

EMS717U/EMS717P

Renewable Energy Sources

Thermodynamics of Energy Conversion

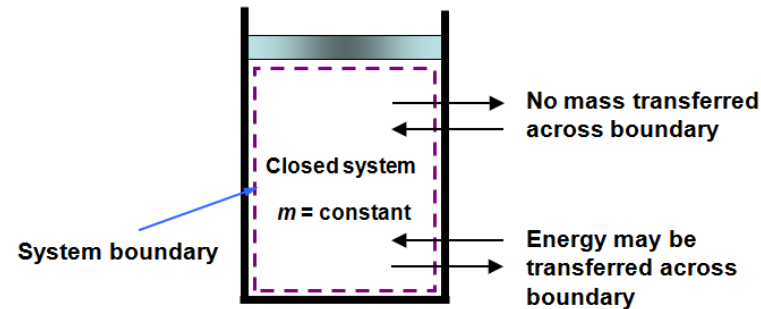
Prof. Huasheng Wang

Laws of thermodynamics

- The first law of thermodynamics

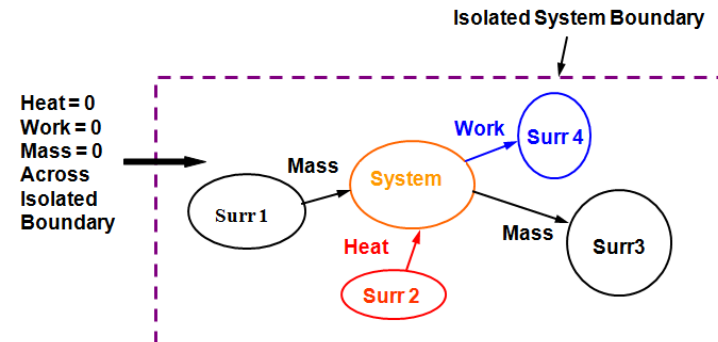
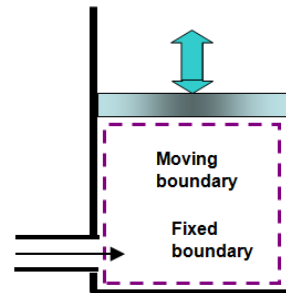
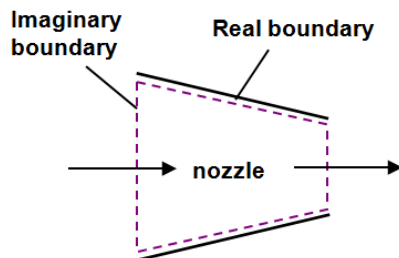
For any process of a stationary system

$$W + Q = E_2 - E_1$$



Steady flow energy equation (SFEE)

$$W + Q = \Sigma[m(h + v^2/2 + gz)]_2 - \Sigma[m(h + v^2/2 + gz)]_1$$



Laws of thermodynamics

- Entropy and the 2nd Law of thermodynamics

Spontaneous (no outside influence) processes lead to increase in entropy. This is the essence of the Second Law of Thermodynamics.

- $dE = 0$
 - $dS \geq 0$
- For an isolated system the internal energy is constant and the entropy can only increase. Entropy is constant for ideal reversible process

Fundamentals

- **Thermodynamic cycle**

Gas power cycles

Carnot cycle

Otto cycle (ideal cycle for spark-ignition engines)

Diesel cycle (ideal cycle for compression-ignition engines)

Stirling cycle

Ericsson cycle

Brayton cycle

Steam power cycles

Rankine cycle

Refrigeration cycles

Vapour-compression refrigeration cycle

Fundamentals

- Energy conversion devices

SFEEs (steady-flow energy equation) for components of cycles

Components	w	q	Δh
Nozzles	0	0	0
Valves	0	0	0
Throttles	0	0	0
Ducts, pipes, diffusers	0	0	0
Boilers	0	$h_{\text{out}} - h_{\text{in}}$	q_{in}
Heat exchangers	0	$ h_{\text{out}} - h_{\text{in}} $	q
Condensers	0	$h_{\text{in}} - h_{\text{out}}$	q_{out}
Evaporators	0	$h_{\text{out}} - h_{\text{in}}$	q_{in}
Combustion chambers	0	$h_{\text{out}} - h_{\text{in}}$	q_{in}
Compressors, pumps, etc.	$h_{\text{out}} - h_{\text{in}}$	0 (?)	w_{in}
Turbines	$h_{\text{in}} - h_{\text{out}}$	0 (?)	w_{out}

Fundamentals

- Thermal efficiency

Thermal efficiency = Desired output/Required input

$$\eta_{\text{Carnot}} \equiv \frac{W_{\text{net}}}{Q_{\text{in}}} = 1 - \frac{Q_{\text{out}}}{Q_{\text{in}}} = 1 - \frac{T_{\text{L}}}{T_{\text{H}}}$$

COP (Coefficient Of Performance)

The performance of refrigerators and heat pumps is evaluated in terms of *coefficient of performance* (COP).

$$COP_{\text{R}} = \frac{\text{Desired output}}{\text{Required input}} = \frac{\text{Cooling effect}}{\text{Work input}} = \frac{Q_{\text{L}}}{W_{\text{net,in}}}$$

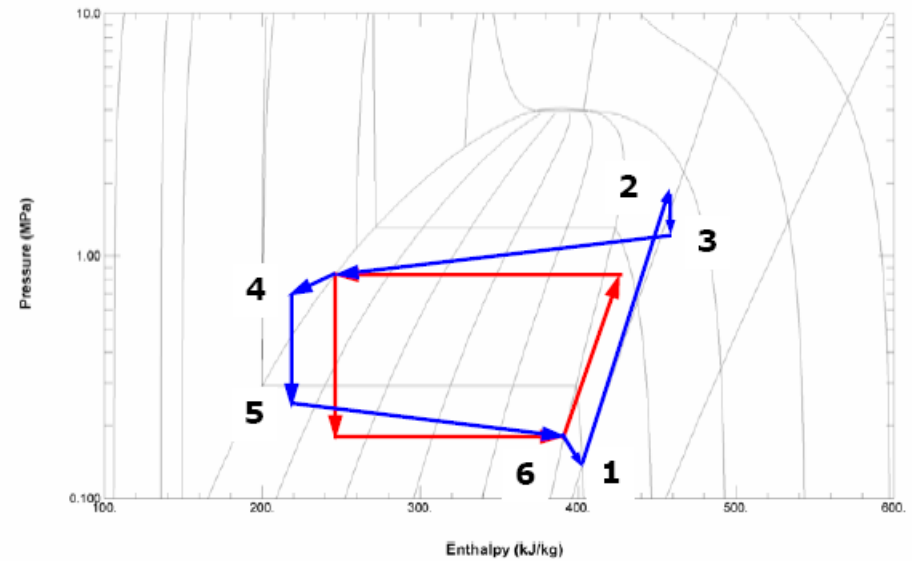
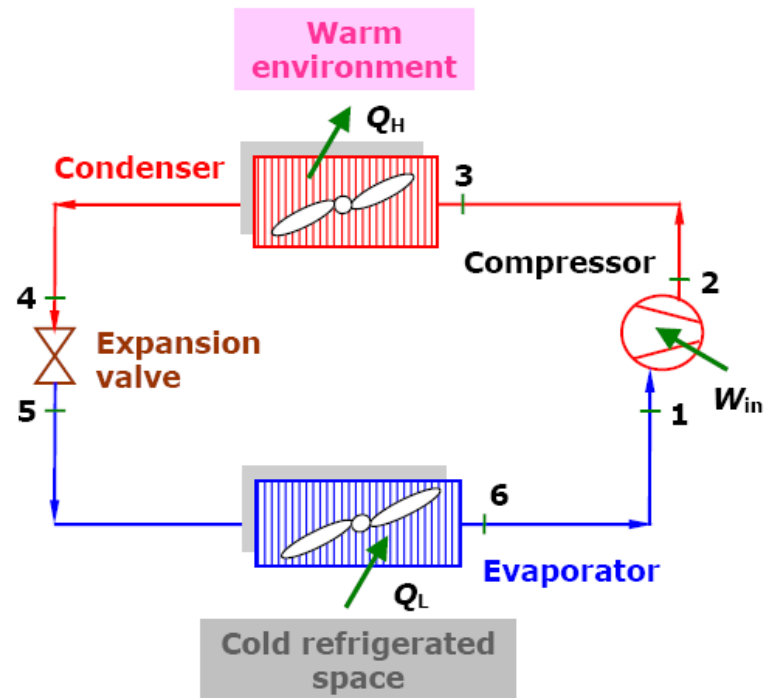
$$COP_{\text{HP}} = \frac{\text{Desired output}}{\text{Required input}} = \frac{\text{Heating effect}}{\text{Work input}} = \frac{Q_{\text{H}}}{W_{\text{net,in}}}$$

Both COP_{R} and COP_{HP} can be larger than 1. Under the same operating conditions, the COPs are related by

$$COP_{\text{HP}} = COP_{\text{R}} + 1$$

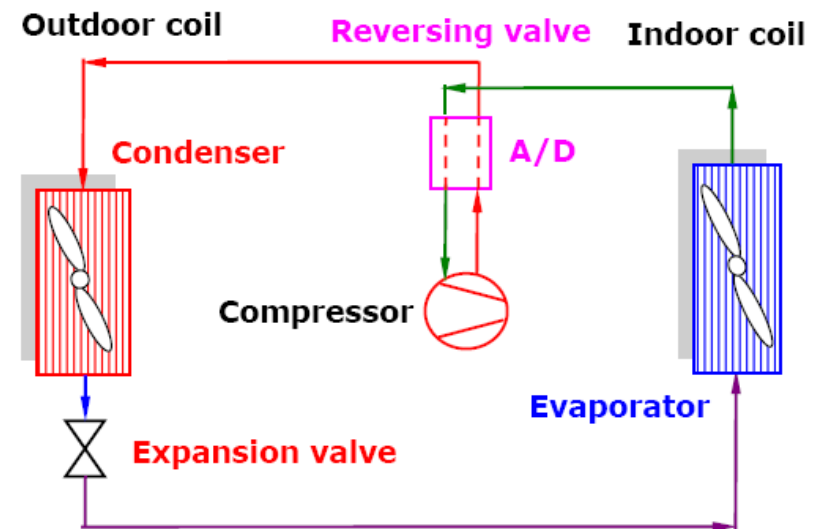
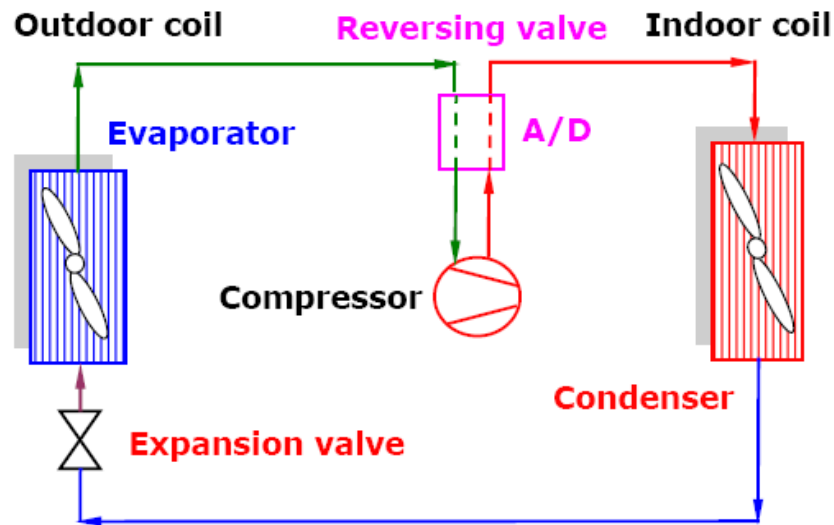
Fundamentals

- Vapour-compression refrigeration cycle



Fundamentals

- Heat pump systems



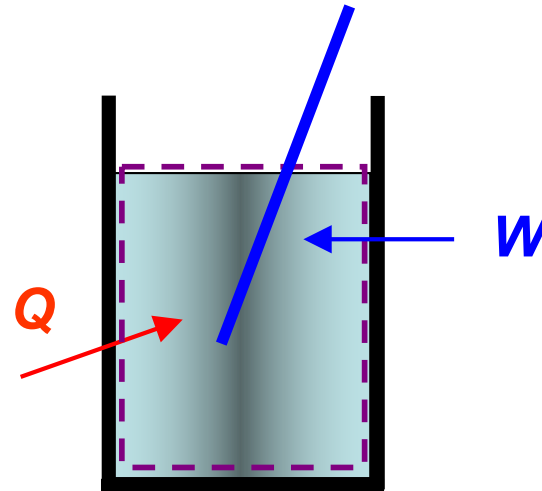
- High pressure/temperature vapour
- High pressure/temperature liquid
- Low pressure/temperature mixture
- Low pressure/temperature vapour

Examples - 1

A vessel containing water is stirred so that an amount of work of 20 J is done on it. The initial and final internal energies of water are 100 J and 105 J respectively. Find the heating during this process.

$$W = +20 \text{ J}$$

$$\begin{aligned} Q &= E_2 - E_1 - W \\ &= (105 - 100 - 20) \text{ J} \\ &= -15 \text{ J} \end{aligned}$$



Note: **negative Q** means the water loses energy by heat in this case.

Examples - 2

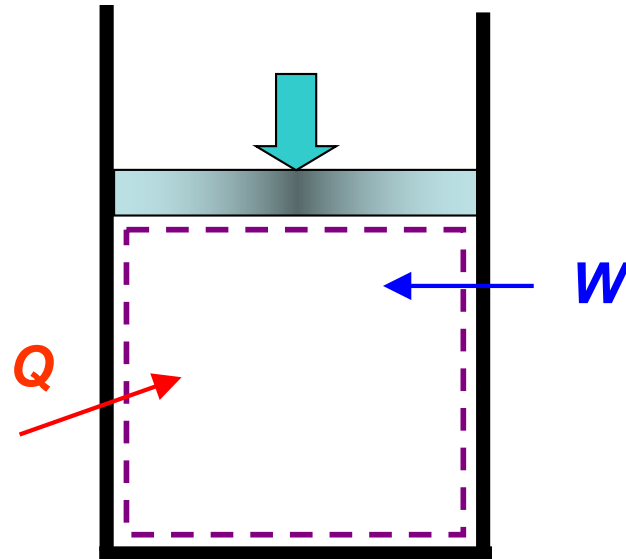
A cylinder containing air is compressed. The work done on the air is 100 J. The initial and final internal energies are equal. Determine the heating during the process.

$$W = 100 \text{ J}$$

$$Q = E_2 - E_1 - W$$

$$= -W$$

$$= -100 \text{ J}$$



Note: **Minus sign** indicates that energy is transferred *from* the system by the heat process or heating is done *by* the air *on* the surroundings.

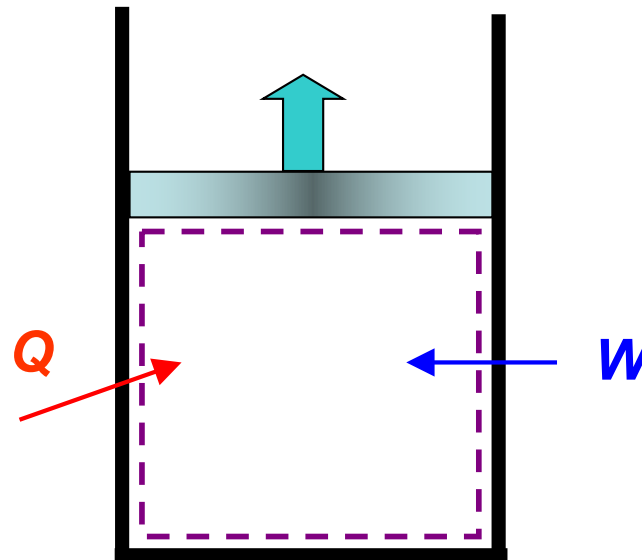
Examples - 3

An amount of gas expands in a cylinder fitted with a piston. During the expansion process the gas does an amount of work equal to 50 J on its environment. The environment is cooler than the gas which also loses 25 J of energy by the heat mode. Find the decrease in internal energy of the gas for this process.

$$W = -50 \text{ J}$$

$$Q = -25 \text{ J}$$

$$\begin{aligned} E_2 - E_1 &= Q + W \\ &= (-25 - 50) \text{ J} \\ &= -75 \text{ J} \end{aligned}$$



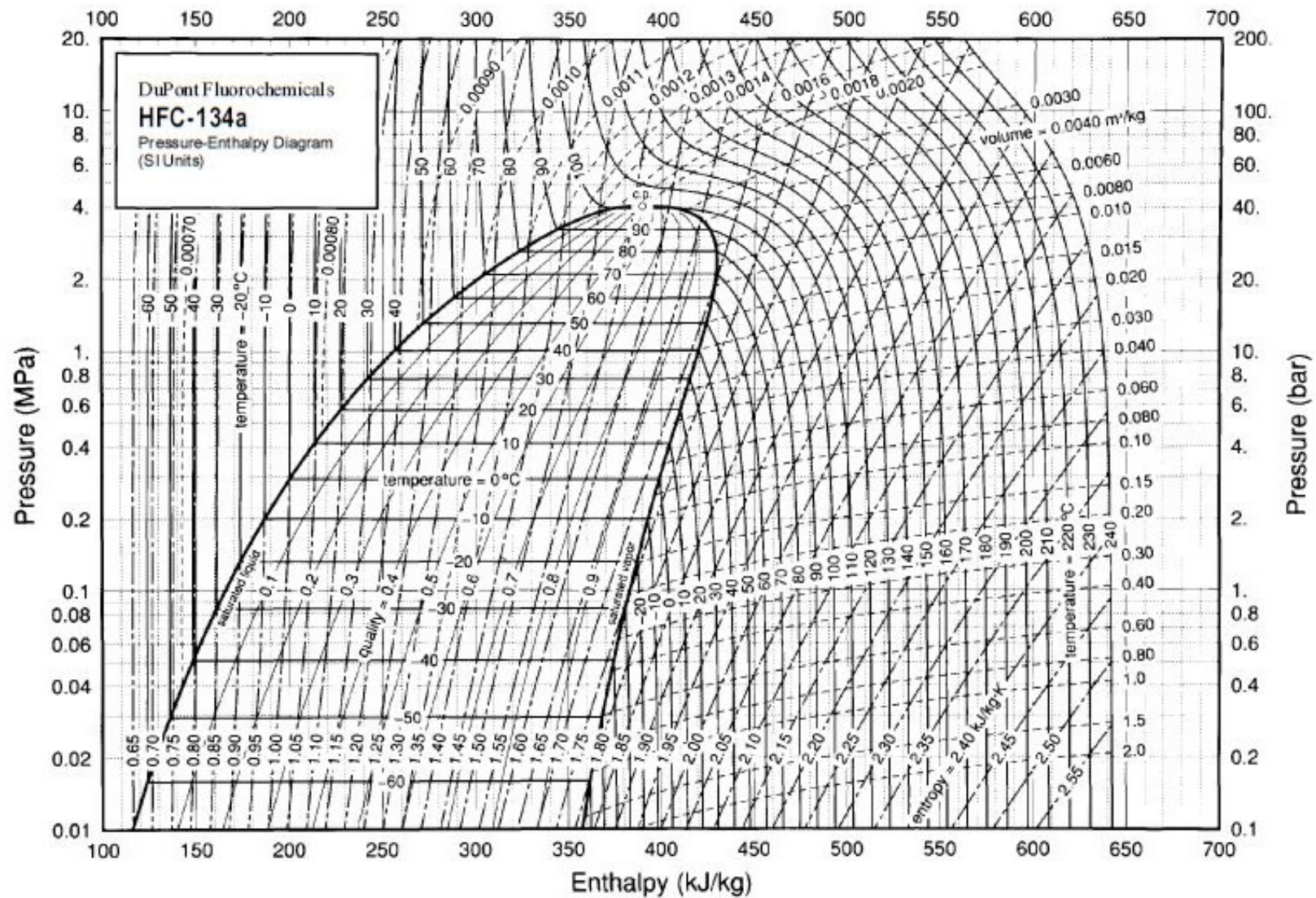
Question asks for **decrease** of internal energy

$$E_1 - E_2 = +75 \text{ J}$$

Problem 2

- a) Draw a diagram of a ground source heat pump for space cooling and heating.
- b) Assume that the working fluid, refrigerant R134a, undergoes in the heat pump an ideal thermodynamics cycle consisting of four processes:
 - i) R134a evaporates at constant temperature of 10 °C in the evaporator to the saturation vapour state;
 - ii) It is adiabatically compressed by the compressor to a high temperature and high pressure vapour state;
 - iii) It is cooled and condensed in the condenser at constant pressure (the condensation temperature is 50 °C) to the saturation liquid state;
 - iv) it expands in the expansion valve through a constant enthalpy process to the evaporation temperature.

The mass flow rate of R134a is 10 g/s. Show the vapour compression cycle on the pressure P – enthalpy h diagram below

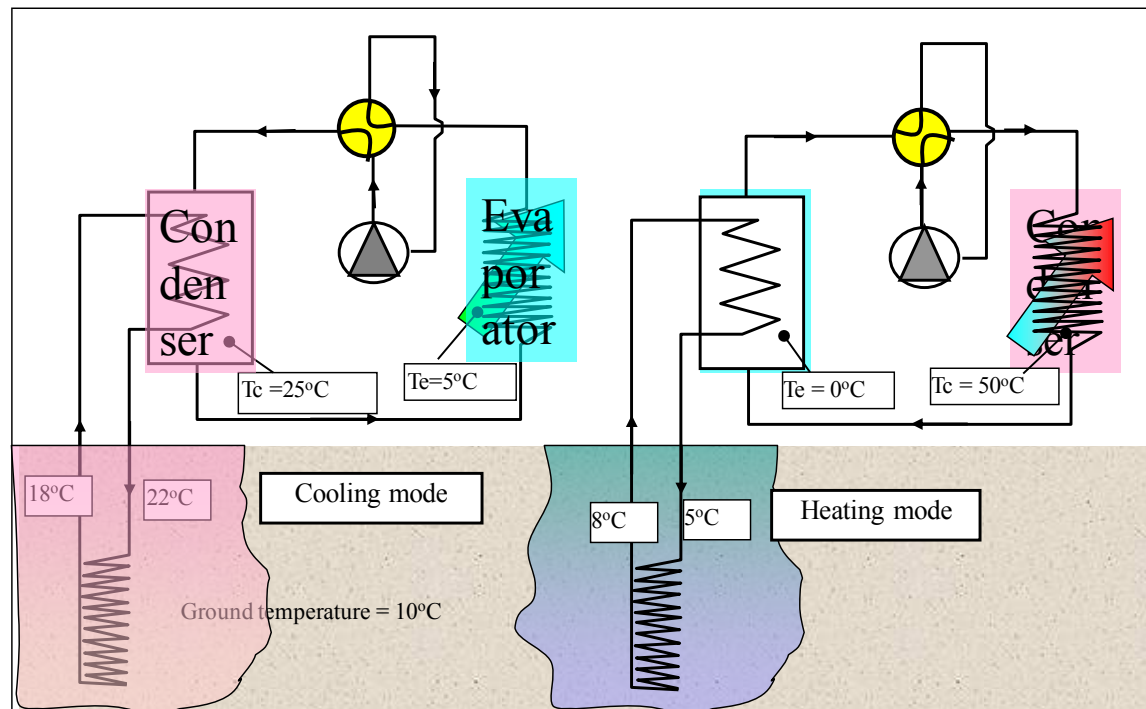


- c) Calculate the heating rate by the condenser, the cooling rate by the evaporator and the work by the compressor.
- d) Define the Coefficient of Performance (COP) for the cooling and heating modes.

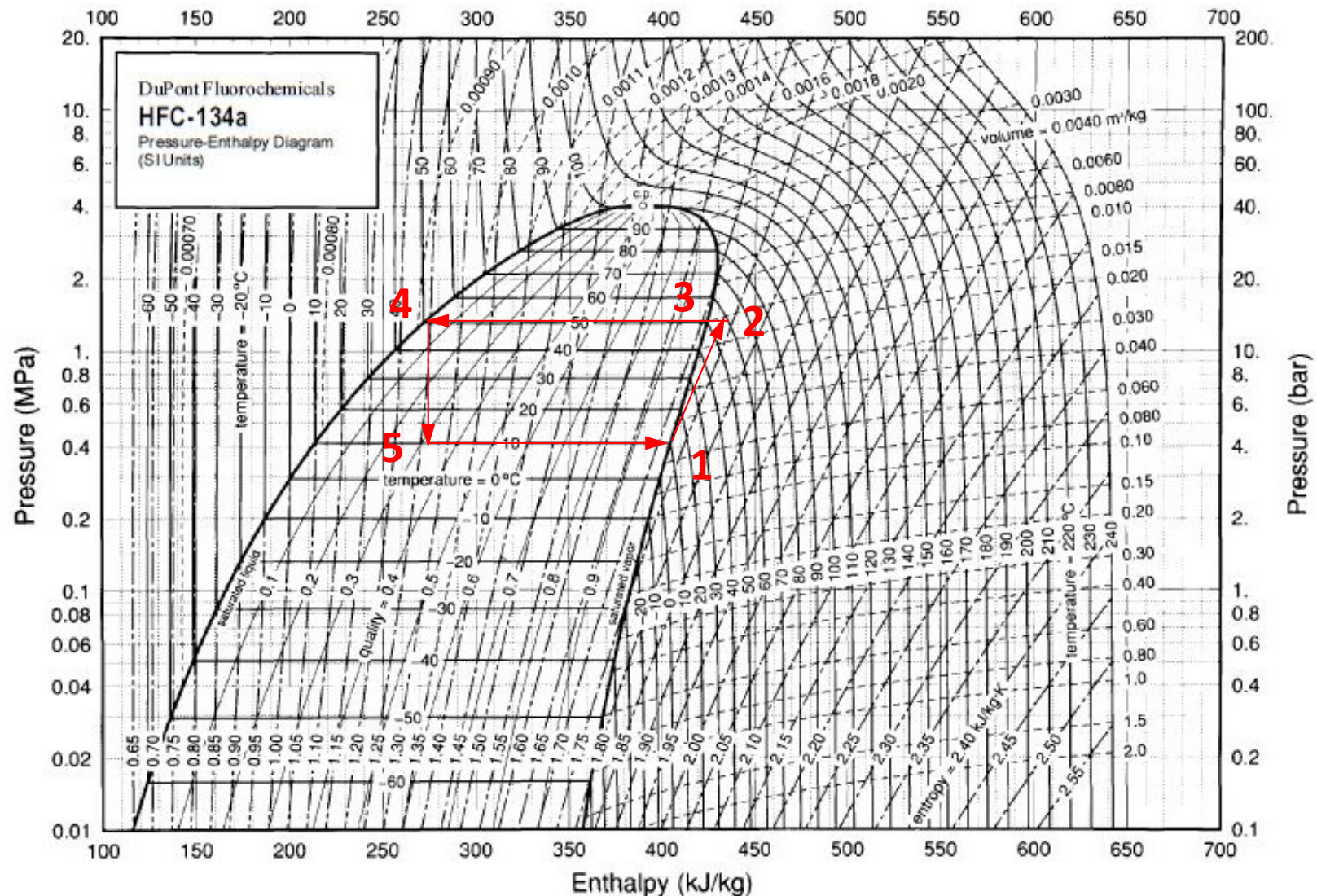
Solution

- a) The figure below shows a schematic of a ground source heat pump. It is a vapour–compression refrigeration cycle using ground heat exchanger as the condenser or evaporator. Since the ground temperature remains constant at about 12 °C through a year, the heat pump provides higher COP.

Ground source heat pumps GSHP



- b) The ideal cycle is shown in P - h diagram below. **State 1** is a saturated vapour state at temperature 10 °C. **State 2** is a superheated vapour state with pressure $P_2 = P_3 = P_4$. **State 3** is a saturated vapour state at temperature 50 °C. **State 4** is the saturated liquid state at temperature 50 °C. **State 5** is a two-phase state with temperature 10 °C and $h_5 = h_4$.



c) From the diagram,

$$T_1 = T_5 = 10\text{ }^{\circ}\text{C}, h_1 = 404.3\text{ kJ/kg}, P_1 = 0.415\text{ MPa}, s_1 = 1.72\text{ kJ/kg K};$$

$$s_1 = s_2 = 1.72, T_2 = 54.0\text{ }^{\circ}\text{C}, h_2 = 428.3\text{ kJ/kg}$$

$$P_3 = P_4 = 1.32\text{ MPa}, h_3 = 423.4\text{ kJ/kg}, h_4 = 271.6\text{ kJ/kg}$$

$$h_5 = h_4 = 271.6\text{ kJ/kg}$$

$$\text{The heating rate} = m (h_2 - h_4) = 0.01 (428.3 - 271.6) = 1567\text{ W}$$

$$\text{The cooling rate} = m (h_1 - h_5) = 0.01 (404.3 - 271.6) = 1327\text{ W}$$

$$\text{The work} = m (h_2 - h_1) = 0.01 (428.3 - 404.3) = 240\text{ W}$$

- d) The Coefficient of Performance (*COP*) for the cooling and heating modes are:

$$COP \text{ for heating} = \frac{\text{the heating rate}}{\text{the work}} = \frac{1567}{240} = 6.53$$

$$COP \text{ for cooling} = \frac{\text{the cooling rate}}{\text{the work}} = \frac{1327}{240} = 5.53$$

Problem Solving Techniques - 1

- **General step-by-step approach**

1. **Problem statement:** list given info and objectives
2. **Schematic:** draw one to indicate relevant info on
3. **Assumptions and approximations:** simplify and justify
4. **Physical laws:** apply relevant laws and principles
5. **Properties:** refer to property relations, tables, property source
6. **Calculations;** verify units, significant figures (“do not copy”)
7. **Conclusion:** reasoning, verification, and discussion; validity, significance, implications, recommendations, limitations
8. **Presentation:** neatness, organization, completeness

Problem Solving Techniques - 2

- **General step-by-step approach**

1. Identify the problem
2. Make appropriate assumptions and approximations
3. Set up a physical model
4. Produce a mathematical model
5. Implement the model into a procedure
6. Verify the model and procedure using a simple case with results available
7. Calculate
8. Analyse results
9. Conclusion

Problem Solving Techniques - 3

- **Conservation of mass, momentum, energy & electrical charge (the first law of thermodynamics)**

$$\left\{ \begin{array}{l} \text{The net gain in} \\ X \text{ by the system} \end{array} \right\} = \left\{ \begin{array}{l} \text{The net amount of } X \\ \text{transported into the system} \end{array} \right\}$$

Problem 1

Water and molten blast furnace slag enter a granulator in which the slag is cooled and vitrified to form a sand-like material. When the inlet flow rate and temperature of the slag are 1400 °C and 6 tonne/min respectively and the water inlet flow rate and temperature are 2400 m³/h and 50 °C respectively, saturated steam at 100 °C is generated and leaves the granulator via a stack. Granulated slag and water leave as slurry from the base of the tank at a temperature of 100 °C. Determine the mass flow rate of the steam (kg/s) leaving the granulator. Take the specific enthalpy of the slag at 1400 °C and 100 °C to be 1670 J/g and 85 J/g respectively. Neglect any heat transfer from the granulator.

Solution - 1

1. Problem statement: list given info and objectives

At the inlet

$$T_{sl1} = 1400\text{ }^{\circ}\text{C}, m_{sl1} = 6\text{ tonne/min}, h_{sl1} = 1670\text{ J/g},$$

$$m_{w1} = 2400\text{ m}^3/\text{h}, T_{w1} = 50\text{ }^{\circ}\text{C},$$

At the outlet

$$T_{v2} = 100\text{ }^{\circ}\text{C}, m_{v2} = ?$$

$$T_{w2} = T_{sl2} = 100\text{ }^{\circ}\text{C (slurry)}, m_{w2} = ?$$

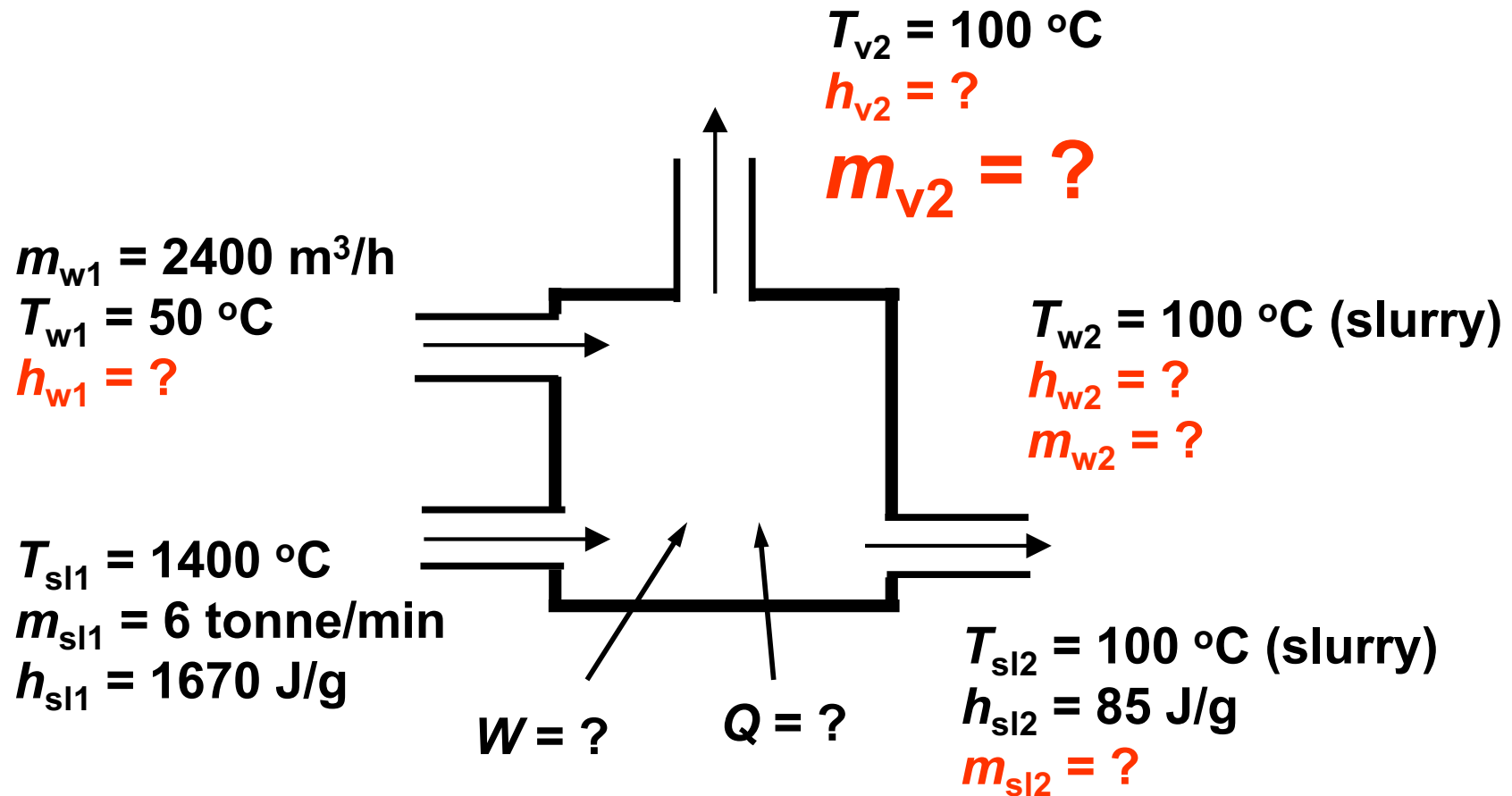
$$T_{sl2} = 100\text{ }^{\circ}\text{C (slurry)}, h_{sl2} = 85\text{ J/g}, m_{sl2} = ?$$

Determine

$$m_{v2} = ?$$

Solution - 2

2. Schematic: draw one to indicate relevant info on



Solution - 3

3. Assumptions and approximations: simplify and justify

- a. Steady state flow?**
- b. Atmospheric pressure?**
- c. Neglect any heat transfer from the granulator?**
- d. Neglect potential energy change?**
- e. Neglect kinetic energy change?**
- f. Work done?**
- g. Slag: molten to sand-like material**
- h. Ideal gas?**
- i. Perfect gas?**
- j. Equilibrium state?**
- k. Assume equilibrium state at inlet and outlet.**

Solution - 4

4. Physical laws: apply relevant laws and principles

Open system

Conservation of mass

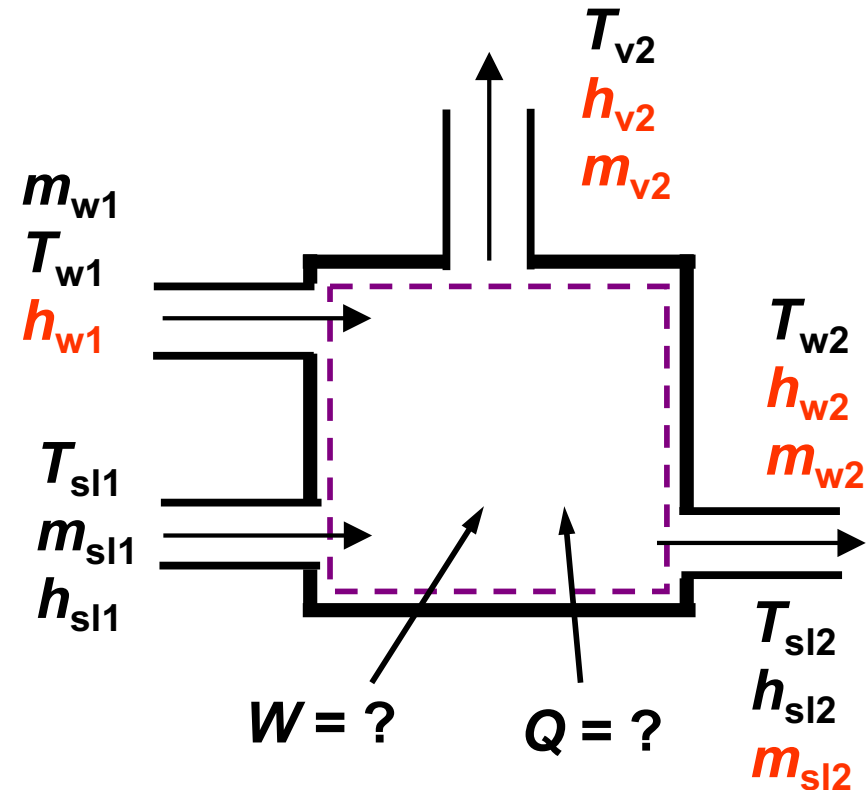
$$m_{sl1} = m_{sl2}$$

$$m_{w1} = m_{w2} + m_{v2}$$

1st law of thermodynamics

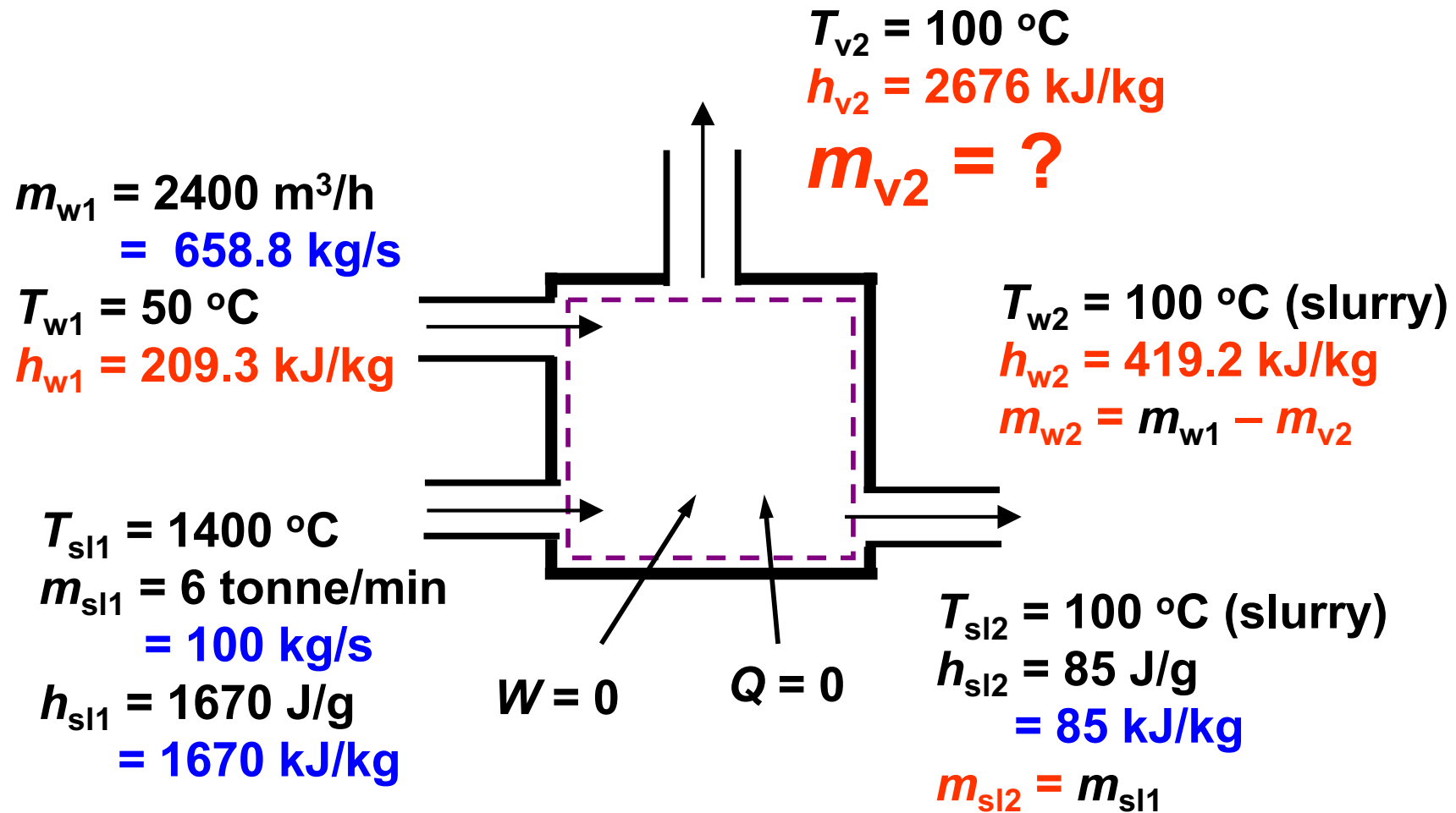
$$W + Q = \Sigma[m(h + v^2/2 + gz)]_2 - \Sigma[m(h + v^2/2 + gz)]_1$$

$$m_{w1}h_{w1} + m_{sl1}h_{sl1} = m_{v2}h_{v2} + m_{w2}h_{w2} + m_{sl2}h_{sl2}$$



Solution - 5

5. Properties: refer to property relations, tables, property source



Solution - 6

6. Calculations: verify units, significant figures (“do not copy”)

$$m_{sl1} = m_{sl2}$$

$$m_{w1} = m_{w2} + m_{v2}$$

$$m_{w1}h_{w1} + m_{sl1}h_{sl1} = m_{v2}h_{v2} + m_{w2}h_{w2} + m_{sl2}h_{sl2}$$

$$m_{w1}h_{w1} + m_{sl1}h_{sl1} = m_{v2}h_{v2} + (m_{w1} - m_{v2})h_{w2} + m_{sl1}h_{sl2}$$

$$m_{v2} = [m_{w1}(h_{w1} - h_{w2}) + m_{sl1}(h_{sl1} - h_{sl2})] / (h_{v2} - h_{w2})$$

$$\begin{aligned} m_{v2} &= [658.8 \times (209.3 - 419.2) + 100 \times (1670 - 85)] \\ &\quad / (2676 - 419.2) \\ &= 8.96 \text{ kg/s} \end{aligned}$$

Solution - 7

2. Schematic: draw one to indicate relevant info on

