

Energy landscapes, structural topologies and rearrangement mechanisms in clusters of dipolar particles

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Clusters of spherical particles with isotropic attraction favour compact structures that maximise the number of energetically optimal nearest-neighbour interactions. In contrast, dipolar interactions lead to the formation of chains with a low coordination number. When both isotropic and dipolar interactions are present, the competition between them can lead to intricate knot, link and coil structures. Here, we investigate how these structures may self-organise and interconvert in clusters bound by the Stockmayer potential (Lennard-Jones plus point dipole). We map out the low-lying region of the energy landscape using disconnectivity graphs to follow how it evolves as the strength of the dipolar interactions increases. From comprehensive surveys of isomerisation pathways, we identify a number of rearrangement mechanisms that allow the topology of chain-like structures to interconvert.

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1 Introduction

The field of chemical topology¹ can be traced back to the first characterisation of interlocked organic rings and the introduction of the term catenane by Wasserman in 1960.² The two rings in a catenane are not chemically bonded, and the topology must therefore be specified to distinguish the structure from the unlinked rings, giving rise to the concept of topological isomers. Wasserman's short report also points out that a single cyclic hydrocarbon of sufficient length may, in principle, exist not only in the form of a simple loop but instead as a topologically distinct knot.

Since this early work, there has been considerable progress both in the theoretical understanding^{3,4} of “mechanically linked” cyclic molecules and in efficient methods for synthesising them.⁵ Knotted topologies have even been discovered in organic synthesis even when originally unexpected.⁶ The prevalence and importance of knots in biological macromolecules has also been recognised. Catenated and knotted forms of circular DNA naturally occur, for which topological isomers are enzymatically interconverted during replication, transcription, and recombination.^{7,8} Knot-like motifs arise in the native structure of a surprisingly large number of proteins. Although proteins are unbranched chains—and therefore have a topology that is formally trivial—considerable insight into the role of

knots can be gained by chemically joining the ends of the protein to make a cyclic topology that contains a genuine topological knot.⁹

Cyclic structures can also emerge from the self-assembly of particles that interact *via* non-covalent interactions. In particular, colloidal particles that carry a dipole moment tend to form chains that can close into loops,¹⁰ and the possibility of knotted topologies in such systems has been pointed out.¹¹ Highly intricate knotted topologies also arise in other types of colloids in a way that is conceptually different—not from chains of particles themselves, but from lines of defects in cholesteric liquid crystals, where twisted nematic order is disrupted by anchoring to spherical particles suspended in the solution.^{12,13} Once again, however, it is the anisotropic interactions between particles that leads to the existence of these lines and the relevance of topological considerations.

Global optimisation calculations have shown that knots, links and coils are energetically favourable structures for clusters of Stockmayer particles, which possess a permanent dipole plus an isotropic soft core and attractive tail.¹⁴ These intricate topologies arise from the competition between the dipole-dipole interactions, which promote chain formation, and the isotropic attraction, which favours compact, highly coordinated structures. The energetically optimal compromise is often a continuous closed-loop chain, which is entwined with itself or with another loop in order to increase the number of nearest-neighbour contacts.

Most of the literature concerning the Stockmayer model has addressed the structure and thermodynamics of the bulk fluid phase. In this context, too, the interplay between chains and compact structures is crucial. van Leeuwen and Smit were

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amongst the first to point out that gas–liquid phase separation is suppressed if the dipolar attraction dominates.¹⁵ ten Wolde *et al.* showed that droplets of Stockmayer particles appear from the vapour phase first by formation of chains and then by collapse of the chains into globules that nevertheless retain a chain-like structure.¹⁶ The more general field of dipolar fluids, including the Stockmayer model, has been summarised in an informative review by Teixeira *et al.*¹⁷ There has been relatively little work on finite clusters bound by the Stockmayer potential,¹⁸ though the model has now been recognised as having applications in self-assembly.¹⁹

The potential for a cluster of N Stockmayer particles takes the form

$$V = \varepsilon \sum_{i < j}^N \left\{ 4 \left[\left(\frac{\sigma}{r_{ij}} \right)^{12} - \left(\frac{\sigma}{r_{ij}} \right)^6 \right] + \frac{\mu^2 \sigma^3}{r_{ij}^3} \left[\hat{\mu}_i \cdot \hat{\mu}_j - \frac{3}{r_{ij}^2} (\hat{\mu}_i \cdot \mathbf{r}_{ij}) (\hat{\mu}_j \cdot \mathbf{r}_{ij}) \right] \right\}, \quad (1)$$

where \mathbf{r}_{ij} is the position vector of particle i with respect to particle j , $\hat{\mu}_i$ is a unit vector along the dipole moment of particle i , ε and σ are the Lennard-Jones (LJ) units of energy and distance, respectively, and μ is a dimensionless parameter that determines the relative strength of the dipolar and LJ components of the potential. As μ increases and the dipolar contribution to the energy dominates, the Lennard-Jones well depth ε is no longer a convenient unit of energy. Here, we will report energies in units of the well-depth ε^* of the full Stockmayer pair potential for parallel head-to-tail dipole vectors at the relevant value of μ .

The structure of the clusters is tuned by the dipole moment strength μ . In the limit of a weak dipole (small μ), compact, icosahedral structures are favoured.²⁰ For strong dipoles the particles behave as dipolar soft spheres, producing chains, rings, and branched clusters, such as those formed by assemblies of ferromagnetic nanoparticles.²¹ It is in the intermediate regime, where competition between the isotropic and directional contributions leads to frustration, that interesting topologies with as many as ten irreducible crossings may arise.¹⁴

The current work builds on these findings concerning energetically optimal structures by making a broader survey of the energy landscape for a selection of cases. The analysis presented here enables us to characterise the overall evolution of the energy landscape between the Lennard-Jones and dipole-dominated limits, and to identify pathways and rearrangement mechanisms by which different morphologies interconvert. Some of the key properties can be observed in the small St_{13} cluster (where St_N denotes an assembly of N Stockmayer particles). The analysis is then extended to more complex rearrangements of knotted morphologies in the St_{21} and St_{38} clusters.

2 Methods

2.1 Optimisation

A Stockmayer particle consists of a single interaction site with an isotropic Lennard-Jones component and an anisotropic point dipole. Since the dipole has cylindrical symmetry, each particle has five degrees of freedom: three translational and two

rotational. In previous work¹⁴ on the global optimisation of Stockmayer particles, the translational coordinates were represented in Cartesians and the orientation of the dipole was described using the polar angle θ and azimuthal angle ϕ .

The present work requires optimisation of first-order saddle points and the calculation of pathways, as well as characterisation of minima on the potential energy surface (PES). For such applications it is more convenient to represent the orientation of the particles using the general angle-axis framework,^{22,23} where the orientation of a rigid body is obtained by rotating a particle from a reference orientation about an axis specified by the rotation vector \mathbf{p} through an angle given by its magnitude $|\mathbf{p}|$. Although this representation introduces a redundant sixth degree of freedom for each particle in this case, it has the advantage that the components of \mathbf{p} can be treated as unbounded variables, unlike θ and ϕ . The redundant degrees of freedom contribute additional zero eigenvalues to the Hessian matrix; the corresponding eigenvectors can be obtained analytically²² and are used for projection while characterising the transition state as described later.

In the case of Stockmayer particles, we arbitrarily took the reference orientation of the dipole along the laboratory-fixed z axis. The unit dipole vector $\hat{\mu}_i$ of a particle can then be obtained from the angle-axis variables \mathbf{p}_i . All angular first and second derivatives of V can then be written with respect to the components of \mathbf{p}_i .

2.2 Calculating pathways

A potential energy landscape can be characterised by mapping out its local minima, which correspond to locally stable inherent structures,²⁴ and their connections *via* first-order saddle points (transition states).

The survey of an energy landscape typically starts from a small selection of energetically low-lying minima obtained from a basin-hopping^{25,26} global optimisation calculation. To find the pathways that connect pairs of these structures,²⁷ we employ the doubly-nudged elastic band method,²⁸ as implemented in the OPTIM package.²⁹ In this approach, a discretised interpolation is first established between the two minima in the full configuration space. The path is then allowed to relax on the PES while the states are kept approximately evenly spaced along the path by springs. Local maxima on the path are then tightly converged to transition states using hybrid eigenvector-following,^{30–32} and the connectivity is determined by approximate steepest-descent minimisations, adding any new minima thus found to the database of structures. It is common for new minima to be identified during this procedure and so the first iteration of attempted connections rarely produces a fully connected path between the minima that were originally specified.

Missing sections on a path are approached in the same way. Candidate pairs of minima to be bridged are selected by repeated application of the missing connection algorithm,²⁷ also implemented in the OPTIM package,²⁹ until a connected sequence of minima and transition states (a discrete path) is found. To characterise complete paths, the minima are considered to be nodes on a graph and the weight of the edge



between two minima is set to zero if they are already directly connected by a single transition state in the database. If no connection is known, the weight is set to a function of the shortest Euclidean distance between the minima.²⁷ Hence, the algorithm identifies breaks in the path that are short in configuration space, making it more plausible that a connection will be found. The Euclidean distance between different Stockmayer clusters was determined from the positions of the particles only, ignoring the orientations of the dipole moments.

2.3 Discrete path sampling

Once an initial path has been established, the stationary point database must be expanded in order to obtain a more complete picture of the landscape. For small systems, a comprehensive survey of the landscape is possible by repeatedly attempting to connect all pairs of minima already in the database by the method described in Section 2.2 until no new minima or pathways emerge. This approach was employed for the St₁₃ clusters.

For clusters even just a few particles larger, it is not feasible to obtain exhaustive lists of stationary points. It is then most informative to obtain a good representation of the regions surrounding the lowest-lying minima of each structural family and a set of kinetically relevant pathways between them. Algorithms for connecting regions of configuration space in this way using discrete path sampling^{33–35} (DPS) have been described in detail in previous work.^{36,37} Consider two regions on the landscape, A and B, which contain the minima we wish to connect. Minima in neither region that nevertheless appear on pathways between A and B minima belong to the intervening region, denoted I. Assuming Markovian dynamics, that local equilibrium is reached within the A and B sets, and that the steady-state approximation applies to minima in the I set, the steady-state rate constants k_{BA}^{SS} and k_{AB}^{SS} can each be written as a sum over discrete paths.³³ In turn, the contribution of a given discrete path can be expressed in terms of the individual rate constants k_{ij} for each successive pair (i, j) of minima along the path.

Since we are concerned with characterising the PES rather than obtaining quantitative values for the rate constants, we estimate the single-step rate constants k_{ij} using a simplified version of harmonic transition state theory, where the ratio of normal mode frequencies at the transition state and the minimum is set to unity. This simplification has the advantage of making it unnecessary to consider the relationship between the vibrational and librational contributions to the normal modes of the Stockmayer particles. To proceed otherwise would require specialisation of the model by choosing a particular moment of inertia for the particles. With this simplification, the expression for the rate constant reduces to

$$k_{ij} = \frac{o_i}{o_{ij}^\ddagger} \exp\left[-(V_{ij}^\ddagger - V_i)/(k_B T)\right],$$

where o_i denotes the order of the point group for minimum i , and the superscript \ddagger is used to refer to the transition state. For the calculation of rate constants, we work at a temperature of $k_B T/\epsilon^* = 1/30$, which is the regime probed by some recent experiments in which rings of magnetic dipolar colloids have been observed.²¹

Using the simplified rate constants for individual steps, the discrete path that makes the largest contribution to k_{BA}^{SS} (the fastest path) can be identified. Attempts are then made to shortcut this path by directly connecting minima that lie on it but are separated by intervening minima. The choice of minima can be made on the basis of their separation along the path or in Euclidean space, or from the height of the barrier that separates them. These schemes guide the exploration of the PES towards kinetically relevant minima and transition states, and are here applied to the more complex landscapes of the St₂₁ and St₃₈ clusters.

It is possible for artificial kinetic traps to arise as the database is generated—that is, for minima to be connected to the A and B regions by large barriers, where smaller barriers exist but have not yet been found, resulting in spuriously small rate constants. As the database grows, we periodically attempt to remove such traps by identifying pairs of minima based on the ratio of the potential energy barrier (from minimum to transition state) to the potential energy difference between the two minima connected by the transition state.³⁷ These pairs are then subjected to reconnection searches as described above. Such ‘untrapping’ cycles were performed for all databases until the low-energy region of the landscape converged.

The methods described above are implemented in the program PATHSAMPLE,³⁸ which is a driver for OPTIM,²⁹ as well as a tool for analysis of the resulting kinetic transition networks.

2.4 Topological characterisation

To determine the topology of a given structure, it is first necessary to trace chains of head-to-tail dipoles within the cluster. An intuitive and robust method for identifying connectivity has been described by Miller and Wales.¹⁴ Beginning with a particle i , the next particle in the chain is the one with the lowest (most favourable) dipole–dipole interaction energy with i , located in the half-space into which the dipole at i points. Similarly, the previous particle is the one with the lowest interaction energy in the complementary half-space. Connections in the chains that this procedure identifies will be referred to here as ‘bonds’.

In some structures, the bonds define a closed-loop chain that incorporates all the particles. If this loop cannot be unravelled into a trivial circle containing no crossings without breaking any bonds, then the structure is a knot. A knot is classified in the Rolfsen notation³⁹ according to the two-dimensional projection of the chain that contains the smallest possible number of crossings. For the structures encountered in this work, the topology can be determined from an arbitrary projection by considering all combinations of splittings at the crossings and thereby evaluating the knot’s Jones polynomial.⁴⁰ The Rolfsen symbol can then be determined from tables.⁴¹

3 Results and discussion

3.1 St₁₃ clusters

For St₁₃, four structures successively become the global minimum as μ is increased from zero: a distorted icosahedron,



with D_3 symmetry; a hexagonal antiprism, with D_{6d} symmetry; stacked rings of six and seven particles, with C_s symmetry; and a planar thirteen-particle ring, with D_{13h} symmetry. The point groups here refer to symmetry operations that map the particle positions onto each other without considering the direction of the dipoles.

Databases of minima and transition states have been constructed for the three values of μ at which the global minimum changes, 1.42, 2.54, and 2.66, to show competition between the different morphologies, as well as at $\mu = 0, 0.2, 1$, and 3.6 to investigate the evolution of the potential energy landscape as a function of dipole strength.

The corresponding disconnectivity graphs are presented in Fig. 1. For $\mu = 0$, the expected single-funnel 'palm tree' motif⁴² is reproduced (Fig. 1a). This nomenclature refers to the fact that the branches of the graph are vertically rather short on the energy scale of the pairwise potential and that they are almost all directly connected to the central stem representing the basin of the global minimum. These features are indicative of

multiple sequences of minima separated by relatively low barriers, converging on the lowest-energy structure.

Introduction of a weak dipole causes a rapid increase in the number of minima on the landscape, owing to the multiplicity of stable arrangements of dipoles for each LJ morphology that are easily interconvertible *via* low barriers (Fig. 1b). Aside from the bifurcation of the branches that these arrangements cause, the form of the graph remains the same. When μ is increased to 1, elements of this splitting are still present (Fig. 1c). Two stable structures with practically identical geometries and energies, but different arrangements of dipoles, lie substantially lower in energy than any other structure. Morphologies that do not exist on the LJ landscape have begun to emerge and, in terms of the optimal pair energy ε^* , the funnel appears to be shallower.

At $\mu = 1.42$, the global minimum changes from the distorted icosahedron to the hexagonal antiprism (Fig. 1d). The disconnectivity graph is still, broadly speaking, a palm tree but now has two low-lying minima that are separated by a substantial barrier. The lowest-energy single-step rearrangement in the

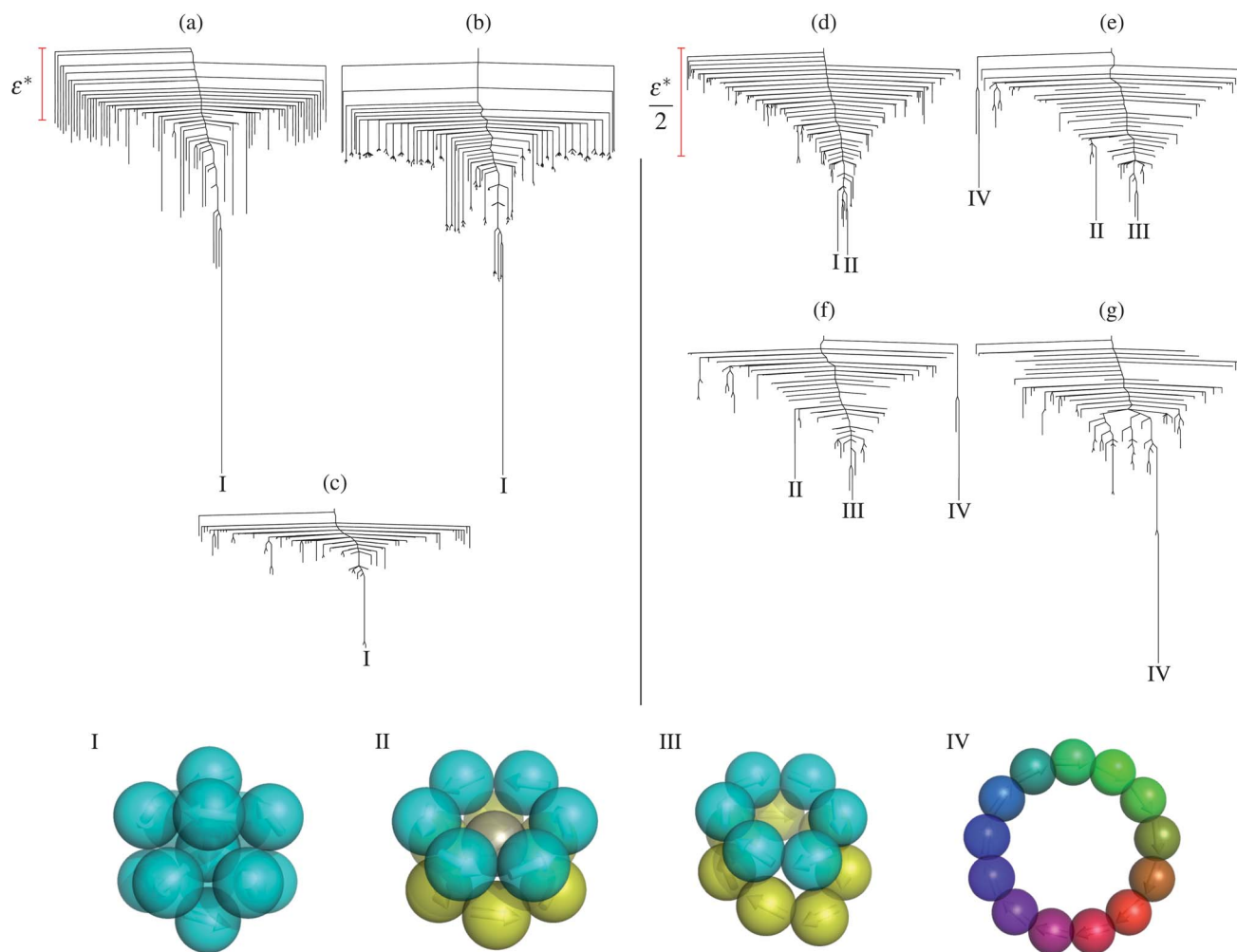


Fig. 1 Upper panel: disconnectivity graphs for St_{13} at $\mu =$ (a) 0; (b) 0.2; (c) 1; (d) 1.42; (e) 2.54; (f) 2.66; (g) 3.6. Branches to the 100 lowest energy minima are shown [except in (b) where 300 are shown for proper comparison with (a)]. Lower panel: global minima for St_{13} . From left to right, the distorted icosahedron, the hexagonal antiprism, stacked rings, and the single ring. Particles are drawn as translucent spheres with arrows pointing along the dipole vector. Different chains are distinguished by their colour. Single chains are depicted with smoothly changing colour.



database that connects the corresponding minima is presented in Fig. 2. The coordinate s in this plot is the integrated distance along the pathway between the initial and final structures in the $3N$ -dimensional Euclidean space of the particle positions. The two portions of a pathway from a transition state to its directly connected minima are defined by the approximate steepest-descent minimisations initiated from the transition state along the unstable mode. The orientations of the dipole are not included in s and pure rotations of dipoles therefore do not contribute to the apparent length of the path. The interconversion mechanism is the well-known diamond-square-diamond rearrangement (DSD), proposed by Lipscomb,⁴³ which describes rearrangements of boranes,⁴⁴ carboranes,⁴⁵ and metal-laboranes.^{46,47} It is also an important mechanism for LJ clusters.⁴⁸ The pathway contains two consecutive DSD rearrangements, which successively cleave both rings of the hexagonal antiprism at diametrically opposite points. To visualise this mechanism it is helpful to think of the hexagonal antiprism as a ring of twelve edge-sharing triangles. The DSD rearrangement rotates two opposite segments, each of four triangles, with respect to one another about a line connecting their centres. In the final structure (Fig. 2d) the two segments now clasp each other in a relative orientation that is orthogonal to where they started (Fig. 2a).

Dipolar 'bonding' adds a new dimension to the DSD rearrangement, illustrated in Fig. 3a. Pairs of dipoles begin aligned along opposite edges of the diamond face. As the long diagonal contracts and the short diagonal elongates, the approaching dipoles reorient so that in the product they are aligned along the new short diagonal. The effect is to 'break' the bonds along the

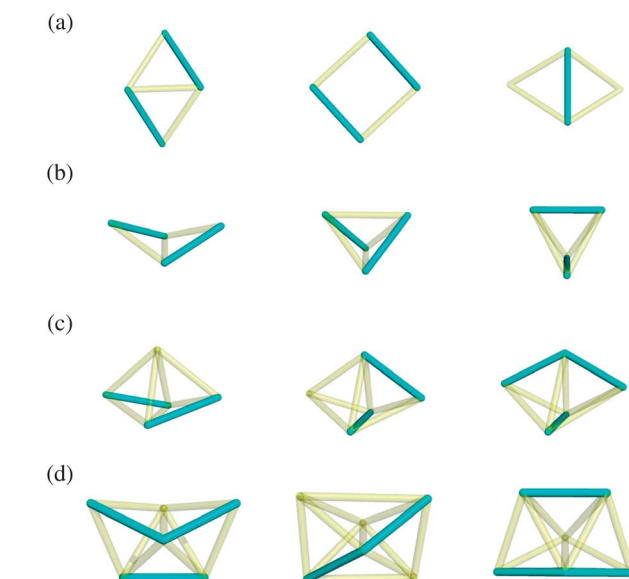


Fig. 3 Rearrangement mechanisms for St_{13} clusters as minimum-transition state-minimum triples: (a) the diamond-square-diamond (DSD) rearrangement; (b) the butterfly-tetrahedron (BT_d) rearrangement; (c) the closed butterfly-double tetrahedron ($BT_dT_d^2$) rearrangement; (d) the double BT_d (BT_dBT_d) rearrangement. Light tubes show the underlying geometry, and dark tubes show dipole bonds.

edges and create a bond along the diagonal of the diamond. In the rearrangement between the distorted icosahedron and hexagonal antiprism of St_{13} , this process breaks one bond in each ring and forms an interchain contact between two of the free ends. Along the pathway there exists a single twelve-particle helix (Fig. 2c) wrapped around the central particle on the downhill path to the distorted icosahedron, which has three closed loops of dipoles.

At $\mu = 2.54$, the global minimum changes again, from the hexagonal antiprism to two stacked rings (Fig. 1e). Compared to the first change of global minimum at $\mu = 1.42$, the landscape at this second change has more double-funnel character, since each of the two low-lying minima has a number of other minima associated with its main branch. In addition, the thirteen-membered planar ring, which is the global minimum in the strong dipole limit, now appears as a metastable structure connected to the main stem of the graph at high energy. The associated region of the energy landscape may be considered as a third funnel.

The pathway connecting the lowest two minima is more complex than at the previous change, containing five transition states, and traversing a variety of links and coils. Starting from the stacked rings (structure III in Fig. 1), the first three steps convert the unlink to a coil, the coil to a link, and finally the link to a different coil. Each of these transformations occurs by the butterfly-tetrahedron rearrangement (BT_d), proposed by Wales *et al.*⁴⁹ It is somewhat similar to Johnson's edge-bridging mechanism⁵⁰ in terms of the centres of mass: a butterfly moiety closes to form a tetrahedron. However, in the context of dipolar particles, this underlying structural change allows recombination of chains, interconverting topological isomers. This

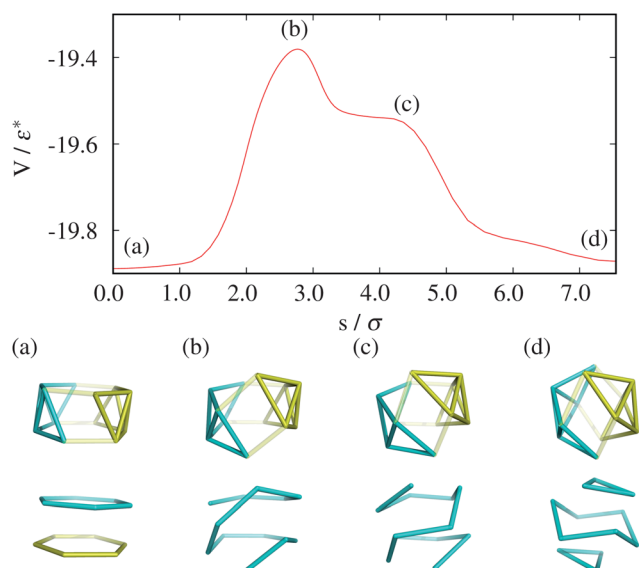


Fig. 2 Upper panel: a path between the hexagonal antiprism and the distorted icosahedron for the St_{13} cluster at $\mu = 1.42$. The graph shows the energy divided by the pair energy, V/ϵ^* , as a function of the integrated path length, s . Lower panel: structures at labelled points on the path: top, in terms of the dipolar particles discussed in the text, and bottom, the dipole network. Note that the shoulder at (c) is not a transition state. (The central particle, which does not participate in dipole bonding, is not shown.)

process is illustrated in Fig. 3b. Similarly to the DSD rearrangement, dipoles begin aligned along opposite edges of the butterfly. In contrast to the DSD process, the long diagonal contracts, but the short diagonal does not elongate significantly. As the incoming particles approach, the dipoles reorient to be along the interparticle vector, and a bond forms along the nascent tetrahedral edge. Simultaneously the dipoles at the hinge vertices align along the opposite edge. In this fashion, parallel chain segments are converted into perpendicular ones, creating a crossing in the chain of dipoles while keeping the number of dipole bonds conserved.

The remaining steps of the interconversion from stacked rings to hexagonal antiprism involve rearrangements related to BT_d . First, a coil is converted to a structure containing a five-membered ring and a seven-membered ring with a particle in the centre (5,1,7). This process is difficult to describe in purely geometrical terms, but insight may be gained by considering which bonds are formed or broken in the process. This analysis allows us to describe what we will call the closed butterfly–double tetrahedron rearrangement ($B_cT_d^2$), illustrated in Fig. 3c. We begin with a face-capped or ‘closed’ butterfly arrangement of particles, where bonds are formed along the same edges as are found in the BT_d process. The butterfly closes around this particle, one wing at a time, forming bonds between the wingtips and the capping particle, and along the hinge. The result can be thought of as two face-sharing tetrahedra. Comparison of Fig. 3c and b reveals the similarity in the overall changes to the dipole network. In this way a particle is released from two face-sharing tetrahedra in the coil, which becomes the central particle in the 5,1,7 ring system.

The final step, in which the 5,1,7 system is converted to the hexagonal antiprism, is a concerted double BT_d , or BT_dT_dB , rearrangement, illustrated in Fig. 3d. In terms of the particle centres, a butterfly and a tetrahedron sharing the hinge edge simultaneously close and open, interconverting. It differs from the other examples in that the butterfly’s dipole bonds are not opposite each other, but adjacent, and the tetrahedra have bonds only along the edge that is broken in the butterfly. In the context of the St_{13} structure, the rearrangement of this fragment effects a particle exchange between the rings. At the transition state of the whole structure, none of the tetrahedral bonds and one butterfly bond for each of the rings exist, forming a twelve-particle helix.

The global minimum in the high-dipole limit is a single thirteen-particle ring. This structure first replaces the stacked rings as the Stockmayer global minimum at $\mu = 2.66$. The disconnectivity graph (Fig. 1f) shows distinct funnels for the two competing structures, but the stacked ring funnel is associated with a larger region of configuration space. As μ increases, and the ring funnel becomes increasingly favourable, the region of configuration space associated with the stacked rings is expected to act as a kinetic trap to structural relaxation.

A pathway containing three transition states, which connects these competing minima, is presented in Fig. 4. The first two steps flatten the ring into an ellipse and twist it about its centre to form a coil. Although much of the bending energy penalty is removed, and favourable contacts are made between particles in

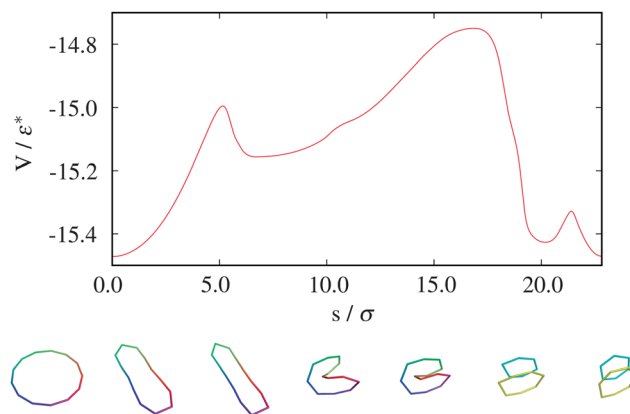


Fig. 4 Upper panel: a path between the ring and stacked rings minima of the St_{13} cluster at $\mu = 2.66$. Lower panel: structures of minima and transition states along the path.

the adjacent chains, the sharp bends at the ends of the ellipse are sufficient to increase its energy with respect to the ring. As the ellipse twists and folds into a coil, it is locked in place by a mechanism reminiscent of a DSD rearrangement, which is illustrated in Fig. 5a. The final step in the path is the now familiar BT_d rearrangement, which interconverts the coil and the stacked rings, removing a crossing from the chain. It is interesting to note that in the previous pathway, at a smaller value of μ , all of the BT_d steps lowered the energy upon forming the more compact tetrahedral site. In the current pathway, the situation is reversed, reflecting the increasing stability of planar moieties as the dipolar contribution to the energy increases.

At large μ , the ring is firmly established as the global minimum. It is separated from the next lowest minimum, the ellipse, by $0.59\epsilon^*$, and from the third lowest energy structure by $0.78\epsilon^*$ at 3.6μ .

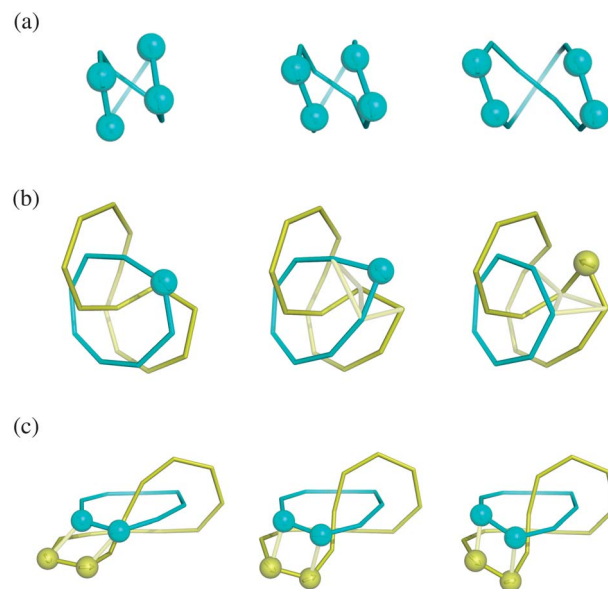


Fig. 5 Chain-based mechanisms: (a) a diamond–square–diamond rearrangement; (b) a budding rearrangement; and (c) a DSD rearrangement that facilitates chains moving past one another.



3.2 St_{21} clusters

The most remarkable structures that appear as global minima for Stockmayer clusters are knotted geometries.¹⁴ The smallest cluster where a knot becomes energetically optimal is St_{21} . The global minimum of this cluster is topologically a trefoil knot over the range $1.7 \leq \mu < 2.9$, beyond which a pair of stacked twelve and thirteen particle rings becomes more favourable. The disconnectivity graph at $\mu = 2.9$ is presented in Fig. 6, and exhibits a pronounced double funnel. As with the graph at $\mu = 2.66$ for St_{13} , the wider funnel is associated with the more compact structure—in this case, the knot.

A low-energy pathway connecting these structures is presented in Fig. 7. The pathway contains eleven transition states, and mostly involves rearrangement of, and particle exchange between, rings in a link. The first such conversion, from an unlink with ten and eleven particle rings to a link with nine and twelve particle rings, is achieved in three transition states. First, the unlink rearranges to a coil by the BT_d mechanism. This coil contracts to a more compact bundle in a similar fashion to the St_{13} ring. The coil itself then twists, facilitated by coupled inter-chain DSD rearrangements, which enable chains to move past one another. Finally, the coil converts to the nine- and twelve-particle link by a second BT_d rearrangement, which conserves a twist in the larger ring (Fig. 7c).

Two further exchanges, to form links of eight- and thirteen-particle rings and then the link of seven- and fourteen-particle rings, occur by a new mechanism, whereby a particle is smoothly transferred between adjacent chains. This ‘budding’ rearrangement is illustrated in Fig. 5b. A three-particle section

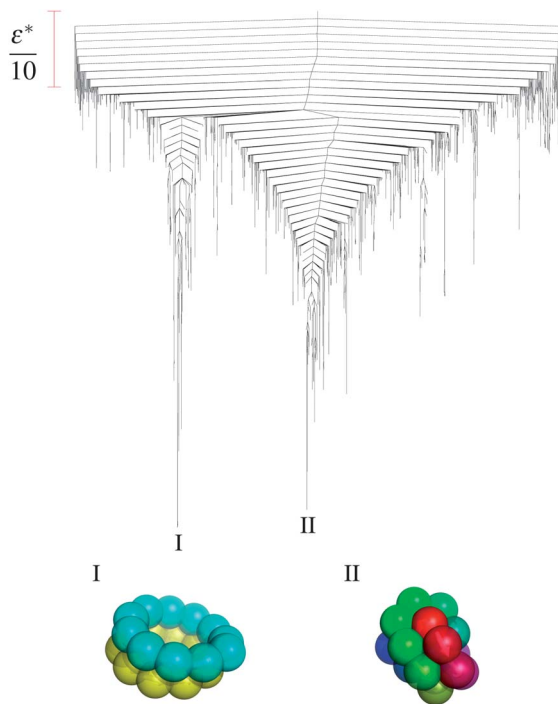


Fig. 6 Upper panel: disconnectivity graph for St_{21} at $\mu = 2.90$. Branches to the 1000 lowest energy minima are shown. Lower panel: structures of the stacked ring (I) and trefoil knot (II) minima.

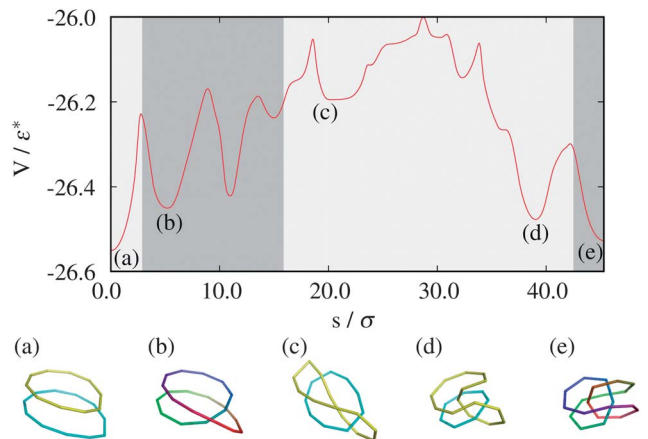


Fig. 7 Upper panel: a pathway between the stacked rings and trefoil knot minima of St_{21} . The figure is shaded according to the number of chains in the structure (light = 1, dark = 2). Lower panel: selected structures from the path: (a) the stacked rings; (b) a coil; (c) a link (9 and 12 particles); (d) a link (7 and 14 particles); and (e) the trefoil knot.

of a chain bends sharply, expelling a particle and forming a new connection between the free ends. The released particle is incorporated into an adjacent chain, which opens as the first one closes. At the transition state, the five sites are arranged in a closed butterfly geometry. The process can be thought of as an edge-bridging particle passing to the opposite edge *via* a face-bridged transition state. The latter exchanges are separated by coiling of the larger ring around the smaller by means of DSD rearrangements, shown in Fig. 5c, and the rings are disposed such that a final BT_d rearrangement produces the trefoil knot.

3.3 St_{38} clusters

The LJ_{38} cluster is unusual in that the global minimum is a truncated octahedron, and not based on an icosahedron.⁵¹ The landscape is double-funneled,⁵² and the lowest energy icosahedral minimum, while significantly higher in energy than the global minimum, is associated with a much larger region of configuration space. As such, the system provides a useful test of global optimisation procedures.

St_{38} exhibits a more complex knotted global minimum than the St_{21} trefoil in the range $1.6 < \mu < 2.2$.¹⁴ This St_{38} knot has eight crossings in its minimal projection and has the topology 8_{19} in Rolfsen's notation.³⁹

The Lennard-Jones truncated octahedron ceases to be the global minimum at $\mu = 0.8$. For values of μ between 0.8 and 1.6, the truncated octahedron and knot—both metastable—are close in energy. Databases of minima and transition states have been constructed at $\mu = 1.22$, where the two morphologies are closest in energy. The disconnectivity graph is presented in Fig. 8. It has the form of a gently sloping single funnel, with a form somewhere between a ‘weeping willow’ and a palm tree.⁵³ (The willow tree has longer vertical branches connected to the main stem, indicating multiple descending pathways towards the global minimum, like the palm tree, but with higher barriers between adjacent minima.) The global minimum has D_{4h} symmetry and is depicted in Fig. 8.



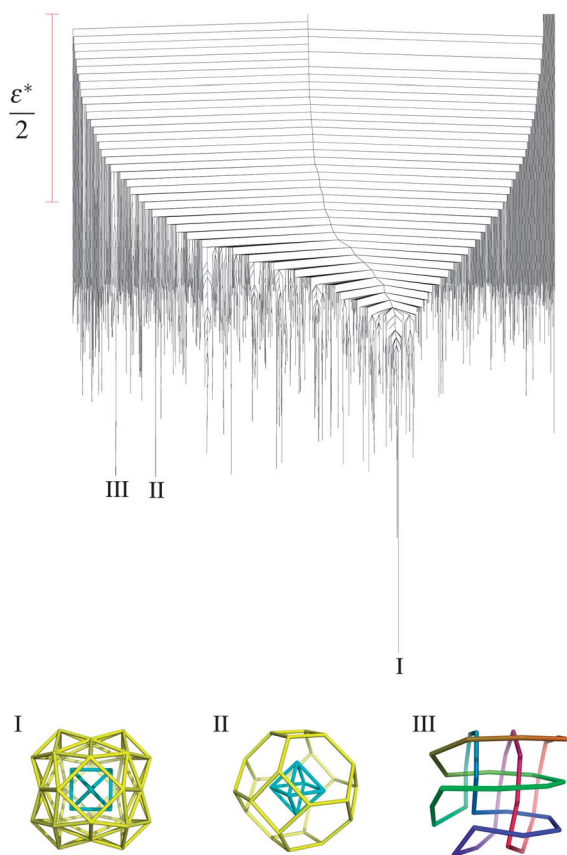


Fig. 8 Upper panel: disconnectivity graph for St_{38} at $\mu = 1.22$. Branches to the 1000 lowest energy minima are shown. Lower panel: the D_{4h} global minimum (I) and the O_h fcc minimum (II), showing the core-shell structure, and the 8_{19} knot minimum, showing the dipole chain.

The truncated octahedron and knot structures can be interconverted by a pathway containing just three transition states. Multiple concerted surface rearrangements occur in the first two steps, an example of which is presented in Fig. 9a. The

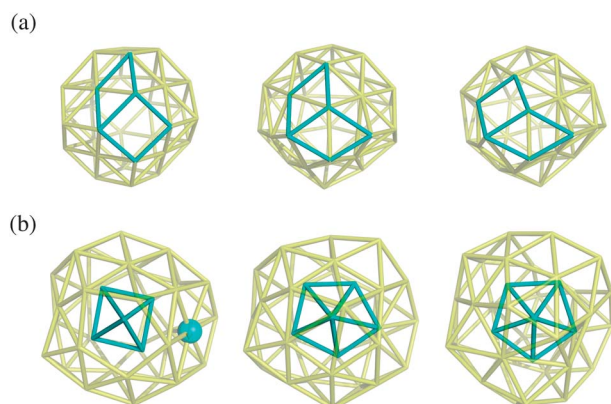


Fig. 9 Steps in the rearrangement of St_{38} from a truncated octahedron (structure II in Fig. 8) to an 8_{19} knot (structure III). (a) A square–diamond–diamond–square (SDDS) rearrangement on the surface of the truncated octahedron (structure II in Fig. 8). (b) A core–shell rearrangement in the final step of the pathway. Darker shading highlights the geometry of the rearranging sites.

square faces of the truncated octahedron minimum permit the previously characterised⁵⁴ square–diamond–diamond–square (SDDS) mechanism to occur. The final step is an inter-shell transfer of a particle from the surface to the core, expanding the octahedron to a pentagonal bipyramid (Fig. 9b). Since we are considering a relatively weak dipole, the absence of dipole-dominated mechanisms, such as BT_d , is not surprising. The 8_{19} knot is still compact, but dipole bonding stabilises the open faces required to arrange the particles in such a way that the dipoles may orient to form a knot.

4 Conclusions

Clusters of spherical particles bound by simple isotropic potentials like Lennard-Jones and Morse are generally dominated by compact—often icosahedral—structures. The anisotropic Stockmayer potential differs from such cases because of the particles' tendency to form chains. The fact that chain-like motifs can readily be identified introduces the new consideration of topology when characterising structure and rearrangements. Since the connectivity of particles within the chains is not fixed (as it is in a polymer), rearrangements may alter the topology.

Some rearrangement mechanisms in Stockmayer clusters resemble those found in clusters bound by other potentials. For example, the diamond–square–diamond rearrangement, which is common in clusters bound by isotropic potentials, also occurs in Stockmayer clusters. However, this familiar mechanism gains additional significance because of the change in orientation of the dipole vectors, which can allow chains to be disconnected in one direction and to become connected in another. Similarly, edge-bridging mechanisms seen in other clusters can change the topology of Stockmayer clusters.

In addition to mechanisms that change the connectivity between chains while preserving the association of each particle with its chain, there are budding processes. Here, a particle is smoothly ejected from one chain and absorbed into another, thereby allowing chains to change in size.

These mechanisms vary in importance with the strength of the dipole moment. A crucial consideration is whether or not the pathway conserves the number of head-to-tail connections between dipoles. The diamond–square–diamond mechanism exchanges two parallel connections for one in an orthogonal orientation, and so is more favourable at lower dipole strengths, where the loss of a dipole–dipole bond does not incur a large energetic penalty. The butterfly–tetrahedron mechanism and some of its variants preserve the number of bonds while interconverting topological isomers, and become more important at higher dipole strengths.

Stockmayer clusters are frustrated systems in which the isotropic Lennard-Jones part of the potential drives the structure towards compact highly coordinated arrangements, while the dipolar interactions favour chain-like motifs. The analysis of structures and rearrangement mechanisms that we have presented here for St_{13} , St_{21} , and St_{38} illustrates a variety of topology-changing events that offer a compromise between the competing terms in the potential.



Experimentally, clusters of colloids with a dipolar component to their interactions are being studied with increasing levels of control, and are being characterised in greater detail. While chain- and loop-based structures currently predominate, it should be possible to explore more compact clusters as well. The knotted topologies and other interesting structures that arise for Stockmayer particles are obtained by adjusting the relative strengths of the dipolar and isotropic parts of the potential. This flexibility can be achieved in suspensions of dipolar colloids by introducing tunable, depletion-induced, isotropic attraction *via* an additional component in the suspension.⁵⁵ The isotropic attraction can also be controlled through the surface chemistry of the colloids.⁴¹ In both cases, the repulsive core of the particles is likely to be considerably harder than the r^{-12} term in eqn (1). However, it is the presence of an isotropic attractive tail and its competition with the dipolar interactions that are crucial in generating a rich selection of topologies.

A number of open questions remain in the context of clusters and molecules with non-trivial topology. These issues include the maximum complexity (number of crossings) that can be attained in a given system, and the precise factors determining the favoured topology where a choice exists. The influence of the interparticle potential and any fixed connectivity between particles will be important in this regard, and these considerations make contact with other knot-forming systems, such as proteins and synthetic organic molecules. The study of idealised model systems, such as the work presented here, should be a fruitful way of gaining insight into the general phenomenon of knot formation.

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References

- H. L. Frisch and E. Wasserman, *J. Am. Chem. Soc.*, 1961, **83**, 3789–3795.
- E. Wasserman, *J. Am. Chem. Soc.*, 1960, **82**, 4433–4434.
- M. L. Mansfield and J. F. Douglas, *J. Chem. Phys.*, 2010, **133**, 044903.
- M. L. Mansfield and J. F. Douglas, *J. Chem. Phys.*, 2010, **133**, 044904.
- G. Barin, R. S. Forgan and J. F. Stoddart, *Proc. R. Soc. A*, 2012, **468**, 2849–2880.
- N. Ponnuswamy, F. B. L. Cougnon, J. M. Clough, G. D. Pantos and J. K. M. Sanders, *Science*, 2012, **338**, 783–785.
- S. Wasserman and N. Cozzarelli, *Science*, 1986, **232**, 951–960.
- D. Meluzzi, D. E. Smith and G. Arya, *Annu. Rev. Biophys.*, 2010, **39**, 349–366.
- A. L. Mallam, J. M. Rogers and S. E. Jackson, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **107**, 8189–8194.
- M. Klokkenburg, R. P. A. Dullens, W. K. Kegel, B. H. Ern  and A. P. Philipse, *Phys. Rev. Lett.*, 2006, **96**, 037203.
- P. Y. Keng, I. Shim, B. D. Korth, J. F. Douglas and J. Pyun, *ACS Nano*, 2007, **1**, 279–292.
- V. S. R. Jampani, M. Skarabot, M. Ravnik, S. Copar, S. Zumer and I. Musevic, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2011, **84**, 031703.
- U. Tkalec, M. Ravnik, S. Copar, S. Zumer and I. Musevic, *Science*, 2011, **333**, 62–65.
- M. A. Miller and D. J. Wales, *J. Phys. Chem. B*, 2005, **109**, 23109.
- M. E. van Leeuwen and B. Smit, *Phys. Rev. Lett.*, 1993, **71**, 3991–3994.
- P. R. ten Wolde, D. W. Oxtoby and D. Frenkel, *J. Chem. Phys.*, 1999, **111**, 4762–4773.
- P. I. C. Teixeira, J. M. Tavares and M. M. Telo da Gama, *J. Phys.: Condens. Matter*, 2000, **12**, R411–R434.
- H. B. Lavender, K. A. Iyer and S. J. Singer, *J. Chem. Phys.*, 1994, **101**, 7856–7867.
- K. Van Workum and J. F. Douglas, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2006, **73**, 031502.
- M. R. Hoare, *Adv. Chem. Phys.*, 1979, **40**, 49–135.
- W.-F. Ding, Z. Li, H. Zhou, B. Zhao, J.-g. Wan, F. Song and G.-H. Wang, *J. Phys. Chem. C*, 2012, **116**, 10805–10813.
- D. Chakrabarti and D. J. Wales, *Phys. Chem. Chem. Phys.*, 2009, **11**, 1970–1976.
- D. J. Wales, *Philos. Trans. R. Soc. London, Ser. A*, 2005, **363**, 357–377.
- F. H. Stillinger and T. A. Weber, *Phys. Rev. A*, 1983, **28**, 2408–2416.
- D. J. Wales and J. P. K. Doye, *J. Phys. Chem. A*, 1997, **101**, 5111–5116.
- D. J. Wales, *GMIN: A program for basin-hopping global optimisation*, <http://www-wales.ch.cam.ac.uk/software.html>.
- J. M. Carr, S. A. Trygubenko and D. J. Wales, *J. Chem. Phys.*, 2005, **122**, 234903.
- S. A. Trygubenko and D. J. Wales, *J. Chem. Phys.*, 2004, **120**, 2082–2094.
- D. J. Wales, *OPTIM: A program for geometry optimisation and pathway calculations*, <http://www-wales.ch.cam.ac.uk/software.html>.
- G. Henkelman and H. J nsson, *J. Chem. Phys.*, 1999, **111**, 7010–7022.
- L. J. Munro and D. J. Wales, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1999, **59**, 3969–3980.
- Y. Kumeda, D. J. Wales and L. J. Munro, *Chem. Phys. Lett.*, 2001, **341**, 185–194.
- D. J. Wales, *Mol. Phys.*, 2002, **100**, 3285–3306.
- D. J. Wales, *Mol. Phys.*, 2004, **102**, 891–908.
- S. A. Trygubenko and D. J. Wales, *Mol. Phys.*, 2006, **104**, 1497–1507.
- J. M. Carr and D. J. Wales, *J. Chem. Phys.*, 2005, **123**, 234901.
- B. Strodel, C. S. Whittleston and D. J. Wales, *J. Am. Chem. Soc.*, 2007, **129**, 16005–16014.
- D. J. Wales, *PATHSAMPLE: A program for generating connected stationary point databases and extracting global kinetics*, <http://www-wales.ch.cam.ac.uk/software.html>.



- 39 D. Rolfsen, *Knots and Links*, Publish or Perish Press, Berkeley, 1976.
- 40 V. F. R. Jones, *Bull. Amer. Math. Soc.*, 1985, **12**, 103–111.
- 41 C. C. Adams, *The Knot Book*, American Mathematical Society, Providence, 2004.
- 42 J. P. K. Doye, M. A. Miller and D. J. Wales, *J. Chem. Phys.*, 1999, **111**, 8417–8428.
- 43 W. N. Lipscomb, *Science*, 1966, **153**, 373–378.
- 44 O. Shameema, B. Pathak and E. D. Jemmis, *Inorg. Chem.*, 2008, **47**, 4375–4382.
- 45 H. Sayin and M. L. McKee, *Inorg. Chem.*, 2007, **46**, 2883–2891.
- 46 A. M. Mebel, K. Morokuma and D. G. Musaev, *J. Am. Chem. Soc.*, 1994, **116**, 3932–3942.
- 47 R. B. King and J. Hagel, *Polyhedron*, 2006, **25**, 3183–3188.
- 48 J. Uppenbrink and D. J. Wales, *J. Chem. Soc., Faraday Trans.*, 1991, **87**, 215–222.
- 49 D. J. Wales, D. M. P. Mingos and Z. Y. Lin, *Inorg. Chem.*, 1989, **28**, 2754–2764.
- 50 B. F. G. Johnson, *J. Chem. Soc., Chem. Commun.*, 1986, 27–30.
- 51 J. Pillardy and L. Piela, *J. Phys. Chem.*, 1995, **99**, 11805–11812.
- 52 J. P. K. Doye, M. A. Miller and D. J. Wales, *J. Chem. Phys.*, 1999, **110**, 6896–6906.
- 53 D. J. Wales, M. A. Miller and T. R. Walsh, *Nature*, 1998, **394**, 758–760.
- 54 D. J. Wales and D. M. P. Mingos, *Inorg. Chem.*, 1989, **28**, 2748–2754.
- 55 S. M. Ilett, A. Orrock, W. C. K. Poon and P. N. Pusey, *Phys. Rev. E: Stat. Phys., Plasmas, Fluids, Relat. Interdiscip. Top.*, 1995, **51**, 1344–1352.

