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# Exact Analysis of Unsteady Convective Diffusion in Herschel-Bulkley Fluid Flow- Application to Catheterised Stenosed Artery

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#### **ARTICLE INFO**

#### **ABSTRACT**

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#### Keywords:

Blood flow; catheter; solute dispersion; Stenosis; Herschel-Bulkley fluid One of the major causes of cardiovascular disease is atherosclerosis or stenosis. This study is designed to improve the current body of knowledge regarding the condition by inserting a long thin tube called a catheter to widen the narrow part in the artery. The study reviewed the effects of catheter radius, yield stress, and power law index on the velocity distribution, and transport coefficients of solute. A mathematical model is deployed to investigate the dispersion of solute in the flow of a Herschel-Bulkley (H-B) fluid in an annulus, whereas the dispersion process is studied using the generalised dispersion model (GDM) by solving the convective diffusion equation. Resultantly, the velocity reduces following an increase in the yield stress, catheter size, and power law index. Meanwhile, the dispersion coefficient exhibits a same behaviour as the aforementioned parameters ascend considerably. The dispersion coefficient alterations occurred rapidly for small values of time and became significantly constant following an increase in the time values. Conclusively, this study can be useful in dispersion of a drug to the affected artery where an abnormal plaque was formed.

## 1. Introduction

The transportation of solute dispersion process has been widely studied due to its extensive application in the field of chemical engineering, physiological fluid dynamics, biomedical engineering, environmental sciences, and pharmacology [1,2]. Studies on the dispersion of solute, such as drug, toxin, or nutrient, in blood flow through a narrow artery having a linear or constricted wall received much attention owing to its significant contribution in understanding the issues in biomedical engineering and cardiovascular mechanics [3-5]. Constriction in an artery occurs because of the accumulation of low-density lipoprotein (LDL) and other macromolecules along the inner lining of the arterial wall. The formation of such lesion or plaque started blocking the artery and reducing the normal blood flow, medically termed as atherosclerosis or stenosis. It is important for clinicians to analyse the rate of dispersion of an injected drug associated with intravenous drug delivery into an affected artery because of its therapeutic nature and also to measure the amount of drug in the system for better efficacy as well as the effectiveness of the delivery [6]. In this situation, a long thin

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tube called a catheter is inserted into the veins to refine the flow. The use of catheters is of immense importance and has become a standard tool for diagnosis and treatment in modern medicine [7]. The catheter is carefully guided to the location at which stenosis occurs and the balloon is then inflated to fracture the fatty deposits and widen the narrowed part of the artery.

Catheterized arteries are now extensively used in medical science to measure various physiological flow characteristics as well as to diagnose and treat various arterial diseases. Numerous researchers have recently evaluated the blood flow in an annulus employed in a catheterised stenosed artery [8-10]. However, these studies did not consider the mass transfer that governs the flow of mass transport in the systemic circulation. The convective diffusion equation controls the mass transfer to the bloodstream [11]. A method known as the generalised dispersion model (GDM) was proposed by Gill and Sankarasubramanian [12] to generate an exact solution of the convective diffusion equation, which is applicable at all times.

Hence, based on the existing literature, there is a lack of studies on the problem of unsteady solute dispersion in blood flow by considering the H-B fluid model in an overlapping catheterised stenosed artery using the GDM. This study extends the works by Sankar and Hemalatha [13] and Abbas *et al.* [14] by investigating the rheological behaviour of blood flow in an overlapping stenosed artery. The investigation of solute dispersion in a non-Newtonian fluid is crucial to yield realistic results that better represent physical problems. This study aims to investigate how reactive solute disperse in the solvent is influenced by physical parameters such as catheter radius, yield stress, and power law index. Specifically, the contributions of the study are twofold, firstly to evaluate the effects of reactive species in an overlapping catheterised stenosed artery using GDM that was only addressed individually in previous studies, and secondly to analyse the rheological behaviour of non-Newtonian fluid in a catheterised stenosed artery.

#### 2. Mathematical Formulation

A catheter is inserted into an artery coaxially, and the artery takes the form of a rigid circular tube. Radius of the circular tube is  $R_0$  and radius of the catheter is given as  $kR_0$ , k < 1. Consider a steady, axially symmetrical, laminar, and fully developed uni-directional flow of blood in the axial direction portrayed as a viscous incompressible non-Newtonian fluid. Herschel-Bulkley (H-B) fluid model is employed to illustrate the blood flow. The entrance, endpoint and distinctive wall effects of the artery can be disregarded given the sufficient length of the arterial segment.

## 2.1 Governing Equations

A cylindrical polar coordinate  $(\bar{r}, \bar{\theta}, \bar{z})$  is used in the study where  $\bar{r}$  and  $\bar{z}$  denote the radial and axial coordinates and  $\bar{\theta}$  is the azimuthal angle. The fluid velocity in  $\bar{r}$  direction is ignored as its magnitude is negligibly small and only accounts in a  $\bar{z}$  direction. Thus,  $\bar{u}_{\bar{r}} = \bar{u}_{\bar{\theta}} = 0$  [15]. Since the pressure gradient is a function of  $\bar{z}$  only and independent of  $\bar{r}$  and  $\bar{\theta}$ , thus  $d\bar{p}/d\bar{z}$  is constant, hence, the momentum equation can be simplified as

$$\frac{d\overline{p}}{d\overline{z}} = -\frac{1}{\overline{r}} \frac{d}{d\overline{r}} (\overline{r} \, \overline{\tau}), \quad \overline{k} \le \overline{r} \le \overline{R} (\overline{z}), \tag{1}$$

where the constant pressure, shear stress, catheter radius and radius of stenosed artery are represented as  $\bar{p}, \bar{\tau}, \bar{k}$  and  $\bar{R}(\bar{z})$ , respectively. The following equation presents the constitutive equation of H-B fluid

$$\eta_{HB} \frac{\partial \overline{u}}{\partial \overline{r}} = \begin{cases}
 |\overline{\tau}|^n \left( 1 - \frac{n\overline{\tau}_y}{|\overline{\tau}|} \right)^n, & \text{if } |\overline{\tau}| \ge \overline{\tau}_y \text{ and } \frac{\partial \overline{u}}{\partial \overline{r}} > 0, \\
 -|\overline{\tau}|^n \left( 1 - \frac{n\overline{\tau}_y}{|\overline{\tau}|} \right)^n, & \text{if } |\overline{\tau}| \ge \overline{\tau}_y \text{ and } \frac{\partial \overline{u}}{\partial \overline{r}} < 0, \\
 0, & \text{if } |\overline{\tau}| < \overline{\tau}_y.
\end{cases} \tag{2}$$

where  $\bar{u}$ ,  $\bar{\tau}_{\bar{y}}$  and n is the axial velocity, yield stress, and power law index, respectively. Meanwhile, the H-B fluid viscosity coefficient is represented by  $\eta_{HB}$  with dimension  $(ML^{-1}T^{-2})^nT$ . The unknown parameters, velocity  $\bar{u}$  and shear stress  $\bar{\tau}$ , can be solved using Eqs. (1) and (2) based on the no-slip boundary conditions for the catheter wall and artery presented below

$$\overline{u} = 0$$
 when  $\overline{r} = \overline{k}$ ,  
 $\overline{u} = 0$  when  $\overline{r} = \overline{R}(\overline{z})$ .

#### 2.2 Non-dimensionalisation

Introducing the following dimensionless variables

$$C = \frac{\overline{C}}{\overline{C}_{0}}, u = \frac{\overline{u}}{u_{0}}, u_{m} = \frac{\overline{u}_{m}}{u_{0}}, r = \frac{\overline{r}}{R_{0}}, R(z) = \frac{\overline{R}(\overline{z})}{R_{0}}, z = \frac{\overline{z}}{R_{0}},$$

$$\tau = \frac{\overline{\tau}}{(p_{0}R_{0}/2)}, \theta = \frac{\overline{\tau}_{y}}{(p_{0}R_{0}/2)}, I_{0} = \frac{\overline{I}_{0}}{R_{0}}, \delta = \frac{\overline{\delta}}{R_{0}}, k = \frac{\overline{k}}{R_{0}},$$

$$(4)$$

where C is the solute concentration, u is the velocity,  $u_m$  is the average velocity,  $u_0 = \frac{R_0^{n+1}p_0^n}{2^n\eta_{HB}}$  is the centreline velocity, r is the radial distance, R(z) is the radius of the stenotic artery, z is the axial distance,  $\tau$  is the shear stress,  $\theta$  is the yield stress,  $l_0$  is the stenosis length,  $\delta$  is the stenosis height, k is the catheter radius and  $\eta_{HB} = \mu \left(\frac{p_0 R_0}{2}\right)^{n-1}$ , where  $\mu$  is the viscosity coefficient for a Newtonian fluid.

## 2.3 Geometry of Stenosis

The geometry of the artery in the dimensionless form can be written as Layek *et al.*, [16] and ZainulAbidin *et al.*, [17]

$$R(z) = \begin{cases} 1 - \frac{3\delta}{2I_0^4} \Big[ 11(z-d)I_0^3 - 47(z-d)^2I_0^2 + 72(z-d)^3I_0 - 36(z-d)^4 \Big], & d \le z \le d+I_0, \\ 1, & \text{Otherwise,} \end{cases}$$
 (5)

where  $\delta$  is the maximum stenosis height occurs at  $d+l_0/3$  and  $d+2l_0/3$ , d is the distance of the stenosis from the inlet, and L is the length of the artery. The geometry of a catheterised artery with an overlapping stenosis under consideration is shown in Figure 1.

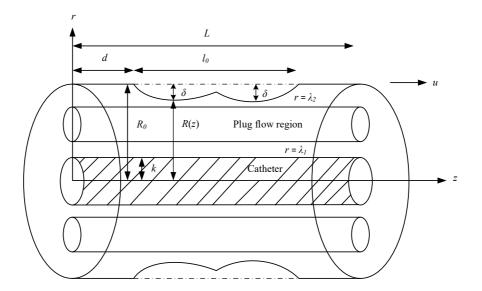


Fig. 1. The geometry of catheterised artery with an overlapping stenosis

# 2.4 Mass Transport

The transport of a reactive solute in the bloodstream in dimensionless form is governed by the convection-diffusion equation [18], which is expressed as

$$\frac{\partial C}{\partial t} + u \frac{\partial C}{\partial z} = \left(\frac{1}{r} \frac{\partial}{\partial r} \left(r \frac{\partial}{\partial r}\right) + \frac{1}{Pe^2} \frac{\partial^2}{\partial z^2}\right) C,$$
(6)

where Pe is the Peclet number, t is the dispersion time,  $D_m$  is molecular diffusivity of the solute which is assumed as constant.

# 2.5 Method of Solution

The momentum Eq. (1) in dimensionless form can be stated as

$$p_{s} = \frac{1}{2r} \frac{d}{dr} (r\tau), \quad k \le r \le R(z), \tag{7}$$

where  $p_{\rm S}$  is the dimensionless pressure gradient in steady state. The constitutive Eq. (2) in dimensionless form is

$$\frac{\partial u}{\partial r} = \begin{cases}
|\tau|^n \left( 1 - \frac{n\theta}{|\tau|} \right), & \text{for } |\tau| \ge \theta \text{ and } \frac{\partial u}{\partial r} > 0, \\
-|\tau|^n \left( 1 - \frac{n\theta}{|\tau|} \right), & \text{for } |\tau| \ge \theta \text{ and } \frac{\partial u}{\partial r} < 0, \\
0, & \text{for } |\tau| < \theta.
\end{cases} \tag{8}$$

Integrating Eq. (7) with respect to r yields

$$\tau = p_s r + \frac{C}{r},\tag{9}$$

where C is the constant of integration. Referring to Eq. (8), there are three different flow regions for  $k \leq r \leq R(z)$ , in which the central core region has a constant velocity and forms the plug flow region. Flow in this plug flow region is not sheared which means blood streamlines are moving at constant velocity because the shear stress is lesser than yield stress. The fluid (blood) will not flow at this region rather it is transported by the fluid particles present in the nearby shear flow region as a solid mass with a constant velocity. For mathematical representation, the plug flow region can be described by  $\lambda_1 \leq r \leq \lambda_2$ , where  $k \leq \lambda_1$  and  $\lambda_2 \leq R(z)$ .  $\lambda_1$  and  $\lambda_2$  are unknown constants to be identified. The three regions are depicted as in Figure 1. From the continuity of the shear stress along the plug flow region boundary, we have

$$-\tau = \tau_y$$
 when  $r = \lambda_1$ , (10)  $\tau = \tau_y$  when  $r = \lambda_2$ .

Using condition (10) in Eq. (9), the unknown constant C is evaluated in terms of  $\lambda_1$  and  $\lambda_2$  as

$$C = -p_s \lambda^2, \tag{11}$$

where

$$\lambda^2 = \lambda_1 \lambda_2. \tag{12}$$

Substitution of Eq. (11) in Eq. (9) delineates the shear stress as

$$\tau = \frac{p_s}{r} \left( r^2 - \lambda^2 \right). \tag{13}$$

Using Eq. (13) and condition (10) yields

$$\lambda_2 - \lambda_1 = \frac{\theta}{\rho} = \varsigma_A, \tag{14}$$

where  $\varsigma_A$  is the width of the plug core region. Velocity expressions for the three regions can be found using Eqs. (13) and (8) as below

$$u^{+}(r) = p_{s}^{n} \left[ \int_{k}^{r} \left( \frac{\lambda^{2} - r^{2}}{r} \right)^{n} dr - n \varsigma_{A} \int_{k}^{r} \left( \frac{\lambda^{2} - r^{2}}{r} \right)^{n-1} dr \right], \text{ when } k \le r \le \lambda_{1},$$

$$(15)$$

$$u_p = u^+(\lambda_1) = u^{++}(\lambda_2) = \text{constant}, \text{ when } \lambda_1 \le r \le \lambda_2,$$
 (16)

$$u^{++}\left(r\right) = p_s^{n} \left[ \int_r^{R(z)} \left(\frac{r^2 - \lambda^2}{r}\right)^n dr - n\varsigma_A \int_r^{R(z)} \left(\frac{r^2 - \lambda^2}{r}\right)^{n-1} dr \right], \text{ when } \lambda_2 \le r \le R\left(z\right), \tag{17}$$

where  $u^+(r)$ ,  $u_p$  and  $u^{++}(r)$  represent the shear flow region from  $k \le r \le \lambda_1$ ,  $\lambda_1 \le r \le \lambda_2$  and  $\lambda_2 \le r \le R(z)$ , respectively. Absence of yield stress  $(\theta = 0)$  will result to  $\zeta_A = 0$ , where both Eqs. (15) and (17) will result to velocity field in the catheterised artery for power law fluid, a finding that is in line with Kapur [19]. To ensure continuous velocity distribution for the entire flow field, below condition needs to be fulfilled

$$u^{+}(r = \lambda_{1}) = u_{p} = u^{++}(r = \lambda_{2}).$$
 (18)

This gives

$$\int_{k}^{\lambda_{1}} \left( \frac{\lambda^{2} - r^{2}}{r} \right)^{n} dr - \int_{\lambda_{2}}^{R(z)} \left( \frac{r^{2} - \lambda^{2}}{r} \right)^{n} dr - n \zeta_{A} \left[ \int_{k}^{\lambda_{1}} \left( \frac{\lambda^{2} - r^{2}}{r} \right)^{n-1} dr - \int_{\lambda_{2}}^{R(z)} \left( \frac{r^{2} - \lambda^{2}}{r} \right)^{n-1} dr \right] = 0.$$
 (19)

Using Eqs. (12) and (14), Eq. (19) can be simplified into integral equation in  $\lambda_1$  as below

$$\int_{k}^{\lambda_{1}} \left( \frac{\lambda_{1} (\lambda_{1} + \zeta_{A}) - r^{2}}{r} \right)^{n} dr - \int_{\lambda_{1} + \zeta_{A}}^{R(z)} \left( \frac{r^{2} - \lambda_{1} (\lambda_{1} + \zeta_{A})}{r} \right)^{n} dr - n\zeta_{A} \left[ \int_{k}^{\lambda_{1}} \left( \frac{\lambda_{1} (\lambda_{1} + \zeta_{A}) - r^{2}}{r} \right)^{n-1} dr - \int_{\lambda_{1} + \zeta_{A}}^{R(z)} \left( \frac{r^{2} - \lambda_{1} (\lambda_{1} + \zeta_{A})}{r} \right)^{n-1} dr \right] = 0.$$
(20)

 $\lambda_1$  in Eq. (20) is solved numerically using Regula-Falsi method whereas the integrals is evaluated via Simpson's 3/8 rule. Once  $\lambda_1$  is known,  $\lambda_2$  can be determined using Eq. (14). According to Gill and Sankarasubramanian [12], the solution of Eq. (6) is computed as a series expansion and is displayed as

$$C(r,z,t) = C_m(z_1,t) + \sum_{i=R(z)}^{\infty} f_i(r,t) \frac{\partial^i C_m(z_1,t)}{\partial z_1^i},$$
(21)

where  $C_m$  is the mean concentration,  $f_i$  is dispersion function and a new axial coordinate moving with the average velocity is  $z_1 = z - u_m t$ . The distribution of  $C_m$  is diffusive as the time starts and thus, the GDM as appropriate functions of time t is given as

$$\frac{\partial C_m}{\partial t} = \sum_{i=1}^{\infty} K_i(t) \frac{\partial^i C_m}{\partial z_i^i} (z_1, t), \tag{22}$$

where  $K_i(t)$  is the dispersion coefficient. Substituting Eq. (21) into Eq. (6) and using Eq. (22) yields the series expansion. By equating the coefficients of  $\partial^i C_m/\partial z_1^i$  and let  $f_1(r,t)=f_{1s}(r)+f_{1t}(r,t)$ , where  $f_{1s}$  is the steady-state and  $f_{1t}$  is the unsteady state, the variable separation method and Bessel function can be used to solve the transient state,  $f_{1t}(r,t)$  of the dispersion function subject to the conditions  $f_{1t}(r,0)=-f_{1s}(r)$  and  $\partial f_{1t}/\partial r=0$ . The solution  $f_{1t}(r,t)$  is numerically computed using Simpson's 3/8 rule and presented as

$$f_{1t}(r,t) = \sum_{m=1}^{\infty} A_m B_0 e^{-(\lambda_m^2 + \gamma^2)t} , \qquad (23)$$

where

$$A_{m} = -\frac{\int_{k}^{R(z)} B_{0} f_{1s}(r) r dr}{\int_{k}^{R(z)} B_{0}^{2} r dr},$$
(24)

$$B_0 = J_1(\lambda_m k) Y_0(\lambda_m r) - Y_1(\lambda_m k) J_0(\lambda_m r)$$
(25)

and the factors  $\lambda_m$  are the solutions of the equation

$$J_{1}(\lambda_{m}R(z))Y_{1}(\lambda_{m}k)-J_{1}(\lambda_{m}k)Y_{1}(\lambda_{m}R(z))=0,$$
(26)

with  $J_0$ ,  $J_1$ ,  $Y_0$  and  $Y_1$  indicate the Bessel functions for first and second kind of order zero and one, respectively. By substituting Eq. (21) into the transport equation Eq. (6) yield a partial differential equation for mean concentration as below

$$\frac{\partial C_{m}}{\partial t} + \left(u - u_{m}\right) \frac{\partial C_{m}}{\partial z_{1}^{1}} - \frac{1}{Pe^{2}} \frac{\partial^{2} C_{m}}{\partial z_{1}^{2}} + \sum_{j=1}^{\infty} \left( \left( \frac{\partial f_{j}}{\partial t} - \frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial f_{j}}{\partial r} \right) \right) \frac{\partial^{j} C_{m}}{\partial z_{1}^{j}} + \left( u - u_{m} \right) f_{j} \frac{\partial^{j+1} C_{m}}{\partial z_{1}^{j+1}} - \frac{1}{Pe^{2}} f_{j} \frac{\partial^{j+2} C_{m}}{\partial z_{1}^{j+2}} + f_{j} \frac{\partial^{j+1} C_{m}}{\partial z_{1}^{j+1} \partial t} \right) = 0.$$
(27)

Substituting Eq. (22) in Eq. (27) and rearranging terms yields

$$\left[K_{1}(t)+(u-u_{m})+\frac{\partial f_{1}}{\partial t}-\frac{1}{r}\frac{\partial}{\partial r}\left(r\frac{\partial f_{1}}{\partial r}\right)\right]\frac{\partial C_{m}}{\partial z_{1}^{1}}+\left[-\frac{1}{Pe^{2}}+(u-u_{m})f_{1}+f_{1}K_{1}(t)+K_{2}(t)\right] + \frac{\partial f_{2}}{\partial t}-\frac{1}{r}\frac{\partial}{\partial r}\left(r\frac{\partial f_{2}}{\partial r}\right)\right]\frac{\partial^{2}C_{m}}{\partial z_{1}^{2}}+\sum_{j=1}^{\infty}\left[-\frac{1}{Pe^{2}}f_{j}+(u-u_{m})f_{j+1}+K_{j+2}(t)+\frac{\partial f_{j+2}}{\partial t}\right] - \frac{1}{r}\frac{\partial}{\partial r}\left(r\frac{\partial f_{j+2}}{\partial r}\right)+\sum_{i=1}^{j+1}K_{i}(t)f_{j+2-i}\frac{\partial}{\partial z_{1}^{j+2}}=0.$$
(28)

For j=1,2,..., we equalize  $\partial^i C_m/\partial z_1^j$  to zero to obtain the infinite system of partial differential equations as below

$$\frac{\partial f_1}{\partial t} - \frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial f_1}{\partial r} \right)_1 + u - u_m + K_1(t) = 0, \tag{29}$$

$$\frac{\partial f_2}{\partial t} - \frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial f_2}{\partial r} \right) + \left[ u - u_m + K_1(t) \right] f_1 + K_2(t) - \frac{1}{Pe^2} = 0, \tag{30}$$

$$\frac{\partial f_{i+2}}{\partial t} - \frac{1}{r} \frac{\partial}{\partial r} \left( r \frac{\partial f_{j+2}}{\partial r} \right) + \left[ u - u_m + K_1(t) \right] f_{j+1} + \left[ K_2(t) - \frac{1}{Pe^2} \right] f_j + \sum_{i=1}^{j+1} K_i(t) f_{j+2-i} = 0, \tag{31}$$

for j = 1, 2, .... with  $f_0 = 1$ .

By multiplying Eqs.(29), (30) and (31) with r followed by integrating the result from k to R(z) imply

$$K_1(t) = -\frac{2}{1-k^2} \int_k^{R(z)} (u - u_m) r dr = 0,$$
 (32)

$$K_{2}(t) = \frac{1}{Pe^{2}} - \frac{2}{1 - k^{2}} \int_{k}^{R(z)} f_{1} \ u \ r \ dr, \tag{33}$$

$$K_{j+2}(t) = -\frac{2}{1-k^2} \int_{k}^{R(z)} f_{j+1} \ u \ r \ dr, \ j=1,2,....$$
 (34)

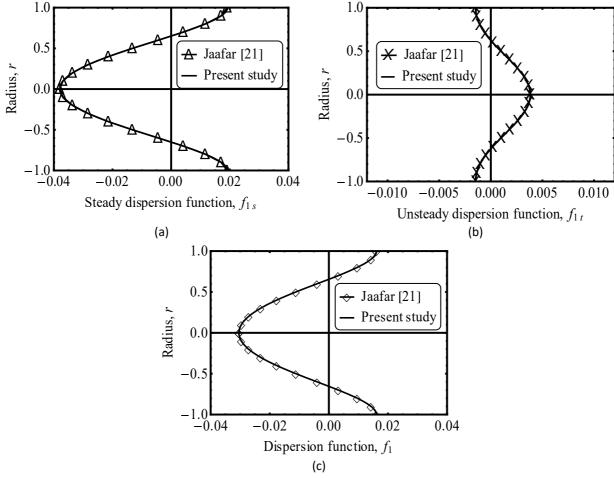
The dispersion coefficient  $K_2(t)$  is an indicator to measure the effectiveness of the solute dispersion in the blood flow. Ramana  $et\ al.$  [20] and Dash  $et\ al.$  [21] evaluated the overall reduction in solute dispersion due to the fluid yield stress at a constant pressure gradient by subtracting  $1/Pe^2$  and multiplying 192 to Eq. (33) resulting to

$$192\left(K_{2}(t) - \frac{1}{Pe^{2}}\right) = -\frac{2}{1 - k^{2}} \int_{k}^{R(z)} f_{1} \ u \ r \ dr. \tag{35}$$

#### 3. Results and Discussion

#### 3.1 Validation

The effect of varying catheter radius, power law index, and yield stress on the velocity profile and dispersion coefficient of solute is analysed, with the following range of parameters: k: 0.1-0.3,  $\theta$ : 0.1-0.3, and n: 0.75-2 [13,22]. The results and data were generated for both comparison and validation purposes using the Mathematica software. Figures 2 (a), 2 (b) and 2 (c) confirmed that the steady dispersion function  $f_{1s}$ , unsteady dispersion function  $f_{1t}$  and dispersion function  $f_1$  of solute obtained in this study is comparable with the findings of Jaafar [23]. For validation purposes, k and  $\delta$  was set to zero while the geometry of the stenosed artery, R(z) was set to one to resemble fluid without stenosis and catheter.



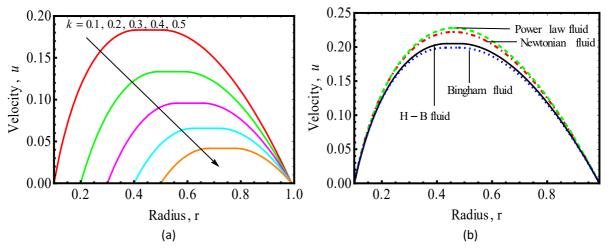
**Fig. 2.** The parameters are fixed at  $n=0.95, \beta=0.1, R(z)=1, \delta=0, k=0$  and  $\theta=0.1$  (a) steady dispersion function  $f_{1s}$ , (b) unsteady dispersion function  $f_{1t}$  with t=0.1 and (c) dispersion function  $f_1$  with t=0.1

## 3.2 Velocity

It is crucial to ensure drug is quickly and efficiently dispersed as soon as possible in order to cure relevant disease. In this case, blood velocity will play a vital role in this direction as it influences the convection and dispersion coefficient [24]. Figure 3 (a) illustrates the variation in velocity distribution with different catheter radius when  $p_s=1$ ,  $\theta=0.1$ , n=0.95,  $l_0=3$ , d=2, z=4 and  $\delta=0.01$ . In the case of constant yield stress, increase in catheter radius will result to decrease in

axial velocity because of the reduction in annular flow in the area between catheter wall and arterial wall.

From the velocity distribution recorded by various fluids in Figure 3 (b), we can see that highest velocity is recorded by power law fluid compare with fluids having yield stress; for example, for n=0.95, power law fluid velocity is higher than Newtonian fluid when n=1,  $\theta=0$ . However, for specific values of k and  $\theta$ , improvement in n results to lessen velocity when  $p_s=1$ ,  $l_0=3$ , d=2, z=4 and  $\delta=0.01$ . Comparing between H-B fluid and Bingham fluid under a similar yield stress, it is shown that as power law index increases from n=0.95 to n=1, the axial velocity slightly decreases. Bessonov  $et\ al.\ [25]$  reported that the erythrocyte concentration is higher near the axis, whereas the platelets are concentrated near the wall. Sankar and Hemalatha [13] stated that results of velocity distribution discovered by Dash  $et\ al.\ [21]$  for Casson fluid are significantly lesser compared to H-B fluid.



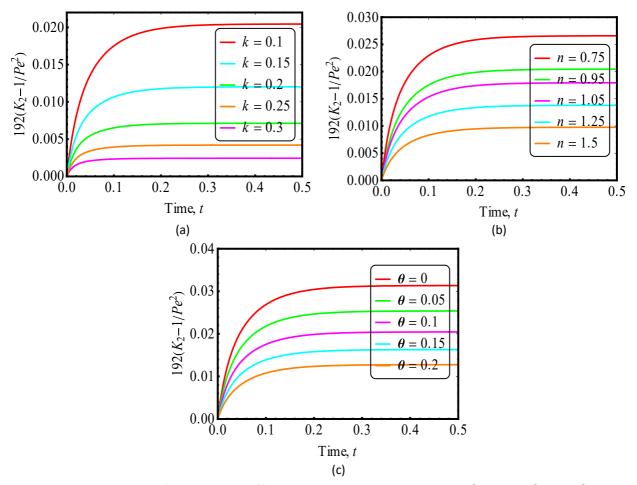
**Fig. 3.** Variation of the velocity distribution when  $p_s=1$ ,  $\delta=0.01$ ,  $l_0=3$ , d=2 and z=4 for (a) different catheter radius k with n=0.95 and  $\theta=0.1$  (b) different fluids with k=0.1

# 3.3 Dispersion Coefficient

Dispersion coefficient  $K_2(t)$  explains the overall dispersion process in terms of simple diffusion process as a function of time. The results when k=0 and  $\theta=0$  reduce to those of [12]. Figure 4 (a) illustrates the variation of dispersion coefficient over time t for various catheter size k when  $n=0.95, p_s=1, \theta=0.1, \delta=0.01, l_0=3, d=2$  and z=4. It is observed that with an increase in the catheter size from k=0.1,0.15,0.2,0.25 to 0.3, the dispersion coefficient reduces significantly. As the catheter size increase, the annular gap between stenosed arterial wall and catheter wall decreases because reduction in annular gap hampers dispersion process. The same behaviour was also noticed by Rao and Desikachar [26] for a Newtonian fluid.

Figure 4 (b) shows the variation of dispersion coefficient over time t for various power law index n when  $p_s=1$ ,  $\theta=0.1$ , k=0.1,  $\delta=0.01$ ,  $l_0=3$ , d=2 and z=4. The dispersion coefficient of the solute diminish as the power law index n increases. As mentioned by Hussain et~al. [27], the power-law index represents the apparent whole blood viscosity. Power law index plays an important role to control viscosity and velocity of a fluid. Physically, the viscosity of the fluid increases as the power-law index increases, hence blood travels faster along the axial distance. When the viscosity increases in the blood flow, the solute movement becomes slower, and hence, the dispersion coefficient decreases.

Figure 4 (c) elucidates the variation of dispersion coefficient over time t for various yield stress  $\theta$  when  $p_s=1, n=0.95, k=0.1, l_0=3, d=2, z=4$  and  $\delta=0.01$ . The dispersion coefficient decreases significantly as  $\theta$  increases from  $\theta=0,0.05,0.1,0.15$  and 0.2. At t=0.4, the dispersion coefficient reaches a steady-state of diffusion when  $K_2(t)=0.083,0.0080,0.0070,0.0065$  and 0.0063, respectively with an increase in  $\theta$ . The dispersion coefficient rises rapidly from t=0 to 0.2, then slowly from t=0.2 to 0.4 and becomes almost constant from t=0.4 to 0.5. It can be seen that the dispersion coefficient  $K_2(t)$  changes quickly for short time scale but it does not change much for large time scale. Specifically for Newtonian fluid ( $\theta=0, n=1$ ), it attains steady-state at time t of around 0.5. Due to yield stress, time recorded will be lower because yield stress corresponds to the non-Newtonian nature of the fluid where improvement in yield stress equals to higher blood viscosity. Apart from that, yield stress is also related to the width of the plug region where improvement in said width will improve yield stress as portrayed in Eq. (14).



**Fig. 4.** Variation of dispersion coefficient  $K_2$  over time t when  $p_s=1$ ,  $\delta=0.01$ ,  $l_0=3$ , d=2 and z=4 for (a) different catheter radius k with n=0.95 and  $\theta=0.1$  (b) different power law index with k=0.1 and  $\theta=0.1$  (c) different yield stress with n=0.95 and k=0.1

# 4. Conclusion

The present study investigated the influence of catheter radius, power law index, and yield stress of the fluid on the solute dispersion process in the cardiovascular system. The important findings are outlined below:

- Upon inserting a catheter into the lumen of the artery, the presence of a plug flow region results in the production of two yield plane locations;
- The velocity reduces following an increase in the yield stress, catheter size, and power law index. The dispersion coefficient exhibits a same behaviour as the aforementioned parameters ascend considerably;
- The dispersion coefficient exhibits two distinctive behaviours which are linear and nonlinear behaviour.

Since this model did not consider the flow pulsatility and the porosity of the arterial tissue, hence, future researchers are recommended to extend this study.

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