

Cite this: *Chem. Sci.*, 2019, 10, 752

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Influence of metal coordination and light irradiation on hierarchical self-assembly processes†

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Smart light-responsive supramolecular materials have been extensively investigated in the past decade, but so far the impact of metal coordination on hierarchical supramolecular structures of light-responsive building blocks has remained nearly unexplored. Herein, we unravel the hierarchical self-assembly of a small π -conjugated azo-containing pyridyl ligand that is able to respond to UV-light and metal complexation. The ligand self-assembles in an antiparallel fashion into long twisted fibers, which are then disassembled upon photoisomerization of the azobenzene groups, resulting in shorter rigid rods with a different packing motif. Complexation of Pd(II) ions enhances the cooperativity of the aggregation and induces a molecular rearrangement into slipped stacks with subsequent formation of long thin fibers. These are then transformed into thinner, shorter rods upon light irradiation. The observed different light-responsiveness, besides clearing up the influence of metal coordination and light irradiation in self-assembly processes, paves the way towards the design of novel supramolecular photochromic systems.

Received 30th August 2018
Accepted 24th October 2018

DOI: 10.1039/c8sc03875a

rsc.li/chemical-science

Introduction

Self-assembly of functional small molecules has become a promising approach to create smart materials¹ that can respond to changes in various external variables such as temperature, concentration, pH and solvent or stimuli such as light,² sound,³ mechanical forces⁴ or cations/anions.⁵ Light-responsive supramolecular assemblies are of particular interest in this regard because of the sensitivity of the molecular shape, size and polarity upon photoisomerization, which enables the modulation of the structural and functional properties associated with the self-assembly pathway.² The incorporation of photochromic moieties such as azobenzenes,⁶ dithienylethenes⁷ and spiropyrans⁸ offers great

potential to create light-responsive supramolecular assemblies. In particular, azobenzenes have been extensively exploited as photoswitchable units in light-responsive host-guest systems,⁹ liquid crystals,¹⁰ vesicles,¹¹ gels,¹² biomaterials¹³ and self-assembled structures of π -conjugated molecules.^{14–16} For the construction of one-dimensional (1D) photoresponsive supramolecular assemblies, the major molecular design is the use of photochromic molecules as single-molecular building blocks.^{14,16} A particularly attractive approach is the oligomerization of photochromic moieties *via* non-covalent interactions, *i.e.* hydrogen bonds, to form supramolecular building blocks, as this provides access to hierarchically organized systems with unique photoresponsive behaviour not achievable by single-molecule based systems.^{15b,c} In this regard, we envisaged that the introduction of a metal to link photochromic units would have a strong impact on the photoresponsive behaviour and hierarchy levels of the assemblies due to the modification in the molecular conformation and packing modes. For example, we have reported that coordination of a Pt(II)/Pd(II) dihalogen metal center to pyridyl-based π -conjugated moieties can induce new non-covalent interaction sites different from the ligand thereby enabling new self-assembly pathways.¹⁷ To date, photoresponsive metallosupramolecular systems have been limited to polymeric materials,¹⁸ metallacycles^{19a} and metal-organic frameworks.^{19b,d} Despite these advances, understanding the influence of metal coordination²⁰ and light-irradiation on hierarchical self-assembly processes remains elusive.^{19c}

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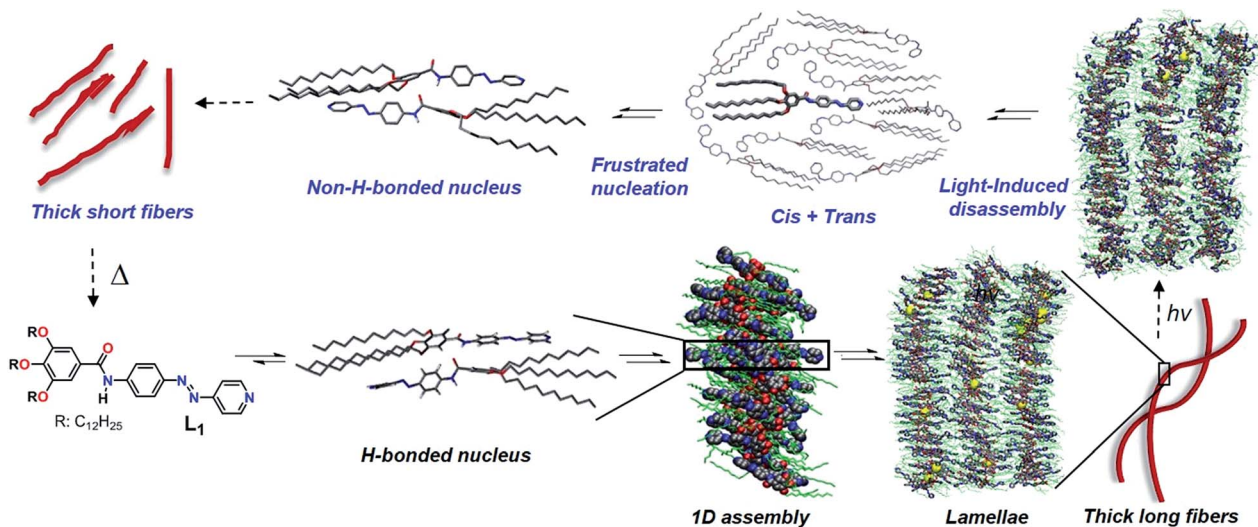
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† Electronic supplementary information (ESI) available: Synthesis and characterization, description of experimental techniques, and additional images. CCDC 1864670. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c8sc03875a





Scheme 1 Proposed hierarchical self-assembly and photoresponsive behavior of L_1 . Structures of monomers and small aggregates were optimized using DFT and dispersion-corrected PM6 calculations, respectively, while structures with large aggregates (270 monomers) were obtained from classical molecular dynamics.

Herein, we unravel the impact of metal complexation and photoisomerization on hierarchical self-assembly processes *via* detailed mechanistic studies by means of combining experimental and theoretical techniques. Our molecular design features a light-responsive azobenzene unit, an amide group for hydrogen bonding, a metal ion-responsive pyridyl ligand and dodecyloxy side chains (L_1 , Scheme 1). The target ligand L_1 and its corresponding Pd(II) complex C_1 (Scheme S1[†]) have been synthesized by modified literature procedures,²¹ as described in the (ESI[†]). Additionally, a related ligand L_2 bearing ethoxy groups as well as its corresponding Pd(II) complex C_2 have been prepared to facilitate crystal growth (Scheme S1[†]).

Results and discussion

Hierarchical self-assembly of L_1

We examined the self-assembly of ligand L_1 in a poor solvent (methylcyclohexane; MCH) by cooling a hot solution ($c = 5 \times 10^{-4}$ M) from 363 K to 283 K with 1 K min^{-1} ramp. The freshly prepared solution showed negligible absorption spectral changes in various good solvents (*i.e.* dichloromethane; DCM), suggesting the molecularly dissolved state (Fig. S1[†]) at 298 K. However, a gradual decrease of the absorption maximum at 363 nm and a concomitant increase of a shoulder at *ca.* 420 nm were observed upon keeping the MCH solution at 283 K over time (Fig. 1a). This, along with the appearance of an isosbestic point at 407 nm, strongly suggests the formation of self-assembled species. A plot of absorbance at 360 nm (A_{360}) *vs.* time reveals that around 80% of the aggregation process is complete within 5 h (Fig. 1a, inset). Likewise, a lower concentrated solution (2.5×10^{-4} M) was also monitored over time at 283 K, but in this case negligible changes were observed, indicating that the critical aggregation concentration is close to 5×10^{-4} M at 283 K. Variable Temperature (VT)-NMR studies of L_1 in MCH- d_{14} ($c = 5 \times 10^{-4}$ M) from 363 to 283 K showed

downfield shifts for the amide protons (H_e) and aromatic protons H_d (*ca.* 1 and 0.15 ppm, respectively, see Fig. 1b), indicating the proximity of an electron-rich environment (*i.e.* N or O atoms from a neighboring molecule) and H-bonding.²² Comparison of the FT-IR spectra of L_1 ($c = 5 \times 10^{-4}$ M) in the monomer (CHCl_3) and aggregated state (MCH) allowed us to assign these hydrogen bonds to N-H \cdots O=C interactions between the amide groups (Fig. S2[†]). On the other hand, the phenyl protons H_f underwent a marked upfield shift upon cooling, strongly suggesting the involvement of the trialkoxyphenyl unit in π -stacking. Interestingly, the pyridine protons ($H_{a,b}$) remain almost unaltered during the cooling process, ruling out the possibility of a face-to-face parallel stacking.^{22a} A noteworthy observation is the sharpening of the proton signals upon decreasing temperature, which can be ascribed to the formation of discrete aggregate species, such as dimers, during the cooling process.²³ This is in agreement with the results obtained from variable-temperature dynamic light scattering (DLS) (Fig. S3; for details see ESI[†]). Attempts to further monitor this process by diffusion-ordered spectroscopy (DOSY) NMR proved unsuccessful due to unreliable data (Fig. S4[†]).

Monitoring this discrete aggregate formation by VT-UV-Vis experiments under identical conditions showed only a slight hyperchromic effect without a clear isosbestic point upon cooling from 363 K to 283 K (Fig. 1a and S5[†]). Though hyper/hypochromic effects can be ascribed to a weak intermolecular interaction of the π -systems on the basis of Kasha's exciton theory,²⁴ the absence of a clear isosbestic point suggests the existence of more than one type of discrete aggregate species, most likely dimers, in equilibrium with the monomer. Time-dependent ^1H NMR studies were then performed immediately after the solution used for VT-NMR studies reached 283 K (Fig. 1b). The observed strong broadening and subsequent disappearance of the signals indicate a further oligomerization of the dimers over time, which is in perfect agreement with the



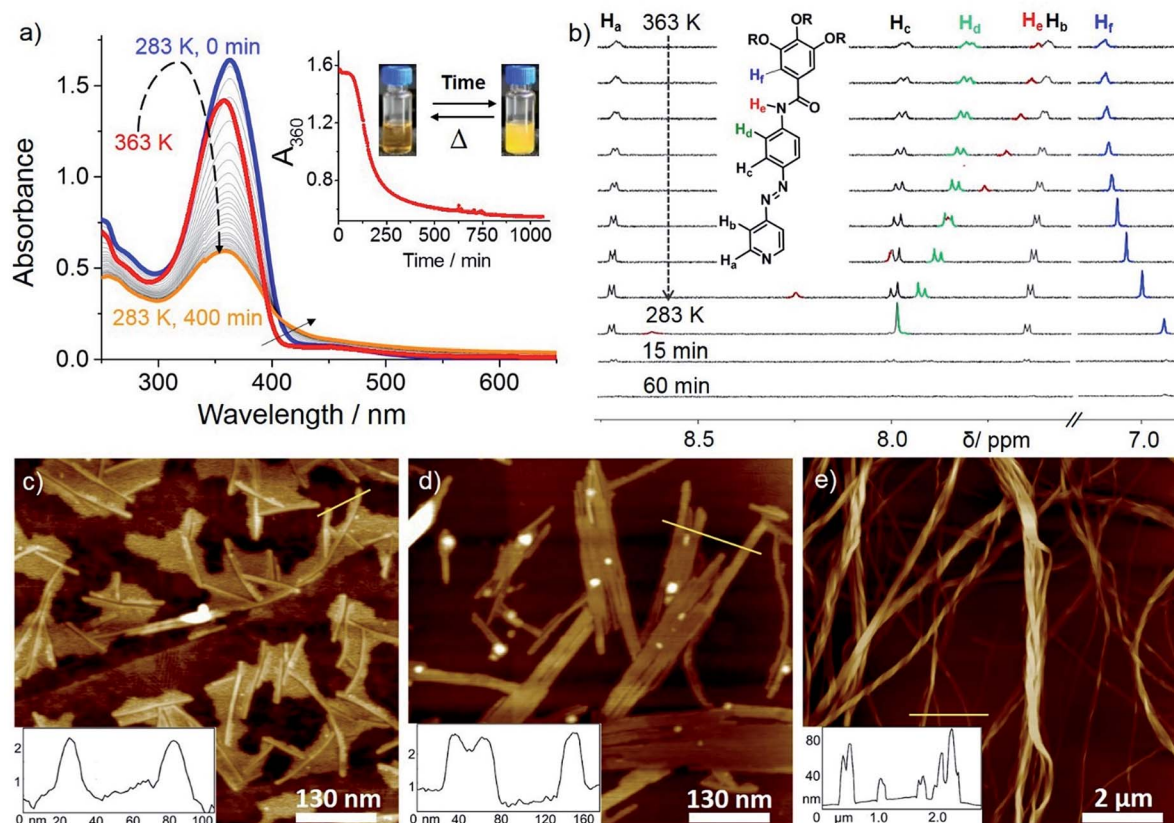


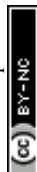
Fig. 1 (a) Temperature-dependent absorption changes of L_1 (5×10^{-4} M, MCH) from 363 K (red line) to 283 K (blue line) followed by time-dependent studies at 283 K for a period of 400 min. Inset: plot of A_{360} vs. time and photographs showing the reversible sol-aggregate formation. (b) Partial VT- 1H NMR spectra of L_1 (5×10^{-4} M, MCH- d_{14}) between 363 and 283 K followed by time-dependent studies at 283 K over a period of 60 min (initial lag is excluded). (c–e) AFM images of self-assembled L_1 in MCH upon keeping the solution at 283 K for (c) 15 min, (d) 30 min and (e) 60 min on HOPG (initial lag is excluded) with cross-section analysis along the yellow lines.

depletion of the monomer band and a concomitant emergence of an aggregate band in time dependent UV-Vis experiments (see Fig. 1a). Time-dependent DLS studies in MCH ($c = 5 \times 10^{-4}$ M) at 283 K (Fig. S3 \dagger) revealed a consistent increase in decay time during the initial 15 min, which supports the formation of oligomers. Over time, a more pronounced aggregation process is evident from the decrease in relative counts over 60 min. Comparison of the 2D COSY and ROESY spectra of an aggregate solution of L_1 (11.7 mM, MCH- d_{14} , 315 K) strongly suggests an H-bonded antiparallel molecular arrangement (see ESI for details, Fig. S6 \dagger) which is in good agreement with the findings observed by previous VT-UV and NMR studies. A related antiparallel arrangement of aromatic groups *via* H-bonds has been previously observed in the solid state for oligomeric zipper complexes bearing aromatic amides.²⁵

The structure of this H-bonded packing optimized *via* the dispersion-corrected PM6 method is shown in Scheme 1. This conformation is appropriate to grow a stable 1D assembly of L_1 (Scheme 1), as revealed by classical Molecular Dynamics (MD) simulations carried out at 300 K and 1 atm (for more details on the MD simulations, see the ESI \dagger). These simulations also reveal that the interactions between the alkyl chains are very important to stabilize the antiparallel stacks,

together with hydrogen bonding. In the MD snapshot shown in Scheme 1, the structure of the simulated supramolecular material is somewhat heterogeneous although in average the monomers tend to stack in an antiparallel fashion driven to a large extent by H-bonding. Also, interdigitation of the alkyl chains facilitates the growth into 2D lamellae (see Scheme 1) and finally long, twisted fibers by shielding the polar pyridine rings from exposing themselves to the surrounding nonpolar medium.

The hierarchical self-assembly of L_1 has also been studied using atomic force microscopy (AFM) imaging by spin-coating MCH solutions of L_1 ($c = 5 \times 10^{-4}$ M, 283 K) onto highly oriented pyrolytic graphite (HOPG) at different time intervals. After keeping the solution for 15 minutes at 283 K (excluding the initial lag), short 1D supramolecular fibers with 2–3 nm in height, 10–15 nm in width (Fig. 1c and S7a \dagger) and a strong tendency to form lamellae were observed, which further grew into 2D lamellar structures with 2–3 nm in height and 50–100 nm in width over a period of 30 minutes (Fig. 1d and S7b \dagger). Subsequent rolling and bundling of the tapes (Fig. S7c \dagger) ultimately results in the formation of thick, long, twisted fibers after an overall equilibration time of 45–60 minutes (Fig. 1e and S7d \dagger). The width of the fibers ranges from 50 to 100 nm,



whereas the length goes up to several microns. The formation of these thicker fibers led to a yellow gel above 50 mM, as confirmed by AFM (Fig. S8†).

Photoisomerization of L_1

Irradiation at 370 nm of the long fibers of L_1 with the azo groups in the *trans* conformation (*trans*- L_1 , $c = 5 \times 10^{-4}$ M) at 283 K led to the disassembly of the fibers, which is attributed to the *trans*-to-*cis* isomerization, as demonstrated by the depletion of the π - π^* transition (363 nm) along with the increase of the n - π^* transition at 450 nm (Fig. 2a, purple line). Some insight into this UV-induced disassembly process could be obtained from MD simulations by inducing an instantaneous *trans*-*cis* isomerization of L_1 within a fiber: a substantial reduction in the number of amide groups involved in short hydrogen-bonding contacts ($d_{O\cdots H} < 2$ Å) was observed, as depicted as yellow spheres in the MD snapshots, and the monomer-monomer distances increased (Scheme 1, top right). These structural changes might explain the initial driving force for the disassembly of the long fibers after light irradiation (*vide infra*). According to UV-Vis results, approximately 80% of the *cis* isomer is formed upon UV irradiation, which reverts progressively to the more stable *trans* isomer over the period of 120 min upon keeping the solution in the dark (blue line in Fig. 2 and S9a†). Monitoring the A_{360} vs. time at 283 K upon the entire back isomerization process allowed us to distinguish an aggregation process that closely resembles the one shown by non-irradiated L_1 (Fig. 1a

and S9a†). Thus, we conclude that the *cis* isomer is a dormant species, whereas aggregation only occurs upon activation to the active *trans* monomer *via* back isomerization (Fig. 2a, inset).^{16,26} These processes (photoisomerization and self-assembly) compete partially at higher concentration (1 mM), where the *cis* form is likewise dormant to aggregation (Fig. S10†).

Closer insight into the influence of *cis*-to-*trans* isomerization on the hierarchical self-assembly of L_1 was provided by time-dependent $^1\text{H-NMR}$ experiments (5×10^{-4} M, MCH- d_{14}) at 283 K. Due to the impossibility of irradiating the sample inside the NMR spectrometer and the longer equilibration time needed compared to UV-Vis, the first recorded $^1\text{H NMR}$ spectrum (denoted as '0 min' in Fig. 2b) was obtained around 15 min after the sample was irradiated for 30 min at 298 K. By this procedure, we established that 40% of the *cis* isomer was present for the first NMR measurement even though back isomerization took place at 298 K prior to the NMR measurement (Fig. S9b-d†). Over a period of around 40 min, complete disappearance of the signals of the *cis* isomer (marked in green with black circles) was concomitant with a slight upfield shift of the N-H proton (H_e), whereas all remaining protons showed insignificant changes. Interestingly, a more significant broadening of the H_e signal and further upfield shift was also observed even after complete *cis*-*trans* conversion (Fig. 2c) around 60 min. This trend was not observed for the *trans* isomer without irradiation (Fig. 1b), and indicates the lack of hydrogen bonding but rather the proximity of a π -surface. We hypothesize

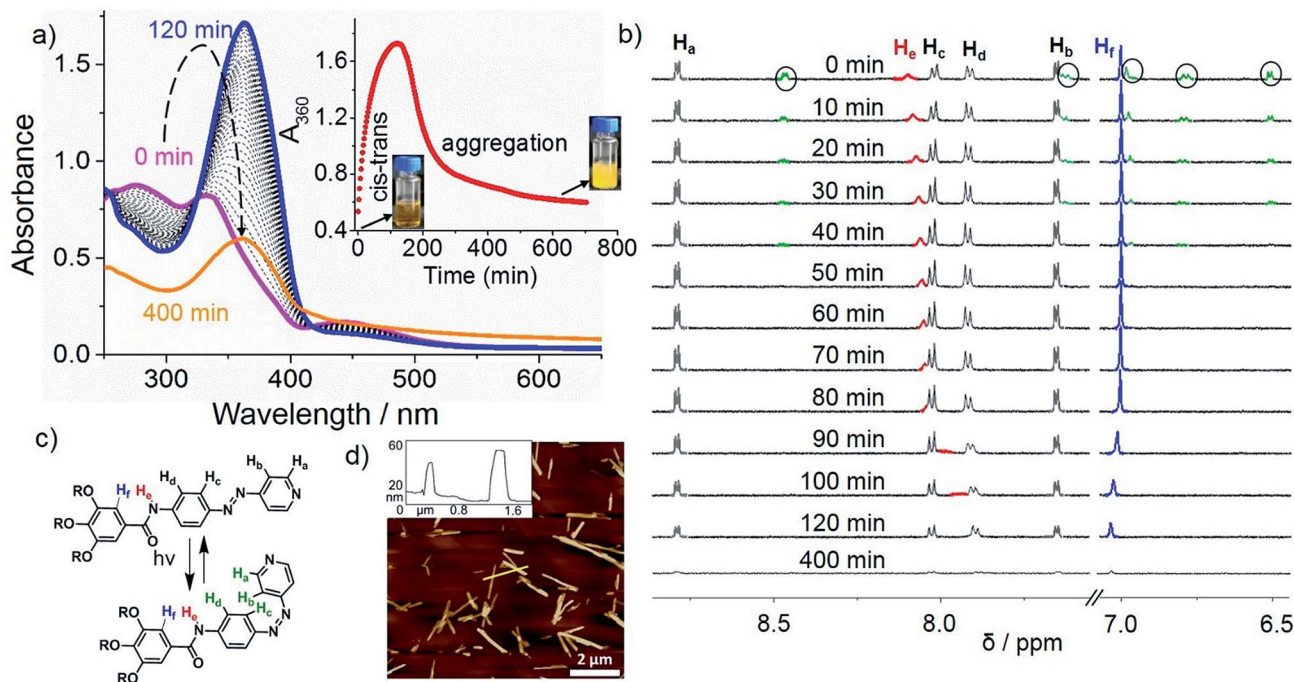


Fig. 2 (a) Time-dependent absorption changes (0–120 min: purple to blue; 120–400 min blue to orange) of L_1 (5×10^{-4} M) in MCH after irradiation for 30 min with a 370 nm LED lamp at 283 K; inset: plot of A_{360} vs. time at 283 K, and photographs showing the irradiated clear L_1 *cis* + *trans* monomer (left) to turbid L_1 *trans* aggregates (right). (b) Time-dependent $^1\text{H-NMR}$ changes at 283 K of L_1 (5×10^{-4} M) after irradiation for 30 min with a 370 nm LED lamp in MCH- d_{14} . Black encircled green peaks correspond to *cis* isomer. (c) Scheme showing reversible *trans*-*cis* isomerization. (d) AFM image of L_1 (5×10^{-4} M) on HOPG upon UV-irradiation for 30 min in MCH and ageing at 283 K for 400 min with corresponding cross-section analysis along the yellow line.



that the high excess of dormant *cis* monomers (80% according to UV-Vis) formed immediately after photoirradiation at 283 K could sequester the *trans*-monomers and prevent them to form antiparallel hydrogen-bonded stacks, leading to a non-H-bonded arrangement different from the non-irradiated pathway (Scheme 1, “*cis* + *trans*” structure).

The above photoisomerization-regulated stepwise aggregation process of L_1 (5×10^{-4} M, MCH) was examined by AFM. The aggregates formed by L_1 in the absence of light at 283 K (Fig. 1e and S11a†) were irradiated with 370 nm UV-light for 30 min, kept at 283 K and finally spin-coated onto HOPG at different time intervals. AFM analysis of the sample kept at 283 K for 140 min upon irradiation (approximately the maximum value observed in the plot A_{360} vs. time, see inset of Fig. 2a) reveals the formation of ill-defined amorphous aggregates (Fig. S11b†) that are most likely the result of a kinetically-driven off-pathway aggregation event. Interestingly, further ageing this solution at 283 K for additional 260 min (total time 400 min after irradiation) showed the transformation of the amorphous short assemblies into photo-reconstructed rod-like structures (Fig. 2d and S11c, d†) that are considerably shorter than those formed without irradiation (Fig. 1e). These results can be explained in terms of a frustrated nucleation event of the active *trans* isomers caused by the presence of sterically hindered, dormant *cis*-monomers, which is supported by a non-H-bonded antiparallel dimerization predicted by PM6 calculations and MD simulations (Scheme 1). The formation of these less organized pre-nuclei dramatically affects the addition of further active *trans* monomers during the subsequent elongation process, leading to a less compact non-H-bonded arrangement that ultimately results in the shortening of the ensembles. These results can be corroborated by the reversible light-induced gel-sol transition of L_1 at 50 mM (Fig. S11f†).

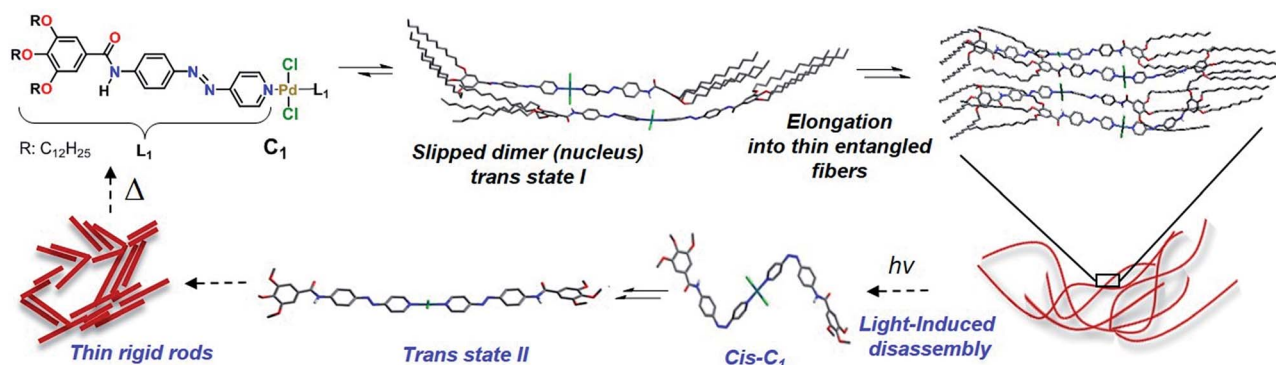
Hierarchical self-assembly of $Pd(L_1)_2Cl_2$

We envisioned that the complexation of metal ions such as Pd(II) by L_1 to yield $Pd(L_1)_2Cl_2$ (C_1 , Scheme 2) would cause a significant impact on the photoresponsive and self-assembly behavior. This metal ion was selected not only based on the expertise of our group in Pd(II)-based assemblies¹⁹ but also due to their preorganized coordination geometry and aggregation

propensity. Unlike *trans*- L_1 , *trans*- C_1 readily self-assembles at room temperature in nonpolar solvents such as MCH (see solvent-dependent UV-Vis studies in Fig. S12†). Fig. 3a shows the spectral changes upon cooling a monomer solution of C_1 in MCH from 363 K to 283 K with 1 K min^{-1} ramp. On cooling, the metal-to-ligand charge transfer (MLCT) transition at 395 nm broadens progressively and decreases in intensity whereas a shoulder at 460 nm becomes apparent (Fig. 3a). This spectral change is suggestive of aggregation. Monitoring the absorbance at 395 nm vs. temperature (T) at four different concentrations yielded non-sigmoidal plots, which could be accurately fitted to the nucleation-elongation cooperative model (Fig. 3a, inset and Fig. S13, S14†).²⁷ According to this model, the formation of a small aggregate (nucleus) is needed to activate the supramolecular growth into fiber-like structures (for an overview of the thermodynamic parameters, see Fig. S13, S14 and Table S1†).

VT-DOSY and DLS measurements of C_1 further support the formation aggregates in solution (Fig. S15 and S16†). ROESY NMR of an aggregate solution (7.5 mM, 358 K) in MCH- d_{14} revealed the presence of five new cross-peaks (highlighted in coloured squares in Fig. 3b) that are absent in COSY studies. For instance, correlation signals between protons H_c and H_f as well as H_d and H_f can be distinguished (Fig. 3b). As these protons are within the vicinity of 5 Å, they should result from an intramolecular coupling. On the other hand, cross-peaks between H_a and H_f (green), H_b and H_c (blue) and H_a and H_d (red) can only be due to intermolecular contacts, as the respective protons are very far from one another (>5 Å). Furthermore, additional intermolecular interactions between $-O-CH_2-$ protons of the alkyl chains (H') and aromatic protons H_{a-d} were also identified (black). This coupling pattern is in agreement with the formation of slipped stacks stabilized by $\pi-\pi$ interactions between the aromatic rings of the ligands as well as $N-H \cdots Cl-Pd$ hydrogen bonding interactions (Fig. 3c), as recently proposed for related π -conjugated Pd(II) complexes.²⁸ The fingerprints associated with these interactions could be also identified by FT-IR measurements in MCH (Fig. S17†).²⁸

This stacking arrangement in solution is in perfect agreement with the molecular packing extracted from X-ray analysis of single crystals grown from DCM/acetonitrile of a nearly identical Pd-azo derivative (C_2) with short ethyl groups. The



Scheme 2 Proposed hierarchical self-assembly and photoresponsive behavior of C_1 . Structures of monomers and small aggregates were optimized using DFT (PBE0/6-31G*/LANL2DZ) and dispersion-corrected PM6, respectively.



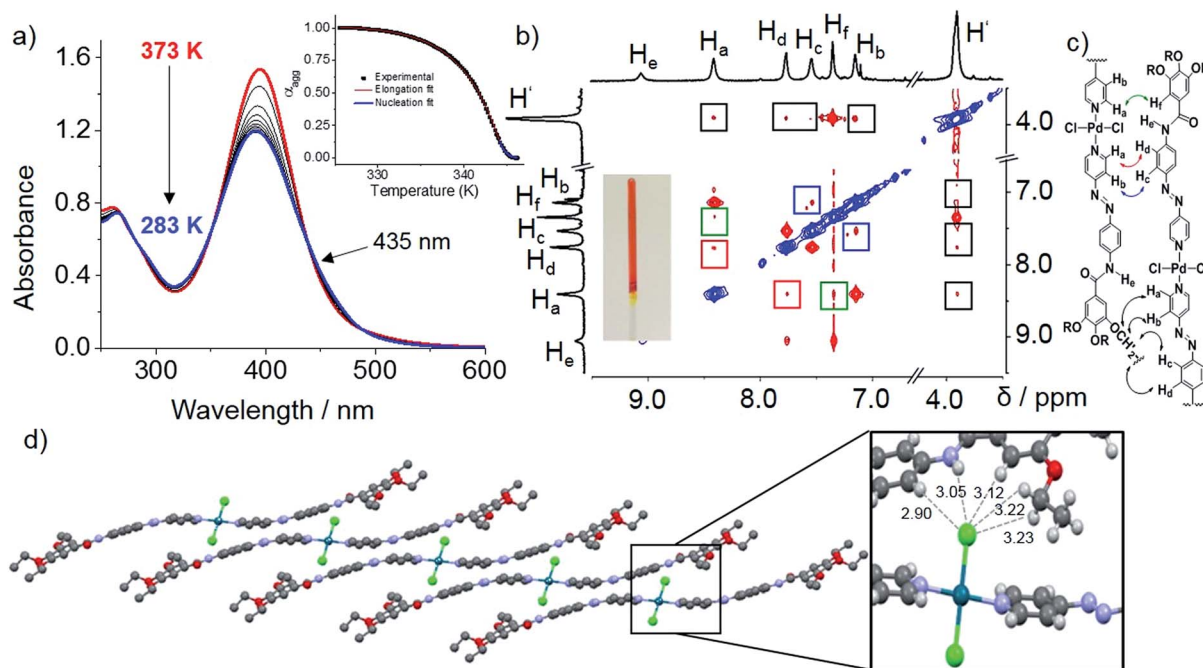


Fig. 3 (a) Temperature-dependent UV-Vis spectra of C_1 at 5×10^{-5} M; inset: plot of α_{agg} vs. T for C_1 at 5×10^{-5} M monitored at 395 nm. (b) COSY and ROESY of C_1 at 7.5 mM in MCH- d_{14} at 358 K and photograph of gel (inset). (c) Plausible molecular arrangement. (d) Packing of C_2 (b-axis) driven by N-H...Cl and C-H...Cl interactions (inset).

crystal structure analysis showed a marked molecular curvature of the azo-based pyridyl-ligands on both sides of the metal center (Fig. S18 and S19[†]). The packing in the crystal structure is mainly driven by three types of cooperative weak interactions: C-H...Cl, N-H...Cl interactions and π - π stacking. In analogy with the packing deduced by ROESY studies, the monomer units are arranged in a slipped fashion driven by a combination of one N-H...Cl and four C-H...Cl intermolecular interactions. Each Cl is interacting with two aromatic protons (H_d and H_f), an NH group of the amide moiety and two polarized methylene groups belonging to the ethoxy chains of a neighboring unit (Fig. 3d, inset). A further growth of the system into layered structures is facilitated by lateral interactions of the formed 1D stacks *via* 4 C-H...Cl and 4 C-H...O interactions. Further, π - π interactions are stabilizing the packing along the *a*-axis (Fig. S20[†]). These overall results highlight the key impact of NH...Cl interactions on slipped stacking stabilization.

In contrast to the free ligand L_1 , C_1 forms considerably thinner and shorter well-defined fibers (5–10 nm in width and 60–150 nm in length) in MCH (Fig. S21[†]). This difference in morphology clearly reflects different molecular packing with distinct intermolecular interactions (π - π , C-H...Cl and N-H...Cl, *vide supra*) compared to the free ligand L_1 . As the NMR signals of C_1 are nearly unidentifiable in pure MCH- d_{14} due to strong aggregation, a solvent mixture with 10% $CDCl_3$ was chosen for further studies (Fig. S22[†]). Prior to the NMR experiments, we confirmed by VT-UV-Vis studies under identical conditions that the addition of 10% $CHCl_3$ does not influence the aggregation behavior of C_1 (Fig. S23[†]). AFM analysis showed the formation of a network of thin entangled fibers further supporting an identical aggregation behavior in pure MCH and

10% $CHCl_3$ -MCH at 5×10^{-5} M (Fig. 4b and S24[†]). The powder X-ray diffraction pattern of a thin film of C_1 showed the formation of a hexagonal columnar structure with the lattice parameter of $a = 3.5$ nm, which is larger than that of L_1 ($a = 2.45$) in a tetragonal columnar structure (Fig. S25[†]). Probably, the well-defined fibers of C_1 visualized by AFM (Fig. 4b and S24[†]) are the elementary structure composed of one-dimensionally stacked C_1 .

Photoisomerization of C_1

Irradiation at 370 nm of the entangled thin fibers of C_1 (MCH : $CHCl_3$ (9 : 1), $c = 5 \times 10^{-5}$ M, see Fig. 4b and S24[†] for AFM images) at 283 K for 30 min caused a small red shift in the absorption maximum and significant hyperchromism (Fig. 4a). Because the UV-Vis spectral changes cannot be explained by usual *trans*-to-*cis* isomerization of azobenzene, we assume this to be due to a photo-induced disassembly *via* formation of *cis*- C_1 followed by a rapid back-isomerization to *trans*- C_1 , which is most likely kinetically trapped. After finishing the UV irradiation for 30 min, we monitored the spectral changes over a period of 800 min while keeping the solution at 283 K (Fig. 4a). A blue shift in the absorption maximum from 397 nm to 391 nm with a small absorption change at around 490 nm was observed with multiple isosbestic points (Fig. 4a). Without any lag, A_{395} started decreasing and reached a plateau after several hours (Fig. 4a, inset), indicating the formation of photo-reconstructed aggregates. Comparison of this spectrum with that corresponding to the aggregation process of C_1 without irradiation (see Fig. S23 and S3a[†]) reveals a shift of the isosbestic point from 435 nm to 460 nm.



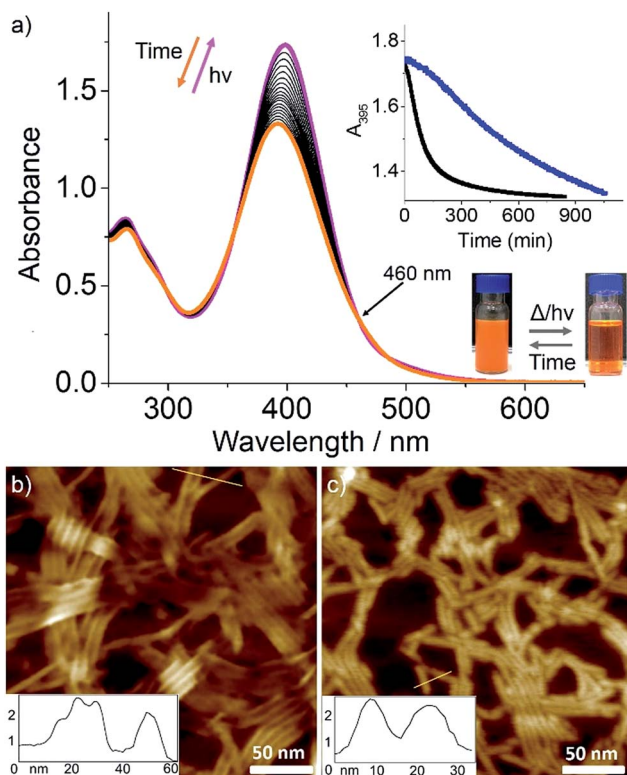


Fig. 4 (a) Time-dependent UV-Vis spectra (5×10^{-5} M MCH/CHCl₃ (90 : 10)) of C₁ upon light irradiation and subsequent quenching to 283 K; inset: plot of A_{395} vs. time before (blue) and after irradiation (black) at 283 K and photographs showing opacity changes upon irradiation for 15 min. AFM images of C₁ aggregates formed under equivalent conditions as for UV-Vis: (b) before and (c) after UV irradiation for 30 min at 283 K with corresponding cross-section analysis along the yellow line.

In order to compare the above photochemically achieved kinetic state with that obtained by quick temperature drop (quenching), we rapidly cooled a hot MCH/CHCl₃ solution of C₁ (5×10^{-5} M) to 283 K and monitored the UV-Vis spectral changes for 1000 min (Fig. S26[†]). These studies showed significant differences compared to the UV-irradiated sample, *i.e.*, a blue shift of the absorption maximum from 398 nm to 394 nm with only one isosbestic point at around 470 nm. A plot of A_{395} vs. time showed a slow decay compared to the irradiated sample (Fig. 4a, inset). These results indicate that the self-assembly of C₁ after the photo-induced disassembly proceeds through a different nucleation-elongation mechanism compared to that from the thermally obtained monomeric C₁.

We next attempted to identify the possible *cis*-C₁ formation by ¹H-NMR measurements. The observed rapid transformation of the initially slightly opaque solution of *trans*-C₁ in 1 : 9 CDCl₃-MCH-d₁₄ at 5×10^{-4} M into a clear solution upon irradiation for 30 min implies a disassembly of *trans*-C₁ aggregates (Fig. 4, inset). However, to our surprise, no resonances corresponding to *cis*-C₁ were identified. Dissociation of the N-H...Cl hydrogen bonds was evident from ¹H-NMR where the amide signal shifts from 9.41 to 9.16 ppm upon irradiation (see Fig. S27[†]). Although nearly complete dissociation of *trans*-C₁ ($c = 9$ mM) aggregates was observed upon irradiation in CDCl₃

(Fig. S28[†]), no signals corresponding to *cis*-C₁ were identified. To justify the UV-induced disassembly of C₁ aggregates, we performed ¹H-NMR experiments of a structurally related OPE-based Pd(II) complex **1** lacking the photo-responsive unit (Fig. S29[†]), which was previously observed to self-assemble *via* N-H...Cl hydrogen bonds.²⁸ In contrast to C₁, irradiation of **1** for 30 min showed negligible NMR shifts (Fig. S29[†]). Accordingly, the disassembly of *trans*-C₁ aggregates upon irradiation is ascribable to the formation of bulky *cis*-C₁.

A plausible explanation for the absence of *cis*-C₁ during the NMR measurements can be related to the change in the excited state dynamics of L₁ upon Pd(II) coordination,^{18c,29} which has been inspected by Density Functional Theory (DFT) and Time-Dependent DFT (TD-DFT) calculations. Initially, irradiation of the *trans*-state I (Scheme 2), in which the carbonyls of each complex are antiparallel to each other, populates a molecular orbital which is antibonding with respect to both azo nitrogens (inset in Fig. S30[†]). The N=N double bond is then broken and rotation around this bond generates the *cis*-C₁ isomer. The rather distorted geometry of the latter (Scheme 2 and Fig. S31[†]) leads to the dissociation of the aggregates. The lowest excited state of *cis*-C₁ is almost resonant with that of *trans*-C₁-II, where carbonyl groups are now oriented parallel to each other (Fig. S32[†]), suggesting a fast *cis*-C₁ → *trans*-C₁-II conversion. The relative orientation of carbonyl groups inside a fiber, namely parallel *vs.* antiparallel, can strongly influence the energy of excited states, whose differences can be as high as 0.7 eV (Fig. S33[†]). As anticipated from the above calculations, the photo-reconstructed C₁ aggregates show appreciable morphological changes compared to the nanostructures before UV irradiation. The AFM images show that the initially formed thin flexible fibers with several μm in length (Fig. 4b and S24, S26[†]) transform into short rods with maximum length of 20–100 nm (Fig. 4c and S34[†]). Supramolecular systems in which all carbonyl groups are pointing in the same direction, like fibers of *trans*-C₁-II, can form giant dipole moments or macrodipoles, influencing the interaction between nearby fibers as well as the final morphology of the material.³⁰ Because the local accumulation of macrodipoles in dense regions containing supramolecular fibers is not thermodynamically favorable,³¹ further growth of fibers of *trans*-C₁-II becomes frustrated, explaining why they are shorter. On the other hand, local accumulation of macrodipoles does not occur in fibers of *trans*-C₁-I because the carbonyls are oriented antiparallel to each other, stabilizing the fibers and allowing them to grow much further (notice that the crystal structure shown in Fig. 3d has also antiparallel amide groups).

Conclusions

We have reported a new small molecule-based supramolecular system (azo-based pyridyl ligand L₁) that undergoes significant changes in its hierarchical self-assembly and aggregate morphology upon response to UV-light irradiation and metal coordination. Combined experimental and theoretical studies allowed us to propose a mechanism for its hierarchical self-assembly behavior. L₁ self-assembles in the absence of



UV-light and metal ions into long twisted fibers driven by hydrogen bonding and π -stacking *via* the formation of an antiparallel dimer species. UV irradiation and subsequent photoisomerization of L_1 leads to a new non-H-bonded packing mode that ultimately results in the formation of shorter rigid rods. Complexation of $PdCl_2$ drastically changes the mode of aggregation from antiparallel to slipped stacks driven by N-H...Cl interactions with subsequent increase in the degree of cooperativity of the supramolecular growth. Finally, these long thin fibers transform into thinner, shorter rods upon UV irradiation *via* a reorganization into a different *trans* conformation of the Pd(II) complex. The stability of short *vs.* long fibers with Pd(II) was rationalized in terms of the accumulation of macro-dipoles, which depend on the relative orientation of carbonyl groups inside the fibers, namely parallel or antiparallel. Our results have allowed us to unravel for the first time the impact of metal coordination and light irradiation on hierarchical self-assembly processes. In our opinion, the present study represents a starting point towards the development of a new class of stimuli-responsive self-assembled materials combining the properties of metal ions and light.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We gratefully acknowledge Dr Mitsuaki Yamauchi (Kwansei Gakuin University) and Dr Xu Lin (Southwest Forestry University) for experimental assistance. We thank the European Commission (ERC StG SUPRACOP 715923) and the Humboldt Foundation (Sofja-Kovalevskaja) for funding. This work was partially supported by KAKENHI grant (26102010 for S. Y.) and the Grant-in-Aid for Scientific Research on Innovative Areas "π-Figuration" (26102001) of the Japanese Ministry of Education, Culture, Sports, Science, and Technology (MEXT). We thank Ana-Maria Krause for X-Ray analysis.

Notes and references

- (a) G. M. Whitesides, J. P. Mathias and C. T. Seto, *Science*, 1991, **254**, 1312–1319; (b) T. F. A. De Greef, M. M. J. Smulders, M. Wolfs, A. P. H. J. Schenning, R. P. Sijbesma and E. W. Meijer, *Chem. Rev.*, 2009, **109**, 5687–5754; (c) T. Aida, E. W. Meijer and S. I. Stupp, *Science*, 2012, **335**, 813–817; (d) J. D. Tovar, *Acc. Chem. Res.*, 2013, **46**, 1527–1537; (e) C. Kulkarni, S. Balasubramanian and S. J. George, *ChemPhysChem*, 2012, **14**, 661–673; (f) S. S. Babu, V. K. Praveen and A. Ajayaghosh, *Chem. Rev.*, 2014, **114**, 1973–2129; (g) C. Rest, R. Kandaneli and G. Fernandez, *Chem. Soc. Rev.*, 2015, **44**, 2543–2572; (h) A. Sandeep, V. K. Praveen, K. K. Kartha, V. Karunakaran and A. Ajayaghosh, *Chem. Sci.*, 2016, **7**, 4460–4467; (i) D. B. Amabilino, D. K. Smith and J. W. Steed, *Chem. Soc. Rev.*, 2017, **46**, 2404–2420; (j) M. Hifsudheen, R. K. Mishra, B. Vedhanarayanan, V. K. Praveen and A. Ajayaghosh, *Angew. Chem., Int. Ed.*, 2017, **56**, 12634–12638; (k) A. Sorrenti, J. LeiraGlesias, A. J. Markvoort, T. F. A. de Greef and T. M. Hermans, *Chem. Soc. Rev.*, 2017, **46**, 5476–5490; (l) E. E. Greciano, B. Matarranz and L. Sánchez, *Angew. Chem., Int. Ed.*, 2018, **57**, 4697–4701.
- (a) S. Yagai, T. Karatsu and A. Kitamura, *Chem.–Eur. J.*, 2005, **11**, 4054–4063; (b) S. Yagai and A. Kitamura, *Chem. Soc. Rev.*, 2008, **37**, 1520–1529; (c) A. J. McConnell, C. S. Wood, P. P. Neelakandan and J. R. Nitschke, *Chem. Rev.*, 2015, **115**, 7729–7793; (d) M. Kathan and S. Hecht, *Chem. Soc. Rev.*, 2017, **46**, 5536–5550; (e) R. Yamakado, M. Hara, S. Nagano, T. Seki and H. Maeda, *Chem.–Eur. J.*, 2017, **23**, 9244–9248; (f) R. Yamakado, M. Hara, S. Nagano, T. Seki and H. Maeda, *Chem. Lett.*, 2018, **47**, 404–407.
- (a) G. Cravotto and P. Cintas, *Chem. Soc. Rev.*, 2009, **38**, 2684–2697; (b) A. Tsuda, Y. Nagamine, R. Watanabe, Y. Nagatani, N. Ishii and T. Aida, *Nat. Chem.*, 2010, **2**, 977–983.
- Z. Chi, X. Zhang, B. Xu, X. Zhou, C. Ma, Y. Zhang, S. Liu and J. Xu, *Chem. Soc. Rev.*, 2012, **41**, 3878–3896.
- (a) Y. Geng, X.-J. Wang, B. Chen, H. Xue, Y.-P. Zhao, S. Lee, C.-H. Tung and L.-Z. Wu, *Chem.–Eur. J.*, 2009, **15**, 5124–5129; (b) Z. Y. Xiao, X. Zhao, X. K. Jiang and Z. T. Li, *Chem. Mater.*, 2011, **23**, 1505–1511; (c) S. Sato, T. Murase and M. Fujita, *Supramolecular Chemistry: From Molecules to Nanomaterials*, John Wiley & Sons, Ltd, Hoboken, 2012; (d) S. Lee, E. H. Brandon, Y. Liu, R. D. James, W. B. David, L. T. Steven and A. H. Flood, *Chem.–Eur. J.*, 2015, **22**, 560–569; (e) B. Qiao, B. E. Hirsch, S. Lee, M. Pink, C.-H. Chen, B. W. Laursen and A. H. Flood, *J. Am. Chem. Soc.*, 2017, **139**, 6226–6233.
- (a) M.-M. Russew and S. Hecht, *Adv. Mater.*, 2010, **22**, 3348–3360; (b) R. Klajn, K. J. M. Bishop and B. A. Grzybowski, *Proc. Natl. Acad. Sci. U. S. A.*, 2007, **104**, 10305–10309; (c) J. W. Fredy, A. Mendez-Ardoy, S. Kwangmettattam, D. Bochicchio, B. Matt, M. C. A. Stuart, J. Huskens, N. Katsonis, G. M. Pavan and T. Kudernac, *Proc. Natl. Acad. Sci. U. S. A.*, 2017, **114**, 11850–11855.
- (a) H. Tian and S. Yang, *Chem. Soc. Rev.*, 2004, **33**, 85–97; (b) J. J. D. de Jong, L. N. Lucas, R. M. Kellogg, J. H. van Esch and B. L. Feringa, *Science*, 2004, **304**, 278–281; (c) S. Yagai, K. Iwai, T. Karatsu and A. Kitamura, *Angew. Chem., Int. Ed.*, 2012, **51**, 9679–9683; (d) S. Yagai, K. Ishiwatari, X. Lin, T. Karatsu, A. Kitamura and S. Uemura, *Chem.–Eur. J.*, 2013, **19**, 6971–6975; (e) S. Yagai, K. Iwai, M. Yamauchi, T. Karatsu, A. Kitamura, S. Uemura, M. Morimoto, H. Wang and F. Würthner, *Angew. Chem., Int. Ed.*, 2014, **53**, 2602–2606; (f) J. T. van Herpt, J. Areephong, M. C. A. Stuart, W. R. Browne and B. L. Feringa, *Chem.–Eur. J.*, 2014, **20**, 1737–1742.
- P. K. Kundu, D. Samanta, R. Leizrowice, B. Margulis, H. Zhao, M. Börner, T. Udayabhaskararao, D. Manna and R. Klajn, *Nat. Chem.*, 2015, **7**, 646–652.
- (a) A. Harada, Y. Takashima and M. Nakahata, *Acc. Chem. Res.*, 2014, **47**, 2128–2140; (b) S. Engel, N. Möller and B. J. Ravoo, *Chem.–Eur. J.*, 2017, **24**, 4741–4748.



- 10 (a) K. Ichimura, *Chem. Rev.*, 2000, **100**, 1847–1874; (b) J. A. Sarah, F. Lancia, S. Iamsaard, B. Matt, T. Kudernac, P. F. Stephen and N. Katsonis, *Angew. Chem., Int. Ed.*, 2017, **56**, 3261–3265.
- 11 (a) A. Samanta and B. J. Ravoo, *Chem.–Eur. J.*, 2014, **20**, 4966–4973; (b) J. Moratz, L. Stricker, S. Engel and B. J. Ravoo, *Macromol. Rapid Commun.*, 2017, **39**, 1700256.
- 12 R. Rajaganesh, A. Gopal, T. M. Das and A. Ajayaghosh, *Org. Lett.*, 2012, **14**, 748–751.
- 13 A. A. Beharry and G. A. Woolley, *Chem. Soc. Rev.*, 2011, **40**, 4422–4437.
- 14 A. Gopal, M. Hifsudheen, S. Furumi, M. Takeuchi and A. Ajayaghosh, *Angew. Chem., Int. Ed.*, 2012, **51**, 10505–10509.
- 15 (a) M. Yamauchi, T. Ohba, T. Karatsu and S. Yagai, *Nat. Commun.*, 2015, **6**, 8936; (b) B. Adhikari, Y. Yamada, M. Yamauchi, K. Wakita, X. Lin, K. Aratsu, T. Ohba, T. Karatsu, M. J. Hollamby, N. Shimizu, H. Takagi, R. Haruki, S.-i. Adachi and S. Yagai, *Nat. Commun.*, 2017, **8**, 15254; (c) B. Adhikari, X. Lin, M. Yamauchi, H. Ouchi, K. Aratsu and S. Yagai, *Chem. Commun.*, 2017, **53**, 9663–9683.
- 16 M. Endo, T. Fukui, S. H. Jung, S. Yagai, M. Takeuchi and K. Sugiyasu, *J. Am. Chem. Soc.*, 2016, **138**, 14347–14353.
- 17 (a) M. J. Mayoral, C. Rest, V. Stepanenko, J. Schellheimer, R. Q. Albuquerque and G. Fernández, *J. Am. Chem. Soc.*, 2013, **135**, 2148–2151; (b) C. Rest, M. J. Mayoral, K. Fucke, J. Schellheimer, V. Stepanenko and G. Fernández, *Angew. Chem., Int. Ed.*, 2014, **53**, 700–705; (c) N. K. Allampally, M. J. Mayoral, S. Chansai, M. C. Lagunas, C. Hardacre, V. Stepanenko, R. Q. Albuquerque and G. Fernández, *Chem.–Eur. J.*, 2016, **22**, 7810–7816; (d) L. Herkert, A. Sampedro and G. Fernández, *CrystEngComm*, 2016, **18**, 8813–8822.
- 18 (a) M. Burnworth, L. Tang, J. R. Kumpfer, A. J. Duncan, F. L. Beyer, G. L. Fiore, S. J. Rowan and C. Weder, *Nature*, 2011, **472**, 334–337; (b) J. R. Kumpfer and S. J. Rowan, *J. Am. Chem. Soc.*, 2011, **133**, 12866–12874; (c) E. Borré, S. Bellemin-Lapponnaz and M. Mauro, *Chem.–Eur. J.*, 2016, **22**, 18718–18721; (d) M. E. Garah, E. Borré, A. Ciesielski, A. Dianat, R. Gutierrez, G. Cuniberti, S. Bellemin-Lapponnaz, M. Mauro and P. Samori, *Small*, 2017, **13**, 1701790.
- 19 (a) X. Yan, J. F. Xu, T. R. Cook, F. Huang, Q. Z. Yang, C. H. Tung and P. J. Stang, *Proc. Natl. Acad. Sci. U. S. A.*, 2014, **111**, 8717–8722; (b) R. D. Mukhopadhyay, V. K. Praveen and A. Ajayaghosh, *Mater. Horiz.*, 2014, **1**, 572–576; (c) E. Borre, J. F. Stumbe, S. Bellemin-Lapponnaz and M. Mauro, *Chem. Commun.*, 2017, **53**, 8344–8347; (d) R. D. Mukhopadhyay, G. Das and A. Ajayaghosh, *Nat. Commun.*, 2018, **9**, 1987.
- 20 (a) M. J. Mayoral and G. Fernández, *Chem. Sci.*, 2012, **3**, 1395–1398; (b) V. W.-W. Yam, V. K.-M. Au and S. Y.-L. Leung, *Chem. Rev.*, 2015, **115**, 7589–7728; (c) T. R. Cook and P. J. Stang, *Chem. Rev.*, 2015, **115**, 7001–7045; (d) K. Li, G. S. Ming Tong, Q. Wan, G. Cheng, W.-Y. Tong, W.-H. Ang, W.-L. Kwong and C.-M. Che, *Chem. Sci.*, 2016, **7**, 1653–1673.
- 21 (a) J. Garcia-Amoros, S. Nonell and D. Velasco, *Chem. Commun.*, 2012, **48**, 3421–3423; (b) A. Florian, M. J. Mayoral, V. Stepanenko and G. Fernández, *Chem.–Eur. J.*, 2012, **18**, 14957–14961.
- 22 (a) X. Lu, Z. Guo, C. Sun, H. Tian and W. Zhu, *J. Phys. Chem. B*, 2011, **115**, 10871–10876; (b) A. Rödle, B. Ritschel, C. Mück-Lichtenfeld, V. Stepanenko and G. Fernández, *Chem.–Eur. J.*, 2016, **22**, 15772–15777.
- 23 J. Gershberg, F. Fennel, T. H. Rehm, S. Lochbrunner and F. Würthner, *Chem. Sci.*, 2016, **7**, 1729–1737.
- 24 M. Kasha, *Radiat. Res.*, 1963, **20**, 55–70.
- 25 H. Adams, P. L. Bernad Jr, D. S. Eggleston, R. C. Haltiwanger, K. D. M. Harris, G. A. Hembury, C. A. Hunter, D. J. Livingstone, B. M. Kariuki and J. F. McCabe, *Chem. Commun.*, 2001, 1500–1501.
- 26 (a) S. Yagai, T. Nakajima, T. Karatsu, K.-i. Saitow and A. Kitamura, *J. Am. Chem. Soc.*, 2004, **126**, 11500–11508; (b) S. Yagai, T. Nakajima, K. Kishikawa, S. Kohmoto, T. Karatsu and A. Kitamura, *J. Am. Chem. Soc.*, 2005, **127**, 11134–11139.
- 27 P. Jonkheijm, P. van der Schoot, A. P. H. J. Schenning and E. W. Meijer, *Science*, 2006, **313**, 80–83.
- 28 A. Langenstroer, Y. Dorca, K. K. Kartha, M. J. Mayoral, V. Stepanenko, G. Fernández and L. Sánchez, *Macromol. Rapid Commun.*, 2018, 1800191.
- 29 (a) T. Yutaka, I. Mori, M. Kurihara, N. Tamai and H. Nishihara, *Inorg. Chem.*, 2003, **42**, 6306–6313; (b) G. Auböck and M. Chergui, *Nat. Chem.*, 2015, **7**, 629–633.
- 30 M. P. Oliveira, H.-W. Schmidt and R. Q. Albuquerque, *Chem.–Eur. J.*, 2017, **24**, 2609–2617.
- 31 C. S. Zehe, J. A. Hill, N. P. Funnell, K. Kreger, K. P. van der Zwan, A. L. Goodwin, H.-W. Schmidt and J. Senker, *Angew. Chem., Int. Ed.*, 2017, **56**, 4432–4437.

