Transport phenomena

3. Forced convection



3.0 Coupling flow with heat/mass exchange



Simple configuration: U uniform



3.1 Cooling flow over a plate



incompressible flow:
$$\nabla \cdot \mathbf{u} = 0 = \frac{\partial u_x}{\partial x} + \frac{\partial u_z}{\partial z} \Rightarrow \frac{U_\infty}{x} \sim \frac{u_z}{\delta_v}$$
$$u_z \sim \frac{\delta_v}{x} U_\infty \ll U_\infty \quad \text{for } x \gg \delta_v$$

velocity profile inside the boundary layer:

$$u_x(x,z) \sim \frac{U_\infty}{\delta_v(x)} z$$

3.1.1 Thermal boundary layer (hypothesis: $\delta_T < \delta_v$)



$$\Rightarrow \delta_T(x) \sim \alpha^{1/3} \nu^{1/6} \left(\frac{x}{U_\infty}\right)^{1/2} \sim \delta_v(x) \left(\frac{\alpha}{\nu}\right)^{1/3} = \delta_v(x) \operatorname{Pr}^{-1/3}$$

Prandtl number

$$\Pr = \frac{\nu}{\alpha}$$

water ~ 7 air ~ 1 Hg ~ $0.02 \Rightarrow$ different behavior glycerin ~ 1000

$$\delta_T < \delta_v~~$$
 ~ OK, if $\Pr > 0.6$

Schmidt number for mass transport

$$Sc = \frac{\nu}{D}$$

3.1.2 Heat transport across the thermal boundary layer



$$\begin{array}{ll} \mbox{Heat flux} & J_T = -\kappa \, \frac{\partial T}{\partial z} \sim \kappa \, \frac{T_\infty - T_s}{\delta_T(x)} \\ \mbox{Average flux} & < J_T > \sim -\frac{1}{L} \int_0^L \kappa (T_\infty - T_s) \frac{U_\infty^{1/2}}{\alpha^{1/3} \nu^{1/6} x^{1/2}} \, dx \end{array}$$

$$\langle J_T \rangle \sim -\kappa \frac{T_{\infty} - T_S}{L} \left(\frac{U_{\infty}L}{\nu}\right)^{1/2} \left(\frac{\nu}{\alpha}\right)^{1/3} = \kappa \frac{T_{\infty} - T_S}{L} \operatorname{Re}^{1/2} \operatorname{Pr}^{1/3}$$

Heat transfer coefficient
$$J_T = -h_T(T_\infty - T_s)$$
 $h_T \sim \frac{\kappa}{\delta_T}$

Nusselt number

$$\mathrm{Nu} = \frac{h_T L}{\kappa} \sim \frac{L}{\delta_T}$$

Forced convection over a plate $Nu \simeq 0.33 \, Re^{1/2} Pr^{1/3}$

Other expressions for different geometries Nu = f(Re, Pr)

Same formalism for mass transport:

Mass transfer coefficient

$$J_m = -h_m(c_\infty - c_s)$$

$$h_m \sim \frac{\kappa}{\delta_c}$$

Sherwood number

$$\mathrm{Sh} = \frac{h_m L}{D}$$

3.2 Practical exemples3.2.1 Drying a sheet in the wind



 $U_{\infty}, r_h, T_{\infty}$

Simplified model

- uniform temperature
- radiative exchanges neglected
- P_{H_2O} at the surface = P_{sat}

relative humidity r_h

$$P_{H_2O_{\infty}} = r_h P_{sat}$$



Vapor pressure



source: dimclim.fr

$$P_{H_2O} \leftrightarrow c_{H_2O}? \qquad PV = nRT \Rightarrow c_{H_2O} = \frac{P_{H_2O}}{RT}$$

(mol.m⁻³)

H₂O molar flux:
$$J_{H_2O} = h c_{sat} (1 - r_h)$$

$$\begin{pmatrix} D \\ L \\ l \end{pmatrix} Sh \sim \frac{D}{L} \left(\frac{U_{\infty}L}{\nu} \right)^{1/2} \left(\frac{\nu}{D} \right)^{1/3}$$
fine to call it "Nu"!

$$J_{H_2O} \sim \frac{D}{L} \left(\frac{U_{\infty}L}{\nu}\right)^{1/2} \left(\frac{\nu}{D}\right)^{1/3} c_{sat}(1-r_h) \quad \text{[mol.m-2.s-1]}$$

if r_h fixed, $T \nearrow \Rightarrow c_{sat} \nearrow \Rightarrow$ higher evaporation

mass flux: $J_{m_{H_2O}} = M_{H_2O} J_{H_2O}$ [kg.m⁻².s⁻¹] volumic flux: $J_{v_{H_2O}} = \frac{1}{\rho_{H_2O}} J_{m_{H_2O}}$ [m.s⁻¹] Numerical estimate: L = 1m $U_{\infty} \sim 5 m/s$

$$\begin{split} D_{H_2O/air} &= 2,6.10^{-5} \,\mathrm{m^{2}/s} \quad \nu_{air} = 1,5.10^{-5} \,\mathrm{m^{2}/s} \\ P_{sat} &= 2.3 \,\mathrm{kPa} \; \Rightarrow \; c_{sat} = 0.9 \,\mathrm{mol/m^{3}} \\ J_{H_2O} &\sim 0.03 \,(1 - r_h) \; [\mathrm{mol.m^{-2}.s^{-1}}] \\ & \text{(with prefactor } 0.33 \;\mathrm{for} \; Sh \,) \\ J_{m_{H_2O}} &\sim 7.10^{-5} (1 - r_h) \; [\mathrm{kg.m^{-2}.s^{-1}}] \\ & J_{v_{H_2O}} &\sim 0.07 \,(1 - r_h) \; [\mathrm{\mu m/s}] \end{split}$$

to evaporate 0.5 kg/m² of water (i.e. 0.5 mm) with r_h = 50%

$$\tau_{evap} \sim \frac{0.5 \cdot 10^{-3}}{2 \times 0.07 \cdot 10^{-6} \times 0.5} \sim 2 \text{h}$$
 (2 faces

probably underestimated: evaporation \Rightarrow cooling down \Rightarrow $c_{sat} \searrow \Rightarrow J_{H_2O} \searrow$

3.2.2 Clay pot cooling



evaporation \Rightarrow latent heat

heat flux at the surface

$$J_{latent} = L_{vap} \, J_{m_{H_2O}}$$

$$J_{latent} \sim 150 \left(1-r_h\right)$$
W/m²

standard fridge ~ 30 to 50 W

3.2.3 Thermal balance



heat flux from the fluid to the surface

$$J_T \sim \frac{\kappa}{L} \left(\frac{U_{\infty}L}{\nu}\right)^{1/2} \left(\frac{\nu}{\alpha}\right)^{1/3} \left(T_{\infty} - T_s\right)$$

heat flux from latent heat

$$J_{latent} \sim L_{vap} M_{H_2O} \frac{D}{L} \left(\frac{U_{\infty}L}{\nu}\right)^{1/2} \left(\frac{\nu}{\alpha}\right)^{1/3} c_{sat}(1-r_h)$$

$$\Delta T \sim L_{vap} M_{H_2O} \frac{D}{\kappa} \left(\frac{\alpha}{D}\right)^{1/3} c_{sat} (1 - r_h) \quad \text{independent of } U_{\infty}$$

Measuring relative humidity

Numerical estimate: $\kappa_{air} = 0.03 \text{ J.m}^{-1}.\text{K}^{-1}$ $\alpha_{air} = 1.9 \text{ 10}^{-5} \text{ m}^{2}/\text{s}$



3.3 Bubbles in a champagne bottle

1294

Langmuir 2002, 18, 1294-1301

Source:

Kinetics of Gas Discharging in a Glass of Champagne: The Role of Nucleation Sites

Gérard Liger-Belair,*,[†] Michèle Vignes-Adler,[‡] Cédric Voisin,[†] Bertrand Robillard,[§] and Philippe Jeandet[†]



3.3.1 Bubbles nucleation

$$\begin{array}{c} P_{atm} \\ \hline P_{atm} \\ \hline P_{cO_2} = P_b \\ cCO_2 \end{array} = P_{atm} + 2\gamma/R \\ bubble \text{ grows if } c_{CO_2} > c_{sat}^{P_b} = c_{sat}^{P_{atm}} \frac{P_b}{P_{atm}} \\ \hline e: \quad \frac{P_{CO_2}^{ini}}{P_{atm}} c_{sat}^{P_{atm}} > \left(1 + \frac{2\gamma}{RP_{atm}}\right) c_{sat}^{P_{atm}} \\ \hline R > \frac{2\gamma}{P_{CO_2}^{ini} - P_{atm}} \\ \hline \end{array}$$

 \Rightarrow heterogenous nucleation

Heterogenous nucleation

 $R_{init} \gg R_{min}$



100 µm

3.3.2 Growth rate



Diffusion process:

$$\delta_c \sim (Dt)^{1/2}$$

$$J_{CO_2} \sim \frac{D}{\delta_c} \left(c_{CO_2}^{ini} - c_{sat}^{P_{atm}} \right)$$

[molecules/m³] in article

$$\frac{d\mathcal{V}}{dt} = 4\pi R^2 \frac{dR}{dt} = J_{CO_2} 4\pi R^2 \frac{1}{\mathcal{N}_a} \frac{RT}{P}$$
$$\frac{dR}{\zeta} \sim \left(\frac{D}{t}\right)^{1/2} \left(c_{CO_2}^{ini} - c_{sat}^{P_{atm}}\right) \frac{1}{\mathcal{N}_a} \frac{RT}{P}$$
volume 1 mole of gas
$$R(t) - R_{ini} \sim \left(c_{CO_2}^{ini} - c_{sat}^{P_{atm}}\right) \frac{1}{\mathcal{N}_a} \frac{RT}{P} (Dt)^{1/2}$$

numerical estimate:

 $c_{CO_2}^{ini} \simeq 1,6.10^{26} \text{ molecules/m}^3$ $c_{sat}^{P_{atm}} \simeq 2,7.10^{25} \text{ molecules/m}^3$ $D_{CO_2} \simeq 1,4.10^{-9} \text{ m}^2/\text{s}$ $R_{detach} \sim 20 \text{ }\mu\text{m} \Rightarrow \tau_{growth} \sim 10 \text{ }\text{ms}$ a bit overestimated: $\tau_{exp} \sim 40 \text{ }\text{ms}$

3.3.2 Rising velocity



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Figure 5. Velocity of ascending champagne bubbles $U(R)(\bigcirc)$, compared with the Stokes velocity (—). *Re* and *Pe* are respectively the Reynolds and Peclet numbers associated with a rising bubble.

3.3.2 Rising velocity





Figure 5. Velocity of ascending champagne bubbles $U(R)(\bigcirc)$, compared with the Stokes velocity (-). *Re* and *Pe* are respectively the Reynolds and Peclet numbers associated with a rising bubble.

Limit low Reynolds number (~OK at the beginning)

$$\frac{4}{3}R^3\Delta\rho g = 6\pi\eta R U$$

$$U\sim \frac{\rho_{liq}gR^2}{\eta}$$

3.3.3 Bubble growth



Mass transfer coefficient?

Flow low *Re* ≠ boundary layer over a plate

in article:

 $\mathrm{Sh} \sim \mathrm{Pe}^{1/3} = \left(\frac{UR}{D}\right)^{1/3} = \left(\frac{UR}{\nu}\right)^{1/3} \left(\frac{\nu}{D}\right)^{1/3}$

 $\mathrm{Sh}\sim\mathrm{Re}^{1/3}\mathrm{Sc}^{1/3}$

for heat exchange, this would be: $\mathrm{Nu}\sim\mathrm{Re}^{1/3}\mathrm{Pr}^{1/3}$

Can we justify this form?



with
$$\delta_T \ll R$$
 $\frac{U}{R} \delta_T \frac{T}{R} \sim \alpha \frac{T}{\delta_T^2}$
 $\delta_T \sim \left(\frac{\alpha R^2}{U}\right)^{1/3} \Rightarrow \operatorname{Nu} \equiv \frac{R}{\delta_T} \sim \left(\frac{UR}{\alpha}\right)^{1/3}$
for mass transport: $\operatorname{Sh} \equiv \frac{R}{\delta_c} \sim \left(\frac{UR}{D}\right)^{1/3}$

3.3.3 Coupling rising dynamics and mass transport

$$J_{CO_2} \sim h_c \left(c_{CO_2}^{ini} - c_{sat}^{P_{atm}} \right) = \frac{D}{R} \operatorname{Sh} \left(c_{CO_2}^{ini} - c_{sat}^{P_{atm}} \right)$$
$$J_{CO_2} \sim \frac{D}{R} \left(\frac{UR}{D} \right)^{1/3} \left(c_{CO_2}^{ini} - c_{sat}^{P_{atm}} \right)$$
$$\frac{d\mathcal{V}}{dt} = 4\pi R^2 \frac{dR}{dt} = J_{CO_2} 4\pi R^2 \frac{1}{\mathcal{N}_a} \frac{RT}{P}$$
$$\frac{dR}{dt} \sim \frac{D}{R} \left(\frac{\rho g R^3}{\eta D} \right)^{1/3} \left(c_{CO_2}^{init} - c_{sat}^{P_{atm}} \right) \frac{1}{\mathcal{N}_a} \frac{RT}{P}$$
$$\frac{dR}{dt} \sim \left(\frac{D^2 g}{\nu} \right)^{1/3} \left(c_{CO_2}^{init} - c_{sat}^{P_{atm}} \right) \frac{1}{\mathcal{N}_a} \frac{RT}{P}$$

numerical estimate:

 $rac{dR}{dt}$ ~ 1.5 mm/s experiment ~ 0.2 mm/s (but after pouring in a glass \Rightarrow $c_{CO_2}^{init}$)



Figure 6. Bubble radius increase vs time for a bubble rising toward the liquid surface. Two typical bubble trains at different steps of gas discharging are considered.