

Gas Miscibility and Interfacial Properties Study
Using the PRIme[™] Method For Statoil

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Project Team

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Confidential

Introduction

Interfacial tension (IFT) is measured by either pendant drop or sessile drop. wettability is determined by either contact angle or degree of adhesion (Note: a contact angle of zero, means the solid surface is totally wetted by the non-drop phase).

In the case of wettability it is important to state which phase is the initiating phase to contact the solid surface. This is important because to wet the surface, the drop must displace the initial phase (initiating) and then bond to the surface through a chemical /physical reaction. Typically, the aqueous phase is the initiating phase and the oil phase are the drops.

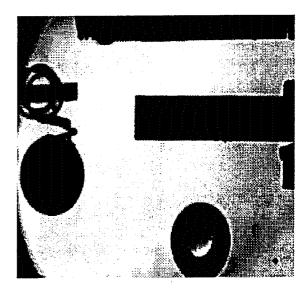


Figure 1

Since the fluids used are complex samples containing a mixture of compounds it should be noted that the image analysis methods used to compute the IFT are complex and produce the best value possible from systems which are of poor optical quality.

The IFT computations are carried out using the axis symmetric drop shape profile (ADSA-P) method. This method gives good results for both pendant and sessile drops. When the drops are of poor quality we use software and methods developed at PRI (Fisher) to find the best estimate of the drop shape co-ordinates to be used in the ADSA-P model.

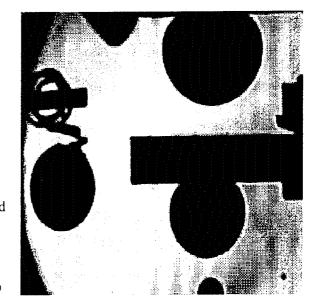


Figure 2

In Figure 1 we show an example of a good pendant drop shape and in Figure 2 we show a good sessile drop shape. On the left side of both images you can see an object connected by a wire to a tube. This is the calibration sphere, it is made of synthetic ruby and has a known dimension (0.47 cm). The sphere in the

image appears as an ellipse. The reason for this is that the images are digitized using a system which uses rectangular pixels with an aspect ratio of 1.25. When rectangular pixels are displayed on a system which uses square pixels the objects appear shorter in the X dimension by this ratio.

When we calculate the physical dimensions of the drops we must convert the X co-ordinates by a factor which contains not only the aspect ratio but a conversion for pixels to centimeters. Because we need both an X and Y calibration factor we must use the co-ordinates from the ruby sphere to compute this information. We make use of the fact that the equation for an ellipse gives us both dimensions at once. We fit the co-ordinates of the sphere to this equation by leastsquares. This gives us the best estimate of the conversion factors and the aspect ratio simultaneously. If the fluids are turbid or have a high molar absorbtivity we must use special image analysis techniques to find the edge co-ordinates for both the sphere and the drops. These methods allow us to calculate IFT values under extreme conditions when the more traditional methods fail.

Results

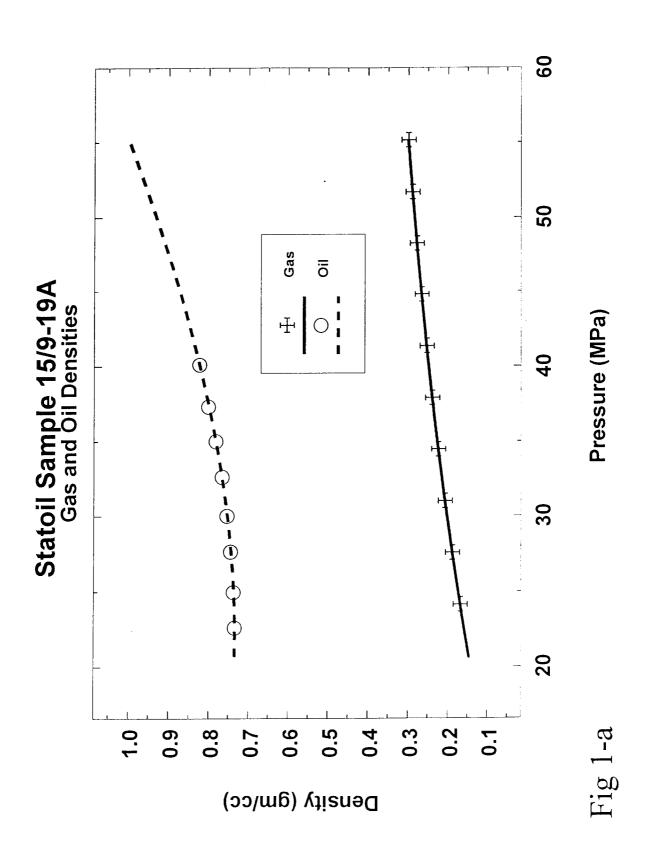
Measurements of IFT were made over a range of pressures using the supplied live oil sample (15/9-19A), using both CO_2 and the supplied hydrocarbon mixed gas. The Figures 1-a to 17-a are the data for the mixed gas whereas the Figures 1-b to 16-b are for the CO_2 .

In the case of the mixed gas we observed that the decline rate for the apparent IFT Vs Pressure did not follow the traditional pattern(Figure 2-a), here we see that there is a jump around the 37MPa mark, this change needs some further discussion.

You will notice that in all of the images at all of the pressures we see some streams of oil dissolving in the gas phase. This represents stripping of the light ends from the oil by the hydrocarbon gas. If the oil has any significant concentration of heavy components these will be enriched in the sample and the apparent IFT will be higher than expected. The discontinuity observed in the graph is the result of sampling rate not any smooth functional relationship, although the lines are calculated by nonlinear leastsquares so that all of the data was used to define the decline slope in order to estimate the intersection point. We feel that the value of this intersection is elevated due to the enriching process. In other words the miscibility process is continuos over pressure rather than rapid at or after a given pressure.

In the case of the CO₂ we see the same effect but magnified. The degree of miscibility is much higher than the mixed hydrocarbon gas, even at the first pressure which is quite near the bubble point. Since the extraction process is rapid and large the drops are distorted in shape and as such the calculated apparent IFT has a much larger error band. So even though the graph indicates a simple straight line (Figure 2-b) we feel that this value is also elevated from what is probably the best estimate of the minimum miscibility pressure. We also observed the same enrichment process in this mixture as well.

The image sequence shown in Figures 8-b to 13-b represent a series of drops appearing at slower and slower rates. This allow for longer and longer extraction periods and as such more and more heavy component left on the tip. This is especially obvious in the sequence starting at 14-b. Here we show the extraction process in more detail, and then allow it to end in a stable higher IFT drop which stays on the tip for some time ending in the shapes shown in Figure 15-b. It is clear that the bulk of the oil is extracted early on and the remaining material is not miscible at these pressures and will not be miscible for some significant increase in pressure.



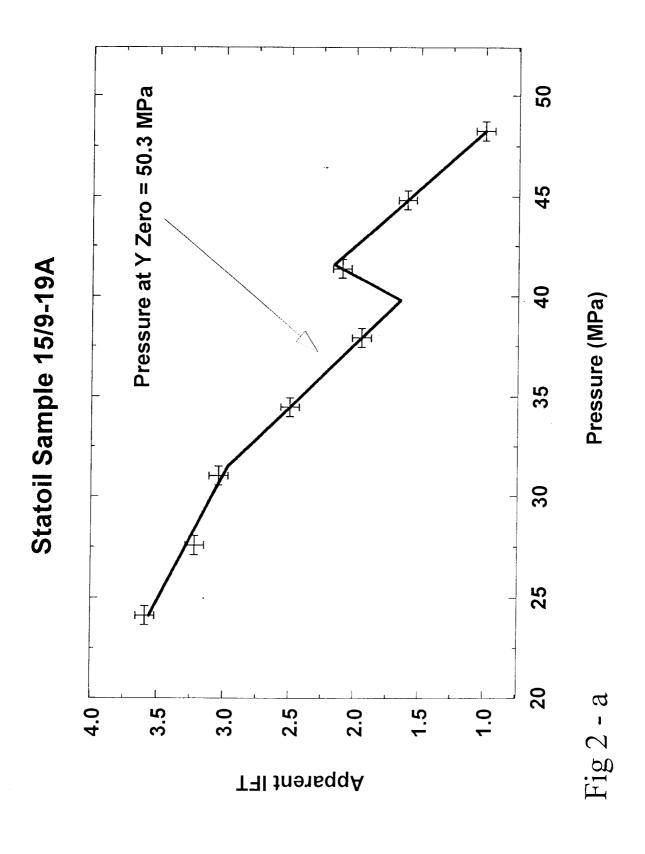
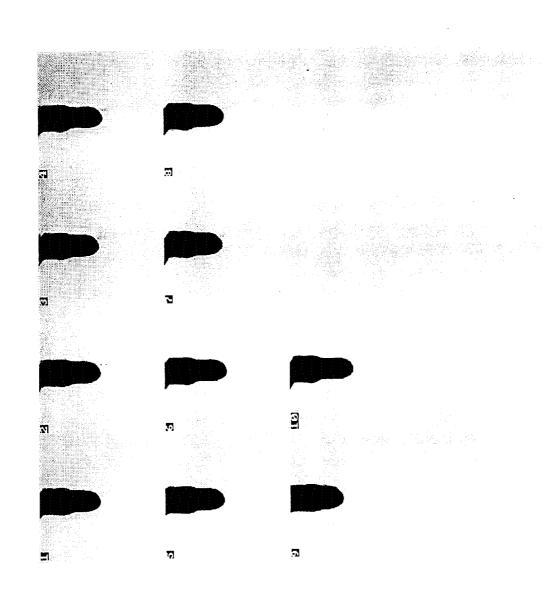


Fig 3 - a

Statoil Sample 15/9-19A Mixed Gas at 27.6 MPa and 110 C



 $\dots = \bigcup_{i \in \{1, \dots, k\} \in \mathcal{K}} (i, k) \cup \bigcup_{i \in \mathcal{K}} (i, k) \cup \bigcup_{i$

Statoil Sample 15/9-19A Mixed Gas at 31.0 MPa and 110 C

Fig 6 - a

Statoil Sample 15/9-19A Mixed Gas at 37.9 MPa and 110 C

Fig 8 - a

Fig 9 - a

Fig 10 - a

Statoil Sample 15/9-19A Mixed Gas at 41.4 MPa and 110 C

Mixed Ga

Fig 11 - a

Statoil Sample 15/9-19A Mixed Gas at 44.8 MPa and 110 C

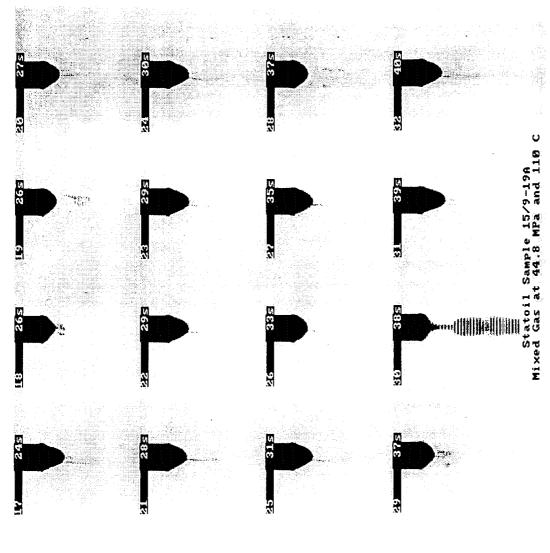


Fig 13 - a

Statoil Sample 15/9-19A Mixed Gas at 44.8 MPa and 110 C

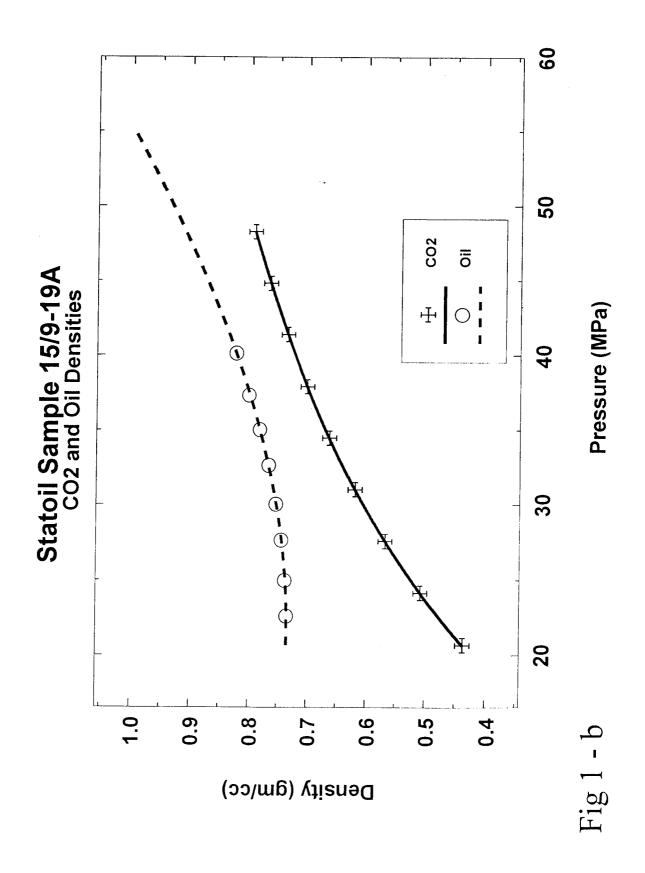
Statoil Sample 15/9-19A Mixed Gas at 48.2 MPa and 110 C

Fig 15 - a

Statoil Sample 15/9-19A Mixed Gas at 48.2 MPa and 110 C

Fig 16 - a

Statoil Sample 15/9-198 Mixed Gas at 48.2 MPa and 110 C



Pressure at Y Zero = 32.1 MPa Statoil Sample 15/9-19A 6 Apparent IFT

Fig 2 - b

Pressure (MPa)

Fig 3 - b

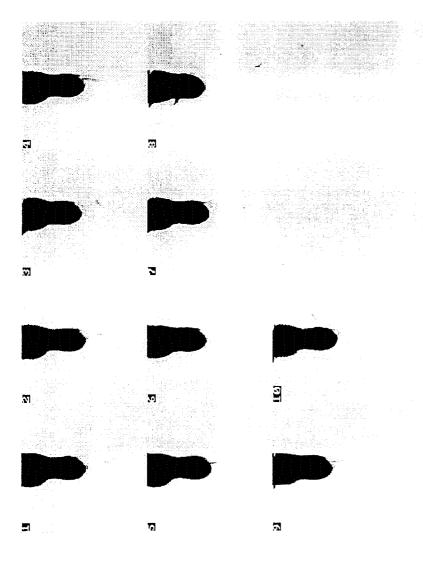
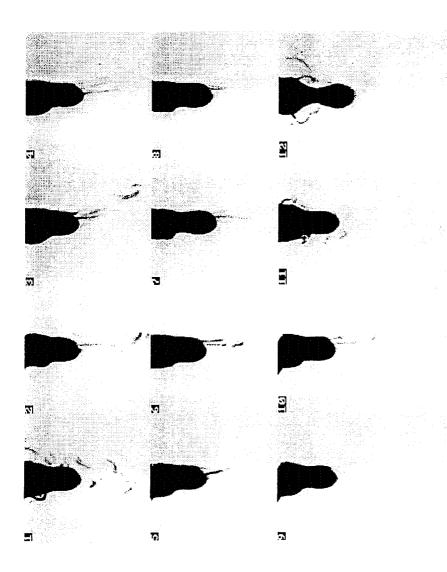
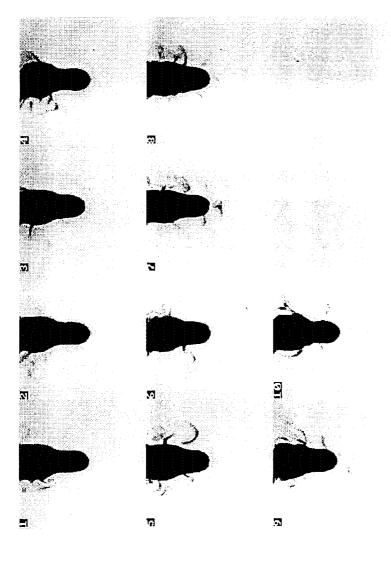


Fig 4 - b



Statoil Sample 15/9-19A CO2 at 24.1 MPa and 110 C



Statoil Sample 15/9-19A CO2 at 25.5 MPa and 110 C

Fig 6 - b

Fig 7 - b

Statoil Sample 15/9-196 CO2 at 31.8 MPa and 118 C

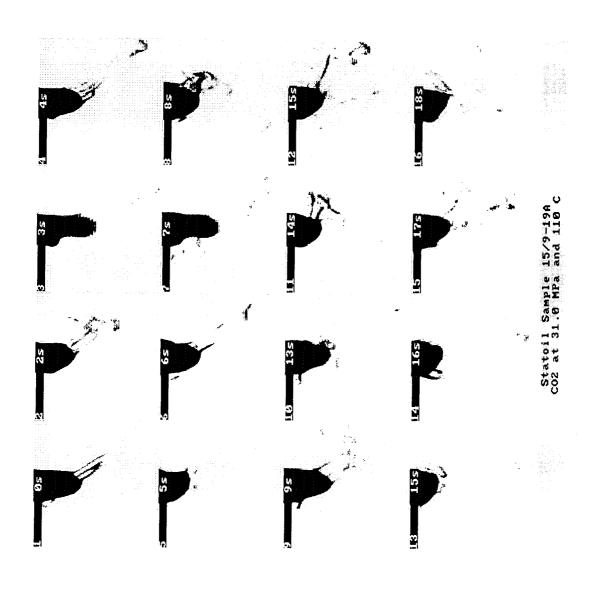


Fig 9 - b

Statoil Sample 15/9-19A CO2 at 31.0 MPa and 110 C

Fig 11 - b

Statoil Sample 15/9-19A CO2 at 31.0 MPa and 110 C

Statoil Sample 15/9-198 CO2 at 31:0 MPa and 110 C

Statoil Sample 15/9-19A CO2 at 31.0 MPa and 110 C

Fig 14 - b

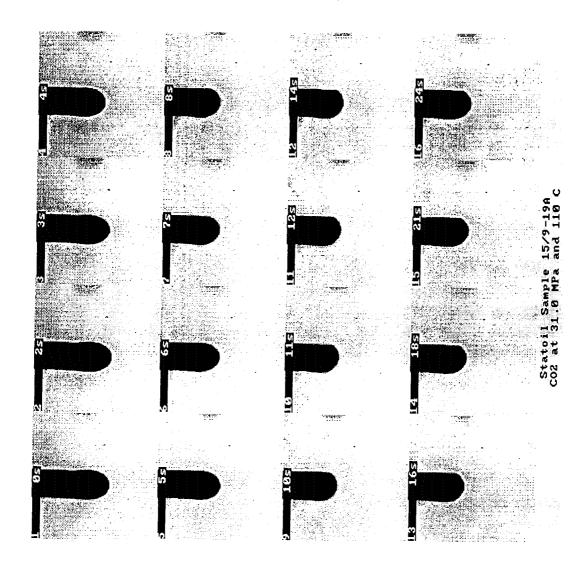


Fig 15 - b

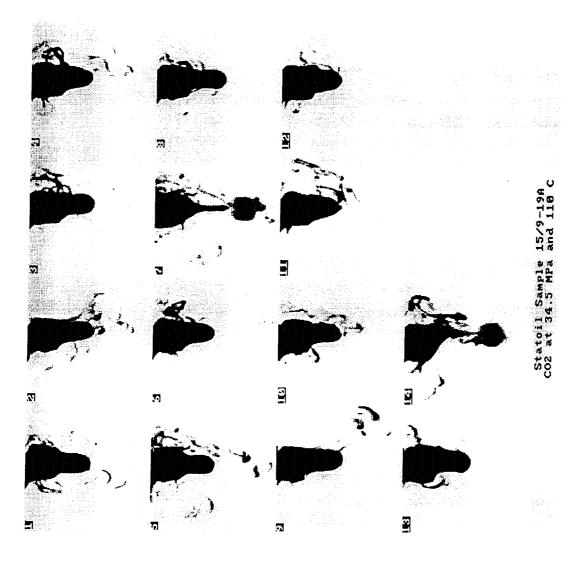


Fig 16 - b