

# Dynamic correlations between susceptibility gradients and $T_2$ -relaxation as a probe for wettability properties of liquid saturated rock cores

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## Abstract

We explore the use of correlations between susceptibility gradients,  $G_0$ , and  $T_2$ -relaxation ( $G_0$ - $T_2$ ), and show how the difference in response for oil and water with respect to  $G_0$  can be used for improved characterization of wettability of the internal surface in porous rock cores.

## Keywords

NMR, diffusion, susceptibility gradients, relaxation, wettability.

## 1. Introduction

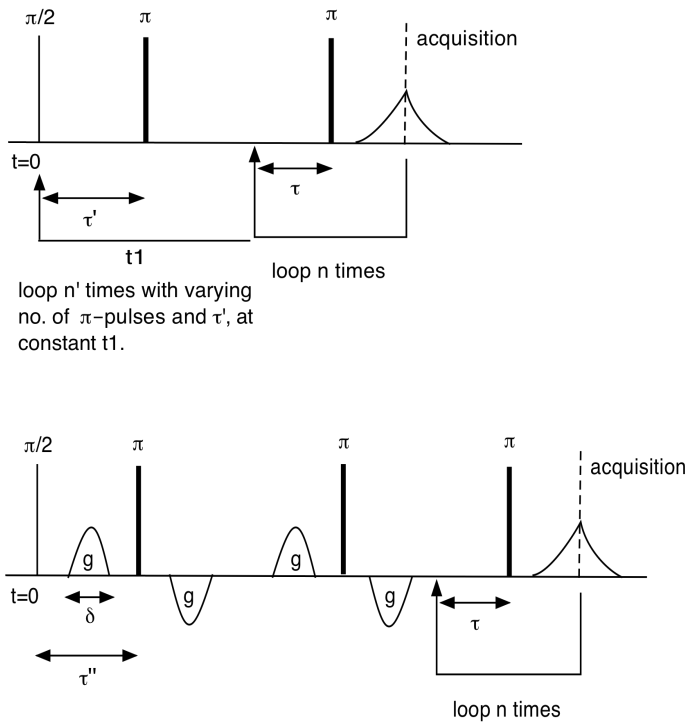
The oil/water wettability is an important property of rock core samples from oil reservoirs. The dynamic behavior (relaxation and diffusion) of the NMR signals from liquids confined in such rock core samples is sensitive to surface interactions and can potentially be used for characterization of wettability properties [1]. The  $T_2$ -distributions of water and oil do however often overlap. The use of diffusion- $T_2$  ( $D$ - $T_2$ ) correlation measurements can be used for a separation of the oil and water signals, increasing the resolution of the measurement [2]. We however explore the use of correlations between susceptibility gradients and  $T_2$ -relaxation ( $G_0$ - $T_2$ ) [3], and show how the difference in response for oil and water with respect to  $G_0$  can be used for improved characterization of wettability [4]. We compare the use of  $G_0$ - $T_2$  correlations to regular  $T_2$  measurements, and to measurements of  $D$ - $T_2$  correlations.

## 2. Materials and Methods

All measurements were performed at 35°C on a Maran DRX 12 MHz spectrometer (Resonance Instruments), with a gradient probe providing a maximum gradient strength of 225 Gauss/cm. The sample studied was a water-wet sandstone Berea rock core (1.5").  $T_2$ ,  $G_0$ - $T_2$ , and  $D$ - $T_2$  measurements were compared in water-saturated ( $S_w$ ) and irreducible water saturated ( $S_{wi}$ ) states. To reach the  $S_{wi}$  state synthetic oil (Marcol82, Exxon Mobile) was flooded through the rock core.

$T_2$  was measured using the regular Carr-Purcell-Meiboom-Gill (CPMG) pulse sequence. For measurement of  $G_0$ - $T_2$  and  $D$ - $T_2$  correlations the sequences given in Fig. 1 were used. Notice that  $G_0$  is never measured directly, but through  $DG_0^2$ . The value of  $\tau'$  was varied (25 steps), between 0.1 and 10 ms and the gradient strength,  $g$ , was varied (25 steps) between 0 and 200 Gauss/cm. The other parameter values were:  $\tau = 0.1$  ms,  $tI = 20$  ms,  $\delta = 3$  ms,  $\tau'' = 3.9$  ms,  $n = 12000$ .

The CPMG data were analyzed using (1D) Inverse Laplace Transformations (ILT). The  $D$ - $T_2$  data were analyzed using 2D-ILT [2], while for the  $G_0$ - $T_2$  data we compared the use of 2D-ILT with a semi-discrete analysis [5] where the  $T_2$ -decays for oil and water were separated based on the difference in response to diffusion in susceptibility gradients. The outline of the semi-discrete analysis is as follows: At a certain  $\tau'$ -value ( $\tau'_c$ ), the faster diffusing signal from water has dephased. For each echo in the  $T_2$ -decays at  $\tau' > \tau'_c$  a linear fit with respect to varying  $\tau'$  can be performed, and can then be extrapolated to find synthetic  $T_2$ -decays for oil at  $\tau' < \tau'_c$ . The synthetic  $T_2$ -decays can be subtracted from the original  $T_2$ -decays to obtain synthetic  $T_2$ -decays for water at varying  $\tau'$ . 1D-ILT can be performed on the synthetic oil and water  $T_2$ -decays, and the obtained signal intensity for each  $T_2$  component vs  $\tau'$  can be used to determine  $DG_0^2$ . For more details we refer to the similar description given in ref. [5].



**Fig. 1:** Pulse sequences used to measure  $G_0$ - $T_2$  [3] (upper figure) and  $D$ - $T_2$  correlation [5] (lower figure).

The echo attenuation for the  $G_0$ - $T_2$  correlation can be expressed using a continuous function (for 2D-ILT analysis) or as discrete functions (for the semi-discrete analysis):

$$M(\tau', 2n\tau) = \iint f(DG_0^2, T_2) e^{-\frac{t_1 + 2n\tau}{T_2}} e^{-\frac{\gamma^2 DG_0^2 \tau'^2 2n\tau}{3}} dT_2 d(DG_0^2) \approx \sum_{i=1}^n \sum_k q_{ik} e^{-\frac{t_1 + 2n\tau}{T_{2ik}}} e^{-\frac{\gamma^2 D_{ik} G_{0ik}^2 \tau'^2 2n\tau}{3}} \quad (1)$$

The index  $i$  indicate a sum over liquids with different dynamic behavior, while index  $k$  indicate a sum over all possible discrete values for  $D$ ,  $G_0$  and  $T_2$ .

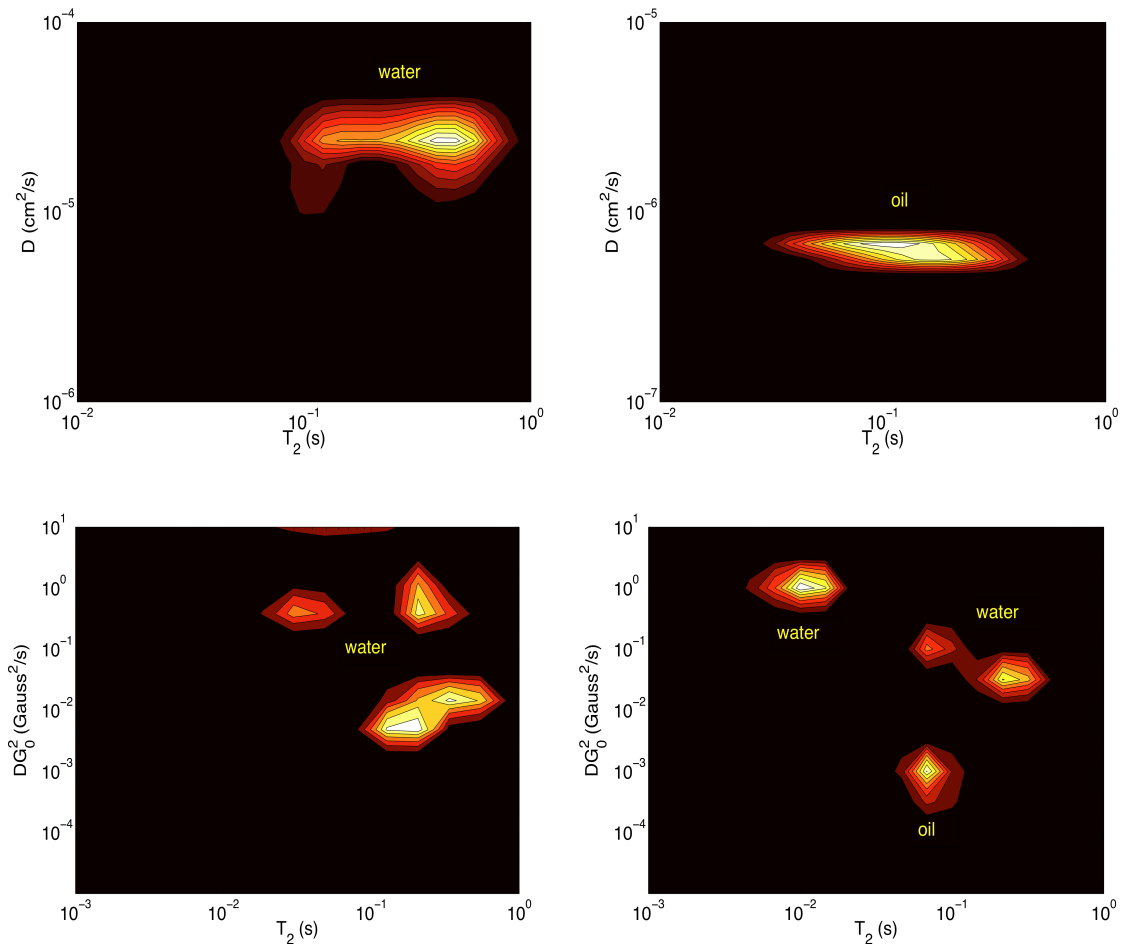
Correspondingly, for the  $D$ - $T_2$  correlation we have:

$$\begin{aligned}
M(g, 2n\tau) &= \iint f(D, T_2) e^{-\frac{4\tau' + 2n\tau}{T_2}} e^{-\left(\frac{16}{\pi^2}\right) \gamma^2 D g^2 \delta^2 \left(\frac{3\tau'}{2} - \frac{\delta}{8}\right)} dT_2 dD \times \int e^{-\frac{\gamma^2 D G_0^2 \tau'^2 4\tau'}{3}} dV \\
&\approx \sum_{i=1}^n \sum_k p_{ik} e^{-\frac{4\tau' + 2n\tau}{T_{2ik}}} e^{-\left(\frac{16}{\pi^2}\right) \gamma^2 D_{ik} g^2 \delta^2 \left(\frac{3\tau'}{2} - \frac{\delta}{8}\right)} \times \int e^{-\frac{\gamma^2 D_{ik} G_{0ik}^2 \tau'^2 4\tau'}{3}} dV
\end{aligned} \quad (2)$$

Notice the last attenuation term in Eq. (2) due to diffusion in susceptibility gradients during the measurement, which is integrated over the total volume of the sample.

### 3. Results and Discussions

The  $T_2$ -values for bulk water and oil were 3000 and 136 ms respectively, while the diffusion coefficient for water at 35°C was  $2.9 \cdot 10^{-5} \text{ cm}^2/\text{s}$ . The CPMG measurements (data not shown) resulted in a broad distribution of  $T_2$ -values in the  $S_w$  state. In the  $S_{wi}$  state  $T_2$ -signals from water and oil were strongly overlapping.

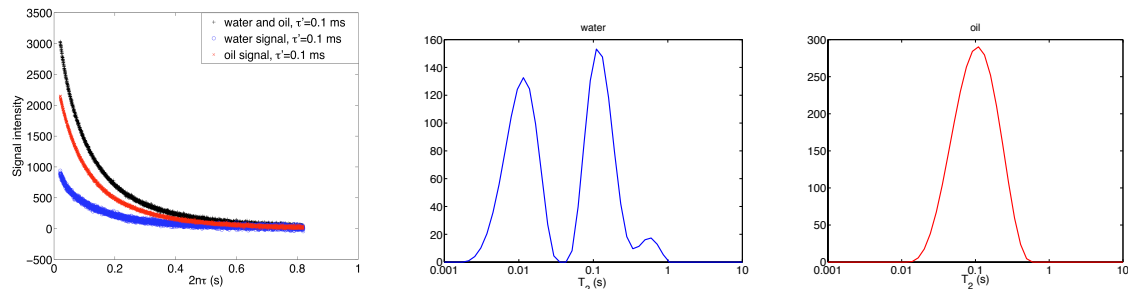


**Fig. 2:**  $D-T_2$  correlation maps (top row), and  $DG_0^2-T_2$  correlation maps (lower row) in  $S_w$  (left column) and  $S_{wi}$  (right column) states. Assignments of the signal originating from water and oil are given.

The  $D-T_2$  maps (Fig. 2, top row) show a narrow distribution of diffusivities for water centered around  $2.3 \cdot 10^{-5} \text{ cm}^2/\text{s}$  in the  $S_w$  state.  $\tau''$  in the PGSE-part was 3.9 ms, resulting in a strong attenuation due to diffusion in susceptibility gradients, so signals with  $T_2 < 100 \text{ ms}$  were therefore lost. Due to the same effect, and since oil displaces bulk water; all signals from water in the  $S_{wi}$  state were lost. The mean diffusivity for the remaining oil was  $6 \cdot 10^{-7} \text{ cm}^2/\text{s}$ . From the  $DG_0^2-T_2$  maps (Fig. 2, lower row), and using the measured diffusion coefficient of

water, the values for  $G_0$  in the  $S_w$  state are in the range 10-200 Gauss/cm. In the  $S_{wi}$  state, the signals from oil and water are separated along the  $DG_0^2$ -dimension. Clearly, oil displaces the bulk water. When normalized with respect to the measured diffusion coefficients,  $G_0$  for the water peaks are 50 and 200, and for the oil peak 30 Gauss/cm.

The dynamic range along the  $DG_0^2$  dimension is larger than what is obtained along the corresponding  $D$ - or  $T_2$ -dimensions, making it more sensitive to surface effects and changes in wettability.



**Fig. 3:** Results obtained from the semi-discrete analysis of the  $G_0$ - $T_2$  data in the  $S_{wi}$  state. Synthetic  $T_2$ -decays for oil and water are plotted together with the  $T_2$ -decay for the total signal (left figure). The other two figures show the resulting  $T_2$ -distributions for water (middle figure) and oil (right figure) obtained using 1D-ILT. All plots were obtained at the minimum  $\tau^2$ -value (0.1 ms).

From the  $DG_0^2$ - $T_2$  correlation map the relative amount of water in  $S_{wi}$  state is 0.6, which is higher than expected. Using the semi-discrete analysis described above, the fraction of water was determined to be 0.29, which is more typical for the  $S_{wi}$  state. The positions of the  $T_2$ -peaks (Fig. 3, middle and right) correspond reasonably well with results obtained in the  $DG_0^2$ - $T_2$  correlation maps. Obtained results from the semi-discrete analysis are summarized in the table below. The values for  $G_0$  of water signals are somehow higher than the values obtained using 2D-ILT, but the overall correspondence is good.

**Table 1:** Values for  $T_2$  and  $G_0$  obtained from the semi-discrete analysis of the  $G_0$ - $T_2$  data in the  $S_{wi}$  state.

	$T_2(1)$ (ms)	$T_2(2)$ (ms)	$G_0(1)$ (Gauss/cm)	$G_0(2)$ (Gauss/cm)
water	30	160	500	80
oil	110	-----	30	-----

## 4. Conclusions

We have shown that  $G_0$ - $T_2$  correlations are potentially a more sensitive wettability probe for liquid saturated rock cores compared to  $D$ - $T_2$  correlations. The separate responses of oil and water to diffusion decay in susceptibility gradients can be used for a separation of the  $T_2$ -decays when they are overlapping in the  $T_2$ -dimension. A semi-discrete analysis of the  $G_0$ - $T_2$  data gave quantitatively more reliable results compared to a 2D-ILT analysis.

## References

- [1] K.-J. Dunn, D.J. Bergman, G.A. Latorraca, *Nuclear Magnetic Resonance. Petrophysical and Logging Applications*, Pergamon, New York (2002).
- [2] M. D. Hurlimann, L. Venkataramanan, *J. Magn. Reson.*, **157**, 31 (2002).
- [3] B. Sun, K.-J., Dunn, *Phys. Rev. E.*, **65**, 051309 (2002).
- [4] J.G. Seland, K.E. Washburn, H.W. Anthonsen, J. Krane, *Phys. Rev. E.*, **70**, 051305 (2004).
- [5] J.G. Seland, G.H. Sørland, H.W. Anthonsen, J. Krane, *Applied Magn. Reson.*, **24**, 41 (2003).