

S. N. E. A. (P)  
DIRECTION EXPLORATION  
DIVISION RECHERCHES et APPLICATIONS en GEOLOGIE  
LABORATOIRE  
ETABLISSEMENT DE PAU

DEX/RAG-Lab.Pau 84/124RP  
/fm

BA 84-6059-1

- 4 JUNI 1984 / b

725.3

15/3-3.

ISOTOPIC STUDY OF CO<sub>2</sub> FROM  
15/3-1, 15/3-3 AND 15/3-4 WELLS  
(offshore Norway)

DEX/RAG/LAB/PAU 84/124RP

M. THOMAS  
May 1984

Work requested by J.P. VERNET

Work order n° 103.100.372

P.J. : 1 table - 1 figure

DIFFUSION LIST :

ELF AQUITAINE NORGE

s/c Conseiller Exploration DIG EUROPE - Paris (12ex.)

Conseiller Exploration DIG EUROPE - Paris

Division Programmes - Paris

Expert régional - Paris

Division Etudes et Synthèses - Paris

Division Orientation Zones Nouvelles - Paris

Division Orientation Zones Nouvelles - Pau

SID MICOULAU (2ex.)

## ABSTRACT

This study deals with the isotopic study of the gases from the 15/3-1, 15/3-3 and 15/3-4 wells from the Norway offshore (see note GEO/LAB/PAU 1003-81) ; the aim of this study being to determine the origin of the low concentrations of CO<sub>2</sub> present in the Bathonian to Oxfordian reservoirs.

Although some hypothesis have to be done before the interpretation, the minor CO<sub>2</sub> gas concentrations trapped in the reservoirs appear to be due to a thermal decomposition of the organic matter, in relation with the hydrocarbon generation. The temperatures to which CO<sub>2</sub> has been generated vary between 115 to 150°C for the 15/3-3 and 15/3-4 samples, and are close to 90°C for the 15/3-1 sample in which the CH<sub>4</sub> generation seems to correspond to a maturation level close to the beginning of the oil window.

6 pages  
1 table  
1 figure



## I - ANALYTICAL RESULTS

Different analytical data have been taken into account in this study, that is to know :

- molar compositions of essentially the  $\text{CH}_4$ ,  $\text{C}_2\text{H}_6$ ,  $\text{C}_3\text{H}_8$  and  $\text{CO}_2$  compounds,
- isotopic compositions of the  $\text{CH}_4$ ,  $\text{C}_2\text{H}_6$  and  $\text{CO}_2$  gases,
- and mean carbon isotopic ratio of the calcitic cementations present in the reservoirs.

Most of these data was already presented in a previous note from SOURISSE and BARADAT (GEO/LAB/PAU 1003/81NI)). The complete synthetical data are reported on the attached table.

It is here to be noted that the 15/3-1 sample in fact corresponds to a mixing of different FIT performed between 4.089 and 4.443m. So, this sample is not very representative and its results should be considered with caution.

## II - COMMENTS OF THE RESULTS

### Origin of the gaseous hydrocarbons :

The following interpretation comes from SOURISSE and BARADAT's report and from the comparison of their data from the 15/3-1 and 15/3-3 samples with those from the 15/3-4 one.

According to these authors, the  $\text{CH}_4$  would correspond to a gas associated with an oil generation in 15/3-3 and to a gas originating from the beginning of the oil window in 15/3-1. The 15/3-4 samples appear as intermediate, but closer to the 15/3-1 gas than to the 15/3-3 one.

On another hand,  $\text{CH}_4$  and  $\text{C}_2\text{H}_6$  appear in 15/3-3 and, but to a lesser extent, 15/3-1 as cogenetic compounds. Such a result also seems to be valuable for the 15/3-4 samples.



These results may be compared to those of the organic matter study of the 15/3-3 well (note GEO/LAB/BSS 0/1857 from CAILLEAUX and ROBERT), for which the DST 2bis condensate has been generated by Callovian/Bathonian source-rocks with degrees of catagenesis corresponding to those of the syngenetical hydrocarbons about 200m below the reservoirs.

Origin of the CO<sub>2</sub> :

The origin of the CO<sub>2</sub> may be only discussed if one considers the isotopic relations which exist between the CO<sub>2</sub> gas and the other carbon-rich compounds, that is to know the gaseous hydrocarbons and the in-situ carbonates. A good thing would be also to consider the isotopic compositions as well of the liquid hydrocarbons as of the kerogen.

For such an isotopic approach, some hypothesis may be done : the most important one being to consider that no late isotopic reequilibration takes place in the reservoirs for that concerns the CO<sub>2</sub> and CH<sub>4</sub> gases. Such an hypothesis reveals correct for the CH<sub>4</sub>, for which the isotopic ratio really denotes its degree of maturity according to the nature of the organic matter. For the CO<sub>2</sub> and taking into account the more important sensibility of the CH<sub>4</sub> isotopic ratios (variations of 80 ‰ ; see figure) to the temperature than the one of the CO<sub>2</sub> (variations of 25 ‰ ), this same hypothesis may be advanced.

So, if one considers the CO<sub>2</sub>-CH<sub>4</sub> system, it is then possible to calculate the temperature for which both compounds were isotopically equilibrated. Such an approach may also be done for the CO<sub>2</sub> gas-CO<sub>2</sub> carbonate system. These results are reported on the attached table. For the 15/3-3 and 15/3-4 samples, the temperatures obtained in the CO<sub>2</sub>-CH<sub>4</sub> system are comprised between 115° and 150°C, that is to say identical to those of the present reservoirs but also close to those required for the hydrocarbon generation in the oil window maturation level. The 15/3-1 sample provides an isotopic equilibration temperature of 90°C. Such a result may also correspond to the temperature which prevailed during the CO<sub>2</sub>-CH<sub>4</sub>

cogenesis if one considers that the  $\text{CH}_4$  comes from a maturation state corresponding to the beginning of the oil window. However, the results from the 15/3-1 sample may be considered with caution due to the mixing of different FIT.

In the  $\text{CO}_2$  gas- $\text{CO}_2$  carbonate system, the temperatures obtained for the 15/3-1 and 15/3-3 samples are too high to imagine these both compounds as being isotopically equilibrated. Only, the FIT 2bis, and to a lesser extent the FIT 1, from 15/3-4 present isotopic ratios which might indicate that the two compounds concerned are isotopically equilibrated in the reservoir conditions ; that is to say  $\text{CO}_2$  may be possibly regarded as originating from a thermal decomposition of the carbonates. This assumption remains however uncertain according to the low carbonate concentrations.

At last, a deep (volcanic or magmatic) origin may not be envisaged for the  $\text{CO}_2$  according to the isotopic equilibrium temperatures deduced from the  $\text{CO}_2$ - $\text{CH}_4$  system which well denote the cogenesis of these both gases.

### III - CONCLUSIONS

In spite of some discrepancies in the results, notably for the 15/3-1 sample, the  $\text{CO}_2$  analysed in the samples studied appears to be due to a thermal decomposition of the organic matter in relation with the hydrocarbon generation. The main data which allow us to arrive to this conclusion are :

- the quite identical molar composition of the total gases of all the samples studied with almost similar low  $\text{CO}_2$  concentrations (the 15/3-1 sample excepted) ;
- the good agreement of the temperature obtained for the  $\text{CO}_2$ - $\text{CH}_4$  isotopic equilibria with those required for the hydrocarbon generation taking into account that  $\text{CH}_4$  rather comes from an oil window maturation state close to those of the actual reservoir ;



- the similarity between these latter temperatures and those of the present reservoirs which are also close to the source-rock horizons ;
- the impossibility of a deep origin ;
- and the possible but only minor incidence (only in 15/3-4) of a CO<sub>2</sub> generation from the thermal decomposition of the carbonates.