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	REPORT ON STABLE ISOTOPES ( ON NATURAL GASES FROM WELL	REV. NO.					
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SUMMARY		DISTRIBUTION					
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#### 1. INTRODUCTION

Two gas samples from well 2/12-1, DST 1 and DST 2 was received March 1987.

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On the samples  $C_1 - C_4$  and  $CO_2$  are quantified, and the  $\delta^{13}C$  value is measured on methane, ethane, propane, the butanes and  $CO_2$ . The  $\delta D$  value is also measured on methane.

### 2. ANALYTICAL PROCEDURE

The natural gases have been quantified and separated into the different gas components by a Carlo-Erba 4200 instrument. This gas chromatograph is equipped with a special injection loop in order to concentrate the samples, in the case of low concentration of the gas components. The hydrocarbon gas components were oxidized in separate CuO-ovens in order to prevent cross contamination. The combustion products  $CO_2$  and  $H_2O$  were frozen into collection vessels and separated.

The water was reduced with zinc metal in a sealed quartz tube to prepare hydrogen for isotopic analysis. The isotopic measurements were performed on a Finnigan Mat 251 and a Finnigan Mat delta mass spectrometer. Our  $\delta^{13}$ C value on NBS-22 is -29.77 ± .06 o/oo PDB.

### 3. RESULTS

The volume composition of the samples is given in Table 1. The results have been normalized to 100%. The stable isotope results are given in Table 2.

Our uncertainty on the  $\delta^{13}C$  value is estimated to be  $\pm 0.3$  o/oo and includes all the different analysis step. The uncertainty on the  $\delta D$  value is likewise estimated to be  $\pm 5$  o/oo.

Sample	IFE no.	с <sub>1</sub> %	с <sub>2</sub> %	с <sub>3</sub> %	i-C <sub>4</sub> %	n-C <sub>4</sub> %	co <sub>2</sub> %	Σc <sub>1</sub> -c <sub>4</sub>	$\frac{\Sigma C_2 - C_4}{\Sigma C_1 - C_4}$	$\frac{i-C_4}{n-C_4}$
DST 1 DST 2	6088 6089				0.62 0.59		2.3 2.2	97.7 97.8	0.24 0.23	0.35 0.33

Table 1 Volume composition of gas samples from well 2/12-1

Table 2 Isotopic composition of gas samples from well 2/12-1

Sample	IFE	C <sub>1</sub>	C <sub>1</sub>	C	C_3	i-C <sub>4</sub>	n-C <sub>4</sub>	CO <sub>2</sub>	
	no.	δ <sup>13</sup> C PDB	ôd Smow	δ <sup>13</sup> C PDB	δ <sup>13</sup> C PDB	δ <sup>1.3</sup> C PDB	δ <sup>13</sup> C PDB	δ <sup>13</sup> C PDB	δ <sup>18</sup> 0 PDB
DST 1	6088	-51.7	-255	-34.8	-30.2	-29.8	-29.8	- 5.5	-11.5
DST 2	6089	-50.9	-250	-34.3	-30.3	-31.5	-30.2	- 6.1	-12.5

#### 4. INTERPRETATION

The isotopic results, Table 2, indicate a mixed gas derived from two or more different sources. Gases derived from the same source but at different maturity levels is also a possibility.

The combined use of the  $\delta^{13}C$  methane and  $\delta^{13}C$  ethane values, Figure 1 (Schoell 1983)<sup>\*</sup> may indicate that the thermogenic methane is mixed with a small component of biogenic methane. The relative light  $\delta^{13}C$  values of methane and ethane may, however, indicate that the organic matter of the source rock is isotopically light.

If the gas components C<sub>1</sub> and C<sub>4</sub> are from the same source, the isotopic separation between  $\delta^{13}C_3$  and  $\delta^{13}nC_4$  indicate a high maturity situation. This may also be in agreement with the  $\delta^{18}$ O-values of the reservoar CO<sub>2</sub>-gas. If this CO<sub>2</sub> is mainly derived from the source rock and

the CO<sub>2</sub> equilibrated with CaCO<sub>3</sub> in the source rock, a high temperature (probably in excess of  $120^{\circ}$ C) or a high maturity is indicated, at least for parts of the reservoir gas. A high maturity situation is, however, not in accordance with the combined use of the  $\delta^{13}$ C and the  $\delta$ D values of methane (Fig. 1, Schoell 1983)\*, where the results suggest a maturity situation in the middle of the oil window.

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# 5. CONCLUSION

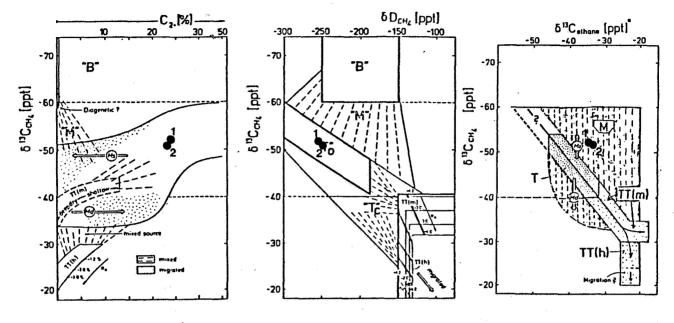
The isotope study of natural gas samples from well 2/12-1 indicates a mixed gas.

The mixed gas is derived from two (or more) different sources or from one source at different maturity levels. The source rock (or rocks) is in this case characterized by isotopically light stable isotopes.

It is in the present case (because of the mixed gas) difficult to use the isotopes to indicate a fixed maturity level of the source rocks. A moderate to high maturity level in the oil window is, however, suggested.

Schoell, M. (1983): Genetic Characterization of Natural Gases, AAPG, December 1983.





1; 2/12-1 DST 1 2: 2/12-1 DST 2

# Figure 1a

Variations of molecular composition in natural gases related to the isotope variations of methane.

# Figure 1b

Carbon and hydrogen isotope variations in methanes.

# Figure 1c

Carbon isotope variations in ethane related to carbon isotope variations in methane.

The principle for the genetic characterization of natural gases is that the primary gases (B-biogenic gas, T-associated gas, TT-non-associated gas) are defined by fields of compositional variations. These primary gases may become mixed and form various mixtures "M" of intermediate composition. "TT(m)" and "TT(h)" are non-associated gases from marine source rocks and coal gases from N.W. Germany, respectively, compositional shifts due to migration are indicated by arrows Md (deep migration) and Ms (shallow migration), respectively. "T<sub>0</sub>" are gases associated with petroleum in an initial phase of formation. "T<sub>1</sub>" are gases associated with condensates. (Schoell 1983).