

# Validation of a low field Rheo-NMR instrument and application to shear-induced migration of suspended non-colloidal particles in Couette flow

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## ABSTRACT

Nuclear magnetic resonance rheology (Rheo-NMR) is a valuable tool for studying the transport of suspended non-colloidal particles, important in many commercial processes. The Rheo-NMR imaging technique directly and quantitatively measures fluid displacement as a function of radial position. However, the high field magnets typically used in these experiments are unsuitable for the industrial environment and significantly hinder the measurement of shear stress. We introduce a low field Rheo-NMR instrument (<sup>1</sup>H resonance frequency of 10.7 MHz), which is portable and suitable as a process monitoring tool. This system is applied to the measurement of steady-state velocity profiles of a Newtonian carrier fluid suspending neutrally-buoyant non-colloidal particles at a range of concentrations. The large particle size (diameter > 200 μm) in the system studied requires a wide-gap Couette geometry and the local rheology was expected to be controlled by shear-induced particle migration. The low-field results are validated against high field Rheo-NMR measurements of consistent samples at matched shear rates. Additionally, it is demonstrated that existing models for particle migration fail to adequately describe the solid volume fractions measured in these systems, highlighting the need for improvement. The low field implementation of Rheo-NMR is complementary to shear stress rheology, such that the two techniques could be combined in a single instrument.

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## 1. Introduction

Transport of suspended non-colloidal solid particles is an important process in many industries, including oil and gas production [1] where the controlled placement of a cement slurry [2] and conveyance of solids by drilling fluids [3] are critical for safe and successful operations. Determining the hydrodynamic properties of non-colloidal large particle (diameter > 200 μm) suspensions is a challenge. These complex fluids are not amenable to measurement in narrow-gap rheometric devices due to particle jamming, and violate the narrow-gap approximation [4] in wide-gap geometries such as the cylindrical Taylor-Couette cell [5], where additionally the local rheology may be a function of radius. Nuclear magnetic resonance rheology (Rheo-NMR) measures the spatial variation in velocity and concentration of a sample under shear [6,7] and gives direct access to local shear rate  $\dot{\gamma}$  in the wide-gap Couette geometry through the relation

$$\dot{\gamma}(r) = r \frac{\partial}{\partial r} \left( \frac{v}{r} \right), \quad (1)$$

where  $v$  is the tangential velocity at radius  $r$  [8]. In the case of liquids containing suspended non-colloidal particles, the sample evolves over time due to shear-induced radial migration of particles, plus centrifugal radial migration and gravity-driven vertical migration depending on the density mismatch between the solid and liquid components [9–13]. The ability of Rheo-NMR to probe inhomogeneous and opaque liquid systems makes it ideal as a tool for routine process monitoring and quality control in the industrial environment. A commercial Rheo-NMR device is available, designed to fit in the vertical bore of a high field superconducting magnet [14]. However, the installation of high field magnets in a factory or remote location is impractical due to the limitations of cost, safety (strong magnetic fields), and maintenance (requirement for cryogenics). Low field permanent magnets, or recently commercialized cryogen-free superconducting magnets, offer lower cost, safer alternatives. Permanent magnet systems also offer the advantage of portability [15]. Low field magnets have been used previously to provide relaxation time measurements of liquid samples in conjunction with shear rheology measurements [16–18]. The usual

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Rheo-NMR experiment lacks a direct measure of shear stress although custom systems have been proposed that combine NMR and torque sensors [19,20], albeit still based on superconducting magnets with limited industrial application. Rheo-NMR has been applied to the study of many complex samples, including shear-banding materials [21–23], polymer solutions [24,25], worm-like micelles [26,27], liquid crystalline materials [19,28–31], and colloidal particle suspensions [32,33]. Shear-induced migration of non-colloidal particles has been studied by NMR imaging in Couette [34–36] and pipe flow [37,38].

The migration of particles in neutrally-buoyant suspensions in non-uniform shear flows has been described with a variety of models. Diffusive-type models describe the migration process directly as a diffusive process, with particle fluxes directed by gradients in shear rate and particle concentration [39,40]. Models of the suspension-balance type describe migration as resulting from gradients in normal stress of the suspension [41,42]. In recent years, researchers have been successful in modeling the behavior of dense suspensions using frictional rheology models [43] similar to those previously used for dry granular flow [44]. These can be shown to be equivalent in some cases to suspension balance models [45], but have been generalized successfully to describe additional physics such as suspensions in yield-stress fluids [46] and compaction beyond the jamming limit [45,38].

In this paper we use a low field permanent magnet imaging system to measure velocity and concentration profiles in a wide-gap Couette cell. The hardware and experimental procedures are described in Section 2. Measurements of large non-colloidal particle suspensions in a Newtonian carrier fluid are demonstrated, and the results compared to high field Rheo-NMR acquisitions at matched shear rates in Section 3. For illustration we compare our experimental data to the model of Morris and Boulay [42]. An assumption of this model, as with the other models described above, is that the suspension is locally Newtonian. However, disregarding issues of particle migration, many suspensions exhibit non-Newtonian behavior even at moderate volume fractions and where inertial and Brownian effects are negligible [47]. We observe such effects in our system and subsequently a departure from the predictions of the suspension-balance model.

## 2. Experimental

### 2.1. Low field Rheo-NMR

The low field NMR system comprised a  $B_0 = 0.25$  T permanent magnet (Laplacian, UK) of SmCo planar disc poles enclosed in a rectangular iron yoke of dimensions  $52 \times 55 \times 48$  cm and approximate mass 150 kg. The field strength corresponded to a resonance frequency of 10.7 MHz for  $^1\text{H}$ . Thermal stability of the magnet was maintained at  $28^\circ\text{C}$  by active heating and cooling. The vertical bore rf resonator had an inner diameter of 53 mm and active damping feedback pre-amplifier (MRF Innovations, UK) for improved response time and signal-to-noise ratio (SNR). The rf power was supplied by a 500 W CW amplifier (Tomco, Australia). Magnet homogeneity was improved by manual adjustment of an 8 channel shim current supply unit (Resonance Research Inc., USA; model MXD-8). Planar gradient coils driven by high-performance audio amplifiers (AE Techtron, USA; model 7224) provided a maximum gradient strength of  $g_{\text{max}} = 0.125$  T m $^{-1}$  ( $x$  and  $y$  axes) and  $g_{\text{max}} = 0.14$  T m $^{-1}$  ( $z$  axis). The water cooled gradient plates maintained the sample temperature at  $25^\circ\text{C}$ . The NMR Experiments were controlled by a DRX-TCP spectrometer (Oxford Instruments, UK) with custom pulse sequences and digital filters [48] optimized for imaging.

Spatially resolved concentration and velocity profiles were obtained using a standard spin-echo imaging sequence [49]. Gaussian rf pulses of duration  $t_p = 1024$   $\mu\text{s}$  were used to provide a vertical slice of 30 mm full-width half-maximum (FWHM) and a horizontal slice of 5 mm FWHM; the rf power was adjusted to provide  $\pi/2$  and  $\pi$  pulses as required. For the velocity encoding, three gradient increments were used ( $-g_v, 0, +g_v$ ) to provide a robust calculation of phase shift. The gradient amplitude was scaled according to the Couette rotation rate to provide the same maximum phase shift in all experiments. The velocity-encoding gradient pulses, placed symmetrically about the refocusing rf pulse, had duration  $\delta = 1.5$  ms and the observation time was  $\Delta = 11.0$  ms. The frequency-encoded read gradient provided a field of view FOV = 59 mm over 256 pixels, such that the spatial resolution was  $\Delta x = 229$   $\mu\text{m}$  with 56 pixels across the Couette gap. The measured profiles extended over the full diameter of the Couette cell, providing antisymmetric velocity components on either side of the rotating center body. The magnitudes of the  $y$  velocities were averaged at the absolute  $|x|$  coordinates (taking  $x = 0$  to be the center of the Couette geometry), to provide  $|\overline{v}_y|$  as a function of radius. A recycle time of 500 ms was used between scans, and 256 repeat scans were summed to improve the SNR, resulting in an acquisition time of approximately 2 min per gradient increment.

A custom Rheo-NMR Couette device was constructed for the low field magnet. The outer cylindrical body (stator) was gun-drilled from a rod of polyamide-imide (Torlon; Solvay, Belgium) to provide a high-precision machined part. The inner cylindrical body (rotor, or “bob”) was machined from polyether ether ketone (PEEK). The Couette geometry had an outer radius of  $r_o = 20.3$  mm and an inner radius of  $r_i = 7.5$  mm, such that the gap was  $r_o - r_i = 12.8$  mm with a gap ratio of  $(r_o - r_i)/r_o = 0.63$ . The total bob length was 252 mm. Rotation of the bob was achieved using a 12 V d.c. motor with the speed determined by manual adjustment of the voltage supply; an optical encoder in the motor was used to confirm the rotation rate. Rotation rates ranged from  $\Omega = 3$ –40 rpm; the rotation rate was limited to a maximum of  $\Omega = 15$  rpm when measuring Newtonian liquids due to the formation of Taylor vortexes at higher shear rates.

### 2.2. High field Rheo-NMR

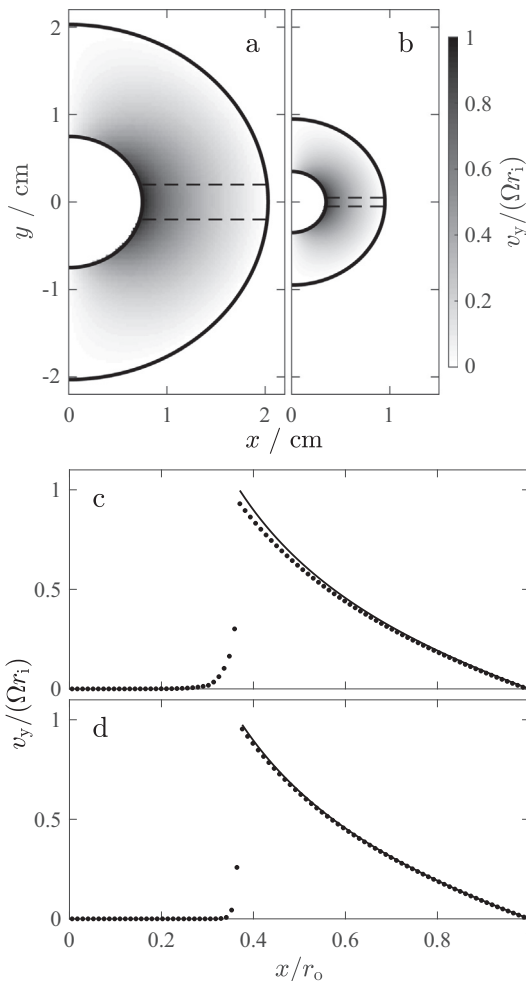
A  $B_0 = 7$  T vertical bore superconducting magnet, corresponding to 300 MHz for  $^1\text{H}$ , was used for the high field Rheo-NMR studies. The imaging experiments were controlled by a Bruker DMX300 spectrometer. A bird-cage rf coil of inner diameter 25 mm was employed. Imaging gradients up to  $g_{\text{max}} = 0.987$  T m $^{-1}$  were available on all three axes. The sample temperature was maintained at  $20^\circ\text{C}$ . A motion-compensated imaging sequence was used with Gaussian rf pulses of duration  $t_p = 512$   $\mu\text{s}$ , providing a vertical slice of 4 mm FWHM and a horizontal slice of 1 mm FWHM. A reliable phase shift measurement was achieved with just two gradient increments (0 and  $+g_v$ ) due to the SNR and stability of the high-field instrument. The velocity-encoding gradient pulses had duration  $\delta = 1.5$  ms and the observation time was  $\Delta = 11.8$  ms. The frequency-encoded read gradient provided a FOV = 28 mm over 256 pixels, such that the spatial resolution was  $\Delta x = 113$   $\mu\text{m}$ , with 53 pixels across the Couette gap. As with the low field data,  $|\overline{v}_y|$  was obtained as a function of radius from the antisymmetric velocity components on either side of the rotating center body. A recycle delay of 6 s was used and 4 repeat scans were summed, with an acquisition time of less than 1 min per profile.

A commercial Rheo-NMR device was used (Magritek, New Zealand and Bruker BioSpin, Germany), designed to fit the vertical bore of the high field magnet. Full details of this device are given in [8]. The radius of the stator  $r_o = 9.5$  mm and the radius of the rotor

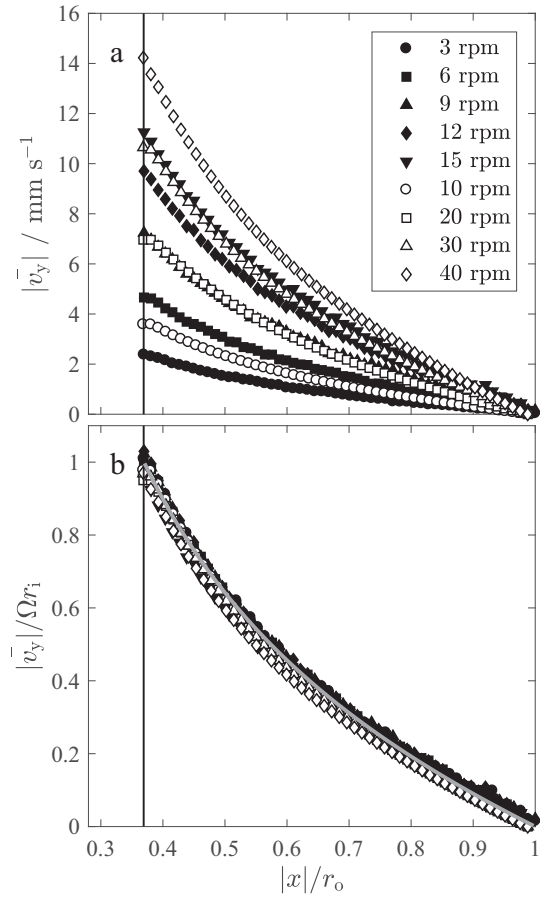
(bob) was  $r_i = 3.5$  mm, providing a Couette gap of  $r_o - r_i = 6.0$  mm and a gap ratio of  $(r_o - r_i)/r_o = 0.63$  (consistent with the custom low field Couette geometry). Rotation rates ranged from  $\Omega = 10$ –40 rpm. Inclusion of a Newtonian marker liquid (water) inside the bob allowed wall slip to be characterized in the high field experiments.

### 2.3. Couette geometry correction

The gap ratio of the two Couette geometries was matched to provide consistent shear rates between the low and high field experiments. In the ideal Rheo-NMR experiment, a thin slice would be selected in the vertical plane so as to be insensitive to the curvature of the inner and outer walls of the Couette cell. Practically, this selected slice must be of finite width and due to the inherently poor SNR available on the low field system it was necessary to select a thick horizontal slice, as detailed in Section 2.1. The relative width of the selected slice compared to the bob diameter is illustrated in Fig. 1a and b for the low and high field Couette



**Fig. 1.** Comparison of Couette geometries in the low and high field magnets. Normalized y-component velocity maps are shown for the (a) low field and (b) high field Rheo-NMR geometries; half of the cell is shown in each case where the tangential velocity component is positive (see gray scale). The velocity maps are calculated using Eq. (2), valid for a Newtonian liquid. The inner and outer radii of the Couette cells are denoted by the solid black lines. The dashed black lines indicate the FWHM of the y-axis Gaussian slice selection. Simulated tangential velocity profiles, calculated by taking a weighted average of velocities over the Gaussian slices, are given by the dotted lines in (c and d) for the low and high field geometries, respectively. Ideal tangential velocity profiles are given by the solid lines in (c and d).



**Fig. 2.** Radial velocity profiles for a Newtonian mixture of water and 2-propanol as (a) absolute velocity and (b) normalized velocity. The filled symbols were obtained at low field, the open symbols at high field. The legend indicates the rotation rate of the inner wall at  $r_i$  (position shown by the solid vertical lines) and applies to both plots. In (b) the theoretical normalized velocity profile from Eq. (2) is shown by the solid black curve.

geometries, respectively. Theoretical y-component velocity maps are overlaid on the Couette cell schematics where

$$\frac{v_y(x, y)}{\Omega r_i} = \frac{x r_i}{r_o^2 - r_i^2} \left( \frac{r_o^2}{x^2 + y^2} - 1 \right), \quad (2)$$

generalized for a Newtonian liquid. The ideal velocity profiles  $v_y(x, 0)/(\Omega r_i)$  are the solid lines in Fig. 1c and d and are independent of Couette geometry due to the matched  $r_i/r_o$  ratios. The velocity profiles we expect to measure will differ from these ideal lines due to averaging over the Gaussian-weighted slices. The expected velocity profiles are simulated here by averaging  $v_y(x, y)$  over the selected (slice) region; these simulated velocity profiles are given by the dotted lines in Fig. 1c and d. Comparing the ideal and simulated velocity profiles we see that a significant underestimate of the velocity near the inner wall will be observed in the low field Couette geometry. An equivalent effect occurs at the outer wall but is considered unimportant due to the low velocity in this region. For the high field Couette geometry, a very small difference is observed between the ideal and simulated velocity profiles. A correction based on the simulations, plus an additional term to account for curvature of the streamlines [51], was applied to all measured velocity profiles.

## 2.4. Materials

A Newtonian mixture of water and 2-propanol was prepared in the ratio 2.65:1 by weight to provide a density of  $\rho_1 = 0.96 \text{ g cm}^{-3}$ . The water was doped with approximately  $1.5 \text{ g dm}^{-3}$   $\text{NiCl}_2$  to provide a typical longitudinal relaxation time of 300 ms, permitting a short recycle delay between repeat scans. The nickel salt was chosen for its lower toxicity compared to other standard doping agents. The carrier fluid density was matched to the density of polyethylene microspheres (Cospheric LLC, USA). These particles were selected for being highly spherical with a narrow diameter distribution in the range 212–250  $\mu\text{m}$  and were sufficiently large that Brownian motion was neglected. Additionally, we estimated a maximum Reynolds number in our experiments at 40 rpm of  $Re \approx 0.1$ , indicating that inertial effects were small (but not necessarily negligible). Particles were added to the liquid to provide solids volume fractions of  $\phi = 0.2, 0.35$ , and  $0.5$  by volume. Precise density matching was achieved by careful adjustment of the water-2-propanol ratio in the carrier fluid at the experimental temperature. During the experiments the Couette cells were sealed to limit evaporation of 2-propanol. Under shear, the particle suspensions were allowed up to 30 min to reach equilibrium before measurements commenced.

## 3. Results and discussion

### 3.1. Low field validation

Measured velocity profiles obtained for a Newtonian mixture of water and 2-propanol are shown in Fig. 2(a). The rotation rate of the bob was restricted in the low field instrument due to the formation of Taylor vortices above approximately  $\Omega = 15 \text{ rpm}$ . All the velocity profiles, whether acquired on the low or high field instrument, showed a consistent decrease from the rotating inner wall to zero velocity at the stationary outer wall, as expected. It should be noted that higher tangential velocities are apparent in the low-field instrument due to the larger bob radius. When the

tangential velocity component is normalized by the rotation speed of the bob, see Fig. 2(b), all the data collapse to a single smooth line (within experimental uncertainty). These normalized velocity profiles agree with the theoretical prediction of velocity from Eq. (2) and demonstrate consistency between the low and high field instruments for a simple Newtonian liquid.

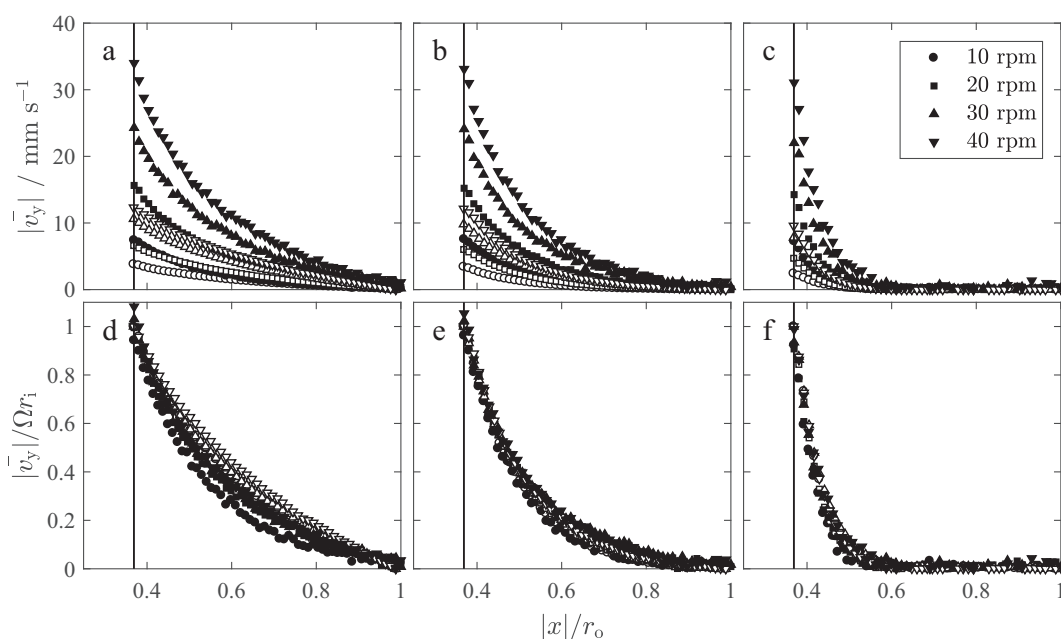
Measured velocity profiles for the polyethylene particles in density-matched liquid are shown in Fig. 3. The bob rotation rates varied between  $\Omega = 10$  and 40 rpm, as indicated by the figure legend. The difference in absolute velocity notwithstanding, the low and high field velocity profiles again collapse to single smooth curves once normalized by the inner wall velocity, see Fig. 3d–f. The shape of the velocity profile is independent of rotation rate at steady-state but depends on solids volume fraction (SVF). At  $\phi = 0.2$ , the  $y$ -velocity component decreases with increasing radius, to zero at the stationary outer wall, and is similar to the behavior of a Newtonian liquid. At  $\phi = 0.35$  the velocity profile is still non-zero but with a higher curvature. At  $\phi = 0.5$  the velocity decreases to zero at a critical radius  $r_c/r_o \approx 0.65 < 1$ , Fig. 3(f). We interpret this region of **near-constant  $\phi(|x| > r_c) \approx 0.52$**  as jamming where the SVF matches or exceeds its maximum packing fraction, i.e.,  $\phi \geq \phi_m$ , as was found in [10]. This is consistent with the fact that the location of the critical radius does not depend on the rotational speed.

Overall, we consider the consistency of the velocity and particle concentration profiles acquired at low and high field to be indicative of their accuracy, and this work demonstrates the feasibility of using low field magnets as quantitative Rheo-NMR instruments.

### 3.2. Model comparison

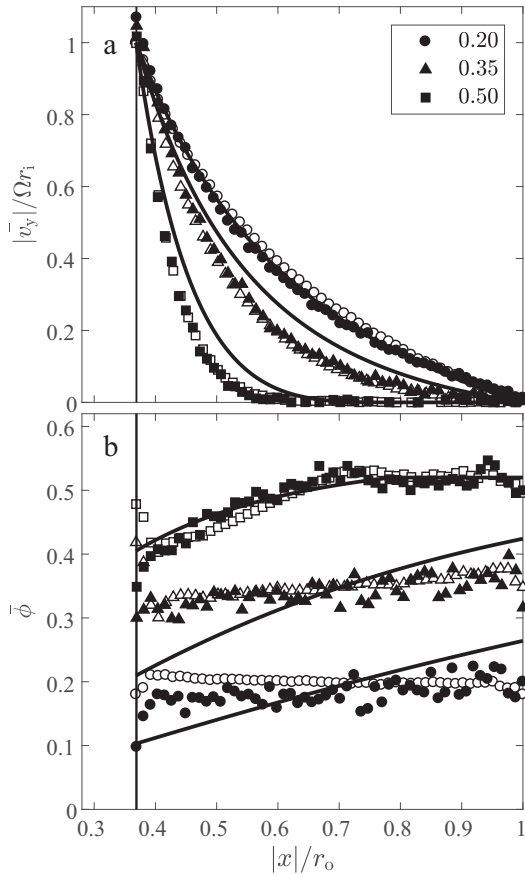
In Fig. 4 we compare the experimental velocity and concentration profiles to the model of Morris and Boulay [42] for steady-state Couette flow. The model describes the radial SVF profile  $\phi(r)$ , arising from a radial momentum balance, implicitly as

$$\frac{\eta_n(\phi)}{\eta_s(\phi)} = A_2 r^{(1+\lambda_2)/\lambda_2}, \quad (3)$$



**Fig. 3.** Steady-state radial velocity profiles for a neutrally-buoyant suspension of polyethylene microspheres in a Newtonian carrier liquid as (a–c) absolute velocity and (d–f) normalized velocity. The solid volume fractions were (a and d)  $\phi = 0.2$ , (b and e)  $0.35$ , and (c and f)  $0.5$ . The filled symbols were obtained at low field, the open symbols at high field. The legend indicates the rotation rate of the inner wall at  $r_i$  (position shown by the solid vertical lines) and applies to all plots.





**Fig. 4.** Steady-state radial (a) velocity and (b) SVF profiles for a neutrally-buoyant suspension of polyethylene microspheres in a Newtonian carrier fluid. The position of the inner rotating wall at  $r_1$  is shown by the solid vertical lines. The scaled velocities and SVF profiles are independent of rotation rate between  $\Omega = 10$  and 40 rpm, and the data shown are an average over the different rotation rates. The legend indicates  $\phi$  and applies to both plots; the filled symbols correspond to the low field experiments and the open symbols correspond to the high field experiments. Predictions from Eqs. (3)–(6) (solid curved lines) are shown for comparison.

with the relative normal and shear viscosities (made dimensionless with the solvent viscosity) given by

$$\eta_n(\phi) = K_n \frac{(\phi/\phi_m)^2}{(1 - \phi/\phi_m)^2} \quad (4)$$

$$\eta_s(\phi) = 1 + 2.5\phi_m \frac{(\phi/\phi_m)}{(1 - \phi/\phi_m)} + K_s \frac{(\phi/\phi_m)^2}{(1 - \phi/\phi_m)^2}, \quad (5)$$

where  $\phi_m = 0.52$  is the maximum packing fraction (as found empirically above).  $K_s = 0.1$ ,  $K_n = 0.75$ , and  $\lambda_2 = 0.8$  are rheological fitting parameters with the values taken from [42] without modification. Note that Eq. (5) matches [50], taking account of a typographical error in [42]. For a given value of the constant  $A_2$ , Eq. (3) may be solved for  $\phi(r)$ ;  $A_2$  is then fixed by specifying the average SVF. Once the SVF profile is obtained in this way, the shear rate is then given by

$$\dot{\gamma}(r) = \frac{C}{r^2 \eta_s(\phi)}, \quad (6)$$

which is integrated to obtain the velocity profile. The wall velocities fix the value of the constant  $C$ .

The velocity profiles predicted by the model match the experiment fairly well, except at higher SVF, see Fig. 4(a). Shortcomings of the suspension-balance model are exposed when we consider

the SVF profiles, see Fig. 4(b). The model profile at  $\phi = 0.5$  shows reasonable agreement, predicting the maximum packing fraction to be reached at a radius  $r_c/r_0 \approx 0.79$ , similar to  $r_c/r_0 \approx 0.65$  found experimentally. The model makes no prediction beyond this radius and in this region, we have set  $\phi = \phi_m$ ; alternate models exist that include compaction in the jammed region, e.g., [45]. The experimentally-measured SVF profiles for the lower values of  $\phi$  do not agree with the model predictions, despite the agreement observed in the velocity profiles. For  $\phi = 0.35$ , the SVF is seen to increase slightly towards  $r_0$ , but not to the extent predicted by the model; for  $\phi = 0.2$  the experimental SVF is relatively constant over the Couette gap.

The discrepancies in the observed and predicted SVF in Fig. 4(b) can be explained by strongly non-Newtonian behavior of the suspension, whereas the model for particle migration assumes locally Newtonian suspension rheology. Fitting a power-law model  $\tau = k\dot{\gamma}^m$  to the experimental velocity profiles, where  $\tau$  is shear stress and  $m$  the power index, we find  $m \approx 0.86$  for  $\phi = 0.20$ , and  $m \approx 0.50$  for  $\phi = 0.35$ . It is clear that any model with a locally-Newtonian rheology for the suspension cannot describe these results. In future work we aim to use our low field Rheo-NMR technology to build improved models for particle suspensions in Newtonian and non-Newtonian carrier fluids.

#### 4. Conclusions

In this paper we introduced a low field NMR system capable of generating quantitative velocity and concentration profiles in a wide-gap Couette geometry. The instrument was validated against conventional high field Rheo-NMR measurements of a Newtonian liquid and neutrally-buoyant non-colloidal spherical particles in a Newtonian carrier fluid at steady state, at matched shear rates. The two instruments provided consistent velocity profiles for the Newtonian liquid, and consistent velocity and particle concentration profiles for the suspended solids. Experimentally, the particulate system jammed at a SVF of  $\phi \approx 0.52$ , and appeared to show strongly non-Newtonian behavior (at roughly constant  $\phi$ ) for the lower volume fractions investigated. This behavior contrasted with model predictions assuming a locally Newtonian system, calling for the design of improved models. The high and low field results remained consistent always, supporting the future use of a low field bench-top instrument for Rheo-NMR studies.

The ability to readily generate robust and quantitative velocity profiles, and hence shear rate profiles, in non-Newtonian liquids using a low field NMR platform has significant advantages, primarily enabling the installation of such a device in the industrial environment. This technology enhances the utility of Rheo-NMR as a process and quality control tool. In future we plan to attach a conventional torque sensor to our Couette cell to provide complementary information from both Rheo-NMR velocity (shear rate) profiles and shear stress rheology.

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