# Introduction to the Natural Gas Processing Plant

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#### **Natural Gas**

- A typical flow scheme for a cryogenic natural gas processing plant is to extract ethane and heavier hydrocarbons from natural gas.
- Methods are shown as an example for rigorous modeling of ethane recovery plant using turbo-expander, refrigeration system, demethanizer and deethanizer columns.



#### **Natural Gas**

- ☐ Mainly composed of methane (>95%)
- Contain significant amounts of nitrogen and carbon dioxide.
- According to CO<sub>2</sub> content, natural gas system is classified into lean or rich gas.

#### **Needs for Studying Gas Plant**

- No special licenses for natural gas processing plants
- Not so much difficult since it is only consisted of fractionators (no reactors).

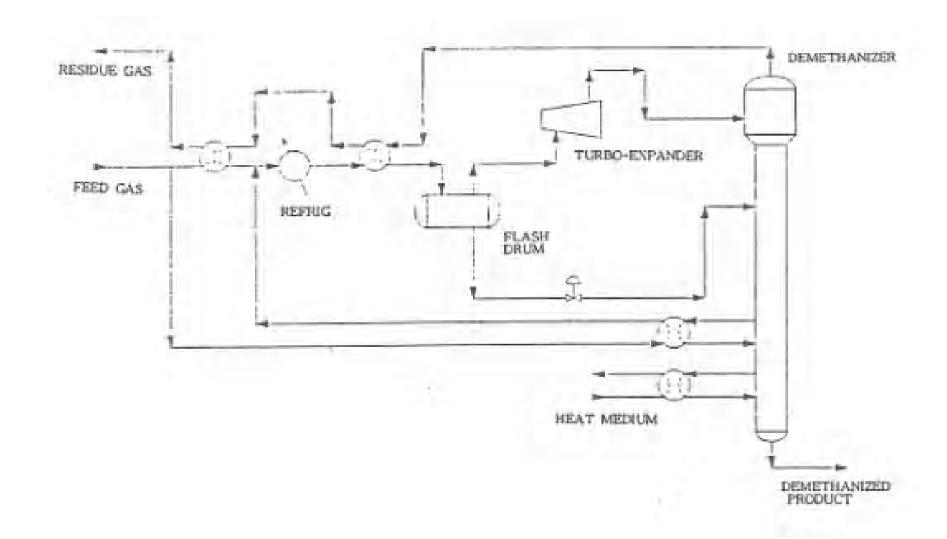


#### **Process Description**

- Gas is dehydrated usually by molecular sieve.
- ☐ Gas is partially chilled by back exchange with residue gas stream.
- ☐ Gas is further chilled by refrigeration system.
- ☐ Gas is partially separated into gas-liquid two phases.
- Gas is further chilled by demethanizer overhead vapor stream.
- Extremely low temperature stream is obtained by letting down the pressure using turbo-expander.
- Ethane and heaviers are obtained by fractionation.



#### **Schematic Diagram of Natural Gas Plant**





#### **Objectives**

- Introduction to typical scheme for the natural gas processing plant
- Application of modified cubic equation of state with an appropriate alpha function and new mixing rules to the natural gas system



#### CO<sub>2</sub>-rich System

Carbon dioxide systems are both of academic and industrial interests because....

#### **Academic Interests**

■ Even though there is no permanent dipole moment in CO₂, large quadrupole moment, which makes it difficult to predict the molecular interactions between CO₂ and other hydrocarbons. (Induced dipole)

#### **Industrial Interests**

It has been also necessary to use CO<sub>2</sub>-rich natural gas according to the increasing demands for usage of natural gas.

### **Typical Natural Gas Composition (Lean gas)**

Component	Mole Percent
Nitrogen	0.106
Carbon dioxide	0.303
Methane	95.600
Ethane	2.361
Propane	0.862
I-Butane	0.195
N-Butane	0.218
Pentane Plus	0.430



### **Dipole Moment**

Component	Dipole Moment
HF	1.910
KBr	9.070
H2O	1.840
NH3	1.470
CO2	0.000
CH4	0.000

### **Quadrupole Moment**

Component	Quadrupole Moment
H2	0.660
C2H2	3.000
C6H6	-3.600
H2O	-3.000
NH3	-1.000
CO	-2.500
CO2	-4.300

### **Why Testing Ternary Systems?**

- ☐ The ultimate goal of future thermodynamics is to predict multi-component phase behavior with pure component experimental information only, but is is still long way to go.
- The objective of modern thermodynamics is to correlate multi-component VLE or LLE with pure and binary experimental data only.

### **Testing Ternary Phase Behavior**

- N₂ and CO₂ systems are very important even though natural gas contains only small amounts.

### Some significant developments in cubic equations of state

- Functional form:
  - To predict liquid phase density accurately
- The alpha forms:
  - To predict pure component vapor pressure vs. temperature accurately
- ☐ The mixing rules:
  - To predict mixture phase behavior more accurately



#### The equations of state formulations

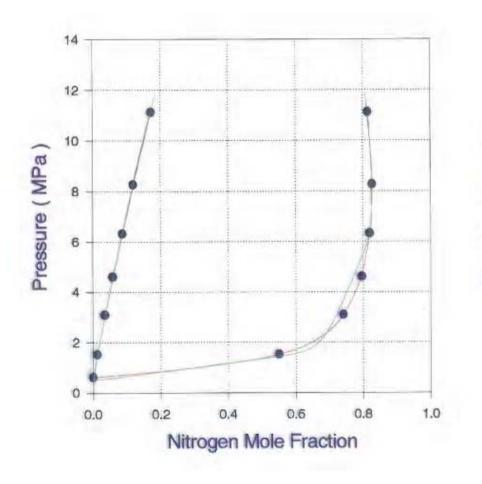
- Functional form is important for the liquid phase density calculations.
- ☐ The van der Waals (1873)
- □ Redlich-Kwong (1949)
- □ Soave-Redlich-Kong (1972)
- Peng-Robinson (1976)

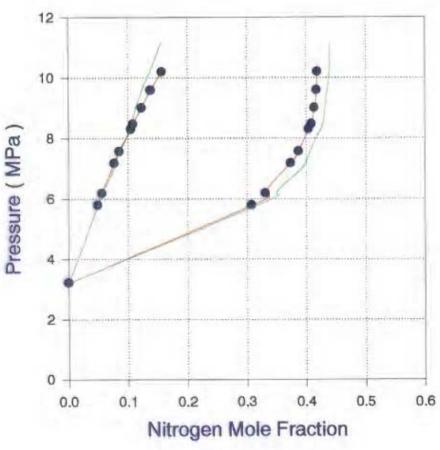
#### **Two-parameter Cubic Equation of State**

- Any two parameter cubic equation of state is notorious for not being able to correlate the liquid density well since the critical compressibility factor is constant for all pure components.
- $\square$  Z<sub>c</sub> in vdw EOS is 0.3750.
- $\Box$  Z<sub>c</sub> in RK EOS is 0.3333.
- $\square$  Z<sub>c</sub> in PR EOS is 0.3074.
- Experimental values for critical compressibility factor are 0.2880 for methane, 0.2840 for ethane and 0.2800 for propane.

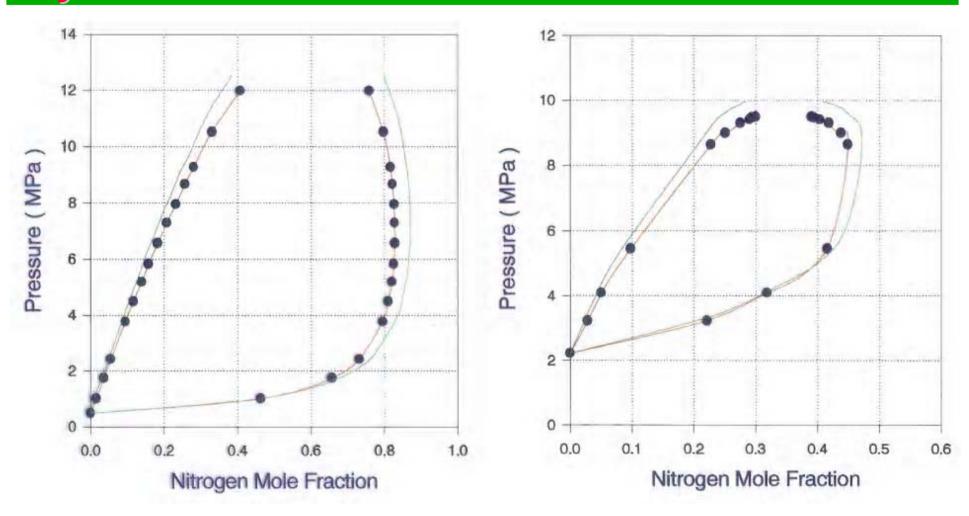


## Comparison of Mixing Rule 1 & 2 for N2+CO2 System at 220K & 270K

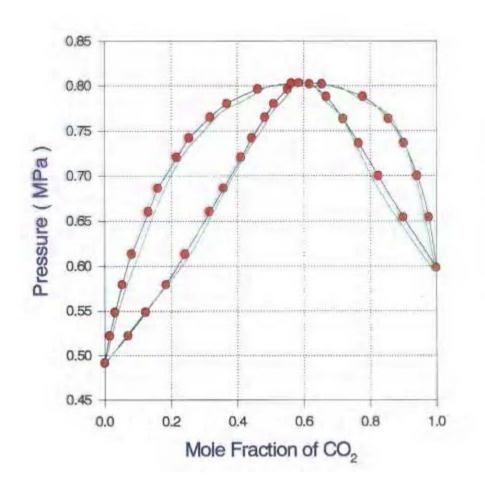


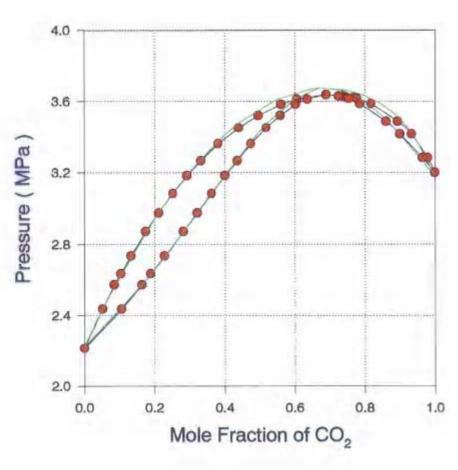


## Comparison of Mixing Rule 1 & 2 for N2+C2 System at 220K & 270K



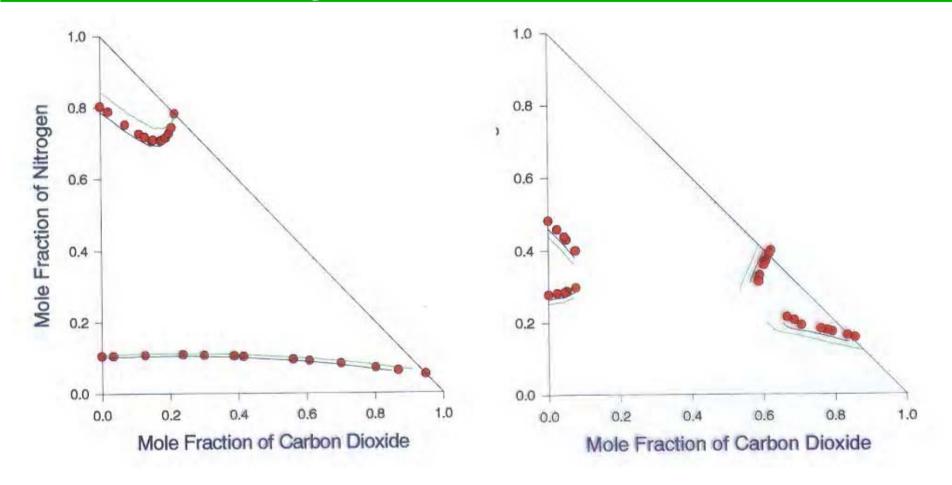
## Comparison of Mixing Rule 1 & 2 for CO2+C2 System at 220K & 270K







## Comparison of Mixing Rule 1 & 2 for CO2+N2+C2 System at 220K & 270K



### The End....

