k - T_b)} \right]^{0.38}$$

 T_c^k : critical temperature

 T_h^k : boiling point at 1 atm

 $\Delta H_{vap}^k(T_b)$ heat of vaporization at T_b^{k-1}

Liquid Mixtures and Two Phase Mixtures

The specific stream enthalpy for liquid mixtures is estimated as a weighted sum of respective mole fractions.

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• For a vapor-liquid mixture with vapor fraction ϕ , the stream enthalpy is given by the weighted sum of the liquid and vapor enthalpies.

$$\Delta H_L(T, x) = \phi \Delta H_V(T, y) + (1 - \phi) \Delta H_L(T, x)$$

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 T_V , P_V specified

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Bubble point temperature

Dew point temperature

Bubble point pressure

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Nonadiabatic flash

Percent vaporization flash

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Adiabatic flash

 Q, P_V specified

Nonadiabatic flash

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We did not consider the flash calculations that involved energy balance. We will consider the adiabatic flash now.

Nonadiabatic/Adiabatic Flash

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- If $\Delta H_{dew} \geq \Delta H_{spec} \geq \Delta H_{bub}$, guess ξ_n (or ϕ).

Perform a flash calculation with ξ_n (or ϕ) and P specified to obtain y_k , x_k and T. Calculate $\Delta H(T) = \phi \Delta H_v + (1-\phi)\Delta H_L$

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 Calculate $\Delta H(T) = \phi \Delta H_v + (1 \phi) \Delta H_L$
- If $f = \Delta H_{spec} \Delta H(T) = 0$ STOP. Otherwise, if f > 0 reguess a higher ξ_n (or ϕ), else guess a lower ξ_n (or ϕ) and go back to previous step.

Consider a 50-50 liquid mixture of benzene and toluene flowing at $100 \ gmol/s$ at $300 \ K$ and $1 \ bar$. If heat is added to this stream at the rate of $860.42 \ kcal/s$, what is the temperature of the mixture.

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• Guess ξ_T and solve for ξ_B , ϕ and T from:

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$$\xi_B = \frac{\alpha_{B/T}\xi_T}{1+(\alpha_{B/T}-1)\xi_T}$$

$$\phi = \frac{50(\xi_B+\xi_T)}{100}$$

$$P_T^0(T) = \frac{P}{\bar{\alpha}}$$

$$\Delta H(T) = \phi \Delta H_V(T) + (1 - \phi) \Delta H_L(T)$$

and compare with $\Delta H_{spec} = 12.862$.

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$-\xi_T$	ξ_B	T	ϕ	$\Delta H(T)$
-0.700	0.851	370.1	0.776	68.439

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0.600	0.786	369.5	0.693	20.876

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0.570	0.765	369.4	0.667	4.895

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0.585	0.776	369.5	0.680	12.993

$$\Delta H(T) = \phi \Delta H_V(T) + (1 - \phi) \Delta H_L(T)$$

and compare with $\Delta H_{spec} = 12.862$.

• Adjust ξ_T till the computed ΔH converges to the specified value.

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Thus, the stream is 68% vaporized at a temperature of 369.5 K.