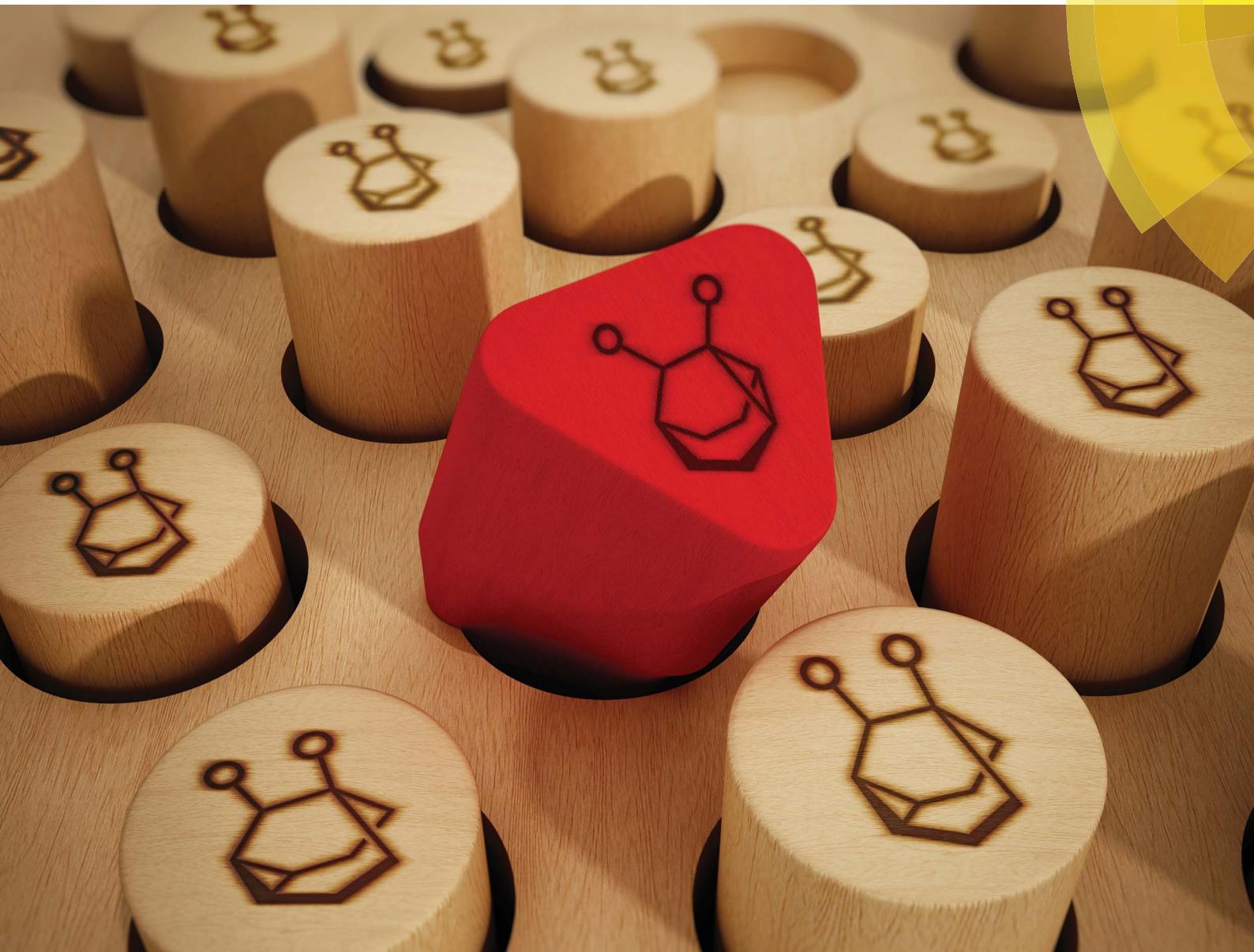


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EDGE ARTICLE

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Shape-selective crystallisation of fluxional carbon cages

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## Introduction

Fluxional carbon cages, such as bullvalene<sup>1</sup> and the barbaralyl cation<sup>2</sup> (Fig. 1a), undergo reversible pericyclic rearrangements on a grand scale.<sup>3</sup> Sequential, low-energy steps interconvert large numbers of degenerate isomers. For example, bullvalene passes back and forth between 1.2 million degenerate valence isomers<sup>1</sup> by strain-assisted Cope rearrangements. Functionalised derivatives, on the other hand, give rise (Fig. 1b) to nondegenerate valence isomers with distinct constitutions.<sup>4</sup> When their C–C bonds trade places with one another, the positions and relative orientations of their substituents are altered, endowing them with dynamic shapes (Fig. 1c) and making them ideal building blocks for adaptive chemistry.<sup>5</sup> Bode *et al.*<sup>6</sup> have taken advantage of these phenomena to design shapeshifting bullvalene sensors—the equilibrium distribution of tetrasubstituted bullvalene isomers is shifted in a characteristic manner when interacting with different guests, either by specific noncovalent interactions<sup>6b</sup> or dynamic covalent bonds.<sup>6c</sup>

Here, we report on the influence of shape over the crystallisation of fluxional carbon cages. Neutral barbaralane derivatives, which equilibrate (Fig. 1b) between sets of two non-equivalent valence isomers<sup>7</sup> in solution, are resolved to single valence isomers upon crystallisation. By analysing NMR spectra of solutions and powdered samples, X-ray crystal structures and

DFT models, we observe that (i) the dynamic isomerism is ‘frozen’ in the solid state and (ii) relatively small differences in the substituents dictate which constitutional isomer is found upon crystallisation. Three of the compounds crystallise as the major solution-phase valence isomers, *i.e.*, their molecular structures in the solid state match those of the thermodynamically favoured species in solution. However, two compounds crystallise (Fig. 1c) as the minor solution-phase valence isomers, achieving more densely packed crystals by adopting molecular structures that fit together more effectively. Size- and shape-matching of these fluxional molecules to their surroundings outweigh the inherent energetic preference of the shapeshifting equilibrium.

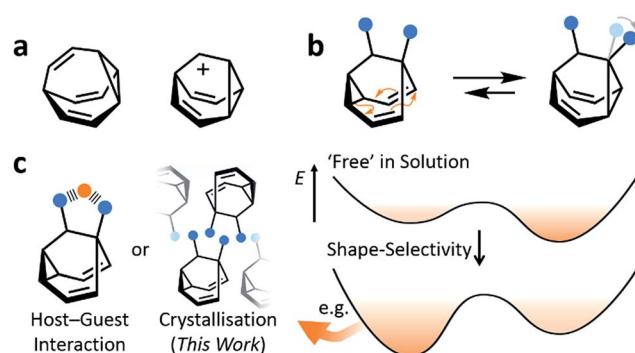


Fig. 1 Structural formulas of (a) bullvalene, the barbaralyl cation and (b) a disubstituted barbaralane, showing its isomerisation. The relative orientations of the substituents, which are represented by blue circles, are altered by the isomerisation. (c) A schematic representation illustrates changes in relative energies of isomers in panel b as a result of a shape-selective intermolecular contact, *e.g.*, through noncovalent bonding with a guest, represented as an orange circle, or by crystal packing.

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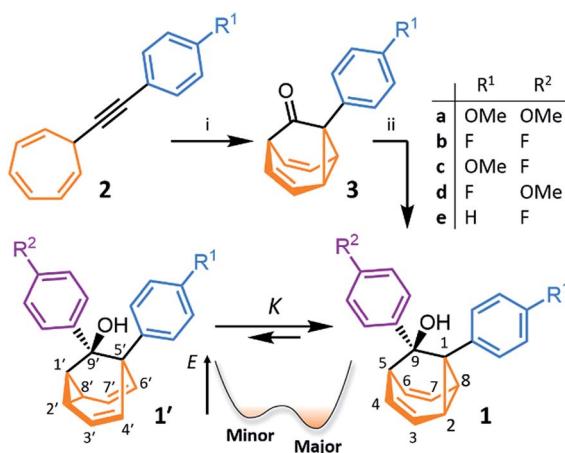
## Results and discussion

We synthesised (Scheme 1) fluxional mixtures of nondegenerate barbaralanes **1**/**1'** in two steps from alkynyl cycloheptatriene precursors **2** (for experimental details see the ESI†). Gold-catalysed cycloisomerisation of **2** proceeds<sup>2c</sup> through barbaralyl cation intermediates, which are oxidised<sup>2d</sup> *in situ* by Ph<sub>2</sub>SO to afford barbaralones **3** in 58–75% yields. Subsequent addition of arylmagnesium bromides to **3** gives barbaralanes **1**/**1'**, each bearing a tertiary alcohol<sup>8</sup> and two aromatic rings. The aromatic ring at the 1/5'-position of each barbaralane breaks the symmetry of the Cope rearrangements, giving rise to the pairs of valence isomers **1** and **1'**. We reasoned that these pairs of nondegenerate valence isomers would be useful compounds for probing the shapeshifting properties of fluxional carbon cages. Compared to the huge numbers of valence isomers accessible to other systems, analysis of the double-well-type potential energy surface (Scheme 1) of **1**/**1'** is more tractable. At the same time, the rearrangement still modulates the molecular shape to some extent—the strain present in the tricyclic cores of **1** and **1'** is slightly different, regulating the relative orientations of the two aromatic rings. Five sets of barbaralanes (**1a–e**/**1a'–e'**) were prepared in which the groups at the *p*-positions of the aromatic rings ( $R^1$  and  $R^2$ ) are varied between OMe, F, and H.

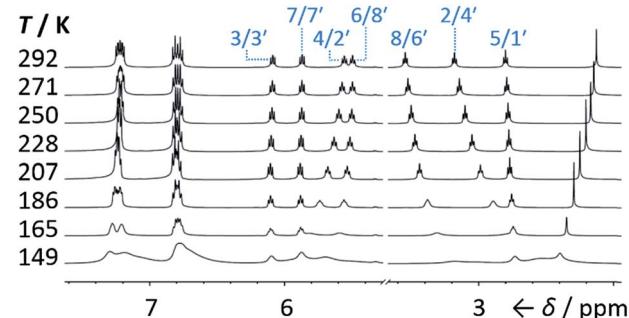
Isomers **1** and **1'** are in fast exchange (Fig. S15–S48†) when observed by solution-phase NMR spectroscopy. Consequently, each pair of exchangeable sites gives rise to one peak. The chemical shift of each peak represents an average of the two discrete chemical environments, weighted by the position of the equilibrium, which enables us to identify the major isomer. For example, the <sup>13</sup>C peak associated with position 2 of **1a** and position 4' of **1a'** appears (Fig. S16†) at 49.6 ppm in CDCl<sub>3</sub> at 298 K, which is closer to the resonance of a divinyl cyclopropane group<sup>8</sup> (~25 ppm) than that of a *cis*-dialkylolfin group<sup>9</sup> (~135 ppm). This observation allows us to assign **1a** as the

major isomer<sup>10</sup> and to estimate an equilibrium constant,  $K$ , of ~3.5, as well as a Gibbs free energy difference,  $\Delta G$ , of ~3 kJ mol<sup>-1</sup>. These conclusions are supported by DFT modelling, which predicts (Table S5†) that **1a** is lower in energy than **1a'** by a margin of ~5 kJ mol<sup>-1</sup> (for calculation details see the ESI†). Further NMR and DFT analyses reveal (Fig. S17–S48 and Table S5†) that  $K$  is only minimally affected by varying the  $R^1$  and  $R^2$  groups at the *p*-positions of the aromatic rings. The substituents are sufficiently remote from the barbaralane core that their electronic and steric differences have little impact—structure **1** remains the thermodynamically favoured solution-phase valence isomer for all derivatives (**1a–e**). In order to probe the equilibria further, we also recorded solution-phase <sup>1</sup>H NMR spectra of bis(4-anisyl)barbaralane **1a**/**1a'** (Fig. S29†) and bis(4-fluorophenyl)barbaralane **1b**/**1b'** (Fig. 2) at low temperatures. The temperature-dependent spectroscopic changes are consistent with the behaviour of two nondegenerate valence isomers interconverting rapidly through a low-lying transition state. For example, the signal for the 8/6' positions of **1b**/**1b'** shifts (Fig. 2) from 3.55 ppm at 292 K to 3.31 ppm at 165 K, moving further towards the chemical shift expected for a cyclopropyl resonance, while the signal associated with 6/8' moves towards the olefinic region, shifting from 5.50 ppm at 292 K to 5.60 ppm at 149 K. Therefore, the equilibrium shifts further in favour of the thermodynamically favoured isomer **1b** at lower temperatures, which would be expected for a Boltzmann distribution of nondegenerate isomers.

All five **1**/**1'** derivatives form crystalline solids, allowing us to analyse (Fig. 3) their solid-state structures by single-crystal X-ray diffraction.† The fluxional mixtures of **1**/**1'** undergo dynamic preferential crystallisation,<sup>11</sup> *i.e.*, just one valence isomer from each pair of fluxional carbon cages crystallises from the mixture.<sup>12,13</sup> The bis(4-anisyl)barbaralane crystallises (Fig. 3a) as the major solution-phase valence isomer **1a**. Similarly, the barbaralanes bearing one 4-fluorophenyl group and one 4-anisyl group each adopt the major solution-phase structures **1c** and **1d** upon crystallisation. Surprisingly, however, we observed the more strained molecular structures in crystals of the bis(4-fluorophenyl)barbaralane **1b'** and its phenyl analogue **1e**.



**Scheme 1** Synthesis of fluxional carbon cages **1**/**1'**. Reagents and conditions: (i) IPrAu(MeCN)BF<sub>4</sub> (5 mol%), Ph<sub>2</sub>SO (2 equiv.), CH<sub>2</sub>Cl<sub>2</sub>, rt, 16 h, 58–75%; (ii) *p*-R<sup>2</sup>C<sub>6</sub>H<sub>4</sub>MgBr, THF, 0 °C to rt, 16 h, 43–96%. A schematic potential energy diagram illustrates the equilibrium,  $K$ , between **1'** and **1**. IPr = 1,3-bis(2,6-diisopropylphenyl)imidazol-2-ylidene.



**Fig. 2** Partial <sup>1</sup>H NMR spectra (500 MHz, CS<sub>2</sub>–CD<sub>2</sub>Cl<sub>2</sub>) of **1b**/**1b'**. As the temperature is decreased, the equilibrium shifts towards the lower-energy isomer **1b**. At temperatures below 186 K, the broadening of peaks indicates the reversible Cope rearrangement is entering the slow-exchange regime. Selected resonances are labelled according to the numbering in Scheme 1.



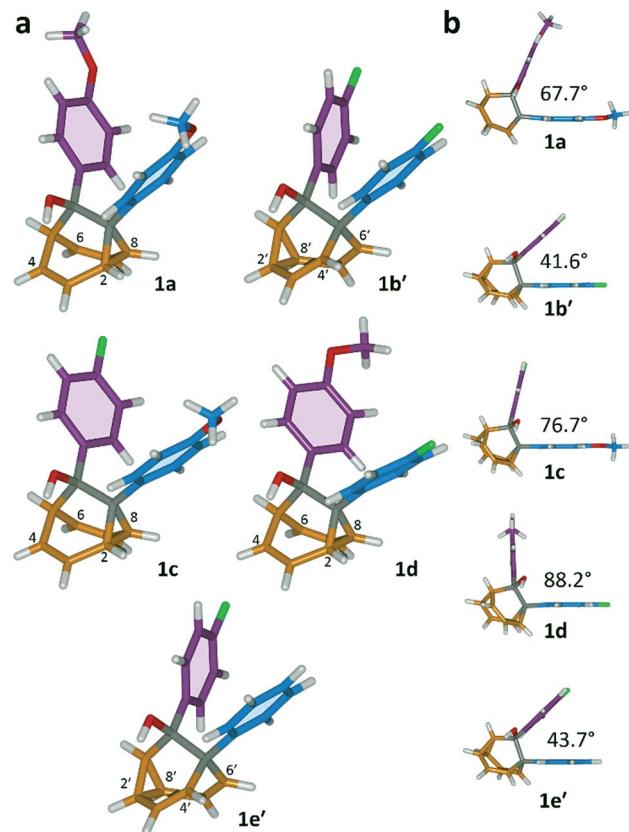


Fig. 3 X-ray crystal structures of **1/1'** shown in stick representations.<sup>†</sup> (a) Whereas the major solution-phase valence isomers are obtained in some cases (**1a**, **1c** and **1d**), the minor solution-phase isomers crystallise preferentially in others (**1b'** and **1e'**). (b) Side-on views show the variation in dihedral angles between the planes of the aromatic rings. Selected interatomic distances at 120 K (Å): **1a**, C2–C8 1.592(2), C4…C6 2.359(2); **1b'**, C2'–C8' 1.592(3), C4'…C6' 2.353(2); **1c**, C2–C8 1.582(2), C4…C6 2.364(2); **1d**, C2–C8 1.586(2), C4…C6 2.367(2); **1e'**, C2'–C8' 1.591(2), C4'…C6' 2.345(2). Oxygen atoms are shown in red, fluorine atoms are green, hydrogen atoms are white, and carbon atoms are blue, purple, orange or grey.

While the *p*-substituents have little influence over the solution-state valence isomerism of the five **1/1'** derivatives, it appears that a subtle change at a position distant from the barbaralane core, such as switching out the OMe group of **1c/1c'** for a F atom (*i.e.*, giving **1b/1b'**), is sufficient to bias crystallisation in favour of another valence isomer.<sup>¶</sup>

Although some fluxional carbon cages<sup>14</sup> are known to become ‘fixed’ structures upon crystallisation, others<sup>15</sup> remain fluxional. Variable-temperature X-ray diffraction (VT-XRD) measurements performed on single crystals of **1a** and **1b'** show (Fig. S42 and S43<sup>†</sup>) negligible changes in C–C distances of bonds involved in Cope rearrangement. For example, the C2–C8 bond length of **1a** only changes from 1.592 Å at 120 K to 1.604 Å at 370 K. The identities of **1** and **1'** isomers in the solid state can also be distinguished by comparing (Fig. 4) their low-temperature solution and ambient-temperature solid-state <sup>13</sup>C NMR spectra. At 149 K, the solution-phase equilibria are biased heavily towards **1a** and **1b'**, so both solution-phase spectra show (Fig. 4a,d) similar patterns of peaks in their alkyl regions. Unlike

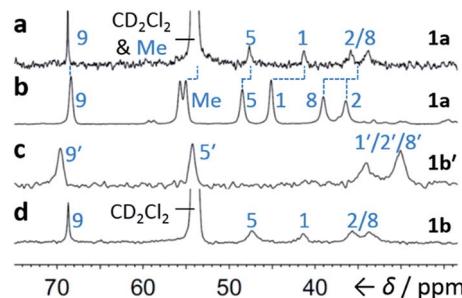


Fig. 4 Comparison of partial <sup>13</sup>C NMR spectra of **1a/1a'** and **1b'/1b'** acquired (a and d) as CS<sub>2</sub>–CD<sub>2</sub>Cl<sub>2</sub> solutions at low temperature (126 MHz, 149 K), and (b and c) as powders at ambient temperature (105 MHz). Peaks are labelled according to the numbering in Scheme 1.

the solid-state NMR (ssNMR) spectrum of the bis(4-anisyl) barbaralane (Fig. 4b), which matches closely with its solution spectra (in keeping with its assignment by X-ray diffraction as structure **1a**), the ssNMR spectrum (Fig. 4c) of the bis(4-fluorophenyl)barbaralane is markedly different. The ssNMR spectrum indicates structure **1b'** is favoured over **1b**. Hence, the ssNMR and VT-XRD data are both consistent with the barbaralanes **1/1'** having fixed constitutions in the solid state.

The barbaralane cage acts as a hinge between its two aromatic substituents, giving (Fig. 3b) the molecules V-shaped structures. Notably, the dihedral angles between the planes of these aromatic rings in the solid state vary substantially between the different derivatives. Compounds **1a**, **1c**, and **1d** have relatively wide dihedral angles  $> 67^\circ$ , whereas the dihedral angles of **1b'** and **1e'** are both  $< 44^\circ$ .<sup>||</sup> Inspection (Fig. 5 and S57–S61<sup>†</sup>) of the solid-state superstructures reveals that these changes in dihedral angles are linked to differences in the crystal packing. Indeed, the packing is dominated by the way in which the V-shaped units fit together, rather than any specific noncovalent interactions. Compounds **1a**, **1c**, and **1d** pack with a preference to arrange their V-shaped structures head-to-tail, while **1b'** and **1e'** are arranged head-to-head, allowing their aromatic groups to interdigitate. We

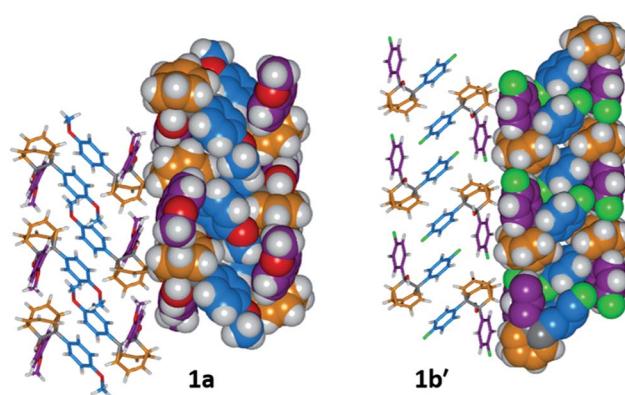


Fig. 5 Solid-state superstructures of **1a** and **1b'** shown in partial stick and space-filling representations and viewed along the crystallographic *b*-axes. The superstructures of **1c** and **1d** (Fig. S59 and S60<sup>†</sup>) are similar to **1a**, whereas the superstructure of **1e'** (Fig. S61<sup>†</sup>) is isostructural with **1b'**. Calculated densities (g cm<sup>−3</sup>): **1a**, 1.33; **1b'**, 1.44; **1c**, 1.35; **1d**, 1.35; **1e'**, 1.39.

attribute changes between these two modes of packing primarily to differences in the sizes of the  $R^1$  and  $R^2$  groups. A 4-anisyl group is too large to allow efficient interdigitation, so the head-to-tail arrangement is favoured in which the aromatic rings are splayed around a neighbouring barbarylone. This geometry is most easily accommodated by the 1-isomer form. Conversely, **1b'** and **1e'** lack 4-anisyl groups, so they are able to achieve more densely packed lattices than the other derivatives (1.39–1.44 vs. 1.33–1.35 g cm<sup>−3</sup>) by adopting head-to-head arrangements. The 4-fluorophenyl and phenyl groups are small enough for the aromatic rings to pack together efficiently, favouring smaller dihedral angles that are more accessible when the barbarylone adapts to the slightly higher energy 1'-isomer. Although there is more strain in the molecular structure, the relatively small ( $\sim 5$  kJ mol<sup>−1</sup>) energy penalty to adopt the structure of the minor solution-state isomer must be overridden by a more favourable lattice energy, making the 1'-isomer favoured in the solid state.<sup>16</sup>

The alcohol function (which might normally be expected to dictate<sup>17</sup> the packing) is overcrowded and electrostatic potential maps show (Fig. S66†) that the aromatic rings are only mildly polarised. In order to search for noncovalent bonding interactions, we calculated the Hirshfeld surfaces<sup>18</sup> of each crystal structure. The Hirshfeld surfaces show (Fig. S67–S72†) that no interatomic distances are significantly closer than the sum of the respective van der Waals radii. For example, the most prominent close contacts present (Fig. 6) in **1d** and **1e'** are weak O–H···C and C–H···C interactions, which each remain within  $\sim 0.2$  Å of the sum of the van der Waals radii.<sup>19</sup> We confirmed this lack of significant, specific noncovalent bonding interactions by performing DFT modelling (CE-B3LYP\*\*) of all the intermolecular interactions in the solid state. These calculations allow us to elucidate the total interaction energies ( $E_{\text{tot}}$ ) between neighbouring molecules based on the individual energetic components of electrostatics ( $E_{\text{ele}}$ ), polarisation ( $E_{\text{pol}}$ ), dispersion forces ( $E_{\text{dis}}$ ) and repulsion ( $E_{\text{rep}}$ ).<sup>20</sup> For example, we examined (Fig. 7) a cluster of 14 neighbouring molecules surrounding a central molecule in the structure of **1e'**. The 14 neighbours account for all of the molecules that come within

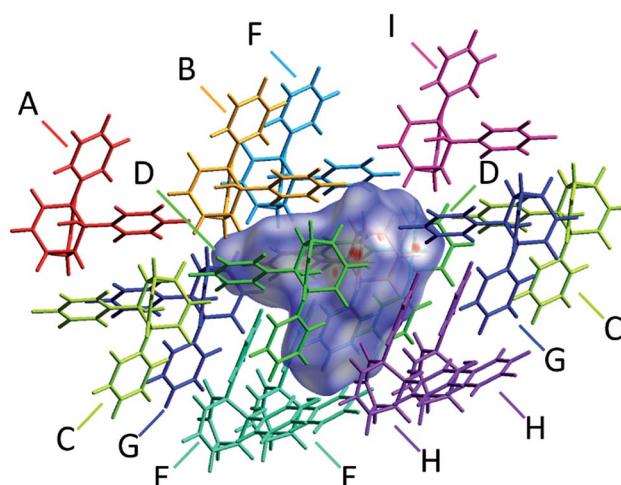


Fig. 7 A section of the solid-state superstructure of **1e'** in which a central molecule (overlaid with its modelled Hirshfeld surface) is surrounded by its 14 nearest neighbours (labelled A–I). Molecules A–I are shown in stick representation. Labels are duplicated for molecules that reside at symmetry-related coordinates relative to the central molecule.

3.8 Å of the central molecule. As would be expected, the pairs of **1e'** molecules that are closest, having the smallest centroid-to-centroid distances ( $R$ ), generally interact most strongly with one another. Molecules in position D and F are separated (Table 1) from the central molecule by  $R < 7$  Å and have interaction energies of  $E_{\text{tot}} = -31.1$  and  $-43.5$  kJ mol<sup>−1</sup>, respectively, which are the strongest pairings present in the structure. Dispersion forces account for the majority of their attractive intermolecular interactions, contributing energies of  $E_{\text{dis}} = -46.8$  kJ mol<sup>−1</sup> for D and  $E_{\text{dis}} = -51.4$  kJ mol<sup>−1</sup> for F. Overall, the modelling for the full series of five 1/1' derivatives shows (Tables S10–S14†) that dispersion forces<sup>21</sup> (which are influenced by the shape of the molecules and their packing) make up the major component of the interactions between molecules in all

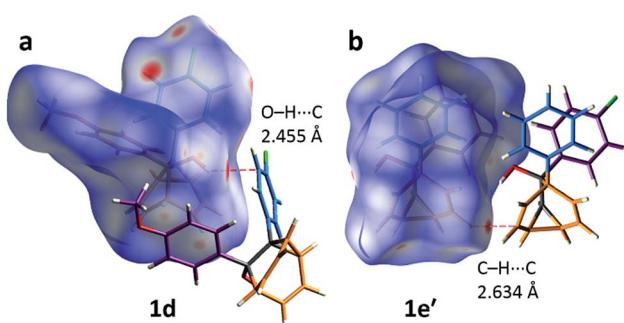


Fig. 6 Calculated Hirshfeld surfaces for the X-ray crystal structures of (a) **1d** and (b) **1e'**, showing the most significant close contacts with neighbouring molecules, which do not correspond to appreciable noncovalent bonding interactions.

Table 1 Calculated intermolecular interaction energies (kJ mol<sup>−1</sup>) for pair of molecules in the solid-state structure of **1e'**

Neighbouring molecule <sup>a</sup>	$R/\text{\AA}^b$	Interaction energies/kJ mol <sup>−1</sup>				
		$E_{\text{ele}}$	$E_{\text{pol}}$	$E_{\text{dis}}$	$E_{\text{rep}}$	$E_{\text{tot}}^c$
A	13.29	2.1	−0.2	−1.3	0.0	1.0
B	7.94	−10.1	−1.4	−31.2	20.3	−26.3
C	11.59	−2.3	−0.4	−3.8	0.6	−5.6
D	6.20	−8.4	−2.4	−46.8	32.8	−31.1
E	8.26	−5.0	−0.9	−30.3	18.7	−20.8
F	6.63	−14.8	−2.8	−51.4	30.9	−43.5
G	9.79	−5.0	−0.8	−16.1	11.7	−12.7
H	7.90	−6.3	−1.1	−31.5	17.9	−23.8
I	11.04	1.1	−0.4	−8.1	1.8	−5.1

<sup>a</sup> The labels A–I correspond to those shown in Fig. 7 and refer to the nearby molecules in the solid state. <sup>b</sup>  $R$  is the centroid-to-centroid distance between the labelled molecule and the central molecule. Centroids are based on the coordinates of all atoms in each molecule and are not weighted by mass. <sup>c</sup>  $E_{\text{tot}}$  is the scaled sum of the individual interaction energy components  $E_{\text{ele}}$ ,  $E_{\text{pol}}$ ,  $E_{\text{dis}}$  and  $E_{\text{rep}}$ .



of the packing motifs, dominating the overall lattice energies in each case. This observation is consistent with our description of these molecules packing in the solid state in a manner that is influenced more by molecular shape than by any specific non-covalent bonding interactions.

## Conclusions

In summary, by investigating the dynamic isomerisation of nondegenerate barbaralanes, we have found that their crystallisation is directed by the shapes of their fluxional isomers. Either the major or the minor solution-phase isomer may be resolved as a result of their assembly in crystal lattices, becoming trapped in the solid state. At present, it has not been possible to discern precisely the shapes available to the most advanced shapeshifting sensors,<sup>6</sup> which encompass hundreds of constitutional isomers and conformationally flexible substituents. Our observations, based on a bistable fluxional carbon cage with rigid substituents, illustrate that shape-selective interactions can override the inherent thermodynamic differences between even closely related valence isomers. Understanding the relationship between shape-selectivity and the energetics of fluxional carbon cages will aid the development of shapeshifting sensors and other applications of adaptive molecules.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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## Notes and references

§ Barbaralanes **1/1'** were prepared and isolated as racemic mixtures. For simplicity, only one enantiomer of each compound is shown.

¶ During our experiments, we did not observe any crystals of solvates or conformational polymorphs. The limited conformational freedom of **1/1'** and limited potential for hydrogen bonding suggest they are unlikely to be highly polymorphic.

|| The differences in dihedral angles coincide with changes to the C1–C9/C5'–C9' distances in the barbaralane core. The averaged C1–C9 distance for **1a**, **1c**, and **1d** is 1.534 Å, whereas the averaged C5'–C9' distance for **1b'** and **1e'** is 1.586 Å, showing that the movement of the divinylcyclopropane motif influences the geometries near the aromatic rings.

\*\* The CE-B3LYP functional was used instead of other DFT functionals, *e.g.*, CAM-B3LYP or MN15-L, because its implementation in CrystalExplorer allows calculation of the individual interaction energy terms in addition to the total energy. It should be noted that CE-B3LYP is different to the original B3LYP functional,

which is not suitable for calculation of weak interactions. It reproduces counterpoise-corrected B3LYP-D2/6-31G(d,p) energies with a mean absolute deviation of about 1 kJ mol<sup>-1</sup> and gives a mean absolute deviation of just 2.5 kJ mol<sup>-1</sup> compared to CCSD(T)/CBS while using much less computational time. See ref. 20.

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