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Computation of shear viscosity of colloidal suspensions by SRD-MD

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The behaviour of sheared colloidal suspensions with full hydrodynamic interactions (HIs) is numerically studied. To this end, we use the hybrid stochastic rotation dynamics-molecular dynamics (SRD-MD) method. The shear viscosity of colloidal suspensions is computed for different volume fractions, both for dilute and concentrated cases. We verify that HIs help in the collisions and the streaming of colloidal particles, thereby increasing the overall shear viscosity of the suspension. Our results show a good agreement with known experimental, theoretical, and numerical studies. This work demonstrates the ability of SRD-MD to successfully simulate transport coefficients that require correct modelling of HIs. © 2015 AIP Publishing LLC. [http://dx.doi.org/10.1063/1.4917039]

I. INTRODUCTION

Colloidal suspensions have a wide range of applications, ranging from commercial products and industrial processes to biological systems. Of particular significance for many of these systems is the ability to tune and optimize the transport properties of the colloids to obtain the desired flow behaviour. Colloidal suspensions are complex fluids and their rheological response is highly sensitive to thermodynamic parameters, flow geometry, and the underlying microscopic interactions.1 Resolving the interplay between all of these variables is therefore of great importance for many technological applications.

A characteristic feature of colloidal suspensions is the huge separation of time scales (of the order of ~106) and length scales (of the order of ~105) between mesoscopic colloids and microscopic fluid. Typical quantities of interest are macroscopic transport coefficients that are defined through the relevant fluctuating microscopic variables such as the local density and velocity.2 Thus, the challenge in understanding the dynamics of colloidal suspensions lies in resolving the huge gap of time and length scales. While experimental studies have revealed several important characteristics of colloidal suspensions, such as shear thinning, shear thickening,3,4 and yield stress,5 probing the underlying molecular-level details is either experimentally impossible or expensive. In addition to this, theoretical models are also limited to suspensions with a simplified definition of the fluid. For instance, the results of Einstein6,7 and Batchelor and Green8 for transport coefficients are already established for dilute and semi-dilute hard-sphere suspensions but there are no unifying empirical equivalents for concentrated suspensions.9–11

Because of the limitations of experimental and analytical approaches, especially for sheared and dense cases, simulations play an increasingly important role in the study of complex fluids. Numerical methods can be used to isolate and analyze the effects of microstructure, composition, geometry, and external perturbation that otherwise cannot be accessed by standard experiments.12 Over the past years, numerical studies have brought significant contributions to our understanding of flow behaviour and dynamics of colloidal suspensions. However, owing to the disparate nature of the length and time scales of mesoscopic systems, the prediction of rheological coefficients from a microscopic standpoint remains an arduous task. Even with the increasing computer power, a full molecular dynamics (MD) treatment of all the interactions present in the system is still computationally unsustainable. Due to the high complexity of these problems, developing an advanced mesoscopic simulation technique that can address the aforementioned challenges is still a subject of ongoing research.

Most methods available to model suspensions either have an idealized description of the fluid or ignore one of the two important aspects of suspension rheology: thermal fluctuations and hydrodynamic interactions (HIs). A traditional numerical tool for colloidal suspensions is Brownian dynamics (BD), where the fluid is treated as a continuum medium and is represented by frictional and random forces. Despite its success in modelling complex systems, the basic formulation of BD does not include hydrodynamic modes.13,14 Incorporating HIs in BD in a simplified way requires the use of hydrodynamic tensors such as the BD-Yamakawa-Rotne-Prager (BD-YRP) tensors.15,16 However, the validity of BD-YRP is constrained to relatively dilute suspensions. In addition, the computational cost increases drastically with increasing number of Brownian particles N since BD-YRP requires tensor evaluation that scales as O(N3) and diagonalization that scales as O(N5).

Alternative tools are being developed to address the gap of modelling HIs in mesoscopic simulations. Examples of these...
techniques are Lattice-Boltzmann (LB),17,18 Dissipative Particle Dynamics (DPD),19 and Stochastic Rotation Dynamics (SRD),20–22 which is otherwise known as Multi-Particle Collision (MPC) dynamics. All of these models employ a similar approach, i.e., to coarse-grain the fluid so that the spatial and temporal scales of the fluid are smaller than those of colloids but larger than its own natural scales. In the LB treatment of the fluid, the Boltzmann equation is fully discretized and solved on a lattice. In its original formulation by Ladd,17,18 thermal fluctuations are not included and artificial Langevin noise terms have to be added to the colloids to restore thermal fluctuations. An improved Fluctuating Lattice-Boltzmann (FLB) model for extended particles with well-defined hydrodynamic radii and without the external Langevin noise has been successfully implemented by Ollila et al.23 This improved method has been introduced and benchmarked for dense and spatially confined systems.

DPD is another popular simulation tool for complex fluid applications because it serves as an augmented version of MD.24 It is a particle-based approach so that the thermal fluctuations are inherently described. However, describing the fluid using DPD can be computationally demanding when dealing with larger systems since the solvent particles interact through pair-wise potentials.

Stochastic rotation dynamics-molecular dynamics (SRD-MD) is a particle-based approach first introduced by Malevanets and Kapral in 1999.20 The simplicity of its algorithm has allowed the derivation of analytic expressions for transport coefficients of pure fluids.24,25 It can successfully reproduce thermal fluctuations and HIs.20 Of equal importance is that SRD-MD is fast and computationally efficient even for cases that require dealing with external shear and high densities.27 Recent studies have also shown that it can successfully predict transport coefficients of colloidal suspensions like diffusion coefficient28–30 and conductivity31 However, the significance of HIs on diffusion coefficient and conductivity is limited if compared with its relevance on viscosity.

Calculating the shear viscosity of a suspension is a model problem for studying the rheological behaviour of mesoscopic systems.1,32 Specifically, there have been plenty of studies of suspension viscosity as a function of the particle concentration. It is a key parameter for investigating experimental phenomena such as phase transitions and yield stress. Moreover, theoretical models with concentration as its parameter are well-known8–8,39 so that it can be used as a benchmark to validate an evolving numerical model. Once validated, one can easily shift to analyze the dependence of shear viscosity on other relevant variables such as shear rate, particle structure, Peclet number (Pe), and Reynolds number (Re).

We limit our study to hard-sphere interactions mainly because their properties are theoretically and numerically well-known as compared to systems with more complex interactions. While it has been shown in Ref. 33 that SRD-MD can be used to calculate viscosity using a Derjaguin-Landau-Verwey and Overbeek (DLVO) potential description for colloid-colloid interaction, we need to justify that the computation of shear viscosity using SRD-MD agrees with existing theoretical predictions and can reproduce results that are comparable with other simulation techniques. The shear viscosity versus volume fraction is a relation that any good numerical method should satisfy. To do this, employing a hard-sphere description of the colloids is sufficient and more practical.

The objective of this paper is to provide a quantitative test on the ability of SRD-MD to model HIs on colloidal suspensions by calculating its shear viscosity. We show that SRD-MD can be used for dilute and concentrated cases. We then compare the results with known theoretical predictions and experimental data. We also propose a method that is different from Refs. 33–35 for calculating shear viscosity. We use a modified version of the SRD-MD stress tensor presented in Ref. 36, a formulation that resembles the virial equation of molecular systems. For this reason, our method of calculation allows for a better characterization of the suspension properties. Moreover, in contrast to Ref. 33 where colloids are treated as point-like particles, we employ a central-force coupling scheme to have a more realistic description of the colloid-fluid dynamics.

The paper is organized as follows. In Sec. II A, we present the system and the simulation procedure. In particular, we describe the MD-treatment of the colloids employed, SRD-treatment of the fluid, and the coupling of the two methods. In Sec. II B, we show how the parameters are chosen to achieve a proper coarse-grained approach. In Sec. II C, we explain how a planar Couette flow is modeled. We also present how the thermostat is implemented, an important point if we want to maintain the correct fluid dynamics while pumping shear. We then proceed to the calculation of relevant quantities. Section III A provides the details of formulating the stress tensors, the main ingredient for obtaining shear viscosity. In Sec. III B, we examine our results for the shear viscosity of the colloidal suspensions and compare them with theoretical equations, numerical models, and experimental data.

II. SIMULATION METHOD

A. The hybrid SRD-MD model

Our system consists of silica spheres (ρc = 2200 kg m⁻³) of radius a_c = 300 nm embedded in water (ρf = 1000 kg m⁻³). The SRD parameters and particle interactions implemented in this work are the same as in Ref. 29. The colloids are described by an inverse-power potential that is commonly used for hard-sphere colloids,28,37

\[
V_{cc} = \begin{cases} 
\epsilon_{cc} \left( \frac{\sigma_{cc}}{r} \right)^{12} & (r < r_{cc}) \\
0 & (r > r_{cc})
\end{cases} 
\]

where \( r \) is the distance between particles, \( \sigma_{cc} = 2a_c = 600 \text{ nm} \) is the interaction parameter between colloids, \( \epsilon_{cc} = 2.5k_B T \) and \( r_{cc} = 2.5\sigma_{cc} \) is the cutoff distance. The mass of the colloid is obtained from \( \rho_c \) and has a value of \( M_c = 2.49 \times 10^{-16} \text{ kg} \).

For the MD part, the positions and velocities of the colloids are updated at each time step \( \Delta t_{\text{MD}} \) using Newton’s equation of motion,

\[
v_i = \frac{dr_i}{dt} \quad \text{and} \quad m_i \frac{dv_i}{dt} = F_i.
\]
The positions and velocities are evaluated using the velocity Verlet algorithm.

The fluid dynamics is described using SRD. The fluid is represented by $N_f$ point particles placed in a simulation box of side length $L$. The SRD process consists of repeatedly streaming and colliding the fluid particles at $\Delta t_{\text{SRD}}$ intervals. During the streaming step, the positions are updated using

$$\mathbf{r}_i(t + \Delta t_{\text{SRD}}) = \mathbf{r}_i(t) + \mathbf{v}_i(t) \Delta t_{\text{SRD}},$$

where $\mathbf{r}_i$ and $\mathbf{v}_i$ are the $i$th particle’s position and velocity, respectively. In the collision step, the simulation box is divided into smaller cells ($L = 32a_0$) so that each cell has a volume of $a_0^3$ and contains an average of $\gamma$ fluid particles. Particle exchanges between cells are allowed but the number of fluid particles in the simulation box is conserved. The collision per cell is performed by rotating the velocities of the particles relative to the center of mass velocity $\mathbf{v}_\text{cm}$ according to

$$\mathbf{v}_i(t + \Delta t_{\text{SRD}}) = \mathbf{v}_\text{cm} + \mathbf{R}(\alpha) [\mathbf{v}_i(t) - \mathbf{v}_\text{cm}],$$

where $\mathbf{R}$ is the rotation matrix. The coarse grained SRD parameters include the cell size $a_0 = a_c/2 = 150$ nm, the rotation angle $\alpha = 90^\circ$, the average number density $\gamma = 5$, and dimensionless mean-free path $\lambda = 0.1$. All of these parameters are chosen to provide a compromise between resolution and computational cost. The mass of the individual particles is obtained from $\rho_f$ and has a value of $m_f = 6.75 \times 10^{-19}$ kg. The unit of time $t_0$, which dictates the values of $\Delta t_{\text{SRD}}$ and $\Delta t_{\text{MD}}$, requires a more thorough discussion and is the subject of Sec. II B.

In the paper by Ihle and Kroll, the authors showed that sorting the fluids fails to satisfy the molecular chaos condition can also be implemented but with an increase of computational cost. Second, the free volume accessible to the fluid particles is now dictated by $\sigma_{c\ell}$ rather than $a_c$. This is defined by $V_f = (32a_0)^3 - N_c \frac{4}{3} \pi \sigma_{c\ell}^3$, where $N_c$ is the number of embedded colloids. In addition, we know from Ref. 29 that the Schmidt number (Sc) of the fluid increases with increasing number of colloids. To keep Sc constant with increasing volume fraction, the number of fluid particles is decreased according to $V_f$, i.e.,

$$N_f = 32^3 \gamma - N_c \frac{4}{3} \pi \left( \frac{\sigma_{c\ell}}{a_0} \right)^3 \gamma.$$

We are now in the position to define the volume fraction occupied by the colloids,

$$\phi = \frac{N_c}{L^3} \frac{4}{3} \pi r_H^3,$$

where $r_H$ is the hydrodynamic radius. This is equal to the hard-sphere radius $a_{hs}$ only in the ideal case when there is no explicit fluid present. Analytical approaches to calculate viscosity often deal with an ideal system, where $a_{hs}$ is well defined. However, defining $r_H$ is difficult in simulations and experiments. A hard-sphere approximation requires an inverse-power potential with $n \to \infty$ in Eqs. (1) and (5) instead of $n = 12$, which will require a more expensive calculation. Moreover, the effective $r_H$ for SRD-MD is dependent on several factors including length scale and shear. This is discussed in Refs. 40 and 45 for slip boundary conditions at the colloid-fluid interface. Experimental measurements of $\phi$ are not simple either. Reference 46 provides an extensive review on the difficulties encountered when measuring $\phi$ in hard-sphere colloids and the unavoidable uncertainties associated with its reported value. To this point, we have defined two radii for the implementation of SRD-MD: $\sigma_{c\ell}/2 = a_c$ for the colloid-colloid interaction and $\sigma_{c\ell} = 0.8a_c$ for the colloid-fluid interaction. In this study, Eq. (7) is evaluated using $\sigma_{c\ell}$ because

inverse-power potential,

$$V_{c\ell} = \begin{cases} \varepsilon_{c\ell} \left( \frac{\sigma_{c\ell}}{r} \right)^{12} & (r < r_{c\ell}) \\ 0 & (r > r_{c\ell}) \end{cases},$$

where $\varepsilon_{c\ell} = 2.5k_BT/a$ and $r_{c\ell} = 2.5\sigma_{c\ell}$. The consequences of this choice of coupling are as follows. First, it is important to choose $\sigma_{c\ell}$ to be slightly less than the colloid radius $a_c$ to avoid unphysical depletion attraction between colloids. Depletion forces are due to spurious interactions that arise from the coarse-grained nature of the fluid when colloids approach each other at short distances. In this work, $\sigma_{c\ell} = 0.8a_c$ is implemented. Subsequently, the fluid particles can slightly penetrate the surface of the colloids. Even when the colloids are in contact, the fluid particles can slide between the colloids, thus, adding lubrication to colloid-colloid interactions. While the MD coupling scheme is more complete in describing colloid-fluid interaction in comparison to the SRD coupling scheme, it does not model angular momentum transfer between colloids and fluid particles leading to an effective slip-boundary condition. The friction coefficient is $\zeta = 4\pi \eta a_c$ for slip-boundary conditions and $\zeta = 6\pi \eta a_c$ for stick-boundary conditions. A stick boundary condition can also be implemented but with an increase of computational cost. Second, the free volume accessible to the fluid particles is now dictated by $\sigma_{c\ell}$ rather than $a_c$. This is defined by $V_f = (32a_0)^3 - N_c \frac{4}{3} \pi \sigma_{c\ell}^3$, where $N_c$ is the number of embedded colloids. In addition, we know from Ref. 29 that the Schmidt number (Sc) of the fluid increases with increasing number of colloids. To keep Sc constant with increasing volume fraction, the number of fluid particles is decreased according to $V_f$, i.e.,

$$N_f = 32^3 \gamma - N_c \frac{4}{3} \pi \left( \frac{\sigma_{c\ell}}{a_0} \right)^3 \gamma.$$
it is the best \( r_H \) approximation for our simulations.\(^{40} \)

Our treatment of volume fraction is similar to the definition used in Ref. 45 for colloid sedimentation velocity, where the volume fraction is defined using \( r_H \) instead of \( \sigma_c \). This is equivalent to viewing the volume fraction from the fluid particle’s point of view instead of the colloid’s point of view. This will be discussed further in Sec. III B. Table I summarizes the number of colloids and fluid particles used for each volume fraction.

<table>
<thead>
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<th>( \phi )</th>
<th>( N_c )</th>
<th>( N_f )</th>
</tr>
</thead>
<tbody>
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<td>0.0513</td>
<td>98</td>
<td>155433</td>
</tr>
<tr>
<td>0.1026</td>
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<tr>
<td>0.1791</td>
<td>342</td>
<td>134501</td>
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<tr>
<td>0.2053</td>
<td>392</td>
<td>130212</td>
</tr>
<tr>
<td>0.2356</td>
<td>450</td>
<td>125236</td>
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<td>550</td>
<td>116657</td>
</tr>
<tr>
<td>0.3142</td>
<td>600</td>
<td>112368</td>
</tr>
<tr>
<td>0.3388</td>
<td>647</td>
<td>108336</td>
</tr>
</tbody>
</table>

**B. SRD mapping**

The mesoscopic nature of colloidal suspensions results to a wide range of time scales. SRD-MD compresses this hierarchy of time scales so that only one time scale can be accurately reproduced. We choose from the two significant time scales for this study: kinetic time scale for hydrodynamic interactions and diffusion time scale for Brownian motion. The kinetic time \( \tau_K = a_c^2/\eta_{\text{H}_2\text{O}} = a_c^2\rho_f/\eta_{\text{H}_2\text{O}} = 9.0 \times 10^{-8} \text{ s} \) and diffusion time \( \tau_D = a_c^2/D_0 = 0.12 \text{ s} \) are 6 orders of magnitudes apart. These are obtained using the viscosity of water \( \eta_{\text{H}_2\text{O}} = 0.001 \text{ Pa s} \) and the diffusion coefficient of a colloid in water \( D_0 = k_B T/(6\pi a_r \eta_{\text{H}_2\text{O}}) = 7.15 \times 10^{-13} \text{ m}^2\text{s}^{-1} \). Mapping to one of the time scales is necessary to efficiently simulate these transport coefficients. SRD-MD gives the liberty of selecting the fastest time scale to optimize the simulation. In this system, mapping to diffusion time scale will bring the value of \( \tau_K \) closer to \( \tau_D \), i.e., \( 6.0 \times 10^{-3} \text{ s} \) instead of \( 9.0 \times 10^{-8} \text{ s} \). However, contracting the time scale results to quantities that are far from laboratory values. For example, the value of shear viscosity will now be \( 1.52 \times 10^{-8} \text{ Pa s} \) for the pure fluid case. When mapped back to kinetic time, the physical value of viscosity is recovered. As mentioned in Ref. 40, the quantitative value is not important as long as the correct regime of the hydrodynamic numbers is obtained and the different time scales are well separated. The diffusion time scale is used in this study because it significantly reduces the computation time by allowing the use of bigger time steps. Hence, longer time scales can be reached at faster rates. Moreover, while the simulated value of shear viscosity is different from its physical value, expressing it in terms of its relative value still preserves all its qualitative features: \( \eta_s = \eta_s/\eta_0 \). This means that we get the same ratio between the suspension viscosity \( \eta_s \) and pure fluid viscosity \( \eta_0 \) regardless of the time scale used.

By adopting the diffusion time scale, the diffusion time can be expressed as a function of the SRD time unit \( t_0 \). A detailed derivation can be found in Ref. 29. Equating the diffusion time in the simulation to the physical value of diffusion time, we get \( t_0 = 7.37 \times 10^{-4} \text{ s} \). Thus, the SRD time step is \( \Delta t_{\text{SRD}} = \lambda t_0 = 7.37 \times 10^{-5} \text{ s} \). If instead the kinetic time scale is used, the resulting unit of time is \( t_0 = 1.12 \times 10^{-8} \text{ s} \). This option significantly reduces the computation time up to \( 10^4 \). The value of \( \Delta t_{\text{MD}} \) is smaller and generally depends on the steepness of the potential. It was found that \( \Delta t_{\text{MD}} = 8\Delta t_{\text{SRD}} \) is sufficient to resolve the correct Newtonian dynamics for the inverse-power potential used in this paper.

Consequently, the value of temperature is also modified. Similar to the transport coefficients, the actual temperature is mapped from experimental scale to SRD scale and vice versa. A system temperature of 293 K is equivalent to a SRD temperature of

\[
T = \frac{a_c^2 m_f}{k_B t_0} = 2.02 \times 10^{-3} \text{ K}. \quad (8)
\]

A detailed mapping procedure for slip-boundary condition can be found in Refs. 29 and 40.

**C. Introducing shear and thermostat**

The initial velocities of the particles obey a Maxwell-Boltzmann distribution. The infinite periodic system is then subjected to a uniform shear in the \( xy \) plane. Shear is applied in the \( y \) direction and the gradient is along the \( x \) direction. The shear rate is defined as

\[
\dot{\gamma} = \frac{\Delta v_y}{\Delta x}, \quad (9)
\]

where \( \Delta v_y \) is the shear velocity. Shear is imposed by using Lees-Edwards boundary conditions (LEBCs)\(^{47} \) which allows the simulation of a planar Couette flow without actual boundaries. The benefit of using this algorithm is that it does not have the numerical instabilities associated with introducing actual walls. The process works by updating the positions and velocities of the particles with the usual periodic boundary conditions for \( y \) and \( z \) directions. However, when a particle crosses the upper or lower boundaries of the simulation box, i.e., \( \Delta x = 0 \) and \( \Delta x = L_x \), its position and velocity are updated with a different rule to sustain the shear. Particles crossing the upper boundary will have an additional velocity of \( +\gamma L_x \) and a position shift of \( +\gamma L_x t \), where \( t \) is the total elapsed time. In contrast, particles crossing the lower boundary will have an additional velocity of \( -\gamma L_x \) and a position shift of \( -\gamma L_x t \). LEBCs are applied for both the colloids and fluid particles.

A shear rate of \( \dot{\gamma} = 100 \text{ s}^{-1} \) is used and gives the system \( \text{Pe} = 12.6 \). Viscosity is measured when the velocity profile is already linear. Fig. 1 shows the velocity profile inside a simulation box for a suspension with \( \dot{\gamma} = 100 \text{ s}^{-1} \) and \( \phi = 0.1534 \).

The introduction of shear results to continuous pumping of external energy into the system. The kinetic energy of the system is kept constant by applying a thermostat. The instantaneous kinetic temperature of the system is defined as...
Velocity profile for a shear rate of $\dot{\gamma} = 100$ s$^{-1}$. A linear profile is observed at 0.2 s.

$$T = \frac{1}{k_B d} \sum_{i=1}^{N} m_i (v_i - u_y)^2,$$

where $d = 3$ is the dimensionality and $u_y$ is the simulated shear velocity. A constant temperature is maintained by using a Monte Carlo scaling thermostat.$^{26,48}$ Several types of thermostats have been employed and Monte Carlo scaling proves to be the most efficient. This method also conserves linear momentum and energy without smearing the velocity profile or changing the viscosity of the fluid. The SRD coupling in Refs. 26 and 33 also changes the temperature of the colloids because they participate to the SRD collision steps. In our case, the thermostat is only applied to the fluid particles. Hence, the interaction of the colloid with the thermal bath happens only during colloid-fluid interactions. The thermostat described has been tested for shear rates between 10 s$^{-1}$ and 500 s$^{-1}$.

An equilibration part consisting of 2000 – 6000 SRD steps is first carried out. This is the part where $T$ is slowly driven to the desired $T$. This approach is needed to ensure that the viscosities are measured with the correct linear velocity profile and at the correct system temperature.

III. CALCULATIONS

A. Stress tensors

The viscosity is obtained from $\eta = \langle \sigma_{yx} \rangle / \dot{\gamma}$, where $\langle \ldots \rangle$ denotes a time average. We use the instantaneous stress tensors from Ref. 36 and adapt it for our system. For the pure fluid case ($\phi = 0$), these are given by

$$\sigma_{\text{kin,}yx} = -\frac{m_f}{V} \sum_{i=1}^{N_f} \dot{u}_i \dot{u}_y - \frac{m_f \gamma \Delta t_{\text{SRD}}}{2V} \sum_{i=1}^{N_f} \dot{u}_i^2,$$  \hspace{1cm} (11)

and

$$\sigma_{\text{col,}yx} = -\frac{1}{V \Delta t_{\text{SRD}}} \sum_{i=1}^{N_f} \Delta \rho_{ix} r'_{ix},$$  \hspace{1cm} (12)

where $V = L^3$ for the pure fluid case and the shorthand notations $\dot{v}_i = v_i(t - \Delta t)$ and $\Delta \rho_{ix} = m_f (\dot{v}_i - \dot{v}_y)$ are used. The apostrophe symbol ($'$) is for quantities measured after LEBCs are imposed. Equations (11) and (12) correspond to the kinetic and collisional contributions, respectively, so that the total stress tensor of the fluid is $\sigma_{yx}^{\text{fluid}} = \sigma_{\text{kin,}yx} + \sigma_{\text{col,}yx}$. Viscosity is calculated for $r'_{ix}$ before and after imposing the grid-shift procedure. We found that an agreement with the known analytical fluid viscosity is obtained when $r'_{ix}$ is measured after the grid-shifting algorithm. We compared them with the analytical predictions for the shear viscosity of pure fluids as a function of SRD parameters.$^{24,25}$

$$\eta_{\text{kin}} = \frac{5\gamma^2 k_1}{(\gamma - 1 + e^{-\gamma})(4 - 2 \cos \alpha - 2 \cos 2\alpha)} - \frac{1}{2} \gamma k_1,$$  \hspace{1cm} (13)

$$\eta_{\text{col}} = k_2 (1 - \cos \alpha) (\gamma - 1 + e^{-\gamma}),$$  \hspace{1cm} (14)

where $k_1 = k_B T \Delta t_{\text{SRD}}/a_0^2$, $k_2 = m_f/(12a \Delta t_{\text{SRD}})$, and $\eta_{\text{tot}} = \eta_{\text{kin}} + \eta_{\text{col}}$. Since we used a small value for $\lambda = 0.1$, the viscosity is dominated by the momentum transfer from particle collisions and not by the streaming of the fluid. At $\alpha = 90^\circ$, the calculated viscosity is $1.5 \times 10^{-3}$ Pa s. The same value of shear viscosity is obtained using other approaches.$^{21,24}$ The viscosity remains constant between 50 s$^{-1}$ and 100 s$^{-1}$. This is expected since experimental results show that shear thickening and shear thinning occur at shear rates beyond these values (see Ref. 49).

A small deviation from the theoretical values is seen at higher angles ($120^\circ$–$150^\circ$). A similar observation is also found when the method proposed in Ref. 24 is used. This small deviation may be due to (1) the increase in the number of degrees of freedom and (2) the small effect of the coarse-grained temperature ($2.02 \times 10^{-3}$ K). This may lead to a small loss of molecular chaos at low temperatures (see Ref. 24).

Because the simulations of the pure fluid case are in good agreement with analytical equations, we have generalized the above approach to obtain the shear viscosity of colloidal suspensions. We use the stress tensor

$$\sigma_{yx}^{\text{colloid}} = -\frac{M_c}{V} \sum_{i=1}^{N_c} \dot{u}'_y \dot{u}_i - \frac{M_c \gamma \Delta t_{\text{SRD}}}{2V} \sum_{i=1}^{N_c} u_i^2,$$

$$-\frac{1}{V} \sum_{i=1}^{N_c} F_y r'_{ix},$$

Since the coupling method used in this paper is MD, the colloid forces are explicitly defined. Using Eqs. (1) and (5), $F_y$ is obtained directly from $\partial (V_{c,c} + V_{c,f})/\partial y$ instead of the pseudo force $\Delta \rho_{iy}$ used in Eq. (12). The total stress tensor of the suspension is calculated from the sum of the contributions from the embedded colloids and suspending fluid,

$$\sigma_{yx} = \sigma_{yx}^{\text{fluid}} + \sigma_{yx}^{\text{colloid}}.$$

Therefore, $\eta_s = \eta_s^{\text{fluid}} + \eta_s^{\text{colloid}}$. The results of five independent simulations are reported and the error bars corresponding to the standard deviations are smaller than the symbols used.
B. Shear viscosity

For the theoretical framework of shear viscosity, we use Einstein’s equation for the dilute case and Krieger’s semi-empirical equation for the concentrated case. The hard-sphere equations are given by

\[ \eta_r = 1 + 2.5\phi \quad \text{for } \phi \rightarrow 0 \quad (17) \]

and

\[ \eta_r = \left(1 - \frac{\phi}{\phi_m}\right)^{-p} \quad \text{for } \phi \rightarrow \phi_m, \quad (18) \]

where \( \phi_m \) is the maximum packing fraction and \( p \) is a parameter that depends on \( \phi_m \) and the shear. The deviation from the linear approximation is due to the increase of the probability of collisions and because hydrodynamic interactions become more significant at higher volume fractions.

Fig. 2 shows how shear viscosity increases with increasing value of \( \sigma_{cf} \). The predicted value using Eq. (17) is also shown for comparison. As \( \sigma_{cf} \) approaches \( a_c \), the calculated shear viscosity moves closer to the analytical result. However, in the limit \( \sigma_{cf} \rightarrow a_c \), depletion attraction starts to occur and the calculated viscosity in this region (\( \sim 0.95a_c \)) is no longer valid. Cluster formations caused by the depletion attraction are observed when \( \sigma_{cf} = a_c \), but are eliminated when \( \sigma_{cf} = 0.8a_0 \). The result of Fig. 2 justifies the choice of using \( \sigma_{cf} \) as the hydrodynamic radius in Eq. (7).

The shear viscosity of the colloids (\( \eta_{\text{colloid}} \)) can also be divided into components owing to the different contributions to the stress tensor. The virial part, described by the last term in Eq. (15) and represented by blue diamonds in Fig. 3, increases because of the increase in the probability of collisions among other colloids when the volume fraction increases. It is also the most dominant contribution to the overall viscosity. The kinetic contribution, described by the first two terms in Eq. (15), is represented by red squares in Fig. 3. It represents the pressure exerted by the colloids on a surface. As expected, it increases with increasing volume fraction. However, it is smaller in comparison to the contribution from the colloid-colloid interactions. The shear viscosity from the colloids is therefore dominated by the momentum exchanges from inter-particle interactions rather than from streaming.

We use the same treatment to analyze what is happening to the shear viscosity of the SRD fluid (\( \eta_{\text{fluid}} \)) as the volume fraction increases (see Fig. 4). The total shear viscosity (black circles) is comprised of the kinetic part (red squares) and collisional part (blue diamonds). We also compare the viscosity of the SRD fluid (\( \eta_{\text{fluid}}(\phi) \)) with the predicted shear viscosity for the pure fluid case (\( \eta_{\text{fluid}}(\phi = 0) \)) given by Eqs. (13) and (14) from Refs. 24 and 25. An average difference of \( \Delta = 2.7 \times 10^{-9} \) Pa s is measured.
and collisional part (blue diamonds). For the collisional contribution, which describes the collisions among fluid particles, no significant change can be observed. In contrast, the kinetic contribution, which describes the streaming of the fluid particles, increases by $\Delta = 2.7 \times 10^{-4}$ Pa s when colloids are present. This is due to the additional momentum imparted by the fluid-colloid interactions when streaming.

Finally, we compare the suspension viscosity with other known results in Fig. 5. For the experimental part, we compared with the data by Segré and van der Werff, where they used undeformable, sterically stabilized spheres. The volume fraction is approximated using the maximum packing fraction of the system, $\phi_m = 0.494$ for Segré and $\phi_m = 0.63$ for van der Werff. It can be observed that our numerical results lie a little below the experimental data of Refs. 53 and 54. This can be due to finite size effects that make the shear rate higher than the expected value.

Our results are comparable with those of Koelman and Hoogerbrugge, where they simulate hard-sphere suspensions under shear flow using DPD. The hard-sphere case is imposed by using the moment of inertia and Euler’s equation of motion for rigid bodies.

The behaviour of hard-spheres suspensions in the absence of HIs was analyzed using BD in Ref. 50. In order to remove Pe and volume fraction dependencies of the shear viscosity, Ref. 50 employed an algorithm, in which after the Brownian position updates, the overlaps are checked and successively removed. By comparing the behaviour of the two systems, HIs increase the viscosity of the suspension. In contrast to the BD where the effect of fluid particles is represented by the frictional and stochastic terms, the fluid description in SRD-MD is more explicit. It is observed that the overall contribution of the fluid decreases with increasing volume fraction since the number of fluid particles decreases with increasing concentration to maintain a constant Sc for the fluid. Even so, the treatment of the fluid by SRD-MD provides lubrication between colloid particles because the fluid can slide between them. This increases the chances of collisions among colloids and the overall friction of the system.

From the theoretical standpoint, the results follow Eq. (17) for low volume fractions and start to diverge near $\phi = 0.15$. For Eq. (18), we used a fitting parameter of $\phi_m = 0.74$ because the highest packing achievable in our simulation box is hexagonal closed packing. Previous theoretical works have shown that $p = 2$ for a variety of situations so that Eq. (18) reduces to $\eta_f = (1 - \phi/0.74)^{-2}$.

IV. CONCLUSIONS

We have demonstrated that SRD-MD can be used to simulate the shear viscosity of colloids. The mapping between physical and simulated values can be done by SRD-MD without losing relevant information. The use of diffusion time scale for computation significantly decreases the simulation time. Shear can also be applied to systems that uses MD as a coupling scheme. A Monte Carlo scaling thermostat is necessary to maintain the correct thermodynamic properties of the fluid.

The choice of $\sigma_{cf}$ in defining the volume fraction also gives the correct viscosity values. A small $\sigma_{cf}$ allows more fluid particles to perturb the flow field surrounding the colloid.

The use of stress tensors for viscosity evaluation provides a better characterization tool than what is previously available as in Ref. 33. We observed that the main contribution to the shear viscosity of the suspension comes from the inter-particle collisions rather than streaming. The contribution of the fluid particles to the stress tensor decreases with increasing concentration. This is a consequence of the coupling scheme used and is necessary in order to preserve the Sc of the fluid. However, their role in facilitating lubrication between colloidal particles aids in the increase of kinetic and collisional viscosities of the colloids, which is not seen in simulations neglecting hydrodynamic effects. Our results are also comparable with other experimental and theoretical studies. While results for hard-sphere suspensions employing other methods, like DPD, have been compared to Eqs. (17) and (18), this is the first time that the relation between shear viscosity and volume fraction is reproduced by SRD-MD. Hence, our study serves as a final test on the ability of SRD-MD to predict transport coefficients and rheological parameters where HIs are significant. Moreover, in comparison to DPD, the SRD treatment of the system is computationally faster especially at intermediate and high-volume fractions. SRD-MD is also a practical tool for studying the rheology of dense suspensions, an important aspect for studying yield-stress and other phenomena that are occurring in the non-Newtonian regime.

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