

TWO-PHASE WELLBORE SIMULATOR AND ANALYSIS OF REINJECTION  
DATA FROM SVARTSENGI, ICELAND

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ABSTRACT

A computer program is developed for the simulation of flowing geothermal wells. The necessary corrections of the salt and CO<sub>2</sub> concentrations on the thermophysical properties of the geothermal fluid are included in the program. The program is tested with the data from the Svartsengi geothermal field in Iceland and the Kizildere geothermal field in Turkey.

The response of the Svartsengi geothermal field to the injection is also investigated. A lumped model, called "the steam model" is used to analyze the effect of injection, and the actual data from the injection test at the Svartsengi field in 1984 is used for that purpose.



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NOMENCLATURE

A	: Cross-sectional area to flow, (m <sup>2</sup> )
A <sub>1</sub>	: Flow area in water zone, (m <sup>2</sup> )
A <sub>2</sub>	: Flow area in feed zone, (m <sup>2</sup> )
B	: Pressure drop factor for skin effect and long term drawdown, (Pa/(kg/s))
c	: Leakage constant for steam model (kg/sec/m)
C	: Turbulence factor, (Pa/(kg/s) <sup>2</sup> )
c <sub>m</sub>	: Specific heat of the rock matrix, (J/kgK)
D	: Diameter of pipe, (m)
(dp) <sub>t</sub>	: Total pressure drop, (Pa)
(dp) <sub>acc</sub>	: Pressure drop due to acceleration, (Pa)
(dp) <sub>fri</sub>	: Pressure drop due to friction, (Pa)
(dp) <sub>pot</sub>	: Pressure drop due to gravitational gradient, (Pa)
dz	: Incremental pipe length, (m)
E	: Energy of fluid, (J/s)
f	: Friction factor
G	: Mass flux, (kg/m <sup>2</sup> s)
g	: Acceleration due to gravity, (m/s <sup>2</sup> )
H	: Enthalpy, (kJ/kg)
h <sub>1</sub>	: Drawdown between two-phase and liquid phase interface, (m)
h <sub>2</sub>	: Drawdown in feed zone, (m)
k	: Permeability of the aquifer, (m <sup>2</sup> )
K	: Thermal conductivity of rock, (W/mK)
m	: Molality of the solution, (moles/kg)
n <sub>L</sub>	: Concentration of carbondioxide in liquid phase, (% by weight)
P	: Pressure, (Pa)
P <sub>S</sub>	: Steam saturation pressure, (Pa)
P <sub>wf</sub>	: Bottom hole flowing pressure, (Pa)
P	: Wellhead pressure, (Pa)
Q	: Heat loss to surroundings, (J/s/m)
q	: Amount of water flowing out of storage in the water zone, (kg/s)
r	: Radial distance, (m)
R	: Universal gas constant, (kJ/kmoleK)
Re	: Reynolds number
r <sub>w</sub>	: Wellbore radius, (m)
S <sub>1</sub>	: Storage in the liquid zone
S <sub>2</sub>	: Storage in the feed zone

T	: Temperature, ( $^{\circ}\text{C}$ )
t	: Time, (sec)
V	: Fluid velocity, (m/s)
W	: Mass flow rate, (kg/s)
x	: Steam quality, (%)
$z_a$	: Aquifer depth, (m)
$z^*$	: Flashing point in the well, (m)

Greek symbols:

$\rho$	: Fluid density, ( $\text{kg}/\text{m}^3$ )
$\rho_1$	: Density of fluid in water zone, ( $\text{kg}/\text{m}^3$ )
$\rho_2$	: Density of fluid in feed zone, ( $\text{kg}/\text{m}^3$ )
$\epsilon$	: Absolute roughness factor of the pipe, (m)
$\mu$	: Dynamic viscosity of fluid, (Pa s)
$\alpha$	: Void fraction
$\lambda$	: Slip factor
$\phi^2$	: Two-phase multiplier
$v$	: Specific volume of fluid, ( $\text{m}^3/\text{kg}$ )
$\kappa$	: Thermal diffusivity of the rock, ( $\text{m}^2/\text{sec}$ )

Subscripts

L	: Liquid phase
m	: Rock matrix
s	: Vapour phase



## 1 INTRODUCTION

### 1.1 Purpose of the study

This report is the final part of a 6 month training at the UNU Geothermal Training Programme, at the National Energy Authority in Reykjavik, Iceland in the summer of 1985.

The author's training programme can be divided into four main sections, namely introductory lectures in various disciplines of geothermal technology, lectures on reservoir engineering for the specialized training in this subject, two weeks field excursion and seminars on the various geothermal fields of Iceland and the final project on the subjects of simulation of flowing wells and response of the Svartsengi field to injection.

The experience and knowledge gained will be helpful to solve not only problems dealt with in this report but a wide range of problems associated with evaluation and utilization of geothermal energy in Turkey.

### 1.2 Statement of the problem

Among the several factors affecting the flow performance of the wells, diameter of the well as well as the reduction in the diameter due to calcite scaling and the drawdown in the reservoir can be mentioned. It is a common practice to develop simulators to study the effects of those factors. Because of the chemical composition of the geothermal fluid, the thermodynamic properties of pure water should be corrected for salt concentration. Another factor which should be taken into account in the two-phase pressure drop calculations is the effect of non-condensable gases.

Reinjection is increasingly becoming an integral part of the design of geothermal projects. The necessity of reinjection mainly arises from three factors, disposal of chemically hazardous geothermal waste water, possibility of pressure recovery of the declining aquifer and more heat extraction from the reservoir rock.

The main problems encountered in geothermal fields in Turkey are the reduction in the yields of the wells due to both scaling and drawdown in the reservoir, and the disposal of waste water. These two problems were selected to provide the writer with a useful experience and knowledge on the methods of solution.

## 2 FLOW PERFORMANCE CALCULATIONS IN GEOTHERMAL WELLS

### 2.1 Introduction

Good production wells in high temperature fields are characterized by high flowrates which make the downhole measurement of the flowing wells impractical. This difficulty makes simulation of the flowing wells important. Therefore, it is necessary to develop and test the flow models for geothermal wells by using limited data which exists for low flowrates.

Wells in high temperature, liquid dominated geothermal reservoirs generally produce two-phase mixtures. In the case of liquid water inflow to the wellbore, the drop in pressure during upflow results in flashing of the fluid in the wellbore. Since the thermodynamic properties of a brine change with the salt concentration, it is necessary to correlate the properties of fluid with respect to salt concentration. One of the constituents in geothermal fluid is non-condensable gas, especially  $\text{CO}_2$ , which causes an increase in the flashing pressure. To analyze the behaviour of flowing wells, a wellbore simulator was developed which takes the effects of salt and  $\text{CO}_2$  concentration into account. This simulator was used to estimate the deliverability curves for two wells in the Svartsengi geothermal field in Iceland. The effect of  $\text{CO}_2$  was studied by using data from the Kizildere geothermal field in Turkey. As a final application, the changes in the output of a well as a result of calcite deposition was also studied.

In the following sections, the derivation of flow equations and corrections for salt concentration and  $\text{CO}_2$  are presented.

## 2.2 Derivation of the equations for fluid flow in a vertical pipe

To describe the fluid flow in a vertical pipe, one can use the equations of conservation of mass, momentum and energy. During the following derivation of equations it is assumed that the flow is homogeneous, steady and one-dimensional.

- Equation for conservation of mass:

$$W = \rho VA \quad (1)$$

- Equation for conservation of momentum:

This equation is generally written from a pressure drop point of view. Total pressure drop is made up of three individual gradients,

$$-(dP)_t = (dP)_{fric} + (dP)_{acc} + (dP)_{pot} \quad (2)$$

Let us define each gradient separately.

### i) Frictional pressure drop

This pressure drop is defined by the Darcy-Weisbach equation as;

$$(dP)_{fric} = \frac{\rho f V^2}{2D} dz \quad (3)$$

The friction factor "f" is a function of the Reynolds Number,  $Re$ , and the relative roughness of pipe and is given by the modified Colebrook's equation as;

$$f = \left[ \left[ -2 \log \left( \frac{\epsilon}{3.7D} \right) + \left( \frac{7}{Re} \right)^{0.9} \right]^2 \right]^{-1} \quad (4)$$

where

$$Re = \frac{\rho VD}{\mu} \quad (5)$$

ii) Acceleration pressure drop

$$(dP)_{acc} = \rho V dV \quad (6)$$

If the mass flux is defined as  $G = W/A$ , and from Eq. 1,  $V = G/\rho$ , then the acceleration pressure drop can be expressed as;

$$(dP)_{acc} = G dV \quad (7)$$

iii) Potential pressure drop

This pressure drop is defined as;

$$(dP)_{pot} = \rho g dZ \quad (8)$$

By using equations 3,7 and 8 the total pressure drop can be written as;

$$-(dP)_t = \frac{\rho f V^2}{2D} dZ + G dV + \rho g dZ \quad (9)$$

- Energy equation:

The total energy equation can be expressed as;

$$dE = W dH + W d \left( \frac{V^2}{2} \right) - W g dZ - Q dZ \quad (10)$$

since there is no energy input for a self-flowing well,  $E=0$ , equation 10 becomes;

$$0 = dH + d \left( \frac{V^2}{2} \right) + g dZ - \frac{Q}{W} dZ \quad (11)$$

The total enthalpy is a function of the enthalpies of each phase,  $H_1$  and  $H_g$ , and the flowing steam quality,  $x$ .

$$H = x H_g + (1-x) H_1 \quad (12)$$



Integration of equation 11 between any two points in the well in an upward direction gives;

$$H_2 = H_1 - 0.5(V_2^2 - V_1^2) - g(Z_2 - Z_1) + \frac{Q}{W} (Z_2 - Z_1) \quad (13)$$

By introducing equation 12, the steam quality at point 2 can be found as;

$$X_2 = \frac{H_1 - 0.5(V_2^2 - V_1^2) - g(Z_2 - Z_1) - H_{1,2} + (Q/W)(Z_2 - Z_1)}{H_{s,2} - H_{1,2}} \quad (14)$$

The heat loss to the surrounding,  $Q$ , will be discussed in a later section.

It is known that a liquid-dominated geothermal reservoir will initially produce undersaturated water at the wellbore sand-face. During the upflow of the fluid along the wellbore, the drop in pressure results in saturated water and the flow becomes a two-phase flow. In order to examine these two different sections, namely single-phase and two-phase, it is necessary to study the above equations at each section separately.

### 2.2.1 Single-phase region

In this section, the fluid density is almost constant which corresponds to the density of the inflow temperature. This constant density results in a constant velocity,  $V_1 = V_2$ , then according to equation 7  $(dP)_{acc} = 0$ .

Now, total pressure drop for single-phase region is reduced to;

$$-(dP) = \frac{\rho f V^2}{2D} dZ + \rho g dZ \quad (15)$$

### 2.2.2 Two-phase region

When the fluid starts flashing in the wellbore, it undergoes different two-phase flow regimes, namely bubble, slug and annular. These flow regimes have been studied by several authors (Ros, 1961; Hagedorn, 1964; Orkiszewski, 1967). In these flow regimes, the vapour and liquid phases travel separately at different velocities. Since the vapour always "slips" past the liquid in vertical flow, equation 9 has to be corrected.

In general the mixture density is presented in terms of actual void fraction defined from the volume occupied by the vapour phase,

$$\rho = \rho_s \alpha + \rho_l (1-\alpha) \quad (16)$$

where

$$\alpha = \frac{A_s}{A} \quad \text{or} \quad (1-\alpha) = \frac{A_l}{A} \quad (17)$$

On the other hand the slip factor,  $\lambda$ , is defined as the velocity ratio,

$$\lambda = \frac{V_s}{V_l} \quad (18)$$

Another important definition for the two-phase region is the steam quality,  $x$ .

$$x = \frac{W_s}{W} \quad \text{or} \quad (1-x) = \frac{W_l}{W} \quad (19)$$

Now, let us define individual pressure drops in the two-phase region.

i) Potential pressure drop

By substituting equation 17 into equation 8

$$(dP)_{pot} = [\alpha\rho_s + (1-\alpha)\rho_l]gdZ \quad (20)$$

ii) Acceleration pressure drop

From the equation for conservation of mass, equation 1, the velocities of each phase can be written as;

$$V = \frac{xW}{\rho_s A_s} \quad (21)$$

and

$$V = \frac{(1-x)W}{\rho_l A_l} \quad (22)$$

On the other hand, the acceleration pressure drop in two-phase region is defined as;

$$(dP)_{acc} = \frac{1}{A} d(W_l V_l + W_s V_s) \quad (23)$$

By substituting the definitions of velocities from equations 21 and 22

$$(dP)_{acc} = \frac{1}{A} d\left(\frac{WW_l(1-x)}{\rho_l A_l} + \frac{WW_s x}{\rho_s A_s}\right) \quad (24)$$

By making further substitutions for rates of mass flow and cross-sectional areas for each phase from equations 17 and 19 the final equation is obtained as;

$$(dP)_{acc} = G^2 d\left(\frac{x^2}{\alpha\rho_s} + \frac{(1-x)^2}{(1-\alpha)\rho_l}\right) \quad (25)$$



iii) Frictional pressure drop

If we introduce the definition of mixture density, equation 16, into equation 3 the frictional pressure drop for the two-phase region can be obtained.

$$(dP)_{ftp} = \frac{fV^2(\alpha\rho_s + (1-\alpha)\rho_l)}{2D}dZ \quad (26)$$

But the difficulty arises here in the evaluation of the two-phase friction factor and velocity of the fluid, because of the definition of the two-phase viscosity and the difference between the velocities of phases due to slip. In order to overcome this difficulty several empirical correlations have been studied.

Martinelli-Nelson (1948) defined an empirical relation to calculate the friction pressure gradient by assuming a ratio, called two-phase multiplier.

$$(dP/dZ)_{ftp} = \phi^2_{lors} \cdot (dP/dZ)_{lors} \quad (27)$$

where  $(dP/dz)_{ftp}$  is the two-phase frictional pressure gradient,  $(dP/dz)_l$  and  $(dP/dz)_s$  are the frictional pressure gradients for the liquid or gas respectively if they are flowing alone in the same tube.

Other correlating parameters have been defined if the total mass is flowing with the physical properties of one of the phases.

$$(dP/dZ)_{ftp} = \phi^2_{lors0} \cdot (dP/dZ)_{lors0} \quad (28)$$

where  $(dP/dz)_{l0}$  and  $(dP/dz)_{s0}$  are the pressure gradients for the total flow of fluid having the liquid or gas physical properties respectively.

In this correlation, frictional pressure drop for the single phase is obtained by assuming that all the flowing fluid is in this phase, then this single phase pressure drop is multiplied by the two-phase multiplier to obtain the frictional pressure drop in the two-phase flow.

In the literature, there are several correlations for a two-phase multiplier. In this report, the correlation given by Chisholm (1972) is used.

$$\phi^2 = 1 + \frac{C}{X} + \frac{1}{X^2} \quad (29)$$

where

$$X^2 = \left( \frac{1-x^2}{x} \right) \cdot \frac{v_l}{v_g} \quad (30)$$

and

$$C = 1 + \frac{xv_g}{xv_g + (1-x)v_l} - \alpha \quad (31)$$

As can be seen in equation 31 the void fraction,  $\alpha$  should be known to calculate the two-phase multiplier. In order to calculate the void fraction the correlation by Armand and Teacher (1959) is used.

$$\alpha = \frac{0.833 + 0.05 \log(P)}{1 + \frac{(1-x)v_l}{xv_g}} \quad P \text{ in bar} \quad (32)$$

This concludes the equations for pressure drop in the two-phase section. The location of the flashing point can be calculated as follows.

$$-(dP)_t = (P_{wf} - P_s) = (dP)_{fri} + (dP)_{pot} \quad (33)$$

From equation 15, by substituting  $dZ = Z_a - Z^*$

$$-(dP)_t = \left( \frac{\rho_f V^2}{2D} + \rho g \right) (Z_a - Z^*) \quad (34)$$

$$Z^* = Z_a - \frac{P_{wf} - P_s}{\rho g + \frac{\rho_f V^2}{2D}} \quad (35)$$

But, the high flowrates and temperatures in geothermal wells, make it difficult to run downhole measurements under normal operating conditions. Therefore, in order to get the value of the bottom hole flowing pressure,  $P_{wf}$ , we should find a relationship between the conditions of static well and flowing well.

During the flow of fluid in the reservoir, the pressure drops from undisturbed reservoir pressure,  $P_a$ , to the bottom hole flowing pressure,  $P_{wf}$ , at the sand-face. This pressure drop in the reservoir is due to the skin effect, the longterm drawdown factor, and the turbulence pressure drop. (Kjaran, 1983) Therefore,  $P_{wf}$  can be written as;

$$P_{wf} = P_a - (BW + CW^2) \quad (36)$$

where B and C are some factors.

In the high flowrate geothermal wells, initially the pressure drop due to turbulence is the dominant factor and equation 36 can be rewritten as;

$$P_{wf} = P_a - CW^2 \quad (37)$$

Since we have some measurements of  $P_{wf}$  for the low flowrates, equation 39 can be solved analytically by taking the slope of  $(P_a - P_{wf})/W$  vs.  $W$  graph as turbulence factor C.

Another way of obtaining the C value is the use of simulators to match the deliverability measurements of the wells since the wellhead pressure is given by,

$$P_o = P_{wf} - (dP)_t = P_a - CW^2 - (dP)_t \quad (38)$$

In this procedure,  $P_a$  is an input and C is the variable to match the measured wellhead pressure for a given flowrate.

### 2.3 Wellbore heat transfer

During the flow of fluid in the wellbore, some heat is transferred by conduction through the rock surrounding the wellbore. This heat transfer to the surrounding has been

studied by Ramey (1962). By assuming that the conductive heat flow is normal to the axis of the well, the equation for heat conduction can be written as;

$$K \frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial T}{\partial r} \right) = \rho_m C_m \frac{\partial T}{\partial t} \quad (39)$$

The boundary conditions are as follows,

$$\begin{aligned} T &\rightarrow T_r & (r &\rightarrow \infty), \\ T &= T_w & (r &= r_w) \end{aligned}$$

where  $T_r$  = undisturbed reservoir temperature, ( $^{\circ}$ K)  
 $T$  = temperature at the well face, ( $^{\circ}$ K)

The conductive heat transfer per unit length of the wellbore is given by;

$$Q = 2\pi r_w K \frac{(T_r - T_w)}{f(t)} \quad (40)$$

where  $f(t)$  is defined as;

$$f(t) = \ln \left( \frac{2\sqrt{Kt}}{r_w} \right) - 0.29 \quad (41)$$

In fact, the boundary condition given for undisturbed reservoir temperature occurs at a certain distance from the wellbore instead of infinity. By introducing this new boundary condition, the solution of the problem becomes more complicated and is given by Carslaw and Jaeger (1959).

#### 2.4 Thermodynamic properties of geothermal fluids

Geothermal fluids are solutions of various types of salts. These are primarily NaCl, KCl, and CaCl<sub>2</sub>, but it is a common practice to use the "equivalent NaCl content" to define the salinity of a geothermal fluid. The "equivalent NaCl content" is the amount of NaCl in solution that will bring the same effect on the properties as the amount of all the salts combined.



For the modelling of the geothermal well, it is necessary to know some thermophysical properties accurately, such as the density, the enthalpy, the entropy, the viscosity and the saturation pressure at a given temperature. To make the corrections on the pure water properties, Michaelides (1981) presented some formulas which are based on "equivalent NaCl content".

One of the effective changes in the properties of pure water with salt content is the depression of saturation pressure at a given temperature,

$$\Delta P = \frac{1.8R(T + 273.15)}{v_s - v_l} \cdot \frac{m}{55.56} \quad \text{kPa} \quad (42)$$

where  $R = (8.314/18) = 0.4619$  kJ/kgK, 55.56 is the moles of water in 1 kg of the substance

$$\text{and } m = \frac{\text{Salinity of water in ppm}}{\text{Molecular weight of NaCl}}$$

The saturation pressure of pure water is given by the following simplified correlation;

$$P(T) = \exp(0.21913E-6T^3 - 0.17816E-3T^2 + 0.0653665T - 4.96087) \quad (43)$$

In his paper Michaelides gives the gas constant as 8.314 kJ/kgK for equation 42, but if the unit analysis is done carefully, it is found that the actual gas constant is the ratio of universal gas constant (8.314 kJ/kmoleK) to the molecular weight of water (18 kg/kmole). The reader should give attention to this change in the formula.

Let us take an example to show the effect of salt on the depression of saturation pressure. If we take the common values from the Svartsengi geothermal field in Iceland,

$$T = 240^\circ\text{C} \text{ and salinity} = 20.000 \text{ ppm}$$

by using equation 43,  $P(T) = 32.93$  bar

$$m = \frac{20.000 \text{ mg/kg}}{58.500 \text{ mg/mol}} = 0.342 \text{ mol/kg}$$

and  $v_s = 0.0595 \text{ m}^3/\text{kg}$ ,  $v_l = 0.0012 \text{ m}^3/\text{kg}$  from the steam tables for pure water.

By substituting the above values into equation 42

$$\Delta P = 45.05 \text{ kPa} = 0.45 \text{ bar}$$

The true saturation pressure therefore reduces to,

$$P = P(T) - \Delta P = 32.93 - 0.45 = 32.48 \text{ bar}$$

For the correlations of density, enthalpy, viscosity, entropy and elevation of saturation temperature, the reader is referred to Michaelides (1981).

### 2.5 Effect of CO<sub>2</sub>

CO<sub>2</sub> provides an important component to the total pressure of geothermal fluid. Presence of CO<sub>2</sub> causes the transition between single-phase and two-phase flow to happen deeper in the wellbore than one would expect by ignoring CO<sub>2</sub>.

The pressure of the vapour phase is the sum of steam and gas partial pressures,

$$P = P_s + P_g \quad (44)$$

The partial pressure of gas can be expressed in terms of the concentration of gas in liquid phase and solubility. The solubility data for CO<sub>2</sub> was fitted to an equation by Sutton (1976).

$$\alpha(T) = \{5.4 - 3.5(T/100) + 1.2(T/100)^2\}E-9, \text{ (1/Pa)} \quad (45)$$

The partial pressure of CO<sub>2</sub> is given by the following formula,

$$P_g = \frac{n_l}{\alpha(T)} \quad (45)$$

In order to show the effect of CO<sub>2</sub> on the flashing pressure of the fluid, let us take an example from the Kizildere geothermal field in Turkey.

The well number KD-6 is producing 15% CO<sub>2</sub> by weight in the steam phase and the wellhead steam quality is measured as 9.47%. The inflow temperature of fluid is 201°C.

In order to get the flashing pressure, the steam saturation pressure and CO<sub>2</sub> partial pressure should be calculated separately.

By using equation 45 the steam pressure can be found.

$$P_s(T) = 15.79 \text{ bar}$$

Now, first of all we should convert the wellhead CO<sub>2</sub> measurement to downhole concentration. By assuming all the CO<sub>2</sub> was in the vapour phase at the wellhead,

$$n_1 = 0.15 \cdot 9.47 = 1.42 \% \text{ by weight}$$

By using equation 45,  $\alpha(T) = 3.21E-9 \text{ 1/Pa}$

$$P_g = \frac{0.0142}{3.21E-9} = 4.42E6, \quad P_a = 44.2 \text{ bar}$$

Therefore, the flashing pressure of this system is;

$$P = 15.8 + 44.2 = 60 \text{ bar}$$

which is far beyond the steam saturation pressure.

After flashing, the CO<sub>2</sub> pressure in the vapour diminishes from its initial value as flashing progresses. Most of the mass of CO<sub>2</sub> is exsolved in the first few weight percent of flashing. Michels (1981) developed a relationship for the CO<sub>2</sub> pressure in a developing vapour phase.

The partial pressure of CO<sub>2</sub> is given accurately enough by the real gas equation,  $PV = znRT$ , and the final form can be expressed as a function of three factors, namely initial

CO<sub>2</sub> partial pressure, the present temperature of the system, and the weight percent of flashing that has developed.

$$P_c = P_c(0) \left\{ 1 + \frac{44 v_{sx}}{\alpha(T)R(T+273.15)} \right\}^{-1} \quad (47)$$

where  $P_c(0)$  = Initial CO<sub>2</sub> partial pressure and 44 is molecular weight of CO<sub>2</sub>.

## 2.6 Applications of the program

The developed computer program can find the main applications in the following subjects:

- i. Prediction of flowing well pressure and temperature profiles.
- ii. Development of deliverability curves for a certain well for different aquifer pressures.
- iii. Determination of the effect of salt and CO<sub>2</sub> concentration on the flashing point of the fluid.
- iv. Determination of the effect of calcite deposition and plugging on production.

These four items were analyzed for two wells in the Svartsengi field in Iceland and one example is given for the effect of CO<sub>2</sub> from the Kizildere field in Turkey. During the application of the computer program, the relative roughness coefficients of the slotted liner, casing and calcite deposition were taken as 1.37 E-4, 4.57 E-5 and 3.047 E-4 m respectively.

The listing of the program is presented in Appendix A.



### 2.6.1 Svartsengi Well No. 4

This well was initially drilled to 1713 m depth, and it was used as a production well until a casing damage occurred in the well in 1980. Presently it is used as an observation well.

The depth of the aquifer is found from the temperature profiles as 1024 m, initial temperature and pressure values were 240°C and 88 bars respectively.

During the application of the computer program, the deliverability curves of the well, with or without calcite deposition, and the pressure profile of the well were obtained. These results are compared with the actual measurements. The output measurements for well 4 in Svartsengi are given in Table 1. As can be seen from Fig. 1, the deliverability curves for the cases with or without calcite are in good agreement with the measured ones. This fit was obtained by using the turbulence coefficient as  $0.0035 \text{ bar}/(\text{kg/s})^2$ .

TABLE 1: Svartsengi well no 4 output measurements

WHP(bara)	Flow(kg/s)	WHP(bara)	Flow(kg/s)
Without calcite		With calcite	
19.0	33.0	18.9	28.0
17.1	59.0	16.8	39.0
14.0	75.0	13.5	49.0
13.2	80.0	11.2	50.0
11.7	85.0	9.2	51.0
		8.9	52.0

Data derived from Fig. 2 was used to see the effect of calcite deposition. The length of deposition was taken as 60 m between the interval 350-410 m and the best fit was obtained by using 11.5 cm as the average diameter of the calcite section. The calcite deposition causes a very high pressure drop in high flowrates and the change in deliverability curve is very abrupt at a particular flowrate. In Svartsengi 4, this flowrate is found as 51 kg/s and with a flowrate higher than this value the flow is choked.

The measured and calculated pressure profiles were also compared at Fig. 3. The calculated pressure profile at 30 kg/s flowrates gives a good fit with the measured one.

As a final application, mass flowrate vs. flashing depth from wellhead was plotted, and the caliper log of the well, Fig. 2, was compared with this plot (Fig. 4). By taking the flashing depth as 410 m from Fig. 2, the corresponding flowrate is found as 31 kg/s from Fig. 4 which was the actual average flowrate of the well before caliper log.

The computer outputs for Svartsengi 4 are given in Appendix B.

#### 2.6.2 Svartsengi Well No. 11

This well was used as an example for the effect of reservoir pressure and casing size on the deliverability curves. The family curves for different reservoir pressures are shown on Fig. 5. The pressure drawdown causes a reduction in the wellhead pressure but this reduction is not of the same order of magnitude as the reduction of reservoir pressure. On the other hand, the shape of the deliverability curves are almost the same for all reservoir pressures, therefore it is possible to find a common formula depending on the reservoir pressure, wellhead pressure and mass flowrate by knowing only one deliverability curve for a reservoir pressure.

As a part of this study, three production tests were carried out and the data is used in the program. The lip pressure method developed by James (1970) was used to test the well. A V-notch was used to measure the low mass flowrates and the enthalpy of fluid is calculated from James's formula which relates mass flow, discharge pipe area, enthalpy of fluid, and lip pressure. For a higher mass flowrate, the well was discharged vertically, direct to the atmosphere. The enthalpy value obtained from low flowrates is used to get the flowrate from James's formula. The results of the test are given in Fig. 5. Since there is about 12 bar pressure reduction in the field from the initial reservoir pressure, due to the production, the

reservoir pressure is taken as 76 bar. The best fit is obtained by using zero turbulence factor.

The effect of casing size on the output curves was also studied and the results obtained for two different wells are plotted in Fig. 6. The main effect of the casing size on the output characteristics of the wells is the decrease in the turbulence pressure drop in the wider wells. The deliverability curve for Svartsengi 11 was obtained by using zero turbulence factor and as can be seen from Fig. 6, the change in flowrate does not affect the wellhead pressure in the wider well as in the narrower well.

### 2.6.3 Kizildere Well No. 6

To see the effect of CO<sub>2</sub> on the output curves, data from Kizildere geothermal field in Turkey was used. The Kizildere field is a hot-water type geothermal field which is producing through six wells. One of the important problems in this field is CaCO<sub>3</sub> scaling in wells which is related not only to the calcium but also to the CO<sub>2</sub> content of the geothermal fluid. High percentage of CO<sub>2</sub> in the geothermal fluid of Kizildere causes early flashing of the fluid because of higher CO<sub>2</sub> partial pressure. The data from Kizildere Well no 6 (KD-6) was used in the program and the flashing depth and the wellhead pressure were examined. The actual data (Tan, 1982) and computer output for this well are given in Table 2. Fig. 7 shows the configuration of casing string and flashing point.

By using zero turbulence pressure drop, the wellhead pressure was obtained as 3.14 bar and the flashing point is at 576 m which shows a good agreement with the measurements.



TABLE 2: Data and computer output for Kizildere well No. 6

NAME OF FIELD: KIZILDERE  
 WELL NAME : RD-6  
 DATE OF CALC : 17-09-85

## SINGLE-PHASE(WATER) SECTION:

Pa (BARS) : 84.000  
 (dp)turb (BARS) : 0.000  
 Pwf (BARS) : 84.000  
 CCO2 (ppm) : 14204.0  
 CNaCl (ppm) : 1200.0

DEPTH(m)	P(bar)	Vcl(f)	Ffactor	D(cm)	DPWAT.	DPOT	DPFRIC	W(kg/s)	T(C)	PFLASH
800.000	79.594	3.034	0.018	21.590	-4.406	-4.242	-0.164	96.00	201.0	59.985
700.000	70.782	3.034	0.018	21.590	-8.812	-8.485	-0.328	96.00	201.0	59.985
665.600	67.750	3.034	0.018	21.590	-3.031	-2.919	-0.113	96.00	201.0	59.985
665.600	67.752	2.366	0.014	24.448	0.002	0.000	0.000	96.00	201.0	59.985
600.000	62.096	2.366	0.014	24.448	-5.656	-5.566	-0.090	96.00	201.0	59.985

## TWO PHASE SECTION:

WL (Kg/s) : 96.000  
 Pwf (BARS) : 84.000  
 H (KJ/Kg) : 856.637  
 Pflash (BARS) : 59.985

DEPTH	P	PC	DPC	D	X(%)	DPT	DPOT	DPACC	DPFRIC	H(J/g)	T(C)	FE	SLIP	VOID TY
575.5	59.99	44.21	0.090	24.45	0.0	0.000	0.000	0.000	0.000	856.6	201.0	0.0141	1.00	0.00 BU
500.0	15.29	2.79	-41.421	24.45	2.5	-44.700	-3.004	-0.156	-0.112	855.9	189.8	0.0141	1.56	0.70 AN
400.0	11.26	1.23	-1.559	24.45	4.6	-4.021	-2.172	-0.070	-0.220	854.9	179.8	0.0141	2.21	0.79 AN
300.0	0.65	0.69	-0.542	24.45	6.6	-2.613	-1.703	-0.057	-0.317	853.0	169.9	0.0141	3.17	0.83 AN
200.0	6.32	0.40	-0.208	24.45	8.9	-2.331	-1.508	-0.070	-0.466	852.7	157.8	0.0141	4.89	0.84 AN
100.0	3.90	0.22	-0.178	24.45	12.1	-2.424	-1.267	-0.204	-0.776	851.2	140.1	0.0141	6.19	0.88 AN
0.0	3.12	0.18	-0.039	24.45	13.5	-0.771	-0.294	-0.105	-0.334	850.5	132.5	0.0141	6.71	0.90 AN

WHP (BAR)-- 3.12

Mass flow rate : 96 kg/s                      Bottom hole pressure : 84 bar  
 Depth : 851 m                                      Temperature : 201 C  
 WHP : 3.0 bar                                      CO2 content of steam : 15 % by weight  
 Salt content : 1200 ppm                              Steam quality : 9.47 %

Table 2. Data and computer output for Kizildere Well no 6

### 3 RESPONSE OF THE SVARTSENGI GEOTHERMAL FIELD TO INJECTION

The Svartsengi geothermal field is a high temperature liquid dominated field with a temperature range 235-240°C and the fluid produced is in composition of two-thirds sea water and one-third fresh water. The geology of the Svartsengi field has been described by Franzson (1983). Reservoir engineering studies in Svartsengi are discussed by Kjaran et al. (1979) and Gudmundsson et al. (1985a, 1985b). The fluid production in Svartsengi has resulted in a drawdown of 135 m, after a fluid production of about 41 E9 kg. The rapid drawdown in the field may affect the following items of the future performance of the field: i) cold water encroachment from surrounding aquifers, and cooling of the formation, ii) decline in the output of production wells with falling reservoir pressure, iii) migration of the flashing zone down the wells and into the formation, iv) sealing of the producing fractures by calcite deposition if the fluid flashes in the formation. Injection would be a solution to the pressure recovery of the field as well as the efficient disposal of waste brine and condensate.

To study the effects of injection in the field two injection and tracer tests were carried out in 1982 and 1984. These tests are discussed by Gudmundsson (1983) and Hauksson (1985). The purpose of this study is to investigate the pressure response of the field to injection and match one year production history of the field by using a lumped-parameter model.

#### 3.1 Analysis of injection data

In the Svartsengi injection test, a brine-condensate mixture was injected for 72 days between 25th July to 4th October, 1984. The mixture was 80 percent flashed brine and 20 percent steam condensate and the injection flowrate was kept at 50 kg/s. The mixture had a pH close to 6.7 and the temperature 80°C. During the injection test, the drawdown was measured as water level in a monitoring well. The critical problem encountered during the injection was the silica scaling in the injection well. It was tried to

avoid this by controlling the pH of the injected fluid and by dilution of the brine with the condensate of the power plant. During the test the condensate concentration could not be kept constant because of the unavailability of adequate amount of condensate from the power plant. As a result the concentration of the condensate decreased down to 10 percent which caused an increase in the rate of silica deposition. One of the proposed solutions to decrease the rate of silica deposition is the injection of CO<sub>2</sub> with the condensate, which produces a weak acid to lower the pH down to 5.5 (Hauksson, T. personal communication).

To see the effect of injection on the drawdown of the field, the rate of production and drawdown data are used for the time period of October 1983 to January 1985. The total rate of production and injection data are shown in Fig. 8 with time, and the net production is the difference between production and injection rates. The water level drawdown is shown in Fig. 9 with time.

The drawdown data shows a linear trend for the first seven months in which the net production rate is almost constant. But after a decrease in production, the rate of drawdown also decreases and during the time of injection the water level starts to increase. The effect of reinjection can also be seen from the late portion of the data when the production rate reaches the initial value. During this period, the change in drawdown shows the similar trend as of the early data. By using these two graphs it can be concluded that the injection can be helpful to recover pressure.

### 3.2 Model description

Fluid extraction and reservoir drawdown in the Svartsengi field developed an increased steam zone after a few years of fluid production. This is evident from Well 10 which changed from being liquid-fed to producing steam only which is the shallowest well among the six main producing wells (424 m deep). Other wells in the field are still liquid-fed. A conceptual model of the Svartsengi reservoir is



shown in Fig. 10. The liquid dominated reservoir is overlain with a steam zone, which increases in size with drawdown (Hauksson, 1985).

Based on the above conceptual model Vatnaskil (1983) developed a lumped parameter model of the field. The schematic representation of the model is given in Fig. 11. The boundaries of the reservoir are impermeable and there is a slightly impermeable barrier between the feed zone and the liquid dominated zone. The drawdown in the feed zone leads to a leakage from the water zone. This leakage results in an increase in the size of the two-phase zone. The balance equations for this model can be written as;

$$W = q + A_2 S_2 \rho_2 \frac{dh_2}{dt} \quad (48)$$

$$q = A_1 S_1 \rho_1 \frac{dh_1}{dt} \quad (49)$$

$$q = c(h_2 - h_1) \quad (50)$$

where  $c$  is a constant.

If we solve these equations for  $h_1$  and  $h_2$  we get;

$$h_1(t) = C_3 \int_0^t m(t-\tau) e^{-\tau/k} d\tau \quad (51)$$

$$h_2(t) = C_1(t) - C_2 \int_0^t m(t-\tau) e^{-\tau/k} d\tau \quad (52)$$

where;

$$C_1 = \frac{1}{(\rho_2 A_2 S_2)^2} \quad (53)$$

$$C_2 = \frac{c}{\rho_2 A_2 S_2} \quad (54)$$

$$C_3 = \frac{C}{A_1 \rho_1 S_1 A_2 \rho_2 S_2} \quad (55)$$

$$K = \frac{1}{C} \frac{A_1 \rho_1 S_1 A_2 \rho_2 S_2}{A_1 \rho_1 S_1 + A_2 \rho_2 S_2} \quad (56)$$

and

$$M(t) = \int_0^k W(\tau) d\tau \quad (57)$$

The same set of production and drawdown data given in Figs. 8 and 9 were used to test the proposed model. The production rate of Well 10 was subtracted from the total field production, because it is producing from the two-phase region. For the best fit of drawdown, the variables in Eq. 54 are found as;

$$K = 125 \text{ days}, C = 5.1 \text{ E-9 m/kg}, C = 2.5 \text{ E-11 m/kg/day}$$

A good fit is obtained for the long term behavior of the drawdown (Fig. 12), but the model was not able to produce the sharp changes in the drawdown. This is due to the nature of the model, which assumes the storage is given by the free surface effect at the boiling surface, which is the long term effect thus neglecting the short term elastic storage. The model also assumes that all the introduced change in the flowrate should reach to the boundaries to be effective on the drawdown. The time required to see the boundary effects in the Svartsengi field was found to be 10-30 days by Kjaran et.al. (1979). Therefore, it is not possible to simulate any changes in the flowrate for a time less than 30 days.



#### 4 SUMMARY

Simulation of flowing wells can be a useful tool for geothermal well analysis. Using the flow model, it is still possible to get information for the conditions where the actual measurements failed. The following parameters can be deduced by the application of a wellbore simulator.

1. The temperature and pressure profiles of flowing wells can be obtained.
2. The change in the performance of the well after scaling development in the well can be simulated.
3. The magnitude of decline in wellhead pressure with aquifer pressure can be found
4. The wellbores with wider production casings give higher flowrates than those with smaller sized wells.

The practical use of the model presented here can be listed as follows:

- a. The flash level of the geothermal fluid can be found within a good accuracy, therefore it gives a valuable data for the cleaning operations of the wells.
- b. The wellhead pressure and mass flowrate history of the wells can be forecasted by using the predicted drawdown of the reservoir itself. This information is used for the planning of the operational life of a geothermal power plant, since it requires some minimum values for both mass flowrate and pressure values in the turbine.

It is found that the decline in the drawdown can be reduced by injection. ReInjection also has merits for the waste disposal problems of geofluid which causes pollution problems. To fit the immediate response of the Svartsengi field to injection, a new model should be developed which can respond to changes in the flowrate over a period of less than a month. In order to see the validity of the steam model the flowrate of injection should be kept constant for a longer time.

ACKNOWLEDGEMENTS

It is a great pleasure for me to express my appreciations to Dr. Snorri Pall Kjaran for his helpful and patient guidance during this study.

I would like to extend my thanks to Mr. Sigurdur Larus Holm who provided the field data and helped in the computer work of this study.

Thanks are due to Mr. Trausti Hauksson who conducted the well test and supplied his experience about the practical aspects of the injection test.

Special acknowledgements are also due to Dr. Ingvar Birgir Fridleifsson for critical reading of this report.

I also wish to extend my gratitude to the United Nations University for the Fellowship, the National Energy Authority of Iceland in the use of the computer and other facilities and the Middle East Technical University for the leave of absence which made this work possible.

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85.09.1179-00

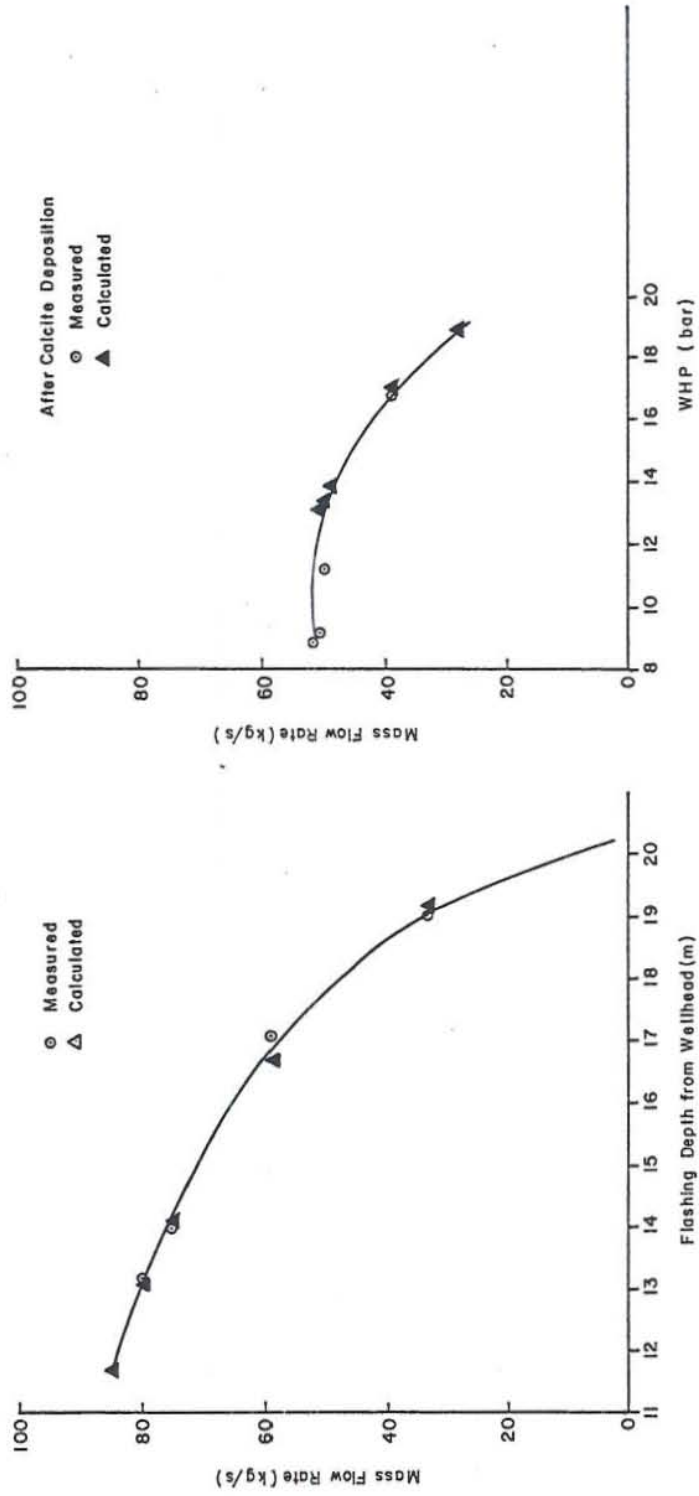


Figure 1. Svartsengi Well no. 4 Output Characteristics

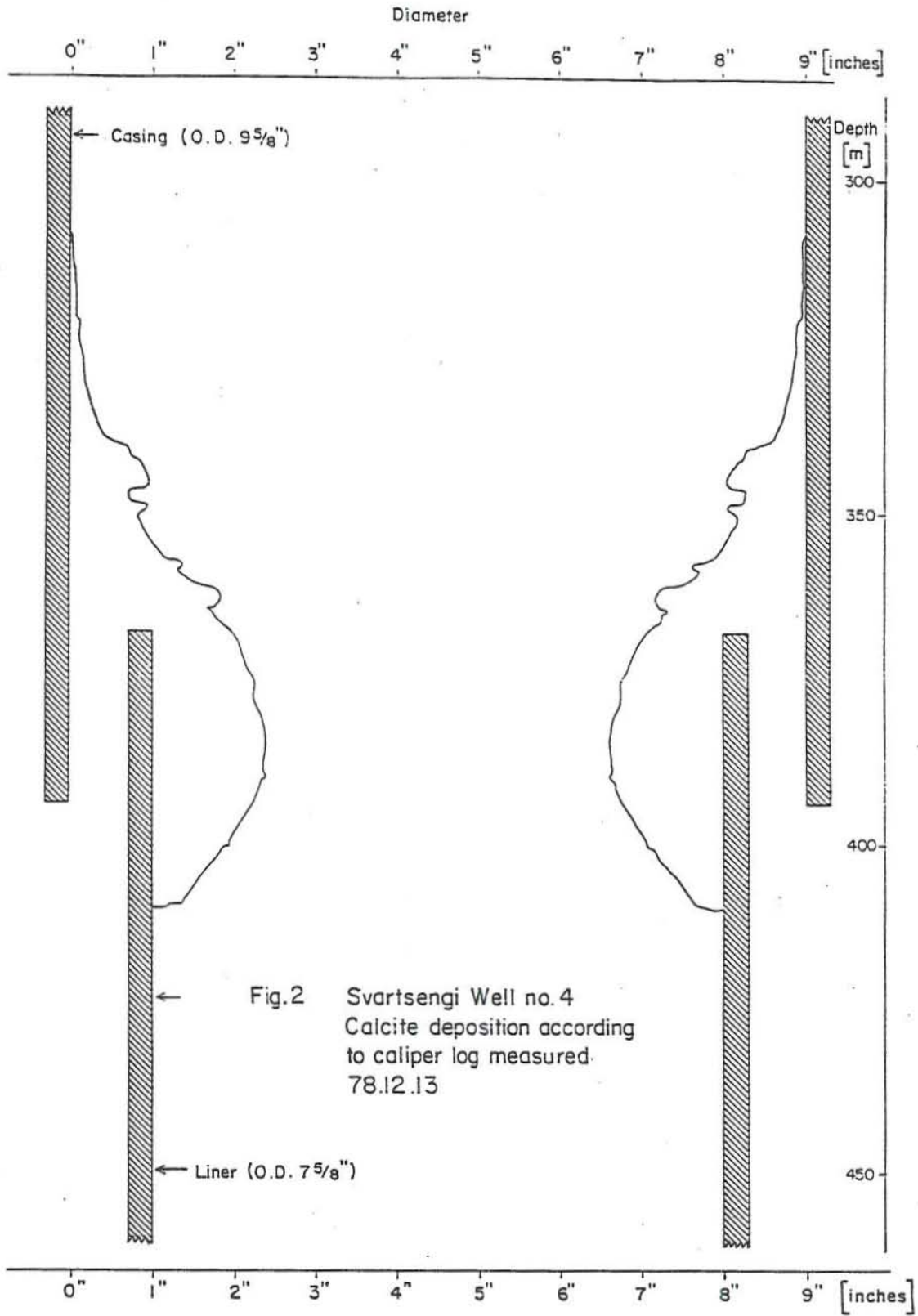
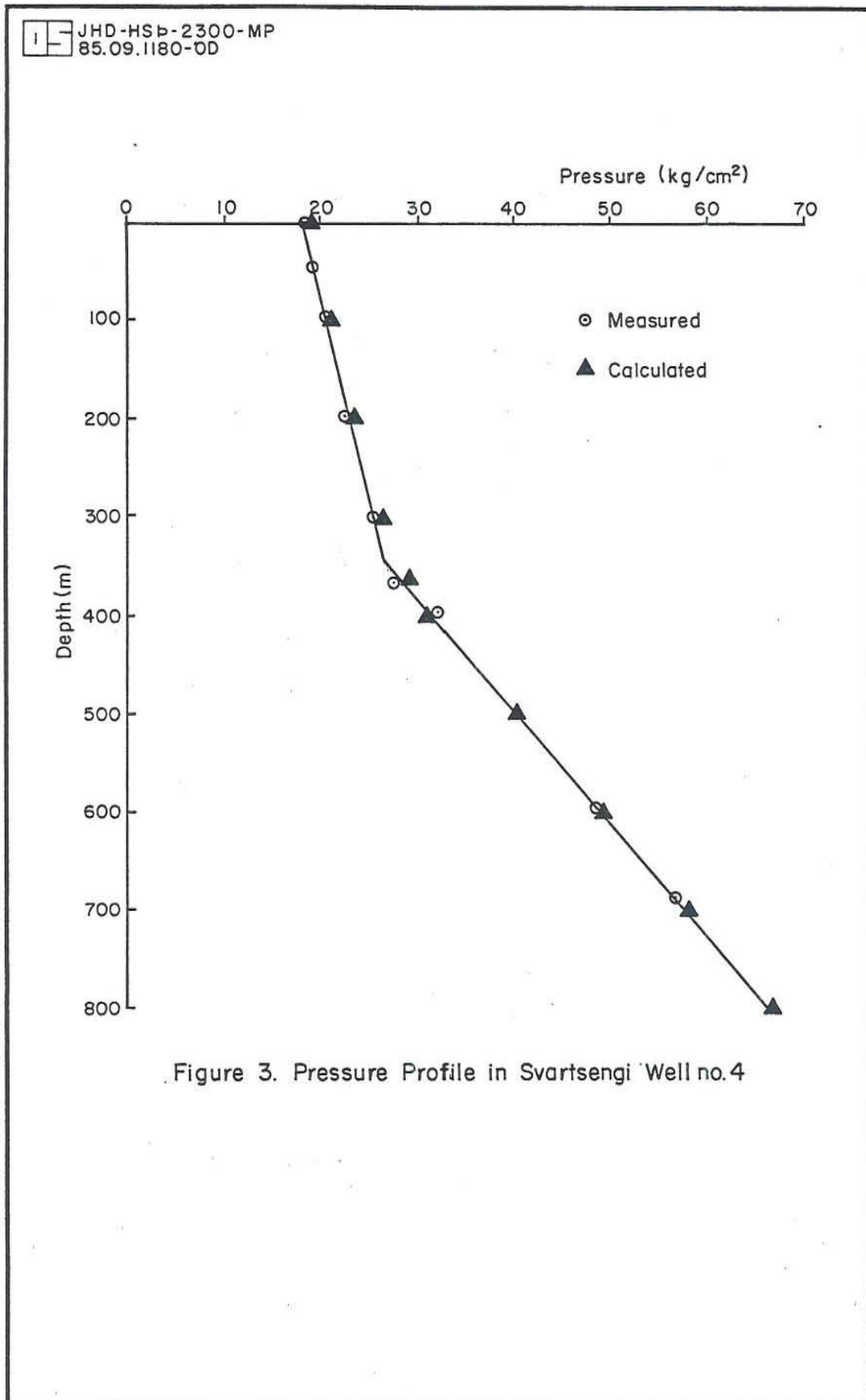


Fig.2 Svartsengi Well no. 4  
 Calcite deposition according  
 to caliper log measured.  
 78.12.13





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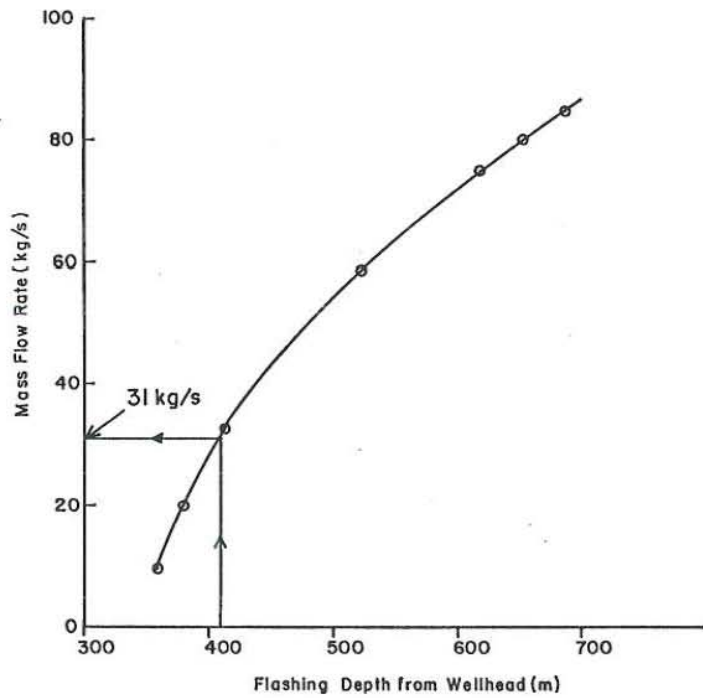


Figure 4. Change in Flashing Depth with Mass Flow Rate

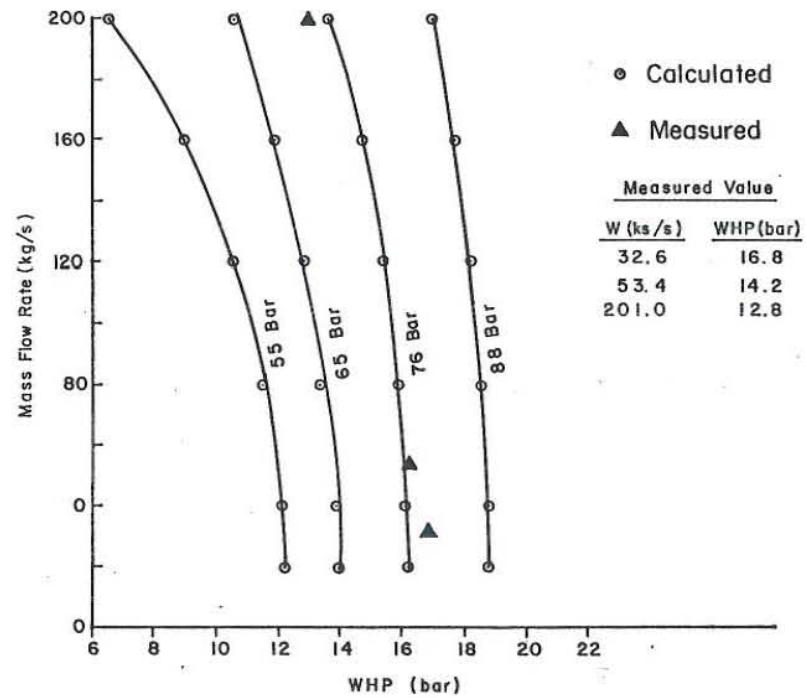


Figure 5. Deliverability curves for SG-II for different reservoir pressures

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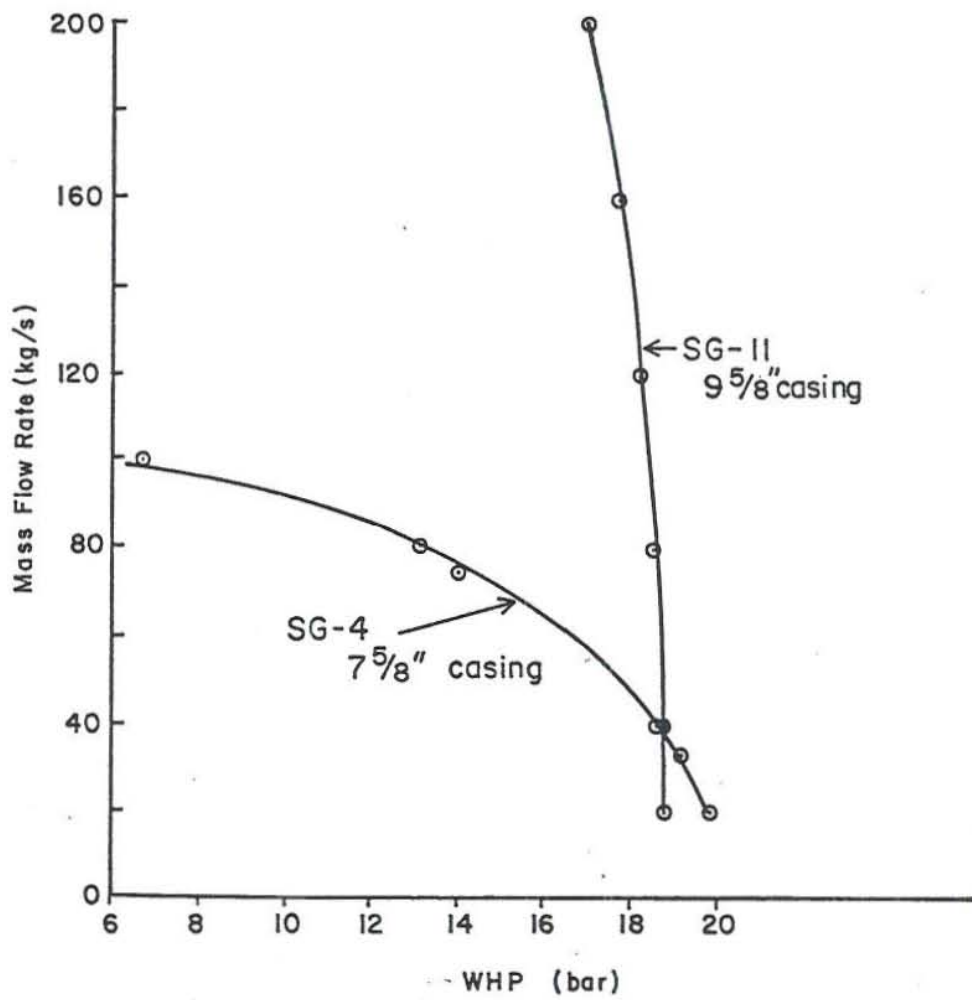


Figure 6. Effect of Casing Size on the Output Curves of geothermal Wells

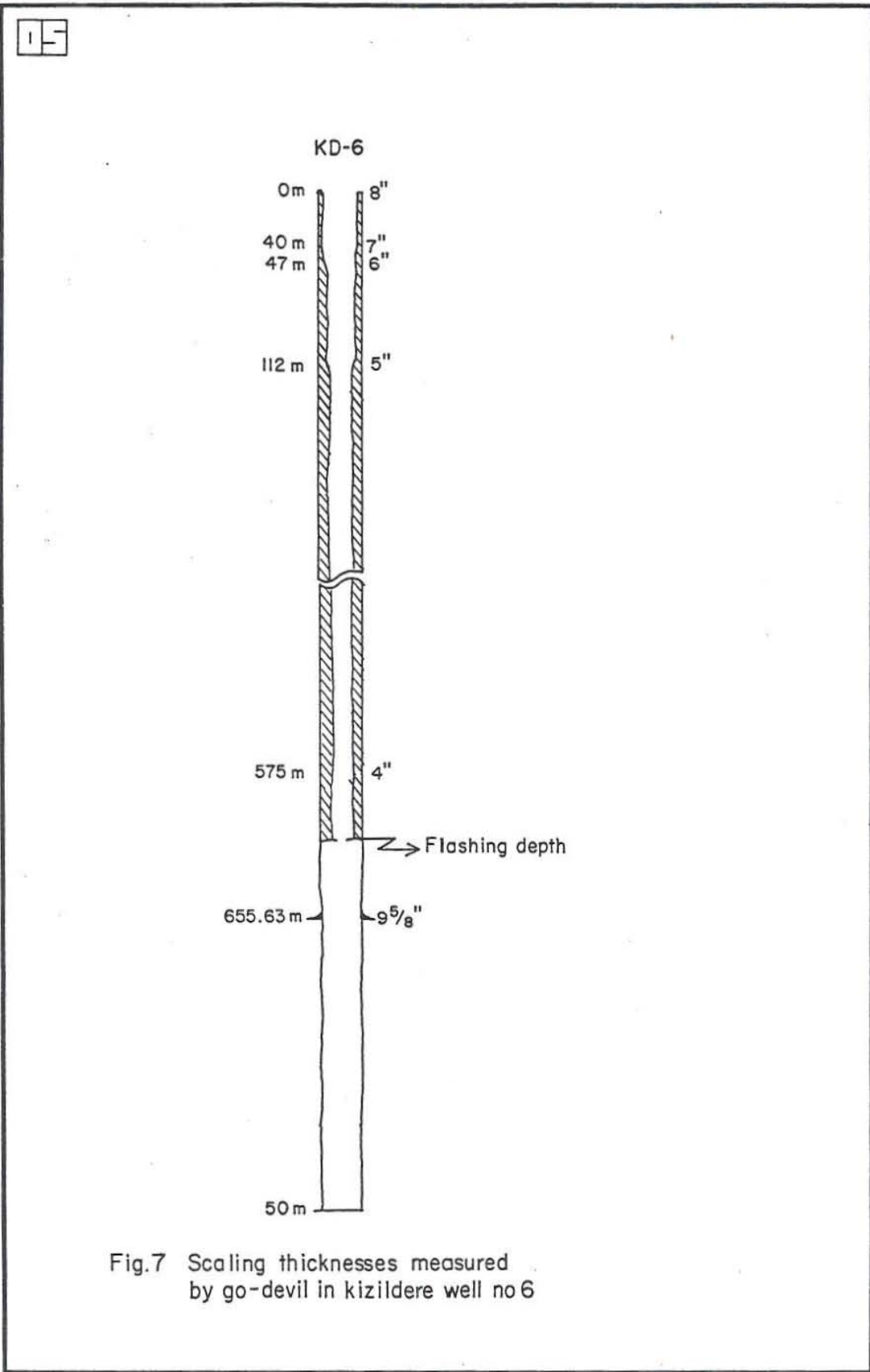


Fig.7 Scaling thicknesses measured by go-devil in kizildere well no 6

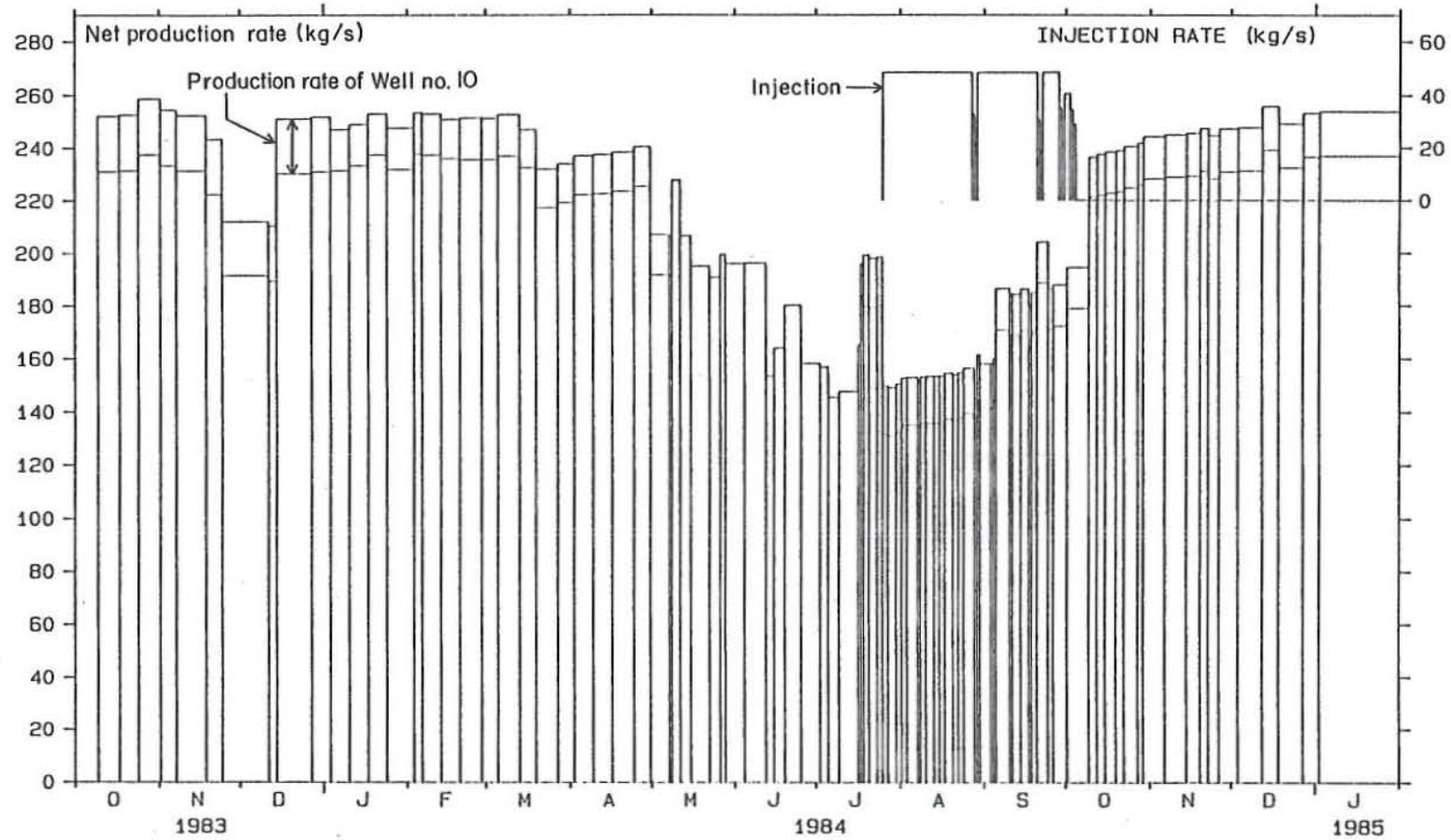
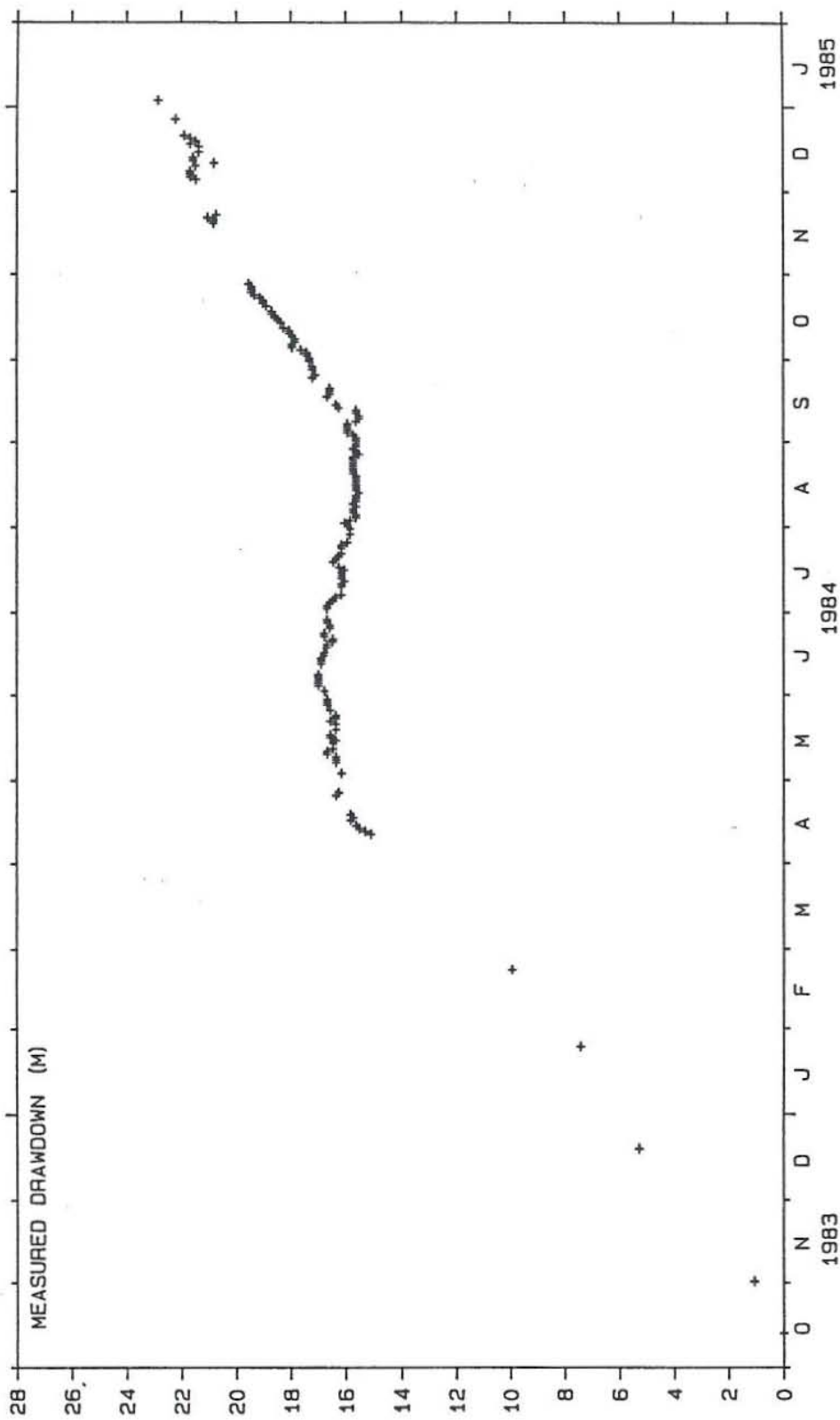


Figure 8. Net Production and Injection rates with time





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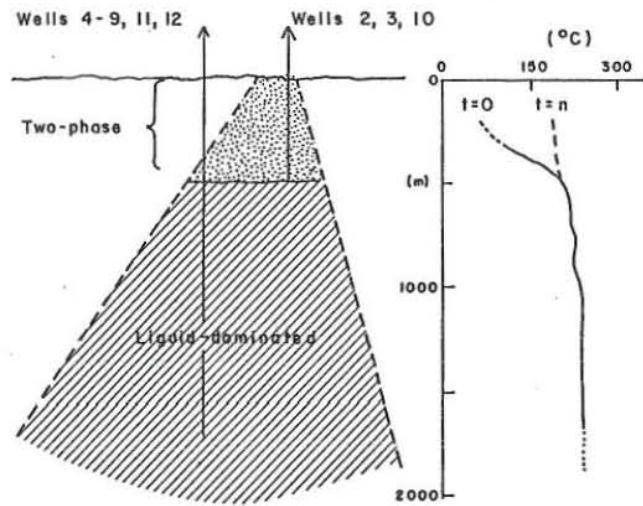


Figure 10. Conceptual Model of the Svartsengi Reservoir

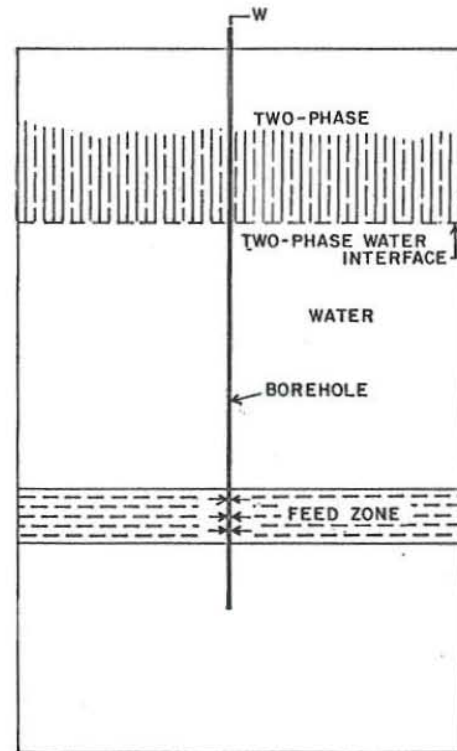


Figure 11. Schematic Representation of Steam Model of the Svartsengi Reservoir

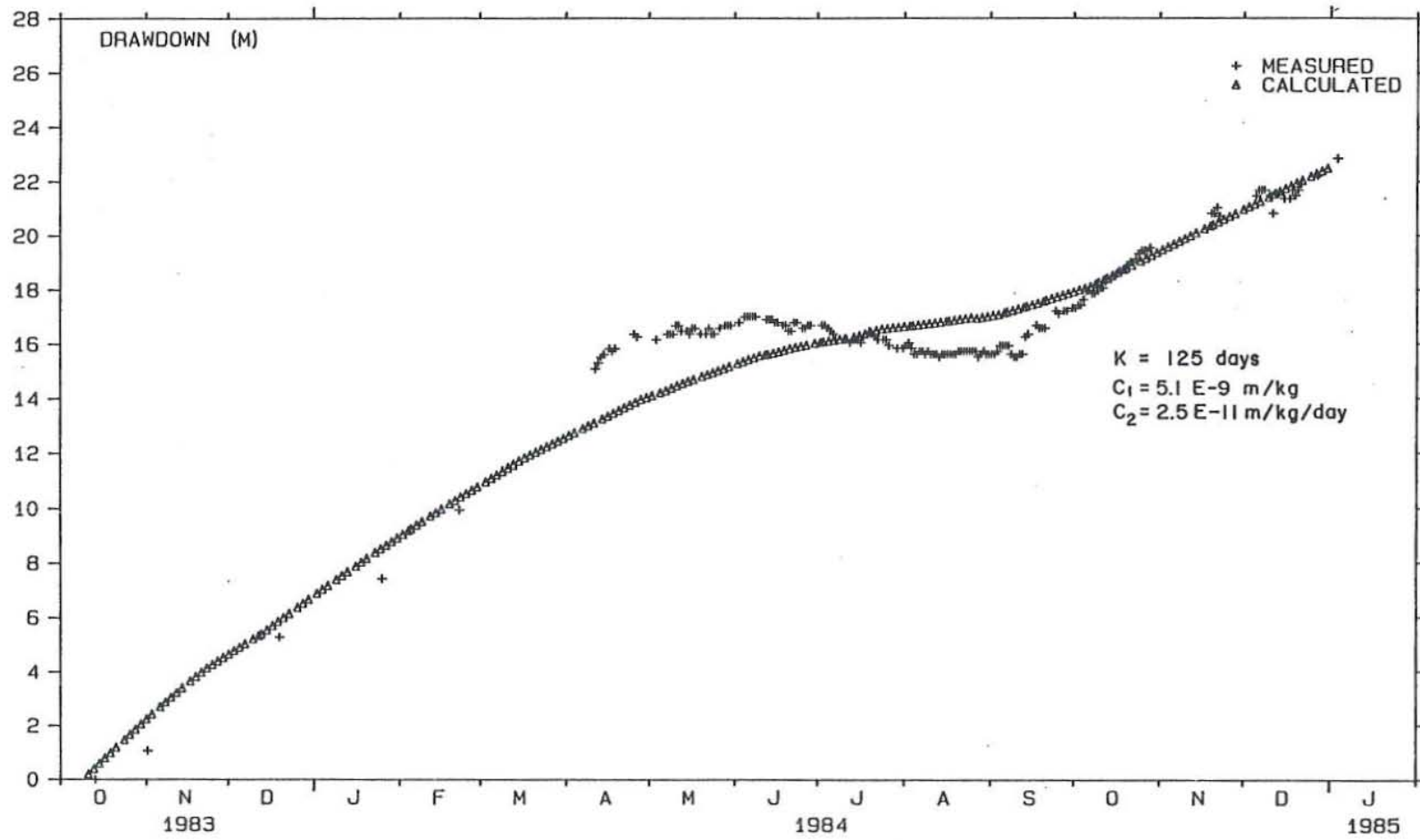


Figure.12. Measured and Fitted Drawdown from Steam Model



```

C-----
C CORRECTION FOR SALT CONCENTRATION
C-----
C CH=CNACL/58500.
C-----
C ENTHALPY OF LIQUID PHASE
C-----
C HL=HLP(T,CM)
C-----
C ENTHALPY OF VAPOR PHASE
C-----
C HG=HGP(T)
C-----
C SPECIFIC VOLUME OF LIQUID PHASE
C-----
C VL=VSH(T, RG, HL)
C-----
C SPECIFIC VOLUME OF VAPOR PHASE
C-----
C VG=VSG(PS,T)
C-----
C THE DIFFERENCE IN THE VAPOR PRESSURE DUE TO SALT
C-----
C CONCENTRATION
C-----
C DPS=(0.8314*(T+273.15)/(VG-VL)*CM/55.56)*1E-2
C-----
C PARTIAL PRESSURE OF CARBON DIOXIDE
C-----
C ALP=(5.4-3.5*(T/100.))+1.2*(T/100.)*2)*1E-9
C-----
C PCO=CO2*1E-11/ALP
C-----
C FLASH PRESSURE (BAR)
C-----
C PSTAR=PS-DPS+PCO
C-----
C FLASHING DEPTH
C-----
C CALL U07VAI(ZA,PA,ZZ,DO,FLAM,FLOW,DZ3,CNACL,IN,T,
C     ITS,PSTAR,VG,VL,DPS,CM,G,ZSTAR)
C-----
C Z=ZSTAR
C-----
C P=PSTAR
C-----
C PSP=PS-DPS
C-----
C PCI=PCO
C-----
C T1=T
C-----
C T2=T1+5
C-----
C J=0
C-----
C N=IN
C-----
C TWO PHASE CALCULATIONS
C-----
C INITIAL CONDITIONS AT TWO-PHASE SECTION
C-----
C HG=HGP(T)
C-----
C HSTAR=HLP(T,CM)
C-----
C HL=HSTAR
C-----
C VSTAR=VSH(T, RG, HL)
C-----
C VSTAR=VSTAR*1E-3
C-----
C VISF=VIS(CM,T)
C-----
C PACC=0.0025
C-----
C Q=HSTAR*1E3-ZSTAR*G
C-----
C TEMPZ=AINT(Z/TYPEZ+1.)*TYPEZ
C-----

```

```

DZ=-(Z-AINT((Z-1.)/DZ1)*DZ1)
IF(N.LE.1) GO TO 20
IF(Z.GT.ZZ(N-1)) GO TO 20
N=N-1
GO TO 10
D=DO(N)/100.
FLAMDA=FLAM(N)
FLOWA=FLOW/(D*D*0.78539816)
REYN=FLOW/(VISF*D*1.27324)
FFACT=((-2*ALOG10(FLAMDA/(3.7*D))+7*REYN**0.9)**2)**2)*E-1
FLOWA2=FLOWA**2
PFLUX=FLOWA2*VSTAR**2
VL=VL*1E-3
V1=VL
V2=VL
V3=VL
VA=VL*1.2
VEFF=VL
POT=0/VL
X=0.
B=0.5*FFACT*FLOWA2/D
D=HSTAR*1E3
ALPHA=0.
IF(.NOT.LTYPE) GO TO 30
WRITE(2,620) FLOW,PA,HSTAR,PSTAR
FORMAT(/' TWO-PHASE SECTION: /19(')/' HL (Kg/s) ',7X,
1=' ,PB.3/'/' Pw (BAR) ',6X,' ,PB.3/'/' H (KJ/Kg) ',
27X,' ,PB.3/'/' Pflash (BAR) ',3X,' ,PB.3/'/'
WRITE(2,670)
FORMAT(' ,T2,DEPTH(M) ',T12,'P(BAR) ',T19,'PC(BAR) ',T27,'DPC(BAR) ',
T136,'D(CM) ',T44,'DRMS ',T52,'DPT ',T59,'DPPOT ',T68,'DPACC ',T77,
2'DPFRC ',T86,'H(J/g) ',T94,'TEMP(C) ',T104,'Ft ',T109,
3'SLIP ',T116,'VOID ',T124,'TYPE '/130(')')
C-----
C NEW CALCULATION STEP
C-----
C IF(N.LE.1) GO TO 40
C IF(Z.GT.ZZ(N-1)) GO TO 40
C M=1
C IGO=1
C GO TO 300
C FLOWA=FLOW
C FLAMDA=FLAM(N)
C D=DO(N)/100.
C VISF=VIS(CM,T)
C REYN=FLOW/(VISF*D*1.27324)
C FFACT=((-2*ALOG10(FLAMDA/(3.7*D))+7*REYN**0.9)**2)**2)*E-1
C FLOWA2=FLOWA**2
C B=0.5*FFACT*FLOWA2/D
C DZ=0
C PFLUX=PFLUX*FLOWA/FLOWA0
C GO TO 70
C FLOWA0=FLOWA
C IGO=2
C IF(-DZ.EQ.0.) GO TO 300
C IF(Z.GT.ZZ(N-1)) GO TO 45
C GO TO 300
C PFLUX=PFLUX

```

```

10
20
620
670
30
35
40
45

```



```

50  DZ--(Z-AINT((Z-1.)/DZ1)*DZ1)
    IF(N.LE.1) GO TO 65
    IF(Z+DZ.GT.ZZ(N-1)) GO TO 70
    DZ--(Z-ZZ(N-1))
65  IF(Z+DZ.GT.ZT) GO TO 70
    DZ--(Z-ZT)
70  U=Q1(Z+DZ)*G
    DP=POT*DZ+FMOM*DZ
    PSP=PSP+DP*1E-5
    PSPF=PSP
    IF(DZ.EQ.0) DP=(X*X*VG/ALPHA+(1-X)**2*VG/
1(1-ALPHA))*(FLOWAO**2-FLOWA**2)
    NINT=0
    XLAST=X
    DDP=DP
    ISTOP=0
    POTO=POT
    FRICO=FRIC
210  NINT=NINT+1
    PDP=P*1E5+DP
    IF(X.EQ.0) GO TO 212
    IF(DZ.EQ.0) GO TO 212
    IF(PDP.GE.0.3116E6) GO TO 211
    DZ1=DZ1/2.
    IF(DZ1.LT.1) GO TO 90
    X=XLAST
    GO TO 50
C-----
C    CALL SUB PROGRAM TO CALCULATE NEW TEMPERATURE
C-----
211  CALL NEWTRM(X, VG, VL, T1, T2, PDP*1E-5, CM, PCO, T)
    HG=HGP(T)
    HL=HLP(T, CM)
    VL=VSW(T, HG, HL)
212  VG=VSG(PSPF, T)
    IF(DP.EQ.0) GO TO 230
    RKIN=0.5*FLOWA2*V4*V4
C-----
C    CALL SUB PROGRAM FOR HEAT LOSS
C-----
    IF(KK.NE.1) GO TO 213
    CALL HEATLOSS(Z, DZ, ITOUT, GRAD1, GRAD2, Z1, T, CON, DIP, D, PTIME, HE)
213  X=(U-RKIN-HL*1E3+HE/FLOW)/((HG-HL)*1E3)
    IF(DZ.EQ.0) GO TO 215
    IF(X.GT.0) GO TO 214
    ALPHA=0.
    V1=VL
    V2=VL
    V3=VL
    V4=VL
    GO TO 216
C-----
C    CORRECTION OF CARBON DIOXIDE PARTIAL PRESSURE
C-----
214  AT=(5.4-3.5*(T/100.))+1.2*(T/100. )**2)*1E-9
    RATIO=(1+(44*1E-3*VG*X)/(AT*8.314*(T+273.15)))**-1
    PC=PCO*RATIO
    DPCO2=PC-PC1
215  ALPHA=U07TP5(PDP*1E-5, X, VL, VG, FLOWA, D, ALPHA)
    C1=ALPHA

```

```

C2=1.-ALPHA
B1=X/C1
B2=(1-X)/C2
V1=1./(C1/VG+C2/VL)
C1=C1*B1
C2=C2*B2
V2=C1*VG+C2*VL
CC1=C1*B1
CC2=C2*B2
V3=CC1*VG+CC2*VL
CC1=CC1*B1
CC2=CC2*B2
V4=SQRT(CC1*VG*VG+CC2*VL*VL)
VEFF=U07TP4(X, ALPHA, VL, VG)
216  POT=G/V1
    FRIC=B*VEFF
    DPPOT=(POT+POTO)*0.5*DZ
    DPFIC=(FRIC+FRICO)*0.5*DZ
    PFLUX=FLOWA2*V3
    DPMOM=PFLUXO-PFLUX
    PSPF=PSP+(DPPOT+DPFRIC+DPMOM)*1E-5
    DPOLD=DP
    DP=DPOT+DPFRIC+DPMOM+(DPCO2-DPCO2O)*1E5
    DDPOLD=DDP
    DDP=DP-DPOLD
    IF(ABS(DDP/DP).LE.PACC) GO TO 230
    GO TO 210
230  IF(DZ.EQ.0) GO TO 232
    P=P+(DPPOT+DPMOM+DPFRIC)*1E-5+DPCO2-DPCO2O
    D1=D1+DPPOT+DPMOM+DPFRIC+(DPCO2-DPCO2O)*1E5
    GO TO 233
232  P=P+DPMOM*1E-5
    D1=D1+DPMOM
233  DPCO2O=DPCO2
    D2=D2+DPPOT
    D3=D3+DPMOM
    D4=D4+DPFRIC
    IF(DZ.NE.0) FMOM=FMOM/DZ
    IF(DZ.EQ.0) FMOM=FMOM*FLOWA/FLOWAO
    Z=Z+DZ
C-----
C    CALL SUB PROGRAM FOR HEAT LOSS
C-----
    IF(KK.NE.1) GO TO 231
    CALL HEATLOSS(Z, DZ, ITOUT, GRAD1, GRAD2, Z1, T, CON, DIP, D, PTIME, HE)
231  X=(U-RKIN-HL*1E3+HE/FLOW)/((HG-HL)*1E3)
    IF(Z.GT.ZT) GO TO 30
90  IF(DZ1.GE.1) GO TO 100
    PCRIT=(HSTAR**1.102*FLOWA)**1.04167*3.029E-13
    WRITE(2,660) PCRIT
660  FORMAT(' JAMES' CRITICAL PRESSURE-',F7.2,' (BARS)')
    GO TO 110
100  IGO=3
    GO TO 300
110  PT=P
    RRETURN
300  IF(.NOT.LTYPE) GO TO 360
C-----
C    DETERMINATION OF FLOW REGIME
C-----

```



```

VGOVL-VG/VL
IF(X.GT.1.) GO TO 306
X1=X/(1-X)
A1A=ALPHA/(1-ALPHA)
IF(X.GT.0) GO TO 302
S=1.
GO TO 304
302 S=X1/A1A*VGOVL
304 QY=A1A/(A1A+VGOVL)
BETA=X1/(X1+1/VGOVL)
GO TO 308
306 S-VGOVL
QX=1.
BETA=1.
R1=V1/V2
R3=V3/V2
R4=V4/V2
REFP=VEFF/V2
HBAQ=X*RG+(1-X)*HL
FR=FLOWA2/G/D*V2*V2
IF(BETA.LT.0.15) GO TO 320
IF(BETA.LT.0.55) GO TO 310
IF(BETA.LT.(-0.0085*FR+0.9962)) GO TO 330
GO TO 340
310 IF(BETA.LT.(-FR*0.02+1.85)) GO TO 330
GO TO 340
320 FLKIND=BUBB
GO TO 350
330 FLKIND=SLUG
GO TO 350
340 FLKIND=ANNU
350 IF(Z.GT.ZA) III=ISTAR
C-----
C CALCULATIONS ARE COMPLETED
C PRESSURE AND TEMPERATURE VALUES AT THE END OF INCREMENT
C-----
D1-D1*1E-5
DPP01=D2*1E-5
DPM01=D3*1E-5
DPRF1=D4*1E-5
D5-D1-(DPP01+DPM01+DPRF1)
IF(J.EQ.0) GO TO 380
C-----
C CALL SUB PROGRAM TO CALCULATE NEW TEMPERATURE
C-----
CALL NEWTEM(X,VG,VL,T1,T2,P,CM,PCO,T)
380 IF(J.EQ.1) PCI=PC
WRITE(2,650) Z,P,PCI,D5,DD(N),X,D1,DPP01,DPM01,DPRF1,HBAQ,
1T,FRAC,T,S,ALPHA,FLKIND
650 FORMAT(' ',T2,4F8.3,F6.2,2P8.3,3F9.4,2X,P7.2,F7.1,P8.4,2F7.3,3X,AB)
IF(2.LE.TEMP2) TEMP2=TEMP2-TYPEZ
IF(M.EQ.1) N=N-1
M=0
D1=0.
D2=0.
D3=0.
D4=0.
J=1
DPC020=0.
GO TO(35,45,110) IGO
END

```

END

C-----  
C SUB PROGRAM FOR TWO PHASE MULTIPLIER  
C-----

```

FUNCTION U07TP4(X,ALPHA,VL,VG)
X2=((1-X)/X)**2*VL/VG
X1=SQRT(X2)
C=1+X*VG/(X*VG+(1-X)*VL)-ALPHA
P2=1+C/X1+1/X2
U07TP4=P2*VL*(1-X)**2
RETURN
END

```

C-----  
C SUB PROGRAM FOR VOID FRACTION  
C-----

```

FUNCTION U07TP5(P,X,VL,VG,FLOWA,D,ALPHA)
FR=(FLOWA*VL)**2/D/9.8067
ALPHA1=(0.833+0.05*ALOG10(P))/(1+(1-X)/X*VL/VG)
BETA=X*VG/(X*VG+(1-X)*VL)
ALPHA2=BETA-0.71*86TA*SQRT(1-BETA)*FR**(-0.045)**(1-P/221.2)
ALPHA=MAX1(ALPHA1,ALPHA2)
U07TP5=ALPHA
RETURN
END

```

C-----  
C SUB PROGRAM FOR LIQUID PHASE ENTHALPY  
C-----

```

FUNCTION HLP(T,CM)
REAL A(4,3)
DATA ((A(I,J),J=1,3),I=1,4) /9633.6,-4080,286.49,166.58,
168.377,-4.6856,-0.90963,-0.36524,-0.0249667,0.179658,-2,
20.71924E-3,-0.49E-4/
ENTHALPY OF MIXING
SUM=0.
DO 3 I=1,4
DO 4 J=1,3
SUM1=0.
SUM1=SUM1+A(I,J)*T**((I-1)*CM**((J-1)
4 SUM=SUM+SUM1
3 SUM=SUM+SUM1
RM=4.184/(1000+58.44*CM)*SUM
C ENTHALPY OF PURE WATER
HM=0.12453E-4*T**3-0.45137E-2*T**4+4.81155*T-29.570
C ENTHALPY OF PURE SALT
HS=0.0715948*(0.83624E-3*T**3+0.16792*T-25.9293)*T
X1=1000/(1000+58.44*CM)
X2=58.44*CM/(1000+58.44*CM)
HLP=X1*HM+X2*HS+CM*RM
RETURN
END

```

C-----  
C SUB PROGRAM FOR VAPOR PHASE ENTHALPY  
C-----

```

FUNCTION HGP(T)
HGP=-0.81257E-2*T**3+3.65228*T+2388.4
RETURN
END

```

```

C-----
C SUB PROGRAM FOR LIQUID PHASE SPECIFIC VOLUME
C-----
FUNCTION VSM(T,HG,HL)
DATA C1,C2,C3,C4,C5/-0.315154,-1.203374E-3,7.48908E-13
1,0.1342489,-3.946963E-3/
TSS-14.9652*(T+273.15)**2/(HG-HL)*CM/55.56
F-374.12-T-TSS
VSM=(3.1975+C1*F**(1./3)+C2*F+C3*F**4)/
1(1+C4*F**(1./3)+C5*F)*1E-3
RETURN
END

```

```

C-----
C SUB PROGRAM FOR VAPOR PHASE SPECIFIC VOLUME
C-----
FUNCTION VSG(P,T)
VSG-1./(100*P/(-0.1296E-2*T*T+0.6325*T+121.05))
RETURN
END

```

```

C-----
C SUB PROGRAM FOR VISCOSITY
C-----
FUNCTION VIS(CM,T)
A=0.3324E-1*CM+0.3624E-2*CM*CM-0.1879*CM**3
B=-0.3961E-1*CM+0.0102*CM*CM-0.702E-3*CM**3
C=20-T
D=C/(96+T)*(1.2378-1.303E-3*C+3.06E-6*C*C+
12.55E-8*C**3)
M=1002*10**0
G=A/B*D
VIS=MT*(10**G)*1E-6
RETURN
END

```

```

C-----
C SUB PROGRAM FOR THE CALCULATIONS IN SINGLE-PHASE
C-----
SUBROUTINE U07VAL(ZA,PA,ZZ,DD,FLAM,FLOW,DZ3,CNACL,IN,T,
ITS,PSTAR,VG,VL,DPS,CM,G,ZSTAR)
DIMENSION ZZ(IN),DD(IN),FLAM(IN)

```

```

C-----
C DENSITY OF WATER
C-----
HROL=DEN(CM,VL)
WRITE(2,1005)
1005 FORMAT(T5,'DEPTH(m)',T19,'P(bar)',T32,'Vel(f)',T43,'Ffactor
1',T55,'D(cm)',T68,'DPWAT.',T80,'DPPOT',T91,'DPRIC',T100,
2'FLOW(kg/s)',T111,'TEMP(C)',T122,'PPLASH'/130('-'//))
N=IN
P=PA
Z=ZA
VISF=VIS(CM,T)
M=1
J=2
DZ=Z-AINT(Z/DZ3)*DZ3
Z=Z-DZ
1010 IF(N.LE.1) GO TO 1020
IF(J.EQ.2) GO TO 1020

```

```

Z=Z-DZ
IF(Z.LT.ZZ(N-1)) GO TO 1060
GO TO 1020
1060 Z=Z+DZ
DZ=Z-ZZ(N-1)
J=1
GO TO 1010
1020 D=DD(N)/100.
P=P*1E5
FLAMDA=FLAM(N)
VVATN=FLOW/HROL/(D*D*0.78539816)
REPT=HROL*VVATN*D/VISF
FFACT=((-2*ALOG10(FLAMDA/(3.7*D)+(7/REPT)**0.9))**2)**-1
DPPOT=-(HROL*G*DZ)
DPRIC=-(FFACT*HROL*VVATN**2/2/D*DZ)
DPVATN=DPPOT+DPRIC
P=P+DPVATN
1030 IF(P.LE.(PSTAR*1E5)) GO TO 1050
P=P*1E-5
DPPOT=DPPOT*1E-5
DPRIC=DPRIC*1E-5
DPVATN=DPVATN*1E-5
1080 WRITE(2,1040) Z,P,VVATN,FFACT,DD(N),DPVATN,DPPOT,DPRIC,
1FLOW,T,PSTAR
1040 FORMAT(BF12.3,F10.2,F9.1,F12.3)
IF(J.EQ.1) GO TO 1070
IF(M.EQ.1) J=0
IF(J.EQ.0) GO TO 1006
DZ=Z-AINT(Z/DZ3)*DZ3
M=1
IF(N.LE.1) GO TO 1090
GO TO 1010
1090 Z=Z-DZ
GO TO 1020
1070 N=N-1
D=DD(N)/100.
FLAMDA=FLAM(N)
VNEW=FLOW/HROL/(D*D*0.78539816)
REPT=HROL*VNEW*D/VISF
FFACT=((-2*ALOG10(FLAMDA/(3.7*D)+(7/REPT)**0.9))**2)**-1
DPVATN=(VVATN-VNEW)**2*HROL/2
P=(P*1E5+DPVATN)*1E-5
DPVATN=DPVATN*1E-5
VVATN=VNEW
DPPOT=0.
DPRIC=0.
M=0
J=3
GO TO 1080
1006 DZ=DZ3
M=0
IF(N.LE.1) GO TO 1090
GO TO 1010
1050 P=P-DPVATN
DPVATN=ABS(DPVATN)
Z=Z+DZ
DZSTAR=(P-PSTAR*1E5)/DPVATN*DZ
ZSTAR=Z-DZSTAR
IN=N
RETURN

```

```

END
C-----
C SUB PROGRAM FOR DENSITY OF LIQUID PHASE
C-----

```

```

FUNCTION DEN(CM,VL)
DATA C1,C2,C3,C4,C5,-167.219,448.55,-261.07,-13.644,13.97/
VL-VL*1E3
P=C1+C2*VL+C3*VL+((C4+C5*VL)+10.224/(3.195-VL)**2)
DEN=(1000+CM*58.44)/((1000+VL+CM*P)*1E3)
RETURN
END

```

```

C-----
C SUB PROGRAM FOR HEAT LOSS CALCULATIONS
C-----

```

```

SUBROUTINE HEATLOSS(Z,DZ,ITOUT,GRAD1,GRAD2,Z1,T,CON,DIF,D,TIME,Q)
DZ=.DZ
IP(Z,LT,Z1) GO TO 1
IF(Z,GT,Z1.AND.(Z-DZZ).LT,Z1) GO TO 3
TR=Z1*GRAD1+(Z-Z1-DZ/2)*GRAD2+ITOUT
GO TO 2
TR-ITOUT+GRAD1*(Z-DZ/2)
1 Q=6.283185*CON*(TR-T)/FOT*DZ
2 RETURN
3 Z11=Z-Z1
Q1=6.283185*CON*(TR1-T)/FOT*Z11
Z2Z=DZ-Z11
TR2=GRAD1*(Z1-Z2Z/2)+ITOUT
Q=Q1+Q2
RETURN
END

```

```

C-----
C SUB PROGRAM TO CALCULATE TEMPERATURE
C-----
SUBROUTINE NEWTRM(X,VG,VL,T1,T2,P,CM,PCO,T)
DATA C1,C2,C3/0.21913E-6,0.17816E-3,0.0653665/
FT(C1,C2,C3)=P-EXP(C1*TA**3-C2*TA**TA+C3*TA-4.96087)+
1(0.8314*(TA+273.15))/(VG-VL)*CM/55.56*1E-2-
2PCO*(1+(44*1E-3*VG*X)/((5.4-3.5*(TA/100.))+
31.2*(TA/100.))**2)*1E-9)*8.314*(TA+273.15))**2-1
T=T1-(T1-T2)*FT(T1)/(FT(T1)-FT(T2))
IF(ABS(T-T1).LE.0.01) RETURN
T2=T1
GO TO 1
END

```

APPENDIX B.1. COMPUTER OUTPUTS FOR SVARTSENGI 4 WITHOUT CALCITE DEPOSITION

NAME OF FIELD: SVARTSENGI  
WELL NAME : SG-4  
DATE OF CALC : 16-09-85

SINGLE-PHASE(WATER) SECTION:

Pa (BARS) = 88.000  
(dp)lamb (BARS) = 3.812  
Pwf (BARS) = 84.188  
CCO2 (ppm) = 500.0  
CNAACL (ppm) = 21000.0

DEPTH(m)	P(Sat)	Vc1(f)	Ffactor	D(cm)	DPMAT	DPFOT	DPFRIC	W(kn/s)	T(C)	RELEASE
1000.000	82.214	1.348	0.018	19.368	-1.975	-1.958	-0.017	33.00	240.0	33.739
900.000	73.986	1.348	0.018	19.368	-8.228	-8.156	-0.072	33.00	240.0	33.739
800.000	65.758	1.348	0.018	19.368	-8.228	-8.156	-0.072	33.00	240.0	33.739
700.000	57.530	1.348	0.018	19.368	-8.228	-8.156	-0.072	33.00	240.0	33.739
600.000	49.302	1.348	0.018	19.368	-8.228	-8.156	-0.072	33.00	240.0	33.739
500.000	41.074	1.348	0.018	19.368	-8.228	-8.156	-0.072	33.00	240.0	33.739

TWO-PHASE SECTION:

WL (KG/S) = 33.000  
Pwf (BARS) = 84.188  
B (KJ/KG) = 1037.213  
PFlash (BARS) = 33.739

DEPTH	P	PC	PPC	D	X(%)	DPF	DPFOT	DPFRIC	B(J/g)	T(C)	PF	SLIP VOID FR
410.9	33.73	1.27	0.90	19.37	0.0	0.000	0.000	0.000	1037.2	240.0	0.0185	1.00
400.0	32.40	0.77	-0.50	19.37	0.4	-1.335	-0.803	-0.018	-0.004	1037.1	238.5	0.0185
362.0	29.78	0.35	-0.41	19.37	1.4	-2.624	-2.165	-0.007	-0.029	1036.7	234.5	0.0185
362.0	29.79	0.35	0.00	24.45	1.4	0.006	0.000	0.006	0.000	1036.7	234.5	0.0148
300.0	27.04	0.19	-0.15	24.45	2.6	-2.746	-2.569	-0.004	-0.013	1036.1	229.4	0.0148
200.0	23.85	0.11	-0.08	24.45	4.2	-3.192	-3.082	-0.005	-0.023	1035.1	222.7	0.0148
100.0	21.34	0.07	-0.03	24.45	5.5	-2.510	-2.443	-0.004	-0.027	1034.2	216.8	0.0148
0.0	19.20	0.05	-0.02	24.45	6.6	-2.139	-2.084	-0.004	-0.031	1033.2	211.4	0.0148

WHP (BAR)= 19.20

where; PV : Type of flow in two-phase,  
BU : Bubbly flow, SL : Slug flow and AN : Annular flow



NAME OF FIELD: SWARTSENGI  
 WELL NAME : SG-4  
 DATE OF CALC : 16-09-85

SINGLE-PHASE(WATER) SECTION:

Pa (BARS) = 88.000  
 (dp)urb (BARS) = 19.688  
 Pwf (BARS) = 68.313  
 CCO2 (ppm) = 500.0  
 CNACL (ppm) = 21000.0

DEPTH(m)	P(bar)	Vel(f)	Pfactor	D(cm)	DPHAT	DPPOF	DPACC	DPFRIC	H(kg/s)	T(C)	PPHASE
1000.000	66.267	3.064	0.018	19.368	-2.046	-1.958	0.000	0.000	1037.2	240.0	0.0183
900.000	57.743	3.064	0.018	19.368	-8.523	-8.156	-0.104	-0.053	1037.0	237.4	0.0183
800.000	49.220	3.064	0.018	19.368	-8.523	-8.156	-0.104	-0.053	1035.0	229.9	0.0183
700.000	40.697	3.064	0.018	19.368	-8.523	-8.156	-0.104	-0.053	1034.6	217.5	0.0183

TWO-PHASE SECTION:

HL (kg/s) = 75.000  
 Pwf (BARS) = 68.313  
 H (kg/kg) = 1037.213  
 Pflash (BARS) = 33.739

DEPTH	P	PC	DFC	D	X(1)	OPT	DPPOF	DPACC	DPFRIC	H(kg/s)	T(C)	FF	SLIP	VOID	TY
613.4	33.74	1.28	0.00	19.37	0.0	0.000	0.000	0.000	0.000	1037.2	240.0	0.0183	1.00	0.00	2U
600.0	31.62	0.59	-0.69	19.37	0.7	-2.124	-1.278	-0.104	-0.053	1037.0	237.4	0.0183	1.14	0.24	SL
500.0	26.30	0.17	-0.42	19.37	3.0	-5.312	-4.381	-0.086	-0.423	1036.0	227.9	0.0183	1.31	0.60	AN
400.0	22.78	0.10	-0.07	19.37	4.7	-3.523	-2.888	-0.064	-0.498	1035.0	220.3	0.0183	1.50	0.70	AN
362.0	21.61	0.08	-0.02	19.37	5.3	-1.169	-0.921	-0.021	-0.212	1034.6	217.5	0.0183	1.58	0.73	AN
362.0	21.68	0.00	0.00	24.45	5.3	0.065	0.000	0.000	0.000	1034.7	217.7	0.0142	1.58	0.73	AN
300.0	20.18	0.07	-0.02	24.45	6.1	-1.499	-1.382	-0.012	-0.090	1034.1	213.9	0.0142	1.71	0.75	AN
200.0	17.98	0.05	-0.02	24.45	7.3	-2.193	-1.993	-0.018	-0.164	1033.1	208.1	0.0142	1.94	0.79	AN
100.0	15.98	0.04	-0.01	24.45	8.5	-2.009	-1.788	-0.017	-0.192	1032.1	202.2	0.0142	2.23	0.81	AN
0.0	14.08	0.03	-0.01	24.45	9.8	-1.895	-1.642	-0.017	-0.227	1031.1	196.5	0.0142	2.60	0.82	AN

WEP (BAR) = 14.08

NAME OF FIELD: SWARTSENGI  
 WELL NAME : SG-4  
 DATE OF CALC : 16-09-85

SINGLE-PHASE(WATER) SECTION:

Pa (BARS) = 88.000  
 (dp)urb (BARS) = 12.184  
 Pwf (BARS) = 75.816  
 CCO2 (ppm) = 500.0  
 CNACL (ppm) = 21900.0

DEPTH(m)	P(bar)	Vel(f)	Pfactor	D(cm)	DPHAT	DPPOF	DPACC	DPFRIC	H(kg/s)	T(C)	PPHASE
1000.000	73.084	2.410	0.018	19.368	-2.012	-1.958	0.000	0.000	59.00	240.0	33.739
900.000	65.420	2.410	0.018	19.368	-8.384	-8.156	-0.228	-0.228	59.00	240.0	33.739
800.000	57.036	2.410	0.018	19.368	-8.384	-8.156	-0.228	-0.228	59.00	240.0	33.739
700.000	48.652	2.410	0.018	19.368	-8.384	-8.156	-0.228	-0.228	59.00	240.0	33.739
600.000	40.269	2.410	0.018	19.368	-8.384	-8.156	-0.228	-0.228	59.00	240.0	33.739

TWO-PHASE SECTION:

HL (kg/s) = 59.000  
 Pwf (BARS) = 75.816  
 H (kg/kg) = 1037.213  
 Pflash (BARS) = 33.739

DEPTH	P	PC	DFC	D	X(1)	OPT	DPPOF	DPACC	DPFRIC	H(kg/s)	T(C)	FF	SLIP	VOID	TY
552.1	33.74	1.28	0.00	19.37	0.0	0.000	0.000	0.000	0.000	1037.2	240.0	0.0184	1.00	0.00	2U
500.0	31.38	0.55	-0.73	19.37	0.1	-2.356	-1.508	-0.066	-0.950	1037.0	237.1	0.0184	1.14	0.25	SL
400.0	26.37	0.17	-0.38	19.37	3.0	-5.017	-4.326	-0.051	-0.263	1036.0	228.0	0.0184	1.31	0.52	SL
362.0	25.00	0.13	-0.04	19.37	3.6	-1.362	-1.200	-0.015	-0.111	1035.6	225.2	0.0184	1.38	0.65	AN
362.0	25.03	0.13	0.00	24.45	3.6	0.031	0.000	0.031	0.000	1035.6	225.3	0.0143	1.37	0.66	SL
300.0	23.23	0.10	-0.03	24.45	4.5	-1.805	-1.714	-0.009	-0.948	1035.0	221.3	0.0143	1.48	0.70	SL
200.0	20.75	0.07	-0.03	24.45	5.8	-2.479	-2.350	-0.012	-0.686	1034.0	215.4	0.0143	1.68	0.75	AN
100.0	18.60	0.05	-0.02	24.45	7.0	-2.151	-2.022	-0.011	-0.400	1033.1	209.7	0.0143	1.90	0.78	AN
0.0	16.56	0.04	-0.01	24.45	8.1	-1.944	-1.805	-0.011	-0.116	1032.1	204.2	0.0143	2.18	0.80	AN

WEP (BAR) = 16.56

APPENDIX 3.3. COMPUTER OUTPUTS FOR SVARTSENGI 4 WITH CALCIUM DEPOSITION

NAME OF FIELD: SVARTSENGI  
WELL NAME : SG-4  
DATE OF CALC : 17-09-85

SINGLE-PHASE(WATER) SECTION:

Pa (BARS) = 88.000  
(dp)urb (BARS) = 2.744  
Pwf (BARS) = 85.256  
CCO2 (ppm) = 500.0  
CMACL (ppm) = 21000.0

DEPTH(m)	P(bar)	Vel(f)	Ffactor	D(cm)	DPDAT	DPDOT	DPFRIC	H(kr/s)	T(C)	PPHASE
1000.000	83.286	1.144	0.018	19.368	-1.970	-1.958	-0.012	28.00	240.0	33.739
900.000	75.073	1.144	0.018	19.368	-8.208	-8.156	-0.052	28.00	240.0	33.739
800.000	66.870	1.144	0.018	19.368	-8.208	-8.156	-0.052	28.00	240.0	33.739
700.000	58.662	1.144	0.018	19.368	-8.208	-8.156	-0.052	28.00	240.0	33.739
600.000	50.454	1.144	0.018	19.368	-8.208	-8.156	-0.052	28.00	240.0	33.739
500.000	42.245	1.144	0.018	19.368	-8.208	-8.156	-0.052	28.00	240.0	33.739
410.000	34.037	1.144	0.018	19.368	-7.387	-7.341	-0.047	28.00	240.0	33.739
400.000	33.855	3.244	0.054	11.500	0.013	0.000	0.000	28.00	240.0	33.739
400.000	33.855	3.244	0.054	11.500	-1.021	-0.915	-0.105	28.00	240.0	33.739

TWO-PHASE SECTION:

HL (kg/s) = 28.000  
Pwf (BARS) = 85.256  
B (kJ/kg) = 1037.213  
Pflash (BARS) = 33.739

DEPTH	P	PC	OPC	D	X(1)	OPT	DPDOT	DPACC	DPFRIC	H(J/m)	T(C)	PE	SLIP	VOID	FV
398.9	33.74	1.29	0.00	11.50	0.0	0.000	0.000	0.000	1037.2	240.0	0.0540	1.00	0.00	80	
350.0	28.79	0.28	-0.10	11.50	1.8	-4.946	-2.797	-0.163	-0.989	1036.7	232.7	0.0540	1.22	0.47	SL
350.0	28.92	0.28	0.00	24.45	1.8	0.073	0.000	0.028	0.000	1036.7	232.3	0.0150	1.21	0.46	SL
300.0	26.75	0.19	-0.09	24.45	2.8	-2.078	-1.966	-0.002	-0.008	1036.2	223.8	0.0150	1.29	0.57	SL
200.0	23.59	0.11	-0.09	24.45	4.3	-3.160	-3.062	-0.003	-0.017	1035.3	222.1	0.0150	1.45	0.68	SL
100.0	21.09	0.07	-0.04	24.45	5.6	-2.505	-2.448	-0.003	-0.024	1034.3	216.2	0.0150	1.63	0.74	SL
0.0	18.94	0.06	-0.02	24.45	6.8	-2.141	-2.096	-0.002	-0.023	1033.3	210.7	0.0150	1.84	0.78	SL

MRP (BAR) = 18.94

NAME OF FIELD: SVARTSENGI  
WELL NAME : SG-4  
DATE OF CALC : 17-09-85

SINGLE-PHASE(WATER) SECTION:

Pa (BARS) = 88.000  
(dp)urb (BARS) = 5.324  
Pwf (BARS) = 82.676  
CCO2 (ppm) = 500.0  
CMACL (ppm) = 21000.0

DEPTH(m)	P(bar)	Vel(f)	Ffactor	D(cm)	DPDAT	DPDOT	DPFRIC	H(kr/s)	T(C)	PPHASE
1000.000	80.695	1.593	0.018	19.368	-1.982	-1.958	-0.024	39.00	240.0	33.739
900.000	72.439	1.593	0.018	19.368	-8.256	-8.156	-0.100	39.00	240.0	33.739
800.000	64.182	1.593	0.018	19.368	-8.256	-8.156	-0.100	39.00	240.0	33.739
700.000	55.926	1.593	0.018	19.368	-8.256	-8.156	-0.100	39.00	240.0	33.739
600.000	47.670	1.593	0.018	19.368	-8.256	-8.156	-0.100	39.00	240.0	33.739
500.000	39.414	1.593	0.018	19.368	-8.256	-8.156	-0.100	39.00	240.0	33.739

TWO-PHASE SECTION:

HL (kg/s) = 39.000  
Pwf (BARS) = 82.676  
B (kJ/kg) = 1037.213  
Pflash (BARS) = 33.739

DEPTH	P	PC	OPC	D	X(1)	OPT	DPDOT	DPACC	DPFRIC	H(J/m)	T(C)	PE	SLIP	VOID	FV
431.3	33.74	1.29	0.00	19.37	0.0	0.000	0.000	0.000	1037.2	240.0	0.0185	1.00	0.00	80	
410.0	31.51	0.57	-0.71	19.37	0.7	-2.224	-1.468	-0.028	-0.222	1037.0	237.3	0.0185	1.14	0.25	SL
410.0	31.37	0.57	0.00	11.50	0.7	0.000	0.000	-0.147	0.000	1037.0	237.0	0.0540	1.14	0.25	SL
400.0	30.15	0.38	-0.19	11.50	1.3	-1.221	-0.559	-0.045	-0.429	1036.9	235.1	0.0540	1.18	0.38	SL
350.0	25.41	0.15	-0.23	11.50	3.4	-4.733	-1.988	-0.161	-2.354	1036.4	226.1	0.0540	1.31	0.62	AN
350.0	25.49	0.15	0.00	24.45	3.4	0.075	0.000	0.075	0.000	1036.4	226.3	0.0147	1.31	0.61	SL
300.0	23.92	0.12	-0.03	24.45	4.2	-1.572	-1.522	-0.003	-0.016	1035.9	222.9	0.0147	1.41	0.66	SL
200.0	21.25	0.08	-0.04	24.45	5.5	-2.657	-2.586	-0.005	-0.036	1034.9	216.7	0.0147	1.58	0.74	SL
100.0	18.98	0.06	-0.02	24.45	6.8	-2.266	-2.197	-0.005	-0.042	1034.9	210.9	0.0147	1.78	0.76	SL
0.0	16.97	0.05	-0.01	24.45	8.0	-2.013	-1.946	-0.004	-0.048	1033.0	205.2	0.0147	2.01	0.79	AN

MRP (BAR) = 16.97