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# Molar Enthalpy Fractal Dimension for Characterizing Shajara Reservoirs of the Permo-Carboniferous Shajara Formation

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## Abstract

The quality and assessment of a reservoir can be documented in details by the application of Molar enthalpy. This research aims to calculate fractal dimension from the relationship among Molar enthalpy, maximum Molar enthalpy and wetting phase saturation and to approve it by the fractal dimension derived from the relationship among capillary pressure and wetting phase saturation. Two equations for calculating the fractal dimensions have been employed. The first one describes the functional relationship between wetting phase saturation as a function of capillary pressure and the fractal dimension. The second equation implies to the wetting phase saturation as a function of capillary pressure and the fractal dimension. Two procedures for obtaining the fractal dimension have been utilized. The first procedure was done by plotting the logarithm of the ratio between Molar enthalpy and maximum Molar enthalpy versus logarithm wetting phase saturation. The slope of the first procedure = 3- Df (fractal dimension). The second procedure for obtaining the fractal dimension was determined by plotting the logarithm of capillary pressure versus the logarithm of wetting phase saturation. The slope of the second procedure = Df -3. On the basis of the obtained results of the fabricated stratigraphic column and the attained values of the fractal dimension, the sandstones of the Shajara reservoirs of the Shajara Formation were divided here into three units.

Keywords: Shajara Reservoirs; Shajara Formation; Molar enthalpy fractal dimension; Capillary pressure fractal dimension

## Introduction

Seismo electric effects related to electro kinetic potential, dielectric permitivity, pressure gradient, fluid viscosity, and electric conductivity was first reported by Frenkel. Capillary pressure follows the scaling law at low wetting phase saturation was reported by Li and Williams. Seismo electric phenomenon by considering electro kinetic coupling coefficient as a function of effective charge density, permeability, fluid viscosity and electric conductivity was reported by Revil and Jardani.

The magnitude of seismo electric current depends on porosity, pore size, zeta potential of the pore surfaces, and elastic properties of the matrix was investigated by Dukhin., *et al.* The tangent of the ratio of converted electic field to pressure is approximately in inverse proportion to permeability was studied by Guan., *et al.* Permeability inversion from seismoelectric log at low frequency was studied by Hu., *et al.* They reported that, the tangent of the ratio among electric excitation intensity and pressure field is a function of porosity, fluid viscosity, frequency, tortuosity, fluid density and Dracy permeability.

A decrease of seismo electric frequencies with increasing water content was reported by Borde., et al. An increase of seismo electric transfer function with increasing water saturation was studied by Jardani and Revil. An increase of dynamic seismo electric transfer function with decreasing fluid conductivity was described by Holzhauer., et al. The amplitude of seismo electric signal increases with increasing permeability which means that the seismo electric effects are directly related to the permeability and can be used to study the permeability of the reservoir was illustrated by Ping., et al. Seismo electric coupling is frequency dependent and decreases expontialy when frequency increases was demonstrated by Djuraev., et al. An increase of permeability with increasing pressure head and bubble pressure fractal dimension was reported by Alkhidir. An increase of geometric relaxation time of induced polarization fractal dimension with permeability increasing and grain size was described by Alkhidir.

### **Materials and Methods**

Sandstone samples were collected from the surface type section of the Permo-Carboniferous Shajara Formation, latitude 26° 52′ 17.4″, longitude 43° 36′ 18″. (Figure 1). Porosity was measured on collected samples using mercury intrusion Porosimetry and permeability was derived from capillary pressure data. The purpose of this paper is to obtain Molar enthalpy fractal dimension and to confirm it by capillary pressure fractal dimension. The fractal dimension of the first procedure is determined from the positive slope of the plot of logarithm of the ratio of Molar enthalpy to maximum Molar enthalpy log  $(ME^{1/4}/ME^{1/4}_{max})$  versus log wetting phase saturation (logSw). Whereas the fractal dimension of the second procedure is determined from the negative slope of the plot of logarithm of log capillary pressure (log Pc) versus logarithm of wetting phase saturation (log Sw).

The molar enthalpy can be scaled as

$$Sw = \left[\frac{ME^{\frac{1}{4}}}{ME^{\frac{1}{4}}_{max}}\right]^{[3-Df]}$$

1

2

Where Sw the water saturation, ME the molar enthalpy in Joule/ mole,  $ME_{max}$  the maximum molar enthalpy in Joule/mole, and Df the fractal dimension

Equation 1 can be proofed from

E=NOM\*ME

Where E the enthalpy in Joule, NOM number of moles, ME the molar enthalpy in Joule/mole

The number of moles can be scaled as

$$NOM = \left[\frac{M}{MM}\right]$$
<sup>3</sup>

Where NOM the number of moles, M the mass in kilo gram, MM the molar mass in kilo gram/mole

Insert equation 3 into equation 2

$$\mathbf{E} = \begin{bmatrix} \frac{\mathbf{M}\mathbf{E} * \mathbf{M}}{\mathbf{M}\mathbf{M}} \end{bmatrix}$$

The mass can be scaled as

$$\mathbf{M} = \begin{bmatrix} \mathbf{F} \\ \mathbf{g} \end{bmatrix}$$
 5

Where M the mass in kilo gram, F the force in Newton, g the acceleration in meter/square second

Insert equation 5 into equation 4

$$\mathbf{E} = \begin{bmatrix} \frac{\mathbf{M}\mathbf{E} * \mathbf{F}}{\mathbf{M}\mathbf{M} * \mathbf{g}} \end{bmatrix}$$
<sup>6</sup>

The acceleration g can be scaled as

$$\mathbf{g} = \begin{bmatrix} \mathbf{EF} \\ \mathbf{ETF} \end{bmatrix}$$

Where g the acceleration in meter/square second, EF the electric filed in volt/meter, ETF the electric transfer function in volt \* square second/square meter

Insert equation 7 into equation 6

$$\mathbf{E} = \begin{bmatrix} \frac{\mathbf{ME} * \mathbf{F} * \mathbf{ETF}}{\mathbf{MM} * \mathbf{EF}} \end{bmatrix}$$
8

The electric field EF can be scaled as

$$\mathbf{EF} = \begin{bmatrix} \mathbf{V} \\ \mathbf{CEK} \end{bmatrix} \qquad 9$$

Where EF the electric in volt/meter, V the velocity in meter/second, and CEK the electro kinetic coefficient in ampere/pascal \*meter

Insert equation 9 into equation 8

$$\mathbf{E} = \begin{bmatrix} \frac{\mathbf{ME} * \mathbf{F} * \mathbf{ETF} * \mathbf{CEK}}{\mathbf{MM} * \mathbf{V}} \end{bmatrix}$$
 10

The velocity V can be scaled as

$$\mathbf{V} = \begin{bmatrix} \mathbf{Q} \\ \mathbf{A} \end{bmatrix}$$

Where V the velocity in meter/second, Q the flow rate in cubic meter/second, and A the area in square meter

11

Insert equation 11 into equation 10

$$\mathbf{E} = \begin{bmatrix} \frac{\mathbf{ME} * \mathbf{F} * \mathbf{ETF} * \mathbf{CEK} * \mathbf{A}}{\mathbf{MM} * \mathbf{Q}} \end{bmatrix}$$
 12

The flow rate Q can be scaled as

$$\mathbf{Q} = \begin{bmatrix} \frac{3.14 * \mathbf{r}^4 * \Delta \mathbf{P}}{8 * \mu * \mathbf{L}} \end{bmatrix}$$
 13

Where Q the flow rate in cubic meter/second, r the pore radius in meter,  $\Delta P$  the differential pressure in pascal,  $\mu$  the fluid viscosity in pascal \* second, L the capillary length in meter

Insert equation 13 into equation 12

$$\mathbf{E} = \begin{bmatrix} \frac{\mathbf{ME} * \mathbf{F} * \mathbf{ETF} * \mathbf{CEK} * \mathbf{A} * \mathbf{8} * \boldsymbol{\mu} * \mathbf{L}}{\mathbf{MM} * \mathbf{Q} * \mathbf{3} \cdot \mathbf{14} * \mathbf{r}^4 * \Delta \mathbf{P}} \end{bmatrix} \quad \mathbf{14}$$

Equation 14 after rearrange of pore radius will become

$$r^{4} = \begin{bmatrix} \frac{ME * F * ETF * CEK * A * 8 * \mu * L}{MM * Q * 3.14 * E * \Delta P} \end{bmatrix} 15$$

The maximum pore radius can be scaled as

$$\mathbf{r}^{4}_{max} = \left[\frac{\mathbf{M}\mathbf{E}_{max} * \mathbf{F} * \mathbf{E}\mathbf{T}\mathbf{F} * \mathbf{C}\mathbf{E}\mathbf{K} * \mathbf{A} * \mathbf{B} * \mathbf{\mu} * \mathbf{L}}{\mathbf{M}\mathbf{M} * \mathbf{Q} * \mathbf{3} \cdot \mathbf{14} * \mathbf{E} * \Delta \mathbf{P}}\right] \quad \mathbf{16}$$

Divide equation 15 by equation 16

$$\left[\frac{r^{4}}{r_{max}^{4}}\right] = \left[\frac{\left[\frac{ME * F * ETF * CEK * A * 8 * \mu * L}{MM * Q * 3.14 * E * \Delta P}\right]}{\left[\frac{ME_{max} * F * ETF * CEK * A * 8 * \mu * L}{MM * Q * 3.14 * E * \Delta P}\right]}\right]$$
17

Equation 17 after simplification will become

$$\left[\frac{r^4}{r_{max}^4}\right] = \left[\frac{ME}{ME_{max}}\right] \qquad 18$$

Take the fourth root of equation 18

$$\sqrt[4]{\left[\frac{r^4}{r_{max}^4}\right]} = \sqrt[4]{\left[\frac{ME}{ME_{max}}\right]}$$
 19

Equation 19 after simplification will become

$$\left[\frac{\mathbf{r}}{\mathbf{r}_{\max}}\right] = \left[\frac{\mathbf{ME}_{4}^{1}}{\mathbf{ME}_{\max}^{1}}\right] \qquad 20$$

Take the logarithm of equation 20

$$log\left[\frac{r}{r_{max}}\right] = log\left[\frac{ME^{\frac{1}{4}}}{ME^{\frac{1}{4}}_{max}}\right] \qquad 21$$

But; log 
$$\left[\frac{\mathbf{r}}{\mathbf{r}_{\max}}\right] = \left[\frac{\log Sw}{3 - Df}\right]$$
 22

Insert equation 22 into equation 21

$$\log\left[\frac{\mathbf{r}}{\mathbf{r}_{\max}}\right] = \log\left[\frac{\mathbf{M}\mathbf{E}^{\frac{1}{4}}}{\mathbf{M}\mathbf{E}_{\max}^{\frac{1}{4}}}\right] \qquad 23$$

Equation 23 after log removal will become

$$Sw = \left[\frac{\frac{ME^{\frac{1}{4}}}{ME^{\frac{1}{4}}_{max}}}\right]^{[3-Df]} \qquad 24$$

Equation 24 the proof of equation 1 which relates the water saturation, molar enthalpy, maximum molar enthalpy and the fractal dimension

The capillary pressure can be scaled as

Where Sw the water saturation, Pc the capillary pressure and Df the fractal dimension.

## **Results and Discussion**

Based on field observation the Shajara Reservoirs of the Permo-Carboniferous Shajara Formation were divided here into three units as described in Figure 1. These units from bottom to top are: Lower Shajara Reservoir, Middle Shajara reservoir, and Upper Shajara Reservoir. Their attained results of the Molar enthalpy fractal dimension and capillary pressure fractal dimension are shown in Table 1. Based on the achieved results it was found that the Molar enthalpy fractal dimension is equal to the capillary pressure fractal dimension. The maximum value of the fractal dimension was found to be 2.7872 allocated to sample SJ13 from the Upper Shajara Reservoir as verified in Table 1. Whereas the minimum value of the fractal dimension 2.4379 was reported from sample SJ3 from the Lower Shajara reservoir as shown in Table 1. The Molar enthalpy fractal dimension and capillary pressure fractal dimension were detected to increase with increasing permeability as proofed in Table 1 owing to the possibility of having interconnected channels.

The Lower Shajara reservoir was symbolized by six sandstone samples (Figure 1), four of which label as SJ1, SJ2, SJ3 and SJ4 were carefully chosen for capillary pressure measurement as proven in Table1. Their positive slopes of the first procedure log of the Molar enthalpy to maximum Molar enthalpy versus log wetting phase saturation (Sw) and negative slopes of the second procedure log capillary pressure (Pc) versus log wetting phase saturation (Sw) are clarified in Figure 2, Figure 3, Figure 4, Figure 5 and Table 1. Their Molar enthalpy fractal dimension and capillary pressure fractal dimension values are revealed in Table 1. As we proceed from sample SJ2 to SJ3 a pronounced reduction in permeability due to compaction was described from 1955 md to 56 md which reflects decrease in Molar enthalpy fractal dimension from 2.7748 to 2.4379 as quantified in table 1. Again, an increase in grain size and permeability was proved from sample SI4 whose Molar enthalpy fractal dimension and capillary pressure fractal dimension was found to be 2.6843 as described in Table 1.

In contrast, the Middle Shajara reservoir which is separated from the Lower Shajara reservoir by an unconformity surface as revealed in Figure 1. It was nominated by four samples (Figure 1), three of which named as SJ7, SJ8, and SJ9 as illuminated in Table1 were chosen for capillary measurements as described in Table 1. Their positive slopes of the first procedure and negative slopes of the second procedure are shown in Figure 6, Figure 7 and Figure 8 and Table 1. Furthermore, their Molar enthalpy fractal dimensions and capillary pressure fractal dimensions show similarities as defined in Table 1.Their fractal dimensions are higher than those of samples SJ3 and SJ4 from the Lower Shajara Reservoir due to an increase in their permeability as explained in table 1.



**Figure 1:** Surface type section of the Shajara Reservoirs of the Permo-Carboniferous Shajara Formation at latitude 26° 52' 17.4" longitude 43° 36' 18".

Formation	Reservoir	Sample	Porosity %	K (md)	Positive slope of the first procedure Slope = 3-Df	Negative slope of the second procedure Slope = Df-3	Molar enthalpy fractal di- mension	Capillary pressure fractal di- mension
Permo-Car- boniferous Shajara Formation	Upper Shajara Reservoir	SJ13	25	973	0.2128	-0.2128	2.7872	2.7872
		SJ12	28	1440	0.2141	-0.2141	2.7859	2.7859
		SJ11	36	1197	0.2414	-0.2414	2.7586	2.7586
	Middle Shajara Reservoir	SJ9	31	1394	0.2214	-0.2214	2.7786	2.7786
		SJ8	32	1344	0.2248	-0.2248	2.7752	2.7752
		SJ7	35	1472	0.2317	-0.2317	2.7683	2.7683
	Lower Shajara Reservoir	SJ4	30	176	0.3157	-0.3157	2.6843	2.6843
		SJ3	34	56	0.5621	-0.5621	2.4379	2.4379
		SJ2	35	1955	0.2252	-0.2252	2.7748	2.7748
		SJ1	29	1680	0.2141	-0.2141	2.7859	2.7859

**Table 1:** Petrophysical model showing the three Shajara Reservoir Units with their corresponding values of Molar enthalpy fractal dimension and capillary pressure fractal dimension.





Figure 2: Log  $(ME^{1/4}/ME^{1/4}_{max})$  & log pc versus log Sw for sample SJ1. Figure 4: Log  $(ME^{1/4}/ME^{1/4}_{max})$  & log pc versus log Sw for sample SJ3



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Figure 6: Log  $(ME^{1/4}/ME^{1/4}_{max})$  & log pc versus log Sw for sample SJ7.



Figure 7: Log  $(ME^{1/4}/ME^{1/4}_{max})$  & log pc versus log Sw for sample SJ8.



Figure 8: Log  $(ME^{1/4}/ME^{1/4}_{max})$  & log pc versus log Sw for sample SJ9.

On the other hand, the Upper Shajara reservoir was separated from the Middle Shajara reservoir by yellow green mudstone as shown in Figure 1. It is defined by three samples so called SJ11, SJ12, SJ13 as explained in Table 1. Their positive slopes of the first procedure and negative slopes of the second procedure are displayed in Figure 9, Figure 10 and Figure 11 and Table 1. Moreover, their Molar enthalpy fractal dimension and capillary pressure fractal dimension are also higher than those of sample SJ3 and SJ4 from the Lower Shajara Reservoir due to an increase in their permeability as simplified in table 1.



Figure 9: Log  $(ME^{1/4}/ME^{1/4}_{max})$  & log pc versus log Sw for sample SJ11.



Figure 10: Log  $(ME^{1/4}/ME^{1/4}_{max})$  & log pc versus log Sw for sample SJ12.





Overall a plot of positive slope of the first procedure versus negative slope of the second procedure as described in Figure 12 reveals three permeable zones of varying Petrophysical properties. These reservoir zones were also confirmed by plotting Molar enthalpy fractal dimension versus capillary pressure fractal dimension as described in Figure 13. Such variation in fractal dimension can account for heterogeneity which is a key parameter in reservoir quality assessment.



Figure 12: Slope of the first procedure versus slope of the second procedure.



Figure 13: Molar enthalpy fractal dimension versus capillary pressure fractal dimension.

## Conclusion

The sandstones of the Shajara Reservoirs of the permo-Carboniferous Shajara Formation were divided here into three units based on Molar enthalpy fractal dimension. The Units from base to top are: Lower Shajara Molar Enthalpy Fractal Dimension Unit, Middle Shajara Molar Enthalpy Fractal Dimension Unit, and Upper Shajara Molar Enthalpy Fractal Dimension Unit. These units were also proved by capillary pressure fractal dimension. The fractal dimension was found to increase with increasing grain size and permeability owing to possibility of having interconnected channels.

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