

CVD TEST - Finish up

BLACK-OIL PVT (Ch. 7)

CVD TEST - What Lab Study "Reports"

$T_R = \text{constant}$

p	y_i ("δ")	$V_{ro}^{CVD} = \frac{V_o}{V_s}$	Z_g	$Z_g(p, y_i)$	$\left(\frac{\Delta n_g}{n_s}\right) (\%)$	$\{ \dots \}$	Z_2
$p_s = p_1$	$= Z_i \text{ Smol-}\%$	0 (PP)					
p_2	6.5	3	}	(Reservoir Gas)	Recovery from p_s	Not Surface Gas	Recovery
\vdots	5.2	12					
p_{N-1}	1.3	40					
p_N		22 19	}				

\uparrow (RF_δ)

Little Importance

Ch. 6

* $\{ x_{ONi}, (M_{n+}, \gamma_{n+})_{ON} \}$ maybe stage N-1

Bleeding process N-1 → N

Z_{wi} = producing wellstream composition of a gas condensate reservoir

Surface Condensate C_{5+}

$\approx y_i^{CVD} (P_R = P_{CVD})$

INDEPENDENT of Rock Relative Permeabilities

$$pV = nRTZ(p)$$

$\uparrow \quad \uparrow \quad \uparrow$
 same ✓ ✓
 80%

constant
A vs B

$$V_A > V_B$$

A: $HCPV_i = const$
 B: $HCPV < HCPV_i$

$$p_A < p_B$$

Darcy Law

$$q_g \propto (p_R - p_{wf})$$

$$q_{gA} < q_{gB}$$

More money/day

Control w/ Choke

(Whitson-Torp SPE paper)

BLACK-OIL PVT PROPERTIES (Ch. 7)

[[4 P-dependent tables]]

SURFACE DENSITIES
(Specific Gravities)
Assumed Constant

$\frac{S}{\rho}$	$\left\{ \begin{array}{l} \delta \\ \text{or} \\ \delta_g \end{array} \right\}$	COMPONENTS
$\frac{S_g}{\rho_g}$	δ_g	Surface Gas ✓
$\frac{S_o}{\rho_o}$	δ_o	Surface Oil ✓

OIL PHASE PVT @ (P, T)

GAS PHASE PVT @ (P, T)

P	R_s	B_o	M_o
S _o			✓

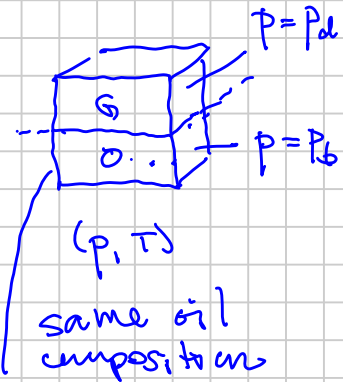
P	r_s	B_{gd}	M_g
S _g			✓

(1) SATURATED PROPERTIES

$p = p_b$ OIL PHASE

$p = p_d$ GAS PHASE

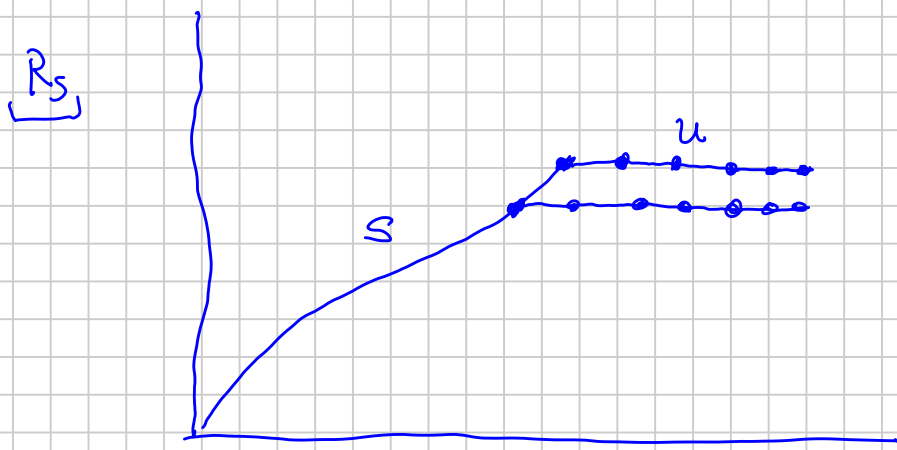
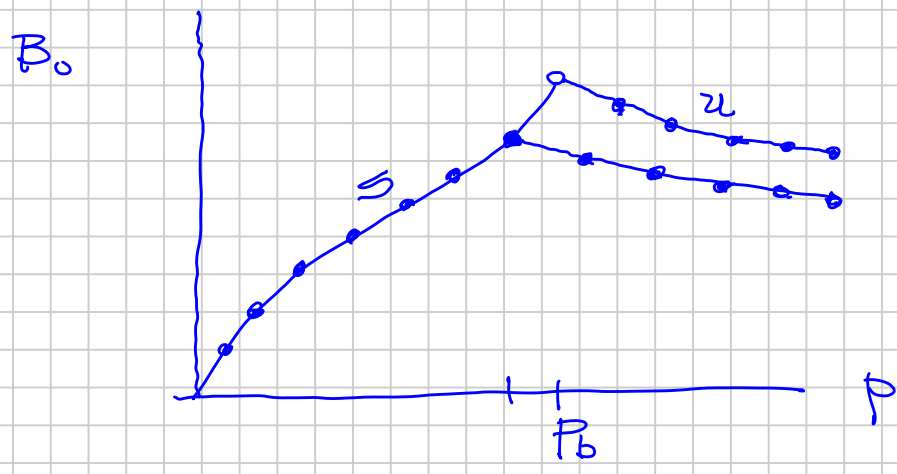
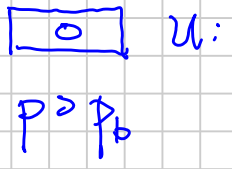
(2) UNDERSATURATED PROPERTIES



"R_s" same ~ "x_i"

p > p_b oil Phase with "R_s" (x_i)

p > p_d GAS Phase with "R_s" (y_i)



Bo PVT Model conserves surface gas component volume (V_g)
 surface oil component volume (V_o)

Ekofisk

Initially V_{gi} ≈ 6 · 10¹² scf

V_{oi} ≈ 6 · 10⁹ STB

all in solution initially
 all free oil

Troll

V_{gi} = 45 · 10¹² scf

V_{oi} = 6 · 10⁹ STB

42 · 10¹² free gas cap
 3 · 10¹² in solution
 0.4 · 10⁹ STB in solution
 0.4 · 10⁹ STB in gas cap
 5.9 · 10⁹ STB free

Surface gas (\bar{g}) can be found in both
reservoir gas and reservoir oil phases

Surface oil (\bar{o}) can be found in both
reservoir gas and reservoir oil phases

At any time (Surface Component Volumetric Material Balance)

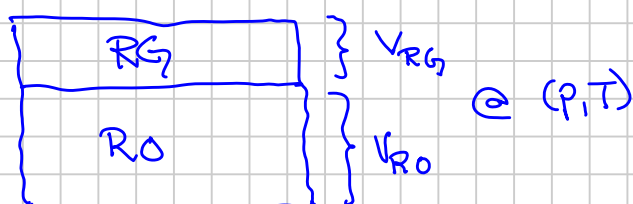
$$\underline{V_{\bar{g}i}} = \underline{V_{\bar{g}gR}} + \underline{V_{\bar{g}oR}} + \underline{V_{\bar{g}p}} - \underline{V_{\bar{g}inj}}$$

$V_{\bar{g}gR}$ = surface gas component in current reservoir
gas phase

$V_{\bar{g}oR}$ = surface gas component in (solution in)
current reservoir oil phase

$V_{\bar{g}p}$ = cumulative produced surface gas component
volume

$V_{\bar{g}inj}$ = cumulative injected surface gas
component volume



$$\underbrace{V_{\bar{a}i}} = \underbrace{V_{\bar{a}gR}} + \underbrace{V_{\bar{a}oR}} + \underbrace{V_{\bar{a}p}} - \underbrace{V_{\bar{a}inj}}_{(=0)}$$

If the surface components truly had invariant mass densities ($\rho_g \neq \rho_o$) then the "Volumetric Material Balance" would also guarantee a "Mass Material Balance".

$\gamma_o \equiv \frac{\rho_o}{\rho_w}$
 $\gamma_g \equiv \frac{\rho_g}{\rho_{air,sc}}$

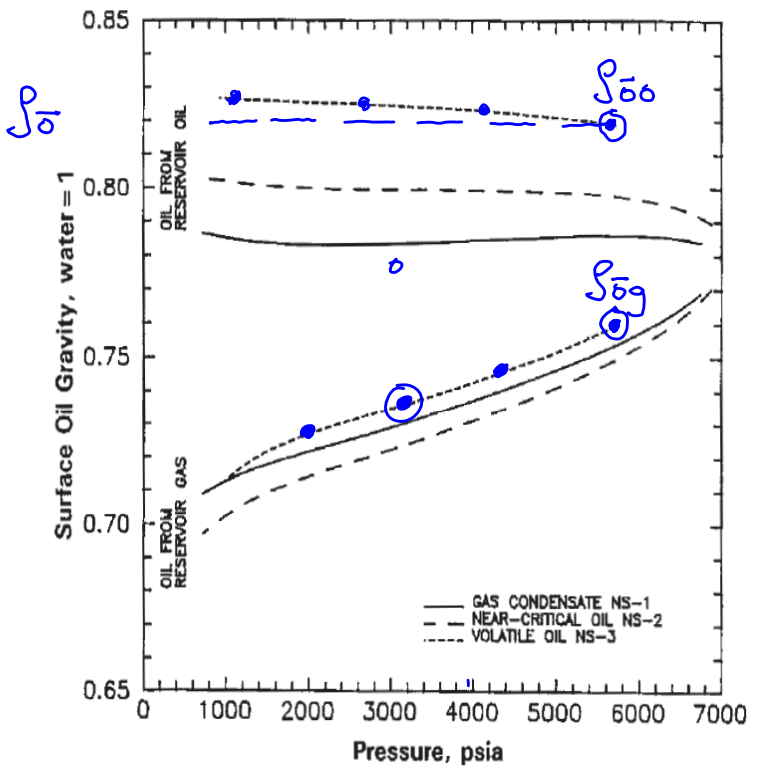
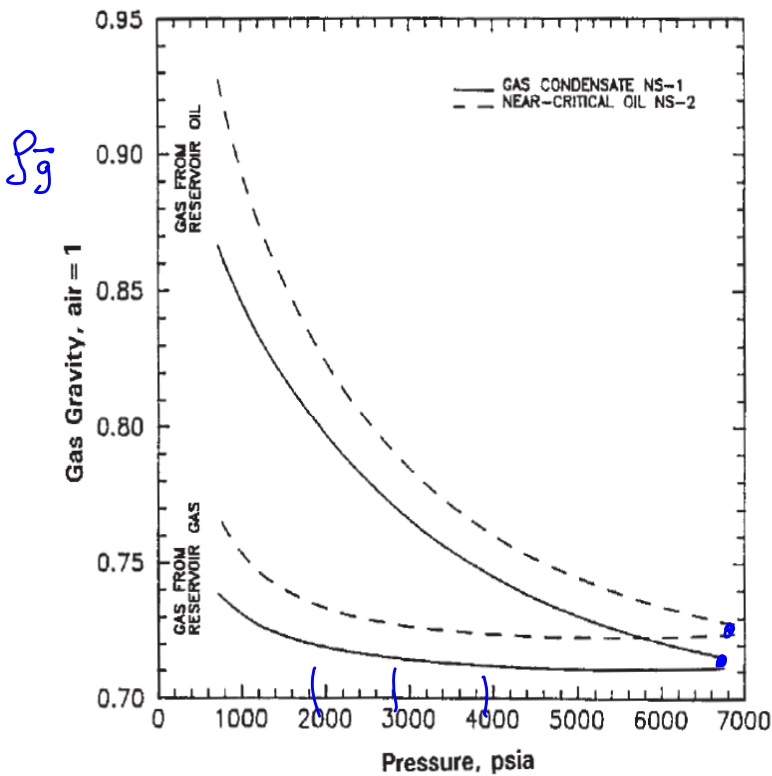


Fig. 7.12—Surface-gas gravities vs. pressure during depletion.

Fig. 7.13—Surface-oil gravities vs. pressure during depletion.

Why do we need, and do we use S_g S_o

Any engineering application (R, T, P) needs 3 PVT quantities (ρ, T) for each phase

$$\textcircled{1} V_{ro} \begin{cases} = 1 & \text{oil} \\ & \text{GAS+OIL} \\ = 0 & \text{GAS} \end{cases} \quad \textcircled{2} \mu \quad \textcircled{3} \rho$$

$$* \rho_o(p) = \frac{\rho_o + \rho_g R_s(p)}{B_o(p)} = \frac{m_o}{V_o} = \frac{m_{\bar{o}} + m_{\bar{g}o}}{V_o}$$

$$* \rho_g(p) = \frac{\rho_g + \rho_o R_s(p)}{B_{gd}(p)} = \frac{m_g}{V_g} = \frac{m_{\bar{g}g} + m_{\bar{o}g}}{V_g}$$

Use $\rho_g \approx \rho_o$ gets the best $\rho_g(p) \approx \rho_o(p)$
 densities

How result