Structural properties of additive binary hard-sphere mixtures

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An approach to obtain the structural properties of additive binary hard-sphere mixtures is presented. Such an approach, which is a nontrivial generalization of the one recently used for monocomponent hard-sphere fluids [S. Pieprzyk, A. C. Brańka, and D. M. Heyes, Phys. Rev. E 95, 062104 (2017)], combines accurate molecular-dynamics simulation data, the pole structure representation of the total correlation functions, and the Ornstein–Zernike equation. A comparison of the direct correlation functions obtained with the present scheme with those derived from theoretical results stemming from the Percus–Yevick (PY) closure and the so-called rational-function approximation (RFA) is performed. The density dependence of the leading poles of the Fourier transforms of the total correlation functions and the decay of the pair correlation functions of the mixtures are also addressed and compared to the predictions of the two theoretical approximations. A very good overall agreement between the results of the present scheme and those of the RFA is found, thus suggesting that the latter (which is an improvement over the PY approximation) can safely be used to predict reasonably well the long-range behavior, including the structural crossover, of the correlation functions of additive binary hard-sphere mixtures.

I. INTRODUCTION

Structural properties of the liquid phase are routinely expressed in terms of pair or structural correlation functions (SCFs) [1–3]. An example is the radial distribution function (RDF) $g(r)$, where $r$ means the interparticle separation. This important structural function can be obtained inter alia from the static structure factor determined through X-ray and neutron scattering experiments [4, 5] and from computer simulations of model liquids.

Because the SCFs are present in many theoretical and experimental treatments, their understanding and mutual relationships are of great importance. In the case of simple liquids, the SCFs have been a subject of intensive studies for decades, and considerable knowledge has been amassed regarding their structure. In the case of multicomponent liquid mixtures, the studies of structural properties, due to their complexity, are much less developed.

Important practical methods for obtaining SCFs are computer simulations and the liquid-state theories based on the Ornstein–Zernike (OZ) relation [1],

\[ h(r_{12}) = c(r_{12}) + \rho \int \text{d}r_3 h(r_{13})c(r_{23}), \]  

where $\rho$ is the number density, $c(r)$ is the direct correlation function (DCF), and $h(r) = g(r) - 1$ is the total correlation function. The subscripts, 1, 2, and 3 denote the positions of three particles, and $r_{ij} = |\mathbf{r}_i - \mathbf{r}_j|$ is the separation between particles $i$ and $j$.

A key role for understanding and describing a liquid structure has been played by various hard-particle models. Among them, the binary hard-sphere (BHS) mixture is of particular relevance as it can be considered as the simplest model for real liquid mixtures [6, 7]. In this paper, a framework allowing us to obtain an accurate representation of the SCFs of additive BHS mixtures is proposed. The method includes molecular dynamics (MD) simulation data, residue theorem analysis, and the OZ relation. The main aim is to obtain a more comprehensive representation of the SCFs for additive BHS, bring to light some new features, and compare results with two analytical predictions. In our approach, the tail parts of the SCFs are taken into account without using any approximate closures. We focus here on the DCFs, which are in general quantities very difficult...
to access from the RDFs due to inherent errors in truncated numerical Fourier transforms [8].

Analytical formulas from the Percus–Yevick (PY) approximation for the DCFs of monocomponent hard-spheres and additive BHS mixtures have been available for many decades, and this can be considered as one of the most important results in the history of statistical-mechanical liquid-state theory. Just one year after Wertheim [9] and Thiele [10] found the exact solution of the (three-dimensional) OZ equation with the PY closure for a monocomponent HS fluid, Lebowitz extended the solution to additive BHS mixtures [11]. In the past two decades, some approximate analytical formulas for the SCFs have been proposed, mainly to reduce limitations of the PY theory. Yuste et al. [12–17] derived analytic approximations, for both the monocomponent hard-sphere (HS) case and additive HS mixtures, based on a generalization of the PY result. Their method, usually referred to as the Rational-Function Approximation (RFA), circumvents the thermodynamic consistency problem of the PY solution. In fact, the RFA can be seen as an augmented PY solution that includes an extra parameter, α_{RFA}, such that the PY form is recovered by the choice α_{RFA} = 0; on the other hand, prescribing the isothermal compressibility and the contact values of the RDFs yields a quadratic (monocomponent HS fluid) or a quartic (additive BHS mixture) equation for α_{RFA}. The RFA method has also been extended to nonadditive HS mixtures [18–20]. Here, we implement the RFA approximation with the Boublík–Grundke-Henderson–Lee–Levesque (BGHLL) contact values [21–23] and the isothermal compressibility corresponding to the Boublík–Mansoori–Carnahan–Starling–Leland (BMCSL) equation of state [21,24]. In this work, those two theoretical approximations (RFA and PY) will be compared with the DCFs obtained from simulation results of the RDFs via our proposed scheme. This in turn will be used to determine the two leading poles of the Fourier transforms of the total correlation functions, and to analyze the structural crossover in additive BHS [25,26]. All of this will allow for an assessment of the performance of the RFA as a valuable tool for exploring different density and/or composition regions of BHS mixtures.

The work is organized as follows. In Sec. II the general theory of the RDF and DCF is covered, focusing especially on the large-wavenumber limit in Fourier space. The monocomponent case is discussed in Sec. III A. In Sec. III B the calculation details for additive BHS mixtures are presented and discussed. The main conclusions are summarized in Sec. IV.

II. STRUCTURAL CORRELATION FUNCTIONS

We consider an additive BHS mixture composed of small (s) and big (b) particles, characterized by the size ratio \( \sigma_s/\sigma_b \leq 1 \), where \( \sigma_s \) and \( \sigma_b \) are HS diameters.

Thus, there are three different separations between particles at contact: the small-small particle separation, \( \sigma_{ss} = \sigma_s \), the small-big particle separation, \( \sigma_{sb} = \frac{1}{2}(\sigma_s + \sigma_b) \), and the big-big particle separation, \( \sigma_{bb} = \sigma_b \). The additive BHS system is defined with the pairwise interaction

\[
\psi_{ij}(r) = \left\{ \begin{array}{ll}
\infty, & 0 < r < \sigma_{ij}, \\
0, & r > \sigma_{ij},
\end{array} \right.
\]

(2)

where \( i, j = s, b \).

The partial packing fractions are defined as \( \eta_i = \frac{\pi}{6} n \rho_i \sigma_i^3 \), where \( \rho_i = N_i / V \) are number densities, \( N_i \) and \( V \) being the number of particles of species \( i \) and the volume of the system, respectively. The total number of particles, number density, and packing fraction are \( N = N_s + N_b \), \( \rho = N/V = \rho_s + \rho_b \), and \( \eta = \eta_s + \eta_b \), respectively.

The focus in this work is on an accurate determination of the structural properties of the additive BHS fluid mixture by exploiting the pole method [1,2,25,27,28]. From the method it follows inter alia that the asymptotic decay of \( h_{ij}(r) \) is determined by the poles of the Fourier transform \( \tilde{h}_{ij}(k) \) with the imaginary part closest to the real axis [25]. Also, the method was recently shown to be a useful means to obtain the entire DCF and the first two poles in the case of the monocomponent HS fluid [28]. Below, we extend this scheme to the additive BHS mixture.

A. Total correlation functions

The representation of \( h_{ij}(r) \) of additive BHS mixtures in terms of the pole structure can be written as [1,2,25,27,28]

\[
h_{ij}(r) = -\Theta(\sigma_{ij} - r) + \Theta(r - \sigma_{ij})
\times \sum_{n=1}^{\infty} \frac{A_{ij}^{(n)}}{r} e^{-\alpha_n r} \sin \left( \omega_n r + \delta_{ij}^{(n)} \right),
\]

(3)

where \( \Theta(x) \) is the Heaviside step function. The damping coefficients \( \alpha_n \) and the oscillation frequencies \( \omega_n \) are common to all the pairs, while the amplitudes \( A_{ij}^{(n)} \) and the phase shifts \( \delta_{ij}^{(n)} \) are specific for each \( h_{ij}(r) \) [27]. It is noteworthy that the Fourier transform,

\[
\tilde{h}_{ij}(k) = 4\pi \int_0^\infty dr r^2 h_{ij}(r) \sin(kr) / kr,
\]

(4)

of the above representation for \( h_{ij}(r) \) can be expressed by the following analytic expression,

\[
\tilde{h}_{ij}(k) = 4\pi \left[ \frac{\sigma_{ij} \cos(\sigma_{ij} k)}{k^2} - \frac{\sin(\sigma_{ij} k)}{k^3} \right]
+ \frac{2\pi}{k} \sum_{n=1}^{\infty} \left[ P_{ij}^{(n)}(\sigma_{ij}, k) - P_{ij}^{(n)}(\sigma_{ij}, -k) \right],
\]

(5)
where

\[
P_{ij}^{(n)}(\sigma_{ij}, k) = \frac{A_{ij}^{(n)} e^{-\alpha n \sigma_{ij}}}{\alpha_n^2 + (\omega_n - k)^2} \left\{ \alpha_n \cos[\delta_i^{(n)} + (\omega_n - k)\sigma_{ij}] - (\omega_n - k) \sin[\delta_i^{(n)} + (\omega_n - k)\sigma_{ij}] \right\}.
\]  

(6)

1. Large-\( k \) limit

The large-\( k \) limit or ‘tail’ of \( \hat{h}_{ij}(k) \) can be obtained from Eq. (4) by expanding in negative powers of \( k \) the functions multiplying the trigonometric functions \( \sin(\sigma_{ij}k) \) and \( \cos(\sigma_{ij}k) \), namely

\[
\hat{h}_{ij}^{\text{tail}}(k) = \cos(\sigma_{ij}k) \sum_{n=1}^{\infty} C_{ij}^{(n)} \frac{k^{2n}}{k^{2n}} + \sin(\sigma_{ij}k) \sum_{n=1}^{\infty} D_{ij}^{(n)} \frac{k^{2n}}{k^{2n+1}}.
\]  

(7)

The first few coefficients \( C_{ij}^{(n)} \) and \( D_{ij}^{(n)} \) are given in Appendix A where it is also shown that those coefficients can be expressed in terms of derivatives of the RDF at contact (i.e., at \( r = \sigma_{ij}^+ \)) as

\[
\begin{align*}
C_{ij}^{(1)} &= 4\pi \sigma_{ij} g_{ij}(\sigma_{ij}^+), \\
C_{ij}^{(2)} &= -4\pi \left[ 2g_{ij}^{''}(\sigma_{ij}^+) + \sigma_{ij} g_{ij}^{''}(\sigma_{ij}^+) \right], \\
D_{ij}^{(1)} &= 4\pi \left[ g_{ij}^{''''}(\sigma_{ij}^+) + \sigma_{ij} g_{ij}^{''''}(\sigma_{ij}^+) \right], \\
D_{ij}^{(2)} &= 4\pi \left[ 3g_{ij}^{''''}(\sigma_{ij}^+) + \sigma_{ij} g_{ij}^{''''}(\sigma_{ij}^+) \right].
\end{align*}
\]

(8a-d)

where single, double, and triple primes represent the first, second, and third derivatives of the RDF, respectively.

2. Small-\( k \) limit

It can be shown from Eq. (5) that \( \hat{h}_{ij}(k) \) is an even function regular at \( k = 0 \), so that its Taylor expansion in powers of \( k \) is

\[
\hat{h}_{ij}(k) = \hat{h}_{ij}^{(0)} + \hat{h}_{ij}^{(2)} k^2 + \hat{h}_{ij}^{(4)} k^4 + \ldots,
\]  

(9)

where the zeroth-order term is

\[
\hat{h}_{ij}^{(0)} = -\frac{4\pi}{3} \sigma_{ij}^3 + 4\pi \sum_{n=1}^{\infty} \frac{A_{ij}^{(n)} e^{-\alpha n \sigma_{ij}}}{(\alpha_n^2 + \omega_n^2)^2} \times \left[ \omega_n (2\alpha_n + \alpha_n^2 \sigma_{ij} + \omega_n^2 \sigma_{ij}) \cos(\delta_i^{(n)} + \omega_n \sigma_{ij}) \\
+ (\alpha_n^2 + \alpha_n^3 \sigma_{ij} - \omega_n^2 + \alpha_n \omega_n^2 \sigma_{ij}) \sin(\delta_i^{(n)} + \omega_n \sigma_{ij}) \right].
\]  

(10)

The coefficients of higher order are similar to the zeroth-order one but have more complex expressions. The coefficients \( \hat{h}_{ij}^{(n)} \) for all \( ij \) are involved in the isothermal compressibility \( \kappa_T \).

B. Direct correlation functions

The OZ relation, Eq. (1), for binary mixtures in Fourier space, has the form

\[
\hat{h}_{ij}(k) = \tilde{c}_{ij}(k) + \sum_{\ell} \rho_{\ell} \tilde{c}_{\ell\ell}(k) \hat{h}_{ij}(k),
\]  

(11)

and in matrix notation,

\[
\hat{h}(k) = \tilde{c}(k) \cdot \left[ 1 + \hat{h}(k) \right],
\]  

(12)

where \( \hat{h}(k) \equiv \sqrt{\rho_s \rho_b} \tilde{h}_{ij}(k), \tilde{c}_{ij}(k) \equiv \sqrt{\rho_s \rho_b} \tilde{c}_{ij}(k), \) and \( I \) is the identity matrix. Thus,

\[
\tilde{c}(k) = \tilde{h}(k) \cdot [1 + \tilde{h}(k)]^{-1} = 1 - [1 + \tilde{h}(k)]^{-1}.
\]  

(13)

More explicitly, one has

\[
\begin{align*}
\tilde{c}_{ss}(k) &= \frac{\tilde{h}_{ss}(k) + \rho_s \left[ \tilde{h}_{ss}(k) \hat{h}_{bb}(k) - \tilde{h}_{sb}^2(k) \right]}{D(k)}, \\
\tilde{c}_{sb}(k) &= \frac{\tilde{h}_{sb}(k)}{D(k)},
\end{align*}
\]

(14a-b)

where

\[
D(k) = 1 + \rho_s \tilde{h}_{ss}(k) + \rho_b \tilde{h}_{bb}(k) \]

\[
+ \rho_s \rho_b \left[ \tilde{h}_{ss}(k) \hat{h}_{bb}(k) - \tilde{h}_{sb}^2(k) \right].
\]  

(15)

For the sake of conciseness, we omit the expression for \( \tilde{c}_{sb}(k) \), which can be obtained from Eq. (14a) by the simple exchange \( s \leftrightarrow b \). Henceforth, we will do the same for any quantity of the form \( \tilde{h}_{ss}(k) \), which can then be obtained from \( \tilde{h}_{ss}(k) \) by the same exchange of indices.

1. Large-\( k \) limit

Taking into account that the absolute value of \( D^{\text{tail}} \equiv D - 1 \) is smaller than 1, we can expand \( D^{-1} = 1 - D^{\text{tail}} + D^{\text{tail}^2} + \ldots \) for large wave number \( k \) and use the tail forms as in Eq. (7). As a consequence, the first few terms in the large-\( k \) limit ("tail") of \( \tilde{c}_{ij}(k) \) are
\[ \tilde{c}_{ss}^{\text{tail}}(k) = \frac{C_{ss}^{(1)}}{k^2} \cos(\sigma_s k) + \frac{D_{ss}^{(1)}}{k^3} \sin(\sigma_s k) + \frac{C_{ss}^{(2)}}{k^2} \cos(\sigma_s k) - \rho_s \frac{C_{ss}^{(1)}}{k^4} \cos(\sigma_s k) - \rho_b \frac{C_{ss}^{(1)}}{k^4} \cos(\sigma_s k) + \cdots, \] 
\[ \tilde{c}_{sb}^{\text{tail}}(k) = \frac{C_{sb}^{(1)}}{k^2} \cos(\sigma_b k) + \frac{D_{sb}^{(1)}}{k^3} \sin(\sigma_b k) + \frac{C_{sb}^{(2)}}{k^2} \cos(\sigma_b k) - \rho_s C_{ss}^{(1)} \cos(\sigma_s k) + \rho_b C_{bb}^{(1)} \cos(\sigma_b k) \] 
\[ \rho_s D_{ss}^{(1)} \sin(\sigma_s k) + \rho_b D_{bb}^{(1)} \sin(\sigma_b k) + \cdots. \] 

\[ \Delta c_{ij}(\sigma_{ij}) = \frac{C_{ij}^{(1)}}{4\pi \sigma_{ij}} = g_{ij}(\sigma_{ij}^+), \] 
\[ \Delta' c_{ij}(\sigma_{ij}) = -\frac{1}{4\pi} \left( \frac{C_{ij}^{(1)}}{\sigma_{ij}^2} + \frac{D_{ij}^{(1)}}{\sigma_{ij}} \right) = g_{ij}'(\sigma_{ij}^+), \] 
\[ \Delta'' c_{ij}(\sigma_{ij}) = \frac{1}{4\pi} \left( \frac{2C_{ij}^{(1)}}{\sigma_{ij}^3} + \frac{2D_{ij}^{(1)}}{\sigma_{ij}^2} - \frac{C_{ij}^{(2)}}{\sigma_{ij}} \right) = g_{ij}''(\sigma_{ij}^+), \] 
\[ \Delta''' c_{ij}(\sigma_{ij}) = \frac{1}{4\pi} \left( -\frac{6C_{ij}^{(1)}}{\sigma_{ij}^4} + \frac{6D_{ij}^{(1)}}{\sigma_{ij}^3} + \frac{3C_{ij}^{(2)}}{\sigma_{ij}^2} + \frac{D_{ij}^{(2)}}{\sigma_{ij}} \right) = g_{ij}'''(\sigma_{ij}^+), \] 

where we have introduced the short-hand notation \( \Delta X(a) = \lim_{r \to a^-} X(r) - \lim_{r \to a^+} X(r) \) and in the last steps use has been made of Eqs. (8). Equations (20) are consistent with the continuity of the indirect correlation functions \( \hat{h}_{ij}(r) = c_{ij}(r) \) and their first three derivatives at \( r = \sigma_{ij} \).

C. Determination of the poles of \( \hat{h}_{ij}(k) \)

From the OZ relation (12) it is straightforward to obtain \( \hat{h}(k) = \frac{1}{1 - \tilde{c}(k)} \) and \( \hat{h}_{ij}(k) \) given by the zeros of the determinant of \( \tilde{c}(k) \), i.e., \( D(k) \equiv [1 - \rho_s \tilde{c}_{ss}(k)][1 - \rho_b \tilde{c}_{bb}(k)] - \rho_s \rho_b \tilde{c}_{sb}(k) = 0 \). By equating real and imaginary parts, the formulas
\[ 1 = \rho_s I_s^{(0)}(\alpha, \omega) + \rho_b I_b^{(0)}(\alpha, \omega) + \rho_s \rho_b I_{sb}^{(0)}(\alpha, \omega), \] 
\[ 1 = \rho_s I_s^{(1)}(\alpha, \omega) + \rho_b I_b^{(1)}(\alpha, \omega) + \rho_s \rho_b I_{sb}^{(1)}(\alpha, \omega), \] 
are obtained. The required integrals \( I_j^{(n)} \) and \( I_{sb}^{(n)} \) are presented in Appendix C. If the DCFs, \( c_{ij}(r) \), are known and decay sufficiently fast, the above equations provide a practical route to obtain the poles of \( \hat{h}_{ij}(k) \).
D. Determination of amplitudes and phases

To determine $h_{ij}(r)$ in Eq. (8), the amplitudes, $A_{ij}^{(n)}$, and phase, $\delta_{ij}^{(n)}$, are required as well. The appropriate prescription for their evaluation was constructed by Evans et al. [27] by application of the residue theorem. In the case of a binary mixture, the contribution to $r h_{ij}(r)$ associated with the pair of poles $k_{\text{pole}} = \omega + i\alpha$ and $-k_{\text{pole}}^* = -\omega + i\alpha$ is $A_{ij} e^{k_{\text{pole}} r} + A_{ij}^* e^{-k_{\text{pole}}^* r}$, the ‘complex amplitudes’ $\tilde{A}_{ij}$ being given by the following expressions:

$$\tilde{A}_{ss} = \frac{k_{\text{pole}} (1 - \rho_b \bar{c}_{bb})}{2\pi \rho_a D'}, \quad \tilde{A}_{sb} = \frac{k_{\text{pole}} \bar{c}_{sb}}{2\pi D'},$$

where

$$\bar{c}_{ss} = -\rho_s (1 - \rho_b \bar{c}_{bb}) \bar{c}_{ss} - \rho_b (1 - \rho_s \bar{c}_{ss}) \bar{c}_{bb} - 2\rho_s \rho_b \bar{c}_{sb} \bar{c}_{sb},$$

$$\tilde{c}_{ij} = \frac{4\pi}{k_{\text{pole}}} \int_0^\infty dr \, r c_{ij}(r) \left[ \cosh(\alpha r) \sin(\omega r) \right. + i \sinh(\alpha r) \cos(\omega r)],$$

$$\tilde{c}_{ij}' = -\tilde{c}_{ij} - \frac{4\pi}{k_{\text{pole}}} \int_0^\infty dr \, r^2 c_{ij}(r) \left[ \cosh(\alpha r) \cos(\omega r) \right. - i \sinh(\alpha r) \sin(\omega r)].$$

Note that $\tilde{c}_{ij} = \tilde{c}_{ij}(k_{\text{pole}}^*)$, $\tilde{c}_{ij}' = \tilde{c}_{ij}'(k_{\text{pole}}^*)$, and $\bar{D}' = \bar{D'}(k_{\text{pole}}^*)$. Provided the DCFs are known and the poles are determined from Eqs. (21), then the amplitudes and phases can be evaluated from the real and imaginary parts of the complex amplitudes $A_{ij} = |A_{ij}| e^{i(\delta_{ij} - \frac{\pi}{2})}$ [so, one obtains e.g., $A_{ij} = 2|A_{ij}|$]. In this way, at least in principle, a contribution of each $n$-pole defined by $\{A_{ij}^{(n)}, \alpha_n, \omega_n, \delta_{ij}^{(n)}\}$ to the $h_{ij}(r)$ may be determined. In practice, the scheme is strongly limited by the accuracy of the DCFs, and even obtaining the contribution of the leading poles becomes a hard task. In this work, it is done for the two leading poles of the additive BHS mixture.

III. RESULTS

In this Section we provide the analysis of the RDF and DCF, with special emphasis on the large-wave-number
limit in Fourier space. First we deal with the monocomponent case and subsequently we consider additive BHS mixtures.

A. Monocomponent HS fluid

The monocomponent case was considered in Ref. 28 and the associated SCFs were determined with the so-called WM-scheme (see Sec. III B), which combines the OZ equation, the residue theorem analysis, and simulation data. Here, for the sake of completeness, we revisit this monocomponent case. Figure 1 shows the DCF at several densities as obtained from the WM-scheme and as given by the PY and RFA theoretical approaches. While the PY approximation predicts well the low-density behavior of the HS fluid, its agreement with the WM-determined DCF deteriorates on increasing density. This known feature of the PY solution is considerably corrected by the RFA. As seen in Fig. 1(a), in the core region \(0 < \rho \sigma < 1\) (where \(\sigma\) is the diameter of the spheres) the DCF points calculated with the WM-scheme follow practically exactly the prediction of the RFA. It is noteworthy that this excellent agreement takes place for all fluid densities, including those close to the freezing region.

The performance of the RFA at larger \(r\)-separations is also very good, as observed from Fig. 1(b). On the other hand, small deviations between the monotonic (RFA) and the oscillatory (WM-scheme) results occur, although they are visible only for high densities. In Fig. 2 the density dependences of the first (\(\Pi_1\)) and second (\(\Pi_2\)) poles determined with the WM-scheme are compared with those obtained from the PY and RFA approaches. The RFA reproduces almost perfectly the density dependence of the first pole, and in this respect it improves upon the outcome of the PY approximation. The results for the second pole [see Fig. 2(b)] demonstrate the non-negligible role of the DCF part for \(r/\sigma > 1\) at sufficiently high densities. More precisely, the departure of the WM-scheme values and the theoretical ones occurs for \(\rho \sigma^3 > 0.7\). It is worth mentioning that Statt et al.’s analysis for a one-component colloidal suspension yielded a plot similar to Fig. 2(a) with simulation and experiment deviating from PY theory at very high packings 26.

B. Additive binary hard-sphere mixtures

In order to obtain accurate DCFs of additive BHS mixtures, the following analytic representation of \(h_{ij}(r)\) in the form of two functional parts is considered,

\[
h^{W, M}_{ij}(r) = \begin{cases} 
-1, & 0 < r < \sigma_{ij}, \\
\frac{\rho_i^{1(2)}}{r} & \sigma_{ij} < r < r_{ij}^{\text{min}}, \\
\frac{\rho_i^{1(M)}}{r} & r > r_{ij}^{\text{min}},
\end{cases}
\]

where

\[
h^{W}_{ij}(r) = \sum_{n=1}^{W} b^{W}_{ij}(n)r^{n-1},
\]

\[
h^{M}_{ij}(r) = \sum_{n=1}^{M} A^{(n)}_{ij} e^{-\alpha r} \sin(\omega r + \delta^{(n)}_{ij}).
\]

As explained above, the parameters \(\{b^{(1)}_{ij}, b^{(2)}_{ij}, \ldots, b^{(W)}_{ij}\}\) and \(\{A^{(1)}_{ij}, \alpha, \omega, \delta^{(1)}_{ij}, \ldots, A^{(M)}_{ij}, \alpha_M, \omega_M, \delta^{(M)}_{ij}\}\) are obtained by a fitting procedure.

The form of \(h^{W}_{ij}(r)\) in Eq. (25a) is fairly arbitrary, but we seek a rather simple function which provides sufficient flexibility at the next stages of the calculation. In this respect, the polynomial form, the Fourier transform of which can be obtained analytically, is a convenient and appropriate form. A suitable choice for \(r_{ij}^{\text{min}}\) is the position of the first minimum of \(h_{ij}(r)\). Also, our tests suggest that, for most studied densities, the optimal choices for \(W\) and \(M\) are in the range of, approximately, 8–15 and in this way the final results are fairly insensitive to the particular values of those parameters. In the results presented below we have usually taken \(W = 15\) and \(M = 10\).

Furthermore, the function \(h^{W, M}_{ij}\) and its first derivative are constrained to be continuous at \(r = r_{ij}^{\text{min}}\), so that the following continuity conditions are imposed in the scheme,

\[
h^{W}_{ij}(r_{ij}^{\text{min}}) = h^{M}_{ij}(r_{ij}^{\text{min}}),
\]

\[
\frac{\partial h^{W}_{ij}(r)}{\partial r} \bigg|_{r_{ij}^{\text{min}}} = \frac{\partial h^{M}_{ij}(r)}{\partial r} \bigg|_{r_{ij}^{\text{min}}} = 0.
\]

Moreover, the contact values proposed, independently, by Boublik 21, Grundke and Henderson 22, and Lee and Levesque 23, are enforced, so that

\[
h^{W}_{ij}(\sigma_{ij}^{+}) = \frac{\rho_i^{1(2)}}{1 - \eta} + \frac{3}{2} \frac{\eta}{(1 - \eta)^2} \frac{\sigma_{ij}}{\mu_2} + \frac{\eta^2}{2} \frac{1}{(1 - \eta)^2} \left[ \frac{\sigma_{ij}}{\mu_3} \right] + \frac{\mu_n}{\rho} \sum \rho_i \sigma_i^{n},
\]

(27)

The Fourier transform of the above representation of \(h^{W, M}_{ij}(r)\) in Eq. (24) is given by the analytic expression

\[
h^{W}_{ij}(k) = 4\pi \left[ \frac{\sigma_{ij} \cos(k \sigma_{ij})}{k^2} - \frac{\sin(k \sigma_{ij})}{k^3} \right] h^{W}_{ij}(k) + h^{M}_{ij}(k),
\]

(28)
where

\[ \tilde{h}_{ij}^W(k) = \sum_{n=0}^{[\frac{W+4}{2}]} 4\pi \sum_{k=2n+1}^{W+2} \frac{1}{k^2} \left[ s_{ij}^{(n)}(\sigma_{ij}) \cos(k\sigma_{ij}) - s_{ij}^{(n)}(r_{ij}^{min}) \right] \]

\[ \times \cos(kr_{ij}^{min}) + \sum_{n=1}^{[\frac{W+2}{2}]} \frac{4\pi}{k^{2n+1}} \left[ t_{ij}^{(n)}(\sigma_{ij}) \sin(k\sigma_{ij}) \right. \]

\[ \left. - t_{ij}^{(n)}(r_{ij}^{min}) \sin(kr_{ij}^{min}) \right] \]

\[ (29a) \]

\[ \tilde{h}_{ij}^M(k) = \frac{2\pi}{h} \sum_{n=1}^{M} \left[ P_{ij}^{(n)}(r_{ij}^{min}, k) - P_{ij}^{(n)}(r_{ij}^{min}, -k) \right]. \]

(29b)

In the upper summation limits of Eq. (29a), \([\cdots]\) denotes the integer part, and the coefficients \( s_{ij}^{(n)}(a) \) and \( t_{ij}^{(n)}(a) \) (with \( a = \sigma_{ij} \) and \( a = r_{ij}^{min} \)) are linear combinations of the coefficients \( b_{ij}^{(n)}(a) \) whose explicit expressions will be omitted here for the sake of simplicity. We recall that in Eq. (29b) \( P_{ij}^{(n)} \) is defined by Eq. (3).

Equations (29) along with the scheme discussed in Sec. 11 can be used as a practical means for determining the SCFs (in particular, the DCFs) of additive BHS mixtures. More explicitly, Eq. (29a) is inserted into Eqs. (13) to obtain an analytic form for \( c_{ij}^{W,M}(k) \) and hence \( c_{ij}^{W,M}(r) \) by the numerical integration described by Eq. (19a). Also, by expanding \( h_{ij}^{W,M}(k) \) for large wavenumbers, we obtain a tail form with the structure of Eq. (7) and hence a tail DCF, \( c_{ij}^{tail}(r) \), with the structure of Eqs. (10). An analytic integration then yields \( c_{ij}^{tail}(r) \) from Eq. (19b). In what follows, as done in the monocomponent HS case, we will refer to this as the WM-scheme.

The set of parameters in Eqs. (29) was determined by a nonlinear fitting procedure based on the minimization of \( |h_{ij}^{W,M}(r) - h_{ij}^{MD}(r)| < 10^{-3} \) for each \( r/\sigma_{ij} \in (1, r^*_c) \), where \( h_{ij}^{MD}(r) \) was obtained from our MD simulations. A choice \( r^*_c = 5 \) was seen to be sufficient for our calculations.

The computation of \( h_{ij}^{MD}(r) \) was performed with the DYNAMO program [29] for the total packing fraction set equal to \( \eta = 0.5 \) and partial packing fractions \( \eta_s = 0.05, 0.1, 0.15, 0.20, 0.22, 0.24, 0.26, 0.28, 0.30, 0.35, 0.40 \); the size ratio was fixed at \( \sigma_s/\sigma_b = 0.648 \). These specific conditions were chosen to compare the results for this BHS system with those that were obtained before by simulation and experiment [20]. The data for \( h_{ij}^{MD}(r)/\sigma_{ij} < r^*_c \) must be obtained from long simulations with a large number of particles \( (N \sim 10^4) \). Only in this way can the finite-size effects and the statistical errors in the simulations be reduced sufficiently. In order to test the \( N \)-dependence and assess those finite size effects, some calculations were carried out for systems of \( N = 2916, 4000, 6912, 8788, \) and \( 16384 \) particles. It was checked that the simulations for the system of 8788 particles were sufficient to obtain reasonably accurate data.

The histogram grid size of \( g_{ij}(r) \) was set to \( \delta r/\sigma_{ij} = 0.01 \), which was found to be an optimal choice. The MD simulations were carried out typically for a total number of \( 2 \times 10^9 \) collisions, and the statistical uncertainty of the \( h_{ij}^{MD}(r) \) function was obtained with the block averaging method [30]. For each density, and in the whole range \( r/\sigma_{ij} < (1, r^*_c) \), the accuracy of \( h_{ij}^{MD}(r) \) was such that the estimated uncertainty was < 10^{-3}, being up to 0.002 near contact for the highest densities and becoming less than 0.0001 at larger particle separations. For large systems, the finite-size effects in the MD calculations of the RDF arise mainly from fixing the particle number, i.e., from the relation between canonical and grand-canonical ensembles. The corrections required to convert data from the MD simulations to the canonical ensemble are of \( O(1/N^2) \) [31, 32], which are negligible here. Also, it was checked for a few densities that the remaining part of the correction factor involving density derivatives was smaller than the obtained data accuracy and therefore could be neglected.

The resulting DCFs, \( c_{ij}^{W,M}(r) \), are shown in Fig. 4 together with the RFA and PY results, in the region \( 0 < r/\sigma_{ij} \). For all three DCFs and all studied partial packing fractions \( \eta_s \), there is very good agreement between the WM-scheme and the RFA results. The agreement with the PY approximation is less satisfactory and deteriorates significantly with increasing packing fraction. Figure 3 is supplemented by Fig. 4, where the DCFs are shown in the region \( r/\sigma_{ij} > 1 \). As in the monocomponent case [see Fig. 1(b)], the WM-scheme shows a (damped) oscillatory behavior, a feature not captured by the RFA. These small deviations between the monotonic (RFA) and the oscillatory (WM-scheme) results are expected to be reflected in the subleading pole density dependence. Also, it is worth noticing that the key region \( r > \sigma_{ij} \) is very well described by the RFA.

1. Determination of the poles of \( h_{ij}(k) \)

The obtained DCFs, \( c_{ij}(r) \), allow for the determination of the leading poles of \( h_{ij}(k) \) by application of the relations in Eqs. (21) and the results for the first two poles \( (\Pi_1 \text{ and } \Pi_2) \) are presented in Fig. 5. The pole \( \Pi_1 \) has \( \omega_1/\sigma_b \gtrsim 2\pi \), which corresponds to a wavelength in the oscillatory decay of \( h_{ij}(r) \) comparable to the diameter of the big spheres, while the pole \( \Pi_2 \) has \( \omega_2/\sigma_b \gtrsim 2\pi/\sigma_s \), corresponding to a wavelength comparable to the diameter of the small spheres. An interesting structural crossover [28, 29] occurs at \( \eta_s \approx 0.29 \), such that the leading pole (i.e., the pole with a smaller value of \( \alpha \)) changes from \( \Pi_1 \) if \( \eta_s < 0.29 \) to \( \Pi_2 \) if \( \eta_s > 0.29 \). The agreement between the WM-scheme and the theoretical predictions is very good, especially in the case of the RFA. In fact, the improvement of the RFA over the PY approximation is quite apparent for the subleading pole (i.e., \( \Pi_1 \) if \( \eta_s > 0.29 \) and \( \Pi_2 \) if \( \eta_s < 0.29 \)). This subleading pole reflects the role of the DCFs in the re-
FIG. 3. Plots of (a) small-small, (b) small-big, and (c) big-big DCFs in the region $0 < r < \sigma_{ij}$ for additive BHS mixtures with a size ratio $\sigma_s/\sigma_b = 0.648$ and a total packing fraction $\eta = 0.5$. The partial packing fractions are, from top to bottom, $\eta_b = 0.05, 0.10, 0.20, 0.30, 0.35$. The open circles represent the results obtained from the WM-scheme, the dashed lines are from the PY approximation, and the solid lines are from the RFA. In the insets, the differences $\Delta_{\text{PY}} = c^{\text{PY}}_{ij}(r) - c^{\text{RFA}}_{ij}(r)$ and $\Delta_{\text{RFA}} = c^{\text{WM}}_{ij}(r) - c^{\text{RFA}}_{ij}(r)$ are shown, those differences increasing with increasing $\eta_b$.

FIG. 4. Plots of (a) small-small, (b) small-big, and (c) big-big DCFs in the region $r > \sigma_{ij}$ for additive BHS mixtures with a size ratio $\sigma_s/\sigma_b = 0.648$ and a total packing fraction $\eta = 0.5$. The partial packing fractions are, from bottom to top, $\eta_b = 0.05, 0.10, 0.20, 0.30, 0.35$. The curves have been shifted vertically for better clarity. The open circles represent the results obtained from the WM-scheme and the solid lines are from the RFA. Note that $c^{\text{PY}}_{ij}(r) = 0$ for $r > \sigma_{ij}$.

gion $r > \sigma_{ij}$ and their subtle but important influence on the long-range structure of additive BHS fluid mixtures. We have checked that the behaviors $c_{ij}(r > \sigma_{ij})$ do indeed have an impact on the subleading pole by considering ‘hybrid’ DCFs given by the WM-scheme for $r < \sigma_{ij}$ and either $c_{ij}(r) = c^{\text{PY}}_{ij}(r) = 0$ or $c_{ij}(r) = c^{\text{RFA}}_{ij}(r)$ for $r > \sigma_{ij}$. From the results for the monocomponent HS fluid (see Fig. 2), it may be expected that the performance of the RFA for the subleading pole deteriorates at higher values of the total packing fraction of the system. Further studies at other conditions and for different BHS systems are needed to be performed.

In Fig. 5, the recent results by Statt et al. reflect quite well the trends of $\Pi_1$ and $\Pi_2$.

The structural crossover phenomenon is illustrated in Fig. 6, where the decay of $h_{ss}(r)$ at $\eta_s = 0.1$ and $\eta_s = 0.4$ is shown. The amplitudes and phases were calculated from the scheme described in Sec. II D. It was also verified numerically that the amplitude and phase relations $A_{ss} A_{bb} = A^2_{sb}$ and $\delta_{ss} + \delta_{bb} = 2\delta_{sb}$ were satisfied. It is observed that the wavelength of the oscillatory decay is close to $\sigma_b$ at $\eta_s = 0.1$, while it is close to $\sigma_s$ at $\eta_s = 0.4$. It is also noteworthy that a leading-pole representation of $h_{ss}(r)$ is already quite good, even for not large distances ($r \approx 2\sigma_b$). A two-pole representation ($M = 2$), including the leading and subleading terms, turns out to be excellent for distances beyond the second maximum.

IV. CONCLUSIONS

In this work we have investigated the SCFs of additive BHS mixtures. A scheme combining accurate MD simulation data, the pole structure representation of the total correlation functions $h_{ij}(r)$, and the OZ equation
monocomponent fluids, a slight discrepancy for higher densities is supposed to be caused by an oscillatory form of the DCFs at separations greater than the sphere diameter. In the case of the BHS mixtures analyzed \((\eta = 0.5)\), however, the agreement is found to be good also for the second pole. Thus, our results indicate that the RFA can predict well the long-range behavior of the SCFs of BHS mixtures, including the structural crossover, thus representing an improvement over the PY approximation.

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FIG. 5. Structure of the two leading poles for additive BHS mixtures with a size ratio \(\sigma_s/\sigma_b = 0.648\) and a total packing fraction \(\eta = 0.5\). Here, \(\alpha\) and \(\omega\) denote the damping coefficient and the oscillation frequency, respectively (see Eq. (3)). Big (green) circles are results from the WM-scheme (for each \(\eta_s, \alpha\) and \(\omega\) were obtained by solving Eqs. (21)), small (red) circles are simulations results from Statt et al. \([26]\), black solid lines are from the RFA, and grey solid lines are from the PY approximation. The horizontal blue double-sided arrow indicates the location of the structural crossover, which takes place near \(\eta_s = 0.29\) and \(\alpha \sigma_b = 2.4\). The left and right poles are denoted as \(\Pi_1\) (first) and \(\Pi_2\) (second), respectively (see main text).

has been developed. It is a nontrivial extension of the approach exploited previously for the monocomponent HS fluid \([28]\).

An important feature of the presented scheme is that some of the calculations can be performed analytically by taking into account the analytical forms for long distances (real space) and wave-numbers (Fourier space). In this way, the DCFs can be determined with great accuracy, thus allowing for calculating the density dependence of the leading poles and hence the decay of the pair correlation functions \(g_{ij}(r)\) of the bulk liquid mixtures.

The obtained results were compared with analytical predictions of the PY and RFA approximations. In the range of the studied densities, a very good agreement between the RFA and the calculated DCFs is found for separations less than \(\sigma_{ij}\). Such an agreement is observed also for the first pole. In the case of the second pole for

FIG. 6. Small-small correlation function \(h_{ss}(r)\) for additive BHS mixtures with \(\sigma_s/\sigma_b = 0.648, \eta = 0.5\), and (a) \(\eta_s = 0.1\) and (b) \(\eta_s = 0.4\). The open (red) circles are the MD data, while the solid (blue) and dot-dashed (black) lines represent the contribution of \(\Pi_1\) and \(\Pi_2\) pole, respectively. Note a change in the type of the leading pole describing the decay of the function at \(\eta_s = 0.1\) (\(\Pi_1\)) and \(\eta_s = 0.4\) (\(\Pi_2\)).
Thus, insertion of Eqs. (16) into Eq. (19b) gives

\[ C_{ij}^{(1)} = 4\pi \sigma_{ij} + 4\pi \sum_{n=1}^{\infty} A_{ij}^{(n)} e^{-\alpha_n \sigma_{ij}} \sin(\omega_n \sigma_{ij} + \delta_{ij}^{(n)}), \] (A1a)

\[ D_{ij}^{(1)} = -4\pi - 4\pi \sum_{n=1}^{\infty} A_{ij}^{(n)} e^{-\alpha_n \sigma_{ij}} \left[ \omega_n \cos(\omega_n \sigma_{ij} + \delta_{ij}^{(n)}) - \alpha_n \sin(\omega_n \sigma_{ij} + \delta_{ij}^{(n)}) \right], \] (A1b)

\[ C_{ij}^{(2)} = -4\pi \sum_{n=1}^{\infty} A_{ij}^{(n)} e^{-\alpha_n \sigma_{ij}} \left[ (\alpha_n^2 - \omega_n^2) \sin(\omega_n \sigma_{ij} + \delta_{ij}^{(n)}) - 2\alpha_n \omega_n \cos(\omega_n \sigma_{ij} + \delta_{ij}^{(n)}) \right], \] (A1c)

\[ D_{ij}^{(2)} = -4\pi \sum_{n=1}^{\infty} A_{ij}^{(n)} e^{-\alpha_n \sigma_{ij}} \left[ (\alpha_n^3 - 3\alpha_n \omega_n^2) \sin(\omega_n \sigma_{ij} + \delta_{ij}^{(n)}) + (\omega_n^3 - 3\alpha_n^2 \omega_n) \cos(\omega_n \sigma_{ij} + \delta_{ij}^{(n)}) \right]. \] (A1d)

Next, taking derivatives in Eq. (3) (for \( r > \sigma_{ij} \)), one finds

\[ g_{ij}^r(r) = \frac{1}{r} \frac{g_{ij}(r)}{r} + \sum_{n=1}^{\infty} A_{ij}^{(n)} r e^{-\alpha_n r} \left[ \omega_n \cos(\omega_n r + \delta_{ij}^{(n)}) - \alpha_n \sin(\omega_n r + \delta_{ij}^{(n)}) \right], \] (A2a)

\[ g_{ij}^r(r) = -2 \frac{g_{ij}(r)}{r} + \sum_{n=1}^{\infty} A_{ij}^{(n)} r e^{-\alpha_n r} \left[ (\alpha^2_n - \omega_n^2) \sin(\omega_n r + \delta_{ij}^{(n)}) - 2\alpha_n \omega_n \cos(\omega_n r + \delta_{ij}^{(n)}) \right], \] (A2b)

\[ g_{ij}^{(n)}(r) = -3 \frac{g_{ij}^{()}(r)}{r} - \sum_{n=1}^{\infty} A_{ij}^{(n)} r e^{-\alpha_n r} \left[ (\alpha_n^3 - 3\alpha_n \omega_n^2) \sin(\omega_n r + \delta_{ij}^{(n)}) + (\omega_n^3 - 3\alpha_n^2 \omega_n) \cos(\omega_n r + \delta_{ij}^{(n)}) \right]. \] (A2c)

Particularizing Eqs. (5) and (A2) to \( r = \sigma_{ij}^+ \), and taking into account Eqs. (A1), it is straightforward to get Eqs. (8).

**Appendix B: First few terms in \( c_{ij}^{\text{tail}}(r) \)**

Let us introduce the mathematical functions

\[ I_n(r, a) = \int_q^\infty dk \frac{\sin(ka) \cos(ka)}{k^n}, \quad J_n(r, a) = \int_q^\infty dk \frac{\sin(ka) \sin(ka)}{k^n}, \quad r, Q > 0. \] (B1)

Thus, insertion of Eqs. (16) into Eq. (19b) gives

\[ c_{ss}^{\text{tail}}(r) = \frac{C_{ss}^{(1)}}{2\pi^2 r} I_1(r, \sigma_s) + \frac{D_{ss}^{(1)}}{2\pi^2 r} J_2(r, \sigma_s) - \rho_s \frac{C_{ss}^{(2)}}{4\pi^2 r} \left[ I_3(r, 2\sigma_s) + I_3(r, 0) \right] \]

\[ -\rho_b \frac{C_{sb}^{(1)}}{4\pi^2 r} \left[ I_3(r, \sigma_s + \sigma_b) + I_3(r, 0) \right] + \frac{D_{ss}^{(2)}}{2\pi^2 r} J_4(r, \sigma_s) - \rho_s \frac{C_{ss}^{(1)}}{4\pi^2 r} \left[ I_3(r, \sigma_s + \sigma_b) + I_3(r, \frac{\sigma_b - \sigma_s}{2}) \right] \]

\[ c_{sb}^{\text{tail}}(r) = \frac{C_{sb}^{(1)}}{2\pi^2 r} I_1(r, \sigma_s) + \frac{D_{sb}^{(1)}}{2\pi^2 r} J_2(r, \sigma_s) + \frac{C_{sb}^{(2)}}{2\pi^2 r} I_3(r, \sigma_s) - \frac{C_{sb}^{(1)}}{4\pi^2 r} \left\{ \rho_s C_{ss}^{(1)} \left[ I_3(r, \sigma_s + \sigma_s) + I_3(r, \frac{\sigma_b - \sigma_s}{2}) \right] \right\} \]

\[ + \rho_b C_{bb}^{(1)} \left[ I_3(r, \sigma_s + \sigma_b) + I_3(r, \frac{\sigma_b - \sigma_s}{2}) \right] \]

\[ - \frac{D_{sb}^{(1)}}{4\pi^2 r} \left\{ \rho_s C_{ss}^{(1)} \left[ J_4(r, \sigma_s + \sigma_s) + J_4(r, \frac{\sigma_b - \sigma_s}{2}) \right] + \rho_b C_{bb}^{(1)} \left[ J_4(r, \sigma_s + \sigma_b) - J_4(r, \frac{\sigma_b - \sigma_s}{2}) \right] \right\} \]

\[ - \frac{C_{sb}^{(1)}}{4\pi^2 r} \left\{ \rho_s D_{ss}^{(1)} \left[ J_4(r, \sigma_s + \sigma_s) - J_4(r, \frac{\sigma_b - \sigma_s}{2}) \right] + \rho_b D_{bb}^{(1)} \left[ J_4(r, \sigma_s + \sigma_b) + J_4(r, \frac{\sigma_b - \sigma_s}{2}) \right] \right\} + \cdots. \] (B2b)
The exact expression for the function $I_1$ is

$$I_1(r, a) = \frac{\pi}{2} \Theta(r - |a|) - \frac{1}{2} \text{Si}(Q(r + a)) - \frac{1}{2} \text{Si}(Q(r - a)),$$  \hspace{1cm} (B3)

where

$$\text{Si}(z) = \int_0^z \frac{\sin t}{t} \, dt$$  \hspace{1cm} (B4)

is the sine integral function.

We observe from Eq. (B3) that $I_1(r, a)$ presents a discontinuity at $r = a$ of zeroth order. More specifically,

$$\lim_{r \to a^+} I_1(r, a) - \lim_{r \to a^-} I_1(r, a) = \frac{\pi}{2}.$$  \hspace{1cm} (B5)

On the other hand, the discontinuities of $J_2(r, a)$, $I_3(r, a)$, and $J_4(r, a)$ at $r = a$ are of first, second, and third order, respectively. Taking into account Eq. (B5), it is possible to obtain from Eqs. (B2) the results displayed in Eqs. (20).

**Appendix C: Integrals $I_1^{(a)}$ and $I_{sb}^{(a)}$**

In this Appendix the integrals required in Eqs. (21) are presented. They are

$$I_1^{(0)} = \frac{4\pi}{\omega} J_{ii}, \quad I_{sb}^{(0)} = -\frac{16\pi^2}{\alpha^2 + \omega^2} \left[ J_{ss} J_{bb} - K_{ss} K_{bb} - J_{sb}^2 + K_{sb}^2 + \frac{\alpha}{\omega} (J_{ss} K_{bb} + J_{bb} K_{ss} - 2 J_{sb} K_{sb}) \right],$$ \hspace{1cm} (C1a)

$$I_1^{(1)} = \frac{4\pi}{\alpha} K_{ii}, \quad I_{sb}^{(1)} = \frac{16\pi^2}{\alpha^2 + \omega^2} \left[ J_{ss} J_{bb} - K_{ss} K_{bb} - J_{sb}^2 + K_{sb}^2 - \frac{\omega}{\alpha} (J_{ss} K_{bb} + J_{bb} K_{ss} - 2 J_{sb} K_{sb}) \right],$$ \hspace{1cm} (C1b)

where

$$J_{ij} = \int_0^\infty dr \, c_{ij}(r) r \cos(\alpha r) \sin(\omega r), \quad K_{ij} = \int_0^\infty dr \, c_{ij}(r) r \sin(\alpha r) \cos(\omega r).$$  \hspace{1cm} (C2)

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[1] J.-P. Hansen and I. R. McDonald, *Theory of Simple Liquids*, 3rd ed. (Academic, London, 2006).
[2] J.-P. Hansen and I. R. McDonald, *Theory of Simple Liquids: with Applications to Soft Matter* (Academic Press, Oxford, 2013).
[3] J. A. Barker and D. Henderson, “What is “liquid”?” Understanding the states of matter,” Rev. Mod. Phys. 48, 587–671 (1976).
[4] N. H. March and M. P. Tosi, *Atomic Dynamics in Liquids* (Macmillan, London, 1976).
[5] D. M. Heyes, “The Lennard-Jones fluid in the liquid-vapour critical region,” Comput. Meth. Sci. Tech. 21, 169–179 (2015).
[6] M. Dijkstra, R. van Roij, and R. Evans, “Phase behavior and structure of binary hard-sphere mixtures,” Phys. Rev. Lett. 81, 2268–2271 (1998).
[7] A. Mulero, ed., *Theory and Simulation of Hard-Sphere Fluids and Related Systems*, Lecture Notes in Physics, Vol. 753 (Springer, Berlin, 2008).
[8] A. Santos, “Finite-size estimates of Kirkwood-Buff and similar integrals,” Phys. Rev. E 98, 063302 (2018).
[9] M. S. Wertheim, “Exact solution of the Percus–Yevick integral equation for hard spheres,” Phys. Rev. Lett. 10, 321–323 (1963).
[10] E. Thiele, “Equation of state for hard spheres,” J. Chem. Phys. 39, 474–479 (1963).
[11] J. L. Lebowitz, “Exact solution of generalized Percus–Yevick equation for a mixture of hard spheres,” Phys. Rev. 133, A895–A899 (1964).
[12] S. B. Yuste and A. Santos, “Radial distribution function for hard spheres,” Phys. Rev. A 43, 5418–5423 (1991).
[13] S. B. Yuste, M. López de Haro, and A. Santos, “Structure of hard-sphere metastable fluids,” Phys. Rev. E 53, 4820–4826 (1996).
[14] B. Yuste, A. Santos, and M. López de Haro, “Structure of multicomponent hard-sphere mixtures,” J. Chem. Phys. 108, 3683–3693 (1998).
[15] C. F. Tejero and M. López de Haro, “Direct correlation function of the hard-sphere fluid,” Mol. Phys. 105, 2999–3004 (2007).
[16] M. López de Haro, S. B. Yuste, and A. Santos, “Alternative Approaches to the Equilibrium Properties of Hard-
Sphere Liquids," in Theory and Simulation of Hard-Sphere Fluids and Related Systems, Lecture Notes in Physics, Vol. 753, edited by A. Mulero (Springer, Berlin, 2008) pp. 183–245.

[17] A. Santos, A Concise Course on the Theory of Classical Liquids, Basics and Selected Topics, Lecture Notes in Physics, Vol. 923 (Springer, New York, 2016).

[18] R. Fantoni and A. Santos, “Nonadditive hard-sphere fluid mixtures: A simple analytical theory,” Phys. Rev. E 84, 041201 (2011)

[19] R. Fantoni and A. Santos, “Multicomponent fluid of nonadditive hard spheres near a wall,” Phys. Rev. E 87, 042102 (2013)

[20] R. Fantoni and A. Santos, “Depletion force in the infinite-dilution limit in a solvent of nonadditive hard spheres,” J. Chem. Phys. 140, 244513 (2014)

[21] T. Boublík, “Hard-sphere equation of state,” J. Chem. Phys. 53, 471–472 (1970)

[22] E. W. Grundke and D. Henderson, “Distribution functions of multi-component fluid mixtures of hard spheres,” Mol. Phys. 24, 269–281 (1972)

[23] L. L. Lee and D. Levesque, “Perturbation theory for mixtures of simple liquids,” Mol. Phys. 26, 1351–1370 (1973)

[24] G. A. Mansoori, N. F. Carnahan, K. E. Starling, and T. W. Leland, Jr., “Equilibrium thermodynamic properties of the mixture of hard spheres,” J. Chem. Phys. 54, 1523–1525 (1971)

[25] R. Evans, R. J. F. Leote de Carvalho, J. R. Henderson, and D. C. Hoyle, “Asymptotic decay of correlations in liquids and their mixtures,” J. Chem. Phys. 100, 591–603 (1994)

[26] A. Statt, R. Pinchaipat, F. Turci, R. Evans, and C. P. Royall, “Direct observation in 3d of structural crossover in binary hard sphere mixtures,” J. Chem. Phys. 144, 144506 (2016)

[27] C. Grodon, M. Dijkstra, R. Evans, and R. Roth, “Decay of correlation functions in hard-sphere mixtures: Structural crossover,” J. Chem. Phys. 121, 7869–7882 (2004)

[28] S. Pieprzyk, A. C. Brańka, and D. M. Heyes, “Representation of the direct correlation function of the hard-sphere fluid,” Phys. Rev. E 95, 062104 (2017)

[29] M. N. Bannerman, R. Sargent, and L. Liee, “Dynamo: A free O(n) general event-driven molecular dynamics simulator,” J. Comput. Chem. 32, 3329–3338 (2011)

[30] M. P. Allen and D. J. Tildesley, Computer Simulation of Liquids (Oxford University Press, Oxford, 2017).

[31] J. J. Salacuse, A. R. Denton, and P. A. Egelstaff, “Finite-size effects in molecular dynamics simulations: Static structure factor and compressibility. I. Theoretical method,” Phys. Rev. E 53, 2382–2389 (1996)

[32] A. Baumkötter and Y. Hiwatari, “Finite-size dependence of the bridge function extracted from molecular dynamics simulations,” Phys. Rev. E 63, 061201 (2001)