Towards a scale-bridging description of ferrogels and magnetic elastomers

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Abstract
Ferrogels and magnetic elastomers differentiate themselves from other materials by their unique capability of reversibly changing shape and mechanical properties under the influence of an external magnetic field. A crucial issue in the study of these outstanding materials is the interaction between the mesoscopic magnetic particles and the polymer matrix in which they are embedded. Here we analyze interactions between two such particles connected by a polymer chain, a situation representative for particle-crosslinked magnetic gels. To make a first step towards a scale-bridging description of the materials, effective pair potentials for mesoscopic configurational changes are specified using microscopic input obtained from simulations. Furthermore, the impact of the presence of magnetic interactions on the probability distributions and thermodynamic quantities of the system is considered. The resulting mesoscopic model pair potentials can be used to economically model the system on the particle length scales. This first coarse-graining step is important to realize simplified but realistic scale-bridging models for these promising materials.

Keywords: magnetic elastomer, ferrogel, scale-bridging, coarse-graining, statistical analysis

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(Some figures may appear in colour only in the online journal)

1. Introduction

Ferrogels and magnetic elastomers are fascinating materials, born by the union of polymeric networks and ferrofluids. Their amazing properties derive from the unique combination of the elastic behavior typical for polymers and rubbers \cite{1} on the one hand, and magnetic effects characteristic of ferrofluids and magnetorheological fluids \cite{2, 3, 4, 5, 6, 7, 8, 9, 10} on the other. One of the most interesting results is that their shape and mechanical properties can be externally controlled by applying a magnetic field \cite{11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21}. A form of tunability, distinguished by reversibility as well as non-invasiveness and based on a magneto-mechanical coupling is one of the most appealing properties of these materials. This makes them excellent candidates for use as soft actuators \cite{22}, magnetic field detectors \cite{23, 24}, as well as tunable vibration and shock absorbers \cite{14, 15}. Moreover, studying their heat dissipation due to hysteretic remagnetization in an alternating external magnetic field might be helpful to understand better the processes during possible applications in hyperthermal cancer treatment \cite{26, 27}.

Typically, these materials consist of cross-linked polymer networks in which magnetic particles of nano- or micrometer size are dispersed \cite{16}. A central role in the coupling of magnetic and mechanical properties is played by the specific interactions between the embedded mesoscopic magnetic particles and the flexible polymer chains filling the space between them. These couplings are responsible for a modified macroscopic elasticity \cite{12, 28, 29}, orientational memory effects...
[30, 31], and reversibility of the magnetically induced deformations [15].

Many theoretical and computational studies have been performed on the topic, using different approaches to incorporate the particle-polymer interaction. Some of them rely on a continuum-mechanical description of both the polymeric matrix and the magnetic component [23, 32, 33]. Others explicitly take into account the discrete embedded magnetic particles, but treat the polymer matrix as an elastic background continuum [34, 35]. Usually in these studies, an affine deformation of the whole sample is assumed. The limitations of such an approach for the characterization of real materials have recently been pointed out [36]. In order to include irregular distributions of particles and non-affine sample deformations, other works employ, for instance, finite-element methods [19, 37–40].

To optimize economical efficiency, a first step is the use of simplified dipole-spring models. In this case, steric repulsion and other effects like orientational memory terms can be included [30, 36, 41–44]. A step beyond the often used harmonic spring potentials can be found in [42] where non-linear springs of finite extensibility are considered.

From all the studies mentioned above it becomes clear that microscopic approaches that explicitly resolve polymer chains are rare [45, 46], and what is particularly missing is a link between such microscopic approaches and the mesoscopic models that only resolve the magnetic particles, not the single polymer chains. In particular, a microscopic foundation of the phenomenological mesoscopic expressions for the model energies should be built up.

The present work is a first step to close this gap. We consider an explicit microscopic description in a first simplified approach: a single polymer chain, discretized through multiple beads, each representing a coarse-grained small part of the polymer, connects two mesoscopic particles. The ends of the polymer chain are rigidly anchored on the surfaces of the two mesoscopic particles, which are spherical, can be magnetized, and are free to rotate and change their distance. From molecular dynamics simulations on the microscopic level, we collect the statistics of the micro-states corresponding to different configurations in the mesoscopic model. Based on these statistics we derive effective mesoscopic pair potentials, inspired by previous achievements for other polymeric systems [47, 48]. The subsequent step in scale-bridging, connecting the mesoscopic picture to the macroscopic description, has been recently addressed [49] for a special class of magnetic polymeric materials.

In the following, we first define and describe our model in section 2. Then, in section 3, we mention the details of the microscopic simulation. In section 4, we further characterize the probability distribution of the mesoscopic variables, connecting it to a wrapping effect in section 5. After that, in section 6, we determine the values of mesoscopic model parameters based on the results of our microscopic simulations. In section 7, we derive an approximated expression for the mesoscopic effective pair potential characterizing the particle configurations. In this way, we build the bridge from the explicit microscopic characterization to the mesoscopic particle-resolved models by averaging over the microscopic details. Last, in section 8, we consider the effect of adding magnetic moments to the particles and, in section 9, the effect of increasing magnetic interaction on the thermodynamic properties, before we draw our final conclusions in section 10. Appendix A addresses the trends in the dependences of the mesoscopic model parameters on varying microscopic system parameters, while appendix B briefly comments on the separability of magnetic interactions in the mesoscopic picture and mesoscopic chain configurations.

2. The system

Our simplified system is composed of two mesoscopic and spherical particles, both of radius $a$, and a polymer chain explicitly resolved by $N = 60$ beads of diameter $\sigma$ and interconnected by harmonic springs. Here we choose the mesoscopic particle radius $a$ to be $5\sigma$.

We consider steric repulsion between all described particles through a WCA potential, which represents a purely repulsive interaction. It is given by

$$V_{\text{WCA}}(r'/\sigma) = \left\{ \begin{array}{ll} 4\varepsilon \left[ (r'/\sigma)^{-12} - (r'/\sigma)^{-6} \right] + \frac{1}{4} & \text{for } r' \leq r_c \\ 0 & \text{otherwise} \end{array} \right.$$  \hspace{1cm} (1)

where $r'$ is the distance between the particle centers, $\varepsilon$ denotes the energy scale of the potential, and $r_c = 2^{1/6}\sigma$ is the cut-off distance. For any combination of large and small particles, $\sigma'$ is chosen as the sum of the radii of the respective particles, which is equivalent to the mean of their respective diameters. In our simulations, we set $\varepsilon = 10k_BT$, where $k_B$ is the Boltzmann constant and $T$ is the temperature of the system. Neighboring beads within the chain are bound by means of a harmonic potential

$$V_H(r') = \frac{k}{2}(r' - r_0)^2,$$  \hspace{1cm} (2)

where we choose the force constant $k = 10k_BT/\sigma^2$. The equilibrium distance $r_0$ is set to match the cut-off distance of the WCA-potential $r_0 = r_c = 2^{1/6}\sigma$.

The ends of the chains are bound via the same harmonic potential to binding sites placed below the surface of the mesoscopic particles, see figure 1. These binding sites are rigidly attached to the mesoscopic particles and follow both, their translational and rotational motion. Thus, when the magnetic particle moves or rotates, the binding site of the polymer chain has to follow, and vice versa. The technical details for the virtual sites mechanism can be found in [50]. We identify the anchoring points of the polymer chain by the vectors $\mathbf{a}_1$ and $\mathbf{a}_2$, respectively (see figure 1). Furthermore, the distance vector between the two mesoscopic particle centers is indicated as $\mathbf{r}$, with magnitude $r$. The angles that the vectors $\mathbf{a}_1$ and $\mathbf{a}_2$ adopt with respect to the connecting vectors $\mathbf{r}$ and $-\mathbf{r}$ are denoted as $\theta_1$ and $\theta_2$, respectively. $\theta_1$ and $\theta_2$ are zenithal angles and they can span the interval $[0, \pi]$. Last, $\phi \in [-\pi, \pi]$ is the relative
azimuthal angle between the projections of $a_1$ and $a_2$ on a plane perpendicular to $r$. It can be used to parametrize the relative torsion between the two particles around the $r$ axis. Let us for brevity introduce the vector $\gamma = (r, \theta_1, \theta_2, \phi)$. Therefore, the $\gamma$-space, where our mesoscopic vector $\gamma$ is defined, is given by $[0, +\infty] \times [0, \pi] \times [0, \pi] \times [-\pi, \pi]$.

Through molecular dynamics simulations (thoroughly described in the next section) we find the probability density $p (\gamma)$ of a certain configuration $\gamma$ of the two mesoscopic particles. We normalize $p (\gamma)$ such that $\int p (\gamma) d\gamma = 1$, with $d\gamma = \sin(\theta_1) \sin(\theta_2) dr d\theta_1 d\theta_2 d\phi$ the $\gamma$-space volume element. Moreover, we are working in the canonical ensemble.

From statistical mechanics [51], we know that the probability density $p (\gamma)$ to find the system in a certain configuration $\gamma$ is $p (\gamma) = \exp[-\beta V (\gamma)]/Z_c$, where $\beta = 1/k_BT$ and $Z_c$ is the partition function of the system. Shifting $V (\gamma)$ by an appropriate constant, we can still reproduce $p (\gamma)$ but simultaneously normalize $Z_c = 1$. Then, since $\ln(Z_c) = 0$,

$$V (\gamma) = -k_BT \ln(p (\gamma))$$

(3)

represents an effective energy of the state $\gamma$ of our mesoscopic two-particle system and corresponds to an effective pair potential on the mesoscopic level, see also [47]. Through the normalization of $Z_c$, we set our reference free energy $F = -k_BT \ln(Z_c)$ equal to zero.

3. Microscopic simulation

To obtain the probability distribution from which the mesoscopic pair potential is derived, we performed molecular dynamics simulations using the ESPResSo software [50, 52]. Because entropic effects are important to capture the behavior of the polymer, the canonical ensemble is employed. This is achieved using a Langevin thermostat, which adds random kicks as well as a velocity-dependent friction force to the particles. For the translational degrees of freedom of each particle, the equation of motion is then given by

$$m_p \ddot{r}(t) = -\zeta \dot{r}(t) + F_c + F_r.$$  

(4)

where $m_p$ is the mass of a particle, $F_r$ is the force due to the interaction with other particles, $F_c$ denotes the random thermal noise, and $\zeta$ is the friction coefficient.

To maintain a temperature of $T$, according to the fluctuation-dissipation theorem, each random force component must have zero mean and variance $2k_BT\zeta$. Furthermore, random forces at different times are uncorrelated. In order to track the orientation of the magnetic nanoparticles, rotational degrees of freedom also have to be taken into account. This is achieved by means of a Langevin equation of motion similar to (4) where, however, mass, velocity, and force are replaced by inertial moment, angular velocity, and torque, respectively [53].

The time step for the integration using the Velocity-Verlet method [54] is $dt = 0.01$. To sample the probability distribution, we record the state of the simulation $\gamma$ every ten time steps. In order to obtain a smooth probability distribution over a wide range of parameters, 34 billion states have been sampled in total, by running many parallel instances of the simulation, summing up to about $10^8$ core hours of CPU time.

Finally, we find the probability distribution by sorting the results of our simulations into a histogram with 100 bins for each variable ($r, \theta_1, \theta_2$, and $\phi$). When a 64-bit unsigned integer is used as data type, this leads to a memory footprint of 800 Mb. Hence, the complete histogram can be held in memory on a current computer. If the resulting numerical version of the effective pair potential as defined in (3) is to be used in a simulation, a smoothing procedure should be employed, especially in parts of the configuration space with a very low probability density. One approach here might be hierarchical basis sets.

In our simulations, we chose the thermal energy $k_BT$ as well as the mass $m_p$, the friction coefficient $\zeta$, and the corresponding rotational quantities to be unity. We measure all lengths in units of $\sigma$ and the energies in multiples of $k_BT$.

4. Description of the probability density

We now consider some aspects of the probability density resulting from the microscopic simulation described in section 3. The entropic role of the microscopic degrees of freedom is considered by assigning to every configuration $\gamma$ a certain probability, given by the number of times it was encountered in the simulation divided by the total number of recorded states. As explained before, $\gamma = (r, \theta_1, \theta_2, \phi)$ corresponds to the set of variables that we use to describe the state of the system on the mesoscopic level.

In calculating the probability density from the microscopic simulations, we must remember the normalizing condition

$$\int p (\gamma) \sin(\theta_1) \sin(\theta_2) dr d\theta_1 d\theta_2 d\phi = 1,$$  

(5)

where $r$ is integrated over $[0, +\infty]$, $\theta_1$ and $\theta_2$ over $[0, \pi]$, and $\phi$ over $[-\pi, \pi]$. Therefore, to obtain the probability density, we have to properly divide the data acquired from the simulations by the factor $\sin(\theta_1) \sin(\theta_2)$. 

Figure 1. A simplified sketch of the geometry of the microscopic system. $r$ is the vector connecting the centers of the mesoscopic particles. $a_1$ and $a_2$ identify the anchoring points of the polymer chain on the surfaces of the particles. $\theta_1$, $\theta_2$, and $\phi$ are the angles that represent the remaining relative rotational degrees of freedom of the system.
It is useful to introduce here the average of a quantity over \( p_i \), defined as \( \langle \gamma \rangle = \int \gamma p_i(\gamma) \, d\gamma \). We can therefore calculate the average value of \( \gamma = \langle \gamma \rangle \), and the covariance matrix \( \Sigma_{\alpha\beta} = \langle (\alpha - \bar{\alpha})(\beta - \bar{\beta}) \rangle \), for \( \alpha, \beta = r, \theta_1, \theta_2, \phi \). We find \( \zeta = (20.20 \sigma, 0.36 \pi, 0.36 \sigma, 0) \). It is more practical to discuss the system in terms of correlation than in terms of covariance. Correlation is defined as \( \zeta_{\alpha\beta} = \Sigma_{\alpha\beta}/\sqrt{\Sigma_{\alpha\alpha} \Sigma_{\beta\beta}} \) (no summation rule in this expression), is dimensionless, and \( \zeta_{\alpha\beta} \in [-1,1] \). Here, we obtain

\[
\zeta = \begin{pmatrix}
1 & -0.341_{\pm 0.028} & -0.356_{\pm 0.029} & 0.083_{\pm 0.011} \\
-0.341_{\pm 0.028} & 1 & -0.006_{\pm 0.006} & -0.083_{\pm 0.010} \\
-0.356_{\pm 0.029} & -0.006_{\pm 0.006} & 1 & -0.083_{\pm 0.011} \\
0.083_{\pm 0.011} & -0.083_{\pm 0.010} & -0.083_{\pm 0.011} & 1
\end{pmatrix}
\]

(6)

where lines and columns refer to \( r, \theta_1, \theta_2, \phi \) in this order. The diagonal elements are unity by definition, because each variable is perfectly correlated with itself. The errors follow from the unavoidable discretization during the statistical sampling procedure in the simulations, where the results have to be recorded in discretized histograms of finite bin size.

We find a strong anticorrelation between \( r \) and \( \theta_{1,2} \), meaning that when the distance between the mesoscopic particles changes they tend to rotate. We will address in detail the background of this behavior in section 5 in the form of the wrapping of the polymer chain around the mesoscopic particles. For angles \( \theta_1 \) and \( \theta_2 \) different from 0 and \( \pi \) it is clear that such a wrapping and corresponding distance changes are likewise induced by modifying the relative torsion of angle \( \phi \). These coupling effects are partly reflected by the remaining non-vanishing correlations which, however, are weaker than the correlations between \( r \) and \( \theta_1, \theta_2 \) by at least a factor 4. The correlations between \( \theta_1 \) and \( \theta_2 \) are very weak and, within the statistical errors, may in fact even vanish. Within the statistical errors, the magnitudes of the correlations \( \zeta_{\theta_1 \theta_2}, \zeta_{\theta_2 \theta_1} \) as well as \( \zeta_{\theta_1 \phi}, \zeta_{\theta_2 \phi} \) agree well with each other, respectively, which reflects the symmetry of the system. Finally, we performed additional microscopic simulations for different sizes of mesoscopic particles and different chain lengths of the connecting polymer. As a general trend, we found that the correlations tend to decrease in magnitude for smaller mesoscopic particles and for longer polymer chains (see appendix A).

As a further step in the analysis of \( p_i \), we can determine the marginal probability density \( \hat{p}_i(\alpha) \) for one of the four mesoscopic parameters \( \alpha = r, \theta_1, \theta_2, \phi \) integrating out the other three, for instance, \( \hat{p}_b(\theta_1) = \int \hat{p}_i(\gamma) \sin(\theta_1) \, d\gamma/d\theta_2 \, d\phi \). This is the total probability density for the variable \( \theta_1 \) to assume a certain value, regardless of the others. Calculations for \( \hat{p}_1(r), \hat{p}_b(\theta_2), \) and \( \hat{p}_b(\phi) \) are analogously performed by integrating out all respective other variables (see figure 2). Of course, \( \hat{p}_i(\alpha) \) is still a normalized probability density since \( \int_{-\infty}^{\infty} \hat{p}_i(\alpha) \, d\alpha = \int \hat{p}_i(\gamma) \, d\gamma = 1 \), where we indicate \( d\alpha = \sin(\theta_1) \, d\theta_1 \), a steep increase in \( \hat{p}_b(\phi) \) at \( r = 11 \sigma \) is to be attributed to the WCA steric repulsion, since at that distance the two particles are in contact. From figure 2 we find that the maximum of \( \hat{p}_b(\theta_1)(i = 1, 2) \) is located at \( \theta_1 = 0 \). The highest probability density for \( \theta_{1,2} = 0 \) is obtained from the microscopic simulations by taking into account the \( \gamma \)-space normalization following from the use of spherical coordinates in (5), see also [47]. Moreover, \( \hat{p}_b(\phi) \) shows a maximum for \( \phi = 0 \), indicating that, as expected, the system does not tend to spontaneously twist around the connecting axis in the absence of further interactions. The presence of a maximum at \( \phi = 0 \) confirms that \( \hat{p}_b \) is an even function of \( \phi \) invariant under the transformation \( \phi \rightarrow -\phi \), as expected from the symmetry of the set-up.

5. Wrapping effect

Before developing an approximate analytical expression for the effective pair potential between the mesoscopic particles, it is helpful to examine in detail the results of the molecular dynamics simulations. In a magnetic gel in which mesoscopic magnetic particles act as cross-linkers [31, 55], two driving mechanisms for a deformation in an external magnetic field are possible. First, in any magnetic gel, the magnetic interactions between the mesoscopic particles lead to attractions and repulsions between them, which directly implies deformations of the intermediate polymer chains. As we would like to examine this mechanism separately, the magnetic interaction was not included explicitly in the simulations described in section 3. Rather, it will be considered later in section 8. Second, due to the anchoring of the polymer chains on the surfaces, rotations of the mesoscopic particles are transmitted to chain deformations. It has been shown in model II of [45] that the second mechanism on its own can lead to a deformation of such a gel in an external magnetic field; if mesoscopic magnetic particles are forced to rotate to align with the field, the polymers attached to their surfaces have to follow. The resulting wrapping of the polymer chains around the particles leads to a shrinking of
the gel. Transferring this to our model system, it could imply an external magnetic field rotating the particles to a state in which the angles $\theta_1$ and $\theta_2$ are non-zero. The effect of induced particle rotations on the interparticle distance is illustrated in figure 3, where the effective pair potential $V_c(r)$ is plotted for the case of $\phi = 0$ and various values of $\theta_1 = \theta_2 = \theta$. i.e. both particles are rotated by the same angle $\theta$ with respect to the connecting vector $\pm r$, respectively. It can be seen that the more the particles are rotated, the more the minimum of the effective pair potential is shifted towards closer interparticle distances corresponding to smaller separation distances $r$.

$V_c(r)$

Figure 3. Plot of the effective mesoscopic pair potential $V_c(r)$ for a situation in which both particles are rotated by the same amount to the angles $\theta_1 = \theta_2 = \theta$ at a torsional angle of $\phi = 0$. It can be seen that the more the particles are rotated out of their equilibrium position, the closer they will approach each other because the minimum of the effective pair potential shifts to smaller separation distances $r$. The irregularities in the effective pair potential for high values of $r$ are attributed to the naturally low sampling of those low-probability configurations.

Figure 2. Marginal probability densities for the single variables. The abscissa has been rescaled; $r$ ranges from 0 to $100\sigma$, whereas the angles range from 0 to $\pi$. $\tilde{p}_{\theta_1}(\theta_1)$ and $\tilde{p}_{\theta_2}(\theta_2)$ are practically indistinguishable due to the symmetry of the set-up. Due to the symmetry of the system under the transformation $\phi \rightarrow -\phi$, here we only plot $\tilde{p}_\phi(\phi)$ from 0 to $\pi$. The maxima of the single-variable densities are located at $r = 17\sigma$ and $\theta_{1,2} = \phi = 0$, respectively. The maxima of $\tilde{p}_{\theta_1}(\theta_1)$ and $\tilde{p}_{\theta_2}(\theta_2)$ at $\theta_{1,2} = 0$ are found from the microscopic simulations after the $\gamma$-space normalization contained in (5) has been taken into account.

$V_c(r)$

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Figure 4. Average distance between the two mesoscopic particles versus the angles $\theta_1$ and $\theta_2$ shown as a color and contour map for the two cases $\phi = 0$ (top) and $\phi = \pi$ (bottom). It can be seen that even by rotating only one of the two particles, the average distance can be reduced considerably. However, the maximum reduction is observed when both particles are rotated. The variation of the torsion angle $\phi$ does not change the main trend but has a moderate influence for intermediate rotations.

To get a more detailed picture, we also consider independent rotations of the two mesoscopic particles. In figure 4, the average distance between the particles is depicted in a contour plot as a function of the angles $\theta_1$ and $\theta_2$. Images are shown for torsion angles of $\phi = 0$ and $\phi = \pi$. It can be seen that quite a decrease in the average distance can already be achieved by rotating only one particle. However, very strong reduction in interparticle distance can only occur when both particles are rotated. The torsion angle $\phi$ does not alter the general trend of reduction of the average distance when the particles are rotated. However, the resulting numbers vary to a certain degree.

6. One-variable effective pair potentials

We now introduce some mesoscopic analytical expressions to model the effective pair potentials $\tilde{\mathcal{U}}_\lambda(\sigma)$ introduced in (7). We will determine functional forms and parameters that can be
used to model strain and torsion energies. Here, we use the term ‘strain’ to denote variations of \( r \), whereas the term ‘torsion’ is used to describe changes in the angles \( \theta_1 \) and \( \theta_2 \) or \( \phi \), depending on the initial orientation of the spheres and relative rotations. The quality of the analytical model expressions will be analyzed by fitting to the corresponding results from the microscopic simulations.

First, we turn to the energetic contributions arising from changes in the interparticle distance \( r \). The corresponding effective energy \( \bar{V}(r) \) obtained from the microscopic data is plotted in figure 5. It can be seen that there are essentially two regimes: up to \( r \approx 11\sigma \) the WCA repulsion between the two mesoscopic particles dominates, whereas, for \( r > 11\sigma \), \( \bar{V}(r) \) shows a smooth behavior and a single minimum arising from the entropic contribution of the polymer chain. Moreover, at \( r \geq 55\sigma \), \( \bar{V}(r) \) shows an irregular, non-smooth behavior. This is attributed to the low sampling rate of this extremely stretched configuration, which has a very low probability to occur in the microscopic simulations (see figure 2). As a first approximation, it is natural to reproduce \( \bar{V}(r) \) by a harmonic expansion for \( r > 11\sigma \),

\[
V_{\text{harm}}(r) = V_0^r + \frac{k_h}{2} (r - r_{0,h})^2. \tag{8}
\]

We derive the mesoscopic parameters \( V_0^r, k_h, r_{0,h} \) by fitting \( V_{\text{harm}}(r) \) in a neighborhood of the minimum to the data \( \bar{V}(r) \) obtained from microscopic simulations. In figure 5 the resulting parameters and the two curves are shown.

Moreover, we have compared \( \bar{V}(r) \) with the following expression for a finitely extensible non-linear elastic potential (FENE) \([42, 56]\),

\[
V_{\text{FENE}}(r) = V_0^r - \frac{K_f r_{\text{max}}^2}{2} \ln \left[ 1 - \left( \frac{r - r_0}{r_{\text{max}}} \right)^2 \right]. \tag{9}
\]

It takes into account that the chain cannot extend beyond a maximal length, since \( V_{\text{FENE}}(r) \) diverges when \( r \rightarrow r_0 + r_{\text{max}} \).

We see that \( d^2V_{\text{FENE}}(r)/dr^2 = K_f \) for \( r = r_0 \) and therefore \( K_f \) represents the elastic constant in a harmonic expansion of this non-linear potential. As for the harmonic expression, we fit \( V_{\text{FENE}}(r) \) to our microscopic data and thus derive the mesoscopic model parameters \( K_f, r_0, r_{\text{max}}, \) and \( V_0^r \) as displayed in figure 6. The agreement between the resulting expression for \( V_{\text{FENE}}(r) \) and \( \bar{V}(r) \) in the regime \( r \geq 11\sigma \) is excellent, especially for the branch of the curve right to the minimum. According to the fit, the maximum extension of the chain occurs for \( r = r_0 + r_{\text{max}} \approx 75\sigma \). In fact, since the radius of the mesoscopic particles is \( 5\sigma \) and each of the 60 beads making up the polymer chain has diameter \( \sigma \), at \( r = 70\sigma \) the polymer chain is completely stretched. A further elongation is of course possible due to the harmonic inter-head interaction and this justifies the result of \( \approx 75\sigma \) for the maximal extension.

Last, we compare the elastic constants \( k_h \) and \( K_f \) obtained from the harmonic and FENE approximation, respectively, as listed in figures 5 and 6. The resulting values of 0.0114k_bT/\sigma^2 and 0.015k_bT/\sigma^2 are in good agreement with each other.

To find a mesoscopic model expansion for the effective pair potential \( \bar{V}_\phi(\theta_1) [\bar{V}_\phi(\theta_2)] \) has a similar behavior, we compare it with the phenomenological expression,

\[
V_{\phi}(\theta_1) = V_0^\phi + D[\cos(\theta_1) - \cos(\theta_0)]^2. \tag{10}
\]

introduced in (3) of [30]. As before, we can derive the mesoscopic parameters \( V_0^\phi, D, \) and \( \theta_0 \) by fitting \( V_{\phi}(\theta_1) \) to the microscopic data represented by \( \bar{V}_\phi(\theta_1) \). The resulting parameters and the comparison between the two curves are shown in figure 7. Although (10) does not perfectly reproduce the one-variable pair potential \( \bar{V}_\phi(\theta_1) \), it appears as a reasonable approximation in the neighborhood of the minimum energy. Moreover, the rather flat behavior of \( \bar{V}_\phi(\theta_1) \) for small \( \theta_1 \) is well represented by \( V_{\phi}(\theta_1) \), which is at leading order proportional to \( \theta_1^4 \).

Finally, we want to find a mesoscopic expression to reproduce the effective pair potential \( \bar{V}_\phi(\phi) \) obtained from the
microscopic data due to relative torsional rotations between the two particles around the connecting vector \( r \). We fit the microscopic data \( V(\theta) \) around the minimum with the expression

\[
V(\phi) = V^0 + \tau [\cos(\phi_0) - \cos(\phi)].
\]  

(11)

It leads to the parameters and fit depicted in figure 8 and is quadratic at lowest order in \( \phi \), \( V(\phi) \approx V^0 + \tau \phi^2 \). The parameter \( \phi_0 \) is redundant and can be absorbed into \( V^0 \), but we leave it for reasons of comparison to the following (12).

It is interesting to compare \( V(\phi) \) with the expression taken from (4) of [30],

\[
V(\phi) = V^0 + \tau [\cos(\phi_0) - \cos(\phi)]^2.
\]  

(12)

The discrepancy between the two curves shown in figure 9 is obvious. The reason becomes clear when we expand (12) to lowest order in \( \phi \), \( V(\phi) \approx V^0 + \tau \phi^4 \). This expression is quartic in the torsion angle \( \phi \), leading to the comparatively flat behavior in the region around the minimum. The last two comparisons suggest that the analytical form of the mesoscopic model pair potential as a function of the azimuthal angle, which acts as a torsion, can be optimized by using a quadratic form as in (11) instead of the quartic one implied by (12) at leading order around the minimum.

Finally, we performed additional microscopic simulations to estimate how the microscopic system parameters affect the mesoscopic model parameters. In particular, the influence of the mesoscopic particle radius \( a \) and the number of beads \( N \) of the connecting polymer chain was investigated. As a trend, we found that the single-variable potentials tend to become stiffer for shorter chains and for bigger mesoscopic particles (see appendix A). Moreover, we observe that the fit of the functional forms in expressions (9)–(11) with the simulation data further improves for increasing chain length and decreasing size of the mesoscopic particles.

So far, we have discussed mesoscopic analytical expressions to approximate the one-variable effective pair potentials

7. Building a coupled effective pair potential

Our goal is to describe the effective interaction between the two mesoscopic particles, coarse-graining the role of the connecting polymer chain. The entropic nature of the polymeric interactions provides a direct route to average out the microscopic degrees of freedom and thus build an effective scale-bridged model. A natural way to proceed would be to find an analytical approximation for \( \rho(\gamma) \) from the microscopic simulations.

Corresponding data curves from the additional microscopic simulations and fitted mesoscopic model curves are summarized in the supplemental material (stacks.iop.org/JPhysCM/27/325105/mmedia).
analytical expression for the mesoscopic pair potentials in the spirit of (3).

As a first approximation we may describe $p_c$, as the simple product of the one-variable probability densities, $\prod_a \tilde{P}_a(\alpha)$. We obtain

$$p_{\text{app}}(\gamma) = \exp \left( -\beta [V_{\text{WCA}}(r/2a) + V_{\text{FENE}}(r) + V_\phi(\theta_1) + V_\phi(\theta_2) + V_\phi(\phi)] \right). \quad (13)$$

when we use (7). Analytical approximations for the pair potentials $\tilde{V}_a(\alpha)$ were derived in (9)–(11) in section 6 together with the fitting parameters listed in figures 6–8. Care must be taken for the term $\tilde{V}_a(r)$, which has to be substituted by $V_{\text{WCA}}(r/2a) + V_{\text{FENE}}(r)$ to take account of the steric repulsion between the mesoscopic particles, see (1). (13) correctly describes some aspects of the system behavior. For instance, the steep variation due to the WCA potential between the mesoscopic particles is well represented in this approximation. However, this description would lead to a distribution with vanishing correlations between the mesoscopic variables. That corresponds to assuming them independent of each other, which omits some important physical aspects, see the wrapping effect in section 5.

To make a step forward and take account of correlations we multiply correction terms to the previous factorized approximation in (13), obtaining the expression

$$p_{\text{app}}(\gamma) \propto p_{\text{app}}^c(\gamma) \times \exp \left( -\gamma - \xi \cdot \Xi \cdot (\gamma - \xi) \right). \quad (14)$$

where the elements of $\xi$, a 4-components vector, and $\Xi$, a $4 \times 4$ symmetric matrix, are free parameters chosen to match the original data. There are at least two possible numerical approaches to find the best $\xi$ and $\Xi$ values. On the one hand, we can simply fit the original probability density (e.g. minimize the squared difference between $p_c$ and $p_{\text{app}}$). On the other hand, we can follow a moment-matching approach, looking for a $p_{\text{app}}$ that has a correlation matrix as close as possible to the original one. We performed a mixed strategy by fitting the expression in (14) to the simulation data $p_c(\gamma)$, using as a criterion for best initialization of the fit an outcome that as closely as possible matches the correlations directly calculated from the simulation data. This fit was performed by minimizing the squared difference between $p_c(\gamma)$ and $p_{\text{app}}(\gamma)$, using the Nelder–Mead algorithm [57] provided by the SciPy library [58]. In this procedure, normalization of the approximated probability density as in (5) is enforced. The best set of parameters $\xi$ and $\Xi$ found is

$$\xi \approx \begin{pmatrix} 23.54a \\ 0.321\pi \\ 0.321\pi \\ 0 \end{pmatrix},$$

$$\Xi \approx \begin{pmatrix} 4.8 \times 10^{-3}a^2 & 0.19/\pi & 0.19/\pi & -1.8 \times 10^{-3}/\pi^2 \\ 0.19/\pi & 1.21/\pi & 1.54/\pi & 1.25/\pi^2 \\ 0.19/\pi & 1.54/\pi & 1.21/\pi & 1.25/\pi^2 \\ -1.8 \times 10^{-3}/\pi & 1.25/\pi & 1.25/\pi & -0.32/\pi^2 \end{pmatrix}. \quad (15)$$

resulting in the correlation matrix

$$q_{\text{app}} \approx \begin{pmatrix} 1 & -0.306 & -0.306 & 0.045 \\ -0.306 & 1 & 0.027 & -0.071 \\ -0.306 & 0.027 & 1 & -0.071 \\ 0.045 & -0.071 & -0.071 & 1 \end{pmatrix}. \quad (17)$$

It would be unrealistic to try to exactly reproduce all properties of $p_c(\gamma)$ through an analytical approximation. Nevertheless, using the resulting expressions from (14)–(16), we can take account of the strong anticorrelation $q_{\theta_1,2}$ between $r$ and $\theta_1,2$, which is the strongest and most important one in the system; compare (6) and (17). For the other elements $q_{\phi,\theta}$, $q_{\theta_1,\phi}$, and $q_{\theta_1,\theta_2}$ we infer that having stronger deviations. However, those correlations are smaller than $q_{\theta_1,\theta_2}$ at least by a factor 4 and therefore carry a smaller amount of information about the overall system behavior.

As a total result and in analogy to (3), we obtain from (14) the optimized analytical expression $V_{\text{app}}$ to model the effective interaction between the mesoscopic particles:

$$V_{\text{app}} = V_{\text{WCA}}(r/2a) + V_{\text{FENE}}(r) + V_\phi(\theta_1) + V_\phi(\theta_2) + V_\phi(\phi) + k_B T \sum_{\alpha \beta} \Xi_{\alpha\beta}(\alpha - \xi_\alpha)(\beta - \xi_\beta). \quad (18)$$

The corresponding expressions and values for the fitting parameters are given by (1), (9)–(11), (15) and (16) as well as figures 6–8. Thus, the effective pair potential is divided into two parts: one-variable and two-variable potentials. The former are the analytical single-variable pair potentials derived in section 6, see (1) and (9)–(11), together with the diagonal $\alpha = \beta$ terms in the double summation of (18). The latter are the off-diagonal $\alpha \neq \beta$ terms and take account, to lowest order, of the correlations between different mesoscopic degrees of freedom. Correlations between $r$ and $\theta_1,2$ are the dominating ones, leading to such physical effects as the wrapping effect introduced in section 5.
8. Impact of magnetic dipole moments

We will now consider how magnetic interactions between the mesoscopic particles modify the physics of the system, in particular the probability densities and other mesoscopic quantities. For this purpose, we assign to each particle a permanent magnetic dipole moment \( \mathbf{m}_i \) \((i = 1, 2)\), in the present work not going into the details of how these could be generated. We introduce these magnetic moments in the state of highest probability density. At vanishing magnetic moments, this is the state of minimal effective energy of the mesoscopic system, i.e. following \( (3) \), the one that has maximum \( p_\gamma(\gamma) \) over all the configurations \( \gamma \). The maximum occurs for \((r^M, \theta_1^M, \theta_2^M, \phi^M) \approx (26\sigma, 0, 0, 0, 0)\), and is displayed in figure 10. When the mesoscopic particles are in this configuration, we assign to them the two magnetic moments \( \mathbf{m}_1 \) and \( \mathbf{m}_2 \) that are parallel to each other and point along the connecting vector \( \mathbf{r} \), i.e. in their orientation of minimal magnetic energy, as depicted in figure 10. To first order, this configuration should leave the angular orientations unchanged when increasing the magnetic interaction, but see also the discussion in section 5. This is just one of the possible orientations that the moments could assume. However, such a configuration could be achieved with a certain probability, for instance, when the sample is cross-linked in the presence of an external magnetic field that aligns the magnetic moments \([59–62]\).

The dipole moments \( \mathbf{m}_1 \) and \( \mathbf{m}_2 \) are assumed to have equal magnitude \( m = |\mathbf{m}_1| = |\mathbf{m}_2| \) and are rigidly fixed with respect to each particle frame. We measure the magnetic moments in multiples of \( m_0 = \sqrt{4\pi\varepsilon_0 T (2a)^3/\mu_0} \), where \( \mu_0 \) is the vacuum magnetic permeability. Then, for each \( \gamma \), we can calculate the magnetic dipole interaction energy between the two moments

\[
V_\text{m}(\gamma) = \frac{\mu_0}{4\pi} \mathbf{m}_1 \cdot \mathbf{m}_2 \cdot r^2 - \frac{3}{r^3} (\mathbf{m}_1 \cdot \mathbf{r})(\mathbf{m}_2 \cdot \mathbf{r})
\]

\[
= \frac{\mu_0 m^2}{4\pi} \frac{-2 \cos(\theta_1^m) \cos(\theta_2^m) + \sin(\theta_1^m) \sin(\theta_2^m) \cos(\phi^m)}{r^3}.
\]

(19)

\( \theta_i^m \) indicates the angle between \( \mathbf{m}_i \) and \( \mathbf{r} \), while \( \phi^m \) is the angle between the projections of \( \mathbf{m}_1 \) and \( \mathbf{m}_2 \) on a plane perpendicular to \( \mathbf{r} \). These quantities can be expressed in the \( \gamma \) variables once the orientations of \( \mathbf{m}_1 \) and \( \mathbf{m}_2 \) with respect to the particle frames are set by the protocol we described above.

For non-magnetic polymer chains, the potential acting on the mesoscopic level \( V(\gamma) \) is separable into a sum of magnetic and non-magnetic interactions \( V(\gamma) = V_m(\gamma) + V_c(\gamma) \). Consequently, magnetic effects do not modify the contributions of the polymer chain to \( V(\gamma) \), as is further explained in appendix B.

This corresponds to a factorization of the probability densities \( p_\gamma(\gamma) \) and \( p_m(\gamma) \), where the latter is defined via the Boltzmann factor

\[
p_m(\gamma) = \exp[-\beta V_m(\gamma)]
\]

in analogy to \( (3) \). Therefore the total probability density becomes

\[
p(\gamma) = \frac{\exp[-\beta V(\gamma)] p_\gamma(\gamma)}{Z(m)} = \frac{p_\gamma(\gamma) p_m(\gamma)}{Z(m)}.
\]

(21)

where \( Z(m) = \int p_\gamma(\gamma) p_m(\gamma) d\gamma \) is the partition function describing the system for non-vanishing magnetic moments. We can calculate averages on the system with magnetic interactions by

\[
\langle \cdots \rangle = \int p(\gamma) p_m(\gamma) d\gamma = \frac{1}{Z(m)} \int p(\gamma) p_m(\gamma) d\gamma.
\]

(22)

The single-variable marginal probability density \( p_\alpha(\alpha) \), \( \alpha = r, \theta_1, \theta_2, \phi \), is again defined as \( p(\gamma) \) integrated over all the \( \gamma \) variables except for \( \alpha \). This is the same procedure as described in section 4 but substituting \( p_\gamma(\gamma) \) with \( p(\gamma) \). As can be seen from figure 11, \( p(r) \) shows an increase of the probability to find the particles close together when \( m \) is increased. A peak builds up at small \( r \) because the magnetic energy tends to \( V_\text{m}(r \to 0) \to -\infty \). Although the magnetic spheres attract each other, a collapse is prevented by the WCA-potential, which becomes effective at \( r \leq 11\sigma \) and for \( r \to 0 \) diverges faster to +\( \infty \) than the magnetic energy to \( -\infty \). For \( m = m_0 \), the presence of a double peak in \( p(r) \) could be connected to a hardening transition of the kind described in \([30]\). Due to the mutual magnetic attraction between the parallel dipoles we expect the average distance \( \langle r \rangle \) to decrease with increasing \( m \), and indeed it does so, as can be seen from the inset in figure 11.

The changes in the angular distributions for \( \theta_1 \) and \( \theta_2 \) due to the magnetic interaction are illustrated in figure 12. For the two angles \( \theta_1 \) and \( \theta_2 \), the distributions \( p_\theta(\theta_1) \) and \( p_\theta(\theta_2) \) are similar, and the behavior for varying \( m \) is approximately the same. Therefore, we only display the results for \( p_\theta(\theta_1) \). The magnetic moments tend to align in parallel along the connecting axis \( \mathbf{r} \), corresponding to their absolute energy minimum. Since we introduced the magnetic moments in that configuration for \( \theta_1 = \theta_2 = 0 \), for any fixed \( \theta_1 \) the minimum of
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Vm, and therefore the maximum of pm, is located at \(\theta_1 = \theta_2 = \pi\) and vice versa. As a consequence, the \(\theta_1 = \theta_2 = 0\) configuration leads to the same magnetic interaction energy as \(\theta_1 = \theta_2 = \pi\). Concerning only magnetic interactions, both configurations show the same probability \(p_m\). Integrating out \(r, \theta_2,\) and \(\phi\) from \(p_m(\gamma)\), the shape of the resulting magnetic probability density for \(\theta_1\) is symmetric around \(\theta_1 = \pi/2\) with one peak at \(\theta_1 = 0\) and one at \(\theta_1 = \pi\). Therefore, coupling magnetic \(p_m\) and non-magnetic \(p_c\) contributions, we find that some probability shifts to higher values of \(\theta_1\) due to the magnetic interactions. However, we find the standard deviation of \(p_m(\theta_1)\) to decrease with increasing \(m\) (see the inset of figure 12), meaning that the particles become less likely to rotate along the \(\theta_1\) (or likewise \(\theta_2\)) direction.

The same calculation for \(p_c(\phi)\) shows that the particles also become less likely to rotate around the connecting vector \(r\). The probability density of \(\phi \approx 0\) rises and sharpens, as we can see in figure 13, and the standard deviation (see the inset of figure 13) decreases, confirming quantitatively the sharpening of \(p_c(\phi)\).

9. Thermodynamic properties

Finally, we provide the connection to the thermodynamics of our canonical system and demonstrate the influence of the magnetic interactions. With the partition function \(Z(m)\) as in (21), we can calculate the overall free energy as \(F(m) = -k_B T \ln(Z(m))\), the internal energy of the system as

\[
U(m) = \langle V(m) \rangle = \frac{\int V(\gamma) r(\gamma) p_m(\gamma) d\gamma}{Z(m)} \tag{23}
\]

with \(V(\gamma)\) defined before (20), and the entropy as \(S(m) = [U(m) - F(m)]/T\).
To test the validity of our effective potential scheme using the coupling approximation described in section 7, we calculate the thermodynamic quantities using both probability densities \( p_r(\gamma) \) and \( p_{app}(\gamma) \) and compare the results. At vanishing magnetic moment, the free energy of the system is the same for both probability densities. This is expected, because it is a direct consequence of the normalization condition: \( Z(m = 0) = Z_c = \int p_r(\gamma) d\gamma = \int p_{app}(\gamma) d\gamma = 1 \) by construction of \( p_{app}(\gamma) \). For non-vanishing magnetic moments, the partition functions obtained from the two probability densities (and thus the corresponding free energies) start to deviate from each other because the integral \( \int p_r(\gamma)p_m(\gamma)d\gamma \) is, in general, different from \( \int p_{app}(\gamma)p_m(\gamma)d\gamma \). With increasing magnetic interaction, the difference between \( p_r(\gamma) \) and \( p_{app}(\gamma) \) becomes more important and, as shown in figure 14, leads to a slightly increasing deviation in the free energies \( F(m) \) calculated in both ways: at \( m \approx 1.224m_0 \) they already differ by \( \sim 5.6\% \). Analogously, we can compare the internal energies of the system, shown in figure 15: at \( m = 0 \) the error due to the probability density approximation is \( \sim 0.12\% \) of the exact value, rising up to \( \sim 0.32\% \) at \( m \approx 1.224m_0 \). A similar deviation follows for the entropy, see figure 16, where, however, the error at \( m \approx 1.224m_0 \) increases to \( \sim 0.26\% \) of the exact value. Overall, however, the relative errors remain small, which confirms the validity and viability of our coarse-grained effective model potential in (18).

To summarize the effect of the magnetic dipoles on the system, we may conclude that the particles are driven towards each other. In other words, the average distance between them decreases (see figure 11) and the distributions for the particle separation and for the angular degrees of freedom sharpen (see figures 11–13). This reflects a decrease in entropy, which becomes possible due to the gain in magnetic interaction energy. Indeed, the calculated entropy decreases with increasing magnetic interactions (see figure 16), which is achieved by the decreasing free energy and internal energy (see figures 14 and 15, respectively).

Figure 16. Entropy \( S(m) = (U(m) - F(m))/T \) as a function of the reduced magnetic moment \( m/m_0 \), calculated once using \( p_r(\gamma) \) from the microscopic simulation (labeled with ‘\( S_c \)’), and once replacing \( p_r(\gamma) \) by \( p_{app}(\gamma) \) (labeled with ‘\( S_{app} \)’).

10. Conclusions

Most of the previous studies on ferrogels describe the polymer matrix as a continuous material [23, 32, 34, 35] or represent it by springs connecting the particles [30, 36, 41–44], but only a few resolve explicitly the polymeric chains [45, 46]. In particular, a link between such microscopic chain-resolved calculations and the expressions for the investigated mesoscopic model energies has so far been missing. We have outlined in the present work a way to connect detailed microscopic simulations to a coarse-grained mesoscopic model.

This manifests a step into the direction of bridging the scales in material modeling. Starting from microscopic simulations considering explicitly an individual polymer chain connecting two mesoscopic particles, we specified effective mesoscopic pair potentials by fitting analytical model expressions. In this way, we were able to optimize a coarse-grained mesoscopic model description on the basis of the input from the explicit microscopic simulation details. Furthermore, we have shown that correlations between the mesoscopic degrees of freedom must be taken into account to provide a complete picture of the physics of the system. Moreover, we have examined the effect of a magnetic interaction, finding a tightening of the system by reducing the average distance between the magnetic particles and by reducing the rotational fluctuations around the configuration of highest probability density.

Our system consisted of only two mesoscopic particles, for which we derived the corresponding effective pair interaction potential. However, using this pair potential in a first approximation, two- and three-dimensional structures can be built up, similarly to elastic network structures generated using pairwise harmonic spring interactions between the particles [36, 43]. Including as a first approach magnetic interactions between neighboring particles only, the different angles between the magnetic moment of a particle and the anchoring points of the polymer chain to its different neighbors must be taken into account. Yet, using the reduced picture of pairwise mesoscopic interactions, it should be possible to reproduce for example previously observed deformational behavior in external magnetic fields for two- and three-dimensional systems [45, 46].

The scope of our approach is a first attempt of scale-bridging in modeling ferrogels and magnetic elastomers. Naturally, the procedure can be improved in many different ways. For example, we would like to study a system with multiple chains connecting the magnetic particles, with anchoring points randomly distributed over the surfaces of the particles. Such a development would eliminate artificial symmetries in the model and be another step towards the description of real systems. Furthermore, to study more realistic systems, interactions between more than two mesoscopic particles would need to be considered. Likewise, also the effect of interactions between next-nearest neighbors connected by polymer chains could be included. The last two points imply a step beyond the reduced picture of considering only effective pairwise interactions between the mesoscopic particles. Finally, via subsequent procedures of scale-bridging from the mesoscopic to the macroscopic level [49], a connection between
Effect of varying microscopic system parameters on the resulting mesoscopic model parameters. In the microscopic simulations, we varied the mesoscopic particle radius \( a \) and the number of beads \( N \) forming the connecting polymer chain. Fits of the single-variable mesoscopic model potentials (9)–(11) to the microscopic simulation data lead to the presented mesoscopic model parameters. (a)–(d), (o)–(q) Model parameters for the FENE potential (9); (e), (f), (Δ) model parameters for the angular \( \theta_r \) and \( \theta_2 \)-potential (10); (g), (h), (Q) model parameters for the torsional \( \phi \)-potential (11). Values for \( \theta_0 \) and \( \phi_0 \) vanished in all cases and are not shown. The data points for \( a/\sigma = 1 \) and \( N/60 = 1 \) correspond to the results presented in figures 6–8.

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Appendix A. Dependence on microscopic system parameters

In sections 6 and 7 we derived an analytical approximation for the effective mesoscopic model potential describing our system. Here, we test how the mesoscopic model parameters depend on the microscopic system parameters. In particular, we performed additional microscopic simulations with varied radius \( a \) of the mesoscopic particles and varied number \( N \) of beads forming the connecting polymer chain.

In each case, we repeated the analysis of sections 4, 6 and 7 by fitting the mesoscopic single-variable model potentials (9)–(11) to the microscopic simulation data. This reveals the trends in the dependences of the coefficients in the mesoscopic model potentials on the parameters \( a \) and \( N \), as depicted in figure A1. From the trend of the resulting model parameters \( K_f \), \( D \), and \( \tau \) in figures A1(c), (e) and (g) we conclude that the potentials tend to become stiffer as the chain becomes shorter or—at least for the rotational degrees of freedom—the mesoscopic particles become larger (see 4 for corresponding fitting curves).

Moreover, we have calculated the trend in the correlations between the mesoscopic degrees of freedom for varying values of \( a \) and \( N \), see figure A2. We find that the magnitude of the correlations decreases with increasing \( N \) or decreasing \( a \). Thus, quite intuitively, the coupling between the \( y \) variables tends to decrease with longer chains or smaller mesoscopic particle sizes.

Appendix B. Separability of the Hamiltonian and consequences for coarse-graining

In section 8 we have included the magnetic interactions analytically on the mesoscopic level. They had not been included in the microscopic simulations. This procedure is possible due to a separability of the magnetic and non-magnetic effects, which as a consequence implies a factorization of the corresponding probability. Thus the contribution of the polymer chain needs to be calculated explicitly. The contribution of any interaction acting solely on the mesoscopic particles can be exactly taken into account separately afterwards.

This argument relies on the separability of the Hamiltonian into a sum of mesoscopic and microscopic parts, as well as on the fact that magnetic interactions affect the mesoscopic part only. We write down the Hamiltonian of the system as

\[
\mathcal{H}(\gamma, \Gamma) = \mathcal{H}_{\text{meso}}(\gamma) + \mathcal{H}_{\text{micro}}(\Gamma) = \sum_{n=1}^{N} [V_{\text{WCA}}^{A,n}(\gamma, \Gamma) + V_{\text{WCA}}^{B,n}(\gamma, \Gamma)] + \sum_{n < n'} V_{\text{WCA}}^{n,n'}(\Gamma)
\]

(4) Corresponding data curves from the additional microscopic simulations and fitted mesoscopic model curves are summarized in the supplemental material (stacks.iop.org/JPhysCM/27/325105/mmedia).
Similarly to the main text, let us here indicate with $\gamma$ the degrees of freedom (now velocities included) of the mesoscopic particles and with $\Gamma$ the ones of the microscopic particles that build up the chain. For brevity we here label the mesoscopic particles by $A$ and $B$ and the microscopic ones by the discrete indices $n, n' = 1, \ldots, N$. Moreover, we denote by $V_P^{\Gamma,\Gamma}(\gamma, \Gamma)$ the corresponding interaction between particles $p$ and $q$, where the explicit expressions of $V_{\text{WCA}}, V_f$, and $V_m$ as described in (1), (2) and (19). $\mathcal{T}_{\text{meso}}(\gamma, \Gamma)$ and $\mathcal{T}_{\text{micro}}(\Gamma)$ indicate, respectively, the kinetic energies of the mesoscopic and microscopic particles. Therefore, $\mathcal{H}(\gamma, \Gamma)$ is separable into

$$\mathcal{H}(\gamma, \Gamma) = \mathcal{H}_{\text{meso}}(\gamma) + \mathcal{H}_{\text{micro}}(\gamma, \Gamma),$$

(B.4)

where $\mathcal{H}_{\text{meso}}(\gamma)$ contains the terms in line (B.1) and $\mathcal{H}_{\text{micro}}(\gamma)$ is composed of the terms in lines (B.2) and (B.3).

Since we work in the canonical ensemble, the physics of the system derives from the partition function

$$\mathcal{Z}(m) = \int e^{-\beta \mathcal{H}(\gamma, \Gamma)} d\gamma d\Gamma = \int e^{-\beta \mathcal{H}_{\text{meso}}(\gamma)} e^{-\beta \mathcal{H}_{\text{micro}}(\gamma, \Gamma)} d\gamma d\Gamma,$$

(B.5)

with $\beta = 1/k_B T$. Coarse-graining means to integrate out the microscopic degrees of freedom $\Gamma$, so we rearrange

$$\mathcal{Z}(m) = \int e^{-\beta \mathcal{H}_{\text{meso}}(\gamma)} \left[ \int e^{-\beta \mathcal{H}_{\text{micro}}(\gamma, \Gamma)} d\Gamma \right] d\gamma = \int e^{-\beta \mathcal{H}_{\text{meso}}(\gamma)} Z_{\text{micro}}(\gamma) d\gamma = \int e^{-\beta V_s^{AB}(\gamma)} e^{-\beta \mathcal{H}_{\text{micro}}(\gamma, \Gamma)} Z_{\text{micro}}(\gamma) d\gamma.$$

(B.6)

The connection to the probability density $p_\mathcal{Z}(\gamma)$ is given by

$$p_\mathcal{Z}(\gamma) = \frac{Z_{\text{micro}}(\gamma) e^{-\beta \mathcal{H}_{\text{micro}}(\gamma, \Gamma)} Z(m = 0)}{Z}. $$

(B.7)

Since the magnetic interactions only affect the mesoscopic particles, see line (B.1), it is solely contained in $\mathcal{H}_{\text{meso}}(\gamma)$. Therefore, the microscopic Hamiltonian $\mathcal{H}_{\text{micro}}(\gamma, \Gamma)$ for a fixed configuration $\gamma$ of the mesoscopic particles does not explicitly depend on the magnetic moments. The magnetic effects do not modify the physics of the polymer chain for a given configuration of the mesoscopic particles. The underlying physical reason is that the polymer chain does not contain magnetic components.

As we can see, the magnetic interactions only appear on the mesoscopic level of the final partition function $Z$. The magnetic interactions are simply included by multiplying on the mesoscopic level with the additional probability factor $\exp(-\beta V_m^{AB}(\gamma))$. The remaining part of the integrand that contains the information drawn from the MD simulations, is not affected.

Explicitly introducing the magnetic moments in the microscopic simulation would therefore not affect the final outcome.

Thus, it is sufficient to determine the effects of the polymer chain through MD simulations and add the magnetic interactions analytically in a subsequent step.

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