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Executive Summary

This report details the data and techniques used to generate representative, usable models of the subsurface fluids involved in the process of injecting and storing CO_2 in the depleted gas condensate field Goldeneye. An accurate representation of the fluids is essential to understanding the full chain of events, from the arrival at the offshore platform of the CO_2 from the capture plant to the ultimate fate of the stored CO_2.

The fluids involved are the original hydrocarbon gas and the injected CO_2. The behaviours of both of these must be properly modelled under the different conditions of temperature and pressure which occur while the original hydrocarbon is being produced to surface and while the CO_2 is being injected. A valid model of the produced Goldeneye gas is necessary in the first instance to enable the simulation model to be checked against actual production history. If the model's forecasts **cannot** be "matched" to the real historical production, it is an indication that some aspect of the model is incorrect. Only correctly history matched models can be taken forward to simulate the CO_2 injection.

Samples of the original Goldeneye hydrocarbon gas were taken when the first discovery or appraisal wells were drilled between the years 1996 and 2000. These were supplied to specialist consultancies, for standard oil and gas industry experimental tests. The results from these have been used to construct models of the fluids, which in turn are input into the simulation models of the Goldeneye field. This process can be relatively straightforward for a fluid such as a "normal" black crude oil such as that found in Shell's Brent field, but it is significantly more difficult to find a way of adequately representing Goldeneye fluid (which exists near the boundary between gas and condensate). To do this requires a detailed model which combines, fully or almost fully, the properties of individual components (each component being made of a different hydrocarbon molecule), and which takes each component back to a fundamental level. Such a model is known as a compositional equation of state (EOS) model, and it is inevitably demanding of computer power. To simplify the situation a little, some individual components can be bundled (or "lumped") together and the bundle treated as an individual component. This is the approach used here. The resulting EOS model of the original Goldeneye gas condensate was used in the final dynamic models of the field.

Another EOS model was built to represent the properties of the injected CO_2 under the conditions of temperature and pressure found at the input to the injection wells, inside the wellbore and in the store.

The properties of Goldeneye gas were first studied before it was brought on stream as a producing field (2004). The most recent study was done during the FEED phase of the Longannet CCS project, when the quality of the pressure, volume and temperature (PVT) data were rechecked and the current EOS models were developed and approved. There have been no further updates to the PVT models since then (March 2010).

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1. Introduction

The Goldeneye hydrocarbon reservoir fluids have been extensively characterised during the hydrocarbon production phase of the field. This PVT characterisation has been updated and extended to facilitate (i) an equation of state description in the reservoir simulation, (ii) a good representation of the properties of CO_2 at storage conditions.

This report details the pressure, volume and temperature (PVT) characterisation that will be used to model the phase behaviour of CO_2 injection into the depleted Goldeneye gas condensate field.

The report includes the following:

- Preparation of a consistent equation of state (EOS) compositional model for Goldeneye gas condensate PVT. This involves the rationalisation of hydrocarbon single components (lumping) to achieve a representative but at the same time manageable fluid characterisation.

- Representation of CO_2 properties through an EOS

- Summary of PVT samples available from Goldeneye

- Discussion of lumping, validation plots (full PVT) and six components (lumped)

- CO_2 phase behaviour representation from Peng-Robinson EOS

Reference to third party software in this document is solely for information purposes to assist in understanding how the work was completed, does not amount to an endorsement of that software nor is any warranty as to its suitability given or implied.

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2. PVT data available

In 1996 Shell discovered the Goldeneye field by drilling Well 14/29a-3 and finding a gas column of 303 ft [92.35 m]. In the following years three appraisal wells were drilled: 1998 Amerada 20/4b-6 (south), 1999 Shell 14/29a-5 (south-east) and 2000 Amerada 20/4b-7 (south-west).
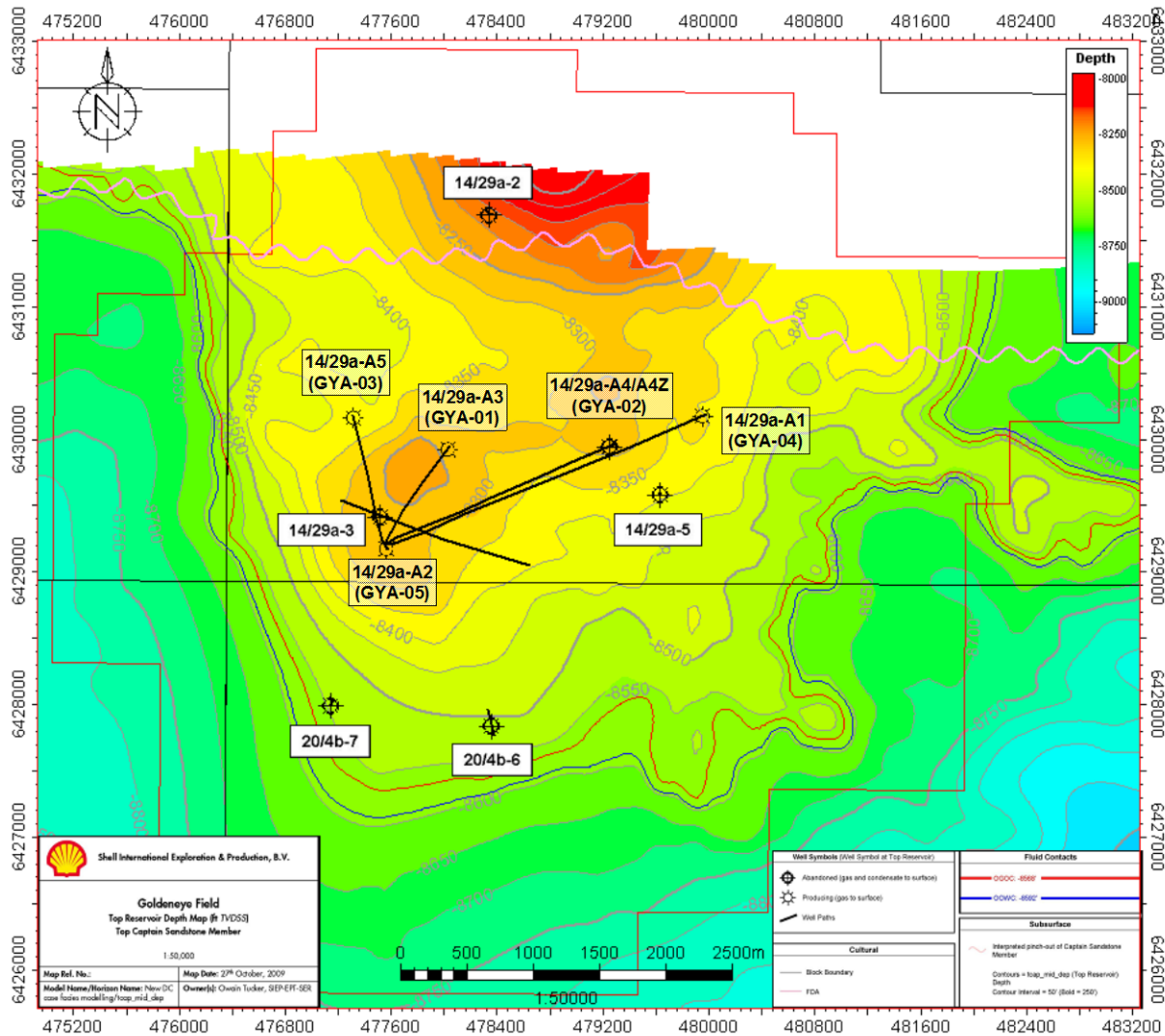
Fig. 2-1 –Goldeneye field top structure map showing well locations

In 2004 five development wells were drilled. The locations of the exploration and development wells are shown in Fig. 2-1.

Fluid samples were taken from all four discovery/appraisal wells during repeat formation tester/modular dynamic tester (RFT/MDT) tests and surface samples from drill stem tests (DST). PVT samples were taken from the test separator during clean up of the five Goldeneye development wells (GYA01 to GYA05). Detailed analysis from two of the wells was carried out, GYA04 and GYA03, including geochemical analysis. Gas and liquid compositions were determined for the other wells. The complete list of PVT samples available from Goldeneye is shown in Table 2-1.

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Table 2-1 History of PVT Characterisations

History of PVT Characterizations

DST

Well	sample	Contents	Characterization
14/29a-3	2336-C1-F	gas	Gas & Liq comp,
	2781-M1-F	liq	RC, CCE, CVD
20/4b-6	1744-M1-F	gas	Comp, FLS, RC
	4309-C1-F	gas	Comp, FLS, RC
	3458-M1-F	liq	Comp, FLS, RC
	1422-M1-F	gas	Comp, FLS, RC
	4380-M1-F	liq	Comp, FLS, RC
(clean up separator samples)			
GYA04	1896-M1-F	gas	Gas & Liq Comp,
	5064-M1-F	liquid	RC, CCE, CVD
			MSST
(clean up separator samples)			
GYA03	3263-M1-F		
	2193-C1-F		
	2188-M1-F		
GYA01	2766-M1-F		Gas & Liq comp
	2191-C1-F		
	0917-M1-F		
GYA02	3329-C1-F		Gas & Liq comp
	0760-M1-F		
GYA05	0966-M1-F		Gas & Liq comp
	2726-C1-F		
	2389-M1-F		

MDT

Well	sample	Contents	Characterization
14/29a-3	313415 (*)	gas & Liquid	FP, FLS, Gas Comp
	120085(*)	oil	FP, CCE, FLS, RC
	120083		drilling fluid & water FP, water analysis
	312104		drilling fluid & water FP, water analysis
	Atmospheric		FP
1429a-3	3452-M1-F(**)	oil	RC, CCE, DV, V
204b-6	0301-G1-F	oil	FLS, Liq comp, fluid prop, CCE, DV, V
	9882-B1-F	water	analysis
	3078-G1-F	water	analysis
14/29a-5	37(*)	gas	FP, Gas Comp
	56(***)	gas	Gas Comp
	554(*)	oil	FP, Comp, FLS, prop, CCE, DV
	550 & 535	water	analysis

FP	Fingerprint analysis
FLS	Flash to standard conditions
CCE	Constant Compostion Exparsion
DV	Differential Vapourisation
CVD	Constant Volume Depletion
V	Viscosity
SF	Separator flash
RC	Recombination
MSST	Multi stage separator test

Comments
(*) Contamination from FP analysis
(**) Transferred from 120085, initial FP analysis showed that the sample was slightly contaminated with drilling fluid base oil.
(***) Low volume of sample, insufficient liquid produced during flash separation for analysis, gas comp not representative
(****) low pressure reported on opening
Samples used for PVT characterisation of Goldeneye

Multi-Stage Separator Test

Well	well head sample	Contents	Characterization	
14/29a-3	1-24 (2846-S1-F)	Gas	Gas Comp	Recombined
	1-26 (3189-S1-F)	Gas	Gas Comp	Multi stage sep test
	1-27 (3615-S1-F)	Gas	Gas Comp	Comp from flash at J/T conditions

Well	separator sample (DST)	Contents	Characterization
14/29a-3	1-3 (1349-C1-F)(****)	Gas	Gas Comp
	1-4 (2674-C1-F)	Gas	Gas Comp
	1-2 (1648-M1-F)	liquid	Liq Comp

Some of these samples were identified as non-representative of Goldeneye fluids. The finger print analysis indicated contamination. There are sufficient representative fluid samples covering the hydrocarbon column and well spread across the field through the development wells (GYA01 to GYA05), to give confidence in the final characterisation.

Fig. 2-2 and Fig. 2-3 below presents recombined gas compositions and phase envelopes from all exploration, appraisal and development wells considered representative.

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Fig. 2-2—Representative Goldeneye recombined gas compositions from exploration, appraisal and development wells.

Fig. 2-3—Representative Goldeneye well phase envelopes from exploration, appraisal and development wells.

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Fig. 2-2 shows all the valid recombined gas composition samples illustrating that compositionally all the gas samples exhibit a high degree of consistency. The phase envelope in Fig. 2-3 of all the valid samples again illustrates the similarity between all the different samples.

Taking this into account and considering that only GYA03 and GYA04 have a complete PVT suite of experiments, the PVT model currently used by Shell for modelling full field and forecasting was generated based on the surface sample of well GYA03 . Both GYA03 and GYA04 data were matched against the experimental constant mass expansion (CME) and constant volume depletion (CVD) data. Well 14/29a-3 was also matched for completeness.

3. PVT Characterisation

The PVT characterisation used for the current Goldeneye field full field dynamic modelling was re-run using a recent release of the PVT modelling software (PVTsim version 17.3 from CALSEP). This was needed in order to facilitate the subsequent PVT lumping step. There were no changes apparent in moving to the newer version of the software.

In order to achieve a coherent fluid characterisation for Goldeneye, a typical workflow for a gas condensate was followed. It involved:

- Normal regression to tune to general phase behaviour (saturation pressure, CVD and CME observations). Not attempting to get a perfect match since the subsequent lumping process would change the match.

- Grouping/lumping components to reduce simulation time while retaining the predictability of the EOS.

- Fine tuning regression choosing high weights on experiments or observations (saturation pressure, retrograde condensate %, etc.) to improve the match of key data, and finally,

- Matching viscosity data while decoupling the rest of the experiments, regressing on the critical volume for each component's contribution to the total viscosity.

EOS used was Peng-Robinson 78 (PR78) Peneloux[1]. The sample was adjusted to a saturation point of 3815 psia [~263 bara] at a reservoir temperature of 181°F [82.78°C]. The following plots compare the results from the tuned EOS model and the experimental data. Note that the reservoir abandonment pressure is in the neighbourhood of 2200 psia [~151 bara].

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[1] Peneloux, A., E. Rauzy., and R. Freze. "A Consistent Correction for Redlich-Kwong-Soave Volumes". Fluid Phase Eq. 8, 7-27 (1982).

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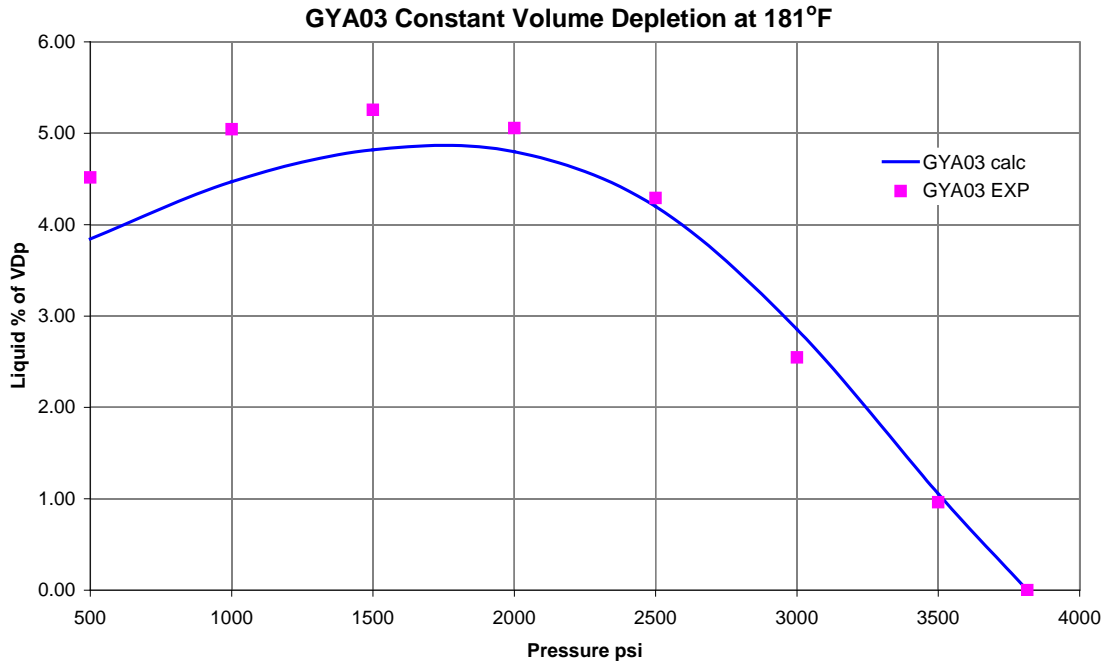
Fig. 3-1—Retrograde condensate %VDp match of CVD data with full characterisation
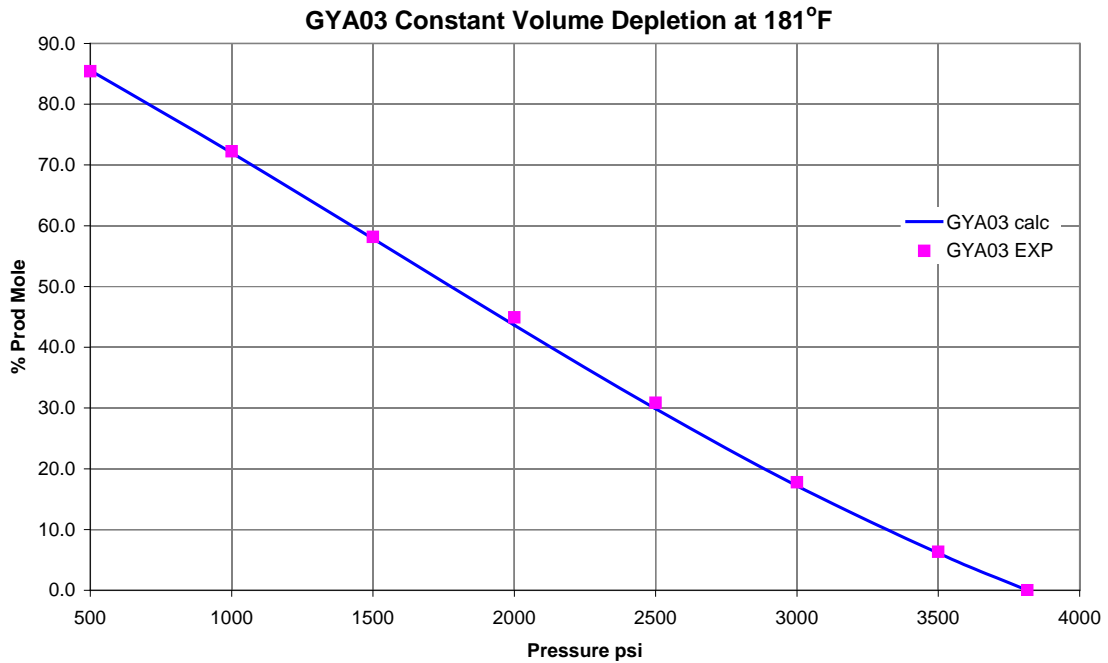
Fig. 3-2—Cumulative % volume of the initial wellstream produced. Match of CVD data with full characterisation

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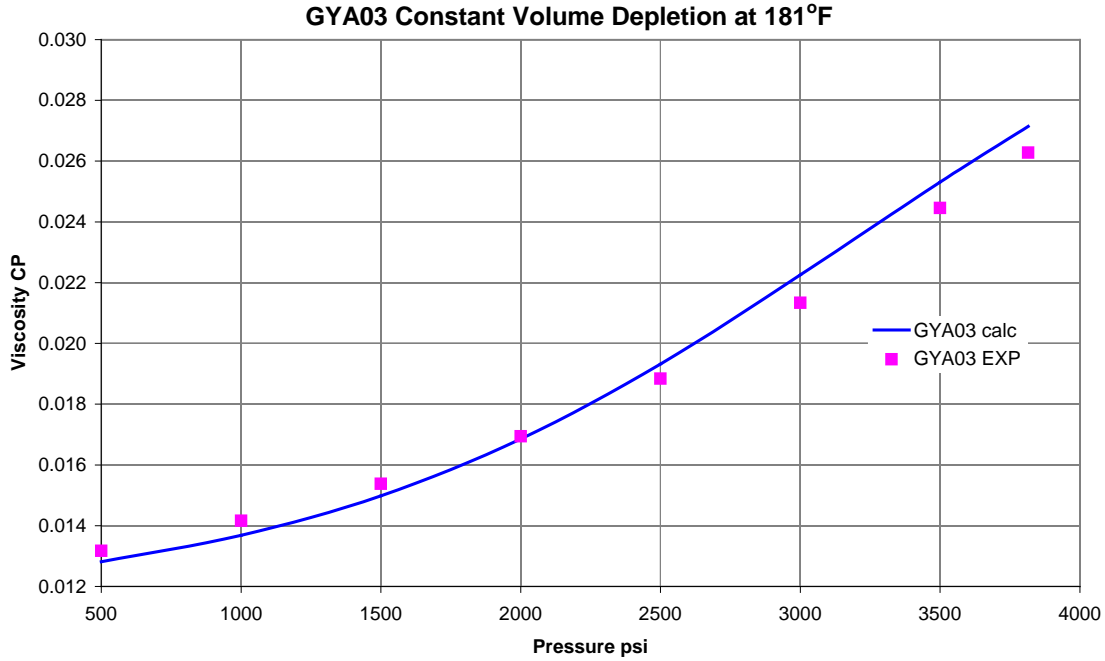
Fig. 3-3—Gas viscosity (calculated) match of CVD data with full characterisation
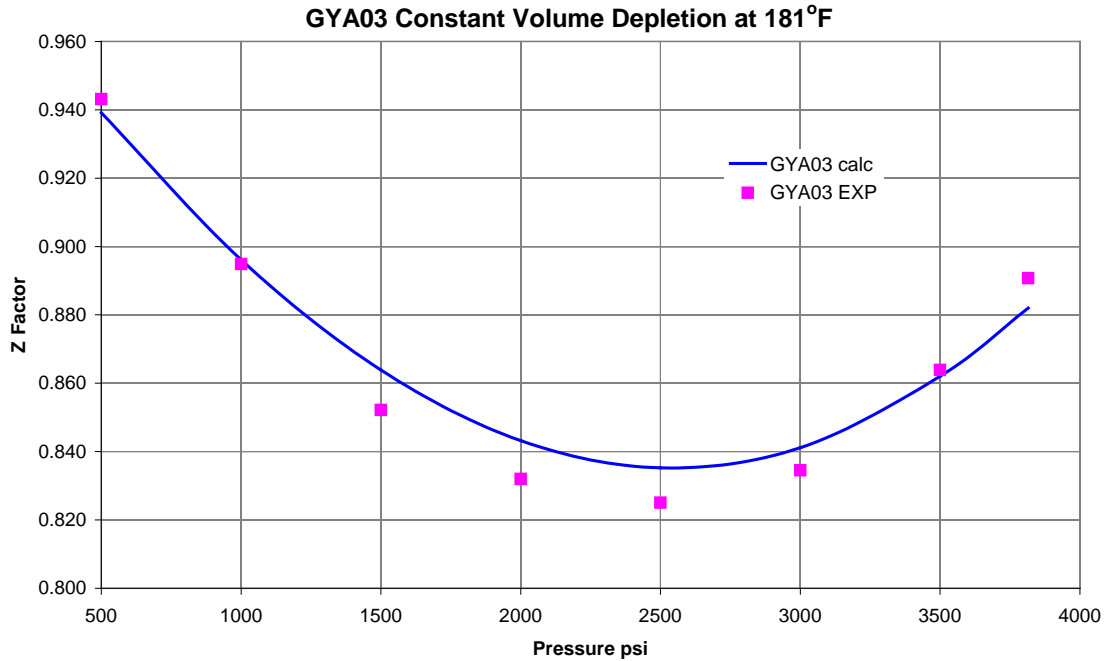
Fig. 3-4—Gas compressibility Z-factor match of CVD data with full characterisation

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Fig. 3-5—Retrograde condensate %VDp match of CME data with full characterisation
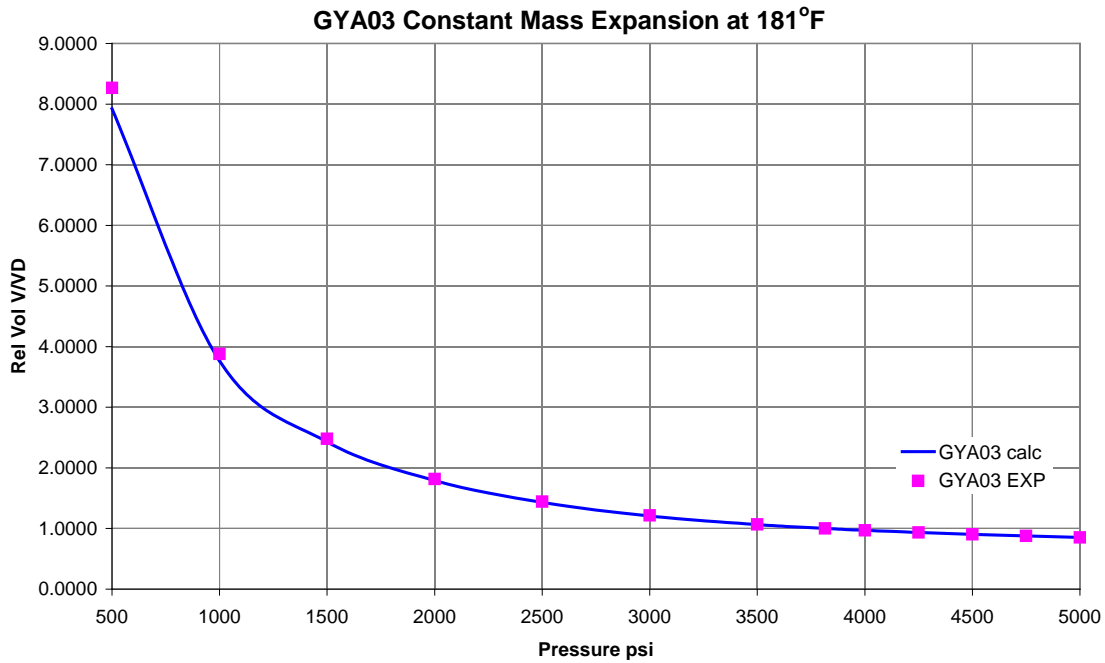
Fig. 3-6—Relative volume match of CME data with full characterisation

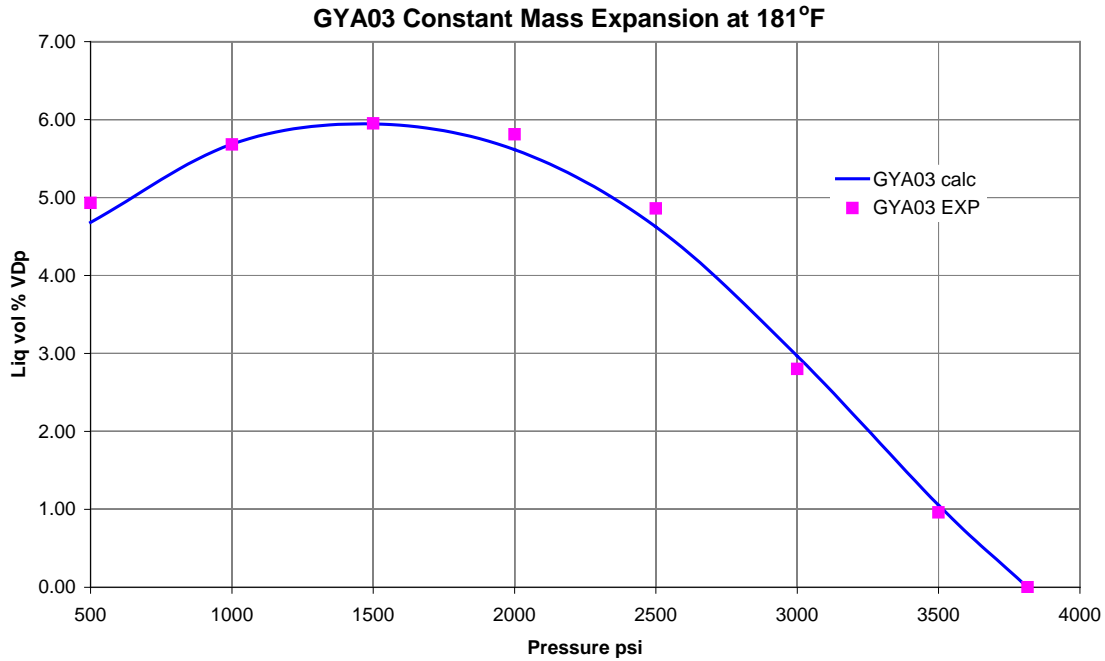Doc. no.: PCCS-05-PT-ZR-0580-00001 – Pressure, Volume and Temperature (PVT) Report   Revision: K02   9

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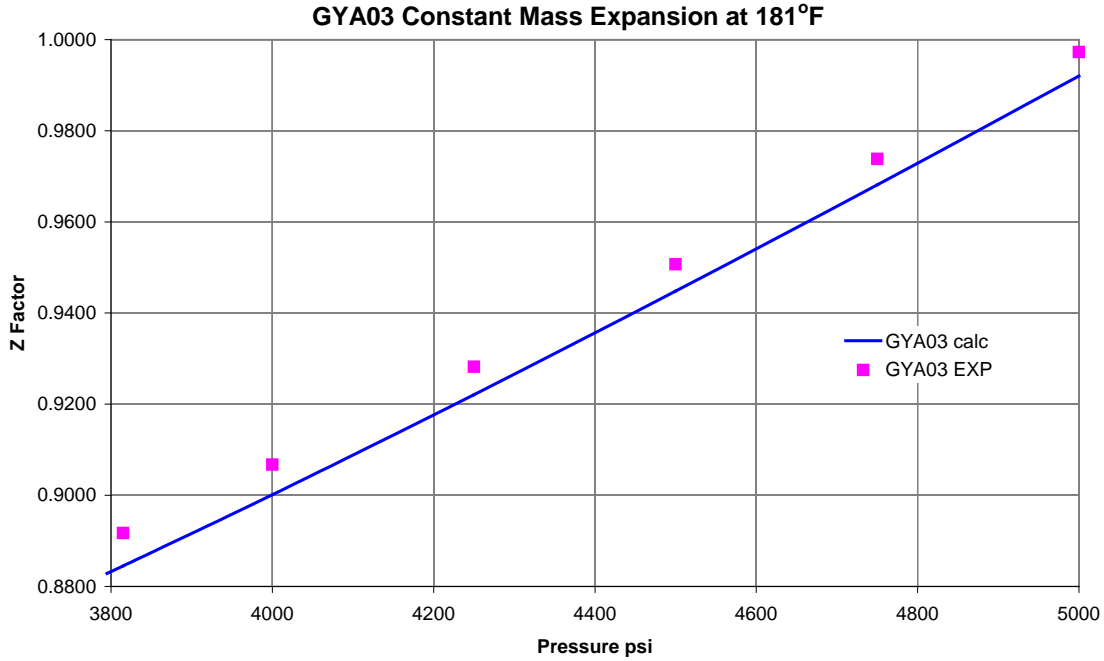
Fig. 3-7—Gas deviation Z-factor match of CME data with full characterisation

The EOS characterisation was modified to improve the predictions of measured data. Nonlinear regression was used to mathematically minimize the difference between PR78P EOS predictions and measured PVT data. Adjustments of EOS parameters such as binary interaction parameters (BIPs) and heavy component critical properties were used. Interaction coefficients between C1 and C7+ pseudo-components were used to improve saturation pressure (Dew Point) representation. A decoupled viscosity experiment regression over critical volume on C7+ pseudo-components was used to match gas viscosity.

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4. PVT lumping scheme

The original calculated recombined fluid composition determined by cryogenic distillation from GYA03 was used for the PR78P EOS calibration; components up to C20+ were included. Generally, if a sufficiently large number of pseudo-components is used to characterise the heavy fraction of a hydrocarbon mixture, a satisfactory prediction of the PVT behaviour by the equation of state can be obtained. However, in compositional models, the cost and computing time can increase significantly with the increased number of components in the system. Therefore, limitations are placed on the maximum number of components that can be used in compositional models and the original components have to be lumped into a smaller number of pseudo-components.

An initial fluid characterisation will typically contain from 13 to 20 components, and sometimes more. A stepwise pseudoisation procedure is recommended in the literature by Curtis Whitson[2], whereby several pseudoised characterisations are developed sequentially (e.g. 15, 12, 10, 7, and 5 pseudo-components). The goal with each pseudoisation is to maintain PVT predictions as close to the original full characterisation as possible. With this stepwise approach, the number of pseudo-components necessary to maintain a required similarity to the original full characterisation is readily determined. Reducing the number of components in a stepwise fashion has three main advantages:

1. It is possible to establish when a further reduction in number of components results in predicted properties that deviate unacceptably from the original N-component characterisation.

2. The procedure usually results in several alternative characterisations with a common basis. One simulation might require more components than another (e.g. radial single-well study versus full-field simulation). Because several characterisations are available, and they are "related" through the original N-component characterisation, more consistency can be expected.

3. Experience has shown that better results are obtained in going from the N-component characterisation to (for example) a 7-component characterisation in several steps, than going from an N-component to a 7-component characterisation in a single pseudoisation.

For the Goldeneye case this was the approach followed. The stepwise pseudoisation procedure recommended by Whitson and presented in the following example, was followed whenever possible.

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[2] Whitson C.H. et al. 1999, Gas Condensate PVT – What's really important and why?, IBC Conference "Optimization of Gas Condensate Fields", London (U.K.)

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Component	EOS22	EOS19	EOS12	EOS10	EOS9	EOS6	EOS4	EOS3
N2	N2	C1N2	C1N2	C1N2	C1N2	C1N2		
CO2	CO2	CO2	CO2	CO2				
C1	C1				CO2C2	C02C2		
C2	C2	C2	C2	C2			C1N2CO2C2-C6	C1N2CO2C2-C6
C3	C3	C3	C3	C3	C3			
IC4	IC4	IC4NC4	IC4NC4	IC4NC4	IC4NC4			
NC4	NC4					C3-C6		
IC5	IC5	IC5NC5	IC5NC5	IC5NC5	IC5NC5			
NC5	NC5							
C6	C6	C6	C6	C6	C6			
C7	C7	C7	C7C8					
C8	C8	C8						
C9	C9	C9	C7C8C9F1F2	C7C8C9F1F2	C7C8C9F1F2	C7C8C9F1F2	C7C8C9F1F2	
C10+	F1	F1	C9F1F2					
	F2	F2						
	F3	F3						C7C8C9F1-F8
	F4	F4	F3-F5					
	F5	F5		F3-F8	F3-F8	F3-F8	F3-F8	
	F6	F6	F6-F8					
	F7	F7						
	F8	F8						
	F9	F9	F9	F9	F9	F9	F9	F9

Fig. 4-1—Example pseudoisation procedure reducing an original EOS characterisation with 22 components to multiple pseudoised characterisations][3]

The original GYA03 composition was described up to C36+ and was reduced down to six pseudo-components. The following figures show the changes in phase behaviour, saturation pressure and retrograde condensate percentage from the CVD experiment, after every grouping step. It is important to see how there are minimal changes in most of the lumping stages and only at the last pseudoisation step, where the fluid characterisation was reduced from eight to six components respectively, is there any significant variation (light blue line in Fig. 4-2). However, it is possible to retune by regression (fine tuning) of the newly created pseudo-components, so that the EOS predictability is maintained after the component reduction (grey line in Fig. 4-2).

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[3] Whitson C.H. et al. 1999, Gas Condensate PVT – What's really important and why?, IBC Conference "Optimization of Gas Condensate Fields", London (U.K.)
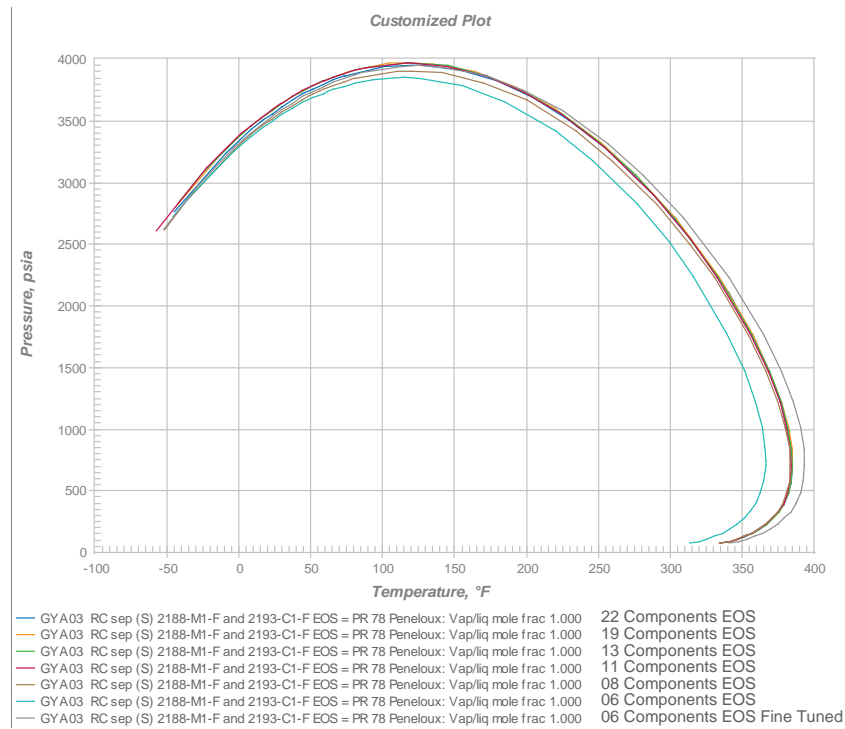
Fig. 4-2—Changes in the phase envelope during the stepwise pseudoisation
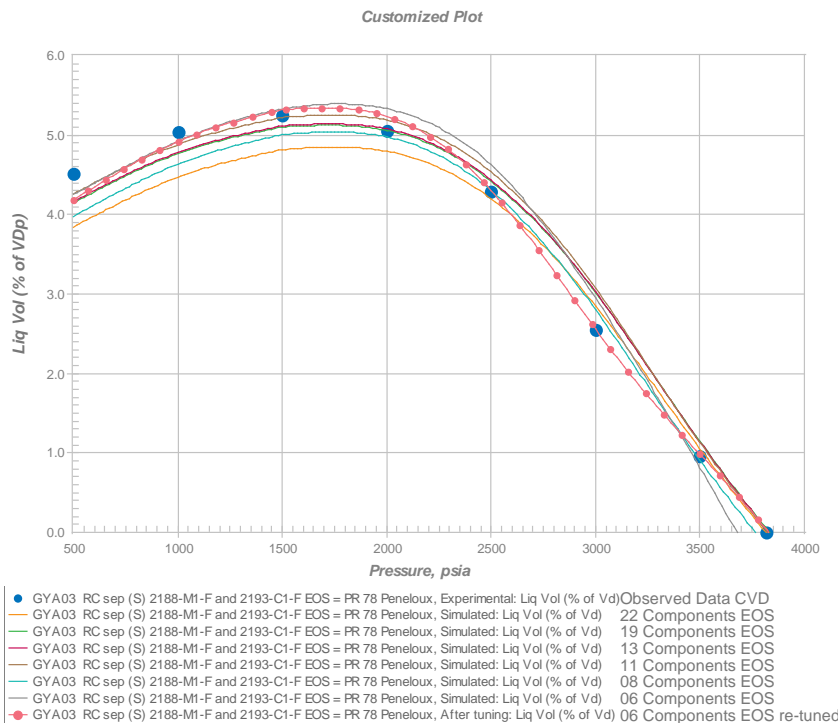
Fig. 4-3—Changes in the retrograde condensate % (Vd) during the stepwise pseudoisation

The fine-tuning of the EOS restored the prediction of the phase behaviour and the changes in the retrograde condensate percentage. Now that the lumping scheme had been implemented with coherent results, it was possible to decouple the viscosity experiment and do a final regression over

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the critical volume for each component's contribution to the total viscosity to match the calculated gas viscosity[4]. The results can be seen in **Fig. 4-4**
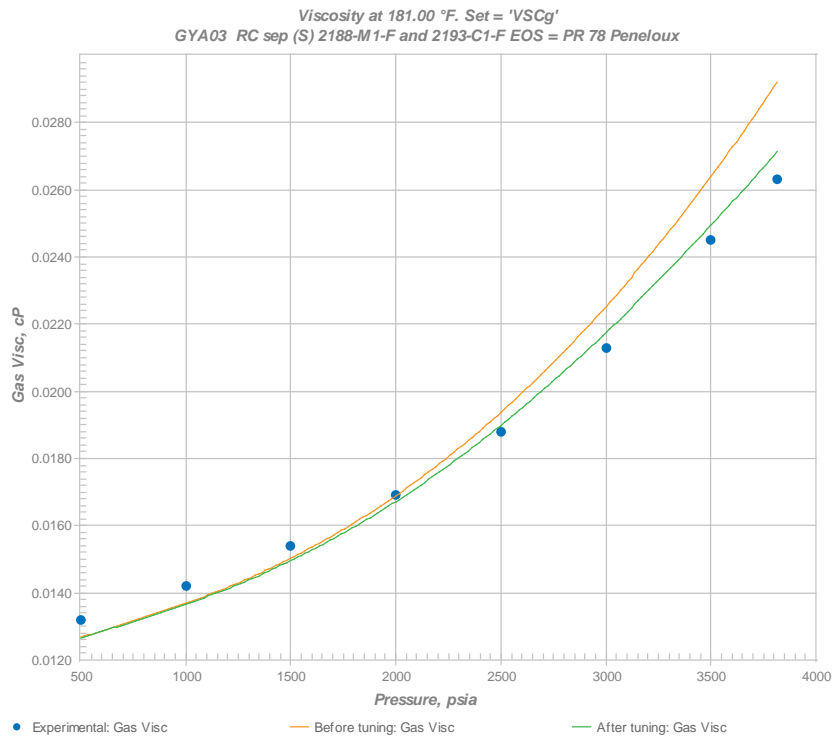
Fig. 4-4—Gas viscosity before and after regressing over the critical volume

Fig. 4-5 shows the mol% of the original GYA03 calculated recombined fluid composition and the resulting pseudoisation lumped into six pseudo-components.

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[4] Gonzalez, Lee and Eakin 1996: "The viscosity of natural gases", J.P.T.

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GYA03 Original Composition

	N2	CO2	C1	C2	C3	iC4	nC4	iC5	nC5	C6	C7	C8	C9	C10	C11	C12	C13	C14	C15	C16	C17-C19	C20-C36
GYA03	0.94	0.37	81.6	6.99	3.2	0.53	1.28	0.42	0.64	1.03	0.91	0.69	0.32	0.29	0.22	0.16	0.12	0.09	0.06	0.05	0.08	0.05

GYA03 Lumped Composition

	CO2	C1N2	C2	C3-C6	C7-C16	C17-C36+
GYA03	0.368	82.509	6.986	7.106	2.898	0.132

Fig. 4-5—GYA03 fluid composition before and after lumping

This lumping scheme is a solution to achieve a manageable fluid characterisation for modelling the displacement processes in Goldeneye CO_2 storage. This characterisation is expected to be applicable in the majority of cases, however, further simplification might be warranted when investigating sensitivities where the mixing of CO_2 and hydrocarbon fluids is insignificant, but where significant increases in the number of grid blocks is required.

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5. CO₂ fluid properties

Depleted oil and gas reservoirs are promising targets for carbon sequestration by direct carbon dioxide (CO_2) injection because of their available volume and proven integrity against leakage. The ability to simulate accurately the supercritical CO_2 and hydrocarbon properties is essential for modelling reservoir processes under injection by CO_2.

Most phenomena related to CO_2 dynamics become apparent with an understanding of the key fluid properties and their dependence on temperature and pressure.
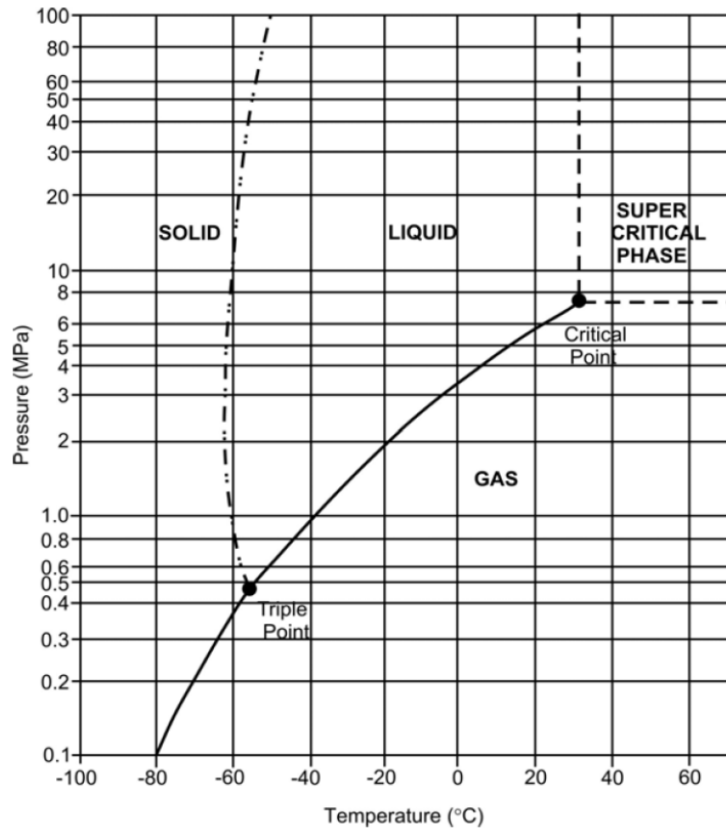

Fig. 5-1.—Phase diagram of CO₂ 5

The critical temperature of CO_2 is 31.1° C and the critical pressure is 73.8 bara [7380 KPa] . At temperatures and pressure above this critical point, CO_2 exists as a supercritical fluid, whereby it has a density similar to a liquid but exhibits gas-like viscosity (and compressibility) (Fig. 5-1).

Pure component CO_2 properties were calculated using the Thermophysical Properties of Fluid Systems from the National Institute of Standards and Technology (NIST)[6]. These properties were later used as observation points to test the predictions of the Peng-Robinson version 1978 EOS in its volume shift corrected Peneloux version.

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[5] Wong, S.: "Module 4 – CO₂ Compression & Transportation to Storage Reservoir". APEC Reference #205-RE-01.3, 2005.

[6] http://webbook.nist.gov/chemistry/fluid

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The standard EOS descriptions used in reservoir simulation were developed for hydrocarbon systems and can be less accurate for modelling CO_2 properties and some other non-hydrocarbon components such as H_2S and SO_2. As a consequence, the default pure component parameters require tuning to assure the best possible match over a range of interest. A set of comparison plots was prepared in order to assess the magnitude of tuning needed.

The analysis was carried out under Goldeneye reservoir conditions at an isothermal reservoir temperature of approximately 181 ºF [82.78 ºC] and for a pressure range that covers the probable abandonment pressure (~2000 psia [~138 bara]) up to reservoir initial conditions of about 3835 psia [~264 bara].

CO_2 physical properties that strongly affect flow and transport are density (ϱ), viscosity (μ), and solubility of CO_2. CO_2 dissolution in brine will be modelled using Henry's Law, whilst density is calculated directly from the EOS, and viscosity in this case calculated from the Lohrenz-Bray-Clark correlation. The following **Fig. 5-2** shows the differences between experimental data from NIST and that estimated in PVTSim through the EOS using the default pure component parameters.
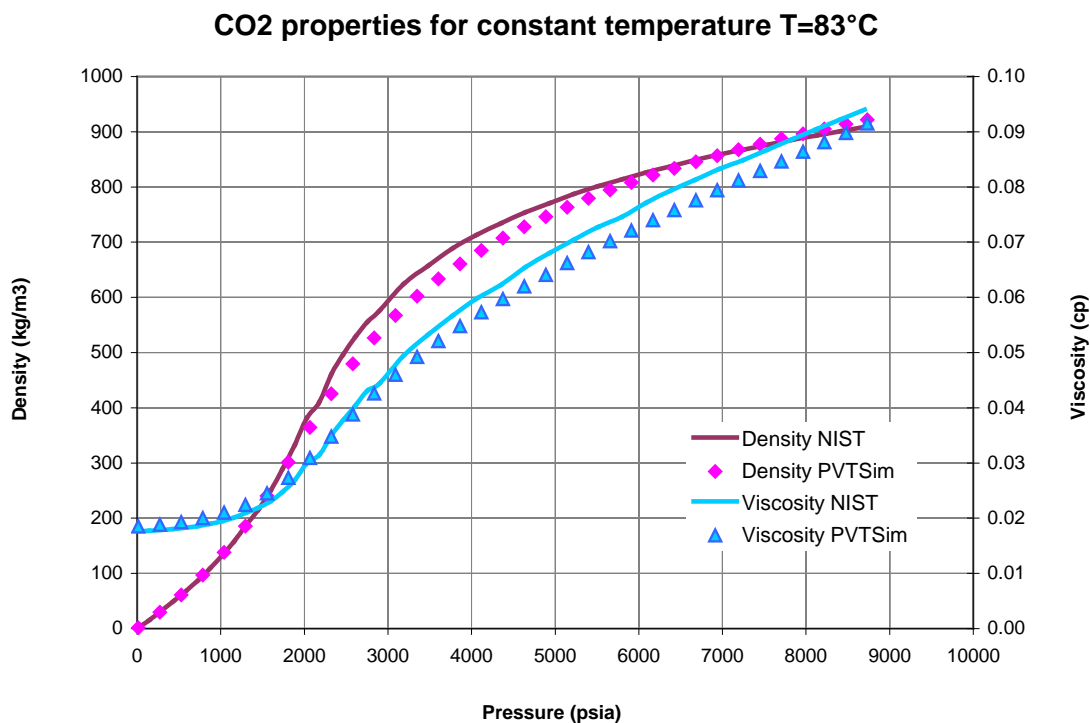
Fig. 5-2—CO₂ Density and Viscosity as a function of pressure. NIST data vs. PVTsim PR78P EOS

The comparison between NIST and PVTSim shows some differences in CO_2 physical properties especially in the range of pressure that Goldeneye will operate (2000 to 3830 psia [138 to 246 bara]). Over this range PVTsim through PR78P EOS overestimates both CO_2 density and viscosity. Fig. 5-3 shows the error percentage between the two sources (PVTsim / NIST) for a range of pressures and temperatures for the default pure component EOS parameters. The light blue thicker line represents 80 ºC, near Goldeneye reservoir temperature.
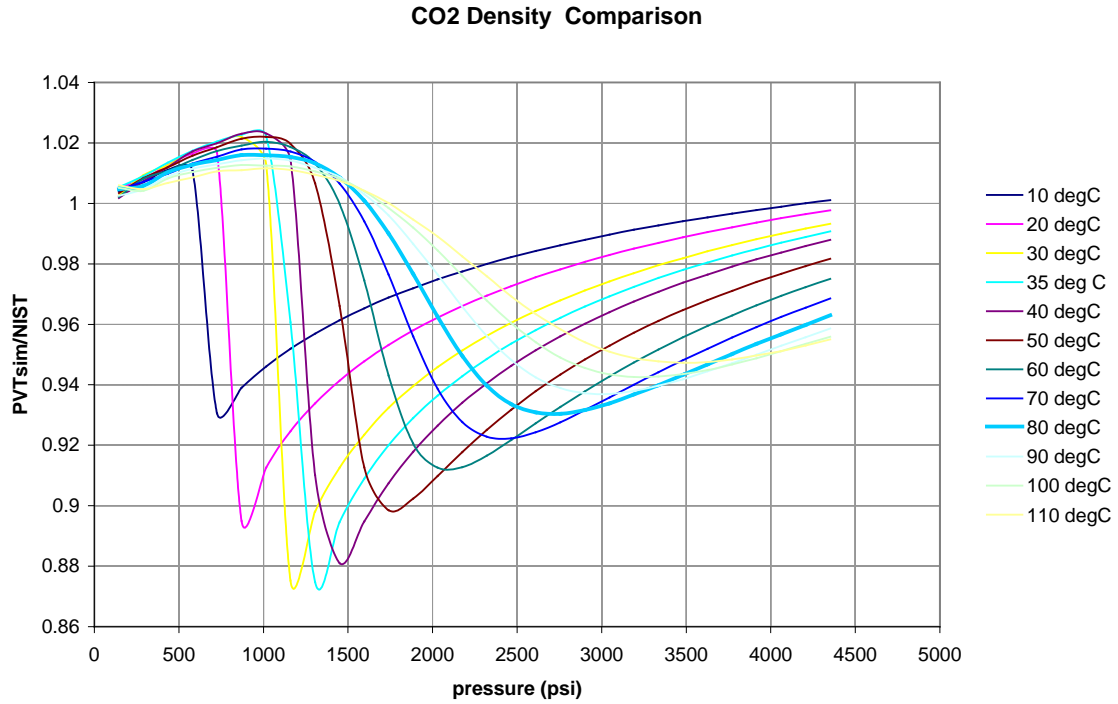
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CO2 Density Comparison

Fig. 5-3—CO$_2$ density error function versus pressure (psia) for a range of temperatures

CO2 Viscosity Comparison

Fig. 5-4—CO$_2$ Viscosity error function versus pressure (psia) for a range of temperatures

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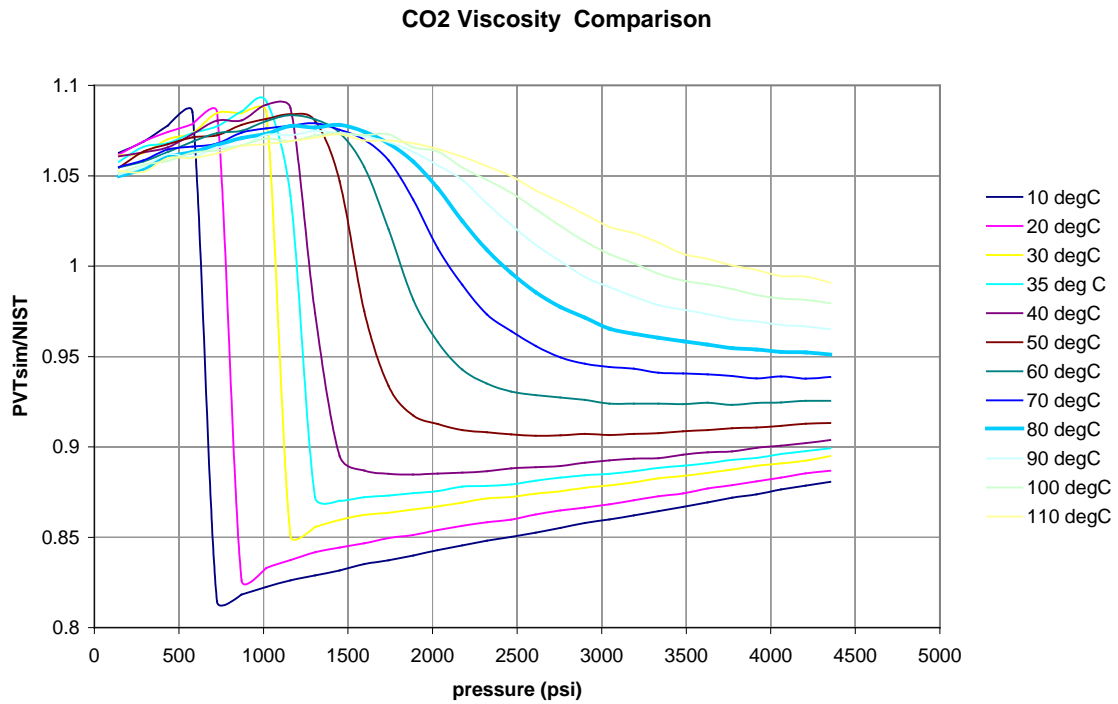
At Goldeneye reservoir conditions, the error in CO_2 density could be as much as 7% while in viscosity only 4.5%.

As previously discussed, issues with EOS predictions of liquid density and other fluid properties, particularly in the vicinity of the critical region, are well documented. The drawback is that the critical compressibility factor takes on a universal critical compressibility of 0.307 for all substances. Consequently, the molar volumes are typically overestimated and hence, densities are underestimated. Peneloux et al. developed a procedure for improving the volumetric predictions of EOS by introducing a volume correction parameter c_i into the equation. This third parameter does not change the vapour-liquid equilibrium conditions determined by the unmodified equation, i.e., the equilibrium ratio K_i, but modifies the liquid and gas volumes. This methodology is known as the volume translation method and was used in the CO_2 density calculation depicted previously.

This methodology has been implemented successfully on previous occasions in the prediction of the behaviour of naturally occurring hydrocarbon systems. However as has been shown previously, CO_2 is a particular component with very specific behaviour for a given set of pressure and temperature conditions. Even though the default values for volume translation for pure component CO_2 have been used, these do not always improve the calculation of density, especially where CO_2 is in a dense phase under critical or super critical conditions.

It was decided to regress over the CO_2 volume shift parameter (C_{pen} after Peneloux) in order to minimize the error between PR78 Peneloux EOS and NIST. Fig. 5-5 shows the impact of C_{pen} values in CO_2 density prediction.
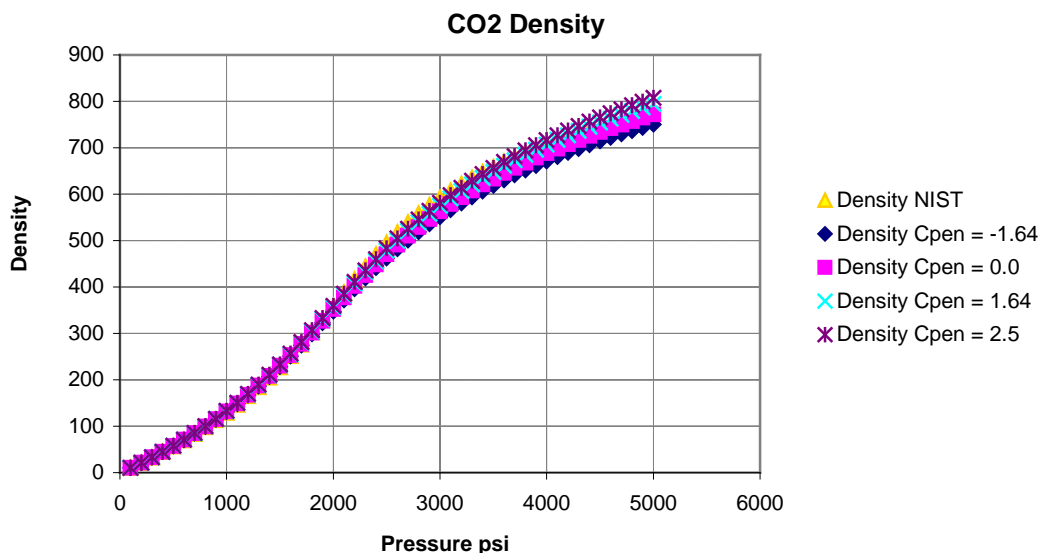
Fig. 5-5—CO₂ density comparison among NIST and different volume shift values (Cpen) for PR78 Peneloux EOS

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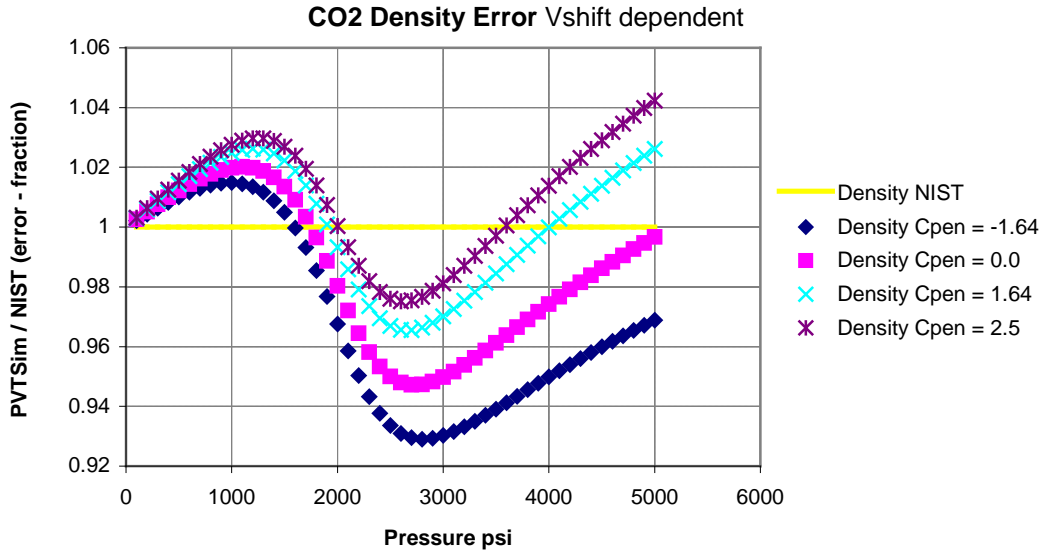
CO2 Density Error Vshift dependent

Fig. 5-6.—CO₂ Density error function versus pressure (psia) for a range of Cpen values

It may be seen that modifying the value of volume shift parameter (C_{pen}) reduces the error between NIST and EOS, but nevertheless there is an error pressure dependency that cannot be corrected completely. Adjusting C_{pen} from –1.64 cm³/mol to a higher value of 2.5 cm³/mol yields a better match against the NIST data, reducing the error to 2.5% in the pressure range of interest (2000 to 3835 psia [138 to 254 bara]). This makes CO_2 slightly less dense than the NIST data. This is a conservative assumption for the CO_2 modelling, meaning that the plume will spread further away in the same proportion.

Viscosity is calculated using the Lohrenz-Bray-Clark correlation which relates it to a fourth-degree polynomial in the reduced density, $\rho_r = \rho/\rho_c$. Sensitivities in the Volume Shift affects both CO_2 density and as a consequence, viscosity of CO_2. **Fig. 5-7** show the impact.
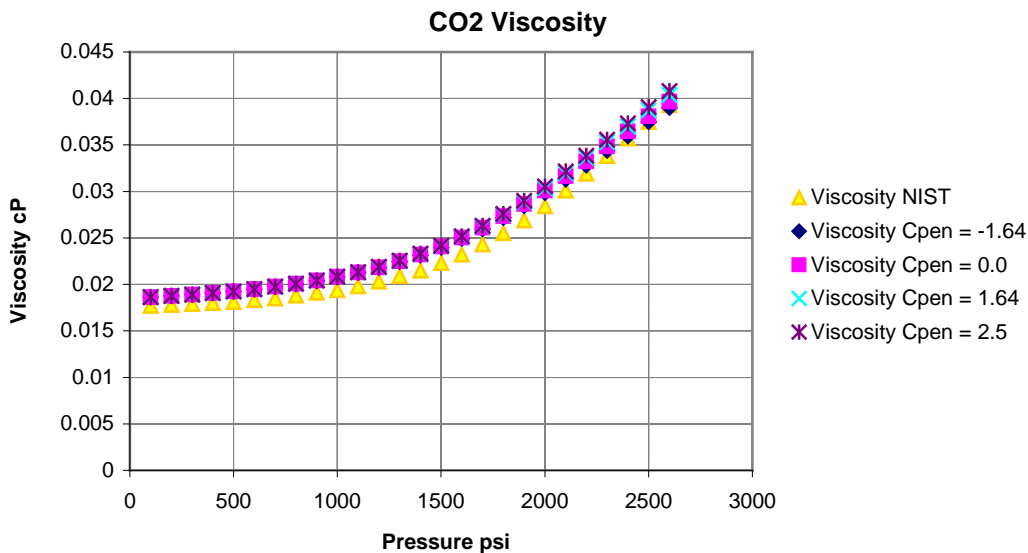
CO2 Viscosity

Fig. 5-7—CO₂ viscosity comparison among NIST and different volume shift values (Cpen) for PR78 Peneloux EOS

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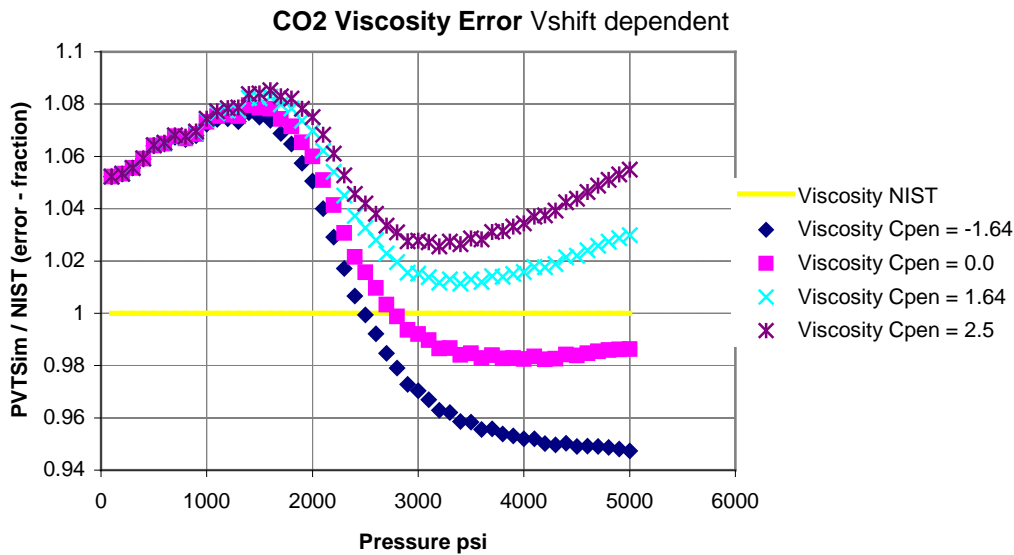
Fig. 5-8—CO₂ Viscosity error function versus pressure (psia) for a range of Cpen values

It may also be seen that modifying the value of volume shift parameter (C_{pen}) reduces the error between NIST and EOS, and the same pressure dependency of the error is seen in viscosity. Adjusting C_{pen} from –1.64 cm³/mol to a higher value of 2.5 cm³/mol will increase the error up to 7.5% at low pressure (2000 psia [138 bara]).

In order to obtain a balance between the error in density and viscosity, a mid value of C_{pen} has been chosen. A value of C_{pen} = 1.64 cm³/mol offers a mid-point between benefits, allowing a reduction of the density error from 7% to 3.2%, while maintaining viscosity error at around 2.5% in average for the pressure range of interest.

It is important to understand that these differences in CO_2 viscosities are small in comparison with the range of uncertainty of other parameters such as the relative permeabilities.

When outputting the gas condensate EOS model from PVTsim in "simulator", the volume shift parameters (C_{pen}) for the CO_2 component are modified as described above.

6. Conclusion

Effective modelling of the Goldeneye hydrocarbon reservoir fluids and the injected CO_2 properties is achievable using a Peng-Robinson EOS representation, lumped into six components. The densities and viscosities are modelled using the Peneloux correlation adjunct to the PR EOS, and the C_{pen} factor has to be tuned to yield correct CO_2 properties at Goldeneye storage conditions.

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7. Abbreviations

BIP	Binary Interaction Parameters, also known as Binary Interaction Coefficients (BIC)
CCS	Carbon, Capture and Storage
CME	Constant Mass Expansion, also known as Constant Composition Expansion (CCE)
CO_2	Carbon Dioxide
C_{pen}	Volume Shift Parameter after Peneloux
CVD	Constant Volume Depletion
D_p	Dew Point
DST	Drill Stem Test
EOS	Equation of State
H_2S	Hydrogen sulphide
LBC	Lohrenz-Bray-Clark
MDT	Modular Dynamic Tester
NIST	National Institute of Standards and Technology
P&T	Pressure and Temperature
PR	Peng-Robinson
PVT	Pressure, Volume and Temperature
RFT	Repeat Formation Tester
SO2	Sulphur dioxide
VD_p	Volume at Dew Point

8. Glossary of Unit Conversions & Well Naming

For the provision of the SI metric conversion factor as applicable to all imperial units in the Key Knowledge Deliverable.

Table 8-1: Unit Conversion Table

Function	Unit - Imperial to SI Metric conversion Factor
Length	1 Foot = 0.3048m Metres 1 Inch = 2.54cm Centimetres 1 Inch = 25.4mm millimetres
Pressure	1 Psia = 0.0690 Bara
Temperature	1°F Fahrenheit = -17.22°C Centigrade
Weight	1lb Pound = 0.45kg Kilogram

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In the text well names have been abbreviated to their operational form. The full well names are given in Table 8-2 below.

Table 8-2 Well Name Abbreviations

Full well name	Abbreviated well name
DTI 14/29a-A3	GYA01
DTI 14/29a-A4Z	GYA02S1
DTI 14/29a-A4	GYA02
DTI 14/29a-A5	GYA03
DTI 14/29a-A1	GYA04
DTI 14/29a-A2	GYA05

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