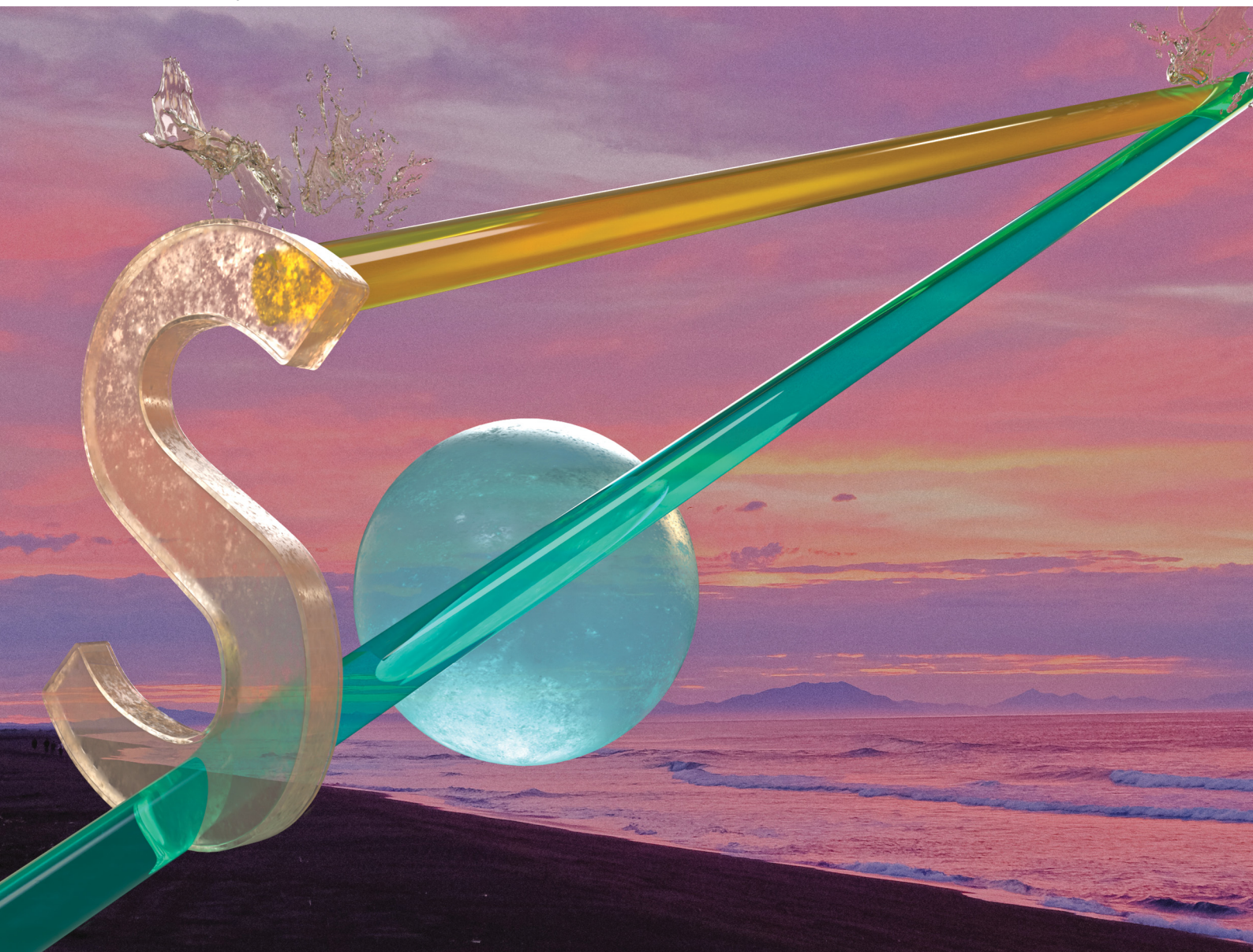


Soft Matter

rsc.li/soft-matter-journal



ISSN 1744-6848

PAPER


Hiroshi Frusawa

Non-hyperuniform metastable states around a disordered hyperuniform state of densely packed spheres: stochastic density functional theory at strong coupling



Cite this: *Soft Matter*, 2021,
17, 8810

Non-hyperuniform metastable states around a disordered hyperuniform state of densely packed spheres: stochastic density functional theory at strong coupling

Hiroshi Frusawa 

The disordered and hyperuniform structures of densely packed spheres near and at jamming are characterized by vanishing of long-wavelength density fluctuations, or equivalently by long-range power-law decay of the direct correlation function (DCF). We focus on previous simulation results that exhibit the degradation of hyperuniformity in jammed structures while maintaining the long-range nature of the DCF to a certain length scale. Here we demonstrate that the field-theoretic formulation of stochastic density functional theory is relevant to explore the degradation mechanism. The strong-coupling expansion method of stochastic density functional theory is developed to obtain the metastable chemical potential considering the intermittent fluctuations in dense packings. The metastable chemical potential yields the analytical form of the metastable DCF that has a short-range cutoff inside the sphere while retaining the long-range power-law behavior. It is confirmed that the metastable DCF provides the zero-wavevector limit of the structure factor in quantitative agreement with the previous simulation results of degraded hyperuniformity. We can also predict the emergence of soft modes localized at the particle scale by plugging this metastable DCF into the linearized Dean-Kawasaki equation, a stochastic density functional equation.

Received 16th July 2021,
Accepted 5th September 2021

DOI: 10.1039/d1sm01052b

rsc.li/soft-matter-journal

1. Introduction

Hyperuniformity is characterized by density fluctuations that decrease to zero at the longest scales.^{1–17} We have observed the hyperuniform states in a variety of complex soft matter systems, including foams, polymer blends, colloidal suspensions and biological tissues (see ref. 1 and 2 for reviews). It has also been found that non-crystalline materials with hyperuniformity have unique physical properties such as high-density transparency and isotropic filtration of elastic or electromagnetic waves.^{1–4} Consequently, considerable attention has been given to disordered hyperuniform materials fabricated at the micro- and nano-scales, because of their potential importance for applications in photonics, electronics, and structural components with novel properties.^{3,4}

Here we focus on computer glasses among the disordered hyperuniform systems. Recent methodological developments allow us to create computer glasses in an experimentally relevant regime,^{1–43} and yet the disordered hyperuniformity at jamming has not always been realized.^{1,2,38–43} The emergence of hyperuniformity depends on the preparation protocols,

partly because of the significantly long computational time that is required to determine the configurations near and at jamming.^{1,2,5–14}

On the one hand, some simulation studies have demonstrated the hyperuniformity in densely packed spheres: the structure factor $S(k)$ in a hyperuniform state exhibits a non-trivial linear dependence on the wavevector magnitude k in the low-wavevector range near and at jamming (*i.e.*, $S(k) \sim k$ ($k \geq 0$)), and the zero-wavevector limit of the structure factor $S(0)$ eventually vanishes at jamming.^{1,2,5–14} These results indicate not only the existence of long-range order, but also the complete suppression of density fluctuations over the system scale.

Meanwhile, other simulation studies near and at jamming^{38–43} provide the non-vanishing structure factor at the zero wavevector. The degradation of hyperuniformity is that either saturation or an upturn is observed for $S(k)$ at the lowest values of k , despite the linear relation above the crossover wavevector k_c .^{38–43}

$$S(k) \sim k \quad (k \geq k_c), \quad \lim_{k \rightarrow 0} S(k) \geq S(k_c). \quad (1)$$

It has also been demonstrated that $S(0)$ is weakly dependent on the density variation.^{38–43}

Laboratory of Statistical Physics, Kochi University of Technology, Tosa-Yamada, Kochi 782-8502, Japan. E-mail: frusawa.hiroshi@kochi-tech.ac.jp



The quantitative difference between the non-hyperuniform and hyperuniform states can be seen from the inverse of the zero-wavevector structure factors. While the non-vanishing values of $S(0)$ due to the incomplete linear-dependence of $S(k)$ are in the range of^{38–43}

$$10^2 < \frac{1}{S(0)} < 10^3, \quad (2)$$

the hyperuniform computer glasses have been characterized by inequality,

$$\frac{1}{S(0)} > 10^4, \quad (3)$$

irrespective of system details.^{1,2,5–14}

In terms of density–density correlation functions in real space, hyperuniformity is a kind of inverted critical phenomenon. It is among the critical phenomena in normal fluids that total correlation functions are long-ranged at critical points, accompanied by the diverging behaviors of density fluctuations and isothermal compressibility, while keeping the direct correlation function (DCF) short-ranged. In contrast, the inverted critical phenomenon is that the hyperuniform DCF is long-ranged in correspondence with the vanishing isothermal compressibility, despite the absence of long-range behavior for the total correlation function.^{1,2,5–14,17}

More concretely, the long-range behavior of the hyperuniform DCF $c(r)$ is described by the power-law as follows:^{1,2,5–7,13,17}

$$c(r) \sim \frac{1}{r^2} \quad (r > \sigma), \quad (4)$$

where $|\mathbf{r}| = r$ and σ denotes the sphere diameter. The long-range decay of $c(r)$ reads $c(k) \sim k^{-1}$ in the Fourier space, which is equivalent to the linear behavior $S(k) \sim k$ due to the following relation between $S(k)$ and $c(k)$:

$$\frac{1}{S(k)} = 1 - nc(k), \quad (5)$$

with n being the spatially averaged density of spheres. Furthermore, in disordered packings of hard spheres, the hyperuniform DCF for $r \leq \sigma$ satisfies another power-law,¹³

$$c(r) \sim \frac{1}{r} \quad (r \leq \sigma), \quad (6)$$

which is divergent at small r .

It follows from eqn (1) and (5) that the non-hyperuniform DCF $c(r)$ at jamming satisfies the scaling relation (4) over a finite range. In other words, the violation of hyperuniformity occurs while maintaining the long-range behavior to a length scale L_c : eqn (4) holds in the range of $\sigma < r \leq L_c$ ^{38–43} where

$$6 \leq \frac{L_c}{\sigma} \leq 10. \quad (7)$$

It is to be noted that the simulation results of hyperuniform systems are also likely to provide the finiteness represented by eqn (7),^{1,2,5–14} for we have computational difficulty obtaining the scaling behavior ($c(r) \sim r^{-2}$) over L_c from the structure

factor, irrespective of whether the computer glass is in a hyperuniform or non-hyperuniform state.

This common feature of the long-range behavior (eqn (4)) in the hyperuniform and non-hyperuniform DCFs raises the question of what causes the difference between eqn (2) and (3). Accordingly, it is the purpose of this study to reveal the underlying mechanism behind the difference between the emergence and degradation of hyperuniformity. To this end, we formulate an analytical form of the non-vanishing zero-wavevector structure factor that satisfies eqn (2) under the condition of eqn (7). A key ingredient of our formulation is the strong-coupling approximation of stochastic density functional theory (DFT)^{44–56} which can consider intermittent fluctuations while fixing the density field at a given distribution of dense packings near and at jamming.

The remainder of this paper consists of two parts. In the former part of Sections II–IV, the problem to be addressed is defined. Section II provides the theoretical background as to why stochastic DFT should be brought into the problem of fluctuation-induced non-hyperuniformity. In Section III, the basic formulation of stochastic DFT is presented, focusing on the definition of metastable states. Then, we find that stochastic DFT allows us to relate the metastable zero-wavevector structure factor $S^*(0)$ to the potential energy λ^* per particle, which will be referred to as the metastable chemical potential. In Section IV, the non-hyperuniform state on the target is specified using Table 1, which shows the classification list of hyperuniform and non-hyperuniform systems.

We see from the system specification that the non-hyperuniformity of our concern requires the short-range cutoff of the metastable DCF $c^*(r)$, as well as a drop in the long-ranged DCF for $r > L_c$. Our primary goal is to derive the short-range cutoff of the DCF by developing the strong-coupling approximation for stochastic DFT.

The latter part of this paper presents the results and discussion regarding the metastable DCF $c^*(r)$. Before entering the main results, Section V compares the stochastic DFT with the equilibrium DFT^{57–74} in terms of $S^*(0)$. It is demonstrated as a preliminary result that the resulting forms of $S^*(0)$ in the equilibrium and stochastic DFTs coincide with each other as far as the Gaussian approximation of stochastic DFT is adopted. Section VI provides the metastable DCF expressed by the Mayer-type function form, hence verifying the cutoff for the metastable DCF $c^*(r)$ inside the sphere. As a consequence,

Table 1 Four types of hyperuniform and non-hyperuniform systems. We investigate the type N1 of non-hyperuniform systems (see Sec. IV for the details)

System	Characterization of the DCF		
	Power-law decay	Magnitude at zero separation	Type
Hyperuniform	Complete	—	H1
	Incomplete	Divergent	H2
Non-hyperuniform	Incomplete	Finite	N1
	Absent	Finite	N2



Table 2 Comparison between the theoretical approaches and results (see also Sec. VIII C). We follow the notation of the type names given in Table 1

Theory		State description	Statics		Dynamics		
DFT type	Approximation	Type	μ or λ^*	DCF	Equation	Short-range	Long-range
Equilibrium	Ramakrishnan-Yussouff	H2	Eqn (31)	Eqn (44) or (56)	Eqn (29)	Frozen	Correlated
Stochastic	Gaussian	H2	Eqn (47)	Eqn (44) or (56)	Eqn (14) or (86)	Frozen	Correlated
Stochastic	Strong-coupling	N1	Eqn (51)	Eqn (52) or (54)	Eqn (14) or (86)	Soft	Correlated

we confirm that $S^*(0)$ satisfies relation (2), instead of eqn (3). In Section VII, the coupling constant γ to represent the strength of interactions is introduced using the hyperuniform DCF $c(r)$, and it is shown that the $1/\gamma$ expansion method becomes equivalent to the virial-type one at the strong coupling of $\gamma \gg 1$. Correspondingly, the interaction term in the metastable chemical potential λ^* is expressed by the above metastable DCF $c^*(r)$. In Section VIII, the stochastic density functional equation clarifies that the short-range cutoff of the metastable DCF $c^*(r)$ leads to the emergence of dynamic softening at the particle scale: the interaction-induced restoring force against density fluctuations around a metastable state vanishes within the scale of spherical diameter σ . The microscopic mechanism behind the soft modes is also discussed in connection with previous simulation results. Furthermore, both Fig. 3 and Table 2 summarize the static and dynamic results for comparing equilibrium DFT, stochastic DFT in the Gaussian approximation, and stochastic DFT in the strong-coupling approximation. The final remarks are given in Section IX.

II. A theoretical background of stochastic DFT

This section is intended to provide a brief overview of theoretical approaches to jammed structures for explaining the relevance of stochastic DFT^{44–56} to the degradation of hyperuniformity.

A. Marginal stability and the free energy landscape

There have been two conceptual approaches to address various issues on computer glasses, including the structure factors that vary depending on the protocols used.^{1,2,5–14,16} One is the ensemble approach to investigate physically relevant packings based on the packing protocol selected. The other method is the geometric-structure approach for the quantitative characterization of single-packing configurations to enumerate and classify the jammed structures.

On the one hand, the ensemble approach has involved the problem that the protocol-dependency of the occurrence frequency of jammed configurations leads to the ambiguity of weighing jammed states.^{1,2,5–17} Recently, however, the protocol-dependency problem is theoretically tackled: the canonical ensemble method is developed for a large number of allowed configurations to resolve the configuration realizability issue.¹⁶

The geometric-structure study, on the other hand, has distinguished three types of jamming for densely packed spheres:^{1,2,11} local, collective and strict jamming. These types of jamming are hierarchical in that local and collective

jammings are prerequisites for collective and strict ones, respectively, as follows: (i) in locally jammed states, a particle cannot translate when the positions of all other spheres in the packing are fixed; (ii) in collectively jammed states, particles prevented from translating are further stable to uniform compression; (iii) strictly jammed packings are stable against both uniform and shear deformations.

The geometric-structure studies on various computer glasses have verified that the hyperuniformity emerges in either strictly or collectively jammed systems having isostaticity.^{1,2,8} Here the isostatic configuration provides a mean contact number $2d$ per particle with d being the spatial dimension, thereby enhancing the mechanical stability. To be noted, however, the mechanical rigidity of jammed packings is a necessary but not sufficient condition for hyperuniformity.

It has been conjectured that any strictly jammed saturated infinite packing of identical spheres is hyperuniform; the conjecture excludes the existence of rattlers or particles that are free to move in a confining cage, by definition of strictly jammed packings.^{1,2,5,12,17} Conversely, it depends on simulation methods and conditions whether dense packings other than the strictly jammed ones, including the isostatic and collectively jammed systems, are hyperuniform or not. The isostatic systems can be destabilized by cutting one particle contact unless disordered packings are strictly jammed. In other words, isostaticity is a critical factor in mechanical marginal stability.^{1,2,12,18–37,75–83}

Recent simulation results have demonstrated that thermal fluctuations in the marginal states are accompanied by the intermittent rearrangements of particles.^{18–37} As a consequence, the marginal systems become responsive to have low-frequency soft modes that are nonphononic and anharmonic. For instance, quasi-localized modes coupled to an elastic matrix create soft spots composed of tens to hundreds of particles undergoing displacements.^{18–37} The low-frequency soft modes exhibit similar behaviors, and the common features of marginal states have been related to the emergence of many local minima in the free-energy landscape.^{26,84–87}

The similarity in anharmonic vibrations suggests that the ensemble of configurations visited by the slow dynamics could reveal the characteristics of marginal stability associated with the free-energy landscape, even though possible configurations depend on a protocol adopted.^{1,2,5–14,16}

B. The free-energy density functional: comparison between stochastic and equilibrium DFTs

For assessing the applicability of density functional approaches to the free-energy landscape in glassy systems, let us compare



stochastic DFT^{44–56} and equilibrium DFT, or classical DFT conventionally used.^{57–74}

Equilibrium DFT,^{57–74} one of the ensemble approaches, has been found relevant to investigate the free-energy landscape.^{84–91} It has been demonstrated that metastable minima determined by equilibrium DFT are not only correlated with the appearance of two-step relaxation and divergence of relaxation time, but are also directly connected with dynamical heterogeneity.^{62–74} In equilibrium DFT, the metastable density profile $\rho^*(\mathbf{r})$ has been approximated by the superposition of narrow Gaussian density profiles centered around a set of points forming an aperiodic lattice. Equilibrium DFT has properly identified the metastable state of a liquid having an inhomogeneous and aperiodic density as a local minimum of the equilibrium free-energy functional with respect to the variation in the width parameter for the above-mentioned Gaussian distribution.^{62–74}

However, the violation of perfect hyperuniformity has been beyond the scope of equilibrium DFT. Recently, the following three scenarios of imperfections have been proposed for demonstrating the degradation of hyperuniformity both theoretically and numerically:¹⁵ (i) uncorrelated point defects, (ii) stochastic particle displacements that are spatially correlated, and (iii) thermal excitations. In this study, we focus on the second scenario (ii) that is related to intermittent particle rearrangements in a contact network, a set of bonds connecting particles which are in contact with each other (see ref. 18–22, 75 and 76 for reviews). The elastic nature of the contact network could be responsible for the above-mentioned second scenario of non-hyperuniformity, or the spatially correlated displacements occurring stochastically; this will be discussed in Sections VIII and IX, based on the results obtained herein.

From stochastic DFT,^{44–56} on the other hand, it is expected that the above-mentioned second scenario (*i.e.*, (ii) stochastic and spatially correlated displacements) could be described in terms of stochastic density dynamics. To see this, a brief review of stochastic DFT is given below.

Stochastic DFT has been used as one of the most powerful tools for describing slowly fluctuating and/or intermittent phenomena, such as glassy dynamics, nucleation or pattern formation of colloidal particles, dielectric relaxation of Brownian dipoles, and even tumor growth (see ref. 44 for a thorough review). The stochastic density functional equation, which has often been referred to as the Dean–Kawasaki equation,^{44,45} forms the basis of stochastic DFT. It has been shown in various systems that the Dean–Kawasaki equation successfully describes the stochastic evolution of the instantaneous microscopic density field of overdamped Brownian particles. Of great practical use is the Dean–Kawasaki equation linearized with respect to density fluctuations around various reference density distributions.^{44,50–52,54–56}

As seen below, stochastic DFT is formulated on the hybrid framework that combines equilibrium DFT and statistical field theory.^{54,55,92} The hybrid framework allows us to investigate the metastable states considering fluctuations as clarified below. In Section VIII, stochastic DFT will also shed light on the dynamical properties of non-hyperuniformity.

III. Basic formulation: stochastic DFT

This section shows that stochastic DFT^{44–56} is available to investigate the zero-wavevector limit of the structure factor $S^*(0)$ in a metastable state. It is not merely a review of the previous formulations,⁵⁴ but rather a revisit for making it clear that stochastic DFT serves as a systematic evaluation of $S^*(0)$: as demonstrated in Sections VI and VII, we can evaluate the extent to which $S^*(0)$ is altered to the stochastic fluctuations around a hyperuniform state in a systematic manner.

First, the constrained free-energy functional $\mathcal{A}[\rho]$ is represented by the hybrid form using the functional and configurational integrals (Section IIIA). Next, we introduce the non-equilibrium excess chemical potential appearing in the stochastic DFT equation (Section IIIB). Third, the metastable state is defined based on stochastic DFT (Section IIIC). Last, the metastable DCF $c^*(\mathbf{r})$ is generated from the metastable chemical potential λ^* , thereby yielding $S^*(0)$ expressed by $c^*(\mathbf{r})$ (Section IIID).

A. Constrained free-energy functional $\mathcal{A}[\rho]$ in connection with the Fokker–Planck equation

Let $\hat{\rho}_N(\mathbf{r}, t) = \sum_{i=1}^N \delta[\mathbf{r} - \mathbf{r}_i(t)]$ be the instantaneous microscopic density of an N -particle system where the position $\mathbf{r}_i(t)$ at time t represents an instantaneous location of the i -th particle. The corresponding distribution functional $P[\rho, t]$ of the density field $\rho(\mathbf{r}, t)$ is defined by

$$P[\rho, t] = \left\langle \prod_{\mathbf{r}} \delta[\hat{\rho}_N(\mathbf{r}, t) - \rho(\mathbf{r}, t)] \right\rangle, \quad (8)$$

where $\langle \mathcal{O} \rangle$ signifies the noise-averaging operation for \mathcal{O} in the overdamped dynamics.

As detailed in Appendix A, $P[\rho, t]$ satisfies the Fokker–Planck equation given by eqn (A1). It follows from the stationary condition $\partial P_{\text{st}}[\rho]/\partial t = 0$ on the Fokker–Planck equation that the distribution functional in a steady state, $P_{\text{st}}[\rho]$, is determined by the free-energy functional $\mathcal{A}[\rho]$ of a given density field $\rho(\mathbf{r}, t)$:

$$P_{\text{st}}[\rho] = \frac{e^{-\mathcal{A}[\rho]}}{\int \mathcal{D}\rho e^{-\mathcal{A}[\rho]}}. \quad (9)$$

We can evaluate the constrained free-energy functional $\mathcal{A}[\rho]$ by introducing a fluctuating potential field ϕ as follows (see Appendix A for the derivation of eqn (10)–(13)):

$$e^{-\mathcal{A}[\rho]} = \int \mathcal{D}\phi e^{-F[\rho, \phi]} \Delta[\rho], \quad (10)$$

where $\Delta[\rho]$ denotes the constraint due to the canonical ensemble:

$$\Delta[\rho] = \begin{cases} 1 & \left(\int d\mathbf{r} \rho(\mathbf{r}) = N \right) \\ 0 & \left(\int d\mathbf{r} \rho(\mathbf{r}) \neq N \right). \end{cases} \quad (11)$$

The functional $F[\rho, \phi]$ in the exponent of eqn (10) is defined using the grand potential as follows:



$$\begin{aligned}
e^{-F[\rho, \phi] + \int d\mathbf{r} \rho(\mathbf{r}) \mu} &= \text{Tr} \prod_i e^{\mu + i\phi(\mathbf{r}_i) - \psi_{\text{dft}}(\mathbf{r}_i)} \prod_{i,j} e^{-v(\mathbf{r}_i - \mathbf{r}_j)} \\
&\times e^{\int d\mathbf{r} \rho(\mathbf{r}) \{\psi_{\text{dft}}(\mathbf{r}) - i\phi(\mathbf{r})\}} \\
&= e^{-\Omega\{\psi_{\text{dft}} - i\phi\} + \int d\mathbf{r} \rho(\mathbf{r}) \{\psi_{\text{dft}}(\mathbf{r}) - i\phi(\mathbf{r})\}},
\end{aligned} \quad (12)$$

where $\text{Tr} \equiv \sum_{N=0}^{\infty} \frac{1}{N!} \int d\mathbf{r}_1 \cdots \int d\mathbf{r}_N$, μ denotes the equilibrium chemical potential, $v(\mathbf{r})$ is the original interaction potential including the hard sphere potential and Lennard-Jones potential, and $\Omega[\psi]$ is the grand potential in the presence of an external field $\psi(\mathbf{r})$. Here it is noted that all the energetic quantities used in this study (e.g., μ , $v(\mathbf{r})$ and $\Omega[\psi]$) are given in $k_B T$ -unit.

As clearly seen from Appendix A, the fluctuating potential field $\phi(\mathbf{r})$ in eqn (12) traces back to the auxiliary field for the Fourier transform of the Dirac delta functional $\prod_{\mathbf{r}} [\delta[\hat{\rho}_N(\mathbf{r}, t) - \rho(\mathbf{r}, t)]]$.^{54,55,92} Also, the functional $F[\rho, \phi \equiv 0]$ in the absence of the ϕ -field corresponds to the intrinsic Helmholtz free energy in the presence of the external field $\psi_{\text{dft}}(\mathbf{r})$. Therefore, the following relation holds:

$$\frac{\delta F[\rho, 0]}{\delta \rho(\mathbf{r})} = \mu - \psi_{\text{dft}}(\mathbf{r}), \quad (13)$$

according to equilibrium DFT.^{58–61}

B. Non-equilibrium excess chemical potential $\lambda_{\text{ex}}[\rho]$

The Fokker-Planck equation for $P[\rho, t]$ is equivalent to the stochastic DFT equation, or the Dean-Kawasaki equation,⁴⁴ which is given by

$$\frac{\partial \rho(\mathbf{r}, t)}{\partial t} = \nabla \cdot \mathcal{D} \rho \nabla \lambda_{\text{ex}}[\rho] + \zeta[\rho, \vec{\eta}], \quad (14)$$

$$\lambda_{\text{ex}}[\rho] = \frac{\delta \mathcal{A}[\rho]}{\delta \rho}, \quad (15)$$

where $\zeta[\rho, \vec{\eta}]$ can be expressed as $\zeta[\rho, \vec{\eta}] = -\nabla \cdot \sqrt{2\mathcal{D}\rho(\mathbf{r}, t)} \vec{\eta}(\mathbf{r}, t)$ using the bare diffusion constant \mathcal{D} and the vectorial white noise field $\vec{\eta}(\mathbf{x}, t)$ defined by the correlation $\langle \eta_i(\mathbf{r}, t) \eta_j(\mathbf{r}', t') \rangle = \delta_{ij} \delta(\mathbf{r} - \mathbf{r}') \delta(t - t')$, and $\lambda_{\text{ex}}[\rho]$ will be referred to as the non-equilibrium excess chemical potential.

Combining eqn (10) and (15), we have

$$\lambda_{\text{ex}}[\rho] = \frac{\int D\phi \frac{\delta F[\rho, \phi]}{\delta \rho} e^{-F[\rho, \phi]} \Delta[\rho]}{\int D\phi e^{-F[\rho, \phi]} \Delta[\rho]}, \quad (16)$$

which further reads

$$\begin{aligned}
\lambda_{\text{ex}}[\rho] &= \overline{\lambda_{\text{ex}}[\rho, \phi]} \\
&\equiv \frac{\int D\phi \lambda_{\text{ex}}[\rho, \phi] e^{-\Delta F_{\text{dft}}[\rho, \phi]}}{\int D\phi e^{-\Delta F_{\text{dft}}[\rho, \phi]}},
\end{aligned} \quad (17)$$

$$\lambda_{\text{ex}}[\rho, \phi] = \frac{\delta F[\rho, 0]}{\delta \rho} + \frac{\delta \Delta F_{\text{dft}}[\rho, \phi]}{\delta \rho} - \lambda_N, \quad (18)$$

where $\Delta F_{\text{dft}}[\rho, \phi]$ signifies the free-energy difference between $F[\rho, \phi]$ and $F[\rho, 0]$, and the Lagrange multiplier λ_N enforces the

number constraint $\int d\mathbf{r} \rho(\mathbf{r}) = N$ and is reduced to the chemical potential μ when considering equilibrium DFT (see Section IIIC).

In the Gaussian approximation of the ϕ -field, we have

$$\begin{aligned}
\Delta F_{\text{dft}}[\rho, \phi] &= F[\rho, \phi] - F[\rho, 0] \\
&= \frac{1}{2} \iint d\mathbf{r} d\mathbf{r}' \phi(\mathbf{r}) w^{-1}(\mathbf{r} - \mathbf{r}') \phi(\mathbf{r}'),
\end{aligned} \quad (19)$$

$$\frac{\delta^2 F[\rho, 0]}{\delta \rho(\mathbf{r}) \delta \rho(\mathbf{r}')} = \overline{\phi(\mathbf{r}) \phi(\mathbf{r}')} = w(\mathbf{r} - \mathbf{r}'), \quad (20)$$

using the density-density correlation function $w^{-1}(\mathbf{r} - \mathbf{r}')$ (see Appendix A6 for details). For the concrete representation of the above propagator $w(\mathbf{r})$, we define the DCF $c(\mathbf{r})$ and the total correlation function $h(\mathbf{r})$ based on equilibrium DFT. The propagator $w(\mathbf{r})$ is expressed as

$$w(\mathbf{r} - \mathbf{r}') = \frac{\delta(\mathbf{r} - \mathbf{r}')}{\rho(\mathbf{r})} - c(\mathbf{r} - \mathbf{r}'), \quad (21)$$

$$w^{-1}(\mathbf{r} - \mathbf{r}') = \rho(\mathbf{r}) \{\delta(\mathbf{r} - \mathbf{r}') + h(\mathbf{r} - \mathbf{r}') \rho(\mathbf{r}')\}. \quad (22)$$

Eqn (21) and (22) manifest that equilibrium DFT is incorporated into stochastic DFT.

It follows from eqn (16)–(18) that

$$\lambda_{\text{ex}}[\rho] = \frac{\delta F[\rho, 0]}{\delta \rho} + \frac{\overline{\delta \Delta F_{\text{dft}}[\rho, \phi]}}{\delta \rho} - \lambda_N, \quad (23)$$

giving

$$\nabla \lambda_{\text{ex}}[\rho] = \nabla \frac{\delta F[\rho, \phi]}{\delta \rho} = \nabla \frac{\delta F[\rho, 0]}{\delta \rho} + \nabla \frac{\overline{\delta \Delta F_{\text{dft}}[\rho, \phi]}}{\delta \rho} \quad (24)$$

because $|\nabla \lambda_N| = 0$. In Section V, we will evaluate the second term on the right-hand side (rhs) of eqn (23), $\overline{\delta \Delta F_{\text{dft}}[\rho, \phi]} / \delta \rho$, following the above-mentioned Gaussian approximation, whereas the strong-coupling approximation developed for the evaluation of $F[\rho, \phi]$ will be presented in Section VII.

C. Comparison with the deterministic DFT equation

It has been proved in various ways that the stochastic DFT equation (eqn (14)) is converted into the deterministic DFT equation⁴⁴ when neglecting the additional free-energy functional $\Delta F_{\text{dft}}[\rho, \phi]$; hence, eqn (24) reduces to

$$\nabla \lambda_{\text{ex}}[\rho] = \nabla \frac{\delta F[\rho, 0]}{\delta \rho} \quad (25)$$

when the last noise term on the rhs of eqn (14) disappears.

Going back to eqn (20) and (21), we find that the Ramakrishnan-Yussouff functional⁵⁷ of the intrinsic Helmholtz free energy $F[\rho, 0]$ is of the following form:



$$F[\rho, 0] = F[n, 0] - \frac{1}{2} \iint d\mathbf{r} d\mathbf{r}' \Delta\rho(\mathbf{r}) c(\mathbf{r} - \mathbf{r}') \Delta\rho(\mathbf{r}') + \Delta F_{\text{id}}[\rho],$$

$$F_{\text{id}}[\rho] = \int d\mathbf{r} \rho(\mathbf{r}) \{\ln \rho(\mathbf{r}) - 1\}, \quad (26)$$

where $\Delta\rho \equiv \rho - n$, $\Delta F_{\text{id}}[\rho] \equiv F_{\text{id}}[\rho] - F_{\text{id}}[n]$, and $n = N/V$ denotes the uniform mean density with V being the system volume. Combining eqn (13) and (26) provides

$$\mu = \ln \rho(\mathbf{r}) + \psi_{\text{dft}}(\mathbf{r}) - \int d\mathbf{r}' c(\mathbf{r} - \mathbf{r}') \Delta\rho(\mathbf{r}'), \quad (27)$$

which reads

$$\rho(\mathbf{r}) = e^{\mu - \psi_{\text{dft}}(\mathbf{r}) + \int d\mathbf{r}' c(\mathbf{r} - \mathbf{r}') \Delta\rho(\mathbf{r}')}, \quad (28)$$

stating that the conventional relation of equilibrium DFT for a prescribed density $\rho(\mathbf{r})$ is satisfied by adjusting the external potential $\psi_{\text{dft}}(\mathbf{r})$ (see also eqn (A7)).

We obtain from plugging eqn (26) into eqn (25) the deterministic DFT equation using the Ramakrishnan–Yussouff functional:

$$\begin{aligned} \frac{\partial \rho(\mathbf{r}, t)}{\partial t} &= \nabla \cdot \mathcal{D} \rho \nabla \frac{\delta F[\rho, 0]}{\delta \rho} \\ &= \mathcal{D} \nabla^2 \rho(\mathbf{r}, t) - \nabla \cdot \mathcal{D} \rho(\mathbf{r}, t) \int d\mathbf{r}' \nabla c(\mathbf{r} - \mathbf{r}') \Delta\rho(\mathbf{r}', t). \end{aligned} \quad (29)$$

Comparison between eqn (14) and (29), or between eqn (24) and (25), indicates the difference between the stochastic and deterministic DFT equations.

D. Defining metastable states based on stochastic DFT

Before considering the metastability condition for the stochastic DFT equation (eqn (14)), we connect the metastable distribution $\rho_{\text{dft}}^*(\mathbf{r})$ determined by equilibrium DFT with the deterministic DFT equation (eqn (29)). In the absence of the external field (*i.e.*, $\psi_{\text{dft}} \equiv 0$), eqn (13) reduces to the metastability condition for equilibrium DFT:

$$\left. \frac{\delta F[\rho, 0]}{\delta \rho(\mathbf{r})} \right|_{\rho=\rho_{\text{dft}}^*} = \mu, \quad (30)$$

where $F[\rho_{\text{dft}}^*, 0]$ becomes equal to the intrinsic Helmholtz free energy defined in equilibrium. Correspondingly, eqn (27) leads to

$$\mu = \ln \rho_{\text{dft}}^*(\mathbf{r}) - \int d\mathbf{r}' c(\mathbf{r} - \mathbf{r}') \Delta\rho_{\text{dft}}^*(\mathbf{r}'). \quad (31)$$

The non-equilibrium excess chemical potential $\lambda_{\text{ex}}[\rho]$ should disappear at ρ_{dft}^* :

$$\begin{aligned} \lambda_{\text{ex}}[\rho_{\text{dft}}^*] &= \left. \frac{\delta F[\rho, 0]}{\delta \rho} \right|_{\rho=\rho_{\text{dft}}^*} - \lambda_N \\ &= \mu - \lambda_N = 0, \end{aligned} \quad (32)$$

implying that the Lagrange multiplier λ_N is correctly identified with the equilibrium chemical potential μ at ρ_{dft}^* , as mentioned above.

The difference between the equilibrium and stochastic DFTs can be clearly seen from plugging eqn (30) into eqn (14) and (29). On the one hand, the deterministic DFT equation (eqn (29)) ensures that eqn (30) is a steady-state condition: we have $\partial \rho_{\text{dft}}^* / \partial t = 0$ because the rhs of eqn (29) vanishes due to $|\nabla \delta F[\rho_{\text{dft}}^*, 0] / \delta \rho_{\text{dft}}^*| = |\nabla \mu| = 0$. On the other hand, the stochastic DFT equation (eqn (14)) for $\rho_{\text{dft}}^*(\mathbf{r}, t)$ becomes

$$\frac{\partial \rho_{\text{dft}}^*(\mathbf{r}, t)}{\partial t} = \nabla \cdot \mathcal{D} \rho_{\text{dft}}^* \nabla \frac{\delta \Delta F_{\text{dft}}[\rho_{\text{dft}}^*, \phi]}{\delta \rho_{\text{dft}}^*} + \zeta[\rho_{\text{dft}}^*, \vec{\eta}], \quad (33)$$

revealing that, in general, $\rho_{\text{dft}}^*(\mathbf{r}, t)$ is not a steady-state distribution in terms of stochastic DFT.

Meanwhile, the metastability condition for the stochastic DFT equation (eqn (14)) is that the metastable excess chemical potential $\lambda_{\text{ex}}[\rho^*]$ given by eqn (23) does not necessarily vanish but has a spatially constant value λ_{ex}^* :

$$\left. \frac{\delta \mathcal{A}[\rho]}{\delta \rho} \right|_{\rho=\rho^*} = \lambda_{\text{ex}}^*. \quad (34)$$

The first term on the rhs of eqn (14) disappears when eqn (34) is satisfied, yielding

$$\left\langle \frac{\partial \rho^*(\mathbf{r}, t)}{\partial t} \right\rangle = \langle \zeta[\rho^*, \vec{\eta}] \rangle = 0, \quad (35)$$

on noise-averaging. A previous study based on stochastic thermodynamics has shown that the heat dissipated into the reservoir is negligible on average when satisfying eqn (34) or eqn (35).⁵⁴

In this study, we thus adopt the metastability condition (34) based on stochastic DFT, instead of eqn (30).

E. The zero-wavevector structure factor $S^*(0)$ in a metastable state defined by eqn (34)

It has been demonstrated near and at jamming that the structure factor $S^*(k)$ in a metastable state can be written as

$$S^*(k) = \frac{1}{N} \langle \rho^*(k) \rangle \langle \rho^*(-k) \rangle, \quad (36)$$

because the structure factor in a frozen state mainly arises from the configurational part which is associated with the averaged positions of arrested particles.⁴² Eqn (32), (34) and (36) imply that $S^*(0)$ is obtained from the metastable chemical potential,

$$\lambda^* = \lambda_{\text{ex}}^* + \lambda_N, \quad (37)$$

in a similar manner to equilibrium DFT as follows:

$$\begin{aligned} \frac{1}{S^*(0)} &= n \left. \frac{\delta \lambda^*}{\delta \rho^*} \right|_{\rho^*=n} \\ &= 1 - n \int d\mathbf{r} c^*(\mathbf{r}) \\ &= -c^*(0) - n \int_{r \geq \sigma} 4\pi r^2 dr c^*(\mathbf{r}) \end{aligned} \quad (38)$$



$$\begin{aligned}
 & -n \int_{r \geq \sigma} 4\pi r^2 dr c^*(\mathbf{r}) h^*(\mathbf{r}) \\
 & \approx -c^*(0) - n \int_{r \geq \sigma} 4\pi r^2 dr c^*(\mathbf{r}),
 \end{aligned} \quad (39)$$

where the metastable DCF $c^*(\mathbf{r})$ is defined by eqn (38) using the metastable chemical potential λ^* and the approximate expression given in the last line of eqn (39) is obtained from the Ornstein–Zernike equation regarding $c^*(\mathbf{r})$ at zero separation $r = 0$ (see Appendix B1 for detailed derivation).

Equilibrium DFT, on the other hand, provides the metastable density distribution $\rho_{\text{df}}^*(\mathbf{r})$ determined by eqn (30). It follows that the metastable structure factor $S^*(k)$ reads

$$S^*(k) = \frac{1}{N} \rho_{\text{df}}^*(k) \rho_{\text{df}}^*(-k), \quad (40)$$

with $\langle \rho^*(k) \rangle$ in eqn (36) being replaced by $\rho_{\text{df}}^*(k)$. Then, we obtain from eqn (26) and (30)

$$\begin{aligned}
 \frac{1}{S^*(0)} &= n \frac{\delta \mu}{\delta \rho_{\text{df}}^*} \Big|_{\rho_{\text{df}}^*=n} \\
 &= 1 - n \int d\mathbf{r} c(\mathbf{r}),
 \end{aligned} \quad (41)$$

confirming that the DCF $c(\mathbf{r})$ determines the zero-wavevector structure factor.

IV. Our aim: non-hyperuniform states on the target

Table 1 classifies the hyperuniform and non-hyperuniform systems into four types for clarifying the non-hyperuniform state to be addressed hereafter. The type H1 in Table 1 signifies the hyperuniform state without requirement for $c^*(0)$ because eqn (4) is completely satisfied (*i.e.*, $L_c \rightarrow \infty$).

Despite the finiteness of the long-range nature, eqn (39) still predicts that the hyperuniformity of type H2 is necessarily observed near and at jamming unless the zero-separation divergence of $c^*(0)$ is avoided. This is because $1/S^*(0)$ diverges due to either the long-range nature or the divergent behavior at zero separation, as found from combining eqn (4) and (39).

To summarize, there are two requirements on the non-hyperuniform DCF $c^*(\mathbf{r})$ of type N1 as follows:

(i) Finiteness of the long-range nature – the non-hyperuniformity requires a drop in the long-ranged DCF for $r > L_c$. Namely, the first requirement is that $c^*(\mathbf{r})$ must decay rapidly to zero for $r > L_c$;^{39–43} otherwise, the second term on the rhs of eqn (39) is divergent.

(ii) Short-range cutoff – as seen from the first term on the rhs of eqn (39), the metastable DCF at zero separation (*i.e.*, $c^*(0)$) must have a finite value even as the densely packed systems approach jamming, which is the second requirement.

Eqn (39) reveals that the zero-wavevector structure factor never vanishes without meeting both of the above requirements. Nevertheless, exclusive attention in previous studies^{39–43} has been paid to the former requirement, and the short-range cutoff of the

metastable DCF (the second requirement (ii)) remains to be investigated.

In reality, the zero-separation DCF tends to have an extremely large value near freezing in repulsive sphere systems; for instance, the Percus–Yevick approximation of hard sphere fluids provides⁹³

$$-k_B T c^*(0) = \frac{\partial P}{\partial n}, \quad (42)$$

suggesting the divergent behavior of $-c^*(0)$ in a frozen state.

Thus, we focus on the emergence of type N1 when investigating the degradation of hyperuniformity. To be more specific, we show theoretically that the non-hyperuniformity of type N1 satisfies

$$c^*(\mathbf{r}) \sim \begin{cases} C & (r = 0) \\ \frac{1}{r^2} & (\sigma < r \leq L_c), \end{cases} \quad (43)$$

though the hyperuniformity of type H2 is incorporated into equilibrium DFT as input:

$$c(\mathbf{r}) \sim \begin{cases} \frac{1}{r} & (0 \leq r \leq \sigma) \\ \frac{1}{r^2} & (\sigma < r \leq L_c), \end{cases} \quad (44)$$

where $c(\mathbf{r})$ is different from the completely hyperuniform DCF in that L_c is supposed to have a finite value. Following equilibrium DFT, eqn (41) and (44) lead to $S^*(0) \sim -c(0) > 10^4$ despite the finiteness of the long-range power-law decay, which is the hyperuniformity of type H2.

We are now ready to address the issues on the non-hyperuniformity of type N1. In what follows, we present a preliminary result obtained in the Gaussian approximation for comparing stochastic and equilibrium DFTs, and subsequently prove in the strong-coupling approximation of stochastic DFT that eqn (44) transforms to eqn (43) as a result of the ensemble average over the fluctuating ϕ -field (see also eqn (17)).

V. Gaussian approximation of stochastic DFT

In the first place, we investigate the free-energy functional difference between the stochastic and equilibrium DFTs when performing the Gaussian approximation given by eqn (19). In the Gaussian approximation, eqn (23) reduces to

$$\begin{aligned}
 \lambda[\rho] &\equiv \lambda_{\text{ex}}[\rho] + \lambda_N \\
 &= \frac{\delta F[\rho, 0]}{\delta \rho(\mathbf{r})} + \frac{1}{2} \frac{\delta}{\delta \rho} \iint d\mathbf{r} d\mathbf{r}' w^{-1}(\mathbf{r} - \mathbf{r}') \phi(\mathbf{r}) \phi(\mathbf{r}').
 \end{aligned} \quad (45)$$



As seen from Appendix A7 for details, we have

$$\begin{aligned} & \frac{\delta}{\delta\rho} \iint d\mathbf{r} d\mathbf{r}' w^{-1}(\mathbf{r}-\mathbf{r}') \phi(\mathbf{r}) \phi(\mathbf{r}') \\ &= \frac{\delta}{\delta\rho} \iint d\mathbf{r} d\mathbf{r}' w^{-1}(\mathbf{r}-\mathbf{r}') \int D\phi \phi(\mathbf{r}) \phi(\mathbf{r}') e^{-\Delta F_{\text{dn}}[\rho, \phi]} \\ & \approx c(0) - h(0). \end{aligned} \quad (46)$$

Combining eqn (26), (37) and (46), eqn (45) reads

$$\begin{aligned} \lambda^* &= \lambda[\rho^*] = \lambda_{\text{ex}}^* + \lambda_N \\ &= \ln \rho^*(\mathbf{r}) - \int d\mathbf{r}' c(\mathbf{r}-\mathbf{r}') \Delta \rho^*(\mathbf{r}') + \frac{1}{2} \{c(0) - h(0)\}, \end{aligned} \quad (47)$$

in a metastable state. We find from eqn (38) and (47)

$$\frac{1}{S^*(0)} = 1 - n \int d\mathbf{r} c(\mathbf{r}) \quad (48)$$

while neglecting $\delta c(0)/\delta\rho$, or the triplet DCF. Comparison between eqn (41) and (48) confirms that no degradation of hyperuniformity is induced by Gaussian potential fluctuations.

To see the correspondence with previous results, it is convenient to transform eqn (47) to

$$\rho^*(\mathbf{r}) = e^{\lambda^* + \int d\mathbf{r}' c(\mathbf{r}-\mathbf{r}') \Delta \rho^*(\mathbf{r}') - \frac{1}{2} \{c(0) - h(0)\}}. \quad (49)$$

Eqn (49) is, on the one hand, of the same form as the previous results obtained from the Gaussian approximation in various ways when $\lambda^* = \mu$.^{94,95} On the other hand, comparison between eqn (28) with $\psi_{\text{dft}} \equiv 0$ and eqn (49) indicates that eqn (49) is identical to the self-consistent equation of $\rho^*(\mathbf{r})$ conventionally used in equilibrium DFT when

$$\lambda^* = \mu + \frac{1}{2} \{c(0) - h(0)\} \quad (50)$$

when $\lambda_N = \mu$ and $\lambda_{\text{ex}} = \{c(0) - h(0)\}/2$. Eqn (50) reveals that stochastic fluctuations create an additional contribution, the second term on the rhs of eqn (50), to the equilibrium chemical potential μ .

VI. Main results and comparison with simulation results

To go beyond the Gaussian approximation, we need to explore an expansion method adequate for strongly correlated sphere systems near and at jamming. One candidate is the virial-type expansion that has proven to be applicable to inhomogeneous ionic fluids at strong coupling.⁹⁶ In the next section, we will verify that the virial-type expansion can apply also to the evaluation of $\lambda_{\text{ex}}[\rho]$ given by eqn (17), hence yielding the metastable DCF $c^*(\mathbf{r})$ other than $c(\mathbf{r})$.

In this section, the obtained form of the metastable DCF $c^*(\mathbf{r})$, which satisfies the relation (43), is presented in advance (Section VIA). Subsequently, the calculated value of $S^*(0)$ is compared with the simulation results (*i.e.*, eqn (2)) on the non-hyperuniform structure factor at jamming (Section VIB).

A. Typical behaviors of the metastable DCF $c^*(\mathbf{r})$

As proved in the next section, the development of the strong-coupling expansion method, or the $1/\gamma$ expansion method, allows us to find the following form of the metastable chemical potential λ^* :

$$\lambda^* = \ln \rho^*(\mathbf{r}) + \frac{w(0)}{2} - \int d\mathbf{r}' c^*(\mathbf{r}-\mathbf{r}') \rho^*(\mathbf{r}'), \quad (51)$$

$$-c^*(\mathbf{r}-\mathbf{r}') = 1 - e^{-v(\mathbf{r}-\mathbf{r}') - w(\mathbf{r}-\mathbf{r}')}. \quad (52)$$

Since the relation $w(0) \gg 1$ holds at jamming, eqn (52) leads to

$$-c^*(0) = 1, \quad (53)$$

regardless of the repulsive potential form of $v(\mathbf{r})$. Eqn (53) reveals that, in general, the metastable DCF $c^*(\mathbf{r})$ given by (52) satisfies the second requirement (or eqn (43)) for the non-hyperuniformity (see requirement (ii) in Section IV). Particularly for hard spheres, the resulting form (52) reads

$$-c^*(\mathbf{r}) = \begin{cases} 1 & (\tilde{r} \leq 1) \\ -c(\mathbf{r}) & (\tilde{r} \gg 1), \end{cases} \quad (54)$$

where $\tilde{r} \equiv r/\sigma$. Eqn (54) meets the above-mentioned non-hyperuniformity requirements given by eqn (43) with $\mathcal{C} = -1$.

While the short-range cutoff is seen in the third term on the rhs of eqn (51), the second term on the rhs of eqn (51) corresponds to the effective self-energy which is divergent due to the power-law behavior expressed by eqn (44). This implies that the effective self-energy term ($= w(0)/2$) offsets the decrease in the interaction contribution due to the short-range cutoff.

Thus, we have obtained various forms of the chemical potential given by eqn (31), (47) and (51) from the equilibrium DFT, the stochastic DFT in the Gaussian approximation, and the stochastic DFT in the strong-coupling approximation, respectively. The above discussions also suggest that the different results of the hyperuniform and non-hyperuniform chemical potentials (*i.e.*, eqn (31) and (51)) are compatible with each other in terms of absolute values.

In Fig. 1, comparison is made between the \tilde{r} -dependencies of $-c(\mathbf{r})$ and $-c^*(\mathbf{r})$ for the repulsive harmonic potential given by

$$v(\mathbf{r}) = \varepsilon(1 - \tilde{r})^2 \Theta(1 - \tilde{r}), \quad (55)$$

where ε controls the interaction strength and $\Theta(x)$ is the Heaviside step function. It is supposed in Fig. 1 that $-c(\mathbf{r})$ is of the following form:

$$-c(\mathbf{r}) = \begin{cases} \frac{\alpha}{\tilde{r}} & (\tilde{r} \leq 1) \\ \frac{\beta}{\tilde{r}^2} & \left(1 < \tilde{r} \leq \frac{L_c}{\sigma}\right) \\ 0 & \left(\tilde{r} > \frac{L_c}{\sigma}\right). \end{cases} \quad (56)$$

To be noted, eqn (56) does not include the delta function $-\frac{1}{4\phi} \delta(\tilde{r}-1)$ due to the isostaticity, a significant negative contribution to $-c(\mathbf{r})$ at $\tilde{r} = 1$.¹³



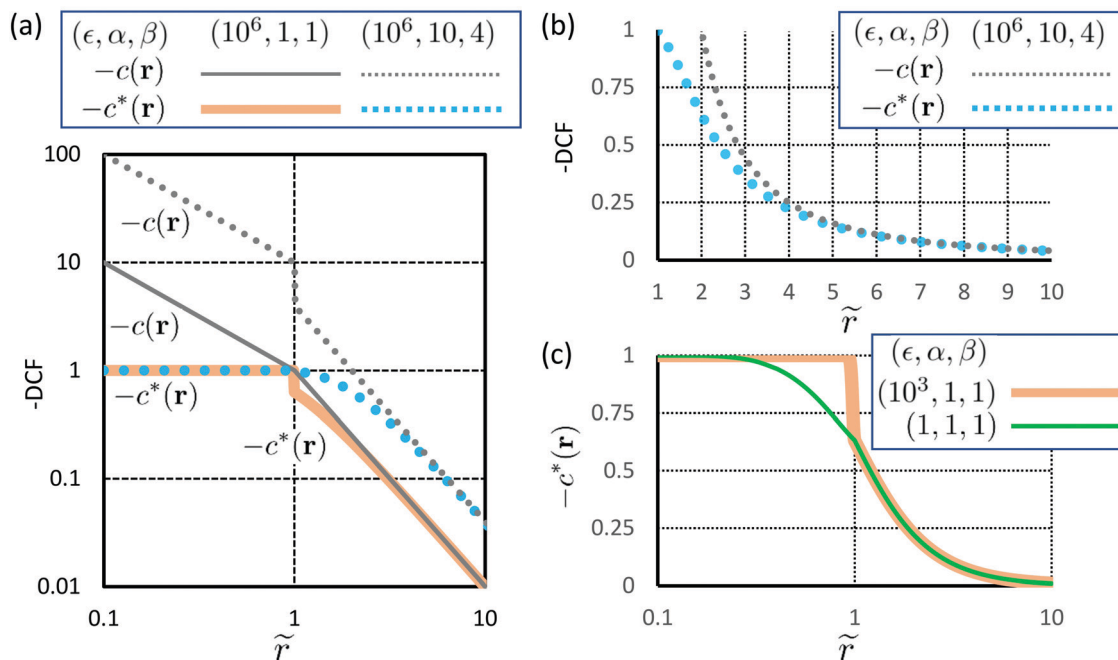


Fig. 1 Comparison between the metastable DCF $c^*(r)$ given by eqn (52) and the hyperuniform DCF $c(r)$ expressed by eqn (56) for the parameter sets of $(\epsilon, \alpha, \beta)$ as follows: while we need to fix two parameters, α and β , for representing the expression (56) of $c(r)$, it is necessary to set not only α and β , but also the parameter ϵ of the original interaction potential $v(r)$ given by eqn (55) for showing the obtained form (52) of $c^*(r)$. (a) A log-log plot of $c(r)$ and $c^*(r)$ which are depicted using the parameter sets as follows: $(\epsilon, \alpha, \beta) = (10^6, 10, 4)$ and $(10^6, 1, 1)$. (b) A linear plot for comparing $c(r)$ and $c^*(r)$ with the parameter set of $(\epsilon, \alpha) = (10^6, 10, 4)$ in more detail. (c) A semi-log plot of $c^*(r)$ when ϵ is decreased from 10^6 to either 10^3 or 1. The metastable DCF $c^*(r)$ is softened with the decrease of ϵ in eqn (55).

Previous simulation studies^{38–43} have indicated the parameter ranges of $\epsilon \geq 10^4$, $\alpha \sim 10^1$ and $0.1 \leq \beta \leq 10^1$ close to jamming. Correspondingly, we consider four sets of parameters in Fig. 1: $(\epsilon, \alpha, \beta) = (10^6, 10, 4)$, $(10^6, 1, 1)$, $(10^3, 1, 1)$ and $(1, 1, 1)$. In Fig. 1(a), the hyperuniform and metastable DCFs, $-c(r)$ and $-c^*(r)$, are depicted for two sets of parameters, $(\epsilon, \alpha, \beta) = (10^6, 10, 4)$ and $(10^6, 1, 1)$, on a log-log plot. We can see from Fig. 1(a) that the potential value of the metastable DCF saturates to unity irrespective of the short-range behavior of $-c(r)$, and that the short-range deviation of $-c^*(r)$ from $-c(r)$ is larger with the increase of α and β . A magnified view for $r \geq \sigma$ is shown in Fig. 1(b), allowing us to make a comparison between $-c(r)$ and $-c^*(r)$ for $(\epsilon, \alpha, \beta) = (10^6, 10, 4)$ in more detail. Fig. 1(b) shows that $-c^*(r)$ converges to $-c(r)$ for $r \gg \sigma$ even when there is an obvious difference in the DCFs at $r = 2\sigma$ between $-c(r = 2\sigma) = \beta/4$ and $-c^*(r = \sigma) = 1 - e^{-\beta/4}$ for $\beta = 4$. Fig. 1(c) compares the profiles of $-c^*(r)$ for $\epsilon = 10^3$ and 1 with α and β being the same value ($\alpha = \beta = 1$) on a semi-log plot. This indicates that the metastable DCF inside the sphere (*i.e.*, $-c^*(r)$ for $r \leq \sigma$) is not changed until the interaction strength represented by the parameter ϵ is reduced considerably (for instance, $\epsilon = 1$ in Fig. 1(c)) far from the jamming values of $\epsilon \geq 10^4$.

B. Comparison with simulation results given by eqn (2) and (7)

It follows from eqn (39), (52), (55) and (56) that the approximate form of the zero-wavevector structure factor $S^*(0)$ is determined

by both the volume fraction f_v of packed spheres and the cutoff length L_c :

$$\frac{1}{S^*(0)} \approx 24f_v\beta \left(\frac{L_c}{\sigma} \right) \quad (57)$$

for $L_c/\sigma \gg 1$; see Appendix B2 for detailed derivation. The first choice to investigate the type-N1 non-hyperuniformity at jamming is to set $L_c/\sigma = 10$ and $f_v = 0.65$, according to the previous simulation results^{38–43} of non-hyperuniform harmonic-core sphere systems. Eqn (57) then becomes $1/S^*(0) = 156\beta$, implying that relation (2) applies to the metastable structure factor:

$$10^2 < \frac{1}{S^*(0)} < 10^3, \quad (58)$$

with $\beta \sim \mathcal{O}[10^b]$ being in a reasonable range of $0 \leq b < 1$.

For validation of the above evaluation, Fig. 2 provides the dependences of $1/S^*(0)$ on L_c/σ in the range of eqn (7) for $\beta = 1, 4$ and 10 with $f_v = 0.65$ being used as before. As seen from Fig. 2, a comparison between the precise result (see eqn (B9) in Appendix B2) and the approximate expression (57) shows that eqn (57) is an acceptable approximation. The precise results depicted by solid lines in Fig. 2 further verify the relation (58) for $1 \leq \beta \leq 10$ in the range of eqn (7) for L_c/σ . Thus, we find that the metastable DCF $c^*(r)$ given by eqn (52), one of the main results in this study, quantitatively explains the previous simulation results on the non-hyperuniformity of type N1.

It is also suggested by Fig. 2 that $\beta \sim 10^{-1}$ leads to $1/S^*(0) < 10^2$ as long as the cutoff of $-c^*(0) = 1$ holds. This result appears



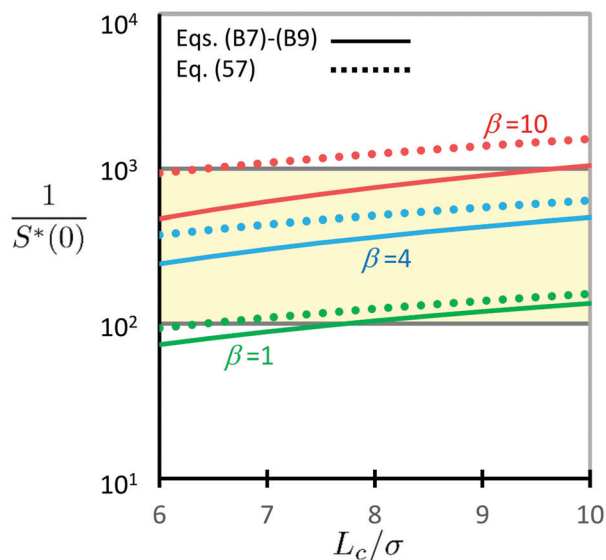


Fig. 2 The last expression in eqn (39) can be calculated analytically when using eqn (52) and (56). The three solid lines depict the analytical result (B9) with eqn (B7) and (B8), or the precise results of the zero-wavevector structure factor $S^*(0)$, for $\beta = 1, 4$ and 10 at $f_v = 0.65$. For comparison, the dotted lines representing the approximate form (57) are also drawn for the same parameter sets: $\beta = 1, 4$ and 10 at $f_v = 0.65$. The yellow area corresponds to the non-hyperuniform range of $1/S^*(0)$ which is given by either eqn (2) or eqn (58).

to contradict previous simulation results¹³ in hyperuniform hard sphere systems where not only the small value of $\beta \sim 10^{-1}$ but also the existence of L_c in the range of eqn (7) have been found. At the same time, however, the divergent relation $-c(\mathbf{r}) \approx 10/\tilde{r}$ ($\tilde{r} < 1$) has been verified for the present hyperuniform hard sphere systems.¹³ Accordingly, the divergent behavior of the hyperuniform DCF $-c(0)$ at zero-separation ensures the hyperuniform relation (3): the relation, $1/S(0) \approx -c(0) > 10^4$, holds even when $\beta \sim 10^{-1}$ and $L_c/\sigma \sim 10^1$, which is exactly the hyperuniform state of the type-H2 in Table 1.

VII. Verification of the main result given by eqn (51) and (52) in the strong-coupling approximation

There are three steps to verify both the metastable chemical potential λ^* and the DCF $c^*(\mathbf{r})$ given by eqn (51) and (52), respectively. First, we define the coupling constant γ and present the free-energy functionals rescaled by γ , suggesting the validity of the strong-coupling expansion method, or the density-expansion method at high density (Section VIIA). Second, the non-equilibrium chemical potential $\lambda[\rho]$ defined by eqn (16)–(18) is calculated for non-interacting spheres at strong coupling (Section VIIB). Third, we connect the $1/\gamma$ expansion, which is equivalent to the density expansion (or the fugacity expansion⁹⁶), with the virial-type term expressed by the Mayer function, thereby proving eqn (51) and (52) (Section VIIC).

A. Rescaled free-energy functionals

We introduce the rescaled propagator $\tilde{w}(\mathbf{r})$ using a coupling constant γ :

$$\tilde{w}(\mathbf{r}) = \frac{w(\mathbf{r})}{\gamma^2}, \quad (59)$$

$$\gamma = e^{\frac{w(0)}{2}}.$$

Since we consider the type-H2 hyperuniform systems as mentioned before, it is found from eqn (21) and (44) that the coupling constant γ is approximated by $\gamma \approx e^{-c(0)/2}$ and becomes extremely large near and at jamming.

We aim to develop the $1/\gamma$ expansion method at strong coupling ($\gamma \gg 1$), provided that γ is extremely large but is finite. In the next subsection, we will show that the virial-type expansion method, the density-expansion method, can be regarded as the $1/\gamma$ expansion method. Before proceeding, we see the γ -dependencies of functionals based on the following criteria:

Criterion 1: $\Delta F_{\text{df}}[\tilde{\rho}, \tilde{\phi}] \sim \gamma^0$,

Criterion 2: $\int d\mathbf{r}' \tilde{w}(\mathbf{r} - \mathbf{r}') \tilde{w}^{-1}(\mathbf{r}' - \mathbf{r}'') = \delta(\mathbf{r} - \mathbf{r}'')$.

Criterion 1 allows us to discern the perturbative terms at strong coupling, in comparison with the rescaled functional $\Delta F_{\text{df}}[\tilde{\rho}, \tilde{\phi}] \sim \gamma^0$, whereas criterion 2 is equivalent to the Ornstein–Zernike equation^{58–61} for rescaled correlation functions, $\tilde{c}(\mathbf{r})$ and $\tilde{h}(\mathbf{r})$, that should be defined to satisfy

$$\tilde{w}(\mathbf{r} - \mathbf{r}') = \frac{\delta(\mathbf{r} - \mathbf{r}')}{\tilde{\rho}(\mathbf{r})} - \tilde{c}(\mathbf{r} - \mathbf{r}'), \quad (60)$$

$$\tilde{w}^{-1}(\mathbf{r} - \mathbf{r}') = \tilde{\rho}(\mathbf{r}) \{ \delta(\mathbf{r} - \mathbf{r}') + \tilde{h}(\mathbf{r} - \mathbf{r}') \tilde{\rho}(\mathbf{r}') \}, \quad (61)$$

consistently with the original definitions given by eqn (21) and (22).

It is found from eqn (19) and (59) that criterion 1 imposes potential rescaling as follows:

$$\phi(\mathbf{r}) = \gamma \tilde{\phi}(\mathbf{r}). \quad (62)$$

On the other hand, it follows from criterion 2, or eqn (59) to (61), that the correlation functions and the density field are necessarily rescaled as

$$c(\mathbf{r}) = \gamma^2 \tilde{c}(\mathbf{r}),$$

$$h(\mathbf{r}) = \gamma^2 \tilde{h}(\mathbf{r}), \quad (63)$$

$$\rho(\mathbf{r}) = \frac{\tilde{\rho}(\mathbf{r})}{\gamma^2},$$

to satisfy criterion 2.

Combining eqn (59) and (62) transforms eqn (19) to

$$\Delta F_{\text{df}}[\tilde{\rho}, \tilde{\phi}] = \frac{1}{2} \iint d\mathbf{r} d\mathbf{r}' \tilde{\phi}(\mathbf{r}) \tilde{w}^{-1}(\mathbf{r} - \mathbf{r}') \tilde{\phi}(\mathbf{r}'), \quad (64)$$

meeting criterion 1. Meanwhile, the rescaled form $F[\tilde{\rho}, 0]$ of the Ramakrishnan–Yussouff free energy functional (26) is



$$F[\tilde{\rho}, 0] = F[\tilde{n}, 0] - \frac{1}{2\gamma^2} \iint d\mathbf{r} d\mathbf{r}' \tilde{\Delta}\rho(\mathbf{r}) \tilde{c}(\mathbf{r} - \mathbf{r}') \tilde{\Delta}\rho(\mathbf{r}') + \Delta F_{\text{id}}[\tilde{\rho}],$$

$$F_{\text{id}}[\tilde{\rho}] = \int d\mathbf{r} \frac{\tilde{\rho}(\mathbf{r})}{\gamma^2} \left\{ \ln \frac{\tilde{\rho}(\mathbf{r})}{\gamma^2} - 1 \right\}, \quad (65)$$

where $\tilde{\Delta}\rho \equiv \tilde{\rho} - \tilde{n}$ and $n \equiv \tilde{n}/\gamma^2$. Comparison between the rescaled functionals, eqn (64) and (65), suggests that the $\tilde{\rho}$ -dependent terms can be treated perturbatively at strong coupling ($\gamma \gg 1$).

B. The non-equilibrium chemical potential of non-interacting spheres at strong coupling

Going back to eqn (12), we develop the $1/\gamma$ expansion method. It is found from eqn (28) that

$$e^{\mu + i\phi(\mathbf{r}_i) - \psi_{\text{dft}}(\mathbf{r}_i)} = \rho(\mathbf{r}) e^{\Delta\psi(\mathbf{r}) + i\phi(\mathbf{r})}, \quad (66)$$

$$\Delta\psi(\mathbf{r}) = - \int d\mathbf{r}' c(\mathbf{r} - \mathbf{r}') \Delta\rho(\mathbf{r}').$$

Also, we shift the fluctuating-potential field from ϕ to φ such that

$$\frac{w(0)}{2} + i\varphi(\mathbf{r}) = \Delta\psi(\mathbf{r}) + i\phi(\mathbf{r}), \quad (67)$$

whose rescaled form is

$$\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r}) = \tilde{\Delta}\psi(\mathbf{r}) + i\gamma \tilde{\phi}(\mathbf{r}), \quad (68)$$

due to the relations (62) and (63). Substituting eqn (68) into eqn (66), we have the rescaled form,

$$e^{\mu + i\phi(\mathbf{r}_i) - \psi_{\text{dft}}(\mathbf{r}_i)} = \rho(\mathbf{r}) e^{\frac{w(0)}{2} + i\varphi(\mathbf{r})} = \frac{\tilde{\rho}(\mathbf{r})}{\gamma} e^{i\gamma \tilde{\varphi}(\mathbf{r})}, \quad (69)$$

considering that $\rho(\mathbf{r}) e^{\frac{w(0)}{2}} = \gamma \rho(\mathbf{r}) = \tilde{\rho}(\mathbf{r})/\gamma$. Moreover, eqn (28), (67) and (69) are arranged to give

$$\begin{aligned} & - \int d\mathbf{r} \rho(\mathbf{r}) \{ \psi_{\text{dft}}(\mathbf{r}) - i\phi(\mathbf{r}) \} \\ &= \int d\mathbf{r} \rho(\mathbf{r}) \{ \ln \rho(\mathbf{r}) + \Delta\psi(\mathbf{r}) + i\phi(\mathbf{r}) - \mu \} \\ &= \int d\mathbf{r} \rho(\mathbf{r}) \left\{ \ln \rho(\mathbf{r}) + \frac{w(0)}{2} + i\varphi(\mathbf{r}) - \mu \right\} \\ &\equiv F_0[\rho, \varphi] - \int d\mathbf{r} \rho(\mathbf{r}) \mu. \end{aligned} \quad (70)$$

Combining eqn (66)–(70), eqn (12) reads

$$\begin{aligned} e^{-F[\rho, \phi = \varphi + i\Delta\psi] + \int d\mathbf{r} \rho(\mathbf{r}) \mu} &= e^{-F_0[\rho, \varphi] + \int d\mathbf{r} \rho(\mathbf{r}) \mu} \\ &\times \text{Tr} \prod_i \frac{\tilde{\rho}(\mathbf{r}_i)}{\gamma} e^{i\gamma \tilde{\varphi}(\mathbf{r}_i)} \prod_{i,j} e^{-v(\mathbf{r}_i - \mathbf{r}_j)}, \end{aligned} \quad (71)$$

which is the functional to be evaluated using the $1/\gamma$ expansion method at strong coupling, $\gamma \gg 1$.

Let us see the non-equilibrium chemical potential $\lambda[\rho]$ in a reference system of non-interacting spheres, prior to formulating the strong-coupling approximation of eqn (71). In the absence of the interaction potential $v(\mathbf{r}_i - \mathbf{r}_j)$, eqn (71) is exactly reduced to the ideal free-energy functional for the non-interacting system:

$$\begin{aligned} F_{\text{non}}[\rho, \varphi] &= F_0[\rho, \varphi] - \int d\mathbf{r} \frac{\tilde{\rho}(\mathbf{r})}{\gamma} e^{i\gamma \tilde{\varphi}(\mathbf{r})} \\ &= \int d\mathbf{r} \tilde{\rho}(\mathbf{r}) \left\{ \frac{1}{\gamma^2} \ln \frac{\tilde{\rho}(\mathbf{r})}{\gamma^2} + \frac{\tilde{w}(0)}{2} + \frac{i\tilde{\varphi}(\mathbf{r})}{\gamma} - \frac{e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})}}{\gamma^2} \right\} \\ &= F_0[\tilde{\rho}, \tilde{\varphi}] - \frac{1}{\gamma^2} \int d\mathbf{r} \tilde{\rho}(\mathbf{r}) e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})}. \end{aligned} \quad (72)$$

It follows that

$$\begin{aligned} \lambda_{\text{non}}[\tilde{\rho}, \tilde{\varphi}] &\equiv \frac{\delta F_{\text{non}}[\rho, \varphi]}{\delta \rho} \\ &= \gamma^2 \frac{\delta F_0[\tilde{\rho}, \tilde{\varphi}]}{\delta \tilde{\rho}} - \frac{\delta}{\delta \tilde{\rho}(\mathbf{r})} \left\{ \int d\mathbf{r} \tilde{\rho}(\mathbf{r}) e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})} \right\} \\ &= \ln \frac{\tilde{\rho}(\mathbf{r})}{\gamma^2} + 1 + \frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r}) \\ &\quad - e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})} - \tilde{\rho}(\mathbf{r}) \frac{\delta e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})}}{\delta \tilde{\rho}(\mathbf{r})}. \end{aligned} \quad (73)$$

We need to perform the average of $\lambda_{\text{non}}[\tilde{\rho}, \tilde{\varphi}]$ over the ϕ -field based on the original definition in addition to relation (68). In the strong-coupling approximation, we obtain from eqn (73)

$$\lambda_{\text{non}}[\rho] = \overline{\lambda_{\text{non}}[\tilde{\rho}, \tilde{\varphi}]} = \ln \rho(\mathbf{r}) + \frac{w(0)}{2}, \quad (74)$$

which corresponds to the non-equilibrium chemical potential $\lambda_{\text{non}}[\rho]$ of non-interacting spheres; see Appendix C2 for the detailed derivation of eqn (74). Eqn (74) implies that

$$F_{\text{non}}[\rho] = F_{\text{id}}[\rho] + \int d\mathbf{r} \rho(\mathbf{r}) \frac{w(0)}{2}. \quad (75)$$

C. Connecting the $1/\gamma$ expansion with the virial-type expansion: derivation scheme of eqn (51) and (52)

In the strong-coupling approximation, the long-range correlations of the shifted fluctuating potential $\varphi(\mathbf{r})$ are maintained:

$$\overline{\varphi(\mathbf{r})\varphi(\mathbf{r}')} = w(\mathbf{r} - \mathbf{r}'), \quad (76)$$

as well as relation (20) for the fluctuating ϕ -potential (see the derivation of eqn (C15) in Appendix C1). Eqn (76) implies that

$$\left| \frac{\tilde{\rho}(\mathbf{r}_i)}{\gamma} e^{i\gamma \tilde{\varphi}(\mathbf{r}_i)} \right| = \frac{\tilde{\rho}(\mathbf{r}_i)}{\gamma}. \quad (77)$$



Hence, the $1/\gamma$ expansion becomes equivalent to the following density expansion (or the fugacity expansion⁹⁶):

$$\begin{aligned} & \text{Tr} \prod_i \frac{\tilde{\rho}(\mathbf{r}_i)}{\gamma} e^{i\gamma\tilde{\phi}(\mathbf{r}_i)} \prod_{i,j} e^{-v(\mathbf{r}_i-\mathbf{r}_j)} \\ &= \sum_{N=0}^{\infty} \frac{1}{N!} \int d\mathbf{r}_1 \cdots \int d\mathbf{r}_N \prod_{i=1}^N \frac{\tilde{\rho}(\mathbf{r}_i)}{\gamma} e^{i\gamma\tilde{\phi}(\mathbf{r}_i)} \prod_{i,j} e^{-v(\mathbf{r}_i-\mathbf{r}_j)} \\ &\approx 1 + \frac{1}{\gamma} \left\{ \int d\mathbf{r} \tilde{\rho}(\mathbf{r}) e^{i\gamma\tilde{\phi}(\mathbf{r})} \right. \\ &\quad \left. + \frac{1}{2\gamma} \int d\mathbf{r}_1 d\mathbf{r}_2 \tilde{\rho}(\mathbf{r}_1) \tilde{\rho}(\mathbf{r}_2) e^{-v(\mathbf{r}_1-\mathbf{r}_2) + i\gamma\tilde{\phi}(\mathbf{r}_1) + i\gamma\tilde{\phi}(\mathbf{r}_2)} \right\} \\ &\equiv 1 + \frac{1}{\gamma} U[\tilde{\rho}, \tilde{\phi}], \end{aligned} \quad (78)$$

where

$$\begin{aligned} U[\tilde{\rho}, \tilde{\phi}] &= \int d\mathbf{r} \tilde{\rho}(\mathbf{r}) e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \hat{\rho}_1(\mathbf{s})} \\ &\quad + \frac{1}{2\gamma} \int d\mathbf{r}_1 d\mathbf{r}_2 \tilde{\rho}(\mathbf{r}_1) \tilde{\rho}(\mathbf{r}_2) e^{-v(\mathbf{r}_1-\mathbf{r}_2) + i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \hat{\rho}_2(\mathbf{s})}. \end{aligned} \quad (79)$$

In eqn (79), we have introduced instantaneous one- and two-particle densities, $\hat{\rho}_1(\mathbf{s}) = \delta(\mathbf{s} - \mathbf{r})$ and $\hat{\rho}_2(\mathbf{s}) = \sum_{i=1}^2 \delta(\mathbf{r} - \mathbf{r}_i)$, for making a distinction between the first and the second terms on the rhs of eqn (79).

Combining eqn (71), (72) and (78) provides

$$F[\tilde{\rho}, \tilde{\phi}] = F_0[\tilde{\rho}, \tilde{\phi}] - \ln \left(1 + \frac{1}{\gamma} U[\tilde{\rho}, \tilde{\phi}] \right), \quad (80)$$

and we define the non-equilibrium chemical potential difference $\Delta\lambda[\rho]$ due to the addition of the interaction potential $v(\mathbf{r})$ as follows:

$$\Delta\lambda[\rho] = \gamma^2 \frac{\delta F[\tilde{\rho}, \tilde{\phi}]}{\delta \tilde{\rho}(\mathbf{r})} - \lambda_{\text{non}}[\rho]. \quad (81)$$

It follows from eqn (72), (74), (76), (79) and (80) that the strong-coupling approximation of eqn (81) leads to

$$\begin{aligned} \Delta\lambda[\rho] &= 1 - \frac{\gamma \delta U[\tilde{\rho}, \tilde{\phi}] / \delta \tilde{\rho}(\mathbf{r})}{1 + U[\tilde{\rho}, \tilde{\phi}] / \gamma} \\ &= 1 - \frac{\gamma \delta U[\tilde{\rho}, \tilde{\phi}]}{\delta \tilde{\rho}(\mathbf{r})} + \frac{\delta U[\tilde{\rho}, \tilde{\phi}]}{\delta \tilde{\rho}(\mathbf{r})} U[\tilde{\rho}, \tilde{\phi}] + \mathcal{O}[\gamma^{-1}] \\ &= 1 - \gamma e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \hat{\rho}_1(\mathbf{s})} \\ &\quad - \int d\mathbf{r}_2 \tilde{\rho}(\mathbf{r}_2) e^{-v(\mathbf{r}_1-\mathbf{r}_2)} e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \hat{\rho}_2(\mathbf{s})} \\ &\quad + \int d\mathbf{r}' \tilde{\rho}(\mathbf{r}') e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \{ \hat{\rho}_1(\mathbf{s}) + \hat{\rho}'_1(\mathbf{s}) \}} + \mathcal{O}[\gamma^{-1}] \\ &= \int d\mathbf{r}' \left\{ 1 - e^{-v(\mathbf{r}-\mathbf{r}') - w(\mathbf{r}-\mathbf{r}')} \right\} \rho(\mathbf{r}') + \mathcal{O}[\gamma^{-2}], \end{aligned} \quad (82)$$

where it is noted that the instantaneous one-particle densities, $\hat{\rho}_1(\mathbf{s}) = \delta(\mathbf{s} - \mathbf{r})$ and $\hat{\rho}'_1(\mathbf{s}) = \delta(\mathbf{s} - \mathbf{r}')$, are unable to coexist at the same

time by definition; see Appendix C for detailed and more precise discussions regarding the derivation of eqn (82). We obtain from eqn (74) and (82)

$$\begin{aligned} \lambda[\rho] &= \lambda_{\text{non}}[\rho] + \Delta\lambda[\rho] \\ &= \ln \rho(\mathbf{r}) + \frac{w(0)}{2} + \int d\mathbf{r}' \left\{ 1 - e^{-v(\mathbf{r}-\mathbf{r}') - w(\mathbf{r}-\mathbf{r}')} \right\} \rho(\mathbf{r}'). \end{aligned} \quad (83)$$

The main result given by eqn (51) and (52) is thus verified.

VIII. Discussions

In this section, we aim to gain insight into the short-range cutoff of the metastable DCF $c^*(\mathbf{r})$ from dynamic aspects. We consider a fluctuating displacement field $\mathbf{u}(\mathbf{r}, t)$ which is related to the density difference, $\nu(\mathbf{r}, t) = \rho(\mathbf{r}, t) - \rho^*(\mathbf{r})$. Since the fluctuating density field $\nu(\mathbf{r}, t)$ obeys the linearized Dean-Kawasaki equation of stochastic DFT,^{44,50–52,54–56} the short-range dynamics of $\mathbf{u}(\mathbf{r}, t)$ can be inferred from the ν -field dynamics. First, we will see that the short-range cutoff of the metastable DCF implies the disappearance of the interaction-induced restoring force against the fluctuating density field $\nu(\mathbf{r}, t)$ (Section VIIIA). Next, the connection of the short-range softening with anharmonic soft modes will be discussed in terms of \mathbf{u} -field dynamics (Section VIIIB). Last, we summarize the results presented so far using Table 2 (Section VIIIC).

A. Dynamic implication for the short-range cutoff of the metastable DCF $c^*(\mathbf{r})$

Expanding the non-equilibrium excess chemical potential $\lambda_{\text{ex}}[\rho]$ around $\rho^*(\mathbf{r})$, the Dean-Kawasaki eqn (14) becomes

$$\begin{aligned} \frac{\partial \rho(\mathbf{r}, t)}{\partial t} &= \frac{\partial \{ \nu(\mathbf{r}, t) + \rho^*(\mathbf{r}) \}}{\partial t} \\ &= \nabla \cdot \mathcal{D} \rho \nabla \left\{ \lambda_{\text{ex}}^* + \int d\mathbf{r}' \frac{\delta \lambda_{\text{ex}}[\rho]}{\delta \rho(\mathbf{r}')} \Big|_{\rho=\rho^*} \nu(\mathbf{r}', t) \right\} + \zeta[\rho, \tilde{\eta}] \\ &= \nabla \cdot \mathcal{D} \rho \nabla \int d\mathbf{r}' \frac{\delta \lambda_{\text{ex}}[\rho]}{\delta \rho(\mathbf{r}')} \Big|_{\rho=\rho^*} \nu(\mathbf{r}', t) + \zeta[\rho, \tilde{\eta}], \end{aligned} \quad (84)$$

$$\begin{aligned} \frac{\delta \lambda_{\text{ex}}[\rho]}{\delta \rho(\mathbf{r}')} \Big|_{\rho=\rho^*} &= \frac{\delta(\mathbf{r} - \mathbf{r}')}{\rho^*(\mathbf{r})} - c^*(\mathbf{r} - \mathbf{r}') - \frac{\delta \lambda_N}{\delta \rho(\mathbf{r}')} \Big|_{\rho=\rho^*} \\ &= \frac{\delta(\mathbf{r} - \mathbf{r}')}{\rho^*(\mathbf{r})} - c^*(\mathbf{r} - \mathbf{r}'), \end{aligned} \quad (85)$$

where $|\nabla \lambda_{\text{ex}}^*| = 0$ has been used in eqn (84). Combining Eqn (84) with eqn (85) leads to the linearized Dean-Kawasaki equation as follows:

$$\begin{aligned} \frac{\partial \nu(\mathbf{r}, t)}{\partial t} &= \mathcal{D} \nabla^2 \nu(\mathbf{r}, t) \\ &\quad - \nabla \cdot \mathcal{D} \rho^* \int d\mathbf{r}' \nabla c^*(\mathbf{r} - \mathbf{r}') \nu(\mathbf{r}', t) + \sqrt{2} \zeta[\rho^*, \tilde{\eta}], \end{aligned} \quad (86)$$

due to the manipulation of the noise term.⁵⁴ Eqn (86) represents the overdamped dynamics of the fluctuating density field $\nu(\mathbf{r}, t)$ around the metastable non-hyperuniform state.



Eqn (86) indicates that the interaction-induced restoring force against the density deviation $\nu(\mathbf{r}, t)$ is given by the sum of $-\nabla c^*(\mathbf{r} - \mathbf{r}')\nu(\mathbf{r}', t)$. Focusing on the short-range contribution to this force, we find that microscopic environments in the hyperuniform and non-hyperuniform states are quite different from each other. While the scaling behavior (6) in a hyperuniform state predicts the divergence of $|\nabla c(\mathbf{r} - \mathbf{r}')| \rightarrow \infty$ in the limit of $|\mathbf{r} - \mathbf{r}'| \rightarrow 0$, the short-range cutoff of the non-hyperuniform DCF $c^*(\mathbf{r})$ creates the opposite situation on the particle scale:

$$|\nabla c^*(\mathbf{r} - \mathbf{r}')| \approx 0 \quad (|\mathbf{r} - \mathbf{r}'| < \sigma), \quad (87)$$

as seen from Fig. 1. The above relation implies that there is no interaction-induced restoring force against the ν -field in the non-hyperuniform states at the particle-scale while preserving the long-range contribution, $-\nabla c^*(\mathbf{r} - \mathbf{r}')\nu(\mathbf{r}', t) = \nabla(\beta/|\mathbf{r} - \mathbf{r}'|^2)\nu(\mathbf{r}', t)$, for $|\mathbf{r} - \mathbf{r}'| \gg \sigma$.

B. Microscopic mechanism behind the appearance of eqn (87)

Eqn (86) for the overdamped Brownian dynamics is insufficient for the descriptor of vibrational modes due to the absence of the inertia term, and yet eqn (87) suggests the emergence of dynamic softening in non-hyperuniform systems. We can learn the microscopic mechanism of soft modes from previous studies on quasicontracts of a contact network, a skeleton of jammed matter.^{18–22,75,76} The previous findings could provide an intuitive understanding of the virial-type expansion at high density as will be seen below.

For isostatic and hyperuniform systems, the packing geometry uniquely defines the contact forces as well as the spatial network structures including void distributions.^{1,2,8–12} Previous studies have shown that the isostatic and hyperuniform states disappear upon relaxing the strict constraints on the size- and spatial-distributions of voids slightly away from jamming.^{1,2,9,10,12} The relative abundance of non-isostatic contacts provides quasicontracts that carry weak forces, thereby creating local excitations with little restoring forces.^{12,19–22,76}

The possible particles forming the quasicontracts include rattlers and/or bucklers.^{12,19–22,76–83} It has been found, for instance, that the bucklers in the d -dimensional space, having $d + 1$ contacts as part of the contact network, are likely to be buckled to generate quasi-localized soft modes observed in the lowest-frequency regime.^{19–22,76–83} The anomalous vibrational modes have been shown to exhibit strong anharmonicities that are accompanied by intermittent rearrangements of particles as follows: opening a weak contact of a buckler (*i.e.*, buckling) yields a disordered core of a few particle scale with a power-law decay of displacements which are coupled to the elastic background of the contact network.^{19–37,76–83}

It is not the center of our concern whether or not the rattlers and/or bucklers significantly contribute to the quasicontracts to degrade the hyperuniformity. It is, however, illuminating to interpret eqn (86) and (87) in terms of quasi-localized soft modes.

Then, let $\mathbf{u}(\mathbf{r}, t)$ be the fluctuating displacement field induced by the fluctuating density field $\nu(\mathbf{r}, t)$. In the first approximation, we have⁹⁷

$$\nu(\mathbf{r}, t) = -\nabla \cdot \{\rho^*(\mathbf{r})\mathbf{u}(\mathbf{r}, t)\}. \quad (88)$$

Combining eqn (54), (56) and (86)–(88), we can verify that the displacement field $\mathbf{u}(\mathbf{r}, t)$ shares common features with that of the quasi-localized soft modes as follows:

- The interaction-induced restoring force of $\mathbf{u}(\mathbf{r}, t)$ is long-ranged in correspondence with recent simulations^{98–101} because of the power-law decay of the metastable DCF $c^*(\mathbf{r})$ represented by eqn (54) and (56).

- Eqn (86)–(88) imply the short-range anharmonicity of $\mathbf{u}(\mathbf{r}, t)$ at the particle scale.

This connection of our theoretical results (particularly, eqn (87)) with the quasi-localized soft modes suggests that the virial-type expansion in a glassy state represents the particle–particle interactions occurring due to the intermittent particle rearrangements.

C. Summarizing the results in comparison with other treatments

The differences in the free-energy density functionals between the equilibrium and stochastic DFTs are summarized as follows:

(i) The density functional $\mathcal{A}[\rho]$ appearing in the metastability eqn (34) represents the free-energy functional of the given density distribution $\rho(\mathbf{r})$, instead of the equilibrium free-energy functional $F[\rho, 0]$. It is a clear advantage over equilibrium DFT that stochastic DFT can make use of the field-theoretic formulation in obtaining $\mathcal{A}[\rho]$.

(ii) The metastability eqn (34) states that the functional derivative $\delta\mathcal{A}[\rho]/\delta\rho$ should yield a spatially constant λ_{ex}^* , which has been referred to as the metastable excess chemical potential. The sum of λ_{ex}^* and the Lagrange multiplier λ_N corresponds to the metastable chemical potential and is reduced to the equilibrium chemical potential (*i.e.*, $\lambda_{\text{ex}}^* + \lambda_N = \mu$) when $\lambda_{\text{ex}}^* = 0$ and $\lambda_N = \mu$ in equilibrium; see also the discussion after eqn (32).

(iii) As found from eqn (30), (41) and (56), the input of the hyperuniform DCF allows equilibrium DFT to predict the hyperuniformity of a metastable state, without the knowledge on the reference density distribution in an amorphous state.

These characteristics of stochastic DFT enable us to evaluate the extent to which fluctuations around the metastable density ρ^* affect the metastable chemical potential λ^* . Actually, we have demonstrated that stochastic DFT is relevant to determine metastable states around a hyperuniform state. Stochastic DFT provides the analytical form of the metastable DCF that has a short-range cutoff inside the sphere while retaining the long-range power-law behavior. We should keep in mind that the long-range hyperuniform behavior is preserved because the Gaussian weight, $e^{-\Delta F_{\text{eff}}}$, for the virial-type expansion premises that non-hyperuniform states considered are located near a hyperuniform state. As confirmed in Section VI, the obtained



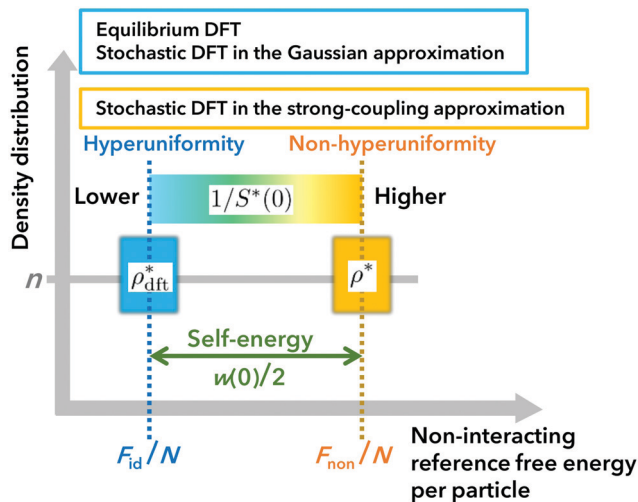


Fig. 3 A schematic comparison of theoretical approaches presented in this study. The spatially uniform density is identically n as shown on the vertical axis, and the hyperuniformity is incorporated into equilibrium DFT by inputting the hyperuniform DCF $c(r)$ given by eqn (56); nevertheless, we have hyperuniform and non-hyperuniform treatments shown in blue and orange, respectively. The different results are due to the distinct values of non-interacting reference free-energy functionals (*i.e.*, $F_{id}[\rho]$ given by eqn (26) and $F_{non}[\rho]$ given by eqn (75)), which is represented by the horizontal axis.

DCF yields the zero-wavevector structure factor in quantitative agreement with previous simulation results^{1,2,38–43} of degraded hyperuniformity.

Moreover, both Fig. 3 and Table 2 summarize the results by comparing the following theoretical approaches discussed so far: the equilibrium DFT using the Ramakrishnan–Yussouff free-energy functional,⁵⁷ the stochastic DFT in the Gaussian approximation (see Section V), and the stochastic DFT in the strong-coupling approximation (see Section VII).

The vertical axis in Fig. 3 shows that the density distribution considered has the same density n on average. The difference is attributed to the inhomogeneous distributions around n : the hyperuniform density distribution $\rho_{dft}^*(\mathbf{r})$, shown in blue, satisfies eqn (3) for the inverse of the zero-wavevector structure factor $1/S^*(0)$, whereas the non-hyperuniform range of density distribution $\rho^*(\mathbf{r})$, shown in orange, satisfies eqn (2). As summarized in Fig. 3, the equilibrium DFT and stochastic DFT in the Gaussian approximation provide hyperuniformity, whereas the stochastic DFT in the strong-coupling approximation provides non-hyperuniformity.

Meanwhile, the transverse axis of Fig. 3 shows that the above two types of theoretical approaches take distinct reference free-energy functionals, as found by comparing F_{id} and F_{non} given by eqn (26) and (75), respectively. In the hyperuniform theories, on the one hand, the ideal free-energy functional F_{id}/N per particle is the reference functional for the evaluation of interaction energy (see eqn (26)), following the conventional treatment of equilibrium DFT,^{57–61} where N denotes the total number of spheres as before. On the other hand, as a reference functional of stochastic DFT in the strong-coupling approximation, we

used the free-energy functional F_{non}/N of a non-interacting system per particle that is larger than the ideal one F_{non}/N by the self-energy $w(0)/2$. It can be stated that a perturbation field theory method becomes more relevant to the evaluation of intermittent fluctuations due to the increase in the reference free energy.

Table 2 presents the more detailed classification of the hyperuniform and non-hyperuniform theories. There are two types of classifications for the above three treatments. One classification is based on the DFT type of whether the dynamical DFT relies on the deterministic equation (eqn (29)) or the stochastic equation (eqn (14)). The former approach represented by eqn (29) is equivalent to equilibrium DFT as clarified at the beginning of Section IIID, whereas the latter eqn (14) forms the basis of the last two stochastic approaches where the additional contribution $\Delta F[\rho, \phi]$ to the intrinsic Helmholtz free energy $F[\rho, 0]$ is to be considered. The other aspects of theoretical classification concern the predictability of non-hyperuniformity especially when the hyperuniformity is incorporated into the DCF (*i.e.*, eqn (44) or (56)) of equilibrium DFT as input. The type specification column in Table 2 indicates that the stochastic DFT in the Gaussian approximation falls into the same category (type H2 defined in Table 1) of the equilibrium DFT in this light.

As confirmed from Table 2, the use of the density-expansion method at strong coupling is indispensable to convert the hyperuniform structure factor at the zero wavevector into the non-hyperuniform one satisfying the simulation results given by eqn (2). The outstanding feature of the non-hyperuniform DCF $c^*(\mathbf{r})$ is the short-range cutoff, thereby predicting the absence of the interaction-induced restoring force for the short-range dynamics (*i.e.*, eqn (87)).

IX. Concluding remarks

For comparison purposes, let us go back to the previous study¹⁵ where the degradation of perfect hyperuniformity in crystals, quasicrystals, and disordered packings has been demonstrated both theoretically and numerically, using the three scenarios of imperfections (see Section II for the list of scenarios). The second scenario (ii), which has been our concern, attributes the violation of hyperuniformity to the stochastic occurrence of spatially correlated displacements.

Combining eqn (76) and the dynamical discussion in Section VIII suggests that the above-mentioned second scenario for the degradation of hyperuniformity is similar to the underlying physics described by the averaged virial-type interaction term, the third term on the rhs of eqn (51), under the long-range-correlated ϕ -field; for it seems plausible that the long-range-correlated potential field arises from the elastic nature of the contact network. From the discussion, we infer that the virial-type degradation of hyperuniformity reflects the intermittent rearrangements of particles and is more likely to be found in the collectively jammed packings allowing for the shear deformations as mentioned before, rather than in the strictly



jammed ones.^{1,2,8} In other words, type H2 defined in Table 1 corresponds to the collectively jammed packings, whereas type H1 corresponds to the strictly jammed packings.

It remains to be seen whether the present formulation can be extended to address the non-hyperuniform behaviors of other measures than the density-density structure factor, which are obtained from various physical quantities including the local number variance for a window¹⁷ and the contact number fluctuations.^{102,103} We also envision that advancing stochastic DFT^{44–56} at strong coupling will pave the way for the realistic description of quasi-localized soft modes induced by the intermittent rearrangements of particles such as bucklers.^{19–22,76–83}

Conflicts of interest

There are no conflicts to declare.

Appendix A: details on the constrained free-energy functional $\mathcal{A}[\rho]$

1. Verification of eqn (9)

The distribution functional $P[\rho, t]$ defined by eqn (8) satisfies the Fokker–Planck equation as follows:^{44,104}

$$\frac{\partial P[\rho, t]}{\partial t} = - \int d\mathbf{r} \frac{\delta}{\delta \rho} \nabla \cdot \mathcal{D} \rho \nabla \left[\frac{\delta}{\delta \rho} + \frac{\delta \mathcal{A}[\rho]}{\delta \rho} \right] P[\rho, t], \quad (\text{A1})$$

from which we find that eqn (9) satisfies the stationary condition $\partial P_{\text{st}}[\rho]/\partial t = 0$. It has also been shown that eqn (A1) is equivalent to the Dean–Kawasaki eqn (14).^{44,104}

2. Definition of $\mathcal{A}[\rho]$

In eqn (A1) as well as in eqn (14), the canonical ensemble is naturally required for the free-energy functional $\mathcal{A}[\rho]$ of a given density ρ because we consider the overdamped dynamics of the densely packed sphere system with the total number N of spheres being fixed. Hence, $\mathcal{A}[\rho]$ is defined using the configurational integral for the canonical ensemble as follows:

$$e^{-\mathcal{A}[\rho]} = \frac{1}{N!} \int d\mathbf{r}_1 \cdots \int d\mathbf{r}_N \prod_{i,j} e^{-v(\mathbf{r}_i - \mathbf{r}_j)} \times \prod_{\mathbf{r}} \delta[\hat{\rho}_N(\mathbf{r}, t) - \rho(\mathbf{r}, t)]. \quad (\text{A2})$$

Yet, equilibrium DFT, a key ingredient in this study, needs to be formulated in the grand canonical system. We therefore write $\mathcal{A}[\rho]$ with the help of the contour integral over a complex variable $z = e^\mu$:^{54,104}

$$e^{-\mathcal{A}[\rho]} = \frac{1}{2\pi i} \oint \frac{dz}{z^{N+1}} \times \left(\text{Tr} \prod_i e^\mu \prod_{i,j} e^{-v(\mathbf{r}_i - \mathbf{r}_j)} \prod_{\mathbf{r}} \delta[\hat{\rho}_N(\mathbf{r}, t) - \rho(\mathbf{r}, t)] \right), \quad (\text{A3})$$

so that the canonical ensemble may be recovered after performing the grand canonical ensemble represented by

$$\text{Tr} \equiv \sum_{N=0}^{\infty} \frac{1}{N!} \int d\mathbf{r}_1 \cdots \int d\mathbf{r}_N.$$

3. Derivation of eqn (13)

We evaluate the underlined term in eqn (A3) with the help of the Fourier transform of the delta functional as follows:

$$\begin{aligned} \text{Tr} \prod_i e^\mu \prod_{i,j} e^{-v(\mathbf{r}_i - \mathbf{r}_j)} \prod_{\mathbf{r}} \delta[\hat{\rho}_N(\mathbf{r}) - \rho(\mathbf{r})] \\ = \int D\psi \text{Tr} \prod_i e^{\mu + i\psi(\mathbf{r}_i)} \prod_{i,j} e^{-v(\mathbf{r}_i - \mathbf{r}_j)} e^{-\int d\mathbf{r} i\psi(\mathbf{r}) \rho(\mathbf{r})} \\ = \int D\psi e^{-\Omega[-i\psi] - \int d\mathbf{r} i\psi(\mathbf{r}) \rho(\mathbf{r})}. \end{aligned} \quad (\text{A4})$$

The ψ -field is separated into the fluctuating potential field $\phi(\mathbf{r})$ and the saddle-point field $i\psi_{\text{dft}}(\mathbf{r})$:

$$\psi(\mathbf{r}) = \phi(\mathbf{r}) + i\psi_{\text{dft}}(\mathbf{r}), \quad (\text{A5})$$

where $\psi_{\text{dft}}(\mathbf{r})$ is determined by the saddle-point equation,

$$\left. \frac{\delta(\Omega[-i\psi])}{\delta\psi(\mathbf{r})} \right|_{\psi=i\psi_{\text{dft}}} = -i\rho(\mathbf{r}). \quad (\text{A6})$$

The functional differentiation on the left-hand side of eqn (A6) provides the density in equilibrium of the system under the external field ψ_{dft} . Denoting the equilibrium density by $\langle \hat{\rho}_N(\mathbf{r}) \rangle_{\text{eq}}$, the saddle-point eqn (A6) implies that

$$\langle \hat{\rho}_N(\mathbf{r}) \rangle_{\text{eq}} = \rho(\mathbf{r}). \quad (\text{A7})$$

The above relation states that a prescribed density $\rho(\mathbf{r})$ is equated with the equilibrium density due to the potential ψ_{dft} along the saddle-point field.

For later convenience, we also introduce the intrinsic Helmholtz free energy $F[\rho, 0]$, the central functional of equilibrium DFT:^{58–61}

$$F[\rho, 0] - \int d\mathbf{r} \rho(\mathbf{r}) \mu \equiv \Omega[\psi_{\text{dft}}] - \int d\mathbf{r} \rho(\mathbf{r}) \psi_{\text{dft}}(\mathbf{r}), \quad (\text{A8})$$

showing that the intrinsic Helmholtz free energy $F[\rho, 0]$ is defined by the first Legendre transform of the grand potential $\Omega[\psi_{\text{dft}}]$ with the saddle-point field $\psi_{\text{dft}}(\mathbf{r})$ being applied. Therefore, the identity (13) is satisfied as well as that in equilibrium DFT.



4. Derivation of eqn (12)

Combining eqn (A4) and (A5), we have

$$\begin{aligned} & \text{Tr} \prod_i e^{\mu} \prod_{i,j} e^{-v(\mathbf{r}_i - \mathbf{r}_j)} \prod_{\mathbf{r}} \delta[\hat{\rho}_N(\mathbf{r}) - \rho(\mathbf{r})] \\ &= \int D\phi \text{Tr} \prod_i e^{\mu + i\phi(\mathbf{r}_i) - \psi_{\text{dft}}(\mathbf{r}_i)} \prod_{i,j} e^{-v(\mathbf{r}_i - \mathbf{r}_j)} \\ & \times \int d\mathbf{r} \rho(\mathbf{r}) \{\psi_{\text{dft}}(\mathbf{r}) - i\phi(\mathbf{r})\} \\ &= \int D\phi e^{-\Omega[\psi_{\text{dft}} - i\phi] + \int d\mathbf{r} \rho(\mathbf{r}) \{\psi_{\text{dft}}(\mathbf{r}) - i\phi(\mathbf{r})\}} \\ &= \int D\phi e^{-F[\rho, \phi] + \int d\mathbf{r} \rho(\mathbf{r}) \mu}, \end{aligned} \quad (\text{A9})$$

when defining

$$\begin{aligned} F[\rho, \phi] - \int d\mathbf{r} \rho(\mathbf{r}) \mu &\equiv \Omega[\psi_{\text{dft}} - i\phi] \\ &- \int d\mathbf{r} \rho(\mathbf{r}) \{\psi_{\text{dft}}(\mathbf{r}) - i\phi(\mathbf{r})\} \end{aligned} \quad (\text{A10})$$

as an extension of eqn (A8). Eqn (A9) and (A10) validate eqn (12).

5. Derivation of eqn (10) and (11)

It follows from eqn (A3) and (A9) that

$$\begin{aligned} e^{-\mathcal{A}[\rho]} &= \frac{1}{2\pi i} \oint \frac{dz}{z^{N+1}} \int D\phi e^{-F[\rho, \phi] + \int d\mathbf{r} \rho(\mathbf{r}) \mu} \\ &= \int D\phi e^{-F[\rho, \phi]} \left(\frac{1}{2\pi i} \oint \frac{dz}{z^{-\int d\mathbf{r} \rho(\mathbf{r}, t) + N+1}} \right) \\ &= \int D\phi e^{-F[\rho, \phi]} \Delta[\rho], \end{aligned} \quad (\text{A11})$$

where

$$\begin{aligned} \Delta[\rho] &\equiv \frac{1}{2\pi i} \oint \frac{dz}{z^{-\int d\mathbf{r} \rho(\mathbf{r}, t) + N+1}} \\ &= \begin{cases} 1 & \left(\int d\mathbf{r} \rho(\mathbf{r}) = N \right) \\ 0 & \left(\int d\mathbf{r} \rho(\mathbf{r}) \neq N \right). \end{cases} \end{aligned} \quad (\text{A12})$$

Eqn (A11) and (A12) verify eqn (10) and (11), respectively.

6. Derivation of eqn (19)

The difference $\Delta F_{\text{dft}}[\rho, \phi] = F[\rho, \phi] - F[\rho, 0]$ between eqn (A8) and (A10) reads

$$\Delta F_{\text{dft}}[\rho, \phi] = \Omega[\psi_{\text{dft}} - i\phi] - \Omega[\psi_{\text{dft}}] + \int d\mathbf{r} i\rho(\mathbf{r})\phi(\mathbf{r}). \quad (\text{A13})$$

The quadratic expansion of $\Omega[\psi_{\text{dft}} - i\phi]$ around the imaginary saddle-point field $i\psi_{\text{dft}}$ yields

$$\Delta F_{\text{dft}}[\rho, \phi] = \frac{1}{2} \int d\mathbf{r} d\mathbf{r}' \frac{\delta^2 \Omega}{\delta \psi(\mathbf{r}) \delta \psi(\mathbf{r}')} \Big|_{\psi=\psi_{\text{dft}}} \phi(\mathbf{r}) \phi(\mathbf{r}') \quad (\text{A14})$$

$$= \frac{1}{2} \int d\mathbf{r} d\mathbf{r}' \phi(\mathbf{r}) w^{-1}(\mathbf{r} - \mathbf{r}') \phi(\mathbf{r}'), \quad (\text{A15})$$

where the basic relationship has been used in equilibrium DFT as follows:

$$\int d\mathbf{r}' \frac{\delta^2 F[\rho, 0]}{\delta \rho(\mathbf{r}) \delta \rho(\mathbf{r}')} \frac{\delta^2 \Omega}{\delta \psi(\mathbf{r}') \delta \psi(\mathbf{r}'')} \Big|_{\psi=i\psi_{\text{dft}}} = \delta(\mathbf{r} - \mathbf{r}''), \quad (\text{A16})$$

as well as the definition (20) of $w(\mathbf{r} - \mathbf{r}')$.

7. Derivation of eqn (46)

It is found from eqn (22) that eqn (46) reads

$$\begin{aligned} & \frac{\delta}{\delta \rho} \int d\mathbf{r} d\mathbf{r}' w^{-1}(\mathbf{r} - \mathbf{r}') \phi(\mathbf{r}) \phi(\mathbf{r}') \\ &= \int d\mathbf{r}' \left[\{w^{-1}(\mathbf{r} - \mathbf{r}')\}' \overline{\phi(\mathbf{r}) \phi(\mathbf{r}')} \right. \\ & \left. + w^{-1}(\mathbf{r} - \mathbf{r}') \frac{\int D\phi \{\phi(\mathbf{r}) \phi(\mathbf{r}')\}' e^{-\Delta F_{\text{dft}}[\rho, \phi]}}{\int D\phi e^{-\Delta F_{\text{dft}}[\rho, \phi]}} \right], \end{aligned} \quad (\text{A17})$$

where

$$\begin{aligned} \{w^{-1}(\mathbf{r} - \mathbf{r}')\}' &\approx \delta(\mathbf{r} - \mathbf{r}') + 2h(\mathbf{r} - \mathbf{r}')\rho(\mathbf{r}') \\ &= \frac{w^{-1}(\mathbf{r} - \mathbf{r}')}{\rho(\mathbf{r})} + h(\mathbf{r} - \mathbf{r}')\rho(\mathbf{r}'), \end{aligned} \quad (\text{A18})$$

neglecting the density dependence of the total correlation function $h(\mathbf{r} - \mathbf{r}')$, and eqn (A17) and (A18) give a precise definition of $\{\phi(\mathbf{r}) \phi(\mathbf{r}')\}'$ appearing in eqn (A17).

In eqn (A17), we use the following approximation:

$$\begin{aligned} \{\phi(\mathbf{r}) \phi(\mathbf{r}')\}' &\equiv \frac{\delta}{\delta \rho(\mathbf{r})} \left\{ \frac{\int D\phi \phi(\mathbf{r}) \phi(\mathbf{r}') e^{-\Delta F_{\text{dft}}[\rho, \phi]}}{\int D\phi e^{-\Delta F_{\text{dft}}[\rho, \phi]}} \right\} \\ &= \frac{\int D\phi \{\phi(\mathbf{r}) \phi(\mathbf{r}')\}' e^{-\Delta F_{\text{dft}}[\rho, \phi]}}{\int D\phi e^{-\Delta F_{\text{dft}}[\rho, \phi]}} \\ & \quad - \overline{\phi(\mathbf{r}) \phi(\mathbf{r}')} \frac{\delta \Delta F_{\text{dft}}[\rho, \phi]}{\delta \rho(\mathbf{r})} \\ & \quad + \overline{\phi(\mathbf{r}) \phi(\mathbf{r}')} \left\{ \frac{\delta \Delta F_{\text{dft}}[\rho, \phi]}{\delta \rho(\mathbf{r})} \right\} \\ &\approx \frac{\int D\phi \{\phi(\mathbf{r}) \phi(\mathbf{r}')\}' e^{-\Delta F_{\text{dft}}[\rho, \phi]}}{\int D\phi e^{-\Delta F_{\text{dft}}[\rho, \phi]}}. \end{aligned} \quad (\text{A19})$$

Accordingly, eqn (A17) reduces to



$$\begin{aligned}
& \frac{\delta}{\delta \rho} \iint d\mathbf{r} d\mathbf{r}' w^{-1}(\mathbf{r} - \mathbf{r}') \phi(\mathbf{r}) \phi(\mathbf{r}') \\
& \approx \int d\mathbf{r}' \left[\left\{ w^{-1}(\mathbf{r} - \mathbf{r}') \right\}' \overline{\phi(\mathbf{r}) \phi(\mathbf{r}')} \right. \\
& \quad \left. + w^{-1}(\mathbf{r} - \mathbf{r}') \left\{ \overline{\phi(\mathbf{r}) \phi(\mathbf{r}')} \right\}' \right] \quad (\text{A20}) \\
& = \frac{1}{\rho(\mathbf{r})} + h(0) - \int d\mathbf{r}' h(\mathbf{r} - \mathbf{r}') \rho(\mathbf{r}') c(\mathbf{r}' - \mathbf{r}) - \frac{w^{-1}(\mathbf{0})}{\rho^2(\mathbf{r})} \\
& = - \int d\mathbf{r}' h(\mathbf{r} - \mathbf{r}') \rho(\mathbf{r}') c(\mathbf{r}' - \mathbf{r}) = c(0) - h(0),
\end{aligned}$$

where

$$\begin{aligned}
& \left\{ \overline{\phi(\mathbf{r}) \phi(\mathbf{r}')} \right\}' = \left\{ w(\mathbf{r} - \mathbf{r}') \right\}' \\
& = \left\{ \frac{\delta(\mathbf{r} - \mathbf{r}')}{\rho(\mathbf{r})} - c(\mathbf{r} - \mathbf{r}') \right\}' \quad (\text{A21}) \\
& \approx - \frac{\delta(\mathbf{r} - \mathbf{r}')}{\rho^2(\mathbf{r})},
\end{aligned}$$

neglecting the density dependence of the DCF, and the Ornstein–Zernike equation has been used in the last equality of eqn (A20).

Appendix B: the zero-wavevector structure factor $S(0)$ represented by the metastable DCF $c^*(\mathbf{r})$

1. Derivation of eqn (39)

We consider the Ornstein–Zernike equation for the metastable correlation functions, $c^*(\mathbf{r})$ and $h^*(\mathbf{r})$, in a uniform state. The Ornstein–Zernike equation at zero separation is expressed as

$$\begin{aligned}
h^*(0) &= c^*(0) + n \int 4\pi r^2 dr c^*(\mathbf{r}) h^*(\mathbf{r}) \\
&= c^*(0) - n \int_{r \leq \sigma} 4\pi r^2 dr c^*(\mathbf{r}) \quad (\text{B1}) \\
&\quad + n \int_{r \geq \sigma} 4\pi r^2 dr c^*(\mathbf{r}) h^*(\mathbf{r}),
\end{aligned}$$

where the relationship $h^*(\mathbf{r}) = -1$ ($0 \leq r \leq \sigma$) has been used in the above second line. Adding the term $-n \int_{r \geq \sigma} 4\pi r^2 dr c^*(\mathbf{r})$ on both sides of eqn (B1), we have

$$\begin{aligned}
& h^*(0) - n \int_{r \geq \sigma} 4\pi r^2 dr c^*(\mathbf{r}) \\
& = c^*(0) - n \int_{r \geq \sigma} d\mathbf{r} c^*(\mathbf{r}) + n \int_{r \geq \sigma} 4\pi r^2 dr c^*(\mathbf{r}) h^*(\mathbf{r}), \quad (\text{B2})
\end{aligned}$$

which reads

$$\begin{aligned}
& -h^*(0) - n \int_{r \geq \sigma} d\mathbf{r} c^*(\mathbf{r}) \\
& = -c^*(0) - n \int_{r \geq \sigma} 4\pi r^2 dr c^*(\mathbf{r}) - n \int_{r \geq \sigma} 4\pi r^2 dr c^*(\mathbf{r}) h^*(\mathbf{r}). \quad (\text{B3})
\end{aligned}$$

2. Derivation of eqn (57)

It is noted that $c^*(\mathbf{r})h^*(\mathbf{r})$ decays far more rapidly than $c^*(\mathbf{r})$ even when the effective correlation functions converge to the hyperuniform ones, $c(\mathbf{r})$ and $h(\mathbf{r})$, for $r > \sigma$. Therefore, eqn (39) becomes

$$\begin{aligned}
\frac{1}{S^*(0)} &\approx -c^*(0) - n \int_{\sigma}^{L_c} 4\pi r^2 dr c^*(\mathbf{r}) \\
&= -c^*(0) - \frac{6f_v}{\pi\sigma^3} \int_{\sigma}^{L_c} 4\pi r^2 dr c^*(\mathbf{r}) \quad (\text{B4}) \\
&= -c^*(0) - 24f_v \int_1^{L_c/\sigma} d\tilde{r} \tilde{r}^2 c^*(\tilde{r}),
\end{aligned}$$

where the relationship $f_v = \pi n \sigma^3 / 6$ between the volume fraction f_v and the spatially averaged density n has been used in the second line of the above equation. For $\tilde{r} = r/\sigma > 1$, combining eqn (52) and (56) provides

$$-c^*(\tilde{r}) = 1 - \exp\left(-\frac{\beta}{\tilde{r}^2}\right), \quad (\text{B5})$$

which will be used in calculating the last term in the last line of eqn (B4). Integration by parts yields

$$\begin{aligned}
-3 \int_1^{L_c/\sigma} d\tilde{r} \tilde{r}^2 c^*(\tilde{r}) &= -[\tilde{r}^3 c^*(\tilde{r})]_1^{L_c/\sigma} + \int_1^{L_c/\sigma} d\tilde{r} \tilde{r}^3 \frac{dc^*(\tilde{r})}{d\tilde{r}} \\
&= -\left(\frac{L_c}{\sigma}\right)^3 c^*\left(\frac{L_c}{\sigma}\right) + c^*(1) + 2\beta \int_1^{L_c/\sigma} d\tilde{r} e^{-\frac{\beta}{\tilde{r}^2}} \\
&= I_1\left(\beta, \frac{L_c}{\sigma}\right) + 2\beta I_2\left(\beta, \frac{L_c}{\sigma}\right), \quad (\text{B6})
\end{aligned}$$

where

$$I_1\left(\beta, \frac{L_c}{\sigma}\right) = -\left(\frac{L_c}{\sigma}\right)^3 c^*\left(\frac{L_c}{\sigma}\right) + c^*(1), \quad (\text{B7})$$

and the change of variable from \tilde{r} to $x = 1/\tilde{r}$ gives

$$\begin{aligned}
I_2\left(\beta, \frac{L_c}{\sigma}\right) &= \int_1^{L_c/\sigma} d\tilde{r} e^{-\frac{\beta}{\tilde{r}^2}} \\
&= \int_{\sigma/L_c}^1 dx \frac{e^{-\beta x^2}}{x^2} \\
&= -\left[\frac{e^{-\beta x^2}}{x}\right]_{\sigma/L_c}^1 - 2\beta \int_{\sigma/L_c}^1 dx e^{-\beta x^2} \quad (\text{B8}) \\
&= \frac{L_c e^{-\beta(\sigma/L_c)^2}}{\sigma} - e^{-\beta} \\
&\quad - \beta \sqrt{\frac{\pi}{\beta}} \left\{ \text{erf}\left(\sqrt{\beta}\right) - \text{erf}\left(\frac{\sqrt{\beta}\sigma}{L_c}\right) \right\}.
\end{aligned}$$

Combining eqn (B2)–(B8), we have

$$\frac{1}{S^*(0)} = -c^*(0) + 8f_v I_1\left(\beta, \frac{L_c}{\sigma}\right) + 16\beta f_v I_2\left(\beta, \frac{L_c}{\sigma}\right). \quad (\text{B9})$$

The approximate form of eqn (B9) for $L_c/\sigma \gg 1$ leads to eqn (57):



$$\frac{1}{S^*(0)} \approx -8f_v \left(\frac{L_c}{\sigma} \right)^3 c^*(L_c/\sigma) + 16\beta f_v \frac{L_c e^{-\beta(\sigma/L_c)^2}}{\sigma} \quad (\text{B10})$$

$$\approx 24f_v \beta \left(\frac{L_c}{\sigma} \right),$$

where we have used the approximations, $-c^*(L_c/\sigma) \approx \beta(\sigma/L_c)^2$ and $L_c e^{-\beta(\sigma/L_c)^2}/\sigma \approx L_c/\sigma$, in the above second line. Fig. 2 shows the L_c/σ -dependencies of $1/S^*(0)$ in order to compare eqn (B9) and (B10).

Appendix C: details on the averaging operation over the ϕ -field in the strong-coupling approximation

1. Shifting the fluctuating-potential field from ϕ to φ in eqn (17): a general formulation and validation of eqn (76)

We first see extra terms when $\Delta F_{\text{dft}}[\rho, \phi]$ is represented by the ϕ -field. Eqn (67) is rearranged to give

$$\phi(\mathbf{r}) = \varphi(\mathbf{r}) + i\Delta\psi(\mathbf{r}) - i\frac{w(0)}{2}, \quad (\text{C1})$$

which further reads

$$\phi(\mathbf{r}) = \gamma\tilde{\phi}(\mathbf{r}) = \gamma\tilde{\varphi}(\mathbf{r}) + i\tilde{\Delta}\psi(\mathbf{r}) - i\frac{\gamma^2\tilde{w}(0)}{2}, \quad (\text{C2})$$

due to the rescaling given by eqn (63) and its associated form,

$$\begin{aligned} \Delta\psi(\mathbf{r}) &= -\int d\mathbf{r}' \gamma^2 \tilde{c}(\mathbf{r} - \mathbf{r}') \frac{\tilde{\Delta}\rho(\mathbf{r}')}{\gamma^2} \\ &= -\int d\mathbf{r}' \tilde{c}(\mathbf{r} - \mathbf{r}') \tilde{\Delta}\rho(\mathbf{r}') \\ &= \tilde{\Delta}\psi(\mathbf{r}). \end{aligned} \quad (\text{C3})$$

Plugging eqn (C2) into eqn (19), we have

$$\Delta F_{\text{dft}}[\rho, \phi = \gamma\tilde{\phi}] = \Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\varphi}] - \frac{1}{\gamma^2} \Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\Delta}\psi] + \frac{i}{\gamma} \mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}]$$

$$\Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\Delta}\psi] = \frac{1}{2} \iint d\mathbf{r} d\mathbf{r}' \tilde{w}^{-1}(\mathbf{r} - \mathbf{r}') \tilde{\Delta}\psi(\mathbf{r}) \tilde{\Delta}\psi(\mathbf{r}'),$$

$$\mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}] = \iint d\mathbf{s} d\mathbf{s}' \tilde{w}^{-1}(\mathbf{s} - \mathbf{s}') \tilde{\varphi}(\mathbf{s}) \tilde{\Delta}\psi(\mathbf{s}'). \quad (\text{C4})$$

because of $\Delta F_{\text{dft}}[\tilde{\rho}, \gamma^2 \tilde{w}(0)/2] = 0$ for the hyperuniform correlation function $h(\mathbf{r})$:

$$\begin{aligned} \Delta F_{\text{dft}}\left[\tilde{\rho}, \frac{\gamma^2 \tilde{w}(0)}{2}\right] &= \frac{w^2(0)}{8} \int d\mathbf{r} \rho(\mathbf{r}) \left[\int d\mathbf{r}' \{ \delta(\mathbf{r} - \mathbf{r}') + h(\mathbf{r} - \mathbf{r}') \rho(\mathbf{r}') \} \right] \\ &= \frac{w^2(0)}{8} \int d\mathbf{r} \rho(\mathbf{r}) \left[1 + \int d\mathbf{r}' h(\mathbf{r} - \mathbf{r}') \rho(\mathbf{r}') \right] \\ &= \frac{Nw^2(0)}{8} S(0) = 0, \end{aligned} \quad (\text{C5})$$

where $S(0) = 1 + \int d\mathbf{r}' h(\mathbf{r} - \mathbf{r}') \rho(\mathbf{r}') = 0$ for the hyperuniform structure factor $S(0)$ at zero wavevector, and it is supposed that $w(0)$ is the position-independent function because of the neglect of the triplet DCF (see also the statement after eqn (48)).

Before proceeding to the ϕ -field averaging operation defined by eqn (17), we clarify the corresponding terms where the strong-coupling approximation needs to be developed. To this end, let $\zeta[\tilde{\varphi}]$ be the general functional given by the sum of the $\tilde{\varphi}$ -independent part ζ_c and the remaining $\tilde{\varphi}$ -dependent contribution $\zeta_r[\tilde{\varphi}]$:

$$\zeta[\tilde{\varphi}] = \zeta_c + \zeta_r[\tilde{\varphi}]. \quad (\text{C6})$$

It is noted that we can validate the following expansion at strong coupling (*i.e.*, $\gamma \gg 1$):

$$\begin{aligned} e^{-\frac{i}{\gamma} \mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}]} &\approx 1 - \frac{1}{\gamma} \left\{ i\mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}] + \frac{1}{2\gamma} \mathcal{E}_0^2[\tilde{\rho}, \tilde{\varphi}] \right\} \\ &\equiv 1 - \frac{1}{\gamma} \mathcal{E}[\tilde{\rho}, \tilde{\varphi}], \end{aligned} \quad (\text{C7})$$

$$\begin{aligned} \mathcal{E}_0^2[\tilde{\rho}, \tilde{\varphi}] &\equiv \iint d\mathbf{r} d\mathbf{r}' \tilde{w}^{-1}(\mathbf{r} - \mathbf{r}') \tilde{\varphi}(\mathbf{r}) \tilde{\Delta}\psi(\mathbf{r}') \\ &\quad \times \iint d\mathbf{s} d\mathbf{s}' \tilde{w}^{-1}(\mathbf{s} - \mathbf{s}') \tilde{\varphi}(\mathbf{s}) \tilde{\Delta}\psi(\mathbf{s}'). \end{aligned}$$

Combining eqn (C4) and (C7) provides the approximate functional $\overline{\zeta}[\tilde{\varphi}]$ that is averaged over the ϕ -field, instead of the $\tilde{\varphi}$ -field, based on the definition of eqn (17) as follows:

$$\begin{aligned} \overline{\zeta}[\tilde{\varphi}] &= \frac{\int D\phi \zeta[\tilde{\varphi}] e^{-\Delta F_{\text{dft}}[\rho, \phi]}}{\int D\phi e^{-\Delta F_{\text{dft}}[\rho, \phi]}} \\ &= \zeta_c + \frac{\int D\tilde{\varphi} \zeta_r[\tilde{\varphi}] e^{-\Delta F_{\text{dft}}[\rho, \phi]}}{\int D\tilde{\varphi} e^{-\Delta F_{\text{dft}}[\rho, \phi]}} \\ &= \zeta_c + \frac{\int D\tilde{\varphi} \zeta_r[\tilde{\varphi}] e^{-\Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\varphi}] + \frac{1}{\gamma^2} \Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\Delta}\psi] - \frac{i}{\gamma} \mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}]}}{\int D\tilde{\varphi} e^{-\Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\varphi}] + \frac{1}{\gamma^2} \Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\Delta}\psi] - \frac{i}{\gamma} \mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}]}} \\ &= \zeta_c + \frac{\int D\tilde{\varphi} \zeta_r[\tilde{\varphi}] \left(1 - \frac{1}{\gamma} \mathcal{E}[\tilde{\rho}, \tilde{\varphi}] \right) e^{-\Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\varphi}]}}{\int D\tilde{\varphi} \left(1 - \frac{1}{\gamma} \mathcal{E}[\tilde{\rho}, \tilde{\varphi}] \right) e^{-\Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\varphi}]}} \\ &= \zeta_c + \frac{\int D\tilde{\varphi} \zeta_r[\tilde{\varphi}] \left(1 - \frac{1}{\gamma} \mathcal{E}[\tilde{\rho}, \tilde{\varphi}] \right) e^{-\Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\varphi}]}}{\int D\tilde{\varphi} e^{-\Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\varphi}]} \left\{ \frac{\int D\tilde{\varphi} \left(1 - \frac{1}{\gamma} \mathcal{E}[\tilde{\rho}, \tilde{\varphi}] \right) e^{-\Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\varphi}]}}{\int D\tilde{\varphi} e^{-\Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\varphi}]}} \right\}} \\ &= \zeta_c + \frac{\int D\tilde{\varphi} \zeta_r[\tilde{\varphi}] \left(1 - \frac{1}{\gamma} \mathcal{E}[\tilde{\rho}, \tilde{\varphi}] \right) e^{-\Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\varphi}]}}{\int D\tilde{\varphi} e^{-\Delta F_{\text{dft}}[\tilde{\rho}, \tilde{\varphi}]} \left(1 - \frac{1}{\gamma} \overline{\mathcal{E}[\tilde{\rho}, \tilde{\varphi}]} \right)}. \end{aligned} \quad (\text{C8})$$

Here, another averaging operation $\overline{\mathcal{E}[\tilde{\rho}, \tilde{\varphi}]}$ appearing in the last



line of eqn (C8) has been introduced for representing

$$\overline{\mathcal{E}[\tilde{\rho}, \tilde{\varphi}]} \equiv \frac{\int D\tilde{\varphi} \mathcal{E}[\tilde{\rho}, \tilde{\varphi}] e^{-\Delta F_{\text{df}}[\tilde{\rho}, \tilde{\varphi}]} \int D\tilde{\varphi} e^{-\Delta F_{\text{df}}[\tilde{\rho}, \tilde{\varphi}]}, \quad (\text{C9})$$

which becomes equivalent to eqn (17) when replacing the ϕ -field with the φ -field; however, the notation of the above average has been altered for revealing that there is a difference between $\overline{\mathcal{E}[\tilde{\rho}, \tilde{\varphi}]}$ and $\overline{\mathcal{E}[\tilde{\rho}, \tilde{\varphi}]}$ as shown by the general form (C8). We obtain from eqn (C7) and (C9)

$$\begin{aligned} \frac{1}{\gamma} \overline{\mathcal{E}[\tilde{\rho}, \tilde{\varphi}]} &= \frac{1}{\gamma} \left\{ \overline{i\mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}]} + \frac{1}{2\gamma} \overline{\mathcal{E}_0^2[\tilde{\rho}, \tilde{\varphi}]} \right\} \\ &= \frac{1}{2\gamma^2} \iiint d\mathbf{r} d\mathbf{r}' d\mathbf{s} d\mathbf{s}' \overline{\tilde{\varphi}(\mathbf{r}) \tilde{\varphi}(\mathbf{s})} \\ &\quad \times \tilde{w}^{-1}(\mathbf{r} - \mathbf{r}') \tilde{\Delta}\tilde{\psi}(\mathbf{r}') \tilde{w}^{-1}(\mathbf{s} - \mathbf{s}') \tilde{\Delta}\tilde{\psi}(\mathbf{s}') \\ &= \frac{1}{\gamma^2} \Delta F_{\text{df}}[\tilde{\rho}, \tilde{\Delta}\tilde{\psi}] \end{aligned} \quad (\text{C10})$$

because of

$$\begin{aligned} \overline{\tilde{\varphi}(\mathbf{r})} &= 0, \\ \overline{\tilde{\varphi}(\mathbf{r}) \tilde{\varphi}(\mathbf{s})} &= \tilde{w}(\mathbf{r} - \mathbf{s}). \end{aligned} \quad (\text{C11})$$

The strong-coupling approximation of the last equality in eqn (C8) further validates that

$$\begin{aligned} \frac{1 - \frac{1}{\gamma} \mathcal{E}[\tilde{\rho}, \tilde{\varphi}]}{1 - \frac{1}{\gamma} \overline{\mathcal{E}[\tilde{\rho}, \tilde{\varphi}]}} &\approx 1 - \frac{1}{\gamma} (\mathcal{E}[\tilde{\rho}, \tilde{\varphi}] - \overline{\mathcal{E}[\tilde{\rho}, \tilde{\varphi}]}) \\ &= 1 - \frac{i}{\gamma} \mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}] - \frac{1}{2\gamma^2} \Delta \mathcal{E}_1[\tilde{\rho}, \tilde{\varphi}] \\ &= 1 - \frac{i}{\gamma} \mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}] + \mathcal{O}[\gamma^{-2}], \end{aligned} \quad (\text{C12})$$

where

$$\begin{aligned} \Delta \mathcal{E}_1[\tilde{\rho}, \tilde{\varphi}] &\equiv \iiint d\mathbf{r} d\mathbf{r}' d\mathbf{s} d\mathbf{s}' \{ \tilde{\varphi}(\mathbf{r}) \tilde{\varphi}(\mathbf{s}) - w(\mathbf{r} - \mathbf{s}) \} \\ &\quad \times \tilde{w}^{-1}(\mathbf{r} - \mathbf{r}') \tilde{\Delta}\tilde{\psi}(\mathbf{r}') \tilde{w}^{-1}(\mathbf{s} - \mathbf{s}') \tilde{\Delta}\tilde{\psi}(\mathbf{s}'). \end{aligned} \quad (\text{C13})$$

Eqn (C12) implies that the above contribution $\Delta \mathcal{E}_1[\tilde{\rho}, \tilde{\varphi}]$ is ignored. Combining eqn (C8) and (C12), we have

$$\begin{aligned} \overline{\zeta[\tilde{\varphi}]} &= \zeta_c + \zeta_r[\tilde{\varphi}] \\ &= \zeta_c + \overline{\zeta_r[\tilde{\varphi}]} - \frac{i}{\gamma} \overline{\tilde{\varphi}[\tilde{\varphi}] \mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}]} + \mathcal{O}[\gamma^{-2}], \end{aligned} \quad (\text{C14})$$

according to the $1/\gamma$ expansion method. Incidentally, it follows from eqn (C14) that

$$\begin{aligned} \overline{\tilde{\varphi}(\mathbf{r}) \tilde{\varphi}(\mathbf{r}')} &= \overline{\tilde{\varphi}(\mathbf{r}) \tilde{\varphi}(\mathbf{r}')} - \frac{i}{\gamma} \overline{\tilde{\varphi}(\mathbf{r}) \tilde{\varphi}(\mathbf{r}') \mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}]} + \mathcal{O}[\gamma^{-2}] \\ &= \tilde{w}(\mathbf{r} - \mathbf{r}') + \mathcal{O}[\gamma^{-2}], \end{aligned} \quad (\text{C15})$$

thereby justifying eqn (76).

In the remaining subsections, we will evaluate eqn (74) and (82) based on the above strong-coupling approximation represented by eqn (C14).

2. Derivation of eqn (74) from eqn (73)

Following the general form (C6), we classify $\lambda_{\text{non}}[\tilde{\rho}, \tilde{\varphi}]$ given by eqn (73) into the $\tilde{\varphi}$ -independent part ζ_c and the $\tilde{\varphi}$ -dependent contribution $\gamma \zeta_1[\tilde{\varphi}]$:

$$\begin{aligned} \lambda_{\text{non}}[\tilde{\rho}, \tilde{\varphi}] &= \zeta_c + \gamma \zeta_1[\tilde{\varphi}], \\ \zeta_c &= \ln \frac{\tilde{\rho}(\mathbf{r})}{\gamma^2} + 1 + \frac{\gamma^2 \tilde{w}(0)}{2}, \end{aligned} \quad (\text{C16})$$

$$\gamma \zeta_1[\tilde{\varphi}] = i\gamma \tilde{\varphi}(\mathbf{r}) - \left\{ \tilde{\rho}(\mathbf{r}) e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})} \right\}',$$

where the definition of $\{\cdot\}'$ is the same as that of eqn (A17). For later convenience, we also introduce an extended form of $e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})}$:

$$\gamma u_{\text{exp}}[\tilde{\varphi}] = e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \int d\mathbf{s} \tilde{\varphi}(\mathbf{s}) \{ \hat{\rho}_1(\mathbf{s}) + m(\mathbf{s}) \}}, \quad (\text{C17})$$

where a test density field $m(\mathbf{s})$ is added to one-particle density $\hat{\rho}_1(\mathbf{s}) = \delta(\mathbf{s} - \mathbf{r})$ that represents a single sphere located at \mathbf{r} . The functional differentiation of $u_{\text{exp}}[\tilde{\varphi}]$ with respect to $m(\mathbf{r})$ offers the benefit of the expression (C17):

$$\left. \frac{\gamma \delta u_{\text{exp}}[\tilde{\varphi}]}{\delta m(\mathbf{s})} \right|_{m=0} = i\gamma \tilde{\varphi}(\mathbf{s}) e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})}, \quad (\text{C18})$$

which is available to calculate the third term on the rhs of eqn (C14).

First, the Gaussian integration over the φ -field yields

$$\begin{aligned} \overline{\gamma \zeta_1[\tilde{\varphi}]} &= - \left\{ \tilde{\rho}(\mathbf{r}) e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})} \right\}' \\ &= - e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})} - \tilde{\rho}(\mathbf{r}) \left\{ \frac{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})}{\delta \tilde{\rho}(\mathbf{r})} \right\}' \\ &= - e^{\frac{\gamma^2 \tilde{w}(0)}{2}} e^{i\gamma \int d\mathbf{s} \tilde{\varphi}(\mathbf{s}) \hat{\rho}_1(\mathbf{s})} \\ &= - e^{\frac{\gamma^2 \tilde{w}(0)}{2}} \frac{\gamma^2}{2} \iint d\mathbf{s} d\mathbf{s}' \tilde{w}(\mathbf{s} - \mathbf{s}') \hat{\rho}_1(\mathbf{s}) \hat{\rho}_1(\mathbf{s}') \\ &= -1, \end{aligned} \quad (\text{C19})$$

because of $\gamma = e^{\gamma^2 \tilde{w}(0)/2}$, $\iint d\mathbf{s} d\mathbf{s}' \tilde{w}(\mathbf{s} - \mathbf{s}') \hat{\rho}_1(\mathbf{s}) \hat{\rho}_1(\mathbf{s}') = \tilde{w}(0)$, $e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})} = 1$, and the similar approximation to eqn (A19).

Next, we investigate the third term on the rhs of eqn (C14), or $\overline{\zeta_1[\tilde{\varphi}] \mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}]}$. The expression (C4) of $\mathcal{E}_0[\tilde{\rho}, \tilde{\varphi}]$ implies the necessity of evaluating the following contribution:

$$\frac{i}{\gamma} \overline{\tilde{\varphi}(\mathbf{s}) \gamma \zeta_1[\tilde{\varphi}]} = - \overline{\tilde{\varphi}(\mathbf{s}) \tilde{\varphi}(\mathbf{r})} - \frac{i}{\gamma} \overline{\tilde{\varphi}(\mathbf{s}) e^{\frac{\gamma^2 \tilde{w}(0)}{2} + i\gamma \tilde{\varphi}(\mathbf{r})}}. \quad (\text{C20})$$



It follows from eqn (C11) and (C18) that eqn (C20) reads

$$\frac{i\overline{\tilde{\phi}(\mathbf{s})\gamma\zeta_1[\tilde{\phi}]}}{\gamma} = -\tilde{w}(\mathbf{s}-\mathbf{r}) - \frac{1}{\gamma} \left(\frac{\overline{\delta u_{\text{exp}}[\tilde{\phi}]}}{\overline{\delta m(\mathbf{s})}} \right) \Big|_{m=0}. \quad (\text{C21})$$

Since we have

$$\overline{\gamma u_{\text{exp}}[\tilde{\phi}]} = e^{\frac{\gamma^2 \tilde{w}(0)}{2} - \frac{\gamma^2}{2} \iint d\mathbf{s} d\mathbf{s}' \tilde{w}(\mathbf{s}-\mathbf{s}') \{m(\mathbf{s}) + \hat{\rho}_1(\mathbf{s})\} \{m(\mathbf{s}') + \hat{\rho}_1(\mathbf{s}')\}} \quad (\text{C22})$$

in the presence of the test field $m(\mathbf{r})$, the relation (C18) reads

$$\begin{aligned} \frac{\overline{\gamma \delta u_{\text{exp}}[\tilde{\phi}]}}{\overline{\delta m(\mathbf{s})}} \Big|_{m=0} &= -\gamma^2 \int d\mathbf{s}' \tilde{w}(\mathbf{s}-\mathbf{s}') \{m(\mathbf{s}') + \hat{\rho}_1(\mathbf{s}')\} \gamma u_{\text{exp}}[\tilde{\phi}] \Big|_{m=0} \\ &= -\gamma^2 \tilde{w}(\mathbf{s}-\mathbf{r}) e^{\frac{\gamma^2 \tilde{w}(0)}{2} - \frac{\gamma^2}{2} \iint d\mathbf{s} d\mathbf{s}' \tilde{w}(\mathbf{s}-\mathbf{s}') \hat{\rho}_1(\mathbf{s}) \hat{\rho}_1(\mathbf{s}')} \\ &= -\gamma^2 \tilde{w}(\mathbf{s}-\mathbf{r}). \end{aligned} \quad (\text{C23})$$

This ensures that the first term on the rhs of eqn (C21) is canceled by the second term:

$$\overline{\tilde{\phi}(\mathbf{s})\gamma\zeta_1[\tilde{\phi}]} = 0, \quad (\text{C24})$$

thereby implying that

$$\overline{\zeta_1[\tilde{\phi}]\mathcal{E}_0[\tilde{\rho}, \tilde{\phi}]} = 0. \quad (\text{C25})$$

Thus, we find from eqn (C14), (C16), (C19) and (C25)

$$\begin{aligned} \lambda_{\text{non}}[\tilde{\rho}, \tilde{\phi}] &= \zeta_c + \overline{\gamma\zeta_1[\tilde{\phi}]}, \\ &= \zeta_c + \overline{\gamma\zeta_1[\tilde{\phi}]} - i\overline{\zeta_1[\tilde{\phi}]\mathcal{E}_0[\tilde{\rho}, \tilde{\phi}]} + \mathcal{O}[\gamma^{-1}] \\ &= \zeta_c + \overline{\gamma\zeta_1[\tilde{\phi}]} + \mathcal{O}[\gamma^{-1}] \\ &\approx \ln \frac{\tilde{\rho}(\mathbf{r})}{\gamma^2} + 1 + \frac{\gamma^2 \tilde{w}(0)}{2} - 1 \\ &= \ln \frac{\tilde{\rho}(\mathbf{r})}{\gamma^2} + \frac{\gamma^2 \tilde{w}(0)}{2}. \end{aligned} \quad (\text{C26})$$

The above last form is equivalent to the target expression (74).

3. Derivation of eqn (82)

Eqn (81) reads

$$\begin{aligned} \Delta\lambda[\tilde{\rho}, \tilde{\phi}] &= 1 + i\gamma\tilde{\phi}(\mathbf{r}) - \frac{\gamma\delta U[\tilde{\rho}, \tilde{\phi}]/\delta\tilde{\rho}(\mathbf{r})}{1 + U[\tilde{\rho}, \tilde{\phi}]/\gamma} \\ &= 1 + \gamma\zeta_1[\tilde{\phi}] + \zeta_2[\tilde{\phi}] + \mathcal{O}[\gamma^{-1}], \end{aligned} \quad (\text{C27})$$

where $\zeta_1[\tilde{\phi}]$ has been given in eqn (C16) and

$$\begin{aligned} \zeta_2[\tilde{\phi}] &= - \int d\mathbf{r}_2 \tilde{\rho}(\mathbf{r}_2) e^{-v(\mathbf{r}_1-\mathbf{r}_2)} \left\{ \tilde{\rho}(\mathbf{r}_1) e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \hat{\rho}_2(\mathbf{s})} \right\}' \\ &\quad + \int d\mathbf{r}' \tilde{\rho}(\mathbf{r}') e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \hat{\rho}'_1(\mathbf{s})} \left\{ \tilde{\rho}(\mathbf{r}) e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \hat{\rho}_1(\mathbf{s})} \right\}', \end{aligned} \quad (\text{C28})$$

where the definition of $\{\dots\}'$ is the same as that of eqn (A17). The non-equilibrium chemical potential difference $\Delta\lambda[\rho]$ is obtained from averaging $\overline{\Delta\lambda[\tilde{\rho}, \tilde{\phi}]}$ over the $\tilde{\phi}$ -field in a similar manner to the strong-coupling approximation adopted in

eqn (C26):

$$\begin{aligned} \Delta\lambda[\rho] &= \overline{\Delta\lambda[\tilde{\rho}, \tilde{\phi}]} \\ &= 1 + \overline{\gamma\zeta_1[\tilde{\phi}]} + \overline{\zeta_2[\tilde{\phi}]} - i\overline{\zeta_1[\tilde{\phi}]\mathcal{E}_0[\tilde{\rho}, \tilde{\phi}]} + \mathcal{O}[\gamma^{-1}] \\ &= 1 - 1 + \overline{\zeta_2[\tilde{\phi}]} + \mathcal{O}[\gamma^{-1}] \\ &\approx - \int d\mathbf{r}_2 \tilde{\rho}(\mathbf{r}_2) e^{-v(\mathbf{r}_1-\mathbf{r}_2)} e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \hat{\rho}_2(\mathbf{s})} \\ &\quad - \int d\mathbf{r}_2 \tilde{\rho}(\mathbf{r}_2) e^{-v(\mathbf{r}_1-\mathbf{r}_2)} \tilde{\rho}(\mathbf{r}_1) \frac{\overline{\delta e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \hat{\rho}_2(\mathbf{s})}}}{\overline{\delta \tilde{\rho}(\mathbf{r}_1)}} \\ &\quad + \int d\mathbf{r}' \tilde{\rho}(\mathbf{r}') e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s})} \left\{ \hat{\rho}_1(\mathbf{s}) \hat{\rho}'_1(\mathbf{s}) \right\}, \end{aligned} \quad (\text{C29})$$

where we have used the results, $\overline{\gamma\zeta_1[\tilde{\phi}]} = -1$ and $\overline{\zeta_1[\tilde{\phi}]\mathcal{E}_0[\tilde{\rho}, \tilde{\phi}]} = 0$, given by eqn (C19) and (C25), respectively,

and $\delta \left\{ e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \left\{ \hat{\rho}_1 + \hat{\rho}'_1 \right\}} \right\} / \delta \rho = 0$ due to the neglect of the density dependence of $e^{-w(0)} = 1/\gamma^2$ as before. It follows from the Gaussian integration when performing the averages in the last equality of eqn (C29) that

$$\begin{aligned} \overline{e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \hat{\rho}_2(\mathbf{s})}} &= e^{-\frac{\gamma^2}{2} \iint d\mathbf{s} d\mathbf{s}' \tilde{w}(\mathbf{s}-\mathbf{s}') \hat{\rho}_2(\mathbf{s}) \hat{\rho}_2(\mathbf{s}')} \\ &= e^{-\gamma^2 \iint d\mathbf{s} d\mathbf{s}' \tilde{w}(\mathbf{s}-\mathbf{s}') \left\{ \delta(\mathbf{s}-\mathbf{r}_1) \delta(\mathbf{s}'-\mathbf{r}_2) + \sum_{i=1}^2 \frac{\delta(\mathbf{s}-\mathbf{r}_i) \delta(\mathbf{s}'-\mathbf{r}_i)}{2} \right\}} \\ &= e^{-\gamma^2 \tilde{w}(\mathbf{r}_1-\mathbf{r}_2) - \gamma^2 \tilde{w}(0)} \\ &= \frac{1}{\gamma^2} e^{-\gamma^2 \tilde{w}(\mathbf{r}_1-\mathbf{r}_2)}, \end{aligned} \quad (\text{C30})$$

$$\frac{\overline{\delta e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \hat{\rho}_2(\mathbf{s})}}}{\overline{\delta \tilde{\rho}(\mathbf{r}_1)}} = \frac{\delta(\mathbf{r}_1-\mathbf{r}_2)}{\tilde{\rho}^2(\mathbf{r}_1)} e^{-\gamma^2 \tilde{w}(\mathbf{r}_1-\mathbf{r}_2)}, \quad (\text{C31})$$

and

$$\begin{aligned} \overline{e^{i\gamma \int d\mathbf{s} \tilde{\phi}(\mathbf{s}) \left\{ \hat{\rho}_1(\mathbf{s}) \hat{\rho}'_1(\mathbf{s}') \right\}}} &= e^{-\frac{\gamma^2}{2} \iint d\mathbf{s} d\mathbf{s}' \tilde{w}(\mathbf{s}-\mathbf{s}') \left\{ \hat{\rho}_1(\mathbf{s}) \hat{\rho}_1(\mathbf{s}') + \hat{\rho}'_1(\mathbf{s}) \hat{\rho}'_1(\mathbf{s}') \right\}} \\ &= e^{-\frac{\gamma^2 \tilde{w}(0)}{2} - \frac{\gamma^2 \tilde{w}(0)}{2}} \\ &= \frac{1}{\gamma^2}, \end{aligned} \quad (\text{C32})$$

respectively. In eqn (C32), it is noted that the cross terms vanish: $\hat{\rho}_1(\mathbf{s}) \hat{\rho}'_1(\mathbf{s}') = \hat{\rho}'_1(\mathbf{s}) \hat{\rho}_1(\mathbf{s}') = 0$ because the one-particle densities, $\hat{\rho}_1(\mathbf{s}) = \delta(\mathbf{s}-\mathbf{r})$ and $\hat{\rho}'_1(\mathbf{s}) = \delta(\mathbf{s}-\mathbf{r}')$, represent the instantaneous densities of the target particle located at different positions of \mathbf{r} and \mathbf{r}' and a single sphere is unable to simultaneously exist at different locations.



Substituting eqn (C30)–(C32) into eqn (C29), we obtain

$$\begin{aligned}\Delta\lambda[\rho] &= -\frac{1}{\gamma^2}\int d\mathbf{r}_2\tilde{\rho}(\mathbf{r}_2)e^{-v(\mathbf{r}_1-\mathbf{r}_2)-w(\mathbf{r}_1-\mathbf{r}_2)} \\ &\quad -e^{-v(0)-\gamma^2\tilde{w}(0)}+\frac{1}{\gamma^2}\int d\mathbf{r}'\tilde{\rho}(\mathbf{r}') \\ &= \frac{1}{\gamma^2}\int d\mathbf{r}'\left\{1-e^{-v(\mathbf{r}-\mathbf{r}')-\gamma^2\tilde{w}(\mathbf{r}-\mathbf{r}')}\right\}\tilde{\rho}(\mathbf{r}')-\frac{e^{-v(0)}}{\gamma^2} \\ &= \int d\mathbf{r}'\left\{1-e^{-v(\mathbf{r}-\mathbf{r}')-w(\mathbf{r}-\mathbf{r}')}\right\}\rho(\mathbf{r}')+\mathcal{O}[\gamma^{-2}],\end{aligned}\quad (\text{C33})$$

hence verifying eqn (82).

Acknowledgements

The author thanks the anonymous referees for their valuable comments and suggestions.

References

- 1 S. Torquato, *J. Chem. Phys.*, 2018, **149**, 020901.
- 2 S. Torquato, *Phys. Rep.*, 2018, **745**, 1.
- 3 J. Kim and S. Torquato, *Proc. Natl. Acad. Sci. U. S. A.*, 2020, **117**, 8764–8774.
- 4 G. J. Aubry, L. S. Froufe-Pérez, U. Kuhl, O. Legrand, F. Scheffold and F. Mortessagne, *Phys. Rev. Lett.*, 2020, **125**, 127402.
- 5 S. Torquato and F. H. Stillinger, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2003, **68**, 041113.
- 6 A. Donev, F. H. Stillinger and S. Torquato, *Phys. Rev. Lett.*, 2005, **95**, 090604.
- 7 A. Donev, S. Torquato and F. H. Stillinger, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2005, **71**, 011105.
- 8 S. Atkinson, F. H. Stillinger and S. Torquato, *Proc. Natl. Acad. Sci. U. S. A.*, 2014, **52**, 18436–18441.
- 9 C. E. Zachary, Y. Jiao and S. Torquato, *Phys. Rev. Lett.*, 2011, **106**, 178001.
- 10 C. E. Zachary, Y. Jiao and S. Torquato, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2011, **83**, 051308.
- 11 S. Atkinson, F. H. Stillinger and S. Torquato, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2013, **88**, 062208.
- 12 Y. Kallus and S. Torquato, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2014, **90**, 022114.
- 13 S. Atkinson, F. H. Stillinger and S. Torquato, *Phys. Rev. E*, 2016, **94**, 032902.
- 14 S. Atkinson, G. Zhang, A. B. Hopkins and S. Torquato, *Phys. Rev. E*, 2016, **94**, 012902.
- 15 J. Kim and S. Torquato, *Phys. Rev. B*, 2018, **97**, 054105.
- 16 G. Zhang and S. Torquato, *Phys. Rev. E*, 2020, **101**, 032124.
- 17 S. Torquato, *Phys. Rev. E*, 2021, **103**, 052126.
- 18 M. van Hecke, *J. Phys.: Condens. Matter*, 2009, **22**, 033101.
- 19 A. J. Liu and S. R. Nagel, *Annu. Rev. Condens. Matter Phys.*, 2010, **1**, 347–369.
- 20 M. Müller and M. Wyart, *Annu. Rev. Condens. Matter Phys.*, 2015, **6**, 177–200.
- 21 V. Lubchenko and P. G. Wolynes, *J. Phys. Chem. B*, 2017, **122**, 3280–3295.
- 22 V. Lubchenko, *Adv. Phys. X*, 2018, **3**, 1510296.
- 23 H. Mizuno, H. Shiba and A. Ikeda, *Proc. Natl. Acad. Sci. U. S. A.*, 2017, **114**, E9767–E9774.
- 24 M. Shimada, H. Mizuno, M. Wyart and A. Ikeda, *Phys. Rev. E*, 2018, **98**, 060901.
- 25 H. Mizuno, M. Shimada and A. Ikeda, *Phys. Rev. Res.*, 2020, **2**, 013215.
- 26 H. Mizuno, H. Tong, A. Ikeda and S. Mossa, *J. Chem. Phys.*, 2020, **153**, 154501.
- 27 M. Shimada, H. Mizuno and A. Ikeda, *Soft Matter*, 2020, **16**, 7279–7288.
- 28 E. Lerner, G. Düring and M. Wyart, *Soft Matter*, 2013, **9**, 8252–8263.
- 29 E. Lerner, G. Düring and E. Bouchbinder, *Phys. Rev. Lett.*, 2016, **117**, 035501.
- 30 D. Richard, K. González-López, G. Kapteijns, R. Pater, T. Vaknin, E. Bouchbinder and E. Lerner, *Phys. Rev. Lett.*, 2020, **125**, 085502.
- 31 A. Moriel, Y. Lubomirsky, E. Lerner and E. Bouchbinder, *Phys. Rev. E*, 2020, **102**, 033008.
- 32 C. Rainone, E. Bouchbinder and E. Lerner, *Proc. Natl. Acad. Sci. U. S. A.*, 2020, **117**, 5228–5234.
- 33 C. Rainone, P. Urbani, F. Zamponi, E. Lerner and E. Bouchbinder, *SciPost Phys.*, 2021, **4**, 008.
- 34 L. Berthier, P. Charbonneau, Y. Jin, G. Parisi, B. Seoane and F. Zamponi, *Proc. Natl. Acad. Sci. U. S. A.*, 2016, **113**, 8397–8401.
- 35 L. Wang, A. Ninarello, P. Guan, L. Berthier, G. Szamel and E. Flenner, *Nat. Commun.*, 2019, **10**, 1–7.
- 36 L. Berthier, G. Biroli, P. Charbonneau, E. I. Corwin, S. Franz and F. Zamponi, *J. Chem. Phys.*, 2019, **151**, 010901.
- 37 X. Tan, Y. Guo, D. Huang and L. Zhang, *Soft Matter*, 2021, **17**, 1330–1336.
- 38 M. Ozawa, L. Berthier and D. Coslovich, *SciPost Phys.*, 2017, **3**, 027.
- 39 A. T. Chieco, M. Zu, A. J. Liu, N. Xu and D. J. Durian, *Phys. Rev. E*, 2018, **98**, 042606.
- 40 L. E. Silbert and M. Silbert, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2009, **80**, 041304.
- 41 Y. Wu, P. Olsson and S. Teitel, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2015, **92**, 052206.
- 42 A. Ikeda and L. Berthier, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2015, **92**, 012309.
- 43 A. Ikeda, L. Berthier and G. Parisi, *Phys. Rev. E*, 2017, **95**, 052125.
- 44 M. te Vrugt, H. Löwen and R. Wittkowski, *Adv. Phys.*, 2020, **69**, 121–247.
- 45 D. S. Dean, *J. Phys. A: Math. Gen.*, 1996, **29**, L613.
- 46 T. Leonard, B. Lander, U. Seifert and T. Speck, *J. Chem. Phys.*, 2013, **139**, 204109.
- 47 B. Kim, K. Kawasaki, H. Jacquin and F. van Wijland, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2014, **89**, 012150.
- 48 H. Jacquin, B. Kim, K. Kawasaki and F. van Wijland, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2015, **91**, 022130.



- 49 N. Bidhoodi and S. P. Das, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2015, **92**, 012325.
- 50 D. S. Dean, B. S. Lu, A. C. Maggs and R. Podgornik, *Phys. Rev. Lett.*, 2016, **116**, 240602.
- 51 V. Démery and D. S. Dean, *J. Stat. Mech.: Theo. Exp.*, 2016, 023106.
- 52 M. Krüger and D. S. Dean, *J. Chem. Phys.*, 2017, **146**, 134507.
- 53 J. F. Lutsko, *Sci. Adv.*, 2019, **5**, eaav7399.
- 54 H. Frusawa, *J. Phys. A: Math. Theo.*, 2019, **52**, 065003.
- 55 H. Frusawa, *Entropy*, 2020, **22**, 34.
- 56 S. Mahdisoltani and R. Golestanian, *Phys. Rev. Lett.*, 2021, **126**, 158002.
- 57 T. V. Ramakrishnan and M. Yussouff, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1979, **19**, 2775.
- 58 R. Evans, *Adv. Phys.*, 1979, **28**, 143–200.
- 59 Y. Singh, *Phys. Rep.*, 1991, **207**, 351–444.
- 60 C. N. Likos, *Phys. Rep.*, 2001, **348**, 267–439.
- 61 J. F. Lutsko, *Adv. Chem. Phys.*, 2010, **144**, 1.
- 62 Y. Singh, J. P. Stoessel and P. G. Wolynes, *Phys. Rev. Lett.*, 1985, **54**, 1059.
- 63 M. Baus and J. L. Colot, *J. Phys. C: Solid State Phys.*, 1986, **19**, L135.
- 64 R. W. Hall and P. G. Wolynes, *J. Chem. Phys.*, 1987, **86**, 2943–2948.
- 65 C. Dasgupta, *Europhys. Lett.*, 1992, **20**, 131.
- 66 R. K. Murarka and B. Bagchi, *J. Chem. Phys.*, 2001, **115**, 5513–5520.
- 67 C. Kaur and S. P. Das, *Phys. Rev. Lett.*, 2001, **86**, 2062.
- 68 K. Kim and T. Munakata, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2003, **68**, 021502.
- 69 P. Chaudhuri, S. Karmakar, C. Dasgupta, H. R. Krishnamurthy and A. K. Sood, *Phys. Rev. Lett.*, 2005, **95**, 248301.
- 70 P. Chaudhuri, S. Karmakar and C. Dasgupta, *Phys. Rev. Lett.*, 2005, **100**, 125701.
- 71 B. S. Gupta, L. Premkumar and S. P. Das, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2012, **85**, 051501.
- 72 L. Premkumar, N. Bidhoodi and S. P. Das, *J. Chem. Phys.*, 2001, **114**, 124511.
- 73 T. Odagaki, *J. Phys. Soc. Jpn.*, 2017, **86**, 082001.
- 74 A. Mondal and S. P. Das, *Prog. Theor. Exp. Phys.*, 2020, 073102.
- 75 R. P. Behringer and B. Chakraborty, *Rep. Prog. Phys.*, 2018, **82**, 012601.
- 76 A. Baule, F. Morone, H. J. Herrmann and H. A. Makse, *Rev. Mod. Phys.*, 2018, **90**, 015006.
- 77 P. Charbonneau, E. I. Corwin, G. Parisi and F. Zamponi, *Phys. Rev. Lett.*, 2015, **114**, 125504.
- 78 S. Franz, G. Parisi, P. Urbani and F. Zamponi, *Proc. Natl. Acad. Sci. U. S. A.*, 2015, **112**, 14539–14544.
- 79 P. Charbonneau, E. I. Corwin, G. Parisi, A. Poncet and F. Zamponi, *Phys. Rev. Lett.*, 2016, **117**, 045503.
- 80 D. Hexner, A. J. Liu and S. R. Nagel, *Phys. Rev. E*, 2018, **97**, 063001.
- 81 S. Franz, A. Sclocchi and P. Urbani, *SciPost Phys.*, 2020, **9**, 012.
- 82 P. Charbonneau, E. I. Corwin, R. C. Dennis, R. D. H. Rojas, H. Ikeda, G. Parisi and F. Ricci-Tersenghi, *Phys. Rev. E*, 2021, **104**, 014102.
- 83 S. A. Ridout, J. W. Rocks and A. J. Liu, 2020, arXiv preprint, arXiv:2011.13049.
- 84 P. Charbonneau, E. I. Corwin, G. Parisi and F. Zamponi, *Phys. Rev. Lett.*, 2012, **109**, 205501.
- 85 P. Charbonneau, J. Kurchan, G. Parisi, P. Urbani and F. Zamponi, *Nat. Commun.*, 2014, **5**, 1–6.
- 86 Q. Liao and L. Berthier, *Phys. Rev. X*, 2019, **9**, 011049.
- 87 C. Artiacco, P. Baldan and G. Parisi, *Phys. Rev. E*, 2020, **101**, 052605.
- 88 F. H. Stillinger, P. G. Debenedetti and S. Sastry, *J. Chem. Phys.*, 1998, **109**, 3983–3988.
- 89 P. G. Debenedetti and F. H. Stillinger, *Nature*, 2001, **410**, 259–267.
- 90 A. Heuer, *J. Phys.: Condens. Matter*, 2008, **20**, 373101.
- 91 M. D. Ediger and P. Harrowell, *J. Chem. Phys.*, 2012, **137**, 080901.
- 92 H. Frusawa, *J. Stat. Mech.: Theo. Exp.*, 2021, 013213.
- 93 D. Henderson, *Condens. Matter Phys.*, 2009, **12**, 127.
- 94 D. Frydel and M. Ma, *Phys. Rev. E*, 2016, **93**, 062112.
- 95 H. Frusawa, *Phys. Rev. E*, 2020, **101**, 012121.
- 96 R. R. Netz, *Eur. Phys. J. E: Soft Matter Biol. Phys.*, 2001, **5**, 557.
- 97 A. Gabrielli, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2004, **70**, 066131.
- 98 J. N. Nampoothiri, Y. Wang, K. Ramola, J. Zhang, S. Bhattacharjee and B. Chakraborty, *Phys. Rev. Lett.*, 2020, **125**, 118002.
- 99 H. Tong, S. Sengupta and H. Tanaka, *Nat. Commun.*, 2020, **11**, 1–10.
- 100 E. Lerner, *J. Chem. Phys.*, 2020, **153**, 216101.
- 101 M. Shimada and E. De Giuli, 2020, arXiv preprint, arXiv:2008.11896.
- 102 D. Hexner, A. J. Liu and S. R. Nagel, *Phys. Rev. Lett.*, 2018, **121**, 115501.
- 103 P. Rissone, E. I. Corwin and G. Parisi, *Phys. Rev. Lett.*, 2021, **127**, 038001.
- 104 H. Frusawa and R. Hayakawa, *J. Phys. A: Math. Gen.*, 2000, **33**, L155.

