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Phase behaviour of polydisperse sticky hard spheres: analytical solutions and perturbation theory

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We discuss the phase coexistence of polydisperse colloidal suspensions in the presence of adhesion forces. The combined effect of polydispersity and Baxter’s sticky-hard-sphere (SHS) potential, describing hard spheres interacting via strong and very short-ranged attractive forces, give rise, within the Percus–Yevick (PY) approximation, to a system of coupled quadratic equations which, in general, cannot be solved either analytically or numerically. We review and compare two recent alternative proposals that have attempted to by-pass this difficulty. In the first, truncating the density expansion of the direct correlation functions, we have considered approximations simpler than the PY one. These \( C_n \) approximations can be systematically improved. We have been able to provide a complete analytical description of polydisperse SHS fluids using the simplest two orders \( C_0 \) and \( C_1 \). Such a simplification comes at the price of a lower accuracy in the phase diagram, but has the advantage of providing an analytical description of various new phenomena associated with the onset of polydispersity in phase equilibria (e.g., fractionation). The second approach is based on a perturbative expansion of the polydisperse PY solution around its monodisperse counterpart. This approach provides a sound approximation to the real phase behaviour, at the cost of considering only weak polydispersity. Although a final determination of the soundness of the latter method would require numerical simulations for the polydisperse Baxter model, we argue that this approach is expected to correctly take into account the effects of polydispersity, at least qualitatively.

1. Introduction

New technological advances in the physico-chemical manipulation of colloidal mixtures have re-ignited the issue of gaining a theoretical understanding of the phase behaviour of polydisperse systems [1]. ‘Polydispersity’ in colloidal solutions means that, due to the production process, suspended macroparticles with the same chemical composition cannot be exactly identical to each other, but, in general, have different sizes, and possibly different surface charges, shapes, etc. In practice, a polydisperse system can be viewed as a mixture with a very large—or essentially infinite—number \( M \) of different species or components, identified by one or several parameters (\( M \) large but finite refers to \emph{discrete polydispersity}, whereas \( M \to \infty \) with a continuous distribution of polydisperse parameters corresponds to \emph{continuous polydispersity}). The present paper considers the discrete polydispersity of spherical colloidal particles, with their diameter being the only polydisperse attribute (size-polydispersity).

When polydispersity is not negligible, the phase behaviour becomes much richer, but determination of the phase transition boundaries requires a much more involved formalism compared with the monodisperse counterpart. In fact, the coexistence condition in terms of intensive variables requires that all phases must have equal temperature, pressure and chemical potentials of the \( M \) components. In the presence of polydispersity, one should thus solve a number of equations of the order of \( M^2 \), a task that is practically impossible for \( M \) large or infinite.

However, the study of phase equilibria can conveniently start from the appropriate thermodynamic potential, which is the Helmholtz free energy \( A \) when the experimentally controlled variables are temperature, volume and number of different colloidal species. In the one-component case, the coexistence condition of equal pressure and chemical potential has a simple geometrical interpretation in terms of free energy density \( a \): the densities of two coexisting phases are determined

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by constructing a double-tangent to a plotted versus particle density. This recipe leads to the well-known Maxwell construction, which connects suitably selected points along a van der Waals (vdW) subcritical isotherm, in order to 'reduce' its unphysical loop to a constant-pressure line characteristic of a first-order phase transition.

In the polydisperse case, significant progress in the very difficult problem of predicting phase equilibria can be obtained for models with truncatable free energies [1]. Here, ‘truncatable’ means that the excess free energy of the polydisperse system turns out to depend only on a finite number of moments of the distribution corresponding to the polydisperse attribute (the diameter \( \sigma \) in the simplest case). For spherical colloids, the excess free energy of the vdW model extended to polydisperse fluids has such a truncatable structure. Due to this property, this vdW theory has often been employed as the simplest model to investigate the effects of polydispersity on the gas–liquid transition [1, 2]. On the other hand, the influence of polydispersity on freezing has been addressed by using the hard-sphere (HS) mixture model, which also admits a truncatable free energy [1] (for the fluid phase, the Boublík–Mansoori–Carnahan–Starling–Leland (BMCSL) [3] expression was employed). It is worth recalling that it is currently believed that size-polydispersity might destabilize crystallization, eventually inhibiting freezing above a certain ‘terminal’ value of polydispersity [1].

The present paper focuses on—and reviews—a number of recent attempts to investigate polydisperse phase equilibria, at least within some approximations, for another prototype model useful for studying colloidal suspensions, namely Baxter’s sticky-hard-spheres (SHS) model [4]. Here the particles are hard spheres with surface adhesion, and the corresponding potential can be obtained as a limit of an attractive potential including steeply repulsive core and short-ranged attractive tail, i.e.

\[
\phi_{ij}^{\text{Baxter SW}}(r) = \begin{cases} 
+\infty, & 0 < r < \sigma_{ij} \equiv (\sigma_i + \sigma_j)/2, \\
-\epsilon_{ij}^{\text{Baxter SW}}, & \sigma_{ij} \leq r \leq R_{ij} \equiv \sigma_{ij} + w_{ij}, \\
0, & r > R_{ij},
\end{cases}
\]

with

\[
\epsilon_{ij}^{\text{Baxter SW}} = k_B T \ln \left( 1 + t_{ij} \frac{\sigma_i}{w_{ij}} \right),
\]

where \( \sigma_i \) is the HS diameter of species \( i \), \( \epsilon_{ij}^{\text{Baxter SW}} > 0 \) and \( w_{ij} \) are the depth and width of the well, respectively, \( k_B \) denotes Boltzmann’s constant, and \( T \) is the temperature. Moreover,

\[
t_{ij} = \frac{1}{12 w_{ij}} \geq 0,
\]

even in the presence of a general solution for this nonlinear algebraic system, the problem of phase coexistence would still remain out of reach in view of the previous remarks.

In a series of recent papers [7–9, 11–13], we attempted to make some progress along two different lines.

First, starting from the density expansion of the cavity function at contact, we considered a sequence of simpler approximations (compared with the PY one) [7–11]. Within the two simplest of these approximations, denoted \( C_0 \) and \( C_1 \) (for reasons that will become clear below), we were able to derive analytically all relevant information regarding structure and thermodynamics, including the phase coexistence, in view of the fact that the corresponding free energy turns out to be truncatable [11]. Due to the simplicity of \( C_0 \) and \( C_1 \), it is however reasonable to expect these approximations to fail at high packing fractions, with a consequent incomplete or even incorrect description of the effects of polydispersity on the phase diagram.

Therefore, in collaboration with Peter Sollich, we recently explored a second approach [12], where the expansion variable (which must be small) is an appropriate polydispersity index. In such a way, we tried to solve the nonlinear algebraic system—involved in the PY result—perturbatively in polydispersity, starting from the monodisperse PY solution.

2. Baxter’s SHS model and PY solution

The SHS model is defined as a limiting case of a particular square-well (SW) model [4], based upon a potential including steeply repulsive core and short-ranged attractive tail, i.e.
where the conventional Baxter parameter $\tau_{ij}$ is an
unspecified increasing function of $T$, and $\tau_{ij}^{-1}$ measures
the strength of the surface adhesion or ‘stickiness’
between particles of species $i$ and $j$.

The ‘sticky limit’ of $\phi_{ij}^{\text{Baxter SW}}(r)$ corresponds to taking
$\eta_{ij} \to 0$. While the SW width goes to zero, its depth $c_{ij}^{\text{Baxter SW}}$
diverges, giving rise to a Dirac delta function in the
Boltzmann factor [4], i.e.
\[
e^{-\rho_0 \phi_{ij}^{\text{SW}}(r)} = \theta(r - \sigma_{ij}) + t_0 \delta_{ij} \delta(r - \sigma_{ij}),
\]
where $\beta = (k_B T)^{-1}$, and $\theta$ and $\delta$ are the Heaviside step
function and the Dirac delta function, respectively.

The advantage of the sticky limit is that one effectively
deals with a single parameter $\tau_{ij}$ for each pair, rather
than a combination of energy and length scales (as
occurs in the square-well model, for which no analytical
solution is known). On the one hand, this particular
limit has the disadvantage of introducing some pathol-
gies into the model, notably in the one-component case
[14]. On the other hand, Baxter’s model represents the
simplest paradigmatic example of a combination of
steep repulsion and short-range attraction which entails
a complete analytical solution in the one-component
case, within a robust approximation such as the PY
closure.

In the multi-component case, the PY solution of the
OZ equation in terms of Baxter’s factor correlation
function reads [5, 6]
\[
g_{ij}(r) = \left\{ \begin{array}{ll}
\left( \frac{1}{2} a_i (r - \sigma_{ij})^2 + (b_i + a_i \sigma_{ij}) \right), & (\sigma_i - \sigma_j)/2 \leq r \leq \sigma_{ij}, \\
(r - \sigma_{ij}) + \Lambda_{ij}, & \sigma_{ij} \leq r \leq \sigma_i, \\
0, & \text{elsewhere},
\end{array} \right.
\]
where the expressions for the parameters $a_i$ and $b_i$ may
be found in [7], while the quantity
\[
\Lambda_{ij} = t_0 y_{ij}(\sigma_{ij}) \sigma_{ij}^2,
\]
which depends on the cavity function at contact $y_{ij}(\sigma_{ij}),$
must be a solution of the following system of quadratic
equations:
\[
\Lambda_{ij} = \alpha_{ij} + \beta_{ij} \sum_m \chi_m \left[ \Lambda_{im} \Lambda_{jm} - \frac{1}{2} (\Lambda_{im} \Gamma_{mj} + \Lambda_{jm} \Gamma_{mi}) \right],
\]
\[i, j = 1, 2, \ldots, M.\]

Here, $\chi_m$ is the molar fraction of the $m$th species
($m = 1, \ldots, M$), and $\alpha_{ij} = t_0 y_{ij}^{\text{HS-PY}}(\sigma_{ij}) \sigma_{ij}^2$, $\beta_{ij} = 12 \rho \Delta y_{ij}(\sigma_{ij})$ ($\rho$ is the total number density), and $\Gamma_{ij} = \sigma_{ij}^4/(1 - \eta)$,
with $\eta$ being the packing fraction [12]. The solution of
these equations for $(\Lambda_{ij})$ is the real bottleneck of the
multi-component PY result, as mentioned in the
Introduction: for large $M$ (and, in particular, for
$M \to \infty$), this calculation is next to impossible,
analytically and numerically.

As a consequence, although the PY closure is commonly believed to be very sound for short-range
potentials (for one-component SHS fluids this was
confirmed by recent numerical simulations [15]), one
has to conclude that, in the multi-component (poly-
disperse) case, the PY solution has very limited practical
usefulness, since its solution scheme cannot be fully
accomplished. This is the reason why other possible
routes have been attempted, as we discuss next.

3. Simplified closures: the class of $C_n$ approximations

A ‘closure’ is a relationship, added to the OZ equation,
between the direct correlation function $c_{ij}(r)$ and
$h_{ij}(r) = g_{ij}(r) - 1$ or $g_{ij}(r) = h_{ij}(r) - c_{ij}(r)$ ($g_{ij}(r)$ being the
radial distribution function) [16].

Let us return to Baxter’s SW model given by
equation (1) (i.e. before the ‘sticky limit’), and consider
the following general class of ‘closures’ [10]:
\[
c_{ij}(r) = \begin{cases}
-1 - \gamma_{ij}(r), & 0 < r < \sigma_{ij}, \\
\gamma_{ij}^{\text{shrink}}(r), & \sigma_{ij} \leq r \leq R_{ij}, \\
0, & r > R_{ij}.
\end{cases}
\]
The expression $c_{ij}(r) = -1 - \gamma_{ij}(r)$ inside the core
($r < \sigma_{ij}$) is exact and dictated by the HS potential. The
form outside the well ($r > R_{ij}$) may then be identified
with the PY approximation,
\[
c_{ij}^{\text{PY}}(r) = f_{ij}[1 + \gamma_{ij}(r)],
\]
since, for Baxter’s potential, the Mayer function,
$f_{ij}(r) = \exp[-\beta \phi_{ij}(r)] - 1$, vanishes for $r > R_{ij}$.

The choice of $c_{ij}^{\text{shrink}}(r)$ inside the well (region which
‘shrinks’ in the sticky limit) defines one particular
closure within the proposed class. Of course,
$c_{ij}^{\text{shrink}}(r) = c_{ij}^{\text{PY}}(r)$ corresponds to the PY approximation.

On the other hand, when $c_{ij}^{\text{shrink}}(r) \neq c_{ij}^{\text{PY}}(r)$, we are in
the presence of mixed closures, which have frequently
appeared in the literature [17]. In order to define
mixed closures simpler than the PY approximation, we
consider the density expansion of the exact direct
correlation function [16], and denote by the
C_\text{n} approximation a truncation of this series to order O(\rho^\text{n}). The simplest two approximations are
\begin{align*}
  c_{ij}^{\text{shrink}}(r) &= f_{ij}(r) \quad (\text{C}_0 \text{ closure}), \\
  c_{ij}^{\text{shrink}}(r) &= f_{ij}(r)[1 + \left( \sum_k \rho_k f_{ik} \ast f_{kj}(r) \right)] \quad (\text{C}_1 \text{ closure}),
\end{align*}

where \( \rho_k \) is the number density of species \( k \), and \( \ast \) denotes convolutive integration [10].

In the ‘sticky limit’ \( R_{ij} \rightarrow \sigma_{ij}^+ \) the well region shrinks, but a ‘memory’ of the approximation chosen for \( c_{ij}^{\text{shrink}} \) remains in the solution of the OZ integral equation. In fact, although all solutions \( q_{ij}(r) \) corresponding to closures belonging to the class given by equation (8) have the same functional form as the PY solution (equation (5)), each closure is characterized by its own approximation to \( y_{ij}(\sigma_{ij}) \), which is involved in the expressions of the parameters \( a_{ij}, b_{ij} \) and \( \Lambda_{ij} \). For instance, the \( \text{C}_0 \) and \( \text{C}_1 \) approximations correspond to
\begin{align*}
  y_{ij}(\sigma_{ij}) &= 1 \quad (\text{C}_0 \text{ closure}), \\
  y_{ij}(\sigma_{ij}) &= 1 + y_{ij}^{(1)}(\sigma_{ij}) \eta \quad (\text{C}_1 \text{ closure}), \quad (11)
\end{align*}

which are, respectively, the zeroth- and first-order truncations of the density expansion for the exact cavity function at contact (see [10] for details).

While a brute-force truncation of the above density expansions leads to analytical expressions sufficiently simple to be applied to the multi-component (polydisperse) case, one should reasonably expect less accuracy, especially in the high-density regime. In the one-component case, we can carefully check this point.

In figure 1, coexistence curves obtained from the \( \text{C}_0 \) and \( \text{C}_1 \) approximations are compared with the PY ones (using both compressibility and energy routes), and with Monte Carlo simulations from [15]. It is apparent how the PY energy route (PYE) yields a rather precise representation of the MC results, unlike the compressibility route (PYC). It is worth noting that the results stemming from the \( \text{C}_1 \) approximation, although rather close to the MC data in the low-density branch, clearly fail to accurately reproduce them for higher densities, as expected.

In spite of their lack of accuracy, the \( \text{C}_0 \) and \( \text{C}_1 \) approximations provide a rather sound basis for obtaining insight into the phase equilibria of polydisperse SHS fluids, since they allow simple analytical, or semi-analytical, treatments.

A first important feature of the \( \text{C}_0 \) and \( \text{C}_1 \) approximations for polydisperse SHS is that the corresponding free energy has a truncatable structure, that is it depends upon a few (four at most) moments of the (discrete) size distribution, \( \xi_v = (\pi/6) \rho \sum \xi_j x_j \sigma_j^3, \) with \( v = 0, 1, 2, 3, \) (A second remarkable fact is that the \( \text{C}_0 \) and \( \text{C}_1 \) approximations are able to describe the so-called fractionation phenomena characteristic in phase equilibria of polydisperse systems. While we refer to a recent review [1] for a detailed description of the increased complexity in the polydisperse phase diagrams, here we just mention two important points. First, fractionation means that the daughter phases, obtained from the demixing of a parent homogeneous phase, need not have the same composition as the parent phase. As a consequence, there is no single coexistence line (‘binodal’) as in the one-component case, but one rather finds a cloud curve, representing the temperature–density dependence line of the low-density majority phase (‘gas’), and a shadow curve, representing the temperature–density dependence of the high-density minority phase (incipient ‘liquid’). While for one-component systems these two curves are identical, for polydisperse systems in general they are not, with the exception of the critical point, where they coincide by definition.

However, in order to apply the \( \text{C}_0 \) and \( \text{C}_1 \) approximations to the multi-component SHS model, we have to tackle a further important problem, that is the definition of the stickiness parameters \( \tau_{ij} \).

4. Size dependence of stickiness parameters

In mixtures, \( \tau_{ij} \) will depend on the particular pair \( i,j \) considered, and should reasonably be expected to be related to the particle sizes. Assuming that we are
dealing only with size-polydispersity, we can always decouple temperature and adhesion as

\[
\frac{1}{\tau_{ij}} = \frac{1}{\tau} \epsilon_{ij} = \frac{1}{\tau} F(\sigma_i, \sigma_j),
\]

(12)

where the last equality stems from the assumption of size-polydispersity and of a purely pairwise potential. Unfortunately, the exact form of the size dependence of these stickiness parameters is still an open problem, due to the lack of experimental and theoretical insights [13]. On the other hand, a few guidelines—based on arguments discussed in [11, 12]—provide, as reasonable and plausible, the following dependencies:

\[
\epsilon_{ij} = F(\sigma_i, \sigma_j) = \begin{cases} 
\frac{\sigma_i^2}{\sigma_0^2}, & \text{Case I}, \\
\frac{\sigma_i}{\sigma_0}, & \text{Case II}, \\
1, & \text{Case IV}, \\
\frac{\sigma_0}{\sigma_{ij}}, & \text{Case V}.
\end{cases}
\]

(13)

Here, \(\sigma_0\) is a characteristic reference length (e.g., the parental mean diameter) and the numbering of the various cases follows the convention of previous work [11, 12].

Figure 2 reports the results of the calculation of the cloud and shadow curves for polydisperse SHS within the simple \(C_0\) approximation. Here and below, polydispersity is measured by an index \(s\), which is the normalized standard deviation of the size distribution. Hence, \(s=0\) corresponds to a mono-disperse case, whereas \(s=0.1\) and \(s=0.3\) indicate moderate and significant polydispersity, respectively. The top panel of figure 2 depicts the results for case I of the size dependence of the stickiness parameters. As \(s\) increases, the coexistence region shrinks, thus suggesting that polydispersity disfavours the phase transition. On the other hand, this trend is markedly case dependent, as illustrated in the bottom panel of figure 2, where the cloud–shadow pairs with polydispersity \(s=0.3\) are displayed for different size dependence cases. It can clearly be seen that, whereas for cases I and V the same trend is observed, case IV seems to suggest a widening of the phase coexistence region (hence favouring the phase transition).

In view of the lack of numerical simulations for polydisperse SHS in order to make a comparison, we have no way, at the present stage, of checking how realistic these results are. On the other hand, we might suspect, based on the comparison in the one-component case, \(C_0\) to fail to provide an accurate representation in the region of low temperatures and high densities. This is the reason why other possible approaches have recently been tested. We now illustrate a different perturbative approach that has proved to be promising in this context.

5. Perturbative treatment of the SHS-PY solution

The main difficulty in dealing with the PY solution for polydisperse SHS stems from the solution of the coupled quadratic system of equations (7). As the one-component case has a well-defined solution, one might then suspect that—for weak polydispersity—a perturbative expansion around this solution might include the main effects of polydispersity. This is, in fact, what happens, as recently shown [12] by exploiting a general perturbation theory due to Evans [18]. The main idea is that, for weak polydispersity, size distributions are narrowly peaked around a mean reference value (\(\sigma_0\) in the present case), and hence all quantities such as

\[
\delta_i = \frac{\sigma_i - \sigma_0}{\sigma_0} \ll 1
\]

(14)

are small. Therefore, one might expand both \(\epsilon_{ij}\), and all quantities appearing in \(\Lambda_{ij}\), in powers of \(\delta_i\). A similar expansion can be performed in the free energy, and hence all thermodynamic quantities can be computed. The entire procedure is described in detail in [1, 12, 18]. Here, we just summarize the main results.

The approximate range of validity of the perturbation expansion can be envisaged by considering the polydisperse HS case where the ‘exact’ BMCSL approximation [3] can be compared with the corresponding perturbative solution based on the one-component (\(s=0\)) counterpart. This is reported in the top-left part of figure 3, where the quantity \(\beta P v_0\) (\(v_0 = \pi \sigma_0^3 / 6\)) is plotted against the packing fraction \(\eta\) for increasing values of polydispersity. It is apparent that the perturbative solution remains rather close to the ‘exact’ polydisperse BMCSL solution even for moderate polydispersity \(s \leq 0.3\), which is the maximum value considered in the remaining part of this work. The remaining plots in figure 3 display the effect of polydispersity on the PY pressure equation of state as obtained from the energy route and for decreasing values of the temperature \(\tau\). In the one-component case \(s=0\), a van der Waals loop starts to appear when we cross the critical temperature \(\tau_c \sim 0.1185\) coming from the high \(\tau\) regime. Obviously, this signals the onset of a liquid–gas phase transition, and the corresponding phase diagram can be obtained by a standard Maxwell construction by connecting appropriate points with the same pressure. In the presence of polydispersity (here represented by choice IV for the size dependence of the stickiness
parameters), the same procedure cannot be applied due to fractionation, as already discussed. Nevertheless, we can clearly see that, as \( s \) increases, the van der Waals loop region (when present) expands, thus suggesting that a phase transition is favoured by the presence of polydispersity. A similar feature occurs for the polydisperse van der Waals model [2] and for the numerical results of the PY compressibility equation of state [19] (note that, in the latter, a gap rather than a loop signals the onset of the transition). A somewhat surprising feature is that, at fixed packing fraction \( \eta \), the pressure decreases with increasing polydispersity less in the presence of adhesion rather than in its absence (i.e. for the HS case). An intuitive plausible interpretation of this feature can be found in [12].

The same perturbative approach allows the determination of the cloud and shadow curves for the various size dependence cases of \( \tau_0^{-1} \). This is reported in figure 4 for cases II, IV (top panel) and I, V (bottom panel). In the first case, the cloud and shadow lines collapse into a single curve, and this can be understood on the basis of the particular scaling properties of the free energy to this order in perturbation theory [12]. In all cases, there is a breakdown of the perturbation theory on approaching the critical point, and this is a known general drawback of Evans’ perturbative scheme. Nevertheless, in all cases and to this order in perturbation theory, there is a tendency of the phase coexistence region to increase with polydispersity, in qualitative agreement with the intuitive picture obtained from figure 3.

It is worth stressing the difference with respect to previous non-perturbative results stemming from the \( C_0 \) solution, where all different cases (with the notable exception of IV) predicted a reduction of the phase coexistence region. While in the \( C_0 \) description we have provided a careful treatment of polydispersity at the
expense of the accuracy of the exploited approximation, in the perturbative description of the PY solution, polydispersity is assumed to be small and hence one might suspect that solutions with large polydispersities cannot fit within this picture. On balance, however, we would favour the latter rather than the former description. An almost correct representation of the one-component counterpart is a necessary requirement for checking the effect of polydispersity, and we are not aware of any physical or experimental system where the effects of polydispersity are so strong that they could not be taken into account, at least at the simplest qualitative level, by the perturbative scheme proposed here.

Along this line, some further proposals have been put forward in [12] to derive a phenomenological BMCSL-like approximation for SHS, which might be regarded as our ‘best and simplest guess’ to the exact phase behaviour of polydisperse SHS. Even on the size dependence of \( \tau_{\eta}^{-1} \) some possible support of the proposed forms may be put forward [12, 13].

6. Conclusions

In this work, we have summarized recent advances in predicting, theoretically, the phase diagram for polydisperse suspensions of colloidal particles with surface adhesion, within the simple description of Baxter’s model. Emphasis was placed on the crucial—unsolved—step required to obtain the multi-component SHS-PY solution, and the proposed recipes to deal with this problem. The first is based on a simplification of the closure. It has the
advantage of allowing a complete analytical analysis on the effects of polydispersity, including fractionation, but has the disadvantage of very questionable accuracy. The second is based on a perturbative method, starting from the energy PY one-component solution, which is known to provide an accurate description of the phase diagram. The drawback of this scheme is that it works for mild polydispersity, but it cannot describe the changes in the critical point region. Notwithstanding these limitations, this novel approach is expected to find practical application in the interpretation of all those phenomena where Baxter’s model and polydispersity both play a role.

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Figure 4. Cloud/shadow pairs from the perturbative results for the PY solution of the SHS Baxter model. In clockwise order, the results for choices II, IV (top) and I, V (bottom) are depicted. In the top two panels, the cloud and shadow curves coincide to this order in perturbation, whereas in the bottom panels they are different. In order to have all pictures on the same scale, the selected value for polydispersity is $s = 0.3$ for models II and IV (top) and $s = 0.1$ for models I and V (bottom). In all cases, the continuous curve represents the monodisperse ($s = 0$) result.