# NMR T<sub>2</sub> CUT OFF: WHICH ONE IS TO BE USED FOR APPLICATION?

by

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#### ABSTRACT

Recent developments in petroleum industry have been witnessing the surge of the use of nuclear magnetic resonance (NMR) log. Despite some remaining problems the NMR technology appears to gain more acceptance as petrophysical tool for evaluating reservoir quality. Comprehensive formation evaluation requires determination of irreducible fluids, movable fluids, and permeability. However, rock heterogeneity introduces complexity in any formation evaluation activities. This can also cause problem for NMR log interpretation. In the presence of clays the most commonly used  $T_2$  cut off values, a constant value throughout a formation, seem to eventually yield inaccurate irreducible water saturation estimates, as well as other output such as permeability. This study focuses at finding a solution for finding the best way of choosing the most representative  $T_2$  cut off value to be used in NMR log interpretation. This is indeed a common pressing problem for heterogeneous formation rocks such as in the case of Tirrawarra sandstones used in this study. The main part of the study is devoted to comparison between the use of single averaged  $T_{2c}$  value and establishment of empirical correlations enabling the provision of  $T_{2c}$  for any level of heterogeneity (i.e. various levels of shaliness). The study however surprisingly shows that, in spite of the theoretical soundness of the empirical correlations established, simple averaging of  $T_{2c}$  values yielded by a reliable method proves itself adequate. This conclusion therefore helps considerably in reducing complexity in NMR log interpretation.

Key words:  $T_{2c}$ , irreducible water saturation, ductile components, empirical correlation, averaged  $T_{2c}$ .

#### I. INTRODUCTION

Nuclear magnetic resonance (NMR) log is at present gaining momentum in its use in petroleum industry. Despite the ever remaining controvercy over its consistencies in measurement accuracy, the method is fast achieving recognition as a petrophysical tool for evaluating various reservoir properties including volumes of movable and irreducible fluids, permeability, and effective porosity. Such properties like irreducible water saturation are acknowledged for its importance in the estimation of hydrocarbon in place and reserves and reservoir productivity.

Earlier studies on the applications of NMR technology in assessing permeability, effective porosity, and irreducible fluid in heterogeneous formations have been carried out by some studies in the past (e.g. Kenyon, 1989; Straley *et al.*, 1991; Kenyon, 1992; Prammer *et al.*, 1996; Coates *et al.*, Coates *et al.*, 1995; 1994; Coates *et al.*, 1997; Oraby *et al.*, 1997; Hassoun *et al.*, 1997; Hodgkins & Howard, 1999). Results of these investigations have indicated that for some particular sandstone samples with significant amount of clay content, a detailed knowledge and understanding of reservoir properties including lithology and mineral composition are essential to apply and to interpret NMR measurement data.

In order to divide effective porosity into movable and irreducible fluid saturations, NMR measurements require a  $T_2$  cutoff, or often expressed as  $T_{2c..}$  In traditional  $T_2$  interpretation, the  $T_{2c}$  is usually taken as a constant value throughout a formation. However, this proved to be impracticable, and in many cases resulting in inaccurate permeability and irreducible water saturation values ( $S_{wirr}$ ). Since this inaccuracy is suspected to be due to the sandstone's nature of heterogeneity in lithology and mineral composition, it was therefore thought that a different approach should be adopted. Musu (2000) and Musu & Widarsono (2007) suggested that for rocks such as shaly sandstones as many as possible  $T_{2c}$  should be taken to accommodate the high heterogeneity level. Accordingly, different  $T_{2c}$  was measured for each of the sample used in the study using which – naturally – better permeability and irreducible water saturation ( $S_{wirr}$ ) were resulted.

In Musu (2000) and Musu & Widarsono (2007) it has been shown that the BVI Error method, a method that is basically an iterative approach, produced the most representative  $T_{2c}$  for individual sample. However, the method does not offer a solution that suggests the choice of  $T_{2c}$  to be applied in practical NMR log interpretation. It is therefore the purpose of this study to find the best way to assign the best  $T_{2c}$ .

## II. DETERMINATION OF T, CUT OFF (T<sub>2</sub>)

The comprehensive study presented in Musu (2000) was carried out on a set of Tirrawarra shaly sandstone samples taken from Cooper Basin, South Australia. In reaching its objectives the study used results from petrographic, NMR, routine core analysis, special core analysis, and mercury intrusion capillary pressure (MICP) data for the sandstone samples.

NMR measurements refer to the response of hydrogen nuclei (protons) to a magnetic field. Hydrogen nuclei are abundant in both water and hydrocarbons stored in formation. NMR tools basically measure the decaying signals of the nuclear spin of protons in the water and hydrocarbons at various times. The signal decaying time required for the protons to complete de-phasing is termed  $T_2$ . The NMR measurements of amplitude versus time are used to calculate the distribution of porosity components ( $T_2$  distribution) as a function of their  $T_2$  times.

 $T_2$  cut off, or  $T_{2c}$ , is the maximum  $T_2$  for part of porosity that is occupied by immovable fluids. It is used to distinguish movable fluid (FFI) from the irreducible or non-movable fluid (BVI). In the  $T_2$  distri-

bution, BVI is all the amplitudes in the spectrum having  $T_2$  values less than the  $T_{2c}$ . In other words,  $T_{2c}$  is the sum of all porosities whose  $T_2$  is less than the  $T_{2c}$ and consequently FFI is the sum of amplitudes with  $T_2$  values greater than  $T_{2c}$ , or the sum of all porosities whose  $T_2$  is greater than  $T_{2c}$ . Therefore, it is obvious that correct determination of the  $T_{2c}$  is very important for estimation irreducible water saturation. The following is brief discussion over the approaches used for  $T_{2c}$  determination. Detailed description of the methods is presented in Musu (2000) and Musu & Widarsono (2007).

# A. $T_{2c}$ by default

For  $T_{2c}$  application in interpretation and analysis, two common  $T_{2c}$  values are often used for sandstone formations. Those  $T_{2c}$  values of 33 ms (Morris *et al.*, 1993; Coates *et al.*, 1999) and 22.6 ms (Numar-Halliburton standard for sandstone) are established based on combination of field experience and extensive laboratory data.

# **B.** $T_{2c}$ direct from spectrometer

 $T_{2c}$  values are usually calibrated in the laboratory using plugs obtained from conventional cores. Measurements are performed by matching the integrated NMR amplitudes up to  $T_{2c}$  in the  $T_2$  spectrum measured for a water saturated-core sample to the volume of water remaining in the core after de-saturating to a desired capillary pressure (Straley *et al.*, 1994).

Under certain formation conditions an average value of core-measured, single or fixed  $T_{2c}$  method is inappropriate. For the Tirrawarra Sandstone sample set used in this study, presence of significant amount of ductile clay and quartz cementation has caused the development of micro-porosity with macro-pores tend to be isolated leading to low permeability.

The average standard  $T_{2c}$  computed from the NMR spectrometer for the samples provided is 42 ms, which corresponds to values of individual samples that range from 18.56 ms to 57.15 ms. This variance indicates that a single  $T_{2c}$  method would not properly represent BVI, which in turn would give incorrect permeability and  $S_{wirr}$ .

# C. $T_{2c}$ using SBVI method

The spectral bulk volume irreducible (SBVI) method is based on the T<sub>2</sub> distribution and has uti-

lized core-measured values of BVI to determine the percentage of amplitude (porosity) associated with each individual T<sub>2</sub> which will remain as capillarybound water (Coates et al., 1997; Santos-Numar, Cooper core sample study, 1999; Oraby et al., 1997). SBVI varies  $T_{2c}$  by measuring the contribution of the irreducible fluid to the overall fluids in each grouping or bin of the T<sub>2</sub> distribution. These measured values are used to calculate the irreducible water,  $S_{wirr}$  in each bin. The  $S_{wirr}$  values of all the bins are then fitted against their  $T_2$  values through the use of a function of

$$S_{wi} = \frac{1}{mT_{2GM} + b} \tag{1}$$

where m is fitting parameter, b is y-intercept, and  $T_{2GM}$  is the geometric mean of the  $T_2$  values (Coates et al., 1997) of the regression line shown in Figure 2. The variable m was determined as the slope of the line relating  $1/S_{wi}$  (core derived) to the geometric mean  $T_2$  ( $T_{2GM}$ ), which gives m of 0.013 for b taken as equal to 1 (sandstone) (Coates et al., 1997). SBVI is then derived by multiplying  $\mathbf{S}_{\text{wirr}}$  and NMR porosity (MPHI). Geometric mean  $T_2^{(m)}(T_{2GM})$  values were obtained from fully brine saturated samples. (Figure 1 shows a very poor fit suggesting that this method



may not be valid in this particular sample set known for their significant amount of ductile and clay components.)

#### **D.** T<sub>2c</sub> using BVI Error method

To ensure the acquisition of representative BVI values, a 'BVI error' method was applied. In the method,  $T_{\gamma_c}$  was estimated through comparing corederived irreducible water volume  $(V_{wirr})$  to NMR cumulative porosity. The core-derived  $V_{wirr}$  itself is actually a multiplication product of core irreducible water saturation  $(S_{wirr})$  and core porosity. It is assumed that this measured irreducible water saturation is a true measure of BVI against which the BVI derived from the NMR can be compared to.  $T_{2c}$  values considered representative are obtained when the BVI values from the NMR data are in the least disagreement or least in error with the corresponding  $V_{wirr}$  values. To observe the  $T_{2c}$ , the absolute difference between the NMR cumulative porosity data and core data is plotted against  $T_2$  on the NMR tabular sheets (for the brine-saturated measurements). In such a plot  $T_{2c}$  for each sample is characterized by a minimum sharp plunge of the cumulative curves (Figure 2). See Musu (2000) for complete discussion regarding procedure.

1000.0

10000.0



#### III. MEASURED NMR RELAXATION TIME (T<sub>2</sub>) DISTRIBUTIONS

NMR relaxation time distributions for fully brinesaturated samples are broad, ranging between less than one and more than 1,000 milliseconds (Figure 3). The distributions commonly encountered are of skewed unimodal type, which usually does not clearly differentiate between inter-granular porosity and micro-porosity. However, unimodal distribution shown by the relaxation time distribution might be interpreted as a combination of two or more sub-populations (Swan & Sandilands, 1995) interpreted as micro-porosity and macro-porosity.

In NMR  $T_2$  measurement, partial saturations were obtained by centrifuging the water-saturated samples. The irreducible water or non-centrifugeable water includes clay bound water and capillary bound water. NMR relaxation time distributions for partially saturated samples are also broad, ranging from zero to nearly 1,000 milliseconds (Figure 3). The distributions commonly encountered are trimodal, which might be associated with small and larger pores. A trimodal distribution with a broad range of  $T_2$  distribution is usually taken as an indication that bound fluid is not only associated with small pores but also with larger pores (FFI). The trimodal curves show that the water is present not only in the small pores but also in the larger pores.

#### IV. DETERMINATION OF IMMOVABLE WATER (BVI)

As mentioned above, movable and immovable water were determined using the  $T_{2c}$  on  $T_2$  relax-

Sample	T <sub>2c</sub> (ms)	MPHI (%)	BVI (%)	FFI (%)	Swirr (%)	Swirr (core) (%)
JM-11	22.6	14.601	2.986	11.615	20.451	64.2
JM-12	22.6	14.527	2.775	11.752	19.102	71.9
JM-26	22.6	13.87	5.213	8.657	37.585	48.7
JM-42	22.6	12.371	2.795	9.576	22.593	59.5
JM-50	22.6	9.459	4.23	5.229	44.719	74
JM-53	22.6	11.75	1.924	9.826	16.374	43

ation curves. In order to correlate BVI with the core derived irreducible water saturation ( $S_{wirr}$ ), the BVIs were then calculated as percentages. The BVI as a percentage is then termed as NMR  $S_{wirr}$ . As reported in Musu (2000),  $S_{wirr}$  values derived using a single and predetermined  $T_{2c}$  values show very poor agreements with the core-derived  $S_{wirr}$  values (Table 1). On the contrary,  $S_{wirr}$  values calculated using representative  $T_{2c}$  values produced through the use of BVI error method and the  $T_{2c}$  measured directly from the NMR machine tend to show very good agreements with the core measurements (Table 2).

The varied  $T_{2c}$  determined by the SBVI method, however, gives weaker correlation between NMR  $S_{wirr}$  and core  $S_{wirr}$ . SBVI results demonstrate inappropriate BVI values. The  $T_{2c}$  determined by SBVI is based on mean geometric  $T_{2c}$ , measured directly from the spectrometer and do not include the effect of reservoir heterogeneity (Oraby et al., 1997). As the representative  $T_{2c}$  values for the samples are known  $S_{wirr}$  can then be estimated reliably.

The comparison between NMR S<sub>wirr</sub> and core S<sub>wirr</sub> values can also be presented graphicaly. Figures 4 through 11 present the plots, respectively, for T<sub>2c</sub> = 22.6 ms (Numar-Halliburton), T<sub>2c</sub> = 33 ms (Morris et al., 1993), varied T<sub>2c</sub> values from spectrometer, average value from spectrometer of T<sub>2c</sub> = 42 ms, varied T<sub>2c</sub> values from SBVI method, average value from SBVI method of T<sub>2c</sub> = 56.6 ms, varied T<sub>2c</sub> values from BVI Error method, and average value from BVI Error method of T<sub>2c</sub> = 66.5 ms.

Table 2Results of irreducible water saturation (Swirr)using multiple representative T20 valuesfrom BVI Error method. Notice the goodagreement between estimated Swirrobserved (core) Swirr (from Musu, 2000)							
Sample	T <sub>2c</sub> (ms)	MPHI (%)	BVI (%)	FFI (%)	Swirr (%)	Swirr (core) (%)	
JM-11	22.6	14.601	8.595	6.006	58.866	64.2	
JM-12	22.6	14.527	9.412	5.115	64.79	71.9	
JM-26	22.6	13.87	6 4 6 6	7 4 0 4	46 619	487	

6.588

7.189

4.766

5.813

2.27

6.984 40.562

53.011

76.002

JM-42

JM-50

JM-53

22.6

22.6

22.6

12.371

9.459

11.75

59.5

74

43

From the plots it can be clearly seen that results from BVI Error method are the best indicated by the very small disagreement when the results are compared to core data (Figure 10). The two predetermined  $T_{2c}$  of 22.6 ms and 33.3 ms appear to have yielded largely inaccurate results. This underlines the urgency of laboratory measured  $T_{2c}$  value for most of cases especially the heterogeneous ones such as Tirrawarra sandstones. The representation of averaged values also seem to have not improved the results either even though their results can still be considered as better than results from the two predetermined  $T_{2c}$  values.

## V. WHICH T<sub>2C</sub> SHOULD BE ADOPTED?

As shown by the plots presented on Figures 4 through 11, the BVI Error method appears to be yielding the best  $S_{wirr}$  estimates if the  $S_{wirr}$  values from core are to taken as reference. However, a question remains over the representative value to be used for interpreting NMR log data. There are some options that can be taken to deal with this question; averaging the available  $T_{2c}$  values, establishing a kind of mathematical function that represents the whole range of  $T_{2c}$  values, choosing one of the  $T_{2c}$  values available, and picking a  $T_{2c}$  value within the range of  $T_{2c}$  values. Among the options, the first option has been performed and the result is presented in Figure 11. The remaining three options are to be tested here.

The effort to establish a mathematical function is started from a consideration about the relevant factor that is linked to  $T_{2c}$ . It was initially thought that it should be porosity and permeability due to their significant influence on  $S_{wirr}$ . However, although this may theoretically and empirically be correct this can be considered as impractical since porosity and permeability are also the output of NMR log. Moreover, wells that are NMR-logged may not have porosity and permeability data from other sources. Another factor that could be considered as important is the major source of heterogeneity itself in the case of Tirrawarra sandstones, presence of ductile components.

The ductile components in the Tirrawarra sandstones are defined as composed by rock fragments, matrix (pore filling fine particles, and clay minerals. These materials cause ductility behaviour in the rocks and serve as the major factor in the creation of isolated pores hence changing effective porosity, per-



Typical NMR relaxation time, as also exhibited by the Tirawarra sandstones used in the study (from Musu, 2000)



meability, and irreducible water saturation. Comprehensive disscussion regarding this matter is presented in Musu & Widarsono (2007). Based on this evidence the  $T_{2c}$  is to be made as a function of contents of ductile components.

Table 3 presents the ductile components for the six core samples used in the study along with the  $T_{2c}$  values from the BVI Error method. Early effort to plot ductile components versus  $T_{2c}$  values proves to be too scattered to yield any reliable correlation. Therefore, a sort of compromise has to be reached. In this light,  $T_{2c}$  values for JM-11 and JM-12 samples appear to be 'too high' and need adjustment. With





Figure 6 Comparison between core S<sub>wirr</sub>.and NMR S<sub>wirr</sub> estimated using varied T<sub>2c</sub> values determined directly using NMR spectrometer















Figure 11 Comparison between core  $S_{wirr}$  and NMR  $S_{wirr}$  estimated using average value of  $T_{2c}$  = 66.5 ms from values determined using BVI Error method



Figure 12 Logarithmic correlation between  $T_{2c}$ (compromised) and ductile component



Figure 13 Linear correlation between T<sub>2c</sub> (compromised) and ductile component



Comparison between core  $S_{wirr}$  and NMR  $S_{wirr}$  estimated using varied but compromised  $T_{2c}$  values determined using BVI Error method (logarithmic correlation)



Figure 15 Comparison between core  $S_{wirr}$  and NMR  $S_{wirr}$  estimated using varied but compromised  $T_{2c}$  values determined using BVI Error method (linear correlation)



Comparison between core  $S_{wirr}$  and NMR  $S_{wirr}$  estimated using average  $T_{2c} = 53.67$  ms from compromised values that are modified from values yielded by BVI Error method





Table 3Ductile components, T2c from BVI Errormethod, and compromised T2c values(from Musu, 2000)							
Sample	Ductile component s (%)	T <sub>2c</sub> (ms)	T <sub>2c</sub> (compromised) (ms)				
JM-11	11	75	50				
JM-12	11	90	52				

32

70

62

70

32

70

60

55

5.4

23.3

12.2

7.17

JM-26

JM-42

JM-50

JM-53



Figure 20 Comparison between core S<sub>wirr</sub>.and NMR S<sub>wirr</sub> estimated using  $T_{2c}$  = 75 ms from BVI Error method









taking into consideration the possible error in measurement and presence of anomalous data the  $T_{2c}$  data was modified to yield the  $T_{2c}$  (compromised) data in Table 3.

Plots between ductile components and  $T_{2c}$  (compromised) values are presented in Figures 12 and 13. The two figures present different correlations, logarithmic on Figure 12 and linear in Figure 13. The correlations are

$$T_{2c} = 22.096 \ln (V_{dc}) + 1.2341$$
 (2)

and

 $T_{2c} = 1.6672 \quad (V_{dc}) + 33.697 \tag{3}$ 

where  $V_{dc}$  is percentage of ductile component.

With the establishment of equations (2) and (3), any  $T_{2c}$  values can be estimated whenever ductile components data becomes abailable. Following the same manner, comparison between NMR S<sub>wirr</sub> and core S<sub>wirr</sub> can be performed (Figures 14 and 15). Comparisons in the two figures appear to be a tendency of more optimistic estimated Swirr, indicated by lower NMR S<sub>wirr</sub> compared to core S<sub>wirr</sub> values.

Comparisons presented in Figures 14 and 15 have also shown that the establishment of general functions of equations (2) and (3) does not seem to help improving the agreement between NMR S<sub>wirr</sub> and core S<sub>wirr</sub>. This is true when they are compared to comparison presented in Figure 11, on which the T<sub>2c</sub> value used is simply the average of T<sub>2c</sub> values from BVI Error method.

In addition to Figures 14 and 15, Figure 16 presents comparison between NMR S<sub>wirr</sub> and core S<sub>wirr</sub> with the use of averaged T<sub>2c</sub> (compromised) value of 53.17 ms. From the figure it can be seen that slight improvement has been made when compared to Figures 14 and 15, and this improvement appears to have made plot in Figure 16 to resemble the plot in Figure 11. Again, up to this point it can be said that, despite the theoretical soundness of the empirical correlations, T<sub>2c</sub> averaging appears to be adequate for yielding representative T<sub>2c</sub>.

For the third option, five plots will be made, which h are comparison between core  $S_{wirr}$  and NMR  $S_{wirr}$ estimated using true  $T_{2c}$  values from BVI Error method: 32, 62, 70, 75, and 90 ms (Table 3). The comparisons are presented in Figures 17 through 21.

From Figures 17 through 21 it is obvious that good agreement between core  $S_{wirr}$  and NMR  $S_{wirr}$  are

shown by NMR S<sub>wirr</sub> values that are estimated using  $T_{2c}$  of 62, 70, and 75 ms. This suggests that the most representative  $T_{2c}$  for the six Tirrawarra sandstone samples used in the study should be around 70 ms, and if an averaged value is to be taken from the three values then it is 68.5 ms (i.e. this also counts as the fourth option,  $T_{2c}$  value picking). Notice the closeness of this value to the overall average  $T_{2c}$  of 66.5 ms (Figure 11).

#### VI. CONCLUSIONS

From the study that has been conducted, several main conclusions can be derived. The conventional use of a standard single NMR relaxation cut off time ( $T_{2c}$ ) values and varied  $T_{2c}$  values directly measured from spectrometer is not appropriate for NMR application on ductile shaly sandstones with high irreducible water saturation. It shows that a fixed or single predetermined  $T_{2c}$  would underestimate the "true" BVI and  $S_{wirr}$  values. BVI Error method appears to be the best method for providing varied  $T_{2c}$  values when compared to  $T_{2c}$  from SBVI and directly from spectrometer.

When the question comes into what  $T_{2c}$  value(s) are to be used for applied NMR log interpretation, comparisons shown using the four options have revealed that even for the case of shaly Tirrawarra sandstones, simple averaging seems to be adequate to yield a reliable single  $T_{2c}$  value. On the other hand, although establishment of empirical correlation may be theoretically more sound, results show that estimated NMR S<sub>wirr</sub> values produced are less accurate compared to simple averaging. Careful  $T_{2c}$  value picking also indicates a result that is close to result from simple averaging.

The overall conclusion of this study is certainly true for the ductile Tirrawarra sandstones used. Results may differ for other cases even though arguably the conclusion may also apply to other sandstones, especially for ones with level of complexities less than the highly heterogeneous Tirrawarra sandstones.

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