Mixed Conductive Soft Solids by Electrostatically Driven Network Formation of a Conjugated Polyelectrolyte

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ABSTRACT: A conjugated polyelectrolyte (poly[2,6-(4,4-bis-potassium butanylsulfonate-4H-cyclopenta-[2,1-b;3,4-b’]-dithiophene)-alt-4,7-(2,1,3-benzothiadiazole)], PCPDTBT-SO3K) assembles into a novel, hierarchical hydrogel structure with all structural evidence indicating dominant electrostatic rather than aromatic or mesogen interactions. PCPDTBT-SO3K forms an entangled polymer mesh, where polymer chains are tied together by ionic cross-links, comprising microgel clusters that percolate to form a macroscopic three-dimensional network. With increasing temperature, ions gain mobility to move toward the exterior of the microgel clusters, dissolving the ionic cross-links and inhibiting network percolation through electrostatic repulsion. While \( \pi-\pi \) stacking interactions may be present in a disorganized fashion, no long-range \( \pi-\pi \) stacking is evident in X-ray scattering. Soft materials based on PCPDTBT-SO3K remain semiconducting and exhibit elevated ionic conductivity at the structural reorganization temperature.

INTRODUCTION

Conjugated polyelectrolytes (CPEs) combine a \( \pi \)-conjugated polymer backbone with pendant ionic groups to create water-soluble semiconductors.1 The aqueous solubility provides an opportunity to form functional hydrogel structures for interfacing with biology.2−4 Modulation of the chemistry of the polymer backbone enables tunability of charge transport, ionization potential, and optical properties. Ionic groups and charge compensating counterions allow control over solubility, charge transport, molecular packing, and solution aggregation.5 Studies to understand the structure−property relationships of CPEs have led to increased chemical diversity and a variety of new synthetic protocols for controlled materials chemistry.6,7 The breadth of developed CPEs has inspired work into their application in biological and chemical sensors8−12 and organic optoelectronic devices, including light-emitting diodes (LEDs),13,14 light-emitting electrochemical cells (LEECs),15 photovoltaic devices,16 and light-emitting transistors (LETs).17 Despite the key role CPEs could play in energy conversion and storage, their solution and hydrogel structures and resulting physicochemical material properties remain largely unknown.

Poly[2,6-(4,4-bis-potassium butanylsulfonate-4H-cyclopenta-[2,1-b;3,4-b’]-dithiophene)-alt-4,7-(2,1,3-benzothiadiazole)], PCPDTBT-SO3K (Figure 1) is an attractive model system for elucidating the structure−property relationships of CPEs. Side-
chain and counterion modifications have allowed these CPEs to be instructive model systems for understanding the factors contributing to thermoelectric performance and electronic conductivity.18,19 In particular because the anionic, narrow-band gap conjugated polyelectrolyte self-dopes in aqueous solution to form a water-soluble, stable, conductive polymer.20 PCPDTBT-SO$_3$K has increased performance in a variety of optoelectronic applications, attributed to increases in interfacial contact and surface coverage between device layers.21—24 CPE hydrogels and solution structures pose an opportunity for improved interfacial contact in bioelectrochemical cells and bioelectronics, particularly in neural interfaces.3 Utilizing the advantageous interfacial modifying properties of CPEs as soft, conductive matrices at the abiotic–biotic interface has increased conversion efficiencies in bioelectrochemical cells through improved contact between the microorganisms and the electrodes.5 However, little is known about the structure of soft conductive CPE matrices.

The interconnection between structure and macroscopic photophysical properties has been explored in neutral conjugated polymers, yet as opposed to CPEs, these are absent of appreciable electrostatic interactions. The hierarchical assembly (i.e., network formation) of these materials in poor and concentrated solutions is driven by aromatic interactions yielding π−π stacked, crystalline sheet-like aggregates embedded in an entangled polymer mesh.25—27 Because CPEs incorporate more complex and interchain interactions, their microstructure should be driven not only by a combination of chain rigidity and aromatic interactions but also by electrostatic and hydrophobic effects in an aqueous solvent. Recent studies have suggested that semidilute solutions28,29 of CPEs might form polyelectrolyte micelles that percolate to yield aggregates at long length scales. The micellar conformation has been suggested to result from the amphiphilic nature of CPEs and presumed backbone π−π stacking interactions. However, the delicate balance of the competitive interactions that result in structure formation remain poorly understood. The electrostatic interactions between charged polymer segments are repulsive, although partially screened by the environment, while π−π and other hydrophobic interactions are net attractive. Ionic cross-linking is also possible, engaging counterions as intermediaries between charged backbone units and representing a net attraction among chain segments. The balance of attractive and repulsive interactions allows for finite sized aggregates that under some conditions can percolate into extended networks.

Herein, we seek a more comprehensive understanding of the interplay between the different molecular interactions on the structure and critical transitions of a CPE hydrogel across a temperature range relevant for use as a processable, conductive, soft matrix. PCPDTBT-SO$_3$K is used to explore the interactions that lead to assembly and network formation of conjugated polyelectrolytes in aqueous solution at moderate concentrations. In particular, by investigating the structure of a PCPDTBT-SO$_3$K hydrogel across a range of temperatures, the electrostatic interactions are found to dominate the structural ordering. Finally, because the percolated microgel pathway within these systems is demonstrated to provide a scaffold for both ionic and electrical conductivity, such conjugated polyelectrolyte hydrogels are well-suited for applications in bioelectronics.

### RESULTS AND DISCUSSION

This work illustrates the relative dominance of electrostatic interactions in determining the microstructure of soft materials formed by PCPDTBT-SO$_3$K in the semidilute concentration regime. A novel, multilength scale structure of PCPDTBT-SO$_3$K hydrogels is proposed, and a mechanism for structural transitions is presented. PCPDTBT-SO$_3$K, a semiflexible polymer, is observed in aqueous solution to form clusters of entangled polyelectrolyte mesh held together by ionic physical cross-links. The PCPDTBT-SO$_3$K hydrogel disorders upon heating due to the dissolution of ionic aggregates and the resulting electrostatic dispersion of clusters from forming a percolated polymer network. Finally, PCPDTBT-SO$_3$K hydrogels are mixed conductors capable of long-range, bulk electron and ion transport required for inclusion in energy conversion devices.

**Solidification and Morphology.** Increasing the concentration of PCPDTBT-SO$_3$K in aqueous solution past a critical threshold results in a well-formed hierarchical microstructure at low polymer fractions. We restrict our study to PCPDTBT-SO$_3$K (M$_w$= 11 000 g mol$^{-1}$, absolute weight-average molecular weight from small-angle neutron scattering (SANS) zero-angle scattering, Figure 3 and Table S2; D = 1.1 by SEC according to PS standards) hydrogels formed from semidilute solution at a concentration of 4.0% w/v in H$_2$O. The microstructure of the material, as illustrated by freeze–fracture scanning electron microscopy (FF-SEM) images in Figures 2a and b, is based on microgel units, where the polymer is assembled into spherical domains of 3–5 µm, which then interconnect to form a bulk three-dimensional network with mesh sizes of 10–20 µm. Noteworthy, the void size is ideal for the diffusion of nutrients and movement of bacteria: a requirement for the development of CPEs as interfacial modifying agents in bioelectrochemical cells.

The spanning of microgel aggregates appears to be a jamming of the particles, giving rise to the macroscopic mechanical properties characteristic of soft materials (e.g., hydrogels). The PCPDTBT-SO$_3$K remains a gel-like elastic solid at 25 °C (i.e., G’ > G’’ over the accessible frequency range), in the sense that it can sustain static loads and store elastic energy over long times. It exhibits an elastic modulus of order 10$^3$ to 10$^5$ Pa and a terminal relaxation time of 100s of seconds which, within the experimentally accessible times, are remarkably similar to traditional cross-linked polymer gels albeit with differences in microstructure. Although physically cross-linked polymer gels and the hydrogel soft material presented here are three-dimensional networks, the former is comprised of polymer chains joined together at a number of covalent connection sites, whereas the latter results from microgel clusters that percolate to form a macroscopic network. Noteworthy, the dynamical frequency sweeps, Figure 2c, are qualitatively consistent with the jamming of microgels, as indicated in the microscopy images and recent work on the jamming transitions of soft polyelectrolyte microgel suspensions.31

The existence of spherical microgels jamming to form a three-dimensional network is incompatible with the solution assembly of conjugated polymer and polyelectrolyte gels highlighted earlier.32 As demonstrated with a combination of electron microscopy and rheology (Figure 2), the polymers are not associating into large crystalline aggregates or wormlike micelles. Thus, it is necessary to further interrogate the role of
aromatic and mesogen driven interactions in the microstructure of CPE semidilute solutions and hydrogels.

**Chain Rigidity Effects.** To elucidate chain rigidity effects on the structure of semidilute solutions of CPEs, it was crucial to understand the chain conformation of PCPDTBT-SO₃K. By operating in the dilute limit where polymer chains can be considered noninteracting, SANS can be used to extract information regarding correlations along a single polymer chain (i.e., form factor).

Conjugated polymers typically adopt configurational states that more closely resemble rigid rods than Gaussian coils. The Wormlike Chain model (WLC), or Kratky–Porod model, provides a useful form factor in describing these semiflexible systems.²²,²³ The WLC model details a chain of contour length, \( L_c \), as a sequence of locally stiff segments of persistence length, \( l_p \). Persistence length is the length over which the chain maintains correlation to itself or, restated, the length over which the flexible cylinder can be considered a rigid rod. A WLC description models backbone stiffness in terms of the Kuhn length, \( b = 2l_p \), as each Kuhn segment can be simulated as freely jointed with each other.

It is useful to compare the expected semiflexible behavior against both rigid rod and Gaussian coil descriptions as a confirmation of the chain behavior and characteristic lengths, highlighting good agreement of the SANS scattering data (Figure 3) with the WLC model across a wide \( q \)-range. PCPDTBT-SO₃K is a semiflexible polymer having a Kuhn length of \((13.4 \pm 0.9)\) nm and a cylinder radius of \((1.4 \pm 0.1)\) nm, as determined from a WLC fit²⁴,²⁵ to the SANS scattering profile for PCPDTBT-SO₃K (dilute solution, 0.3% w/v, D₂O) presented in Figure 3.

PCPDTBT-SO₃K is thus significantly more rod-like than classical polymers.²⁶ It is also several times stiffer than poly(3-alkylthiophenes), P3ATs, due to the absence of syn–anti isomerism between thiophene units.²⁷ Although chain stiffness...
has historically been associated with the fused aromatic ring structure of the backbone and the electrostats of the pendant ionic groups, PCPDTBT-SO₃K has a persistence length similar to neutral conjugated polymers such as poly(2-methoxy-5-(2'-ethylhexyloxy)-p-phenylenevinylene), MEH-PPV, and polyfluorene, PF, even with the high density of charges along the polymer. Thus, electrostatic repulsions appear unimportant in dictating the stiffness of a chain as long as the backbone is conjugated. These comparisons allow coarse-graining of PCPDTBT-SO₃K as a semiflexible polymer, likely due to the finite backbone torsion and nonzero bond angles between donor (cyclopentadithiophene, CPDT) and acceptor (benzo-thiadiazole, BT) monomer units.

The semiflexible conformation, rather than rigid rod, is suggestive of the unimportance of chain rigidity effects compared to electrostatics when determining the structure and length scales of interactions of the soft solid. This is further illustrated by calculating critical concentrations in the Gaussian coil and rigid rod limits, in the absence of additional interactions. In the Gaussian coil limit, the size of the Gaussian coil and rigid rod limits, in the absence of additional interactions. The size of the Gaussian coil used is determined by the SANS data, , where is Avogadro’s number, and the radius of gyration used was determined by the SANS fit to the Debye model (Figure 3 and Table S1).

Conversely, in the rigid rod model, the overlap concentration depends on the volume accessible to a rotating cylinder, where cylinder length , was determined by the SANS fit to the rigid rod model (Figure 3, Table S3). In this rigid rod limit, it is also useful to calculate the critical concentration for the isotropic to nematic transition, where is the diameter of the rod determined from the SANS fit to the rigid rod model (Table S3). Considering the chain length and stiffness for the PCPDTBT-SO₃K chains studied yield remarkably similar overlap concentrations in both the Gaussian and rigid rod limits and, comparable to the polymer concentration investigated, the chain interactions of a solution of PCPDTBT-SO₃K (4.0% w/v, H₂O) are anticipated to be characteristic of the semidilute regime. However, the hydrogels are sufficiently below the predicted isotropic to nematic transition, and thus presumably the mesogen interactions are of little structural or thermodynamic consequence.

Aromatic Interactions. Upon examining the local interactions of the PCPDTBT-SO₃K chains in their hydrogel environment via wide-angle X-ray scattering (WAXS), we observe the striking lack of diffraction peaks or stacking (Figure 4). Any local interchain contacts, including π-orbital overlap, must be weak or relatively disordered. Further, the absence of π-stacking or regularly ordered, close packed interchain correlations suggest the CPEs are not behaving according to the proposed micellar model as amphiphilic polymers that associate hydrophobic conjugated backbones to arrange hydrophilic ion side-chains out toward the bulk solvent. The free energy penalty associated with the loss of entropy from water ordering in the solvation shell around the polymer (i.e., hydrophobic effect) appears less than the electrostatic energy penalty that would be required to associate the ionic groups so closely together in an ordered micellar arrangement. However, while not the dominant driving force for network formation, unstructured aromatic interactions likely contribute to the overall structure and conductivity pathways.

Electrostatic Interactions. Instead of π-π stacking and chain stiffness effects dominating the structural ordering, electrostatics appear to be the primary driving force for solution assembly and solidification. The local correlations evident upon heating PCPDTBT-SO₃K hydrogels (4.0% w/v, H₂O) (Figure 4) are associated with electrostatic assembly, particularly given the relatively broad peaks that are classically a signature of ionic interactions. The broad peak centered at corresponds to the presence of approximately 1.6 nm ionic clusters. Polarizability contrast between the conjugated polymer backbone, pendant ionic charges, counterions, and bulk aqueous solvent result in a nonuniform dielectric contrast, causing the ions to aggregate in semidilute solution. The aggregation of multiple ions serves to physically cross-link the chains together in entangled polyelectrolyte mesh clusters. With increasing temperature, the ionic aggregates dissociate, effectively un-cross-linking the polymer chains and disordering the microstructure. This greater solubilization of polymer chains causes a concomitant rise in intensity of a broad amorphous solvation peak. The dissociation of ionic cross-links and increase in polymer solvation occur over a narrow temperature range, 40 to 45 °C, agreeing well with additional scattering and conductivity measurements (Figures 7 and 8).

These local interactions complement the interpretation of larger length scale correlations. Figure 5 presents SANS profiles for 4.0% w/v PCPDTBT-SO₃K in D₂O at two temperatures, 25
and 75 °C, where elastic and viscous behavior are observed, respectively. To extract correlation lengths and other relevant structural parameters, the data are fit to a scattering function. A semiempirical scattering function, Broad Peak or Polyelectrolyte model, is used to describe systems that exhibit electrostatically driven assembly, such as semidilute polyelectrolyte solutions with low levels of added salt.44,45 While PCPDTBT-SO3Khydrogels (4.0% w/v, D2O) at 25 °C (blue ●, −, −) and 75 °C (red ●, −, −). Outside of high-q data points, error bars are smaller than markers.

Figure 5. SANS data (●), fits to our conjugated polyelectrolyte model (−), and fits to the polyelectrolyte model (−) for PCPDTBT-SO3K hydrgels (4.0% w/v, D2O) at 25 °C (blue ●, −, −) and 75 °C (red ●, −, −). Outside of high-q data points, error bars are smaller than markers.

While the upick at low-q, corresponding to the onset of power law behavior, is often interpreted as a form of aggregation, it is a standard feature of ionic polymer solutions at low salt concentrations that reveals the length scale correlations of the three-dimensional microstructure. This has been demonstrated with the Polyelectrolyte model for semidilute low-salt polyelectrolyte solutions exhibiting ordered regions of high polymer density coexisting with disordered regions of low polymer content4,47 and is believed to be related to the attractive electrostatic correlations, independent of backbone hydrophobicity or solvation.46

The clusters are compact, collapsed structures, characterized by a Flory-like exponent of 0.39 ± 0.003 that percolates through the disordered low polymer concentration solution forming a network. The microgels are composed of an entangled polyelectrolyte mesh with correlations revealed by the peak present in the mid-q region of the scattering pattern. In the semidilute regime, these polymer chains are interacting, thus yielding interchain correlations but are not dominated by entanglements as in the melt or concentrated solution. This interchain correlation peak at q* is believed to be the well-known polyelectrolyte peak, although there is no direct evidence presented here. The polyelectrolyte peak is caused by local chain rigidity derived from like-charge repulsion of highly charged polymer chains and is seen in various polyelectrolyte systems through light, X-ray, and neutron scattering.4,47,53–55 For PCPDTBT-SO3K at 25 °C, there exists a statistical correlation length, ℘chian, of (13.7 ± 0.2) nm separating the polymer chains and an electrostatic screening length, ξchain, of (4.9 ± 0.4) nm around which each chain can be felt.

The correlation length, ℘chain, can be compared to the scaling relationship4,56 for the correlation length of a polyelectrolyte incorporating electrostatic and entropic chain conformation effects:

\[ \xi \approx f^{-2/\nu} \left( \frac{L_{\text{chain}}}{b} \right)^{-1/7} (bC)^{-1/2} \]

where n is the low-q scaling exponent, m is the polyelectrolyte scaling exponent, p is the ionic aggregate scaling exponent, \( L_{\text{chain}} \) is the electrostatic screening length of the interchain correlation, \( L_{\text{ion}} \) is the electrostatic screening length of the ionic aggregate correlations, q* is the interchain correlation peak position, qion is the ionic aggregate correlation peak position, and \( I_{\text{inc}} \) is the background incoherent scattering. Real space correlation distances are extracted from these peak positions by \( \xi = 2\pi/q \). A, B, and C are scaling factors, dependent on the concentration and scattering length density difference associated with each feature. The exponents n, m, and p capture the breadth of the power law and Lorentzian-like functions, where m is inversely related to the Flory exponent (\( \nu \)) for polymer systems, i.e., m = 1/\( \nu_{\text{chain}} \), where \( \nu = 0.5 \) corresponds to the θ condition. The Flory exponent corresponds to collective behavior, highlighting the conformation due to the relative importance of polymer–polymer or polymer–solvent interactions. Thus, m need not be constrained to the \( \nu = 1.3 \) result from PCPDTBT-SO3K in dilute solution (0.3% w/v, D2O), which only corresponded to correlations along a single chain and highlighted the persistence of the polymer. Similarly, the inverse of n can be used as a Flory-like exponent (\( \nu_{\text{microgel}} \)) to envision the conformation of the microgels.

Parameters extracted by fitting the Conjugated Polyelectrolyte model, eq 4, to the scattering functions of PCPDTBT-SO3K (4.0% w/v, D2O) at both temperatures are included in the Supplementary Information. Calculated structural parameters are summarized in Table 1.

Table 1. Structural Parameters of PCPDTBT-SO3K (4.0% w/v) at 25 and 75 °C

<table>
<thead>
<tr>
<th>Parameter</th>
<th>25 °C</th>
<th>75 °C</th>
</tr>
</thead>
<tbody>
<tr>
<td>( \nu_{\text{microgel}} )</td>
<td>0.39 ± 0.003</td>
<td>0.39 ± 0.003</td>
</tr>
<tr>
<td>( \nu )</td>
<td>0.48 ± 0.04</td>
<td>0.45 ± 0.05</td>
</tr>
<tr>
<td>( \xi_{\text{chain}} ) (nm)</td>
<td>13.7 ± 0.2</td>
<td>12.8 ± 0.3</td>
</tr>
<tr>
<td>( \xi_{\text{ion}} ) (nm)</td>
<td>4.9 ± 0.4</td>
<td>4.3 ± 0.3</td>
</tr>
<tr>
<td>( L_{\text{chain}} ) (nm)</td>
<td>6.6 ± 1.3</td>
<td>6.4 ± 1.2</td>
</tr>
<tr>
<td>( L_{\text{ion}} ) (nm)</td>
<td>2.0 ± 0.8</td>
<td>1.7 ± 1.2</td>
</tr>
</tbody>
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where $f$ is the fraction of charged monomers, $l_B$ is the Bjerrum length, the length at which electrostatic interaction energy is equal to the thermal energy $k_B T$, $b$ is the monomer size, and $C$ is the polymer concentration. Taking into account counterion condensation, the polyelectrolyte chain and condensed counterions are equivalent to a polyelectrolyte with charges $l_B$ apart. Because the fraction of charged monomers is experimentally controlled as $f = b/l_B$, the resulting correlation length is given by

$$\xi \approx \left( \frac{l_B}{b} \right)^{1/7} (bC)^{-1/2}$$

which agrees well with the experimentally determined mesh size and corroborates the presented description of the CPE hydrogel as a polyelectrolyte in semidilute solution. The larger value of $L_{chain}$ compared to other polyelectrolytes such as PSS is likely due to both the finite thickness of the PCPDTBT-SO_3K chain and the increased backbone stiffness from the fused ring conjugated backbone.

These scaling relationships (eqs 5 and 6) not only give qualitative agreement with the hierarchical structure of the PCPDTBT-SO_3K hydrogels presented here but also provide insights into the molecular design of CPEs. Compared to traditional polyelectrolytes, CPEs allow these parameters ($f$, $b$, and $l_B$) to span a greater breadth of values. Owing to the combination of highly polarizable backbone, pendant ionic groups, and nonpolar alkyl side chain linkers, CPEs are soluble in a range of solvent conditions, typically with larger Bjerrum lengths, allowing larger correlation lengths. Additionally, the monomer size in CPEs is typically also much larger (~1.2 nm for PCPDTBT backbones) and can be tuned easily through the addition of different donor or acceptor moieties. Control of these two parameters should allow design of CPEs with precisely tunable polyelectrolyte mesh spacing. Further, due to the interplay of these two parameters, CPEs significantly delay the onset of counterion condensation as the Bjerrum length is increased, enabling dissociated ions and electrostatic behavior in low dielectric solvents.

Ion–ion correlations or ionic aggregates present in the entangled polyelectrolyte mesh appear at progressively shorter length scales. This weak, secondary correlation appears as a shoulder of the more pronounced polyelectrolyte correlation in the scattering function of Figure 5. The ionic aggregates (with scattering contrast including both the pendant ionic sulfonate groups and potassium counterions) are correlated with a characteristic spacing, $\xi_{ion}$ (6.6 ± 1.3) nm of between aggregates, which is on order of the chain length rather than the monomer scale. As such, it is believed these ionic aggregates help to tie separate chains together as dynamic physical cross-links. The screening length, $L_{ion}$ of (2.0 ± 0.8) nm is interpreted as a characteristic length scale for electrostatic screening between ions and is related to the dielectric constant of the medium, the concentration of ionic species, and the number of ions associating in each ionic aggregate.

As the temperature is increased from 25 to 75 °C, the overall hierarchical structure (Figure 6) remains intact, but there is a compaction of both the entangled polyelectrolyte mesh and ionic correlations (Table 1). The percolated three-dimensional network of clustered microgels shows only minor changes, a surprising result as the macroscopic mechanical properties shift from elastic to viscous response. The Flory-like exponents characterizing the cluster morphology and interchain correlations remain effectively constant, demonstrating no detectable change in conformation. The correlation and screening lengths show compaction for the interchain correlations: $\xi_{chain}$ falls from (13.7 ± 0.2) to (12.8 ± 0.3) nm and $L_{chain}$ decreases from (4.9 ± 0.4) to (4.3 ± 0.3) nm. While the ion correlations follow similar behavior, it is hard to make quantitative statements as relatively weak scattering results in high uncertainty of the fits.

X-ray scattering studies aid in investigating the ionic interactions, as improved signal intensity results from an increased electron density contrast between the ionic moieties relative to the polymer and solvent, and better temperature resolution is experimentally accessible in understanding the structural transitions. Figure 7 summarizes the small-angle X-ray scattering (SAXS) profiles for PCPDTBT-SO_3K (4.0% w/v, H_2O) from 25 to 75 °C.

Strikingly, the disordering of the microgel morphology begins at the reorganization temperature determined by WAXS, at which the dissociation of ionic cross-links and increase in polymer solvation occurs: 45–50 °C. Nonetheless, the microgels continue to lose structure with further heating. The polyelectrolyte peak, as probed by SAXS, is more diffuse and at lower-q than from SANS, which we attribute to differences in scattering length density probing slightly different correlations. While it appears that the polyelectrolyte peak shifts to higher q, indicating the collapse of the entangled polyelectrolyte mesh network, the predominant effect is the collapse in intensity of the peak, due to a loss of correlations. This apparently reflects increased solvation of the CPE chains and associated counterions, which are then able to access more conformations, resulting in a smearing of the network structure. Further, the thermoresponsive character is reversible, highlighting an equilibrated system; the structure and properties (i.e., rheology, conductivity) can be recovered given sufficient equilibration at any temperature.
The ions' charge transport is dominated by electronic conductivity, with temperature increased (40°C) to 45°C, ionic cross-links and release of unbound counterions. At sharply increases as the ions gain mobility due to dissociation of delocalized aromatic interactions and the extremely high (96%) π-orbital overlap where there are interchain connections so that holes can hop between adjacent chains.

In designing new mixed conductor hydrogels based on CPEs, there are several important principles that can be elucidated or anticipated from this investigation. Increasing the polymer concentration should lead to higher conductivities due to increased contact of the electrodes with the electroactive microgel cluster and increased percolation of the microgel clusters. Manipulating the solvent dielectric properties could allow for conduction outside the clusters and affect the formation of ionic aggregates. Strategies to incorporate π−π stacking through processing, materials design, or solvent control could have important effects on transport.

**CONCLUSIONS**

PCPDTBT-SO3K, a high-mobility CPE, undergoes solution assembly and solidification driven primarily by electrostatics, rather than π−π stacking, forming a three-dimensional network with a novel, hierarchical structure (Figure 6). A percolated network exists at large distances, composed of microgel clusters that are aggregated at ambient temperatures but become electrostatically stabilized at higher temperatures as ions gain mobility out of the ionic aggregates toward the surface of the microgel. The interiors of these microgel clusters are entangled polyelectrolyte meshes that behave predominantly like classical polyelectrolytes in the semidilute regime without added salt but additionally incorporate unstructured aromatic interactions and π−π stacking, cross-links. The organization of CPEs in this regime is free from π−π stacking, with implications for CPE hydrogels and solution assemblies relying predominantly on intrachain charge transfer. Future work should be aimed at elucidating mechanisms for control of ionic aggregates, percolation of the microgel network, and π−π stacking, especially in added salt regimes.

**EXPERIMENTAL SECTION**

**Materials.** Milli-Q ultra pure water (18.2 MΩ cm at 25°C) was used for the preparation of all aqueous samples and in all synthetic procedures. All chemicals, unless otherwise noted, were commercially available and used as received from Sigma-Aldrich. Deuterated solvents were obtained from Cambridge Isotope Laboratories.

PCPDTBT-SO3K was synthesized according to literature procedures. The molecular weight and dispersity of PCPDTBT-SO3K were determined by gel permeation chromatography in DMF of its derivative, resulting from exhaustive ion exchange with excess tetrabutylammonium bromide, followed by dialysis.

**Molecular Characterization.** 1H NMR spectra were recorded at either 500 MHz (Bruker AVANCE500) or 600 MHz (Varian VNMRS), and 13C NMR spectra were recorded at either 126 MHz (Bruker AVANCE500) or 150 MHz (Varian VNMRS) in D2O or DMSO-d6. Chemical shifts are reported in ppm referenced to residual solvent peaks. Gel permeation chromatography (GPC) was conducted on a Waters instrument using a refractive index detector and Viscotek Visco Gel I-Series columns (IMBHMW-3078, IMBLMW-3078), DMF with 0.1% LiBr at 25°C was used as the mobile phase with a flow rate of 1 mL/min. Reported molecular weight and dispersity (Đ) values are relative to polystyrene (PS) equivalents as calibrated with narrow PS Standards (Agilent). Mass analysis was performed using electrospray
ionization methods (ESI) using a Waters Micromass QTOF mass spectrometer. Elemental analysis using the Dumas method was performed on a Control Equipment Corp. instrument (CEC 440HA).

**Electron Microscopy.** FF-SEM experiments were conducted on a FEI XL30 microscope operating at an acceleration voltage of 10.0 kV. Freeze-fractured samples were prepared with a Quorum Technologies Polaron PP2000T cryo-stage. The stage was cooled to the temperature of liquid nitrogen. Then, a small volume of solution was added and rapidly cooled by liquid nitrogen, allowing vitrified amorphous water, avoiding ice crystalization and preserving the native structure of the solution. The samples were put under vacuum at continuous cooling with liquid nitrogen and fractured with a microtome knife. The surface was replicated by coating with Pt at an angle of 90°.

**Rheology.** Rheological measurements were taken on an AR-G2 rheometer using 20 mm diameter stainless steel parallel plates with a 200 µm gap. Sample environment was controlled by using a Peltier-controlled lower plate for temperature control and covering the fixture to prevent ambient humidity from affecting the measurements. Dynamic strain sweeps were performed at 1 and 10 rad s⁻¹ to determine the linear viscoelastic regime, where the upper limit of the linear viscoelastic regime is defined as the critical strain (γc), below which the dynamic modulus remains invariant. The γc for the PCPDTBT-SO₃K hydrogel (4.0% w/v, H₂O) at 25 °C was approximately 5% (Figure S2). All further measurements were taken at 1% strain to remain within the linear viscoelastic regime.

**X-ray Scattering.** X-ray scattering experiments were performed on a diffractometer at the national neutron source (NIST) in Oak Ridge, TN. Neutron scattering experiments were performed at LANL in Los Alamos, NM. Ultrasonic measurements were taken on a Control Equipment Corp. instrument (CEC 440HA). The authors declare no competing financial interest.

**Electrochemical Impedance Spectroscopy.** Impedance measurements were taken on a Biologic VSP-300 potentiostat using an in-plane configuration. Sample was deposited onto a glass substrate with evaporated gold contacts. The electroactive area of the sample was controlled through a layer of Kapton tape, with a well of defined area. The lateral dimensions were measured with a digital micrometer, and the thickness of the gel sample was calculated from the volume deposited. The impedance measurements were performed from 7 MHz to 1 Hz with a sinusoidal amplitude of 100 mV, using an INSTEAD stage (±0.1 °C accuracy) to control the temperature of the sample. The conductivity was monitored until stabilized at each temperature. The presented data are from a single cycle of monotonic heating from 25 to 75 °C but are reproducible upon significant equilibration (up to 24 h from 75 °C back to 25 °C).

Given the interactions between electronic and ionic charge carriers, especially in an inhomogeneous dielectric continuum (i.e., the hierarchical three-dimensional network structure, composing PCPDTBT-SO₃K chains, counterions, and solvent both inside the microgels and in the void spaces), it is not currently possible to fit the data to an appropriate equivalent circuit model. To obtain estimates of the ionic and electronic conductivities, the frequency independent plateaus (Figure S6) of impedance were converted to conductivity according to

\[
\sigma = \frac{l}{IZ^2tw}
\]

where \(IZ^2\) is the total impedance, \(l\) is the thickness of the sample, \(w\) is the lateral sample width, and \(I\) is the distance between electrodes and taken as the DC conductivities. The total impedance of the high frequency plateau is taken as the electronic resistance and is used to calculate the electronic conductivity. The impedance of the low frequency plateau is taken as the sum of ionic and electronic resistances. Subtracting the electronic contribution to this impedance, the ionic conductivity can be calculated according to eq 7.

Noteworthy, this approach likely significantly underestimated the conductivities by neglecting contact resistance between the sample and the electrodes, which would be present in both plateaus in the Bode plots (Figure S6).

## ASSOCIATED CONTENT

### Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.chemmater.7b05303.

Experimental details, characterization data, and SANS fit parameters and theoretical background (PDF)

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### Notes

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References


