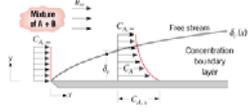


# Introduction to Convection: Mass Transfer

## The Concentration Boundary Layer

- Features
  - A consequence of evaporation or sublimation of species A from a liquid or solid surface across which a second fluid species B is flowing.
  - A region of the flow characterized by species fluxes and concentration gradients.
  - A region between the surface and free stream whose thickness  $\delta_c$  increases in the flow direction.
  - Why does  $\delta_c$  increase in the flow direction?
  - Manifested by a surface species flux  $N_{A,s}^*$  and a convection mass transfer coefficient  $h_m$ .
  - What is the heat transfer analog to Fick's law?



$$\delta_c \rightarrow \frac{C_{A,s} - C_A(y)}{C_{A,s} - C_{A,\infty}} = 0.99$$

$$N_A^* = -D_{AB} \left. \frac{\partial C_A}{\partial y} \right|_{y=0}$$

*Fick's Law*

$$N_{A,s}^* = -D_{AB} \left. \frac{\partial C_A}{\partial y} \right|_{y=0}$$

$$N_{A,s}^* = h_m (C_{A,s} - C_{A,\infty})$$

$$h_m = \frac{-D_{AB} \partial C_A / \partial y|_{y=0}}{C_{A,s} - C_{A,\infty}}$$

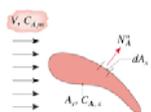
Concentration Boundary (cont.)

- Definitions

Term	Variable	Units
Species molar flux	$N_{A,s}^*$	kmol/s·m <sup>2</sup>
Species molar rate	$N_{A,s}$	kmol/s
Species mass flux	$n_{A,s}^*$	kg/s·m <sup>2</sup>
Species mass rate	$n_{A,s}$	kg/s
Species molar concentration	$C_A$	kmol/m <sup>3</sup>
Species mass concentration (density)	$\rho_A$	kg/m <sup>3</sup>
Species molecular weight	$\mathcal{M}_A$	kg/kmol
Convection mass transfer coefficient	$h_m$	m/s
Binary diffusion coefficient <sup>1</sup>	$D_{AB}$	m <sup>2</sup> /s

<sup>1</sup> Table A.8

- Convection Calculations



**Species Molar Flux:**

$$N_{A,s}^* = h_m (C_{A,s} - C_{A,\infty})$$

**Species Mass Flux:**

$$n_{A,s}^* = \mathcal{M}_A N_{A,s}^* \quad \rho_A = \mathcal{M}_A C_A$$

$$n_{A,s}^* = h_m (\rho_{A,s} - \rho_{A,\infty})$$

**Total Transfer Rates:**

$$N_{A,s} = \bar{h}_m A_s (C_{A,s} - C_{A,\infty})$$

$$n_{A,s} = \bar{h}_m A_s (\rho_{A,s} - \rho_{A,\infty})$$

**Average Mass Transfer Coefficient:**

$$\bar{h}_m = \frac{1}{A_s} \int_{A_s} h_m dA_s$$

Vapor Concentration

### Species Vapor Concentration or Density

- At a Vapor/Liquid or Vapor/Solid Interface

The vapor concentration/density corresponds to saturated conditions at the interface temperature  $T_s$ .

$$C_{A,s} = C_{A,sat}(T_s)$$

$$\rho_{A,s} = \rho_{A,sat}(T_s) = \mathcal{M}_A C_{A,sat}(T_s)$$

Assuming perfect gas behavior, the concentration/density may be estimated from knowledge of the saturation pressure.

$$C_{A,s} \approx \frac{p_{A,sat}(T_s)}{\mathcal{R}T_s} \approx \frac{p_{A,s}}{\mathcal{M}_A}$$

The concentration may also be directly determined from saturation tables. E.g., from Table A.6 for saturated water,

$$\rho_{A,s} = v_g^{-1}(T_s) = \mathcal{M}_A C_{A,s}$$

Vapor Concentration (cont.)

- Free Stream Conditions

- The free stream concentration/density may be determined from knowledge of the vapor pressure,  $p_{A,\infty}$ , assuming perfect gas behavior.

$$C_{A,\infty} \approx \frac{p_{A,\infty}}{\mathcal{R}T_\infty} \approx \frac{p_{A,\infty}}{\mathcal{M}_A}$$

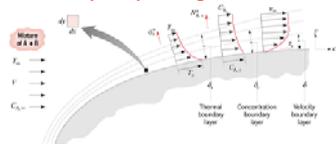
- For water vapor in air, the free stream concentration/density may be determined from knowledge of the relative humidity,  $\phi_\infty$ .

$$\phi_\infty = \frac{p_{A,\infty}}{p_{A,sat}(T_\infty)} \approx \frac{C_{A,\infty}}{C_{A,sat}(T_\infty)} = \frac{\rho_{A,\infty}}{\rho_{A,sat}(T_\infty)}$$

For dry air,

$$\phi_\infty = 0$$

### Species Boundary Layer Equation and Similarity



- Species boundary layer approximation:

$$\frac{\partial C_A}{\partial y} \ll \frac{\partial C_A}{\partial x}$$

- Species equation for a non-reacting boundary layer:

$$u \frac{\partial C_A}{\partial x} + v \frac{\partial C_A}{\partial y} = D_{AB} \frac{\partial^2 C_A}{\partial y^2}$$

What is the physical significance of each term?

Is this equation **analogous** to another boundary layer equation?

- Dimensionless form of the species boundary layer equation:

$$C_A^* \equiv \frac{C_A - C_{A,\infty}}{C_{A,s} - C_{A,\infty}}$$

$$u^* \frac{\partial C_A^*}{\partial x^*} + v^* \frac{\partial C_A^*}{\partial y^*} = \frac{1}{Re_L Sc} \frac{\partial^2 C_A^*}{\partial y^{*2}}$$

$$Sc \equiv \frac{\nu}{D_{AB}} \rightarrow \text{the Schmidt number}$$

How may the Schmidt number be interpreted?

- Functional dependence for a prescribed geometry:

$$C_A^* = f(x^*, y^*, Re_L, Sc)$$

$$h_m = \frac{-D_{AB} \partial C_A / \partial y|_{y=0}}{C_{A,s} - C_{A,\infty}} = -\frac{D_{AB}}{L} \frac{(C_{A,s} - C_{A,\infty}) \partial C_A^*}{(C_{A,s} - C_{A,\infty}) \partial y^*} \Big|_{y^*=0} = \frac{D_{AB}}{L} \frac{\partial C_A^*}{\partial y^*} \Big|_{y^*=0}$$

The dimensionless local convection mass transfer coefficient is then

$$Sh \equiv \frac{h_m L}{D_{AB}} = \frac{\partial C_A^*}{\partial y^*} \Big|_{y^*=0} = f(x^*, Re_L, Sc)$$

$Sh \rightarrow$  local **Sherwood number**

What is the functional dependence of the average Sherwood number?

### Analogies

- Heat and Mass Transfer Analogy**

From **analogous forms** of the dimensionless boundary layer **energy and species equations**, it follows that, for a prescribed geometry and equivalent boundary conditions, the functional dependencies of Nu and Sh are equivalent.

$$Nu = f(x^*, Re, Pr)$$

$$Sh = f(x^*, Re, Sc)$$

Since the Pr and Sc dependence of Nu and Sh, respectively, is typically of the form  $Pr^n$  and  $Sc^n$ , where n is a positive exponent ( $0.30 \leq n \leq 0.40$ ),

$$\frac{Nu}{Pr^n} = f(x^*, Re) = \frac{Sh}{Sc^n}$$

$$\frac{hL/k}{Pr^n} = \frac{h_m L / D_{AB}}{Sc^n}$$

$$\frac{h}{h_m} = \frac{k}{D_{AB}} \frac{Pr^n}{Sc^n} = \frac{k}{D_{AB}} Le^{-n} = \rho c_p Le^{1-n}$$

$$Le = \frac{\alpha}{D_{AB}} \rightarrow \text{the Lewis number}$$

Analogies (cont.)

- Reynolds Analogy**

$$\text{For } dp^*/dx^* = 0$$

$$C_f \frac{Re}{2} = Nu$$

$$\text{and } Pr = Sc = 1,$$

$$Nu = Sh$$

it follows that

$$\frac{C_f}{2} = \frac{Sh}{Re} = St_m$$

where

$$St_m \equiv \frac{Sh}{Re Sc} = \frac{h_m}{\nu} \rightarrow \text{the mass transfer Stanton number}$$

- Modified Reynolds Analogy**

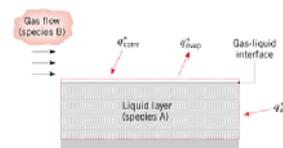
$$\frac{C_f}{2} = St_m Sc^{2/3} \equiv j_m \quad 0.6 < Sc < 300$$

Colburn j factor for mass transfer

- Applicable to laminar flow if  $dp^*/dx^* \sim 0$ .

- Generally applicable to turbulent flow without restriction on  $dp^*/dx^*$ .

### Evaporative Cooling



- The term **evaporative cooling** originates from association of the latent energy created by evaporation at a liquid interface with a reduction in the thermal energy of the liquid. If evaporation occurs in the absence of other energy transfer processes, the thermal energy, and hence the temperature of the liquid, must decrease.

- If the liquid is to be maintained at a fixed temperature, energy loss due to evaporation must be replenished by other means. Assuming convection heat transfer at the interface to provide the only means of energy inflow to the liquid, an energy balance yields

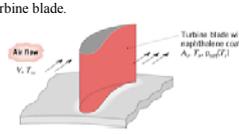
$$q_{conv}^* = q_{evap}^*$$

$$h(T_\infty - T_s) = n''_A h_{fg} = h_m [\rho_{A,sat}(T_s) - \rho_{A,\infty}] h_{fg}$$

$$(T_\infty - T_s) = \underbrace{\left( \frac{h_m}{h} \right)}_{\text{Steady-state Cooling}} \underbrace{\left[ \frac{\rho_{A,sat}(T_s) - \rho_{A,\infty}}{\rho_{A,sat}(T_s) - \rho_{A,\infty}} \right]}_{\text{Obtained from heat/mass transfer analogy}} h_{fg}$$

- With radiation from the interface and heat addition by other means,
 
$$q''_{conv} + q''_{add} = q''_{evap} + q''_{rad}$$

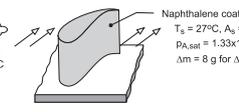
**Problem 6.60:** Use of the **naphthalene sublimation technique** to obtain the average convection heat transfer coefficient for a gas turbine blade.



**KNOWN:** Surface area and temperature of a coated turbine blade. Temperature and pressure of air flow over the blade. Molecular weight ( $M_A = 128.16 \text{ kg/kmol}$ ) and saturation vapor pressure of the naphthalene coating. Duration of air flow and corresponding mass loss of naphthalene due to sublimation.

**FIND:** Average convection heat transfer coefficient.

**SCHEMATIC:**



$T_s = 27^\circ\text{C}$ ,  $A_s = 0.05 \text{ m}^2$   
 $\rho_{A,sat} = 1.33 \times 10^{-4} \text{ bar}$   
 $\Delta m = 8 \text{ g}$  for  $\Delta t = 30 \text{ min}$

Problem: Naphthalene (cont.)

**ASSUMPTIONS:** (1) Applicability of heat and mass transfer analogy, (2) Negligible change in  $A_s$  due to mass loss, (3) Naphthalene vapor behaves as an ideal gas, (4) Solid/vapor equilibrium at surface of coating, (5) Negligible vapor density in freestream of air flow.

**PROPERTIES:** Table A-4, Air ( $T = 300\text{K}$ ):  $\rho = 1.161 \text{ kg/m}^3$ ,  $c_p = 1007 \text{ J/kg}\cdot\text{K}$ ,  $\alpha = 22.5 \times 10^{-6} \text{ m}^2/\text{s}$ . Table A-8, Naphthalene vapor/air ( $T = 300\text{K}$ ):  $D_{AB} = 0.62 \times 10^{-5} \text{ m}^2/\text{s}$ .

**ANALYSIS:** From the rate equation for convection mass transfer, the average convection mass transfer coefficient may be expressed as

$$\bar{h}_m = \frac{n_A}{A_s (\rho_{A,s} - \rho_{A,\infty})} = \frac{\Delta m / \Delta t}{A_s \rho_{A,s}}$$

where

$$\rho_{A,s} = \rho_{A,sat}(T_s) \approx \frac{M_A \rho_{A,sat}}{RT_s} = \frac{(128.16 \text{ kg/kmol})(1.33 \times 10^{-4} \text{ bar})}{(8.314 \text{ J/mol}\cdot\text{K})(300 \text{ K})} = 6.83 \times 10^{-4} \text{ kg/m}^3$$

Hence,

$$\bar{h}_m = \frac{0.008 \text{ kg} / (30 \text{ min} \times 60 \text{ s/min})}{0.05 \text{ m}^2 (6.83 \times 10^{-4} \text{ kg/m}^3)} = 0.13 \text{ m/s}$$

Using the heat and mass transfer analogy with  $n = 1/3$ ,

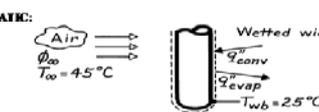
$$\bar{h} = \bar{h}_m \rho \text{Pr}^{1/3} = \bar{h}_m \rho \text{Pr}^{2/3} = 0.130 \text{ m/s} (1.161 \text{ kg/m}^3)^{2/3} \times 1007 \text{ J/kg}\cdot\text{K} (22.5 \times 10^{-6} / 0.62 \times 10^{-5})^{2/3} = 359 \text{ W/m}^2\cdot\text{K}$$

**COMMENTS:** The naphthalene sublimation technique is used extensively to determine convection coefficients associated with complex flows and geometries.

Problem: Wet-bulb Thermometer

**Problem 6.74:** Use of wet and dry bulb temperature measurements to determine the relative humidity of an air stream.

**SCHEMATIC:**



**ASSUMPTIONS:** (1) Perfect gas behavior for vapor, (2) Steady-state conditions, (3) Negligible radiation, (4) Negligible conduction along thermometer.

**PROPERTIES:** Table A-4, Air (308K, 1 atm):  $\rho = 1.135 \text{ kg/m}^3$ ,  $c_p = 1007 \text{ J/kg}\cdot\text{K}$ ,  $\alpha = 23.7 \times 10^{-6} \text{ m}^2/\text{s}$ ;

Table A-6, Saturated water vapor (298K):  $v_g = 44.25 \text{ m}^3/\text{kg}$ ,  $h_{fg} = 2443 \text{ kJ/kg}$  (318K);  $v_g = 15.52 \text{ m}^3/\text{kg}$ ;

Table A-8, Air-vapor (1 atm, 298K):  $D_{AB} = 0.26 \times 10^{-4} \text{ m}^2/\text{s}$ ,  $D_{AB} (308\text{K}) = 0.26 \times 10^{-4} \text{ m}^2/\text{s} \times (308/298)^{3/2} = 0.27 \times 10^{-4} \text{ m}^2/\text{s}$ ,  $Le = \alpha/D_{AB} = 0.88$ .

Problem: Wet-bulb Thermometer

**ANALYSIS:** Dividing the energy balance on the wick by  $\rho_{A,sat}(T_w)$ ,

$$\frac{T_\infty - T_s}{\rho_{A,sat}(T_s)} = h_{fg} \left[ \frac{\bar{h}_m}{h} \right] \left[ \frac{\rho_{A,sat}(T_s)}{\rho_{A,sat}(T_\infty)} - \frac{\rho_{A,\infty}}{\rho_{A,sat}(T_\infty)} \right]$$

With  $[\rho_{A,\infty} / \rho_{A,sat}(T_\infty)] \approx \phi_\infty$  for a perfect gas and  $h_m$  obtained from the heat and mass transfer analogy,

$$\phi_\infty = \frac{\rho_{A,sat}(T_s)}{\rho_{A,sat}(T_\infty)} - \frac{\rho c_p}{Le^{2/3} \rho_{A,sat}(T_\infty) h_{fg}} (T_\infty - T_s)$$

where

$$\frac{\rho_{A,sat}(T_s)}{\rho_{A,sat}(T_\infty)} = \frac{v_g(T_\infty)}{v_g(T_s)} = \frac{15.52}{44.25} = 0.351$$

$$\rho_{A,sat}(T_\infty) = (15.52 \text{ m}^3/\text{kg})^{-1} = 0.064 \text{ kg/m}^3$$

Hence,

$$\phi_\infty = 0.351 - \frac{1.135 \text{ kg/m}^3 (1007 \text{ J/kg}\cdot\text{K})}{(0.88)^{2/3} (0.064 \text{ kg/m}^3) (2.443 \times 10^6 \text{ J/kg})} (45 - 25) \text{ K}$$

$$\phi_\infty = 0.351 - 0.159 = 0.192$$