

A solution is obtained for the distribution of a trace component in a Oldroyd liquid for various forms of the pressure function: pulsed, stepwise, and periodic.

The model of [1, 2] for the dispersion of a material in a flow of viscous liquid is applicable also to laminar and turbulent flows provided that the diffusion processes are fairly prolonged. The use of this model is restricted to the range $W_m R/D \geq 6.9$ for the time factor, but this is not always obeyed for real processes (particularly ones of nonstationary type). In [3, 4], this restriction does not apply; they deal with nonstationary diffusion for any instant. For a rheologically complex medium [5] for which the relaxation time is $\lambda = 10^{-3} - 10^0$ sec, the duration of the diffusion processes are comparable with the transient-response times. Under those conditions, the diffusion processes will be dependent not only on the rheological ones [6] but also on the relaxation characteristics.

We consider the distribution of an impurity in a flow of Oldroyd liquid [7] with relaxation behavior:

$$\left(1 + \lambda_1 \frac{\partial}{\partial t}\right) \frac{\partial \tau}{\partial t} = v \left(1 + \lambda_2 \frac{\partial}{\partial t}\right) \frac{\partial \dot{\gamma}}{\partial t}. \quad (1)$$

The one-dimensional equation of motion in a circular tube then takes the form

$$\left(1 + \lambda_1 \frac{\partial}{\partial t}\right) \frac{\partial \omega}{\partial t} = \left(1 + \lambda_1 \frac{\partial}{\partial t}\right) f(t) + v \left(1 + \lambda_2 \frac{\partial}{\partial t}\right) \left(\frac{\partial^2 \omega}{\partial r^2} + \frac{1}{r} \frac{\partial \omega}{\partial r}\right). \quad (2)$$

The function $f(t) = (-1/\rho)(\partial p/\partial x)$ is dependent on time. We assume that $f(t)$ is one of the commoner functions: $f_1(t) = A\delta(t)$ is a Dirac delta function ($k = 1$), $f_2(t) = AH(t)$ a Heaviside step function ($k = 2$), and $f_3(t) = A \cos \omega t$ a harmonic function ($k = 3$). Then the velocity distribution at any instant is defined as

$$\omega(r, t) = \sum_{i=1}^{\infty} a_i^k(t) J_0(r, \xi_i) \quad (k = 1, 2, 3). \quad (3)$$

Here

$$\begin{aligned} a_i^1 &= \frac{2}{R\xi_i J_1(R, \xi_i) \lambda_1 (\alpha - \beta)} [(1 + \lambda_1 \alpha) \exp(\alpha t) - (1 + \lambda_1 \beta) \exp(\beta t)]; \\ a_i^2 &= \frac{2A}{R\xi_i J_1(R\xi_i) \lambda_1 (\alpha - \beta)} \left[\frac{\exp(\alpha t)}{\alpha} - \frac{\exp(\beta t)}{\beta} + \frac{\alpha - \beta}{\alpha\beta} + \right. \\ &\quad \left. + \lambda_1 (\exp(\alpha t) - \exp(\beta t)) \right]; \\ a_i^3 &= \frac{2A}{R\xi_i J_1(R\xi_i) \lambda_1 (\alpha - \beta)} \left[\frac{(1 + \lambda_1 \alpha) (\alpha \exp(\alpha t) - \alpha \cos \omega t)}{\alpha^2 + \omega^2} + \right. \\ &\quad \left. + \frac{\omega \sin \omega t}{\alpha^2 + \omega^2} - \frac{(1 + \lambda_1 \beta) (\beta \exp(\beta t) - \beta \cos \omega t + \omega t + \omega \sin \omega t)}{\beta^2 + \omega^2} \right] / 2\lambda_1; \\ \beta &= [-(1 + v\lambda_2 \xi_i^2) - \sqrt{(1 + v\lambda_2 \xi_i^2)^2 - 4\lambda_1 v \xi_i^2}] / 2\lambda_1; \\ \alpha &= [-(1 + v\lambda_2 \xi_i^2) + \sqrt{(1 + v\lambda_2 \xi_i^2)^2 - 4\lambda_1 v \xi_i^2}] / 2\lambda_1; \end{aligned}$$

and ξ_i are the roots of the equation for the Bessel function $J_0(R\xi_i) = 0$. The mean instantaneous flow speed is

$$W_m(t) = \frac{2}{R^2} \int_0^R \omega(r, t) r dr = \frac{2}{R} \sum_{i=1}^n \frac{a_i^b(t)}{\xi_i} J_1(R\xi_i). \quad (4)$$

If a trace of material is injected into the flow at the initial instant, one can determine the concentration $c(t, x, r)$ at any instant by using the usual equation of convective diffusion [8]:

$$\frac{\partial c}{\partial t} + \omega(r, t) \frac{\partial c}{\partial x} = \frac{D_0}{r} \frac{\partial}{\partial r} \left(r \frac{\partial c}{\partial r} \right). \quad (5)$$

The following is the balance equation for matter in a coordinate system moving at the mean flow speed $W_m(t)$:

$$\pi R^2 \frac{\partial c_m}{\partial t} + \frac{\partial Q}{\partial x} = 0, \quad c_m = c_m(t, x) = \frac{2}{R^2} \int_0^R r c dr, \quad (6)$$

$$Q = 2\pi \int_0^R [\omega(r, t) - W_m(t)] r c dr. \quad (7)$$

We can take Taylor's assumption as correct for the instantaneous values of the time and write (5) as

$$[\omega(r, t) - W_m(t)] \frac{\partial c_m}{\partial x} = \frac{D_0}{r} \frac{\partial}{\partial r} \left(r \frac{\partial c}{\partial r} \right); \quad (8)$$

The left side of (8) is dependent on the variable r , and it is therefore simple to integrate it in order to determine the concentration profile in terms of the mean concentration over the flow section $\partial c_m / \partial t$; we get

$$c(t, x, r) = c_0(t, x) + \frac{\partial c_m}{\partial x} \int_0^r \frac{dr'}{D_0 r'} \int_0^{r'} [\omega(r, t) - W_m(t)] r dr. \quad (9)$$

From (9) we transform (3) to

$$Q = -\pi R^2 D \frac{\partial c_m}{\partial x}, \quad (10)$$

where the effective diffusion coefficient D is determined as

$$D = \frac{2}{D_0 R^2} \int_0^R \frac{dr'}{r} \left[\int_{r'}^R [\omega(r, t) - W_m(t)] r dr \right]^2. \quad (11)$$

Use of the velocity patterns of (3) and (4) for flows in relaxing systems in (11) for various pressure functions enables one to find D ; here one has to incorporate the condition for orthogonality of the eigenfunctions $J_0(\xi_i, r)$ and use the major properties of Bessel functions [9]:

$$\int_0^R r^2 J_1(r\xi_i) dr = \frac{2R}{\xi_i^2} J_1(R\xi_i), \quad \int_0^R r J_1(r\xi_i) J_1(r\xi_j) dr = \frac{R^2}{2} [J_1(R\xi_i)]^2.$$

It is characteristic that the effective diffusion coefficient is dependent not only on time but also on the relaxation characteristics: the relaxation time and the retardation time:

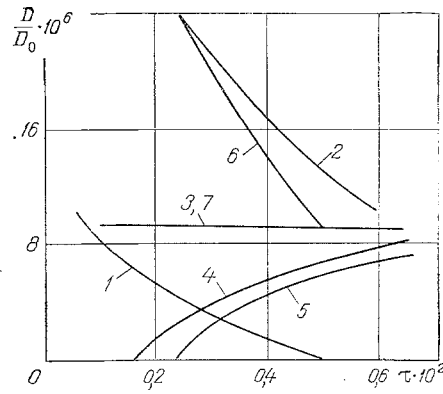


Fig. 1. Variation in the relative effective diffusion coefficient with time for various pressure functions: pulse (1); stepped ($\lambda_1 \neq \lambda_2 \neq 0$) (2), and ($\lambda_1 = \lambda_2 = 0$) (3), sinusoidal ($\omega = 1 \text{ sec}^{-1}$, $\lambda_1 \neq \lambda_2 \neq 0$) (4), ($\omega = 1 \text{ sec}^{-1}$, $\lambda_1 = \lambda_2 = 0$) (5), ($\omega = 0.01 \text{ sec}^{-1}$, $\lambda_1 \neq \lambda_2 = 0$) (6), ($\omega = 0.01 \text{ sec}^{-1}$, $\lambda_1 = \lambda_2 = 0$) (7).

$$D = \frac{2}{D_0 R^2} \left\{ \sum_{i=1}^{\infty} \sum_{j=1}^{\infty} a_i^h(t) a_j^h(t) J_1(\xi_i R) J_1(\xi_j R) \left[\frac{R^2}{4\xi_i^2 \xi_j} - \frac{2}{\xi_i \xi_j^3} - \frac{2}{\xi_i^3 \xi_j} \right] + \sum_{i=1}^{\infty} [a_i^h(t)]^2 \frac{R^2}{2\xi_i^2} J_1(\xi_i R) \right\} \quad (12)$$

or

$$D = \frac{2}{D_0 R^2} \left\{ \sum_{i=1}^{\infty} \sum_{j=1}^{\infty} a_i^h(t) a_j^h(t) \left[\frac{R^2}{4\xi_i^2 \xi_j^2} - \frac{2}{\xi_i \xi_j^3} - \frac{2}{\xi_i^3 \xi_j} \right] + \sum_{i=1}^{\infty} [a_i^h(t)]^2 \frac{R^2}{2\xi_i^2} \right\}. \quad (13)$$

In the limit where the relaxation behavior of the liquid is so slight by comparison with the duration of the transient state, we have $\lambda_1 \rightarrow 0$, $\lambda_2 \rightarrow 0$, $\lambda_1(\alpha - \beta) \rightarrow 1$, $\alpha \rightarrow -\nu\mu^2/R^2$, $\beta \rightarrow -\infty$, $\mu_i = \xi_i R$, and the coefficients $a_i^h(t)$ in (13) take the form

$$\begin{aligned} a_i^1 &= 2A \exp[-\nu\mu_i R^{-2}t/\mu_i], \\ a_i^2 &= \frac{2A}{\mu_i} \left[\frac{\exp(-\nu\mu_i t/R^2)}{-\nu\mu_i^2/R^2} - \frac{R^2}{\nu\mu_i^2} \right], \\ a_i^3 &= \frac{2A}{\mu_i} \left[\{(-\nu\mu_i^2/R^2) \exp(-\nu\mu_i^2/R^2) t + (\nu\mu_i^2/R^2) \cos \omega t + \right. \\ &\quad \left. + \omega \sin \omega t\} / \{(\nu\mu_i^2/R^2)^2 + \omega^2\} \right]. \end{aligned}$$

These expressions with (10) show that the effective diffusion coefficient very rapidly acquires a steady-state value for $\lambda_1 = \lambda_2$ and is dependent not only on time.

If the pressure in the system changes stepwise in accordance with the Heaviside function, then for $t \rightarrow \infty$ we get from (13) an expression for the effective diffusion coefficient [1].

We made quantitative estimates of the effects of the relaxation characteristics on the diffusion processes by calculating for the flow with parameters $\nu = 4.8 \text{ cm}^2/\text{sec}$, $\lambda_1 = 2.4 \text{ sec}$; $\lambda_2 = 0.6$; $D_0 = 10^{-5} \text{ cm}^2/\text{sec}$; $R = 0.2 \text{ cm}$; $l = 10^3 \text{ cm}$; the amplitude value of the pressure difference for the three functions was $A = \Delta P_0 = 10^5 \text{ Pa}$.

Figure 1 shows the numerical results as the relative value of the effective diffusion coefficient D/D_0 in terms of the dimensionless time $\tau = D_0 t/R^2$; there is a maximal increase in

the mixing rate at the start for the pulse function, with a subsequent fall to zero (curve 1). If a Heaviside function applies, a steady-state value of the mixing rate is established in a relaxing system for a certain time interval (curve 2). For a liquid not having relaxation characteristics, the steady-state values are attained almost instantly (curve 3). A similar situation is observed for a sinusoidal pressure function at frequencies $\omega < 0.01 \text{ sec}^{-1}$ (curves 6 and 7). At frequencies $\omega \geq 1 \text{ sec}^{-1}$, the rheological characteristics have less effect on the establishment of the steady-state value (curves 4 and 5).

Therefore, the diffusion processes in a flow of relaxing liquid are dependent on the hydrodynamic characteristics and the relaxation ones. One can vary these characteristics to adjust the diffusion in a relaxing medium.

NOTATION

W_m , mean flow velocity; $w(r, t)$, instantaneous flow velocity; R , tube radius; x , coordinate along axis; D_0 , molecular diffusion coefficient; D , effective diffusion coefficient; t , time; λ_1, λ_2 , relaxation and retardation times; ρ , density; ν , kinematic viscosity; τ , shear stress; $\dot{\gamma}$, shear rate; c , concentration; c_0 , axial concentration; c_m , mean concentration; Q , flow rate; $\partial P / \partial x$, pressure gradient; J_0 , zero-order Bessel function; J_1 , Bessel function of first order, real argument; ξ_1 , roots of Bessel function; A , oscillation amplitude; ω , frequency; δ , Dirac delta function; H , Heaviside function.

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