

# UNSTEADY CONVECTIVE DIFFUSION IN VISCOELASTIC FLUID FLOWING THROUGH A TUBE

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**Abstract** The longitudinal dispersion of passive contaminant molecules released in unsteady viscoelastic fluid flowing through a tube is examined. A finite difference implicit scheme is adopted to solve the Aris integral moment equations arising from the unsteady convective-diffusion equation for all time period. Here it is shown how the injected material spread due to the shear effect in the viscoelastic fluid caused by the combined action of periodic flow and lateral diffusion about its mean position, centre of gravity of mass moves and the mean concentration distribution approaches to Gaussianity, when the contaminant is initially uniform over the cross-section of the tube and the Péclet number is large. The analysis reveals that for viscoelastic fluid the dispersion coefficient changes cyclically with a double frequency period and reaches asymptotically a stationary state after a certain time as in the case of a Newtonian fluid, and its increases with the viscoelastic parameter. Further, it shows that there is a remarkable similarity between the mean concentration distribution of solute in a Newtonian and non-Newtonian fluid.

**Keywords:** longitudinal dispersion, non-Newtonian fluid, viscoelastic fluid, oscillatory flow, finite-difference scheme

## 1. INTRODUCTION

The study of dispersion of contaminants in a fluid is a part of Environmental Fluid Mechanics which has become essential in controlling pollution of rivers, environment, etc. due to the release of contaminating materials from different sources. Now-a-days this study, which gives an insight into dispersion phenomena of passive contaminants in solvents, has primary importance in industrial and technological fields.

In his classic paper, Taylor [1] pointed out that in longitudinal dispersion of soluble matter in a moving fluid, solute is more slowly dispersed by molecular or turbulent diffusion alone than the dispersion due to 'shear effect' caused by combined effects on convection and lateral diffusion. Aris [2] subsequently proposed an idea of moment method in solving the model removing the restrictions imposed by Taylor and studied the asymptotic behaviour of second moment about the mean. Barton [3] resolved certain technical difficulties in Aris' method and obtained the solutions of second and third moment equations of the distribution of solute which are valid for all time. All the work mentioned above was based on steady flow.

The longitudinal dispersion of a solute in time-dependent flow due to periodic pressure gradient in an infinite tube was first studied by Aris [4] using his method of moments, and his analysis was limited to asymptotically large time after the injection of the solute. Mukherjee and Mazumder [5] extended the Aris-Barton theory for studying the all-time evolution of the second central moment of dispersion of passive contaminant in the shearing current due to the combined effect of steady and periodic flows within a conduit of uniform cross-section. The solution was based on the method of separation of variables, which depends on a certain eigenvalue problem with a discrete spectrum of eigenvalues. Mazumder and Das [6] extensively studied the effect of boundary absorption on the axial dispersion of contaminant cloud released in pulsatile tube flow. Using a numerical scheme they computed the effective diffusivity for different values of absorption parameter for all time periods. The problem of dispersion phenomenon in time-dependent flows within

conduits has been studied by Chatwin [7], Smith [8], Jimenez and Sullivan [9], Yasuda [10], Mazumder and Das [11] and others.

Fan and Hwang [12] first extended the analysis of Taylor to the dispersion of solute in non-Newtonian (Ostwald-de Waele) fluid and they showed that Taylor's method can be applied in the case of non-Newtonian fluid. Fan and Wang [13] also studied Taylor's analysis of dispersion in the flow of a Bingham plastic and an Ellis model fluid and showed that Aris' modification of Taylor's analysis can be applied in the case of non-Newtonian fluid. Using the Taylor conceptual model, Erdoğan [14] studied the dispersion in non-Newtonian flow at low flow rates. He showed that the longitudinal dispersion coefficient depends on the ratio of the yield stress to the wall stress. When the ratio is zero it is equivalent to a Newtonian fluid. For a ratio greater than zero, the dispersion coefficient first increases and then decreases with the increase of the ratio, and the longitudinal dispersion coefficient becomes zero when the value of the ratio reaches 1. He also studied the effect of a Bingham plastic and Ostwald-de Waele fluids on dispersion and observed that initially the dispersion coefficient increases upto a certain value and then it decreases with an increase in the non-Newtonian parameter. Dispersion in Eyring and Reiner-Philippoff model fluids has been studied by Ghoshal [15] using Taylor's analysis but he did not discuss anything in dispersion phenomenon as in his study some integrals were un-evaluated. Shah and Cox [16] and Gupta and Mazumder [17] solved the problem completely and showed that the dispersion coefficient decreases with the increase of the Eyring model parameter. However, all the investigations on non-Newtonian fluid models mentioned above were asymptotically valid for a large time after the injection of the solute. Subramanian and Gill [18] explored the generalized dispersion model valid for all time period to study the spreading of solutes in non-Newtonian fluid (Ostwald-de Waele), but they confined their analysis to the steady laminar flow in a tube. They found that the dispersion coefficient at any given time decreases with increasing the non-Newtonian parameter.

To the best of our knowledge, the dispersion of a solute in oscillating flow of a viscoelastic fluid has not been studied in the literature. Our main objective of the present paper is to explore the dispersion phenomenon of a solute in Maxwell linear model of viscoelastic fluid (Bird *et al.* [19]) within a tube when the flow is oscillatory due to the periodic pressure gradient. More precisely, it is shown for all time period how the spreading of tracers is influenced by the combined action of characteristic relaxation time and the frequency of oscillation about the mean position, the center of mass of slug moves and the behaviour of mean concentration distributions approaches to normality, when the contaminant is initially uniform over the cross-section of the tube and the Péclet number is large. The motivation of the study of viscoelastic oscillatory flows stems mainly from the important application, namely the dispersion of tracers in pulsatile blood streams and the mass transfer in polymer solutions. In particular, a water-solution of polyacrylamide is a well-known viscoelastic fluid.

## 2. MATHEMATICAL FORMULATION

If the convected derivative is written in full then the constitutive equation of the Maxwell model of a viscoelastic fluid which is a superposition of the Hookean solid and Newtonian liquid with zero retardation time is given by (Oldroyd [20]),

$$\tau_{ij} + t_0 \left( \frac{\partial \tau_{ij}}{\partial t} + u^k \tau_{i,j,k} + u_i^k \tau_{kj} + u_j^k \tau_{ik} \right) = -\mu (u_{i,j} + u_{j,i}) \quad (1)$$

Here  $t_0$  is a characteristic relaxation time for the fluid,  $u_i^j$  is the velocity in the  $i$ th direction,  $\mu$  is the coefficient of viscosity of the fluid, and  $\tau_{ij}$  is the shear stress.

Consider an oscillatory fully developed incompressible laminar flow of the above fluid in a straight circular pipe with uniform cross section of radius  $R$ . As the flow is induced by the periodic pressure difference along the axial direction, the flow will be unidirectional. So the velocity has only axial component  $u_z$ , depending on radial coordinate  $r$  and time  $t$ . Also the stresses have  $r$ - $z$  component  $\tau_{rz}$  only. Taking this into account further simplification is

possible if we take the linearized equation as well as if we consider slow motion of the fluid. So the proper invariant derivative in equation (1) reduces to the partial time derivative.

Neglecting the convective term  $u' \cdot \nabla u'$  of the momentum equation, the linearized equation of motion in two-dimensional cylindrical form is given by, (see Bittl *et al.* [19])

$$\rho \frac{\partial u'_z}{\partial t'} = -\frac{\partial p'}{\partial z'} - \frac{1}{r'} \frac{\partial}{\partial r'} (r' \tau_{rz}) \quad (2)$$

and linearized Maxwell model is

$$\tau_{rz} + t_0 \frac{\partial \tau_{rz}}{\partial t'} = -\mu \frac{\partial u'_z}{\partial r'} \quad (3)$$

where  $\rho$  is the density of the fluid,  $p'$  is the pressure and  $t_0 = \mu/G$ ,  $G$  is the modulus of shear rigidity.

Eliminating  $\tau_{rz}$  from equations (2) and (3) we have

$$\rho t_0 \frac{\partial^2 u'_z}{\partial t'^2} + \rho \frac{\partial u'_z}{\partial t'} = -t_0 \frac{\partial^2 p'}{\partial t' \partial z'} - \frac{\partial p'}{\partial z'} + \mu \frac{1}{r'} \frac{\partial}{\partial r'} \left( r' \frac{\partial u'_z}{\partial r'} \right) \quad (4)$$

The boundary conditions are,

$$u' = 0 \quad \text{at} \quad r' = R \quad (5)$$

and the velocity at the center of the pipe is finite.

As the flow is oscillatory, velocity and pressure of the fluid can be written as of the following form,

$$u'_z = \text{Real} \{ u_0(r) e^{i\omega t'} \}, \quad -\frac{\partial p'}{\partial z'} = b_0 \text{Real} \{ e^{i\omega t'} \} \quad (6)$$

Using the expression of (6), the solution of equation (4) subject to the boundary condition (5) for periodic flow is:

$$u'_z = \text{Real} \left[ \frac{b_0}{i\omega\rho} \left( 1 - \frac{J_0(kr')}{J_0(kR)} \right) e^{i\omega t'} \right] \quad (7)$$

where,  $k^2 = -i\omega\rho(1 + i\omega t_0)/\mu$ , and  $\omega$  is frequency of oscillation when the local fluid motion is a sinusoidal function.

The velocity from equation (7) can be written in dimensionless form as:

$$u = -\text{Real} \left[ \frac{i}{\alpha} \left\{ 1 - \frac{J_0(rk_1)}{J_0(k_1)} \right\} e^{i\alpha St} \right], \quad (8)$$

where  $k_1 = \sqrt{-i\alpha(1 + \alpha T_0)}$ ,  $\alpha = \omega R^2/\nu$  is the dimensionless oscillation Reynolds number or frequency parameter,  $S = \nu/D$  is the Schmidt number,  $T_0 = \nu t_0/R^2$  is the dimensionless elasticity parameter,  $u_z = u'_z/U$  is axial velocity in which  $U = R^2 b_0/\mu$  is the time averaged velocity, and  $r = r'/R$  is the dimensionless radial distance. If the dimensionless elasticity number  $T_0$  is zero, the velocity equation (8) corresponds to the oscillatory Newtonian fluid flow through a tube (Uchida [21], Schlichting [22]). The oscillating flow of a viscoelastic fluid is characterized by dimensionless number  $\alpha$ ,  $S$  and  $T_0$ . Here  $\alpha = \omega R^2/\nu$  is a measure of the ratio of time necessary for shear-wave propagation across the tube section to the period of oscillation or the ratio of pipe radius to the Stokes-layer thickness. The Schmidt number  $S$  is the ratio of viscous diffusion and molecular diffusion; and  $\alpha S$  is the measure of the ratio of the characteristic time of transverse diffusion to the period of oscillation. The dimensionless viscoelastic parameter  $T_0 (= t_0/(R^2/\nu))$  is the ratio of the fluid relaxation time to the time taken for shear-wave propagation over the cross-section of the tube. Figure 1(a-d) presents the velocity profiles against the frequency parameter  $\alpha$  in different flow phases for various values of  $T_0$ , and  $r = 0.5$ . For a Newtonian fluid ( $T_0 = 0$ ), velocity decreases with increase in  $\alpha$ , whereas for a viscoelastic fluid ( $T_0 > 0$ ), it increases upto a certain  $\alpha$ , then decreases (Khabakhpasheva *et al.* [23]) for the phases  $\alpha St = 0, \pi$ . The increase of velocity with viscoelastic parameter  $T_0$  seems to be due to the presence of elasticity in the liquid (Beard and Walters [24]). For phases  $\alpha St = \pi/2$ , and  $3\pi/2$ , it is oscillatory in nature upto a certain

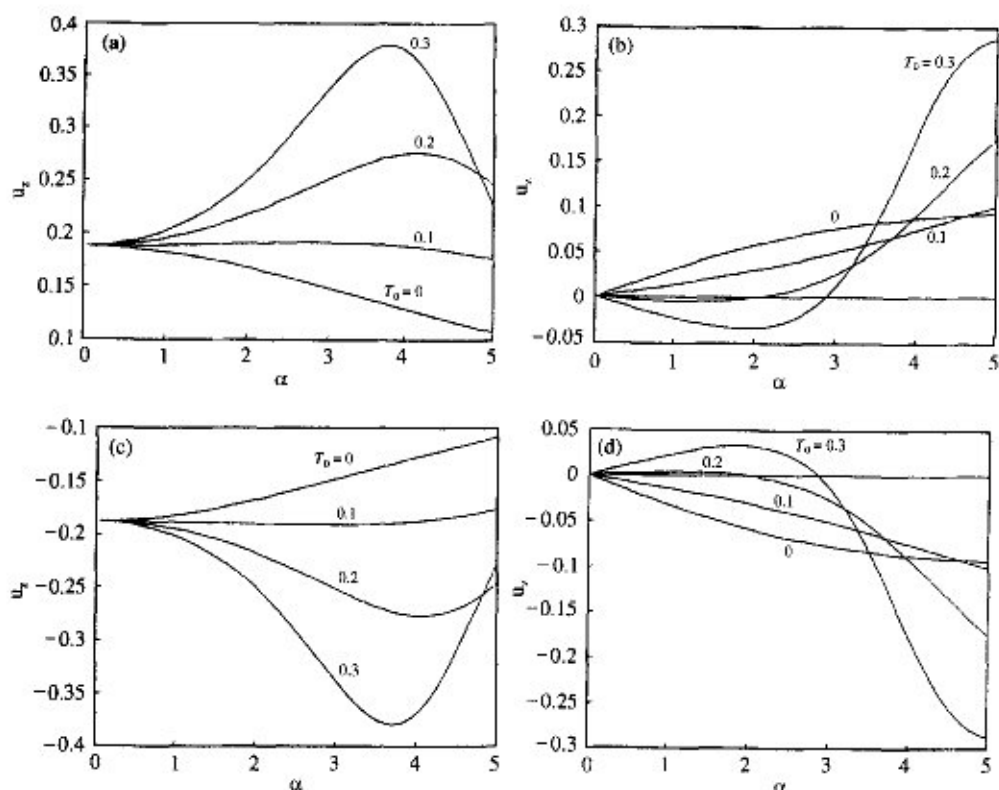


Fig. 1. Oscillatory velocity  $u_x$  of a viscoelastic fluid against frequency parameter  $\alpha$  for different phase: (a)  $\alpha St = 0$ , (b)  $\alpha St = \pi/2$ , (c)  $\alpha St = \pi$ , (d)  $\alpha St = 3\pi/2$ , at  $r = 0.5$ .

value of  $\alpha$ , then it increases with increase in  $T_0$  at large  $\alpha$ . It may be mentioned here that the velocity profile for a pulsating Poiseuille flow of a non-Newtonian fluid studied by Rajagopal and Sciuba [25] is somewhat consistent with the present velocity profile for a small value of frequency parameter. It is also observed from the figure that the velocity profile in the flow phases is more flattened with deceleration than in the phases with acceleration.

If a slug is released in the above mentioned periodic flow in a tube, the concentration  $C(t, r, z)$  of the solute, with constant molecular diffusivity  $D$ , satisfies the non-dimensional convective-diffusion equation of the form

$$\frac{\partial C}{\partial t} + \text{Pe} u_x \frac{\partial C}{\partial z} = \left[ \frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial C}{\partial r} \right) + \frac{\partial^2 C}{\partial z^2} \right] \quad (9)$$

where  $z = z'/R$  and  $\text{Pe} = RU/D$ ,  $u_x = u_x'/U$ ,  $t = Dt'/R^2$ ,  $r = r'/R$ . Here  $u_x$  is the periodic velocity due to imposed periodic pressure gradient,  $U$  is the reference velocity, and  $\text{Pe}$ , the Péclet number is the ratio of the characteristic time of the diffusion process ( $R^2/D$ ) to the convective process ( $R/U$ ).

The initial and boundary conditions are

$$\left. \begin{aligned} C(0, r, z) &= \delta(z), \\ \frac{\partial C}{\partial r} &= 0 \text{ at } r = 1, \\ C &\text{ finite at all points,} \\ z^m \frac{\partial^m C}{\partial z^m} &\rightarrow 0 \text{ as } |z| \rightarrow \infty \text{ for } m, n = 0, 1, 2, \dots, \\ \frac{1}{\pi} \int_0^1 \int_0^{2\pi} \int_{-\infty}^{\infty} r C(0, r, z) \, dr \, d\theta \, dz &= 1, \end{aligned} \right\} \quad (10)$$

where  $\delta(z)$  is the Dirac delta function.

Following the Aris [2] method of moments, the  $p$ th integral moment of the concentration distribution can be defined as

$$C_p = \int_{-\infty}^{\infty} z^p C(t, r, z) dz, \quad (11)$$

and

$$M_p = \bar{C}_p = \frac{1}{2\pi} \int_0^{2\pi} d\theta \int_0^1 r C_p(t, r) dr. \quad (12)$$

So using equations (11) and (12), the diffusion equations (9) and (10) can be written in the form of  $C_p$  and  $M_p$  separately which are

$$\frac{\partial C_p}{\partial t} - \frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial C_p}{\partial r} \right) = p \text{Pe } u_z(t, r) C_{p-1} + p(p-1) C_{p-2}, \quad (13a)$$

with

$$C_p(0, r) = 1 \quad \text{and} \quad \frac{\partial C_p}{\partial r} = 0 \quad \text{at} \quad r = 1, \quad (13b)$$

and

$$\frac{dM_p}{dt} = p \text{Pe } \overline{u_z(t, r) C_{p-1}} + p(p-1) \overline{C_{p-2}}, \quad (14)$$

with

$$M_p(0) = 1, \quad \text{for } p = 0, \quad \text{and} \quad M_p = 0 \quad \text{for } p > 0,$$

overbar denotes the cross-sectional mean.

The  $p$ th central moment of concentration distribution about the mean can be written as,

$$v_p(t) = \frac{1}{\pi} \frac{\int_0^1 \int_0^{2\pi} \int_{-\infty}^{\infty} r(z - z_g)^p C(0, r, z) dr d\theta dz}{\int \int \int C dv}, \quad (15)$$

where

$$z_g = \frac{\int \int \int z C dv}{\int \int \int C dv} = \frac{M_1}{M_0}$$

is the centroid or first moment of the solute which measures the location of the centre of gravity of the slug with the mean velocity of the viscoelastic fluid initially located at the source, and  $M_0$  represents the total mass of inert solute in the whole volume of the tube.

Putting  $p = 2, 3, 4$  in (15) we get the central moments as,

$$\left. \begin{aligned} v_2(t) &= \frac{M_2}{M_0} - z_g^2 \\ v_3(t) &= \frac{M_3}{M_0} - 3z_g v_2 - z_g^3 \\ v_4(t) &= \frac{M_4}{M_0} - 4z_g v_3 - 6z_g v_2^2 - z_g^4 \end{aligned} \right\} \quad (16)$$

Here  $v_2$  represents the variance of the distribution of solute about the centre of the slug. Third ( $v_3$ ) and fourth ( $v_4$ ) central moments represent the symmetry and peakedness of the distribution of the slug about its mean ( $z_g$ ), respectively.  $M_i$ s are calculated from equations (13a) and (14).

### 3. NUMERICAL PROCEDURE

Because of the complexity of the analytical solution of the moment equation (13a) with the expression of  $u_z$  given by equation (8) subject to the initial and boundary conditions

specified in (13b), it is solved numerically. A finite-difference scheme based on the Crank-Nicholson implicit method has been adopted to study this problem. A marching technique for time has been used in this equation as the initial condition is specified and the derivatives along the marching direction have been replaced by backward differencing with second order accuracy, whereas the third-order accuracy in central differencing along the radial direction has been used for the diffusion term, and the resulting difference scheme becomes implicit. The scheme has been discussed in detail in the work of Mazumder and Das [6, 11]. The derivatives and all the other terms have been written at the mesh point  $(i + 1, j)$  where  $i = 0$  corresponds to the time  $t = 0$  and  $j = 0$  to the axis of the pipe  $r = 0$ . The mesh point  $(i, j)$  indicates a point where  $t = \Delta t \times i$  and  $r = \Delta r \times j$ ,  $\Delta t$  and  $\Delta r$  are the increments of  $t$  and  $r$ , respectively. The discretized equations are a system of linear algebraic equations with a tridiagonal coefficient matrix.

$$P_j C_p(i - 1, j + 1) - Q_j C_p(i + 1, j) + R_j C_p(i + 1, j - 1) = S_j \quad (17)$$

where

$$P_j = \frac{1}{2(\Delta r)^2} \left( \frac{1}{2j} + 1 \right),$$

$$Q_j = \frac{1}{\Delta t} + \frac{1}{(\Delta r)^2},$$

$$R_j = \frac{1}{2(\Delta r)^2} \left( \frac{1}{2j} - 1 \right),$$

and

$$\begin{aligned} S_j = & \frac{1}{\Delta t} C_p(i, j) + \frac{1}{4j(\Delta r)^2} \{C_p(i, j + 1) - C_p(i, j - 1)\} + \frac{1}{(\Delta r)^2} \{C_p(i, j + 1) \\ & - 2C_p(i, j) + C_p(i, j - 1)\} + \frac{\rho Pe}{4} \{u(i + 1, j) + u(i, j)\} \{C_{p-1}(i + 1, j) \\ & + C_{p-1}(i, j)\} + \frac{n(n-1)}{2} \{C_{p-2}(i + 1, j) + C_{p-2}(i, j)\} \end{aligned}$$

and the matrix elements.

The finite-difference form of the initial and boundary conditions are.

$$\begin{aligned} C_p(1, j) &= \begin{cases} 1 & \text{for } p = 0 \\ 0 & \text{for } p \geq 1 \end{cases} \\ C_p(i + 1, 1) &= C_p(i + 1, -1) \end{aligned} \quad (18)$$

at the axis, and

$$C_p(i + 1, N + 1) = C_p(i + 1, N - 1) \quad (19)$$

at the boundary for  $p \geq 0$ ,  $N$  is the value of  $j$  at the boundary.

This tridiagonal coefficient matrix has been solved by the method of the Thomas algorithm with the help of prescribed initial and boundary conditions. The integration for calculation of  $M_p$  from equation (14) has been performed employing Simpson's one-third rule. Initially, velocity is computed from equation (8). Using the values of velocity  $u_z$  at the grid  $(i + 1, j)$  in equation (13a), the concentration  $C_p$  is calculated at each grid point of computational domain. Finally, with the help of Simpson's one-third rule,  $M_p$  is evaluated using the values of  $u_z$  and  $C_p$  in the corresponding point. The values of the variables can be calculated for all time iteratively in the marching direction. The present scheme is linearly stable for a finite value of  $n$ , where  $n = \Delta t / (\Delta r)^2 = 0.01$  because of its implicitness. For

frequency parameter  $\alpha = 0.5, 4.0$  in oscillatory flow, mesh size has been taken as  $\Delta t = 0.00001$  and  $\Delta r = 0.01$ , and it gives a good accuracy and no considerable difference in results for smaller mesh size.

#### 4. DISCUSSION OF RESULTS

In order to validate the numerical scheme, the values of  $C_1$  and  $M_1$  have been calculated analytically using the moment method and numerically using the present scheme. The first moment of concentration distribution of solute  $C_1$  has been studied from equation (13a). So the diffusion equation for  $C_1$  is

$$\frac{\partial C_1}{\partial t} - \frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial C_1}{\partial r} \right) = \text{Pe } u_c C_0$$

This is a one-dimensional diffusion equation having a source term  $\text{Pe } u_c C_0$ . The first moment  $C_1$  can easily be solved using the method given by Mukherjee and Mazumder [5] and the expression for  $C_1$  is given by,

$$C_1(r, t) = \text{Real} \left[ \frac{2i\text{Pe } k_1 J_1(k_1)}{\alpha} \sum_j \frac{J_0(\alpha_j r)}{(k_1^2 - \alpha_j^2)(\alpha_j^2 + i\alpha S) J_0(\alpha_j)} \{e^{i\alpha_j S t} - e^{-\alpha_j^2 r}\} - \frac{\text{Pe}}{\alpha^2 S} \left\{ 1 - \frac{2J_1(k_1)}{k_1 J_0(k_1)} \right\} \{e^{i\alpha S t} - 1\} \right], \quad (20)$$

where  $J_0, J_1$  are the 1st kind Bessel functions of zeroth and first order respectively,  $\alpha_j$ s are the zeros of  $J_1$ .

The values of  $C_1$  calculated from the numerical scheme as well as from the analytical method given by Mukherjee and Mazumder [5] have been compared and it is found that the analytical and numerical values are in good agreement. The variation of distribution of  $C_1$  against  $r$  for different phase values has been plotted in Fig. 2 for  $T_0 = 0.1, 0.3$  and

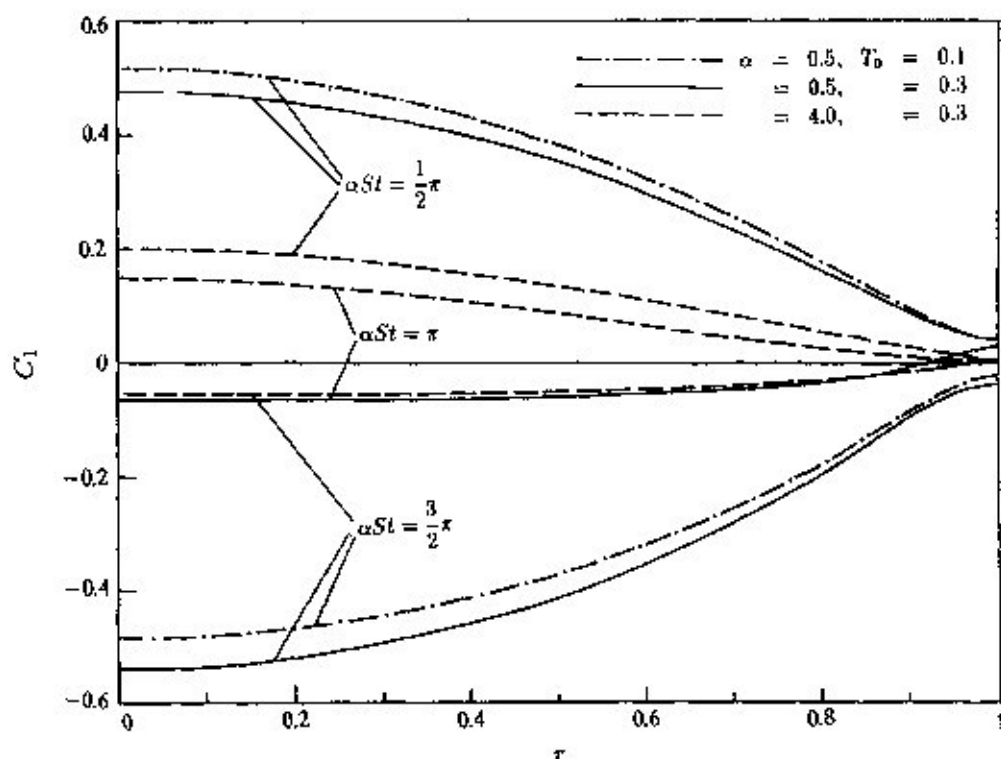


Fig. 2. First order moment  $C_1$  against radius  $r$  for different phases, when  $\text{Pe} = S = 1000$ .

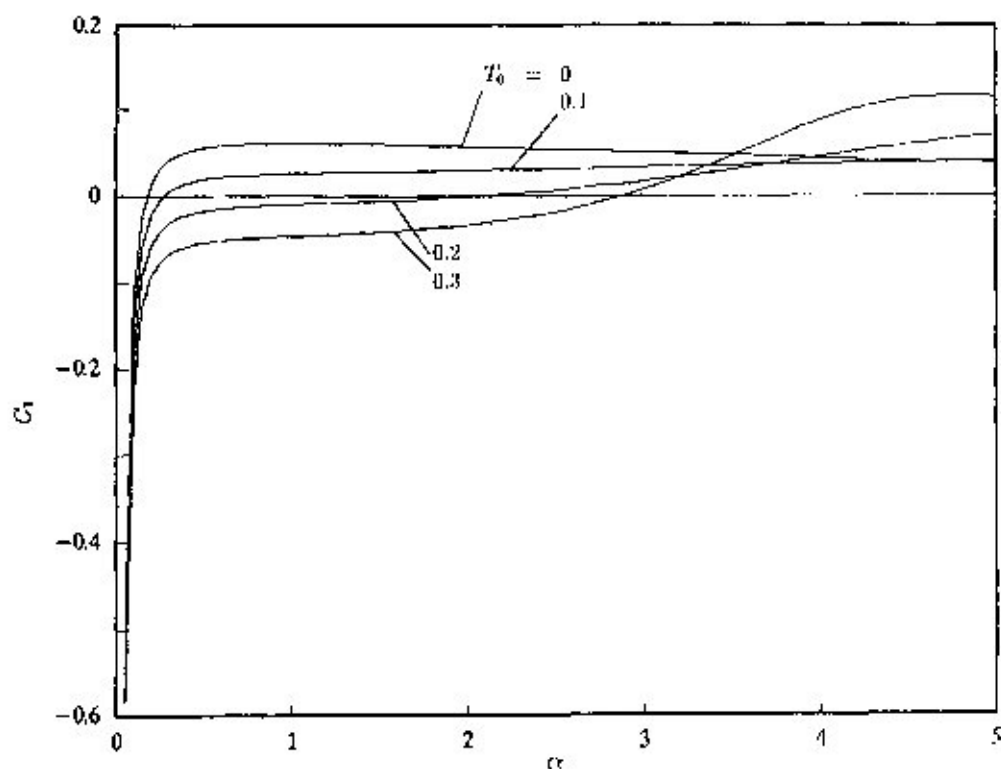


Fig. 3. First order moment  $C_1$  against frequency parameter  $\alpha$ , when  $Pe \rightarrow S = 1000$  and  $r = 0.5$  for  $\alpha St = \pi$ .

$\alpha = 0.5, 4.0$ .  $C_1$  reveals its periodicity with flow phases because of the oscillatory nature in velocity  $u_z$ . Figure 3 shows the variation of  $C_1$  against  $\alpha$  for different values  $T_0$ ,  $r = 0.5$ , and  $\alpha St = \pi$ . It is interesting to note that in the case of a Newtonian fluid ( $T_0 = 0$ ) the effect of the frequency parameter  $\alpha$  on  $C_1$  is not significant, whereas with the increase of the visco-elastic parameter  $T_0 > 0$ ,  $C_1$  decreases for a certain range of  $\alpha$ , then it increases.

The first moment  $M_1$  indicates the mean concentration distribution over the cross-section of the tube. Putting  $p = 1$ , the equation of  $M_n$  gives,

$$\frac{dM_1}{dr} = Pe \overline{u_z C_0}$$

with  $M_1(0) = 0$

Similar to  $C_1$ ,  $M_1$  can be obtained as

$$M_1(z) = -\frac{Pe}{\alpha^2 S} \text{Real} \left[ \left\{ 1 - \frac{2J_1(k_1)}{k_1 J_0(k_1)} \right\} \{ e^{i\alpha St} - 1 \} \right] \quad (21)$$

The mean longitudinal displacement  $Z_g (= M_1/M_0)$  of the solute moving with the mean periodic velocity of the solvent mainly depends on  $\alpha$ ,  $T_0$  and  $z$ . Figure 4(a, b) shows the displacement of centroid ( $z_g$ ) for different values of  $T_0$  and  $\alpha = 0.5, 4.0$ . It is seen that the centroid of the slug moves cyclically with the oscillatory nature of the flow, and it changes asymptotically over a period. For low frequency of oscillation ( $\alpha = 0.5$ ), the amplitude of oscillation decreases in the first part of the period and increases in the second part with the increase in elastic parameter  $T_0$ , whereas for a large frequency ( $\alpha = 4.0$ ), the amplitude of oscillation increases in the both parts [Fig. 4(b)]. For both low and high values of frequency, it is seen that there is a phase shift which decreases with an increase in  $T_0$ . It is also observed that the amplitude of positive pulsation is more prominent than that of negative pulsation for high frequency.

Owing to the complexity of analytical solution of equations (13a)–(14) for  $p = 2$ , subject to the initial and boundary conditions, we have solved numerically, and hence the variance



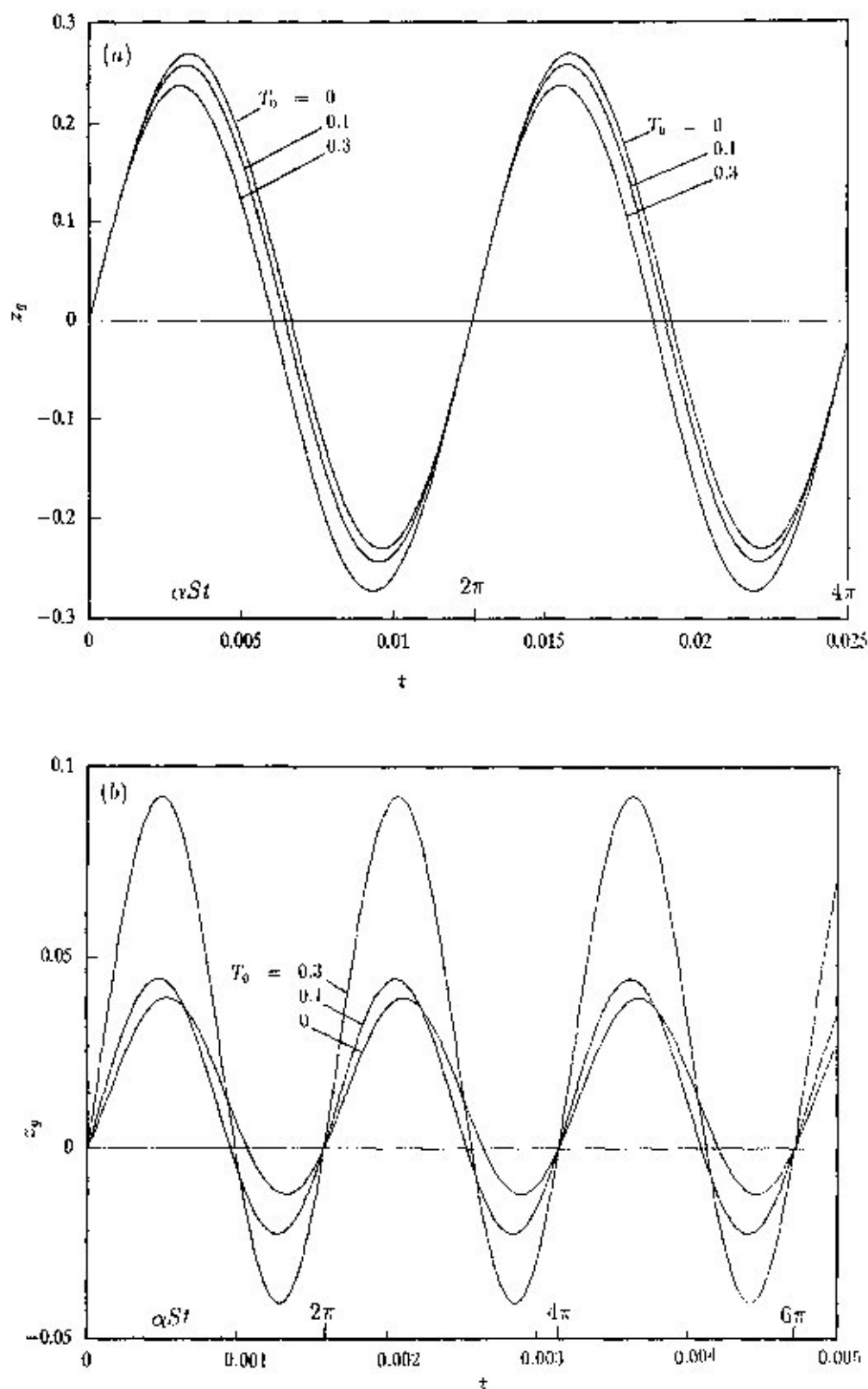


Fig. 4. The centroid displacement  $z_y$  due to oscillatory flow, when  $Pe = S = 1000$  for (a)  $\alpha = 0.5$  and (b)  $\alpha = 4.0$ .

$v_2$  and dispersion coefficient due to shear effect have been evaluated. The plots of second central moment  $v_2$  (variance) of the longitudinal concentration distribution against dispersion time  $t$  for  $T_0 = 0, 0.1, 0.3$ ,  $Pe = S = 10^3$  have been presented in Fig. 5 when  $\alpha = 0.5$  and in Fig. 6 when  $\alpha = 4.0$ . It is essentially the dispersion due to both the longitudinal diffusion

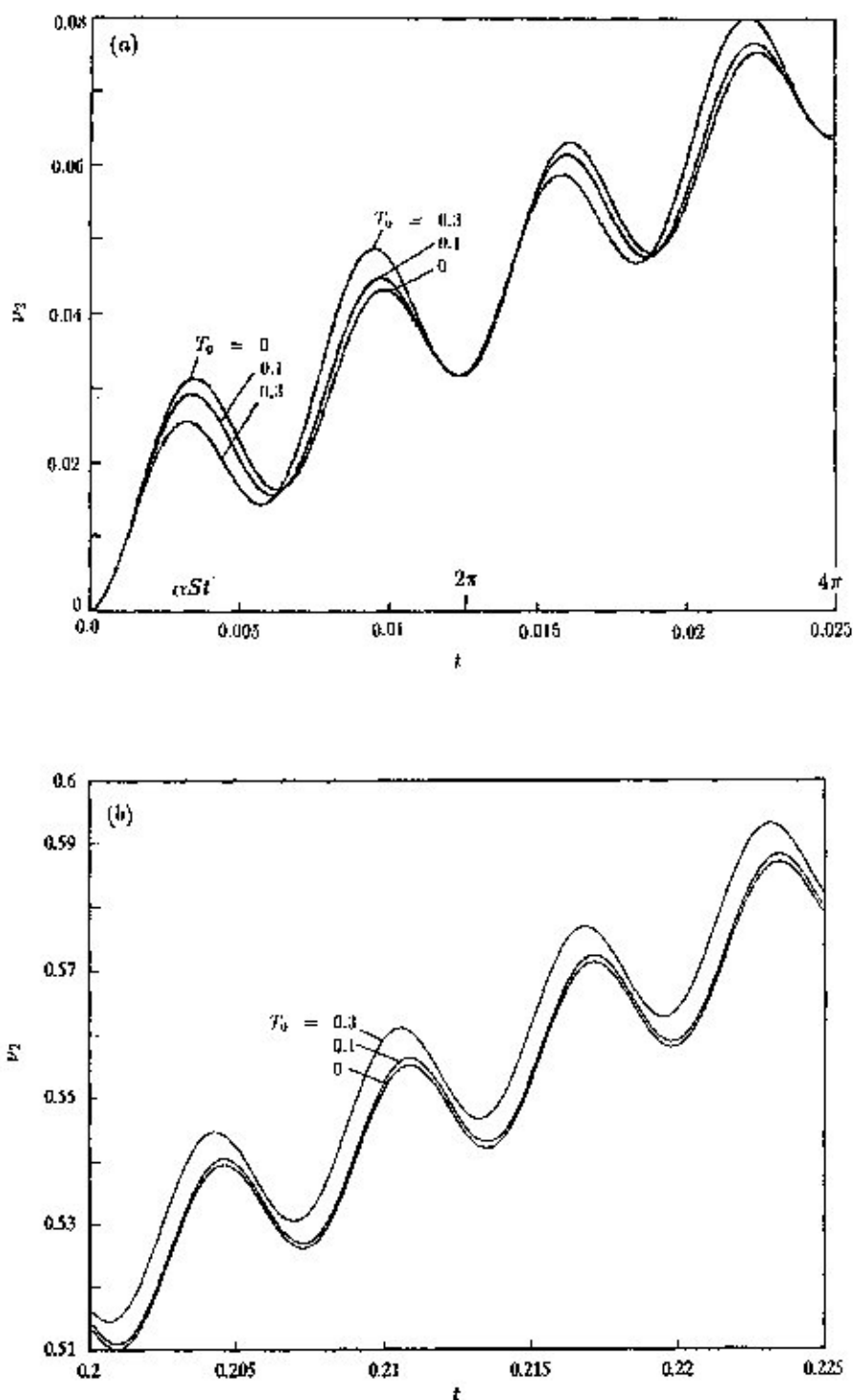
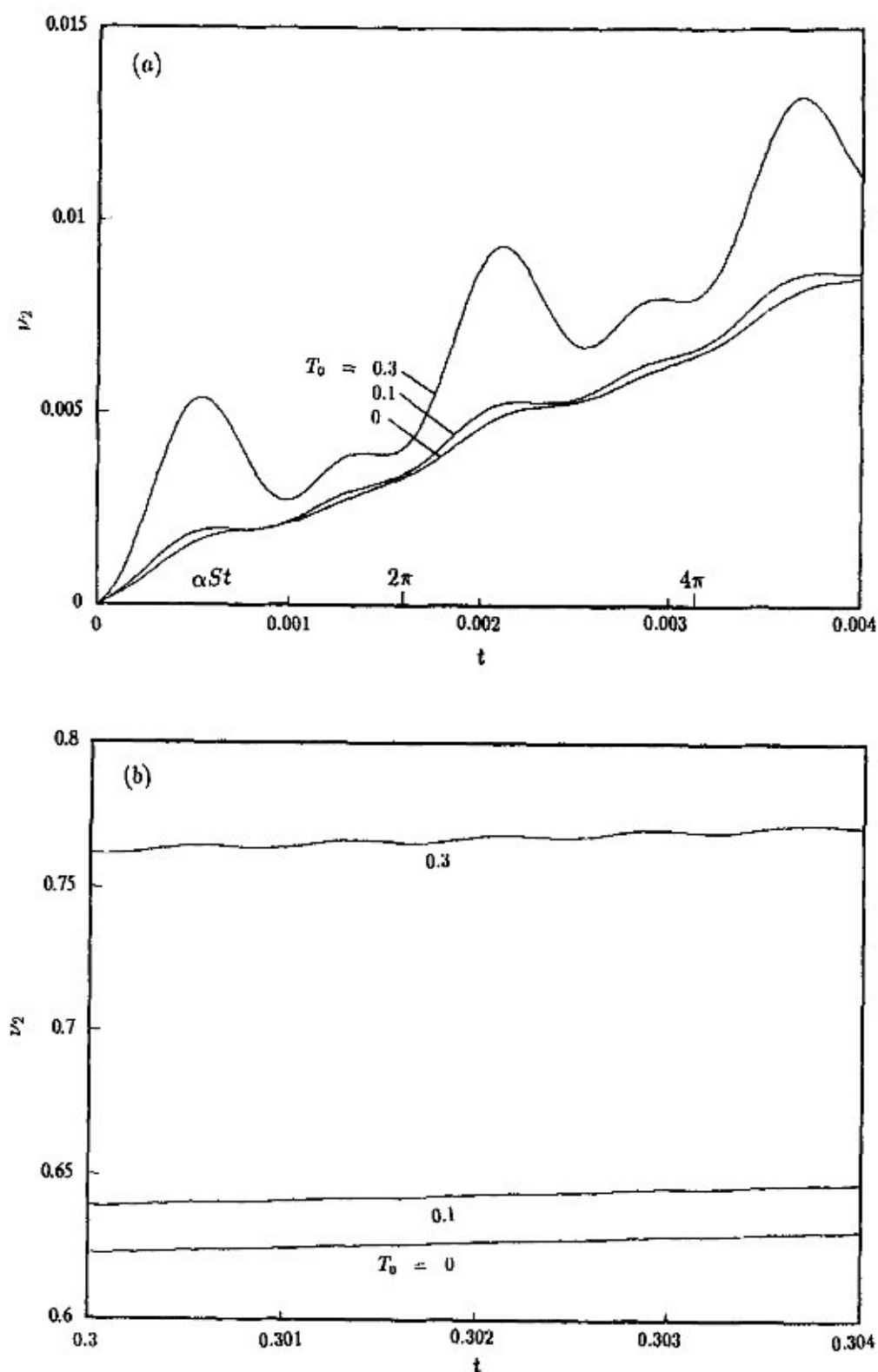


Fig. 5. The temporal variation of variance  $v_2$  for (a) small time, (b) large time, when  $Pe = S = 1000$ ,  $\alpha = 0.5$ .

and interaction of the periodic current and lateral diffusion. At low frequency of oscillation, Fig. 5 shows that for a given value of  $T_0$ , the variance increases with time in a wavy pattern. In a complete period, variance decreases in the first part of oscillation, and then increases in the second part with increase in relaxation time  $T_0$  [Fig. 5(a)], but this behaviour completely diminishes for large time where variance increases at a fairly uniform rate with  $T_0$  [Fig. 5(b)]. The increase of variance ( $v_2$ ) about the mean ( $z_0$ ) with  $T_0$  can directly be related to the increase of fluid velocity with elasticity in fluid. It also reveals that with an increase of  $\alpha$  (i.e. increase of frequency of oscillation) the variance of distribution decreases. For  $\alpha = 0.5$

Fig. 6. As Fig. 5 but  $\alpha = 4.0$ .

the wavy nature of the variance ( $v_2$ ) of the solute distribution is almost the same for all time period, whereas for  $\alpha = 4.0$  (Fig. 6) it shows double frequency oscillation in a complete period upto a certain time, and it reaches asymptotically a steady state for large time. Further, it may be mentioned here that the amplitude of variance increases for all time period with the relaxation time  $T_0$ .

Aris, in his method of moment, found out that the rate of change of variance is proportional to the sum of the molecular diffusion coefficient and the Taylor diffusion coefficient. Therefore, according to Aris [2] the rate of growth of variance is defined as:

$$\frac{dV_2}{dt} = 2 + 2Pe^3 D_*(S, \alpha, T_0, t) \quad (22)$$

where  $D_*$  is the apparent dispersion coefficient depending on parameters  $S$ ,  $\alpha$ ,  $T_0$  and  $t$ . The first term on the right hand side of the above expression represents the longitudinal diffusion, whereas the second term represents the interaction between the convection and lateral diffusion. Therefore, the apparent dispersion coefficient  $D_*$  is discussed. Taking  $Pe = S = 10^3$ , the variation of  $D_*$  against dispersion time  $t$  has been presented for various values of  $T_0$  in Fig. 7 for  $\alpha = 0.5$  and Fig. 8 for  $\alpha = 4.0$ . The variation of  $D_*$  in oscillatory flow changes cyclically with a double frequency and reaches a stationary state after a certain time, which is related to the cross-sectional mixing time. The amplitudes of oscillation of  $D_*$  during the first and second half of a complete period of oscillatory flow are almost symmetrical for low frequency (Fig. 7), whereas for the high frequency parameter,  $D_*$  is more significant during the first half of the period than the second one (Fig. 8). However, this situation completely stabilizes after a certain time and then the solute disperses at a fairly uniform rate (Mazumder and Das [6], Yasuda [10]). The apparent dispersion coefficient  $D_*$  in the low frequency of oscillation reaches a stationary state earlier than for the high frequency. The dispersion coefficient  $D_*$  changes cyclically with time even in the stationary state.

The longitudinal dispersion of solute strongly depends on the visco-elastic parameter  $T_0$ . The effect of  $T_0$  for a small value of  $\alpha$  is small compared to that for a large value of  $\alpha$ . Figure 7(a, b) shows the increment in phase lag of  $D_*$  with increase of  $T_0$  due to a decrease of the modulus of elasticity. Initially, at the low frequency of oscillation the amplitude of  $D_*$  decreases during the first half of the period of oscillation and then increases in the second half with increase in  $T_0$ ; at large time [Fig. 7(b)] it becomes stable with the flow. Figure 8(a, b) shows the variation of  $D_*$  with time when  $\alpha = 4.0$  for different  $T_0$ . Initially, a double-frequency oscillation is observed and the difference between two consecutive oscillations increases with increase in  $\alpha$ , i.e. increase in frequency of oscillation, and for this it takes a larger time to reach the steady state. The amplitude of the oscillation increases with increase in  $T_0$  which is comparable with the study made by Erdoğan [14] for small values of a non-Newtonian parameter in asymptotically large time.

Once the central moments  $v_2, v_3, v_4, \dots$ , are known, it is possible to compute the mean concentration distribution  $C_m(t, z)$  deviated from Gaussianity in terms of Hermite polynomial representation (Chatwin [26], Andersson and Berglin [27]) and is given by:

$$C_m(t, z) = M_0(t) e^{-x^2} \sum_{n=0}^{\infty} a_n(t) H_n(x) \quad (23)$$

where

$$M_0(t) = \iiint C \, dv, \quad x = \frac{z - z_k}{(2v_2)^{1/2}}, \quad z_k = \frac{M_1}{M_0}$$

and  $H_n$  the Hermite polynomials, satisfy the recurrence relation with  $H_0(x) = 1.0$  as

$$H_{i+1}(x) = 2xH_i(x) - 2iH_{i-1}(x), \quad i = 0, 1, 2, \dots \quad (24)$$

The coefficients  $a_i$  are

$$a_0 = 1/(2\pi v_2)^{1/2}, \quad a_1 = a_2 = 0, \quad a_3 = 2^{1/2} a_0 \beta_2 / 24, \quad a_4 = a_0 \beta_3 / 96.$$

Coefficient of skewness  $\beta_2 = v_3/v_2^{3/2}$ , and that of kurtosis  $\beta_3 = v_4/v_2^2 - 3$  represent the degree of symmetry and peakedness of the distribution of solute respectively. These indicate basically the nature of the distribution and the deviations from Gaussianity. If the distribution is exactly Gaussian, both coefficients will be zero.

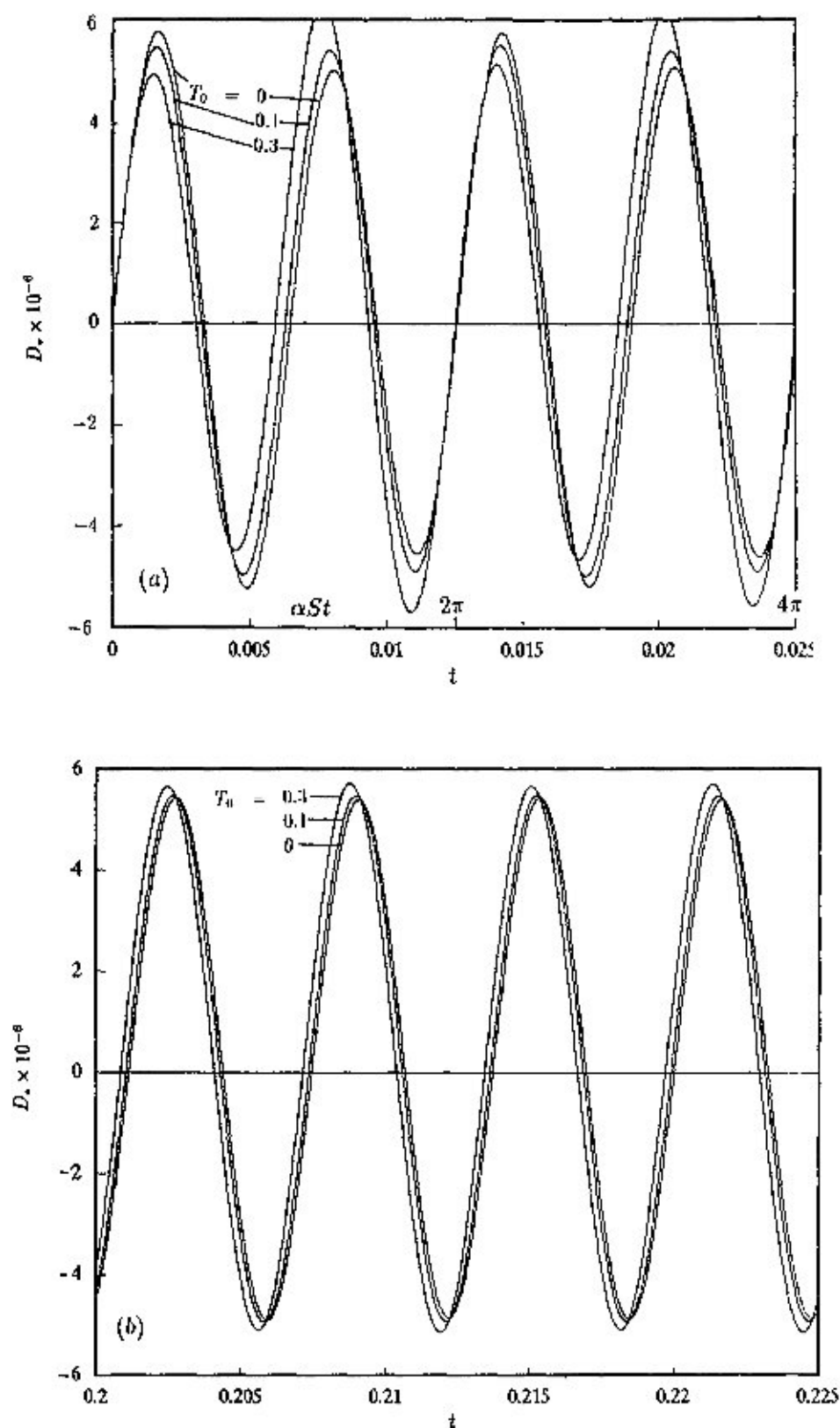
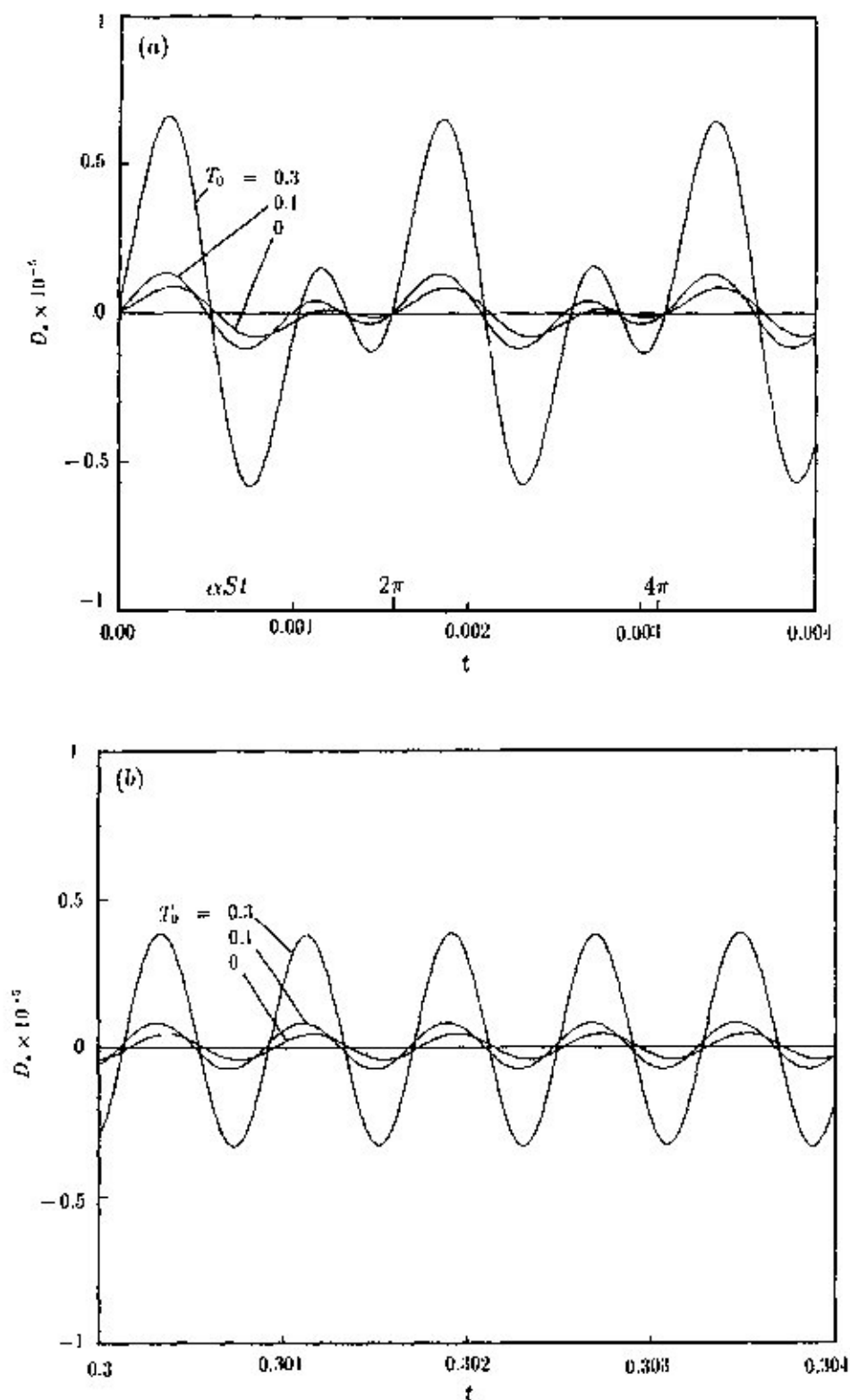


Fig. 7. The dispersion coefficient  $D_s$  for (a) small time, (b) large time, when  $Pc = S = 1000$ ,  $\alpha = 0.5$ .

Table 1 shows the variation of  $\beta_2$  and  $\beta_3$  with the frequency of oscillation  $\alpha$ , viscoelastic parameter  $T_0$  and the dispersion time  $t$ . It is seen from the table that there is a small deviation in  $\beta_2$  from zero, which increases with the increase of  $T_0$ . The variation for  $T_0$  becomes oscillatory for large values of  $\alpha$  which is clear from the table when  $\alpha = 4.0$ .  $\beta_3$

Fig. 8. As Fig. 7 but  $\alpha = 4.0$ .

increases with increase of  $T_0$  as well as  $\alpha$ . With increase of time,  $\beta_2$  decreases steadily for small  $\alpha$  and for large values of  $\alpha$  it decreases in oscillatory nature. But  $\beta_3$  decreases steadily for all  $\alpha$  with increase of time  $t$ . So it is revealed that if we increase the non-Newtonian parameter, i.e. increase the elastic property in fluid, it takes much time to reach the Gaussianity.

Table 1. Variation of coefficients of skewness ( $\beta_2$ ) and kurtosis ( $\beta_3$ )

$T_0$	$t$	$\alpha = 0.5$		$\alpha = 4.0$	
		$\beta_2$	$\beta_3$	$\beta_2$	$\beta_3$
0.0	0.25	0.00343	-1.60	0.00041	-0.011
0.1	0.25	0.00356	0.0025	0.00096	0.0031
0.3	0.25	0.00388	0.0028	0.00734	0.0443
0.3	0.20	0.00444	0.0033	0.01610	0.0514
0.3	0.15	0.00508	0.0040	0.01630	0.0604

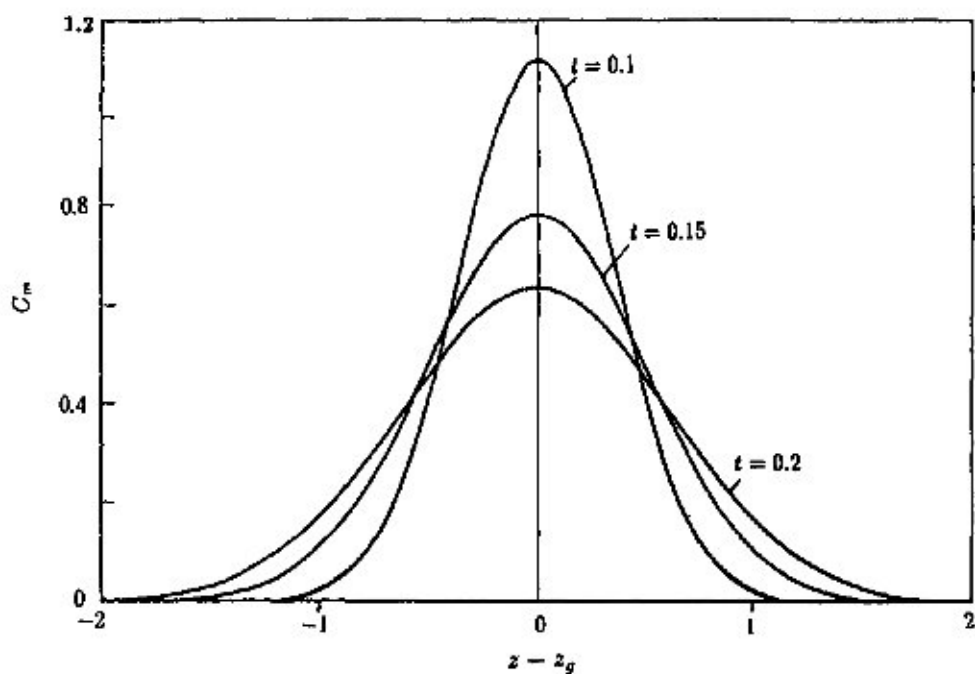


Fig. 9. The mean concentration distribution  $C_m$  along the pipe for  $Pe = S = 1000$ ,  $T_0 = 0.3$  and  $\alpha = 0.5$ .

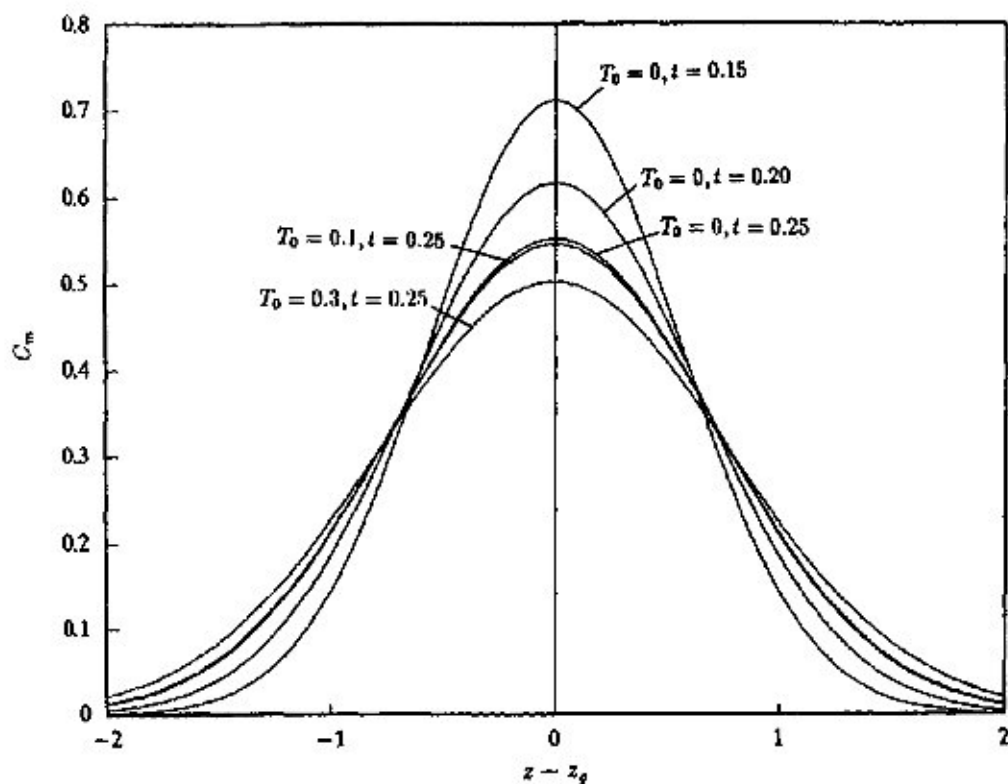


Fig. 10. The mean concentration distribution  $C_m$  along the pipe for  $Pe = S = 1000$ , and  $\alpha = 4.0$ .

In Figs 9 and 10,  $C_m(t, z)$  has been plotted against axial distance  $(z - z_g)$  for different values of time  $t$  and  $T_0$  when  $\alpha = 0.5$  and 4.0, respectively. From the figures it is observed that the peak of the concentration distribution gradually decreases with increase of dispersion time  $t$  as well as increase of relaxation time  $T_0$ , which implies the distribution gradually tends to become flat. It is also seen that the mean concentration distribution due to unsteady flow of viscoelastic fluid is essentially symmetrical. From the above observation it is noted that there is a remarkable similarity between the mean concentration distribution of solute in an unsteady non-Newtonian (viscoelastic) fluid and in periodic flow of a Newtonian fluid discussed by Mazumder and Das [6]. Mean concentration distribution profiles show how fast the slug's centre of gravity moves, how it disperses due to shear effects and how the distribution deviates from the Gaussianity due to the viscoelastic parameter.

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#### REFERENCES

1. G. I. Taylor, *Proc. R. Soc. Lond.* **A219**, 186 (1953).
2. R. Aris, *Proc. R. Soc. Lond.* **A235**, 67 (1956).
3. N. G. Barton, *J. Fluid Mech.* **126**, 205 (1983).
4. R. Aris, *Proc. R. Soc. Lond.* **A259**, 370 (1960).
5. A. Mukherjee and B. S. Mazumder, *Acta Mechanica* **74**, 107 (1988).
6. B. S. Mazumder and S. K. Das, *J. Fluid Mech.* **239**, 523 (1992).
7. P. C. Chatwin, *J. Fluid Mech.* **71**, 513 (1975).
8. R. Smith, *J. Fluid Mech.* **114**, 379 (1982).
9. C. F. Jimenez and P. J. Sullivan, *J. Fluid Mech.* **142**, 57 (1984).
10. H. Yasuda, *J. Fluid Mech.* **148**, 383 (1984).
11. B. S. Mazumder and S. K. Das, *Acta Mechanica* **80**, 151 (1989).
12. L. T. Fan and W. S. Hwang, *Proc. R. Soc. Lond.* **A283**, 576 (1965).
13. L. T. Fan and C. B. Wang, *Proc. R. Soc. Lond.* **A292**, 203 (1966).
14. M. J. Frdojgan, *I & EC Fundamentals* **6**, 463 (1967).
15. S. Ghoshal, *Chem. Engng Sci.* **26**, 185 (1971).
16. S. N. Shah and K. J. Cox, *Chem. Engng Sci.* **29**, 1282 (1974).
17. A. S. Gupta and B. S. Mazumder, *Int. J. Heat Mass Transfer* **20**, 341 (1977).
18. R. S. Subramanian and W. N. Gill, *Can. J. Chem. Engng* **54**, 121 (1976).
19. R. B. Bird, W. E. Stewart and E. N. Lightfoot, *Transport Phenomena*. Wiley, New York (1960).
20. J. G. Oldroyd, *Proc. R. Soc. Lond.* **A200**, 523 (1950).
21. S. Uchida, *Z. Angew. Math. Phys.* **7**, 403 (1956).
22. H. Schlichting, *Boundary-Layer Theory*. McGraw-Hill (1979).
23. E. M. Khabakhpasheva, V. I. Popov, A. N. Kekalov and E. S. Mikhailova, *J. Non-Newtonian Fluid Mech.* **289** (1989).
24. D. W. Beard and K. Walters, *Proc. Camb. Phil. Soc.* **60**, 667 (1964).
25. K. R. Rajagopal and E. Scibba, *Maths Computers Simulation* **26**, 276 (1984).
26. P. C. Chatwin, *J. Fluid Mech.* **43**, 321 (1970).
27. B. Andersson and T. Berglin, *Proc. R. Soc. Lond.* **A377**, 251 (1981).