Equation of state of sticky-hard-sphere fluids in the chemical-potential route

René D. Rohrmann
Instituto de Ciencias Astronómicas, de la Tierra y del Espacio (ICATE-CONICET), Avenida España 1512 Sur, 5400 San Juan, Argentina

Andrés Santos
Departamento de Física, Universidad de Extremadura, Badajoz, E-06071, Spain
(Dated: January 16, 2014)

In this paper we use the µ-route to evaluate the EOS of the three-dimensional sticky-hard-sphere (SHS) model introduced by Baxter [14]. In this fluid, impenetrable particles of diameter σ interact through a square-well potential of infinite depth and vanishing width. The study of this pair potential model has two advantages. First, it admits an exact analytical solution to the Ornstein–Zernike equation with the PY closure [14], its thermodynamic properties being described in terms of two simple parameters, the packing fraction η and a stickiness parameter α [13]. Second, the SHS model has proved to provide an excellent starting point for the study of colloidal systems with short-rang attraction [16, 20], interactions between protein molecules in solution [21], and other interesting applications [22, 23].

We will exploit the known exact solution of the PY integral equation for both single and multicomponent SHS fluids [14, 24] to obtain the EOS through the µ-route and compare the outcome with the three standard routes (virial, energy, and compressibility) and with Monte Carlo (MC) simulations [25]. As we will see, the µ-route EOS in the PY approximation changes with the choice of the prescription followed to switch-on the extra particle to the rest of the system. Despite this, the spread is typically much smaller than the one existing among the different routes. Interestingly enough, the µ-route generally provides the best results, including the gas-liquid transition properties of the fluid.

The paper is organized as follows. In Sec. II, we give...
the mathematical formulation of the \( \mu \)-route for SHS fluids. In Secs. III and IV we use the known exact and PY solutions of the SHS system in one and three dimensions, respectively, to derive the \( \mu \)-route EOS of those systems. Section IV is reserved for discussions of the results. The relevant calculations are presented in a series of Appendices.

II. THE CHEMICAL POTENTIAL ROUTE

We consider a \( d \)-dimensional system of volume \( V \) containing \( N = \rho V \) spherical particles of diameter \( \sigma \) with surface adhesion. The SHS interaction potential \( \phi(r) \) between two particle with centers separated at a distance \( r \) is defined by

\[
e^{-\beta \phi(r)} = \Theta(r - \sigma) + \alpha \sigma \delta(r - \sigma),
\]

where \( \beta = 1/k_B T, k_B \) and \( T \) being the Boltzmann constant and the absolute temperature, respectively. The dimensionless parameter \( \alpha \) measures the strength of surface adhesion (“stickiness”). The equality in (2.1) must be interpreted as the limit of an increasingly deeper (\( \epsilon \to \infty \)) and narrower (\( \Delta \to 0 \)) square-well potential of depth \( \epsilon \) and (relative) width \( \Delta \) with

\[
\alpha = e^{\beta \Delta}
\]

kept constant. The stickiness parameter \( \alpha \) is related to the “Baxter temperature” \( \tau \) \( \frac{12}{\alpha} \) by \( \tau = 1/12\alpha \). The pure HS model is recovered from Eq. (2.1) in the limit \( \alpha \to 0 \).

We now include into the system an additional particle (the “solute”). Its interaction with any other particle in the fluid (the “solvent”) is given by

\[
e^{-\beta \phi_\xi(r)} = \Theta(r - \xi \sigma) + \alpha_\xi \sigma \delta(r - \xi \sigma),
\]

where \( \xi \) plays the role of a coupling parameter and \( \alpha_\xi \) is a continuous function of \( \xi \) encoding the strength of the solute-solvent attractive force. It runs from \( \alpha_\xi = 0 \) at \( \xi = 0 \) to \( \alpha_\xi = \alpha \) at \( \xi = 1 \).

For large \( N \), the excess chemical potential \( \mu^{ex} = \mu - \mu^{id} \), \( \mu^{id} \) being the contribution of the corresponding ideal fluid, can be written as follows \( 2.4 \) \( 12 \) \( 13 \)

\[
- \beta \mu^{ex} = \ln \frac{Q^{(\xi = 1)}_{N+1}}{Q^{(\xi = 0)}_{N+1}},
\]

where

\[
Q^{(\xi)}_{N+1} = \frac{1}{V^{N+1}} \int \mathrm{d}r^N \int \mathrm{d}r_0 \ e^{-\beta \Phi^{(\xi)}_{N+1}(r^N, r_0)}
\]

is the configurational integral of \( N \) solvent particles plus one solute particle with a coupling parameter \( \xi \). Here, \( \mathbf{r}^{N+1} = \{ \mathbf{r}^N, \mathbf{r}_0 \} \), where \( \mathbf{r}^N \) refers to all the translational coordinates of the \( N \) solvent particles and \( \mathbf{r}_0 \) refers to the coordinates of the solute particle. Furthermore,

\[
\Phi^{(\xi)}_{N+1}(\mathbf{r}^N, \mathbf{r}_0) = \frac{1}{2} \sum_{i \neq j} \phi(r_{ij}) + \sum_{i = 1}^{N} \phi_\xi(r_{0i})
\]

is the total potential energy, \( r_{ij} \) being the distance between particles \( i \) and \( j \). Hence,

\[
Q^{(\xi)}_{N+1} = \frac{1}{V^{N+1}} \int \mathrm{d}r^N \ e^{-\beta \Phi_\xi(r^N)} \int \mathrm{d}r_0 \ \prod_{i = 1}^{N} e^{-\beta \Phi_\xi(r_{0i})},
\]

(2.7)

where \( \Phi_\xi(r^N) \) denotes the solvent potential energy.

For convenience, we decompose the right-hand side of Eq. (2.7) into two separate contributions,

\[
- \beta \mu^{ex} = \ln \frac{Q^{(\xi = 1)}_{N+1}}{Q^{(\xi = 0)}_{N+1}} + \int_0^1 \mathrm{d}\xi \ \partial \ln Q^{(\xi)}_{N+1} \frac{\partial \phi_\xi(r_{0i})}{\partial \xi}.
\]

(2.8)

Henceforth, for the sake of simplicity, we adopt \( \sigma = 1 \). The two contributions in Eq. (2.8) are worked out as follows. First, with (2.3), we write

\[
\prod_{i = 1}^{N} e^{-\beta \phi_\xi(r_{0i})} = \prod_{i = 1}^{N} \Theta(r_{0i} - \xi) + \alpha_{\xi} \sum_{i = 1}^{N} \delta(r_{0i} - \xi) \prod_{j \neq i} \Theta(r_{0j} - \xi) + (\alpha_{\xi})^2 \sum_{i, j} \delta(r_{0i} - \xi) \delta(r_{0j} - \xi)
\]

\[
\times \prod_{k \neq i, j} \Theta(r_{0k} - \xi) + O(\alpha_{\xi}^3).
\]

(2.9)

For \( \xi < \frac{1}{2} \), the \( N \) surfaces defined by \( r_{0i} = \xi \) \( (i = 1, \ldots, N) \) do not overlap, so that the condition \( r_{0i} = \xi \) implies \( r_{0j} > \xi \) \( \forall j \neq i \). As a consequence, the integrals in (2.7) of order two or higher in \( \alpha_{\xi} \) vanish. On the other hand, integration of \( \prod_{i = 1}^{N} \Theta(r_{0i} - \xi) \) over \( r_0 \) gives the free volume of the solute particle,

\[
\int \mathrm{d}r_0 \ \prod_{i = 1}^{N} \Theta(r_{0i} - \xi) = V - N \Omega_{\xi}, \quad \xi < \frac{1}{2},
\]

(2.10)

where \( \Omega_{\xi} = \frac{\pi d^2}{2}(1 + d/2)|\xi|^d \) is the volume of a \( d \)-sphere of radius \( \xi \). Furthermore,

\[
\int \mathrm{d}r_0 \ \delta(r_{0i} - \xi) \prod_{j \neq i} \Theta(r_{0j} - \xi) = \Sigma_{\xi}, \quad \xi < \frac{1}{2},
\]

(2.11)

\( \Sigma_{\xi} = \partial \Omega_{\xi}/\partial \xi = d \Omega_{\xi}/\xi \) being the surface of a \( d \)-sphere of radius \( \xi \). Therefore,

\[
\int \mathrm{d}r_0 \ \prod_{i = 1}^{N} e^{-\beta \phi_\xi(r_{0i})} = V - N \Omega_{\xi} + N \alpha_{\xi} \Sigma_{\xi}, \quad \xi < \frac{1}{2}.
\]

(2.12)

With this result, Eq. (2.7) yields

\[
Q^{(\xi)}_{N+1} = [1 - \rho \Omega_{\xi} (1 - d \alpha_{\xi})] Q_N, \quad \xi < \frac{1}{2},
\]

(2.13)
where
\[ Q_N = \frac{1}{V_N} \int \text{d}r_N e^{-\beta \Phi_N(r^N)} \]  
(2.14)
is the configurational integral of the solvent. As shown in Appendix A, the case \( \xi = \frac{1}{2} \) is singular if \( d \neq 2 \). This difficulty can be overcome by the choice \( \alpha_2 = 0 \). Therefore, taking into account that \( Q_{N+1}^{(0)} = Q_N \), and taking the limit \( \xi \to \frac{1}{2} \) in Eq. (2.13), the first term on the right-hand side of Eq. (2.8) becomes

\[ \ln Q_{N+1}^{(0)} / Q_{N+1} = \ln (1 - \eta), \]  
(2.15)

where
\[ \eta \equiv \rho \Omega_2 \]  
(2.16)
is the packing fraction.

For \( \xi > \frac{1}{2} \) one must follow another strategy because integration over \( \mathbf{r}_0 \) in Eq. (2.7) depends upon the coordinates of all the solvent particles. In this case, we consider

\[ \frac{\partial \beta^{-\Phi_N(r^N)} \partial \xi}{\partial \xi} = e^{-\beta \Phi_N(r^N)} \sum_{i=1}^N \frac{\partial e^{-\Phi_N(r^{(i)})}}{\partial \xi} \prod_{j \neq i} e^{-\beta \Phi_N(r^{(j)})}, \]

(2.17)

The solute-solvent RDF is expressed as

\[ g_\xi(r_0) = \frac{V^{-(N-1)}}{Q_{N+1}^{(0)}} \int \text{d}r_2 \cdots \int \text{d}r_N e^{-\beta \Phi_N(r^{(N+1)})}. \]

(2.18)

It follows from Eqs. (2.7), (2.17), and (2.18) that

\[ \frac{\partial \ln Q_{N+1}^{(0)}}{\partial \xi} = \frac{V^{-(N-1)}}{Q_{N+1}^{(0)}} \int \text{d}r^{(N+1)} \frac{\partial e^{-\beta \Phi_N(r^{(N+1)})}}{\partial \xi} \]

\[ = \frac{1}{V_2} \sum_{i=1}^N \int \text{d}r_0 \int \text{d}r \, y_\xi(r_0) \frac{\partial e^{-\beta \Phi_N(r_0)}}{\partial \xi}, \]

(2.19)

\[ \ln Q_{N+1}^{(0)} / Q_{N+1} \]

and in the last step of Eq. (2.14) we have used spherical coordinates.

Finally, inserting Eqs. (2.16) and (2.19) into Eq. (2.8), we obtain

\[ \frac{\partial \beta^{-\Phi_N(r)} \partial \xi}{\partial \xi} = \frac{\partial \beta^{-\Phi_N(r)} \partial \xi}{\partial \xi} = \frac{\partial (\alpha_2 - 1) \xi}{\partial \xi} \delta \left( r - \xi \right) - \alpha_2 \xi \frac{\partial \delta \left( r - \xi \right)}{\partial r}. \]  
(2.23)

Thus,

\[ M_\xi(\eta, \alpha) = \frac{\partial (\alpha_2 - 1) \xi}{\partial \xi} y_\xi(\xi) + \alpha_2 \xi y_\xi(\xi). \]  
(2.24)

We may now derive the compressibility factor \( Z \equiv \beta \rho / \rho \) (\( \rho \) being the pressure) in the \( \mu \)-route. The familiar thermodynamic relation

\[ \frac{\partial \eta}{\partial \rho} = \rho \frac{\partial \mu}{\partial \rho}, \]  
(2.25)

can be expressed as

\[ \frac{\partial |\eta(Z - 1)|}{\partial \eta} = \eta \frac{\partial (\beta \mu^{ex})}{\partial \eta}, \]  
(2.26)

and with Eq. (2.22) we obtain

\[ Z^{(\mu)}(\eta, \alpha) = \frac{-\ln(1 - \eta)}{\eta} - d \int_0^1 d\xi \left[ M_\xi(\eta, \alpha) \right] \]  
(2.27)

This constitutes the EOS of \( d \)-dimensional SHS obtained from the coupling parameter procedure. To have a more explicit expression for \( M_\xi \), we note that, according to Eq. (2.23),

\[ \ln Q_{N+1}^{(0)} / Q_{N+1} \]

**III. STICKY HARD RODS. EXACT RESULTS**

As a test of the correctness of Eq. (2.22), we prove in this section that it leads to the exact equation of state for the one-dimensional system (\( d = 1 \)). In that case, Eq. (2.22) reduces to

\[ \beta \mu^{ex}(\eta, \alpha) = -\ln(1 - \eta) - 2 \eta \int_0^1 d\xi M_\xi(\eta, \alpha). \]  
(3.1)

with

\[ M_\xi(\eta, \alpha) = \frac{\partial (\alpha_2 - 1) \xi}{\partial \xi} y_\xi(\xi) + \alpha_2 \xi y_\xi(\xi). \]  
(3.2)
As shown in Appendix C

\[ y_{\xi}(\xi) = \frac{\beta p/\eta}{1 + \beta \rho \alpha_{\xi} \xi}, \quad y'_{\xi}(\xi) = -\frac{\eta (\beta p)^{2}/\eta}{1 + \beta \rho \alpha_{\xi} \xi}. \] (3.3)

Thus, Eq. (3.2) may be written in the form

\[ M_{\xi}(\eta, \alpha) = -\frac{\beta p}{\eta} + \frac{1}{\eta} \frac{\partial}{\partial \xi} \ln(1 + \beta \rho \alpha_{\xi} \xi). \] (3.4)

Then, Eq. (3.1) becomes

\[ \beta \mu_{\mathrm{ex}}(\eta, \alpha) = -\ln(1 - \eta) + \beta p - 2 \ln(1 + \beta \rho \alpha). \] (3.5)

This result is exact and does not depend on the explicit form of \( \alpha_{\xi} \) in the interval \( \frac{1}{2} < \xi < 1 \). Making use of Eq. (C.4), it is straightforward to check that Eq. (2.26) is indeed satisfied.

**IV. STICKY HARD SPHERES. PERCUS–YEVICK THEORY**

The excess chemical potential for three-dimensional SHS fluids \( (d = 3) \) is obtained from Eqs. (2.22) and (2.24) as

\[ \beta \mu_{\mathrm{ex}}(\eta, \alpha) = -\ln(1 - \eta) - 24 \eta \int_{\frac{1}{2}}^{1} \frac{1}{\eta} \frac{\partial}{\partial \xi} M_{\xi}(\eta, \alpha), \] (4.1)

\[ M_{\xi}(\eta, \alpha) = \frac{\partial(\alpha_{\xi} - 1) \xi^{2} y_{\xi}(\xi)}{\partial \xi} + \alpha_{\xi} \xi \frac{\partial[y_{\xi}(\xi)]}{\partial r} \bigg|_{r=\xi}. \] (4.2)

The associated \( \mu \)-route compressibility factor is given by [see Eq. (2.27)]

\[ Z(\mu)(\eta, \alpha) = -\frac{\ln(1 - \eta)}{\eta} - 24 \eta \int_{\frac{1}{2}}^{1} \frac{1}{\eta} \frac{\partial}{\partial \xi} M_{\xi}(\eta, \alpha) \]

\[ - \int_{0}^{1} \frac{1}{\eta} \frac{\partial}{\partial t} M_{\xi}(\eta t, \alpha_{t}). \] (4.3)

The evaluation of Eq. (4.2) requires the contact values of the solute-solvent cavity function \( y_{\xi}(\xi) \) and its derivative \( \partial_{r} y_{\xi}(\xi) \). These may be obtained using the PY approximation for an SHS binary mixture (see Appendix D). In particular, \( y_{\xi}(\xi) \) and \( y'_{\xi}(\xi) \) are given by Eqs. (D.16) and (D.17), respectively.

For an explicit evaluation of (4.1)–(4.3) we need to specify the \( \xi \)-dependence of \( \alpha_{\xi} \) (within the constraints \( \alpha_{\xi} = 0 \) and \( \alpha_{1} = \alpha \)). In this paper, we shall consider results based on three representative prescriptions:

\[ \alpha_{\xi} = \begin{cases} (2 \xi - 1)^{2} \alpha, & \text{(A)}, \\ (2 \xi - 1) \alpha, & \text{(B)}, \\ \sqrt{2 \xi - 1} \alpha, & \text{(C)}. \end{cases} \] (4.4)

These three protocols are depicted in Fig. 1. In all of them, the solute-solvent stickiness monotonically grows from zero to the solventsolvant value as the solute diameter \( (2 \xi - 1) \) grows from zero to the solvent diameter \( (\sigma = 1) \). At a given solute diameter, the strength of the solute-solvent attraction increases when going from A to C.

Since the PY integral equation is an approximate theory, a common RDF is expected to yield different EOSs depending on the route followed. In the case of the \( \mu \)-route, as will be seen below, an extra source of thermodynamic inconsistency arises: the EOS depends on the choice for the protocol \( \alpha_{\xi} \).

**A. Virial expansion**

A standard method of examining different approximations in statistical mechanics is to compare the successive terms in the virial expansion of the compressibility factor. For SHS fluids,

\[ Z(\eta, \alpha) = 1 + \sum_{j=2}^{\infty} b_{j}(\alpha) \eta^{j-1}. \] (4.5)

The virial coefficients \( b_{j}(\alpha) \) in the virial, energy, compressibility, and chemical-potential routes can be respectively evaluated from Eqs. (B.4), (B.10), (B.8), and (4.3).

All the routes in the PY approximation yield the exact second and third virial coefficients:

\[ b_{2}(\alpha) = 4 - 12 \alpha, \] (4.6)

\[ b_{3}(\alpha) = 10 - 60 \alpha + 144 \alpha^{2} - 96 \alpha^{3}. \] (4.7)
The exact fourth virial coefficient is a sixth-degree polynomial in $\alpha$ [26], i.e.,

\[ b_4(\alpha) = b_4^{HS} + \sum_{i=1}^{6} b_{4,i} \alpha^i, \tag{4.8} \]

where the numerical coefficients are given by the first row of Table I. The PY predictions for $b_4(\alpha)$ depend on the thermodynamic route. They have the structure of Eq. (4.8), except that $b_{4,6} = 0$. The corresponding numerical coefficients are displayed in Table I. Note that the energy route is unable to fix the HS EOS, so that the coefficients $b_{4,i}^{HS}$ remain undetermined in that route. All the coefficients $b_{4,i}$ derived from the $\mu$-route are rational numbers although a limited number of digits is shown in Table I. Note that the three protocols of the $\mu$-route are rational numbers, i.e., $b_{\mu} = 0$. The corresponding numerical coefficients are displayed in Table I. Note that the energy coefficients $b_{4,2}$ and $b_{4,5}$ depend on the choice of $\alpha_\xi$. For an arbitrary function $\alpha_\xi$, they are

\[ b_{4,2}^{(\mu)} = 837 \left[ 1 + \frac{12}{31} \int_{\frac{3}{4}}^{1} d\xi \left( \xi^2 - \frac{1}{4} \right) \frac{\xi^2 \alpha_\xi^2}{\alpha^2} \right], \tag{4.9} \]
The three protocols in Eq. (4.3) have the common form 
\[ \alpha = (2\xi - 1)^q \alpha \]  
with \( q = 0.5, 1, \) and \( A \) respectively. Taking \( q > 0 \) as a free parameter and using Eqs. (4.9)–(4.12), it is possible to find the optimal value of \( q \) that makes \( b_4^{(\mu)}(\alpha) = b_4^{\text{exact}}(\alpha) \) for a given value \( \alpha > 0.282 \). For instance, the optimal values are \( q = 0.199, 1.208, 2.076, 3.702, \) and \( 4.997 \) for \( \alpha = 0.3, 0.5, 0.7, 0.9, \) and \( 1 \), respectively. For \( \alpha < 0.282 \) the mathematical solutions of \( b_4^{(\mu)}(\alpha) = b_4^{\text{exact}}(\alpha) = 0 \), but these are nonphysical values violating the condition \( \alpha > 0 \).

Figure 2 compares the exact \( b_4(\alpha) \) with various PY routes, where the Carnahan–Starling (CS) [5] value \( b_4^{\text{HS}} = 18 \) has been taken in the case of the energy route. For low and moderate stickiness (\( \alpha \lesssim 0.35 \)), small deviations occur among all PY solutions and a good agreement with the exact values is found. For \( \alpha \gtrsim 0.35 \), however, larger discrepancies occur, with the energy and virial routes showing the most extreme deviations with respect to the exact solution. The \( \mu \)-route predictions lie between the virial and the compressibility ones, becoming closer to the exact values as a softer stickiness prescription is used (protocol A).

Although, to the best of our knowledge, the fifth virial coefficient is not exactly known, it is worthwhile comparing the different PY predictions for it. They have the polynomial structure

\[ b_5^{(\mu)}(\alpha) = b_5^{\text{HS}} + \sum_{i=1}^{7} b_{5,i} \alpha^i, \]

the coefficients being presented in Table II. Again, all the coefficients are rational numbers. The dependence of \( b_5(\alpha) \) on the stickiness parameter is shown in Fig. 3 (with the CS choice \( b_5^{\text{HS}} = 28 \) for the energy route). As expected, the influence of the thermodynamic route on \( b_5(\alpha) \) is stronger than in the case of \( b_4(\alpha) \). The general shapes of \( b_5 \) in the \( \mu \)- and compressibility routes are intermediate between those in the virial and energy routes.

### B. Weakly sticky limit

As a complement of the virial expansion [1.5], it is of interest to examine the leading terms in the series expansion

\[ Z(\eta, \alpha) = Z_{\text{HS}}(\eta) + \sum_{i=1}^{\infty} Z_i(\eta) \alpha^i \]

of the compressibility factor in powers of the stickiness parameter. Obviously, the zeroth-order coefficient in the \( \alpha \)-expansion is just the compressibility factor of the pure HS system. Equation (4.14) can be interpreted as a high-temperature expansion.

Making use of the results of Appendix D in Eqs. (1.4), (1.10), (1.8), and (1.3), the first-order and second-order coefficients from the different routes can be derived. The results are displayed in Table III. Interestingly, one has \( Z_1^{(v)}(\eta) = Z_1(\eta) \), thus generalizing the results \( b_4^{(v)} = b_4^{(e)} \) and \( b_5^{(v)} = b_5^{(e)} \) observed in Tables I and II. This reinforces that the “natural” extension of the energy route to HS fluids is the virial equation of state [27, 28].

The HS EOS \( Z_{\text{HS}}^{(\mu)}(\eta) \) was already derived in Ref. [12]. We observe that \( Z_{\text{HS}}^{(\mu)}(\eta) \) is protocol-independent. This generalizes to any order in density the behavior observed for \( b_4^{(\mu)} \) and \( b_5^{(\mu)} \) in Tables I and II respectively. The

\[ b_4^{(\mu)} = -2646 \left[ 1 + \frac{24}{49} \int_{\frac{1}{2}}^{1} d\xi \left( \xi^2 + \frac{\xi^2 \alpha \xi}{6 \alpha} - \frac{3}{8} \frac{\xi^2 \alpha^2}{\alpha^2} \right) \right], \]

\[ b_5^{(\mu)} = 4104 \left[ 1 - \frac{3}{19} \int_{\frac{1}{2}}^{1} d\xi \left( 1 - \frac{4 \xi^2 \alpha \xi}{3 \alpha} \frac{\xi^2 \alpha^2}{\alpha^2} \right) \right], \]

\[ b_4^{(\mu)} = -2160 \left( 1 + \frac{2}{5} \int_{\frac{1}{2}}^{1} d\xi \frac{\xi^4 \alpha^3}{\alpha^3} \right). \]
TABLE III. Expressions for $Z_{HS}(\eta)$ and the coefficients $Z_1(\eta)$ and $Z_2(\eta)$ [cf. Eq. (4.14)].

|               | $Z_{HS}(\eta)$ | $Z_1(\eta)$ | $Z_2(\eta)$ |
|---------------|----------------|--------------|--------------|
| **PY-$\mu_A$** | $\frac{1+2\alpha+3\eta^2}{(1-\eta)^2}$ | $\frac{-12\eta(1+2\alpha)}{(1-\eta)^2}$ | $\frac{36\eta^2(2\alpha+2+3\eta)}{(1-\eta)^4}$ |
| **PY-$\mu_B$** | Undetermined | $\frac{-27\ln(1-\eta)}{(1-\eta)^3}$ | $\frac{-648\ln(1-\eta)}{(1-\eta)^4}$ |
| **PY-$\mu_C$** | $\frac{1+\alpha+\eta^2}{(1-\eta)^3}$ | $\frac{-27\ln(1-\eta)}{3(1-\eta)^2}$ | $\frac{27(1+\eta)}{(1-\eta)^3}$ |

expression of $Z_2(\eta)$ for arbitrary $\alpha$ is

$$Z_2(\eta) = -540 \frac{\ln(1-\eta)}{\eta} - \frac{18(30 - 105\eta + 122\eta^2 - 77\eta^3)}{(1-\eta)^4}$$

$$-3888 \int_{\frac{1}{2}}^{1} \frac{1}{1} \left( \xi - \frac{1}{2} \right) \left( \xi - \frac{1}{6} \right) \frac{\ln(1-\eta)}{\eta} d\xi$$

$$+ \frac{13\eta - \frac{3}{2} \eta^2}{(1-\eta)^4} \alpha^2. \quad (4.15)$$

The coefficients $Z_1(\eta)$ and $Z_2(\eta)$, relative to the virial-route predictions, obtained from various PY routes are shown in Fig. 3. The discrepancies grow as density increases. In particular, the largest inconsistencies occur between the energy and compressibility routes. On the other hand, the $\mu$-route (which exhibits a moderate dependence on the protocol in the case of $Z_2$) deviates only slightly from the virial route, especially in the case of $Z_2$.

C. Finite density and stickiness

After having examined the low-density and low-stickiness (or high-temperature) regimes, we now consider the full non-perturbative regime. The density dependence of the reduced pressure $\eta Z^{(\mu)}$ is plotted in Fig. 5 for different values of $\alpha$ and for the three protocols $[4,4]$. Of course, in the limit of zero stickiness ($\alpha = 0$), the choice of the protocol becomes irrelevant and one recovers the EOS $Z_{HS}(\eta)$ (see Table III) corresponding to the PY theory in the $\mu$-route [12]. As $\alpha$ increases, the pressure decreases with respect to the HS value and the influence of the protocol is practically negligible up to $\alpha \approx 0.5$. For higher stickiness, however, the values of $Z^{(\mu)}$ are increasingly sensitive to the protocol chosen. We observe that, as expected on physical grounds, the stronger the relative stickiness $\alpha \xi / \alpha$, the smaller the pressure.

The three higher values of $\alpha$ in Fig. 5 correspond to the gas-liquid critical values $\alpha_{c}^{(v)}$, $\alpha_{c}^{(e)}$, and $\alpha_{c}^{(c)}$ predicted by the PY approximation in the virial, energy, and compressibility routes, respectively (see below). In fact the kink in $Z^{(\mu)}$ at $\alpha = \alpha_{c}^{(c)}$ and $\eta = \eta_{c}^{(c)} \approx 0.12132$ reflects the fact that $(\eta_{c}^{(c)}, \alpha_{c}^{(c)})$ is the critical point for the existence of real solutions of the PY equation (see Appendix B).

In Fig. 6 we compare MC simulations at $\alpha = \frac{\eta}{\eta}$ with PY predictions from various routes (virial, energy, compressibility, and chemical potential). Since, as discussed before, the energy route leaves the “integration constant” $Z_{HS}(\eta)$ undetermined, henceforth the CS EOS $\frac{Z_{CS}(\eta)}{Z_{HS}(\eta)} = \frac{1 + \eta + \eta^2 - \eta^3}{(1-\eta)^3} \quad (4.16)$

will be taken to complete the determination of $Z^{(c)}(\eta, \alpha)$ via Eq. (B8), despite the fact that the choice $Z_{HS}(\eta) = Z^{(v)}$ would be more consistent [27, 28]. The virial and compressibility data have been obtained from Eqs. (B4) and (B10), respectively. In all the cases, use has been made of the PY solution detailed in Appendix D.
FIG. 6. (Color online) Reduced pressure \( \eta Z \) as a function of the packing fraction for SHS fluids at \( \alpha = \frac{\pi}{2} \approx 0.556 \) (\( \tau = \frac{3}{20} = 0.15 \)). The curves correspond to PY results from various routes as indicated on the plot. Open circles represent MC calculations [22].

FIG. 7. (Color online) Phase diagram of the SHS fluid showing the gas-liquid coexistence curves from PY solutions in the virial (−), energy (−−−), compressibility (−−−), and chemical-potential (−−−−) routes. Results in the \( \mu \)-route are based on prescription A. MC simulation data taken from Ref. [23] are shown with error bars. The critical points are indicated with circles.

We observe that in the low-density range (\( \eta \lesssim 0.15 \)) all PY routes and simulation data agree very well. For higher densities, the curves corresponding to the three different protocols of the \( \mu \)-route remain rather close in comparison with those from the virial, energy, and compressibility routes, which show a larger spread. In the range 0.2 \( \lesssim \eta \lesssim 0.4 \), the \( \mu \)-route gives the best fits to the simulation data. In the same region, \( Z^{(v)} \) and \( Z^{(e)} \) overestimate the simulation values, while \( Z^{(e)} \) underestimates them. Up to \( \eta \approx 0.4 \), one has \( Z^{(v)} < Z^{(\mu e)} < Z^{(\mu A)} < Z^{(\mu A)} < Z^{(c)} \). Finally, there is a rather strong disagreement of all the PY routes at high densities, 0.4 \( \lesssim \eta \lesssim 0.5 \), where the simulation data exhibit lower pressure values than the theoretical ones. The compressibility route shows the largest deviations from MC results on the whole range of studied densities.

D. Gas-liquid transition

The two highest values of \( \alpha \) in Fig. 5 show a domain of mechanical instability, \( \partial(\eta Z)/\partial \eta < 0 \) (i.e., \( \partial p/\partial \rho < 0 \)), which indicates a phase transition according to the \( \mu \)-route in the various protocols analyzed. As is well known, the SHS fluid has a (metastable) fluid-fluid transition which was early predicted in the compressibility [14] and energy [24] routes of the PY approximation. Critical values of the parameters \( \alpha \) and \( \eta \) (or \( \tau \) and \( \rho \)) can be determined by the conditions \( \partial p/\partial \rho = 0 \) and \( \partial^2 p/\partial \rho^2 = 0 \). Results concerning various PY routes are summarized in Table IV, where they can be compared to the ones obtained by Miller and Frenkel [23] using MC simulations.

As is known, the compressibility route produces a gross underestimation of the critical density. This quantity is much better approximated by the virial route (deviation of 5%) and, especially, the \( \mu \)-route (deviations of 4%, 1.2%, and 0.6% for protocols A, B, and C, respectively). On the other hand, the critical value of the stickiness parameter evaluated from the virial and the compressibility routes differ significantly from those predicted by numerical experiments. For this parameter, the energy route (deviation of 4%) and the \( \mu \)-route (deviations of 7%, 10%, and 13% for A, B, and C, respectively) give the best results. It must be remarked that the critical point obtained from \( Z^{(c)} \) is quite sensitive to the choice of \( Z_{HS} \). If, instead of the CS EOS [14,18], the more con-
sistent choice \( Z_{\text{HS}} = Z_{\text{HS}}^{(\mu)} \) is used, then no SHS critical point is predicted by the energy route. In conclusion, the parameters of the critical points obtained from the \( \mu \)-route show the best global agreement with simulations.

We have also computed coexistence curves with the familiar equal-area Maxwell construction that is applicable when \( \partial p/\partial \rho < 0 \). Figure 4 displays the coexistence curve and the location of the critical point derived by various PY routes and from computer evaluations 23. For simplicity, in the case of the \( \mu \)-route only results from protocol A are shown. As may be seen in Fig. 4 the curves obtained from the virial and compressibility routes differ substantially from computer evaluations. On the contrary, the agreement is reasonably good for the energy and \( \mu \)-routes. As already seen from Table IV the critical Baxter temperature \( \tau \), predicted by the energy route is more accurate than the one obtained from the \( \mu \)-route. However, the general shape of the coexistence curve (both the gas and the liquid branches) is better described by the \( \mu \)-route.

V. CONCLUSIONS

In this paper we have studied the SHS fluid using the concept of a partially coupled particle, whereby the interaction potential connecting this particle (the solute) to all other particles in the fluid (the solvent) is regulated by a charging parameter \( \xi \) which varies from \( \xi = 0 \) (no interaction) to \( \xi = 1 \) (full interaction). With this method, first introduced by Onsager 1 and subsequently developed by Kirkwood 2 and other authors 3,4, we have derived the chemical potential of \( d \)-dimensional SHS fluids in terms of the contact values of the solute-solvent cavity function \( y_\xi(\xi) \) and its first derivative \( y'_\xi(\xi) \) [see Eqs. (2.24) and (2.22)].

The procedure requires \( y_\xi(\xi) \) and \( y'_\xi(\xi) \) in the range \( \frac{1}{2} < \xi < 1 \), where the effective diameter \( 2(\xi - 1) \) of the coupled particle varies between zero to the solute diameter \( \sigma = 1 \). Thus, the explicit evaluation of the EOS of the fluid in the \( \mu \)-route requires the structural functions of the corresponding binary system in the infinite dilution limit. While this may represent a practical disadvantage with respect to the other standard routes (which only require the RDF of the one-component fluid), it nicely complements them. As is well known, the natural variables of the free energy \( F \) are the temperature \( (T) \), the volume \( (V) \), and the number of particles \( (N) \). The internal energy, the pressure and the isothermal compressibility are directly related to \( \partial F/\partial T \), \( \partial F/\partial V \), and \( \partial^2 F/\partial V^2 \), respectively. Therefore, the chemical potential, being related to \( \partial F/\partial N \), completes the picture.

The \( \mu \)-route also requires a prescription for the solute-solvent interaction potential, i.e., the dependence of the solute-solvent stickiness \( \alpha_{\xi} \) on the coupling parameter \( \xi \) must be specified. Here, in order to avoid the effects of trimer particle configurations, we have selected prescriptions with \( \alpha_{\xi} = 0 \) at \( \xi = \frac{1}{2} \). The resulting EOS, Eq. (2.20), is exact if the correct contact values \( y_\xi(\xi) \) and \( y'_\xi(\xi) \) are used, regardless of the explicit form of \( \alpha_{\xi} \) in the range \( \frac{1}{2} < \xi \leq 1 \). This is the case of the one-dimensional sticky fluid, in which case the exact EOS is recovered from the \( \mu \)-route (see Sec. III).

We have also applied the \( \mu \)-route to three-dimensional SHS fluids in the PY approximation. In this case, since the associated RDF is only approximate, the \( \mu \)-route EOS is influenced by the choice of the \( \alpha_{\xi} \)-protocol. On the other hand, this thermodynamical inconsistency becomes small in comparison with the spread of results obtained from the other three standard routes (virial, energy, and compressibility). When compared with available simulation data 22, the \( \mu \)-route EOS exhibits a general better agreement than those evaluated from the other routes. The gas-liquid phase transition has also been analyzed. The \( \mu \)-route provides the best prediction for the critical density, being only improved by the energy route in the prediction of the critical stickiness parameter. To put this latter fact in perspective, it is important to remark that, as usually done, the energy route has been complemented ad hoc by the accurate CS EOS for the HS fluid. As for the coexistence curve, the best global agreement with simulation data is obtained from the \( \mu \)-route. In addition, comparison of the fourth virial coefficient with exact results shows that the best performance corresponds to the \( \mu \)-route with a slower switching-on of stickiness (protocol A).

To conclude, we expect that the results presented in this paper may contribute to place the \( \mu \)-route on the same footing as the other three conventional routes. This is especially important in the case of mixtures 13, where the chemical-potential concept fits in a more natural way. Regarding the PY approximation, it is interesting to note that, whereas in the case of the HS fluid 12,13 the best behavior corresponds to the compressibility route (followed by the \( \mu \)-route), the inclusion of an attractive part in the interaction potential seems to favor the \( \mu \)-route as the most advantageous one.

ACKNOWLEDGMENTS

R.D.R. is grateful to the Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET, Argentina) by financial support through Grants Nos. PIP 112-200801-01474 and PIP 114-201101-00208. The research of A.S. has been supported by the Spanish Government and by the Junta de Extremadura (Spain) through Grants Nos. FIS2010-12587 and GRU10158, respectively, both partially financed by Fondo Europeo de Desarrollo Regional (FEDER) funds.
Appendix A: The limit $\xi \to \frac{1}{2}$

In contrast to the situation with $\xi < \frac{1}{2}$, a solute particle with $\xi = \frac{1}{2}$ allows spatial configurations where the solute and two solvent particles are simultaneously touching each other. Such configurations have a non-zero statistical weight in the evaluation of Eq. (2.7) through the term of order $\alpha_2^2$ in Eq. (2.9), unless $\alpha_2^2 = 0$. Such a term is

$$Q_{N+1}^{(\frac{1}{2},2)} = \frac{\alpha_2^2}{4N^{N+1}} \int dr^N e^{-\beta \Phi_N(r^N)} \int \prod_{i \neq j}^{N} \delta \left( r_{ij} - \frac{1}{2} \right)$$

$$\times \delta \left( r_{0i} - \frac{1}{2} \right) \prod_{k \neq i,j} \Theta \left( r_{0k} - \frac{1}{2} \right)$$

$$= \frac{\alpha_2^2 N(N-1)}{4V^{N+1}} \int dr^N e^{-\beta \Phi_N(r^N)}$$

$$\times \int \prod_{i \neq j}^{N} \delta \left( r_{ij} - \frac{1}{2} \right) \delta \left( r_{0i} - \frac{1}{2} \right) , \quad (A1)$$

where we have taken into account that, if $r_{0i} = r_{0j} = \frac{1}{2}$, one has $r_{0k} > \frac{1}{2}$ $\forall k \neq i,j$. Next, $r_{01} = r_{02} = \frac{1}{2}$ is compatible with $r_{12} \geq 1$ only if $r_{12} = 1$. Thus, using spherical coordinates,

$$Q_{N+1}^{(\frac{1}{2},2)} = \frac{\alpha_2^2}{8} \int \prod_{i \neq j}^{N-1} \delta \left( r_{ij} - \frac{1}{2} \right)$$

$$\times \prod_{k \neq i,j} \Theta \left( r_{0k} - \frac{1}{2} \right)$$

$$= \frac{\alpha_2^2}{8} \int dr^N e^{-\beta \Phi_N(r^N)} \delta (r_{12} - 1)$$

$$\times \int dr \delta (r - 1) g(r) , \quad (A2)$$

where

$$g(r) = \frac{V^{-(N-2)}}{Q_N} \int \delta_r d3 \cdot \int \delta_r e^{-\beta \Phi_N(r^N)} \quad (A3)$$

is the solvent RDF. Finally, introducing the solvent cavity function

$$y(r) \equiv g(r) e^{\beta \phi(r)} \quad (A4)$$

and using Eq. (2.9), we obtain

$$Q_{N+1}^{(\frac{1}{2},2)} = \frac{\alpha_2^2}{2} d^2 2^{d-3} \eta^2 y(1) Q_N \lim_{r \to 1} \left[ \Theta(r - 1) + \alpha \delta (r - 1) \right] , \quad (A5)$$

where $\eta$ is defined by Eq. (2.10). This contribution is singular by a two-fold reason. First, the Heaviside function implies that the result depends on whether $\xi \to \frac{1}{2}$ from below or from above. Second, and more importantly, the delta function gives a divergent term. Both singularities are avoided by the choice $\alpha_2^2 = 0$.

Appendix B: Virial, energy, and compressibility routes

For systems of particles interacting through two-body central forces, the thermodynamic functions can be evaluated in terms of the RDF $g(r)$. In particular, the pressure $p$, the excess internal energy per particle $u^e$, and the isothermal susceptibility $\chi$ are given by

$$p = \rho kT - \frac{\rho^2}{2d} \int dr \frac{\partial \phi(r)}{\partial r} r g(r) , \quad (B1)$$

$$u^e = \rho \int dr \phi(r) g(r) , \quad (B2)$$

$$\chi = k_B T \left( \frac{\partial \rho}{\partial r} \right) T = 1 + \rho \int dr [g(r) - 1] . \quad (B3)$$

Equations (B1), (B2), and (B3) are usually known as the pressure (or virial), energy, and compressibility equations, respectively.

For SHS fluids, the compressibility factor, $Z \equiv p/\rho kT$, can be expressed from Eqs. (B1) and (2.1) in terms of the contact values of the cavity function and its radial derivative $y^r(r)$ as

$$Z(v, \alpha) = 1 + 2^{d-1} \eta \{ y(1) - \alpha [dy(1) + y'(1)] \} . \quad (B4)$$

Here, the superscript $v$ specifies that the compressibility factor proceeds from the virial equation.

In turn, the excess of internal energy per particle is related with the compressibility factor as follows:

$$\rho \frac{\partial u^e}{\partial \rho} = -k_B T \frac{\partial Z}{\partial T} . \quad (B5)$$

For SHS fluids, the changes of variables $\rho \to \eta$ and $T \to \alpha$ yield

$$\eta \frac{\partial u^e / \epsilon}{\partial \eta} = \alpha \frac{\partial Z}{\partial \alpha} , \quad (B6)$$

where we have taken into account that, according to Eq. (2.2), $\partial T = -\epsilon \alpha / k_B T \partial \alpha$. Moreover, the excess energy can be expressed from Eq. (2.1) in terms of the cavity function using Eqs. (A11) and (2.11):

$$\frac{u^e - \epsilon}{\epsilon} = -d 2^{d-1} \eta \alpha g(1) . \quad (B7)$$

Integration of Eq. (B6) with (B7) yields the compressibility factor in the energy route as

$$Z^{(c)}(\eta, \alpha) = Z_{HS}(\eta) - d 2^{d-1} \eta \int_0^\alpha \frac{\partial [\eta y(1)]}{\partial \eta} \frac{\partial Z}{\partial \eta} , \quad (B8)$$

where $Z_{HS}(\eta)$ is the compressibility factor for pure HS (that here remains undetermined) and in the integrand $y(1)$ is a function of $\eta$ and $\alpha'$.

As for the compressibility route, taking into account that $\chi^{-1} = (\partial [\eta Z / \partial \eta])_T$ and introducing the moments

$$H_n = \int_0^\infty dr r^n h(r) \quad (B9)$$

of the total correlation function $h(r) = g(r) - 1$, one can find

$$Z^{(c)}(\eta, \alpha) = \int_0^1 \frac{dt}{\chi(\eta t)} = \int_0^1 \frac{dt}{1 + d 2^d \eta H_{d-1}(\eta t)} \quad (B10)$$
Appendix C: RDF of sticky hard rods

The exact solution of one-dimensional (d = 1) fluids with nearest-neighbor interactions is well known \[6, \ 10, \ 32, \ 34\]. In the case of an infinitely diluted solute particle in a solvent, the Laplace transform of the RDF \(g_\xi(r)\) is given by

\[
G_\xi(s) = \int_0^\infty dr \ e^{-rs}g_\xi(r) \quad (C1)
\]

of the RDF \(g_\xi(r)\) is given by

\[
G_\xi(s) = \frac{1}{\frac{\rho}{1 - \Psi(s + \beta p)}/\Psi(\beta p)} \quad (C2)
\]

where \(\Psi(s)\) and \(\Psi_\xi(s)\) are the Laplace transforms of \(e^{-\beta\phi(r)}\) (solvent-solvent interaction) and \(e^{-\beta\phi_\xi(r)}\) (solvent-solute interaction), respectively.

In the particular case of sticky hard rods, use of Eqs. 2.31 and 2.28 gives

\[
\Psi(s) = \left(\frac{1}{s + \alpha}\right) e^{-s}, \quad \Psi_\xi(s) = \left(\frac{1}{s + \alpha_\xi}\right) e^{-s\xi}. \quad (C3)
\]

Moreover, the exact equation of state is

\[
\beta p = \frac{\sqrt{1 + 4\alpha\gamma/(1 - \eta)} - 1}{2\alpha}. \quad (C4)
\]

Expansion of the right-hand side of Eq. (C2) in powers of \(e^{-s}\) allows one to obtain \(g_\xi(r)\) in the shells \(0 < r < 1 + \xi, 1 + \xi < r < 2 + \xi, \ldots\) In particular, if \(r < 1 + \xi,\)

\[
g_\xi(r) = \frac{\alpha_\xi\delta(r - \xi) + e^{-\beta p(r - \xi)}\Theta(r - \xi)}{\eta(\alpha_\xi + 1/\beta p)}, \quad r < 1 + \xi. \quad (C5)
\]

Taking into account Eqs. 2.3 and 2.20, one has

\[
y_\xi(r) = \frac{e^{-\beta p(r - \xi)}}{\eta(\alpha_\xi + 1/\beta p)}, \quad \xi \leq r < 1 + \xi. \quad (C6)
\]

From here one easily gets Eq. 3.3.

Appendix D: Solution of the PY equation for SHS

In this Appendix we summarize the main results obtained from the exact solution of the PY integral equation for SHS. The reader is referred to Refs. 14, 15, 24, 35, 39 for further details.

1. Solvent properties

The PY solution is expressed in terms of the Laplace transform

\[
G(s) = \int_0^\infty dr \ e^{-rs}rg(r) \quad (D1)
\]

of \(rg(r)\). Such a solution is

\[
s^2e^G(s) = \frac{L_0 + L_1s + L_2s^2}{\eta/2 - 12\eta\psi_2(s)L_0 + \psi_1(s)L_1 + \rho(0)L_2}, \quad (D2)
\]

where \(\psi_n(s) = \frac{1}{s^{n+1}} \sum_{m=0}^n \frac{(-s)^m}{m!} - e^{-s}\). \quad (D3)

The quantities \(L_0, L_1,\) and \(L_2\) are given as functions of \(\eta\) and \(\alpha\) by

\[
L_0 = \frac{1 + 2\eta}{(1 - \eta)^2} - \frac{12\eta}{1 - \eta}L_2, \quad L_1 = \frac{1 + \eta/2}{(1 - \eta)^2} - \frac{6\eta}{1 - \eta}L_2, \quad L_2 = \frac{1 - (1 - 2\alpha)\eta - K}{24\alpha(1 - \eta)\eta}, \quad (D4)
\]

where

\[
K = \sqrt{(1 - \eta)(1 - \eta(1 - 24\alpha + 48\alpha^2)))} + 72\alpha^2\eta^2. \quad (D6)
\]

The large-s behavior of \(G(s)\) provides the contact values of \(y(r)\) and \(y'(r)\). The results are

\[
y(1) = \frac{L_2}{\alpha}, \quad (D7)
\]

\[
y'(1) = -\frac{9\eta(1 + \eta)}{2(1 - \eta)^3} + \frac{12\eta(1 + 5\eta)}{(1 - \eta)^2}L_2 - \frac{12\eta(1 + 11\eta)}{1 - \eta}L_2 + 144\eta^2 L_2^3. \quad (D8)
\]

Insertion of these expressions into Eq. (B4) (with \(d = 3\)) gives the virial equation

\[
Z^{(c)} = \frac{1 + 2\eta + 3\eta^2}{(1 - \eta)^2} + 18\alpha \eta^2(1 + \eta)/(1 - \eta) - \frac{12\eta}{1 - \eta}\left[1 + 3\eta + 4\eta(1 + 11\eta)\right]L_2 + 48\eta^2\left[1 + \alpha + \frac{1 + 11\eta}{1 - \eta}\right]L_2^2 - 576\eta^3\alpha L_2^3. \quad (D9)
\]

Analogously, insertion of Eq. (D7) into Eq. (B8) yields an analytical expression for \(Z^{(c)} - Z_{HS}\).

The moment \(H_2\) of the total correlation function [cf. Eq. (B9)] can be obtained from the small-s behavior of \(G(s)\) as \(s^2G(s) = 1 + H_1 s^2 - 2H_2 s^3 + O(s^4)\). Inserting the resulting expression of \(H_2\) into \(\chi = 1 + 24\eta H_2\) [cf. Eq. (B3)], one obtains

\[
\frac{1}{\chi} = \frac{[1 + 2\eta - 12\eta(1 - \eta)L_2]^2}{(1 - \eta)^4}. \quad (D10)
\]

The compressibility factor \(Z^{(c)}\) by the compressibility route is readily obtained in analytical form by application
of Eq. (B10). For conciseness, the explicit expressions of $Z^{(c)}$ and $Z^{(e)}$ will be omitted here.

As a consequence of the square root present in $K(\eta, \alpha)$ [cf. Eq. (D6)], the PY solution is not physically meaningful if $\alpha > \alpha^{(c)}(\eta) \equiv 2\sqrt{2} \eta/(1-\eta) \simeq 0.535$ (or $\tau < 1/12\alpha^{(c)} = 0.09763$) and $\eta_+(\alpha) < \eta < \eta_+(\alpha)$, where

$$
\eta_\pm(\alpha) = \frac{1 - 12\alpha + 24\alpha^2 \pm 6\alpha \sqrt{2} - 16\alpha(1 - \alpha)}{1 - 24\alpha + 12\alpha^2}.
$$

(D11)

In the limit $\alpha \to \alpha^{(c)}$ one has $\eta_+ \to \eta^{(c)} = (3\sqrt{2} - 4)/2 \simeq 0.1213$. It can be easily checked that the right-hand side of Eq. (D11) vanishes at $(\eta, \alpha) = (\eta^{(c)}, \alpha^{(c)})$. This implies that $(\eta^{(c)}, \alpha^{(c)})$ is the critical point in the compressibility route.

2. Solute-solvent RDF

From the exact solution of the PY equation for an SHS binary mixture $^{24, 39}$ one can take the limit where one of the species (the solute) is infinitely dilute. As a result, the Laplace transform

$$
G_\xi(s) = \int_0^\infty dr \ e^{-rs} rs_\xi(r)
$$

(D12)
of $rs_\xi(r)$ is given by

$$
s^2 e^{s\xi} G_\xi(s) = \frac{L_0 + L_1^2 s + L_2^2 s^2}{1 - 12\eta \psi_2(s)L_0 + \psi_1(s)L_1 + \psi_0(s)L_2}.
$$

(D13)

where

$$
L_1^\xi = \frac{\xi + \eta(2\xi - 3/2)}{(1-\eta)^2} - \frac{6\eta(2\xi - 1)}{1-\eta} L_2,
$$

(D14)

$$
L_2^\xi = \left( \frac{1}{\alpha z_\xi} + \frac{6\eta}{1-\eta} - 12\eta L_2 \right)^{-1} L_1^\xi.
$$

(D15)

From the large-$s$ behavior of $G_\xi(s)$ we can obtain the contact values of $y_\xi(r)$ and $y_\xi'(r)$ as

$$
y_\xi(\xi) = \frac{L_2^\xi}{\alpha z_\xi^2}.
$$

(D16)

$$
\xi y_\xi'(\xi) = 12\eta L_2^\xi \left[ 3\eta (L_0 - 2L_1 + 2L_2)^2 - L_0 + L_1 \right]
$$

$$
+ L_0 + 6\eta L_1^\xi (L_0 - 2L_1 + 2L_2) - y_\xi(\xi).
$$

(D17)

ACKNOWLEDGMENTS

The work of R.D.R. has been supported by the Consejo Nacional de Investigaciones Científicas y Técnicas (CONICET, Argentina) through Grant No. PIP 112-200801-01474. A.S. acknowledges support from the Spanish government through Grant No. FIS2010-16587 and the Junta de Extremadura (Spain) through Grant No. GR10158, partially financed by Fondo Europeo de Desarrollo Regional (FEDER) funds.
[29] R. O. Watts, D. Henderson, and R. J. Baxter, Adv. Chem. Phys. 21, 422 (1971).
[30] R. Balescu, Equilibrium and Nonequilibrium Statistical Mechanics (Wiley, New York, 1974).
[31] J. A. Barker and D. Henderson, Rev. Mod. Phys. 48, 587 (1976).
[32] J. L. Lebowitz and D. Zomick, J. Chem. Phys. 54, 3335 (1971).
[33] A. Santos, Phys. Rev. E 76, 062201 (2007).
[34] See also A. Santos, “Radial Distribution Function for Sticky Hard Rods”,

http://demonstrations.wolfram.com/RadialDistributionFunction wolfram Demonstrations Project.
[35] S. B. Yuste and A. Santos, J. Stat. Phys. 72, 703 (1993).
[36] S. B. Yuste and A. Santos, Phys. Rev. E 48, 4599 (1993).
[37] B. Barboy, Chem. Phys. 11, 357 (1975).
[38] B. Barboy and R. Tenne, Chem. Phys. 38, 369 (1979).
[39] A. Santos, S. B. Yuste, and M. López de Haro, J. Chem. Phys. 109, 6814 (1998).