

**PHYSICAL HYDRODYNAMIC PROPULSION MODEL STUDY FOR
CREEPING VISCOUS FLOW THROUGH A CILIATED POROUS TUBE**

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Abstract: The present investigation focuses on a mathematical study of creeping viscous flow induced by metachronal wave propagation in a horizontal ciliated tube containing porous media. Creeping flow limitations are imposed i.e. inertial forces are small compared with viscous forces and therefore very low Reynolds number ($Re \ll 1$) is taken into account. The wavelength of metachronal wave is also considered as very large for cilia movement. The physical problem is linearized and exact solutions are developed for the differential equation problem. Mathematica software is used to compute and illustrate numerical results. The influence of slip parameter and Darcy number on velocity profile, pressure gradient and trapping of bolus are discussed with the aid of graphs. It is found that with increasing magnitude of slip parameter the trapped bolus inside the streamlines increases in size. The study is relevant to biological propulsion of medical micro-machines in drug delivery.

Keywords: *Viscous fluid; Ciliated tube; Porous medium; Darcy number; Permeability; Low Reynolds number; Hydrodynamic slip; Biomimetic propulsion.*

Nomenclature

a radius of tube

c wave speed

Da Darcy number

F flow rate in moving frame

\bar{h} height of tube

K biological medium permeability

\bar{P} pressure in the moving frame

\bar{p} pressure in the fixed frame

ΔP pressure rise

Q flow rate in the fixed frame

\bar{r}, \bar{z}	<i>Cylindrical coordinates in the fixed frame</i>
\bar{R}, \bar{Z}	<i>Cylindrical coordinates in the moving frame</i>
t	<i>time</i>
\bar{w}, \bar{u}	<i>Velocity components in the fixed frame</i>
\bar{W}, \bar{U}	<i>Velocity components in the moving frame</i>
\bar{Z}_0	<i>Reference position of particle</i>

Greek

λ	<i>wavelength</i>
α	<i>eccentricity of the elliptical motion</i>
α_1	<i>hydrodynamic slip parameter</i>
β	<i>wave number</i>
ε	<i>cilia length</i>
λ	<i>wavelength</i>
μ	<i>dynamic viscosity of fluid</i>
ρ	<i>density of fluid</i>

1. INTRODUCTION

Biological propulsion is increasingly attracting the interest of engineers owing to diverse applications in medicine, aerospace, naval and even geological systems. Many mathematical models have been proposed for different organisms and micro-organisms at different length scales and Reynolds numbers. A most lucid review of the subject has been given by Wu [1] in which 18 orders of magnitude have been identified for the range of flows studied from microscopic organisms to large marine mammals (e.g. the blue whale). At the microscopic scale and in particular in the context of embryological systems, cilia and flagellar propulsion are of great interest. This area of biological hydrodynamic propulsion has also motivated significant interest for many decades, as elaborated by Wu *et al.* [2]. Cilia and flagella lengths may span from a few microns to more than 2 mm in the case of some insect sperm flagella. Hence, cilia and flagella essentially have no difference, although they have been ascribed different

terminologies prior to a proper biological examination of their structures. Cilia have been established as beating with a *whip-like asymmetric mechanism* which comprises both an effective stroke and a recovery stroke. Moreover, when many cilia function collectively, fluid dynamic interactions may induce beating out-of-phase, and this manifests in the generation of metachronal waves, and exacerbated hydrodynamics flow. The specific metachrony is termed symplectic (or antiplectic) when the metachronal wave is in the same (or opposite) direction as the effective stroke. These features have recently attracted attention in biomimetics and bio-inspired engineering systems, notably in medical micro-swimmers which offer tremendous potential in nano-medicine, drug-delivery and so forth, as described by Feng and Cho [3]. Commonly, cells own one or two long flagella. Conversely ciliated cells have many short cilia. For instance, the mammalian spermatozoa has a single flagellum, the unicellular green alga *Chlamydomonas* has two flagella, and the unicellular protozoan *Paramecium* is covered with a few thousand cilia, these are used for both locomotion and nutrition. In mammals, many epithelial cells are ciliated in order to sweep materials across the tissue surface. For instance, considerable numbers of cilia (more than $10^7/\text{mm}^2$) cover the surfaces of mammalian respiratory passages (the nose, pharynx, and trachea), here they dislodge and expel particulate matter that collects in the mucus secretions of these tissues. The motion of cilia and flagella is controlled by the Stokes equations (linear) with no-slip boundary conditions on their surfaces and vanishing fluid disturbance at infinity [4]. The structure of cilia, factors which affect cilia activities, movement of cilia and flagella and the coordination of the beating of cilia have been addressed analytically by Sleight [5]. Lardner and Shack [6] examined cilia effects on transport flow rates in the ductus efferentes of the male reproductive tract. A mathematical model to represent the microstructure of ciliated organisms was developed by Blake [7]. In continuation of the studies on cilia movement and its importance in various fields of research, Wu [8] theoretically investigated the fluid mechanism of cilia motion. Brennen [9] presented an oscillatory thin shear layer (boundary layer) theory for cilia movement. Sleight and Aiello [10] studied water movement by cilia. Agarwal and Uddin [11] investigated the fluid flow with variable viscosity by cilia transport; Blake [12] developed a spherical envelope approach for cilia motion. Miller [13] focused on the movement of Newtonian fluids sustained by mechanical cilia oscillation; Barton and Raynor [14] presented an analytical approach for cilia induced mucous flow; Smith *et*

al. [15] addressed viscoelastic fluid flow effects on cilia transport. Dauplain *et al.* [16] studied computationally the hydrodynamics of cilia movement from a fluid-structure interaction (FSI) viewpoint, specifically simulating *ctenophore Pleurobrachia pileus* marine micro-organisms (quasi-spherical geometry) for multiple flexible cilia with 3-dimensional models. Khaderi and Onck [17] presented three dimensional computations for cilia movement. Khaderi *et al.* [18] further studied the non-reciprocal motion due to beating of artificial cilia. Another interesting communication by Khaderi *et al.* [19] reports on magnetically-actuated artificial cilia for microfluidic propulsion.

The above biological propulsion simulations were restricted to purely fluid media. In recent years significant interest has also developed in *transport phenomena in biological porous media*. Many organs in the human body comprise permeable materials including kidneys, lungs, tissue and the human skin [20]. Porous media hydrodynamics further arises in circulation of cerebrospinal fluid in the brain. In biomedical engineering, flow in porous media is integral to the fabrication of artificial physiological organs including artificial kidneys, gastric tract and lungs [21]. Almost invariably, biological materials (tissue etc) are heterogeneous and exhibit strongly anisotropy. These complex constitutive properties originate from the cellular microstructure. Although intrinsically the porous media arising in different organs of the body are anisotropic, approximations can be made to simulate transport through them. A popular approach is the Darcy model which has been used for microvasculature via networks of one-dimensional segments, whereas the capillary bed is often approximated as a spatially-averaged Darcy compartment. Darcy's law and its modification assumes that the rate at which a fluid flows through a permeable substance per unit area is equal to the product of the *permeability* (a property of the material through which the fluid is flowing) and the pressure drop per unit length of flow, divided by the viscosity of the fluid. A porous medium is generally defined in continuum mechanics [22] as a medium which contains a number of small holes distributed throughout the matter. Khaled and Vafai [23] have shown that convective flow models for porous media are accurate for modelling hemodynamics in tumours and muscles and in particular mimic quite well the impeded transport of physiological fluids in vessels clogged with fatty cholesterol plaques and clots. Interstitial fluid mechanics also lends itself well to porous media transport since the tissue may be approximated as a medium of dispersed cells

separated by connective voids, the latter permitting percolation of nutrients, minerals and other substances to cells within the tissue. Darcy's law however neglects the friction within the fluid and exchange of momentum between the fluid and solid phases. Modifications of Darcy models have therefore been developed for some time. A good example of flow with boundary condition at the interfaces of a porous medium was presented by Saffman [24]. Most physiological vessels feature porous media characteristics and considering this fact many works have been reported in literature. An interesting transport problem in porous media (of the moving boundary type) is *peristalsis*, and arises in intestinal fluid dynamics. Representative studies in this area have been communicated by for example Miyamoto *et al.* [25] who studied the two-dimensional laminar flow in a circular porous tube and considering a small water absorption or secretion in the intestinal perfusion experiment whereas Jeffrey *et al.* [26] discussed the flow fields generated by peristaltic reflex in isolated guinea pig ileum. Elshehawey *et al.* [27] developed another model for peristaltic transport through an asymmetric porous medium channel and focused the application to intra uterine fluid motion in a sagittal cross-section of the uterus. Other recent investigations on peristaltic flow of Newtonian fluid and non-Newtonian fluids have been presented and span many multi-physical aspects of the subject including power law fluids [28], magnetohydrodynamics [29] and Maxwell rheological pumping through porous media [30].

A careful inspection of the scientific literature has revealed however that the creeping flow induced by metachronal wave propagations of viscous fluid in a porous ciliated tube is yet to be studied mathematically in the literature. Porous media offer a well-tested mechanism for flow control in hydrodynamic propulsion. The present work aims to therefore address this topic for the first time. Analytical solutions are developed for the linear partial differential equations generated. Linearization is achieved by neglecting inertial forces relative to viscous forces, using the low Reynolds number approximation from hydrodynamic lubrication theory. Numerical results are presented graphically and the influence of key geometric and hydromechanical parameters (e.g. Darcy number) on transport characteristics is elaborated in detail. The simulations presented herein further elucidate the mechanism of creeping viscous flow induced by cilia motion and are relevant to gastric fluid mechanics and also biomimetic devices exploiting surface modifications for optimized propulsion [31-35].

2. MATHEMATICAL MODEL

Let us consider incompressible Newtonian flow in a ciliated tube with hydrodynamic slip at the wall. The tube interior wall is ciliated with metachronal waves and the flow is generated due to the collective beating of cilia. We select a cylindrical coordinate system (\bar{R}, \bar{Z}) , where the \bar{Z} - axis is orientated along the center line of the tube and \bar{R} - axis is normal to it. Cilia deform in a wave-like fashion with a propagation velocity, c , along the wall of the outer tube. Keeping in view the geometry of the metachronal wave pattern, it is assumed that the envelope of cilia tips can be expressed mathematically in the following form:

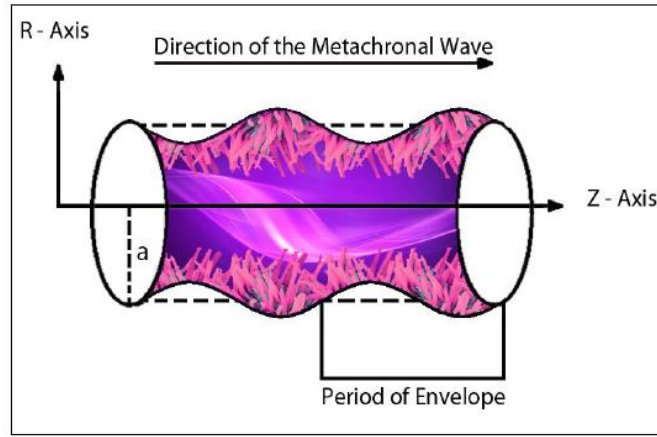


Fig. 1. Geometry of the problem

Fig. 1: Schematic of the biological propulsion problem

The envelopes of the cilia tips can be expressed mathematically [1, 9] as follows:

$$\begin{aligned}\bar{R} = \bar{H} = \bar{f}(\bar{Z}, \bar{t}) &= a + a\varepsilon \cos\left(\frac{2\pi}{\lambda}(\bar{Z} - c\bar{t})\right), \\ \bar{Z} = \bar{g}(\bar{Z}, \bar{Z}_0, \bar{t}) &= a + a\varepsilon\alpha \sin\left(\frac{2\pi}{\lambda}(\bar{Z} - c\bar{t})\right),\end{aligned}\quad (1)$$

where a denotes the mean radius of the tube, ε is the non-dimensional measure with respect to the cilia length, λ and c are the wavelength and wave speed of the metachronal wave respectively, \bar{Z}_0 denotes the reference position of the particle and α is a measure of the eccentricity of the elliptical motion. The velocities of the transport fluid are essentially those

induced by the cilia tips, and may be defined as:

$$\begin{aligned}\bar{W} &= \left. \frac{\partial \bar{Z}}{\partial \bar{t}} \right|_{\bar{z}_0} = \frac{\partial \bar{g}}{\partial \bar{t}} + \frac{\partial \bar{g}}{\partial \bar{Z}} \frac{\partial \bar{Z}}{\partial \bar{t}} = \frac{\partial \bar{g}}{\partial \bar{t}} + \frac{\partial \bar{g}}{\partial \bar{Z}} \bar{W}, \\ \bar{U} &= \left. \frac{\partial \bar{R}}{\partial \bar{t}} \right|_{\bar{z}_0} = \frac{\partial \bar{f}}{\partial \bar{t}} + \frac{\partial \bar{f}}{\partial \bar{Z}} \frac{\partial \bar{Z}}{\partial \bar{t}} = \frac{\partial \bar{f}}{\partial \bar{t}} + \frac{\partial \bar{f}}{\partial \bar{Z}} \bar{W}.\end{aligned}\quad (2)$$

Using Eqn. (2) in Eqn. (1) , we get

$$\begin{aligned}\bar{W} &= \frac{-\frac{2\pi}{\lambda} [\varepsilon \alpha a c \cos(\frac{2\pi}{\lambda})(\bar{Z} - c\bar{t})]}{\left[1 - \frac{2\pi}{\lambda} \left\{ \varepsilon \alpha a \cos(\frac{2\pi}{\lambda})(\bar{Z} - c\bar{t}) \right\}\right]}, \\ \bar{U} &= \frac{\frac{2\pi}{\lambda} [\varepsilon \alpha c \sin(\frac{2\pi}{\lambda})(\bar{Z} - c\bar{t})]}{\left[1 - \frac{2\pi}{\lambda} \left\{ \varepsilon \alpha a \cos(\frac{2\pi}{\lambda})(\bar{Z} - c\bar{t}) \right\}\right]}.\end{aligned}\quad (3)$$

In the fixed coordinates (\bar{R}, \bar{Z}) , the flow in the tube is *unsteady*. It becomes steady in a *wave frame* (\bar{r}, \bar{z}) moving with the same speed as the wave moves in the \bar{Z} – direction. The governing equations for the viscous flow through porous medium in moving frame is expressed as:

Continuity equation:

$$\frac{1}{\bar{R}} \frac{\partial (\bar{R}\bar{U})}{\partial \bar{R}} + \frac{\partial \bar{W}}{\partial \bar{Z}} = 0, \quad (4)$$

R-direction momentum equation:

$$\begin{aligned}\rho \left[\frac{\partial \bar{U}}{\partial \bar{t}} + \bar{U} \frac{\partial \bar{U}}{\partial \bar{R}} + \bar{W} \frac{\partial \bar{U}}{\partial \bar{Z}} \right] &= -\frac{\partial \bar{P}}{\partial \bar{R}} + \mu \frac{\partial}{\partial \bar{R}} \left[2 \frac{\partial \bar{U}}{\partial \bar{R}} \right] + \mu \frac{2}{\bar{R}} \left(\frac{\partial \bar{U}}{\partial \bar{R}} - \frac{\bar{U}}{\bar{R}} \right) \\ &+ \mu \frac{\partial}{\partial \bar{Z}} \left[\left(\frac{\partial \bar{U}}{\partial \bar{R}} + \frac{\partial \bar{W}}{\partial \bar{Z}} \right) \right],\end{aligned}\quad (5)$$

Z-direction momentum equation:

$$\begin{aligned}\rho \left[\frac{\partial \bar{W}}{\partial \bar{t}} + \bar{U} \frac{\partial \bar{W}}{\partial \bar{R}} + \bar{W} \frac{\partial \bar{W}}{\partial \bar{Z}} \right] &= -\frac{\partial \bar{P}}{\partial \bar{Z}} + \mu \frac{\partial}{\partial \bar{Z}} \left[2 \frac{\partial \bar{W}}{\partial \bar{Z}} \right] + \mu \frac{1}{\bar{R}} \frac{\partial}{\partial \bar{R}} \left[\bar{R} \left(\frac{\partial \bar{U}}{\partial \bar{Z}} + \frac{\partial \bar{W}}{\partial \bar{R}} \right) \right] \\ &- \frac{\mu}{K} \bar{W},\end{aligned}\quad (6)$$

The transformations between the two frames are:

$$\bar{r} = \bar{R}, \quad \bar{z} = \bar{Z} - c\bar{t}, \quad \bar{u} = \bar{U}, \quad \bar{w} = \bar{W} - c, \quad \bar{p}(\bar{z}, \bar{r}) = \bar{P}(\bar{Z}, \bar{R}, \bar{t}) \quad (7)$$

The governing equations for the regime can be written via these transformations as follows:

$$\frac{1}{\bar{r}} \frac{\partial(\bar{r}\bar{u})}{\partial\bar{r}} + \frac{\partial\bar{w}}{\partial\bar{z}} = 0, \quad (8)$$

$$\begin{aligned} \rho \left[\bar{u} \frac{\partial\bar{u}}{\partial\bar{r}} + \bar{w} \frac{\partial\bar{u}}{\partial\bar{z}} \right] &= -\frac{\partial\bar{P}}{\partial\bar{r}} + \mu \frac{\partial}{\partial\bar{r}} \left[2 \frac{\partial\bar{u}}{\partial\bar{r}} \right] + \mu \frac{2}{\bar{r}} \left(\frac{\partial\bar{u}}{\partial\bar{r}} - \frac{\bar{u}}{\bar{r}} \right) \\ &+ \mu \frac{\partial}{\partial\bar{z}} \left[\left(\frac{\partial\bar{u}}{\partial\bar{r}} + \frac{\partial\bar{w}}{\partial\bar{z}} \right) \right], \end{aligned} \quad (9)$$

$$\begin{aligned} \rho \left[\bar{u} \frac{\partial\bar{w}}{\partial\bar{r}} + \bar{w} \frac{\partial\bar{w}}{\partial\bar{z}} \right] &= -\frac{\partial\bar{P}}{\partial\bar{z}} + \mu \frac{\partial}{\partial\bar{z}} \left[2 \frac{\partial\bar{w}}{\partial\bar{z}} \right] + \mu \frac{1}{\bar{r}} \frac{\partial}{\partial\bar{r}} \left[\bar{r} \left(\frac{\partial\bar{u}}{\partial\bar{z}} + \frac{\partial\bar{w}}{\partial\bar{r}} \right) \right] \\ &- \frac{\mu}{K} (\bar{w} + c), \end{aligned} \quad (10)$$

Corresponding boundary conditions are as defined in Ellahi et al. [31], Nadeem and Sadaf [35].

$$\frac{\partial\bar{w}}{\partial\bar{r}} = 0, \quad \text{at} \quad r = 0, \quad (11)$$

$$w = -c - c \frac{2\pi\epsilon\alpha\beta\cos(2\pi z)}{1 - 2\pi\epsilon\alpha\beta\cos(2\pi z)} - \frac{K}{\alpha_1^*} \frac{\partial\bar{w}}{\partial\bar{r}}, \quad \text{at } r = \bar{h}(z).$$

Where $c \frac{2\pi\epsilon\alpha\beta\cos(2\pi z)}{1 - 2\pi\epsilon\alpha\beta\cos(2\pi z)}$ is cilia factor.

Introducing the following non-dimensional variables, which are defined in the nomenclature:

$$r = \frac{\bar{r}}{a}, \quad z = \frac{\bar{z}}{\lambda}, \quad w = \frac{\bar{w}}{c}, \quad u = \frac{\lambda\bar{u}}{ac}, \quad p = \frac{a^2\bar{p}}{c\lambda\mu}, \quad \beta = \frac{a}{\lambda}, \quad D_a = \frac{K}{a^2}, \quad \alpha_1 = \frac{\alpha_1^*}{a}. \quad (12)$$

Making use of these variables in Eqns. (9,10), and using the assumptions of low Reynolds number and long wavelength, the non-dimensional governing equations can be written as:

$$\frac{\partial p}{\partial r} = 0, \quad (13)$$

$$\frac{dp}{dz} = \frac{1}{r} \frac{\partial}{\partial r} \left(r \frac{\partial w}{\partial r} \right) - \frac{1}{D_a} (w+1), \quad (14)$$

The non-dimensional boundary conditions on the ciliated porous walls are prescribed as:

$$\frac{\partial w}{\partial r} = 0, \quad \text{at} \quad r = 0, \quad (14a)$$

$$w = -1 - \frac{2\pi\varepsilon\alpha\beta\cos(2\pi z)}{1 - 2\pi\varepsilon\alpha\beta\cos(2\pi z)} - \frac{\sqrt{D_a}}{\alpha_1} \frac{\partial w}{\partial r}, \quad \text{at } r = h(z) = 1 + \varepsilon \cos(2\pi z). \quad (14b)$$

3. ANALYTICAL SOLUTIONS

Integrating Eqn. (13) and using boundary conditions (14 a, b), *axial velocity* is obtained as:

$$w(r, z) = -1 - D_a \frac{dp}{dz} + \frac{\alpha_1 \left(D_a \frac{dp}{dz} - \frac{2\pi\varepsilon\alpha\beta\cos(2\pi z)}{1 - 2\pi\varepsilon\alpha\beta\cos(2\pi z)} \right) I_0\left(\frac{r}{\sqrt{D_a}}\right)}{I_1\left(\frac{h}{\sqrt{D_a}}\right) - I_0\left(\frac{h}{\sqrt{D_a}}\right)}, \quad (15)$$

The volumetric flow rate (F) is defined as:

$$F = 2\pi \int_0^h r w dr. \quad (16)$$

Integrating the above expression for flow rate, we have:

$$\frac{dp}{dz} = \frac{I_1\left(\frac{h}{\sqrt{D_a}}\right) \left((2\pi\varepsilon\alpha\beta\cos(2\pi z) - 1)(F + \pi h^2) - 2\pi\alpha_1\sqrt{D_a} h n \right) - (F + \pi h^2) I_0\left(\frac{h}{\sqrt{D_a}}\right)}{\pi D_a h \left(h I_0\left(\frac{h}{\sqrt{D_a}}\right) - \left(h - 2\alpha_1\sqrt{D_a} \right) I_1\left(\frac{h}{\sqrt{D_a}}\right) \right)} \quad (17)$$

Flow rate in moving and fixed frames are related as follows:

$$Q = F + \frac{1}{2} \left(1 + \frac{\varepsilon^2}{2} \right). \quad (18)$$

The pressure rise (ΔP) is readily found by integrating the pressure gradient w.r.t. z over the interval $[0, 1]$ as follows:

$$\Delta P = \int_0^1 \frac{dp}{dz} dz. \quad (19)$$

The *stream function* in the wave frame (obeying the Cauchy-Riemann equations,

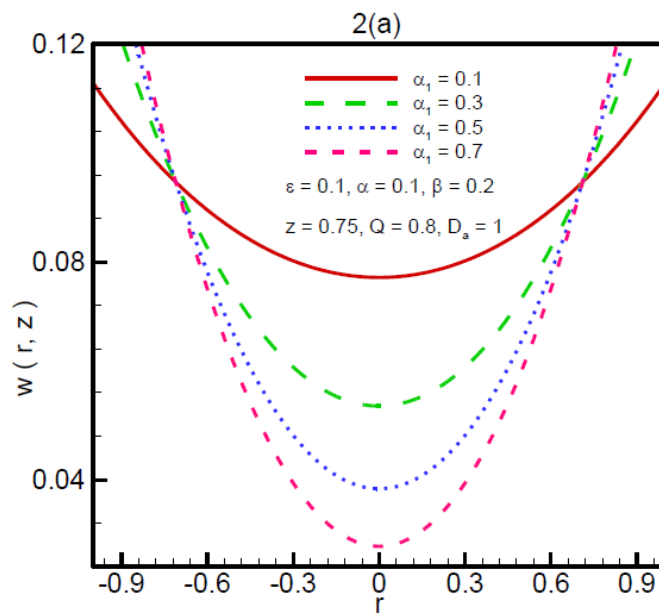
$u = \frac{\partial \psi}{\partial y}$ and $v = -\frac{\partial \psi}{\partial x}$) may be computed by using Eqn. (18). Visualization of streamlines is

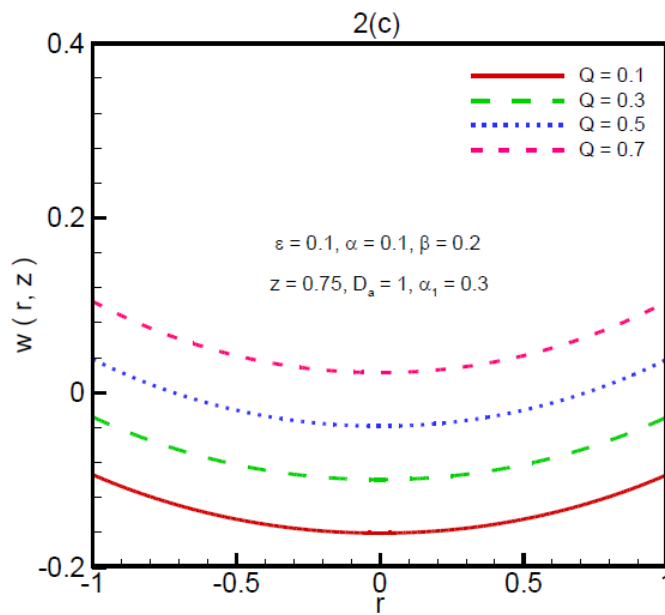
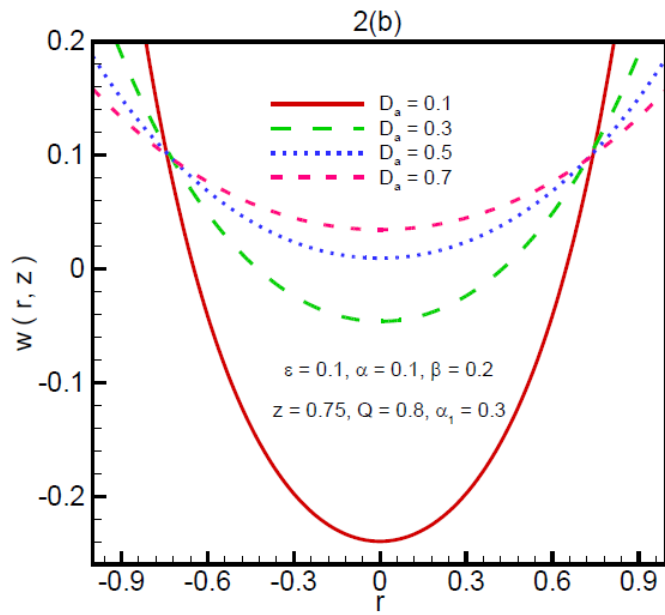
achieved with Mathematica symbolic software.

4. NUMERICAL RESULTS AND DISCUSSION

In this section, the effect of several physical parameters i.e. slip parameter (α_1) and Darcy number (Da) on velocity profile, flow rate, pressure gradient and stream lines are elaborated with the help of Figs. (2-5).

Figs. 2 represent the change in the velocity of the fluid with respect to certain changes in the constraints.





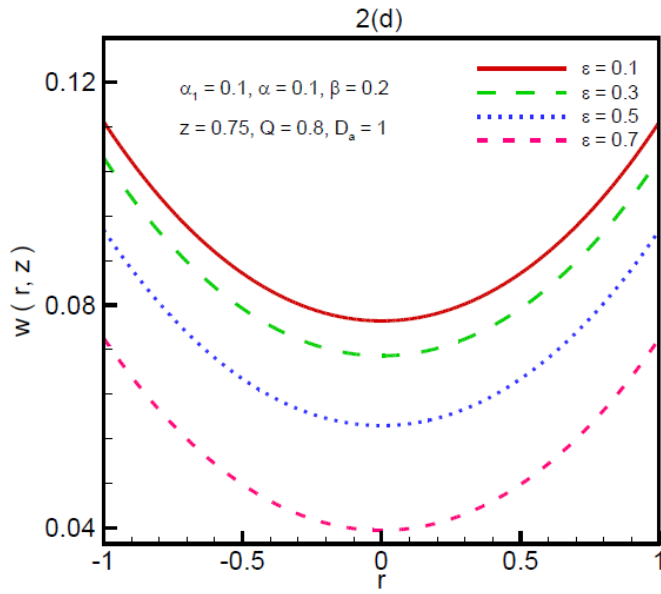
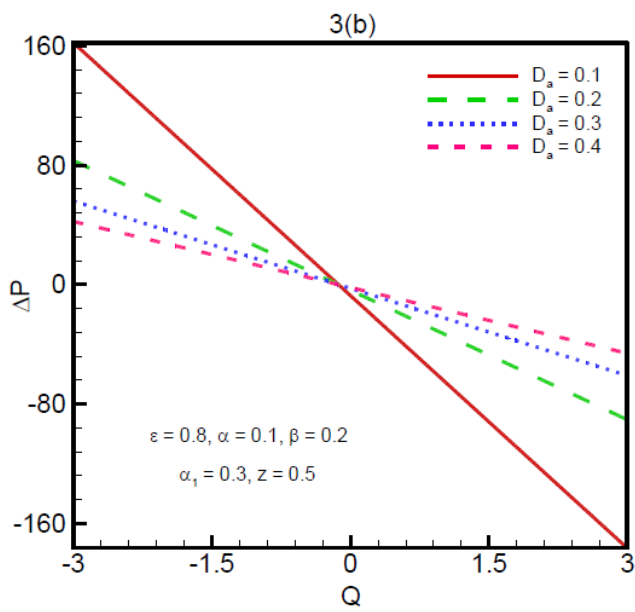
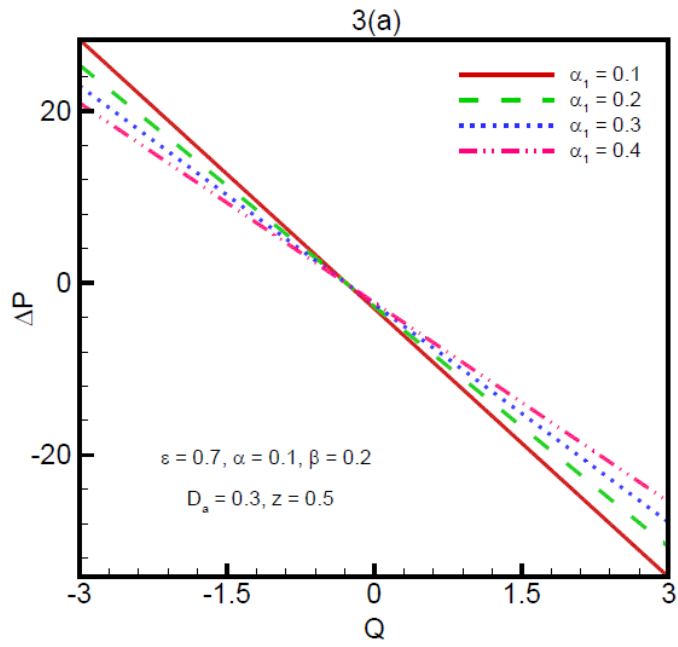


Fig. 2. Velocity profile (axial velocity vs. radial coordinate) for (a) Slip parameter slip parameter α_1 . (b) Darcy number D_a . (c) Flow rate Q . (d) Cilia length parameter ϵ .



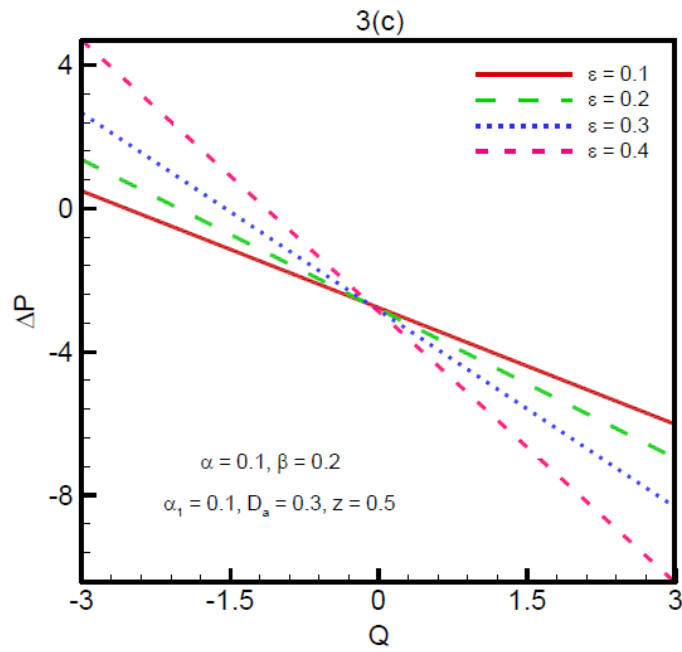
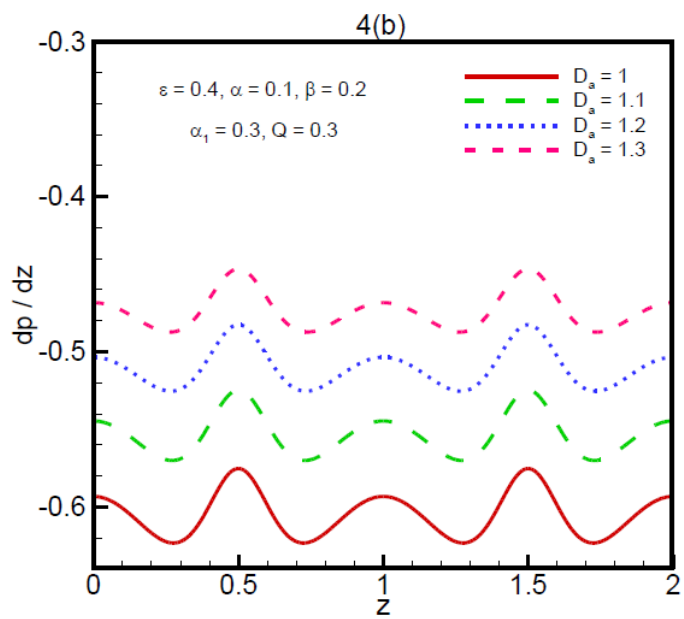
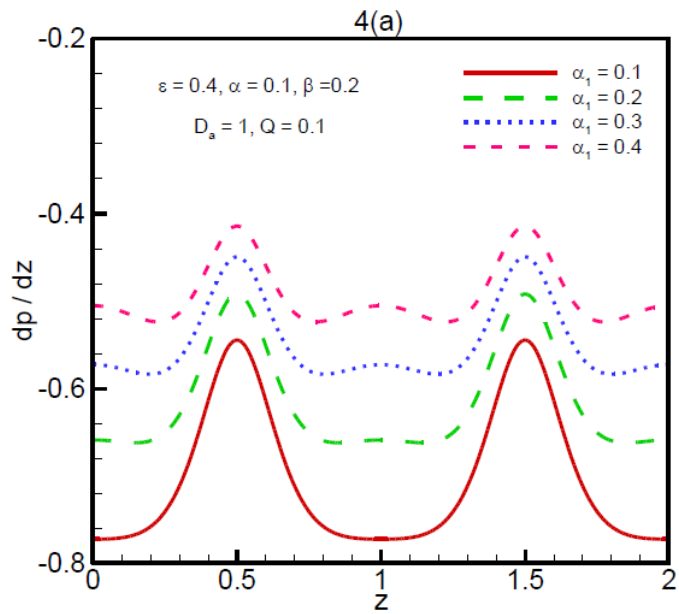


Fig. 3. Pressure rise versus flow rate for (a) Slip parameter slip parameter α_1 . (b) Darcy number D_a . (c) Cilia length parameter ε .



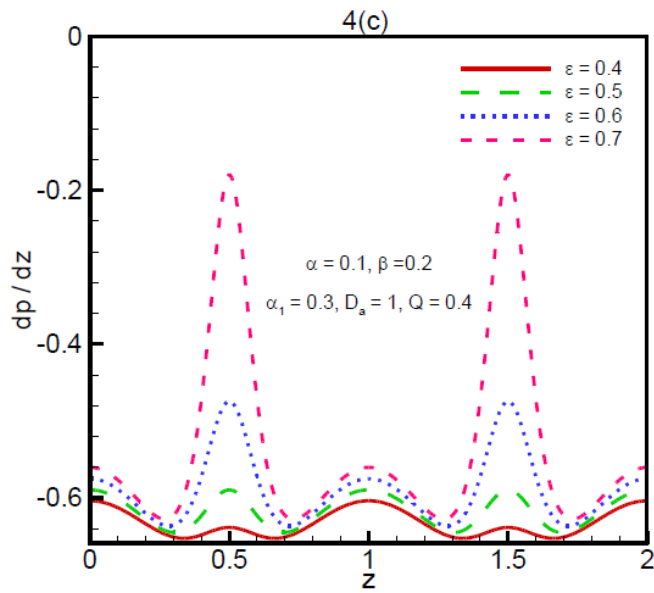
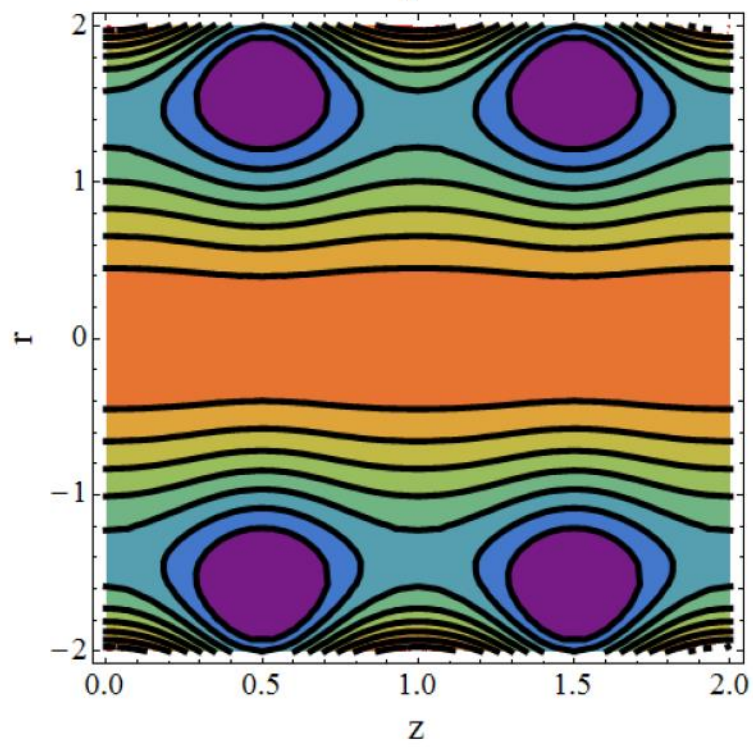
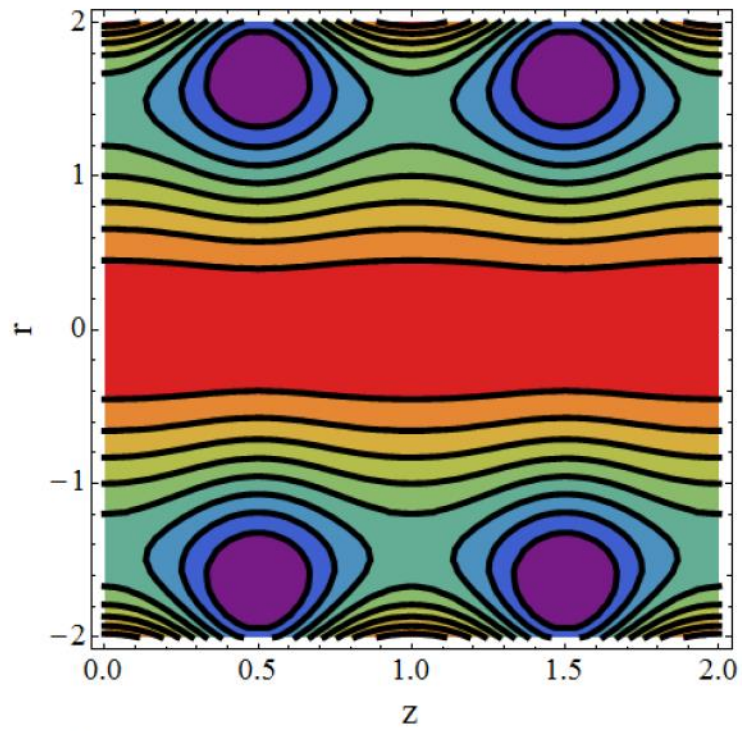


Fig.4. Pressure gradient vs. axial coordinate for (a) Slip parameter slip parameter α_1 . (b) Darcy number D_a . (c) Flow rate Q . (d) Cilia length parameter ε .



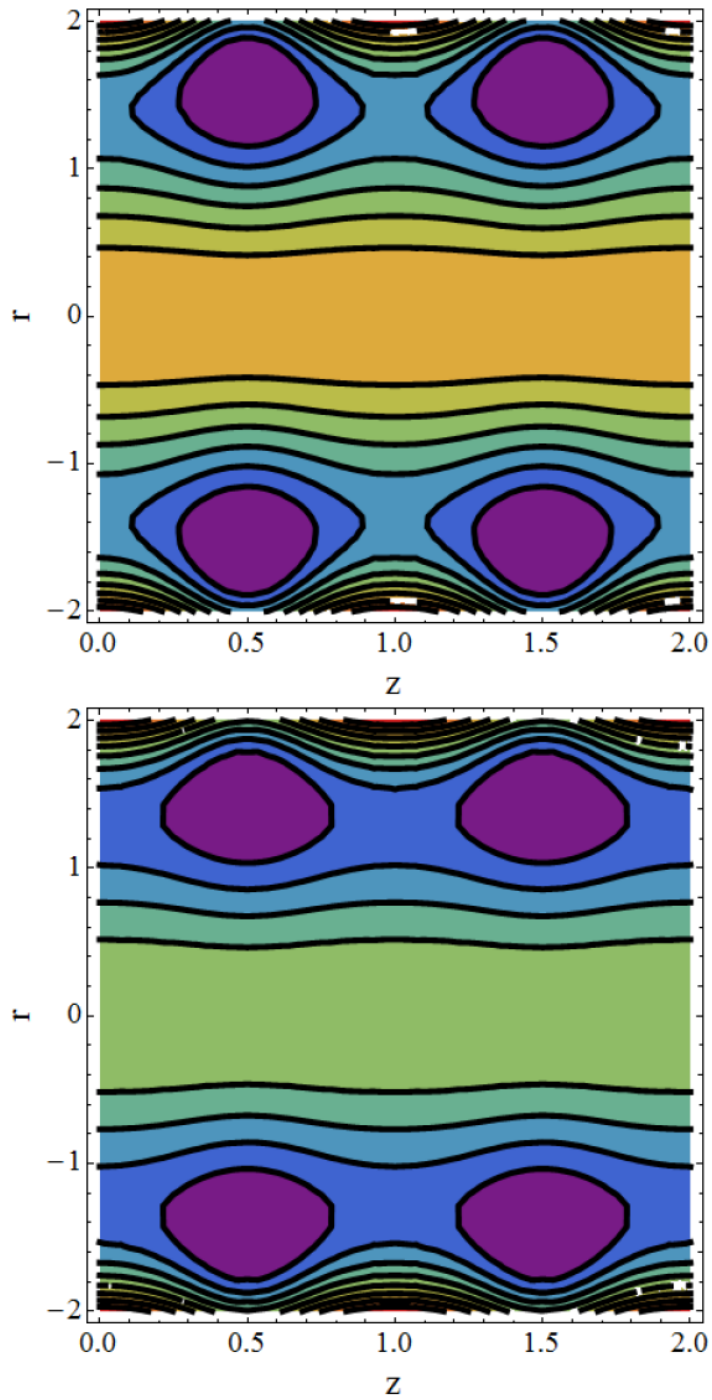


Fig. 5. Streamlines for the velocity profile for slip parameter slip parameter $\alpha_1 = 0.1, 0.11, 0.12, 0.15$ respectively.

The *parabolic* nature of the velocity profile is clearly evident from these figures. Fig. 2a shows that velocity is inversely proportional to slip parameter (α_1). Greater slip at the wall is found to counteract axial momentum development and manifests in a drop in axial velocity. Fig. 2b depicts the effects of Darcy number on velocity profile and it is observed that the velocity is directly proportional to permeability of the medium. Greater permeability corresponds to a significant decrease in Darcian impedance i.e. the drag force associated with solid fibers in the regime. Fig. 2c represents the influence of flow rate on velocity profile and it is apparent that if flow rate is increasing, velocity also increases. Propulsion is therefore enhanced with greater flow rates. Fig.2d reveals that velocity profile diminishes with increasing cilia length parameter (ϵ). Cilia spacing and length influences the viscous resistance per cilium and thereby the axial flow. The latter is assisted with greater cilia length and this aids in pressure rise in the lower channel half space. The introduction of extra energy to the flow at the lower wall however must be compensated for by an extraction at the upper wall, and these features are also related to synchronicity of beating cilia. The special case of $\epsilon = 0$ implies vanishing cilia and absence of a metachronal wave- in this scenario the flow is a purely peristaltic mechanism due to flexibility of the walls, and is therefore not considered here.

Figs. 3 illustrate the pressure rise evolution with flow rate for various parameters. A linear relation between flow rate and pressure is observed and there are three regions of pumping (i) pumping region ($\Delta P > 0$) (ii) free pumping ($\Delta P = 0$) and (iii) augmented pumping region ($\Delta P < 0$). The effect of slip parameter on pressure rise is shown in Fig.3a and it is evident that pressure diminishes with rise of slip parameter in the pumping region whereas the converse response is computed in the augmented pumping region. The impact of Darcy number on pressure rise is illustrated in fig.3b and exerts a similar effect as with the slip parameter. From fig 3c, it is seen that pressure rise is an increasing function of amplitude in the pumping region whereas it is decreasing function in augmented pumping region.

Figs. 4a-c depict the pressure gradient distributions with axial coordinate for various hydromechanical and geometric parameters. Inspection of the figures reveals that pressure gradient increasing with small changes in slip parameter and Darcy number (α_1, D_a) and also

cilia length parameter (ε). In all cases, the maxima (peaks) surface at $z = 0.5, 1.5$, however the relative minima change with modification in parameter values.

Figs 5a-d present the streamline visualizations for the influence of slip parameter. These streamlines constitute a family of curves which are tangential to the velocity vector at any instant of the flow. These show the direction of the fluid element at any point in time. The plots generally demonstrate that with increasing the magnitude of slip parameter (α_1), the trapped bolus inside the streamlines increases in size whereas the number of bolus decreases. In much of a digestive tract such as the human gastrointestinal tract, smooth muscle tissue contracts in sequence to produce a peristaltic wave, which propels a ball of food (called a bolus while in the esophagus and upper gastrointestinal tract and chyme in the stomach. So trapped bolus are enclosed streamlines. From the graphs we can see that near the tube walls more trapped bolus occurs because there is some fluid resistance and at center fluid moves easily and free stream occurs at the center of the tube. So that's why at the center of the tube no trapped bolus occurs all the bolus are near the walls of the tube.

Evidently in summary the geometric and hydrodynamic parameters result in non-trivial modifications in the propulsion characteristics and these warrant further exploration in the future.

5. CONCLUDING REMARKS

In this paper, a mathematical study has been conducted for creeping flow induced by a metachronal wave generated by beating of cilia tips in viscous propulsion through a porous medium-filled tube. The study has been motivated by further expounding biological propulsion mechanisms e.g. motion induced by cilia tips in uro-dynamics, breathing in the human respiratory system, sperm movement in reproductive system. Furthermore the study is pertinent to biomimetic propulsion mechanisms exploiting surface treatments via artificial cilia. The work it is envisaged will further stimulate much-needed laboratory investigations which would serve to garner vital experimental data. Although confined to viscous (Navier-Stokes) fluids and viscous-dominated porous media transport (Darcy model), some useful deductions can be made from the computations, namely:

1. Velocity is a decreasing function of slip parameter whereas it is an increasing function of Darcy number.

2. Greater flow rate elevates axial velocity whereas the latter decreases with increasing axial coordinate.
3. Pressure rise reduces with increasing the effects of slip velocity and permeability (higher Darcy number) in the pumping region whereas the reverse trend is observed in the augmented pumping region.
4. Axial coordinate and Darcy number exert the opposite effect on the pressure rise.
5. The size of trapped bolus of stream line increases with increasing the magnitude of slip parameter.

Acknowledgement:

Authors are thankful to the reviewer for useful suggestion to improve the manuscript.

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