

# Heavy-organic particle deposition from petroleum fluid flow in oil wells and pipelines

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**Abstract:** Suspended asphaltenic heavy organic particles in petroleum fluids may stick to the inner walls of oil wells and pipelines. This is the major reason for fouling and arterial blockage in the petroleum industry. This report is devoted the study of the mechanism of migration of suspended heavy organic particles towards the walls in oil-producing wells and pipelines. In this report we present a detailed analytical model for the heavy organics suspended particle deposition coefficient corresponding to petroleum fluids flow production conditions in oil wells. We predict the rate of particle deposition during various turbulent flow regimes. The turbulent boundary layer theory and the concepts of mass transfer are utilized to model and calculate the particle deposition rates on the walls of flowing conduits. The developed model accounts for the eddy diffusivity, and Brownian diffusivity as well as for inertial effects.

The analysis presented in this paper shows that rates of particle deposition (during petroleum fluid production) on the walls of the flowing channel due solely to diffusion effects are small. It is also shown that deposition rates decrease with increasing particle size. However, when the process is momentum controlled (large particle sizes) higher deposition rates are expected.

**Key words:** Asphaltene, Brownian deposition coefficient, diffusivity, diamondoids, heavy organic particles, paraffin/wax, particle deposition, petroleum fluid, prefouling behavior, production operation, suspended particles, turbulent flow

## 1 Introduction

A common problem faced by the oil industry is the deposition of heavy organics inside production wells, storage vessels, and transfer pipelines. Our studies and experiences have indicated that heavy organic deposition is one of the major factors that increase the cost of production and transportation of petroleum fluids (Mansoori, 1988; Carpentier et al, 2007). Furthermore, miscible flooding of petroleum reservoirs by lean gas, carbon dioxide, natural gas, and other high pressure injection fluids has become an economically viable technique for petroleum production. Introduction of a miscible fluid in petroleum reservoirs, will, in general, produce a number of alterations in petroleum fluid flow and phase behavior and reservoir rock characteristics. One such alteration is the heavy organic precipitation, flocculation and deposition (asphaltenes, diamondoids, etc.), which in most of the observed instances result in plugging or wettability reversal in the conduits (Escobedo and Mansoori, 1995a; 1995b; Branco et al, 2001; Mousavi-Dehghani et al, 2004; Mansoori et al, 2007).

In our recent reports we presented the various causes and effects of phase behavior of petroleum fluids containing heavy organic fractions and their depositions (Mansoori, 2009a; 2009b). It is always preferred to prevent heavy organics deposition during petroleum fluid flows. In cases when heavy organics precipitation can not be prevented we need to understand the fluid flow behavior which is less likely to cause fouling of a conduit. In these cases there is a need for understanding how the precipitated and flocculated particles suspended in the oil will behave under certain flow conditions. This motivated the research presented in this report. Our main objective is to study the behavior of suspended heavy organic particles during flow conditions.

As a preliminary step, the tendency of the crude oil to form solid particles, whenever a miscible solvent is injected into the petroleum reservoir, must be determined. This may be accomplished by using existing experimental techniques (Mousavi-Dehghani et al, 2004; Mansoori et al, 2007) together with predictive models and packages (Branco et al, 2001). These combined experimental/predictive approaches have proven to be a useful tool for design of production and transportation schemes for crude oils prone to heavy organics precipitation, flocculation and deposition.

The study of the behavior of suspended heavy-organic particles during flow conditions has been focused on the production well since flow through a pipeline is only a special

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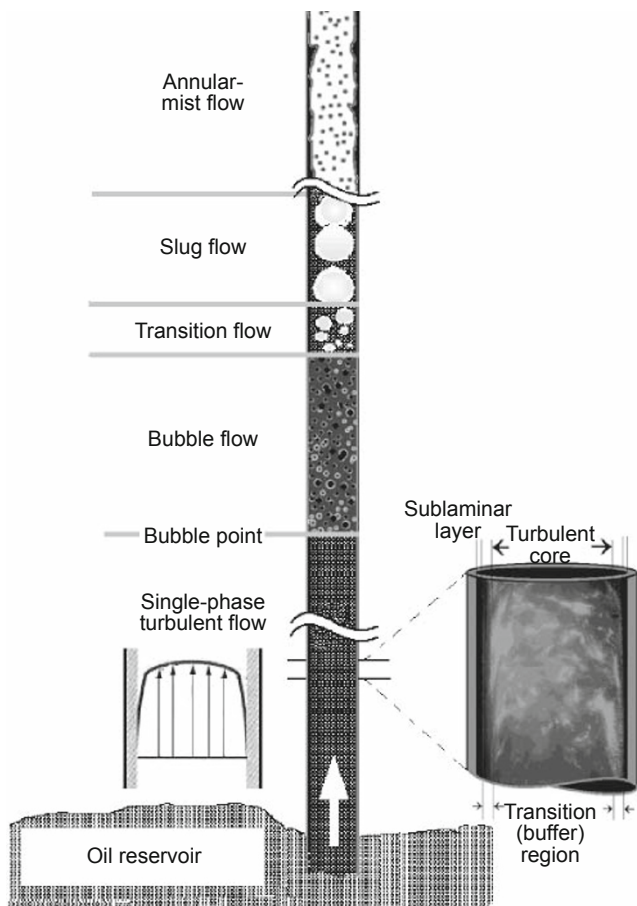
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case of this more general one. A typical production well may be divided into two distinct sections (See Fig. 1):

(I) The pressure region above the bubble point (single-phase) is the emphasis of the present report. Understanding of deposition of particles in this region is especially important for particles which enter into the oil well from the reservoir.

(II) The analysis of the region below the bubble-point pressure (two-phase flow) is the subject of our ongoing research.



**Fig. 1** Illustration of velocity distribution and different flow regimes in the prefoiling single-phase turbulent flow condition in the oil well

In this report we present an analytical model for the heavy organic suspended particle deposition coefficient prior to deposition corresponding to petroleum fluids flow in producing wells condition. In our previous publications (Escobedo and Mansoori, 1995a; 1995b; 2010) we reported various segments of this analytical model some of which contained typographical errors. For the sake of comprehensiveness and to provide the readership with the details of the algebraic manipulations involved we report here the details of the model in its entirety.

## 2 Development of the analytical model

The theory presented here is for a turbulent petroleum fluid system of constant density and viscosity. For petroleum fluid flow in wells this is the case for the region above the bubble pressure where only the liquid phase is the medium for the suspended particles.

Substantial work has been done by many researchers on the topic of particle deposition on the walls of channels or pipes in turbulent flow by many researchers (Lin et al, 1953; Laufer, 1954; Friedlander and Johnstone, 1957; Beal, 1970; Chen and Ahmadi, 1997; Derevich and Zaichik, 1988; Johansen, 1991). The model presented here is a combination of the work performed by the authors mentioned above, modified to be applicable to the deposition of heavy organic particles in turbulent petroleum fluid flow. A key assumption in the development of this model is that fully developed petroleum turbulent flow has a structure as proposed by Lin et al. From experimental observations, they proposed a generalized velocity distribution for turbulent flow of fluids in pipes comprised of three main regions:

- A sublaminar (wall) layer  $0 \leq r^* \leq 5,$
- A buffer layer  $5 \leq r^* \leq 30,$
- A turbulent core  $30 \leq r^*,$

where  $r^* = D_i(V_{avg}\sqrt{f/2} / \nu)$  is a dimensionless distance measured from the wall. This dimensionless distance is a function of the inner pipe diameter,  $D_i$ , fluid average velocity,  $V_{avg}$ , the Fanning friction factor,  $f$ , and the fluid kinematic viscosity,  $\nu$ . The model presented here is for a system of constant density and viscosity. Therefore, it is applicable only to a single phase petroleum fluid flow. This theory can be extended to the region below the bubble point pressure (gas-liquid slug flow, etc.) using reliable expressions for petroleum viscosity and density versus pressure and temperature. The assumption of constant viscosity and constant density is justified since density changes are not appreciable until the bubble point pressure is reached inside the well (or tubing). It is also assumed that suspended particles are of uniform diameter, and that, due to their rather low concentration we can neglect particle-particle interactions. We assume the thickness of the boundary layer is quite small compared to the radius of the pipe as a result we can neglect the wall curvature effects in our model.

We start with reporting the equation which is used to describe the particle flux,  $N_p$ , in terms of the diffusivities and the concentration gradient (Von Karman, 2004), i.e.,

$$N_p = (D^B + D^E) \frac{dC^p}{dr} \tag{1}$$

where  $D^B$  is the Brownian diffusivity;  $D^E$  is the eddy diffusivity;  $C^p$  is the particle concentration; and  $r$  is the radial distance.

The Brownian diffusivity is defined by the following equation:

$$D^B = \frac{k_B T}{3\pi d_p \mu} \tag{2}$$

where  $k_B$  is the Boltzmann constant ( $k_B=1.38066 \times 10^{-23}$  J/K);  $T$  is the absolute temperature;  $d_p$  is the particle diameter; and  $\mu$  is the liquid viscosity of the suspending medium (petroleum fluid). Equation (1) is subject to the boundary condition:

at  $r = S_d$   $C^p = C_{S_d}^p$   
 where  $C_{S_d}^p$  is the particle concentration at  $r = S_d$  and  $S_d$  is the particle "stopping distance" measured from the wall. A particle needs to diffuse only within one stopping distance

from the wall, and from this point on, due to the particle momentum, it would coast to the wall. For small particles the stopping distance is small compared with the boundary layer thickness and consequently diffusion dominates. The proposed correlation for the particle stopping distance is (Friedlander and Johnstone, 1957):

$$S_d = \frac{0.05\rho_p V_{avg} d_p^2 \sqrt{f/2}}{\mu} + \frac{d_p}{2} \quad (3)$$

where  $\rho_p$  is the density of particles;  $V_{avg}$  is the average velocity of petroleum fluids; and  $f$  is the Fanning friction factor.

Equation (1) may be integrated following the procedure for the calculation of temperature drop across a composite wall. We will find the concentration profiles from point to point across the boundary layer. That is, we will calculate the concentration differences through the sublaminal layer, the buffer region and the turbulent core. By adding these concentration differences we can find the overall particle flux in terms of the average and wall concentrations. Before we integrate Equation (1) for  $C^p$ , we need to have expressions for  $N_p$  and  $D^E$  as functions of the radial distance ( $r$ ) for each of the three main regions in the oil well and pipeline, i.e. sublaminal layer, buffer layer and turbulent core.

### 1) Sublaminal Layer

Johansen (1991) proposed the following correlation to express the eddy diffusivity as a function of radial distance ( $r$ ) for the sublaminal layer:

$$D^E = \nu \left( \frac{r^*}{11.15} \right)^3 \quad \text{for } r^* \leq 5 \quad \text{or} \quad \frac{D_{S_d^*-5}^E}{\nu} = \left( \frac{r^*}{11.15} \right)^3 \quad (4)$$

In this equation  $\nu$  is the kinematic viscosity of the flowing petroleum fluid,  $r^*$  is the dimensionless radial distance and  $D_{S_d^*-5}^E$  represents the sublaminal layer ( $r^* \leq 5$ ) eddy diffusivity.

The particle molar flux,  $N_p$ , is assumed to vary linearly from the wall to the center line of the channel, as proposed by Beal (1970):

$$N^p = N_o^p \left( 1 - \frac{2r^*}{D_i^*} \right) \quad (5)$$

In this equation  $N_o^p$  is the particle flux at the wall;  $r^*$  is a dimensionless radial distance and  $D_i^*$  is the dimensionless inner well (or tubing) diameter, both defined by the following equations:

$$r^* = r \left[ \frac{V_{avg} \sqrt{f/2}}{\nu} \right] \quad (6)$$

$$D_i^* = D_i \left[ \frac{V_{avg} \sqrt{f/2}}{\nu} \right] \quad (7)$$

where  $D_i$  is the inner diameter of the well (or tubing);  $V_{avg}$  is the average fluid velocity;  $f$  is the Fanning friction factor; and  $\nu$  is kinematic viscosity of the flowing fluid,  $m^2/s$ . Let us also define the dimensionless stopping-distance,  $S_d^*$  by the

following equation:

$$S_d^* = S_d \left[ \frac{V_{avg} \sqrt{f/2}}{\nu} \right] \quad (8)$$

For the sublaminal layer Equation (1) may be integrated following the procedure for the calculation of temperature drop across a composite wall. We will find the concentration profiles from point to point across the boundary layer. That is, we will calculate the concentration differences through the sublaminal layer, the buffer region and the turbulent core. By adding these concentration differences we can find the overall particle flux in terms of the average and wall concentrations. Note that Equation (4) is only valid for dimensionless radial distances smaller than 5, which is the limit of the sublaminal layer.

Introducing all the new dimensionless variables and the expressions for  $N$  and  $D^E$  into Equation (1), considering that

$dr = dr^* \left[ \frac{\nu}{V_{avg} \sqrt{f/2}} \right]$  we get:

$$N_p = N_o^p \left( 1 - \frac{2r^*}{D_i^*} \right) = \left[ \frac{D^B}{\nu} + \left( \frac{r^*}{11.15} \right)^3 \right] V_{avg} \sqrt{f/2} \frac{dC^p}{dr^*} \quad (9)$$

subject to the following boundary conditions:

$$\begin{cases} \text{at } r^* = S_d & C^p = C_{S_d}^p \\ \text{at } r^* = 5 & C^p = C_5^p \end{cases}$$

Rearranging Equation (9), integrating and applying the above boundary conditions we arrive at the following integral form:

$$C_5^p - C_{S_d}^p = \frac{N_o^p}{V_{avg} \sqrt{f/2}} \left[ \frac{11.15 N_{Sch}^{2/3}}{3} F_1 - \frac{2(11.15)^2 N_{Sch}^{1/3}}{3 D_i^*} F_2 \right] \quad (10)$$

In the above equation  $N_{Sch}$  is the Schmidt number defined as:

$$N_{Sch} \equiv \frac{\nu}{D^B} \quad (11)$$

and  $F_1$  and  $F_2$  are defined by the following expressions:

$$F_1 = \frac{1}{2} \ln \left[ \frac{(1+5\phi)^2}{(1-5\phi)^2} \right] - \frac{1}{2} \ln \left[ \frac{(1+S_d^*\phi)^2}{(1-S_d^*\phi)^2} \right] + \quad (12)$$

$$\sqrt{3} \tan^{-1} \left( \frac{10\phi-1}{\sqrt{3}} \right) - \sqrt{3} \tan^{-1} \left( \frac{2S_d^*\phi-1}{\sqrt{3}} \right)$$

$$F_2 = \frac{1}{2} \ln \left[ \frac{(1-5\phi)^2}{(1+5\phi)^2} \right] - \frac{1}{2} \ln \left[ \frac{(1-S_d^*\phi)^2}{(1+S_d^*\phi)^2} \right] + \quad (13)$$

$$\sqrt{3} \tan^{-1} \left( \frac{10\phi-1}{\sqrt{3}} \right) - \sqrt{3} \tan^{-1} \left( \frac{2S_d^*\phi-1}{\sqrt{3}} \right)$$

where for simplicity we have defined:

$$\phi \equiv \frac{N_{Sch}^{1/3}}{11.15} \tag{14}$$

Equations (10-14) describe the transport of suspended particles in the sublaminal layer to the wall in terms of the concentration difference between the limits  $r^* = S_d^*$  (dimensionless stopping distance) and  $r^* = 5$  (limit of the sublaminal layer).

**2) Buffer Layer**

The next step is the calculation of the particle flux between the concentration at  $r^* = 5$  and  $r^* = 30$  (limit of the buffer layer). The eddy diffusivity expression for the buffer layer is assumed to be:

$$D^E = \nu \left( \frac{r^*}{11.4} \right)^2 - 0.1923 \text{ for } 5 \leq r^* \leq 30 \text{ or}$$

$$\frac{D_{S_d^*-30}^E}{\nu} = \left( \frac{r^*}{11.4} \right)^2 - 0.1923 \tag{15}$$

Integration of Equation (1), using Equation (15) for  $D_{S_d^*-30}^E$  gives:

$$C_{30}^P - C_5^P = \frac{N_o^P}{V_{avg} \sqrt{f/2}} \left[ 11.4\theta \cdot F_3 \right] - \frac{11.4^2}{D_i^*} \ln \left[ \frac{11.4^2 - (30\theta)^2}{11.4^2 - (5\theta)^2} \right] \tag{16}$$

with

$$F_3 = \frac{1}{2} \ln \left[ \frac{11.4 - 30\theta}{11.4 + 30\theta} \right] - \frac{1}{2} \ln \left[ \frac{11.4 - 5\theta}{11.4 + 5\theta} \right] \tag{17}$$

$$\theta \equiv \left[ \frac{N_{Sch}}{(0.1923 \cdot N_{Sch} - 1)} \right]^{1/2} \tag{18}$$

Equation (16) describes the particle transport in terms of the concentration difference between the limits of the buffer layer.

**3) Turbulent core**

The following step is the calculation of the particle transport rate in the turbulent core in terms of the difference between the concentration at  $r^* = 30$  (upper limit of the buffer layer) and the bulk concentration (average conc.). The eddy diffusivity for the turbulent core is taken to be (Johansen, 1991):

$$D^E = 0.4r^* \nu \text{ for } r^* \geq 30 \text{ or } \frac{D_{30-avg}^E}{\nu} = 0.4r^* \tag{19}$$

If we assume that at  $V = V_{avg}$  we have  $C^P = C_{avg}^P$ , then we can integrate Equation (1) using Equation (19) to obtain the following expression for  $C_{avg}^P$ ,

$$C_{avg}^P - C_{30}^P = \frac{N_o^P}{V_{avg} \sqrt{f/2}} \times \left\{ \left( 2.5 + \frac{12.5}{D_i^* N_{Sch}} \right) \ln \left( \frac{1 + 0.4r_{avg}^* N_{Sch}}{1 + 0.4r^* N_{Sch}} \right) - \frac{5r_{avg}^*}{D_i^*} + \frac{5r^*}{D_i^*} \right\} \tag{20}$$

In this expression  $r_{avg}^*$  is the dimensionless radial distance (measured from the wall) where  $V = V_{avg}$ . Equation (20) describes the particle transport in terms of the concentration difference between the bulk (average) and the upper limit of the buffer layer.

So far, we have expressions for the three different regions (sublaminal/wall layer, buffer layer, and turbulent core). Now they may be added together to obtain an expression for  $N_o$  in terms of  $C_{avg}^P, C_{S_d^*}^P$ , average fluid velocity,  $S_d^*$ , and physical parameters of the system.

Until now, only dimensionless stopping distances ( $S_d^*$ ) with values less than 5 have been considered. However, for particles large enough  $S_d^*$  could be greater than 5. If so, then the preceding analysis is not valid under these conditions. This difficulty may be overcome if Equation (1) is integrated between the limits

$$C^P = C_{S_d^*}^P \text{ at } r^* = S_d^*$$

and

$$C^P = C_{30}^P \text{ at } r^* = 30,$$

using the eddy diffusivity correlation for the buffer layer as expressed by equation (15).

Introducing Equation (15) into Equation (1) and integrating using the assumptions noted previously, we get:

$$C_{30}^P - C_{S_d^*}^P = \frac{11.4N_o}{V_{avg} \sqrt{f/2}} \left\{ \theta F_4 - \frac{11.4}{D_i^*} \ln \left[ \frac{11.4^2 - (30\theta)^2}{11.4^2 - (S_d^*\theta)^2} \right] \right\} \tag{21}$$

Such that,

$$F_4 = \frac{1}{2} \ln \left[ \frac{11.4 - 30\theta}{11.4 + 30\theta} \right] - \frac{1}{2} \ln \left[ \frac{11.4 - S_d^*\theta}{11.4 + S_d^*\theta} \right] \tag{22}$$

and  $\theta$  is defined by Eq. (18).

For  $S_d^* = 5$  then Equations (21) and (22) reduce to Equations (16) and (17). Equation (13) still applies to the turbulent core.

**For particles with a dimensionless stopping distance  $0 \leq S_d^* < 5$**

We add Equations (10), (16), and (20), to obtain an expression for the mass transfer (transport) coefficient defined as  $N_o^P / (C_{avg}^P - C_{S_d^*}^P) = K$  with the dimension of velocity in cm/sec. The expression for the deposition coefficient obtained is:

$$K = V_{avg} \sqrt{f/2} \left\{ \frac{11.15^3 \phi^2}{3} F_1 - \frac{11.15^3 \phi}{1.5D_i^*} F_2 + 11.4\theta F_3 + F_5 \right\}^{-1} \tag{23}$$

Parameter  $F_1, F_2$  and  $F_3$  appearing in this equation are the same as defined previously by Equations (12), (13), and (17), respectively and parameter  $F_5$  is given below:

$$F_5 = -\frac{11.4^2}{D_i^*} \ln \left[ \frac{11.4^2 - (30\theta)^2}{11.4^2 - (5\theta)^2} \right] + \left( 2.5 + \frac{12.5}{D_i^* 11.15^2 \phi^2} \right) \ln \left[ \frac{1 + 0.4(11.15^2) r_{avg}^* \phi^2}{1 + 12(11.15^2) \phi^2} \right] + \frac{150 - 5r_{avg}^*}{D_i^*} \tag{24}$$

**For particles with a dimensionless stopping distance  $5 \leq S_d^* < 30$ :**

By combining Equations (20) and (21) and solving for  $K$  we get:

$$K = V_{avg} \sqrt{f/2} / \{11.4\theta F_4 + F_5\} \tag{25}$$

Parameters  $F_4$  and  $F_5$  appearing in this equation are the same as previously defined by Equations (22) and (24), respectively.

**Inertial effects**

In the above analysis we have derived analytical expressions for the mass deposition coefficient for different particle sizes in terms of the dimensionless stopping distance  $S_d^*$ . Next we must account for the inertial effects. We use the following expression to account for inertial effects (Beal, 1970):

$$K^{inert} = \frac{N_0^p}{C_{avs}^p} = \frac{K \cdot p \cdot \mathcal{G}}{K + p \cdot \mathcal{G}} \tag{26}$$

where  $K^{inert}$  is the deposition coefficient (mass transfer coefficient which contains the inertial effects);  $N_0^p$  is the particle flux at the wall (as previously defined);  $p$  is the particle sticking factor (taken equal to unity);  $\square$  is the average velocity of the particles near the walls, and it consists of two parts:

$$\mathcal{G} = \mathcal{G}_{im} + \mathcal{G}^B \tag{27}$$

where  $\mathcal{G}_{im}$  is the particle velocity component due to fluid motion and  $\mathcal{G}^B$  is the component due to particle Brownian motion,

$$\mathcal{G}^B = \left( \frac{k_B T}{3\pi m} \right)^{1/2} \tag{28}$$

with

$$k_B = \frac{1}{4} V_{avg} \sqrt{f/2} \left( \mathcal{G}_{im}^* (d^*/2) + \mathcal{G}_{im}^* (S_d^*) \right)$$

where  $m$  is the particle mass;  $\mathcal{G}_{im}^* (d^*/2)$  is the particle velocity at a dimensionless radial distance  $d^*/2$ ;  $d^*$  is the dimensionless particle diameter; and  $\mathcal{G}_{im}^* (S_d^*)$  is the particle velocity at a dimensionless radial distance  $S_d^*$  (dimensionless stopping distance). The quantity  $\mathcal{G}_{im}^*$  can be calculated using the following correlation proposed by Laufer (1954):

$$\mathcal{G}_{im}^* = 0.05r^* \quad \text{for } 0 \leq r^* \leq 10$$

$$\mathcal{G}_{im}^* = 0.5r^* + 0.0125(r^* - 10) \quad \text{for } 10 \leq r^* \leq 30$$

The particles can be anywhere between  $r^* = d^*/2$  and  $r^* = S_d^*$ . Considering this we are to choose the appropriate

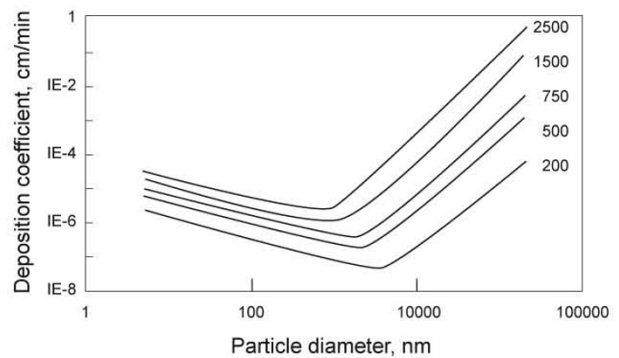
expression for  $\mathcal{G}_{im}^*$ .

The analysis for particle deposition onto the walls of a flowing channel from turbulent fluid streams is concluded by taking into account the inertial effects as in Equation (25). At this point all the phenomena influencing the deposition rate (Brownian diffusivity, eddy diffusivity, and inertial effects) have been taken into account.

**3 Predictions for particle deposition in wells and tubings**

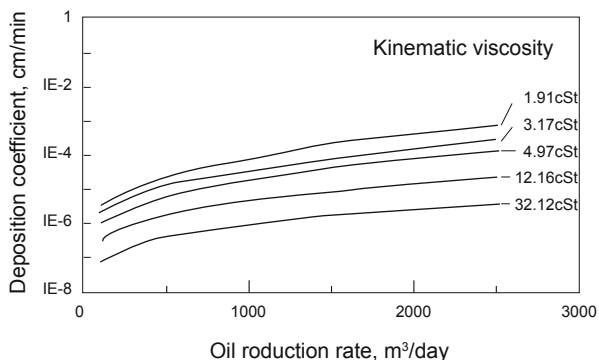
In our previous publications (Escobedo and Mansoori, 1995a; 1995b; 2010) we compared the predictions of the proposed model with the available experimental data and the other models. Since experimental data for particle deposition from turbulent petroleum fluid flows were scarce we used the deposition data from aerosols to verify our model. As it was shown, the results of our analysis for particle deposition from turbulent fluid streams were in very good agreement with the experimental deposition rates for aluminum and iron particles in air. The predictions of the present model showed a better agreement with the mentioned experimental data than the models proposed earlier. We believe our model is suitable for *a priori* predictions of the particle deposition coefficient from turbulent petroleum fluid flow.

Fig. 2 shows the predicted deposition coefficients for particles as a function of particle diameter suspended in a light petroleum fluid with 30.21 °API gravity (corresponding to 0.875 SG) and the kinematic viscosity of 11 centi-Stokes (cSt) at various production rates. Fig. 3 shows the predicted deposition coefficients values for petroleum fluids with different kinematic viscosities (and at different °APIs) and containing suspended particles of 1,000 nm in diameter.



**Fig. 2** Effect of particle size on the deposition coefficient for a 30.21 °API petroleum fluid with a kinematic viscosity of 11 cSt at various production rates in m<sup>3</sup>/day

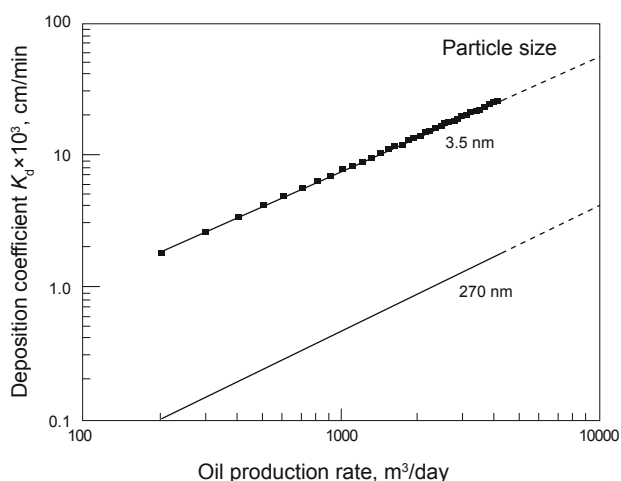
The particle sizes analyzed ranged from 5 to 200,000 nm. According to Fig. 2, deposition coefficients are generally small except at high production rates and for very large particles. Note a minimum after which the deposition coefficient increases more rapidly with increasing particle diameter. This is due to the fact that at the minimum point the deposition process is momentum controlled. According to Fig. 3, particle deposition from turbulent petroleum fluid flow is very small when the diameter of the suspended particles



**Fig. 3** Effect of petroleum production rate on the deposition coefficient of 1 micron (1000 nm) diameter suspended particles in petroleum fluids of various kinematic viscosities ranging from 1.91 - 32.12 cSt

is less than 1,000 nm. Fig. 2 indicates a decrease in the deposition coefficients with increasing kinematic viscosity and an increase in the deposition coefficient with increasing production rate. The lighter the petroleum fluid, the higher the probability of having particle deposition.

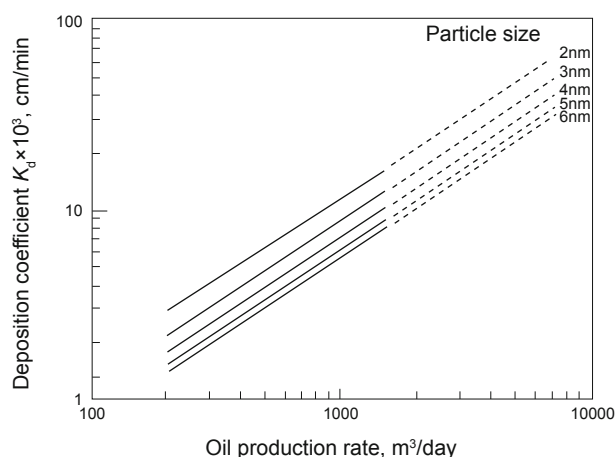
Fig. 4 shows the predicted deposition coefficients for particles as a function of the crude oil production rate for two extreme particle sizes, 3.5 nm (corresponding to asphaltene steric-colloidal particles in original crude oil) and 270 nm (corresponding to flocculated asphaltene aggregates). As it was mentioned the present analysis is for the single-phase region of the well-tubing (region before the bubble point pressure is reached). One can notice from this figure the very small values of deposition coefficients for 3.5 nm particles, for the volumetric flow rates of up to 3,000 m<sup>3</sup>/day. Fig. 4 also indicates that deposition coefficients increase with increasing velocities. However, when the oil production rate is about 10,000 m<sup>3</sup>/day, deposition coefficient is roughly 0.06 cm/min. Therefore we would expect sizeable amounts of deposition at high production rates.



**Fig. 4** Predicted deposition coefficients as a function of the crude oil production rates. It is calculated for particles with a diameter of 3.5 nm suspended in a 37.8 °API petroleum fluid

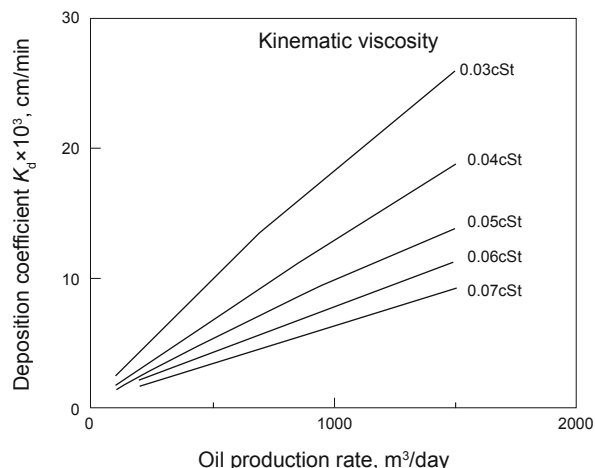
With the idea in mind of finding another factor that could increase the tendency of the particles to deposit, we examined the effect of particle size on deposition coefficient.

This is shown in Fig. 5 where we can notice that deposition coefficients decrease with increasing particle size due to the fact that Brownian diffusion is inversely proportional to the particle size of the diffusing species. However, with increasing particle size (or mass) the momentum increases, so that particle momentum effects become more important. It is expected a minimum deposition rate for a given particle size, beyond which the process is essentially momentum controlled, resulting in higher deposition rates. For a radius of 270 nm, we still notice very small values (see Fig. 4), this means that heavy organic particle size must become much larger in order to exhibit high deposition rates.



**Fig. 5** Effect of particle size on deposition coefficient for a 37.8 °API petroleum fluid

We performed model predictions varying the kinematic viscosity of the crude oil to study the effect of this parameter on the deposition coefficient. This was done because kinematic viscosity decreases with increasing temperature, and because it is a function of the API gravity of the crude oil. Fig. 6 shows the predicted values, and we can see a decrease in the deposition coefficient with increasing kinematic viscosity. This means that the lighter the petroleum fluid is the higher the probability of having heavy organic deposition will be. However, these predicted values are still very small.



**Fig. 6** Effect of kinematic viscosity on deposition coefficient

## 4 Conclusions

The model developed for the particle deposition onto the walls of a pipe from a turbulent petroleum fluid stream in oil wells and pipelines was used to predict the deposition coefficient from turbulent flow production operations. The effect of particle size on the deposition coefficient was investigated, finding that when the deposition process is diffusion-controlled (particles with a diameter < 1,000 nm) the predicted values are very small. However, when the deposition process is momentum controlled (particles with a diameter > 1,000 nm) the predicted values for the deposition coefficient increase more rapidly with increasing particle diameter. We also investigated the effect of petroleum fluid kinematic viscosity on the deposition coefficient. We found that the deposition coefficient decreases with increasing petroleum fluid kinematic viscosity. For kinematic viscosity of 12.16 cSt the predicted deposition coefficient is negligible for suspended particles of 1,000 nm. We also found that the deposition coefficient increases with increasing production rate. This is due to the fact that at larger production rates the amount of eddy diffusion is bigger. The proposed model can be used for various cases of the behavior of heavy organics particle deposition from turbulent petroleum flows so long as the particles are neutral, their sizes are stable, there are no particle-particle interactions and there are no phase transitions occurring in the flow. However, in cases where such changes are occurring in the system this model will require appropriate modifications as presented and applied elsewhere (Mansoori G A, ASPHRAC: A comprehensive package of computer programs and database which calculates various properties of petroleum fluids containing heavy organics. [www.uic.edu/~mansoori/ASPHRAC\\_html](http://www.uic.edu/~mansoori/ASPHRAC_html)).

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