

# ME-662 CONVECTIVE HEAT AND MASS TRANSFER

A. W. Date  
Mechanical Engineering Department  
Indian Institute of Technology, Bombay  
Mumbai - 400076  
India

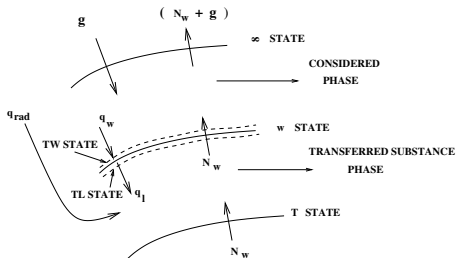
## LECTURE-34 REYNOLDS FLOW MODEL

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- 1 Reynold Flow Model - Definitions
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# Definitions - 1 - L34( $\frac{1}{14}$ )

- 1 This model postulates fictitious mass flux ( $g$ ) called Reynolds flux in the  $\infty$  - state
- 2 Model develops algebraic relations that are related to real transport phenomena by reference to ( $g$ )
- 3 Model develops  $N_w = g \times B$  relation.
- 4 The objective is to relate  $g$  to heat transfer coefficient  $h_{N_w \rightarrow 0}$



- 1 TW - state is just inside the Considered phase
- 2 TL - state is just inside the Neighbouring phase
- 3 In the T - state, temperature and compositions are uniform

## Definitions - 2 - L34( $\frac{2}{14}$ )

- ① **Mass Transfer Flux** is sum of all species transferred across the interface. Thus

$$N_w = \sum_j N_{j,w} = \sum_k N_{k,T}$$

where **species j and k need not be the same** .

- ②  $q_w$  is **positive** when flowing towards the interface from the considered phase.
- ③  $q_l$  is **positive** when flowing away from the interface in the Neighbouring phase
- ④ Therefore,  $(q_w - q_l) = N_w (h_{TW} - h_{TL})$  where  $h_{TW}$  and  $h_{TL}$  are **enthalpies of the transferred substance at the TW and TL states** . When  $(q_w - q_l) \neq 0$ , **phase-change occurs** .
- ⑤  $q_{rad}$  **from Considered phase to the interface** is positive. It is accounted in the Neighbouring phase for convenience.

# The Hypothesis - L34( $\frac{3}{14}$ )

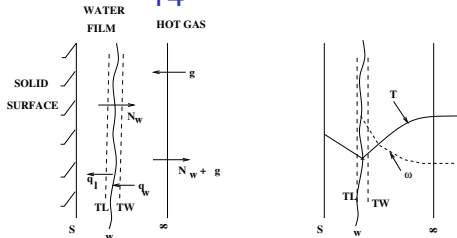
- 1 In different applications with or without chemical reaction, complex processes occur in the considered phase
- 2 To avoid solution of any differential equations, it is assumed that the complex processes can be effectively simulated by
  - 1 A fictitious flux ( $g$ ) through the  $\infty - \infty$  plane towards the interface carrying with it properties of the  $\infty$  - state .
  - 2 A fictitious flux ( $N_w + g$ ) through the  $\infty - \infty$  plane away from the interface carrying with it properties of the  $w$  - state .
- 3 The magnitude of ( $g$ ) is not affected by presence of gradients of  $\omega_j$  or  $T$  or turbulence, radiation etc in the considered phase
- 4 The  $g$  - flux is also not affected by direction of  $N_w$
- 5 The  $g$ -flux is supposed to produce same effect at the interface that the real flow is likely to produce

# An Illustration

## Evaporative Cooling - L34( $\frac{4}{14}$ )

Consider a solid surface ( $T_s$ ) to be protected from hot gases ( $T_\infty > T_s$ ) and ( $\omega_{v,\infty} < \omega_{v,w}$ ) by a thin water film. The Reynolds Hypothesis states that a  $g$ -flux of hot gases towards the interface together with ( $N_w + g$ ) flux of cooler moist air away from the interface will produce the effect:  $N_w > 0$ ,  $q_w > 0$  and  $q_l < 0$ .

The real flow will also produce precisely these effects.



In this sense, the  $g$ -flux is considered capable of responding to the mass-fraction and temperature gradients as well as turbulence effects in the considered phase as shown in the right figure.

# Momentum Transfer - L34( $\frac{5}{14}$ )

- 1 Consider control volume between  $\infty$ - and w- states. Then

$$\text{Rate of Momentum Flux In} = g U_{\infty}$$

$$\text{Rate of Momentum Flux Out} = \tau_w + (N_w + g) u_w = \tau_w$$

- 2 Hence,

$$\frac{g U_{\infty}}{\tau_w} = 1 = \frac{N_w}{\rho V_w} \text{ or}$$

$$N_w = g \times \frac{\rho v_w U_{\infty}}{\tau_w} = g \times \frac{v_w / U_{\infty}}{\tau_w / (\rho U_{\infty}^2)}$$

$$N_w = g \times B_f \rightarrow B_f = \frac{v_w / U_{\infty}}{(C_{f,x}/2)}$$

- 3 Further  $g = \tau_w / U_{\infty} = \rho U_{\infty} (C_{f,x}/2)$

# Single Phase Conv HT - 1 - L34( $\frac{6}{14}$ )

- 1 Here, CV between T- and  $\infty$ -states is considered. Then

$$\begin{aligned}\text{Rate of Heat Flux In} &= g h_{\infty} + N_w h_T \\ \text{Rate of Heat Flux Out} &= (N_w + g) h_w\end{aligned}$$

- 2 Equating and rearranging and taking  $T_{ref} = 0$

$$N_w = g \times B_h \rightarrow B_h = \frac{h_{\infty} - h_w}{h_w - h_T} = \frac{(c_p T)_{\infty} - (c_p T)_w}{(c_p T)_w - (c_p T)_T}$$

where  $N_w = \rho_w V_w$

- 3 If specific heats are equal

$$B_h = \frac{T_{\infty} - T_w}{T_w - T_T}$$



# Single Phase Conv HT - 2 - L34( $\frac{7}{14}$ )

- ① Further, CV between  $\infty$ - and w-states is considered. Then

$$\begin{aligned}\text{Rate of Heat Flux In} &= g h_{\infty} + N_w h_w \\ \text{Rate of Heat Flux Out} &= (N_w + g) h_w + q_w\end{aligned}$$

- ② Equating and rearranging  $g = q_w / (h_{\infty} - h_w)$   
③ If specific heats are equal

$$g = \frac{q_w}{c_p (T_{\infty} - T_w)} = \frac{h_{cof,v_w}}{c_p}$$

Thus,  $g$  is related to  $h_{cof,v_w}$ .

# Inert MT without HT - L34( $\frac{8}{14}$ )

- 1 Here, in all states, temperatures are equal ( $T_T = T_w = T_\infty$ )
- 2 Consider CV between T- and  $\infty$  - states. Then

$$\begin{aligned}\text{Rate of mass Flux In} &= g \omega_{V,\infty} + N_w \omega_{V,T} \\ \text{Rate of Mass Flux Out} &= (N_w + g) \omega_{V,w}\end{aligned}$$

- 3 Equating and rearranging

$$N_w = g \times B_m \quad \rightarrow \quad B_m = \frac{\omega_{V,\infty} - \omega_{V,w}}{\omega_{V,w} - \omega_{V,T}}$$

where for a pure liquid in the T-state,  $\omega_{V,T} = 1$ .  
 $\omega_{V,w}$  is evaluated from equilibrium at  $T_w$

# Inert MT with HT - 1 - L34( $\frac{9}{14}$ )

- 1 Here, let  $T_\infty \neq T_w \neq T_T$ . Then
- 2 Consider, CV between  $\infty$ - and T-states is considered. Then

$$\text{Rate of Heat Flux In} = g h_{m,\infty} + N_w h_{m,T}$$

$$\text{Rate of Heat Flux Out} = (N_w + g) h_{m,w}$$

$$\text{Equating : } N_w = g \times B_{mh} \quad \text{where} \quad B_{mh} = \frac{h_{m,\infty} - h_{m,w}}{h_{m,w} - h_{m,T}}$$

$$h_m = \omega_v h_v + (1 - \omega_v) h_a$$

$$h_v = c_{p,v} (T - T_{ref}) + \lambda_{ref}, \quad h_a = c_{p,a} (T - T_{ref})$$

- 3 Making  $Le = 1$  assumption,  $B_{mh} = B_m$  from which  $\omega_w \sim h_w (T_w)$  relationship is iteratively calculated.

## Inert MT with HT - 2 - L34( $\frac{10}{14}$ )

- ① Now, consider CV between  $\infty$ - and w-states. Then

$$q_w + (g + N_w) h_{m,w} = g h_{m,\infty} + N_w h_{m,TW}$$
$$N_w = g \times B_{mh} \quad \text{where} \quad B_{mh} = \frac{h_{m,\infty} - h_{m,w}}{h_{m,w} - h_{m,TW} + q_w/N_w}$$

- ② But,  $(q_w - q_l) = N_w (h_{TW} - h_{TL})$ . Hence

$$B_{mh} = \frac{h_{m,\infty} - h_{m,w}}{h_{m,w} - h_{m,TL} + q_L/N_w}$$

If  $T_w = T_{bp}$ ,  $q_L = 0$  and  $h_{m,w} - h_{m,TL} = h_{fg}$ .

- ③ If radiation is included then  $N_w h_{m,T} + q_L + q_{rad} = N_w h_{TL}$ .  
Hence

$$B_{mh} = \frac{h_{m,\infty} - h_{m,w}}{h_{m,w} - h_{m,T} - q_{rad}/N_w}$$

# MT with HT and SCR - 1 - L34( $\frac{11}{14}$ )

- 1 Here, we have 3 species:  $fu$ ,  $O_2$  and  $Pr$  with stoichiometric coefficient  $r_{st} = \text{kg of } O_2 / \text{kg of fuel}$ .
- 2 Consider CV between  $\infty$ - and T-states. Then

$$\text{For fuel} \quad g \omega_{fu,\infty} + \omega_{fu,T} N_w - \omega_{fu,w} (g + N_w) = - |R_{fu}|$$

$$\text{For } O_2 \quad g \omega_{O_2,\infty} + \omega_{O_2,T} N_w - \omega_{O_2,w} (g + N_w) = - |R_{O_2}|$$

- 3 But,  $R_{O_2} = r_{st} R_{fu}$ . Hence, dividing the 2nd Eqn by  $r_{st}$  and subtracting from 1st Eqn, we have

$$N_w = g \times B_\phi \rightarrow B_\phi = \frac{\phi_\infty - \phi_w}{\phi_w - \phi_T} \rightarrow \phi = \omega_{fu} - \frac{\omega_{O_2}}{r_{st}}$$

- 4 By the same reasoning, it is also possible to define

$$\phi = \omega_{fu} + \frac{\omega_{pr}}{1 + r_{st}} = \frac{\omega_{pr}}{1 + r_{st}} + \frac{\omega_{O_2}}{r_{st}}$$

# MT with HT and SCR - 2 - L34( $\frac{12}{14}$ )

- ① Invoking the **energy conservation principle** and assuming  $c_{p,k} = c_{pm}$ , we have shown that

$$h_m = \sum_k \omega_k h_k = c_{pm}(T - T_{ref}) + \omega_{fu} \Delta h_c = c_{pm}(T - T_{ref}) + \frac{\omega_{O_2}}{r_{st}} \Delta h_c$$

- ② Consider CV between  $\infty$ - and w-states. Then

$$g h_{m,\infty} + N_w h_{m,TW} = (g + N_w) h_{m,w} + q_w$$

- ③ Taking the 2nd definition for  $h_m$ , we have  $N_w = g \times B_{mh}$  where **taking**  $T_{ref} = T_w$

$$B_{mh} = \frac{h_{m,\infty} - h_{m,w}}{h_{m,w} - h_{m,TW} + q_w/N_w}$$

$$B_{mh} = \frac{c_{pm}(T_\infty - T_w) + \Delta h_c (\omega_{O_2,\infty} - \omega_{O_2,w})/r_{st}}{\Delta h_c (\omega_{O_2,w} - \omega_{O_2,TW})/r_{st} + q_w/N_w}$$

# MT with HT and SCR - 3 - L34( $\frac{13}{14}$ )

- ① For a **volatile liquid fuel** or **transpiration cooling** by a **combustible gas** ( e. g.  $H_2$  ),  $\omega_{O_2,w} = 0$  and since the transferred substance does not contain  $O_2$ ,  $\omega_{O_2,TW} = 0$ . Hence,

$$B_{mh} = \frac{c_{pm} (T_\infty - T_w) + \Delta h_c \omega_{O_2,\infty} / r_{st}}{q_w / N_w}$$

- ② Now, for a liquid fuel at its **boiling point** (  $T_w = T_{bp}$  ),  $q_w - q_l = N_w (h_{m,TW} - h_{m,TL}) = N_w \lambda_{fu}$ . Hence,

$$B_{mh} = \frac{c_{pm} (T_\infty - T_{bp}) + \Delta h_c \omega_{O_2,\infty} / r_{st}}{\lambda_{fu} + q_L / N_w}$$

- ③ For an atomised tiny liquid droplet, if  $T_T \simeq T_{bp}$ ,  $q_L = 0$

# MT with HT and ACR - L34( $\frac{13}{14}$ )

- 1 When it is difficult to ascertain mass-fractions of compounds in different states, it is preferred to use conserved property  $\eta_\alpha = \sum_k \eta_{\alpha,k} \omega_k$  where  $\eta_{\alpha,k} = M_\alpha / M_k$ .
- 2 Consider CV between  $\infty$ - and T-states. Element balance gives

$$g \eta_{\alpha,\infty} + N_w \eta_{\alpha,T} = (g + N_w) \eta_{\alpha,w} \quad \text{or}$$
$$N_w = g \times B_m \quad \rightarrow \quad B_m = \frac{\eta_{\alpha,\infty} - \eta_{\alpha,w}}{\eta_{\alpha,w} - \eta_{\alpha,T}}$$

- 3 It is usually convenient to combine different  $\eta_\alpha$  into a new conserved property variable  $\Phi$  as was shown in lecture 32 ( slide 18 )



# Summary - L34( $\frac{14}{14}$ )

- 1 Thus, in all cases, we have shown that

$$N_w = g \times B \rightarrow B = \frac{\Psi_\infty - \Psi_w}{\Psi_w - \Psi_T}$$

where  $\Psi = \omega_v, \eta_\alpha, h_m$  and  $\Phi$ .

- 2 Any linear combinations of  $\Psi$  are also solutions
- 3 The result  $N_w = g \times B$  from the Reynolds flow model is different from the result  $N_w = g^* \ln(1 + B)$  for 1D Stefan flow model ( diffusion ) and Couette flow model ( convection ).
- 4 The result from the model correctly identifies  $B_f$  in momentum transfer and shows that  $g = h_{cof, v_w} / c_p$  in single phase Conv HT.
- 5 The Reynolds flow model result will be employed to provide **interface boundary conditions** to the **Boundary Layer flow model** in the next lecture.