THE IDEAL-GAS EQUATION

- Any equation that relates the pressure, temperature, and specific volume of a substance is referred to as an equation of state
- Property relations that involve other properties of a substance at equilibrium states are also referred to as equations of state
- The simplest and best known equation of state is the ideal-gas equation of state, which predicts the P-v-T behavior of a gas

or

- · Any gas that obeys this relationship is referred to as an ideal gas
- In order to use this equation, all values

• The gas constant R is is unique for each gas and is determined from:

- R_{II} is the universal gas constant and M is the molar mass
- The values for R can be found for common gases in Table A-1 and A-2 of the textbook

Substance	R, kJ/kg⋅K
Air	0.2870
Helium Argon	2.0769 0.2081
Nitrogen	0.2968
	• • •

$$R_{u} = \begin{cases} 8.31447 \text{ kJ/kmol} \cdot \text{K} \\ 8.31447 \text{ kPa} \cdot \text{m}^{3}/\text{kmol} \cdot \text{K} \\ 0.0831447 \text{ bar} \cdot \text{m}^{3}/\text{kmol} \cdot \text{K} \\ 1.98588 \text{ Btu/lbmol} \cdot \text{R} \\ 10.7316 \text{ psia} \cdot \text{ft}^{3}/\text{lbmol} \cdot \text{R} \\ 1545.37 \text{ ft} \cdot \text{lbf/lbmol} \cdot \text{R} \end{cases}$$

 The mass of a system is equal to the product of its molar mass M and the mole number N

$$m = MN$$

 It follows that the ideal gas equation of state can be written in several different forms:

$$PV = mRT$$

 $PV = NR_uT$
 $Pv = RT$

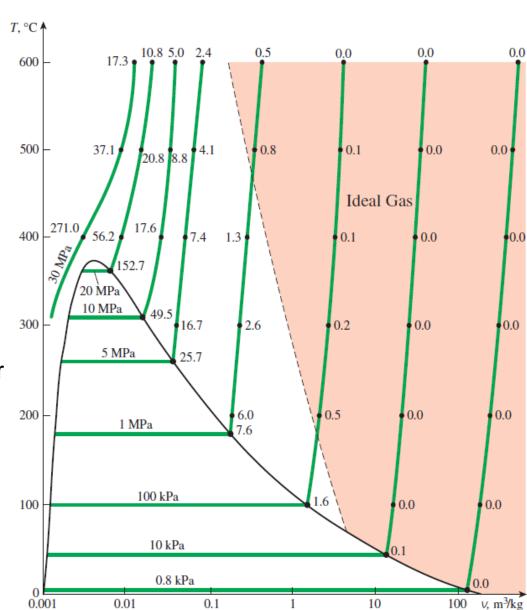
It follows that for a fixed mass:

$$\frac{PV}{T} = Constant$$

- An ideal gas is an imaginary substance that obeys the ideal gas relationship
- It has been experimentally observed that the ideal-gas relation closely approximates real gases at *low densities*
- Low densities are encountered when a gas is at ______

 or _____
 - Familiar gases such as air, nitrogen, oxygen, hydrogen, helium, argon, krypton and carbon dioxide can <u>often</u> be treated as an ideal gas with negligible error
- However, dense gases such as <u>water vapor</u> in steam power applications and <u>refrigerant vapor</u> should not be treated as ideal gases -> _____

- At pressures below 10 kPa, water vapor can be treated as an ideal gas, regardless of its temperature, with negligible error (less than 0.1 percent).
- At higher pressures, however, the ideal gas assumption yields unacceptable errors, particularly in the vicinity of the critical point and the saturated vapor line.
- In some instances, such as airconditioning applications, the water vapor in the air can be treated as an ideal gas since the pressure of the water vapor is very low
- However, for this class, <u>for</u>
 <u>properties of water and R-134a</u>
 <u>always</u>



• The deviation of a gas from ideal-gas behavior can be easily accounted for by introducing the compressibility factor, Z:

· where

o v_{ideal} can be calculated from the ideal-gas relationship:

$$v_{ideal} = \frac{RT}{P}$$

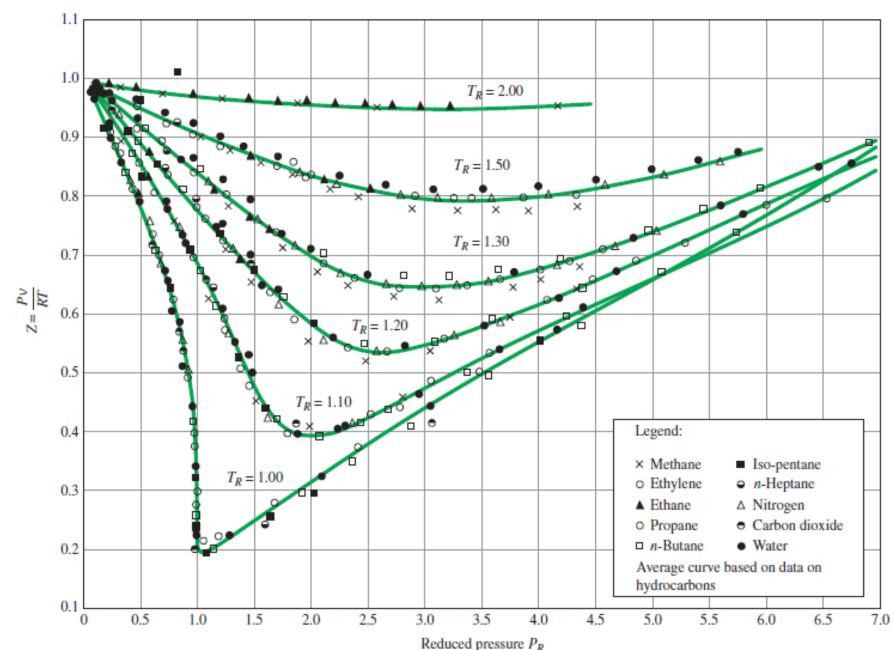
 The farther Z is from a value of 1, the more the gas deviates from ideal gas behavior

- It has been mentioned previously that the ideal-gas equation is followed by gases at low pressure and high temperatures
 - For example, at -100°C and atmospheric pressure, air can be treated as an ideal gas with under 1% error
- We define what counts as a "high" or "low" temperature / pressure by introducing a normalization
 - The reduced pressure is defined as:

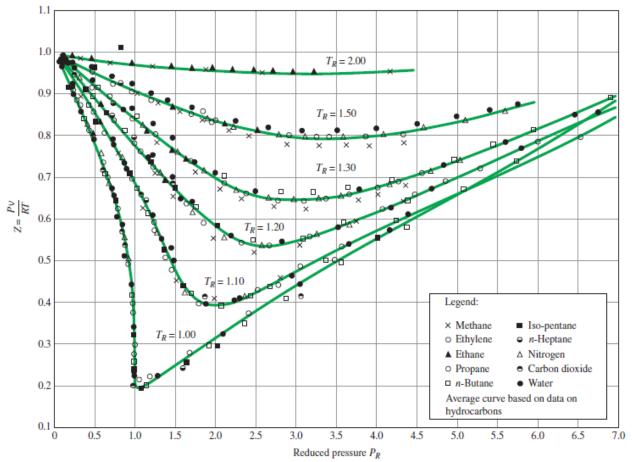
The reduced temperature is defined as:

- For the reduced pressure and temperature, <u>absolute units must be used!</u>
- The Z factor for all gases is approximately the same at the same reduced pressure and temperature

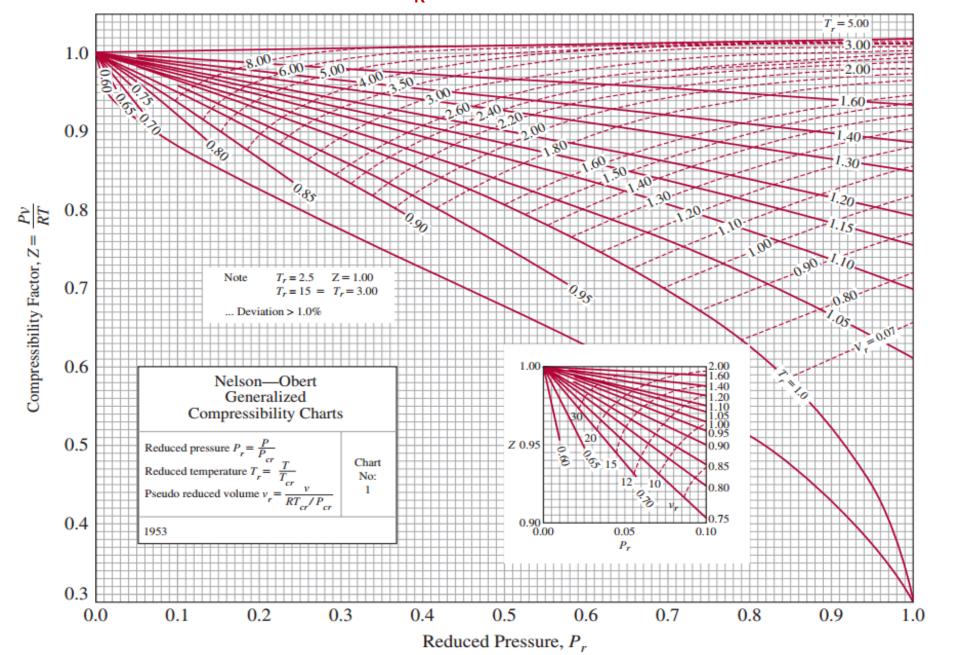
GENERALIZED COMPRESSIBILITY CHART – EXPERIMENTAL DATA



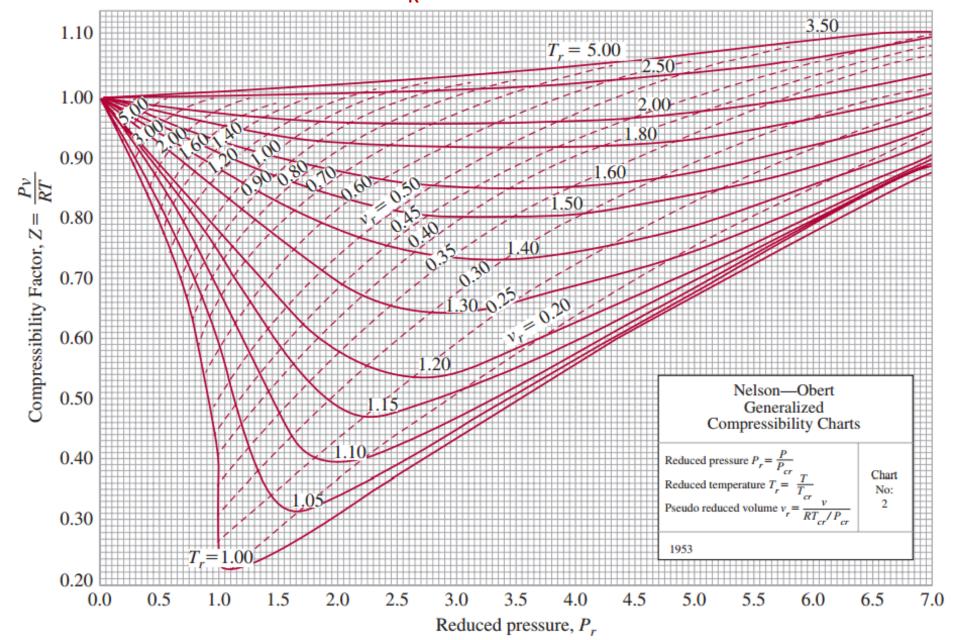
- The following observations can be made from the generalized compressibility chart
 - At very low pressures (P_R______), gases behave as an ideal gas regardless of temperatures (*in general – not always*)
 - At high temperatures (T_R ______), ideal gas behavior can be assumed with good accuracy regardless of pressure (in general not always)



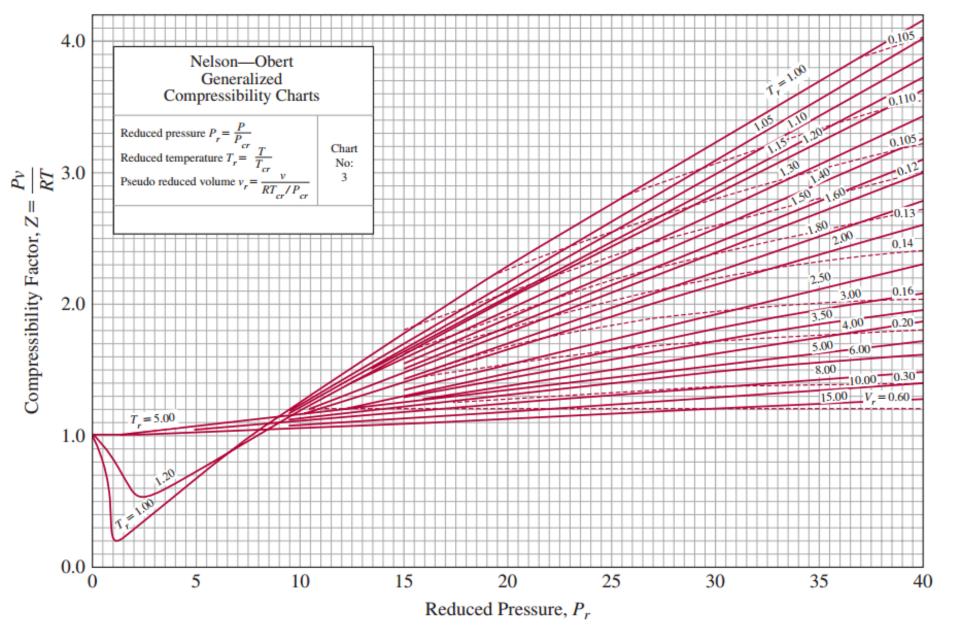
COMPRESSIBILITY FACTOR FOR $P_R = 0$ to 1



COMPRESSIBILITY FACTOR FOR $P_R = 0$ to 7



COMPRESSIBILITY FACTOR FOR $P_R = 0$ to 40



In order to use the compressibility chart when P and v or T and v are given instead
of P and T, a pseudo-reduced specific volume is defined as:

- Unlike the definitions of P_R and T_R , v_R is related to both P_{CR} and T_{CR}
- Lines of constant v_R are added to the compressibility chart to enable use of v_R

Example 5-1:

Determine the specific volume of nitrogen gas at 10 MPa and –123°C using (a) the ideal-gas equation (b) the compressibility chart. Determine the error involved with (a) and (b) by comparing to the experimentally measured value of 0.002388 m³/kg.

• Example 5-2:

Methane at 1450 psia and 80°F is heated at constant pressure until its volume has increased by 80%. Determine the final temperature using (a) the ideal gas equation and (b) the compressibility factor.

Example 5-3:

5 kg of Ethylene undergoes a constant pressure process from an initial pressure and temperature of 4.1 MPa and 310 K, respectively, to a final temperature of 775 K. Determine the change in the total internal energy of the system if 2000 kJ of heat is transferred into the system.

OTHER EQUATIONS OF STATE

- Other equations of state have also been formulated, although they are generally more complex:
 - Van der Waals Equation of State

$$\left(P + \frac{a}{V^2}\right)(V - b) = RT$$
 $a = \frac{27R^2T_{\text{cr}}^2}{64P_{\text{cr}}}$ $b = \frac{RT_{\text{cr}}}{8P_{\text{cr}}}$

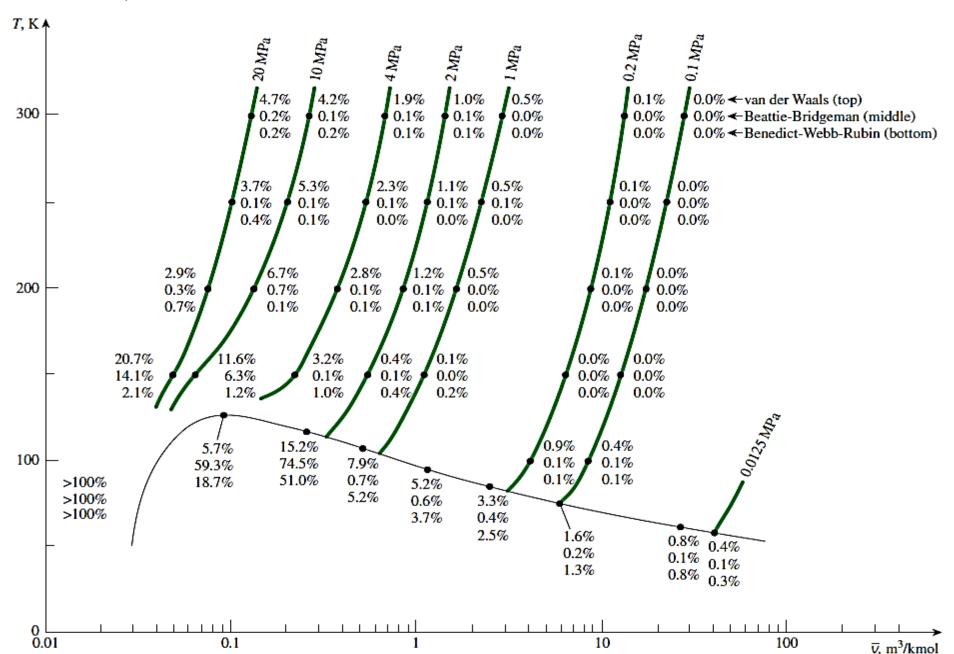
Beattie-Bridgeman Equation of State

$$P = \frac{R_u T}{\overline{v}^2} \left(1 - \frac{c}{\overline{v} T^3} \right) (\overline{v} + B) - \frac{A}{\overline{v}^2}$$
$$A = A_0 \left(1 - \frac{a}{\overline{v}} \right) B = B_0 \left(1 - \frac{b}{\overline{v}} \right)$$

Benedict-Webb-Rubin Equation of State

$$P = \frac{R_u T}{\overline{V}} + \left(B_0 R_u T - A_0 - \frac{C_0}{T^2} \right) \frac{1}{\overline{V}^2} + \frac{b R_u T - a}{\overline{V}^3} + \frac{a \alpha}{\overline{V}^6} + \frac{c}{\overline{V}^3 T^2} \left(1 + \frac{\gamma}{\overline{V}^2} \right) e^{-\gamma/\overline{V}^2}$$

OTHER EQUATIONS OF STATE - COMPARISON FOR NITROGEN



COMPRESSIBILITY - CORRECTION FOR H AND U

