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'Sacrificial' supramolecular assembly and pressure-induced polymerization: toward sequence-defined functionalized nanothreads

Herein, aryl/perfluoroaryl (Ar/ArF) stacking pre-organizes reactants for precise C—C bond formation through pressure-induced polymerization. The resultant polymer is a one-dimensional alternating structure featuring an sp^3 -rich backbone, and pendant -OH groups. Such rigid backbones are theorized to possess the strength of diamond, the chemical versatility of graphene, and a large insulating band gap. Our approach relies on a retrosynthetic supramolecular design to engineer an alternating sequence embedded in the reactant that transfers completely to the product with high fidelity.

As featured in:



See Elizabeth Elacqua *et al.*,
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Scheme 1 Synthesis of (A) benzene-derived nanotreads¹⁴ and (B) Ar/ArF-derived nanotreads.¹⁵ Approach (C) uses Ar–OH/ArF–OH co-crystals, wherein a functional group bias is introduced that, along with the sacrificial synthons, effectively decreases the required reaction pressure.¹⁶

dimensional poly(adamantane)-like structure featuring an sp^3 -rich backbone.¹⁴ Such sp^3 -based backbones are theorized to possess the strength of diamond,²² the chemical versatility of graphene, and large insulating band gaps.²³ In contrast to simple molecules (*e.g.*, benzene) under pressure, polymerization of this co-crystal produced ordered nanotreads even under fast compression. Despite an electronically-biased pre-organization, the pressure needed for reaction was still quite high, *ca.* 28 GPa.

Here, we report a functionalized nanotread arising from pressure-induced polymerization of an Ar/ArF co-crystal comprised of phenol and pentafluorophenol (Scheme 1). Our

design features a hydroxy substituent, that, in addition to allowing close-packed π - π distances, accesses crystalline nanotreads at markedly lower pressures than those needed for supramolecular designs.¹⁵ The lower reaction pressure is attributed to the electronically activating –OH substituent on the ring system, which facilitates participation of the Ar component (phenol) as a diene to promote polymerization with a lower activation barrier. Importantly, compression of Ar–OH/ArF–OH provides an experimental glimpse at reaction mechanism: prominent C=O and C=C character is observed through IR spectroscopy, suggestive of a keto-enol tautomerization that can only occur after a cycloaddition.

Our motivation lies in developing precisely sequenced and functionalized nanotread polymers using a supramolecular process. In studies to attain alternating copolymer threads, we realized that ‘sacrificial’ supramolecular synthons would provide a strategic entry point toward A-B-type polymers. Given noncovalent interactions typically *template* solid-state reactions while remaining intact, we designate our design – the use of sp^2 -enriched Ar/ArF supramolecular synthons – as sacrificial owing to the solid-state design elements being *consumed* throughout the duration of the reaction. These synthons additionally guide the establishment of a well-defined sp^3 -rich polymer backbone.

Results and discussion

Our design required engineering a 1:1 phenol : pentafluorophenol co-crystal to confirm orientation of the Ar/ArF units and exogenous –OH groups prior to compression. Thus, equimolar phenol and pentafluorophenol were crystallized from hexanes through slow cooling. Molecular and crystalline arrangement of the co-crystal was confirmed using single-crystal X-ray diffraction (XRD), in close similarity to a reported deuterated structure.²⁴ As expected, the structure of the co-crystal is dictated by uniaxial π - π stacking quadrupole-type interactions of the Ar/ArF complex (Fig. 1),²⁵ with the

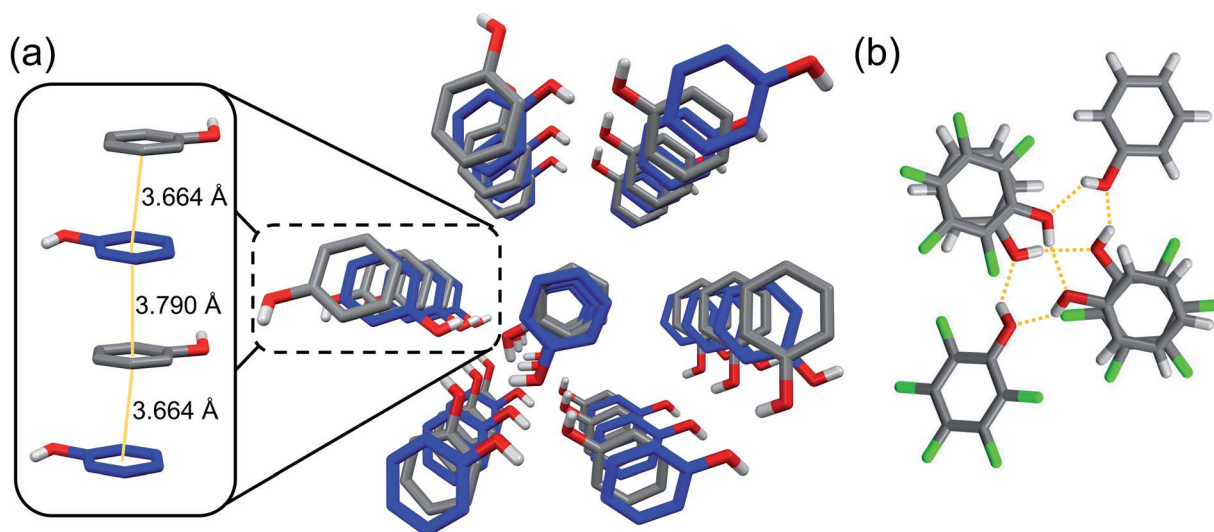


Fig. 1 View of (a) Ar–OH (blue)/ArF–OH (grey) co-crystal along the *a*-axis with the left inset illustrating π - π stacking distance of an Ar/ArF slip-stack along the *b*-axis, and (b) hydrogen-bonding hexameric unit.



functional groups oriented to favor stable hydrogen bonds.²¹ The alternating arrangement of short π - π stacking distances are preorganized for an alternating copolymerization to occur. The co-crystal molecular structure is further guided by the presence of solely (Ar) O-H \cdots O (ArF) and (ArF) O-H \cdots O (Ar) hydrogen bonds between electronically opposed aryl groups (Ar/ArF). The hydrogen bonds ($d_{O\cdots O} = 2.7$ – 2.8 Å) exist entirely between adjacent stacks to form hexameric assemblies (Fig. 1b).

Given high onset pressures ranging from 18–20 GPa for nanothreads derived from benzene and naphthalene/octafluoronaphthalene (Scheme 1), we sought to include an external substituent off of the ring that would realize a lower reaction pressure. We hypothesized that the phenol/pentafluorophenol co-crystal might electronically activate the aryl unit, thus favorably lowering the activation energies required for cycloaddition. The proposed co-crystal affords, in principle, 36 isomers of the [4 + 2] cycloaddition dimer, for which we analyzed the respective energies and activation barriers (see ESI†).²⁶ With phenol acting as the diene, the average activation barrier is 59 kcal mol⁻¹. When pentafluorophenol acts as the diene, the calculated activation barrier increases to 61 kcal mol⁻¹. The determined activation barrier for our Ar/ArF co-crystal is significantly lower than that for benzene-based threads (68 kcal mol⁻¹), suggestive of a lessened onset pressure. Interestingly, we found that the four lowest energy isomers suggest that the C–C bonds within a dimer are *meta* and *para* to the hydroxyl group of the ArF unit. These positions are the most likely to exhibit electron-rich character through resonance and are abundant in the precursor crystal as probable reaction sites.

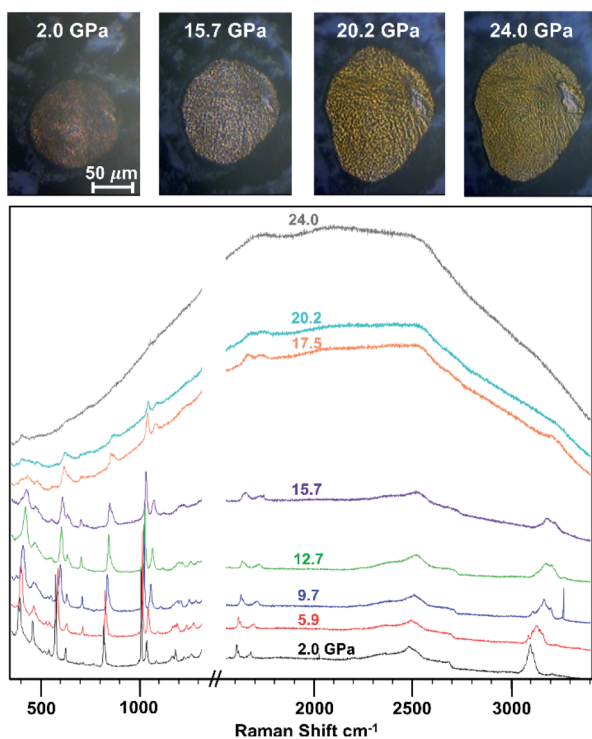


Fig. 2 *In situ* Raman spectral overlay depicting 1:1 phenol : pentafluorophenol co-crystal throughout compression. The top images are taken with the Raman microscope during compression.

Upon loading the co-crystal in a diamond anvil cell, the powder sample was slowly compressed and analyzed using *in situ* Raman microscopy (Fig. 2). At 5.9 GPa, we observed the appearance of peaks at 3160 cm⁻¹, characteristic of sp³-hybridized C–H stretching and/or a phase-induced new sp²-hybridized C–H mode. At 5.9 GPa, we additionally noted a decrease in the ring breathing mode at 1007 cm⁻¹, most likely owing to the formation of pressure-induced metastable adducts such as oligomeric threads present below 15 GPa.

Between 15.7 GPa and 17.5 GPa, further reaction occurred, as the sample began to appear yellow-orange, the photoluminescent (PL) background increased, and the gasket deformed (Fig. 2). In contrast, none of these signature characteristics were observed in compression attempts of the individual components (Fig. S2†). Instead, compression yielded mainly amorphous character, with higher reaction pressures being required to observe significant changes. In the co-crystal, an increase in PL background, change in C–H stretching modes, and decrease in ring breathing modes all occurred below 18 GPa. These combined observations confirmed that the lowered onset pressure in comparison to both benzene and the naphthalene:octafluoronaphthalene co-crystal can be attributed to the –OH functional groups favorably providing an electronic bias upon reaction.

The final product was analyzed through X-ray diffraction. The observed six-fold diffraction pattern (Fig. 3) suggests that nanothreads were formed with quasi-hexagonal packing. The diffraction pattern indicated *d*-spacings of 6.72 Å for one Friedel pair and 6.75 Å for two others.²⁷ We attribute the distinct *d*-spacings to different orientations of the exogeneous hydroxyl groups around the thread axis. Since the width and subsequent packing of the Ar–OH/ArF–OH co-crystal thread is likely determined by the ArF–OH ring, the small difference in *d*-spacings is consistent with the ArF ring being approximately circular due to similar C–F and C–OH bond lengths; this supposition is examined further in the simulations of packing described below.

A diffraction simulation of the co-crystal at ambient pressure along the Ar/ArF stacks illustrates a similar pattern in that two

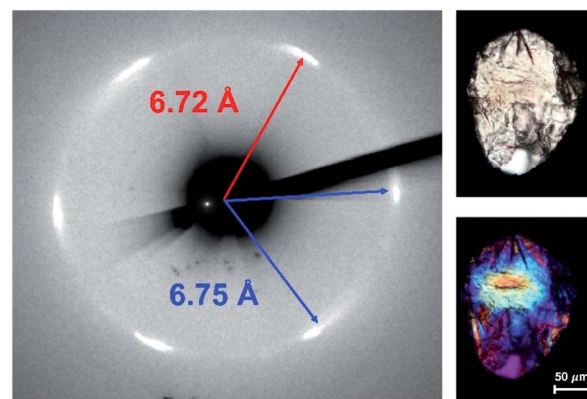


Fig. 3 (Left) Hexagonal six-fold X-ray diffraction pattern obtained for the functionalized nanothread; and (right) 50 \times images within a stainless-steel gasket, illustrating crystallinity along with (bottom) a polarized light micrograph using a 530 nm waveplate.



Friedel pairs have similar d -spacings, here 5.83 Å, with an additional d -spacing of 5.78 Å for the other. The increased d -spacing in the product suggests polymerization from an sp^2 supramolecular-bound co-crystal composed of molecular rings to an sp^3 -rich architecture,¹⁶ necessitating an expansion due to rigid covalent constraints. In addition, the functionalized and crystalline nanothread product exhibits birefringence when viewed under polarized light (Fig. 3), suggestive of optical anisotropy.

In designing simulations of how functionalized nanothreads may pack, the combinatorial complexity of kinetic pathways (*i.e.*, several possible [4 + 2] cycloaddition products) motivated a strategy of extracting general lessons from illustrative chiral and achiral thread categories. We therefore investigated the packings of nanothreads based on two archetypes: chiral polytwistane and achiral (3,0)-type²⁸ backbone, both adapted to the alternating Ar-OH/ArF-OH-derived stack. These structures were packed into crystals and relaxed; d -spacings were then calculated (see ESI†). For both types, many isomers exist, reflecting different positions of the -OH groups for polytwistane, as well as eclipsed, gauche, and conserved hexameric hydrogen-bonding pattern arrangements for the (3,0)-type threads (Fig. 4).

Before addressing the detailed results, we note that the ~ 1.1 Å increase in d -spacing from benzene-derived nanothreads to the Ar-OH/ArF-OH-derived threads (*i.e.*, from 5.6 Å to 6.7 Å) corresponds to an increase of about 1.3 Å in the interthread center-to-center separation. The change from C-H to C-F corresponds to

a 0.24 Å increase in bond length and a 0.37 Å increase in the van der Waals radius of an F-atom compared to H. Doubling this ~ 0.6 Å expansion yields a center-to-center dilation of 1.2 Å, as close to the observed 1.3 Å as would be expected for a rough empirical estimation. A chiral structure, such as the helical polytwistane, cannot easily assume a consistently anisotropic interthread hydrogen-bonding pattern, thus, the simulated d -spacings are closely grouped in magnitude.

Achiral structures such as thread (3,0), in contrast, can naturally assume either near-isotropic (*e.g.*, eclipsed B) or anisotropic (*e.g.*, eclipsed A, gauche) interthread bonding geometries. The experimental observation of a near-hexagonal symmetry with consistent anisotropy in d -spacings rules out an achiral structure with anisotropic inter-thread bonding such as eclipsed A or gauche (absent sufficient structural defects). The information available at present is not definitive enough to decide a specific thread backbone and substituent pattern therein.

Given the successful compression, we sought to understand more about the final nanothread molecular structure and elucidate any mechanistic details that might result from the -OH units being incorporated. We, thus, studied the IR spectroscopic features of the nanothread product in comparison to the starting co-crystal. Prior to compression, the experimental IR spectrum featured prominent modes (Table S5†), including a sharp and strong signal at 3418 cm^{-1} indicative of the O-H \cdots O hydrogen bonding, and sp^2 C-H stretching around 3200 cm^{-1} (Fig. 5, black

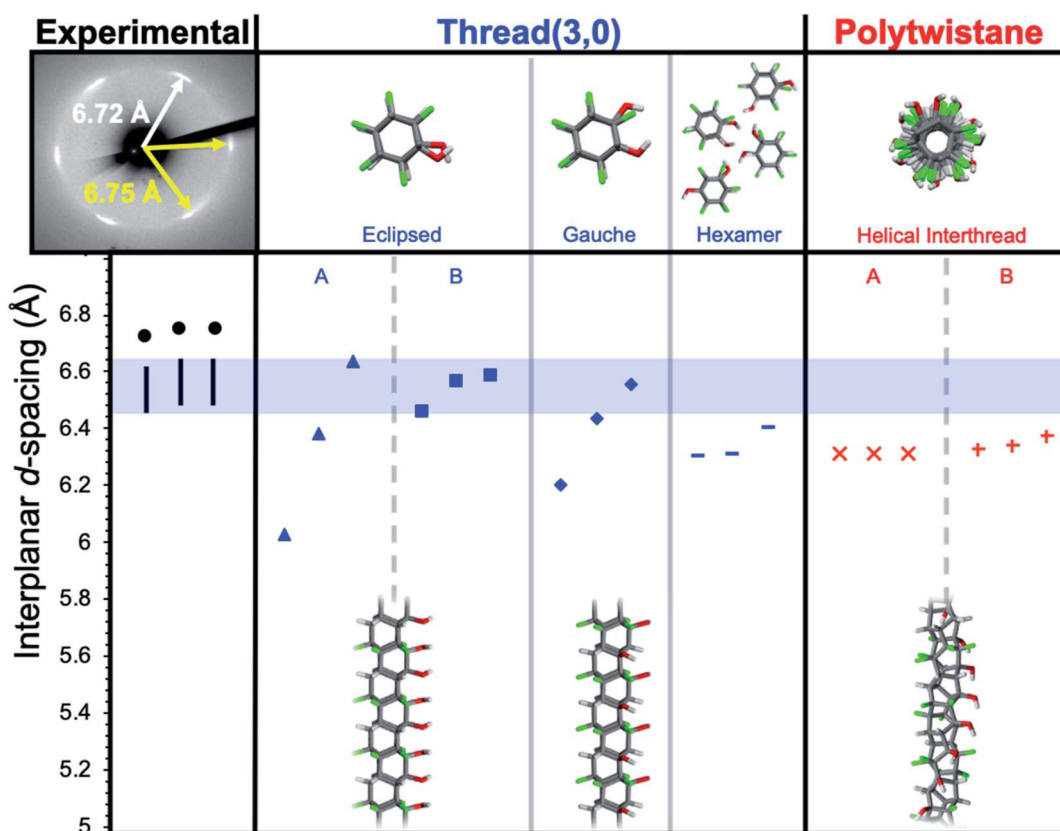


Fig. 4 Comparison of interplanar d -spacings for simulated and experimental X-ray diffraction patterns along the co-crystal a -axis. Simulations account for 1–3% subtraction for thermal expansion.



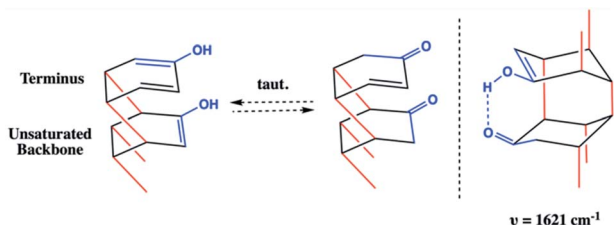


Fig. 5 Comparison IR spectra of precursor co-crystal at ambient pressure (black) and final nanothread (red).

trace). These modes were in agreement with a simulation of the co-crystal at ambient pressure (Fig. S4†).

In contrast, the IR spectrum of the final nanothread product (Fig. 5) highlighted a larger degree of hydrogen bonding, evidenced by the broadened O–H stretching between 3200–3700 cm^{-1} . The nanothread product also featured sp^3 C–H and sp^2 C–H stretching centered around 2800 cm^{-1} . Additional modes corresponding to both C–F and C–H bending and stretching in the region of 1000–1400 cm^{-1} were observed, as expected. In addition, the IR spectrum features a significant C=O signal centered around 1700 cm^{-1} , as well as a sharp C=C stretching peak.

Whereas the C=C signal could result from unreacted co-crystal, the two combined vibrational modes are more likely the result of rapid tautomerization of isolated enolic intermediates to their more prominent keto forms (Scheme 2). Each successful keto–enol tautomerization could, thus, turn an electronically-activated ‘hot’ diene into a deactivated, α,β -unsaturated ketone (akin to a dienophile) or an isolated alkene within the thread structure. Evidence of such a structural detail can also be observed in the IR spectrum. Signals around 1620 cm^{-1} are commonly observed in 1,3-diketone species wherein an enolic form predominates at one position,^{29,30} giving



Scheme 2 Schematic depicting representative cycloadducts capable of (left) facilitating keto–enol tautomerization and (right) intramolecular enol-assisted hydrogen-bonding. H- and F-atoms have been removed for clarity.

rise to a cyclic intramolecular hydrogen-bonding vibration. The strong shoulder on the C=O stretching at 1621 cm^{-1} could suggest the analogous diketone pattern that can arise in proximal rings upon cycloaddition to favor the formation of an enolic H-bond donor within the thread backbone (Scheme 2). Given these signatures of tautomerization are only plausible after an addition reaction that retains some degree of sp^2 character, the IR spectral features – combined with the lowering of reaction pressure through installation of a functional group able to induce polarization – suggest that a [4 + 2] Diels–Alder-like cycloaddition is a key step within initiation/propagation events leading to nanothread formation.

Conclusions

In conclusion, we demonstrate that a functional Ar/ArF-based co-crystal undergoes an electronically-activated pressure-induced polymerization and results in a well-defined crystalline nanothread polymer, as evidenced by a hexagonal X-ray diffraction pattern. Our investigations have highlighted three salient points: (i) the thread-forming polymerization is tolerant of exogenous functional groups, such as –OH; (ii) the combination of ‘sacrificial’ supramolecular assembly and intact hydrogen bonding leads to a lower activation barrier for the reactants; and (iii) isolation of the product highlights key features: the presence of C=O in concert with intense C=C signals in the IR spectrum suggests tautomerization of –OH groups after a [4 + 2] cycloaddition as a mechanistic lynchpin in nanothread formation. The combined results suggest that the polymerization is influenced by electronic substituents which act to reduce energetic barriers for initiation of the cycloaddition-based polymerization, thus lowering the required reaction pressure.³¹ Future work will look to pinpoint the effect of electronic bias on polymerization to gain further insights into mechanism, functional group tolerance, and precise theoretical structure elucidation to provide the intuition for designs amenable to post-polymerization functionalization.³²

Experimental section

Synthesis of functionalized nanothreads

Compression of the Ar/ArF co-crystal was conducted in a symmetric diamond anvil cell (DAC) with a 400 μm culet while monitoring the pressure-induced polymerization using *in situ* Raman spectroscopy. The solid co-crystal was cooled in the fridge, ground using a mortar and pestle, and placed on the lower diamond within a stainless-steel gasket. A ruby chip was added to the sample chamber to monitor the pressure through fluorescence. The DAC was closed to form a powder sample by applying 2.0 GPa of pressure. A typical compression proceeded over the course of 16 hours to a maximum pressure of 23 GPa. The compression rate slowed upon the incidence of a photoluminescent (PL) background that indicated polymerization occurring at around 11 GPa to 2–3 GPa h^{-1} . The compression rate was slowed further above 17 GPa to less than 1 GPa h^{-1} . Decompression progressed over the course of 16–18 hours with similar rates on the descent.



Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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- We define the activation barrier to be the necessary energy to initiate formation of an sp^3 -hybridized carbon, typically through dimer formation, concomitantly exposing a reactive diene.
- A powder CeO_2 standard was utilized before diffraction collection and witnessed a FWHM of 0.3 Å. We thus have reasoned that the 6.72 Å and 6.75 Å d -spacings are not symmetrically equivalent, but we cannot rule out that the two recorded 6.75 Å Friedel pairs are or are not symmetrically equivalent to one another by analytical precision.
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- Reduction of the reaction pressure would allow for the use of pressure apparatus with larger volume output. Diamond anvil cells (DACs), such as those used in these studies, operate with a sample area equating to 100 μm , thus affording nanograms of material. The lessened amount of product from DACs precludes conventional analysis including TGA and DSC, as well as post-polymerization functionalization.
- Our current efforts are focused on scalable syntheses (e.g., by using a Paris–Edinburgh press that can afford mg quantities) to further interrogate thermal properties and post-synthetic modification.

