Universality in Driven and Equilibrium Hard Sphere Liquid Dynamics

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We demonstrate that the time evolution of the van Hove dynamical pair correlation function is governed by adiabatic forces that arise from the free energy and by superadiabatic forces that are induced by the flow of the van Hove function. The superadiabatic forces consist of drag, viscous, and structural contributions, as occur in active Brownian particles, in liquids under shear and in lane forming mixtures. For hard sphere liquids, we present a power functional theory that predicts these universal force fields in quantitative agreement with our Brownian dynamics simulation results.

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The van Hove function is arguably one of the most fundamental correlators that characterize the dynamical pair structure of a liquid on the microscopic scale [1,2]. It measures the probability of finding two particles at a distance \( r \), where the particles are randomly chosen, but with a time lapse of duration \( t \) between the two position measurements. Both the motion of the same particle and spatiotemporal correlations of two distinct particles are captured. Significant physical insights into the dynamics of simple and complex systems could be gained from studying their van Hove function. Examples thereof include cage formation in systems 

accounting for the observed reduction of the interparticle correlations [32] compared to benchmark data from Brownian dynamics (BD) computer simulations. This trend persists even when choosing Rosenfeld’s fundamental measure theory [33–36] as an excellent approximation for the (hard sphere) free energy functional. Accounting for the observed reduction of particle mobility at increased density requires empirical adjustments to the DDFT framework [6–8,27].

Power functional theory (PFT) [37] provides formally exact test particle dynamics [28], albeit very little explicit knowledge of the crucial superadiabatic force contributions [37,38], i.e., those beyond DDFT, had originally been available [28]. In BD simulation work, it was shown that the superadiabatic forces that govern the van Hove function are both significant in magnitude and nontrivial in their spatial and temporal structure [39]. In a variety of non-equilibrium systems, different superadiabatic force types were identified as providing the key mechanisms for prominent physical effects, such as the emergence of viscous and structural forces in BD flow [40–42], motility-induced phase separation in active Brownian particles [43,44], spontaneous lane formation in counterdriven mixtures [45], and memory-induced motion reversal [46].

Here we show that the identical types of superadiabatic forces that rule the behavior of these driven systems determine, qualitatively and quantitatively, the van Hove function and hence the intrinsic equilibrium dynamics. That the same form of superadiabatic forces applies across such a wide range of different physical situations indicates that the microscopic liquid dynamics are governed by universal mechanisms. Besides the conceptual importance of this finding, it allows concrete crossfertilization between results obtained for apparently very different systems.
Within the dynamical test particle limit, the van Hove function is expressed as a time-dependent one-body density profile \( \rho(r, t) \); here \( r \) indicates position and \( t \) time. Often one splits into self and distinct parts: 
\[
\rho(r, t) = \rho_{\text{self}}(r, t) + \rho_{\text{dist}}(r, t).
\]
At the initial time, the test (“self”) particle is taken to be at the origin, and the distinct particles are distributed according to the (static) pair correlation function \( g(r) \) of the bulk liquid (as prescribed by Percus’ static test particle limit [47]). Hence, the initial conditions at \( t = 0 \) are \( \rho_{\text{self}}(r, 0) = \delta(r) \) and \( \rho_{\text{dist}}(r, 0) = \rho_b g(r) \), where \( \delta(\cdot) \) indicates the Dirac delta function, \( \rho_b \) is the bulk fluid number density, and \( r = |r| \). Figure 1(a) and (b) depicts an illustration. The dynamics of the van Hove function are associated with time-dependent one-body self and distinct currents, \( J_{\text{self}}(r, t) \) and \( J_{\text{dist}}(r, t) \), respectively. The total van Hove current is the sum \( J = J_{\text{self}} + J_{\text{dist}} \).

A continuity equation holds for each species:
\[
\frac{\partial \rho^\alpha}{\partial t} = -\nabla \cdot J^\alpha, \quad \alpha = \text{self, dist}
\]
where \( \rho^\alpha \) labels the two different species, and \( \nabla \) indicates the derivative with respect to \( r \). The partial one-body currents \( J^\alpha(r, t) \) arise from free diffusion and from internal interactions. Hence, the one-body force density balance relation is

\[
J^\alpha(r, t) = \frac{F^\alpha}{\rho^\alpha}, \quad \alpha = \text{self, dist}
\]

The ideal one-body force density is \( \frac{1}{2} k_BT \nabla \rho^\alpha \) and \( F^\alpha_{\text{ideal}} = \frac{1}{2} k_BT \nabla \rho^\alpha \).

FIG. 1. Two-body dynamics of the hard sphere fluid. (a) Illustration of the van Hove dynamical two-body correlation function in a bulk liquid of hard spheres of diameter \( \sigma \) at time \( t = 0 \) and at \( t > 0 \) (b). (c)–(g) Results for the dynamical decay of the two-body structure of a bulk liquid of hard spheres at packing fraction 0.35 at times \( t = 0.1\tau \) (left column), 0.3\( \tau \) (middle column), and 0.6\( \tau \) (right column) and as a function of the scaled distance \( r/\sigma \). (c) Total (\( \rho \)), self (\( \rho_{\text{self}} \)), distinct (\( \rho_{\text{dist}} \)), and differential (\( \rho_{\Delta} \)) parts of the van Hove function, as indicated. The results from BD simulation (symbols) of the time evolution and from MC simulation (lines) of the corresponding adiabatic state coincide on the scale of the plot. (d) Self part of the superadiabatic force density \( F_{\text{self}}^\alpha \) as obtained from BD (symbols) and from PFT (solid line); also shown is the adiabatic self force density \( F_{\text{self}}^\alpha \) from MC simulation (blue symbols) and DFT (blue line). The ideal self force density \( -k_BT \nabla \rho_{\text{self}} \) is shown as a reference. (e) Distinct part of the superadiabatic force density \( F_{\text{dist}}^\alpha \) as obtained from BD and from PFT. (f) Differential superadiabatic (drag) force density \( G_{\text{sup}}^\alpha \) as obtained from BD (symbols) and from PFT (black line), and differential adiabatic force density \( G_{\text{ad}}^\alpha \) as arising from the adiabatic self correction. (g) Species-independent superadiabatic force field \( f_{\text{sup}} \) as obtained from BD and from PFT, along with the theoretical viscous (\( f_{\text{visc}} = -\rho^{-1} \partial \rho_{\text{visc}} / \partial v \)) and structural contributions (\( f_{\text{struc}} = -\rho^{-1} \partial \rho_{\text{struc}} / \partial v \)), where \( f_{\text{sup}} = f_{\text{visc}} + f_{\text{struc}} \). For the sake of clarity, the results in (d)–(g) at \( t = 0.3\tau \) and 0.6\( \tau \) are multiplied by factors of 2, 4, 8, or 16 as indicated.
\[ \gamma J_\alpha = -k_B T \nabla \rho_\alpha + F^\alpha_{\text{int}}, \]  

(1)

where \( \gamma \) is the friction constant against the static background, \( k_B \) is the Boltzmann constant, \( T \) indicates absolute temperature, and \( F^\alpha_{\text{int}}(r,t) \) is the internal force density distribution that acts on species \( \alpha \). No external forces act in the bulk system, and the time-dependent (non-equilibrium) situation is solely introduced by the initial conditions \( \rho_\alpha(r,0) \).

The internal force density consists of adiabatic \( (F^\alpha_{\text{ad}}) \) and superadiabatic \( (F^\alpha_{\text{sup}}) \) contributions [37,38], according to the sum \( F^\alpha = F^\alpha_{\text{ad}} + F^\alpha_{\text{sup}} \). Here \( F^\alpha_{\text{ad}}(r,t) \) is the force density distribution in an equilibrium (“adiabatic”) system that is defined to possess one-body density profiles \( \rho^\alpha_{\text{ad}}(r) \) that are identical to those in the dynamical system at time \( t \): \( \rho^\alpha_{\text{ad}}(r) \equiv \rho^\alpha_{\text{ad}}(r,t) \). This adiabatic construction is performed at each point in time, and hence \( \rho^\alpha_{\text{ad}}(r) \) depends parametrically on \( t \). The interparticle interaction potential in the adiabatic system is identical to that in the original dynamical system. The density distributions \( \rho^\alpha_{\text{ad}}(r) \) in the adiabatic system are stabilized by species-dependent external potentials \( V^\alpha_{\text{ext}}(r) \), which are guaranteed to exist in the adiabatic system due to the Mermin-Evans map of classical density functional theory (DFT) [2,29].

The adiabatic force density can be obtained by direct sampling in the adiabatic system [48] or, as we do here, from the force balance in the adiabatic system:

\[ F^\alpha_{\text{ad}} = k_B T \nabla \rho^\alpha_{\text{ad}} + \rho^\alpha_{\text{ad}} \nabla V^\alpha_{\text{ext}}, \]  

(2)

where all quantities on the right-hand side are known. In practice, we use a variant of the custom flow iterative method [48], where we sample the density profile at each iteration step using Monte Carlo (MC) simulation and adjust the external potentials \( V^\alpha_{\text{ext}}(r) \) accordingly [38] until the sampled density profiles in the adiabatic system match the dynamical (“target”) density profiles \( \rho^\alpha_{\text{ad}}(r,t) \).

Within classical density functional theory, the adiabatic internal force density acting on species \( \alpha \) is given by \( F^\alpha_{\text{ad}}(r,t) = -\rho_\alpha(r,t) \nabla F_{\text{exc}}/\delta \rho_\alpha(r,t) \), where \( F_{\text{exc}} \) is the excess (over ideal gas) intrinsic Helmholtz free energy functional. For the case of hard spheres, Rosenfeld’s fundamental measure theory [33–36] constitutes an excellent approximation for \( F_{\text{exc}} \). We furthermore use the “quenched” approach by Stopper et al. [8], where a self-correction is applied in order to account for the fact that the self density profile represents a single particle sharply (rather than a grand ensemble average). This approach avoids having to use canonical decomposition [49,50] in order to generate results that are specific to fixed particle number.

Within PFT, the superadiabatic force density is obtained from a functional derivative of the superadiabatic excess free power functional \( P^\text{exc}_t \) according to

\[ F^\alpha_{\text{sup}}(r,t) = -\frac{\delta P^\text{exc}_t}{\delta \rho_\alpha(r,t)}, \]  

(3)

where the derivative is taken at fixed density profiles, and the species-resolved one-body velocity profile is \( \mathbf{v}_\alpha(r,t) = J_\alpha(r,t)/\rho_\alpha(r,t) \). As an approximation, we use a functional that consists of drag [43–45], viscous [40–42,45,46], and structural [41,42,44] contributions, \( P^\text{exc}_t = P^\text{drag}_t + P^\text{visc}_t + P^\text{struc}_t \), according to

\[ P^\text{exc}_t = \frac{C_{\text{drag}}}{2} \int dr \rho_{\text{self}} \rho_{\text{dist}} \left( \mathbf{v}_{\text{self}} - \mathbf{v}_{\text{dist}} \right)^2 \]
\[ + \int dr dr' \int_0^t dt \nu n_3 (\nabla \cdot \mathbf{v})(\nabla' \cdot \mathbf{v}') K_{\text{visc}} \]
\[ - \int dr dr' \int_0^t dt' (n_3' \mathbf{v}')^2 (\nabla \cdot \mathbf{J}) K_{\text{struc}}, \]  

(4)

where \( C_{\text{drag}} \) is a constant and the kernels \( K_{\text{exc}}(\Delta \mathbf{r}, \Delta t) \), where \( e = \text{visc}, \text{struc} \), depend on the relative spatial and temporal distances \( \Delta \mathbf{r} = \mathbf{r} - \mathbf{r}' \) and \( \Delta t = t - t' \); the local packing fraction \( n_3(r,t) \) is obtained by convolution with \( \rho(r,t) \) [33–36]; \( \nabla \) indicates the derivative with respect to \( r' \), and we use the shorthand \( n_3' \equiv n_3(r',t) \) and \( \mathbf{v}' \equiv \mathbf{v}(r',t) \). Here the total microscopic velocity profile is \( \mathbf{v} = \mathbf{J}/\rho \). We use the diffusing memory form [46] for \( K_{\text{exc}}(\Delta \mathbf{r}, \Delta t) \), which consists of a product of a constant \( C_e \) that controls the overall strength, an exponential decay with decay time constant \( \tau_e \), and a diffusing Gaussian with diffusion constant \( D_e \). Explicitly the form is

\[ K_{\text{exc}}(\Delta \mathbf{r}, \Delta t) = C_e \exp[-\Delta \mathbf{r}^2/(4D_e \tau_e)] / (4\pi D_e \tau_e)^{3/2}. \]  

(5)

The derivative Eq. (3), when applied to Eq. (4), yields an explicit expression for \( F^\alpha_{\text{sup}} \), which we evaluate below, using BD data for \( \rho_\alpha \) and \( \mathbf{v}_\alpha \) as input. We choose the following set of parameters. The drag strength is \( C_{\text{drag}} = 2.2 \gamma \sigma^5 \). The values for viscous memory kernel are identical to those used in Ref. [46]: \( C_{\text{visc}} = 5.8 k_B T / (\sigma^3 \tau) \), \( D_{\text{visc}} = 5.6 \sigma^2 / \tau \), and \( \tau_{\text{visc}} = 0.02 \tau \). The structural memory kernel has \( C_{\text{struc}} = 0.42 k_B T \sigma^2 / \tau^2 \), \( D_{\text{struc}} = 0.25 \sigma^2 / \tau \), and \( \tau_{\text{struc}} = 0.8 \tau \). Here the natural (Brownian) timescale is \( \tau = \gamma \sigma^2 / (k_B T) \), where \( \sigma \) is the hard sphere diameter. For standard colloids with diameter \( \sigma = 1 \mu m \) dispersed in water at \( T = 20^\circ \text{C} \), using the Stokes-Einstein form for \( \gamma = 3 \pi \eta \sigma / 2 \) gives \( \gamma = 9.44 \times 10^{-13} \text{ kg s}^{-1} \), which yields \( \tau = 2.33 \text{ s} \). The corresponding memory times are \( \tau_{\text{visc}} = 0.046 \text{ s} \) and \( \tau_{\text{struc}} = 1.87 \text{ s} \), i.e., values that are well inside of an experimentally accessible range. Using larger colloids [6,51] scales up the values for the memory times accordingly; the particles used, e.g., for the (two-dimensional) system of Ref. [6] are of size \( \sigma = 4.04 \mu m \), which correspondingly upscales the values for both memory times by a factor of 4.
In order to gain further insight into the nature of the relevant forces, we follow Ref. [45] and rewrite the internal forces that act in a binary mixture as consisting of a nonselective (“differential”) force field (\(f_{\text{int}}\)) and a selective (“total”) force density (\(G_{\text{int}}\)) such that the self and distinct internal force density distributions, respectively, can be expressed as

\[
F_{\text{int}}^{\text{self}} = \rho_{\text{self}} f_{\text{int}} + G_{\text{int}}, \quad F_{\text{int}}^{\text{dist}} = \rho_{\text{dist}} f_{\text{int}} - G_{\text{int}}.
\]

Using the new fields \(f_{\text{int}}\) and \(G_{\text{int}}\) in the partial force density balance Eq. (1) leads to equations of motion for the total and for the “differential” motion,

\[
\gamma \nabla = -k_B T \nabla \ln \rho + f_{\text{int}},
\]

\[
\gamma J_\Delta = -k_B T \nabla \rho_\Delta + \rho_\Delta f_{\text{int}} + 2G_{\text{int}},
\]

where the differential van Hove current is \(J_\Delta = J_{\text{self}} - J_{\text{dist}}\), and the differential van Hove function is \(\rho_\Delta = \rho_{\text{self}} - \rho_{\text{dist}}\). Solving the linear set of Eq. (6) yields

\[
f_{\text{int}} = F_{\text{int}}/\rho, \quad G_{\text{int}} = (\rho_{\text{dist}} F_{\text{int}}^{\text{self}} - \rho_{\text{self}} F_{\text{int}}^{\text{dist}})/\rho.
\]

which allows one to obtain results for \(f_{\text{int}}(r, t)\) and \(G_{\text{int}}(r, t)\) [from the correlators on the right-hand sides of Eq. (9)]. Due to the linearity of the transformations Eq. (9), splitting into adiabatic and superadiabatic contributions applies according to \(f_{\text{int}} = f_{\text{ad}} + f_{\text{sup}}\) and \(G_{\text{int}} = G_{\text{ad}} + G_{\text{sup}}\).

Figure 1(c) presents results for the van Hove function of hard spheres. Shown are the self and the distinct parts, \(\rho_{\text{self}}(r, t)\) and \(\rho_{\text{dist}}(r, t)\), at three different representative times: \(t/\tau = 0.1, 0.3, 0.6\). The results are obtained using event-driven BD computer simulations [52]. We use \(N = 1090\) particles in a three-dimensional simulation box of size \(15 \times 10 \times 10 \sigma^3\). The sampling is based on \(10^8\) time steps of size \(10^{-3} \tau\), and hence an overall simulation time of \(10^7 \tau\). Appropriate filling of histograms of particle pair distances yields results for the van Hove function.

At the early time \(t = 0.1\tau\) (first column of Fig. 1), the van Hove function has moderately decayed compared to its initial condition. Over the course of time, cf. the results for \(0.3\tau\) (middle column) and \(0.6\tau\) (right column), the self part broadens and its height correspondingly decreases. The initial correlation hole in the distinct van Hove function is gradually being filled. Besides these transport processes, the initially pronounced oscillations at distances \(r \lesssim \sigma\) decay.

We demonstrate the agreement of adiabatic and dynamical density profiles in Fig. 1(c) by showing the MC simulation results obtained from equilibrium sampling of the adiabatic state, i.e., of the system in which the external potential \(V^{\text{self}}(r)\) acts on the (single) self particle and \(V^{\text{ad}}(r)\) acts on the remaining \(N - 1\) particles. (As \(N\) is large enough, we do not expect that finite size effects are relevant.) Apart from very small numerical artifacts, clearly the agreement of the dynamical and adiabatic density profiles is excellent. Hence, we trust the results for the adiabatic force densities (presented below) obtained via Eq. (2).

Besides the self and distinct parts, we also show results for the total van Hove function \(\rho = \rho_{\text{self}} + \rho_{\text{dist}}\) and the differential van Hove function \(\rho_\Delta = \rho_{\text{self}} - \rho_{\text{dist}}\) in Fig. 1(c). Clearly the spatial structuring of \(\rho\) is much reduced upon disregarding the self-distinct labeling. Nevertheless, as all particles in the system are ultimately identical, and the self-distinct labeling was introduced for mere bookkeeping purposes, one might wonder whether the physically most relevant phenomena are revealed or are rather hidden by the labeling.

In Fig. 1(d), we show results for the different contributions to the self force density. As a reference, we plot the ideal contribution \(-k_B T \nabla \rho_{\text{self}}\), which tends to spread the self peak in time. Here positive (negative) values of force fields and force densities indicate the outward (inward) direction. The adiabatic force density counteracts the ideal part and hence tends to stabilize the self density peak. The DFT results agree very satisfactorily with the MC data at all times considered. The superadiabatic self force density supports the effect of \(F^{\text{self}}\), but it has longer range and larger magnitude at later times. Except for a slight overestimation at time \(0.1\tau\), the PFT reproduces this effect very well and hence provides a mechanism for the slowing down of the dynamics.

The contributions to the distinct force density, shown in Fig. 1(e), show more complex, oscillatory behavior at both earlier times. The ideal force density is again directly related, via the spatial derivative, to the distinct density, shown in Fig. 1(c). The oscillations of the distinct density profiles are hence imprinted into the ideal force density and their effect is to homogenize the density. As is the case for the self part, the adiabatic force density counteracts this effect and hence tends to stabilize the density oscillations. The DFT results for \(F^{\text{dist}}\) are very satisfactory, with some underestimation inside the core \(r \lesssim \sigma\). The superadiabatic distinct force density has complex spatial features. It tends to slow down the decay of the spatial structure. At early times, the magnitude is smaller than that of \(F^{\text{dist}}\), but this relationship changes at later times where \(F^{\text{dist}}\) becomes dominant. Again, up to some deviations inside of the core, the PFT describes \(F^{\text{dist}}\) in very good agreement with the BD data.

In Fig. 1(f), we show results for the contributions to the differential force density \(G_{\text{int}}\) [as defined in Eq. (9)], which is relevant for the differential equation of motion, Eq. (8). Within the PFT, we can clearly identify that \(G_{\text{sup}}\) is solely due to the drag effect, i.e., the friction generated by the interflow of the self and distinct components. This result is relevant for the hard sphere dynamics at long times. Taking only the drag force as the dominant internal effect and
balancing it with ideal diffusion, the long time self diffusion coefficient follows as $D_L = k_B T/(\gamma + \rho_b C_{\text{drag}})$. Within this approximation, we obtain $D_L = 0.38\sigma^2/\tau$ (\(= 0.16 \mu m^2/s\) for $\sigma= 1 \mu m$ and $\tau= 2.33$ s as above), which is in very reasonable agreement with our bare simulation result of $D_L = 0.32\sigma^2/\tau$ (\(= 0.14 \mu m^2/s\)). In contrast, the motion of the total van Hove function, for which we show the relevant force fields in Fig. 1(g), is due to both compressional viscosity and structural forces, with both complex spatial and temporal behavior, which are well captured by the PFT. Crucially, while the details of the superadiabatic force fields vary depending on the type of dynamical situation considered, regarding these as arising from a kinematic functional Eq. (4) reveals their universal characteristics.

In conclusion, we have traced the mechanisms that govern the time evolution of the van Hove function for hard spheres by identifying three different and universal types of nonequilibrium force contributions, all of which have been shown previously to be relevant across a broad spectrum of nonequilibrium and driven systems. The forces are due to (i) drag of the tagged ("self") particle against the surrounding fluid of distinct particles, (ii) volume (or "bulk") viscosity due to the correlation shells undergoing compressional-expansional flow, and (iii) structural nonequilibrium effects that stabilize the spatial liquid structure against decay. The power functional approximation generates all three types of nonequilibrium force fields in quantitative agreement with Brownian dynamics computer simulation results. Our results hence demonstrate intimate interrelationships between equilibrium and nonequilibrium hard sphere properties.

It would be interesting to investigate in future work the relationship of our findings to Rosenfeld’s excess entropy scaling [53,54], as advanced by Truskett and his coworkers [55,56], to the nonequilibrium Ornstein-Zernike framework [57,58], and to the findings of Dyre and coworkers on universality across systems with different interparticle interaction potentials [59,60].

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