In vitro analysis of multiple blood flow determinants using red blood cell dynamics under oscillatory flow
**In vitro** analysis of multiple blood flow determinants using red blood cell dynamics under oscillatory flow†

Ziya Isiksacana,b Murat Serhatlioglua and Caglar Elbuken†a,b

The flow behavior of blood is determined mainly by red blood cell (RBC) deformation and aggregation as well as blood viscoelasticity. These intricately interdependent parameters should be monitored by healthcare providers to understand all aspects of circulatory flow dynamics under numerous cases including cardiovascular and infectious diseases. Current medical instruments and microfluidic systems lack the ability to quantify these parameters all at once and in physiologically relevant flow conditions. This work presents a handheld platform and a measurement method for quantitative analysis of multiple of these parameters from 50 µl undiluted blood inside a miniaturized channel. The assay is based on an optical transmission analysis of collective RBC deformation and aggregation under near-infrared illumination during a 1 s damped oscillatory flow and at stasis, respectively. Measurements with blood of different hemo-rheological properties demonstrate that the presented approach holds a potential for initiating simultaneous and routine on-chip blood flow analysis even in resource-poor settings.

**Introduction**

Blood flow-related disorders are responsible for a quarter of all deaths across the world.1 Interdependent hemo-rheological parameters, comprising mainly deformation and aggregation of red blood cells (RBC) and blood viscoelasticity, are the determinants of blood flow in vascular networks.3,4 Acquired or hereditary diseases affecting one parameter cause alterations in others, such as hyperviscosity syndrome as a result of anemia originated in decrease in RBC deformation.5 Another example is that storage-induced lesions like RBC stiffness cause alterations in multiple parameters, paving the way for severe complications following blood transfusion.6,7 Presently, these parameters are rarely, if not at all, measured in clinics due to the drawbacks of using several benchtop instruments, requiring highly skilled and diversified workforce, lab space, high sample volume, operation time, and a large budget.8 An all-at-once measurement approach also remains elusive thus far in the microfluidics community. There are only a few studies analyzing two parameters at a time based on image processing.9,10 This manuscript addresses a novel point-of-care solution for the rapid analysis of multiple hemo-rheological parameters (RBC deformation, RBC aggregation, and blood/plasma viscoelasticity) from an undiluted blood sample without requiring cleanroom fabricated chips or microscopy setups.

Human RBCs are anucleated discoid cells of 6–8 µm diameter with a 10 nm viscoelastic membrane. They constitute around 2/3 of all body cells and 1/2 of blood volume, amounting to an RBC count of 20–30 trillion in the circulatory system of a healthy adult.11,12 Deformability of RBCs is essential for oxygen delivery through capillary networks. Numerous pathological changes in the RBC membrane, skeleton, and interior cause RBC stiffness, which potentially results in capillary blockage (malaria, diabetes, and sickle cell disease) and insufficient oxygen supply to tissues.13,14

To date, most RBC characterization methods focus on a single aspect of these cells at a low throughput, obscuring an understanding of the overall flow characteristics. Cell deformation is quantified by micropipette aspiration,15 atomic force microscopy,16 flow cytometry,17 and optical tweezers.18 There is also an extensive literature on microfluidic techniques for RBC deformability. The operating principles of platforms include cell blockage,19 transit (velocity, margination, and sorting),20 and impedance analysis21 for the characterization of malaria-infected or hardened RBCs. These platforms have not yet found a ground in clinics possibly due to costly chip fabrication, requirement for sample...
RBC aggregation is the formation of face-to-face RBC clumps at low shear rates and stasis. This mechanism is reversible and affected by cellular properties (surface charge and deformability), medium properties (viscosity and protein level), and flow properties (shear rate). The shear rate of blood flow in healthy circulation is above 10 s⁻¹ where RBCs do not form aggregates. However, clinical conditions slowing down blood flow and affecting medium/cell properties (like elevated fibrinogen concentration) cause aggregation that may lead to hyperviscosity or vascular occlusion. Such conditions include various viral infections, inflammations, multiple myeloma, and diabetes. Microfluidic platforms based on electrical conductivity, optical analysis, or microscopy imaging can further elucidate. More specifically, a complete analysis of RBC deformation and aggregation in an undiluted blood sample. The assay principle is based on the optical transmission of hemo-rheological parameters and help diagnose unhealthy blood flow. Herein, we report a portable device for the analysis of multiple parameters from 50 µl blood in 3 min. The assay principle is based on the optical transmission analysis of RBC deformation and aggregation in an undiluted blood sample. The test starts off by sample introduction to a rigid, straight channel of a disposable polycarbonate cartridge. This is followed by a 1 s reciprocating sample motion with gradually decreasing stroke distances, initiated by an integrated displacement pump. The sample is illuminated in the near-infrared band with a wavelength of 830 nm where hemoglobin absorption is minimum. Optical transmission through the sample is recorded during flow (14 s) and then at stasis (126 s) to optically quantify RBC deformation and aggregation, respectively. Medium viscosity, medium elasticity, RBC stiffness, and RBC aggregability are varied one at a time to obtain blood samples with different hemo-rheological properties, and the measured optical signals are shown to be responsive to corresponding alterations. The analysis is carried out in a channel environment that mimics vascular flow: confined, directional, pulsatile movement of blood at shear rates and RBC volume fractions comparable to physiological levels. Finally, the portability of the analyzer (200 g), use of small sample volume (50 µl), short assay time (3 min), and low-cost disposable cartridges (US$ 0.02) are noteworthy for point-of-care adaptation of the presented novel technique.

### Experimental section

#### Blood collection

Ethical approval was obtained from the Ethics Committee of Bilkent University (Ankara, Turkey), and all experiments were performed in compliance with the institutional guidelines. Informed consent was obtained from the human participant.

10 ml of fresh whole blood with 47% RBC volume fraction was acquired from a healthy male volunteer into a sodium citrate containing vacuum tube. The blood was centrifuged and washed three times with 1× phosphate-buffered saline (PBS), keeping the RBC volume fraction constant. Following blood withdrawal, the sample was kept at 25 °C and used within 6 h.

#### Suspending medium preparation

(See Table 1) Blood plasma has a relaxation time of 0.1 ms. Hence, a plasma replacement medium with a relaxation time of 0.1 ms was prepared by dissolving 0.0085% w/v polyethylene oxide polymer (PEO; $M_w = 600$ kDa, Sigma-Aldrich) in 1× PBS (ESI Note 1). Glycerol (Sigma-Aldrich) was mixed with the plasma replacement solution at three concentrations (5%, 10%, 20% v/v) to obtain three media for viscosity analysis. PEO was dissolved in 1× PBS at three concentrations (0.05%, 0.1%, 0.2%)

<table>
<thead>
<tr>
<th>Suspending Medium</th>
<th>Solvent</th>
<th>PEO in solvent (w/v)</th>
<th>$\eta$ [mPa s]</th>
<th>$\lambda$ [ms]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Medium 1</td>
<td>5% Glycerol &amp; 95% PBS</td>
<td>0.0085%</td>
<td>0.98</td>
<td>0.099</td>
</tr>
<tr>
<td>Medium 2</td>
<td>10% Glycerol &amp; 90% PBS</td>
<td>0.0085%</td>
<td>1.14</td>
<td>0.099</td>
</tr>
<tr>
<td>Medium 3</td>
<td>20% Glycerol &amp; 80% PBS</td>
<td>0.0085%</td>
<td>1.53</td>
<td>0.099</td>
</tr>
<tr>
<td>Medium 4</td>
<td>100% PBS</td>
<td>0.05%</td>
<td>1.40</td>
<td>0.312</td>
</tr>
<tr>
<td>Medium 5</td>
<td>100% PBS</td>
<td>0.1%</td>
<td>1.57</td>
<td>0.490</td>
</tr>
<tr>
<td>Medium 6</td>
<td>100% PBS</td>
<td>0.2%</td>
<td>2.41</td>
<td>0.768</td>
</tr>
</tbody>
</table>
0.2% w/v) to obtain three additional media for elasticity analysis.

**Blood sample preparation**

(See Table S2† for summary) Healthy RBCs have a shape recovery time of 0.15–0.30 s following deformation.13,34 Six different blood samples (H-samples 1 to 6; H for healthy) were prepared by suspending healthy RBCs (47% v/v) in media 1 to 6, respectively. To obtain RBCs with stiffened cell membranes, the healthy RBCs suspended in 1× PBS (47% v/v) were treated with 0.025% v/v glutaraldehyde (GA) solution, which was prepared in 1× PBS from a 25% w/v stock GA (Sigma-Aldrich). 0.025% v/v GA fixation increases RBC rigidity around 4 times.35 Then, six additional blood samples (S-samples 1 to 6; S for stiffened) were prepared by suspending stiffened RBCs (47% v/v) in media 1 to 6, respectively. 70 kDa Dextran (Sigma-Aldrich) was added to three blood samples (healthy RBCs suspended in 1× PBS, 47% v/v) at different concentrations to change RBC aggregability (A-samples 1 to 3; A for aggregated). Dextran concentrations for three suspending media were 10 mg ml$^{-1}$, 20 mg ml$^{-1}$, and 40 mg ml$^{-1}$, respectively.

**Rheological measurements**

A rotational rheometer (MCR 301, Anton Paar) was used for the viscosity ($\eta$) measurement of the six suspending media (Fig. S1†). The measuring unit has a 50 mm diameter cone plate geometry (CP50-1).

Constant shear rate mode was chosen, and the temperature was set to 25 °C. A 760 µl sample was used for each measurement. The shear rate was swept from 10 to 1000 s$^{-1}$. The relaxation times ($\lambda$) of the suspending media were calculated using the Zimm estimation (ESI Note 1†). All rheological values are given in Table 1.

**Pump characterization**

The pressure generated by the pump was characterized using a pressure sensor (40PC015 G, Honeywell). A silicone tube was connected to the pump, and the pressure was recorded and sampled using a data acquisition card (DAQPad-6015, National Instruments).

**Image analysis**

A 1 s video recording of the blood sample inside a channel was taken using an inverted optical microscope (Zeiss Axio Vert) equipped with a high-speed camera (Phantom Miro 2) at 2.5× magnification. The recording rate was 2800 frame per second. Images were obtained from the video using the camera software.

**Cartridge fabrication**

The cartridge had a single straight fluidic channel (depth ($d$), width ($w$) = 1 mm; length ($l$) = 50 mm) with one inlet and outlet. It was designed using CAD software (AutoCAD 2007). For initial prototyping and channel optimization, the cartridges were fabricated out of poly(methyl methacrylate) (PMMA) using CO2 laser ablation.36 Then, a metal mold was fabricated using CNC machining, and the transparent polycarbonate rigid cartridges were mass-fabricated by injection molding following industrial standards.

**Measurement platform**

Fig. 1 shows the schematic illustration and photograph of the portable device that consists of three main elements: integrated pump, optical detection unit (light-emitting diode and phototransistor), and microprocessor (also see Fig. S2† for workflow). Following a 50 µl sample introduction into the single channel of the cartridge and cartridge insertion into a 3D printed cartridge holder at room temperature, an airtight fluidic connection was made between the channel outlet and the pump. With the start of the test, the pump mechanically moves an actuator pin forward and backward with 0.5 s$^{-1}$ frequency for a duration of 14 s (ESI Movie 1†). Each forward pin movement generates 12.5 mbar pressure driving the sample forward inside the channel with 5 mm stroke distance and initiates a 1 s damped oscillatory flow (Fig. 2a). Meanwhile, an 830 nm wavelength near-infrared light-emitting diode (LED;
Vishay) illuminates the blood sample, and an NPN phototransistor (Vishay) collects the light transmitting through the sample in the form of an electrical current. The near-infrared band of the electromagnetic spectrum at which the LED works minimizes the photon absorption by hemoglobin (proteins confined in RBCs) and therefore enhances the transmission through the blood sample. RBC deformation during a flow reduces RBCs’ surface area to volume ratio giving photons more space to pass through the sample.\textsuperscript{37,38} Transmission intensity, that is, measured electrical current, is higher for RBCs deformed at shear flow compared to the disaggregated suspended RBCs at stasis. Likewise, individually suspended RBCs result in lower transmission intensity at stasis, and as they aggregate, the intensity level increases.\textsuperscript{25,39} Hence, we quantified (I) RBC deformation during flow and (II) RBC aggregation at stasis by measuring the transmission intensity (electrical current) using the phototransistor. The current output ($I_c$ [mA]) passes through an analog circuit consisting of a transimpedance amplifier, a lowpass filter, and an amplifying differentiator to obtain a voltage output ($V_{out}$ [mV]) such that
\begin{equation}
V_{out} = c_1 - c_2 dI_c/dt
\end{equation}
where $c_1$ and $c_2$ are constants. While the current output quantifies RBC deformation, the voltage output gives an understanding of the rate of RBC deformation at specific time intervals of optical measurement. The resulting analog voltage signal is digitized using the microprocessor and analyzed using MATLAB (R2018, MathWorks). Assay parameters such as LED intensity, test duration, and pressure can be tuned with a microprocessor (IOIO OTG, SparkFun).

**Results and discussion**

**Optical analysis of RBC movements in a fluidic channel**

During the test, the displacement pump moves the actuator pin forward and backward for 14 s. Fig. 2a depicts one cycle of...
the pressure generated by the pump. The pump action consists of a 1 s push state and a 1 s pull state (ESI Movie 1†). At the start of the cycle ($t_0$), the pin is in contact with an elastic membrane that makes an airtight connection to the channel outlet. In the push state, the pin pushes the membrane inward for 60 ms compressing the air within the system to generate an increasing pressure with a maximum value of 12.5 mbar at $t_1$. Following $t_1$, the pressure decreases and finally attenuates to zero at the end of 1 s ($t_4$). For the push state ($t_0$ to $t_4$), the initial pin movement ($t_0$ to $t_1$: $\Delta t_1 = 60$ ms) initiates a 1 s oscillatory flow of the sample within the channel (ESI Movie 2†). We are interested in the push state for the optical measurements. In the pull state, the pin withdraws, letting the membrane retract to its undeformed position.

To understand the flow dynamics, a microscope is used to obtain a side-view video recording of the blood sample inside the channel (not necessary for routine use of the system). Fig. 2b shows the side-view images of a 1 s oscillatory flow during a push state (ESI Movie 2†) at an observation window near the sample receding edge (also see Fig. S3†). The sample is stationary at $t_0$. Then, the pin pushes the adaptor for 60 ms, driving the sample forward, reaching a stroke distance of 5 mm at $t_1$ ($\Delta t_1 = 60$ ms). During this bulk motion, a small layer of fluid remains attached to the channel walls due to adhesion between the walls and fluid. Hence, although the pin stops at $t_1$, the bulk fluid moves backward for 1 mm until $t_2$ ($\Delta t_2 = 42$ ms) because of the cohesive forces between the fluids on the wetted walls and bulk. Again, the wetting on the advancing edge (not shown) causes a small forward flow until $t_3$ ($\Delta t_3 = 36$ ms). Finally, the fluid moves backward very slowly (mean velocity: 1.2 mm s$^{-1}$) for 860 ms until $t_4$. There are apparently four phases of sample flow. The sample has a different velocity in each phase. This results in different wall shear rates (phase-1: 480 s$^{-1}$; phase-2: 174 s$^{-1}$; phase-3: 66 s$^{-1}$; phase-4: 7.2 s$^{-1}$) that we approximately calculated using eqn (2), which is used for a rectangular channel profile ($d/w \leq 1$), where $\dot{\gamma}$ and $Q$ are the wall shear rate and flow rate, respectively.$^{40}$

$$\dot{\gamma} = 6Q/wd^2 \tag{2}$$

Microscopy images of the oscillatory flow in Fig. 2b reveal the differences in flow rates, and hence wall shear rates, for each phase. The shear rate determines the shear stress experienced by RBCs that, as a result, deform longitudinally to accommodate the stress. During the experiments, we measure physical changes in collective RBC deformation in the 1 s push states by illuminating the blood sample and collecting the transmitted light using the phototransistor. Fig. 2c illustrates an example of the current signal measured directly from the phototransistor during a 1 s oscillatory flow as well as the corresponding voltage signal obtained after analog processing (amplification and differentiation) of the current signal. As in Fig. 2b, the current signal consists of four distinct phases. The signal is concave-up with an increasing positive slope (phase-1), followed by a continuation of the rise with a decreasing slope (phase-2). After $t_2$ where the current is at maximum, the signal is concave-down with a decreasing negative slope (phase-3), and it goes back to the initial level (phase-4). The distinction between current levels in these phases, viz. RBC deformation in these phases, can be more clearly understood in the voltage signals that represent the rate of deformation. The voltage signal forms two peaks (AMP$_{1,2}$) at the end of phase-1 and phase-3, respectively. It has a zero crossing when the current is at maximum ($t_2$), and it settles down to the initial voltage level at the end of phase-4 ($t_4$).

The dense population of RBCs in the channel, because of the clinically relevant RBC volume fraction and the channel depth, hinders the observation of individual RBCs under a microscope. Based on the microscopy observation of the bulk sample shown in Fig. 2b and the corresponding optical transmission analyses in Fig. 2c, a 1 s four-phase oscillatory flow is schematically summarized in Fig. 2d. ($t = t_0$) RBCs are arbitrarily suspended in the medium. Phase 1: ($t_0 < t < t_1$) The sample moves forward. The RBCs align with the Poiseuille flow and deform longitudinally in response to increasing shear stress. The current level increases because the RBCs are deformed, allowing more light transmission. ($t = t_1$) Pump pressure is at a maximum. The RBCs have the largest rate of deformation. Hence, the voltage output forms a peak, AMP$_1$. Phase 2: ($t_1 < t < t_2$) The wetting effect on the receding edge moves the sample backward. The RBCs align with the flow and deform longitudinally. The current level increases due to RBC deformation. The shear stress generated by the wetting effect (0.15 Pa) is smaller than the stress generated by the pump (0.43 Pa). Therefore, deformation is smaller at $t_1 < t < t_2$ than at $t_0 < t < t_1$. ($t = t_2$) The RBCs have the highest backward deformation. The voltage output reaches the zero crossing. Phase 3: ($t_2 < t < t_3$) The wetting effect on the advancing edge moves the sample forward. The RBCs start regaining their original shape due to fading shear stress. The current level decreases when the deformation decreases and the second largest rate of deformation is attained at $t_3$ where the voltage output forms another peak, AMP$_2$. Phase 4: ($t_3 < t < t_4$) RBCs move backward very slowly until their complete stop and shape recovery at $t = t_4$. The current level gradually falls to the initial level.

### Change in medium properties

Viscoelastic plasma is a suspending medium for RBCs, and its rheology is a key determinant of overall blood flow. Specifically, medium viscoelasticity determines the generation of viscous and elastic stresses applied on RBCs, thereby determining RBC deformation during flow. Viscous stresses facilitate RBC deformation, whereas elastic stresses suppress it. Hence, an increase in medium viscosity or a decrease in medium elasticity enhances RBC deformation$^{41,42}$ and thus optical transmission. We prepared suspending media with different viscoelastic properties (Table 1) to confirm this. Firstly, three media were prepared using varying concentrations of glycerol during the plasma replacement (media 1, 2, and 3). Using these media, three blood samples (H-samples 1, 2, and 3) were prepared from healthy, deformable RBCs (47% v/v).
Fig. 3a shows the average optical signal corresponding to seven repetitive 1 s oscillatory flows (Fig. S4a†) for these samples. Fig. 3b compares the optical signal parameters (AMP1 and AMP2) with the glycerol concentration. AMP1,2 increase with the increasing medium viscosity, while t2 (Fig. S5a†) remains the same. As demonstrated in Fig. 2b, in phase-1, the pump generates an increasing pressure to drive the sample forward. In the meantime, shear stress (τ) applied on the RBCs deforms the cells. Given that the shear rate (γ) is a function of flow rate (Q) and is the same for all the samples during phase-1 (eqn (2)), an increase in medium viscosity (η) increases the shear stress as follows:

\[ \tau = \eta \dot{\gamma} = \eta \cdot 6Q/wd^2 \]  \hspace{1cm} (3)

The increasing shear stress in phase-1 accordingly deforms the RBCs more, which eventually increases light transmission and AMP1. Therefore, given the same elasticity, the more viscous the medium, the higher the rate of deformation and AMP1 value.

Pressure is the dominant factor in determining the flow rate in phase-1 and phase-2, given the short time scales of these phases (Δt1 = 60 ms, Δt2 = 40 ms). Hence, t2 only negligibly increases with the increasing medium viscosity.

The RBCs are in a deformed state at t2. During phase-3, they lose their deformation due to fading shear stress (Fig. 2b), and they partially recover their undeformed shape at t3. The more viscous the medium, the higher the stress change (eqn (3)). This consequently results in a higher rate of deformation leading to a higher intensity change. Hence, a higher AMP2 is measured in higher viscosity media.

Secondly, three media are prepared using varying concentrations of PEO in PBS (media 4, 5, and 6). Fig. 3c shows the average optical signal for seven repetitive 1 s oscillatory flows (Fig. S4b†) for the samples again with healthy RBCs (H-samples 4, 5, and 6). Fig. 3d compares the characteristic signal maxima and minima with the polymer concentration. AMP1 decreases with the increasing medium elasticity, while AMP2 increases. t2 also decreases with the increasing medium elasticity (Fig. S5b†). The relative importance of elastic and viscous forces of a viscoelastic medium can be represented by the Weissenberg number (Wi) as43

\[ Wi = \lambda \dot{\gamma} \]  \hspace{1cm} (4)

There are non-linear elastic forces in a viscoelastic fluid expressed in terms of normal stress differences, predominantly the first normal stress (N1) that increases with the polymer concentration:44

\[ N_1 = 2\lambda \eta \dot{\gamma}^2 \]  \hspace{1cm} (5)

N1 induced in the flow is at the normal direction to the RBC membrane,45,46 and the existence of N1 suppresses RBC deformation during phase-1. Therefore, AMP1 decreases with the increasing polymer concentration. t2 is different for each sample. The polymers are stretched by the stress generated in phase-1, and they store elastic stress, the magnitude of which is proportional to the polymer concentration. They discharge this energy and relax in phase-2 (viscoelastic recovery). More
elastomeric samples recover their initial state faster, explained by the creep-recovery response of the Kelvin model as:

\[ \epsilon(t) = Ce^{-E/\eta}t \]  

where \( \epsilon, E, \eta, \) and \( C \) are the strain, sample elasticity, sample viscosity, and initial-state constant, respectively (ESI Note 2†). Therefore, \( t_2 \) is the lowest at the highest polymer concentration, namely the highest \( E \).

The effective relaxation times for all the media are less than 1 ms (Table 1), so the polymers are already relaxed prior to phase-3. Therefore, the media act only in a viscous manner (Newtonian) in phase-3. As in the previous discussion for AMP2 in Fig. 3b, AMP2 increases with the increasing medium viscosity.

Having demonstrated the relationship between the optical parameters and the medium viscosity (Fig. 3b) or medium elasticity (Fig. 3d), governing equations relating individual optical parameters to overall medium viscoelasticity can be derived using the viscosity (\( \eta \)) and relaxation time (\( \lambda \)) values presented in Table 1. Fig. 4a compares \( k_1 = [\eta/\lambda] \) to optically measured AMP1, and Fig. 4b compares \( k_2 = [\eta/\lambda] \) to AMP2 (recalling that all media act only in a viscous manner in phase-3). Using the regression analysis, two linear equations are obtained as:

\[ k_1 = k_{1,0} + k_{1,1} \text{AMP1} \]  

\[ k_2 = k_{2,0} + k_{2,1} \text{AMP2} \]  

where \( k_{1,0}, k_{1,1}, k_{2,0}, \) and \( k_{2,1} \) are constants with the values of 93.1, -7.8, 6.3, and 0.7, respectively. This analysis demonstrates that using these equations and the parameters obtained from an optical measurement, medium properties can be calculated as an alternative to the conventional rotational rheometer. In addition, since AMP1 and AMP2 are measures of RBC deformation, if the medium properties are already known, these parameters can be employed to obtain the deformation during flow.

**Change in RBC membrane stiffness**

Healthy RBCs are deformable due to the viscoelastic membranes that enclose the cellular interior. Numerous clinical conditions such as malaria can decrease RBC deformability. We prepared six blood samples (S-samples 1-to-6) by suspending glutaraldehyde-fixed stiffened RBCs inside the six suspending media. Fig. 5 shows the optical measurements for these samples. The trends seen for the H-samples in Fig. 3b&d are also observed for the S-samples in Fig. 5b&d for the same reasons. Briefly, given the same elasticity, AMP1,2 increase with the increasing medium viscosity (Fig. 5b) while \( t_2 \) remains the same (Fig. S5c†). In addition, AMP1,2 decrease with the increasing medium elasticity (Fig. 5d) while \( t_2 \) increases (Fig. S5d†). We also observe that due to the compromised deformability of the stiffened RBCs, AMP1 and AMP2 are lower for the S-samples compared to those values in the H-samples (Fig. S6†).

**RBC aggregation analysis at stasis**

RBC aggregation is the main factor determining blood flow at low shear rates. This microfluidic device can also quantify the RBC aggregation of blood. RBCs form aggregates at stasis in a suspending medium containing inducer molecules like dextran, which is a highly water-soluble polysaccharide. The aggregation can be enhanced with increasing dextran concentration, depending on the molecular weight and concentration interval.

Four media were prepared using varying dextran concentrations (0, 10, 20, and 40 mg ml\(^{-1}\)) in PBS in which RBC aggregation and solution viscosity\(^{48}\) increase linearly with the dextran concentration whereas solution elasticity is negligibly affected.\(^{49,50}\) Then, RBCs (47% v/v) were suspended in these media. Fig. 6 shows the corresponding optical transmission measurements, \( V_{out} \). Initially, the pump operates for 14 s, moving the sample back and forth. The generated pressure creates a disaggregating shear rate (phase-1: 480 s\(^{-1}\)), which leads to complete disaggregation of the RBCs. At the time the flow ceases, individual RBCs are dis-

![Fig. 4](image-url)  
**Fig. 4** Comparing optical signal amplitudes to medium properties for the six blood samples with healthy RBCs. (a) The scatter plot and the linear fit comparing AMP1 to \( k_1 = [\eta/\lambda] \). The correlation is 88% using linear regression. (b) The scatter plot and the linear fit comparing AMP2 to \( k_2 = [\eta/\lambda] \). The correlation is 98% using linear regression.
aggregated and dispersed within the channel, resulting in maximum scattering and minimum light transmission. Following the complete disaggregation, the RBCs start to aggregate in the absence of a shear rate until their complete aggregation at the end of 140th s. The maximum final transmission levels are obtained at the completion of the aggregation phase. The relative transmission level at the 60th s is taken as a measure of RBC aggregation, namely the aggregation index (AI) where AI = \[ V_{60th}^{out} - V_{14th}^{out} \]. Fig. 6 inset compares AIs for different samples. The correlation between AI and dextran concentration is calculated as 97% using linear regression, confirming the enhancement of aggregation with dextran addition and the ability of our device to quantify it.

Although this manuscript reports the first proof-of-concept study of measuring multiple blood flow determinants using an optical measurement principle on an entirely portable device, it has several limitations that should be addressed in upcoming studies. Firstly, we herein focused on altering one property of the blood sample at a time and observed the resulting changes in the optical signals. For future studies, more than one blood property needs to be swept simultaneously to investigate whether multiple hemorheological parameters can be extracted from the signals. After multiple parameters are quantified from a single measurement, the next step is performing blind tests using clinical samples. Success in clinical trials depends substantially on the availability of conventional equipment (flow cytometer, rheometer, and aggregometer) for benchmarking as well as correlating the blood flow abnormalities to disease states. In addition, a built-in temperature controller unit is needed to keep blood samples at body temperature throughout the test.
Conclusion

We have presented a portable lab-on-a-chip device and a novel measurement method for the determination of multiple hemo-rheological parameters, one at a time, in disposable cartridges from 50 µl undiluted blood. The operation principle is based on a 3 min optical transmission analysis of collective RBC movements (deformation and aggregation) in a straight single channel under infrared illumination. A unique four-phase damped oscillatory flow mechanism was generated jointly by a displacement pump and wetting effect. The analyses were performed in a miniaturized channel environment that mimicked vascular flow. Optical transmission through the blood sample was recorded during the flow and at stasis to quantify RBC deformation and aggregation, respectively. Blood samples with different medium viscoelasticity and RBC stiffness were employed to experimentally demonstrate that (I) viscous stresses facilitate RBC deformation and (II) elastic stresses suppress it. This presented technique is promising to remove the financial and practical drawbacks associated with the state-of-the-art hemo-rheological analysis methods. Such a sample-in result-out microfluidic approach can eventually make hemo-rheological monitoring accessible in clinics and research labs.

Author contributions

Z. I and C. E conceived and designed the experiments. Z. I. and M. S. performed the experiments. All authors discussed the results. Z. I. prepared the figures. Z. I. and C. E wrote the paper.

Conflicts of interest

There are no conflicts to declare.

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