

Thermodynamics Lecture Series

Assoc. Prof. Dr. J.J.

Gas Mixtures – Properties and Behaviour

Applied Sciences Education Research
Group (ASERG)

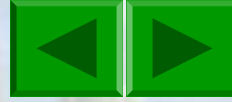
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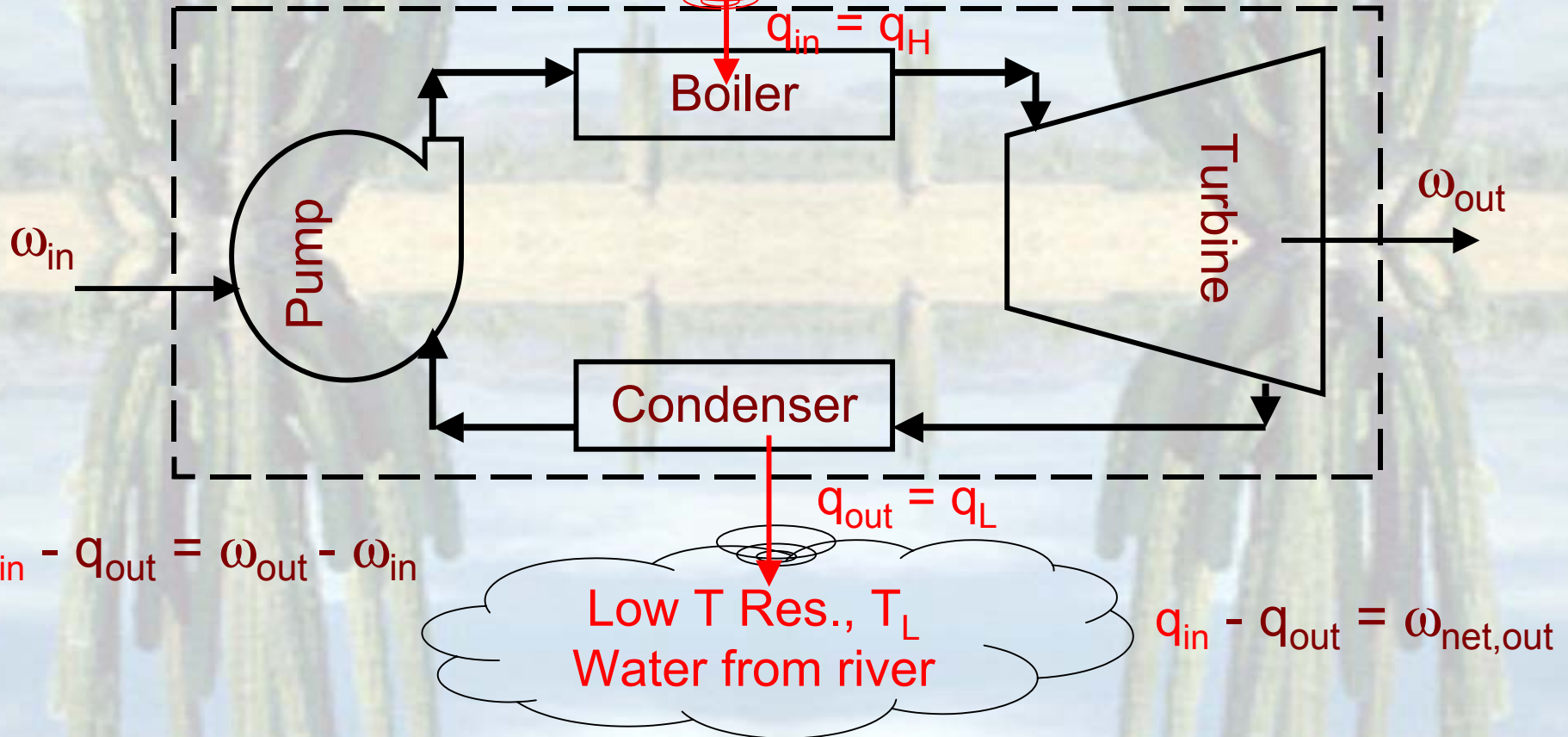
<http://www.uitm.edu.my/faculties/fsg/drjjl.html>

Review – Steam Power Plant



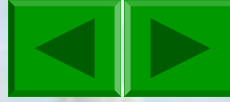
Working fluid:
Water

High T Res., T_H
Furnace



A Schematic diagram for a Steam Power Plant

Review - Steam Power Plant



Working fluid:
Water

High T Res., T_H
Furnace

$$q_{in} = q_H$$

Purpose:
Produce work,
 W_{out} , ω_{out}

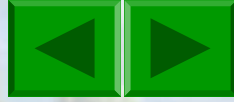
Steam Power Plant

$$\omega_{net,out}$$

$$q_{out} = q_L$$

Low T Res., T_L
Water from river

An Energy-Flow diagram for a SPP



Review - Steam Power Plant

Thermal Efficiency for steam power plants

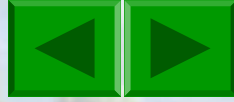
$$\eta = \frac{\textit{desired output}}{\textit{required input}} = \frac{\omega_{net, out}}{q_{in}}$$

$$\eta = \frac{\omega_{net, out}}{q_{in}} = \frac{q_{in} - q_{out}}{q_{in}} = 1 - \frac{q_{out}}{q_{in}} = 1 - \frac{q_L}{q_H}$$

$$\eta_{rev} = 1 - \frac{T_L}{T_H}$$

For real engines, need to find q_L and q_H .





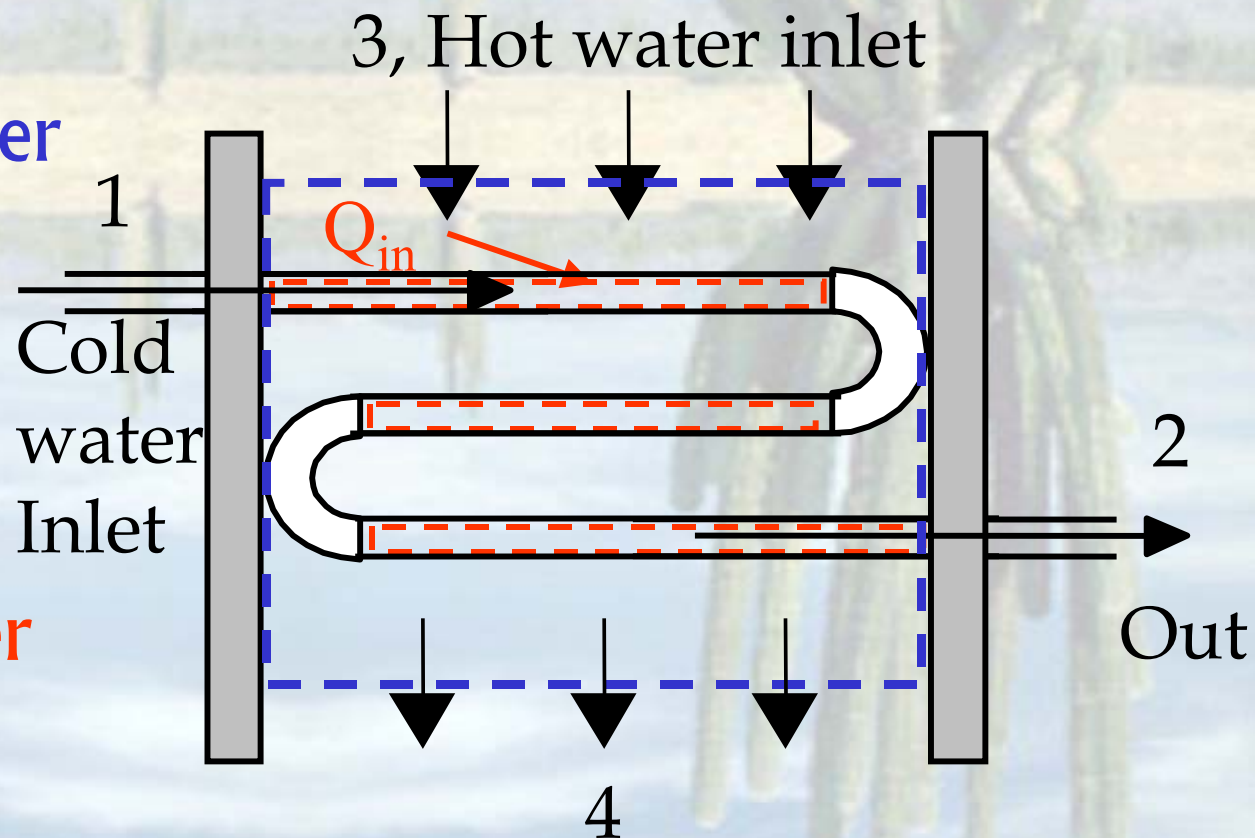
Review - Entropy Balance

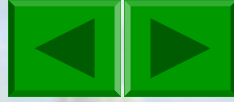
Entropy Balance – Steady-flow device

Heat exchanger

Case 1 – blue border

Case 2 – red border





Review - Entropy Balance

Entropy Balance –Steady-flow device

Heat exchanger: energy balance;

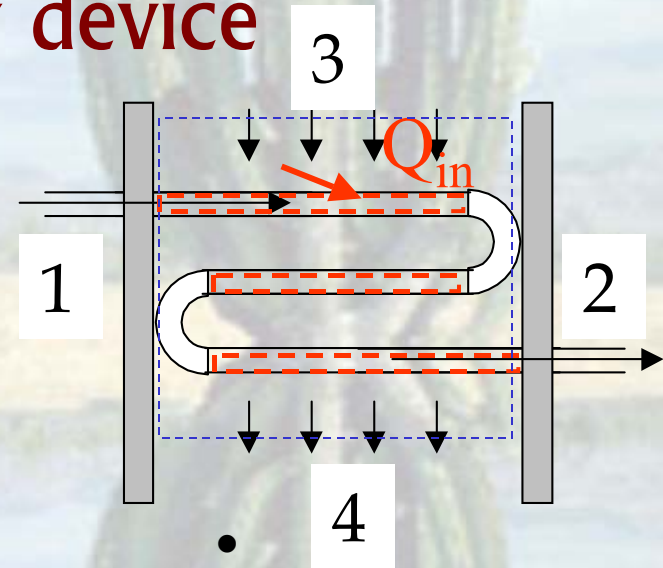
where $\dot{m}_4 = \dot{m}_3$ $\dot{m}_2 = \dot{m}_1$

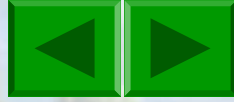
Assume $\Delta ke_{\text{mass}} = 0$, $\Delta pe_{\text{mass}} = 0$

Case 1 $\dot{Q}_{in} - \dot{Q}_{out} + \dot{W}_{in} - \dot{W}_{out} = (\dot{m} \vartheta)_{\text{exit}} - (\dot{m} \vartheta)_{\text{inlet}}$, kW

$$0 = \dot{m}_4 h_4 - \dot{m}_3 h_3 + \dot{m}_2 h_2 - \dot{m}_1 h_1, \text{ kW}$$

$$\dot{m}_4 (h_4 - h_3) = \dot{m}_2 (h_1 - h_2), \text{ kW}$$





Review - Entropy Balance

Entropy Balance –Steady-flow device

Heat exchanger: energy balance;

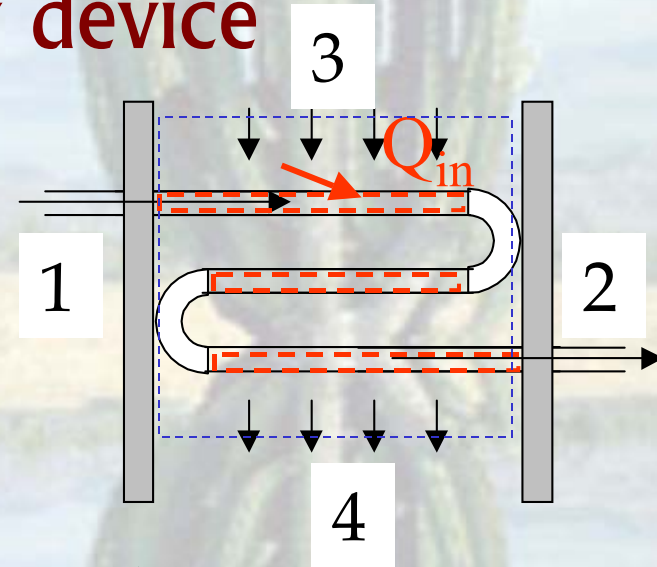
where $\dot{m}_4 = \dot{m}_3$ $\dot{m}_2 = \dot{m}_1$

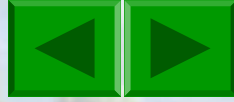
Assume $\Delta ke_{\text{mass}} = 0$, $\Delta pe_{\text{mass}} = 0$

Case 1 $\dot{m}_4 (h_4 - h_3) = \dot{m}_2 (h_1 - h_2)$, kW

Case 2 $\dot{Q}_{in} - \dot{Q}_{out} = \dot{m}_2 \theta_2 - \dot{m}_1 \theta_1$, kW

$$\dot{Q}_{in} - 0 = \dot{m}_2 (h_2 - h_1), \text{ kW}$$





Review - Entropy Balance

Entropy Balance – Steady-flow device

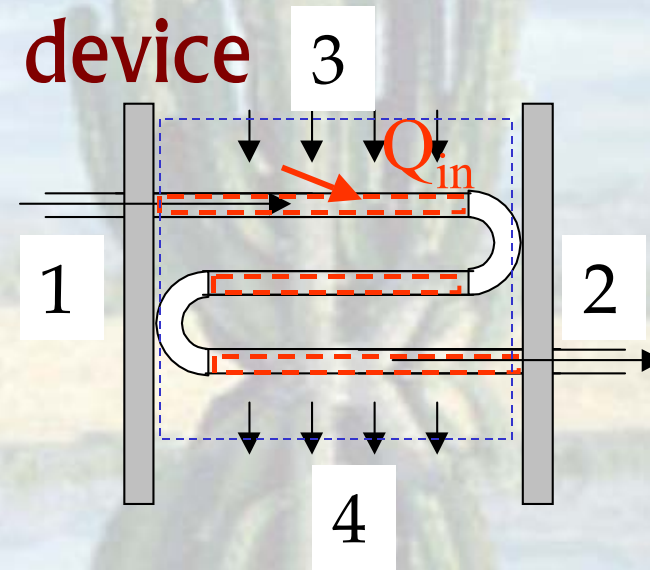
Heat exchanger:
Entropy Balance

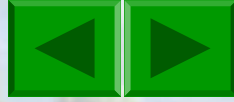
where $\dot{m}_4 = \dot{m}_3$ $\dot{m}_2 = \dot{m}_1$

Case 1

$$\dot{S}_{gen} = 0 - 0 + \dot{m}_4 s_4 - \dot{m}_3 s_3 + \dot{m}_2 s_2 - \dot{m}_1 s_1, \quad \frac{kW}{K}$$

$$\dot{S}_{gen} = \dot{m}_4 (s_4 - s_3) + \dot{m}_2 (s_2 - s_1), \quad \frac{kW}{K}$$

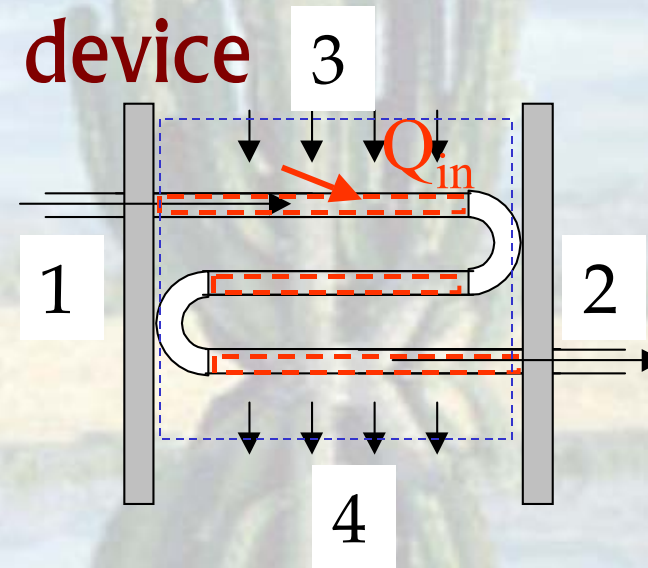




Review - Entropy Balance

Entropy Balance – Steady-flow device

Heat exchanger:
Entropy Balance

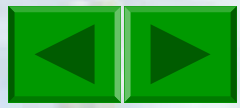


where $\dot{m}_4 = \dot{m}_3$ $\dot{m}_2 = \dot{m}_1$

Case 2

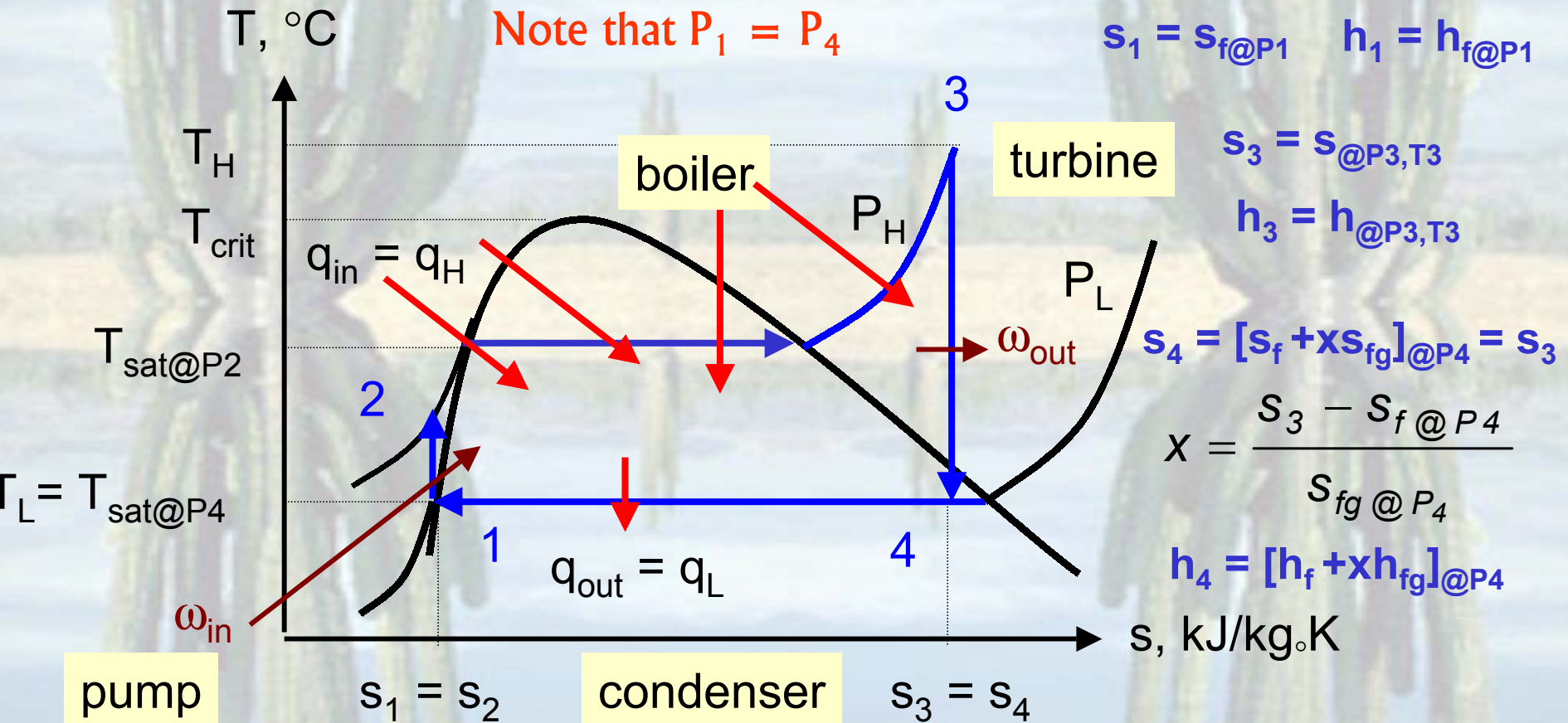
$$\dot{S}_{gen} = \frac{\dot{Q}_{out}}{T_{out}} - \frac{\dot{Q}_{in}}{T_{in}} + \dot{m}_2 s_2 - \dot{m}_1 s_1, \quad \frac{kW}{K}$$

$$\dot{S}_{gen} = 0 - \frac{\dot{Q}_{in}}{T_{in}} + \dot{m}_2 (s_2 - s_1), \quad \frac{kW}{K}$$

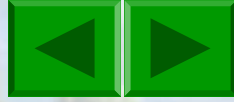


Vapor Cycle – Ideal Rankine Cycle

T- s diagram for an Ideal Rankine Cycle



$$h_2 = h_1 + v_2(P_2 - P_1); \text{ where } v_2 \cong v_1 = v_{f@P1}$$



Review – Ideal Rankine Cycle

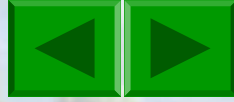
Energy Analysis

Efficiency

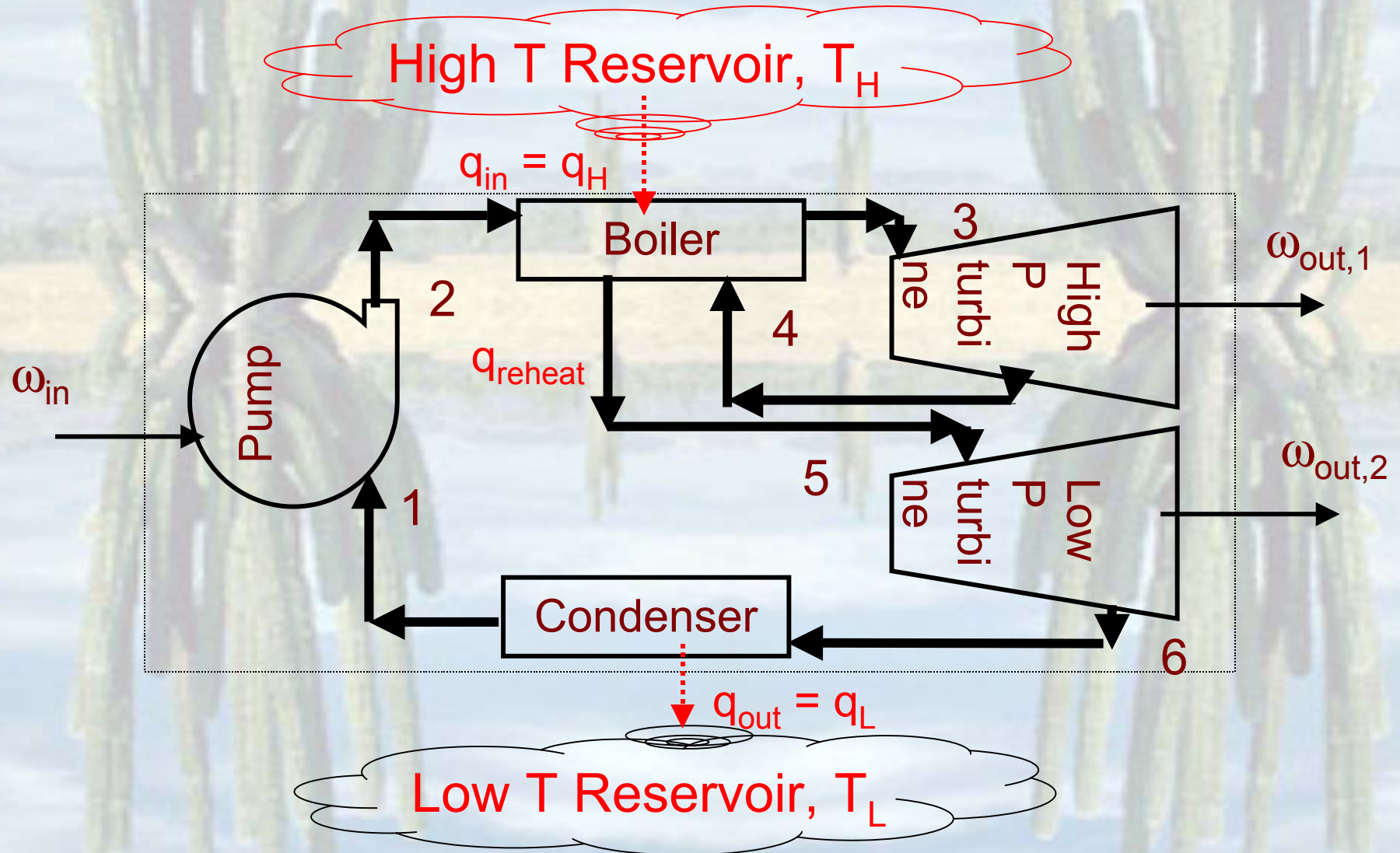
$$\eta = \frac{\omega_{net,out}}{q_{in}} = \frac{q_{in} - q_{out}}{q_{in}} = \frac{h_3 - h_2 - (h_4 - h_1)}{h_3 - h_2}$$

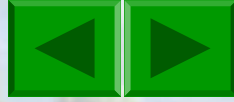
$$\eta = \frac{\omega_{net,out}}{q_{in}} = \frac{\omega_{out} - \omega_{in}}{q_{in}} = \frac{h_3 - h_4 - (h_2 - h_1)}{h_3 - h_2}$$

$$\eta = \frac{h_3 - h_4 - h_2 + h_1}{h_3 - h_2}$$



Review – Reheat Rankine Cycle



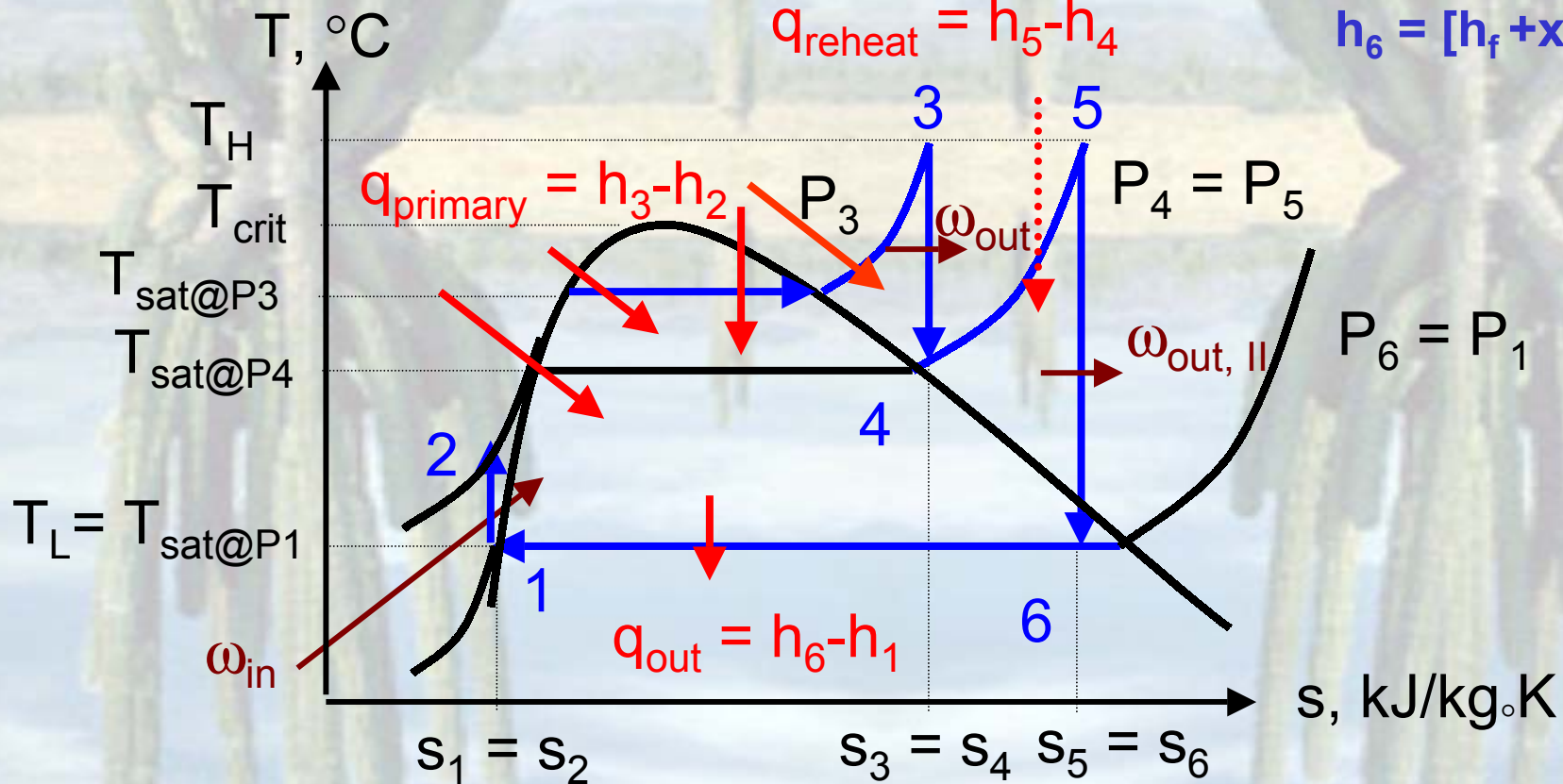


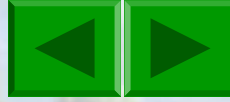
Review – Reheat Rankine Cycle

Reheating increases η and reduces moisture in turbine

$$s_6 = [s_f + x s_{fg}]_{@P_6} \text{ Use } x = 0.896 \text{ and } s_5 = s_6$$

$$h_6 = [h_f + x h_{fg}]_{@P_6}$$





Review – Reheat Rankine Cycle

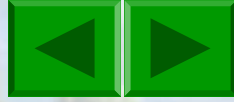
Energy Analysis

$$q_{in} = q_{primary} + q_{reheat} = h_3 - h_2 + h_5 - h_4 \quad q_{out} = h_6 - h_1$$

$$\omega_{net,out} = \omega_{out,1} + \omega_{out,2} = h_3 - h_4 + h_5 - h_6$$

$$\eta = \frac{\omega_{net,out}}{q_{in}} = \frac{q_{in} - q_{out}}{q_{in}} = \frac{h_3 - h_2 + h_5 - h_4 - (h_6 - h_1)}{h_3 - h_2 + h_5 - h_4}$$

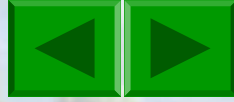
$$\eta = \frac{\omega_{net,out}}{q_{in}} = \frac{\omega_{out1} + \omega_{out2}}{q_{in}} = \frac{h_3 - h_4 + h_5 - h_6}{h_3 - h_2 + h_5 - h_4}$$



Gas Mixtures – Ideal Gases

Vapor power cycles – Rankine cycle

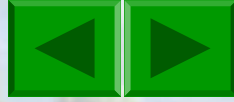
- Water as working fluid
 - ❖ cheap
 - ❖ Easily available
 - ❖ High latent heat of vaporisation, h_{fg} .
- Use property table to determine properties



Gas Mixtures – Ideal Gases

Non-reacting gas mixtures as working fluid

- Properties depends on
 - ❖ Components (constituents) of mixtures
 - ❖ Amount of each component
 - ❖ Volume of each component
- Pressure each component exerts on container walls
- Extended properties may not be tabulated
- Treat mixture as pure substances
- Examples: Air, CO_2 , CH_4 (methane),
 C_3H_8 (Propane)



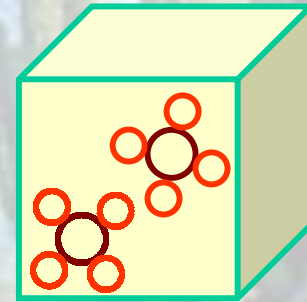
Gas Mixtures – Ideal Gases

Ideal Gases

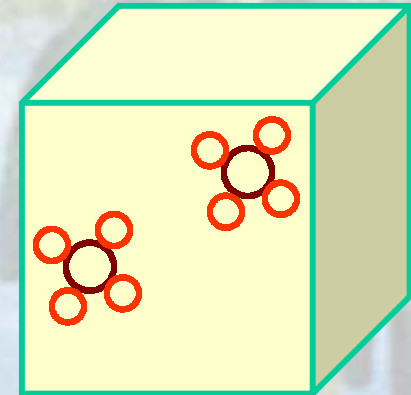
- Low density (mass in 1 m^3) gases
Molecules are further apart

- Real gases satisfying condition

$P_{\text{gas}} \ll P_{\text{crit}}; T_{\text{gas}} \gg T_{\text{crit}}$,
have low density and can be
treated as ideal gases

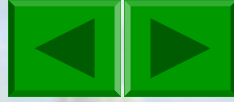


High density



Low density

Molecules far apart

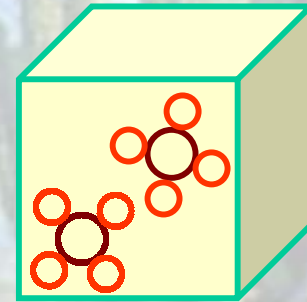


Gas Mixtures – Ideal Gases

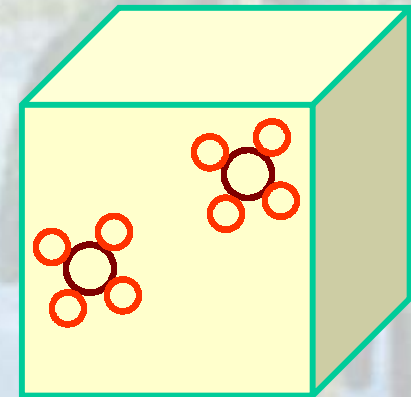
Ideal Gases

➤ **Equation of State** - P- v -T behaviour

$Pv = RT$ (energy contained by 1 kg mass) where v is the specific volume in m^3/kg , R is gas constant, $kJ/kg \cdot K$, T is absolute temp in Kelvin.

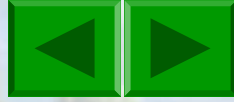


High density



Low density

Molecules far apart



Gas Mixtures – Ideal Gases

Ideal Gases

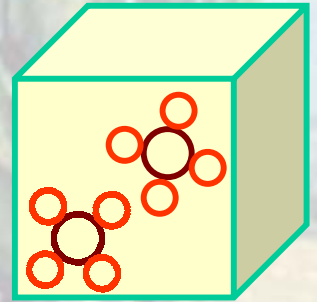
➤ **Equation of State** - P-v-T behaviour

$Pv = RT$, since $v = V/m$ then,

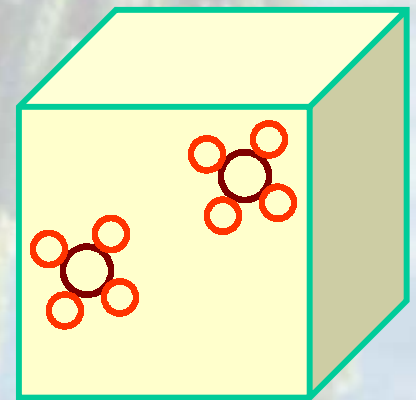
$P(V/m) = RT$. So,

$PV = mRT$, in $\text{kPa}\cdot\text{m}^3 = \text{kJ}$.

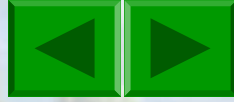
Total energy of a system.



High density



Low density



Gas Mixtures – Ideal Gases

Ideal Gases

➤ **Equation of State** - P-v-T behaviour

$$PV = mRT = NMRT = N(MR)T$$

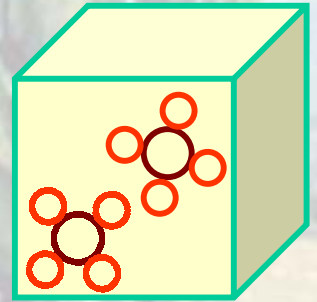
Hence, can also write $PV = NR_u T$
where

N is no of kilomoles, kmol,

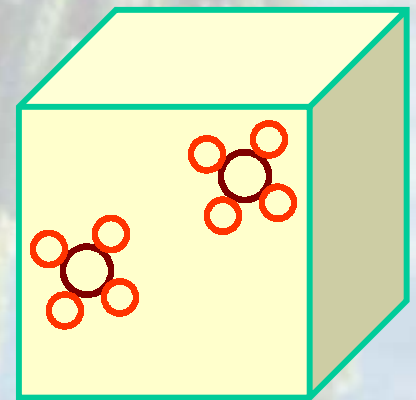
M is molar mass in kg/kmole and

R_u is universal gas constant; $R_u = MR$.

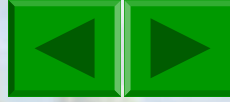
$$R_u = 8.314 \text{ kJ/kmol.K}$$



High density



Low density



Gas Mixtures – Ideal Gases

Ideal Gases

➤ Equation of State for mixtures

$$P_{\text{mix}} v_{\text{mix}} = R_{\text{mix}} T_{\text{mix}}, \quad P_{\text{mix}} V_{\text{mix}} = m_{\text{mix}} R_{\text{mix}} T_{\text{mix}}$$

$$P_{\text{mix}} V_{\text{mix}} = N_{\text{mix}} R_u T_{\text{mix}} \quad \text{where} \quad m_{\text{mix}} = M_{\text{mix}} N_{\text{mix}}$$

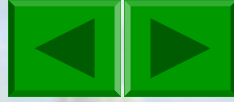
R_{mix} is apparent or mixture gas constant, kJ/kg.K,

T_{mix} is absolute temp in Kelvin, N_{mix} is no of kilomoles,

M_{mix} is molar mass of mixture in kJ/kmole and

R_u is universal gas constant; $R_u = MR$.

$$R_u = 8.314 \text{ kJ/kmol.K}$$



Gas Mixtures – Ideal Gases

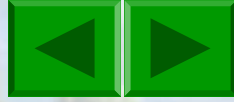
Composition of gas mixtures

- Specify by **mass (gravimetric analysis)**
or **volume (volumetric or molar analysis)**

mass = Molar mass * Number of kilomoles

Mass is $m = MN$, in kg

Number of kilomoles is $N = \frac{m}{M}$, in kmole



Gas Mixtures – Composition by Mass

Gravimetric Analysis

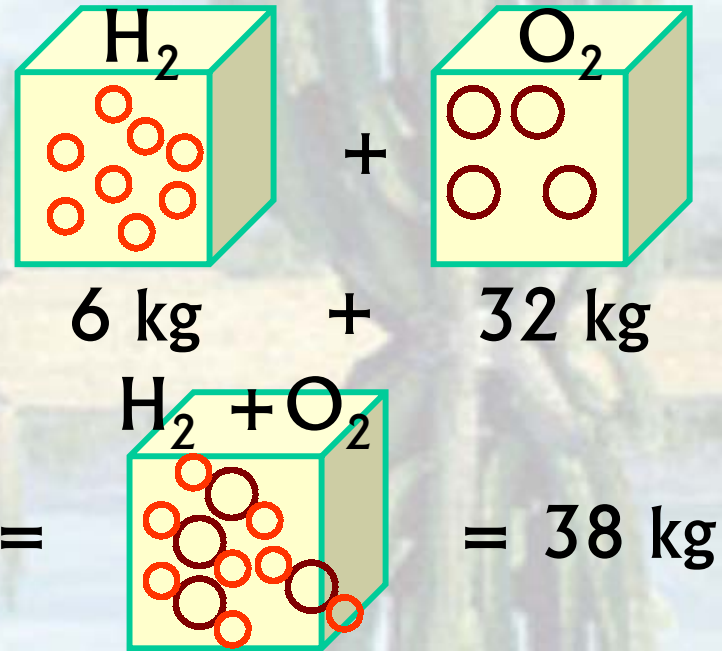
- Composition by weight or mass
- Mass of components add to the total mass of mixtures

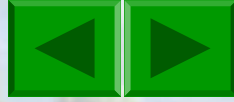
$$m_{mix} = m_{H_2} + m_{O_2}$$

Mass fraction of components

$$mf_{H_2} = \frac{m_{H_2}}{m_{mix}} = \frac{6}{38} = 0.1579 = 15.8\%$$

$$mf_{O_2} = \frac{m_{O_2}}{m_{mix}} = \frac{32}{38} = 0.8421 = 84.2\%$$





Gas Mixtures – Composition by Moles

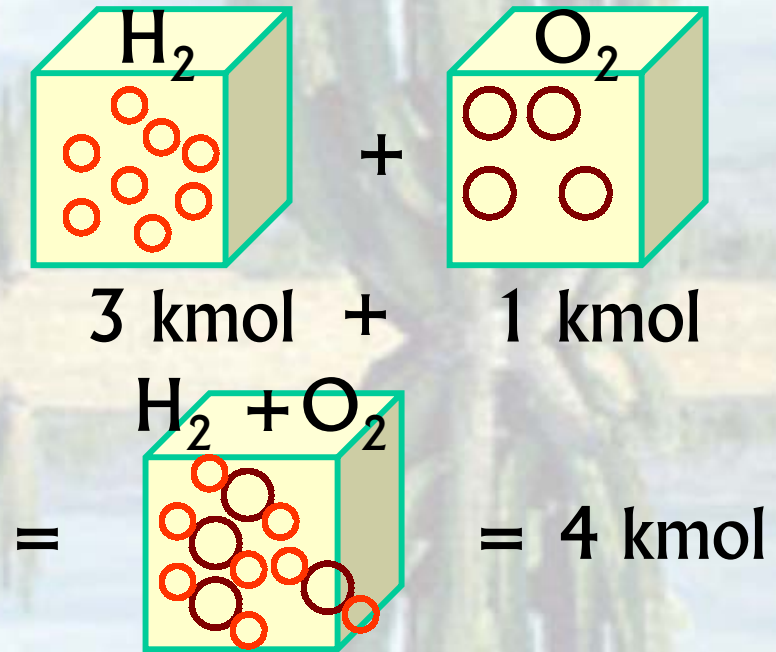
Volumetric Analysis

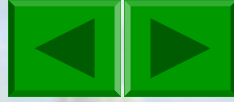
- Composition by kilomoles
- Number of kilomoles of components add to the total number of kilomoles of mixtures

$$N_{mix} = N_{H_2} + N_{O_2}$$

Number of kilomoles is $N = \frac{m}{M}$

Hence,
$$\left(\frac{m}{M}\right)_{mix} = \left(\frac{m}{M}\right)_{H_2} + \left(\frac{m}{M}\right)_{O_2}$$





Gas Mixtures – Composition by Moles

Volumetric Analysis

Mole fraction of components

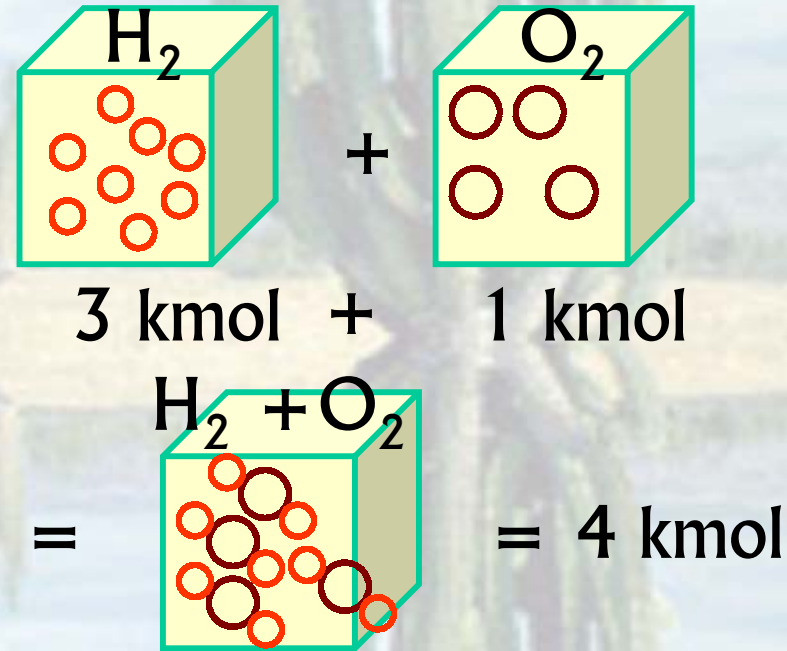
$$N_{H_2} = \frac{6 \text{ kg}}{2 \text{ kg/ kmol}} = 3 \text{ kmol}$$

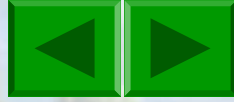
$$N_{O_2} = \frac{32 \text{ kg}}{32 \text{ kg/ kmol}} = 1 \text{ kmol}$$

Hence

$$y_{H_2} = \frac{N_{H_2}}{N_{mix}} = \frac{3}{4} = 0.75 = 75 \%$$

$$y_{O_2} = \frac{N_{O_2}}{N_{mix}} = \frac{1}{4} = 0.25 = 25 \%$$





Gas Mixtures – Composition by Moles

Composition Summary

Gravimetric Analysis

$$m_1 + m_2 = m_{mix}$$

$$mf_1 + mf_2 = 1 \text{ or } 100\%$$

where $mf_i = \frac{m_i}{m_{mix}}$

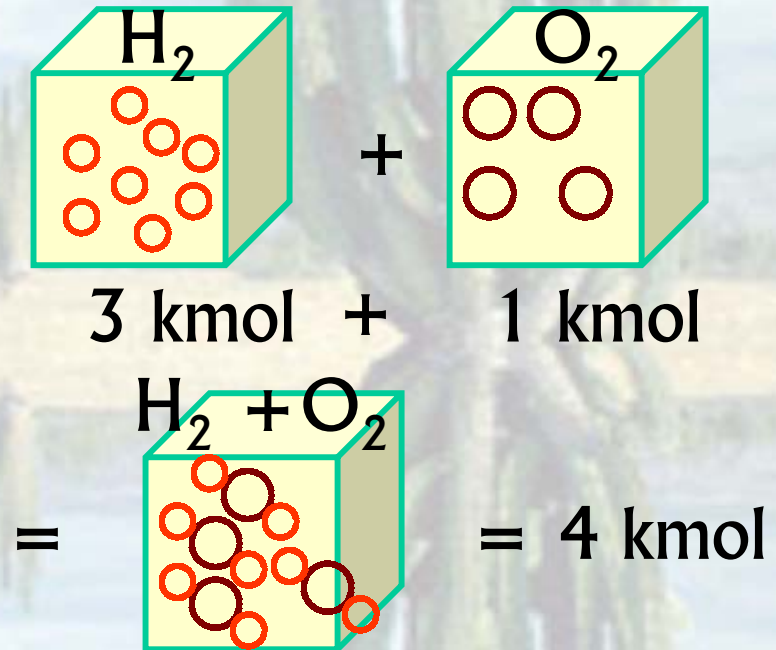
Volumetric Analysis

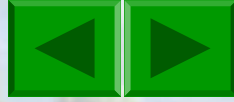
$$N_1 + N_2 = N_{mix}$$

$$y_1 + y_2 = 1 \text{ or } 100\%$$

where

$$y_i = \frac{N_i}{N_{mix}}$$

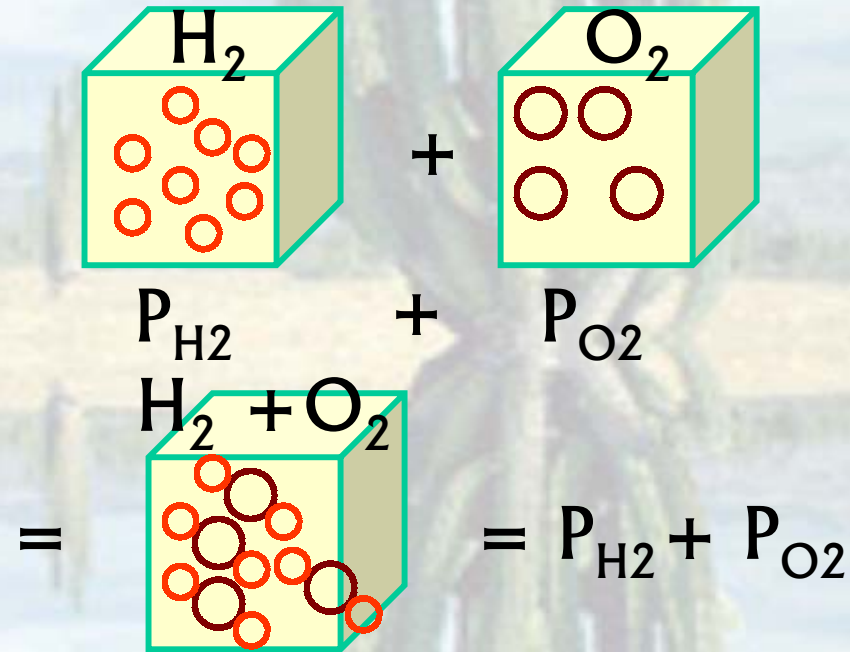




Gas Mixtures – Additive Pressure

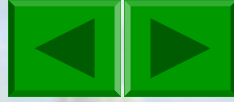
Dalton's Law

➤ The total pressure exerted in a container at volume V and absolute temperature T , is the sum of component pressure exerted by each gas in that container at V, T .



$$P_{mix} = P_1 + P_2$$

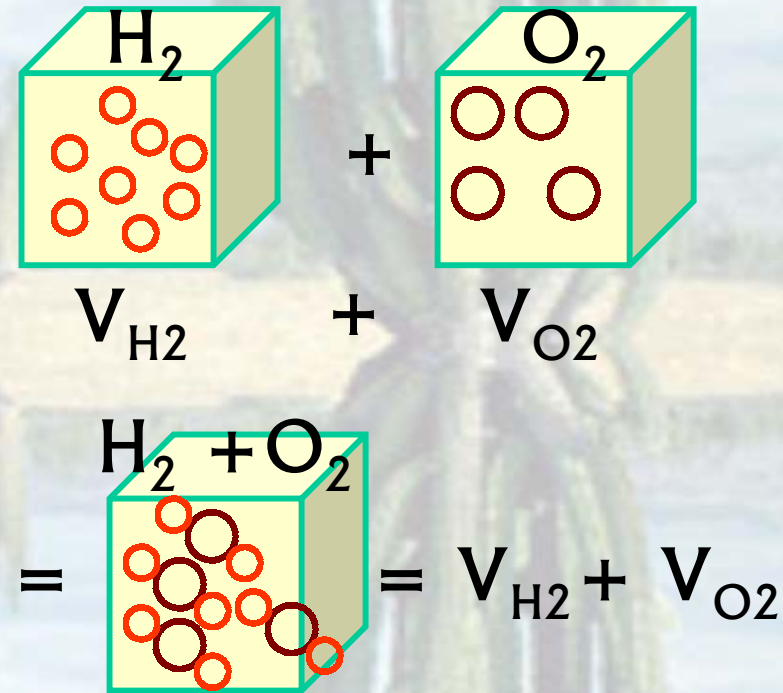
$$P_{mix} = \sum_{i=1}^k P_i (V_{mix}, T_{mix}); k \text{ is total number of components}$$



Gas Mixtures – Additive Volume

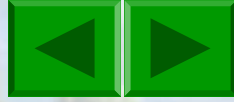
Amagat's Law

➤ The total volume occupied in a container at pressure P_{mix} and absolute temperature T_{mix} , is the sum of component volumes occupied by each gas in that container at P_{mix} , T_{mix} .



$$V_{mix} = V_1 + V_2$$

$$V_{mix} = \sum_{i=1}^k V_i (P_{mix}, T_{mix}); k \text{ is total number of components}$$



Gas Mixtures – Pressure Fraction

Partial Pressure

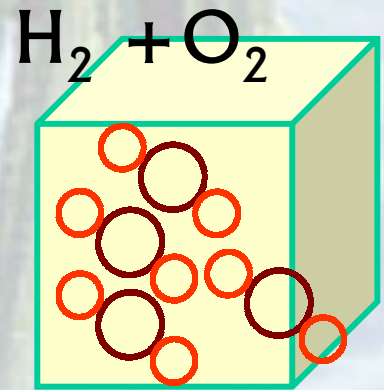
Since $P_{mix} V_{mix} = N_{mix} R_U T_{mix}$; $P_1 V_{mix} = N_1 R_U T_{mix}$

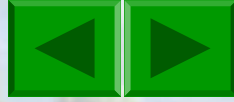
➤ The pressure fraction for each gas inside the container is

$$\frac{P_1}{P_{mix}} = \frac{N_1 R_u \frac{T}{V}}{N_{mix} R_u \frac{T}{V}} = \frac{N_1}{N_{mix}} = y_1$$

Hence the partial pressure is $P_1 = y_1 P_{mix}$

In general, $P_i = y_i P_{mix}$





Gas Mixtures – Volume Fraction

Partial Volume

Since $P_{mix} V_{mix} = N_{mix} R_U T_{mix}$; $P_{mix} V_1 = N_1 R_U T_{mix}$

➤ The volume fraction for each gas inside the container is

$$\frac{V_1}{V_{mix}} = \frac{N_1 R_u \frac{T}{P}}{N_{mix} R_u \frac{T}{P}} = \frac{N_1}{N_{mix}} = y_1$$

Hence the partial volume is

$$V_1 = y_1 V_{mix}$$

In general, $V_i = y_i V_{mix}$

