Properties of patchy colloidal particles close to a surface: A Monte Carlo and density functional study

Nicoletta Gnan,1, a) Daniel de las Heras,2 José Maria Tavares,3 Margarida M. Telo da Gama,4 and Francesco Sciortino1

1Dipartimento di Fisica, Università di Roma “La Sapienza”, Piazzale A. Moro 2, 00185 Roma, Italy
2Theoretische Physik II, Physikalisches Institut, Universität Bayreuth, D-95440 Bayreuth, Germany
3Instituto Superior de Engenharia de Lisboa, Rua Conselheiro Emídio Navarro, P-1590-062 Lisbon, Portugal
4Departamento de Física, Faculdade de Ciências da Universidade de Lisboa, Campo Grande, P-1749-016 Lisbon, Portugal and Centro de Física Teórica e Computacional da Universidade de Lisboa, Avenida Professor Gama Pinto 2, P-1649-003 Lisbon, Portugal

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We investigate the behavior of a patchy particle model close to a hard-wall via Monte Carlo simulation and density functional theory (DFT). Two DFT approaches, based on the homogeneous and inhomogeneous versions of Wertheim’s first order perturbation theory for the association free energy are used. We evaluate, by simulation and theory, the equilibrium bulk phase diagram of the fluid and analyze the surface properties for two isotherms, one of which is close to the liquid side of the gas-liquid coexistence curve. We find that the density profile near the wall crosses over from a typical high-temperature adsorption profile to a low-temperature desorption one, for the isotherm close to coexistence. We relate this behavior to the properties of the bulk network liquid and find that the theoretical descriptions are reasonably accurate in this regime. At very low temperatures, however, an almost fully bonded network is formed, and the simulations reveal a second adsorption regime which is not captured by DFT. We trace this failure to the neglect of orientational correlations of the particles, which are found to exhibit surface induced orientational order in this regime. © 2012 American Institute of Physics. [http://dx.doi.org/10.1063/1.4746428]

I. INTRODUCTION

Colloidal particles with patterned surfaces—better known as patchy particles1–4—have been studied extensively in recent years owing to their ability to self-assemble in a rich number of cluster, gel, glassy, and crystalline phases. Understanding how the surface pattern influences the self-assembly mechanism is crucial to a bottom-up strategy for designing new materials where the desired macroscopic behavior is encoded in the microscopic properties of the building-blocks.5

Patchy particles represent a valuable model system for investigating and understanding the behavior of more complex constituents such as amphiphilic molecules, colloidal clays, proteins, and DNA nano-assemblies.6–10 New concepts, as equilibrium gel,5 optimal network density,11 and empty liquid have arisen from the study of the phase diagrams12,13 of patchy particles with limited valence.14,15 These models emphasize the role of the number of bonds between the particles in determining the equilibrium as well as the static and dynamic behavior of the system in and out of equilibrium.16

Homogeneous patchy particle fluids are described satisfactorily by Wertheim’s first order thermodynamic perturbation theory17–20 that provides an expression for the free energy per particle with \( n_p \) patches, which are treated independently. Less known are the properties of patchy particle fluids in confined geometries. Understanding the behavior of patchy particles close to surfaces has direct impact on a number of different applications which require patchy particles to self-assemble in confined geometries. We recall for instance the templated self-assembly technique21 where confined geometries are used to orient bulk structures or to induce the formation of novel morphologies. Moreover recent studies have focused on effective forces between colloids generated by confined critical patchy particles aiming to control colloid stability.22 Finally, such systems may be studied with small-angle neutron scattering23 and atomic force microscopy24 and thus a quantitative microscopic description of patchy particles in confined geometries is highly desirable.

The description of confined associating fluids has been addressed in the past. Density functional theory (DFT) based on a perturbation of the inhomogeneous hard-sphere (HS) fluid was used to describe the structure of inhomogeneous associating fluids. An early example is the work of Segura et al.25 for particles with four patches close to a hard wall where the weighted density approximation (WDA) of Tarazona26,27 was combined with Wertheim’s theory to obtain a DFT description of associating fluids. In the original DFT formulation both homogeneous and inhomogeneous versions of Wertheim’s first order perturbation theory were used to account for the association free energy of the particles and it was found that the homogeneous theory—where the law of mass action is identical to that of the bulk system with the

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a)nicoletta.gnan@roma1.infn.it.
density replaced by the averaged local density—yields satisfactory results over the whole range of parameters. By con-
trast, the inhomogeneous version of Wertheim’s theory was found to overestimate badly the layering of the particles near
the wall. The reasons for this discrepancy are not fully under-
stood but they appear to be related to the difficulty of imple-
menting Wertheim’s law of mass action in the inhomogeneous
system. This approach was extended to mixtures of associat-
ing and neutral equi-sized hard-spheres.28

A second approach for the same model—particles with
four patches—was developed by Yu and Wu.29 They employed
a modified fundamental measure theory (FMT)30 based on
Rosenfeld’s functional for inhomogeneous hard-sphere fluids
and an inhomogeneous version of Wertheim’s free energy for
association. Yu and Wu30 employed an inhomogeneous ver-
sion of the law of mass action which was shown to give
comparable results to the homogeneous theory of Segura et al. for
intermediate densities. Yu and Wu’s theory, however, is more
accurate at the highest densities and can be extended to binary
mixtures of particles with different sizes. More importantly, it
reveals that the inhomogeneous version of Wertheim’s first
order perturbation theory depends crucially on the general-
zation of the law of mass action for inhomogeneous patchy
particle systems.

Both theories were applied to systems where bonding is
not fully developed, i.e., at temperatures that are not too low,
and the relation between surface and bulk properties has not
been investigated. In this paper, we focus on the self-assembly
of patchy particles with \( n_p = 3 \) at a planar hard wall, and ex-
tend our calculations to the region where a fully bonded opti-
nal network develops. We have simulated the system by fix-
ing the density and scanning the temperature down to very low
temperatures. We compare the simulation results to the results of
two density functional theories: a homogeneous WDA ap-
proach similar to that of Segura et al. and the homogeneous
FMT approach of Yu and Wu, which were shown to be equiva-
 lent for hard-spheres in the range of densities considered here.
For the three-patch particle model the theories give similar
results at moderate to high temperatures, by contrast to the
low-temperature regime, relevant to the formation of arrested
states, where both theories break down. We trace this failure
to the assumption of independent patches and stress the need
for the development of density functionals that account for the
directionality of the bonding, which plays a crucial role in the
low-temperature regime.

The paper is organized as follows: in Sec. II we describe
the model and the simulation techniques. Moreover, a brief
comparison of the results of Wertheim’s theory and the simu-
lation results for the bulk fluid are presented and discussed.
In Sec. III, we describe the two density functional approaches
and in Sec. IV we discuss their validity and limitations, with
emphasis on the low-temperature regime.

II. BACKGROUND

A. Model and simulation methods

Patchy colloidal particles are modeled as hard spheres of
diameter \( \sigma \) with \( n_p = 3 \) equidistant bonding sites on the par-
ticle equator (see the inset of Fig. 1). The interaction between
two patches \( V_{ij} \) belonging to particles \( i \) and \( j \) is given by the
Kern-Frenkel potential:31

\[
V_{ij} = V_{SW}(|\vec{r}_{ij}|) G(\vec{r}_{i}, \vec{r}_{j}),
\]

where \( \vec{r}_{ij} \) is the vector between the centers of particles \( i \) and \( j \),
and \( \vec{r}_{i} \) is the unit vector from the center of particle \( i \) to the cen-
ter of one patch on its surface. \( V_{SW} \) is a square well potential,

\[
V_{SW}(|\vec{r}_{ij}|) = \begin{cases}
\infty & \text{if } |\vec{r}_{ij}| < \sigma, \\
-\epsilon & \text{if } \sigma \leq |\vec{r}_{ij}| \leq \sigma + \delta, \\
0 & \text{otherwise,}
\end{cases}
\]

and \( G \) is the angular part:

\[
G(\vec{r}_{ij}, \vec{r}_{i}, \vec{r}_{j}) = \begin{cases}
1 & \text{if } \vec{r}_{i} \cdot \vec{r}_{j} > \cos(\theta_{\text{max}}), \\
-\vec{r}_{ij} \cdot \vec{r}_{j} > \cos(\theta_{\text{max}}), \\
0 & \text{otherwise.}
\end{cases}
\]

The interaction energy between sites \( \epsilon \) sets the energy scale.
The spatial range \( \delta \) and the angle \( \theta_{\text{max}} \) control the volume
available for bonding, \( v_b \), which is

\[
v_b = \frac{\pi \sigma^3}{3}[(1 + \delta/\sigma)^3 - 1][1 - \cos(\theta_{\text{max}})]^2.
\]

We fix the parameters \( \delta = 0.119\sigma \) and \( \cos\theta_{\text{max}} = 0.895 \),
fulfilling the single bond per patch condition assumed in
Wertheim’s first order perturbation theory.

We perform Gibbs ensemble Monte Carlo (GEMC)32
simulations to locate the gas-liquid coexistence line and
grand-canonical Monte Carlo (GCMC) simulations33 to es-
imate the critical point. In the GEMC method, a Monte Carlo
(MC) step consists on average of 4000 roto-translation at-
ttempts, 400 particle swap attempts and one volume change.
About 300 particles in a volume \( V = 2880\sigma^3 \) were simulated.
In the GCMC method, we consider boxes with \( L = 6\sigma \) to

![FIG. 1. Phase diagram of patchy particles with valence \( n_p = 3 \). Symbols are
results from MC simulations. Squares depict the gas-liquid coexistence line.
Dashed lines indicate the isochores investigated in this study. Patchy particles
are modeled as hard spheres with 3 equidistant bonding sites (patches) on the
particle equator (inset).](http://jcp.aip.org/doi/abs/10.1063/1.4735255)
The bonding probability model, the bonding contribution can be expressed in terms of the distribution of the Ising universality class.34, 35

The surface properties are investigated by canonical ensemble MC simulations, at different temperatures and densities, by introducing a surface in the middle of the simulation box and periodic boundary conditions. The surface is modeled by a planar hard wall acting on the particles through a hard-core repulsion, i.e., the interaction between the particles and the surface is purely entropic. The density profiles close to the wall are calculated at two bulk densities, at temperatures $T$ above and below the percolation line.

B. Wertheim theory

Within Wertheim’s theory, the free energy for a homogeneous system is the sum of two contributions. The free energy of the hard-sphere reference system and the bonding contribution $F_{\text{bond}}(\rho, T)$ which arises from considering certain graphs in the Mayer expansion.36 For the present single-component model, the bonding contribution can be expressed in terms of the bonding probability $p_b$ (fraction of bonded sites) as

$$\beta F_{\text{bond}} / N = n_p \ln(1 - p_b) + \frac{1}{2} n_p p_b,$$

where $\beta = 1/kT$ with $k$ the Boltzmann constant. Assuming that all sites have the same probability of bonding, Wertheim’s theory predicts that $p_b$ is determined by the law of mass action:

$$\frac{p_b}{(1 - p_b)^2} = \rho n_p \Delta,$$

where $\Delta$ is the equilibrium constant of the “reaction” (bonding) between two patches and $\rho$ is the density. To evaluate $\Delta$ for the present model, we assume that the radial distribution function of the reference HS fluid $g_{\text{HS}}(r)$ is approximated by:

$$g_{\text{HS}}(r) = (A_0 + A_1) + A_1(r/\sigma - 1),$$

where

$$A_0 = \frac{1 - 0.5\eta}{(1 - \eta)^3} + \frac{4.5\eta(1 + \eta)}{(1 - \eta)^3},$$

$$A_1 = \frac{-4.5\eta(1 + \eta)}{(1 - \eta)^3}. $$

As a result, $\Delta = 4\pi / \chi^3 \left[ \frac{(1 + \delta)^3}{3} - A_0 + \frac{(1 + \delta)^4}{4} - A_1 \right] \times [\exp(\beta \varepsilon) - 1].$ where $\delta = 0.5(1 - \cos(\theta_{\text{max}}))$ is the fraction of surface covered by the patch.

C. Bulk behavior

Figure 1 shows the phase diagram obtained via GCMC and GEMC simulations and compares it with the theoretical results from Wertheim’s theory. The figure also shows the FH percolation line. The high density phase is always percolated, in the sense that there is a non-zero probability of finding an infinite cluster that contains almost all the particles. The percolation line (i.e., the line that separates percolated from non-percolated states) intercepts the binodal on the low-density phase, near the critical point. As shown in Fig. 1, the percolation line starts on the left of the critical point and the temperature increases monotonically with the density. In Fig. 2 we plot the bonding probability as a function of temperature for different densities. The agreement between Wertheim’s theory and Monte Carlo simulations is quite good both for the phase diagram and for the bonding probability.

III. DENSITY FUNCTIONAL THEORY

As usual in density functional theory, we split the Helmholtz free-energy functional into the ideal and excess parts:

$$F[\rho(\vec{r})] = F_{\text{id}}[\rho(\vec{r})] + F_{\text{exc}}[\rho(\vec{r})],$$

where $\rho(\vec{r})$ is the number density. Here and in what follows, we assume that the single-particle distribution function $\rho(\vec{r})$ depends on the spatial but not on the orientational coordinates.
Note that the arrangement of patches on the equator breaks the rotational symmetry of the particles, and therefore states with a preferred orientation of the particles cannot be ruled out. In fact, this is a crude approximation that will break down at low temperatures, as we will discuss later.

The ideal part is given by

$$\beta \mathcal{F}_{id}[\rho(\mathbf{r})] = \int d^3 r \rho(\mathbf{r}) [\ln(\rho(\mathbf{r}))V] - 1, \quad (13)$$

where $V$ is the thermal volume. The integral is over the volume $V$.

$\mathcal{F}_{exc}$ contains the excluded volume interactions between HS and the bonding free energy due to bond formation between the particles. We have used two different approximations for $\mathcal{F}_{exc}$: a modified version 41 of the local WDA introduced by Segura et al. 25 and the FMT for associating fluids developed by Yu and Wu. 29

In the WDA method, the HS and the association term in the free-energy per particle are evaluated for a homogeneous system at the same weighted density. The latter is calculated for a fluid of HS. 41 The Carnahan-Starling 36 approximation is employed for the HS contribution, while the association free energy is given by Wertheim’s first-order perturbation theory for a homogeneous system (Eq. (5)).

In the FMT proposed by Yu and Wu, 29 Wertheim’s free-energy functional for the inhomogeneous system is used and thus the two contributions are treated separately. The HS reference fluid is described by the Rosenfeld FMT approach while the association contribution is given by an appropriate inhomogeneous Wertheim’s term. A detailed description of both methods is found in Appendices A and B, respectively.

**IV. RESULTS**

We focus on the surface properties of the fluid at two bulk densities ($\rho_b \sigma^3 = 0.70$ and $\rho_b \sigma^3 = 0.40$) in contact with a neutral hard-wall, and consider different temperatures (see Fig. 1). The main results are reported in Figs. 3 and 4.

The density profiles calculated from MC simulations for $\rho_b \sigma^3 = 0.70$, are plotted together with FMT and WDA theoretical results in Fig. 3. The limiting case of HS is also included for reference (panel (a) of Fig. 3). Panels (b)–(d) of Fig. 3 show the results for three different reduced temperatures $T$ above and below the percolation line. The main difference between the associating fluid (finite temperature) and the reference HS fluid ($T \to \infty$) is the reduction in the adsorption of particles near the wall. This effect can be understood as follows: at very high temperatures almost all the particles are fully unbonded, and there is a strong adsorption of particles near the wall due to entropic reasons (gain in configurational entropy). As we decrease the temperature, the level of association between particles increases. The probability of bonding at distances of order $\sigma$ from the wall is lower than at larger distances (the wall is neutral). The contribution to the bonding free energy of the particles near the wall is lower than the contribution due to the other particles, resulting in a reduction of the adsorption of patchy particles at the wall when compared to the adsorption of hard-spheres (see the contact density, i.e., the value of the density at $z/\sigma = 0.5$, in panel (f) of Fig. 3).

The WDA provides a fairly good description of the density profiles down to $kT/\epsilon = 0.15$, i.e., well inside the percolation region (see Fig. 1). At this temperature WDA seems slightly more accurate than FMT, providing a good estimate of the contact density. The agreement is lost at $kT/\epsilon = 0.10$ and below (Figs. 3(d) and 3(e)) where both theories underestimate the contact density and do not account for the strong fluid layering on increasing the distance from the wall. At this density, we find that patchy particles are adsorbed on the wall at all $T$ (although the adsorption decreases as the temperature decreases). No changes with further cooling are expected at lower temperatures since $p_b(kT/\epsilon = 0.08) \approx 0.98$, meaning.
FIG. 4. Density profiles as a function of the distance from the wall at different temperatures for $\rho_{b} \sigma^3 = 0.40$: (a) $T \to \infty$, (b) $kT/\varepsilon = 0.30$, (c) $kT/\varepsilon = 0.20$, (d) $kT/\varepsilon = 0.15$, (e) $kT/\varepsilon = 0.12$, (f) $kT/\varepsilon = 0.10$, and (g) $kT/\varepsilon = 0.08$. The full circles are Monte Carlo simulation results. Lines are density functional theory results: FMT (red solid line) and WDA (dashed blue line). (h) Contact density as a function of temperature.
that the liquid has reached an almost fully bonded configuration and thus the structural properties become essentially $T$-independent.\textsuperscript{4,42}

In Fig. 4, panels (a)–(g), we plot the density profiles for $\rho_0\sigma^3 = 0.40$. Panel (a) illustrates the HS fluid corresponding to $T \rightarrow \infty$. At $kT/e = 0.3$ (b) and $kT/e = 0.2$ (c) both, FMT and WDA, are in reasonable agreement with the MC simulation results. FMT yields slightly better results than WDA for $kT/e = 0.2$. Again, the two density profiles indicate an adsorption of particles near the wall. The adsorption, due to the excluded volume, is small compared to the adsorption of HS (Fig. 4(a)). Figure 4(d) illustrates the system at $kT/e = 0.15$ (slightly below the percolation threshold). The MC and FMT density profiles are almost uniform. Patchy particles are slightly desorbed from the wall, indicating the cancellation between the effects due to excluded volume and association. At this temperature WDA predicts a desorption of particles very close to the wall, but it also predicts the formation of a more or less well-defined layer of particles at approximately $z/\sigma = 1$ from the wall. Panels (e)–(g) illustrate the low temperatures: $kT/e = 0.12$ (e), $kT/e = 0.10$ (f), and $kT/e = 0.08$ (g). At these $T$ the association of particles is very high (see the fraction of unbonded sites in Fig. 6), and the energy of bonding dominates the behavior of the system. As a result, FMT predicts a strong desorption of particles from the wall. However, the simulation shows the opposite behavior; there is a well-defined layer of particles near the wall. The layer grows and approaches the wall as the temperature decreases. This desorption-adsorption crossover is also reflected in the contact density depicted in panel (h) of Fig. 4. WDA predicts the presence of a layer of particles at approximately $z \approx \sigma$ for low $T$. Nevertheless, the peak heights do not vary significantly with $T$, by contrast to the simulation results.

The overall $T$ dependence of the contact density at $\rho_0\sigma^3 = 0.40$ is significantly different from the system at $\rho_0\sigma^3 = 0.70$. Indeed, for $\rho_0\sigma^3 = 0.40$, the contact density changes continuously from the HS adsorption limit to the low $T$ desorption, while in the $\rho_0\sigma^3 = 0.70$ system only adsorption is present. To rationalize this behavior we recall the thermodynamics of the bulk system, and in particular the gas-liquid coexistence. In limited valence systems, the liquid side of the gas-liquid coexistence is almost vertical in the $T - \rho$ plane. The associated density provides a quantification of the so-called optimal network density, i.e., the density at which particles in the liquid (actually a gel at low $T$) are not stressed. At the same time, the small value of the density of the gas-phase indicates that the coexisting pressure is rather small. The contact density, at a hard-wall, is a direct measure of the pressure in the system. Hence, close to the coexisting liquid branch, the contact density decreases as $T$ decreases, resulting in the reported desorption. On increasing $\rho$ at constant low $T$, the formation of an extended network of bonds causes the increase of stresses in the system and the pressure increases. This results in a significantly large value of the contact density, driving the adsorption phenomenon.

To pin down the origin of the discrepancies between the DFT results and MC simulations which build up on cooling, we plot in Fig. 5(a) the uniaxial order parameter profile $S_N(z)$ for the system at $\rho_0\sigma^3 = 0.40$. This is defined as $S_N(z) = \int d\Omega h(\hat{\Omega}, z) P_2(\cos(\theta))$ where $h(\hat{\Omega}, z)$ is the orientational distribution function at distance $z$ from the wall, $P_2(\cos(\theta))$ is the second Legendre polynomial and $\theta$ is the
angle between the unit vector \( \mathbf{u} \) normal to the wall and the unit vector \( \mathbf{p} \) normal to the plane containing the patches. \( S_{\theta}(z) \) provides information on the orientation of particles as a function of the distance \( z \) from the wall. As shown in Fig. 5, \( S_{\theta}(z) \) grows significantly close to the wall on cooling. The value of \( S_{\theta}(z) \) signals a crossover from an isotropic fluid to a nematic-like phase near the wall. To highlight the particles orientation close to the wall, we show in Fig. 5(b) the probability distributions \( P(\theta, z) \), confirming that at low \( T \) the particles are oriented near the wall, with the plane containing the patches parallel to the wall. This geometry maximizes the bonding probability by moving the patches away from the neutral wall. The average orientational angle close to the wall decreases continuously with \( T \) and it should approaches \( \theta = 0 \) for perfect order. We note indeed that the finite bonding volume allows for a flexibility in the orientation of the particles, contributing to a small but non-zero average angle even in highly bonded conditions close to the wall. At high \( T \) the particles are randomly oriented (see Figs. 5(a) and 5(b)) and \( P(\theta, z) \sim \sin(\theta) \). Finally, we note that far from the wall (Fig. 5(c)) the particles undergo a continuous change from a state with preferred orientation to an isotropic one.

We stress that both DFT approaches neglect the orientational order that develops in the system at low \( T \), since the free-energy depends on the number density but not on the orientation of the particles.

Finally, we comment on the difference between the results of the two DFT approaches at low \( T \) for \( \rho_b \sigma^3 = 0.40 \). Even though the agreement with the numerical results is poor in both cases, WDA describes qualitatively the layering of the particles near the wall at low \( T \) while this feature is absent in the FMT results. Such layering is related—in part—to the orientation of the particles (which is neglected in both DFTs) in order to minimize the fraction of unbonded sites \( X(z) \) (i.e., \( 1 - \rho_b \)) of particles near the wall. Figure 6 shows \( X(z) \) along \( z \) as predicted by FMT (solid lines) and WDA (dashed lines). Symbols are the results from MC simulations. At low \( T \), \( X(z) \) obtained from the homogeneous Wertheim theory (WDA) exhibits a trend similar to the MC results, while \( X(z) \) calculated using the inhomogeneous Wertheim theory (FMT) increases sharply near the wall. The WDA \( X(z) \) follows the density profile, increasing over the bulk value when \( \rho(z) > \rho_b \), which appears to describe more accurately the \( z \) dependence of \( X \). A somewhat related but more drastic failure of an inhomogeneous version of Wertheim theory was reported by Segura et al.\(^{25} \)

V. CONCLUDING REMARKS

In this work we have studied the properties of a fluid of patchy particles with \( n_p = 3 \) patches. We have located the gas-liquid coexistence line, the critical point and the percolation line, providing a full characterization of the thermodynamics and structure of the fluid phases. We have also shown that, for this patchy model, Wertheim’s theory for homogeneous fluids is accurate. We have then studied the surface properties of this fluid in contact with a neutral hard-wall using MC simulation and two DFT approaches.

We have investigated the behavior of the system at two different densities, one close to the liquid branch of the coexistence curve and one about 70% higher, for several \( T \), covering the structural change of the system from a monomer solution to an almost fully bonded network state. We have shown that at low \( \rho_b \), the wall adsorbs particles at high \( T \) which desorb as \( T \) is lowered. The physical mechanism responsible for the adsorption-desorption crossover is understood in terms of the proximity to the gas-liquid coexistence curve. Indeed, at low \( T \), close to the liquid branch, the liquid coexists with a gas at very low density and the pressure is small. Since the contact density at the hard wall is a measure of this pressure, at small \( \rho_b \) desorption must occur. On increasing \( T \) the pressure increases and for \( T \) above the percolation threshold the particles are adsorbed at the wall. Such crossover is not observed at the higher \( \rho_b \) where the density at the wall is always larger than the bulk density. Indeed, even at low \( T \), the price to pay for the formation of a distorted network leads to an increase of the pressure and hence to a large contact density, even when \( \rho_b \to 1 \).

Not surprisingly, the description using density functional theory is consistent with simulations and with the results reported in Refs. 28 and 29 at high and intermediate \( T \), but it fails at low \( T \). We have traced this failure to the inadequacy of describing the orientational degrees of freedom of the particles. Indeed, close to the wall, the particles are oriented in such a way that the plane containing the bonding sites is almost parallel to the wall. It appears that the orientational degrees of freedom and an appropriate coupling to the density profile need to be taken into account in future work in order to describe the structure of associating fluids (and gels) close to a hard wall at low temperatures.

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APPENDIX A: WEIGHTED DENSITY FUNCTIONAL

The weighted density functional approach is based on the idea that the free energy of the inhomogeneous system, characterized by the single-particle density \( \rho(\vec{r}) \), may be written in terms of the free energy of a homogeneous fluid with an effective density that is evaluated through an appropriate averaging procedure. The approach was proposed by Tarazona \(^{27}\) and was used and modified by several authors.\(^{30, 43, 44}\)

The excess free energy functional of the inhomogeneous fluid is written as

\[
F_{\text{exc}}[\rho] = \int d\vec{r} \rho(\vec{r}) f[\rho; \vec{r}],
\]  
(A1)

where \( f[\rho; \vec{r}] \) is the local excess free energy functional per particle. \( f[\rho; \vec{r}] \) can be written as a function of the density \( \bar{\rho} \), a functional of the single-particle density, which satisfies

\[
f[\rho; \vec{r}] = f(\bar{\rho}(\vec{r})),
\]  
(A2)

where \( f(\bar{\rho}(\vec{r})) \) is the local free-energy density of the homogeneous system and \( \bar{\rho}(\vec{r}) \) is the weighted density defined by

\[
\bar{\rho}(\vec{r}) = \int d\vec{r}' \rho(\vec{r}') \omega(\vec{r} - \vec{r}'; \bar{\rho}(\vec{r})).
\]  
(A3)

In Eq. (A3) \( \bar{\rho}(\vec{r}) \) is the difference between the density and its weighted counterpart and \( \omega \) is the weight function satisfying the constraint

\[
\int d\vec{r}' \omega(\vec{r} - \vec{r}'; \bar{\rho}(\vec{r})) = 1.
\]  
(A4)

The weight function, which encodes the non-local character of the functional, is related via nonlinear differential equations to the direct correlation function of the inhomogeneous fluid. An approximation to the weight function is obtained by requiring that the second functional derivative of the excess free-energy

\[
c^{(2)}(r_1, r_2; \rho) = -\frac{\beta \delta^2 F_{\text{exc}}[\rho]}{\delta \rho(r_1) \delta \rho(r_2)}
\]  
(A5)

gives an accurate description of the correlations of the homogeneous fluid.

Different WDA functionals result from different approximations for the weight function. We use a modified version of the WDA proposed by Tarazona, developed by Kim et al.\(^{41}\) In this approximation

\[
\bar{\rho}(\vec{r}) = \int d\vec{r}' \rho(\vec{r}') \omega(\vec{r} - \vec{r}'; \rho_b),
\]  
(A6)

where \( \rho_b \) is the bulk density. Following Tarazona, \( \omega(\vec{r}, \rho) \) is expanded in powers of \( \rho \) in order to reduce the computational effort:

\[
\omega(r, \rho) = \omega_0(r) + \omega_1(r) \rho + \omega_2(r) \rho^2. \tag{A7}
\]

The same is done for \( \bar{\rho}(\vec{r}) \):

\[
\bar{\rho}(\vec{r}) = \rho_0(\vec{r}) + \rho_1(\vec{r}) \bar{\rho}(\vec{r}) + \rho_2(\vec{r}) \bar{\rho}(\vec{r})^2,
\]  
(A8)

where

\[
\bar{\rho}(\vec{r}) = \rho_0(\vec{r}) + \rho_1(\vec{r}) \rho_b + \rho_2(\vec{r}) \rho_b^2,
\]  
(A9)

with

\[
\rho_i(\vec{r}) = \int d\vec{r}' \rho(\vec{r}') \omega_i(\vec{r} - \vec{r}') \quad i = 0, 1, 2.
\]  
(A10)

The grand potential functional

\[
\beta \Omega[\rho] = \beta F[\rho] + \beta \int d\vec{r} (V_{\text{ext}}(\vec{r}) - \mu) \rho(\vec{r})
\]  
(A11)

is minimized with respect to variations of \( \rho(\vec{r}) \):

\[
\frac{\beta \delta \Omega[\rho]}{\delta \rho(\vec{r})} = \frac{\beta \delta F[\rho]}{\delta \rho(\vec{r})} - \beta (\mu - V_{\text{ext}}(\vec{r})) = 0,
\]  
(A12)

which yields the equation for the equilibrium density profile,

\[
\frac{\beta \delta F[\rho]}{\delta \rho(\vec{r})} = \beta \mu - \beta V_{\text{ext}}(\vec{r}).
\]  
(A13)

The first derivative of the excess free energy functional is the single-particle direct correlation function \( c^{(1)}(\vec{r}) \) and putting together Eqs. (13) and (A1)–(A3) we obtain

\[
\beta \mu - \beta V_{\text{ext}}(\vec{r}) = \ln \rho(\vec{r}) - c^{(1)}[\vec{r}; \rho],
\]  
(A14)

with

\[
c^{(1)}[\vec{r}; \rho] = -\beta f(\bar{\rho}(\vec{r})) - \beta \int d\vec{r}' \rho(\vec{r}') f' \left( \bar{\rho}(\vec{r}') \right) \frac{\delta \bar{\rho}(\vec{r}')}{\delta \rho(\vec{r})},
\]  
(A15)

and

\[
\frac{\delta \bar{\rho}(\vec{r}')}{\delta \rho(\vec{r})} = \omega(\vec{r} - \vec{r}', \bar{\rho}(\vec{r}'))
\]  
(A16)

\[
+ \omega(\vec{r} - \vec{r}', \rho_b) \int d\vec{r}'' \rho(\vec{r}'') \omega(\vec{r}' - \vec{r}'', \bar{\rho}(\vec{r}'')).
\]  
(A17)

The chemical potential \( \mu \) is evaluated from the homogeneous version of Eq. (A15):

\[
\beta \mu = \ln \rho_b - c^{(1)}(\rho_b)
\]  
(A18)

\[
= \ln \rho_b - \beta f(\rho_b) - \beta \rho_b f'(\rho_b).
\]  
(A17)

Finally, combining Eqs. (A14) and (A17), we obtain the density profile:

\[
\rho(\vec{r}) = \rho_b \exp[-\beta V_{\text{ext}}(\vec{r}) + c^{(1)}[\vec{r}; \rho] - c^{(1)}(\rho_b)]. \tag{A18}
\]

The excess free energy is the sum of the free energy of HS given by the Carnahan-Starling approximation\(^{36}\) and the bonding contribution given by Eq. (5).

At a planar hard-wall, the external field is \( V_{\text{ext}} = 0 \) for \( z > 0 \) and infinite otherwise, and the density profile depends only on the distance \( z \) from the wall:

\[
\rho(z) = \rho_b \exp[c^{(1)}(z; \rho_b) - c^{(1)}(\rho_b)] \quad z > \sigma/2,
\]  
\[
= 0 \quad z < \sigma/2. \tag{A19}
\]
The weight functions \( \omega_i(\vec{r}) \) for this system, defined in Ref. 41, are

\[
\omega_0(z) = \begin{cases} 
\frac{3}{4\pi\sigma}, & |z| < \sigma, \\
0, & \text{otherwise}, 
\end{cases}
\]  

\( \text{(A20)} \)

\[
\omega_1(z) = 0.21\sigma^2 - 1.49z^2 - 0.18 \left( \frac{z^2}{\sigma} \right)^2 + 1.36 \frac{|z|^3}{\sigma}, \quad z < \sigma 
\]

\[
= -0.11\sigma^2 + 2.9z^2 + 0.29 \left( \frac{z^2}{\sigma} \right)^2 - 1.81\sigma|z| 
\]

\[
-1.6 \frac{z^3}{\sigma} \quad \sigma \leq z \leq 2\sigma, 
\]

\[
= 0 \quad \text{otherwise}, 
\]

\( \text{(A21)} \)

\[
\omega_2(z) = \begin{cases} 
\frac{10\pi^2}{576}\sigma(\sigma^4 - 12(\sigma z^2) - 5z^4 + 16\sigma|z|^3), & |z| < \sigma, \\
0, & \text{otherwise}, 
\end{cases}
\]  

\( \text{(A22)} \)

APPENDIX B: FUNDAMENTAL MEASURE DENSITY FUNCTIONAL

A second approach, uses a geometry-based density-functional to describe the excluded volume between spheres as well as the inhomogeneous associating free-energy. It was proposed by Yu and Wu \(^{29}\) to describe inhomogeneous mixtures of hard-spheres (HS) with an arbitrary set of interaction sites. In the following we consider a single-component fluid of HS with \( n_p \) identical patches, in which case, the excess free-energy reads:

\[
\beta F_{\text{excl}}(\rho(\vec{r})) = \int d^3r \left( \beta f_{\text{HS}}[n_a(\vec{r})] + \beta f_{\text{bond}}[n_a(\vec{r})] \right), 
\]  

\( \text{(B1)} \)

where \( f_{\text{HS}} \) is the reduced excess free-energy of the fluid of HS, and \( f_{\text{bond}} \) is the free-energy arising from the association of the particles. Both quantities depend on a set of weighted densities \( \{n_a(\vec{r})\} \) (see later).

1. Hard sphere fluid

The excluded volume interactions between HS are described by the Rosenfeld functional:\(^{30}\)

\[
\beta f_{\text{HS}} = -n_0 \ln(1 - n_3) + \frac{n_1n_2 - \hat{n}_{v2} \cdot \hat{n}_{v2}}{1 - n_3} + \frac{(n_3)^2 - 3n_2\hat{n}_{v2} \cdot \hat{n}_{v2}}{24\pi(1 - n_3)^2},
\]

\( \text{(B2)} \)

where we have dropped the spatial dependence of the weighted densities for convenience. \( n_a(\vec{r}) \) are convolutions of the density with the weight functions \( w_a(\vec{r}) \), which are related to geometrical properties of the particles:

\[
n_a(\vec{r}) = w_a(\vec{r}) * \rho(\vec{r}),
\]

\( \text{(B3)} \)

where * denotes the three-dimensional convolution \( h(\vec{r}) * g(\vec{r}) = \int d^3x h(x)g(\vec{x} - \vec{r}) \). For HS the weight functions are:

\[
w_3(\vec{r}) = \Theta(R_S - |\vec{r}|),
\]

\( \text{(B4)} \)

\[
w_2(\vec{r}) = \delta(R_S - |\vec{r}|),
\]

\( \text{(B5)} \)

\[
w_1(\vec{r}) = w_2(\vec{r})/(4\pi R_S),
\]

\( \text{(B6)} \)

\[
w_0(\vec{r}) = w_2(\vec{r})/(4\pi R_S^2),
\]

\( \text{(B7)} \)

\[
\tilde{w}_{v2}(\vec{r}) = w_2(\vec{r})/|\vec{r}|,
\]

\( \text{(B8)} \)

\[\delta(\cdot)\] is the Dirac-delta distribution and \( \Theta(\cdot) \) is the Heaviside step function. \( R_S = \sigma/2 \) is the sphere radius. In planar geometry the one particle distribution function depends only on the normal distance from the wall, \( \xi \). The weight functions are obtained integrating over the lateral coordinates,

\[
w_a(\xi) = \int dx \int dy w_a(\vec{r}).
\]

\( \text{(B10)} \)

The resulting weight functions are

\[
w_3(\xi) = \pi \left( R_S^2 - \xi^2 \right) \Theta(R_S - |\xi|),
\]

\( \text{(B11)} \)

\[
w_2(\xi) = 2\pi R_S \Theta(R_S - |\xi|),
\]

\( \text{(B12)} \)

\[
w_1(\xi) = w_2(\xi)/(4\pi R_S),
\]

\( \text{(B13)} \)

\[
w_0(\xi) = w_2(4\pi R_S^2),
\]

\( \text{(B14)} \)

\[
\tilde{w}_{v2} = 2\pi \xi \Theta(R_S - |\xi|)/\xi,
\]

\( \text{(B15)} \)

\[
\tilde{w}_{v1} = w_{v2}/(4\pi R_S),
\]

\( \text{(B16)} \)

with \( \xi \) the unit vector normal to the wall.

2. Wertheim’s inhomogeneous free-energy

The bulk free energy of a fluid of particles with \( n_p \) identical sites (Eq. (5)) can be rewritten in terms of the bulk fraction of unbonded sites \( X_b \equiv 1 - p_b \) as\(^{17-20}\)

\[
\beta f_{\text{bond}} = n_p \rho_b \left( \ln X_b - X_b \right) + \frac{1}{2} \left( X_b - \frac{1}{2} \right),
\]

\( \text{(B17)} \)

with \( \rho_b \) the bulk density, and \( X_b \) the bulk fraction of unbonded sites, related to the thermodynamic variables through the law of mass action:

\[
X_b = (1 + n_p \rho_b X_b \Delta)^{-1}.
\]

\( \text{(B18)} \)

Yu and Wu\(^{29}\) generalized the bulk free energy, Eq. (B17), to inhomogeneous systems by including a new factor \( \xi = 1 - \hat{n}_{v2} \cdot \hat{n}_{v2}/n_3^2 \) that incorporates the vectorial weight.
densities into the associating part of the free energy. The bonding free energy for inhomogeneous fluids reads

\[ \beta f_{\text{bond}}[\rho_a(\vec{r})] = n_p n_0(\vec{r}) \xi(\vec{r}) \left( \ln X(\vec{r}) - \frac{X(\vec{r})}{2} + \frac{1}{2} \right). \]  

(B19)

where \( X(\vec{r}) \) is the fraction of unbonded sites at position \( \vec{r} \) given by the modified law of mass action:

\[ X(\vec{r}) = \left( 1 + n_p n_0(\vec{r}) \xi(\vec{r}) X(\vec{r}) \Delta(\vec{r}) \right)^{-1}. \]  

(B20)

The interaction between two sites determines \( \Delta \). For the Kernel-Frenkel potential (where orientational and translational degrees of freedom are decoupled) \( \Delta \) is given in terms of \( g_{\text{HIS}}(\vec{r}) \), the pair correlation function of the reference HS fluid, and \( f_M \), the Mayer function:

\[ \Delta(\vec{r}) = \int d^3r g_{\text{HIS}}(\vec{r}) f_M. \]  

(B21)

where \( f_M = \exp(\beta \varepsilon) - 1 \), and the integral is over the bonding volume \( v_b \). Following Yu and Wu29 we use a modified contact value for the pair correlation function:

\[ g_{\text{HIS}}(\vec{r}) = \frac{1}{1 - n_3} + \frac{\sin 2 \zeta}{4(1 - n_3)} + \frac{\sigma^3 (n_2)^2}{72(1 - n_3)^2}. \]  

(B22)

Assuming that the pair correlation function is constant over the bonding volume, we approximate Eq. (B21) by

\[ \Delta(\vec{r}) = v_b g_{\text{HIS}}(\vec{r}) f_M. \]  

(B23)

Finally, we minimize the grand potential,

\[ \Omega[\rho] = F[\rho] - \mu \int d^3r \rho(\vec{r}), \]  

(B24)

to obtain the equilibrium density profiles. \( \mu \) is the chemical potential and the integrals are computed using the trapezoidal rule and a step size \( \Delta z = 0.01 \sigma \). We use a standard conjugated-gradients method to minimize \( \Omega \).