

Improved Gaussian self-consistent method—applications to homopolymers with different architectures in dilute solution

Edward G. Timoshenko^{a)}

Theory and Computation Group, Department of Chemistry, University College Dublin, Belfield, Dublin 4, Ireland

Yuri A. Kuznetsov

Centre for High Performance Computing Applications, University College Dublin, Belfield, Dublin 4, Ireland

(Received 16 April 2002; accepted 21 June 2002)

A version of the Gaussian self-consistent (GSC) method, which avoids the use of the Edwards' virial expansion, is presented. Instead, the mean energy is evaluated directly via a convolution of the attractive-part of the pair-wise nonbonded potential with the Gaussian trial radial distribution function. The hard-sphere repulsion is taken into account via a suitably generalized Carnahan–Starling term. Comparison of the mean-squared inter-monomer distances and radius of gyration, as well as of the mean energy, between the results from the GSC calculations and Monte Carlo (MC) simulation in continuous space are made across the coil-to-globule transition for isolated ring, open and star homopolymers of varied lengths and flexibility. Importantly, both techniques utilize the same polymer model so that the data points could be directly superimposed. A surprisingly good overall agreement is found between these GSC and MC results. Caveats of the Gaussian technique and ways for going beyond it are also discussed. © 2002 American Institute of Physics. [DOI: 10.1063/1.1500357]

I. INTRODUCTION

The Gaussian self-consistent (GSC) method represents a quite general, albeit relatively simple, theoretical framework for studying the equilibrium, dynamics and kinetics of conformational changes in polymer solutions. One of its most attractive features is that it can be applied to virtually any type of heteropolymer,^{1–3} with any specific interaction terms involving, e.g., chain stiffness^{4,5} or charges.⁶ The connectivity of the chain can be arbitrary: an open polymer,⁷ a ring,⁸ a star,⁹ or a dendrimer,¹⁰ whether in the limit of a single chain¹¹ or at finite concentrations.¹² Moreover, not only the radius of gyration of the polymer or the total density can be computed, but much finer details of the conformational structure, such as the mean-squared distances between monomers or radial distributions at a given time, are available. Such a versatility naturally comes at the cost of certain limitations and inaccuracies intrinsic to the technique. However, these are well known and manageable, although often largely overstated in the popular view. Some of these, such as the Gaussian shape of the radial distribution function (RDF), are quite unavoidable at the level of the GSC theory, but others, such as the traditional use of the Edwards' virial expansion,^{13,14} can indeed be surpassed as we shall demonstrate in the current work. This would permit us for the first time to compare observables between the predictions of the GSC technique and Monte Carlo (MC) simulation in continuous space based

on the same model for given values of thermodynamic parameters throughout the range of the coil-to-globule transition.

Historically, perhaps the first of the equilibrium versions of the Gaussian method in application to a single homopolymer chain were independently proposed by Edwards *et al.*¹⁵ and des Cloizeaux.¹⁶ The approach of the work by Edwards¹⁵ relied on the following idea. One would like to approximate the coordinates of the real homopolymer chain, X_i , governed by the Gibbs distribution with the exact Hamiltonian H , via a “trial” Gaussian chain with the monomer positions $X_i^{(0)}$, governed by a simpler trial Hamiltonian $H^{(0)}$. By assuming that the quantities $\Delta X_i = X_i - X_i^{(0)}$ and $\Delta H \equiv H - H^{(0)}$ are small and hence by applying the Taylor expansion to the equilibrium Gibbs averages, in the first order, one requires that the mean-squared radius of gyration obeys

$$\langle R_g^2 \rangle_0 = \langle R_g^{(0)2} \rangle_0,$$

$$\text{or equivalently } \langle R_g^{(0)2} H^{(0)} \rangle_0 - \langle R_g^{(0)2} \rangle_0 \langle H^{(0)} \rangle_0 = 0. \quad (1)$$

Such a theory predicts the correct mean-field swelling exponent $\nu_F = 3/5$ for the repulsive coil, only missing the subtle renormalization group correction. It also gives appropriately $\nu_\theta = 1/2$ for the ideal coil, and $\nu_g = 1/3$ for the globule. Moreover, this approach can be further extended to kinetics by considering time dependent analogues of Eq. (1).^{18,17,18}

A seemingly more refined theory proposed by des Cloizeaux¹⁶ uses $N-1$ (where N is the degree of polymerization) variational variables $X_q^{(0)}$, which are the normal modes of the monomer positions for a ring chain $X_i^{(0)}$. This technique is based on the standard Gibbs–Bogoliubov varia-

^{a)} Author to whom correspondence should be addressed.

Web: <http://darkstar.ucd.ie>; electronic mail: edward.timoshenko@ucd.ie

tional principle with a quadratic diagonal trial Hamiltonian $H^{(0)}$. The equilibrium corresponds to the minimum of the trial free energy, $\mathcal{A}_{\text{trial}} = \mathcal{A}^{(0)} + \langle H - H^{(0)} \rangle_0$, where $\mathcal{A}^{(0)}$ is the free energy associated with $H^{(0)}$. Expressed in the Edwards' form the extremum conditions for the trial free energy are precisely the following $N-1$ equations

$$\langle \mathbf{X}_q^2 \rangle_0 = \langle \mathbf{X}_q^{(0)2} \rangle_0,$$

$$\text{or equivalently } \langle \mathbf{X}_q^{(0)2} H^{(0)} \rangle_0 - \langle \mathbf{X}_q^{(0)2} \rangle_0 \langle H^{(0)} \rangle_0 = 0. \quad (2)$$

Unfortunately, this theory, although more accurate for polymers around the theta-point and indeed for the globule, is known to have a deficiency in that it predicts the exponent $\nu = 2/3$ instead of the mean-field value of $\nu_F = 3/5$ for the swollen coil in the thermodynamic limit $N \rightarrow \infty$. For this reason, some suspicion arose concerning the validity of the theory by des Cloizeaux, hampering further efforts in this direction. It should be emphasised, however, that for finite values of N the resulting theory does give quite reasonable numerical results fairly close to $\nu_F = 3/5$. In fact, the numerically fitted value of the swelling exponent tends to be somewhere in between the two above theoretical predictions. In a way, this is not an unusual situation, even for the more elaborate integral equations theories of molecular fluids used in attempts to deduce the Flory exponent.^{19,20} More importantly, this well known deficiency of the theory by des Cloizeaux becomes irrelevant when considering the theta-point or the poor solvent conditions.

Therefore, activity on further extending and improving the GSC theory has persisted, notably in Milan by Allegra and Ganazzoli *et al.*,^{2,3,5,10,21-23} in Saclay by Orland and Garel *et al.*,^{6,17,18} as well as in our Group in Dublin,^{1,4,7,8,11,12} and some collaborations between these have emerged.⁹ Furthermore, extensions of the GSC theory to kinetics, as well as to essentially any polymeric system, have been also achieved. For the former end, one proceeds from the Langevin equation and approximates the exact stochastic ensemble via a linear trial one. We refer the reader to our papers in Refs. 1, 9, and 11, for the ultimate formulation of the GSC theory in terms of the real space variables. Such a version of the GSC method avoids the limitations of the earlier normal modes formulations since it can distinguish the frustrated states of heteropolymers with spontaneously broken kinematic symmetries associated with the connectivity structure.

Despite the relative success of the most recent form of the GSC method, the theoretical situation is not as yet fully satisfactory and further efforts along these lines are required. First, one would like to be able to use the actual molecular interaction parameters rather than the coefficients in the Edwards' virial expansion, which are somewhat obscure in their meaning. Second, the convergence of such a virial expansion in the dense liquid globular state is rather problematic, being at best that of an asymptotic series. Third, despite the conventional view, the three-body term is unable by itself to fully withstand a catastrophic pair-wise collapse of monomers onto each other in the attractive regime for heteropolymers. In Refs. 1 and 11 an *ad hoc* self-interaction energy term has been added to tackle this problem, but it is merely a

tweak of the model. Finally, the effective three-body and higher order virial terms make the current theory numerically inefficient due to a scaling law involving a high power in N for the time expense per step, despite the fact that the original model only contains two-body interactions.

Therefore, in the current work we shall rid ourselves of relying on the virial representation of the Hamiltonian entirely. Indeed, the exact molecular two-body potential can be averaged over the Gaussian RDF directly. However, the Gaussian function does not vanish at the origin, yielding a divergence from the repulsive part of the Lennard-Jones pairwise potential. One simple alternative is to use the latter potential with a hard core part, as in Ref. 24, in which case one has to cutoff the integration at the hard-sphere diameter, or more generally, within the excluded volume area.

To include the effect of the hard-sphere repulsion one can in principle use any of the standard reference techniques developed for the hard sphere liquids, e.g., the Percus–Yevick expression for RDF, or the Carnahan–Starling (CS) free energy formula.^{25,26} We prefer to use the latter due to its practical simplicity and accuracy when compared to molecular simulations. The Carnahan–Starling equation, which was originally derived for a single component hard-sphere liquid, has been naturally extended to mixtures. It is quite popular in the liquid matter literature at present (see, e.g., Ref. 27). Its extension to polymeric fluids is somewhat less obvious though. Here we propose to express the partial packing coefficient for each of the monomers via an integral of the Gaussian trial RDF over the excluded volume area. The resulting GSC equations involve well tractable, albeit somewhat more complicated than before, expressions, since they now depend on the exact shapes of the molecular potentials.

Although we shall present the resulting technique in its most general form, which is applicable to any type of polymeric system, our numerical analysis in this work will be restricted by homopolymers of three different architectures—ring, open and star—across the coil-to-globule transition. This is done to enable us to make a comparison of the results from the present GSC theory with the MC data obtained in our previous paper in Ref. 24.

II. MODEL

The current coarse-grained polymer model is based on the following Hamiltonian (energy functional)^{4,9,24,28} in terms of the monomer coordinates, \mathbf{X}_i :

$$\begin{aligned} \frac{H}{k_B T} = & \frac{1}{2\ell^2} \sum_{i \neq j} \kappa_{ij} (\mathbf{X}_i - \mathbf{X}_j)^2 \\ & + \frac{1}{2\ell^2} \sum_{i \approx j \approx k} \lambda_{ijk} (\mathbf{X}_i + \mathbf{X}_k - 2\mathbf{X}_j)^2 \\ & + \frac{1}{2} \sum_{ij, i \neq j} (U_{ij}^{(lj)}(|\mathbf{X}_i - \mathbf{X}_j|) + U_{ij}^{(\text{cou})}(|\mathbf{X}_i - \mathbf{X}_j|)). \end{aligned} \quad (3)$$

Here the first term represents the connectivity structure of the polymer with harmonic springs of a given strength κ_{ij} , in-

roduced between any pair of connected monomers (which is denoted by $i \sim j$). The second term represents the bending energy penalty given by the square of the local curvature with a characteristic stiffness λ_{ijk} between any three consecutively connected monomers (which is denoted by $i \approx j \approx k$) in the form of the Kratky–Harris–Hearst. Below we shall prefer to rewrite the first two terms in the following equivalent form:

$$U_{ij}^{(ij)}(r) = \begin{cases} +\infty, & r < r_i^{(0)} + r_j^{(0)} \\ U_{ij}^{(0)} \left(\left(\frac{r_i^{(0)} + r_j^{(0)}}{r} \right)^{12} - \left(\frac{r_i^{(0)} + r_j^{(0)}}{r} \right)^6 \right), & r > r_i^{(0)} + r_j^{(0)}, \end{cases} \quad (5)$$

where there is also a hard core part with the monomer radii $r_i^{(0)}$, and where $U_{ij}^{(0)}$ are the dimensionless strengths of the interactions. The Coulomb interaction potential similarly is

$$U_{ij}^{(\text{cou})}(r) = q_i q_j \frac{l_B}{r} \exp(-r/l_D), \quad (6)$$

$$l_B \equiv \frac{Q_0^2}{4\pi\epsilon_0 k_B T},$$

where l_D is the Debye screening length, ϵ_0 is the dielectric permittivity of vacuum, $Q_i = Q_0 q_i$ are the charges, and l_B is the Bjerrum length.

III. METHOD

A. Equations of the GSC method

The main objects in the GSC method are the mean-squared distances between monomers

$$\mathcal{D}_{ij}(t) \equiv \frac{1}{3} \langle (\mathbf{X}_i(t) - \mathbf{X}_j(t))^2 \rangle. \quad (7)$$

Note that our convention includes the factor of 1/3 here and hence in the later definition of the mean-squared radius of gyration according to the tradition. This allows us to rid of such factors from the radial distribution function (RDF) and numerous averages over it.

The GSC method is based on replacing the stochastic ensemble for \mathbf{X}_i with the exact Hamiltonian in the Langevin equation of motion onto the trial ensemble $\mathbf{X}_i^{(0)}(t)$ with a trial Hamiltonian $H^{(0)}(t)$. The latter is taken as a generic quadratic form with the matrix coefficients, which are called the time-dependent effective potentials

$$H^{(0)}[\mathbf{X}(t)] = \frac{1}{2} \sum_{ij} V_{ij}(t) \mathbf{X}_i(t) \mathbf{X}_j(t). \quad (8)$$

Then one requires that the inter-monomer correlations satisfy the condition

$$\langle \mathbf{X}_i(t) \mathbf{X}_j(t) \rangle_0 = \langle \mathbf{X}_i^{(0)}(t) \mathbf{X}_j^{(0)}(t) \rangle_0, \quad (9)$$

$$\frac{H^{(\text{bond})}}{k_B T} = \frac{1}{2} \sum_{ij} U_{ij}^{(\text{bond})} (\mathbf{X}_i - \mathbf{X}_j)^2. \quad (4)$$

Finally, the third and fourth terms represent pair-wise non-bonded interactions between monomers such as the van der Waals and Coulomb forces. We can adopt the Lennard-Jones form of the former potential

which means that the trial ensemble well approximates the exact one as far as monomer correlations are concerned. This procedure yields expressions for the effective potentials via the instantaneous mean energy

$$V_{ij}(t) = -\frac{2}{3} \frac{\partial \mathcal{E}[\mathcal{D}(t)]}{\partial \mathcal{D}_{ij}(t)}, \quad (10)$$

and hence the mean-squared distances themselves satisfy the self-consistent equations.⁹ These, in the absence of the hydrodynamic interaction, are simply

$$\frac{\zeta_b}{2} \frac{d}{dt} \mathcal{D}_{ij}(t) = -\frac{2}{3} \sum_k (\mathcal{D}_{ik}(t) - \mathcal{D}_{jk}(t)) \times \left(\frac{\partial \mathcal{A}[\mathcal{D}(t)]}{\partial \mathcal{D}_{ik}(t)} - \frac{\partial \mathcal{A}[\mathcal{D}(t)]}{\partial \mathcal{D}_{jk}(t)} \right). \quad (11)$$

Here ζ_b is the friction coefficient of a monomer, and the instantaneous free energy has the same functional expression via the instantaneous $\mathcal{D}_{ij}(t)$ as it has at equilibrium. Extension to the preaveraged hydrodynamic approximation is quite straightforward also and it is discussed in Ref. 1.

The stationary limit of these equations produces the equations for the minimum of the free energy, which are the same as those derived from the Gibbs–Bogoliubov variational principle with a generic quadratic trial Hamiltonian. Although in this paper we shall only be concerned with the equilibrium properties, the numerical solution of Eq. (11), applied until the stationary limit is reached, presents by far the most efficient technique for finding the global free energy minimum. This, based on the fifth order adaptive step Runge–Kutta integrator,¹ was used for obtaining the results from the GSC technique in this work.

It should be also noted that the systems studied here possess a large number of kinematic symmetries for \mathcal{D}_{ij} , and hence for V_{ij} , matrices coming from their symmetricity and from the equivalence of any monomer in a ring, or any arm in a star homopolymers. Thus, the computational expenses per step in our calculations are of order $t_c \sim NF$, where $N/2 \lesssim F \lesssim N^2/2$ is the total number of independent elements in the matrix \mathcal{D}_{ij} . These symmetries significantly reduce the computational times compared to MC for an equivalent sys-

tem, where such symmetries only appear in the observables after averaging over the statistical ensemble. For comparison, the computational expenses per step in MC are of order $t_c \sim N\Delta t S$, where $\Delta t \sim N^2$ is the number of MC steps needed to ensure a good statistical independence between measurements, and where S is the number of measurements needed for sampling of observables. Typical values of S should be of order of $10^4 - 10^6$ for a good accuracy in the present case.²⁴ Moreover, kinetic iteration of the GSC equations towards the equilibrium is also significantly faster than the equivalent equilibration procedure in the MC. For example, for a ring polymer of $N=150$ units in the good solvent, $U^{(0)}=1$, the GSC method turns out to be ≈ 1000 times faster²⁹ than MC for gaining the same data, whereas for an open chain of the same length, which possesses much fewer kinematic symmetries, GSC is ≈ 200 times faster than MC.

B. Hard-sphere contribution

Carnahan and Starling have devised a simple but rather accurate equation of state for hard sphere liquids²⁶ in terms of the packing coefficient η , yielding the following free energy:

$$\frac{\mathcal{A}^{(\text{CS})}}{k_B T N} = F^{(\text{CS})}(\eta) = \frac{\eta(4-3\eta)}{(1-\eta)^2}, \quad (12)$$

$$\eta = N \frac{v_0}{V},$$

where v_0 and V are the volumes of the hard sphere and of the whole system, respectively. This is obtained from an interpolation formula for the virial coefficients based on several of them known exactly (see more detailed discussions in, e.g., Ref. 25).

To extend these ideas to polymeric fluids, we would have to distinguish the packing coefficients for individual monomers. In Eq. (12) we can write $N = \sum_i$; the volume of the sphere v_0 is equal to the 1/8th of its excluded volume; and the inverse volume is equal to the RDF of the ideal reference system, $1/V = g_{\text{ideal}}^{(2)}$. Thus, we can similarly express the packing coefficient for the i th monomer as a sum over all other monomers of the 1/8th of the integral over the excluded volume of the monomer i of the RDF for the Gaussian reference system

$$g_{ij}^{(2)}(\mathbf{r}) = \frac{1}{(2\pi\mathcal{D}_{ij})^{3/2}} \exp\left(-\frac{\mathbf{r}^2}{2\mathcal{D}_{ij}}\right), \quad (13)$$

yielding finally

$$\eta = N \frac{v_0}{V} \rightarrow \eta_i = \sum_{i \neq j} \frac{a}{8} \int_{|\mathbf{r}| < r_i^{(0)} + r_j^{(0)}} d\mathbf{r} g_{ij}^{(2)}(\mathbf{r}). \quad (14)$$

One may note that the Gaussian distribution does not possess a well defined volume beyond which it vanishes. Thus, to account for this we can, in principle, include a multiplicative parameter a , which should be once and for all chosen to match the data best, but this should, in any case, be fairly close to the unity. Therefore, the total hard-sphere contribution will be

$$\frac{\mathcal{A}^{(\text{hs})}}{k_B T} = N F^{(\text{CS})}(\eta) \rightarrow \sum_i F^{(\text{CS})}(\eta_i). \quad (15)$$

C. Free energy in the GSC method

The total mean energy includes both the bonded and the pair-wise nonbonded interactions via

$$\frac{\mathcal{E}^{(\text{int})}}{k_B T} = \frac{1}{2} \sum_{ij} \left(3 U_{ij}^{(\text{bond})} \mathcal{D}_{ij} + \int_{r > r_i^{(0)} + r_j^{(0)}} d\mathbf{r} g_{ij}^{(2)}(\mathbf{r}) [U_{ij}^{(lj)}(\mathbf{r}) + U_{ij}^{(\text{cou})}(\mathbf{r})] \right). \quad (16)$$

Note that here we integrate only beyond the excluded volume as the hard-sphere contribution will be included explicitly via the Carnahan–Starling term.

Given that the conformational Gaussian entropy has been calculated by us in Appendix B of Ref. 1, the total free energy can be summarized as follows:

$$\mathcal{A} \equiv \mathcal{E} - T\mathcal{S}, \quad \mathcal{E} = \mathcal{E}^{(\text{int})} + \mathcal{A}^{(\text{hs})}, \quad (17)$$

$$\mathcal{E}^{(\text{int})} = \mathcal{E}^{(\text{bond})} + \mathcal{E}^{(lj)} + \mathcal{E}^{(\text{cou})}, \quad \mathcal{S} \equiv \mathcal{S}^{(\text{gau})},$$

where its various terms are given by

$$\frac{\mathcal{E}^{(\text{bond})}}{k_B T} = 3 \sum_{i < j} U_{ij}^{(\text{bond})} \mathcal{D}_{ij}, \quad (18)$$

$$\frac{\mathcal{E}^{(lj)}}{k_B T} = \sum_{i < j} U_{ij}^{(0)} E^{(lj)}[y_{ij}], \quad (19)$$

$$\frac{\mathcal{E}^{(\text{cou})}}{k_B T} = \sum_{i < j} \frac{l_B q_i q_j}{r_i^{(0)} + r_j^{(0)}} E^{(\text{cou})}[y_{ij}, (r_i^{(0)} + r_j^{(0)})/l_D], \quad (20)$$

$$\frac{\mathcal{S}^{(\text{gau})}}{k_B} = \frac{3}{2} \ln \det R^{(N-1)}, \quad (21)$$

$$\frac{\mathcal{A}^{(\text{hs})}}{k_B T} = \sum_i F^{(\text{CS})}(\eta_i), \quad (22)$$

$$F^{(\text{CS})}(\eta_i) = \frac{\eta_i(4-3\eta_i)}{(1-\eta_i)^2},$$

and the arguments of these functions are defined as

$$\eta_i = a \sum_{j \neq i} F^{(n)}[y_{ij}], \quad (23)$$

$$y_{ij} = \frac{\sqrt{D_{ij}}}{r_i^{(0)} + r_j^{(0)}}, \quad (24)$$

$$R_{ij} = \frac{D_i + D_j}{2} - \mathcal{R}_g^2 - \frac{D_{ij}}{2}, \quad (25)$$

$$\mathcal{D}_i \equiv \frac{1}{N} \sum_j D_{ij}, \quad \mathcal{R}_g^2 \equiv \frac{1}{2N^2} \sum_{ij} D_{ij}.$$

$$E^{(lj)}[y] \equiv \frac{1}{y^3} \sqrt{\frac{2}{\pi}} \int_1^\infty x^2 dx \exp\left(-\frac{x^2}{2y^2}\right) \left(\frac{1}{x^{12}} - \frac{1}{x^6}\right)$$

$$= \frac{\sqrt{2/\pi} \exp\left(-\frac{1}{2y^2}\right) (y - y^3 + 3y^5 + 300y^7 - 210y^9) - \operatorname{erfc}\left(\frac{1}{y\sqrt{2}}\right) (1 + 315y^6)}{945y^{12}}, \quad (27)$$

$$E^{(\text{cou})}[y, k] \equiv \frac{1}{y^3} \sqrt{\frac{2}{\pi}} \int_1^\infty x^2 dx \exp\left(-\frac{x^2}{2y^2}\right) \frac{\exp(-kx)}{x} = \exp\left(-\frac{1}{2y^2} - k\right) \left(\frac{1}{y} \sqrt{\frac{2}{\pi}} - \frac{2k}{\pi} \exp\left(\frac{(1+ky^2)^2}{2y^2}\right) \operatorname{erfc}\left(\frac{1+ky^2}{y\sqrt{2}}\right)\right). \quad (28)$$

Here we have used the standard definitions for the error functions

$$\operatorname{erf}(z) \equiv \frac{2}{\sqrt{\pi}} \int_0^z dx \exp(-x^2), \quad (29)$$

$$\operatorname{erfc}(z) \equiv 1 - \operatorname{erf}(z).$$

Note also that for a large z , numerically, one has to use the truncated asymptotic expansion in calculating $E^{(\text{cou})}$ to avoid divergences

$$\operatorname{erfc}(z) \exp(z^2) \approx \frac{1}{z\sqrt{\pi}} \left(1 - \frac{1}{2z^2} + \frac{3}{4z^4} - \frac{15}{8z^6} + \dots\right). \quad (30)$$

IV. RESULTS

Here we shall restrict ourselves by the case of homopolymers, so that in Eq. (3) all nonzero bonded interaction constants are equal: $\kappa_{ij} = \kappa$, $\lambda_{ijk} = \lambda$, as well as all nonbonded interaction parameters are identical in Eq. (5): $U_{ij}^{(0)} = U^{(0)}$, $r_i^{(0)} \equiv d/2$. We also choose the hard-sphere diameter d equal to the length ℓ defined in Eq. (3), as in Ref. 24. Moreover, henceforth we shall use the mean energy $\mathcal{E}^{(\text{int})}$ expressed in units of $k_B T$ and the mean-squared distances \mathcal{D}_{ij} and the mean-squared radius of gyration $3\mathcal{R}_g^2$ expressed in units of ℓ^2 .

First, to understand the influence of the adopted Carnahan–Starling term, we shall look at the case of a ring homopolymer with varied values of the spring constant in the good athermal solvent, $U^{(0)} = 0$. In Table I we compare values of the mean energy $\mathcal{E}^{(\text{int})}$ between MC and GSC with two choices of the multiplicative parameter a in Eq. (23) for the packing coefficient, namely $a = 0.9$ and $a = 1.0$. Likewise, in

Finally, the functions of y_{ij} are

$$F^{(\eta)}[y] \equiv \frac{1}{(2y)^3} \sqrt{\frac{2}{\pi}} \int_0^1 x^2 dx \exp\left(-\frac{x^2}{2y^2}\right) \\ = \frac{y \operatorname{erf}\left(\frac{1}{y\sqrt{2}}\right) - \sqrt{2/\pi} \exp\left(-\frac{1}{2y^2}\right)}{8y}, \quad (26)$$

Table II we present the data for the mean-squared radius of gyration $3\mathcal{R}_g^2$. One can see that the results from the GSC theory simply coincide with those from MC as $\kappa \rightarrow 0$. Yet, the agreement in the energy is somewhat better for the theory with $a = 0.9$ than for that with the naive choice $a = 1.0$, and that is how exactly the particular value $a = 0.9$ has been chosen by us. A typical relative error in the energy is less than 1.5 percent for $a = 0.9$ and is under 4 percent for $a = 1.0$. The agreement for the mean-squared radius of gyration is somewhat less impressive—the relative error steadily increases with κ , being better for $a = 0.9$, but never exceeding a few dozen percent.

One can comment on the reason why a value $a < 1$ produces a somewhat better agreement with MC. To facilitate this discussion, RDFs from the GSC and MC techniques are exhibited in Fig. 1. In Ref. 24 we have discussed the role of the “correlation hole” at small separations beyond the excluded volume area, which is also evident for $r \leq 15$ in the main part of Fig. 1. As this feature is absent in the Gaussian theory [see Eq. (13)], the resulting packing coefficient values in Eq. (23) are overestimated. Thus, reducing the parameter a permits us to effectively lower the packing coefficient and this can be done only once because the correlation hole effect is expressed in terms of the function of the dimensionless variables

$$\hat{g}_{ij}^{(2)}(\hat{r}) \equiv \mathcal{D}_{ij}^{3/2} g_{ij}^{(2)}(r), \quad (31)$$

$$\hat{r} \equiv \frac{r}{\mathcal{D}_{ij}^{1/2}}.$$

Next, we would like to analyze the dependence on the degree of polymerization, N , in the good solvent, $U^{(0)} = 0$, (the second block in Tables I and II). The mean energy in this case comes entirely from the bonded interactions and hence the relative energy deviation of GSC from MC does not in-

TABLE I. Comparison of the mean energy values $\mathcal{E}^{(int)}$ (in $k_B T$ units) for different homopolymers from the MC simulation (second column) based on the data set of Ref. 24 and from the GSC theory with the parameter $a = 0.9$ (third column) and $a = 1.0$ (fourth column). The fifth and sixth columns contain the relative deviation $\delta = (\mathcal{E}^{(int)}(GSC)/\mathcal{E}(MC) - 1) \times 100\%$ for these two cases given as a percentage. The model parameters, which are suppressed within the tables, such as, e.g., $\lambda, U^{(0)}$, are equal to zero.

System	MC	GSC, $a = 0.9$	GSC, $a = 1.0$	$\delta_{a=0.9}, \%$	$\delta_{a=1.0}, \%$
Ring, $N = 100$					
$\kappa = 0.01$	148.6	148.58	148.59	0	0
$\kappa = 0.1$	151.21	151.22	151.51	0	0.2
$\kappa = 0.2$	155.24	155.52	156.22	0.2	0.6
$\kappa = 0.5$	168.3	170.03	172.0	1.0	2.2
$\kappa = 2.0$	232.4	232.1	239.0	-0.1	2.8
Ring, $\kappa = 1$					
$N = 50$	94.76	96.08	97.99	1.4	3.4
$N = 100$	190.4	193.1	196.9	1.4	3.4
$N = 200$	381.5	386.9	394.4	1.4	3.4
$N = 300$	572.6	580.6	591.9	1.4	3.4
Open, $\kappa = 1$					
$N = 150$	284.2	287.1	292.7	1.0	3.0
$N = 200$	378.7	383.9	391.3	1.4	3.3
Semiflexible ring					
$\lambda = 1, \kappa = 1$					
$N = 50$	106.6	104.9	107.3	-1.6	0.7
$N = 100$	213.3	210.2	215.1	-1.4	0.8
$N = 200$	426.8	420.9	430.7	-1.4	0.9
$N = 300$	640.2	631.6	646.2	-1.3	0.9
Semiflexible ring					
$\lambda = 5, \kappa = 1$					
$N = 50$	110.1	110.5	113.2	0.4	2.8
$N = 100$	218.6	220.5	225.7	0.9	3.2
$N = 200$	436.2	440.9	451.4	1.1	3.5
$N = 300$	653.9	661.4	677.0	1.1	3.5
Star, $N/f = 50, \kappa = 1$					
$f = 3$	286.5	295.1	301.2	3.0	5.1
$f = 6$	575.1	596.5	608.8	3.7	5.9
$f = 9$	864.8	902.1	920.8	4.3	6.5
$f = 12$	1156.	1211.	1236.	4.7	6.9
Globule of a ring					
$U^{(0)} = 6, \kappa = 1$					
$N = 100$	-426.7	-287.2	-211.9	-32.7	-50.3
$N = 200$	-1000.	-722.6	-540.3	-27.7	-46.0

crease with N . This means simply that the values of $\mathcal{D}_{i,i+1}$ match with those from MC quite well. As for the radius of gyration, the disagreement increases slowly with N , i.e., the mean-squared distances $\mathcal{D}_{i,i+k}$ are increasingly more overestimated by GSC as compared to MC for large k . Based on these results, we can determine the swelling exponent of the ring coil by fitting $3\mathcal{R}_g^2$ via Eq.,

$$3\mathcal{R}_g^2 = b^2 N^{2\nu}, \tag{32}$$

using our data in the range $N = 50 - 500$. The resulting pre-factor b^2 and the exponent ν in all three cases are

$$b^2 = \begin{cases} 0.223 \pm 0.005 & \text{MC} \\ 0.180 \pm 0.003 & \text{GSC, } a = 0.9, \\ 0.184 \pm 0.003 & \text{GSC, } a = 1.0 \end{cases} \tag{33}$$

$$\nu = \begin{cases} 0.610 \pm 0.006 & \text{MC} \\ 0.648 \pm 0.002 & \text{GSC, } a = 0.9, \\ 0.651 \pm 0.001 & \text{GSC, } a = 1.0. \end{cases} \tag{34}$$

Thus, the increase in the deviation of $3\mathcal{R}_g^2$ in the GSC method from MC indeed comes entirely from the overestimation of the swelling exponent. This is believed to reach the value $\nu = 2/3 = 0.666 \dots$ asymptotically. However, in the present range of N both GSC and MC give higher apparent exponent values than the most accurate renormalization group result up-to-date,³⁰ $\nu = 0.5882 \pm 0.0011$. The GSC results for ν are only slightly sensitive on the value of a , being in a relatively small overestimation over the MC results. This is related to the Gaussian shape of $g_{ij}^{(2)}$ in the GSC theory, whereas a stretched exponential tail, $\exp(-B\hat{r}^\delta)$ (see the range $r \geq 40$ in Fig. 1), contributes most to $3\mathcal{R}_g^2$ in MC. We may note also that the GSC estimate for ν is fairly close to the result $\nu = 0.635$ from the integral equations approach based on a complicated closure of the Born-Green-Yvon hierarchy in Ref. 20.

When considering open flexible homopolymers ($\kappa = 1, \lambda = 0$) in the good solvent ($U^{(0)} = 0$), the general behavior of $\mathcal{E}^{(int)}$ and $3\mathcal{R}_g^2$ in the third block of Tables I and II is very similar to that of a ring: a good and almost

TABLE II. Comparison of the mean-squared radius of gyration values $3\mathcal{R}_g^2$ (in ℓ^2 units) for different homopolymers from the MC simulation (second column) based on the data set of Ref. 24 and from the GSC theory with the parameter $a=0.9$ (third column) and $a=1.0$ (fourth column). The fifth and sixth columns contain the relative deviation $\delta=(\mathcal{R}_g^2(\text{GSC})/\mathcal{R}_g^2(\text{MC})-1)\times 100\%$ for these two cases given as a percentage.

System	MC	GSC, $a=0.9$	GSC, $a=1.0$	$\delta_{a=0.9}$, %	$\delta_{a=1.0}$, %
Ring, $N=100$					
$\kappa=0.01$	2497.	2503.6	2504.6	0.3	0.3
$\kappa=0.1$	273.57	275.18	277.8	0.6	1.5
$\kappa=0.2$	153.56	157.61	160.81	2.6	4.7
$\kappa=0.5$	82.68	91.04	94.69	10.1	14.5
$\kappa=2.0$	45.98	60.70	64.84	32.0	41.0
Ring, $\kappa=1$					
$N=50$	26.66	28.90	30.32	8.4	13.7
$N=100$	61.45	70.30	74.16	14.4	20.7
$N=200$	141.4	172.3	182.7	21.8	29.2
$N=300$	232.4	292.0	310.3	25.6	33.5
Open, $\kappa=1$					
$N=150$	185.7	221.4	234.3	19.2	26.2
$N=200$	253.5	321.6	341.0	26.9	34.5
Semiflexible ring					
$\lambda=1, \kappa=1$					
$N=50$	37.91	38.18	40.71	0.7	7.4
$N=100$	86.07	92.97	99.44	8.0	15.5
$N=200$	193.6	222.9	239.0	15.1	23.5
$N=300$	305.1	371.5	398.8	21.8	30.7
Semiflexible ring					
$\lambda=5, \kappa=1$					
$N=50$	78.05	60.60	65.58	-22.3	-16.0
$N=100$	210.8	161.0	175.8	-23.6	-16.6
$N=200$	499.8	382.5	419.0	-23.5	-16.2
$N=300$	861.3	622.0	681.5	-27.8	-20.8
Star, $N/f=50, \kappa=1$					
$f=3$	146.2	176.4	186.6	20.6	27.6
$f=6$	185.2	239.0	253.1	29.0	36.7
$f=9$	208.7	278.0	294.5	33.2	41.1
$f=12$	228.7	308.1	326.5	34.7	42.8
Globule of a ring					
$U^{(0)}=6, \kappa=1$					
$N=100$	6.126	6.431	7.384	5.0	20.5
$N=200$	9.368	9.872	11.31	5.4	20.7

N -independent agreement for $\mathcal{E}^{(\text{int})}$ and a slow increase with N in the relative error for $3\mathcal{R}_g^2$. In Fig. 2 we plot the mean-squared distances from the end monomer \mathcal{D}_{0k} versus the chain index k . Up to $k\sim 10$, the agreement of both GSC curves with the MC data is nearly perfect, whereas both GSC curves increasingly overestimate the MC data for larger k . This is consistent with the overestimation of $3\mathcal{R}_g^2$ by the GSC method, dominated by large k contributions. Notably, the effect of changing the parameter a is rather weak on this scale.

Further, let us investigate the effect of increasing the chain stiffness λ from a value corresponding to a fairly flexible ring $\lambda=1$ (see Fig. 3 and the fourth block in Tables I and II) to that of a semistiff ring $\lambda=5$ (see Fig. 4 and the fifth block in Tables I and II). The fairly flexible case gives the energies in a very good agreement with the MC data, albeit the theory with $a=0.9$ is somewhat less accurate accidentally. However, the agreement of $3\mathcal{R}_g^2$ in the GSC theory with MC is even better here than for the corresponding flexible coil. The semiflexible case also gives the energies in a good agreement with the MC data, while $3\mathcal{R}_g^2$ tends to be underestimated by the GSC method. Plots of the mean-

squared distances \mathcal{D}_{0k} versus the chain index k in Figs. 3 and 4 match nearly perfectly up to $k\sim 10$ as well, diverging for larger k . The MC curve is lower (higher) than the GSC curves for $\lambda=1$ ($\lambda=5$) in accord with the tables data. Thus, overall, the increase of $3\mathcal{R}_g^2$ with the chain stiffness λ is more rapid in MC than in the GSC theory. Indeed, conformations of a stiff chain in MC become those of a rigid ring (or rod for an open chain) with increasing λ . However, the Carnahan–Starling equation was deduced in the assumption of a total three-dimensional (3D) isotropy, when its influence would be weaker than in an effective one-dimensional (1D) projection.

Next, let us bring our attention to the effect of changing the topology of the chain. Thus, in the sixth block of Tables I and II we present $\mathcal{E}^{(\text{int})}$ and $3\mathcal{R}_g^2$ for the flexible ($\lambda=0$) stars with the arm length $N/f=50$ in the good solvent ($U^{(0)}=0$). The relative errors in $\mathcal{E}^{(\text{int})}$ and $3\mathcal{R}_g^2$ increase with the number of arms f steadily, again the energy values being more close between MC and GSC. The mean-squared distances \mathcal{D}_{0k} from the core monomer for the largest star with $f=12$ arms are plotted in Fig. 5. Clearly, the agreement be-

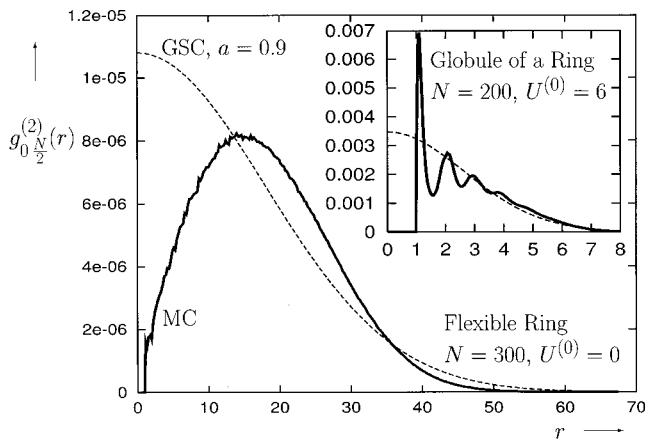


FIG. 1. Plots of the half-ring radial distribution functions $g_{0,N/2}^{(2)}(r)$ (in ℓ^{-3} units) from MC (solid thick lines) and the GSC theory with $a=0.9$ (thin dashed lines) vs the radial separation r (in ℓ units) for the flexible ring homopolymers with $\lambda=0$ and $\kappa=1$. The main part of the figure corresponds to the good solvent $U^{(0)}=0$ and the polymer size $N=300$, whereas the inset corresponds to the poor solvent $U^{(0)}=6$ and the polymer size $N=200$.

tween GSC and MC is the worst of all previously considered cases here. Even the values of \mathcal{D}_{0k} do not match for small k because the core monomer is strongly affected by the very pronounced correlation hole effect in MC.²⁴ However, \mathcal{D}_{ij} between monomers within same arms and away from the core are naturally closer between GSC and MC, just as for the open chain in Fig. 2. Clearly, the divergence of the GSC and MC curves does not increase with k after $k \sim 10$ and the curves have rather similar overall shapes.

It is interesting to analyze the structure of the collapsed globule now. Thus, in Fig. 6 the mean-squared distances \mathcal{D}_{0k} are plotted versus k for a flexible ring homopolymer in the globular state, $U^{(0)}=6$. Overall shapes of the GSC and MC curves are quite similar, reflecting the compactness of the globule. The discrepancy between GSC and MC at small k is present systematically. GSC generally overestimates the values of all \mathcal{D}_{0k} , but the theory with $a=0.9$ manages to come

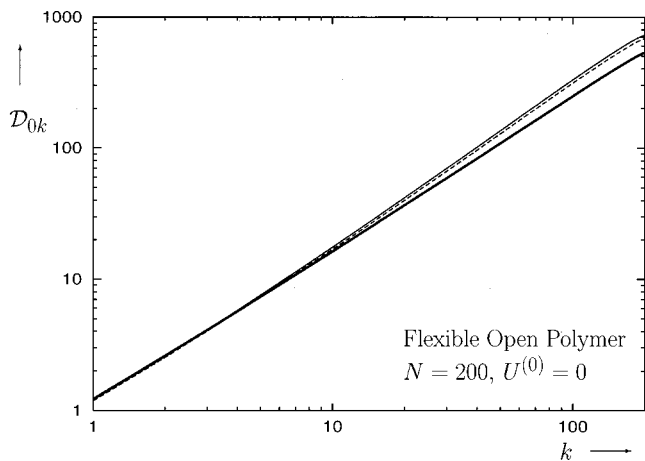


FIG. 2. The mean-squared distances \mathcal{D}_{0k} (in ℓ^2 units) of an open flexible homopolymer with $N=200$, $\lambda=0$, and $\kappa=1$ in the good solvent, $U^{(0)}=0$, vs the chain index k . Here and below the solid thick lines correspond to the MC data, the solid thin lines to the GSC theory with $a=1.0$, and the dashed lines to the GSC theory with $a=0.9$.

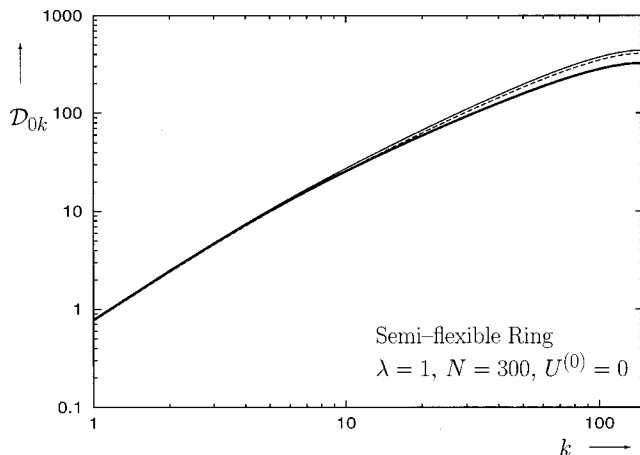


FIG. 3. The mean-squared distances \mathcal{D}_{0k} (in ℓ^2 units) of a fairly flexible, $\lambda=1$, ring homopolymer with $N=300$ and $\kappa=1$ in the good solvent, $U^{(0)}=0$ vs the chain index k .

to a nearly correct limit of the globule size for $k^* \geq 20$ (see Ref. 24), whereas the theory with $a=1.0$ is less accurate. The data for $3\mathcal{R}_g^2$ in the last block of Table II thus show a much better agreement than before, but the energy values in Table I have a significantly larger discrepancy between GSC and MC. The GSC theory noticeably underestimates the negative Lennard-Jones energy contribution between all pairs of monomers. This, however, is to be expected given that the shape of RDF from MC²⁴ has a very tall first liquid-like peak (see the inset of Fig. 1). Since the Lennard-Jones interaction is rather short-ranged, the first peak gives a predominant negative contribution to the mean energy in Eq. (16). As the GSC theory has merely an effective smooth “interpolating” Gaussoid in $g_{ij}^{(2)}$ in that area (see the inset of Fig. 1), the resulting negative energy contribution is significantly smaller in such a theory.

We can also remark that the scaling for the swelling exponent in the globule is correct in the GSC theory. Indeed, the maximal compression is reached when $\eta \lesssim 1$, therefore, by considering $y_{ij} \rightarrow \infty$ in Eqs. (23), (24), and (26) we indeed obtain, $\mathcal{D}_{ij} \sim (r_i^{(0)} + r_j^{(0)})^2 N^{2/3}$.

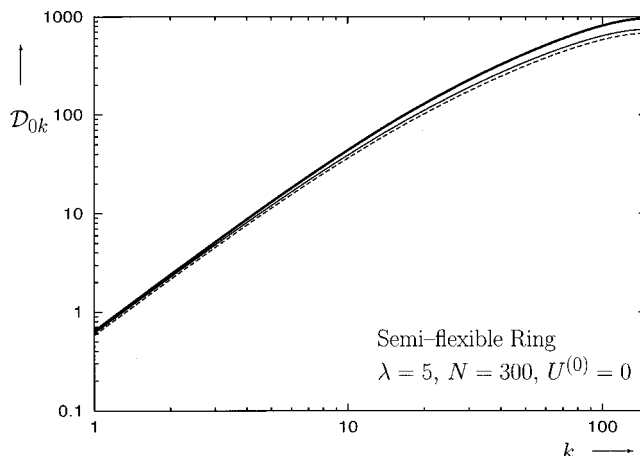


FIG. 4. The mean-squared distances \mathcal{D}_{0k} (in ℓ^2 units) of a semiflexible, $\lambda=5$, ring homopolymer with $N=300$ and $\kappa=1$ in the good solvent, $U^{(0)}=0$ vs the chain index k .

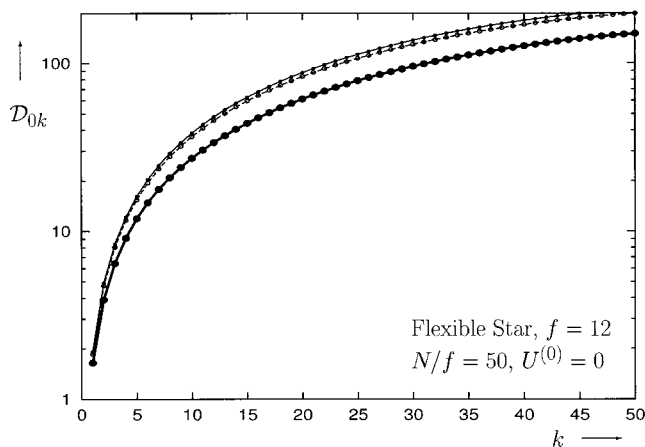


FIG. 5. The mean-squared distances \mathcal{D}_{0k} (in ℓ^2 units) from the core monomer of a flexible, $\lambda=0$, homopolymer star with $f=12$ arms, $\kappa=1$, and the arm length $N/f=50$ in the good solvent, $U^{(0)}=0$ vs the chain index k .

Next, we would like to look at the plots of the mean-squared radius of gyration, $3\mathcal{R}_g^2$, and of the mean energy, $\mathcal{E}^{(int)}$ across the coil-to-globule transition. These are depicted in Figs. 7 and 8, respectively. The following points can be made. First of all, the shapes of these curves are quite similar for MC and GSC with both values of a . Second, the coil-to-globule transition is continuous in all three cases, with the energy slope changing noticeably at around the theta-point. Third, the transition occurs at a somewhat higher value of the attraction constant $U^{(0)}$ in the GSC theory than in the MC simulation. This can be explained by the underestimation of the Lennard-Jones attraction energy in the globule discussed above. Last, $3\mathcal{R}_g^2$ of the globule is partly overestimated by the GSC method with $a=1$, since the value $U^{(0)}=6$ is much closer to the point of the coil-to-globule transition for the theory with $a=1$ than that with $a=0.9$ or MC.

Finally, to understand the N -dependence of the coil-to-globule transition in Fig. 9 we present the plots of the specific energy slope, $N^{-1}d\mathcal{E}^{(int)}/dU^{(0)}$, versus $U^{(0)}$ for the flexible rings of different sizes. These curves nearly coincide in the repulsive coil region, starting to diverge from a value of $U^{(0)} \approx 1.2$. The region of the transition, where the quantity

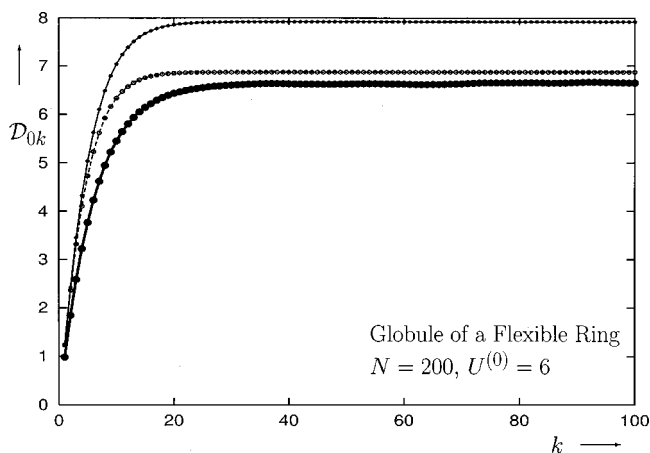


FIG. 6. The mean-squared distances \mathcal{D}_{0k} (in ℓ^2 units) for the globule of a flexible, $\lambda=0$, ring homopolymer with $N=200$, $\kappa=1$, and $U^{(0)}=6$ vs the chain index k .

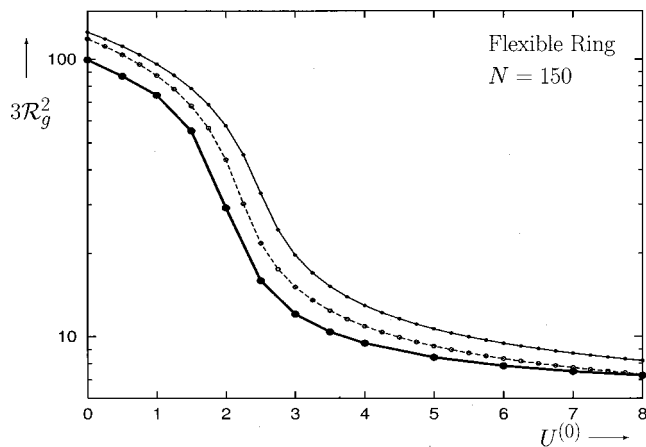


FIG. 7. The mean-squared radius of gyration $3\mathcal{R}_g^2$ (in ℓ^2 units) of a flexible, $\lambda=0$, homopolymer ring with $N=150$ and $\kappa=1$ vs the dimensionless degree of the Lennard-Jones attraction, $U^{(0)}$, across the coil-to-globule transition.

$N^{-1}d\mathcal{E}^{(int)}/dU^{(0)}$ experiences the most dramatic drop, becomes increasingly narrower with increasing polymer size N . Moreover, since the magnitude of the overall change in the specific energy slope also increases with N , the coil-to-globule transition becomes “sharper” with N in the GSC theory, consistent with the MC simulation data and the transition being of second order.^{13,14} Note also, that the theta-point, which we may define, e.g., as the point of the maximal change in $N^{-1}d\mathcal{E}^{(int)}/dU^{(0)}$, shifts towards lower values of $U^{(0)}$ with increasing N .

V. CONCLUSION

In this paper we have developed a version of the Gaussian self-consistent (GSC) technique which does not rely on the virial-like expansion of the Hamiltonian in terms of powers of the density, $\int d\mathbf{r}\rho(\mathbf{r})^L$. As a result, it is now possible to apply the new method to practically any polymer model involving conventional molecular interactions.

Thus, we have been able to compare the mean spatial characteristics and the energy values between the results

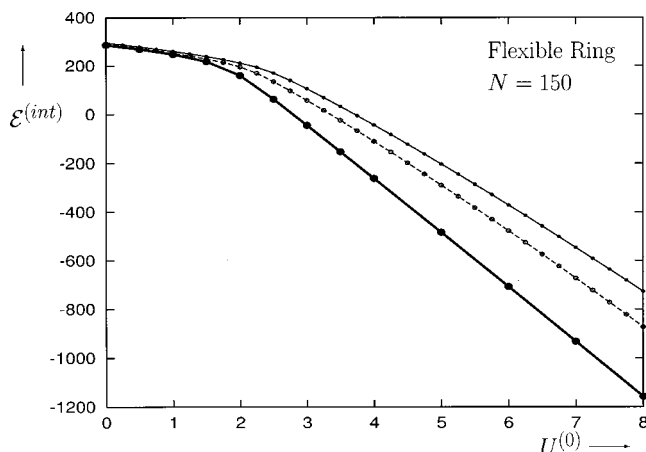


FIG. 8. The mean energy $\mathcal{E}^{(int)}$ (in $k_B T$ units) of a flexible, $\lambda=0$, homopolymer ring with $N=150$ and $\kappa=1$ vs the degree of the Lennard-Jones attraction, $U^{(0)}$, across the coil-to-globule transition.

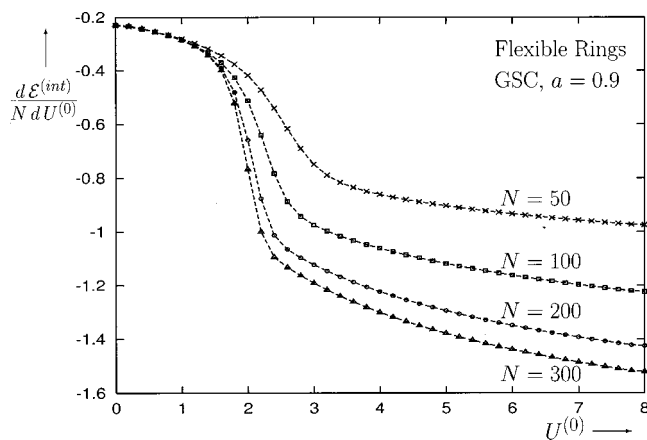


FIG. 9. Plots of the specific energy slope, $N^{-1} d\mathcal{E}^{(int)}/dU^{(0)}$ (in $k_B T$ units) of flexible, $\lambda=0$, homopolymer rings with $\kappa=1$ vs the degree of the Lennard-Jones attraction, $U^{(0)}$, across the coil-to-globule transition for different polymer sizes $N=50, 100, 200, 300$ (from top to bottom). These are obtained from the GSC theory with $a=0.9$.

from the GSC theory and Monte Carlo (MC) simulation based on precisely the same model, which includes the harmonic bonded and the Lennard-Jones pair-wise interactions. This comparison has been performed for three types of macromolecular architectures of an isolated chain: a ring, an open polymer, and a star. We have done this also across the range of the coil-to-globule transition, as well as for different degrees of polymerization and degrees of flexibility. Naturally, the GSC theory agrees with the second order nature of the coil-to-globule transition for flexible homopolymers.

Importantly, the speed of numeric computation is much faster²⁹ in the GSC method than in the equivalent MC simulation for obtaining the same data, particularly so for systems possessing an extra kinematic symmetry, such as for rings or stars.

Overall, the agreement in the shapes of the curves and many of the particular numerical values of observables between GSC and MC is better than one could have anticipated given the relative simplicity of the GSC technique. Where any significant level of deviation does occur, it has been identified as either related to the correlation hole effect at small separations, or to the stretched Gaussian behavior at large separations, in the radial distribution function (RDF).²⁴

In particular, for the repulsive coil, the energy and the mean-squared distances between near monomers along the chain are quite accurate in the GSC theory as compared to the MC data (with a typical deviation of several percent), but the distances between remote monomers, and hence the radius of gyration, are overestimated by the GSC method (with a typical deviation of a few dozen percent). This is a well known drawback of such a theory, related to the overestimation of the Flory swelling exponent for long chains, due to the fact that the RDF here does not have a stretched exponential behavior at large separations. On the contrary, for a rather stiff coil, the GSC theory underestimates the radius of gyration. This may be due to shortcomings of the hard sphere Carnahan–Starling term. For the collapsed globule, on the other hand, the distances and the radius of gyration are quite

accurate in the GSC theory, although the mean energy is less so because of the lack of a sharp liquidlike peak in the RDF.

To make the agreement of GSC and MC better, one has to finally overcome the most restrictive feature of the method—the Gaussian shape of RDF itself. One possible way of doing this is to take a linear superposition of the Gaussian trial functions, thereby permitting “stretching” of the Gaussoid. This should be sufficient for curing the problem with the swelling exponent of the repulsive coil in the GSC theory. Work along these lines is currently in progress. The main difficulties in doing this are in the considerable mathematical complications when calculating the non-Gaussian conformational entropy of the chain, as well as in the added numerical complexity, since a radial mesh for RDF would have to be introduced. Nevertheless, we intend to resolve these issues and hope to present a more accurate, and at last a non-Gaussian self-consistent theory in the near future.

ACKNOWLEDGMENTS

The authors are grateful for interesting discussions by Professor F. Ganazzoli, Professor H. Orland, Dr. G. Raos, Dr. T. Garel, and to R. Connolly for his help. The support of the Enterprise Ireland international collaboration Grant Nos. IC/2001/074 and BC/2001/034, as well as the IRCSET basic research grant SC/02/226 are also acknowledged.

- ¹E. G. Timoshenko, Yu. A. Kuznetsov, and K. A. Dawson, *Phys. Rev. E* **57**, 6801 (1998).
- ²F. Ganazzoli, *J. Chem. Phys.* **108**, 9924 (1998).
- ³F. Ganazzoli, *J. Chem. Phys.* **112**, 1547 (2000).
- ⁴Yu. A. Kuznetsov and E. G. Timoshenko, *J. Chem. Phys.* **111**, 3744 (1999).
- ⁵F. Ganazzoli, R. La Ferla, and G. Allegra, *Macromolecules* **28**, 5285 (1995).
- ⁶R. R. Netz and H. Orland, *Eur. Phys. J.: Appl. Phys.* **8**, 81 (1999).
- ⁷Yu. A. Kuznetsov, E. G. Timoshenko, and K. A. Dawson, *J. Chem. Phys.* **104**, 3338 (1996).
- ⁸E. G. Timoshenko, Yu. A. Kuznetsov, and K. A. Dawson, *J. Chem. Phys.* **102**, 1816 (1995).
- ⁹F. Ganazzoli, Yu. A. Kuznetsov, and E. G. Timoshenko, *Macromol. Theory Simul.* **10**, 325 (2001).
- ¹⁰F. Ganazzoli and R. La Ferla, *J. Chem. Phys.* **113**, 9288 (2000); F. Ganazzoli, R. La Ferla, and G. Raffaini, *Macromolecules* **34**, 4222 (2001).
- ¹¹Yu. A. Kuznetsov and E. G. Timoshenko, *Il Nuovo Cimento* **20D (12bis)**, 2265 (1998).
- ¹²E. G. Timoshenko and Yu. A. Kuznetsov, *J. Chem. Phys.* **112**, 8163 (2000).
- ¹³M. Doi and S. F. Edwards, *The Theory of Polymer Dynamics* (Oxford Science, New York, 1989).
- ¹⁴J. des Cloizeaux and G. Jannink, *Polymers in Solution* (Oxford Science, New York, 1990).
- ¹⁵S. F. Edwards and P. Singh, *J. Chem. Soc., Faraday Trans. 2* **75**, 1001 (1979).
- ¹⁶J. des Cloizeaux, *J. Phys. (France)* **31**, 715 (1970).
- ¹⁷E. Pitard and H. Orland, *Europhys. Lett.* **41**, 467 (1998).
- ¹⁸E. Pitard, *Eur. Phys. J. B* **7**, 665 (1999).
- ¹⁹H. H. Gan and B. C. Eu, *J. Chem. Phys.* **99**, 4084 (1993); **99**, 4103 (1993).
- ²⁰M. P. Taylor and J. E. G. Lipson, *J. Chem. Phys.* **104**, 4835 (1996); **106**, 5181 (1997).
- ²¹G. Allegra and F. Ganazzoli, *J. Chem. Phys.* **83**, 397 (1985).
- ²²G. Allegra, E. Colombo, and F. Ganazzoli, *Macromolecules* **26**, 330 (1993).
- ²³G. Raos, G. Allegra, and F. Ganazzoli, *J. Chem. Phys.* **100**, 7804 (1994).
- ²⁴E. G. Timoshenko, Yu. A. Kuznetsov, and R. Connolly, *J. Chem. Phys.* **116**, 3905 (2002).

- ²⁵J.-P. Hansen and I. R. McDonald, *Theory of Simple Liquids* (Academic, London, 1990).
- ²⁶N. F. Carnahan and K. E. Starling, *J. Chem. Phys.* **51**, 635 (1969).
- ²⁷V. Talanquer and D. W. Oxtoby, *Faraday Discuss.* **112**, 91 (1999); M. Robles and M. L. de Haro, *Phys. Chem. Chem. Phys.* **3**, 5528 (2001).
- ²⁸E. G. Timoshenko and Yu. A. Kuznetsov, *Colloids Surf., A* **190**, 135 (2001).
- ²⁹Benchmarks were performed on an AMD Athlon MP 1900+ with the following run time parameters. For the ring: MC: $S=50\,000$, $\Delta t=8N^2$, resulting in the total run time $t_{\text{total}}\approx 1000$ min; GSC: $n_{\text{iterations}}\approx 5000$, resulting in $t_{\text{total}}\approx 1$ min. For the open chain: MC: $S=50\,000$, $\Delta t=40N^2$, resulting in $t_{\text{total}}\approx 5000$ min; GSC: $n_{\text{iterations}}\approx 18\,000$, resulting in $t_{\text{total}}\approx 25$ min.
- ³⁰R. Guida and J. Zinn-Justin, *J. Phys. A* **31**, 8103 (1998).