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Efimov-DNA Phase diagram: three stranded DNA on a cubic lattice

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We define a generalised model for three-stranded DNA consisting of two chains of one type and a third chain of a different type. The DNA strands are modelled by random walks on the three-dimensional cubic lattice with different interactions between two chains of the same type and two chains of different types. This model may be thought of as a classical analogue of the quantum three-body problem. In the quantum situation it is known that three identical quantum particles will form a triplet with an infinite tower of bound states at the point where any pair of particles would have zero binding energy. The phase diagram is mapped out, and the different phase transitions examined using finite-size scaling. We look particularly at the scaling of the DNA model at the equivalent Efimov point for chains up to 10000 steps in length. We find clear evidence of several bound states in the finite-size scaling. We compare these states with the expected Efimov behaviour.

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I. INTRODUCTION

One of the strange results in quantum mechanics is the formation of an infinite number of bound states in a three-particle system when any two would have given a zero-energy bound state. This result goes by the name of Efimov effect[1–6]. It has recently been argued that the classical analogue of the Efimov effect is the formation of triple-stranded DNA at the melting point of duplex DNA[7]. In this paper we introduce a generalised three-stranded DNA model and examine its phase diagram using the flatPERM Monte Carlo method. Our model consists of a simple extension of the usual Gaussian-Chain model of DNA[8]. In the standard DNA model, the configurations of two identical random walk chains of given length, joined at a common origin, are considered where the only energy comes from base-pairings (common-visited sites) occurring at the same contour length from the common origin along both chains[9].

In our model we label the two chains from the standard model as type **B**, and introduce a third chain, which we label type **A**. We denote the interaction strength by dimensionless variables (i.e. absorbing the factor k_BT into the interaction strength, where k_B is the Boltzmann constant and T is the temperature) ε_1 for a base-pairing interaction between the type **A** chain and either of the type **B** chains, and ε_2 for base-pairing interaction between the two chains of type **B**. These are the only interactions included; there are no three-chain interactions other than those generated between the chains in pairs. The model is shown for strands of length of 14 on the square lattice in Fig 1.

Our purpose in this paper is to examine the full phase diagram for the model defined through the partition function

$$Z_N(\varepsilon_1,\varepsilon_2) = \sum_{\Omega_{3,N}} g_{3,N}(n_1,n_2)\exp(n_1\varepsilon_1 + n_2\varepsilon_2), \tag{1}$$

where we have denoted by $\Omega_{3,N}$ the set of configurations of three random walks of length N. Here, $g_{3,N}(n_1,n_2)$ is the number of configurations with exactly n_1 interactions between the type **A** chain with either of the type **B** chains ($n_1 \in [0,2N]$) and exactly n_2 interactions between the chains of type **B**. The attractive contact energies are taken as ε_1 and ε_2 for **A**-**B** and **B**-**B** pairs. Note that the temperature has been absorbed in the definition of these contact energies, so that $\varepsilon_{1,2}$ are dimensionless. In other words, the contact energies are given by $k_BT\varepsilon_{1,2}$. We also examine the scaling of the free-energy at the three-stranded DNA

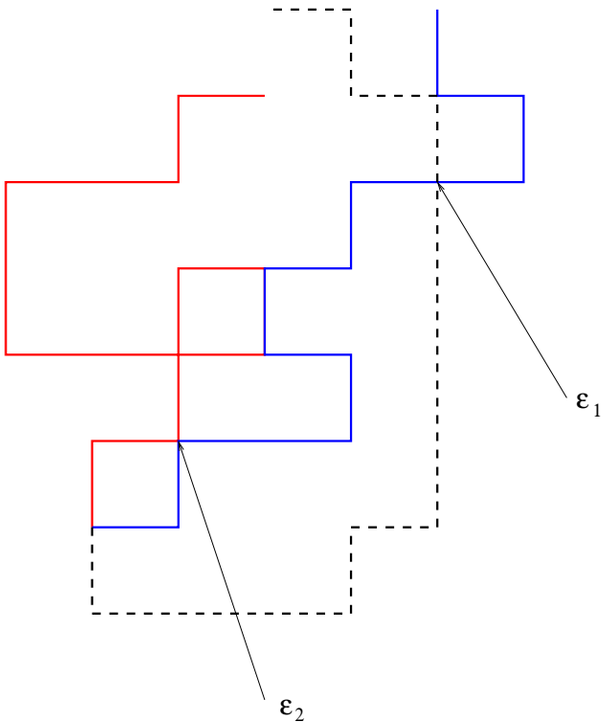


FIG. 1. The type A chain is shown as a dashed line, whilst the type B chain are shown as solid lines. The only two interactions occur at equal contour distances from the common origin. All other contacts do not give rise to interactions.

equivalent to the Efimov point.

The Efimov point is located where the inter-chain interactions are the same, and at a value where any two chains are at the two-chain binding transition.

II. CONNECTION BETWEEN DNA MODEL AND THE EFIMOV EFFECT

The formal connection between DNA melting and the quantum problem can be established as follows[7]. Take three gaussian polymers with native base-pairing interaction, i.e., two monomers on two chains interact if and only if they have the same contour length index

measured from a predetermined end. The Hamiltonian is

$$\mathsf{H} = \frac{H}{k_B T} = \sum_{i=1,2,3} \frac{1}{2} \int_0^N \left(\frac{\partial \mathbf{r}_i(s)}{\partial s} \right)^2 ds + \sum_{i<j} \int_0^N V(\mathbf{r}_i(s) - \mathbf{r}_j(s)) ds, \tag{2}$$

where $\mathbf{r}_i(s)$ is the position coordinate of a monomer (or base) at contour length s. The first term represents the elastic energy or the connectivity of the chain as a polymer, while the second is the interaction between two monomers at the same contour length s (native base pairing of DNA). Like a Hydrogen bond, the range of interaction of V is taken to be small. The partition function is then given by

$$Z = \int \mathcal{D}\mathcal{R} \exp(-\mathsf{H}), \tag{3}$$

where the integral represents the sum over all configurations as a path integral.

If we now do an imaginary transformation $s = it$, then the partition function changes to

$$G = \int \mathcal{D}\mathcal{R} \, \exp(iS), \tag{4a}$$

$$\text{with, } S = \int dt L, \tag{4b}$$

$$\text{and, } L = \sum_{i=1,2,3} \frac{1}{2} \left(\frac{\partial \mathbf{r}_i(t)}{\partial t} \right)^2 - \sum_{i<j} V_{ij}, \tag{4c}$$

as if L represents the Lagrangian of three particles with pairwise interaction $V_{ij} = V(\mathbf{r}_i(t) - \mathbf{r}_j(t))$, with t as real time. In this path-integral representation, G now describes the quantum propagator of three particles with short-range, pairwise interactions. The key point in this exact transformation is the native base pairing of DNA (monomers with the same index) that got translated into the same time interaction in the quantum picture. In the $N \to \infty$ limit, the groundstate energy of the quantum problem maps onto the free energy of the DNA problem.

Double-stranded DNA undergoes a melting transition, as temperature is increased, or as the strength of the potential in Eq. (2) is changed. The melting point corresponds to the critical strength of the potential in the quantum problem above, in which a bound state occurs in a short-range potential in three dimensions. The bound state energy is related to the width a of the wave function, with $E \sim \hbar^2/2ma^2$, so that $E \to 0$ implies $a \to \infty$[10].

At the critical value of zero energy bound state, Efimov argued that three particles will produce a long-range effective interaction $-1/r^2$ which leads to a tower of bound states with energies

$$E_n = \gamma^n E_0, E_0 < 0, \text{ where } \gamma \approx \frac{1}{(22.7)^2}. \tag{5}$$

This is the Efimov effect.

The quantum fluctuations arise from the paths in the classically forbidden regions which are outside the potential well. In the DNA picture, these are the regions on the chains where the hydrogen bonds are broken by thermal fluctuations. A portion of the duplex with broken hydrogen bonds will be called a bubble. The bubbles are characterized by two lengths, $\xi_\parallel$, the fluctuation in the number of bonds broken, and ξ the corresponding length scale for the spatial size, with the relation

$$\xi \sim \xi_\parallel^\nu, \tag{6}$$

where ν is the polymer size exponent. For Gaussian polymers (random walks) $\nu = 1/2$.

The melting transition of the DNA at temperature $T = T_c$, where the hydrogen bonds of the duplex DNA are cooperatively broken, is described by the free energy per unit length

$$f \sim \frac{k_B T_c}{\xi_\parallel}. \tag{7}$$

For a continuous transition, as one finds from exact solutions or from the Poland-Scheraga arguments, we may take $f \sim (|T - T_c|/T_c)^{2-\alpha}$ at least for $T < T_c$, so that $\xi_\parallel \sim (|T - T_c|/T_c)^{\alpha-2}$. This transition, like many other critical points, shows continuous scale invariance in the sense that under a scale transformation $r \to br$, the free energy scales as $f \to b^{-2} f$ for any b.

Let us now consider two strands of DNA — let us call them A, B — separated by a distance R much larger than the hydrogen bond distance so that they do not form any doublet. Now we add a third chain C that can pair with both A and B with the same bond energy. If the temperature is close to the melting point of any pair, the large bubbles will allow C to make contacts with both A and B, resulting in an attraction between the latter two. This fluctuation induced interaction is given by an R-dependent free energy, modifying Eq. (7) to

$$\Delta f \sim -\frac{k_B T_c}{\xi_\parallel} \mathcal{F}(R/\xi), \tag{8}$$

where the scaling function $\mathcal{F}(x)$ should be such that Eq.(8) makes sense for $\xi_{\|}, \xi \to \infty$. By using Eq. (6), we then require $\mathcal{F}(x) \sim x^{-1/\nu}$ so as to cancel $\xi_{\|}$. At the critical point for C, we then get

$$\Delta f(R) \sim -\frac{1}{R^2}, \tag{9}$$

where the gaussian chain value $\nu = 1/2$ has been used.

The above long-ranged inverse-square interaction is at the heart of the Efimov effect, but it is obtained here via the DNA mapping. For DNA, this interaction would lead to a bound phase of three strands at the melting point of the duplex DNA. Consequently, the three chain complex will melt at a temperature higher than T_c.

There are two aspects of the Efimov effect. One is the formation of a three-particle bound state for potentials where two would not have formed a bound state. The second one, more subtle, is the formation of the Efimov tower precisely at the critical potential of zero energy bound-state for a pair, corresponding to the breaking of the continuous scale invariance of the critical point to a discrete scale invariance[11,12].

III. PHASE DIAGRAM

In this section we present numerical results for the phase diagram, where we examine the different transition lines and phases. In order to determine the phase diagram, we have used the flatPERM method[13] to stochastically enumerate (or partially enumerate) the coefficients of the relevant partition functions.

For the full model, we can define the partition function Z_N through:

$$Z_N(\varepsilon_1, \varepsilon_2) = \sum_{\Omega_{3,N}} g_{3,N}(n_1, n_2) \exp(n_1\varepsilon_1 + n_2\varepsilon_2), \tag{10}$$

where we have denoted by $\Omega_{3,N}$ the set of configurations of three random walks of length N. $g_{3,N}(n_1, n_2)$ is the number of configurations with exactly n_1 interactions between the chain of type **A** with either of the type **B** chains ($n_1 \in [0, 2N]$) and exactly n_2 interactions between the type **B** chains.

Having good estimates of $g_{3,N}(n_1, n_2)$ then allows densities and fluctuations in energy to be calculated directly. Suppose we have a quantity $X(n_1, n_2)$, which we also calculate during

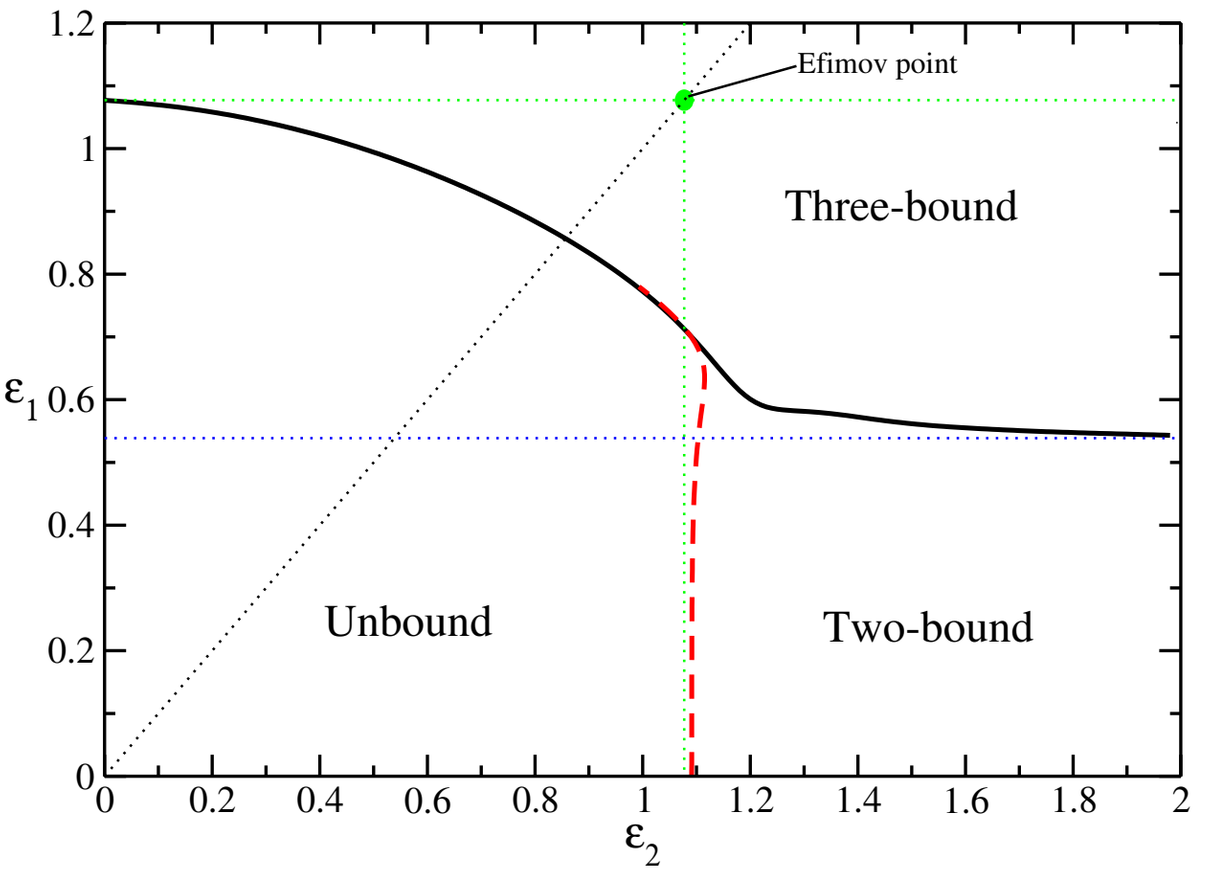


FIG. 2. Phase diagram in the ϵ_1-ϵ_2 plane. The dotted horizontal and vertical lines at $\epsilon_c^{(2)} = 1.07726...$ are the two-chain melting lines. The horizontal line at 0.5357... is the expected transition line for peeling of A from tightly bound BB pair. The solid line represents the transition to the three chain bound state, see text for details on the nature of the transition. The $\epsilon_1 = \epsilon_2$ line meets the two-chain melting lines at the Efimov point (green disk). There is a region where three chains are bound though no two should have been bound. This region is the Efimov-DNA region.

the flatPERM calculation, then the average is calculated:

$$\langle X \rangle = \frac{\sum_{\Omega_{3,N}} X(n_1, n_2) g_{3,N}(n_1, n_2) e^{n_1\varepsilon_1 + n_2\varepsilon_2}}{Z_N(\varepsilon_1, \varepsilon_2)}. \tag{11}$$

In particular, we can calculate the average number of contacts $\langle n_i \rangle$ and the corresponding fluctuations $\Delta_i = \frac{1}{N}\left(\langle n_i^2 \rangle - \langle n_i \rangle^2\right)$, with $i = 1, 2$.

The average number of contacts is expected to scale as[8]

$$\langle n_i \rangle \sim N^{\phi_i}, \tag{12}$$

where $\phi_i = 0$ in the unbound phase and $\phi_i = 1$ in the bound phase, and taking a potentially non-trivial value at the transition. This behaviour enables the setting up of a phenomenological renormalisation group method[16] using the function

$$\varphi_{i,N,N'} = \frac{\log(\langle n_i \rangle_N / \langle n_i \rangle_{N'})}{\log(N/N')}. \tag{13}$$

Estimates for the critical values of ε_1 may be calculated looking for crossings of the $\varphi_{i,N,N'}$ keeping ε_2 fixed (and the other way round). These crossings give estimates of ϕ_i at the transition. Logically, one uses $\varphi_{1,N,N'}$ to calculate the critical values $\varepsilon_1^\star(\varepsilon_2)$ (and vice-versa). The solid black line in the phase diagram in Fig 2 is calculated from $\varphi_{1,N,N/2} = \varphi_{1,N/2,N/4}$ and the red dashed line using $\varphi_{2,N,N/2} = \varphi_{2,N/2,N/4}$ with $N = 200$.

The phase diagram consists of three distinct phase transition lines that join at a multicritical point and separate out three phases: unbound, two-bound and three-bound, corresponding to the number of chains involved in the bound states.

A. Unbound/Two-Bound Phase boundary

When $\epsilon_1 = 0$ the type **A** chain does not interact at all with the two type **B** chains, and the transition as ε_2 is decreased is the standard two chain DNA melting transition at $\varepsilon_2 = \varepsilon_c^{(2)}$, where $\varepsilon_c^{(2)} = 1.07726\cdots$ is the 2-chain binding transition[8].

As ε_1 is increased, we can view the situation of the 2-chain complex adsorbing to the type **A** chain. Whilst the number of contacts is small ($\phi_1 = 0$), the chain type **B** will not affect the two-chain binding transition, and we would expect $\varepsilon_{2,c}(\varepsilon_1) = \varepsilon_c^{(2)}$ to remain constant until the third chain binds at the multicritical point. In Fig 2 the discrepancy between the estimated line (dashed) and expected transition is due to finite size effects. On the transition line we expect $\phi_1 = 0$ and $\phi_2 = 1/2$, which is consistent with the results found for chains up to $N = 200$.

B. Two-bound/Three-bound boundary

As $\varepsilon_2 \to \infty$, the two type **B** chains become tightly bound and behave as one Gaussian chain. Each contact with the type **A** chain is a double contact, and we would thus expect a binding transition when $\varepsilon_1 = \varepsilon_c^{(2)}/2$. As ε_2 is lowered, whilst the two type **B** chains

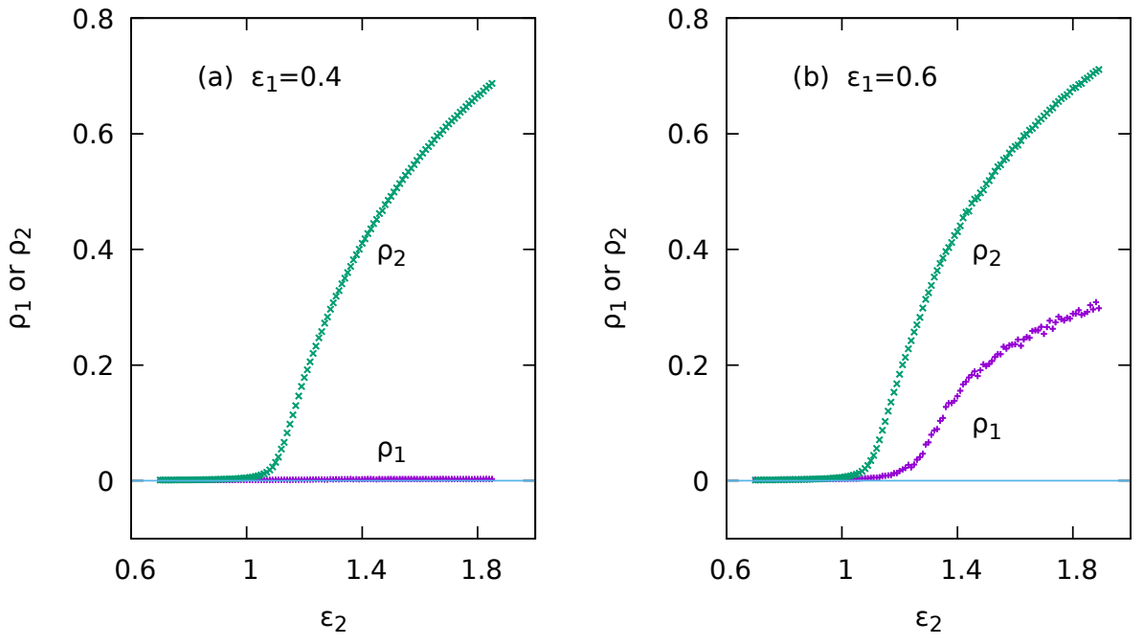


FIG. 3. Plots of ρ_1, ρ_2 vs ϵ_2 for (a) $\epsilon_1 = 0.4$, and (b) $\epsilon_1 = 0.6$ for chains of length 2000. In (a) we see the Melting of BB at $\epsilon_c^{(2)}$ with decreasing ϵ_2, while A remains unbound. In (b) we see different melting points for BB (at $\epsilon_c^{(2)}$), and A-BB triplet.

remain bound, they will start containing bubbles. Now, when the type **A** chain comes into contact with the bound duplex, the number of contacts will sometimes be with one chain and sometimes with 2, making it harder to bind. This will have the effect of elevating the critical temperature, or making it harder to bind, such that $\varepsilon_{1,c}(\varepsilon_2) > \varepsilon_c^{(2)}/2$. As the $\epsilon_2 \to \epsilon_c^{(2)}$, the phase transition line merges with the 2-chain binding transition at the multi-critical point. Along this line we expect $\phi_1 = 1/2$, as this is a standard type binding between two random walks (at least for large ε_2, and we see no evidence of a change in behaviour before the multi-critical point) and $\phi_2 = 1$, since the two type **B** chains are bound. This is also borne out by the numerical results. In Figure 3 we show the plots of ρ_1 and ρ_2 as a function of ϵ_2 for two values of ε_1. When $\varepsilon_1 = 0.4$ (Fig. 3 (a)) we see that the density ρ_1 remains zero, whilst ρ_2 becomes non-zero as the phase boundary is crossed. When $\varepsilon_1 = 0.6$ (Fig. 3 (a)) ρ_1 becomes non-zero later than ρ_2, as we cross successively the unbound/two-bound phase boundary and the two-bound/three-bound phase boundary.

C. Unbound/Three-bound boundary

We first consider the case where $\varepsilon_2 = 0$. When $\varepsilon_2 = 0$, the two type **B** chains do not see each other, they only see the type **A** chain. At the critical interaction $\varepsilon_1 = \varepsilon_c^{(2)}$ each type **B** chain binds with the chain of type **A** independently. This transition can be seen as two independent events. The chain of type **A** binds the other two into a triplex-bound state. This bound state ensures that the two chains of type **B** remain close to each other. The number of interactions n_1 is the sum of the number of interactions with each of the chains of type **B**, and these contacts will be decorrelated between the two chains. It is clear then that $\langle n_1 \rangle \sim N^{1/2}$, giving $\phi_1 = 1/2$.

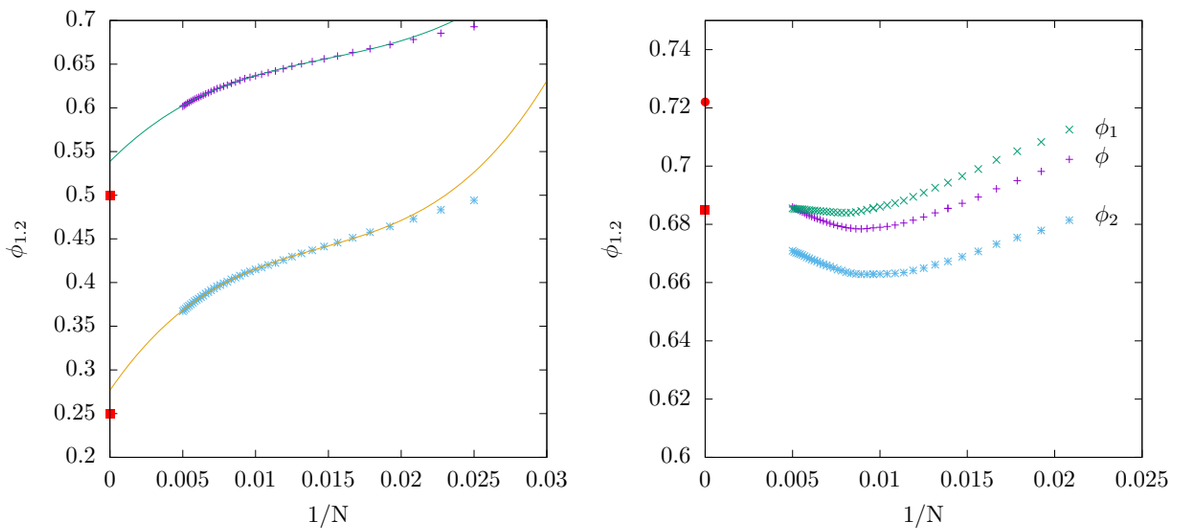
Estimates of ϕ_1 and ϕ_2 are shown in Figure 4, extrapolated by fitting to a quadratic function. It can be seen that ϕ_1 could reasonably extrapolate to 1/2, and ϕ_2 to a non-zero value, possibly 1/4, which is consistent with there being a bound triplet state, where the two type **B** chains are bound through the intermediary of type **A** chain.

We looked at the free energy for the triplet state at this point, and found it to be the same form as the free energy for the two-chain DNA model at its melting point (shown in Fig. 7) but twice as large.

This is interesting, since it is clear that when $\varepsilon_1 = \varepsilon_2$ the three chains are equivalent, and $\phi_1 = \phi_2$, which indicates that the point $\varepsilon_1 = \varepsilon_c^{(2)}, \varepsilon_2 = 0$ is different in nature from the rest of the line. This is understandable, because the type **B** chains are already bound by the type **A** chain, and so a small change in ε could reasonably make a big change. In Fig. 5 we look at the densities of the interactions $\rho_1 = n_1/N$ and $\rho_2 = n_2/N$ as a function of ε_1 for $\varepsilon_2 = 0.6$ and $\varepsilon_2 = 1$. In both cases we can see that the two densities become non-zero at the same time, indicating clearly that the phase above the transition is a triplet phase.

Fig 4 (b) shows ϕ_1 and ϕ_2 for $\varepsilon_2 = 0.7$. The two ϕ seem as if they may reasonably give the same limiting value (around 0.68), which is bigger than 1/2. We compare to the plot with $\varepsilon_1 = \varepsilon_2$ (so $\phi_1 = \phi_2$ by construction). Here we seem to have a different limit, leading to the possibility (unverified) that $\phi_{1,2}$ may vary along the unbound/3-bound phase boundary.

There is another possibility, which is that the line is weakly first order, which is the prediction of studies on hierarchical lattices[7]. It is difficult for the length of chains considered to tell the difference between a smooth variation of the density, or a small jump which might develop only for very long chains.



(a) $\varepsilon_2 = 0$ (b) $\varepsilon_2 = 0.7$ and $\varepsilon_1 = \varepsilon_2$

FIG. 4. In figure (a) we show plots of ϕ_1 and ϕ_2 for $N \leq 200$ plotted against $1/N$. Figure (b) shows ϕ_1 and ϕ_2 for $\varepsilon_2 = 0.7$ and $\phi = \phi_1 = \phi_2$ for $\varepsilon_1 = \varepsilon_2$.

As ε_2 is increased, the type **B** chains will tend to bind more, which has the effect of making it easier for the type **A** chain to bind, which lowers the value of ε_1 required to maintain the triple-bound state. Along the whole of this phase transition line, the two type **B** chains are nevertheless held together by the action of the the type **A** chain. This stops when $\varepsilon_2 = \varepsilon_c^{(2)}$, and the **B**-chains can bind in their own right. This occurs at the multi-critical point.

D. The multi-critical point

The multi-critical point location can be identified by looking at the crossings for φ_i defined in Eq. 13 along the line $\varepsilon_2 = \varepsilon_c^{(2)} = 1.07726\cdots$. The value of $\phi_1 = 0$ is expected along this line until the multicritical point, where it will be expected to take on a new value, indicating the adsorption of the type **A** chain to the type **B** chains. Likewise, the value $\phi_2 = 1/2$ is expected along this line, but may or may not take on a new value at the multicritical point. In Fig. 6 we show plots of $\varphi_{1,2}$ against ε_1 showing crossings at the estimated location of the multicritical point. We show $\varphi_{1/2,N,N/2}$ for chains chains of lengths from $N = 20$ to $N = 200$ in steps of 20. In the same figure we show the variation of the estimate for the value of ε_1

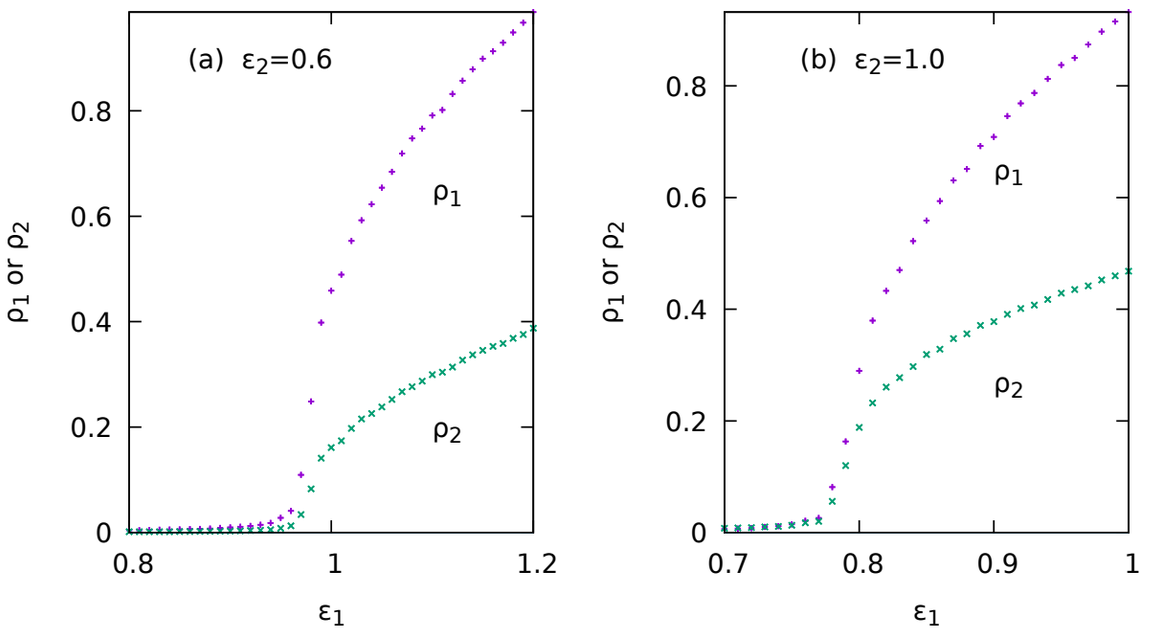


FIG. 5. Efimov-DNA. Plots of ρ_1, ρ_2 vs ϵ_1 for (a) $\epsilon_2 = 0.6$, and (b) $\epsilon_2 = 1.0$ for chains of length 2000. We see both BB and AB pairings at the same $\epsilon_1 < \epsilon_c^{(2)}$ for $\epsilon_2 = 0.6$ (a) and $\epsilon_2 = 1.0$ (b). The transition is between the unbound and the ABB triplet phases. The transition takes place in the region where any pair would have been in the unbound phase.

at the multi-critical point, which we estimate to be at $\varepsilon_1 = 0.71(5), \varepsilon_2 = 1.07726....$

IV. FINITE-SIZE SCALING

The distinctive feature of the Efimov effect is the occurrence of the Efimov constant that determines the geometric scaling of the energy levels, viz., γ in Eq. (5). Although γ is not universal, it is still a characteristic number for the effect, and the value Efimov determined for fermions is $\gamma = (22.7)^{-2}$. The analogy with DNA seems to provide a different way of having analogue behaviour, using a polymer-based Monte Carlo approach, in particular here we use the flatPERM method introduced by Prellberg and Krawczyk[13].

For this purpose, we evaluated the free energy for three chains with the same interaction $\epsilon_1 = \epsilon_2 = \epsilon_c^{(2)}$ so that any pair would be at its critical point. The free energy has been evaluated for lengths up to 10,000.

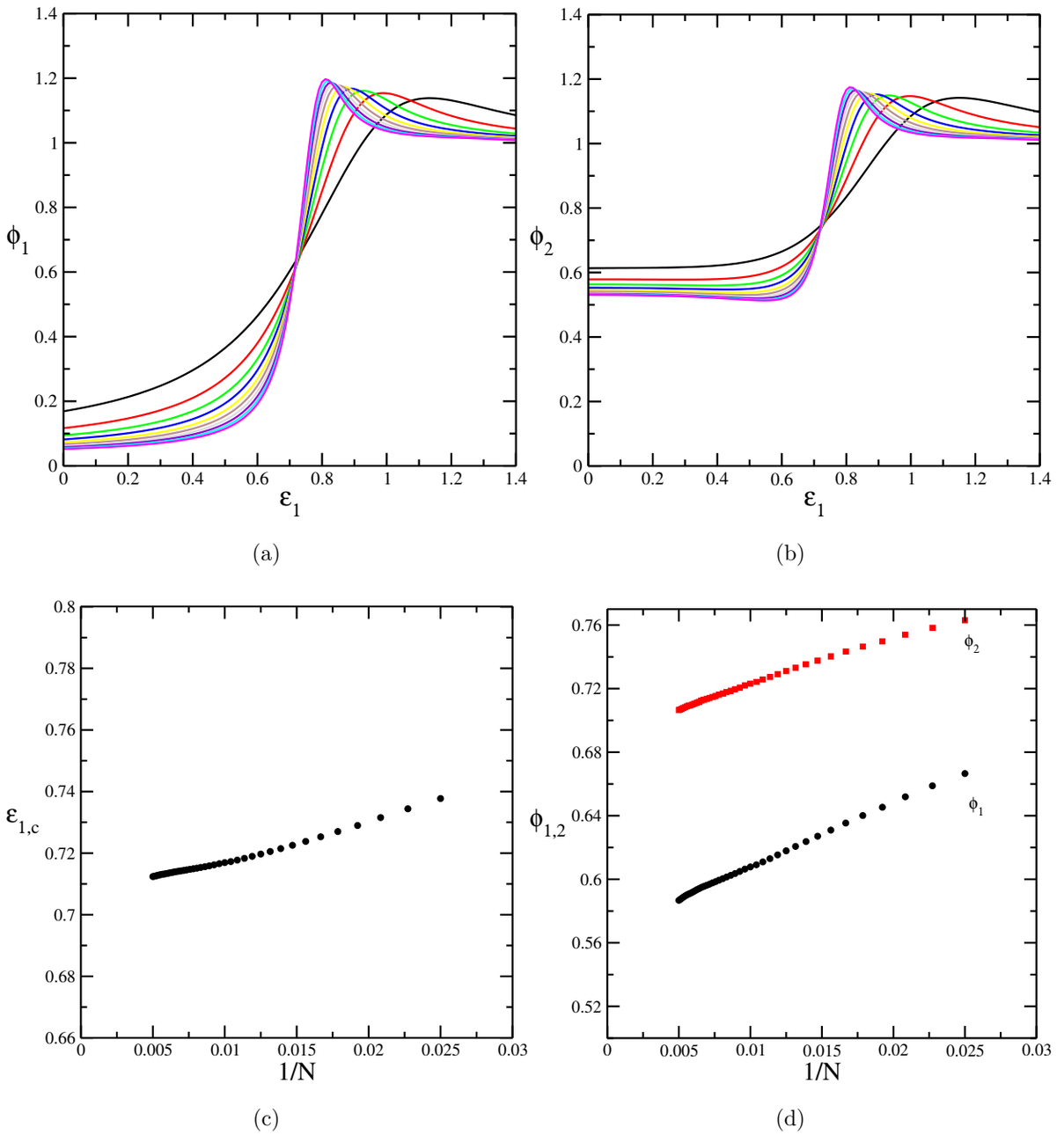


FIG. 6. Figures (a) and (b) show plots of $\varphi_{1/2}$ vs ε_1. The crossings give finite-size estimates of the critical coupling. Figure (c) shows these estimates plotted against $1/N$ and Figure (d) shows estimates of the ϕ values.

To test the quality of the numerical data obtained from PERM, we check the nature at $\epsilon = \epsilon_c^{(2)} = 1.07726$. In fact, the simulations done at this point will not give critical point data, as there is still a small shift $\delta T \neq 0$ due to finite size effects.

Let us first derive the expected two-chain and three-chain scaling behaviour in order to compare with our flatPERM data.

For a contact interaction, $V(\mathbf{r}_i(t) - \mathbf{r}_j(t)) = v_0 \delta(\mathbf{r}_i(t) - \mathbf{r}_j(t)), v_0 < 0$ in Eq. (2), standard dimensional analysis tells us $[s] = [L]^2$, and $[v_0] = [L]^{d-2}$, where $[L]$ denotes the dimension of length. For $d > 2$, the two chain melting is described by a Renormalization group fixed point $u^* = 2\pi\varepsilon$, where $\varepsilon = 2 - d$ and u is the renormalized dimensionless coupling constant with the bare value $u_0 = v_0 L^{\varepsilon}$. In $d = 3$, the melting point is the unstable fixed point $u^* = -2\pi$ ($\epsilon = -1$). At this fixed point, we associate the exponent for length ξ as

$$\xi \sim |\Delta T|^{-\phi}, \quad \phi = \frac{1}{|\varepsilon|}, \tag{14}$$

where $\Delta T = \frac{v_0 - v_c}{v_c}$ is the deviation from the melting point, so that in $d = 3$,

$$\xi_{\|} \sim |\Delta T|^{-2}. \tag{15}$$

Here ξ is the length in space while $\xi_{\|}$ is along the chain (number of bases).

The partition function is that of two or three interacting chains, free at one end but tied together at the other. This configuration goes by the name of "survival" partition functions of "vicious walkers". At the unstable fixed point, the behaviour of the finite length p-chain partition function is of the form

$$Z_N^{(p)} \sim \mu^N N^{-\psi_p}, \tag{16}$$

where the exponent ψ_p is given by[14]

$$\psi_p = \eta_p/2, \tag{17}$$

$$\eta_p = \begin{cases} \varepsilon & \text{for } p = 2, \text{(exact)} \\ 3\varepsilon + 3\ln(3/4)\,\varepsilon^2 + O(\varepsilon^3) & \text{for } p = 3, \end{cases} . \tag{18}$$

We now use it for $d = 3$, i.e., $\varepsilon = -1$. At this fixed point, we find $\psi_2 = -1/2$, for two chains at the melting point. For the critical contribution to the three-chain free energy, a direct sum gives $\psi_3 = -1.9315$ which we approximate as -2.

Combining Eqs. (15) and (18), the scaling for a small δT is (see also Ref[8])

$$Z_N^{(2)} \sim (2d)^{2N} N^{1/2}\, \mathcal{G}(\delta T\, N^{1/2}), \tag{19}$$

For a small fixed δT, $\mathcal{G}(x) \approx a' + b'x$. Therefore, with $\mu = 2d$ (for a random walk on a d-dimensional cubic lattice), we have

$$F_N^{(2)} \equiv \ln\left(\frac{Z_N^{(2)}}{\mu^{2N}}\right) \approx a + |\psi_2| \ln N + bN^{1/2}. \tag{20}$$

For the three chain case, the critical point contribution to the free energy

$$F_N^{(3)} \equiv \ln\left(\frac{Z_N^{(3)}}{\mu^{3N}}\right) \approx a_3 + |\psi_3| \ln N + b_3 N^{1/2}, \tag{21}$$

without the Efimov effect. The Efimov tower (Eq. (5)) would lead to a different class of terms of the type

$$F_{\text{efi}} \sim N|E_0| + \qquad c_j e^{-Nk_j}, \tag{22}$$

where k_j's come from the energy gaps of the tower. The absolute sign for E_0 is required because we are actually writing the expression for $\ln Z$. Combining the two different contributions we have a form

$$F_N^{(3)} = f_3 N + |\psi_3| \ln N + b_3 N^{1/2} + a_3 + \qquad c_j \exp(-Nk_j). \tag{23}$$

Note that, even though $F_N^{(2)}$ has no $O(N)$ term, a linear term in $F_N^{(3)}$ with $f_3 > 0$ is a signature of the Efimov effect.

We use this RG based formula to fit numerically calculated free energy.

V. COMPARISON WITH FLATPERM DATA

We determined the free energies of the 2-chain and the 3-chain systems at the two chain melting point $\epsilon_c^{(2)} = 1.07726$) for lengths up to 10,000 for 10^8 iterations

Fig. 7 shows a good fit of Eq. 3 to the critical point data, with parameters as noted in the caption. The good fit is an assurance of the good quality of data.

Armed with the success for two chains, we try to fit to equation 23. The best fit to the data was provided by the following form:

$$F_3 = f_3 N + 2\ln N + b_3 N^{0.5} + a_3, \tag{24}$$

$$\delta F = c_0 e^{-kN} + c_1 e^{-\gamma kN} + c_2 e^{-\gamma^2 kN} + c_3 e^{-\gamma^3 kN}. \tag{25}$$

with f_3, a_3, b_3 and the c_i as fitting parameters to the free energy data for three chains at ε_c.

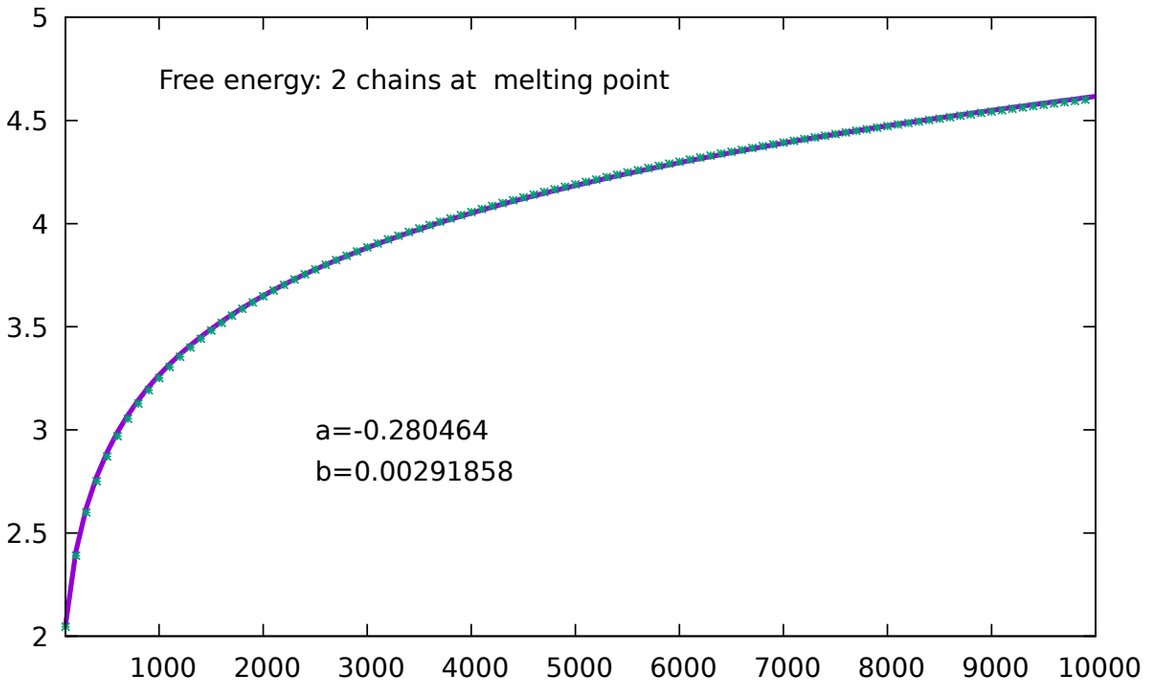


FIG. 7. Length dependence of the critical point free energy for two chains. RG functional form, Eq. 20, (solid line) fits the data from PERM evaluated at $\epsilon = 1.07726$ (blue dots), with $a = 0.280464, b = 0.00292$.

A fit to F_3, given by Eq. 24, with reduced $\chi^2 = 0.053$ gave $f_3 = 0.26182, b_3 = -0.14524, a_3 = -7.9429$ with errors in the last digit, as estimated by the fitting program of GNUPLOT. Fitting to $F_3 + \delta F$ (Eqs. 24, 25) gives a fit with the parameters reported in Table I. The two fits are shown in Fig. 8. To see the difference in fit, we need to look at short chain lengths, and the exponential terms are required to ensure a good fit with a $\gamma \approx 0.107$.

f_3	b_3	a_3	c_0	c_1
0.261660	-0.121678	-8.781304	4.012730	2.78526
c_2	c_3	γ	k	
4.165424	2.727346	0.107447	2.036299	

TABLE I. Best fit parameters, which lead to the fit shown in Fig 8.

Whilst caution should always be taken to not over-interpret a numerical fit, this fit seems

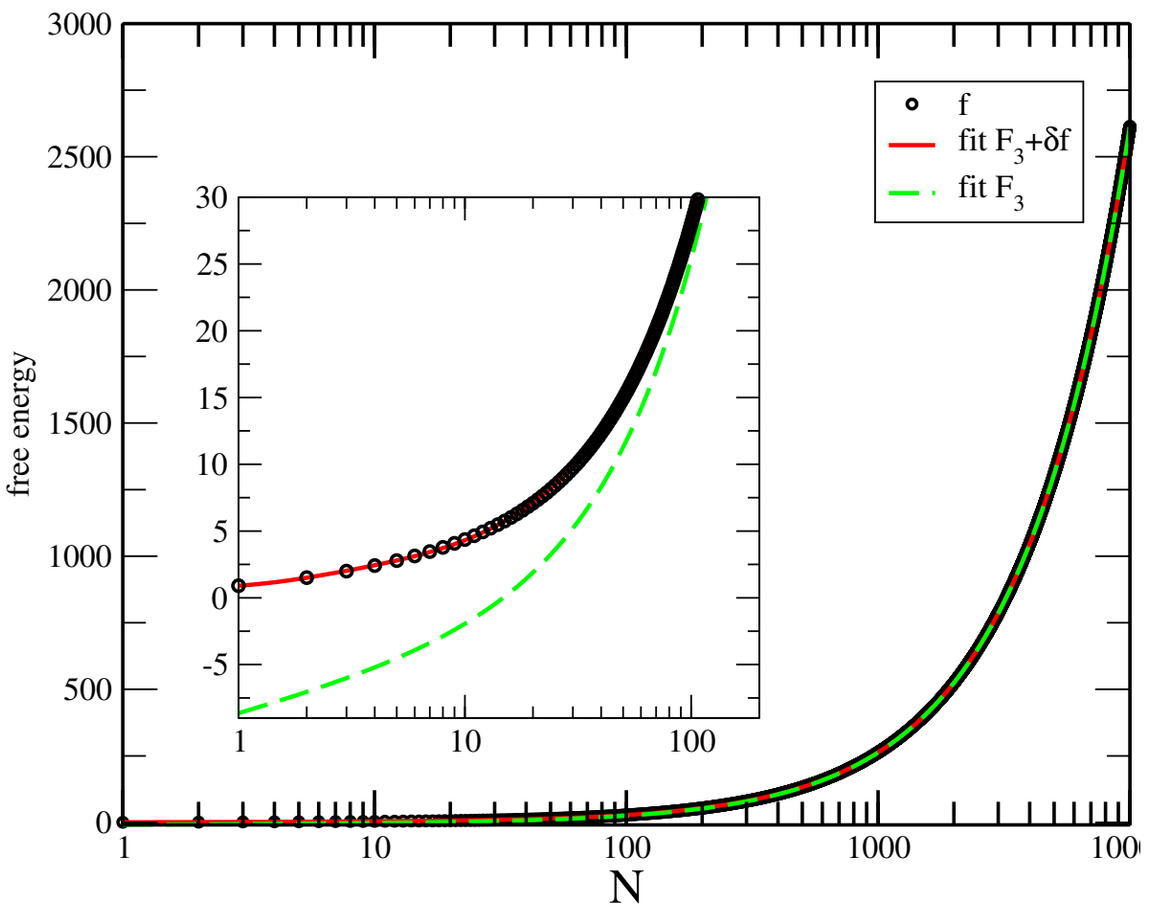


FIG. 8. Length dependence of 3-chain free energy. The circles are the finite-size data points obtained for the free energy. The dashed (green) line is a fitting using Eq. (24) whilst the solid (red) line is found by fitting to $F_3 + \delta F$.

to indicate convincingly the existence of at least three bound states, but the energy gaps, determined by $\delta E_i = \gamma^i k$ does not fit the Efimov tower prediction, where we would have expected $\delta E_i = (1 - \gamma^i) f_3$, so whilst we confirm that we have multiple bound states in the three-bound phase, where any two chains would not be bound, we do not confirm the Efimov tower. This could be either an indication that the behaviour is different, or rather the finite walk aspect of our investigation does not allow us to see all the possible bound states. We will pick up this point in the discussion.

VI. DISCUSSION

In this article we presented results for the DNA analogue of Efimov-type scaling consisting of three-stranded DNA modelled by three random walkers starting from a common origin, and interacting only when two walks meet an equal number of steps from the origin, and the generalised model is studied for its phase diagram in ε_1 and ε_2.

At the equivalent point to the Efimov point ($\varepsilon_1 = \varepsilon_2 = 1.07726$) we find three bound states, and excellent fitting to the free energy from length 0 to 10000, but these states don't follow the Efimov tower structure expected. Whilst the different energy states give the finite-size scaling behaviour of our system, it is not guaranteed that all will contribute. In the quantum system, our calculation would be equivalent to looking at the short-time evolution of the particles from a specific initial state, rather than a study of the time-evolution operator in general, which would be stationary in time. In the quantum system, we are looking at the eigenvalues of the time evolution operator, which in our case would correspond to a transfer matrix which adds a step to each of the DNA molecules. This can be seen as three interacting partially directed walks in 3+1 dimensions. In a future work we will look at this transfer matrix to see if it is possible to extract the eigenvalue structure directly.

DATA AVAILABILITY

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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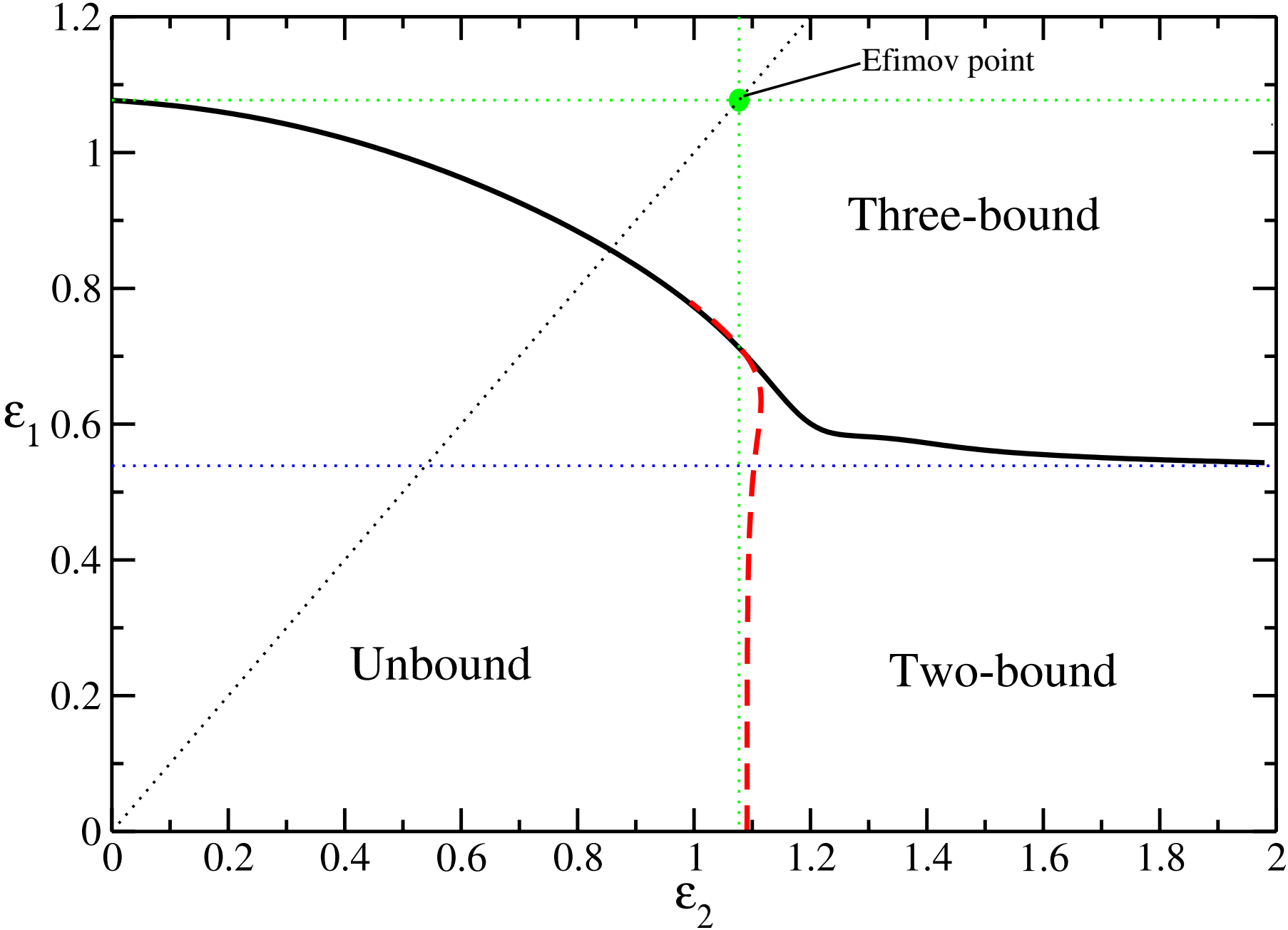
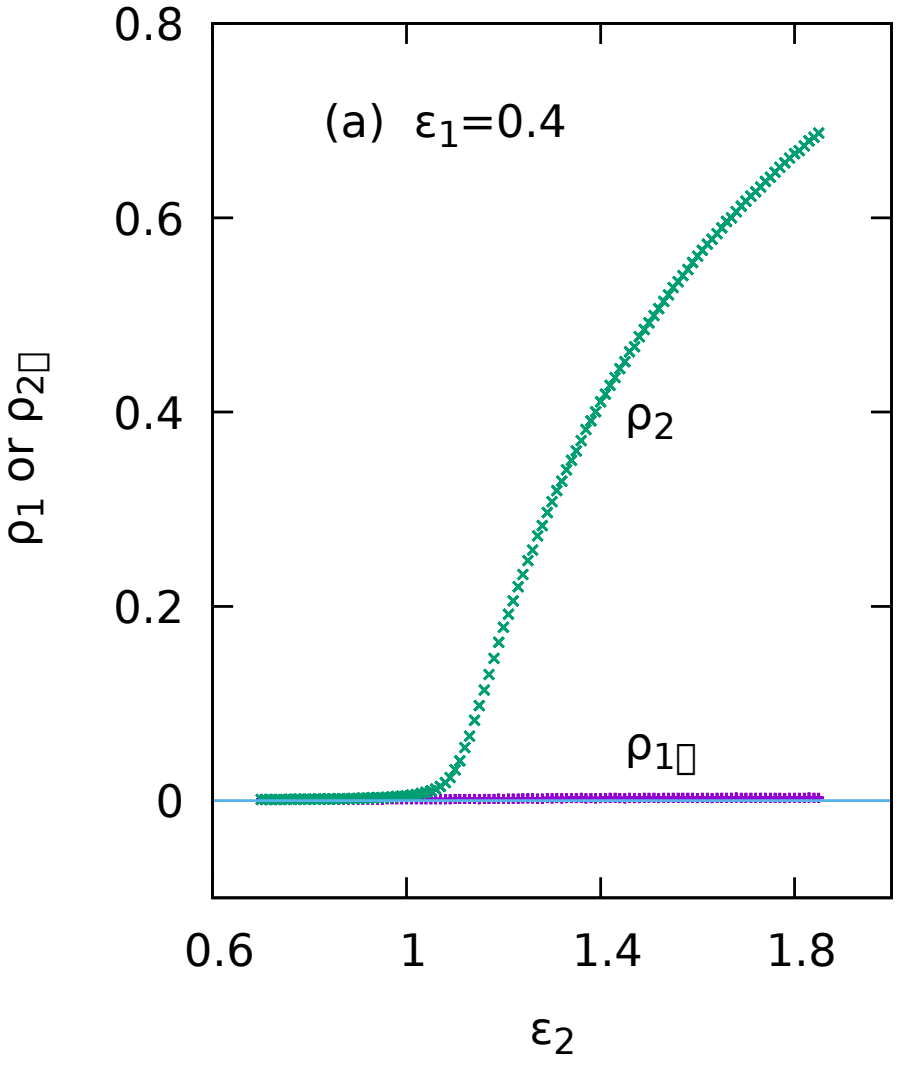
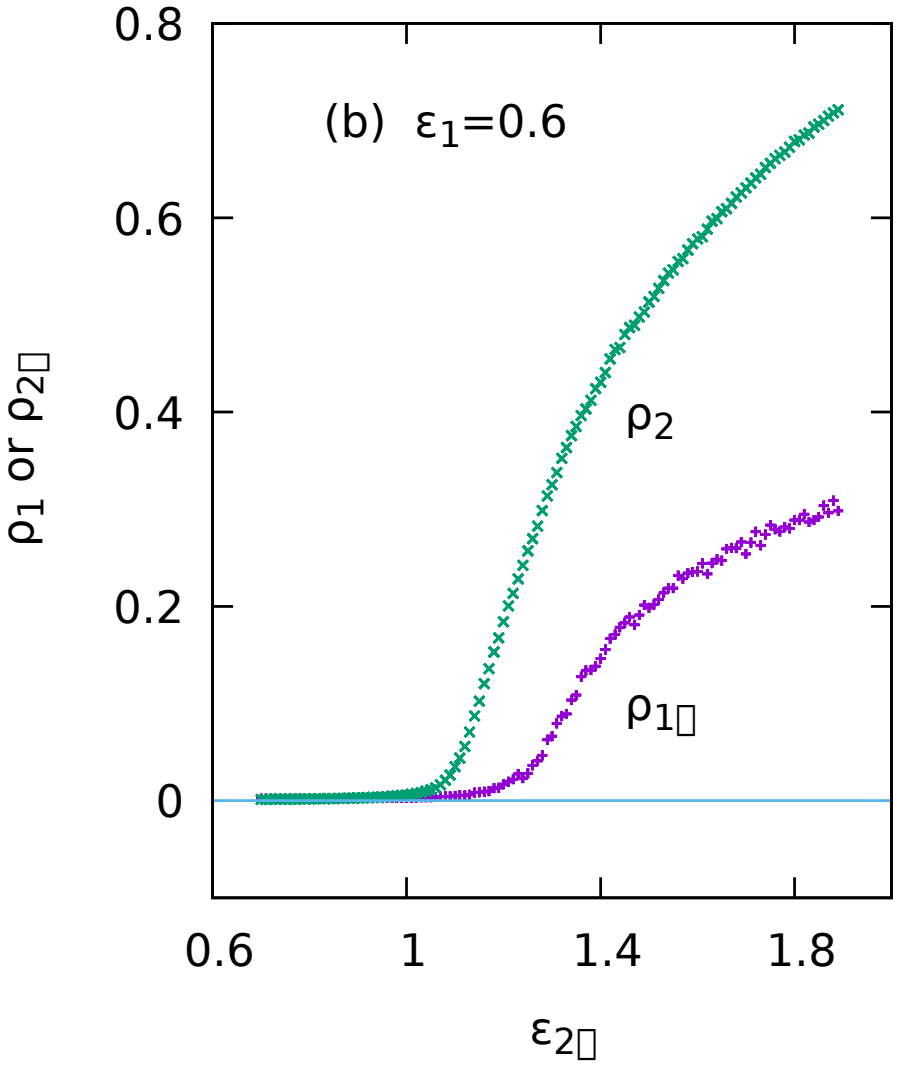
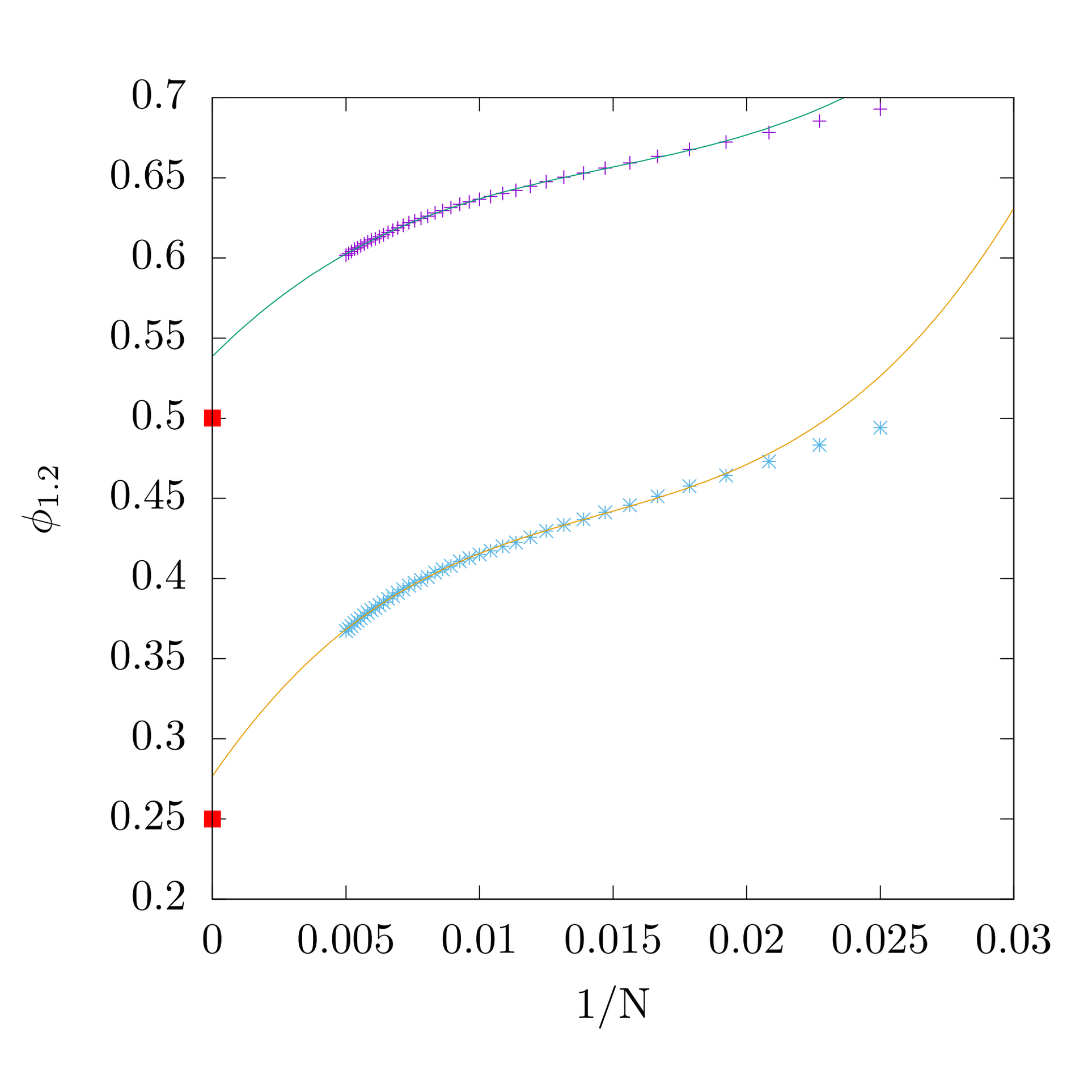
ε_1

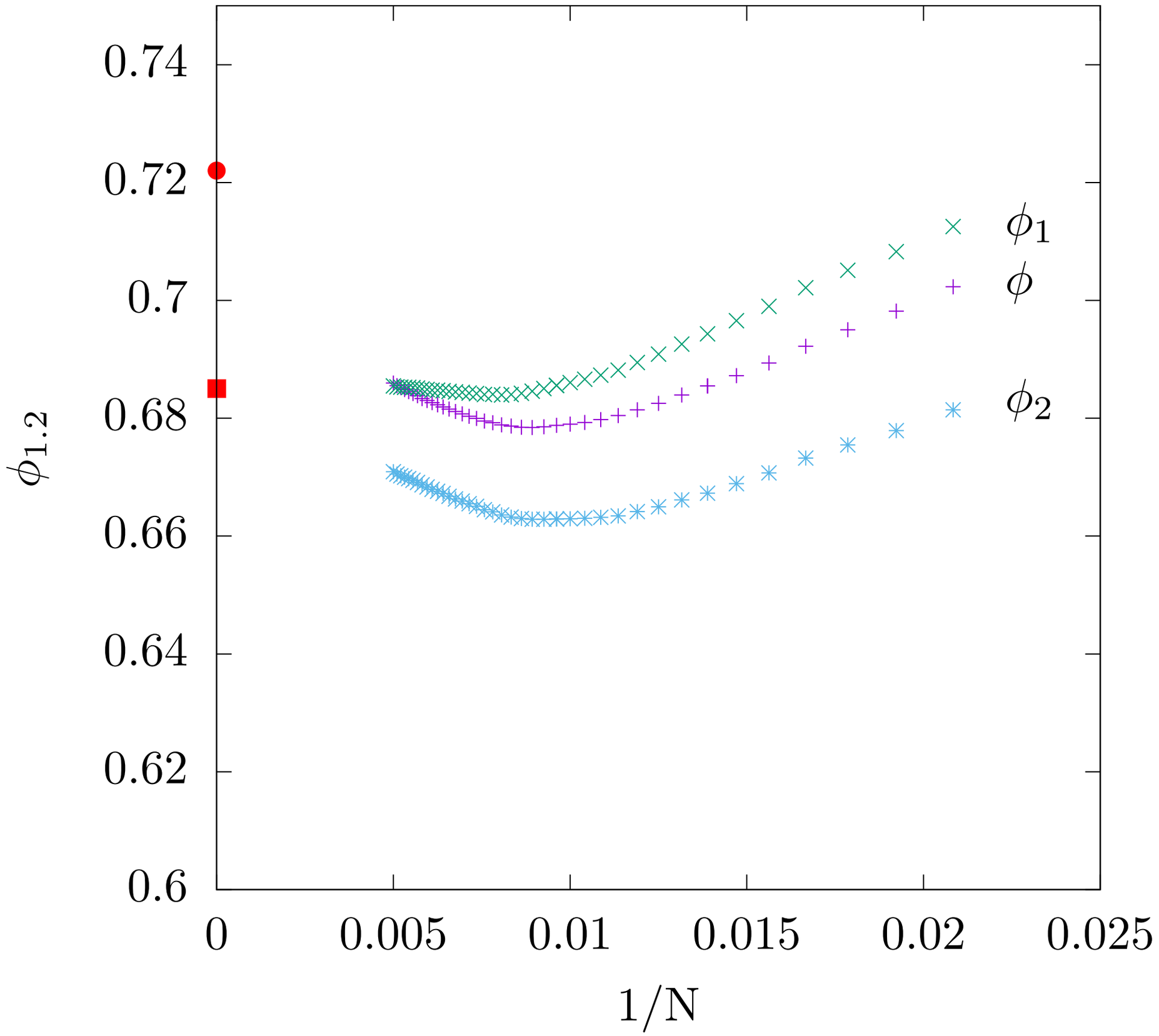
ε_2

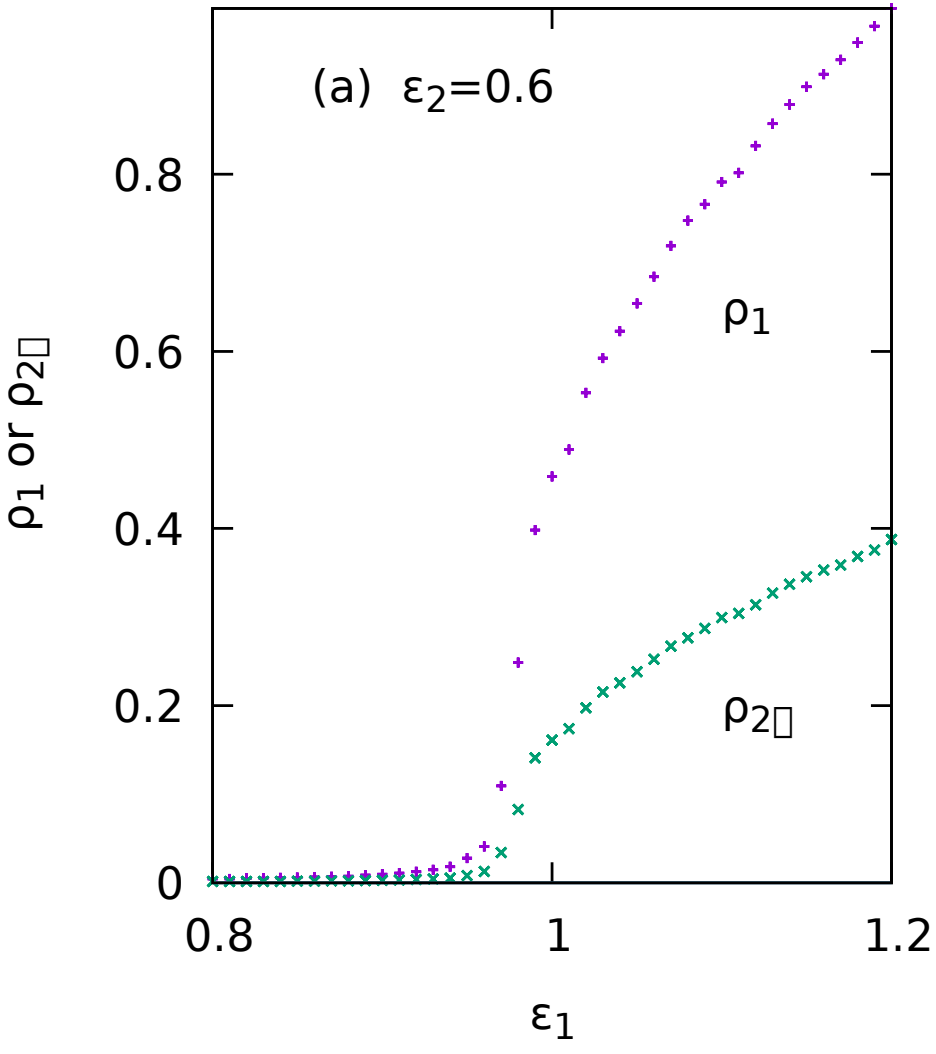

Efimov point
Three-bound
Unbound
Two-bound
ε_1
ε_2
0
0.2
0.4
0.6
0.8
1
1.2
1.4
1.6
1.8
2

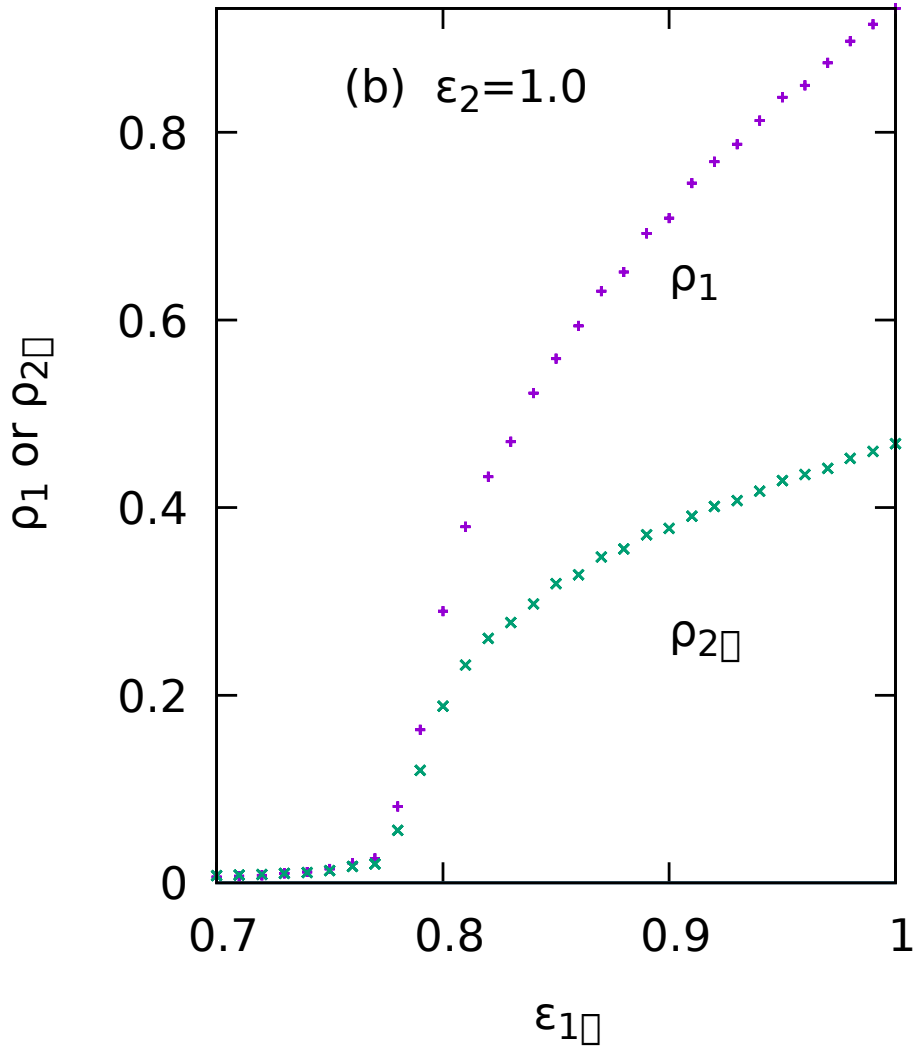

(a) ε1=0.4
ρ2
ρ1□
0.8
0.6
0.4
0.2
0
0.6
1
1.4
1.8
ε2
ρ1 or ρ2□

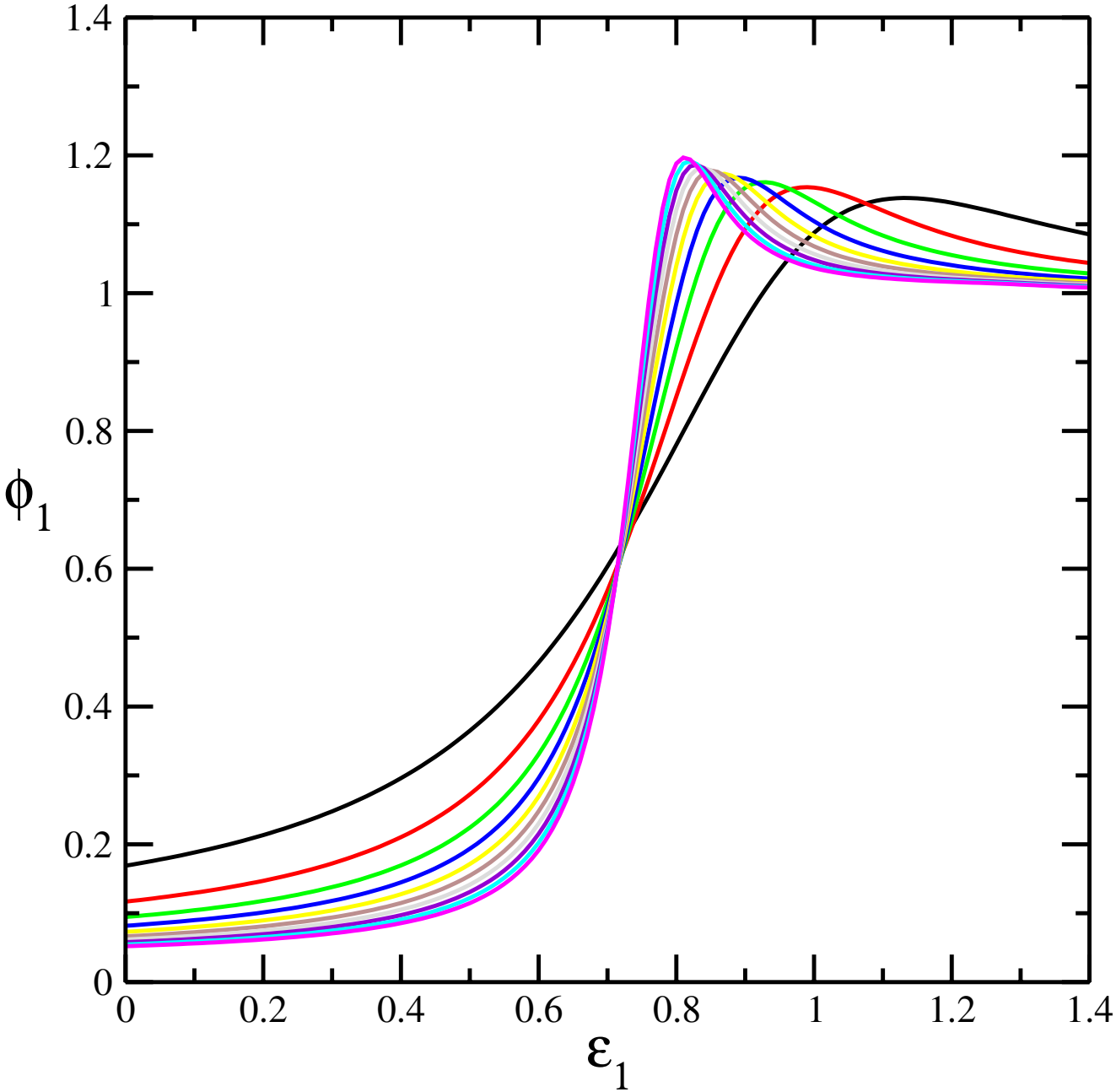

(b) ε1=0.6
ρ2
ρ1□
0.8
0.6
0.4
0.2
0
0.6
1
1.4
1.8
ε2□
ρ1 or ρ2□

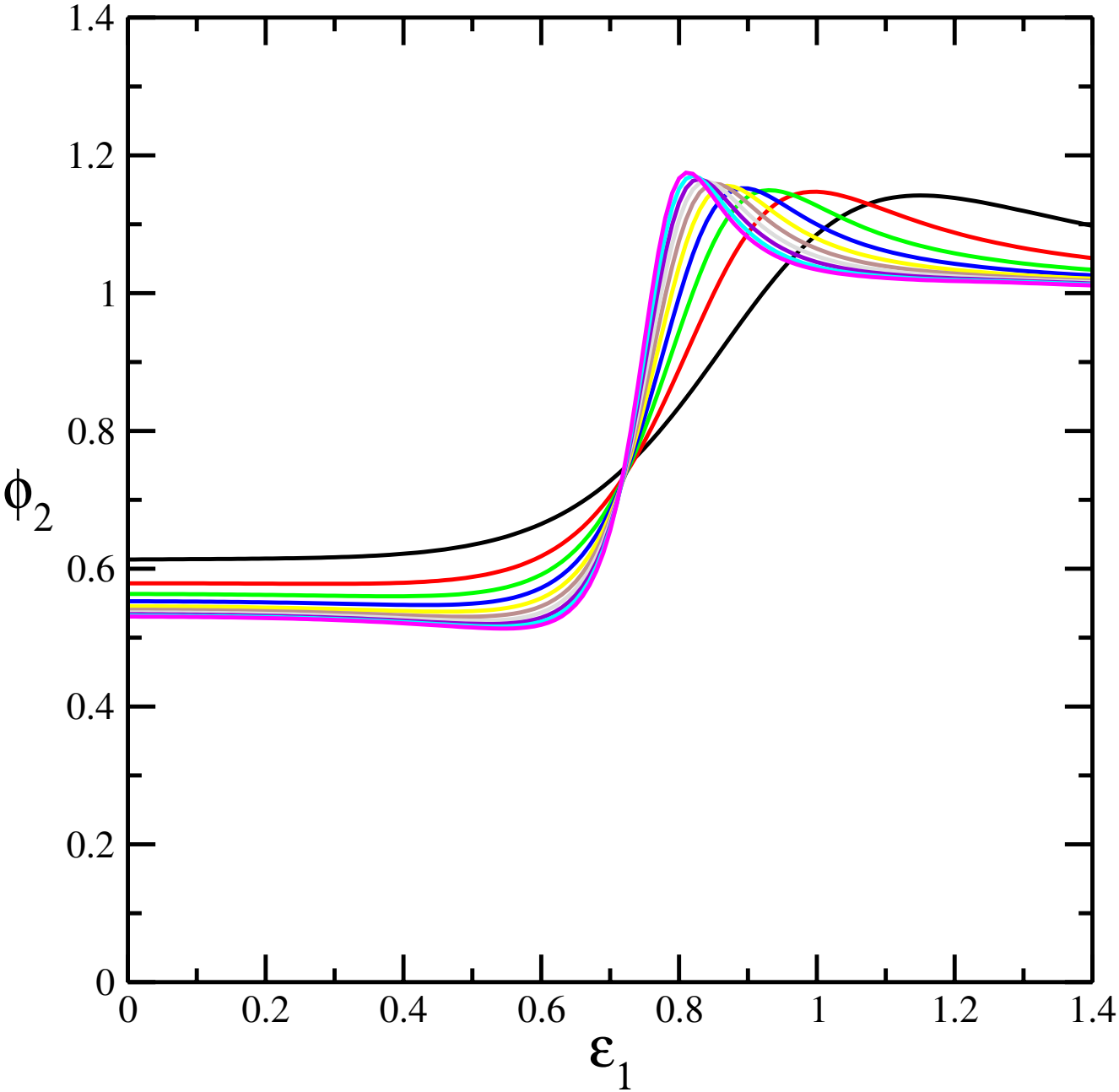

0.7
0.65
0.6
0.55
0.5
0.45
0.4
0.35
0.3
0.25
0.2
$\phi_{1.2}$
0
0.005
0.01
0.015
0.02
0.025
0.03
1/N

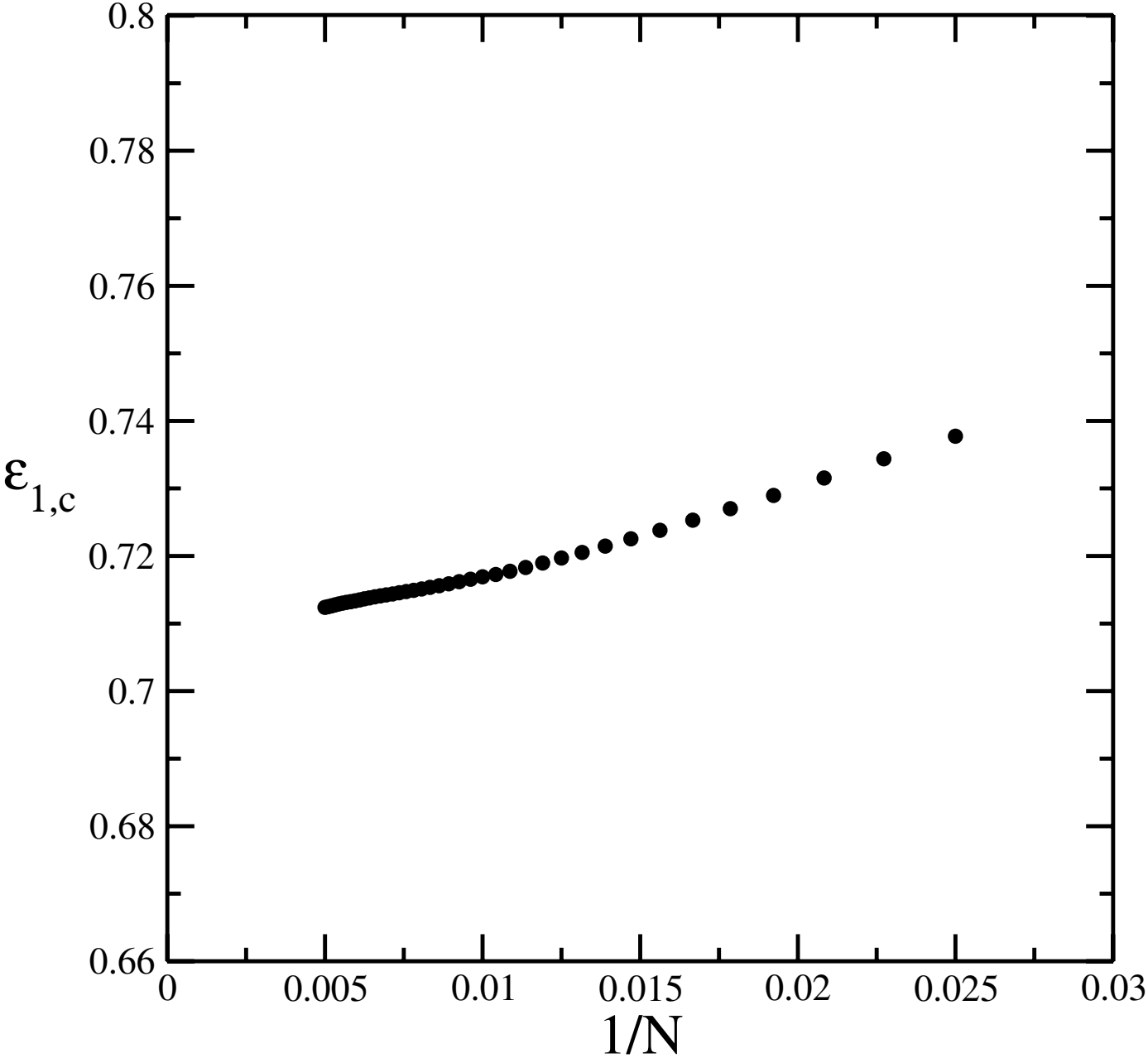

0.74
0.72
0.7
0.68
0.66
0.64
0.62
0.6
$\phi_{1,2}$
0
0.005
0.01
0.015
0.02
0.025
1/N
ϕ_1
ϕ
ϕ_2

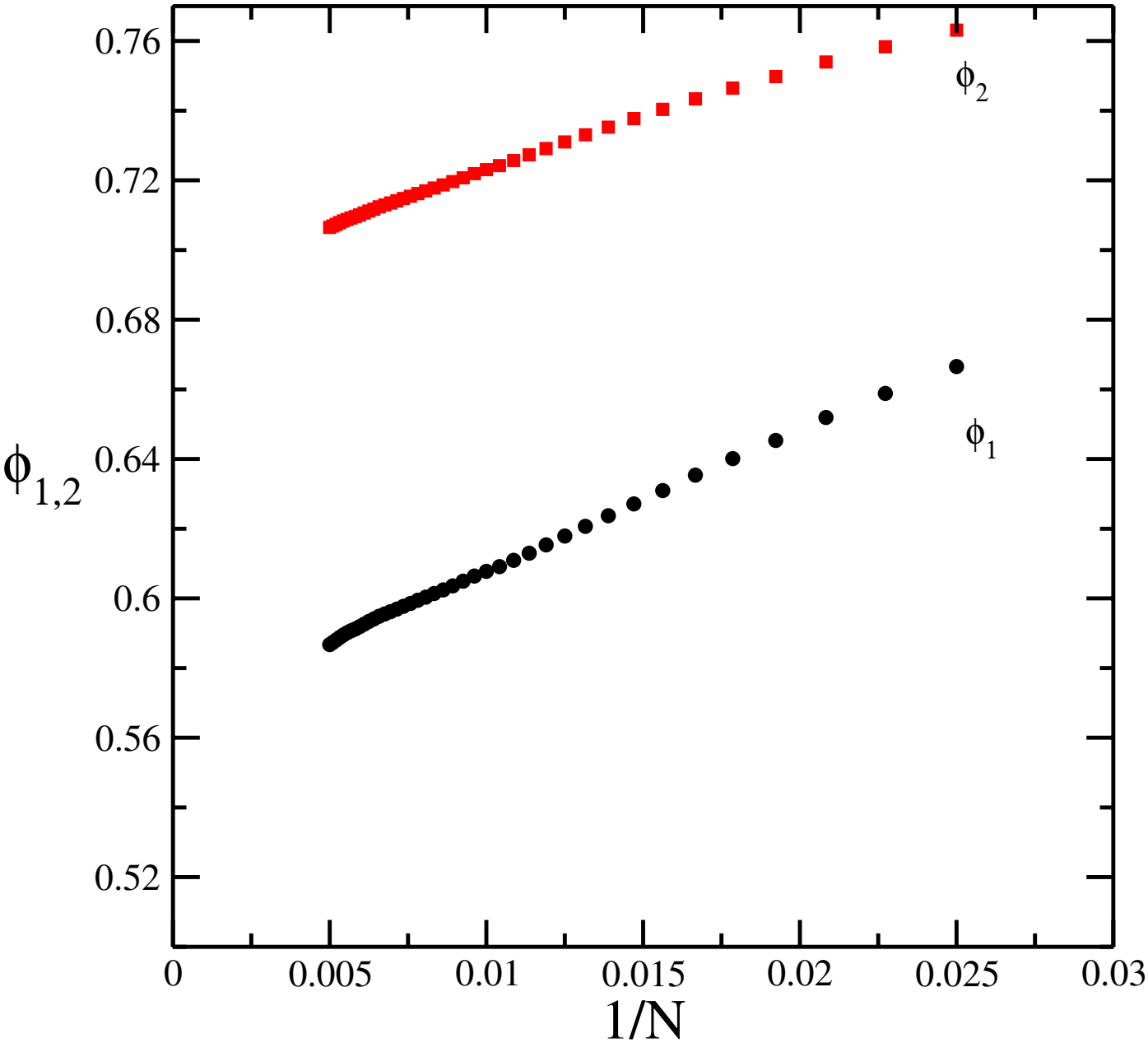

(a) ε2=0.6
ρ1
ρ2□
ρ1 or ρ2□
ε1
0
0.2
0.4
0.6
0.8
0.8
1
1.2

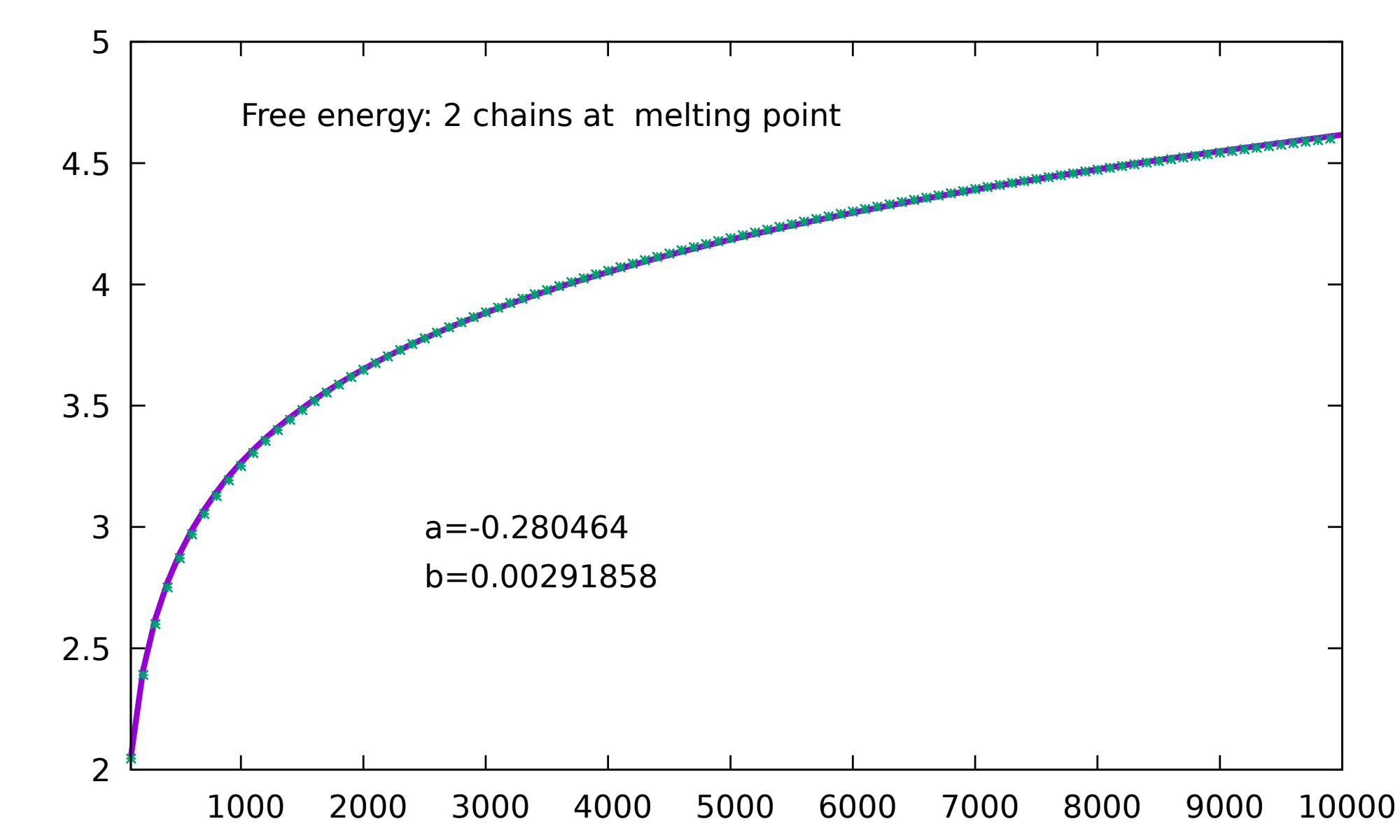

(b) ε2=1.0
ρ1
ρ2□
ρ1 or ρ2□
ε1□
0
0.2
0.4
0.6
0.8
0.7
0.8
0.9
1


1.4
1.2
1
0.8
0.6
0.4
0.2
0
ϕ_1
0
0.2
0.4
0.6
0.8
1
1.2
1.4
ε_1


1.4
1.2
1
0.8
0.6
0.4
0.2
0
ϕ_2
0
0.2
0.4
0.6
0.8
1
1.2
1.4
ε_1


0.8
0.78
0.76
0.74
0.72
0.7
0.68
0.66
$\varepsilon_{1,c}$
0
0.005
0.01
0.015
0.02
0.025
0.03
1/N


0.76
0.72
0.68
0.64
0.6
0.56
0.52
$\phi_{1,2}$
ϕ_2
ϕ_1
0
0.005
0.01
0.015
0.02
0.025
0.03
1/N


Free energy: 2 chains at melting point
a=-0.280464
b=0.00291858
5
4.5
4
3.5
3
2.5
2
1000
2000
3000
4000
5000
6000
7000
8000
9000
10000