

# TOPIC II - GAS PROPERTIES

## Perfect/Ideal gases ( Lecture 1/2)

Establish laws for  $p$ ,  $V$ ,  $T$ ,  $U$  relationships

### Contents:

1. Definition of a Property
2. Perfect (Ideal) Gas Law
3. Internal Energy

Recall **state postulate** – two independent properties determine equilibrium state.

E.g.  $p$ ,  $T$  allow us to find  $\rho$ ,  $u$ ,  $h$ ,  $s$

**equilibrium** - all driving forces acting on system are balanced

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### Perfect/Ideal gases ( Lecture 1/2)

#### 1) Definition of a Property

Describes a state of equilibrium –  
independent of path

**independent properties** can be combined

$$\rho = m / V \quad (1)$$

$$\bar{v} = V / m = 1 / \rho \quad (2)$$

**extensive** properties ( $V$ ,  $m$ , ...) – depend on  
system size

**intensive** properties ( $T$ ,  $p$ ,  $\bar{v}$ , ...) do not.  
Specific properties refer to a unit mass. Three  
other intensive properties:-

$u$  – internal energy (today)

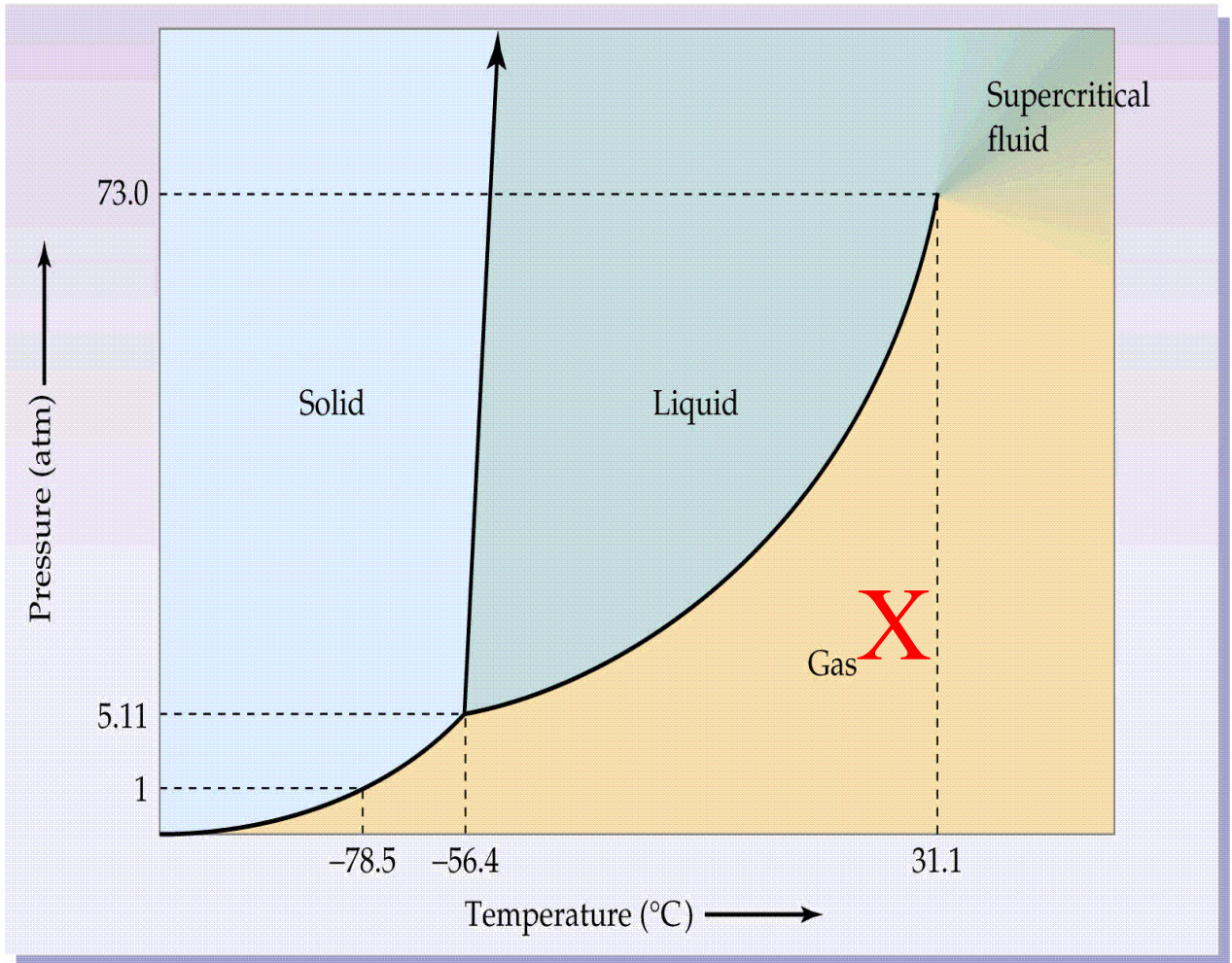
$h$  – enthalpy (next lecture)

$s$  – entropy (much later)

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## Perfect/Ideal gases ( Lecture 1/2)

Conditions that interest us:



Perfect gases where molecule-to-molecule forces are very small. (See location X).

Air is usually far from triple and critical point.

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## Perfect/Ideal gases ( Lecture 1/2)

### 2. Perfect (Ideal) Gas Law

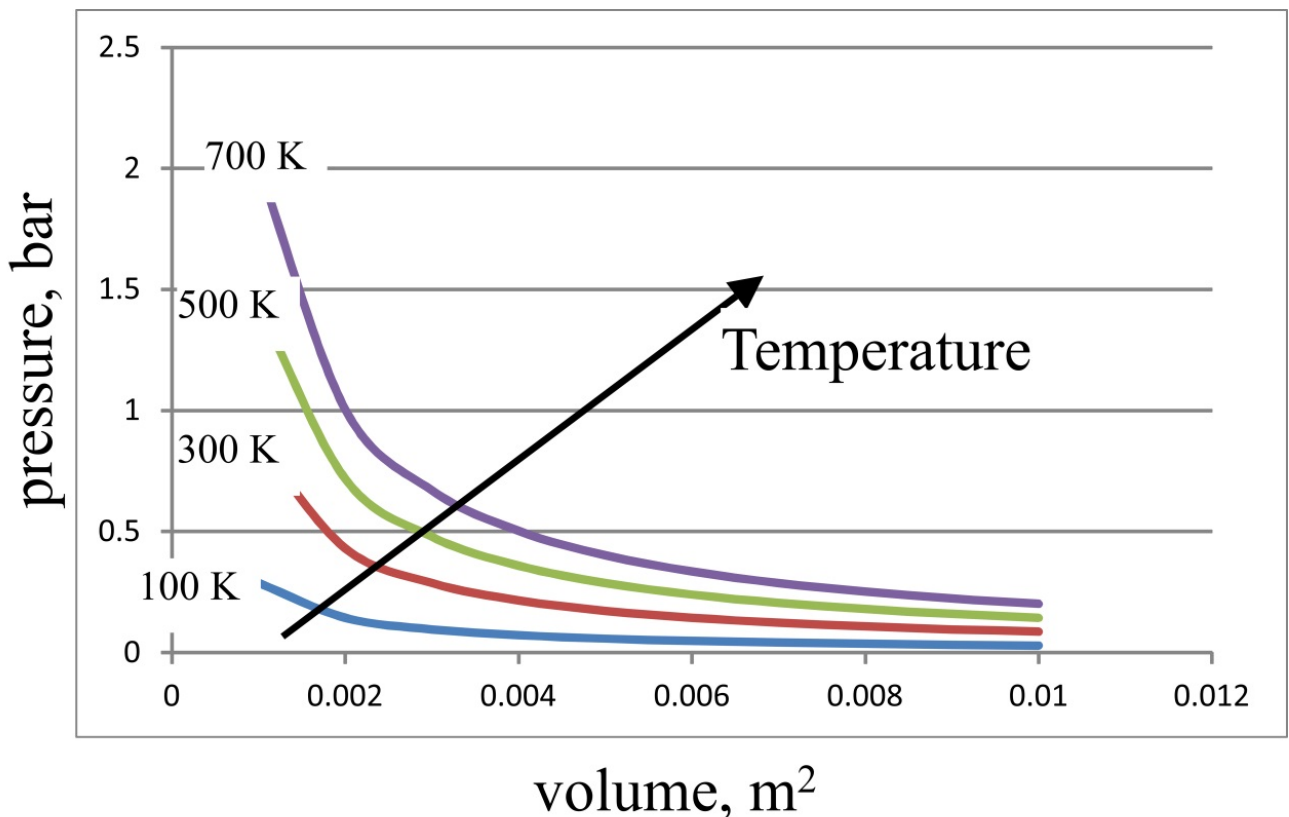
Gives ideal (or perfect) gas law.

Engineering forms:

$$p \bar{v} = R T \quad (3)$$

$$pV = mRT \quad (3b)$$

$$\frac{p_1 V_1}{T_1} = \frac{p_2 V_2}{T_2} \quad (3c)$$



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### Perfect/Ideal gases ( Lecture 1/2)

$R$  is the specific gas constant ( $\text{kJ kg}^{-1} \text{K}^{-1}$ ),  
following from universal gas constant  
( $\text{kJ kmol}^{-1} \text{K}^{-1}$ )

$$R = \frac{\tilde{R}}{m} \quad (4)$$

If the amount of substance (in kmol) is

$$n = m / \tilde{m}$$

then

$$pV = n \tilde{R} T \quad (5)$$

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### Perfect/Ideal gases ( Lecture 1/2)

*Example, air density at atmospheric pressure and 25°C*

$$R = \frac{\tilde{R}}{\tilde{m}} = \frac{8.314 \text{ kJ kmol}^{-1} \text{ K}^{-1}}{29 \text{ kg kmol}^{-1}} = 0.287 \text{ kJ kg}^{-1} \text{ K}^{-1}$$
$$\rho = \frac{1}{\sqrt{RT}} = \frac{p}{RT} = \frac{(1.013 \text{ bar} \times 100 \text{ kN m}^{-2} \text{ bar}^{-1})}{0.287 \text{ kJ kg}^{-1} \text{ K}^{-1} \times (273 + 25) \text{ K}} =$$
$$1.184 \frac{\text{kN m}^{-2}}{\text{kJ kg}^{-1}} = 1.184 \text{ kg m}^{-3}$$

Conversion factors yield engineering units (kN, kJ etc). E.g.

$$\frac{100 \text{ kN m}^{-2}}{1 \text{ bar}} \equiv 1$$

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### Perfect/Ideal gases ( Lecture 1/2)

#### 3. Internal Energy

Properties of steam (non-ideal) are tabulated against  $T$ ,  $p$ . Air properties against  $T$  only.

**Ideal gas** => perfect gas &  $U = f(T)$  only

$$U = m c_v (T - T_o) \quad (6)$$

$$u = U/m = c_v (T - T_o) \quad (6b)$$

$T_o$  is a **datum temperature**.

proportionality  
constant

If  $U$  is measured in a rigid vessel heat addition is

$$Q_v = U_2 - U_1 = m c_v (T_2 - T_1) \quad (7)$$

provided  $V = \text{constant}$  and  $W = 0$

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### Perfect/Ideal gases ( Lecture 1/2)

**Definition:** The **specific heat capacity** is the quantity of energy required to raise the temperature of one unit mass by one degree of temperature. Term  $c_v$  refers to such energy addition at constant volume.

It has units of  $\text{kJ kg}^{-1} \text{K}^{-1}$ .

**air**  $c_v = 0.718 \text{ kJ kg}^{-1} \text{K}^{-1}$  or  $c_v = 2.5 R$

Because  $U$  is a property:

$c_v$  can always be used to get  $U$

$Q$ ,  $W$  then follow from NFEE

$c_v$  only yields  $Q$  when  $W = 0$

Note – energy can cross boundaries as:

.....



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#### Conclusions:

This year – focus on pure substances with no change of phase.

Ideal– weak intermolecular forces and hence  
 $pV = m R T$  and  $U = m c_v (T - T_o)$

Engineers tend to prefer mass quantities to molar quantities ( $R$  vs  $\tilde{R}$  ,  $m$  vs  $n$ )

Definition of specific heat capacity – depends on process (today we considered constant volume)

Think of  $c_v$  as an indicator of internal energy.