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Hydrodynamic characterization within a spinner flask and a rotary wall vessel for stem cell culture

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HIGHLIGHTS

- CFD simulations of a spinner flask and a rotary wall vessel were performed.
- Increasing impeller speed reduced the Kolmogorov length scales.
- The Kolmogorov length scales were strongly correlated with cell aggregate sizes.
- The rotary wall vessel was designed to minimize shear stress distributions.
- The rotary vessel reduced shear stress distribution by half than the spinner flask.

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ABSTRACT

Stirred suspension culture is becoming a popular method for expanding human pluripotent stem cells (hPSCs). While stirring generates adequate fluid motions to lift the cells and facilitates mass transfers (of nutrients, dissolved gases, and metabolic wastes), excessive stirring could impose hydrodynamic forces deleterious for the growth of the cells. In this study, computational fluid dynamics (CFD) simulations were performed to first investigate hydrodynamic characteristics of fluid flows in a spinner flask, a common stirred suspension culture vessel used in laboratories. Flow patterns and distributions of shear stresses and the Kolmogorov length scales at varying impeller speeds were obtained. Comparison of the Kolmogorov length scales and sizes of hPSC aggregates, measured in the authors' previous experimental study, showed a strong correlation between the two. In addition to the spinner flask which generated complex and transient turbulent flows, this study investigated a newly developed rotary wall vessel that had been designed to produce laminar, circular Couette flows in order to control shear stress. CFD simulations revealed significantly more uniform and homogeneous flows compared to those in the spinner flask, suggesting that the rotary wall vessel is a suitable culture vessel to investigate roles of shear stress on hPSCs in suspension.

1. Introduction

Human pluripotent stem cells (hPSCs), including human embryonic stem cells (hESCs) and induced pluripotent stem cells (hiPSCs), have the potential to transform the future of cell therapies thanks to their ability to expand indefinitely and differentiate into any cell type [1–3]. However, significant biological and engineering challenges remain to be addressed before stem cell mediated therapies can be routinely employed to treat patients in a safe and economical manner. For

example, engineering a scalable culture system that can promote maintenance of pluripotency and effective differentiation uniformly in a controllable manner is necessary to meet the anticipated clinical demands for hPSCs and their derivatives [4,5].

hPSCs are routinely cultured as two-dimensional (2D), adherent monolayers in tissue culture dishes and flasks. However, this mode of culture is labor intensive and costly in reagents and materials. In addition, manual handling of the culture is susceptible to human errors, contamination, and batch-to-batch variations in both cell quality and

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Nomenclature**Symbols**

C_s	Smagorinsky constant
G	Filter function
P	Static pressure [Pa]
Re	Reynolds number
S_{ij}	Strain rate tensor [s^{-1}]
t	Time [s]
T	Period of revolution [s]
u	Fluid velocity [m/s]
x	Coordinate vector [m]

Greek Symbols

δ_{ij}	Kronecker delta
Δ	Filter width
ε	Turbulent dissipation rate [m^2/s^3]
η	Kolmogorov length scale [m]

μ	Fluid dynamic viscosity [$kg\ m^{-1}\ s^{-1}$]
μ_t	Subgrid scale turbulent viscosity [$kg\ m^{-1}\ s^{-1}$]
ρ	Fluid density [kg/m^3]
τ	Shear stress [Pa]
τ_{ij}^{SGS}	Subgrid scale stress tensor [Pa]
Ω	Impeller speed [rpm]
Ω_i	Inner wall speed [rpm]
Ω_o	Outer wall speed [rpm]

Abbreviations

CFD	Computational Fluid Dynamics
hESCs	Human embryonic stem cells
hiPSCs	Human induced pluripotent stem cells
hPSCs	Human pluripotent stem cells
LES	Large Eddy Simulation
RMS	Root mean square
rpm	Revolutions per minute
SGS	Subgrid-scale

quantity. These limitations have encouraged laboratories to investigate alternative modes of stem cell culture that are expected to be more amenable to scaling and automation [6–8].

A promising alternative to monolayer culture for hPSC expansion is suspension culture [5,9–14]. Multiple accounts have demonstrated that undifferentiated hPSCs can be expanded in stirred suspension culture

[6,15–17]. Expansion using such a platform, however, has multiple challenges such as variations of microenvironmental conditions within the culture volume and unclear links between culture parameters and stem cell fate [17]. In particular, spinner flasks, a common stirred suspension culture system, have large temporal and spatial distributions of shear stress, which can promote heterogenous cell aggregate

Fig. 1. Spinner flask. (a) Corning ProCulture spinner flask. (b, c, d) Horizontal and vertical cross-sections of the computational domain used in this study. Measurements are in mm, unless otherwise noted.

formation and apply nonhomogeneous forces that may promote spontaneous differentiation and compromise pluripotency [12]. There is also the risk of damaging cells from too high of shear stresses and turbulent eddies [13,18].

Current stirred suspension culture systems are empirically optimized for application-specific purposes without knowing the entire effect of physical forces on the cells. hPSCs can sense and respond to biophysical cues in their microenvironments [19,20]. However, the molecular mechanisms by which cells engage in mechanotransduction are not fully understood. Understanding how fluid shear stresses and other physical forces affect stem cell behavior such as protein expression and folding, contractile activity, proliferation, maintenance of pluripotency or differentiation would enable new designs of scalable stem cell culture systems that utilize fluid shear as a key input parameter. With such an ambitious goal, we recently reported our initial experimental study where we investigated effects of varying impeller speeds on undifferentiated hPSCs using Corning ProCulture spinner flasks [21]. Interestingly, varying impeller speeds (0, 40, 60, 80, and 100 rpm) resulted in significant differences in stem cell aggregate size, population growth, maintenance of pluripotency, spontaneous differentiation, and protein kinase expressions.

The primary objective of the current study is to characterize the complex fluid flows within the spinner flask by reproducing the experimental conditions (e.g., culture volume and impeller speeds) using powerful Computational Fluid Dynamics (CFD) simulations. CFD simulations have been previously employed to understand the hydrodynamic characteristics within bioreactors [13,14,22–24]. In particular, researchers from CSIRO Australia have successfully employed

Large Eddy Simulation (LES) to study the flow field within stirred suspension culture systems, including the Corning ProCulture spinner flask, and experimentally validated the simulation results using particle image velocimetry (PIV) [13,14]. Here we take a similar LES approach to simulate complex flow fields corresponding to our previous report [21] and compute key flow properties such as shear stress and the Kolmogorov length scale [25,26]. We compared the distribution of the Kolmogorov length scale to that of stem cell aggregate size which we measured in our previous study [21]. Finally, we performed CFD simulations of a newly developed custom rotary wall vessel, which features a pair of concentric rotating cylindrical walls designed to produce a laminar, circular Couette flow, resulting in a narrow and controlled distribution of shear stress.

2. Methods

This paper studied the hydrodynamics characteristics inside two different stem cell culture vessels. The first culture vessel chosen for this study was a Corning ProCulture spinner flask featuring a magnet-driven impeller and three baffles located on the vessel to promote mixing (Fig. 1). Although the standard working volume of this spinner flask is 125 mL, a 50 mL working volume was used to be consistent with our previous experimental study [21]. The second culture vessel was a rotary wall vessel designed to impose more uniform shear stress on the stem cells (Fig. 2). The inner radius, the outer radius, and the length are 30 mm, 37.5 mm and 94 mm, respectively, resulting in a working volume of 150 mL (Fig. 2). The commercial CFD solver, ANSYS CFX Version 18 (ANSYS Inc., Canonsburg, Pennsylvania, USA), was

Fig. 2. Rotary wall vessel. (a) The entire platform of the rotary wall vessel culture system. (b) Close view of the cylindrical vessel. (c, d) Cross sections of the vessel. Measurements are in mm.

Fig. 3. The computational grids for the spinner flask and the rotary wall vessel. (a) Unstructured grid used for the spinner flask, showing computational elements and inflation layers. Half of the entire domain (360°) is shown. (b) Structured grid used for the rotary wall vessel showing densely clustered elements near the walls. Half of the entire domain (full cylinder) is shown.

Fig. 4. Contour of instantaneous velocity magnitude on a vertical plane through the center of spinner flask for an impeller speed of $\Omega = 60$ rpm. The left side includes the velocity vectors to show different eddies in the flow.

Fig. 5. Contour of instantaneous shear stress on a vertical plane through the center of spinner flask for an impeller speed of $\Omega = 60$ rpm.

employed to solve governing equations (Section 2.3). The fluid inside both devices is considered Newtonian and incompressible.

2.1. Spinner flask

An unstructured grid was generated using ANSYS Meshing. The computational grid included 3.2 million elements (Fig. 3a). The computational domain was divided into a rotating region close to the impeller which was surrounded by a stationary region. Inflation layers were used close to the walls and impeller to capture the boundary layer. The interaction between the rotating and stationary regions was accounted for by using a transient rotor-stator technique (also known as a sliding mesh technique). The simulation was carried out for impeller speeds of 40, 60, 80, and 100 rpm. Due to the complex geometry of the

impeller and the baffles, the flows for these rotational speeds were considered turbulent. Therefore, LES was employed to capture the turbulent field. A high-resolution scheme, the second-order backward Euler scheme, and a second-order upwind scheme were used for the advection term, the temporal terms, and turbulence modeling, respectively. The Gauss's divergence theorem and finite-element shape functions were utilized to evaluate the control volume gradients. The residual target was that the normalized root mean square (RMS) errors for each conservation balance over the entire mesh fell to less than 10^{-5} . The time step was chosen in a way that the impeller rotated 1° each time step. LES was initialized with the results of steady-state $k-\omega$ (SST) turbulence modeling simulations. All simulations were run for ten revolutions, and only data for the last five revolutions were collected for the averaging process.

Fig. 6. Distribution of time-averaged shear stresses within spinner flask for impeller speeds of $\Omega = 40, 60, 80$ and 100 rpm.

Fig. 7. Boxplots of time-averaged shear stresses inside the spinner flask for impeller speeds of $\Omega = 40, 60, 80$ and 100 rpm.

2.2. Rotary wall vessel

Due to the geometrical simplicity of the rotary wall vessel, a structured grid was generated using ANSYS Meshing. The computational grid included 425,000 hexahedral elements (Fig. 3b). Depending on the simulation case, the inner or outer wall rotated with a rotational speed of 40, 60, 80, or 100 rpm. Because of the Reynolds numbers ($Re = \rho R_o^2 \Omega_0 / \mu \approx 6600$ and $16,500$ for 40 rpm and 100 rpm, respectively), the flow inside the rotary wall vessel was considered laminar [27]. The high-resolution scheme and the second-order backward Euler scheme were used for the advection term and the temporal terms, respectively. The Gauss's divergence theorem and finite-element shape functions

were utilized to evaluate the control volume gradients. The residual target was that the normalized RMS errors for each conservation balance over the entire mesh fell to less than 10^{-5} . The time step was set to 0.01 s. The simulations were initialized with a zero value for all velocity components. All simulations were run for ten revolutions and only data for the last five revolutions were collected for the averaging process.

2.3. Governing equations

As mentioned earlier, LES was utilized to capture the turbulence field inside the spinner flask. The rationale behind LES is to separate the large scale and the small scale in the flow. The larger scale turbulent motions are solved directly, and the influence of smaller scales are taken into account by appropriate subgrid-scale (SGS) models [28]. The flow variables (velocity vector u_i and static pressure p) are decomposed into supergrid and subgrid scale components as follows:

$$u_i = \langle u \rangle_i + u_i' \quad (1)$$

$$p = \langle p \rangle + p' \quad (2)$$

To obtain the supergrid components, the following filtering operation is used [29]:

$$\langle u \rangle_i = \int_D G(\mathbf{x} - \mathbf{x}') u_i(\mathbf{x}') d\mathbf{x}' \quad (3)$$

$$\langle p \rangle = \int_D G(\mathbf{x} - \mathbf{x}') p(\mathbf{x}') d\mathbf{x}' \quad (4)$$

where integration is over the entire flow domain D , and the specified filter function, G , determines the scale of resolved eddies and satisfies the normalized condition [29]:

$$\int_D G(\mathbf{x} - \mathbf{x}') d\mathbf{x}' = 1 \quad (5)$$

Fig. 8. Distribution of time-averaged Kolmogorov length scale within spinner flask for impeller speeds of $\Omega = 40, 60, 80$ and 100 rpm.

Fig. 9. Boxplots of time-averaged Kolmogorov length scale inside the spinner flask for impeller speeds of $\Omega = 40, 60, 80$ and 100 rpm.

The filtered governing equations for LES are obtained by filtering the continuity equation and the Navier-Stokes equations as follows [14,29]:

$$\frac{\partial \langle u \rangle_j}{\partial x_j} = 0 \quad (6)$$

$$\rho \left(\frac{\partial \langle u \rangle_i}{\partial t} + \frac{\partial \langle u \rangle_i \langle u \rangle_j}{\partial x_j} \right) = -\frac{\partial \langle p \rangle}{\partial x_i} + \frac{\partial}{\partial x_j} \left[\mu \left(\frac{\partial \langle u \rangle_i}{\partial x_j} + \frac{\partial \langle u \rangle_j}{\partial x_i} \right) \right] - \frac{\partial \tau_{ij}^{SGS}}{\partial x_j} \quad (7)$$

where ρ is the fluid density, μ is the fluid dynamic viscosity, and τ_{ij}^{SGS} is the SGS stress tensor. τ_{ij}^{SGS} is modeled using the Boussinesq hypothesis as [30]

$$\tau_{ij}^{SGS} - \frac{1}{3} \delta_{ij} \tau_{kk} = -\mu_t \left(\frac{\partial \langle u \rangle_i}{\partial x_j} + \frac{\partial \langle u \rangle_j}{\partial x_i} \right) \quad (8)$$

where μ_t is the subgrid scale turbulent viscosity. The Smagorinsky kernel is used to obtain μ_t [31]:

$$\mu_t = \rho (C_s \Delta)^2 \sqrt{2 \langle S \rangle_{ij} \langle S \rangle_{ij}} \quad (9)$$

where C_s is the Smagorinsky constant and $C_s = 0.1$ was used in this study [29]. $\Delta = (\Delta x \Delta y \Delta z)^{1/3}$ is the filter width, and $\langle S \rangle_{ij}$ is the supergrid strain rate tensor defined as

$$\langle S \rangle_{ij} = \frac{1}{2} \left(\frac{\partial \langle u \rangle_i}{\partial x_j} + \frac{\partial \langle u \rangle_j}{\partial x_i} \right) \quad (10)$$

Previous studies have shown that the effects of hydrostatic pressure on cells are negligible compared to the effects of shear stress [32,33]. Therefore, the current study focused on shear stress, which was calculated as follows [14,29]:

(a) H9-hESC

(b) RIV9-hiPSC

Fig. 10. Photomicrographs of day 7 cell aggregates grown under different impeller speeds in spinner flasks. (a) H9 human embryonic stem cells. (b) RIV9 human induced pluripotent stem cells. Scale bars: 250 μm .

Fig. 11. Comparison of cell aggregate size (black diamond) data [21] and the Kolmogorov length scale (red square) data from CFD simulation for different impeller speeds in spinner flask for H9-hESCs and RIV9-hiPSCs (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article).

$$\tau(\mathbf{x}, t) = (\mu + \mu_t) \sqrt{\langle S \rangle_{ij} \langle S \rangle_{ij}} \quad (11)$$

As mentioned in Sections 2.1, all simulations were run for ten revolutions and the data from the last five revolutions were used for the averaging purpose. Accordingly, the time-averaged shear stress over the five revolution was computed as

$$\bar{\tau}(\mathbf{x}) = \frac{1}{5T} \int_{5T}^{10T} \tau(\mathbf{x}, t) dt = \frac{1}{5T} \int_{5T}^{10T} (\mu + \mu_t) \sqrt{\langle S \rangle_{ij} \langle S \rangle_{ij}} \quad (12)$$

where T is the period of one impeller revolution.

3. Results and discussion

3.1. Spinner flask

Fig. 4 shows the magnitude of instantaneous velocity vector on a vertical plane through the center of spinner flask for an impeller speed of $\Omega = 60$ rpm. Several clockwise and counterclockwise vortices were

observed inside the flask. However, the number and location of these eddies varied with time. As expected, the velocity had the maximum value at the tip of the impeller. The flow below the impeller, however, has a lower velocity than the surrounding regions. Cells can become trapped in this region and grow into large cell aggregates. The spinner flasks have an indentation in the center bottom specifically to minimize this stagnation.

Fig. 5 shows the corresponding instantaneous shear stress on a vertical plane through the center of spinner flask for an impeller speed of $\Omega = 60$ rpm. Regions of high shear stresses were observed close to the impeller tip and the bottom of the flask. In these regions, the shear stress exceeds 100 mPa. High shear stress and small eddies in these regions could damage cells [34].

To get an overall picture of shear stress inside the spinner flask, the shear stress distributions across the computational domain averaged over 5 revolutions at different impeller speeds were plotted in Fig. 6. As expected, increasing the impeller speeds moved the distribution towards higher shear stresses. For instance, at 40 rpm, the shear stress

Fig. 12. Boxplots of time-averaged shear stresses inside the rotary wall vessel with rotation of the inner or outer wall.

mostly varied between 5–20 mPa, while for 100 rpm it varied mainly from 15–100 mPa. To get better insight, data were analyzed, and different descriptive statistics parameters such as average, standard deviation, median, Q_1 , and Q_3 were calculated (Fig. 7).

Fig. 7 shows the boxplots for time-averaged shear stresses inside the spinner flask for impeller speeds of $\Omega = 40, 60, 80$ and 100 rpm. These boxplots were plotted using the *matplotlib* library in Python. For better visualization, outlier data were not included in the boxplots. The outliers were neglected by setting *showfliers* parameter as *False* [35]. The outliers were defined as the data outside $[(Q_1 - 1.5 \text{ IQR}), (Q_3 + 1.5 \text{ IQR})]$ range, where Q_1 and Q_3 are the first and third quartiles and IQR is $Q_3 - Q_1$. The median shear stress increased linearly with the impeller speed. For instance, increasing impeller speed from 40 rpm to 100 rpm raised the median of shear stress from 11 mPa to 35 mPa. The mean shear stresses were 15, 25, 36 and 46 mPa for $\Omega = 40, 60, 80$ and 100 rpm, respectively. The standard deviations were 11, 19, 29, and 38 mPa for $\Omega = 40, 60, 80$ and 100 rpm, respectively.

The turbulent flow inside the spinner flask can be described using Kolmogorov's theory of isotropic turbulence, which suggests that the energy generated by impeller rotation is transmitted to the fluid by the generation of large eddies [36]. These large eddies split into smaller eddies due to inertial and friction forces. When the eddy size reaches a threshold (known as the Kolmogorov length scale), the viscous forces become predominant over the inertial forces [37]. The Kolmogorov

length scale, η is proportional to $(\mu^3/\rho^3\varepsilon)^{1/4}$, where ε is the turbulent dissipation rate [14]. In order to address the role of small scales of flow on cell growth, the Kolmogorov length scales were computed.

The distribution of Kolmogorov length scales inside the spinner flask is shown in Fig. 8. Increases in impeller speeds shifted the Kolmogorov length scale distribution toward smaller values. This shift in distribution was due to higher dissipation rate at higher impeller speeds. The Kolmogorov length scale ranged from 100 μm to $\sim 400 \mu\text{m}$ at 40 rpm while it ranged from 40 μm to $\sim 210 \mu\text{m}$ for 100 rpm.

The boxplots for the time-averaged Kolmogorov length scale inside the spinner flask for impeller speeds of $\Omega = 40, 60, 80$ and 100 rpm are shown in Fig. 9. The Kolmogorov length scale decreased with impeller speed. For example, increasing impeller speed from 40 rpm to 100 rpm decreased the mean Kolmogorov length scale from 264 μm to 149 μm .

The relationship between eddy size and cell aggregate size has been discussed in several studies [18,34]. Sen et al. calculated the minimum Kolmogorov length scale based on the maximum energy dissipation rate near the tip of impeller [34]. However, the shear stress inside the spinner flask is not uniform. The current study calculated the volumetric distribution of Kolmogorov length scale rather than its maximum value.

Fig. 10 shows photomicrographs of the day 7 aggregates of H9-hESCs and RIV9-hiPSCs at different impeller speeds in Corning ProCulture spinner flasks with a working volume of 50 mL [21]. These images show that the cell aggregate size decreased with increasing impeller speeds. Mean cell aggregate size for both H9-hESCs and RIV9-hiPSCs for different impeller speeds were compared with the Kolmogorov length scale in Fig. 11. Smaller eddies led to smaller mean cell aggregate size. Also, the standard deviation was shown to indicate how wide the Kolmogorov length scale and cell aggregate size varied throughout the flask. At lower impeller speeds, the cell aggregates were larger due to lower shear stresses, but some of the aggregates might be broken up near the tip of the impeller. Hence, the 40-rpm condition had the largest standard deviation among the tested impeller speeds. Fig. 11 also shows that the mean cell aggregate size for both H9-hESCs and RIV9-hiPSCs did not exceed the corresponding mean Kolmogorov length scale for any impeller speed. The distributions of the cell aggregate size reflect the trend observed in those of the Kolmogorov length scale. Although further investigation is needed, it is possible that the Kolmogorov length scale controlled the cell aggregate sizes.

As stated before, the above CFD simulations were conducted using a working volume of 50 mL, corresponding to our previous experimental study [21]. Additional CFD results with the standard working volume of the Corning ProCulture spinner flask (125 mL) are provided in the

Fig. 13. Streamlines and contour of instantaneous axial velocity within the space between the inner and outer walls. (a) The inner wall rotated at $\Omega_i = 60$ rpm, and the outer wall was stationary ($\Omega_o = 0$ rpm). (b) The inner wall was stationary ($\Omega_i = 0$ rpm), and the outer wall rotated at $\Omega_o = 60$ rpm.

Fig. 14. Distribution of time-averaged shear stresses within the rotary wall vessel for different outer wall speeds ($\Omega_o = 40, 60, 80$ and 100 rpm) with a stationary inner wall.

Fig. 15. Boxplots of time-averaged shear stresses inside the rotary wall vessel for different outer wall speeds ($\Omega_o = 40, 60, 80$ and 100 rpm).

supplementary information (Figs. S1–S4).

3.2. Rotary wall vessel

As shown in Section 3.1, the flow inside spinner flasks is transient and heterogeneous. Consequently, cells experience a wide range of shear stresses and Kolmogorov length scales. As discussed above, it is possible that the flow characteristics can impact cell expansion and differentiation. Therefore, having a uniform flow with homogeneous shear stress is expected to enhance the controllability and efficiency of both expansion and differentiation. In order to impose more uniform shear stress on stem cells, a rotary wall vessel that generates steady and reasonably uniform shear distribution was developed in collaboration with Synthecon, Inc. (Houston, TX) (Fig. 2).

Fig. 12 shows the boxplots of time-averaged shear stresses within the rotary wall vessel for two scenarios. In the first scenario, the inner wall rotated with a rotational speed of 60 rpm and the outer wall and sidewalls were stationary ($\Omega_i = 60$ rpm, $\Omega_o = 0$ rpm). In the second scenario, the outer wall and the sidewalls rotated at 60 rpm and the inner wall was kept stationary ($\Omega_i = 0$ rpm, $\Omega_o = 60$ rpm). The mean shear stresses for both cases were about 28 mPa, while the standard deviation was significantly different. Rotating the inner wall led to a wider range of variation in the shear stress. For instance, the standard deviation for the first scenario ($\Omega_i = 60$ rpm, $\Omega_o = 0$ rpm) was 29 mPa,

while this value for second scenario ($\Omega_i = 0$ rpm, $\Omega_o = 60$ rpm) was just 9 mPa.

In order to answer why the shear stress was remarkably different and understand the flow pattern differences between the two cases, the streamlines and contour of instantaneous axial velocity in the gap between the two cylinders were plotted in Fig. 13. In the first scenario (Fig. 13a), there were numerous regions of low and high axial velocity, with its value reaching 20 mm/s, while in the second scenario (Fig. 13b), the regions of low and high axial velocity formed near the sidewalls, with its value reaching only 2 mm/s. This major difference in the flow pattern was due to the direction of centrifugal forces. In the first scenario, the centrifugal force was larger than the local pressure gradient. Therefore, the flow was unstable, and several Taylor vortices [38] were created inside the space, but in the second scenario, the centrifugal force was less than the local pressure gradient and the flow was stable. According to these results, rotating the outer wall leads to a more uniform and homogeneous flow. Therefore, it is easier to control the shear stress and consequently cell expansion and differentiation in this environment.

Due to the flow stability observed in the second scenario (Fig. 13b), we decided to keep the inner wall stationary and rotate the outer wall with different rotational speeds. Figs. 14 and 15 show the histograms and the boxplots of shear stresses inside the rotary wall vessel for outer wall speeds of $\Omega_o = 40, 60, 80$ and 100 rpm, respectively. The mean shear stresses were 19, 27, 36 and 44 mPa for $\Omega_o = 40, 60, 80$ and 100 rpm, respectively. The standard deviation did not vary significantly with rotational speed.

Comparisons of shear stresses between the spinner flask (Figs. 6 and 7) and the rotary wall vessel (Figs. 14 and 15) showed that while the average shear stresses at certain rotating speeds were almost identical between the two vessels, the standard deviations were significantly different. For instance, the mean shear stresses at 60 rpm were around 25–27 mPa for both vessels but the standard deviation was 53 % less in the rotary wall vessel compared to the spinner flask (9 mPa vs. 19 mPa). Across the board, cell aggregates would experience a narrower range of shear stress in rotary wall vessel. Therefore, we believe the newly developed rotary wall vessel offers a better platform to investigate the roles of controlled mechanical forces, particularly shear stress, which is a key objective in our future study.

4. Conclusion

CFD simulations were carried out to characterize the hydrodynamics inside a spinner flask and a rotary wall vessel. The shear stress and Kolmogorov length scale distributions inside the cell culture vessels were examined for different operation speeds. The main conclusions are summarized as follows:

- Increasing the impeller speeds reduces the Kolmogorov length scale inside the spinner flask due to higher dissipation rates.
- The mean cell aggregate size for both H9-hESC and RIV9-hiPSC cells did not exceed the mean Kolmogorov length scale for any impeller speed. Although further investigation is needed, it is possible that the Kolmogorov length scale controls (but does not necessarily limit) the cell aggregate size.
- Rotating the inner cylinder creates Taylor vortices and secondary flow inside the rotary wall vessel which significantly increases the shear stress variations. This unwanted secondary flow can be minimized if the outer wall is rotated while the inner wall remains stationary.
- Cells would experience almost identical mean shear stresses inside both rotary wall vessel and spinner flask, but the shear stress variations would be much less in the rotary wall vessel compared to the spinner flask.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary data

Supplementary material related to this article can be found, in the online version, at doi:<https://doi.org/10.1016/j.bej.2020.107533>.

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Online Supplementary Material

Hydrodynamic characterization within a spinner flask and a rotary wall vessel for stem cell culture

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Figure S1: Contours of instantaneous velocity magnitude on a vertical plane through the center of spinner flask for impeller speeds of $\Omega = 40, 60, 80$ and 100 rpm. The left side of each panel includes the velocity vectors to show different eddies in the flow. Standard working volume of 125 mL.

Figure S2: Contours of instantaneous shear stress on a vertical plane through the center of spinner flask for impeller speeds of $\Omega = 40, 60, 80$ and 100 rpm. Standard working volume of 125 mL.

Figure S3: Distribution of time-averaged shear stresses within spinner flask for impeller speeds of $\Omega = 40, 60, 80$ and 100 rpm. Standard working volume of 125 mL.

Figure S4: Distribution of time-averaged Kolmogorov length scale within spinner flask for impeller speeds of $\Omega = 40, 60, 80$ and 100 rpm. Standard working volume of 125 mL.