SIFAT VOLUMETRIK FLUIDA MURNI

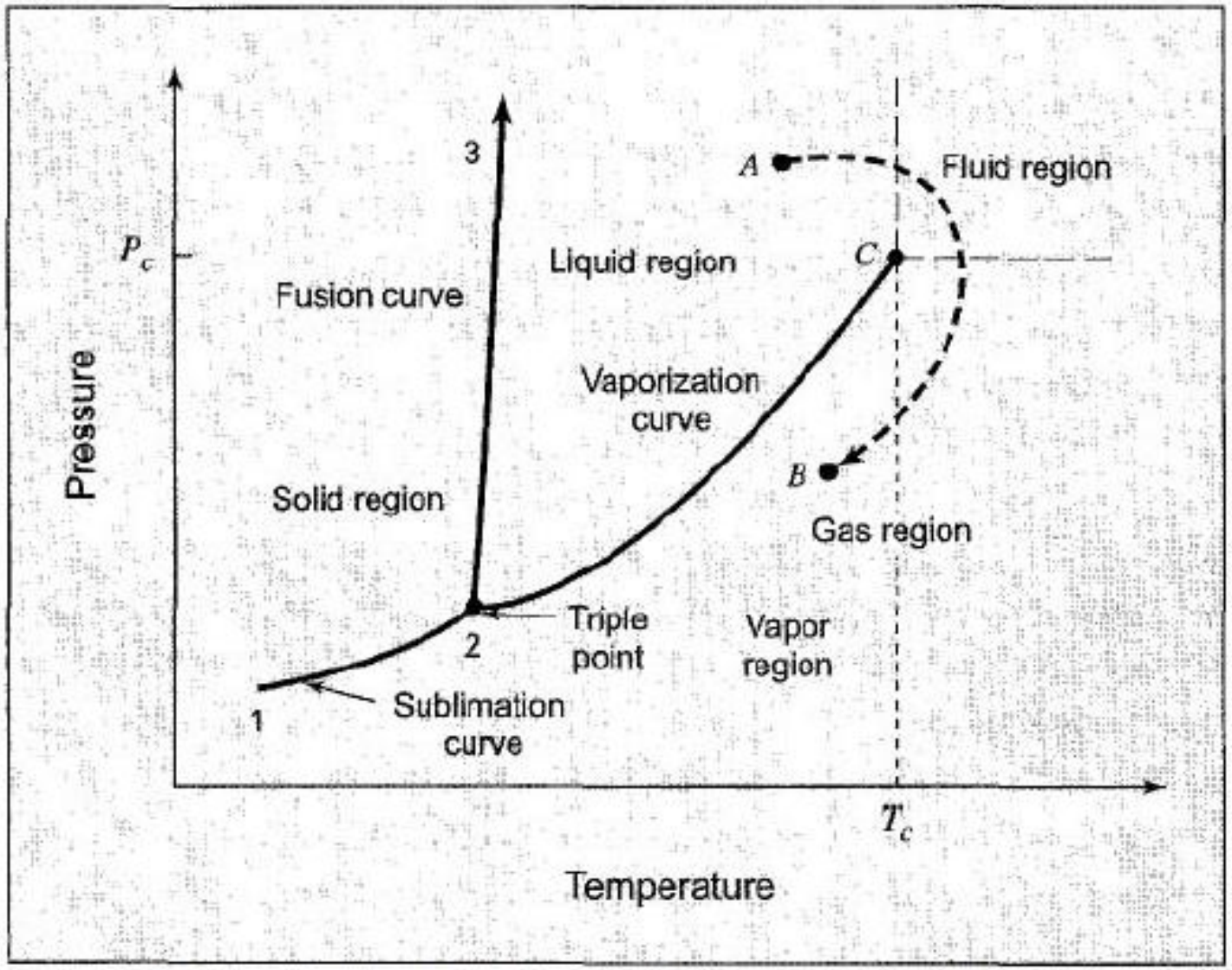


Figure 3.1 PT diagram for a pure substance

Kaidah Fase Gibbs

Pada Kesetimbangan Fase dan tidak ada reaksi kimia, berlaku :

$$F = 2 - \pi + N$$

Dengan :

F = derajat kebebasan

π = jumlah fase

N = jumlah komponen

$$F = 2 - \pi + N$$

Contoh:

Air (cair) setimbang dengan uapnya

$F = 2 - 2 + 1 = 1$

Pada kurva penguapan :

$\pi = 2$, $N = 1 \rightarrow$ $F = 2 - 2 + 1 = 1$

Jadi pd kurva penguapan kita bisa memilih T tertentu, atau P tertentu. Tapi ingat hanya 1 pilihan T saja atau P saja, yg lain mengikuti saja.

Derajat kebebasan pada titik tripel:

$\pi = 3,\ N = 1 \rightarrow \quad F = 2 - 3 + 1 = 0$

Jadi Titik Tripel sudah tertentu.

Persamaan Keadaan
(Equation of State = EOS)

adalah :

Persamaan yang menunjukkan hubungan antara P, V dan T suatu zat

atau

$$f(P, V, T) = 0$$

Khas untuk tiap zat

Gas Ideal

- ✓ EOS berbentuk : $PV = RT$

  atau $Z = PV/RT$ dan $Z = 1$

  dengan

  Z = faktor kompresibilitas

  P = pressure, Pa

  V = volume molar, m^3/mol

  T = temperature absolut, K

  R = konstanta gas umum,

  $= 8{,}314\ m^3\ Pa\ mol^{-1}\ K^{-1} = 8{,}314\ J\ mol^{-1}\ K^{-1}$

  Untuk satuan lain, lihat appendix A SVNA

- ✓ U = f(T) Energi dalam hanya fungsi suhu
- ✓ Berlaku pada tekanan sangat rendah

Contoh :

Hitung densitas gas Nitrogen (g/cm^3) pada tekanan 1 atm dan suhu 30^oC? (Asumsi: Gas ideal)

Jawab: ??

Hubungan antar Sifat pada Gas Ideal

$$Cv = \left(\frac{\partial U}{\partial T}\right)_V = \frac{dU}{dT} = Cv(T)$$

$$H = U + PV = U(T) + RT = H(T)$$

$$Cp = \left(\frac{\partial H}{\partial T}\right)_P = \frac{dH}{dT} = Cp(T)$$

$$Cp = \frac{dH}{dT} = \frac{dU}{dT} + \frac{d(RT)}{dT} = Cv + R \rightarrow \quad \boxed{Cp = Cv + R}$$

$$dU = Cv\, dT \qquad \Delta U = \int Cv\, dT$$

$$dH = Cp\, dT \qquad \Delta H = \int Cp\, dT$$

Perhitungan Proses dengan Persamaan Gas Ideal

PROSES ?

Proses-proses yg akan dibahas (untuk sistem tertutup), a.l. :

1. Proses Isotermal
2. Proses Isobarik
3. Proses Isokhorik
4. Proses Adiabatik

Proses Isotermal (ΔT = 0)

$\Delta U = Cv\ \Delta T = 0$ ingat $\left(\frac{\partial U}{\partial T}\right)_V = Cv$

$\Delta H = Cp\ \Delta T = 0$ ingat $\left(\frac{\partial H}{\partial T}\right)_P = Cp$

$$\Delta U = Q + W = 0$$

$$Q = -W = \int_{V_1}^{V_2} P\, dV \qquad \text{(ingat} \quad P = \frac{RT}{V})$$

$$Q = -W = \int_{V_1}^{V_2} \frac{RT}{V}\, dV = RT \ln \frac{V_2}{V_1}$$

$$\text{(ingat } \frac{P_1 V_1}{T_1} = \frac{P_2 V_2}{T_2} \text{ dan } T_2 = T_1 \text{ sehingga } \frac{V_2}{V_1} = \frac{P_1}{P_2})$$

$$Q = -W = RT \ln \frac{P_1}{P_2} = -RT \ln \frac{P_2}{P_1} = RT \ln \frac{V_2}{V_1} \qquad (\text{T tetap})$$

Proses Isobarik

$\Delta U = \int Cv\, dT$ dan $\Delta H = \int Cp\, dT$

$Q = \Delta H = \int Cp\, dT$ (P tetap)

$W = -\int P\, dV = -P\int dV = -\int R\, dT = -R\int dT$

$= -R\,(T_2 - T_1) = -P\,(V_2 - V_1)$

Cek dgn Hk I : $\Delta U = Q + W$

$Cv\, dT = Cp\, dT - R\, dT$

$Cv = Cp - R \rightarrow Cp = Cv + R$

Proses Isokhorik ($\Delta V=0$)

$\Delta U = \int Cv\, dT$ dan $\Delta H = \int Cp\, dT$

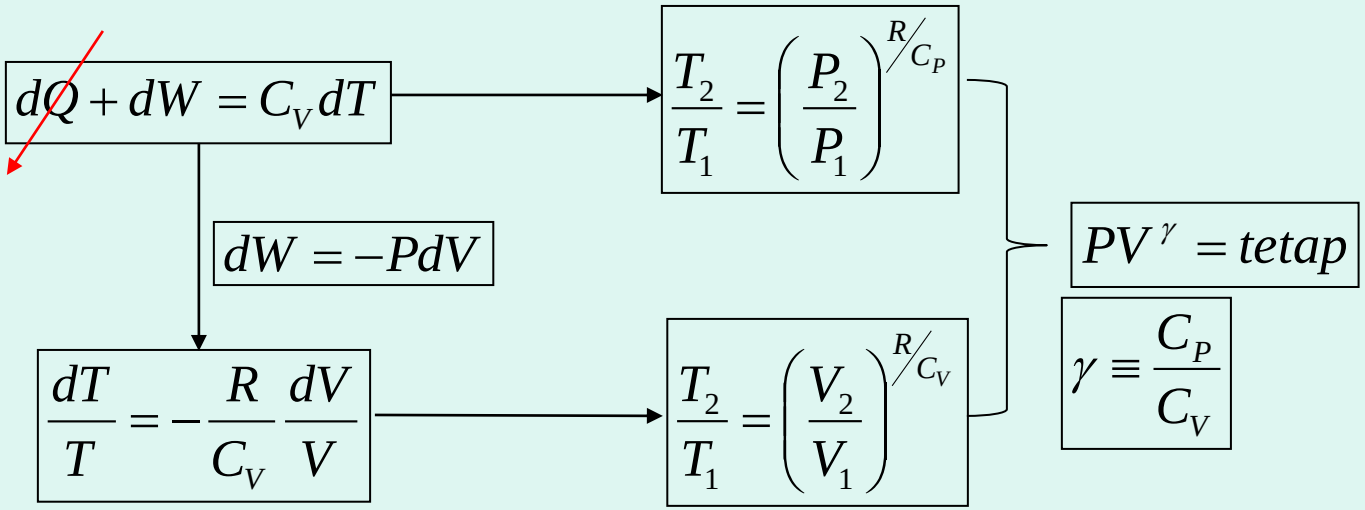
$Q = \Delta U = \int Cv\, dT$ (V tetap)

$=0$

$W = -\int P\, dV = 0 \rightarrow W = 0$

Cek dgn Hk I : $\Delta U = Q + W \rightarrow \Delta U = Q$

Proses Adiabatik (Q=0)



$$T\,V^{\gamma-1} = tetap \rightarrow T_1 V_1^{\gamma-1} = T_2 V_2^{\gamma-1}$$

$$T\;P^{(1-\gamma)/\gamma} = tetap$$

$$P\,V^{\gamma} = tetap$$

γ = 1,67 pada gas monoatomik (He, Ne, Ar dsb)

γ = 1,4 pada gas diatomik (H_2, N_2, O_2)

γ = 1,3 pada gas poli atomik sederhana (CO_2, SO_2, NH_3, CH_4)

Proses Adiabatik (lanjutan)

dari Hk I: $\Delta U = Q + W \rightarrow W = \Delta U$

$$W = \int Cv\, dT = Cv\, \Delta T \qquad (Cv = \frac{R}{\gamma - 1})$$

$$= \frac{R}{\gamma - 1}\Delta T = \frac{RT_2 - RT_1}{\gamma - 1} = \frac{P_2V_2 - P_1V_1}{\gamma - 1}$$

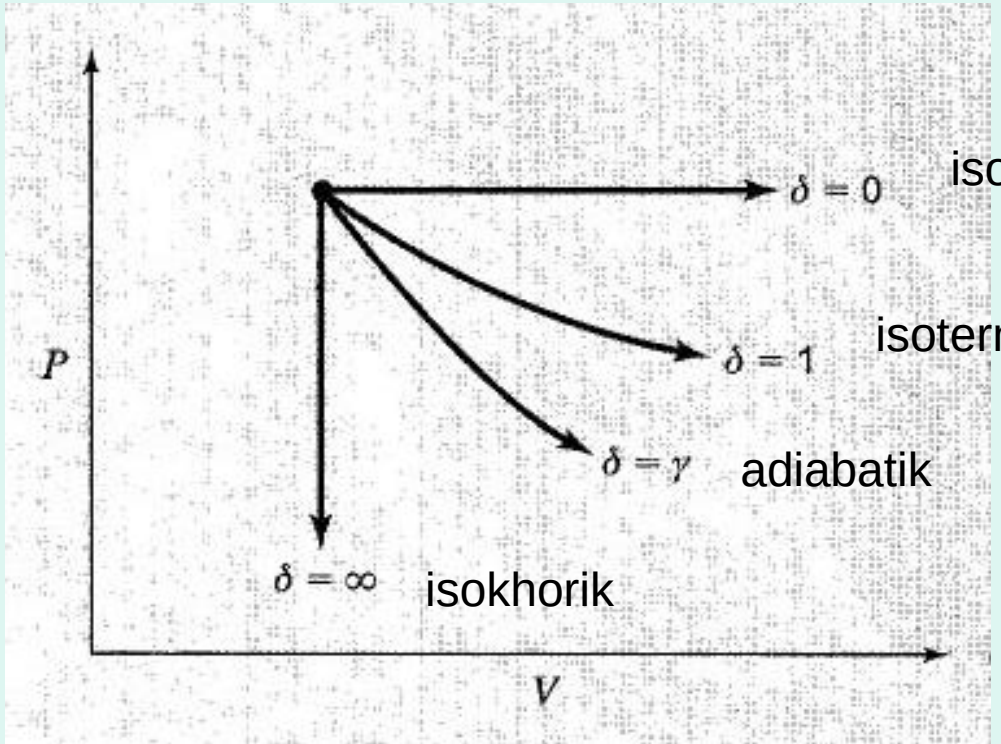
$$W = \frac{P_1V_1}{\gamma - 1}\left[\left(\frac{P_2}{P_1}\right)^{\frac{\gamma-1}{\gamma}} - 1\right] = \frac{RT_1}{\gamma - 1}\left[\left(\frac{P_2}{P_1}\right)^{\frac{\gamma-1}{\gamma}} - 1\right]$$

Proses Politropik

"turning many ways" → δ = konstan

$$T\,V^{\delta-1} = tetap \rightarrow T_1 V_1^{\delta-1} = T_2 V_2^{\delta-1}$$
$$T\;P^{(1-\delta)/\delta} = tetap$$
$$P\,V^{\delta} = tetap$$

$$W = \frac{P_1 V_1}{\delta - 1}\left[\left(\frac{P_2}{P_1}\right)^{\frac{\delta-1}{\delta}} - 1\right] = \frac{RT_1}{\delta - 1}\left[\left(\frac{P_2}{P_1}\right)^{\frac{\delta-1}{\delta}} - 1\right]$$



Gas Non Ideal

Z = PV/RT Z ≠ 1 (Z = faktor kompresibilitas)

EOS untuk gas non ideal:

1. Bentuk Virial

   a. $Z = 1 + B'P + C'P^2 + D'P^3 + \ldots$

   b. $Z = 1 + \frac{B}{V} + \frac{C}{V^2} + \frac{D}{V^3} + \ldots$

B', C', D' dst dan B, C, D dst : koefisien virial

2. Bentuk Kubik (Cubic EOS) (Pangkat Tiga)

a.l.

- Persamaan Van der Waals :

$$P = \frac{RT}{V-b} - \frac{a}{V^2}$$

Dalam bentuk polinomial :

$$V^3 - \left(b + \frac{RT}{P}\right)V^2 + \frac{a}{P}V - \frac{ab}{P} = 0$$

$$a = \frac{27}{64}\frac{R^2 T_c^{\,2}}{P_c} \qquad b = \frac{1}{8}\frac{RT_c}{P_c}$$

Tc = suhu kritik

Pc = tekanan kritik

- Persamaan Redlich-Kwong

$$P = \frac{RT}{V-b} - \frac{a}{V(V+b)\sqrt{T}}$$

$$a = 0{,}42748\,\frac{R^2 T_c^{2,5}}{P_c} \qquad b = 0{,}08664\,\frac{RT_c}{P_c}$$

(Pada buku SVNA mrpk parameter ψ dan Ω)

❑ Persamaan Soave-Redlich-Kwong

$$P = \frac{RT}{V-b} - \frac{a(T)}{V(V+b)}$$

$$a(T_c) = a_c = 0{,}42748\,\frac{(RT_c)^2}{P_c}$$

$$b = 0{,}08664\,\frac{RT}{P_c}$$

$$a(T) = a_c \cdot \alpha(T)$$

$$T_r = \frac{T}{T_c}$$

$$m = 0{,}480 + 1{,}574\omega - 0{,}176\omega^2$$

$$\alpha(T) = \left(1 + m\left(1 - \sqrt{T_r}\right)\right)^2$$

ω = acentric factor (lihat tabel) → faktor bentuk molekul
seberapa dekat dengan bentuk bola

- Pers. bentuk kubik lain masih banyak lagi, tetapi cukup 3 pers. di atas sebagai contoh.
- Persamaan bentuk kubik membentuk kurva sbb:

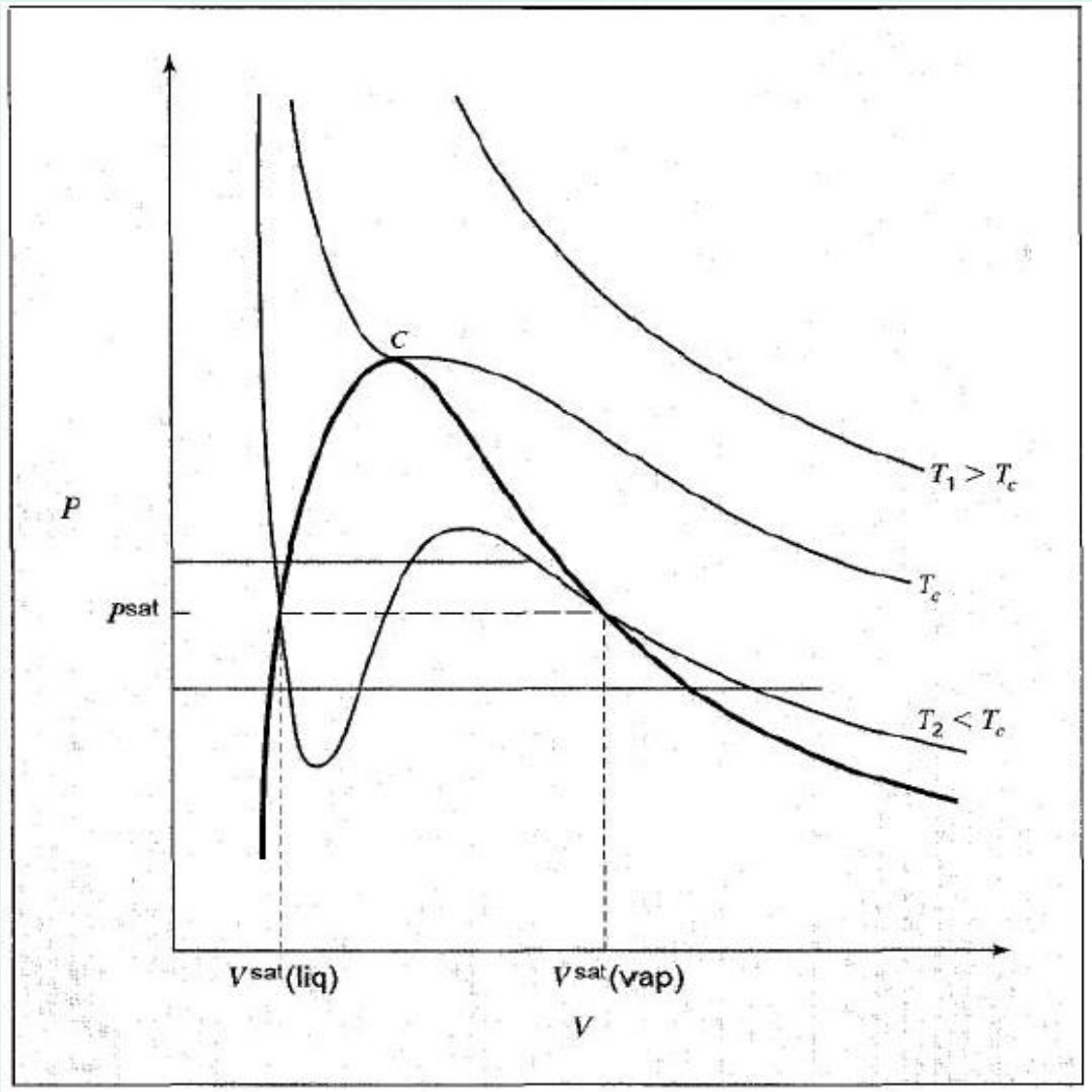


Figure 3.12 Isotherms as given by a cubic equation of state

Application of the virial equations

- Differentiation:

$$\left(\frac{\partial Z}{\partial P}\right)_T = B' + 2C'P + 3D'P^2 + \ldots \qquad \left(\frac{\partial Z}{\partial P}\right)_{T;P=0} = B'$$

- the virial equation truncated to two terms satisfactorily represent the PVT behavior up to about 5 bar

$$Z = 1 + B'P \qquad Z = 1 + \frac{BP}{RT} = 1 + \frac{B}{V} \longrightarrow$$

At low pressure

- the virial equation truncated to three terms provides good results for pressure range above 5 bar but below the critical pressure

$$Z = 1 + B'P + C'P^2 \qquad Z = 1 + \frac{B}{V} + \frac{C}{V^2}$$

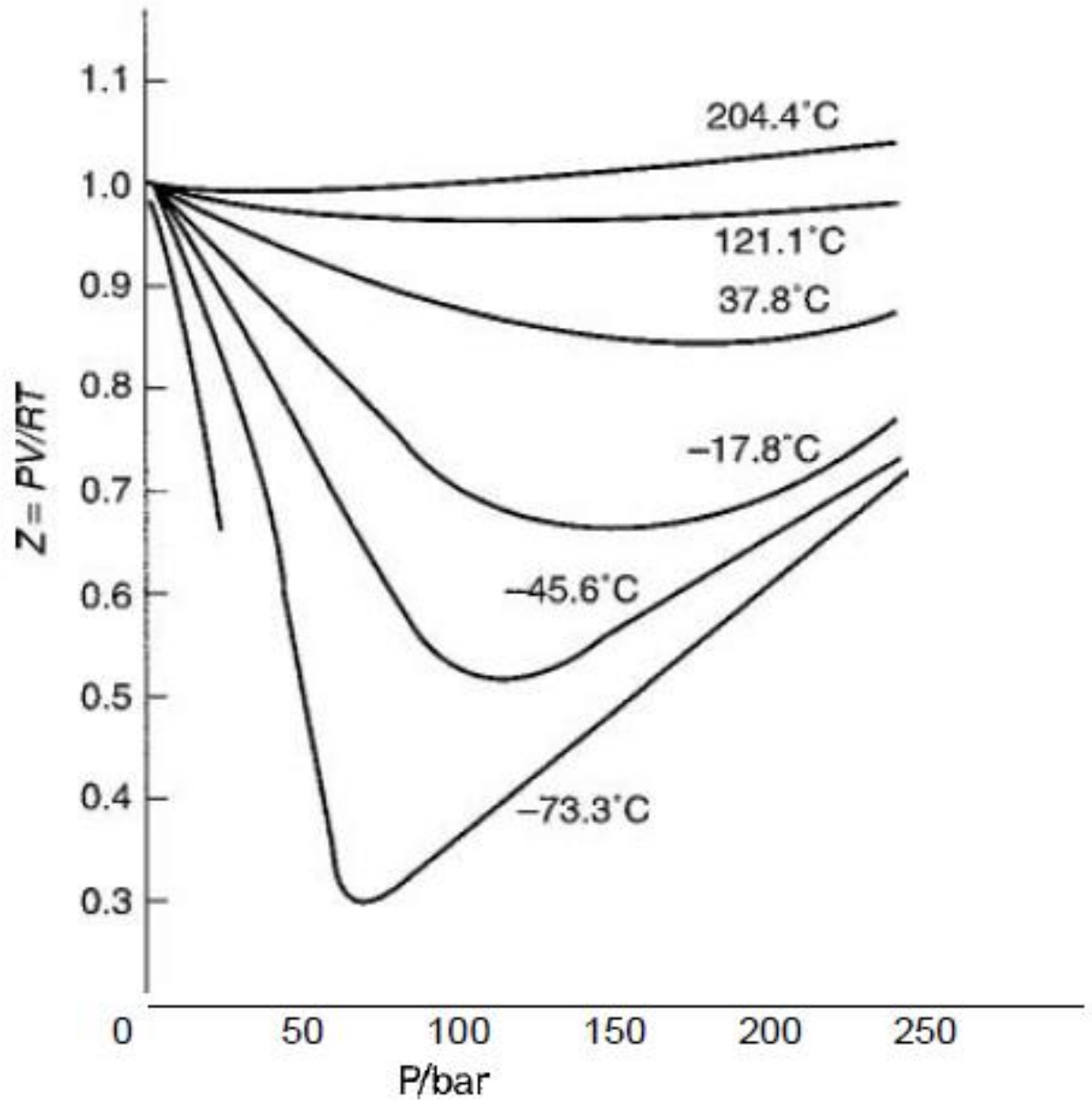


Figure 3.10 Compressibility-factor graph for methane

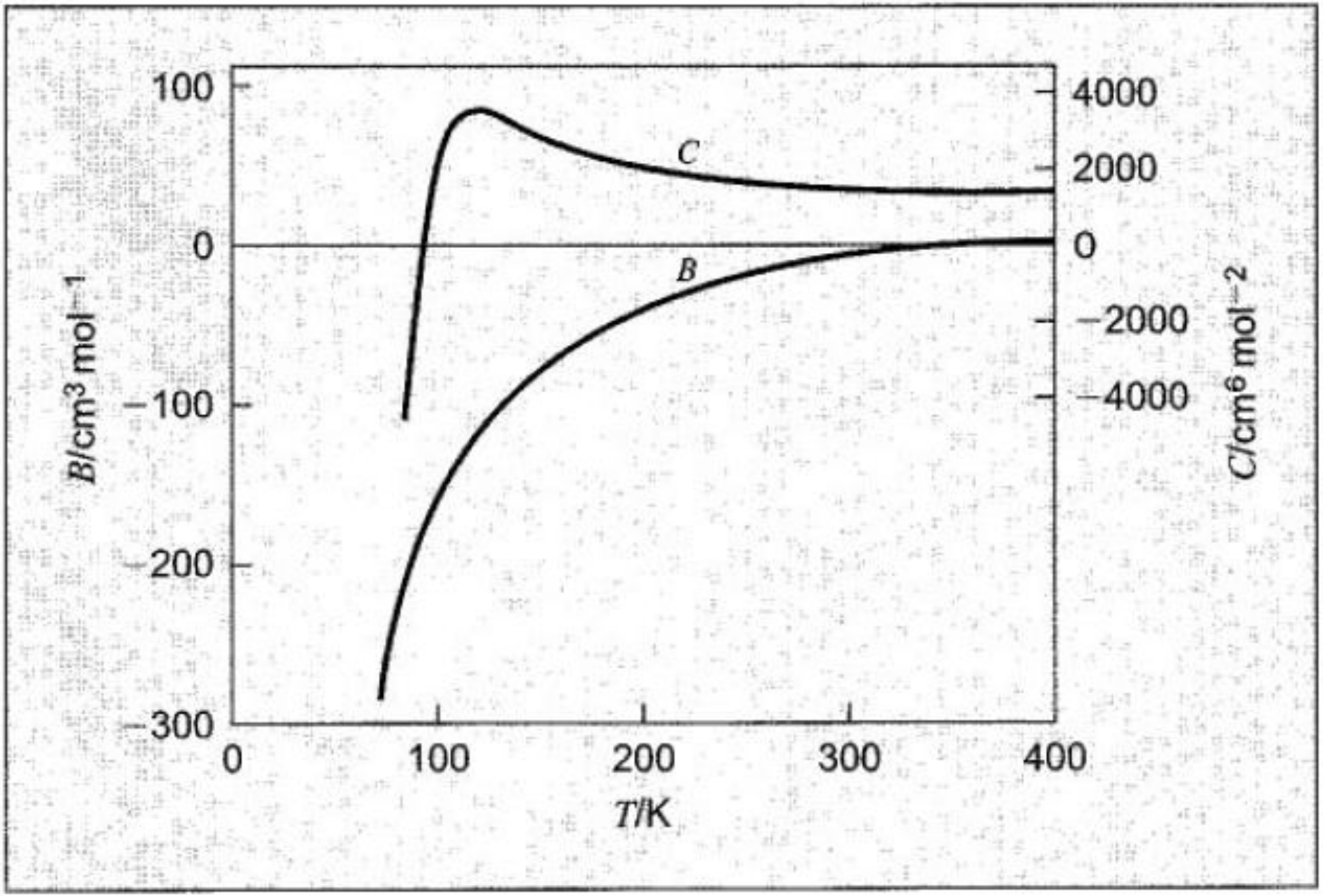


Figure 3.1 1 Density-series virial coefficients B and C for nitrogen

Reported values for the virial coefficients of isopropanol vapor at 200°C are:
B = -388 cm^3/mol and C = -26000 cm^6/mol^2. Calculate V and Z for isopropanol vapor at 200 °C and 10 bar by (1) the ideal gas equation; (2) two-term virial equation; (3) three-term virial equation.

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(1) For an ideal gas, Z = 1:

$$V = \frac{RT}{P} = \frac{83.14 \times 473.15}{10} = 3934\, cm^3/mol$$

(2) two-term virial equation:

$$V = \frac{RT}{P} + B = 3934 - 388 = 3546\, cm^3/mol \longrightarrow Z = \frac{PV}{RT} = 0.9014$$

(3) three-term virial equation:

$$V_{i+1} = \frac{RT}{P}\left(1 + \frac{B}{V_i} + \frac{C}{V_i^2}\right) = 3934\left(1 + \frac{-388}{3934} + \frac{-26000}{(3934)^2}\right) = 3539\, cm^3/mol$$

1st iteration

Ideal gas value

...

After 5 iterations $V_4 \sim V_5 = 3488\, cm^3/mol \longrightarrow Z = \frac{PV}{RT} = 0.8866$

Generalized correlations for gases (Korelasi Umum)

- Pitzer correlations for the compressibility factor:

  - $Z^0 = F^0 (T_r, P_r)$ $\quad\quad$ $Z = Z^0 + \omega Z^1$
  - Simple linear relation between Z and ω for given values of T_r and P_r.
  - Of the Pitzer-type correlations available, the Lee/Kesler correlation provides reliable results for gases which are nonpolar or only slightly polar (App. E, table E.1 – E.4)).
  - Only tabular nature (disadvantage)

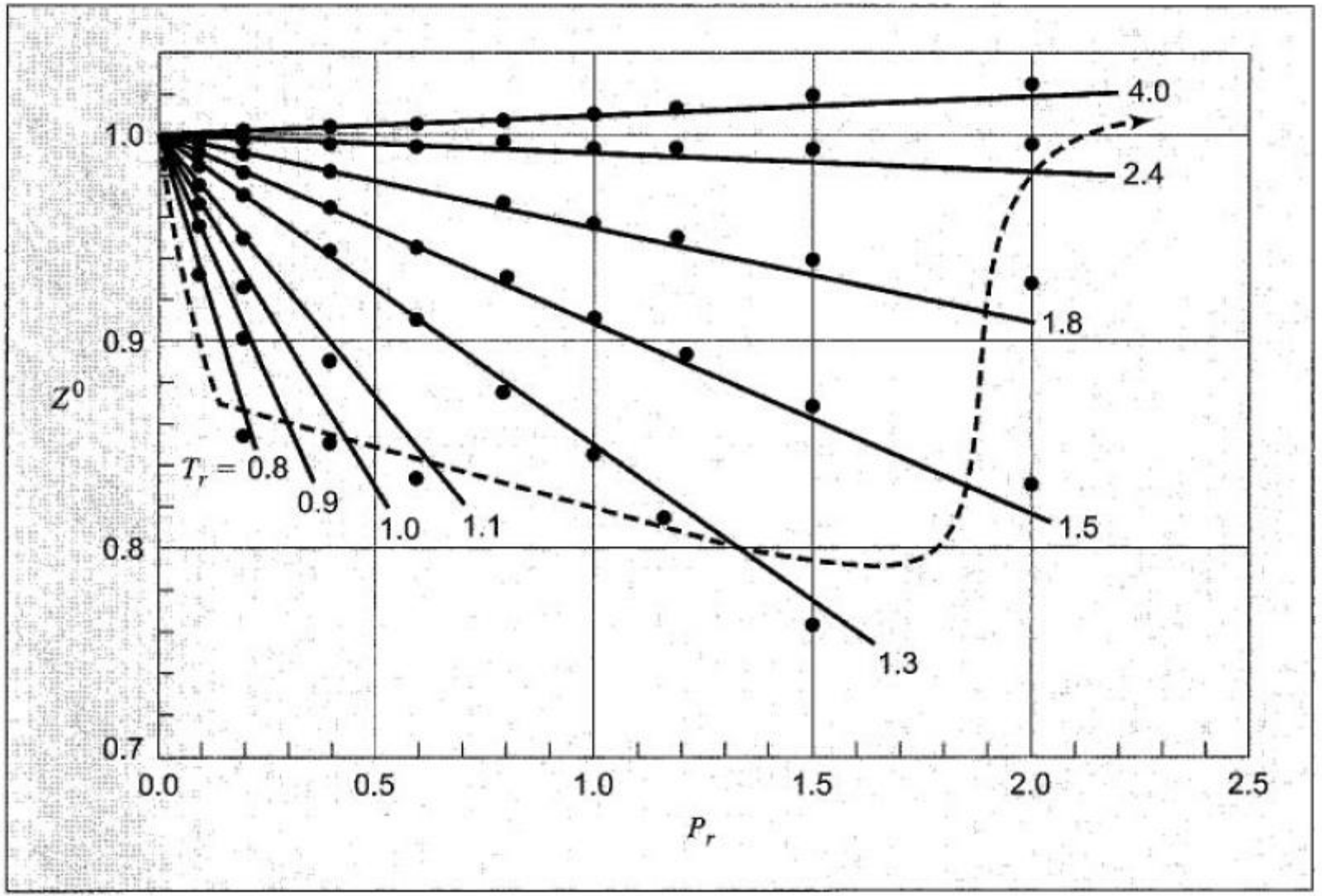

Figure 3.15 Comparison of correlations for Z^0. The virial-coefficient correlation is represented by the straight lines; the Lee/Kesler correlation, by the points. In the region above the dashed line the two correlations differ by less than 2%

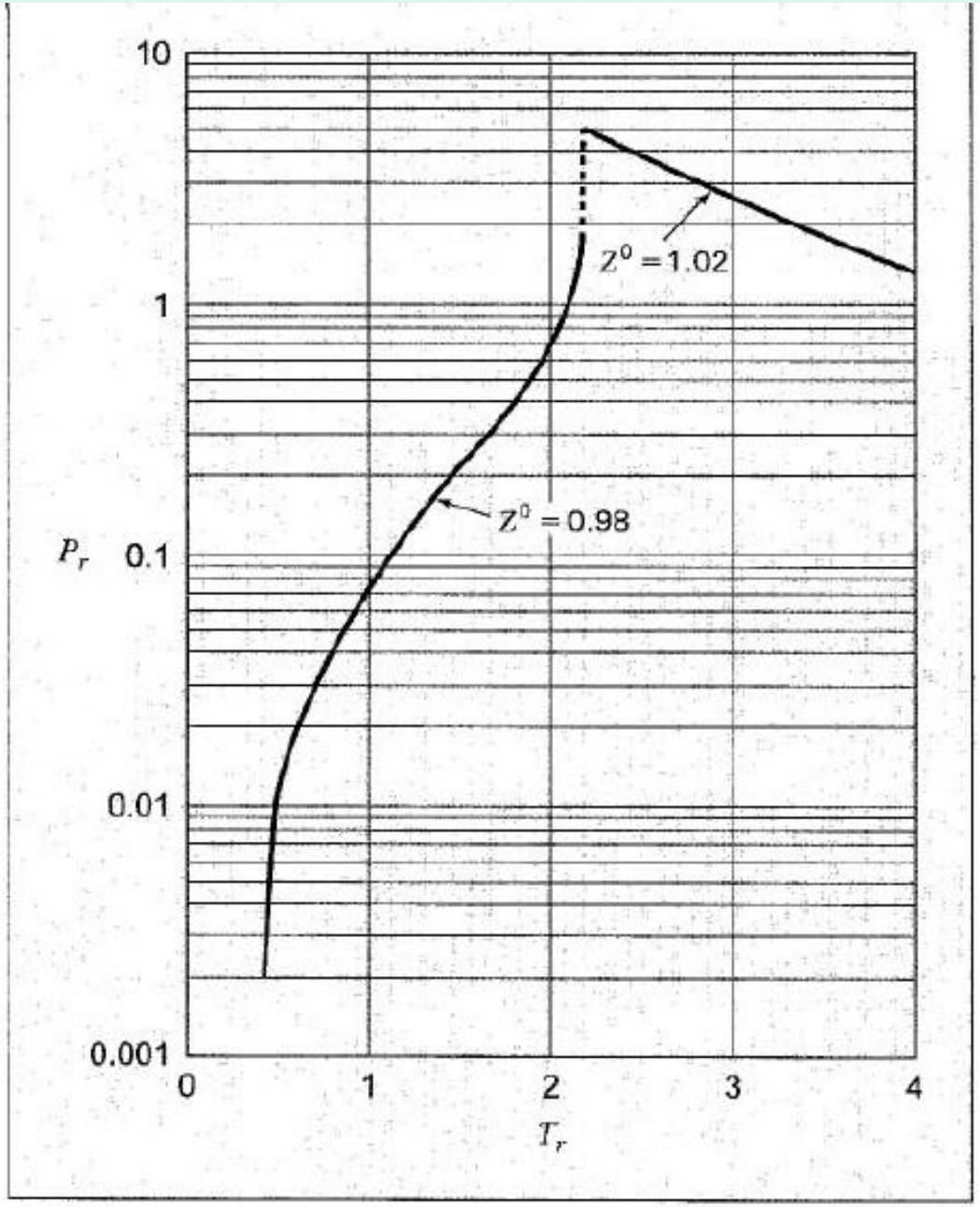

Figure 3.16 Region where Z^0 lies between 0.98 and 1.02, and the ideal-gas equation is a reasonable approximation

- Selain tersedia dalam bentuk tabel, metode Lee Kesler juga menyediakan bentuk grafik

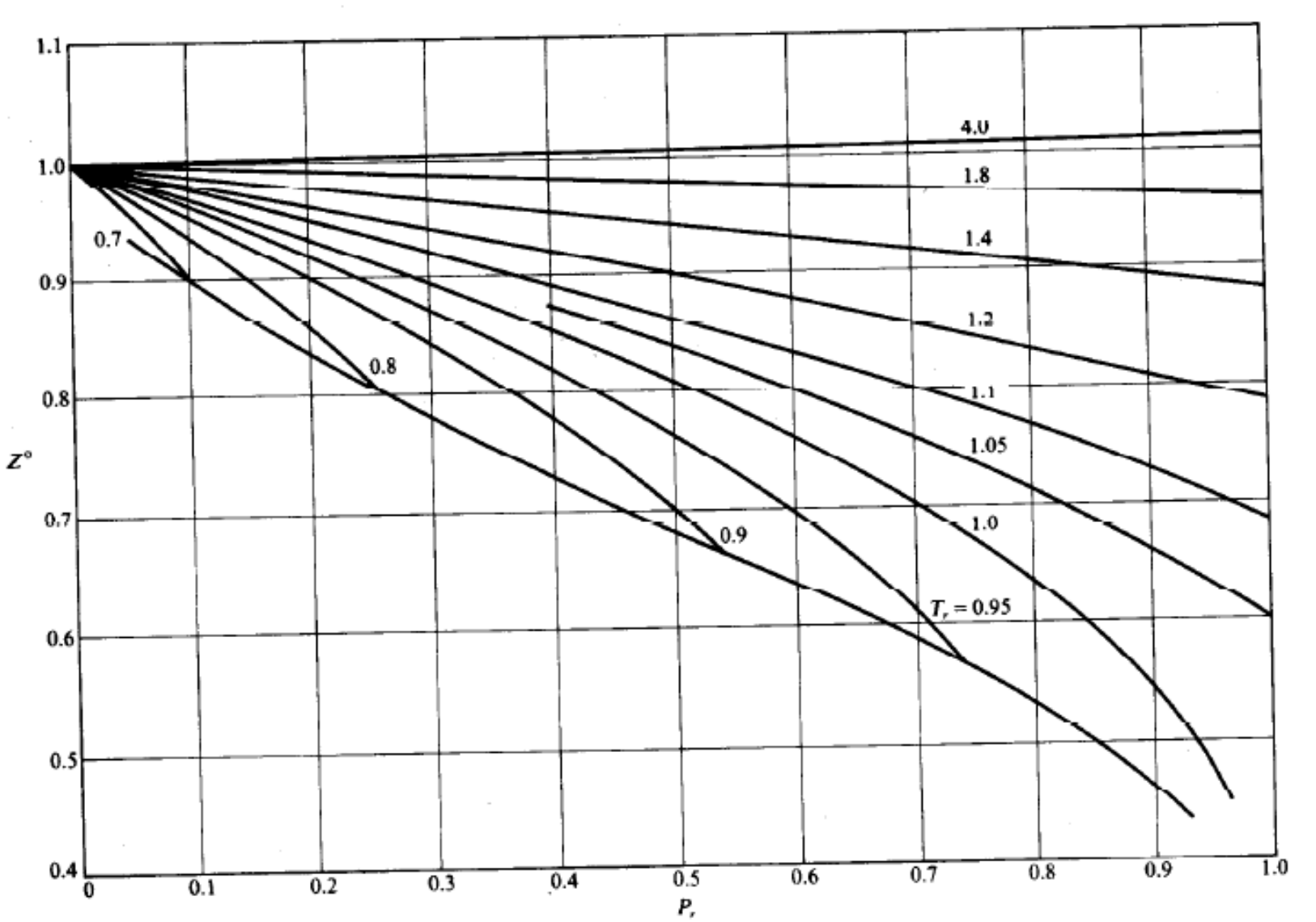

Figure 3.12 Generalized correlation for Z^0, $P_r < 1.0$. *(Based on data of B. I. Lee and M. G. Kesler, AIChE J.,* **21**: *510-527, 1975.)*

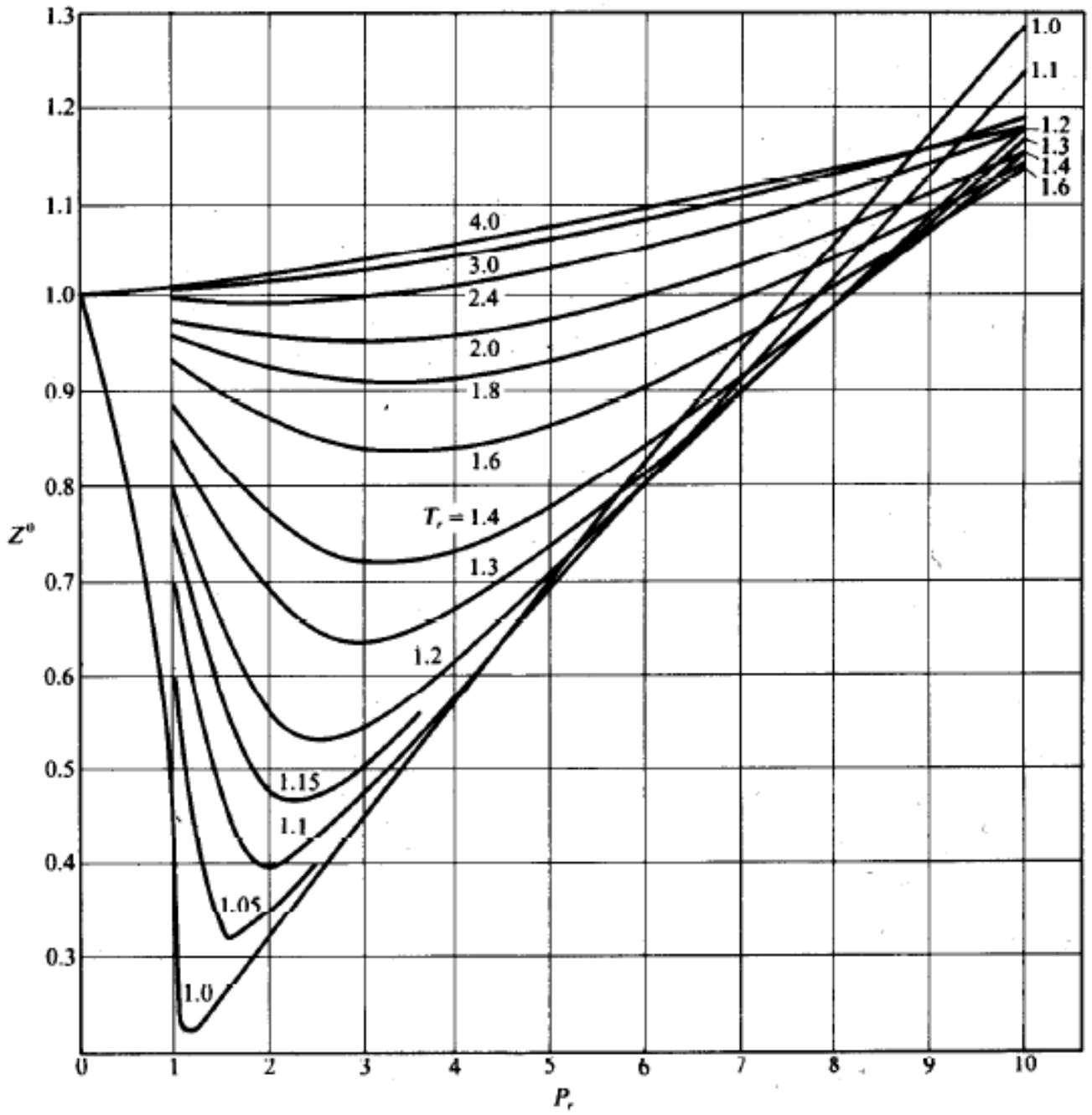

Figure 3.13 Generalized correlation for Z^0, $P_r > 1.0$. *(Based on data of B. I. Lee and M. G. Kesler,* ibid.*)*

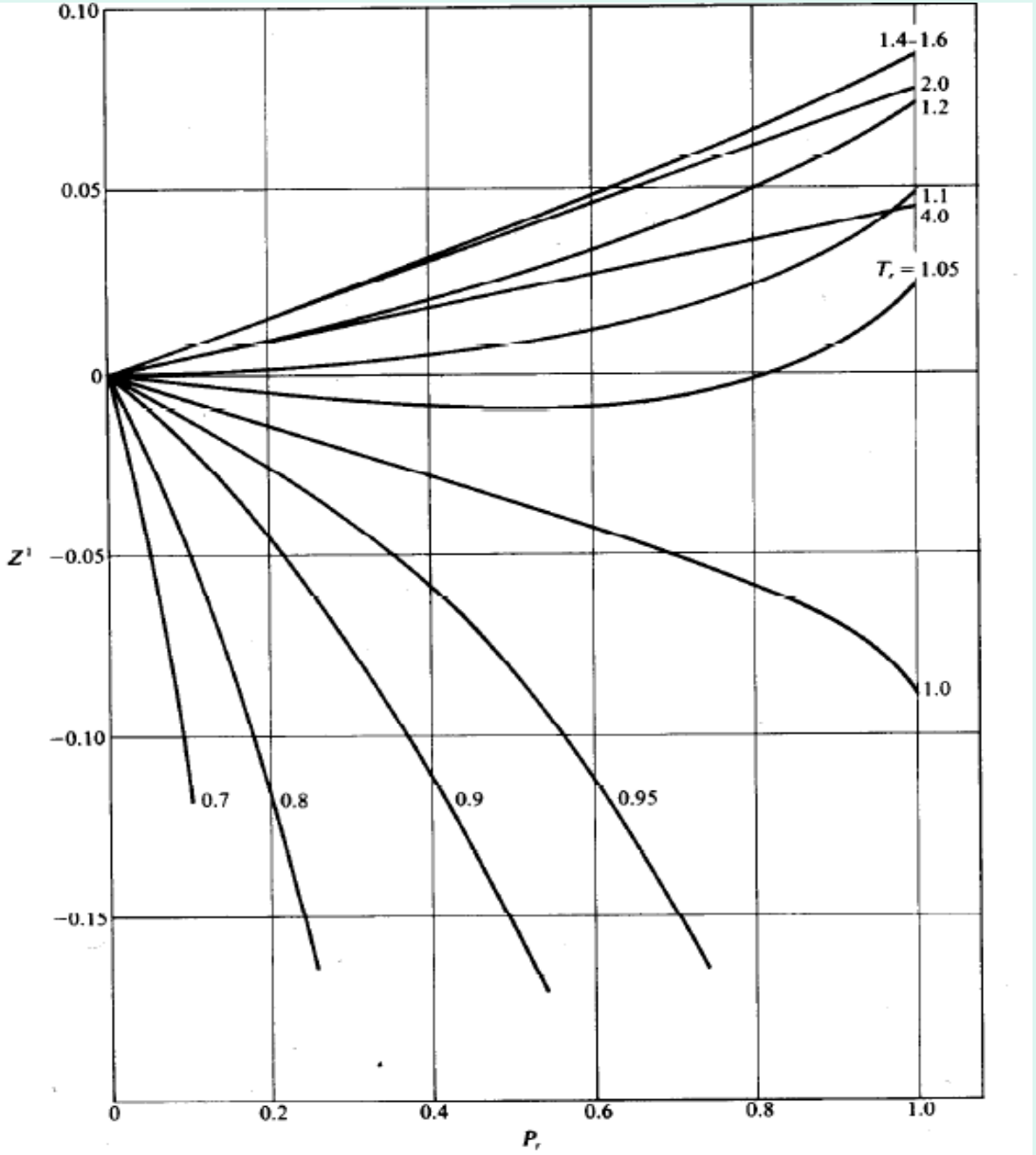

Figure 3.14 Generalized correlation for Z^1, $P_r < 1.0$. *(Based on data of B. I. Lee and M. G. Kesler,* ibid.*)*

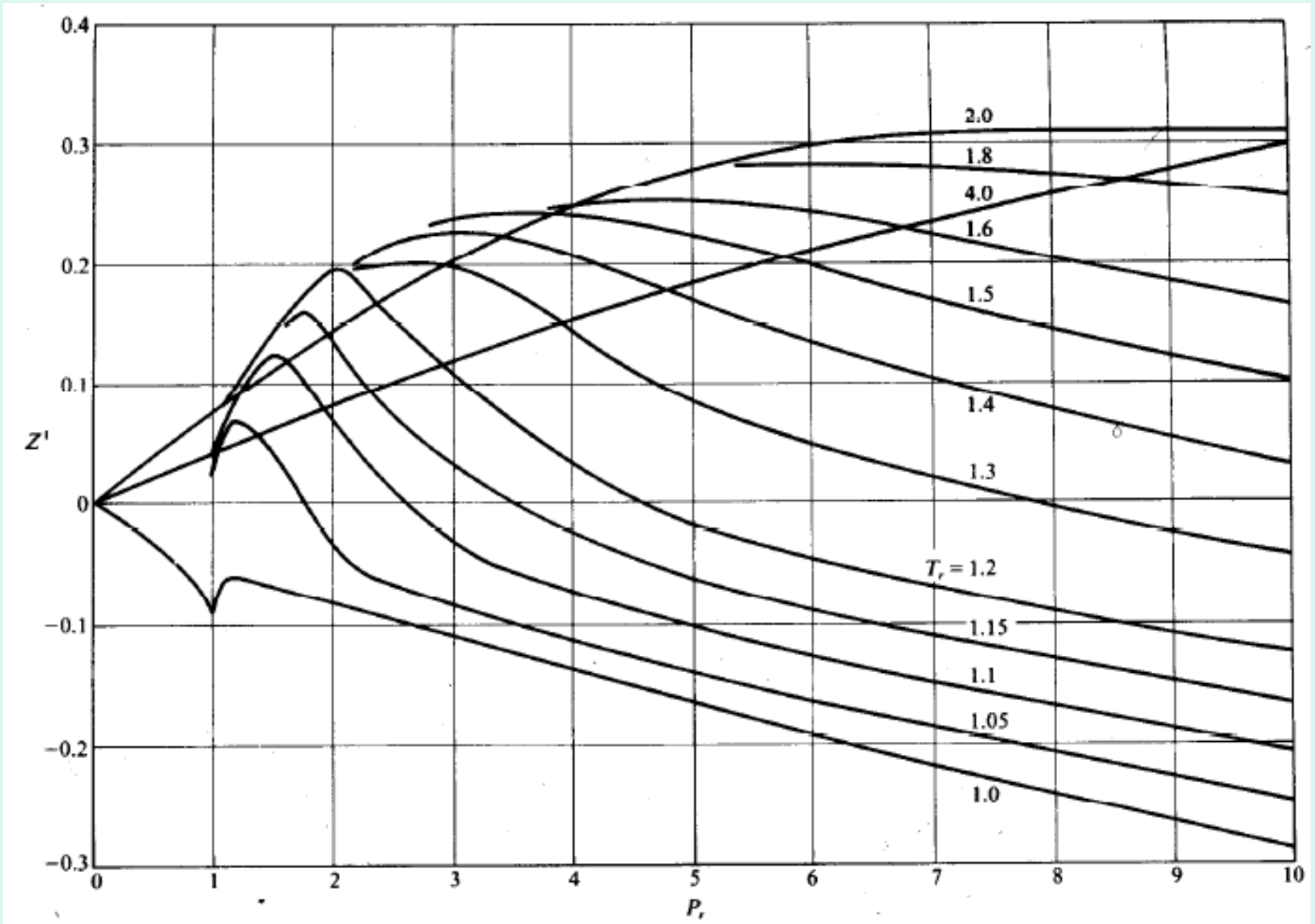

Figure 3.15 Generalized correlation for Z^1, $P_r > 1.0$. *(Based on data of B. I. Lee and M. G. Kesler,* ibid.*)*

Pitzer correlations for the 2nd virial coefficient

- Correlation:

$$Z = 1 + \frac{BP}{RT} = 1 + B^0 \frac{P_r}{T_r} + \omega B^1 \frac{P_r}{T_r} \quad \longleftrightarrow \quad Z = Z^0 + \omega Z^1$$

$$Z^0 = 1 + B^0 \frac{P_r}{T_r} \qquad Z^1 = B^1 \frac{P_r}{T_r}$$

$$B^0 = 0.083 - \frac{0.422}{T_r^{1.6}}$$

$$B^1 = 0.139 - \frac{0.172}{T_r^{4.2}}$$

  - Validity at low to moderate pressures
  - For reduced temperatures greater than $T_r \sim 3$, there appears to be no limitation on the pressure.
  - Simple and recommended.
  - Most accurate for nonpolar species.

Determine the molar volume of n-butane at 510K and 25 bar by, (1) the ideal-gas equation; (2) the generalized compressibility-factor correlation; (3) the generalized virial-coefficient correlation.

(1) The ideal-gas equation

$$V = \frac{RT}{P} = 1696.1 \frac{cm^3}{mol}$$

(2) The generalized compressibility-factor correlation

$$T_r = \frac{510}{425.1} = 1.200 \qquad P_r = \frac{25}{37.96} = 0.659$$

the acentric factor → $\omega = 0.200$

the Lee/Kesler correlation ↓

$$Z^0 = 0.865 \qquad Z^1 = 0.038 \rightarrow Z = Z^0 + \omega Z^1 = 0.873$$

$$V = \frac{ZRT}{P} = 1480.7 \frac{cm^3}{mol}$$

(3) The generalized virial-coefficient correlation

$$T_r = \frac{510}{425.1} = 1.200 \qquad B^0 = 0.083 - \frac{0.422}{T_r^{1.6}} \qquad B^1 = 0.139 - \frac{0.172}{T_r^{4.2}}$$

$$Z = 1 + B^0 \frac{P_r}{T_r} + \omega B^1 \frac{P_r}{T_r} = 0.879$$

$$V = \frac{ZRT}{P} = 1489.1 \frac{cm^3}{mol}$$

What pressure is generated when 1 (lb mol) of methane is stored in a volume of 2 $(ft)^3$ at 122°F using (1) the ideal-gas equation; (2) the Redlish/Kwong equation; (3) a generalized correlation .

(1) The ideal-gas equation

$$P = \frac{RT}{V} = \frac{0.7302(122+459.67)}{2} = 212.4\ atm$$

(2) The RK equation

$$T_r = \frac{581.67}{343.1} = 1.695$$

$$a(T) = \Psi \frac{\alpha(T_r) R^2 T_c^2}{P_c} = 453.94\ \frac{atm}{ft^6}$$

$$b = \Omega \frac{RT_c}{P_c} = 0.4781\ ft^3$$

$$P = \frac{RT}{V-b} - \frac{a(T)}{V(V+b)} = 187.49\ atm$$

(3) The generalized compressibility-factor correlation is chosen (high pressure)

$$P = \frac{ZRT}{V} = \frac{Z(0.7302)(122+459.67)}{2} = 212.4Z\ atm$$

$$P_r = \frac{P}{45.4} = \frac{Z}{0.2138}$$

$$T_r = \frac{581.67}{343.1} = 1.695$$

→ Z starts at Z = 1 and converges on Z = 0.890

$$P = 189.0\ atm$$

A mass of 500 g of gases ammonia is contained in a 30000 cm^3 vessel immersed in a constant-temperature bath at 65°C. Calculate the pressure of the gas by (1) the ideal-gas equation; (2) a generalized correlation .

$$V = \frac{V^t}{n} = 1021.2 \frac{cm^3}{mol}$$

(1) The ideal-gas equation

$$P = \frac{RT}{V} = 27.53\, bar$$

(2) The generalized virial-coefficient correlation is chosen (low pressure, $P_r \sim 3$)

$$T_r = \frac{338.15}{405.7} = 0.834 \longrightarrow B^0 = 0.083 - \frac{0.422}{T_r^{1.6}} \qquad B^1 = 0.139 - \frac{0.172}{T_r^{4.2}}$$

$$P_r \sim \frac{27.53}{112.8} = 0.244$$

the acentric factor $\longrightarrow \omega = 0.253$

$$Z = 1 + \left(B^0 + \omega B^1\right)\frac{P_r}{T_r} = 1 - 0.541\frac{P_r}{T_r}$$

$$P = \frac{ZRT}{V} = 23.76\, bar$$

Generalized correlations for liquids

- The generalized cubic equation of state (low accuracy)
- The Lee/Kesler correlation includes data for subcooled liquids
  - Suitable for nonpolar and slightly polar fluids
- Estimation of molar volumes of saturated liquids
  - Rackett, 1970:

$$V^{sat} = V_c Z_c^{(1-T_r)^{0.2857}}$$

- Generalized density correlation for liquid (Lydersen, Greenkorn, and Hougen, 1955):

$$\rho_r \equiv \frac{\rho}{\rho_c} = \frac{V_c}{V}$$

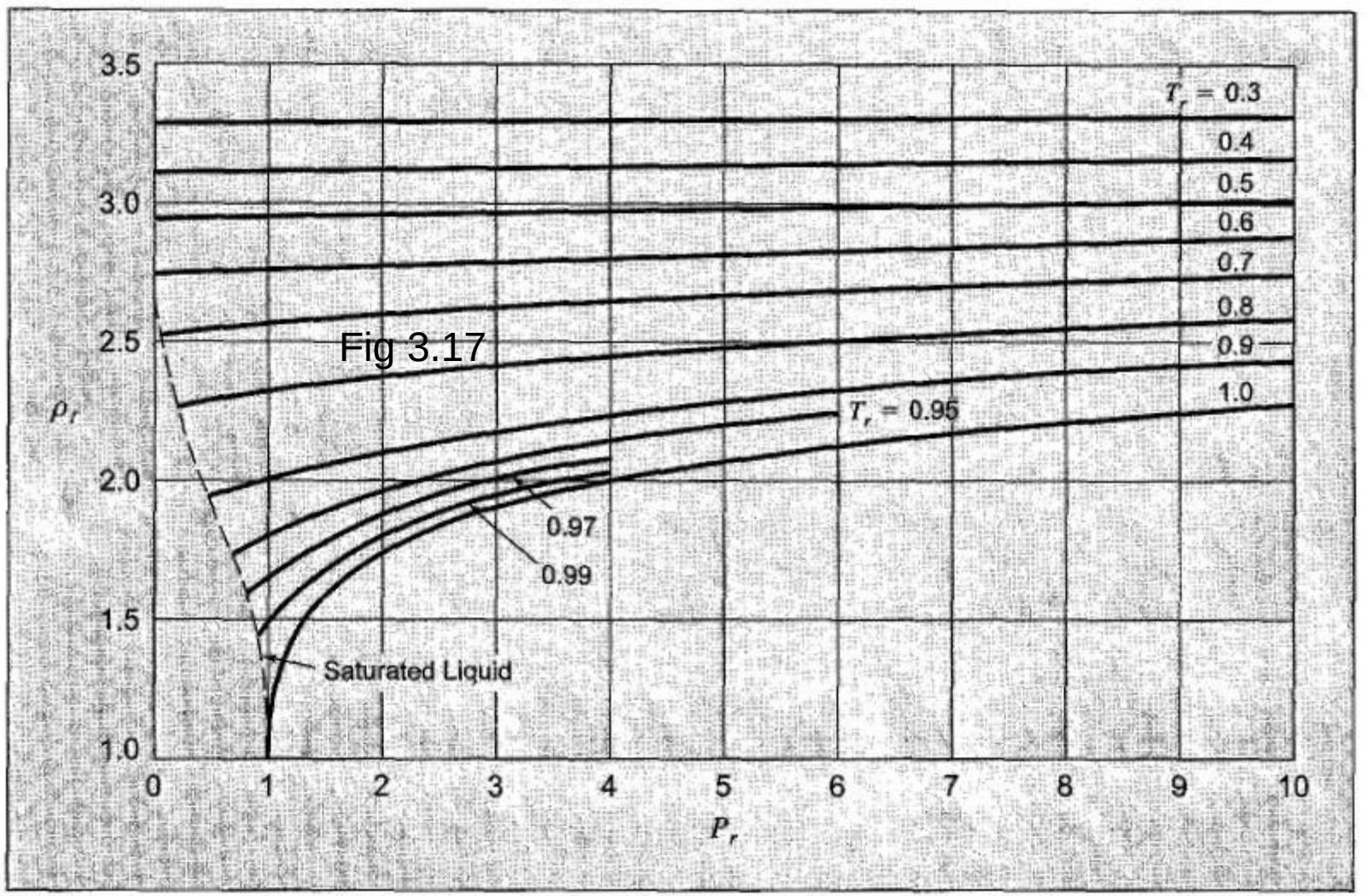

***Figure* 3.17** Generalized density correlation for liquids

$$V_2 = V_1 \frac{\rho_{r1}}{\rho_{r2}}$$

For ammonia at 310 K, estimate the density of (1) the saturated liquid; (2) the liquid at 100 bar

(1) Apply the Rackett equation at the reduced temperature

$$T_r = \frac{310}{405.7} = 0.7641 \qquad V_c = 72.47 \qquad Z_c = 0.242$$

$$V^{sat} = V_c Z_c^{(1-T_r)^{0.2857}} = 28.33 \frac{cm^3}{mol}$$

(2) At 100 bar

$$P_r = \frac{100}{112.8} = 0.887$$

$$T_r = \frac{310}{405.7} = 0.7641$$

Fig 3.17 →

$$\rho_r = 2.38$$

$$V = \frac{V_c}{\rho_r} = 30.45 \frac{cm^3}{mol}$$

$$V_2 = V_1 \frac{\rho_{r1}}{\rho_{r2}} = V^{310\,K} \frac{\rho_{r1,310\,K,saturated\ liquid}}{\rho_{r2,100\,bar}} = 29.14 \frac{2.34}{2.38} = 28.65 \frac{cm^3}{mol}$$

Terimakasih