## CONDUCTIVITY ANISOTROPY IN SHALE-FREE SANDSTONE

W. David Kennedy ExxonMobil Exploration Co. David C. Herrick Baker Atlas

## Abstract

The existence of conductivity anisotropy has implications for formation evaluations using Archie's model for relating formation water saturation to formation resistivity. The bulk anisotropy of sedimentary rocks can arise from the interleaving of rock units having differing electrical properties. For special core analysis whole core is often selectively plugged in attempting to obtain homogeneous samples in the supposition that measurements taken on homogeneous samples will yield values representing typical reservoir properties. It is an article of faith that heterogeneous rocks composited from components similar to the homogeneous samples will exhibit physical properties predictably intermediate between the properties of the homogeneous samples. This is a reasonable belief that is demonstrably true for certain scalar parameters of the formation, for example porosity. Unfortunately, the directiondependent properties of the formation do not behave in this intuitive manner. For example, it can be shown in a model rock that the values of Archie's porosity exponent m and saturation exponent n for a composite rock can have values very different from its constituents. The exact values of m and n for composite rocks depend not only upon the values of the constituent components but also upon the relative volume fractions of the components, and the orientation at which the conductivity measurements are made. The theoretical results are supported by the general variability in the resistivities of samples having similar porosities in the experiment that determines m, manifest as the commonly observed "scatter" in the formation resistivity factor-porosity plots, and as anomalously low values of saturation exponents  $(n \approx 1)$  often observed in aeolian sandstones.

Neglecting the existence of these effects (as has been, and is still, done) must result in a false sense of the accuracy of formation evaluations, as well as an unwarranted lack of confidence in results from the laboratory when they are in conflict with naïve preconceptions of how formation properties should behave in the aggregate. A theoretical understanding of this issue must, at a minimum, improve estimates of uncertainty in formation evaluations. There is a definite (but as yet unrealized) promise that these effects can be accounted for and properly weighted in formation evaluations based upon triaxial induction logging instrument responses.

### Introduction

The electrical properties of reservoir rocks have been historically important because for most occurrences of hydrocarbons resistivity anomalies are as close to the

direct detection of hydrocarbons as any measurement made from the borehole provides. Archie (1942) showed how to relate the resistivity of a reservoir rock to the volume of fluids contained in the rock. Archie's relationship is empirical—its predictive power depends upon a knowledge of curve fitting parameters known as the porosity and saturation exponents, m and n respectively. These are determined in a laboratory from corederived samples of the reservoir, and the resulting values are taken as typical for the entire suite of rocks in the reservoir. In the absence of core-measured values for Archie parameters the nominal values are taken as equal to two, as a rule of thumb—slightly reduced for softer or younger rocks, slightly increased for harder or older rocks. There is a long-standing recognition of the problem of reservoir heterogeneity at the core scale. As a result, when special core analysis for Archie parameters is an objective, cores are typically plugged where they appear to be as visually homogeneous, and hence isotropic, as possible. On the other hand, in practice it is not easy to find sedimentary rocks, even at the coreplug scale, that are actually visually homogeneous and isotropic. This raises the question of whether the measured values actually provide the desired parameter values, and if they do not, do the Archie parameters determined from heterogeneous plugs represent an intuitively reasonable average of the properties of their constituents, leading to reasonable predictions of average water saturations, or are there unsuspected pitfalls in using them for interpretation?

A related question is why Archie's formation resistivity factor F is so poorly determined by the data used to infer it. That is, why do plots of porosity-formation resistivity factor data, used to determine m, exhibit so much scatter? Similar plots of water saturation-resistivity index data used for the determination of n are usually very well determined. Of course, the latter values are determined from a more controlled experiment, performed as it is on a single core plug. The conventional answer to the question of why there should be so much scatter in the corresponding porosity-formation resistivity factor plot—that the plugs used are taken from different places along the core, having differing porosities, with more-or-less differing depositional and diagenetic histories—no doubt has elements of truth, but is too vague to argue.

The discovery in the 1980s that saturation exponents could exhibit "anomalously" low values, close to or even less than one, was unexpected. At first the results of laboratory measurements were not accepted as being correct. Eventually it was understood that the laboratory

results were not in error, and low n values, especially in aeolian sandstones, became accepted (LaTorraca and Hall, 1991). In other circumstances the saturation exponent can be anomalously large, say four or greater (Herrick and Kennedy, 1996). My purpose here is to show that the range of values observed for Archie's m and n can be understood as a manifestation of anisotropy in shale-free sandstones.

## **Theoretical Development**

Consider a laminated sand reservoir comprising alternating sand layers that differ in their petrophysical properties (figure 1.a). For simplicity let us consider only two distinct sand types in the discussions to follow. The resulting two-component model is readily generalizable. For purposes of this study we consider each of the two sands to be internally, or locally, isotropic (figure 1.b). We also assume that the Archie (e.g., m and n) and other petrophysical parameters (e.g., porosity  $\varphi$ , water saturation  $S_w$ , etc.) of the individual layers are known, regardless that in actual practice the layers may be below the resolution limit for special core analysis. For specificity we assign an index "1" to the first component of the two-component model, and assign to it properties appropriate to a "fine-grained" component, and an index "2" to the second component, which is assigned properties appropriate to a "coarsegrained" component. The presumed consequence of grain-size variation is that, all other conditions being identical, the fine-grained rock, with its greater mineral surface area exposed to brine, will have its initial porosity diminished more rapidly, and therefore to a greater degree, by diagenesis (e.g., cementation, compaction, etc.) than the otherwise similar coarse-grained rock. In other words, the fine-grained component is assumed to have a lower porosity than the coarsegrained component in this model.

The bulk horizontal conductivity  $\sigma_h$  of this medium as computed from the volume-weighted average of the conductivities of the two sand fractions,  $\sigma_1$  and  $\sigma_2$ , where both of the sands are locally homogeneous and isotropic, is

$$\sigma_h = \beta \sigma_1 + (1 - \beta)\sigma_2 \tag{1}$$

where  $\beta$  (0.0 <  $\beta$  < 1.0) is the volume fraction of the fine-grained sand component. The bulk vertical conductivity  $\sigma_v$  is computed as the volume-weighted average of the reciprocal conductivities, viz.,

$$\frac{1}{\sigma_{\nu}} = \frac{\beta}{\sigma_{1}} + \frac{(1-\beta)}{\sigma_{2}} \ . \tag{2}$$
 We wish to examine certain petrophysical properties,

We wish to examine certain petrophysical properties, specifically Archie's parameters m, n, the formation resistivity factor F, and the resistivity index I, of a composite rock comprising layers below the limit of logging instrument resolution that therefore exhibits average (or bulk) properties that differ from the properties of the individual constituents.

You may note that the volume-weighted average (or bulk) porosity  $\overline{\varphi}$  of the sand-package assembled from the two component sands is

$$\varphi = \beta \varphi_1 + (1 - \beta) \varphi_2 \tag{3}$$

and that the volume-weighted average brine content  $\varphi S_w$  (figure 1.b) for the package will be

$$\overline{\varphi S_{w}} = \beta \varphi_{1} S_{w_{1}} + (1 - \beta) \varphi_{2} S_{w_{2}}.$$
(4)

Now define  $\overline{S}_w$  by  $\overline{S}_w \overline{\varphi} = \overline{\varphi S}_w$ . From this definition and equation (4), the average (or bulk) water saturation will be

$$\overline{S}_{w} = \frac{\beta \varphi_{1} S_{w_{1}} + (1 - \beta) \varphi_{2} S_{w_{2}}}{\overline{\varphi}}, 
= \frac{\beta \varphi_{1}}{\beta \varphi_{1} + (1 - \beta) \varphi_{2}} S_{w_{1}} + \frac{(1 - \beta) \varphi_{2}}{\beta \varphi_{1} + (1 - \beta) \varphi_{2}} S_{w_{2}}.$$
(5)

 $\overline{S}_w$  is also a volume weighted average; the first term on the right side of (5) is the fraction of the porosity that belongs to the first porosity component multiplied by the water saturation in that component, while the second term on the right side of (5) is the fraction of the porosity that belongs to the second porosity component multiplied by the water saturation in that component. Thus, in a sense (5) is a volume-weighted average of the brine content of the separate components. We will show that while  $\sigma_h$ ,  $1/\sigma_v$ ,  $\overline{\varphi}$ , and  $\overline{S}_w$  and a few other petrophysical properties of the composite medium behave in this "intuitively obvious" manner for averages, the important properties of averaged m and n are more complicated.

## Direction-dependent formation factors: $F_{\parallel}$ and $F_{\perp}$

In a model rock composited from two components having porosities  $\varphi_1$  and  $\varphi_2$  present in the volume fractions  $\beta$  and  $1-\beta$ , the parameter  $\beta$  is required explicitly in terms of the average porosity given in equation (3). This is

$$\beta = \frac{\varphi_2 - \overline{\varphi}}{\varphi_2 - \varphi_1}, \quad \text{where} \quad \varphi_1 \le \overline{\varphi} \le \varphi_2 \quad (6)$$

and where we have assumed for specificity that the fine-grained component has a lower porosity than the coarse-grained component. Archie's "law," usually expressed in terms of resistivity as  $R_t = R_w FI = R_w \cdot 1 / \varphi^m \cdot 1 / S_w^n$  where all of the symbols have their conventional meanings, is expressed in terms of conductivity as  $\sigma_t = \sigma_w \varphi^m S_w^n$  where the subscripts and exponents have their conventional meanings and conductivity  $\sigma = 1/R$  is reciprocal resistivity. When hydrocarbons are not present in the formation, by convention the subscript t is changed to 0. The parameters corre-

<sup>\*</sup>If  $\varphi_1$  and  $\varphi_2$  are distributed in layers thinner than can be resolved by a logging instrument,  $\overline{\varphi}$  corresponds to a "porosity" logging instrument response.

sponding to one or the other component are distinguished by the use of subscripts, 1 and 2.

Consider the case where  $S_w = 1.0$ . In equation (1) substituting  $\sigma_1 = \sigma_w \ \varphi_1^{m_1}$  and  $\sigma_2 = \sigma_w \ \varphi_2^{m_2}$ , and dividing by the connate water conductivity  $\sigma_w$ , results in

$$\frac{\sigma_h}{\sigma_w} = \frac{\sigma_{\parallel}}{\sigma_w} = \beta \varphi_1^{m_1} + (1 - \beta) \varphi_2^{m_2} = \frac{R_w}{R_{\parallel}} = \frac{1}{F_{\parallel}}$$
 (7)

where the subscript  $\parallel$  (read "parallel") indicates properties sampled parallel to laminations. In flat lying beds this component may happen to be "horizontal" and often is given a subscript h, but not all laminations are oriented in this simple way. The " $\parallel$ " notation avoids possible ambiguity in the interpretation of "horizontal." Thus

$$F_{\parallel} = \frac{1}{\beta \varphi_{1}^{m_{1}} + (1 - \beta)\varphi_{2}^{m_{2}}} \equiv \frac{1}{\overline{\varphi}^{m_{\parallel}}}$$
 (8)

where the definition of  $m_{\parallel}$  at the right end of (8) is made by analogy with the corresponding Archie relationship.  $F_{\parallel}$  and  $\overline{\varphi}$  are obtained from laboratory measurements made in the usual way.

Making the same substitutions in equation (2) will lead to

$$\frac{1}{\sigma_{v}} = \frac{1}{\sigma_{\perp}} = \frac{\beta}{\sigma_{w} \varphi_{1}^{m_{1}}} + \frac{1 - \beta}{\sigma_{w} \varphi_{2}^{m_{2}}} \tag{9}$$

or

$$F_{\perp} = \frac{\beta \varphi_2^{m_2} + (1 - \beta)\varphi_1^{m_1}}{\varphi_1^{m_1} \varphi_2^{m_2}} \equiv \frac{1}{\overline{\varphi}^{m_{\perp}}}$$
(10)

where the subscript  $\perp$  (for perpendicular, read "perp") indicates properties sampled perpendicular to bedding planes or laminations, avoiding ambiguity in the use of the word "vertical." Again, the definition at the right end is suggested by the analogous Archie relationship. In equations (6)–(10), notice that  $\overline{\varphi}$  is a scalar property while  $F_{\parallel}$ ,  $m_{\parallel}$ ,  $F_{\perp}$ , and  $m_{\perp}$  are all direction dependent.

## Archie's porosity exponent m

In terms of average properties the Archie relations for the bulk medium are direction dependent. In the equations expressing the relations, if one side is direction dependent, then of course, both sides must be direction dependent. That is, considering equations (8) and (10), if  $F_{\parallel}$  and  $F_{\perp}$  are direction dependent but  $\varphi$  (and  $\overline{\varphi}$ ) and  $S_w$  (and  $\overline{S}_w$ ) are not direction dependent, then the saturation exponent n and porosity exponent m, as the only parameters remaining on the right side of the equations, necessarily have to be the direction-dependent factors in Archie's relations connecting resistivity to porosity and water saturation. For example for the horizontal direction, in terms of conductivity, from (1) we have

the composite conductivity parallel to the bedding planes

Equation (11) will provide the definition of the porosity and saturation exponents parallel to the bedding planes.

In the case of  $S_{w_1} = S_{w_2} = 1.0$  the relation between the bulk horizontal conductivity and the sand fractions would be

$$\sigma_{l_{\parallel}} = \beta \, \sigma_{1} + (1 - \beta) \, \sigma_{2} = \sigma_{w} \overline{\varphi}^{m_{\parallel}} \, \overline{S}_{w}^{n_{\parallel}} = \sigma_{w} \left( \beta \, \varphi_{1}^{m_{1}} \, S_{w_{1}}^{n_{1}} + (1 - \beta) \, \varphi_{2}^{m_{2}} \, S_{w_{2}}^{n_{2}} \right).$$
(11)

$$\sigma_{0_{\parallel}} = \beta \sigma_w \, \varphi_1^{m_1} + (1 - \beta) \, \sigma_w \varphi_2^{m_2} = \sigma_w \overline{\varphi}^{m_{\parallel}}. \quad (12)$$

Thus

$$\overline{\varphi}^{m_{\parallel}} = \beta \, \varphi_1^{m_1} + (1 - \beta) \, \varphi_2^{m_2} = \frac{\sigma_{0_{\parallel}}}{\sigma_{u_1}},$$
 (13)

leading finally to a solution for  $m_{\parallel}$ 

$$m_{\parallel} = \frac{\ln(\beta \, \varphi_1^{m_1} + (1 - \beta) \, \varphi_2^{m_2})}{\ln \overline{\varphi}}.$$
 (14)

 $m_{\parallel}$  applies to conductivity measurements that are made parallel to the bedding planes. Similarly

$$\frac{1}{\sigma_{\perp}} = \frac{1}{\sigma_{w} \overline{\varphi}^{m_{\perp}} \overline{S}_{w}^{n_{\perp}}} = \frac{\beta}{\sigma_{w} \varphi_{1}^{m_{1}} S_{w_{1}}^{n_{1}}} + \frac{1 - \beta}{\sigma_{w} \varphi_{2}^{m_{2}} S_{w_{2}}^{n_{2}}} . (15)$$

Proceeding with a solution for this equation for  $m_{\perp}$ , for the case when  $S_{w_1} = S_{w_2} = 1.0$ 

$$m_{\perp} = \frac{\ln \left( \frac{\varphi_{1}^{m_{1}} \varphi_{2}^{m_{2}}}{\beta \varphi_{2}^{m_{2}} + (1 - \beta) \varphi_{1}^{m_{1}}} \right)}{\ln \overline{\varphi}^{m_{\perp}}}$$
(16)

where  $m_{\perp}$  applies to conductivity measurements that are made perpendicular to the bedding planes.

## Variation of m in a two component sandstone

Figures 2 and 3 illustrate a hypothetical and nearcanonical experiment of determining the porosity exponent parameter in a resistivity experiment. The resistivity of an ensemble of brine-saturated rocks of varying porosity is plotted as shown on a log-log plot. The end members of the ensemble shown have porosities between 5 and 40 percent. These values are chosen to represent the entire porosity range between a point near the percolation threshold and a point near the disaggregation limit. At the percolation threshold the conducting brine becomes a discontinuous phase. At the disaggregation limit the rock-forming mineral becomes a discontinuous phase. Since these represent the hypothetical limiting cases, results using them should encompass all measurements taken on "real" rocks. This obviously means that when observed in actual data, the effects

will typically be smaller than those presented for limiting cases.

In the Archie experiment, values of formation resistivity factor (=  $R_0/R_w$ ) are plotted against  $\varphi$  on a log-log grid and a line is fitted through the resulting cloud of points (the points are omitted from the figures). The canonical value of the slope of the line is 2. Figure 3 illustrates the more realistic situation where the slope departs from the canonical value. In younger, softer, higher porosity rocks the value of the porosity exponent tends to be less than 2. In older, harder, lower porosity rocks the value of the porosity exponent tends to be greater than 2. Figure 3 shows the cases of a line determined on two ensembles of samples, one having a porosity exponent of 1.8 and the other having a porosity exponent of 2.3. The canonical line with slope m=2 is shown for reference. I now propose a gedanken\*. Construct a laminated rock comprising two components: one component has 5 percent porosity and lies on the trend for m = 2.3 while the other component has 40 percent porosity and lies on the trend for m = 1.8. Imagine the relative proportions of these two components to vary over the entire range of possible volume fractions. What will be the resulting porosity exponent for the composite rock? The slope of the dashed line in figure 3 connecting the two end members offers a possible answer to this question. Is it a correct answer?

To resolve the issue analytically, begin with the consideration of the simpler case of the composite rock with end members that are both limiting elements of what we shall call the "canonical ensemble". That is, consider two sandstone components having differing porosities of 5 and 40 percent but having the same porosity exponent, m = 2. If we composite the gedanken rock from these components, many properties of the composite will be volume-weighted averages of the end members. Assuming this to be true of the porosity exponent, then it seems reasonably "obvious" that the porosity exponent of the composite rock will lie on the canonical m = 2 line. This may be an "obvious" assumption to adopt, but it is not correct.

Figure 4 shows the results of using equations (8) and (10) to compute the formation resistivity factors parallel and perpendicular to the bedding planes. The heavy dashed line marks m=2 on the plot. The coordinates of the end member components are circled. As fractional volume is varied between the end members, the formation resistivity factor measured parallel to the bedding planes follows the path traced by the lower solid curve. Measured perpendicular to the bedding planes, the formation resistivity factor follows the path traced by the upper dash-dot curve. An envelope (the two dotted lines) is drawn from the "brine point" (i.e.,  $\varphi = 1.0$ , F =

When the end members of the composite are modeled by the more realistically chosen porosity exponents illustrated in figure 3, the composite porosity exponent satisfies 1.5 < m < 4.75.

Because in practice the measurement of m is always attempted parallel to the bedding planes, the range of variability will typically be less than the maximum ranges indicated above. It is natural to question whether rocks composited of end members, each having m = 2, could be said to have a variation in porosity exponent limited to the range 1.6 < m < 2.0. This is true to the degree that measurements are successfully aligned with the bedding planes. But, considering the model illustrated in figures 2, 4, and 5, for a misalignment of 30°, depending upon the relative volume fractions of the components, it can be shown that  $m \approx 2.2$  (for  $\beta \approx 0.1$ ) or  $m \approx 1.7$  (for  $\beta \approx 0.9$ ). Our intuition, founded as it is in linear variations, is not a useful guide. These estimates are supported by computations based on the analysis given in the Appendix.

Of course, when the end members of the composite are more similar in porosity than  $\varphi_1 = 0.05$  and  $\varphi_2 = 0.40$ , the effects of fractional composition and measurement alignment are diminished. But, that is not to say that there are no practical consequences.

The gedanken strongly suggests that the customary procedure of selectively acquiring the core plugs used in special core analysis in the most homogeneous units visible in the whole core, if it is entirely successful, will not lead to correct predictions of the porosity exponents that are representative of nearby laminated rocks composited from the homogeneous components if simple, volume-weighted arithmetic averaging is used in the prediction. On the other hand, given the difficulty in identifying such homogeneous units, a possible cause (in addition to many other causes that have been postulated) for the often-observed "scatter" in these data is a failure to control the experiment for the variability of laminations among the plugs in a set of samples.

<sup>1.0)</sup> through the upper and lower tangents to the functions. The result shows that depending on the orientation of the sample and the volume fraction of the components, the composite rock can have porosity exponents that vary from 1.6 < m < 4.0. Figure 5 plots the same data in a more dramatic diagram that shows the porosity exponent plotted against the volume fraction of the rock components. From this we see that the average porosity exponent of the composite rock does not conform to our naïve notion of the expected behavior of this rock property.

thought experiment

# Direction-dependent resistivity indices: $I_{\parallel}$ and $I_{\perp}$

We now consider the conductivity behavior of the composite rock for cases where  $S_w < 1.0$ . The resistivity index is defined as

$$I = \frac{R_t}{R_0} = \frac{\sigma_0}{\sigma_t} = S_w^{-n} \,. \tag{17}$$

By analogy with the resistivity index for an isotropic medium

$$I_{\parallel} = \frac{\sigma_{0_{\parallel}}}{\sigma_{t_{\parallel}}} = \frac{\beta \varphi_{1}^{m_{1}} + (1 - \beta)\varphi_{2}^{m_{2}}}{\beta \varphi_{1}^{m_{1}} S_{w_{1}}^{n_{1}} + (1 - \beta)\varphi_{2}^{m_{2}} S_{w_{2}}^{n_{2}}} \equiv \frac{1}{\overline{S}_{w}^{n_{\parallel}}}$$
(18)

where the definition at the right end of (18) is made by analogy with the corresponding Archie relationship. Similarly

$$I_{\perp} = \frac{\sigma_{0_{\perp}}}{\sigma_{t_{\perp}}} = \frac{\beta \varphi_{2}^{m_{2}} S_{w_{2}}^{n_{2}} + (1 - \beta) \varphi_{1}^{m_{1}} S_{w_{1}}^{n_{1}}}{\varphi_{1}^{m_{1}} S_{w_{1}}^{n_{1}} \varphi_{2}^{m_{2}} S_{w_{2}}^{n_{2}}} \equiv \frac{1}{\overline{S}_{w}^{n_{\parallel}}}$$
(19)

where, again, the definition at the right end of (19) is suggested by the analogous Archie relationship. Note that  $\overline{S}_w$  is a scalar property while  $I_{\parallel}$ ,  $n_{\parallel}$ ,  $I_{\perp}$ , and  $n_{\perp}$  are all direction dependent.

# Archie's saturation exponent n

Given  $m_{\parallel}$  and  $m_{\perp}$ , bulk values for  $n_{\parallel}$  and  $n_{\perp}$  can be solved for. Starting again with equation (1)

$$\sigma_{\parallel} = \beta \sigma_1 + (1 - \beta) \sigma_2 \tag{1}$$

and substituting the expressions for the conductivities of  $\sigma_1$ ,  $\sigma_2$ , and  $\sigma_{\parallel}$  results in

$$\sigma_{w}\overline{\varphi}^{n_{\parallel}}\overline{S}_{w}^{n_{\parallel}} = \beta\sigma_{w}\varphi_{1}^{n_{1}}S_{w_{1}}^{n_{1}} + (1-\beta)\sigma_{w}\varphi_{2}^{n_{2}}S_{w_{2}}^{n_{2}}. \quad (20)$$

The solution for  $n_{\parallel}$  is

$$n_{\parallel} = \frac{\ln \left( \frac{\beta \sigma_{w} \varphi_{1}^{m_{1}} S_{w_{1}}^{n_{1}} + (1 - \beta) \sigma_{w} \varphi_{2}^{m_{2}} S_{w_{2}}^{n_{2}}}{\overline{\varphi}^{m_{\parallel}}} \right)}{\ln \overline{S}_{w}} . \quad (21)$$

Note that  $\varphi_1$ ,  $\varphi_2$ ,  $m_1$ , and  $m_2$  are fixed parameters of the two rock components chosen for the particular model. Similarly  $S_{w_1}$ ,  $S_{w_2}$ ,  $n_1$ , and  $n_2$  are likewise

fixed for the rock components chosen.  $S_{w_1}$  and  $S_{w_2}$  are, however, not independent since both are functions of capillary pressure, so a model relating  $S_{w_1}$  and  $S_{w_2}$  is required to apply this equation. The independent variable in the model actually used is normalized height above "free" water in a reservoir. Otherwise, the only variable is  $\beta$ , the volume fraction of the components.

Similarly for the perpendicular component, beginning with equation (2)

$$\frac{1}{\sigma_1} = \frac{\beta}{\sigma_1} + \frac{(1-\beta)}{\sigma_2},\tag{2}$$

and following the same series of steps that led to equation (21) gives

$$\overline{\varphi}^{m_{\perp}} \overline{S}_{w}^{n_{\perp}} = \frac{\varphi_{1}^{m_{1}} S_{w_{1}}^{n_{1}} \varphi_{2}^{m_{2}} S_{w_{2}}^{n_{2}}}{\beta \varphi_{2}^{m_{2}} S_{w_{2}}^{n_{2}} + (1 - \beta) \varphi_{1}^{m_{1}} S_{w_{1}}^{n_{1}}}.$$
 (22)

This is easily solved for  $n_L$ ,

$$n_{\perp} = \frac{\ln \left( \frac{1}{\overline{\varphi}^{m_{\perp}}} \frac{\varphi_{1}^{m_{1}} S_{w_{1}}^{n_{1}} \varphi_{2}^{m_{2}} S_{w_{2}}^{n_{2}}}{\beta \varphi_{2}^{m_{2}} S_{w_{2}}^{n_{2}} + (1 - \beta) \varphi_{1}^{m_{1}} S_{w_{1}}^{n_{1}}} \right)}{\ln \overline{S}_{w}} . \quad (23)$$

# A model for water saturation and height above free water level in the reservoir

To compute values for  $n_{\parallel}$  and  $n_{\perp}$  it is necessary to evaluate the water saturations of the two sand fractions in the model at the same height above the free water level in the reservoir. In other words, a model for the saturation-height function is required. For generality we introduce a function for computing water-saturation at each height in the reservoir, expressed in terms of the parameter h/H, where H is the thickness of the oil column and h/H is the fractional height above the "free water level". We use

$$S_{w} = \begin{cases} 1 & h < h_{0} \\ 1 - \left(1 - S_{w_{irr}}\right) \left(1 - e^{-\alpha \left(\frac{h}{H} - \frac{h_{0}}{H}\right)}\right) & h > h_{0} \end{cases}$$
 (24)

to model the "saturation-height" function. In this model  $S_{w_{irr}}$  has its usual loosely defined meaning as a pseudo-asymptote. If  $h_0$  is the height above the hydrocarbon-water contact at which hydrocarbon can displace brine from the largest pores in the rock,  $h_0/H$  corresponds to an "entry" pressure for those points, and  $\alpha$  is a "decay constant" that controls the rate at which water saturation diminishes as h/H increases. This function is a qualitative model for observed capillary pressure curves. The sand fractions in the model are assigned different representative values of  $\alpha$ ,  $S_{w_{irr}}$ , and  $h_0/H$  in order to closely mimic actual capillary pressure data. Making the substitutions

$$S_{w} = \begin{cases} 1 & h < h_{0_{1}} \\ 1 - \left(1 - S_{w_{irr}}\right) \left(1 - e^{-\alpha_{1}\left(\frac{h}{H} - \frac{h_{0_{1}}}{H}\right)}\right) & h > h_{0_{1}} \equiv S_{w_{1}}(h/H) \end{cases}$$
Figure 7 illustrates the relation of the conventional resistivity index plot (shown on the left) and the axesexchanged saturation-height plot (shown on the right). Since  $S_{w}$  depends upon  $h/H$ , the end-member water saturations for the resistivity index plot will vary with  $h/H$ , although the variation may well be small far from the hydrocarbon-water contact. The average water saturation is a function of the volume fraction of the components of the two-component system. For  $\beta = 1$  the

$$S_{w} = \begin{cases} 1 & h < h_{0_{2}} \\ 1 - \left(1 - S_{w_{irr}}\right) \left(1 - e^{-\alpha_{2} \left(\frac{h}{H} - \frac{h_{0_{2}}}{H}\right)}\right) & h > h_{0_{2}} \equiv S_{w_{2}}(h/H) \end{cases}$$

$$(25.b)$$

effectively changes the variable from two different water saturations to a single value of h/H at any given height in the reservoir. Armed with this model,  $n_{\parallel}$  and  $n_{\perp}$  can be computed as functions of  $0 < \beta < 1$  and 0 <h/H < 1. We emphasize that the function used in (24) is specified for convenience only. Any similar function that may be deemed more "realistic" can be used without changing the effects on the Archie parameters of the composite rock, or the results and conclusions to be drawn therefrom.

Note that I versus  $S_w$  has meaning only where  $S_w < 1$ . The relationship of  $S_{w_lirr}$  to  $S_{w_2irr}$  will depend on the saturation height function for the two components. We consider the usual case where the higher porosity component will typically exhibit a lower water saturation than the lower porosity component. The most succinct notation for expressing the expected relationship is  $S_{w_0}$  $\leq S_{w_i} \leq 1$ ; i.e., the water saturation in the fine-grained (subscript 1), component is greater than, or at most equal to, the water saturation in the coarse-grained (subscript 2), component, and less than, or at most equal to, a fractional volume of 1.0. Accordingly the domain of  $n_{\parallel}$  and  $n_{\perp}$  will be taken as, in succinct notation,  $h_{02}/H < h_{01}/H < 1$ . This is to say that the saturation exponent is not defined, nor even definable, until  $S_w$ departs from  $S_w = 1.0$ .

## Illustrating the variation of n in a two-component sandstone

Figure 6 illustrates the saturation-height functions for two sandstone samples. For this plot water saturation is usually presented on the abscissa and height above the hydrocarbon-water contact on the ordinate. The scales in figure 6 are exchanged to make figure 7 more readily interpretable. That is, we will plot h/H on the horizontal axis and two functions of water saturation, representing the two sand fractions, on the vertical axis. The diagram is illustrating two sands with two different entry pressures, indicated by differing values of h/H, and two different "irreducible" water saturations.

saturations for the resistivity index plot will vary with h/H, although the variation may well be small far from the hydrocarbon-water contact. The average water saturation is a function of the volume fraction of the components of the two-component system. For  $\beta = 1$  the saturation-height function is identical to one end member; at  $\beta = 0.0$  the saturation-height function is identical to the other end member; for  $0.0 < \beta < 1.0$  the saturation-height function is a smooth average of the end component saturation-height functions weighted by the porosity volume fractions of the components.

Figure 8 illustrates four cases of the "parallel" and "perpendicular" resistivity index behavior for sandstone end members having various values of porosity, and porosity and saturation exponents. The figures show clearly that "parallel" saturation exponents  $n \approx 1$  are a possible consequence of a rock's being composited from laminated sandstone fractions (figures 8.a and 8.d). Even when all the end member Archie properties (i.e.,  $\varphi$ , m, n) are equal (or approximately so), the difference in capillary pressure properties is sufficient to induce a strong variation of apparent saturation exponent as a function of volume fraction of the components (figures 8.b and 8.c).

Figure 8 shows a series of "snapshots" corresponding to a single value of h/H. Figures 9.a and 9.b are complete representations of n for all values of h/H, as well as all values of the fractional volumes of the components, shown in a representation similar to the presentations of m in figure 5. But whereas m depends only upon pore geometry, n depends also upon brine geometry. Obviously, while pore geometry varies with position, otherwise it is a constant property of the rock. However, brine geometry changes as water saturation varies making the saturation exponent not only a function of position, but also a function of water saturation. The surfaces shown in figures 9.a and 9.b bound the volume that contains all the possible values that could be measured for n in the composite medium given the parameters assumed for the end members. The plotting facility did not support the simultaneous plotting of figures 9.a and 9.b on the same graph; however, figure 9.b fits atop figure 9.a along the line defined by n = 2. In the model of figure 9, 1.25 < n < 7 approximately. Since the axes of core plugs are usually selected as close to parallel to the bedding planes as possible, n for core plugs in this rock would tend to be measured as 1.25 < n < 2.00 approximately, depending upon the fractional composition and the height above the hydrocarbon-water contact.

### Conclusion

Layered media composited from laminations having differing physical properties give rise to anisotropic physical properties at scales larger than the laminations (Maxwell, 1871). Formulas for the conductivity of such composite media are at least 80 years old in the geophysical literature (Schlumberger, 1920). However, only relatively recently have the implications for formation evaluation parameters begun to be recognized (Herrick and Kennedy, 1996). Using the methods of Maxwell and Schlumberger, in this report we have shown that simple laminated rocks composited from layers of locally isotropic, well-behaved sandstones, can exhibit Archie parameters that seem counterintuitive and that are not simple interpolations between the Archie parameters of the component fractions. The fact is that the correct values of the Archie parameters for the composited rocks comprise the components of tensors, and are not simple volume-weighted averages of the values of the components.

Subsequent to Archie's formulation (1942) of a relationship between formation resistivity and hydrocarbon saturation there followed several decades of vigorous refinement of the understanding of this relationship. However, a thorough consideration of anisotropy appears never to have been an objective of these studies. This lack of concern may have followed from an unexpressed<sup>†</sup> recognition that (in vertical wells) even where anisotropy exists, logging instruments respond directly only to its "horizontal" component,  $\sigma_h$  (ignoring that there might be two different horizontal components,  $\sigma_{\rm x}$ and  $\sigma_{\nu}$ ), cores are plugged horizontally when analysis of their Archie parameters is an objective, and it was generally believed that while shales were demonstrably anisotropic, sands should be isotropic, at least approximately. Horizontal drilling and logging have shown such a view of reservoir rocks to be too simple, and it is a current industry goal to understand the effects of anisotropy both on logging instrument responses and also on the industry-standard techniques of formation evaluation. The effects reported upon herein have indicated that there is a definite effect in layered rocks. Since all clastic rocks are layered to a greater or lesser degree, evidence of these has doubtless been observed but has gone unremarked upon.

The practicality of these results depends upon whether the effects are significant, not for end members selected to have extreme property values for theoretical purposes, but for ensembles of rocks encountered in actual petroleum reservoirs observed using standard techniques. The resistivity index experiment—covering, as

it does, all water saturations from  $S_w = 1.0$  down to  $S_{w_{irr}}$ —naturally observes the entire range of the  $S_w$  parameter from one saturation end member to the other. The variation in  $n_{\parallel}$  predicted by the two-component model is totally supported by observations in rocks (LaTorraca and Hall, 1991).

On the other hand, porosity variations in real reservoirs are typically less than the extremes that we have used as limiting-case end members. What will be the effect of taking variation more typical of actual reservoirs? Let us first recognize that if the pure porosity end members of the composite are never observed because they do not occur in quantities thick enough to be sampled as "homogeneous" plugs, then the porosity variation observed will be less than the porosity variation between the actual end members. In other words, it is possible that the scale of our observations is too coarse to sample the porosity variation that actually exists. In this situation the fact that bulk porosity is observed to vary over a certain range is not equivalent to a statement regarding the variation between the true end members of the composite, which will be larger. To identify this condition core plugs would have to be examined on a scale as small as the visible laminations in the plugs. We are not arguing that every reservoir, or any reservoir, has actual porosity end members near 5 porosity units (pu) and 40 pu. We are saying that the mere fact that porosity varies between two "apparent" limiting values in a certain suite of observations on core plugs should not necessarily be interpreted as the observed porosity values representing the porosity end member values in those plugs.

There should be observable consequences of a limited observable  $\beta$  range. Recall that the formation resistivity factor is often defined as  $F = a/\varphi^m$ , where a is the "intercept" of a line least squares fitted to observed formation resistivity factor-porosity data with the vertical axis corresponding to  $\varphi = 100$  porosity units (pu). For example, in the well known "Humble" formula, a =0.62, m = 2.15 (Winsauer et al., 1952). Specifically, a < 0.621 for the Humble formula. For other data sets a > 1. We illustrate in figure 10 that for a case where the reservoir is laminated with typically low  $\beta$  (higher porosity), a line fit through data collected at random in such a reservoir will exhibit a > 1. Conversely, for a reservoir with typically high  $\beta$  values (lower porosity), a < 1. Thus, use of conductivity anisotropy theory explains the origin of intercepts  $a \neq 1$  in terms of sand lamination volume fraction distribution. We do not claim that effects of anisotropy and skewed volume fraction distribution of conductive sand components in laminated reservoirs is the sole cause for  $a \ne 1$ , but do suggest that it is at least one heretofore unrecognized cause.

т

<sup>&</sup>lt;sup>†</sup> The so called "paradox of anisotropy" was discussed by Schlumberger (1920), seven years prior to the invention of well logging.

Figure 11 illustrates the effect on porosity exponents of a smaller range of porosity variation in the end members. Not surprisingly the effect diminishes as the end members approach one another. Obviously the effect would vanish for end members chosen to have the same porosity and the same Archie exponent for they would then be electrically identical. A study of many cases (not shown) reveals that the details of the porosity exponent function depend not only upon the separation in porosity units but also upon where the end members are located on the porosity axis. The effects of laminations are likely to be a consistently smaller source of uncertainty in m than in n, especially if the core plug axes are substantially parallel to the planes bounding the laminations. However, in formation analysis a conscious and explicit choice of whether to ignore, or to attempt to account for, anisotropy effects on log responses and parameter selection should be made on a case by case basis.

This analysis has shown that small errors in orientation of core plugs lead to correspondingly small errors in the determination of m values in special core analysis. The close analogy of the relations of F to m and I to n suggests that if  $m \approx m_{\parallel}$  for small departures of the core plug axis from parallel to laminations, then  $n \approx n_{\parallel}$  should also be true. The analysis has likewise shown that a large amount of variation in these values, depending on the volume fraction of the constituent components, is legitimate. This variation has gone largely unexplained in the past, or else has been attributed to measurement error in special core analysis.

The first representatives of tri-axial induction logging instruments capable of separately resolving the components of the conductivity tensor are now being employed in the field. Whereas heretofore formation resistivity was considered to be a single number  $R_t$ , using these instruments it will be resolved into two  $(R_{\parallel}, R_{\perp})$ , or even three  $(R_x, R_y, R_z)$  numbers at each point in the formation. Thus, the implications for formation evaluation of the variation in Archie's parameters m and n (or, we should perhaps say "Archie's tensors "m and n") with the composition and orientation of sand laminations—or indeed, whether there are any unrecognized implications—clearly needs to be evaluated. Such an evaluation is the next step in our program to understand and quantify the electrical behavior of reservoir rocks, as used in formation evaluation, as thoroughly as possible.

Also of note is a feature of the use of power laws in the description of the relationships connecting formation resistivity factors and porosity, and resistivity index and water saturation. As we have calculated them for this report, the values of  $m_{\parallel}$  and  $m_{\perp}$  at a point in the log  $\varphi$ -log F plane is the slope of a line connecting that point

with  $(\varphi = 1, F = 1)$  having a formula of the form  $m = -\log F / \log \varphi$ . As suggested by figures 3 and 10, if the  $\log \varphi - \log F$  function is not a power law containing the point (1,1) the slope of its graph would be better defined as  $m = d(\log F)/d(\log \varphi)$ . The latter (re)definition would require a lengthy study and explanation of its consequences. Such a study is beyond the scope of this research. But although the numerical values of m (and n) would be changed by such a redefinition, the fundamental conclusions of this study regarding the genesis of anisotropy in shale-free sand would remain unchanged. It is interesting to note that Winsauer's (1952) method implicitly finds a line that is parallel to a tangent to a curve that could be imagined to describe the trajectory of a data set in  $\log \varphi - \log F$  space.

We close this report with a remark on the generality of these results. The use of a two-component composite in this study is engendered solely by its being the simplest non-trivial case. Clearly real composites can be, and are, multicomponented. The two-component analysis generalizes in the obvious way, but there is insufficient space in this article to show the explicit development.

# Acknowledgment

The suggestions for enhancement of the content of this report offered by Drs. Hezhu Yin, Dale Fitz, Quinn Passey, and other ExxonMobil colleagues are greatly appreciated and contributed to making this a more thorough and readable report. We appreciate ExxonMobil management's granting their permission to communicate these findings to the greater petrophysics community.

# References

Archie, G. E., 1942, The electrical resistivity log as an aid in determining some reservoir characteristics, *Petroleum Technology*, Vol. 5, No. 1, T.P. 1422, 8 p.

Herrick, D. C., Kennedy, W. D., 1996, Electrical properties of rocks: Effects of secondary porosity, laminations, and thin beds, *Transactions* of the Society of Professional Well Log Analysts, paper C.

Kennedy, W. D., Herrick, D. C., Yao, T., 2001, Calculating Water Saturation in Electrically Anisotropic Media, *Petrophysics*, Vol. 42, No. 2, 118–136.

LaTorraca, Gerald A., Hall, Connie G., 1991, Observations of rock fabric controls on the electrical properties of sandstones, *Transactions* of the Society of Core Analysts, SCA Conference Paper 9116.

Maxwell, James C., 1871, in *A Treatise on Electricity and Magnetism*, Volume I (Third Edition, Preface by J. J. Thompson) in the section "On Stratified Conductors", article 321 appearing at the bottom of page 446.

Schlumberger, C., 1920, Etude sur la prospection électrique du sous-sol, Paris, Gauthier-Villars, p. 40.

Winsauer, W. O., Shearin, H. M., Jr., Masson, P. H., Williams, M., 1952, Resistivity of brine-saturated sands in relation to pore geometry, AAPG Bulletin, Vol. 38, No. 2, 253-277.

## **Appendix: Application to Tensor Archie Theory**

Kennedy et al. (2001) showed that Archie's law for isotropic rocks generalizes into the form

$$\hat{\sigma}_t = \sigma_w \varphi^{\hat{\mathbf{m}}} S_w^{\hat{\mathbf{n}}} \tag{A.1}$$

where the conductivity tensor on the left,  $\hat{\sigma}_t$ , is reflected on the right side by the "porosity" tensor  $\varphi^{\hat{\mathbf{m}}}$  and the "saturation" tensor  $S_w^{\hat{\mathbf{n}}}$  . Note that  $\varphi$  and  $S_w$ themselves are scalars, but raised to the matrix powers the resulting quantities are tensors. We could guess that  $\varphi^{\hat{\mathbf{m}}}$  and  $S_w^{\hat{\mathbf{n}}}$  are tensor quantities simply because the left hand side of (A.1) is a tensor, thereby requiring whatever is on the right side to also be a tensor quantity. Tensors are distinguished from other matrices by their properties under coordinate transformations and it can be shown by means of such transformations that  $\hat{\mathbf{m}}$ and  $\hat{\mathbf{n}}$  are themselves tensors in their own right. Kennedy et al. contemplated that the principal components of the conductivity tensor would be available through laboratory measurements, and we have shown that for transversely isotropic media comprised of locally isotropic, laminated sands, the principal components can be theoretically predicted if the volume fraction and Archie parameters of the individual components are known. Thus

$$\hat{\mathbf{m}}_{\mathrm{D}} = \begin{bmatrix} m_{\parallel} & 0 & 0 \\ 0 & m_{\parallel} & 0 \\ 0 & 0 & m_{\perp} \end{bmatrix}, \quad \hat{\mathbf{n}}_{\mathrm{D}} = \begin{bmatrix} n_{\parallel} & 0 & 0 \\ 0 & n_{\parallel} & 0 \\ 0 & 0 & n_{\perp} \end{bmatrix}, \text{ (A.2)}$$

where  $m_{\parallel}$ ,  $m_{\perp}$ ,  $n_{\parallel}$ , and  $n_{\perp}$  are given by the formulas developed in the text for a two-component system.

These tensor components are functions of the fractional volumes of the constituents, and indicate the results of experiments parallel and perpendicular to bedding planes, but it is natural to question what effect the orientation might have if the measurements are skewed with respect to these directions. Is the effect large or small?

One way to approximate an answer is graphically. For example, in figure 5 the solid (lower) line represents

measurements taken at 0° with respect to the bedding planes, while the dashed (upper) line represents measurements taken at 90° with respect to the bedding planes. Perhaps measurements at intermediate angles might be expected to be intermediate in some fashion. So a grid of parameter lines at 10°, 20°, ..., 80° could be sketched between the solid and dashed lines using some interpolation scheme. The only physics applied would be an assumption that the function of angle, whatever it may be, is smoothly varying. The theme of this article is that such a method may not work well. On the other hand, the problem is somewhat, if not entirely, amenable to a theoretical analysis.

Consider. In unbounded media the component of the Archie parameters in various directions could be obtained directly from the principal component forms (i.e., equations (A.2) and their tensor properties. Thus at a relative deviation of  $\delta$ 

$$\hat{\mathbf{m}} = \hat{\mathbf{R}} \hat{\mathbf{m}}_{\mathrm{D}} \hat{\mathbf{R}}^{\mathrm{T}} = \begin{bmatrix} m_{xx} & m_{xy} & m_{xz} \\ m_{yx} & m_{yy} & m_{yz} \\ m_{zx} & m_{zy} & m_{zz} \end{bmatrix}$$
(A.3)

where  $\hat{\mathbf{R}}^T$  is the transpose of  $\hat{\mathbf{R}}$  and

$$\hat{\mathbf{R}}(\delta) = \begin{bmatrix} \cos \delta & 0 & -\sin \delta \\ 0 & 1 & 0 \\ \sin \delta & 0 & \cos \delta \end{bmatrix}, \quad (A.4)$$

and a similar expression would hold for  $\hat{\mathbf{n}}$ . Unfortunately, in the laboratory things are not so simple.

Figure 12 illustrates that in the unbounded medium there is no doubt that the current density J has a component parallel  $J_{\parallel E}$  (read "J parallel to E") to, and a component perpendicular  $J_{\perp E}$  (read "J perpendicular to E") to, the applied electric field E; i.e., linear superposition holds for the current density field. However, as figure 12 also illustrates, the  $J_{\perp E}$  component is constrained by the physical boundary of a core plug. Since the current density component perpendicular to the applied field cannot circulate, as suggested in figure 12 only the  $J_{\parallel E}$  component is non-zero, and the effective conductivity of the medium is given by the ratio  $J_{\parallel E}/|E|$ . This pretty picture is spoiled by end effects that are not amenable to elementary analysis. We believe such effects will be small for core plugs with a large aspect ratio; i.e., for "long" plugs. End effects could not be neglected for short plugs, and for plugs having an aspect ratio near unity a resort to numerical methods would be necessary to determine the magnitude of the current traversing the plug.

Т

The case of the "long" core plug is susceptible to analysis. Consider a hypothetical medium, of the kind we have been discussing, composited of two components.

The medium is therefore anisotropic at a scale larger than its layer thicknesses. We want to consider two core plugs cut parallel and perpendicular to the tensor principal axes, and a set cut every one degree between the parallel (call this x-directed) and the perpendicular (call this z-directed) principal components of the conductivity tensor. To each of these samples we propose to apply an electric field of 1 volt per meter. For the sample cut at an angle  $\delta$  with respect to the horizontal principal axis, the electric field components are  $E_x = \cos \delta$  and  $E_z = \sin \delta$ . The current density components are then  $J_x =$  $\sigma_x E_x = \sigma_x \cos \delta$  and  $J_z = \sigma_z E_z = \sigma_z \sin \delta$ . In general, because the medium is anisotropic, this current density vector J is not parallel to the applied electric field E. The angle  $\psi$  between **J** and **E** is given from elementary vector analysis as

$$\cos \psi = \frac{\mathbf{E} \cdot \mathbf{J}}{|\mathbf{E}||\mathbf{J}|} = \frac{\sigma_x E_x^2 + \sigma_z E_z^2}{\sqrt{E_x^2 + E_z^2} \sqrt{(\sigma_x E_x)^2 + (\sigma_z E_z)^2}}$$
(A.5)

The current that can flow through the core plug that we are considering is constrained to flow parallel to the plug axis. This will be the component of current density that is parallel to **E**. This component is just the projection of **J** onto **E** given by  $J_{\parallel E} = |\mathbf{J}|\cos \psi$ . Given that  $|\mathbf{E}| = 1$ , the result is simply  $J_{\parallel E} = \sigma_x \cos^2 \psi + \sigma_z \sin^2 \psi$ .

From Ohm's law in field theory form, viz.,  $\mathbf{J} = \hat{\sigma} \mathbf{E}$ , we assert that  $\sigma_{\parallel \mathbf{E}} = J_{\parallel \mathbf{E}} / |\mathbf{E}| = J_{\parallel \mathbf{E}}$ , since  $|\mathbf{E}| = 1$  by design.

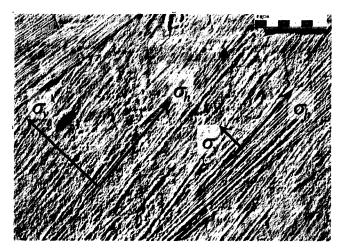


FIG. 1.a. Sand peel of a shale-free slip face of a West Texas dune. The laminations are due to grain size variations. The scale is in centimeters. The coordinates on the left are taken as the parallel and perpendicular (to bedding) components of the conductivity tensor for the case of  $S_w = 1$ . For  $S_w < 1$  the perpendicular conductivity is reduced more than the parallel conductivity, suggested by the coordinate frame on the right.

By an extension of Archie's definition  $\sigma_{\parallel \rm E} = \sigma_w \overline{\varphi}^{m_{\parallel \rm E}}$ . This can be solved for  $m_{\parallel \rm E}$  resulting in

$$m_{\parallel E} = \frac{\ln \sigma_{\parallel E} - \ln \sigma_{w}}{\ln \overline{\varphi}} . \tag{A.6}$$

Now  $m_{\parallel E} = m_{\parallel E}$  ( $\beta$ ,  $\delta$ ), depending upon  $\beta$  through the conductivity and average porosity, and upon  $\delta$  through the angle that the axis of the core plug makes with the planes of lamination. This is the formula that we have used to compute the approximate parameter curves in figure 13. The parameters are exact in the limit of a very long core plug, otherwise they are approximate. Their direct connection to the physics of this problem makes them a more useful heuristic device than a mere interpolation would have provided. For example, it can be seen in figure 13 that the line at 10° from the horizontal is essentially not distinguishable from the horizontal parameter line itself. This means that slight errors in choosing the direction of a plug axis will have a negligible effect on measurements made on the core. On the other hand, the shape of the intermediate curves is quite different than would be obtained by an interpolation.

A similar, albeit more complicated, analysis would yield the  $n_{\parallel E}$  surfaces that would partition the volume bounded by  $n_{\parallel}$  and  $n_{\perp}$  shown in figures 9.

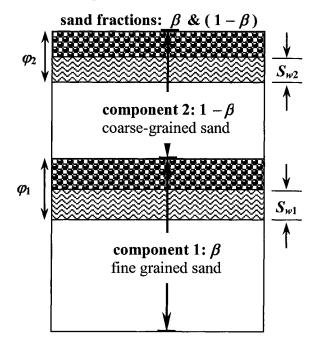


FIG. 1.b. This cartoon illustrates the components of a laminated sand package. The porosity and fluid fraction of each component has been lumped for purposes of easy illustration. The hydrocarbon fraction is coded as black spheres.

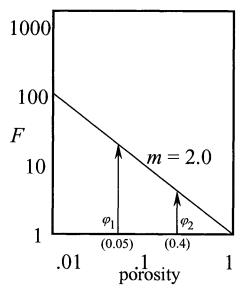


FIG. 2. Consider a sand package comprising layers of two distinct types of sand. Both sands fall along the m=2 line of the formation resistivity factor-porosity plot. In other words, although the porosity varies, the Archie parameters are the same for each type of rock. Obviously, the porosity of the composite rock is the volume-weighted average of the two components. What is porosity exponent m for the composite?

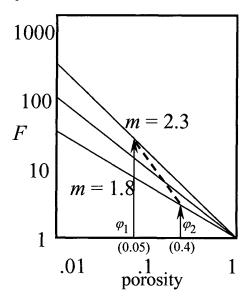


FIG. 3. In this case the Archie m parameter varies with the porosity, as it might be expected to do in some real rocks. The low porosity fraction is assigned m = 2.3; the higher porosity fraction is assigned m = 1.8. In an anisotropic rock composited from these two fractions, we might expect the slope of the formation resistivity factor-porosity line to vary linearly between the end points. However, the actual variation is more interesting.

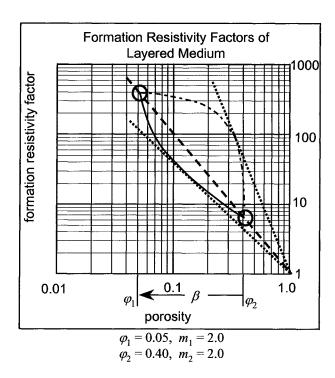
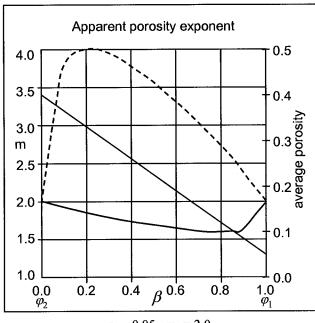


FIG. 4. A hypothetical reservoir exhibiting a formation resistivity factor - porosity function with a slope of m = 2. The locus of points with m = 2 is indicated by the heavy dashed line. The porosity end members have  $\varphi = 0.05$  (presumed finegrained) and  $\varphi = 0.40$  (presumed coarse-grained), respectively. These end members are circled. The direction of increasing  $\beta$  and the end points of the  $\beta$ scale are shown for reference. For a laminated sand built from the end members in various proportions, how does m vary? The lower (solid) curve shows the locus of points for the parallel component of the formation resistivity factor as the fractional volume of the components varies from 0 to 100 percent. The upper curve (dash-dot) is the locus of points for the perpendicular component of the formation resistivity factor as the fractional volume of the components varies from 0 to 100 percent. The dotted lines bound the envelope of m values. From the slopes of the dotted lines we find that  $1.6 \le m \le 4.0$ approximately.



 $\varphi_1 = 0.05, \ m_1 = 2.0$  $\varphi_2 = 0.40, \ m_2 = 2.0$ 

FIG. 5. A comparison of the porosity exponent m measured in the horizontal and vertical directions. Although the sand package is composited from individual sands with m=2, the composite has m values quite different from 2. The solid curve is for the component of m measured parallel to bedding planes, and the dashed curve is the component of m measured perpendicular to bedding planes. The average porosity is shown on the right scale. It is 5 percent on the right, 40 percent on the left, and varies linearly for intermediate composite compositions.

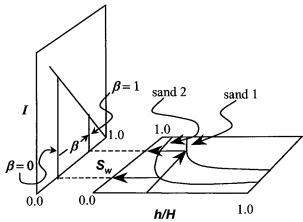


FIG. 7. The resistivity index experiment for a rock composited of two constituents having water saturations (as well as porosities) that differ. The water saturations are coupled through a capillary pressure relationship, and depend upon the height in the reservoir of the samples in question.

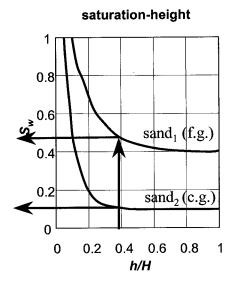


FIG 6. A saturation height function, plotted with the conventional axes exchanged. The height function is normalized by the thickness of an oil column measured from the free water level, plotted on the horizontal axis. Water saturation is plotted on the vertical axis. Note that  $S_{wlirr} = 0.40$  for the fine-grained (f.g.) component;  $S_{w2irr} = 0.10$  for the coarse-grained (c.g.) component. The illustration shows that for sand components with different capillary pressure curves, the same height above free water h/H in the reservoir corresponds to  $(S_{w1} = 0.50;$ different water saturations  $S_{w2} = 0.10$ ).

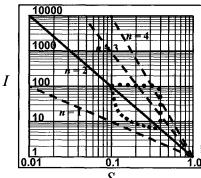


FIG. 8.a  $\varphi_1 = 0.10, m_1 = 2.0, n_1 = 2.0, S_{w1} = 0.40$  $\varphi_2 = 0.45, m_2 = 2.0, n_2 = 2.0, S_{w2} = 0.10$ 

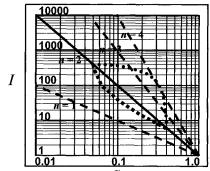


Fig. 8.c  $S_w$   $\varphi_1 = 0.44, m_1 = 2.0, n_1 = 2.0, S_{w1} = 0.40$  $\varphi_2 = 0.45, m_2 = 2.0, n_2 = 2.0, S_{w2} = 0.05$ 

# 

FIG. 8.b  $S_w$   $\varphi_1 = 0.44$ ,  $m_1 = 2.0$ ,  $n_1 = 2.0$ ,  $S_{w1} = 0.40$  $\varphi_2 = 0.45$ ,  $m_2 = 2.0$ ,  $n_2 = 2.0$ ,  $S_{w2} = 0.10$ 

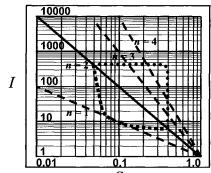


FIG. 8.d  $\varphi_1 = 0.10, \ m_1 = 2.0, \ n_1 = 2.0, \ S_{w1} = 0.40$  $\varphi_2 = 0.45, \ m_2 = 2.0, \ n_2 = 2.0, \ S_{w2} = 0.05$ 

# **Parallel Saturation Exponent**

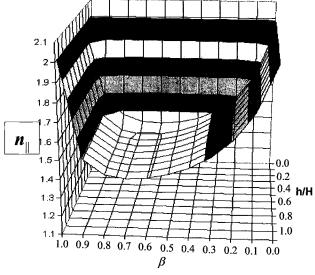


FIG. 9.a. The Archie saturation exponent for the composite medium measured in a direction parallel to bedding planes between the components of the composite. If one of the components retains a high water saturation while the other has most of its water displaced by hydrocarbons, the composite medium can have low saturation exponent values, once thought to be anomalous, but shown here to follow directly from the properties of the medium.

## **Perpendicular Saturation Exponent**

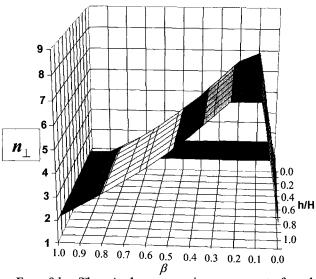


FIG. 9.b. The Archie saturation exponent for the composite medium measured in a direction perpendicular to bedding planes between the components of the composite. If one of the components retains a high water saturation while the other has most of its water displaced by hydrocarbons, the composite medium can have high saturation exponent values, once thought to be anomalous, but shown here to follow directly from the properties of the medium.

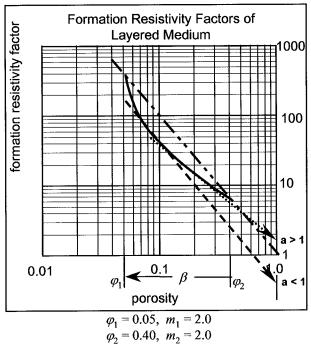


FIG. 10. The figure illustrates the effect of a limited range of porosity variation in a laminated reservoir. If the high porosity end member predominates, a least squares fit of a line to the data will have an intercept a > 1. Conversely, if the low porosity end member predominates, then the least squares line will have a < 1. These possibilities arising in a laminated sand reservoir suggest a physical cause for observations where  $a \ne 1$ .

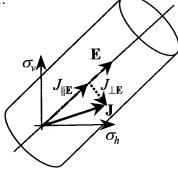


FIG. 12. In the unbounded medium there is no doubt that the current density **J** has components parallel  $(J_{\parallel E})$ and perpendicular  $(J_{\perp E})$  to the applied electric field E; i.e., linear superposition holds for the current density field. However, the  $J_{\perp E}$  component is constrained by the physical boundary of a core plug. Since the current density component perpendicular to the applied field cannot circulate, only the  $J_{\parallel E}$  component is non-zero, and the effective conductivity of the medium is given by the ratio  $J_{|\mathbf{E}|}/|\mathbf{E}|$  . This pretty picture is spoiled by end effects that are not amenable to easy analysis. I believe such effects will be small for core plugs with a large aspect ratio. End effects could not be neglected for short plugs, and for plugs having an aspect ratio near unity a resort to numerical methods would be necessary to determine the magnitude of the current traversing the plug, and consequently m and n.

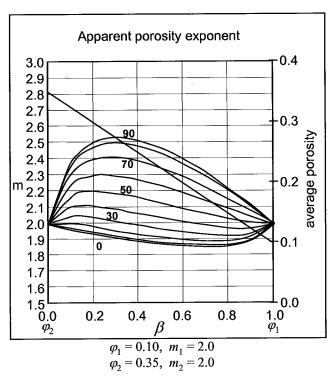


FIG. 11. The end member porosities used to develop the theory of Archie's parameters in a laminated, composite medium were at the extremes of the possible variations in porosity. In this figure a diminishing of the effect is recorded as end member porosities are chosen to lie more closely together. Archie's original data from the Nacatosh sandstone exhibits a variation of approximately  $0.10 < \varphi < 0.35$ .

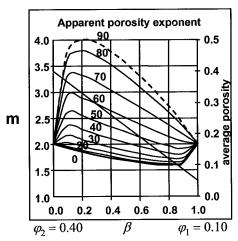


FIG. 13. If the value of the porosity exponent depends upon the relative fractional volumes of the constituents and whether the measurements are performed parallel to, or perpendicular to, the bedding planes, what is the effect of a misalignment of the core plug axis? The chart suggests that small misalignments (less than 10 degrees) will induce correspondingly small errors. The curves on this chart are not universally applicable but depend upon a number of assumptions explained in the text. The parameters in this model are the same as those used in figure 5.