Heat Exchanges/Theory

The double-pipe heat exchanger



cocurrent

Call the hot stream h and the cold stream c and determine all coefficients w.r.t the outside diameter of inside pipe.

Usually, we need to figure out the required heat transfer area for a given job, given a flowrate and a temperature difference.



Performing an energy balance on each stream separately, we get

$$2\pi r_0 dz$$

$$\uparrow$$

$$dq = w_c C p_c dt_c = -w_h C p_h dt_h = U_0 |_z (t_h - t_c) |_z dA_0 |_z$$

$$\uparrow \qquad \uparrow \qquad \uparrow$$
temp & position temp & temp & position position position

Usually, w_c , Cp_c , w_h , Cp_h , U_0 are temperature and position dependent.

To simplify, assume mass flow, and Cp_c , are independent of temperature hence, position. Then integrate inside equations.

$$w_{c} Cp_{c} (T_{c,2} - T_{c,1}) = -w_{h} Cp_{h} (T_{h,2} - T_{h,1}) = Q$$

$$w_{c} C p_{c} dt_{c} = U_{0} (t_{h} - t_{c}) dA_{0}$$

$$-w_{h} C p_{c} dt_{h} = U_{0} (t_{h} - t_{c}) dA_{0}$$

$$\rightarrow \frac{dt_{c}}{t_{h} - t_{c}} = \frac{U_{0} dA_{0}}{w_{c} C p_{c}}$$

$$\rightarrow -\frac{dt_{h}}{t_{h} - t_{c}} = \frac{U_{0} dA_{0}}{w_{h} C p_{h}}$$

if we add these two

$$-\left(\frac{d(t_h-t_c)}{(t_h-t_c)}\right) = U_0 dA_0 \left(\frac{1}{w_c C p_c} + \frac{1}{w_h C p_h}\right)$$

This equation relates the Δt 's to position for U_0 independent of position, integrate from inlet (1) to outlet (2)

$$-\ell n \left(\frac{\left(t_{h,2} - t_{c,2} \right)}{\left(t_{h,1} - t_{c,1} \right)} \right) = U_0 A_0 \left(\frac{1}{w_c C p_c} + \frac{1}{w_h C p_h} \right)$$

$$\frac{1}{w_h C p_h} = \frac{t_{h,2} - t_{h,1}}{-Q}$$
$$\frac{1}{w_c C p_c} = \frac{t_{c,2} - t_{c,1}}{Q}$$

Hence

and

$$\frac{-\ell n \left(\frac{\left(t_{h,2} - t_{c,2} \right)}{\left(t_{h,1} - t_{c,1} \right)} \right) = \frac{U_0 A_0}{Q} \left(\left(t_{c,2} - t_{c,2} \right) - \left(t_{h,1} - t_{h,1} \right) \right) \\ = -\frac{U_0 A_0}{Q} \left(\left(t_{h,2} - t_{c,1} \right) - \left(t_{h,2} - t_{c,1} \right) \right)$$

13 - 2

$$U_{0} A_{0} \frac{\left(t_{h,2} - t_{c,2}\right) - \left(t_{h,1} - t_{c,1}\right)}{\ell n \frac{\left(t_{h,2} - t_{c,2}\right)}{\left(t_{h,1} - t_{c,1}\right)}} = Q = U_{0} A_{0} \Delta t_{\ell m}$$
$$\Delta t_{\ell m} = \frac{\left(t_{h,2} - t_{c,2}\right) - \left(t_{h,1} - t_{c,1}\right)}{\ell n \frac{\left(t_{h,2} - t_{c,2}\right)}{\left(t_{h,1} - t_{c,1}\right)}}$$

Linear Temperature Dependence of U_0 on Temperature Let $U_0 = a + b \Delta t$

and insert this in the balance equation

$$w_{c} Cp_{c} dt_{c} = U_{0} (t_{h} - t_{c}) dA_{0}$$

$$-w_{h} Cp_{c} dt_{h} = U_{0} (t_{h} - t_{c}) dA_{0}$$

$$\frac{dt_{c}}{t_{h} - t_{c}} = \frac{U_{0}}{w_{c} Cp_{c}} dA_{0}$$

$$-\frac{dt_{c}}{t_{h} - t_{c}} = \frac{U_{0}}{w_{h} Cp_{h}} dA_{0}$$

$$\frac{d(t_{h} - t_{c})}{(t_{h} - t_{c})} = U_{0} dA_{0} \left(\frac{1}{w_{c} Cp_{c}} + \frac{1}{w_{h} Cp_{h}}\right)$$

$$\frac{d(t_{h} - t_{c})}{U_{0} (t_{h} - t_{c})} = dA_{0} \left(\frac{1}{w_{c} Cp_{c}} + \frac{1}{w_{h} Cp_{h}}\right)$$

$$\frac{\Delta t_{2} - \Delta t_{1}}{Q}$$
Let
$$U_{0} = a + b(t_{h} - t_{c})$$

$$t_{h} - t_{c} = \Delta t$$

$$\int_{\Delta t_1}^{\Delta t_2} \frac{d(\Delta t)}{(a+b\Delta t)\Delta t} = \frac{\Delta t_2 - \Delta t_1}{Q} A_0$$

$$\begin{bmatrix} \frac{1}{a} \end{bmatrix} \ell n \frac{\Delta t}{a+b\Delta t} \Big|_{\Delta t_1}^{\Delta t_2} = \begin{bmatrix} \frac{1}{a} \end{bmatrix} \ell n \frac{\Delta t}{U_0} \Big|_{\Delta t_1}^{\Delta t_2}$$

$$-\frac{1}{a} \ell n \frac{(U)_1(\Delta t)_2}{(U)_2(\Delta t)_1} \text{ and}$$

$$\frac{1}{a} \ell n \frac{U_{0_1} \Delta t_2}{U_{0_2} \Delta t_1} = \frac{\Delta t_2 - \Delta t_1}{Q} A_0$$

in which,
$$U_{0_1} = a + b(t_h - t_c)_1; U_{0_2} = a + b(t_h - t_c)_2$$

= $a + b \Delta t_1$; $U_{0_2} = a + b \Delta t_2$

13 - 3

$$U_{0_{1}}\Delta t_{2} = a\Delta t_{2} + b\Delta t_{1}\Delta t_{2}$$

$$U_{0_{2}}\Delta t_{1} = a\Delta t_{1} + b\Delta t_{1}\Delta t_{2}$$
subtract
$$U_{0_{1}}\Delta t_{2} - U_{0_{2}}\Delta t_{1} = a(\Delta t_{2} + \Delta t_{1})$$

$$\frac{U_{0_{1}}\Delta t_{2} - U_{0_{2}}\Delta t_{1}}{\Delta t_{2} + \Delta t_{1}} = a$$

$$\frac{\Delta t_2 - \Delta t_1}{\tilde{U}_{0_1} \Delta t_2 - U_{0_2} \Delta t_1} \ell n \frac{U_{0_1} \Delta t_2}{U_{0_2} \Delta t_1} = \frac{\Delta t_2 - \Delta t_1}{Q} A_0$$

$$Q = \frac{U_{0_1} \Delta t_2 - U_{0_2} \Delta t_1}{\ell n \frac{U_{0_1} \Delta t_2}{U_{0_2} \Delta t_1}} A_0$$



shown -1 shell pass -1 tube concurrent heat exchanger

Benefits – cost, space, pressure drop

consists of bundle of tubes through which one fluid passes the other fluid flows through the shell. Baffles are used to direct the flow of one shell fluid to improve heat transfer.

- 1) If one fluid fouls it should be routed through tubes.
- 2) If one fluid is at high pressure—should go through tubes to avoid cost of high pressure

shell.

- 3) Corrosive material should go through tubes to avoid costs associated with shell construction.
- 4) More viscous fluid should be directed through shell ΔP is low.
- 5) If major resist:lite is in tube side, a 2 tube pass, one shell pass may be better.

In the 1-D analysis we get a similar result to the double-pipe exchanger

$$Q = U_0 A_0 \left[\frac{\left(t_{2,i} - t_{2,0} \right) - \left(t_{1,i} - t_{i,0} \right)}{\ell n \frac{t_{2,i} - t_{2,0}}{t_{1,i} - t_{1,0}}} \right]$$

where A_0 is the total outside area of all the tubes, and U_0 is the overall heat transfer coefficient.

How to evaluate U_0 ? — The tube side fluid is no problem, but to evaluate shell side.

Heat transfer coefficient for flow past tube bundles



 s_t — transverse pitch

 s_{ℓ} — longitudinal pitch

if $\gtrsim 10$ rows of tubes in bundle and

 $\gtrsim 10$ tubes in each row, neglect entrance, end effects

and

$$Nu = F \left(\text{Re}, \text{Pr}, \mu_b / \mu_0, S_\ell, S_t \right)$$
$$S_\ell = s_\ell / D \qquad S_t = s_t / D$$

experimental correlations show that

$$Nu = \left(0.4 N_{\rm Re}^{1/2} + 0.2 N_{\rm Re}^{2/3}\right) N_{\rm Pr}^{0.4} \left(\frac{\mu_b}{\mu_0}\right)^{.14}$$

$$N_{\text{Re}} = \frac{D_p G}{\mu_b (1 - \varepsilon)}$$

$$\Pr = \frac{Cp}{\mu_b k_b}$$

$$N_{Nu} = \frac{h D_p}{k_b} \left(\frac{\varepsilon}{(1 - \varepsilon)}\right)$$
range
$$N_{\text{Re}} = 1 - 40,000 \qquad \frac{\mu_b}{\mu_0} \sim .1 - 5$$

$$\Pr \sim .7 - 800 \qquad \varepsilon \sim .42 - .65$$

 ε — void fraction = $\frac{\text{volume of tubes}}{\text{total shell volume}}$

$$a_v - \frac{\text{area for heat transfer}}{\text{unit volume}}$$

 D_p - hydraulic radius= 3/2 D tube

To evaluate h_{outside} using this tube bundle correlation, we need to estimate the average mass flow rate *G*.

If the diameter of the shell is D_s and the distance between baffles is L_b , then an average mass velocity is

$$G_{av} = \frac{4\,\dot{m}}{\pi\,D_s L_b}$$

 \dot{n} — mass flow rate of shell fluid and the Reynolds number is

$$\operatorname{Re} = \frac{D_p G_{av}}{\mu (1 - \varepsilon)} = \frac{4 D_p \dot{m}}{\pi \mu D_s L_b (1 - \varepsilon)}$$

 ε — void fraction and in terms of parameters

$$\operatorname{Re} = \frac{6D \,\dot{m}}{\pi \mu \, D_s L_b \left(1 - \varepsilon\right)}$$
$$Nu = \left(.4 \, N^{1/2} + .2 \, N^{2/3}\right) \operatorname{Pr}^4 \left(\frac{\mu_b}{\mu_0}\right)^{.14}$$

Fouling of heat transfer surfaces:

One of the most difficult aspects of heat exchange design is the reduction in heat transfer rates due to deposition of "scale"—insoluble inorganic compounds—on the tube wall. The magnitude depends on the nature and thickness. It is usually reported in terms of a film heat transfer coefficient for the scale which is referred to as the fouling factor

$$h_s = \{ \text{film heat transfer for scale} \}$$

$$\frac{1}{h_s}$$
 = fouling factor

to determine U_0 , we assume all the resistance add in series:

$$\frac{1}{U_0} = \frac{1}{h_I} = \frac{D_0}{D_i} + \frac{1}{h_{s,i}} \left(\frac{D_0}{D_i}\right) + \frac{D_0}{2k} \quad \ln \frac{D_0}{D_i} + \frac{1}{h_{s,0}} + \frac{1}{h_{II}}$$

As the scale increases, the overall U_0 decreases, leading to poor performance and cleaning becomes necessary.



Observations

- 1. wire temperature rises smoothly to $\sim 300^{\circ}$
- 2. temperature jumps to $\sim 1800^{\circ}$
- 3. decrease heat flux to °570 wire temp at a much lower heat flux
- 4. jumps back down in temperature

Explanation

Different types of boiling

Nucleate Boiling	Transition	Film Boiling
high heat transfer rate	intermediate unstable	low heat transfer rate

Nucleate boiling occurs directly at solid surface

Need to nucleate bubbles of gas somehow, but interface costs energy.

 $\underbrace{P_{s}}_{P_{A}} \leftarrow \text{surface tension} \quad A \text{ pressure drop exists across any interface} \\ \text{due to surface tension.}$

$$\Delta P = 2H\sigma = \frac{2\sigma}{R}$$

$$H =$$
 mean curvature of interface $=\frac{1}{2} \left(\frac{1}{R_1} + \frac{1}{R_2}\right)$

capillary rise in a tube: $\rho_{\ell}gh - \rho_{r}gh = \frac{2\sigma}{R}$



Note that as $R \to 0$, $\Delta P \to \infty$ and to achieve the excess pressure, we need to achieve superheat.

extreme

\rightarrow case	homogeneous nucleation — bubbles of gas form spontaneously by molecular aggregation
$\stackrel{\rm most}{\rightarrow}$	heterogeneous nucleation — bubbles form at existing gas pockets, dust, dirt, etc.
common	

Surface irregularities also are heterogeneous nuclei

Film Boiling

Heat flux causes sufficient boiling that a vapor film covers surface, slows down heat transfer.

$$k_{water} = .35 \frac{Btu}{hr \text{ ft}^2 \, ^\circ F}$$
$$k_{vapor} = .0183 \frac{Btu}{hr \text{ ft}^2 \, ^\circ F}$$

at least a factor of 20!



Boiling Summary

- 1) Heat flux is highest when change of phase occurs
- 2) Nucleate Boiling highest rates, bubbles form directly at surface
- 3) Film Boiling intermediate rates, bubbles form at vapor-liquid interface

Condensation

Condensation is used to heat solids in the same way boiling is often used to cool solids.

Similarly also, these are two types of condensation:

1) Dropwise \leftrightarrow Nucleate

2) Film \leftrightarrow Film

- 1) Dropwise —condensate doesn't wet surface. forms drops
- 2) Film —condensate wets surface, forms liquid film