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To cite this version:

Alberto Parola, Davide Pini, Luciano Reatto. (SI)The smooth cut-off Hierarchical Reference Theory of fluids. Molecular Physics, 2009, 107 (04-06), pp.503-522. 10.1080/00268970902873547. hal-00513278ff

HAL Id: hal-00513278 <https://hal.science/hal-00513278v1>

Submitted on 1 Sep 2010

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The smooth cut-off Hierarchical Reference Theory of fluids

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We provide a comprehensive presentation of the Hierarchical Reference Theory (HRT) in the smooth cut-off formulation. A simple and self-consistent derivation of the hierarchy of differential equations is supplemented by a comparison with the known sharp cut-off HRT. Then, the theory is applied to a hard core Yukawa fluid (HCYF): a closure, based on a mean spherical approximation ansatz, is studied in detail and its intriguing relationship to the self consistent Ornstein-Zernike approximation (SCOZA) is discussed. The asymptotic properties, close to the critical point are investigated and compared to the renormalization group results both above and below the critical temperature. The HRT free energy is always a convex function of the density, leading to flat isotherms in the two-phase region with a finite compressibility at coexistence. This makes HRT the sole liquid-state theory able to obtain fluid-fluid phase equilibrium in homogeneous systems without resorting to the Maxwell construction. The way the mean field free energy is modified due to the inclusion of density fluctuations suggests how to identify the spinodal curve. Thermodynamic properties and correlation functions of the HCYF are investigated for three values of the inverse Yukawa range: $z = 1.8$, $z = 4$ and $z = 7$ where Monte Carlo simulations are available. The stability of the liquid-vapor critical point with respect to freezing is also studied.

PACS numbers: 64.60.F-, 61.20.Gy, 64.60.A-, 05.70.Fh

I. INTRODUCTION

Our understanding of the liquid-vapor phase transition was boosted, in the seventies, by the introduction in a statistical physics framework, of genuinely field theoretical methods: long wavelength effective actions and Renormalization Group (RG) techniques [1]. The microscopic justification of these coarse grained approaches has been the subject of several studies in the eighties [2–4] paving the way to quantitative investigations of the phase diagrams of simple liquids and mixtures [5].

is in the two-phase regar was a minit compressioning at ocksascie. The met consistence interesting to the Maxwell construction. The way the mean field free energy is shown to density the technomic density the theory able t In particular, the Hierarchical Reference Theory of fluids (HRT) [4, 6] can be considered as a successful attempt at developing the RG methods in the framework of a liquid-state theory. HRT is based on the microscopic hamiltonian of the fluid, but its RG-like structure endowes it with a number of features that are not shared by other theories of liquids: specifically, a non-trivial description of critical behavior that incorporates universality and scaling and, even more remarkably, a treatment of first-order phase transitions that does not violate the convexity of the free energy required by thermodynamic stability, thereby yielding rigorously flat pressure vs. density isotherms whenever different phases coexist at equilibrium. The region of phase coexistence is therefore straightforwardly obtained by HRT as the locus of diverging compressibility, and does not have to be recovered a posteriori by enforcing thermodynamic equilibrium via the Maxwell construction, that is instead necessary in other theories of homogeneous liquids, and may turn out to be quite hard to implement, especially in the critical region or in the case of binary fluids.

Notwithstanding these qualities, the original formulation of HRT leaves room for improvement. Specifically, its most serious deficiency consists in the fact that it fails to predict the discontinuity of the inverse compressibility expected in systems with a scalar order parameter, when the coexistence curve is crossed at any noncritical temperature. In fact, HRT predicts a diverging compressibility not only inside the coexistence region, but also on its boundary, so that the coexistence and spinodal curves coincide [7]. Besides preventing one from recovering the correct behavior of the compressibility at coexistence, this deficiency also forbids the possibility for the system to survive in a single phase inside the coexistence region as a metastable state — an occurrence that would be allowed to states located between the coexistence and the spinodal. One might think of this as a minor problem, since the very concept of a spinodal curve has little to do with equilibrium thermodynamics. However, the instance of metastability and its breakdown via the process of spinodal decomposition are experimentally relevant, especially in the context of complex fluids. These systems, often characterized by short range, tunable potentials and/or competing interactions, now offer the possibility to tailor phase diagrams and thermophysical properties [8] giving rise to phase boundaries of different topologies. Non-equilibrium states play a major role in these systems, being rather ubiquitous in colloidal physics. In particular, the relevance of spinodal decomposition for the occurrence of attractive gel phases has recently been pointed out [9].

Here we present a new formulation of HRT, the smooth cut-off HRT, that is free from the aforementioned shortcoming. We will detail the physical motivations at the basis of this generalization, together with the technical implementation of the smooth cut-off HRT to a rather flexible class of models: the Yukawa fluids. For this class of

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potentials, the smooth cut-off HRT allows also to implement exactly the requirement that the particles be mutually impenetrable because of the hard-core part of the interaction, usually referred to as the core condition. A numerical study of the resulting equations allows to test the accuracy of the theory via the comparison with available numerical simulations.

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II. THE SMOOTH CUT-OFF HRT EQUATIONS

Following the program previously outlined, we consider a gas of classical particles in dimension d, interacting via a spherically symmetric two body potential $v(r)$ written as the sum of a reference part and a tail $w(r)$:

$$
v(r) = v_R(r) + w(r) \tag{1}
$$

Then we define a sequence of intermediate systems interacting with potential $v_t(r)$ parametrized by $t \in (0,\infty)$ which interpolates between $v_R(r)$ (for $t = 0$) and $v(r)$ (as $t \to \infty$). This procedure is able to capture the RG spirit whenever $v_t(r)$ effectively suppresses the Fourier components of $w(r)$ with $k < Q + e^{-t}$. This effect can be obtained by different definitions of $v_t(r)$. Here we concentrate on the following class of parametrizations [12]:

$$
v_t(r) = v_R(r) + w_t(r)
$$

= $v_R(r) + [w(r) - \psi(t) e^{-dt} w(r e^{-t})]$ (2)

where $\psi(t)$ is a monotonically decreasing switching function, to be specified later, vanishing as $t \to \infty$ and satisfying $\psi(0) = 1$. The Fourier transform of $w_t(r)$ is given by:

$$
\tilde{w}_t(k) = \tilde{w}(k) - \psi(t) \tilde{w}(k e^t)
$$
\n(3)

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For a short range interaction characterized by an attractive tail with Fourier components $\tilde{w}(k)$, negligible beyond a certain wave vector Q^* , the intermediate potential $\tilde{w}_t(k)$ coincides with $\tilde{w}(k)$ for $k > Q^* e^{-t}$ while $\psi(t) < 1$ inhibits the Fourier components at $k < Q^*$. Therefore the definition (2) does satisfy the requirements discussed in the introduction, gives rise to a smooth, regular interaction at all t's and may represent the starting point for a microscopic implementation of the RG approach.

First order perturbation theory [16] provides the exact expression for the change in the free energy density of this model when t is varied:

$$
\frac{dA_t}{dt} = -\beta \frac{\rho^2}{2} \int d\mathbf{r} \, g_t(r) \, \frac{dw_t(r)}{dt} \tag{4}
$$

where A_t is minus the excess free energy density divided by k_BT , $g_t(r)$ is the radial distribution function of the intermediate system and $\beta = 1/k_BT$. This equation relates the change in the free energy with the change in the interaction but requires the knowledge of the two body correlations. An analogous equation for the pair correlations can be obtained starting from the formal diagrammatic expansion of the free energy functional in powers of the interaction, as derived in Ref. [6]:

$$
\frac{d}{dt}\tilde{c}_{t}^{(2)}(k) = -\beta \int \frac{d\mathbf{q}}{(2\pi)^{d}} \left[\frac{1}{2}\tilde{c}_{t}^{(4)}(\mathbf{k}, -\mathbf{k}, \mathbf{q}, -\mathbf{q}) + \tilde{c}_{t}^{(3)}(\mathbf{k}, -\mathbf{q}, \mathbf{q} - \mathbf{k}) \tilde{F}_{t}^{(2)}(|\mathbf{k} - \mathbf{q}|) \tilde{c}_{t}^{(3)}(-\mathbf{k}, \mathbf{q}, \mathbf{k} - \mathbf{q}) \right] \left[\tilde{F}_{t}^{(2)}(q) \right]^{2} \frac{d\tilde{w}_{t}(q)}{dt}
$$
\n(5)

where $\tilde{c}_{t,j}^{(n)}(\mathbf{k}_1 \dots \mathbf{k}_n)$ is the *n*-particle direct correlation function (including the ideal gas contribution) defined in terms of the n^{th} functional density derivative of the total free energy of the partially interacting system [6]. In particular, $\tilde{c}_{t}^{(2)}(k)$ is related to the usual direct correlation function $\tilde{c}_{t}(k)$ by

$$
\tilde{c}_t^{(2)}(k) = -\frac{1}{\rho} + \tilde{c}_t(k).
$$

The Ornstein-Zernike equation connects $\tilde{c}_{t}^{(2)}(k)$ to $\tilde{F}_{t}^{(2)}(k)$, which equals the structure factor $S_{t}(k)$ multiplied by ρ :

$$
\tilde{F}_t^{(2)}(k) = \rho S_t(k) = \frac{\rho}{1 - \rho \tilde{c}_t(k)} = \rho + \rho^2 \int d\mathbf{r} e^{i\mathbf{k} \cdot \mathbf{r}} \left[g_t(r) - 1 \right].
$$

tring from the formal diagrammatic expansion of the free energy further dual of the free energy further (6):
 $\frac{d\mathbf{q}}{2\pi\gamma^d} \left[\frac{1}{2} \hat{c}_i^{(4)} (\mathbf{k}, -\mathbf{k}, \mathbf{q}, -\mathbf{q}) + \hat{c}_i^{(3)} (\mathbf{k}, -\mathbf{q}, \mathbf{q} - \mathbf{k}) \hat{F}_i^{(2)} (\math$ Equations (4,5) are the first two terms of an exact hierarchy of differential equations for the free energy and the many body direct correlation functions of the model. Together with the definition of $w_t(r)$ (2) this hierarchy defines the smooth cut-off formulation of HRT. It is interesting to notice that when Eq. (5) is evaluated at vanishing external momentum $(k = 0)$, it becomes equivalent to the second density derivative of the HRT equation for the free energy (4). An analogous correspondence takes place throughout the full hierarchy, each differential equation being related to the density derivative of the previous one. This coincidence is not accidental, because the direct correlation functions are just functional derivatives of the free energy, and the functional derivative evaluated at zero momentum reduces to the partial derivative with respect to the uniform density. This shows that the first HRT equation (4) does contain information on the structure of the full hierarchy by virtue of the exact relationship between the free energy and the direct correlation function implied by the compressibility sum rule:

$$
\int d\mathbf{r} c_t(r) = \frac{\partial^2 A_t}{\partial \rho^2}.
$$
\n(6)

In order to exploit the connection of the HRT equations with the RG theory, it is convenient to rescale appropriately the set of direct correlation functions. As previously discussed, the switching parameter t provides the wavevector scale of density fluctuations $Q \div e^{-t}$. At criticality, according to general scaling arguments [11], the expected behavior of the n-particle direct correlation function (also known as one particle irreducible n-point function) is then given by

$$
\tilde{c}_t^{(n)}(\mathbf{q}_1 \dots \mathbf{q}_n) = -e^{\left[\frac{n}{2}(d-2+\eta)-d\right]t} u_t^{(n)}(\mathbf{q}_1 e^t \dots \mathbf{q}_n e^t)
$$
\n⁽⁷⁾

where η is the critical exponent related to the anomalous dimension [1]. The scaling form which characterizes correlations at the critical point is recovered provided the function $u_t^{(n)}(\mathbf{x}_1 \dots \mathbf{x}_n)$ tends to a finite limit as $t \to \infty$. By substituting Eq. (7) into the full (infinite) set of HRT equations, it is easy to show that the explicit dependence on the parameter t disappears from the equations for the scaling functions $u_t^{(n)}$ if the switching function is given by $\psi(t) = e^{-(2-\eta)t}$. With this choice of $\psi(t)$, the full hierarchy, rescaled through the definition (7), admits a fixed point solution providing the formal link with RG methods and allowing for a microscopic derivation of the known results on

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scaling laws and critical exponents [12] in full agreement with the known dimensionality expansion [1]. Note however that the scaling form is expected to hold only at small wavevectors, i.e. at large t , implying that the HRT equations provide a microscopic implementation of the RG procedure for any choice of $\psi(t)$ characterized by the asymptotic behavior:

$$
\lim_{t \to \infty} e^{(2-\eta)t} \psi(t) = \psi_{\infty} \tag{8}
$$

with ψ_{∞} finite and non zero.

A. Sharp vs. smooth cut-off procedure

En manguator (14); 1 are ocquate or metrinecate or particular and the full
tend manguary and ϕ or ϕ/ν way $\phi/\gamma/\nu$ interpolates between the efferince system and the full
the σ' of potential $\delta w_Q = w_Q - a_Q - w_Q$ includ The original implementation of HRT [4] was based on a different switching-on of the attractive interaction, inspired by the momentum shell integration RG [11]. The sequence of intermediate systems are defined by an attractive part of the potential $w_Q(r)$ whose Fourier components are set to zero below a cut-off wave vector Q. By varying Q from infinity to zero $v_Q(r) = v_R(r) + w_Q(r)$ interpolates between the reference system and the fully interacting one. In this case, however, the "slice" of potential $\delta w_Q \equiv w_{Q-dQ} - w_Q$ included at each step is characterized by a finite Fourier transform $\tilde{w}(Q)$ in an infinitesimal domain $(Q - dQ, Q)$. Such a singular form of δw_Q has profound consequences on the structure of the HRT equations: now, first order perturbation theory is not sufficient to evaluate the change in the free energy when Q is infinitesimally decreased and a resummation of the ring diagrams is required [4]. Therefore, the HRT equations in the sharp cut-off formulation do not coincide with Eqs. (4,5): in particular, the momentum integrations are limited to the shell $q = Q$ suggesting a decoupling of the long wavelength properties of the fluid model. In the critical region, the sharp cut-off HRT hierarchy acquires a form independent of the interparticle interaction, leading in a natural way to the concept of universality of the critical behavior. This appealing feature is not shared by the smooth cut-off hierarchy where the proof of universality requires a more elaborate analysis of the equations [12]. On the other hand, the intermediate potentials $w_O(r)$ in the sharp cut-off formulation display unphysical long range oscillatory tails as a consequence of the discontinuity in Fourier space. This circumstance brings about some difficulty in the formulation of an approximate closure to the infinite hierarchy of differential equations, due to the absence of an underlying physical intuition on the properties of fluid models characterized by such a class of potentials. The simple Ornstein-Zernike (OZ) closure to the first equation of the hierarchy adopted in all the previous investigations of HRT, although provides non classical critical exponents and scaling law together with flat isotherms at coexistence, has been proved to induce some artificial feature in the description of first order phase transitions leading to the divergence of the compressibility at the phase boundary, i.e to the coincidence of the spinodal with the binodal line [7]. Here we will show how the same class of OZ closures instead provides a satisfactory description of the phase diagram of simple fluids within the smooth cut-off formulation of the HRT equation for the free energy (2,4,8).

III. A SIMPLE CLOSURE FOR YUKAWA POTENTIALS

The OZ closure we are going to study (named HRT-OZ) amounts to parametrize the radial distribution function for the intermediate system $g_t(r)$ appearing in the first equation of the HRT hierarchy (4). Here we will focus on a simple yet flexible class of interactions: the hard core plus Yukawa potentials defined as the sum of a pure hard core term of diameter σ and an attractive Yukawa tail of inverse range z

$$
w(r) = -\epsilon \frac{e^{-z(r-\sigma)}}{r}
$$
 (9)

In the following, σ and ϵ/k_B will be taken as units of length and temperature respectively (i.e. we set $\sigma = 1, \epsilon = k_B$).

As previously noticed, in order to ensure that a memory of the full structure of the HRT hierarchy is contained in the free energy equation (4), the compressibility sum rule (6) has to be satisfied within the parametrization. A simple way to implement such a requirement is to adopt an approximate form of the correlation functions coming from some accurate liquid state theory suitably generalized in order to allow for the consistency between free energy and correlations (6). In order to take advantage of the analytical form available for the correlation functions of a Yukawa fluid, we have chosen to use, as a supporting liquid state theory, the Mean Spherical Approximation (MSA) defined by the usual core condition $(g_t(r) = 0$ for $r < 1$) supplemented by the explicit form of the direct correlation function outside the core:

$$
c_t(r) = -\beta (w_t(r) + \lambda_t w(r))
$$

= -\beta(1 + \lambda_t) w(r) + \beta \psi(t) e^{-dt} w(r e^{-t}) \qquad \text{for} \quad r > 1 \qquad (10)

where λ_t is implicitly defined by the compressibility sum rule (4) in terms of the excess free energy A_t . The radial distribution function and the direct correlation function are connected via the OZ equation. Within MSA the direct correlation function is always analytic in k^2 at small k, even at the critical point, implying that the critical exponent η vanishes within this class of approximations. For the HCYF, Eq. (10) implies a direct correlation function written as the sum of *two* Yukawa's of different ranges z_i and amplitudes K_i :

$$
c_t(r) = K_1 \frac{e^{-z_1(r-1)}}{r} + K_2 \frac{e^{-z_2(r-1)}}{r}
$$
\n(11)

outside the particle core (for i.e. $r > 1$), with

$$
z_1 = z
$$

\n
$$
z_2 = ze^{-t}
$$

\n
$$
K_1 = \frac{1+\lambda_t}{T}
$$
\n(12)

$$
K_2 = -\frac{e^{z(1-e^{-t})}}{T} \psi(t) e^{-(d-1)t}
$$
\n(13)

 $K_1 = \frac{r_1 + \alpha_1}{T}$
 $K_2 = -\frac{e^{z(1-e^{-t})}}{T} \psi(t) e^{-(d-1)t}$

the MSA integral equation can be solved analytically for this class of

set of algebraic equations [17]. Within HRT-OZ, this set is coupled ibility

itelligy sum In three dimensions, the MSA integral equation can be solved analytically for this class of parametrizations and the solution reduces to a set of algebraic equations [17]. Within HRT-OZ, this set is coupled to the evolution equation (4) via the compressibility sum rule (6) which implicitly defines the parameter λ_t present in K_1 . The explicit form of the equations, together with a detailed description of the numerical scheme adopted for their resolution, can be found in Appendix A. Here we remark that, as discussed in detail in Appendix A, the parametrization (10) also allows to improve the performance of the MSA (and then of the HRT-OZ) in the high density region, where the hard sphere direct correlation function is known to display a non negligible tail outside the particle core. If this additional contribution to $c_t(r)$ is represented as a Yukawa tail of the same inverse range z as that of the attractive part of the potential, its presence is equivalent to a redefinition of the parameter λ_t for $t = 0$, whose value is determined by Eq. (6) if the Carnahan-Starling expression for the hard sphere free energy is used.

A. HRT-OZ vs. SCOZA

The Self consistent Ornstein Zernike Approximation (SCOZA) is a well known liquid state theory which proved successful in the determination of phase diagrams in Yukawa fluids [18]. Originally, SCOZA has been formulated as a generalization of the MSA expression for the direct correlation function outside the core:

$$
c(r) = -\beta \lambda w(r) \tag{14}
$$

where the parameter λ is determined by requiring thermodynamic consistency between the internal energy and the compressibility routes to thermodynamics. In terms of the excess free energy density (divided by $-k_BT$) A, this condition reads:

$$
\frac{\partial A}{\partial \beta} = -\frac{\rho^2}{2} \int d\mathbf{r} \, g(r) \, w(r) \tag{15}
$$

$$
\frac{\partial^2 A}{\partial \rho^2} = \int d\mathbf{r} c(r) \tag{16}
$$

The two exact expressions (15,16), together with the core condition, $g(r) = 0$ for $r < 1$, and the parametrization (14) for $r > 1$, give rise to a partial differential equation for the free energy as a function of ρ and β . The close similarity between Eqs. (15,16) and (4,6) suggests that SCOZA can be somehow related to the smooth cut-off formulation of HRT. In fact, by defining the special sequence of intermediate potentials $w_t(r)$ as:

$$
w_t(r) = t w(r) \tag{17}
$$

the HRT-OZ equations (4,6) exactly reproduce SCOZA, showing that the two theories differ just by the switchingon procedure of the attractive part of the interaction: while SCOZA tunes the amplitude of $w(r)$, HRT changes simultaneously both its amplitude and range. This relationship is not peculiar to the Yukawa potential but holds for every choice of the interaction. Note that the consistency condition between free energy and internal energy (15) is generally violated within the HRT-OZ approximation: in order to enforce Eq.

(15) a further adjustable parameter in $c_t(r)$ is therefore required, as observed in Ref. [19] where a unification scheme between the sharp cut-off HRT-OZ and SCOZA was investigated.

The long range repulsive tail introduced by HRT during the integration, allows to inhibit phase separation at all finite t's thereby mimicking the RG approach. In order to emphasize the key role played by the weak long range repulsive tail added to the physical interaction in the smooth cut-off HRT, let us consider the simple MSA for a Yukawa fluid, which, at low temperatures, is known to display a region inside the spinodal where no solution exists. However, as shown in Fig. 1, if the physical attractive Yukawa potential is supplemented by a long range repulsive Yukawa contribution of the form (2), the spinodal curve disappears and a solution of the MSA equation exists in the whole phase diagram. When $t \to \infty$, the repulsive term in the potential becomes vanishingly small: the isotherm becomes flat and the compressibility diverges in the whole region inside the spinodal of the single Yukawa, thereby enforcing the convexity of the free energy. Therefore, cutting off long wave length density fluctuations by the addition of a small, long ranged repulsive tail appears to be an easy way to implement one of the most important features of equilibrium statistical mechanics within an approximate liquid state theory. Note however that within MSA, this kind of "regularization" of the physical two body interaction leads to the collapse of the spinodal and coexistence curve.

On the basis of these observations, the smooth cut-off HRT-OZ approach is then expected to provide a convex free energy: in the following Section we will show that HRT-OZ enjoys other relevant physical properties of first and second order phase transitions.

IV. ASYMPTOTIC ANALYSIS

For the parametrology metascular cases of the conduct of HRT-OZ approach is then experience observations, the smooth cut-off HRT-OZ approach is then experience observations.
 FV. ASYMPTOTIC ANALYSIS
 FORMALYSIS
 FORM As already noticed, the critical properties of the theory are determined by long wavelength fluctuations which come into play at the last stages of the t-evolution, i.e. in the limit $t \to \infty$. The full HRT-OZ equations, in the critical region and at large t's considerably simplify: remarkably, the formal structure of the asymptotic HRT-OZ equation is in fact identical to that obtained by a much simpler closure to the first equation of the hierarchy (4) based on the Random Phase Approximation (RPA), i.e. neglecting the core condition. In fact, according to the critical phenomena paradigm, short range correlations should not affect the physics at long wavelengths which dominates in the critical region. And this property is explicitly satisfied within HRT-OZ.

In the simplified RPA form, the direct correlation function reads

$$
c_t(r) = c_R(r) - \beta \left(w_t(r) + \lambda_t w(r) \right) \tag{18}
$$

for all r's. As usual, the parameter λ_t is determined by the compressibility sum rule (6). Appendix B contains the details of the asymptotic analysis of (4) under both closures (18) and (10). Here we simply discuss the physical contents of the result. It is convenient to first define a modified free energy density (in units of k_BT) A_t which encompasses both the ideal gas term (A^{id}) and the mean field contribution due to that part of the interaction not already included in A_t :

$$
\mathcal{A}_t = A_t + A^{id} - \beta \frac{\rho^2}{2} [\tilde{w}(0) - \tilde{w}_t(0)] \tag{19}
$$

The evolution equation (4) under the closure (18) for a three dimensional Yukawa fluid in the $t \to \infty$ limit and in the critical region becomes (see Appendix B):

$$
\frac{\partial \mathcal{A}_t}{\partial t} = e^{-3t} \frac{z^3}{4\pi} \left[1 + p^{-1/2} \frac{2 - p - \sqrt{p}\sqrt{1 + x}}{\sqrt{p + x + 2\sqrt{p}\sqrt{1 + x}}} \right]
$$
(20)

where

$$
x = -\frac{z^2}{4\pi\beta\psi_{\infty}}e^{2t}\frac{\partial^2 \mathcal{A}_t}{\partial \rho^2} \tag{21}
$$

is a renormalized inverse compressibility and the non universal parameter $p < 1$ is the ratio between the curvature of the potential and of the direct correlation function divided by ψ_{∞} (8), as shown in Eqs. (67,79). It is convenient to remove an analytic contribution and a regular pre-factor in Eq. (20) by a suitable rescaling of the free energy $\mathcal{A}_t \to \Psi_t$ and of the density $\rho \to \varphi$ as detailed in Eq. (69):

$$
\rho - \rho_c = \varphi \left(\frac{z^5}{(4\pi)^2 \beta \sqrt{p}}\right)^{1/2}.
$$
\n(22)

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The resulting evolution equation reads:

$$
\frac{\partial \Psi_t}{\partial t} = -e^{-3t} U(x) = -e^{-3t} \frac{2 - p - \sqrt{p}\sqrt{1 + x}}{\sqrt{p + x + 2\sqrt{p}\sqrt{1 + x}}}
$$
(23)

where (21) is now expressed as

$$
x = e^{2t} \frac{\partial^2 \Psi_t}{\partial \varphi^2} \tag{24}
$$

Notice that, according to the definition (21), the variable x may attain a finite limit (x^*) for $t \to \infty$ only if the compressibility diverges, meaning that the fully interacting system is either at the critical point or in the two phase region. The asymptotic Eq. (23) has been numerically integrated for an initial condition at $t = 0$ of the Landau-Ginzburg type, i.e. appropriate for a self-interacting scalar field theory:

$$
\Psi_0(\varphi) = \Psi_0(0) + r\varphi^2 + u\varphi^4 \tag{25}
$$

ppropriate for a self-interacting scalar field theory:
 $\Psi_0(\varphi) = \Psi_0(0) + r\varphi^2 + u\varphi^4$

ins information on the range of the potential, this effective model is defined by the mean field critical value ρ 0 is assumed t Besides p , which retains information on the range of the potential, this effective model is defined by two parameters: r is a measure of temperature (shifted by the mean field critical value) while $u > 0$ is assumed to be state independent. The typical behavior of the rescaled inverse compressibility $x(\varphi, r)$ during the "time" evolution determined by Eq. (23) is shown in Fig. 2 for $p = 0.5$, $u = 0.1$ and three states on the critical isochore $\varphi = 0$: above, below and very close to the critical temperature r_c . Above r_c the divergence of x shows that the inverse compressibility attains a finite, positive value, as expected in supercritical fluids. Close but still above the critical point, the evolution of x displays a plateau at $x^* < 0$ before diverging again. Instead, for $r < r_c$ the evolution crosses x^* and asymptotically sets to a finite, negative value $x₅$ implying that the inverse compressibility vanishes. Inside the coexistence curve the fluid is therefore characterized by flat isotherms, i.e. the Maxwell construction is recovered via the inclusion of long wavelength fluctuations. Such a behavior of the "time" evolution suggests the possible presence of two distinct fixed points of the evolution equation, x^* and $x<$, governing the critical and subcritical region respectively (while the "high temperature fixed point" is located at $x = +\infty$).

A. Critical behavior

The singular behavior of the thermodynamic properties in the critical region can be obtained from the asymptotic form of the evolution equation (23) following the fixed point analysis developed in the RG theory. As a first step we eliminate from Eq. (23) the explicit dependence on the evolution parameter t by a t-dependent rescaling of the density and free energy density:

$$
\zeta \equiv e^{\frac{t}{2}} \varphi
$$

\n
$$
\Gamma_t(\zeta) \equiv \left[\Psi_t(\varphi) - \Psi_t(0)\right] e^{3t}
$$
\n(26)

Notice that the rescaled inverse compressibility x (21) is simply expressed in terms of Γ_t by $x = \frac{\partial^2 \Gamma_t}{\partial \zeta^2}$. The evolution equation for the "renormalized" quantities now acquires a typical RG structure:

$$
\left[\frac{\partial}{\partial t} + \frac{1}{2}\zeta \frac{\partial}{\partial \zeta} - 3\right] \Gamma_t(\zeta) = U\left(\frac{\partial^2 \Gamma_t}{\partial \zeta^2}\right)_{\zeta=0} - U\left(\frac{\partial^2 \Gamma_t}{\partial \zeta^2}\right) \tag{27}
$$

where $U(x)$ is given by Eq. (23). Equation (27) admits a non trivial fixed point: i.e. a solution $\Gamma^*(\zeta)$ independent of t, regular and not identically zero on the whole real axis $\zeta \in (-\infty, \infty)$. A fixed point corresponds to a diverging compressibility at $\varphi = 0$, leading to $x^* = \frac{d^2\Gamma^*}{d\zeta^2}|_{\zeta=0} \sim -0.2848...$ (for $p = 0.5$), which perfectly agrees with the position of the plateau in the evolution shown in Fig. 2 for $r \sim r_c$, and therefore identifies the critical point. Linearizing the evolution equation (27) around $\Gamma^*(\zeta)$ we can investigate the stability properties of the fixed point: letting $\Gamma_t(\zeta) = \Gamma^*(\zeta) + \delta \Gamma_t(\zeta)$ and keeping terms up to first order in $\delta \Gamma_t$ we find solutions of the form $\delta \Gamma_t(\zeta) = e^{\Lambda t} \gamma(\zeta)$ where Λ is the eigenvalue and $\gamma(\zeta)$ is the corresponding eigenfunction. As usual in this context we find two unstable eigenmodes, characterized by positive eigenvalues [6] governing displacements from the critical point along the critical isotherm and isochore. The first is odd in ζ and can be expressed as $\gamma(\zeta) = \frac{d\Gamma^*}{d\zeta}$ with $\Lambda_o = 1/2$, while the other, even in ζ, must be computed numerically. The RG structure of the HRT-OZ evolution equation (27) shows that scaling

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and hyperscaling are correctly reproduced by the theory. Due to the known relation between eigenvalues and critical exponents [6]

$$
\delta = 1 + \frac{2}{\Lambda_o}
$$

$$
\gamma = \frac{2}{\Lambda_e}
$$
 (28)

is whose results do depend on the cut-off function
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 **For Allie Colling to depend on the cut-off function [20]. Note however,

exponents are corr** we find $\delta = 5$, as implied by scaling when the correlation exponent η vanishes, while the other critical exponents follow from the numerical determination of Λ_e and are shown in Table I for few choices of the parameter p: HRT-OZ displays non classical critical exponents weakly dependent on p. It is in fact apparent that, contrary to the sharp cut-off HRT-OZ approach, here the asymptotic equation (23) does not acquire a universal form but retains memory of the microscopic model through the value of the parameter p . Therefore the critical properties of the three dimensional fluid within our approximation will generally violate universality and both critical exponents and scaling laws will depend on the precise value of p. This situation closely resembles the case of non perturbative approximations to the RG flow equations, whose results do depend on the cut-off function [20]. Note however that, as shown in Ref. [12, 13], the critical exponents are correct to first order in a $\epsilon = 4 - d$ expansion of the HRT-OZ equation (4) and, within a systematic ϵ −expansion of the full HRT hierarchy, the dependence on p disappears and universality is indeed recovered. The critical exponents can be also evaluated by numerical integration of the asymptotic evolution equation (23) close to the critical point with no reference to any RG procedure. The results, shown in Fig. 3 for the case $p = 0.5$ and $u = 0.1$ conform to the expectation of the fixed point analysis previously outlined both above and below the critical temperature.

The renormalized equation (27) also allows to determine other relevant properties of the critical regime. In Table I we show the compressibility amplitude ratio defined as $U_2 = C_+/C_-$ where

$$
\chi(r,\varphi(r)) = C_{\pm} |\Delta r|^{-\gamma} \tag{29}
$$

 χ is the isothermal compressibility and $\Delta r = r - r_c$. The + sign corresponds to $r > r_c$ and $\varphi(r) = 0$, while the $-$ sign refers to $r < r_c$ and $\varphi(r)$ along the coexistence curve. Note that in the numerical solution of the evolution equation, the value of the inverse compressibility on the binodal, i.e on the point of discontinuity, is not directly available and must be obtained by extrapolation of the results at neighboring mesh points. This limits the possibility to get arbitrarily close to the critical point and to obtain accurate estimates of the amplitude ratio U_2 .

The scaling form of the equation of state of the model can be also obtained in the form:

$$
\chi^{-1} = |\varphi|^{\delta - 1} W\left(\frac{\Delta r}{|\varphi|^{1/\beta}}\right) \tag{30}
$$

The numerical results for $p = 0.5$ are shown in Fig. 4 together with a polynomial approximation of the "exact" scaling function from Ref. [21]. Other choices of p do not modify appreciably the good agreement between HRT-OZ and RG results.

B. Below the critical temperature

As previously noticed (see also Ref. [13]), within the smooth cut-off HRT-OZ scheme, the inverse compressibility develops a singularity for $t \to \infty$ at any $r < r_c$: it attains a finite limit on the binodal while vanishes identically inside the two phase region. The way χ^{-1} drops to zero below r_c is governed by a low temperature fixed point:

$$
x_{\leq} = \lim_{t \to \infty} e^{2t} \frac{\partial^2 \Psi}{\partial \varphi^2}
$$

defined by the divergence of $U(x)$ in Eq. (23) and explicitly given by $x₀ = p - 2\sqrt{p}$. The agreement between the formal analysis and the numerical results shown in Fig. 2 fully confirms this interpretation.

It is interesting to examine more closely the HRT-OZ evolution of the inverse compressibility below the critical temperature in order to understand how the inclusion of long wavelength fluctuations drives such a behavior, which perfectly matches the expectations for the case of a scalar order parameter. In Figs. 5a,5b we contrast the evolution of two typical isotherms above and below r_c by showing the inverse compressibility for four values of the switching parameter t, as obtained by the numerical integration of Eq. (23). At $t = 0$ the initial condition is a smooth function of the density which reproduces the Landau-Ginzburg mean field structure. Below the mean field critical temperature

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EVALUAT SURFACAL ALGONS The universal consideration of the equal of the boundary of the metastable phases inside the coexistence curve. "
In feature of the volution shown in Fig. 5b suggests a way to discrimina
distable $(r = 0)$ it displays a large unstable region, corresponding to the presence of a Van der Waals loop. In general, the inclusion of density fluctuations, governed by the parameter t , leads to a gradual increase of the inverse compressibility thereby reducing the width of the unstable region. Above the true critical temperature this eventually leads to the complete suppression of the instability (see Fig. 5a). Instead, below r_c , long wavelength fluctuations severely affect the shape of the isotherms: i) χ^{-1} , when negative, flattens as a function of density and approaches zero at large t; ii) The isotherm sharpens at the boundaries of the interval where χ^{-1} < 0 developing a discontinuity, which defines the binodal; *iii*) At large length-scales the coexistence region widens again and in a small density interval close to the binodal the compressibility becomes positive before dropping to zero. The origin of the flattening and sharpening of the HRT-OZ isotherm has been analyzed in detail in Ref. [13] where the smooth cut-off HRT-OZ has been studied for a pure Φ^4 field theory. As anticipated, the smooth cut-off prescription overcomes one of the main problems of the previous implementations of HRT-OZ [7], providing finite compressibility at coexistence. This is in agreement with the field-theoretical analysis of Ref. [22] showing that the coincidence of binodal and spinodal is a peculiarity of the sharp cut-off scheme, while finite compressibility is expected for every choice of the cut-off function (or "smooth cut-off regulator"). In this framework, we may investigate whether it is possible to locate the boundary of the metastable phases inside the coexistence curve. The spinodal curve is not rigorously defined within equilibrium statistical mechanics, the free energy being a convex function in the whole phase diagram, but a special feature of the evolution shown in Fig. 5b suggests a way to discriminate between instability and metastability. In a metastable state we expect that density fluctuations on a large but finite range do not drive the system towards phase separation, i.e. χ^{-1} remains positive until t does not exceed a (large) crossover value t_s . Only when density fluctuations on a macroscopic scale are included, does the first order transition take place. In contrast, in the truly unstable region χ^{-1} the inverse compressibility is negative throughout the evolution. The curves dividing the regions of positive and negative χ^{-1} in the (φ, t) plane at different temperatures are shown in Fig. 6, together with the extent of the metastable region obtained via the criterion previously outlined. According to this definition, the shape of the spinodal line in a φ vs. Δr diagram appears to be characterized by the same critical exponent β of the coexistence curve, as shown in Fig 3 . The value t^* of the evolution parameter t where the width of the unstable region has a minimum, diverges when the critical point is approached. This corresponds to the divergence of a characteristic length-scale $R_c \div e^{t^*}$, which may be identified as the critical droplet radius on the spinodal line, according to the standard droplet model picture of nucleation [23]. This divergence follows a power law consistent with the scaling $R_c \div \xi \div |\Delta r|^{-\nu}$ with $\nu = \gamma/2$ (see Table I).

V. NUMERICAL RESULTS AND PHASE DIAGRAMS

Here we show some results obtained by the numerical integration of the full smooth cut-off HRT-OZ equation as derived in Appendix A (62). We systematically investigated the HCYF at three values of the inverse range parameter $z: z = 1.8$, which mimics the usual Lennard-Jones potential, $z = 4$ and $z = 7$ which are more appropriate for modeling colloidal suspensions. The numerical solution of the partial differential equation has been performed by a full implicit predictor-corrector finite difference method on a density mesh of 2000 points. The estimated numerical error cannot be appreciated on the scale of the figures.

A. Spinodal and binodal

The coexistence curve can be immediately read-off the numerical solution of the equation, for in the two phase region the inverse compressibility vanishes. The spinodal curve instead is evaluated following the criterion discussed in Section (IV-B) and illustrated in Fig. 6. The results are shown in Figs. 7,8,9 together with the coexistence curves obtained by Monte Carlo simulations and SCOZA.

The HRT-OZ coexistence curve agrees very well with the Monte Carlo data for $z = 1.8$ and $z = 4$ while some deviation appears at $z = 7$ where the MSA closure, which underlies the present implementation of the smooth cut-off HRT-OZ, is less accurate. As expected, the coexistence curve becomes flatter by reducing the range of the attractive interaction, as testified by the different scale of the temperature axis in the three cases. Note that the SCOZA results are always close to the HRT-OZ binodal, with some deviation in the critical region where the HRT-OZ coexistence line is flatter. Again, the largest discrepancy between HRT-OZ and SCOZA occurs at high z , however, even in this case, the two critical temperatures differ by less than 1%.

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B. The liquid-solid phase boundary

The fluid-solid phase equilibrium of the HCYF has been widely studied, mainly because it allows one to describe the qualitative change in the shape of the phase diagram brought about by changing the attraction range. As is well known, as the range is narrowed, i.e., as z is increased, one goes from the situation where the phase diagram has both a stable fluid-fluid critical point and a fluid-fluid-solid triple point, to that where the critical point moves into the fluid-solid coexistence region, thereby making the fluid-fluid transition metastable with respect to freezing, and there is no triple point [25, 26]. The former case is found in atomic and molecular fluids, while the latter is characteristic of many colloidal systems, including protein solutions [27]. The disappearance of a stable fluid-fluid transition is in fact a general feature of short-ranged attractive interactions, but the Yukawa potential was, together with the Asakura-Oosawa one [28], among the first to be considered in this respect.

HRT is not a theory of freezing, but of course it can be used to predict the equilibrium fluid-solid phase diagram, if the free energy of the solid is provided by some different procedure. We have resorted to the simplest and most adopted one [25, 28–31], based on Barker and Henderson perturbation theory [32]. This approach was originally developed for the fluid phase, but it can be straightforwardly employed in the solid one as well. According to it, the Helmholtz free energy of a solid of particles interacting via a hard-sphere plus tail potential is given by:

$$
A_s = A_s^R - \frac{1}{2} \beta \rho^2 \int d\mathbf{r} \, g_s^R(r) \, w(r) + A_2 + \mathcal{O}(\beta^3) \,, \tag{31}
$$

where A_s is minus the excess free energy per unit volume of the solid divided by $k_B T$, A_s^R is the corresponding quantity for the reference hard-sphere solid, $g_s^R(r)$ is the radial distribution function of the hard-sphere solid averaged over the solid angle, and $w(r)$ is the potential tail, i.e., the attractive Yukawa in the present case. Following previous investigations [28, 31], Eq. (31) has been truncated at the second-order term A_2 , which has been estimated by the so-called "macroscopic-compressibility" approximation, also suggested by Barker and Henderson [32]:

$$
A_2 \simeq \frac{1}{4} \beta^2 \rho^2 \chi_s^R \int d\mathbf{r} \, g_s^R(r) \, w^2(r) \,, \tag{32}
$$

where χ_s^R is the isothermal compressibility of the hard-sphere solid divided by the ideal-gas value. Comparison with Monte Carlo simulations on depletion potentials [33] has shown that, although the estimate of A_2 given by Eq. (32) deviates quite substantially from its exact value, which would involve knowledge of the three- and four-particle distribution function of the reference solid, the sum of A_1 and A_2 given by Eqs. (31,32) is nevertheless remarkably close to the simulation results for the full free energy A_s , unless the interaction is very short-ranged. In fact, the truncated expansion (31) has proved to be more accurate in the solid than in the fluid [31], of course provided the solid in study has the same lattice structure as the reference one.

⁻³¹), based on Barker and Henderson perturbation theory [32]. This
id phase, but it can be straightforwardly employed in the solid one as
y of a solid of particles interacting via a hard-sphere plus tail potentia
 $A_s = A_s$ Equations (31,32) were already used together with the sharp cut-off version of HRT-OZ to study the fluid-solid equilibrium of non-additive hard-sphere mixtures, including the Asakura-Oosawa case [34]. They were also applied together with SCOZA to both the HCYF considered here [35] and a two-Yukawa fluid with competing attractive and repulsive interactions [36]. As in these studies, the Helmholtz free energy A_s^R has been obtained by integrating with respect to density the equation of state of the hard-sphere solid given by Hall [37], starting from a density $\rho_0 = 0.736 \rho_{cp}$, where $\rho_{cp} = \sqrt{2}\sigma^{-3}$ is the density at close packing, and setting the integration constant by the result $-A_s^R/\rho_0 = 5.91889$ determined via numerical simulation by Frenkel, also reported in [35]. The radial distribution function of the hard-sphere solid $g_s^R(r)$ has been described by the parametrizations developed by Kincaid and Weis [38] starting from Weis' Monte Carlo simulations [29] and by Choi, Ree, and Ree [39] on the basis of their own simulation results. These parametrizations share the same functional form, and differ only for the specific dependence of the parameters on the average density. In both cases, we used the data for the neighbor distance and coordination number of the face-centered cubic lattice reported in [40]. The integrals in Eqs. (31,32) were truncated at $r_c = 8\sigma$, and a long-range correction was added by setting $g_s^R(r) \equiv 1$ for $r > r_c$. Such a correction is actually irrelevant for the case discussed below.

The fluid-solid phase boundary has been determined by equating the pressure and chemical potential of the solid given by Eqs. (31,32) to those of the fluid as predicted by the smooth cut-off HRT-OZ. As expected, we found that, as the inverse range z is increased, the freezing line becomes wider, and for large enough z it lies above the fluid-fluid coexistence curve. Figure 10 shows the phase diagram for the case of marginal stability, such that the freezing line is tangent to the fluid-fluid coexistence curve at the critical point. This separates the stable fluid-fluid transition regime from the metastable one. We have reported the results obtained using both Kincaid and Weis [38] and Choi and Ree [39] radial distribution function $g_s^R(r)$. The two sets of results are nearly superimposed, save for some small deviations along the melting line. In both cases we obtain $z = 5.6$ as the value of marginal stability. This result is very close to that given by SCOZA, $z = 5.7$, via the same recipe used here to get Carnahan-Starling thermodynamics

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in the fluid phase, see Sec. III, and is similar to the value $z = 6$ determined by Hagen and Frenkel [25] via Gibbs Ensemble Monte Carlo simulations. In the case of SCOZA, using the hard-sphere direct correlation function $c_R(r)$ given by Waisman [41] instead of fixing its range a priori gave $z = 6.05$ [35], in closer agreement with the simulation result. It is likely that a similar trend would be shown by the smooth cut-off HRT-OZ with the same $c_R(r)$.

C. Equation of state and thermodynamics

The equation of state of the HCYF can be easily obtained by integrating the HRT-OZ differential equation. The pressure divided by k_BT is compared to Molecular Dynamics simulations in Figs. 11 and 12 for $z = 1.8$ and $z = 4$ respectively. The agreement is very good showing that HRT-OZ is able to provide a consistent picture of thermodynamics throughout the phase diagram of the fluid.

Finally, in Fig. 13 we plot the specific heat as a function of temperature at the critical density for the three cases we have investigated $z = 1.8$, $z = 4$ and $z = 7$. The expected λ -line can be clearly seen in the HRT-OZ data. Note the different growth of C_V at low temperature among the three cases.

D. Correlation functions

Although the Hierarchical Reference Theory has been especially devised for investigating the thermodynamical properties of fluids, it also allows for the study of correlation functions through the link between thermodynamics and correlations implied by the closure relation discussed in Section III (6,10). However, due to the MSA-like structure of the adopted scheme, we cannot expect an accuracy of the quality found for the thermodynamics. In Figs. 14-15 we compare the HRT-OZ results with Monte Carlo simulations for different values of z. The HRT-OZ result considerably underestimates the contact value of $g(r)$, especially at large z or at low density, as a consequence of the "linear" relationship between $c(r)$ and the potential outside the core radius.

As a side comment, we remark that the structure of correlations in the two phase region is known to be analytically related to the correlation functions of the two pure phases: liquid (L) and vapor (V). According to the theory of first order phase transitions [44], the density correlations at coexistence are linear combinations of the density correlations for the two pure phases, leading to the following analytical structure of the radial distribution function at coexistence:

$$
g(r) = \left(\frac{\rho_L}{\rho}\right)^2 \frac{\rho - \rho_V}{\rho_L - \rho_V} \left[g_L(r) - 1\right] + \left(\frac{\rho_V}{\rho}\right)^2 \frac{\rho_L - \rho}{\rho_L - \rho_V} \left[g_V(r) - 1\right] + \frac{\rho(\rho_L + \rho_V) - \rho_L \rho_V}{\rho^2}
$$
(33)

z = 1.0, z = 4 and z = 1. 1 me expected λ -men can be casary seen in
 $\sigma \leq T$ i.e., z = 4 and z = 1. me expected λ -men cases.
 D. Correlation functions

archical Reference Theory has been especially devised for in This shows that in the two phase region, the radial distribution function $q(r)$ tends at large distances to a non trivial limit, which differs from the expected uncorrelated result $q(r) \rightarrow 1$. Such a behavior is in fact consistent with the divergence of the isothermal compressibility in the two phase region, via the compressibility sum rule: $k_B T \rho \chi_{red} = 1 + \rho \int d\mathbf{r} (g(r) - 1)$. Remarkably, the smooth cut-off HRT-OZ is able to capture this peculiar structure of the two body correlation. In fact, it can be shown [15] that the asymptotic limit of $g(r)$ predicted by HRT-OZ precisely reproduces the last term of Eq. (33). However, further analysis shows that the closure adopted in this work is not able to describe the full structure of correlations in the two phase region expected on the basis of the exact result (33): a spurious long range decay in $g(r)$ appears inside the coexistence curve.

VI. CONCLUSIONS

In this work we have analyzed a simple Ornstein-Zernike closure to the Hierarchical Reference Theory of fluids in the smooth cut-off formulation for hard core Yukawa fluids. Among the main advantages of this approach we can recall a consistent representation of the critical point, as customary in the HRT scheme, including non classical critical exponents and scaling laws, together with a correct description of the first order phase boundary. Contrary to the usual sharp cut-off HRT-OZ, here the inverse compressibility shows the expected discontinuity on the binodal which also allows to define the critical exponent γ' describing the divergence of the isothermal compressibility when approaching the critical point from the low temperature side. The expected long distance limit of two body correlations in the two phase region is also correctly reproduced by HRT-OZ. An estimate of the spinodal line can be extracted by studying the evolution of the inverse compressibility when density fluctuations of larger and larger wavelength are included. The thermodynamics of the HCYF agree well with simulations both near and far from the critical point: coexistence curves and equations of state can be predicted with high accuracy by HRT-OZ.

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Despite such a remarkable performance of HRT-OZ, there is still room for significant improvement. Nowadays the most challenging applications of liquid state theory concern the so called complex fluids, often characterized by extremely short range attractive interactions and/or more structured potentials with both attractive and repulsive tails [8]. The application of HRT to this class of systems requires further work: the closure to the exact HRT hierarchy considered up to now is clearly not adequate to describe complex fluids, for it assumes a linear relation between the space dependence of the direct correlation function and the two body interaction. Moreover the rather artificial splitting between a short range "reference" part of the potential $v_R(r)$ and a long range "perturbation" $w(r)$, although customary in liquid state theory, limits the applicability of the method to potentials where repulsive and attractive contribution can be easily identified. The HRT-OZ description of correlations, already for moderately short range interactions, shown in Figs. 14-15, is not fully satisfactory and should be improved. In Section III we pointed out a natural way to generalize the adopted closure, which paves the way to more elaborate, flexible and general approximation schemes able to combine the virtues of the HRT approach and the accuracy of more sophisticated liquid state theories. We also mention a straightforward generalization of the smooth cut-off HRT method to binary mixtures, which is potentially very useful in the framework of complex fluids.

beneuary very seental une naturewors of comparison and, the matter of which is a critical exponential point of view, few problems still remain open, and should be a comparison to comparison the more control with its propo From a more fundamental point of view, few problems still remain open, and should be addressed in future investigations. The HRT-OZ critical exponents are always affected by a systematic error (which is of about 10% for γ) clearly related to the adopted Ornstein-Zernike structure of correlations which implies $\eta = 0$. This feature does not allow to apply the HRT-OZ approach in two dimensions. In order to go beyond this approximation, it is necessary to include, at least in some partial form, the effects of fluctuations on the momentum dependence of two body correlations: an investigation of approximation schemes to the second equation of the HRT hierarchy (5) should be considered in the future. Hopefully, such a generalization will also provide a fully consistent picture of correlations in the two phase region.

Finally, even within the present approximation scheme we did not address the subtle problem of the analytic structure of the free energy near the coexistence curve. According to the droplet model [23], later confirmed by an exact result for the Ising model [45], an essential singularity is present at coexistence: the nth derivative of the free energy with respect to the chemical potential should scale as $n!^{3/2}$ in three dimensions. Some preliminary numerical evidence suggests that an essential singularity is indeed present also within HRT-OZ but determining the precise scaling requires some further analytical and numerical effort.

We are grateful to Nigel Wilding, Dino Costa and Giuseppe Pellicane for providing simulation results of the radial distribution function of the HCYF.

Appendix A: The HRT-OZ equations and their numerical solution

Here we discuss the formal structure of the smooth cut-off HRT-OZ equations (4,6) specialized to the three dimensional $(d = 3)$ Yukawa fluid (11,12,13). We follow the notation of Ref. [46] (hereafter referred to as KJ) for the solution of the MSA equations. By substituting the form (11) into the evolution equation (4) we get:

$$
\frac{dA_t}{dt} = 2\pi\rho^2 \left[C_1 \hat{g}_t(s) + C_2 \partial_s \hat{g}_t(s) \right]_{s=z_2}
$$
\n(34)

Here $\hat{g}_t(s)$ is the Laplace transform of $r g_t(r)$:

$$
\hat{g}_t(s) = \int_0^\infty r g_t(r) e^{-sr} dr \tag{35}
$$

and the coefficients are defined as

$$
C_1 = \frac{e^z}{T} e^{-2t} (2\psi(t) - \dot{\psi}(t))
$$
\n(36)

$$
C_2 = \frac{z}{T} e^z e^{-3t} \psi(t) \tag{37}
$$

where dots represent derivatives with respect to the evolution parameter t. According to Eq. $(KJ-15)$

$$
\hat{g}_t(s) = \frac{s\tau(s)e^{-s}}{1 - 12\eta q(s)}\tag{38}
$$

where η is the packing fraction $\eta = \pi \rho/6$. The functions $\tau(s)$ and $q(s)$ are given by Eqs. (KJ-16,KJ-17,KJ-18) in terms of (z_1, z_2) and the parameters $(a, b, \beta_1, \beta_2, d_1, d_2)$ which satisfy the equations (KJ-10,KJ-11,KJ-12,KJ-13). This

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complicate set of six non-linear equations in six unknowns allows to find the MSA parameters as a function of the input variables: $(\rho, z_1, z_2, K_1, K_2)$. However, our specific problem is slightly different because K_1 is actually unknown, depending on the consistency parameter λ_t . Instead, we would like to express $\hat{g}_t(s)$ in terms of (ρ, z, z_2, K_2) and the compressibility which appears into Eq. (6). Luckily, the inverse compressibility is easily expressed in terms of the parameter a by

$$
\rho A_t'' = \rho \int d\mathbf{r} c_t(r) = 1 - a^2 \tag{39}
$$

where a prime represents differentiation with respect to ρ . Therefore, in order to integrate the HRT-OZ differential equation (34) we should solve, for each t, the five equations $(KJ-10,KJ-11,KJ-12)$ and the second of the pair of equations (KJ-13) obtaining the five parameters $(b, \beta_1, \beta_2, d_1, d_2)$ in terms of $(\rho, z_1, z_2, A''_t, K_2)$. Then, by substituting these expressions into Eq. (38) we can evaluate the right hand side of Eq. (34) which becomes a partial differential equation (PDE) for the free energy density $A_t(\rho)$. Unfortunately, an analytic expression for the parameters $(b, \beta_1, \beta_2, d_1, d_2)$ is not available and then we decided to adopt a different procedure.

As a first step it is convenient to change the parametrization of the MSA equations because, during the HRT evolution, the inverse range of the potential $z_2 = ze^{-t}$ becomes extremely small giving rise to singularities in the parameters $(a, b, \beta_1, \beta_2, d_1, d_2)$. We first introduce $\tilde{\beta}_i$ and Δ_i defined by:

$$
\tilde{\beta}_i = \beta_i z_i^{-2} \tag{40}
$$

$$
d_i z_i^3 = -12\eta + \Delta_i \tag{41}
$$

together with

$$
k_2 = K_2 z_2^{-4} \tag{42}
$$

which attains a finite limit for $t \to \infty$. Then we note that the equations depend on the particular combination

$$
\gamma = b - 12 \eta \sum_i \tilde{\beta}_i
$$

m we decided to adopt a different procedure.
 For Peer Review the Conservant Conservation of the MSA equations is convenient to change the parametrization of the MSA equations is range of the potential $z_2 = ze^{-t}$ become which will be used in place of b. The new set of parameters is then given by $(a, \gamma, \tilde{\beta}_1, \tilde{\beta}_2, \Delta_1, \Delta_2)$. Actually only the renormalization of $\tilde{\beta}_2$ and Δ_2 are necessary but, for formal convenience, we adopted the same definition also for $\tilde{\beta}_1$ and Δ_1 . The Laplace transform of the radial distribution function is written as

$$
\hat{g}(s) = \frac{1}{s^2} + \frac{N(s)}{D(s)}
$$
\n(43)

where the ideal gas term has been written explicitly. The key quantities $N(s)$ and $D(s)$ are expressed in terms of the parameters by the following set of definitions:

$$
\Sigma = \sum_{i} \left[-f_4(z_i) \left(-12\eta + \Delta_i \right) + f_1(z_i) \right] \tilde{\beta}_i \frac{z_i}{z_i + s} \tag{44}
$$

$$
\Sigma_0 = -\sum_i \tilde{\beta}_i \frac{z_i}{z_i + s} \tag{45}
$$

$$
\Gamma = \sum_{i} \tilde{\beta}_i \Delta_i \frac{1}{z_i + s} \tag{46}
$$

$$
\tau(s) = a + (a + \gamma)s + s^2 \Gamma - 12\eta s \Sigma_0 \tag{47}
$$

$$
q(s) = f_5(-s)\,\tau(s) \tag{48}
$$

$$
P = \Sigma - \frac{a + \gamma}{24} - \frac{s}{24} \Gamma + \frac{1}{2} \eta \Sigma_0 + q(s)
$$
\n(49)

$$
N(s) = \tau(s)f_2(-s) + \Gamma - a - \gamma - s\Gamma + 12\eta\Sigma_0 + 12\eta P \tag{50}
$$

$$
D(s) = a + \gamma s - 12\eta (s^2 P + s \Sigma_0)
$$
\n(51)

where we introduced the notation $f_n(x)$ for:

$$
f_n(x) \equiv x^{-n} \left[e^x - \sum_{j=0}^{n-1} \frac{x^j}{j!} \right]
$$

which attains a finite limit as $x \to 0$. The two equations KJ-12 and the single equation KJ-13 now become, respectively:

$$
z_1^3 - \Delta_1 = 12\eta z_1^2 \frac{N(z_1)}{D(z_1)}
$$
\n(52)

$$
z_2^3 - \Delta_2 = 12\eta z_2^2 \frac{N(z_2)}{D(z_2)}\tag{53}
$$

$$
k_2 z_2 = \tilde{\beta}_2 D(z_2) \tag{54}
$$

while the first two equations KJ-10,KJ-11 are:

$$
\gamma = 12\eta \left[-\frac{a}{8} - \frac{\gamma}{6} - \sum_{i} \tilde{\beta}_i \Delta_i f_3(z_i) + 12\eta \sum_{i} \tilde{\beta}_i z_i f_4(z_i) + \sum_{i} \tilde{\beta}_i z_i f_1(z_i) \right]
$$
(55)

$$
1 - a = 12\eta \left[-\frac{a}{3} - \frac{\gamma}{2} - \sum_{i} \tilde{\beta}_{i} \Delta_{i} f_{2}(z_{i}) + 12\eta \sum_{i} \tilde{\beta}_{i} z_{i} f_{3}(z_{i}) + \sum_{i} \tilde{\beta}_{i} z_{i} e^{z_{i}} \right]
$$
(56)

By writing explicitly the ideal gas contribution to the radial distribution function (43) the evolution equation (34) acquires an additional term which precisely generates the mean field part of the free energy. This term can be absorbed into the definition of a modified free energy density A_t :

$$
-a = 12\eta \left[-\frac{a}{3} - \frac{7}{2} - \sum_{i} \tilde{\beta}_{i} \Delta_{i} f_{2}(z_{i}) + 12\eta \sum_{i} \tilde{\beta}_{i} z_{i} f_{3}(z_{i}) + \sum_{i} \tilde{\beta}_{i} z_{i} e^{z_{i}} \right]
$$
(56)
the ideal gas contribution to the radial distribution function (43) the evolution equation (34)
at rem which precisely generates the mean field part of the free energy. This term can be absorbed
a modified free energy density A_{i} .

$$
A_{t} = A_{t} + A_{id} + \frac{\rho^{2}}{2} \left[\tilde{\phi}(0) - \tilde{\phi}_{t}(0) \right]
$$

$$
= A_{t} + A_{id} + \frac{2\pi e^{z}}{Tz^{2}} \rho^{2} \psi(t)
$$
(57)
the Fourier transform of $\phi(r) = -\beta w(r)$. Therefore, the evolution equation of the modified free

$$
\frac{\partial A_{t}}{\partial t} = 2\pi \rho^{2} \left[C_{1} \frac{N(s)}{D(s)} + C_{2} \frac{D(s)\partial_{s}N(s) - N(s)\partial_{s}D(s)}{D(s)^{2}} \right]_{s=z_{2}}
$$

ew variable

$$
\frac{\partial^{2} \frac{\partial^{2}}{\partial t}}{\partial t^{2}} e^{2t} \left[(2\psi(t) - \dot{\psi}(t)) z_{2}^{2} \frac{N(s)}{D(s)} + \psi(t) z_{2}^{3} \frac{D(s)\partial_{s}N(s) - N(s)\partial_{s}D(s)}{D(s)^{2}} - \dot{\psi}(t) \right]_{s=z_{2}}
$$
(58)
idies with the right hand side of the evolution equation (34), which in fact simply becomes

$$
\frac{\partial A_{t}}{\partial t} = z_{2}^{2} \left[u_{t} + \frac{2\pi e^{z}}{Tz^{4}} \rho^{2} e^{2t} \dot{\psi}(t) \right]
$$
(59)
apressibility equation (39) is expressed in terms of the modified quantities as:

$$
-\rho A_{t}^{''} = a^{2} - \rho \frac{4\pi e^{z}}{Tz^{4}} z_{2}^{2} e^{2t} \psi(t)
$$
(60)
and (60) we finally get the evolution equation for the bare dimensionless inverse compressibility

$$
2a\dot{a} = -\rho z_{
$$

where $\phi(k)$ denotes the Fourier transform of $\phi(r) = -\beta w(r)$. Therefore, the evolution equation of the modified free energy density is

$$
\frac{\partial \mathcal{A}_t}{\partial t} = 2\pi \rho^2 \left[C_1 \frac{N(s)}{D(s)} + C_2 \frac{D(s)\partial_s N(s) - N(s)\partial_s D(s)}{D(s)^2} \right]_{s=z_2}
$$

Now we define the new variable

$$
u_t \equiv 2\pi \rho^2 \frac{e^z}{Tz^4} e^{2t} \left[\left(2\psi(t) - \dot{\psi}(t) \right) z_2^2 \frac{N(s)}{D(s)} + \psi(t) z_2^3 \frac{D(s)\partial_s N(s) - N(s)\partial_s D(s)}{D(s)^2} - \dot{\psi}(t) \right]_{s=z_2}
$$
(58)

which basically coincides with the right hand side of the evolution equation (34) , which in fact simply becomes

$$
\frac{\partial \mathcal{A}_t}{\partial t} = z_2^2 \left[u_t + \frac{2\pi e^z}{T z^4} \rho^2 e^{2t} \dot{\psi}(t) \right] \tag{59}
$$

Analogously, the compressibility equation (39) is expressed in terms of the modified quantities as:

∂A^t

$$
-\rho \mathcal{A}_t'' = a^2 - \rho \frac{4\pi e^z}{T z^4} z_2^2 e^{2t} \psi(t)
$$
\n(60)

Combining Eqs. (59) and (60) we finally get the evolution equation for the bare dimensionless inverse compressibility a^2 :

$$
2a\dot{a} = -\rho z_2^2 u_t^{\prime\prime} \tag{61}
$$

In summary, we start from the definition (58). We differentiate with respect to t, which is also contained in z_2 and in the six parameters $(\gamma, \tilde{\beta}_i, \Delta_i, a)$. The derivative of the first five parameters are calculated by differentiating the five equations (52-56) and solving the linear problem in terms of \dot{a} . Finally, the derivative \dot{a} is expressed in terms of u''_t via Eq. (61) . This procedure provides a closed equation for u_t which must be coupled to the five algebraic non-linear equations (52-56) and to the definition (58) in order to formally express the six parameters in terms of u_t . An analytic solution of the set of equations (52-56,58) is clearly impossible, but we can implement a Raphson-Newton iterative scheme at each step in t, which in fact converges very quickly. The resulting evolution equation for the independent variable $u_t(\rho)$ has the general structure

$$
\frac{\partial u_t}{\partial t} = g_1(u_t, t) + g_2(u_t, t) \frac{\partial^2 u_t}{\partial \rho^2}
$$
\n(62)

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which is a quasi-linear PDE in two dimensions and it is stable provided $g_2 \geq 0$. This condition must be checked during the numerical integration and depends on the specific definition of the cut-off function $\psi(t)$. In this work we have considered two different choices for the function $\hat{\psi}(t)$: In the first part of the integration $(0 < t < t^*)$ we used

$$
\psi(t) = (1 + t/t_0)^{-2} \tag{63}
$$

where t_0 is a parameter. In the second part $(t > t^*)$ we have chosen

$$
\psi(t) = (\cosh t)^{-2} \tag{64}
$$

The transition point t^* is obviously defined by the equation $1 + t^*/t_0 = \cosh t^*$.

as appreciable deviations from the Verte-Weis form $|47|$. A better channel weight constant and the verte-time of this additional Yukawa equal to z, i.e. the inverse range of this additional Yukawa equal to z, i.e. the in The initial condition at $t = 0$ corresponds to a vanishing attractive tail $w_t(r)$ and can be obtained by substituting the analytic solution of the MSA equation for a hard sphere liquid, equivalent to the Percus-Yevick (PY) solution, into the definition of u_t (58). This form of the reference correlations is however rather inaccurate at high density, where the PY solution shows appreciable deviations from the Verlet-Weis form [47]. A better choice is to parametrize the hard sphere direct correlation function of the hard sphere liquid outside the core as a Yukawa tail. For convenience we decided to fix the inverse range of this additional Yukawa equal to z , i.e. the inverse range of the physical attractive part of the potential which defines the original problem. The amplitude of this additional contribution to the direct correlation function is fixed by requiring that the spatial integral of $c(r)$ is consistent with the Carnahan-Starling equation of state [16] via the compressibility sum rule (39). This choice is in fact equivalent to setting the initial value of the parameter λ_t to a density dependent non vanishing limit. The numerical solution of the PDE also requires two boundary conditions at the minimum and maximum density: at $\rho = 0$ u_t vanishes due to the definition (58) while at $\rho = \rho_{max} \sim 1$ the MSA approximation, corresponding to $\lambda_t = 0$, provides a reasonable approximation. The PDE has been solved by a finite difference fully implicit predictor-corrector algorithm [48] with stability and accuracy checks during the numerical integration. The parameter t_0 in the cut-off function has been chosen of order $1/z$ in order to preserve stability at all steps.

Appendix B: The asymptotic form of the HRT-OZ equations for a Yukawa potential

Let us first consider the simpler RPA based closure (18) of the HRT-OZ evolution equation. By substituting Eq. (18) into (4) and by rescaling the momentum variable $q \equiv ke^t$, we see that for $t \to \infty$ the direct correlation function $\tilde{c}_t(k)$ is evaluated in the $k \to 0$ limit and can be therefore expanded:

$$
\tilde{c}_t(k) \sim \frac{\partial^2 A_t}{\partial \rho^2} - b k^2 - \psi_\infty e^{-2t} \tilde{\phi}(ke^t)
$$
\n(65)

where b attains a finite limit in the $t \to \infty$ also at the critical point and is usually close to the range squared of the Yukawa tail while ψ_{∞} is given by Eq. (8). Notice that even in the $k \to 0$ limit, the direct correlation function does depend on the precise form of the two body interaction $\dot{\phi}(q)$ which, in our case is just the three dimensional Fourier transform of the Yukawa potential divided by $k_B T: \tilde{\phi}(q) = 4\pi\beta/(z^2 + q^2)$. After some algebra, the evolution equation for the modified free energy (19) acquires the following form in three dimensions $(d = 3)$:

$$
\frac{\partial \mathcal{A}_t}{\partial t} = \frac{e^{-3t}}{2} \int \frac{d^3q}{(2\pi)^d} \frac{2\tilde{\phi}(q) - \tilde{\phi}'(q)}{\left(-e^{2t} \frac{\partial^2 \mathcal{A}_t}{\partial \rho^2} + bq^2\right) \psi_{\infty}^{-1} + \tilde{\phi}(q)}\tag{66}
$$

Let us introduce the parameter

$$
p = \frac{bz^4}{4\pi\beta\psi_{\infty}}\tag{67}
$$

measuring the ratio between the curvature of the potential and of the direct correlation function (times ψ_{∞}). Then we rescale the inverse compressibility via Eq. (21) leading to

$$
\frac{\partial \mathcal{A}_t}{\partial t} = e^{-3t} \frac{z^3}{2\pi^2} \int_0^\infty dy \, \frac{y^2 (1+2y^2)}{[1+y^2][1+(1+y^2)(x+py^2)]} \tag{68}
$$

The integral is well defined provided the denominator does not vanish in the integration domain. Taking into account that $p > 0$ we must have $x + 1 > 0$ and either $p + x > 0$ or $p + x < 0$ and $x > p - 2\sqrt{p}$. The latter condition may be

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satisfied only for $p < 1$. This integral is performed analytically by contour integration with the result (20) , valid for all p and x satisfying the above conditions.

It is now natural to rescale the physical quantities (free energy and density) in order to get rid of the prefactors in Eq. (20):

$$
\mathcal{A}_t(\rho) = \frac{z^3}{4\pi} \left[-p^{-1/2} \Psi_t(\varphi) + \frac{1 - e^{-3t}}{3} \right]
$$

$$
\rho - \rho_c = \varphi \left(\frac{z^5}{(4\pi)^2 \beta \sqrt{p}} \right)^{1/2} \tag{69}
$$

In terms of the new variables Ψ_t and φ we simply get

$$
x = e^{2t} \frac{\partial^2 \Psi_t}{\partial \varphi^2}
$$

$$
\frac{\partial \Psi_t}{\partial t} = -e^{-3t} U(x)
$$
 (70)

where

$$
U(x) \equiv \frac{2 - p - \sqrt{p}\sqrt{1 + x}}{\sqrt{p + x + 2\sqrt{p}\sqrt{1 + x}}}
$$
\n(71)

which coincides with Eq. (23). Differentiating the latter definition with respect to t, implicitly contained in $x(t, \varphi)$, we have

$$
\frac{\partial U}{\partial t} = \frac{dU}{dx} \left[2x - e^{-t} \frac{\partial^2 U}{\partial \varphi^2} \right]
$$
\n(72)

The equation is now written in quasi-linear form. We just have to formally obtain the coefficients by inverting the definition (71) and evaluating dU/dx .

By defining the auxiliary quantity $y = \sqrt{p(1 + x)}$ all the ingredients appearing in (72) are given by:

$$
x = \frac{y^2}{p} - 1 \tag{73}
$$

$$
y = \frac{U^2(1-p) + (2-p)^2}{Up^{-1/2}\sqrt{U^2 - p + 4} + U^2 - p + 2}
$$
\n(74)

$$
\frac{dU}{dx} = -\left[1 + \frac{p}{2y}\right]D^{-3} \tag{75}
$$

$$
D = \frac{4-p}{2U + \sqrt{p(U^2 - p + 4)}}\tag{76}
$$

 $\frac{\partial \Psi_t}{\partial t} = -e^{-3t} U(x)$
 $U(x) = \frac{2-p-\sqrt{p}\sqrt{1+x}}{\sqrt{p+x+2\sqrt{p}\sqrt{1+x}}}$
Eq. (23). Differentiating the latter definition with respect to *t*, implies $\frac{\partial U}{\partial t} = \frac{dU}{dx} \left[2x - e^{-t}\frac{\partial^2 U}{\partial \varphi^2}\right]$
written in quasi-linear form. W The asymptotic analysis of the evolution equation with the full MSA-based closure (10), which includes the core condition, is more involved. By inspection of the equations for the MSA parameters (52-56), it is natural to guess that γ , $\tilde{\beta}_1$, $\tilde{\beta}_2$, Δ_1 , Δ_2 are all finite in that limit Actually, outside the coexistence curve, $\Delta_2 \rightarrow 0$, as shown by the numerical solution of the equations. In particular, in the critical region, we may assume that $\Delta_2 \to 0$ and analyze the equations in the limit $z_2 \to 0$, while a/z_2 and Δ_2/z_2 are finite. To leading order in a and z_2 , the equations KJ-10, KJ-11 and the first of the KJ-12 just depend on $\tilde{\beta}_1$, Δ_1 and $\gamma_0 = \gamma + 12\eta\tilde{\beta}_1$. Therefore, the two equations (53,54) provide the limiting values of $\tilde{\beta}_2$ and Δ_2/z_2 :

$$
\tilde{\beta}_2 = (12\eta)^{-1} \left[-(a/z_2 + \gamma_0) + \sqrt{(a/z_2 + \gamma_0)^2 + 24\eta k_2} \right]
$$
\n(77)

$$
\Delta_2 = -12\eta \frac{\tilde{\beta}_1 \Delta_1/z_1 - \gamma_0 (1 + \eta/2) + \tilde{\beta}_1 (12\eta - \Delta_1) f_4(z_1) + \tilde{\beta}_1 f_1(z_1)}{\sqrt{(a/z_2 + \gamma_0)^2 + 24\eta k_2}} z_2 \tag{78}
$$

where k_2 has been defined in (42). When these expressions are substituted into the definition of u_t (58) the singular contribution to u_t becomes

$$
u_t^{sing} \propto z_2 \frac{\tilde{\beta}_2 \gamma_0 - 4k_2}{\sqrt{(a/z_2 + \gamma_0)^2 + 24\eta k_2}}
$$

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besides an additive constant. Precisely at the critical point we expect that $a \sim z_2$, while inside the coexistence curve the ratio a/z_2 will tend to the value defined by the vanishing of the denominator: $(a/z_2 + \gamma_0)^2 + 24\eta k_2 = 0$ (recall that $k_2 < 0$) leading to a divergence of u_t^{sing}/z_2 as $t \to \infty$ and to a finite limiting value of Δ_2 . By comparing this expression with that obtained for a pure RPA (71) , it is possible to identify the effective RPA parameters p and x as:

$$
p = \frac{\gamma_0^2}{24\eta|k_2|} \tag{79}
$$

$$
1 + x = \frac{a^2}{24\eta |k_2| z_2^2}
$$
\n(80)

The requirement $0 < p < 1$ then gives $0 < \gamma_0^2 < 24\eta |k_2|$ which should be verified in the region where phase transitions take place.

- [1] See for instance J. Zinn-Justin Quantum Field Theory and Critical Phenomena Oxford University Press, Oxford (2002).
- [2] M. E. Fisher in *Critical Phenomena*, Lecture Notes in Physics, Vol. 186 edited by F.J.W. Hahne, Springer, Berlin (1982). [3] J. Hubbard and P. Schofield, Phys. Lett. A, 40, 245 (1972).
- [4] A. Parola and L. Reatto, Phys. Rev. Lett. 53, 2417 (1984); Phys.Rev. A 31, 3309 (1985).
- [5] M. Tau, A. Parola, D. Pini, and L. Reatto Phys. Rev. E 52, 2644 (1995); D. Pini, M. Tau, A. Parola, and L. Reatto, Phys. Rev. E 67, 046116 (2003).
- [6] A. Parola and L. Reatto, Adv. Phys. 44, 211 (1995).
- [7] A. Parola, D.Pini, and L. Reatto, Phys. Rev. E 48, 3321 (1993).
- [8] See for instance J. L. Barrat and J. P. Hansen Basic Concepts for Simple and Complex Fluids, Cambridge University Press, Cambridge (2003).
- [9] P. J. Lu, E. Zaccarelli, F. Ciulla, A. B. Schofield, F. Sciortino, and D. A. Weitz, Nature, 453, 499 (2008).
- [10] P. C. Hemmer, J. Math. Phys. 5, 75 (1964); J. L. Lebowitz et al. J. Math. Phys. 6, 1282 (1965).
- [11] K. G. Wilson and J. B. Kogut, Phys. Rep. C 12, 75 (1974).
- [12] A. Parola, J. Phys. C **26**, 5071 (1986).
- [13] C. D. Ionescu, A. Parola, D. Pini, and L. Reatto, Phys. Rev. E 76, 031113 (2007).
- [14] J. M. Caillol, Mol. Phys. 104, 1931 (2006).
- [15] A. Parola, D. Pini, and L. Reatto, Phys. Rev. Lett. 100, 165704 (2008).
- [16] J. P. Hansen and I. R. McDonald, *Theory of Simple Liquids*, $3rd$ ed., Academic Press, London (2006).
- [17] J. S. Høye, G. Stell, and E. Waisman, Mol. Phys. **32**, 209 (1976).
- [18] D. Pini, G. Stell, and N. B. Wilding, Mol. Phys. **95**, 483 (1998).
- [19] J. S. Høye, A. Reiner, Phys. Rev. E 75, 041113 (2007).
- [20] J. Berges, N. Tetradis, and C. Wetterich, Phys. Rep. **363**, 223 (2002).
- [21] A. Pelissetto and E. Vicari, Phys. Rep. 368, 549 (2002).
- $[22]$ A. Bonanno and G. Lacagnina, Nuclear Phys. B 693, 36 (2004) .
- [23] M. E. Fisher, Physics (N.Y.) 3, 255 (1967); W. Klein, Phys. Rev. B 21, 5254 (1980).
- $[24]$ Y. Duda, A. Romero-Martínez, and P. Orea, J. Chem. Phys. 126 , 224510 (2007) .
- [25] M. H. J. Hagen and D. Frenkel, J. Chem. Phys. **101**, 4093 (1994).
- [26] E. Lomba and N. G. Almarza, J. Chem. Phys. 100, 8367 (1994).
- [27] D. Rosenbaum, P. C. Zamora, and C. F. Zukoski, Phys. Rev. Lett 76, 150 (1996); N. Asherie, A. Lomakin, and G. B. Benedek, Phys. Rev. Lett. 77, 4832 (1996).
- Zinn-Justin Quantum Field Theory and Critical Phenomena Oxford Universide Rehenomena, Lecture Notes in Physics, Vol. 186 eilticed Phenomena Oxford Universidento, Phys. Rev. Lett. 54, 2417 (1984); Phys.Rev. A 31, 3309 (198 [28] A. P. Gast, W. B. Russell, and C. K. Hall, J. Colloid Interface Sci. 96, 1977 (1983); A. P. Gast, W. B. Russell, and C. K. Hall, J. Colloid Interface Sci. 109, 161 (1986).
- [29] J. J. Weis, Mol. Phys. **28**, 187 (1974).
- [30] C. E. Hecht and J. Lind, J. Chem. Phys. 64, 641 (1976); J. M. Kincaid, G. Stell, and E. Goldmark, J. Chem. Phys. 65, 2172 (1976).
- [31] M. Dijkstra, J. M. Brader, and R. Evans, J. Phys.: Condens. Matter 11, 10079 (1999).
- [32] J. A. Barker and D. Henderson, J. Chem. Phys. 47, 2856 (1967).
- [33] B. Rotenberg, J. Dzubiella, J. P. Hansen, and A. A. Louis, Mol. Phys. 102, 1 (2004).
- [34] F. Lo Verso, D. Pini, and L. Reatto, J. Phys.: Condensed Matter 17, 771 (2005).
- [35] G. Foffi, G. D. McCullagh, A. Lawlor, E. Zaccarelli, K. A. Dawson, F. Sciortino, P. Tartaglia, D. Pini, and G. Stell, Phys. Rev. E 65, 031407 (2002).
- [36] D. Pini, A. Parola, and L. Reatto, J. Phys.: Condensed Matter 18, S2305 (2006).
- [37] K. R. Hall, J. Chem. Phys. 57, 2252 (1972).
- [38] J. M. Kincaid and J. J. Weis, Mol. Phys. **34**, 931 (1977).
- [39] Y. Choi, T. Ree, and F. H. Ree, J. Chem. Phys. 95, 7548 (1991).
- [40] J. O. Hirschfelder, C. F. Curtis, and R. B. Bird, The Molecular Theory of Gases and Liquids, John Wiley & Sons, New York, 1954.

- [41] E. Waisman, Mol. Phys. **25**, 45 (1973).
- [42] E. Garnett, L. Mier-Y-Terán, and F. Del Río et al., Mol. Phys. 97, 597 (1999).
- [43] D. Costa and G. Pellicane, unpublished.
- [44] M. E. Fisher, Rep. Progr. Phys. 30, 615 (1967).
- [45] S. N. Isakov, Commun. Math. Phys. 95, 427 (1984).
- [46] J. Konior and C. Jedrzejek, Mol. Phys. 48, 219, (1983).
- [47] L. Verlet and J. J. Weis Phys. Rev. A 5, 939 (1972).
- [48] F. W. Ames, Numerical Methods for Partial Differential Equations, Academic, New York, (1977).

FIG. 1: Dimensionless inverse compressibility $S_t(0)^{-1}$ of a $z = 1.8$ Yukawa fluid within MSA below the critical temperature $(T = 1)$. The potential is given by (2) with $\psi(t) = (\cosh t)^{-2}$ (see Eq. 64) at $t = 2$ (dotted line) and $t = 10$ (full line). Vertical dashed lines bracket the coexistence region: here the $t = 10$ isotherm is smaller than 10^{-7} . The standard MSA result for a single Yukawa coincides with the full line outside coexistence while no solution exists inside the coexistence region.

FIG. 2: Evolution of the scaled inverse compressibility (24) on the critical isochore $\varphi = 0$ for $p = 0.5$, $u = 0.1$ and three temperatures: above $(r = -0.30810)$, below $(r = -0.30811)$ and very close to the critical temperature $(r = -0.3081037)$. obtained by numerical integration of Eq. (23). The dotted line is the critical fixed point $(x^* = -0.2848...)$ obtained by standard RG analysis of Eq. (27). The dashed line is the low temperature fixed point $x₀ = p - 2\sqrt{p} = -0.9142...$ (see Ref. [13]).

FIG. 3: Left panel: Log-Log plot of the inverse compressibility vs. reduced temperature for $p = 0.5$ and $u = 0.1$. Upper points are the HRT-OZ results below the critical temperature in the homogeneous phase, extrapolated to the coexistence curve. Lower points are on the critical isochore $\varphi = 0$ above r_c . Right panel: Log-Log plot of the coexistence curve (squares) and of the spinodal (triangles) for $p = 0.5$ and $u = 0.1$. Lines mark the asymptotic power law behavior with the critical exponents quoted in Table I: $\gamma = 1.335$ and $\beta = 0.334$.

FIG. 4: Scaling form of the equation of state $W(x)$ as defined in Eq. (30) for $p = 0.5$. Symbols: HRT-OZ results obtained by numerical integration of Eq. (23). Full line: polynomial approximant of the exact scaling function [21]. The metric factors fixing the units of $r - r_c$ and φ are determined so as to locate the coexistence curve at $\Delta r = -|\varphi|^{1/\beta}$ and $\chi^{-1} = |\varphi|^{\delta-1}$ at r_c .

FIG. 5: Snapshots of the inverse compressibility as a function of density at four values of the evolution parameter. From bottom to top: $t = 0, t = 0.5, t = 1.75$ and $t \to \infty$. Panel a refers to $r = -0.27$ above the critical temperature r_c , panel b to $r = -0.34$ below r_c .

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FIG. 6: Boundaries of the region defined by a negative compressibility as a function of the evolution parameter t for $p = 0.5$, $u = 0.1$. Curve a refers to $r = -0.27 > r_c$, curve b corresponds to $r = -0.34 < r_c$ and curve c to $r = -0.308105$ \lesssim_c . Full dots mark the coexistence region, empty dots the spinodal. The dashed lines are guides to the eye. The width of the unstable region has a minimum at the near-critical temperature at $t = 8.5$ and $\varphi = 0.0024$.

FIG. 7: Coexistence curve and spinodal of the HCYF for $z = 1.8$: HRT-OZ (full circles), SCOZA (line) and Monte Carlo [18] (error bars). The HRT-OZ spinodal is also shown (crosses).

FIG. 8: Coexistence curve and spinodal of the HCYF for $z = 4$. Notation as in Fig. 7. The Monte Carlo simulations are taken from Ref. [24].

FIG. 9: Coexistence curve and spinodal of the HCYF for $z = 7$. Notation as in Fig. 7. The Monte Carlo simulations are taken from Ref. [24].

FIG. 10: Fluid-solid phase diagram of the HCYF for $z = 5.6$. The fluid and solid phases have been described by HRT and perturbation theory respectively (see Eqs. (31,32)). Full circles: HRT fluid-fluid coexistence curve. The asterisk marks the position of the fluid-fluid critical point. Squares: freezing and melting lines obtained using the hard-sphere solid radial distribution function $g_s^R(r)$ by Kincaid and Weis [38]. Crosses: same as above using the $g_s^R(r)$ by Choi, Ree, and Ree [39]. Lines are a guide to the eye.

FIG. 11: Pressure divided by k_BT as a function of density for $z = 1.8$ at three temperatures: $T = 1, T = 1.5$ and $T = 2$ (from bottom to top). Line: HRT-OZ results. Full points: Molecular Dynamics data form Ref. [42]. The two open dots mark the boundaries of the HRT-OZ coexistence curve at the lowest temperature, characterized by a flat portion of the isotherm.

FIG. 12: Pressure divided by k_BT as a function of density for $z = 4$ at four temperatures: $T = 0.7$, $T = 1$, $T = 1.5$ and $T = 2$ (from bottom to top). Line: HRT-OZ results. Full points: Molecular Dynamics data form Ref. [42].

FIG. 13: Constant volume specific heat as a function of temperature at the critical density. From left to right $z = 1.8$, $z = 4$ and $z = 7$. The critical temperature is identified by a vertical dashed line.

FIG. 14: Radial distribution function of the HCYF for $z = 1.8$ at $T = 1.5$ and $\rho = 0.4$. Full line: HRT-OZ results. Points: Monte Carlo simulation from [18].

FIG. 15: Radial distribution function of the HCYF for $z = 4$ at $T = 0.5$ and $\rho = 0.816$. Full line: HRT-OZ results. Points: Monte Carlo simulation from [43].

Tables

TABLE I: HRT-OZ estimates of the critical exponents and compressibility amplitude ratio in three dimensions for few values of the parameter $p < 1$ compared to the exact values [21] obtained by extrapolation of high-temperature series expansions.