

# Calculation the Irreducible water saturation $S_{wi}$ and determination Capillary pressure curve $P_C$ from Well Log data

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**Abstract.** Calculation of irreducible water saturation  $S_{wi}$  and determination Capillary pressure curve  $P_C$  is very important in the Oil and gas exploration and production. The complex reservoirs always represent a quite challenge to geologist and engineers to calculate  $S_{wi}$  [1]. Capillary pressure curves are usually determined in the laboratory in core analysis and only can perform when we known the irreducible water saturation  $S_{wi}$ . It is very difficult in the fact.

This research gives a method: Calculate  $S_{wi}$  and determine of  $P_C$  curve from obtain's data set  $(S_{wi}, P_C)_j$  with:  $j = 1 \dots p$  (which is easily to collect for every reservoirs). The method of this study can use for both the carbonate reservoirs and the Sandstone reservoirs. These reservoirs consist of 90% oil and gas in the world

The declared actual testing result from data in variety of diefferent huge oil fields around the world which are found on website and PVEP data has affirmed the appropriateness of this method.

## 1. Introduction

Calculation of irreducible water saturation  $S_{wi}$  is an important step in 3 D reservoir modeling studies. The irreducible water saturation  $S_{wi}$  distribution will dictate the original oil in place (STOIP) estimation and influence to the subsequent steps in establishment of dynamic modeling. The complex reservoirs always represent a quite

challenge to geologist and engineers to calculate  $S_{wi}$  [1].

The capillary phenomemna occurs in porous media when more immiscible fluids are present in the pore space [2]. In the interface between the two phases, Capillary pressure is defined as the difference in pressure between the wetting and nonwetting phases [2]:

$$P_c = P_{nw} - P_w \quad (1)$$

where  $P_w$  is wetting phase capillary and  $P_{nw}$  is non wetting phase capillary.

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Because the gravity forces are balanced by the Capillary forces, so that Capillary pressure at a point in the reservoir can be estimated from the height above the oil-water contact and the difference in fluid densities. For the oil-water media, we have:

$$P_c = (\rho_w - \rho_o)gh \quad (2)$$

Capillary forces are reflected by Capillary pressure curves affect the recovery efficiency of oil displaced by water, gas or different chemicals. thus, Capillary pressure functions are need for performing reservoir simulation studies of the different oil recovery processes. In [3] and more other studies suggest methods to plot  $P_c$  curves by empiricalism and analyse the relationships between it and other parameters.

Interpretation of Capillary pressure curves may yield useful informations regarding the petrophysical properties of rocks and the fluid rock interaction.. Relative permeability, absolute permeability and pore distribution to the nonwetting and wetting phases can obtain from the Capillary pressure curves.

This research gives a method: From obtain's data set  $(S_w; P_c)_j$  with:  $j=1...p$ , we calculate  $S_{wi}$  and determine function of  $P_c$  curve, determine three constants:  $S_{wi}$ ,  $a$ ;  $b$ .

The object of this study is both the carbonate reservoirs and the Sandstone reservoirs. These reservoirs consist 90% reservoir oil and gas in the world. Verification for both these reservoirs

The most important result of this study is calculation of the irreducible water saturation  $S_{wi}$  and plot  $P_c$  curve as the graph of a continuous function from  $(S_w; P_c)_j$  data, which is easily to collect in the fact. On the

other hand, the measurement of it in the laboratory by core analysis is very difficult, expensive and time-consuming.

The declared actual testing result from data in variety of different huge oil fields around the world which are found on website and PVEP data has affirmed the appropriateness of this method

*Nomenclature:*

$S_{wi}$ : irreducible water saturation

Well log data :

$P_c$ : Capillary pressure

## 2. The theoretical basic of the method

### 2.1. The empirical method:

Capillary pressure curves are usually determined in the laboratory in core analysis by the mercury injection method. The determination of Capillary pressure using reservoir fluids is usually done by the restored – state method or using a centrifuge, and according to the correlation coefficient to obtain the reservoir pressure [4].

### 2.2. The method of this study:

Capillary pressure curves are presented by the equation [5]:

$$P_c = \frac{a}{(S_w - S_{wi})^b} \quad (3)$$

Where :  $a, b, S_{wi}$  are three constants with  $0 < a < 1$  and  $0 < b < 2$

Formula (3) is established empirically and used for more studies. But the empirical establishment of this function is very difficult.

It requires to know the value  $S_{wi}$ , which is hard to implement and also sufficient measurement data is not simple to obtain in practice. Nevertheless, the measurement to gather data set from 8 to 10 values is feasible in every reservoir.

This research offers a method: From obtain's data set  $(S_w; P_c)_j$  with:  $j=1...p$ , we calculate  $S_{wi}$  and determine function of  $P_c$  curve, determine three constants:  $S_{wi}, a, b$ , and plot the graph of (3).

**3. Calculate the Irreducible water saturation  $S_{wi}$  and determination Capillary pressure curve  $P_c$  from Well Log data :**

The capillary pressure curves are represented by the equation

$$P_c = \frac{a}{(S_w - S_{wi})^b} \tag{3}$$

The problem is that: From the collection data:  $(S_w; P_c)_j$  with:  $j=1...p$  we calculate  $S_{wi}$ ,  $P_c$  curve, determine three constants:  $S_{wi}$  and  $a, b$ , plot the graph of function by (3).

Taking common logarithms of both sides of (3) gives:

$$\lg P_c = \lg a - b \lg(S_w - S_{wi}).$$

Denote:  $y = \lg P_c; A = -b; B = \lg a$  we have:  $y = Ax + B$ .

Consider  $S_{wi}$  is constant, perform the linear regression analysis to determine  $A$  and  $B$  (to infer  $a, b$ ).

In order to minimize the mean squared error :  $F = \sum_{j=1}^p [y_j - (Ax_j + B)]^2 \rightarrow \min$ .

Differentiate function  $F$  with respect to arguments  $A$  and  $B$  we have:

$$\begin{cases} \frac{\partial F}{\partial A} = -2 \sum_{j=1}^p [y_j - (Ax_j + B)] \cdot x_j = 0 \\ \frac{\partial F}{\partial B} = -2 \sum_{j=1}^p [y_j - (Ax_j + B)] = 0 \end{cases}$$

or: 
$$\begin{cases} (\sum_{j=1}^p 1) \cdot B + (\sum_{j=1}^p x_j) \cdot A = \sum_{j=1}^p y_j \\ (\sum_{j=1}^p x_j) \cdot B + (\sum_{j=1}^p x_j^2) \cdot A = \sum_{j=1}^p x_j y_j \end{cases}$$

in the matrix form:

$$\begin{pmatrix} \sum_{j=1}^p 1 & \sum_{j=1}^p x_j \\ \sum_{j=1}^p x_j & \sum_{j=1}^p x_j^2 \end{pmatrix} \cdot \begin{pmatrix} B \\ A \end{pmatrix} = \begin{pmatrix} \sum_{j=1}^p y_j \\ \sum_{j=1}^p x_j y_j \end{pmatrix} \dots \tag{4}$$

Using the liner regression method represents in [5] to solve equation (4), we find out  $A$  and  $B$ . The constants  $a, b$  are calculated as following:

$$a = 10^B; b = -A$$

Obviously that:  $S_{wi} \leq \min(VtS_w)$ , we give consecutively:  $S_{wi} = 0; \lambda, 2\lambda, \dots, (n-1)\lambda$  with  $\lambda = 0.0025$  and:  $n\lambda = \min(VtS_w)$

For every value :  $S_{wi} = 0; \lambda, 2\lambda, \dots, (n-1)\lambda$  we calculate  $F$ , then choose  $F$  min is the smallest value in the series  $n$  values of  $F$ , we determine immediately three constants:  $S_{wi}$  and  $a, b$  in (3). With 3 parameters  $S_{wi}$  and  $a, b$ , we plot the graph of (3)

Programming by the MATLAB language .

*Application's conditions*

Differentiate (3) we have :

$$P'_c = \frac{-ab}{(S_w - S_{wi})^{b+1}} < 0$$

so function  $P_c$  is

degenerated and non uniform continuous on  $(S_{wi}; 1]$ ; thus application's conditions is the collection data:  $(S_w; P_c)_j$  with:  $j=1...p$  must be unit value and monotono degreeing. It is mean that : Data  $(S_w; P_c)_j$  must satisfy condition:

$$\text{If } (S_w)_k < (S_w)_j, \text{ then } (P_c)_k > (P_c)_j.$$

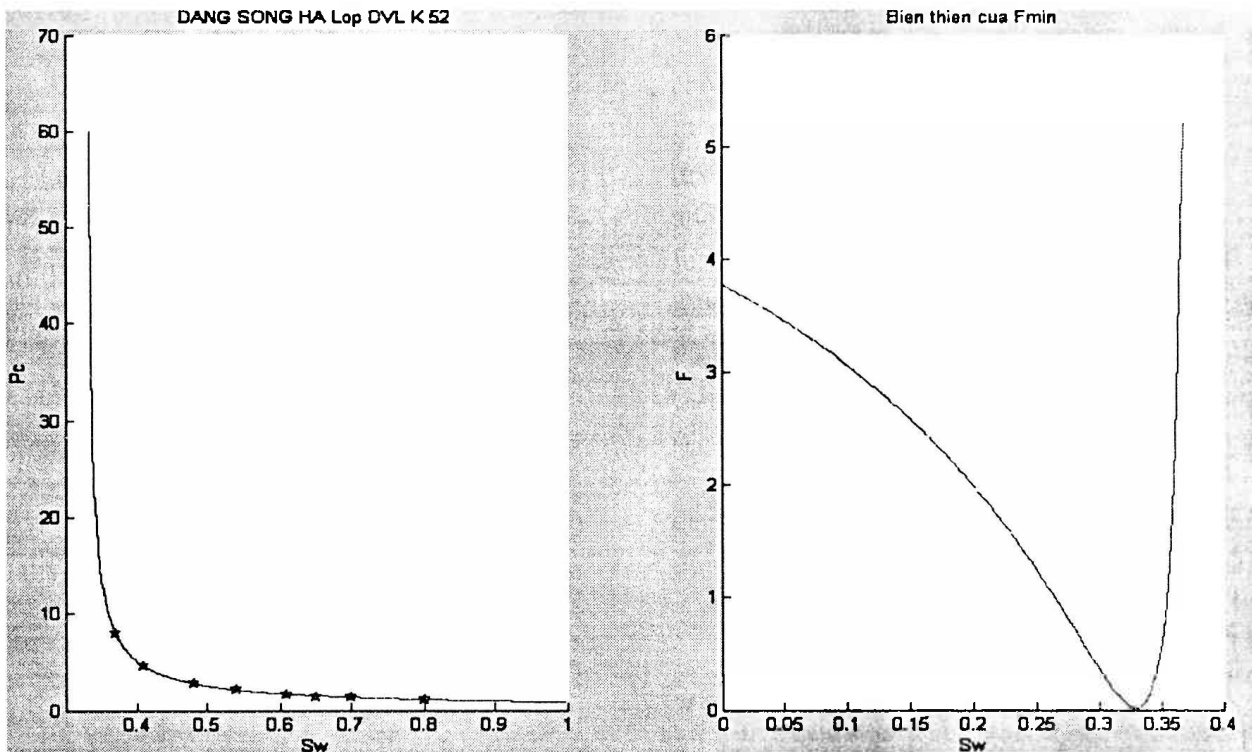
This condition is satisfified easily.

*Notice:*

1) The value  $S_{wi}$  calculates by this study usually smaller than the measured value a little,

becausee  $S_{wi}$  calculates while  $P_c \rightarrow +\infty$  and  $S_{wi}$  measured value with  $P_c$  only big enough.

2) The accuracy of the calculation result can be evaluated by analising and interpretation the constants  $a; b$  and  $S_{wi}$  (exemple: satisfy condition:  $0 < a < 1; 0 < b < 2$ ) and the variable behaviour of the function  $F$ . In the MATLAB Programming we plot the graph  $F-S_w$ . Consider the theory and testing on the practical data, we see that: The  $F$  curve reflects the accuracy of calculation result. The result is good if the minimum of function is reflected clearly., It is mean that function  $F$  decreases quickly to the minimum and increases quickly as the following figure (on the right):



*Verification:*

1) **On the Internet:** Consist of 6 Samples from the big oil and gas fields in the world. Reader can find them in [1,4,6] on the Internet.

Sample 1:

VtPc=[8.00 4.56 2.78 2.15 1.64 1.40 1.30 1.15];

VtSw=[0.37 0.41 0.48 0.54 0.61 0.65 0.70 0.80];

Sample 2:

VtPc=[0.867 1.16 1.45 1.73 2.02 2.31 2.89 7.8 21.7];

VtSw=[0.90 0.80 0.70 0.60 0.50 0.45 0.40 0.35 0.30];

Sample 3:

VtPc=[0.15 0.30 0.50 1 2 4 7 10];

VtSw=[1 0.982 0.883 0.771 0.698 0.664 0.648 0.639];

Sample 4:

VtPc=[0.15 0.30 0.50 1 2 4 7 10];

VtSw=[1 0.917 0.815 0.699 0.616 0.581 0.566 0.560];

Sample 5:

VtPc=[0.15 0.30 0.50 1 2 4 7 10];

VtSw=[0.984 0.942 0.868 0.786 0.72 0.687 0.673 0.665];

Sample 6:

VtPc=[0.15 0.30 0.50 1 2 4 7 10];

VtSw=[0.983 0.929 0.823 0.708 0.648 0.617 0.603 0.596];

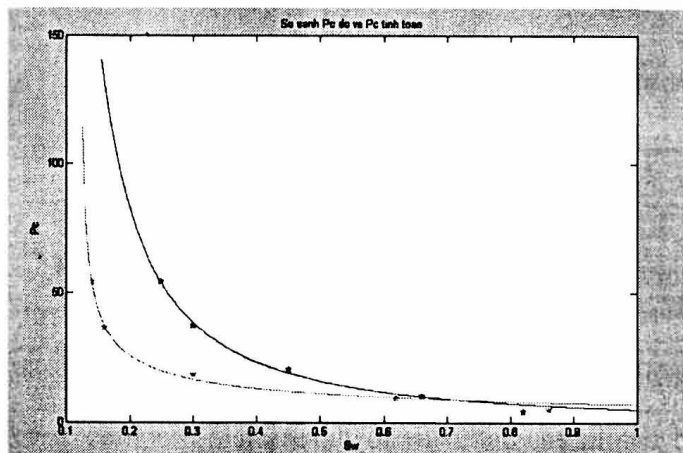
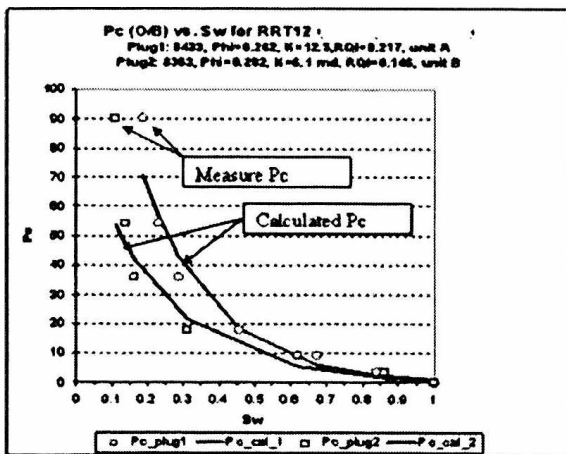
Calculation results from the samples:

Sample	Results in the Internet $S_{wi}$	Results of this study		
		$S_{wi}$	a	B
1	0.330	0.3300	0.5925	0.8085
2	0.300	0.2850	0.6012	0.8476
3	0.607	0.6050	0.0555	1.5359
4	0.539	0.5375	0.0803	1.2703
5	0.633	0.6325	0.0401	1.6138
6	0.575	0.5725	0.0631	1.3509

Comparison:

According to the data from “Calculations of Fluid Saturations from Log-Derived J-Functions in Giant Complex Middle-East

Carbonate Reservoir”[1] We calculate and compare: The result of this study is plotted by MATLAB on the right, the result of the author [4] on the left in the following figure:



#### 4. Results and Discussions

1) The fact that objective testing data in range of the reservoirs in the world which is found on website and data from PVEP, the appropriateness of parameter  $a$ ,  $b$  and variation of  $F$  curve is good has shown that: This research method is used for determining  $S_{wi}$  and  $P_c$  curve establishment. Good variation of  $F$  curve proves that mathematical basic about minimum condition of friction is its derivation equals zeros is entrusted theoretical foundation

2) The most important result of this study is the calculation of the irreducible water saturation  $S_{wi}$  from  $(S_w; P_c)_j$  data. The influence of  $S_{wi}$  on  $P_c$  is mentioned detailly in [3] and other documents, but none of them has mentioned about the calculation of  $S_{wi}$  from  $P_c$ . The calculation of  $S_{wi}$  from  $(S_w; P_c)_j$  data is reasonable. Value  $S_{wi}$  determines  $(S_w; P_c)$  then from  $(S_w; P_c)$ , value  $S_{wi}$  can be found out by solving reverse mathematical problem which is used frequently in geology

3) As well as other problem in Petrol and geology, the calculation for  $S_{wi}$  and establishment for curve  $P_c$  in this research should not be reviewed separately but analytical comparison  $S_{wi}$  of this study with other parameters as Permeability  $K$ , porosity  $\phi$  of the reservoir. Thus, this study is an approaching way together with other ones make a solution for significant as well as difficult problem in petrol exploration and geology investigation

#### 5. Conclusions

1. The empirical method only can plot the empirical capillary pressure curves but can not calculate irreducible water saturation  $S_{wi}$  and two parameters  $a, b$

2. The method of this study can calculate  $S_{wi}$ , two parameters  $a, b$  and plot the graph of (3). Not only  $S_{wi}$  but also two parameters  $a; b$  have their petrophysical meaning. In the reservoirs we usually have more data sets  $(S_w; P_c)_j$ , so we have more data sets  $\{S_{wi}; a; b\}$  respectively. Analysis, comparison data sets  $\{S_{wi}; a; b\}$  may yield useful informations regarding the reservoir

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#### References

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- [5] Noaman El-Khatib: *Development of a Modified Capillary Pressure J-Function*, KingSaud University
- [6] Crain's Petrophysical Handbook—Capillary pressure

## Appendix

### 1. Programm calculation for $S_{wi}$ and establishment for curve $P_c$

```

VtSw=[0.37 0.41 0.48 0.54 0.61 0.65 0.70 0.80]; %thay 2 dong nay khi can
VtPc=[8.00 4.56 2.78 2.15 1.64 1.40 1.30 1.15]; % thay 2 dong nay khi can
x=min(VtSw); l=round((x/0.0025));
p=length(VtSw);
Kqua =zeros(1,4); for n=1:l
    Swi=0.0025*(n-1);
    Kqua(n,1)=Swi;
    Hso=zeros(2,2);
    Nghiem=zeros(2,1);
    Tudo=zeros(2,1);
    for j=1:p
        Sw=VtSw(j);
        Pc=VtPc(j);
        Hso(1,2)=Hso(1,2)+log10(Sw-Swi);
        Hso(2,2)=Hso(2,2)+(log10(Sw-Swi))^2;
        Tudo(1,1)=Tudo(1,1)+log10(Pc);
        Tudo(2,1)=Tudo(2,1)+log10(Sw-Swi)*log10(Pc);
    end
    Hso(1,1)=p;
    Hso(2,1)=Hso(1,2);
    Nghiem=inv(Hso)*Tudo;
    a1=Nghiem(1,1); a=10^a1;    Kqua(n,2)=a;
    b1=Nghiem(2,1); b=-b1;    Kqua(n,3)=b;

    Fmin=0;
    for j=1:p
        Sw=VtSw(j);
        Pc=VtPc(j);
        Fmin= Fmin+[Pc-(a/((Sw-Swi)^b))]^2;
    end
    Kqua(n,4)= Fmin;
end

Fmin=Kqua(1,4);
for n=2:l
    x=Kqua(n,4);
    if (x<Fmin)
        Fmin=x;
    end
end
for n=1:l
    x=Kqua(n,4);
    if (x==Fmin)
        Swi=Kqua(n,1);
        a=Kqua(n,2);
        b=Kqua(n,3);
    end
end
end

```

```

% Hien thi
disp('Gia tri Swi ='); disp(Swi);
disp('          Bang Ket qua ');
disp('          Swi          a          b          Fmin'); disp(Kqua);
disp('Gia tri a =');disp(a);
disp('Gia tri b =');disp(b);
cach= (a/200)^(1/b);
dau= Swi+cach;
u=[dau:0.0025:1]; v=u;
m=length(u);
for j=1:m
    Sw= u(j);
    v(j)=a/((Sw-Swi)^b);
end
figure
subplot(1,2,1);
title('DANG SONG HA Dai hoc Mo Dia chat'); xlabel('Sw'); ylabel(' Pc ');
hold on :
plot(u,v, 'r');
hold on :
plot(VtSw,VtPc, 'pb');
hold on :

u=Kqua(:,1);v=Kqua(:,4);

subplot(1,2,2);
plot(u,v, 'b');
title('Bien thien cua Fmin');
xlabel('Saturation');ylabel('F');

```

## 2. Some data set

```

VtSw=[ 0.90    0.80    0.70    0.60    0.50    0.45    0.40    0.35    0.30]; %
1 VtPc=[ 0.867    1.16    1.45    1.73    2.02    2.31    2.89    7.8    21.7];
%0.578

VtSw=[1  0.904    0.782    0.640    0.518    0.478    0.447    0.409];
VtSw=[1  0.982    0.883    0.771    0.698    0.664    0.648    0.639 ];
Swi= 0.6050 a=0.0555 b=1.5359
VtSw=[1  0.917    0.815    0.699    0.616    0.581    0.566    0.560];
Swi=0.5375 a=0.0803 b=1.2703
VtSw=[0.983    0.929    0.823    0.708    0.648    0.617    0.603    0.596];%
Swi=0.5725 a=0.0631 b=1.35.9
VtSw=[1  0.967    0.808    0.706    0.568    0.517    0.483    0.438];
%VtSw=[0.952    0.930    0.865    0.740    0.633    0.594    0.555    0.499];
%VtPc=[ 2      4      8      15      35      70      120      200];

MtSw = [      1    0.982    0.883    0.771    0.698    0.664    0.648    0.639;
         1    0.917    0.815    0.699    0.616    0.581    0.566    0.560;
         0.984    0.942    0.868    0.786    0.72    0.687    0.673    0.665;
         0.983    0.929    0.823    0.708    0.648    0.617    0.603    0.596];];
VtPc = [ 0.15    0.30    0.50    1      2      4      7      10];

```