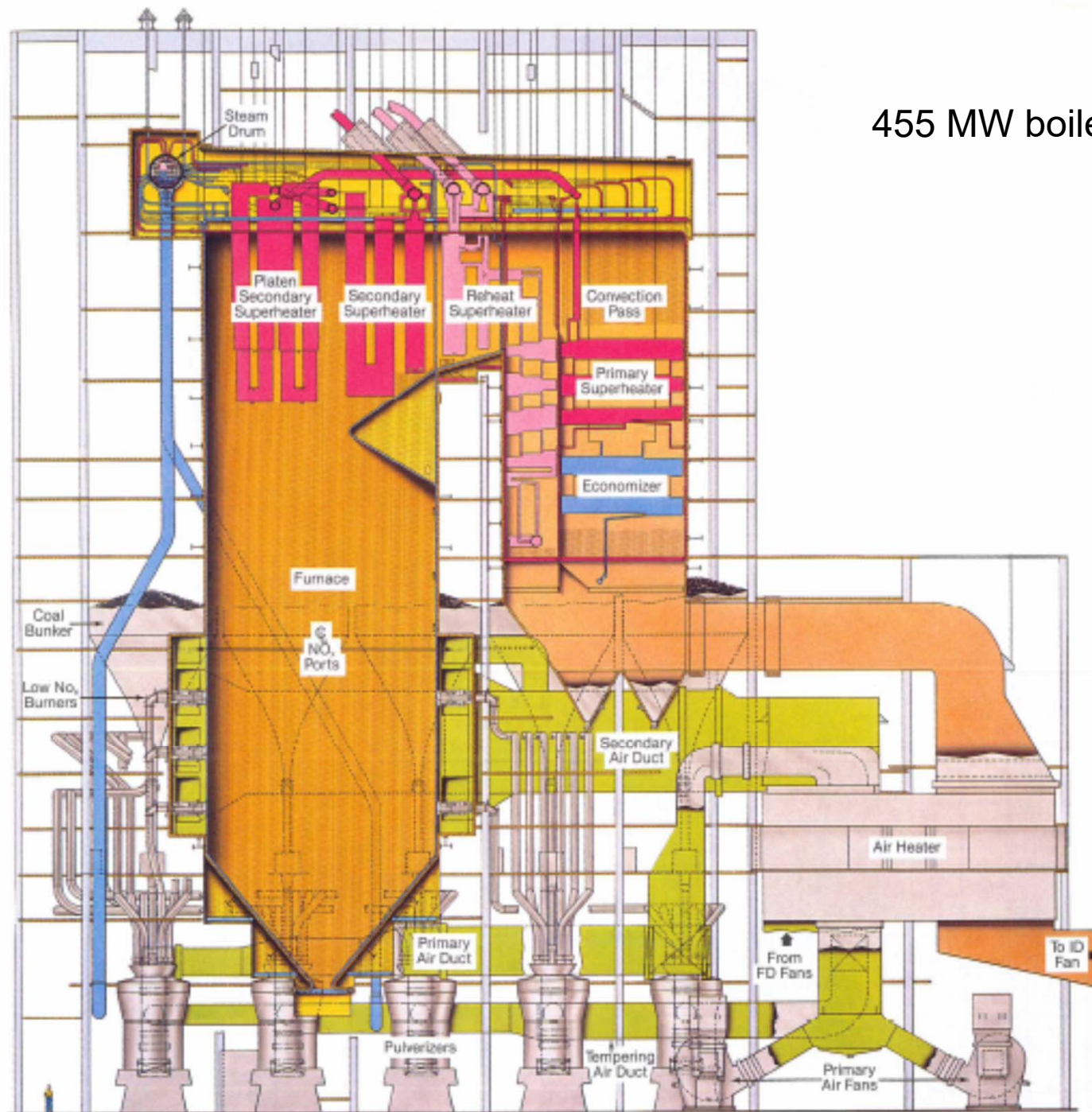


From Steam, Babcock & Wilcox, 1992, Plate 1.



455 MW boiler





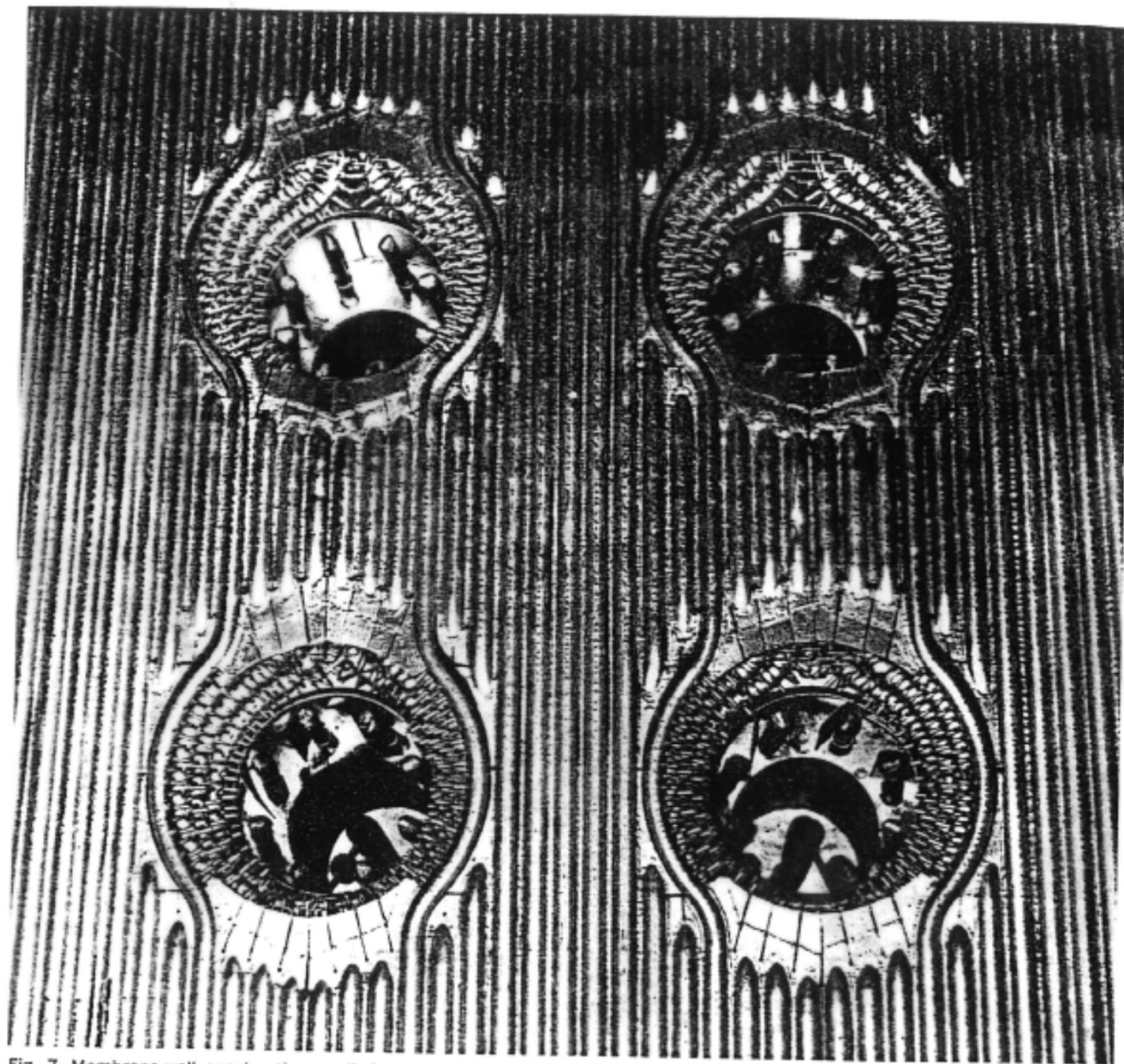
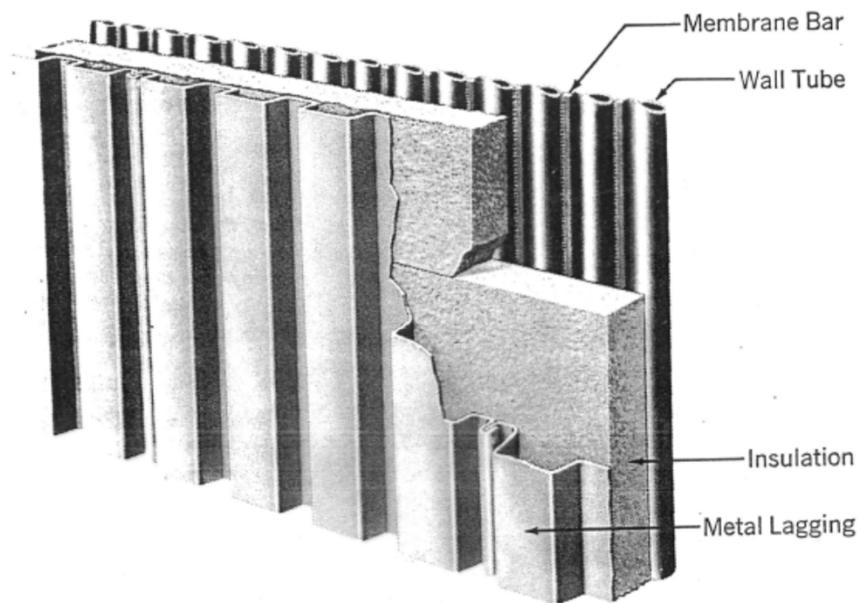
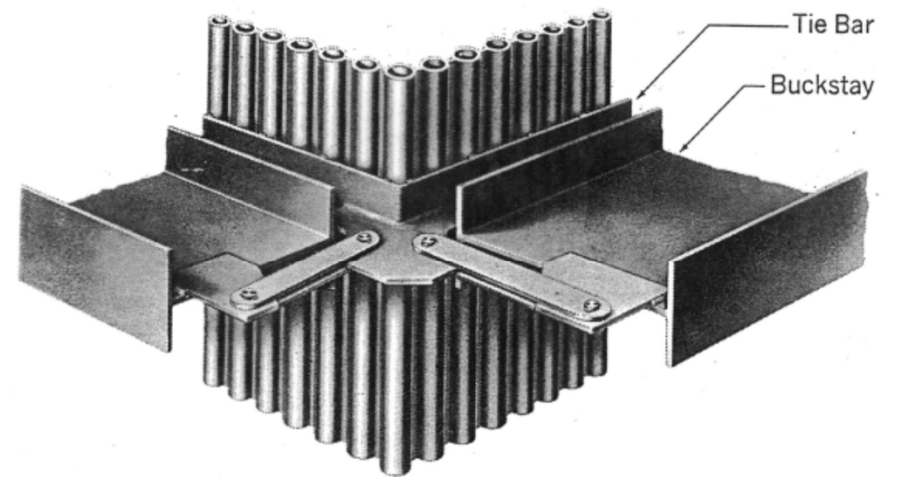


Fig. 7 Membrane-wall construction applied to a burner wall.



**Fig. 2** Membrane wall construction.



**Fig. 6** Tie bar and buckstay arrangement at corner of furnace.

# Heat Exchanger (6 ft diameter)



# High Pressure Heat exchanger



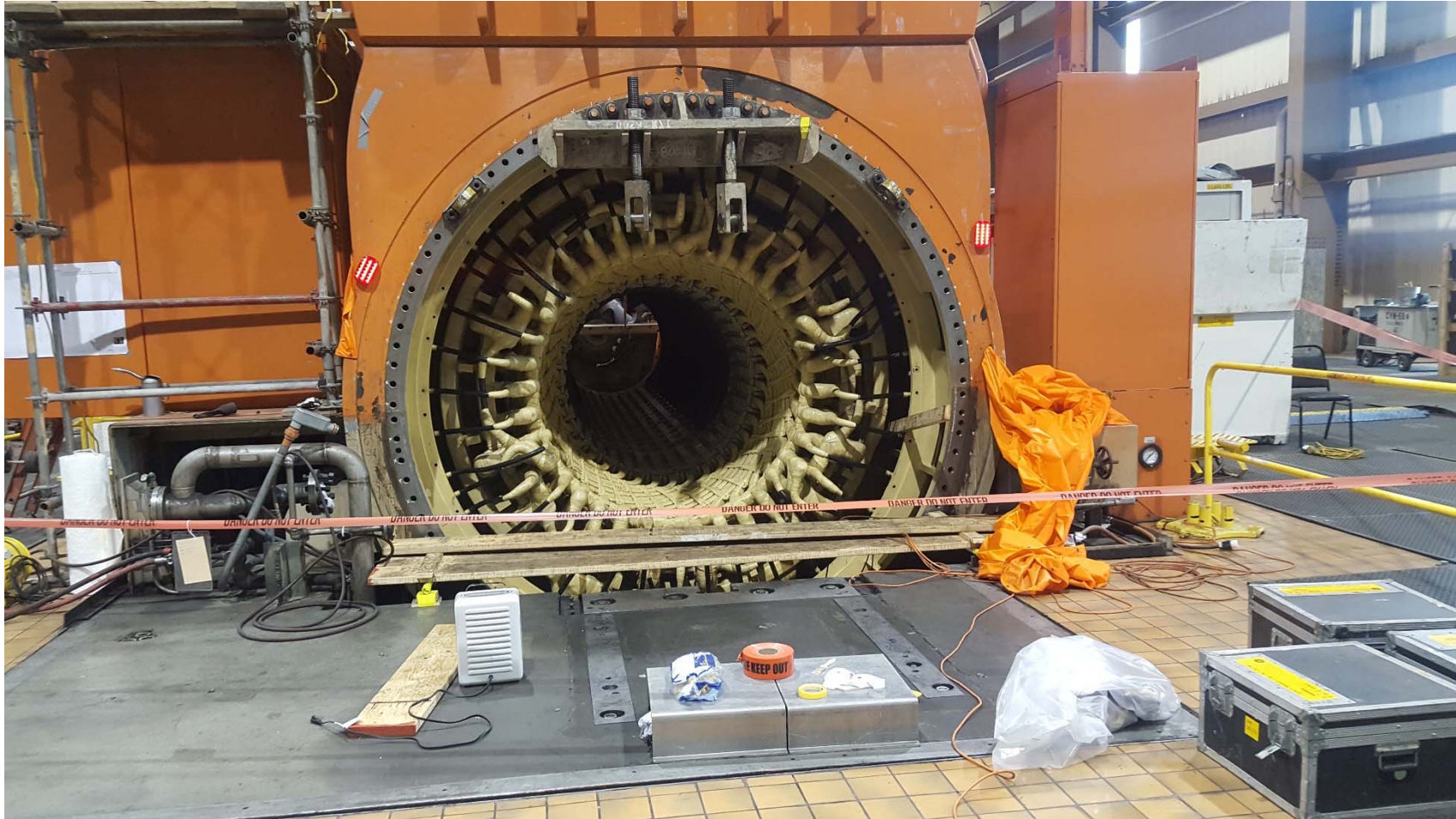


# Turbine





# Generator



# Spinning Part of Generator





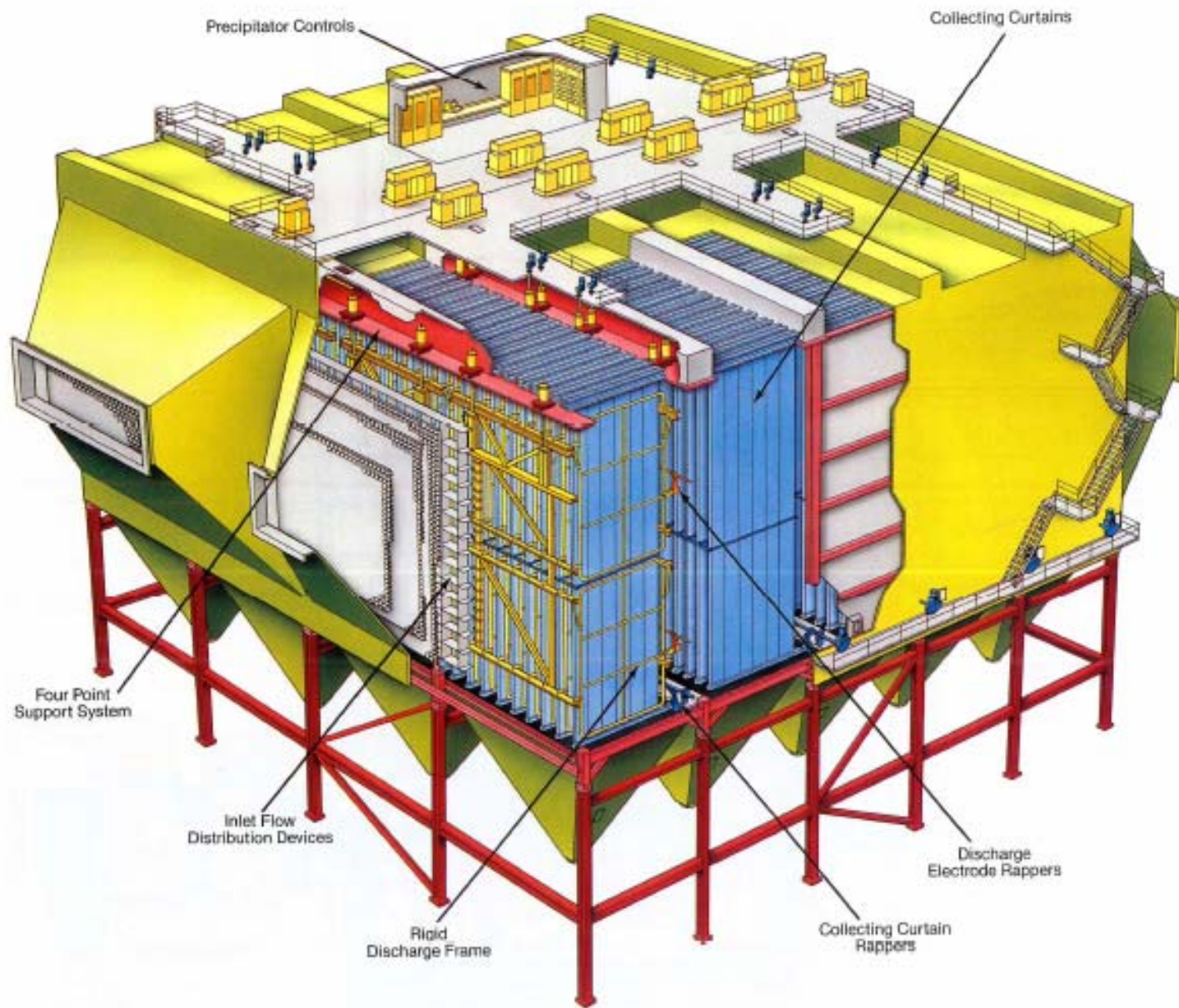
# Water Pipe to Cooling Towers

## 350,000 gal/min!!!

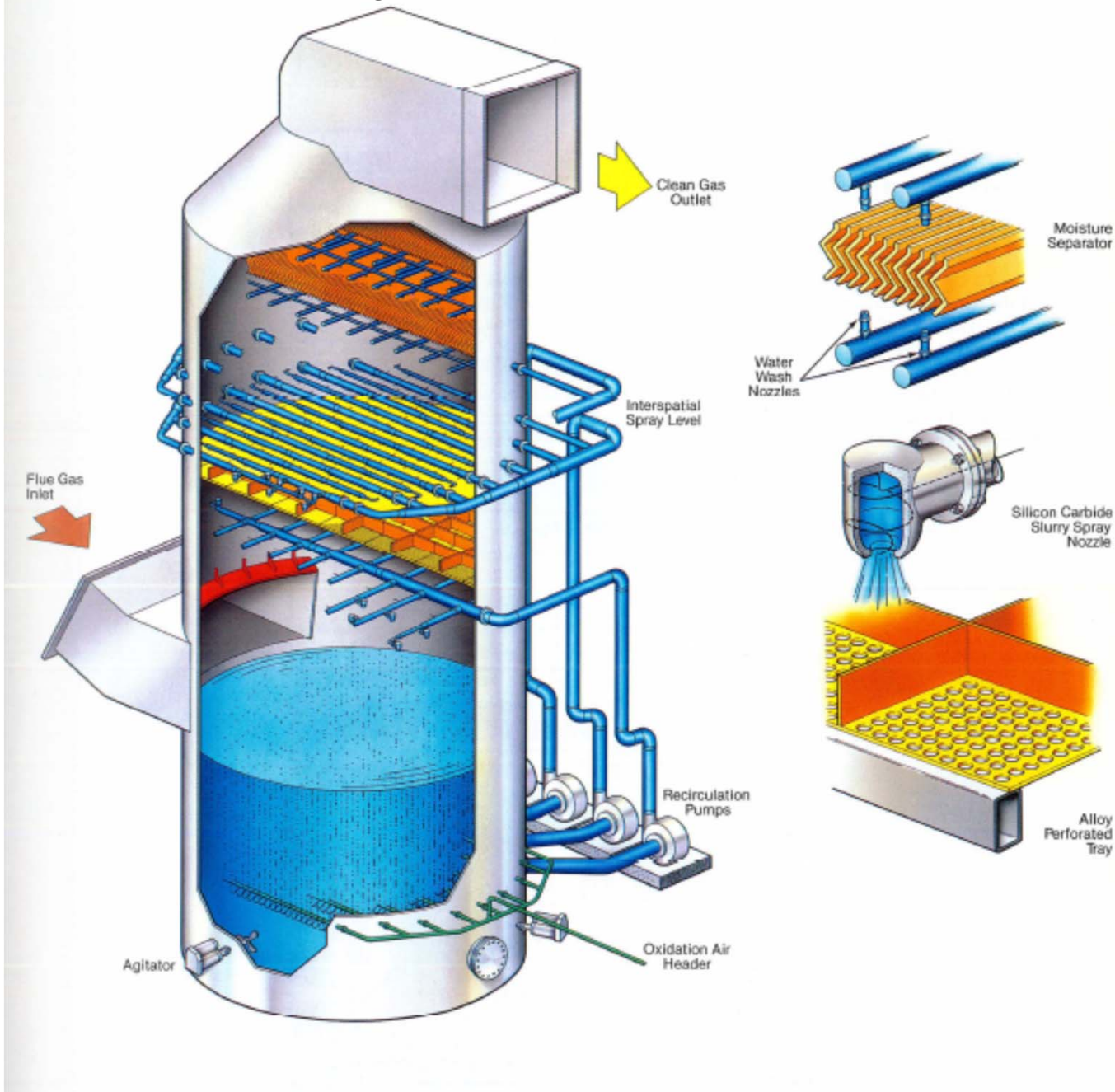




## Electrostatic Precipitator



## Wet flue gas desulfurization scrubber



# Homework Hints from Web Page

- 14-6. Wow- This was a hard problem. The author's answer key had lots of mistakes. The solubilities are not in the book, but are listed in Perry's Chemical Engineering Handbook. To be consistent, let's use the following solubilities:  
0.002 kg  $\text{CaCO}_3$ /100 kg liquid  $\text{H}_2\text{O}$   
0.003 kg  $\text{CaSO}_3$ /100 kg liquid  $\text{H}_2\text{O}$   
I found that sometimes drawing a unit separately, with the inlet and outlet streams, along with the different phases, helped me to sort through this problem. See the TA's to check answers.  
On 14.6e, the second part of this problem is somewhat ambiguous. What is really wanted is the mass fraction of each component in the wet solids.
- 14-7. I found it a lot easier to do this problem if I calculated the heat of formation of coal. The high heating value of the coal is given, which is the negative of the heat of combustion assuming that the hydrogen goes to liquid water. All reactants and products are 25 C for this value of the heat of combustion. Since the heat of combustion is the total enthalpy of the products minus the total enthalpy of the reactants, and the total enthalpies at 25 C are merely the weighted sums of the heats of formation, the only unknown in this equation is the heat of formation of coal. Assume a basis of 100 kg of dry coal, calculate the products, and then calculate the weighted sum of enthalpies, and finally determine the heat of formation of the coal.
- 14-10. The only way that this problem makes sense is if the heat input to the boiler is really the heating value of the coal going into the boiler. This is the way that the regulations are written.
- 14-14c. From the flow rate of methane, multiply the flow rate by the heat of combustion (i.e., the high heating value). This is an approximate heat addition rate. Then divide by the high heating value of the coal.



# Individual Scrubber

## Slurry in

CaCO<sub>3</sub>(s)  
Inert(s)  
Sat'd CaCO<sub>3</sub>(aq)  
Sat'd CaSO<sub>3</sub>(aq)  
H<sub>2</sub>O (liq)  
CaSO<sub>3</sub> · ½H<sub>2</sub>O(s)

## Air Out

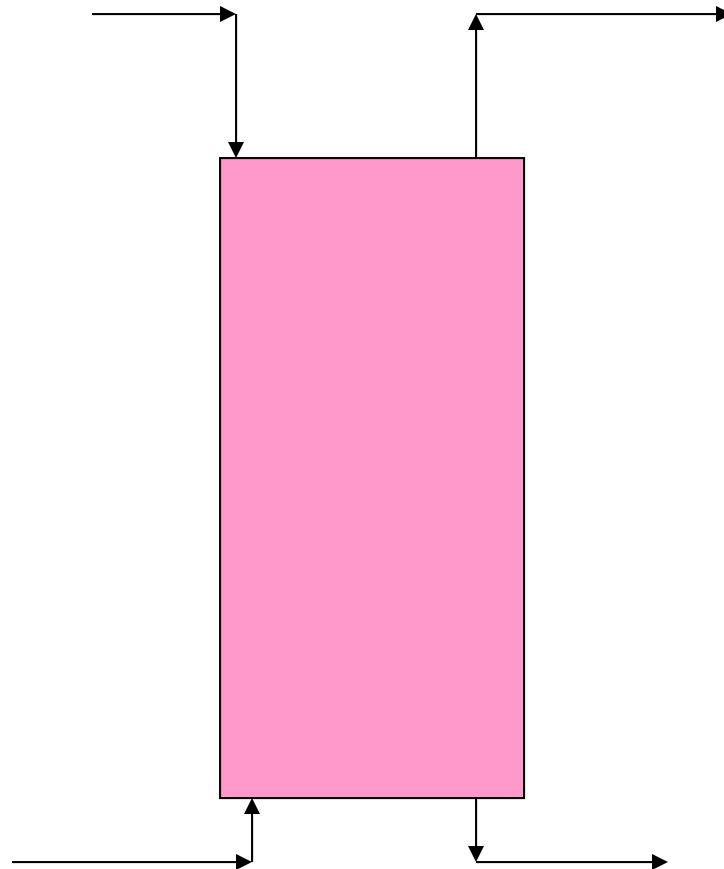
O<sub>2</sub>  
N<sub>2</sub>  
10% of SO<sub>2</sub> gas in  
H<sub>2</sub>O (sat'd vapor)  
CO<sub>2</sub> + Extra CO<sub>2</sub>

## Air In

O<sub>2</sub>  
N<sub>2</sub>  
H<sub>2</sub>O (gas)  
CO<sub>2</sub>  
SO<sub>2</sub> gas  
Fly ash

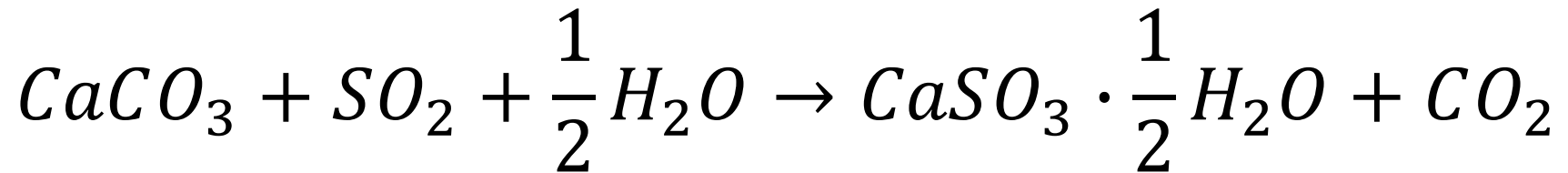
## Slurry Out

CaCO<sub>3</sub>(s)  
Inert(s)  
Sat'd CaCO<sub>3</sub>(aq)  
Sat'd CaSO<sub>3</sub>(aq)  
H<sub>2</sub>O (liq)  
CaSO<sub>3</sub> · ½H<sub>2</sub>O(s)  
Fly ash



# SO<sub>2</sub> Reaction

---



$$\dot{n}_{SO_2,removed} = \dot{n}_{SO_2,in} - \dot{n}_{SO_2,out} \quad (= \dot{\xi})$$

$$\dot{n}_{CO_2,formed} = \dot{n}_{SO_2,removed}$$

$$\dot{n}_{CaCO_3,consumed} = \dot{n}_{SO_2,removed}$$

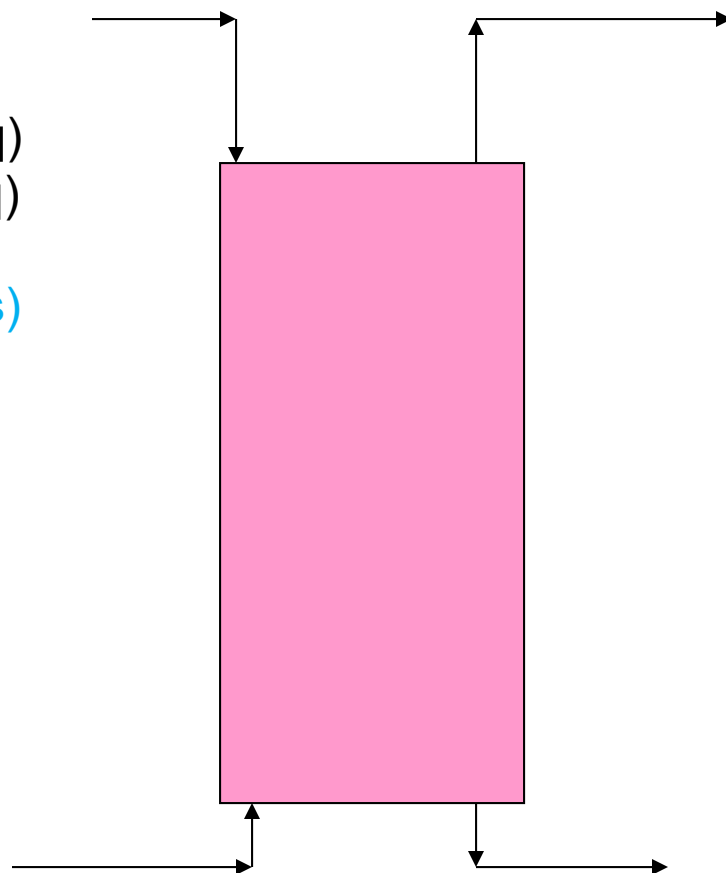
$$\dot{n}_{CaSO_3-hydrate,formed} = \dot{n}_{SO_2,removed}$$

$$\dot{n}_{H_2O,consumed} = \frac{1}{2} \dot{n}_{SO_2,removed}$$

## H<sub>2</sub>O Balance on Individual Scrubber

### Slurry in

CaCO<sub>3</sub>(s)  
Inert(s)  
Sat'd CaCO<sub>3</sub>(aq)  
Sat'd CaSO<sub>3</sub>(aq)  
H<sub>2</sub>O (liq)  
CaSO<sub>3</sub> · ½H<sub>2</sub>O(s)



### Air Out

O<sub>2</sub>  
N<sub>2</sub>  
10% of SO<sub>2</sub> gas in  
H<sub>2</sub>O (sat'd vapor)  
CO<sub>2</sub> + Extra CO<sub>2</sub>

### Air In

O<sub>2</sub>  
N<sub>2</sub>  
H<sub>2</sub>O (gas)  
CO<sub>2</sub>  
SO<sub>2</sub> gas  
Fly ash

### Slurry Out

CaCO<sub>3</sub>(s)  
Inert(s)  
Sat'd CaCO<sub>3</sub>(aq)  
Sat'd CaSO<sub>3</sub>(aq)  
H<sub>2</sub>O (liq)  
CaSO<sub>3</sub> · ½H<sub>2</sub>O(s)  
Fly ash



# H<sub>2</sub>O Balance on a Scrubber

---

$$\begin{array}{l} \dot{n}_{H_2O(g),in} + \dot{n}_{H_2O(liq),in} + 0.5\dot{n}_{CaSO_3-hydrate,in} \\ \text{From boiler} \quad \text{From recycled} \quad \text{From recycled} \\ \text{exhaust} \quad \text{slurry} \quad \text{slurry} \\ = \dot{n}_{H_2O(g),out} + \dot{n}_{H_2O(liq),out} + 0.5\dot{n}_{CaSO_3-hydrate,out} \\ \text{To stack} \quad \text{In slurry} \quad \text{In slurry} \end{array}$$

Values in green and blue are unknown, but -----

$$0.5\dot{n}_{CaSO_3-hydrate,out} - 0.5\dot{n}_{CaSO_3-hydrate,in} = \dot{n}_{CaSO_3-hydrate,formed}$$

Since  $\dot{n}_{CaSO_3-hydrate,formed}$  is known (previous slide), the water balance now has only one unknown:

$$\dot{n}_{H_2O(liq),out}$$

## A note on treating the mass of liquid with solubilities

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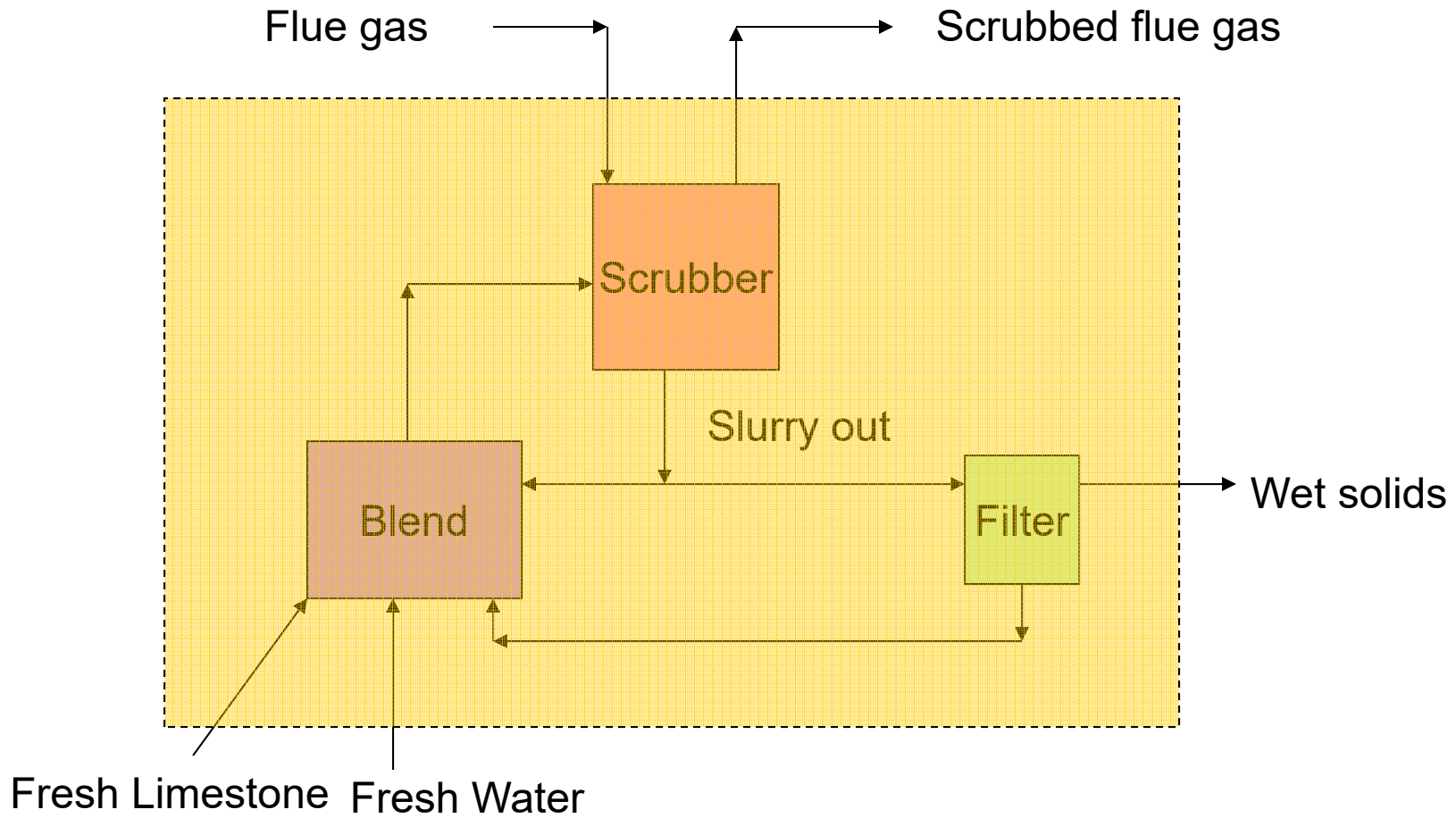
$$\dot{m}_{liq} = \dot{m}_{H_2O,liq} + \dot{m}_{CaCO_3,aq} + \dot{m}_{CaSO_3,aq}$$

$$\dot{m}_{CaCO_3,aq} = \frac{0.002}{100} \dot{m}_{H_2O}$$

$$\dot{m}_{CaSO_3,aq} = \frac{0.003}{100} \dot{m}_{H_2O}$$

$$\dot{m}_{liq} = \left( 1 + \frac{0.002}{100} + \frac{0.003}{100} \right) \dot{m}_{H_2O}$$

14.6 (e)





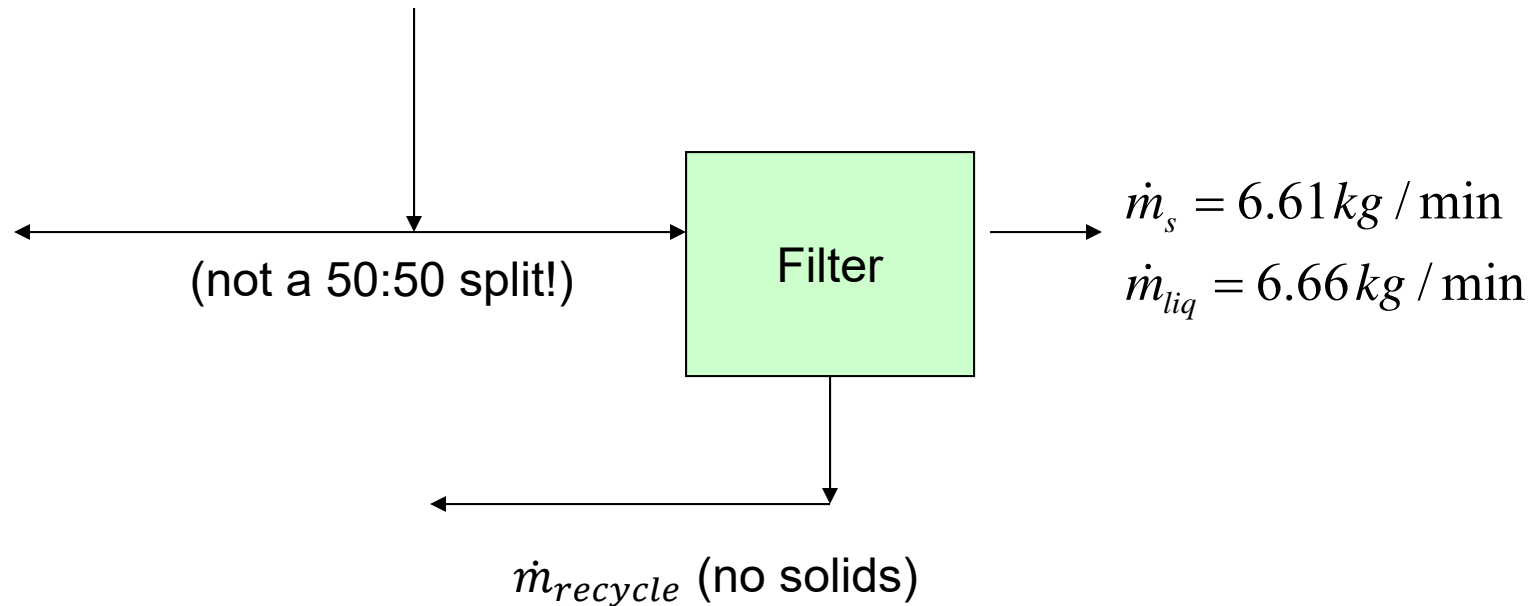
# Solids Out in Filter Cake

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- All inerts (solid) from limestone → inerts in solid in filter cake
- All unreacted  $\text{CaCO}_3$  in with fresh limestone → wet filter cake
- All  $\text{SO}_2$  removed → wet filter cake ( $\text{CaSO}_3$  hydrate)
- All flyash in with flue gas → filter cake
- Sum of all of above =  $\dot{m}_{\text{solids}}$   
+ small amount of aqueous  $\text{CaSO}_3$  hydrate
- Need to guess  $\dot{m}_{\text{H}_2\text{O},\text{liq}}$  and use solver to ensure that
$$\frac{\text{Total mass of liquid out}}{\text{Total mass out}} = 0.502$$

14.6 (f)

S/L ratio = 0.1115 (from 14.6c)



So  $\frac{\dot{m}_s}{\dot{m}_{liq} + \dot{m}_{recycle}} = 0.1115$  (combination of outputs from filter has same S/L ratio as the inlet)

Rearranging,

$$\dot{m}_{recycle} = \frac{\dot{m}_s}{0.1115} - \dot{m}_{liq}$$