

Boron Nitride-Doped Polyphenylenic Organogels

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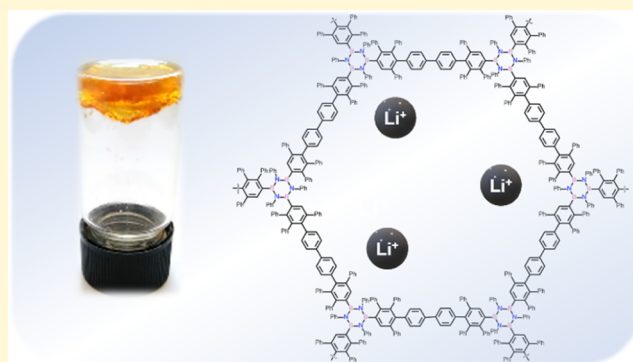


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Supporting Information

ABSTRACT: Herein, we describe the synthesis of the first boron nitride-doped polyphenylenic material obtained through a [4 + 2] cycloaddition reaction between a triethynyl borazine unit and a biscyclopentadienone derivative, which undergoes organogel formation in chlorinated solvents (the critical jellification concentration is 4% w/w in CHCl_3). The polymer has been characterized extensively by Fourier-transform infrared spectroscopy, solid-state ^{13}C NMR, solid-state ^{11}B NMR, and by comparison with the isolated monomeric unit. Furthermore, the polymer gels formed in chlorinated solvents have been thoroughly characterized and studied, showing rheological properties comparable to those of polyacrylamide gels with a low crosslinker percentage. Given the thermal and chemical stability, the material was studied as a potential support for solid-state electrolytes, showing properties comparable to those of polyethylene glycol-based electrolytes, thus presenting great potential for the application of this new class of material in lithium-ion batteries.



INTRODUCTION

In the last decades, polyphenylene-based materials have been the subject of extensive research efforts.^{1,2} This interest is related to the many different potential applications of these materials, from precursors in graphene synthesis^{3,4} to antennae systems,^{5–7} proton exchange membranes,^{8,9} optoelectronic devices,¹⁰ and self-assembled systems.¹¹ Structural modifications of the polyphenylene-based frameworks by heteroatom-doping^{12,13} or peripheral addition of functional groups^{14–16} are currently the main strategies to enrich the chemical and physical properties of polyphenylenic materials. Polymeric phenylene-based materials are principally prepared by exploiting [4 + 2] cycloaddition reactions with CO extrusion.^{17,18} Particularly, the pioneering work of Müllen and co-workers showed how a large variety of controlled structures, tailored for a given application, can be prepared through an appropriate combination of specifically designed dienes and alkyne precursors.¹⁹

We contributed to the field by preparing discrete borazino-doped oligophenylenes following the [4 + 2] cycloaddition with CO extrusion synthetic route using opportunely functionalized ethynyl- and tetraphenylcyclopentadienone-bearing borazine derivatives (Figure 1b).²⁰ The replacement of a phenyl ring with a borazine ring proved to be particularly effective in breaking the π -conjugation, causing a blue shift in the emission envelopes compared to those of the all-carbon congeners.^{20,21} Borazino-doped oligophenylenes can also be prepared by exploiting a condensation reaction approach, either condensing previously formed chloroborazoles²² via a

silicon/boron exchange (Figure 1a)²³ or by using a mixture of *para*-phenylenediamine and aniline with BCl_3 , followed by the addition of an aryl lithium derivative (Figure 1b).²¹ When only arylendiamine derivatives are used, boron nitride (BN)-doped polymeric materials can be obtained in which chloro-borazole units are formed (Figure 1a).^{24,25}

In this case, the moisture-sensitive nature of the chloro-borazole units limits the manipulation of these materials under normal ambient conditions and consequently limits the versatility of their applications. The presence of BN (or BO) bonds proved to increase the affinity of the materials toward polar or charged species,^{24,26} which is particularly important when engineering materials for gas adsorption/storage applications^{24,25} and solid-state electrolytes (SSEs).^{27,28} Moreover, the insertion of the thermally stable BN bonds has frequently led to the use of BN polymers as pre-ceramic/hybrid materials.^{29–31} With the objective of preparing porous functional borazine-doped materials, this paper tackles the challenge of engineering borazino-bearing polyphenylene-based polymeric structures. When devising the preparation and structure of these materials, we must consider the susceptibility of the BN core toward moisture.^{32,33} Up to

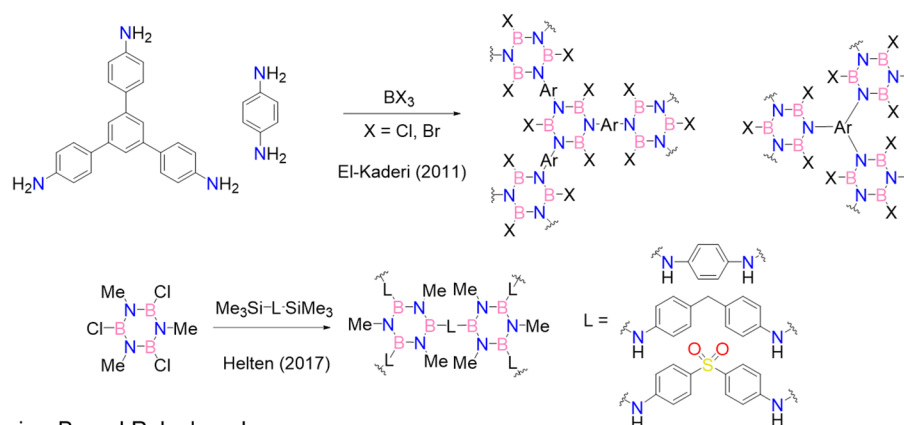
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a) Borazine-Based Polymers



b) Borazine-Based Polyphenylenes

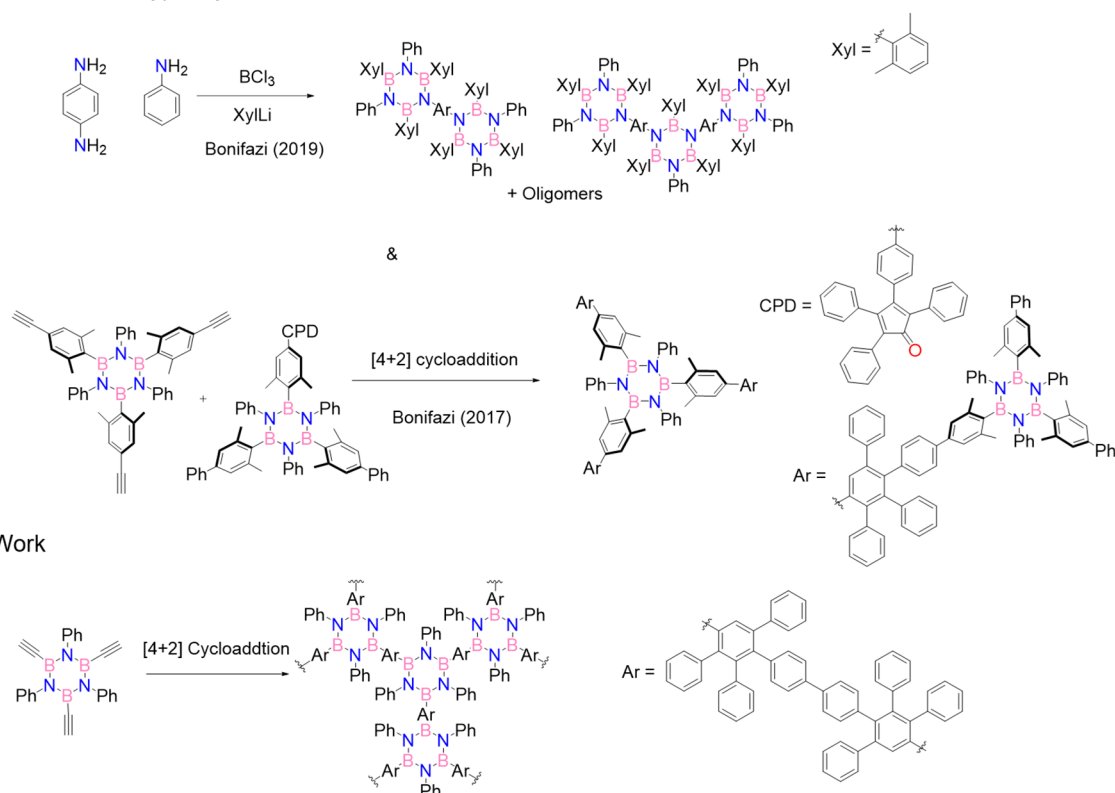


Figure 1. From molecular to macromolecular borazino-phenylenes.

now, we have circumvented this problem by preparing borazine precursors bearing 2,6-disubstituted aryl moieties at the B-position, with the 2,6-substituents sitting atop the boron atoms, shielding them from possible hydrolytic reaction (Figure 1b).^{32,33} On the one hand, although this approach has been effective in providing robust borazine linkers for preparing porous metal–organic frameworks,³⁴ the preparation of sterically protected borazine precursors is synthetically demanding and could be a limiting factor in expanding the chemical space of these BN precursors. On the other hand, the preparation of borazino-doped polyphenylenic materials by exploiting borazine formation as the polymerization reaction would require the use of strongly reactive BCl_3 and BBr_3 boron sources.^{24,25} Moreover, it would be very difficult to functionalize the B-atoms with protecting aryl substituents in a later step due to the generally low solubility of the final materials

and the high steric hindrance surrounding the B-atoms.²¹ Based on these considerations, we concluded that a desirable synthetic approach would involve a borazine precursor that, through a high-yielding reaction, could polymerize and simultaneously place sterically shielding moieties atop the boron centers. This line of thought led us to conjecture that the *B,B',B''*-triethynyl-*N,N',N''* triphenyl borazine scaffold,³⁵ known to be relatively stable under normal ambient conditions,³⁶ could serve as a suitable precursor to undergo polymerization through [4 + 2] cycloaddition with CO extrusion when reacted with a suitable biscyclopentadienone (Figure 1c). If a tetraphenylcyclopentadienone scaffold is used, formation of the tetra-substituted B-aryl moiety would place a phenyl ring atop the boron centers, making the borazine ring inert toward hydrolysis reaction. Embracing this synthetic strategy, we prepared a *B,B',B''*-triethynyl-*N,N',N''* triphenyl

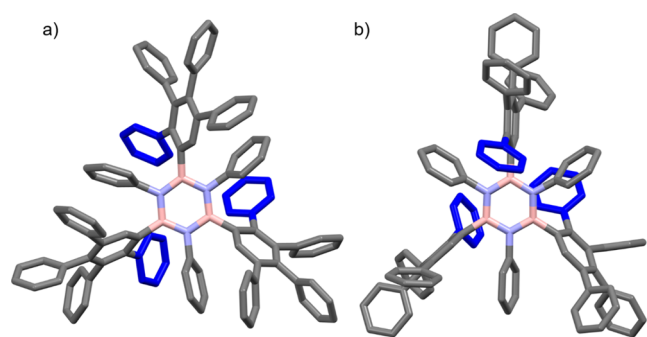


Figure 2. Molecular structures of cycloadduct isomers *cc*-3 (a) and *ct*-3 (b) in their crystal structures determined from single-crystal XRD, with space groups $P6_3$ and $P1$, respectively.

Supporting Information, a powder XRD study of the bulk sample of cycloadduct 3 prepared directly from the chemical reactions shown in Scheme 1, followed by crystallization from CH_2Br_2 , indicates that the only detectable crystalline phase is the $P1$ phase of the CH_2Br_2 solvate of 3, which contains the *ct* isomer. We note that other crystalline phases (e.g., containing the *cc* isomer) may also be present but are below the detection limit, which is estimated to be several percent given the relatively poor signal-to-noise level of the powder XRD data. This study confirms that the bulk sample of cycloadduct 3 prepared directly from the chemical reactions shown in Scheme 1 contains predominantly the *ct* isomer, fully consistent with the evidence from the multidimensional NMR analysis discussed above.

To exploit the cycloaddition reaction to prepare borazine-doped polyphenylenic polymers, we have designed a suitable dimeric diene unit, terminating with two tetraphenylcyclopentadienone moieties (molecule 6, Scheme 1). The first step of the synthesis involved a double Sonogashira-type cross-coupling reaction between phenylacetylene and 4,4'-diiodobiphenyl to yield bis-alkyne derivative 4 in quantitative yield.

Oxidation of molecule 4 in the presence of KMnO_4 gave tetracarbonyl derivative 5 in 97% yield, which was in turn reacted with 1,3-diphenylacetone to produce the desired dimeric cyclopentadienone 6 in good yield (52%). The structure of biscyclopentadienone 6 was confirmed via NMR spectroscopy and high resolution mass spectroscopy (HRMS) (Figures S27–S29).

The cycloaddition reaction between borazine 2 and biscyclopentadienone 6 in dry and degassed Ph_2O at 220 °C yielded a red gel. After precipitation and purification of the solid, the resulting reddish material was reacted with phenylacetylene in Ph_2O at 230 °C to end-cap the unreacted cyclopentadienone units. Purification of the resulting pale-yellow solid with various sonication–centrifugation cycles using different solvents (petroleum ether/ CH_2Cl_2 , acetone, MeOH, and Et_2O) gave the desired **BN-polymer** in 52% yield. Given the insolubility of the product in common organic solvents, characterization was carried out by thermogravimetric analysis (TGA), attenuated total reflectance infrared spectroscopy (ATR–IR), and solid-state NMR techniques. TGA measurements showed that the solid is thermally stable up to 400 °C (Figure S30), trapping ca. 10–20% of the solvent (see mass loss at around 100 °C). ATR–IR analysis of both cycloadduct 3 and **BN-polymer** (Figure S31) showed strong signals at ca. 1355 and 1321 cm^{-1} assigned to BN bond stretching. Notably, no signals related to $\text{C}=\text{O}$ bonds were detected, suggesting the full conversion of the end-capping reaction. High-resolution solid-state ^{13}C NMR spectra recorded using the $^1\text{H} \rightarrow ^{13}\text{C}$ CPMAS technique for precursor 2, reference cycloadduct 3, and the **BN-polymer** material are shown in Figure 3. The solid-state ^{13}C NMR spectrum of precursor 2 has isotropic peaks at 84 and 101 ppm due to the ethynyl moieties and isotropic peaks in the range 125–150 ppm due to aromatic environments. The fact that the ethynyl peaks are absent from the solid-state ^{13}C NMR spectra of the cycloadduct 3 and **BN-polymer** samples is consistent with our hypothesis of complete conversion of the cycloaddition

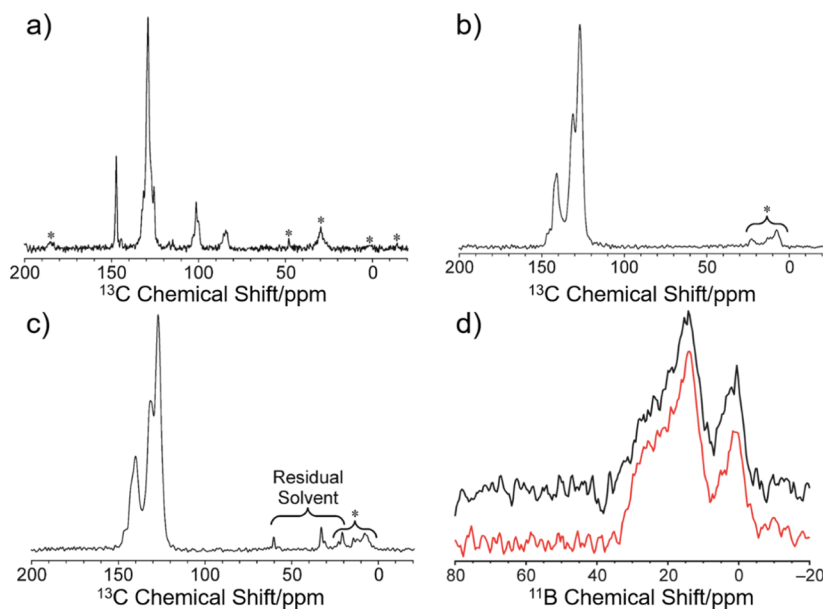


Figure 3. Solid-state $^1\text{H} \rightarrow ^{13}\text{C}$ CPMAS NMR spectra recorded for (a) borazine precursor 2, (b) reference cycloadduct 3, and (c) **BN-polymer**. Peaks marked with asterisks are spinning sidebands. (d) One-dimensional projections of the solid-state ^{11}B MQMAS NMR spectra onto the direct dimension for cycloadduct 3 (red) and **BN-polymer** (black).

reaction. The solid-state ^{13}C NMR spectra of the cycloadduct **3** and **BN-polymer** materials contain multiple overlapped peaks between 120 and 150 ppm, corresponding to the aromatic ^{13}C environments in these materials. The fact that the solid-state ^{13}C NMR spectra of cycloadduct **3** and **BN-polymer** are virtually identical in this region (see the overlay of the spectra in Figure S36) suggests that the local structure around the central aromatic ring (including the conformations of the substituents) is very similar in these materials. Low-intensity peaks between ca. -15 and 60 ppm are assigned to spinning sidebands and, in the case of **BN-polymer**, also to the residual solvent (assigned as Et_2O and MeOH), as annotated on the spectra in Figure 3b,c.

Solid-state ^{11}B MQMAS NMR spectra⁴¹ recorded for samples of cycloadduct **3** and the **BN-polymer** are shown in Figure S37, and one-dimensional projections of these spectra onto the direct dimension are shown in Figure 3d. The one-dimensional projections in Figure 3d were obtained from the ^{11}B MQMAS NMR spectra by summation of all data between 39 and 49 ppm along the indirect dimension (vertical in Figure S37; see details in the Supporting Information). In the one-dimensional projections in Figure 3d, the shapes of the peaks are very similar for cycloadduct **3** and the **BN-polymer**, strongly suggesting that cycloadduct **3** and the **BN-polymer** have similar values of the quadrupolar parameters and isotropic chemical shifts for the ^{11}B environments. Again, we may infer from this observation that the local structures in the vicinity of the central B_3N_3 rings (including the conformations of the substituents bonded to the B atoms) are very similar in cycloadduct **3** and the **BN-polymer**.

Gel Formation and Rheological Studies. As observed during the preparation of the polymer in Ph_2O , **BN-polymer** shows a tendency to form gels. By suspending **BN-polymer** in different solvents, it was observed that chlorinated solvents (such as CHCl_3 and CH_2Cl_2) give gels upon sonication with a critical jellification concentration of 4 wt % for CHCl_3 . Low-field ^1H NMR (LF-NMR) studies (Table 1) were then carried

concentration, an increasing number of relaxation times (T_{2i} values) are detected, reflecting a higher heterogeneity of the jellifying network and suggesting the presence of meshes of different sizes (with small T_{2i} times corresponding to small cavity size). From the rheological point of view, we analyzed the 4 wt % gel by short stress sweep (SSS) and frequency sweep (FS) measurements. From SSS measurements (Figure S39), the presence of an elastic modulus G' much greater than the viscous modulus G'' further confirms the formation of a jellified material.

FS measurements (Figure 4) along with Maxwell best fitting (solid lines) allowed an estimation of the shear modulus (G), the cross linking (ρ_x), and the mesh size (ξ_{RHEO}) of the gel, giving $G = 667 \pm 54$ Pa, $\rho_x = 2.7 \times 10^{-7} \pm 2.2 \times 10^{-8}$, and $\xi_{\text{RHEO}} = 23.0 \pm 0.6$ nm. The value of G is comparable to that of a polyacrylamide gel with a low crosslinking percentage,⁴² which is in line with the low cross-linking value obtained from the measurement. The mesh size is consistent with a mesoporous material; however, the isotherm of N_2 adsorption at 77 K, measured on the xerogel, indicates a macroporous/non-porous material (Figure S45) with a calculated apparent Brunauer–Emmett–Teller surface area of $62 \text{ m}^2/\text{g}$, which suggests that the porous structure collapses upon drying.

BN-Polymer as Support in SSEs. The ability to form gels, the thermal stability, and the presence of polar BN bonds (see, e.g., the case for boroxines^{27,28}) make this material a good candidate as the support component in Li^+ -containing SSEs. Thus, **BN-polymer**/TEGDME (tetraethylene glycol dimethyl ether)/ LiClO_4 -based SSEs were prepared following the process shown in Figure 5 and tested (the technical details are reported in the Supporting Information). **BN-polymer** was first suspended in CH_2Cl_2 (2 mL) under ultra-sonication at 35°C for 2 h followed by vigorous stirring at r.t. for 12 h. Then, the TEGDME solution in CH_2Cl_2 (0.5 mL) was added and the obtained mixtures stirred at r.t. for 12 h. Finally, the LiClO_4 solutions at different concentrations (15, 20, and 25 wt %) in THF (0.5 mL) were added to the **BN-polymer**/TEGDME suspensions and ultra-sonicated at 35°C for 2 h followed by vigorous stirring at r.t. for 48 h so that homogeneous suspensions were obtained. The SSEs were deposited by solution cast on stainless steel disks adapted for Swagelok cells and dried at r.t. for 18 h followed by drying under vacuum at 70°C for 2 h.

The ion transport behavior of the SSEs prepared in this way was investigated by electrochemical impedance spectroscopy at r.t., using TEGDME/15 wt % LiClO_4 liquid electrolyte as the reference (Figures 6a, S44 and Table 2). Solid electrolytes containing **BN-polymer** showed a decrease in ionic conductivity with increasing salt content, with the highest ionic conductivity of about $1.51 \times 10^{-5} \text{ S cm}^{-1}$ for the sample containing 15 wt % of LiClO_4 (Figure 6b and Table 2). The bulk electrolyte resistance value (R_b) increases with increasing amount of LiClO_4 incorporated into the **BN-polymer**/TEGDME matrix due to a decrease in the available coordination sites of TEGDME by ion pairing, which is consistent with the ether crown ion chelation mechanism.^{43–45} This could be due to the distance between dissociated ions becoming too small and thus enabling the recombination into neutral ion pairs. This in turn leads to the formation of large aggregates of ions that reduce the segmental motion of ethylene oxide sites in the network, thus hampering the polymer motion (which is essential for fast ion conduction).^{46–51}

Table 1. Values of T_2 from LF-NMR for Gels Formed with Various Concentrations of **BN-Polymer** in CHCl_3

BN-polymer wt %	T_{21} (ms)	A_1 (%)	T_{22} (ms)	A_2 (%)	T_{23} (ms)	A_3 (%)	T_{24} (ms)	A_4 (%)
0%	2531	100						
2%	2373	81	1350	19				
4%	1927	73	981	18	108	9		
8%	1537	51	635	20	133	11	23	17

out to study the jellification behavior at three different concentrations to obtain data within the sub-gelated (2%) to super-gelated (8%) systems. In this analysis, the relaxation time T_2 was considered since it depends not only on the magnetic field strength (B_0) and temperature but also on the presence of solid surfaces such as those typical of a polymeric gel. The relaxation process is faster with increasing concentration of the “solid” component, which is reflected in a smaller T_2 . In the case of non-homogeneous systems, where meshes of different sizes can exist, the average T_2 depends on the relaxation time arising from each different mesh size present in the gel. In general, the ^1H relaxation is faster with decreasing mesh size.

As expected, T_2 is observed to decrease as the polymer concentration increases (Table 1). When moving from 2 to 8%

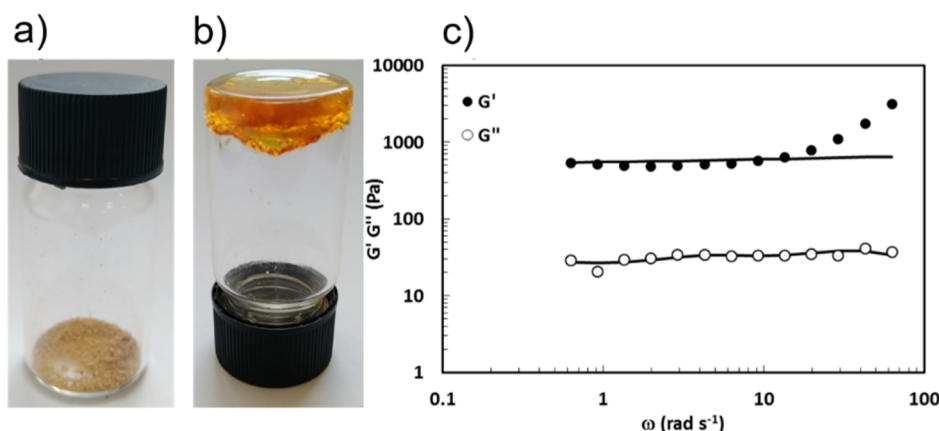


Figure 4. (a) Xerogel and (b) organogel of BN-polymer; (c) FS plot (circles) and Maxwell fitting (solid lines) for the 4 wt % gel in CHCl_3 .

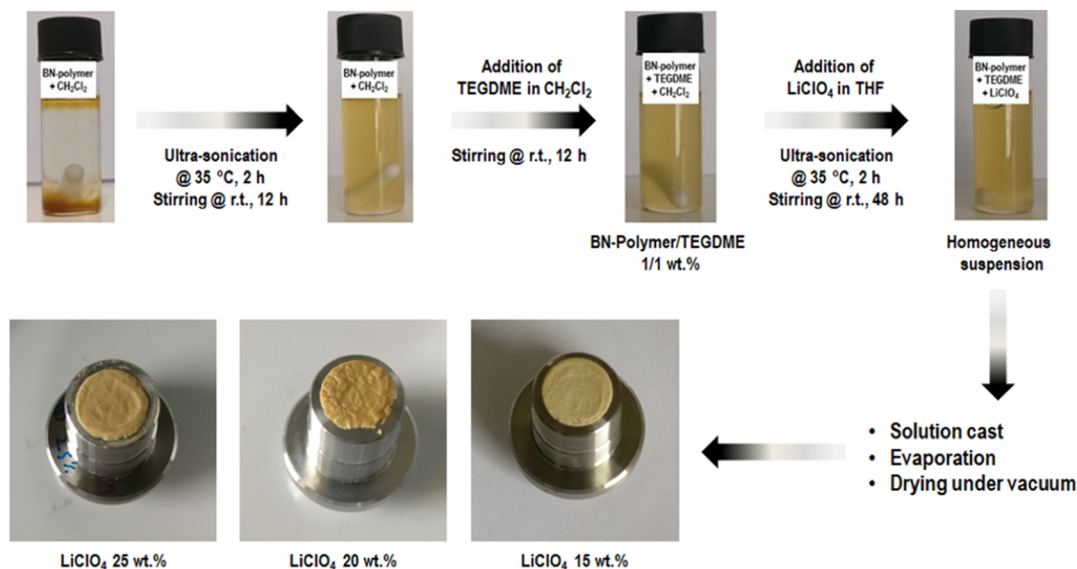


Figure 5. Process to prepare the BN-polymer/TEGDME/ LiClO_4 -based SSEs.

The estimated values of E_a are listed in Table 3.⁵² The lower the value of activation energy in the system, the higher the ionic conductivity at r.t. and vice versa.⁵³ At low temperatures, the BN-polymer-based electrolyte exhibits a lower ionic conductivity than that of the liquid TEGDME/15 wt. LiClO_4 system. However, it presents a similar trend compared to the liquid TEGDME/15 wt. LiClO_4 system at high temperatures. The lower activation energy for ion migration at high temperatures is probably due to the amorphous nature of the polymer, which promotes more free volume in the polymer electrolyte system upon increasing temperature.^{54,55} Furthermore, the wettability of BN-polymer pores by TEGDME increases at high temperatures, leading to a softer network, and TEGDME chains act as a transport medium.⁵⁶ In addition, the interaction of lithium salts with TEGDME and the presence of BN polar bonds help to weaken the ionic interaction in the salt and thus improve the charge carriers mobility.^{52,57} Since conductivity is influenced by temperature, σ was studied as a function of T for pure TEGDME, liquid TEGDME/15 wt % LiClO_4 , and BN-polymer/TEGDME/15 wt % LiClO_4 (Figure 6c). The ionic conductivity of the BN-polymer/TEGDME/15 wt % LiClO_4 electrolyte increases up to $2.7 \times 10^{-4} \text{ S cm}^{-1}$ at 373 K. Figure 6c shows a linear variation in plots of $\log(\sigma)$ versus $10^3/T$, suggesting an Arrhenius-type thermally activated

process:⁵⁸ $\sigma = \sigma_0 \exp(-E_a/kT)$ where σ is the conductivity, E_a is the activation energy associated with conduction, k is the Boltzmann constant, T is the temperature, and σ_0 is a preexponential term.

To investigate how the morphological changes of the BN-polymer/TEGDME/ LiClO_4 -based electrolytes could affect the performance, the films were subjected to scanning electron microscopy (SEM) analysis (Figure S41). Pure BN-polymer (see Figure S41a) has a “sponge-like” structure containing cavities with an average diameter in the micrometer range (1–20 μm), whereas no pores are visible in the BN-polymer/TEGDME/ LiClO_4 composite (Figure S41b–d). For the 15 wt % LiClO_4 sample, the higher conductivity could be attributed to the relatively homogenous structure, implying a good dispersion of all components in the electrolyte film (Figure S41b). However, the 20 wt % LiClO_4 sample presents spherical aggregates with an average diameter in the micrometer range, which could be ascribed to the lithium salt not completely dissolved in the polymer matrix (Figure S41c). Increasing the amount of the Li^+ salt to 25 wt % leads to a further decrease in the conductivity of the SSE. This could be attributed to the bigger aggregates formed in the polymer matrix leading to a microstructure that is interspersed with big pores (Figure S41d).

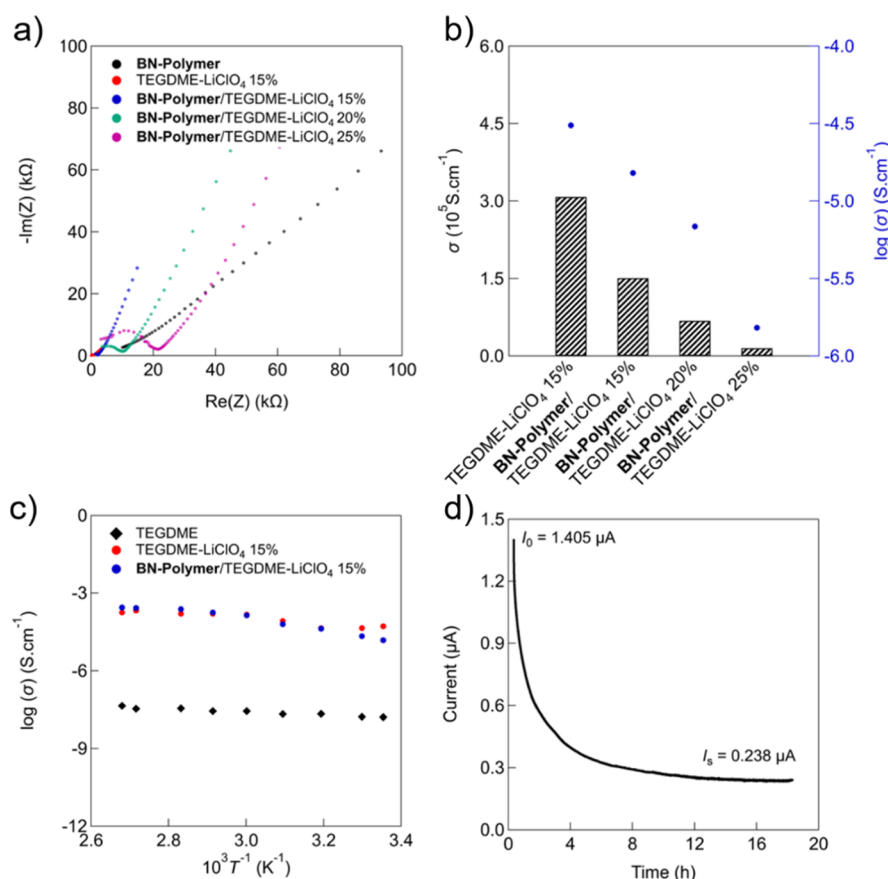


Figure 6. (a) Nyquist plots of pure **BN-polymer**, TEGDME/15 wt % LiClO_4 , and **BN-polymer**/TEGDME/ x LiClO_4 ($x = 15, 20$, or 25 wt % of LiClO_4) at r.t. (b) Ionic conductivity at r.t. of TEGDME/15 wt % LiClO_4 reference and **BN-polymer**-based SSEs as a function of LiClO_4 content. (c) Arrhenius plots of ionic conductivity vs reciprocal absolute temperature for TEGDME, liquid TEGDME/15 wt % LiClO_4 , and **BN-polymer**/TEGDME/15 wt % LiClO_4 electrolytes. (d) Current vs time profile of a symmetrical $\text{Li}/\text{BN-polymer}/\text{TEGDME}/15 \text{ wt } \% \text{LiClO}_4/\text{Li}$ cell configuration after applying a DC voltage of 10 mV on the cell for determining the Li^+ transfer number.

Table 2. Characteristic Properties of the Prepared SSEs at r.t. as a Function of Lithium Salt Content; t : Thickness, A : Surface Area of the Specimen, R_b : Bulk Electrolyte Resistance, and σ : Ionic Conductivity

sample	wt %			t (cm)	A (cm^2)	R_b (Ω)	σ (S cm^{-1})
	BN-polymer	TEGDME	LiClO_4				
0	100			0.018	1.33		
1		85	15	0.02159	1.33	526	3.08×10^{-5}
2	42.5	42.5	15	0.02451	0.79	2060	1.51×10^{-5}
3	40	40	20	0.03	0.79	5580	6.80×10^{-6}
4	37.5	37.5	25	0.036	0.79	20586	1.50×10^{-6}

Table 3. Room Temperature (25°C) and 100°C Values of Ionic Conductivity as Well as Activation Energy Values for the TEGDME, Liquid TEGDME/15 wt % LiClO_4 , **BN-Polymer**/TEGDME/15 wt % LiClO_4 Electrolytes

sample	σ (S cm^{-1}) 25°C	σ (S cm^{-1}) 100°C	E_a (eV) $25\text{--}100^\circ\text{C}$
pure TEGDME	1.62×10^{-8}	4.43×10^{-8}	0.12
liquid TEGDME/15 wt % LiClO_4	3.08×10^{-5}	1.18×10^{-4}	0.20
BN-polymer /TEGDME/15 wt % LiClO_4	1.51×10^{-5}	2.75×10^{-4}	0.38

The thermal stability of the **BN-polymer**/TEGDME/ LiClO_4 SSEs was studied by TGA (measured under a N_2 atmosphere at a heating rate of $10^\circ\text{C min}^{-1}$ from r.t. to 1000°C , Figure S42a) and differential scanning calorimetry (DSC), (measured under a N_2 flow at a ramp rate of $10^\circ\text{C min}^{-1}$, Figure S42b) on both the single components and the films. As expected, **BN-polymer**/TEGDME/ LiClO_4 samples display the same thermal behavior as that of the single components. A

gradual weight loss of about 5% is observed during the initial heating up to 100°C , which can be attributed to the loss of moisture present in the electrolytes. The first significant weight loss (about 45–55%) is due to the evaporation or degradation of TEGDME at 150°C , followed by a gradual loss of about 5% between 300 and 450°C due to LiClO_4 . The complete decomposition of the sample takes place between 500 and 600°C with a weight loss of about 75%, in line with the TGA data

on the pristine **BN-polymer** (Figure S30). The DSC results (Figure S42b) for **BN-polymer**/TEGDME/LiClO₄ in the range 30–250 °C show no endothermic peak associated with melting, which is a further demonstration of the amorphous nature of the polymer and the polymer–salt systems. The absence of a temperature of glass transition suggests the high rigidity of **BN-polymer**, possibly related to the blocking of segmental motion, which would be consistent with the low conductivity. The electrochemical stability of the highest conducting SSE system was also studied with linear sweep voltammetry (LSV) and cyclic voltammetry (CV) using a stainless steel working electrode and Li metal as the reference and counter electrodes. The SSE system proved to be electrochemically stable between 1.84 and 3.70 V versus Li/Li⁺ with no decomposition of any of the components in this potential region (Figure S43a). The CV responses for seven cycles (Figure S43b) present redox peaks around 1.55 and at 3.86 V corresponding to anodic oxidation and cathodic reduction, respectively. While no changes in the redox peak voltages are observed during the cycles, the overlapping of the sweeps indicates that the charge-transfer reaction at the interface between the solid electrolyte and Li metal is reversible. From the CV study, the range of electrochemical stability goes from 2.00 to 3.60 V, which is in very good accordance with LSV results (Figure S43a). Finally, the Li-ion transference number (t_{Li^+}) was evaluated following the potentiostatic polarization method. A symmetric Li/**BN-polymer**/TEGDME/15 wt % LiClO₄/Li cell configuration was used in this study, and a DC polarization with 10 mV potential was applied until the current reached a steady state (Figure 6d). The initial total current decreases with time from 1.405 μA and reaches a steady state value of 0.238 μA after 18 h. The interfacial bulk resistance increases from 2260 to 21,713 Ω through polarization. Thus, the **BN-polymer**/TEGDME/15 wt % LiClO₄ electrolyte presents a transference number of about 0.2, which is in the range of those observed for polyethylene glycol (PEG)-based electrolytes.⁵⁶

CONCLUSIONS

In conclusion, we have reported on a unique BN-doped gel material obtained through a polymerization reaction exploiting the [4 + 2] cycloaddition with CO extrusion between B-ethynyl-substituted borazine and a suitable biscyclopentadiene derivative. Considering that only a few examples of [4 + 2] cycloadditions are reported on alkynes directly bonded to boron atoms,^{59–61} to the best of our knowledge, this is the first example of a Diels Alder cycloaddition for a B-alkynyl functionalized borazine. Solid-state ¹³C NMR, solid-state ¹¹B NMR, and Fourier-transform infrared (FT-IR) spectroscopic characterization were used to confirm the formation of a BN-doped polymeric material, which was characterized by comparison with the monomeric unit. The new BN-doped polyphenylenic material efficiently forms organogels in chlorinated solvents, producing the only example to date of a borazine-doped gel. Rheological investigations confirmed the formation of the gel and allowed us to estimate the shear modulus, the mesh size, and the crosslinking density in the gel. Since the organogel material has high thermal and chemical stability, it was integrated as a support component in a SSE for lithium-ion batteries. In particular, the SSE with **BN-polymer**/TEGDME/15 wt % LiClO₄ composition exhibits an Arrhenius behavior and a r.t. ionic conductivity of $1.51 \times 10^{-5} \text{ S cm}^{-1}$ with a Li⁺ transference number of about 0.2, which is

comparable to that of PEG-based electrolytes. Furthermore, the electrical conductivity studies suggest an enhanced performance of the 15 wt % SSE at higher temperatures. This result represents a stepping stone toward future potential applications of BNC materials in the field of lithium-ion batteries.

EXPERIMENTAL SECTION

Instrumentation. Thin-layer chromatography was conducted on pre-coated aluminum sheets with 0.20 mm Merk Millipore silica gel 60 with fluorescent indicator F254. Column chromatography was carried out using Merck Gerduran silica gel 60 (particle size 40–63 μm). Melting points were measured using a Gallenkamp apparatus in open capillary tubes and have not been corrected. NMR spectra were recorded with a Bruker Fourier 300 MHz spectrometer equipped with a dual (¹³C, ¹H) probe, a Bruker AVANCE III HD 400 MHz NMR spectrometer equipped with a Broadband multinuclear (BBFO) SmartProbe, or a Bruker AVANCE III HD 500 MHz spectrometer equipped with Broadband multinuclear (BBO) Prodigy CryoProbe. ¹H spectra were obtained at 500, 400, or 300 MHz, ¹³C spectra were obtained at 75, 100, or 126 MHz, and ¹¹B were obtained at 128 or 224 MHz; all spectra were obtained at r.t. if not otherwise stated. Chemical shifts were reported in ppm according to tetramethylsilane using the solvent residual signal as an internal reference (CDCl₃: $\delta_{\text{H}} = 7.26 \text{ ppm}$, $\delta_{\text{C}} = 77.16 \text{ ppm}$; CD₂Cl₂: $\delta_{\text{H}} = 5.32$, $\delta_{\text{C}} = 54.00$, C₆D₆: $\delta_{\text{H}} = 7.16 \text{ ppm}$, $\delta_{\text{C}} = 128.6$). Coupling constants (*J*) were given in hertz. Resonance multiplicity was described as s (singlet), d (doublet), t (triplet), dd (doublet of doublets), dt (doublet of triplets), td (triplet of doublets), q (quartet), m (multiplet), and bs (broad signal). Carbon spectra were acquired with a complete decoupling for the proton. LF NMR was performed at 20 °C with a Bruker Minispec mq 20 (0.47 T, Germany). The determination of the average water protons transverse (spin–spin) relaxation time (T_{2m}) was performed according to the CPMG sequence (Carr–Purcell–Meiboom–Gill) $\{90^\circ[-\tau - 180^\circ - \tau(\text{echo})]_n - T_R\}$ with a 8.36 μs wide 90° pulse, $\tau = 250 \mu\text{s}$, and T_R (sequences repetition rate) equal to 5 s. Solid-state NMR ¹H → ¹³C CPMAS NMR spectra were recorded with a Bruker Avance III HD spectrometer at 9.4 T [Larmor frequencies: 400.2 MHz (¹H), 100.6 MHz (¹³C)] using ramped CP for the borazine precursor 2, the reference 3, and the **BN-polymer** material, with magic angle spinning frequencies of 10, 12, and 12 kHz, respectively. ¹¹B MQMAS NMR spectra were recorded with a Bruker Avance III HD spectrometer at 9.4 T [Larmor frequency: 128.4 MHz (¹¹B)] using a four-pulse split- t_1 sequence with the indirect dimension scaled to have the same contribution from the isotropic shift as the direct dimension (please refer to the Supporting Information for more details). Powder XRD data for the crystallized sample of 3 (prepared by diffusion of MeOH into a solution of 3 in CH₂Br₂) were recorded at 21 °C on a Bruker D8 diffractometer (Ge-monochromated CuK α 1 radiation; transmission mode; Vantec detector covering 3° in 2 θ ; 2 θ range, 4° to 50°; step size, 0.016°; data collection time, 119.5 hr). IR spectra were recorded with a Shimadzu IR Affinity 1S FTIR spectrometer in ATR mode with a diamond mono-crystal. Mass spectrometry: (i) high-resolution ESI mass spectra were obtained with a Waters LCT HR TOF mass spectrometer in the positive or negative ion mode. (ii) High-resolution MALDI mass spectra were obtained with a Bruker Autoflex speed MALDI-TOF instrument; the sample was prepared with a 1:1 ratio of sample to the matrix DCTB (15 mg/mL) in CH₂Cl₂; all these analyses were carried out at Cardiff University. Rheological measurements were performed using a stress-controlled rotational rheometer (Haake Mars Rheometer, 379–0200 Thermo Electron GmbH, Karlsruhe, Germany) equipped with parallel plate geometry (PP35, $\phi = 35 \text{ mm}$, serrated surfaces to avoid slippage at the wall). The measuring device was kept at 10 °C inside a glass bell at saturated conditions to avoid evaporation effects. SSEs. TGA was performed using a TGA/SDTA-851 instrument from METTLER TOLEDO at a heating rate of 10 °C min^{−1} from room temperature to 1000 °C under N₂ flow. DSC was performed with a METTLER TOLEDO DSC-1 STAR^c system instrument under a N₂ flow at a

ramp rate of 10 °C min⁻¹ in the temperature range of 30–250 °C. Powder XRD was performed with a STOE StadiP X-ray diffractometer with Cu K α_1 radiation (λ = 1.5418 Å) from 2θ = 1.5 up to 60° with a 0.02° increment using an operation voltage and current of 40 kV and 40 mA, respectively. SEM images were obtained with a FEI XL30 FEG equipped with a Raith laser interferometer-controlled stage operated at 10 kV and the Elphy Plus software. Electrochemical measurements of the optimized films of BN-polymer/TEGDME/LiClO₄-based SSEs were studied using a Biologic Science Instruments VMP-300 potentiostat analyzer.

Materials and Methods. *Synthesis.* Chemicals were purchased from Sigma-Aldrich, Acros Organics, TCI, Apollo Scientific, Alfa Aesar, and Fluorochem and were used as received. Solvents were purchased from Fisher Scientific, while deuterated solvents were from Eurisotop and Sigma-Aldrich. THF and toluene were dried on a Braun MB SPS-800 solvent purification system and further dried over activated 4 Å molecular sieves. Diphenyl ether (Ph₂O) was dried over 4 Å molecular sieves. Aniline was distilled from CaH₂ under reduced pressure and stored away from light in a N₂ atmosphere. Aniline was left on CaH₂ overnight prior to distillation. Low-temperature baths were prepared using different solvent mixtures depending on the desired temperature: −84 °C with ethyl acetate/liq. N₂, −10 °C with ice/NaCl, and 0 °C with ice/H₂O. Anhydrous conditions were achieved by drying Schlenk tubes or two-neck flasks by flaming with a heat gun under vacuum and purging with N₂. The inert atmosphere was maintained using N₂-filled balloons equipped with a syringe and needle that was used to penetrate the silicon stoppers used to close the flask's necks. Additions of liquid reagents were performed using plastic syringes. Degassing of solutions was performed using the freeze–pump–thaw procedure: solutions were frozen using liquid N₂ and kept under vacuum for 10 min before thawing. Alternatively, degassing was performed by bubbling N₂ in the reaction solution under sonication for at least 10 min. Molecular sieves (4 Å) were activated by heating at 165 °C under vacuum overnight and by further heating with heat gun under vacuum immediately before use. All reactions were performed in dry conditions and under inert atmosphere unless otherwise stated. All procedures for the synthesis are reported in the [Supporting Information](#). SSE. Lithium perchlorate LiClO₄ (battery grade) was obtained from Acros and used without further purification. TEGDME from Sigma-Aldrich was dried under vacuum at 70 °C for 48 h before use. CH₂Cl₂ and THF (spectroscopy grade) were purchased from VWR and were used as received.

■ ASSOCIATED CONTENT

SI Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.chemmater.2c01766>.

Details of experimental methodologies and NMR, crystallographic, electrochemical, and rheological characterization of compounds ([PDF](#))

Crystal structure of 2 ([CIF](#))

Crystal structure of *ct*-3 ([CIF](#))

Crystal structure of *cc*-3 ([CIF](#))

Crystal structure of 7 ([CIF](#))

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The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Notes

The authors declare no competing financial interest.

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■ REFERENCES

- (1) Müllen, K. Evolution of graphene molecules: Structural and functional complexity as driving forces behind nanoscience. *ACS Nano* **2014**, *8*, 6531–6541.

- (2) Hammer, B. A. G.; Müllen, K. Dimensional evolution of polyphenylenes: Expanding in all directions. *Chem. Rev.* **2016**, *116*, 2103–2140.
- (3) Narita, A.; Feng, X.; Müllen, K. Bottom-up synthesis of chemically precise graphene nanoribbons. *Chem. Rec.* **2015**, *15*, 295–309.
- (4) Simpson, C. D.; Mattersteig, G.; Martin, K.; Gherghel, L.; Bauer, R. E.; Räder, H. J.; Müllen, K. Nanosized Molecular Propellers by Cyclodehydrogenation of Polyphenylene Dendrimers. *J. Am. Chem. Soc.* **2004**, *126*, 3139–3147.
- (5) Liu, D.; De Feyter, S.; Cotlet, M.; Stefan, A.; Wiesler, U. M.; Herrmann, A.; Grebel-Koehler, D.; Qu, J.; Müllen, K.; De Schryver, F. C. Fluorescence and intramolecular energy transfer in polyphenylene dendrimers. *Macromolecules* **2003**, *36*, 5918–5925.
- (6) Weil, T.; Reuther, E.; Müllen, K. Shape-Persistent, Fluorescent Polyphenylene Dyads and a Triad for Efficient Vectorial Transduction of Excitation Energy. *Angew. Chem., Int. Ed.* **2002**, *41*, 1900–1904.
- (7) Chen, L.; Honsho, Y.; Seki, S.; Jiang, D. Light-harvesting conjugated microporous polymers: Rapid and highly efficient flow of light energy with a porous polyphenylene framework as antenna. *J. Am. Chem. Soc.* **2010**, *132*, 6742–6748.
- (8) Adamski, M.; Skalski, T. J. G.; Britton, B.; Peckham, T. J.; Metzler, L.; Holdcroft, S. Highly Stable, Low Gas Crossover, Proton-Conducting Phenylated Polyphenylenes. *Angew. Chem., Int. Ed.* **2017**, *129*, 9186–9189.
- (9) Miyake, J.; Taki, R.; Mochizuki, T.; Shimizu, R.; Akiyama, R.; Uchida, M.; Miyatake, K. Design of flexible polyphenylene proton-conducting membrane for next-generation fuel cells. *Sci. Adv.* **2017**, *3*, No. eaao0476.
- (10) Li, C.; Liu, M.; Pschirer, N. G.; Baumgarten, M.; Müllen, K. Polyphenylene-based materials for organic photovoltaics. *Chem. Rev.* **2010**, *110*, 6817–6855.
- (11) Huang, Y.; Mai, Y.; Yang, X.; Beser, U.; Liu, J.; Zhang, F.; Yan, D.; Müllen, K.; Feng, X. Temperature-Dependent Multidimensional Self-Assembly of Polyphenylene-Based “Rod-Coil” Graft Polymers. *J. Am. Chem. Soc.* **2015**, *137*, 11602–11605.
- (12) Helten, H. B=N Units as Part of Extended π -Conjugated Oligomers and Polymers. *Chem.—Eur. J.* **2016**, *22*, 12972–12982.
- (13) Bonifazi, D.; Fasano, F.; Lorenzo-Garcia, M. M.; Marinelli, D.; Oubaha, H.; Tasseroul, J. Boron–nitrogen doped carbon scaffolding: organic chemistry, self-assembly and materials applications of borazine and its derivatives. *Chem. Commun.* **2015**, *51*, 15222–15236.
- (14) Hammer, B. A. G.; Wu, Y.; Fischer, S.; Liu, W.; Weil, T.; Müllen, K. Controlling Cellular Uptake and Toxicity of Polyphenylene Dendrimers by Chemical Functionalization. *ChemBioChem* **2017**, *18*, 960–964.
- (15) John, H.; Bauer, R.; Espindola, P.; Sonar, P.; Heinze, J.; Müllen, K. 3D-hybrid networks with controllable electrical conductivity from the electrochemical deposition of terthiophene-functionalized polyphenylene dendrimers. *Angew. Chem., Int. Ed.* **2005**, *44*, 2447–2451.
- (16) Oubaha, H.; Demitri, N.; Rault-Berthelot, J.; Dubois, P.; Coulembier, O.; Bonifazi, D. Photoactive Boron-Nitrogen-Carbon Hybrids: From Azo-borazines to Polymeric Materials. *J. Org. Chem.* **2019**, *84*, 9101–9116.
- (17) Hou, I. C. Y.; Hu, Y.; Narita, A.; Müllen, K. Diels-Alder polymerization: A versatile synthetic method toward functional polyphenylenes, ladder polymers and graphene nanoribbons. *Polym. J.* **2018**, *50*, 3–20.
- (18) Hizal, G.; Tunca, U.; Sanyal, A. Discrete macromolecular constructs via the Diels-Alder “click” reaction. *J. Polym. Sci.* **2011**, *49*, 4103–4120.
- (19) Wiesler, U. M.; Berresheim, A. J.; Morgenroth, F.; Lieser, G.; Müllen, K. Divergent synthesis of polyphenylene dendrimers: The role of core and branching reagents upon size and shape. *Macromolecules* **2001**, *34*, 187–199.
- (20) Marinelli, D.; Fasano, F.; Najjari, B.; Demitri, N.; Bonifazi, D. Borazino-Doped Polyphenylenes. *J. Am. Chem. Soc.* **2017**, *139*, 5503–5519.
- (21) Dosso, J.; Marinelli, D.; Demitri, N.; Bonifazi, D. Structural Properties of Highly Doped Borazino Polyphenylenes Obtained through Condensation Reaction. *ACS Omega* **2019**, *4*, 9343–9351.
- (22) Riensch, N. A.; Deniz, A.; Köhl, S.; Müller, L.; Adams, A.; Pich, A.; Helten, H. Borazine-based inorganic–organic hybrid cyclomatrix microspheres by silicon/boron exchange precipitation polycondensation. *Polym. Chem.* **2017**, *8*, 5264.
- (23) Ayhan, O.; Eckert, T.; Plamper, F. A.; Helten, H. Poly-(iminoborane)s: An Elusive Class of Main-Group Polymers? *Angew. Chem., Int. Ed.* **2016**, *55*, 13321–13325.
- (24) Reich, T. E.; Behera, S.; Jackson, K. T.; El-Kaderi, H. M.; Jena, P. Highly selective CO₂/CH₄ gas uptake by a halogen-decorated borazine-linked polymer. *J. Mater. Chem.* **2012**, *22*, 13524–13528.
- (25) Reich, T. E.; Jackson, K. T.; Li, S.; Jena, P.; El-Kaderi, H. M. Synthesis and characterization of highly porous borazine-linked polymers and their performance in hydrogen storage application. *J. Mater. Chem.* **2011**, *21*, 10629–10632.
- (26) Bagheri, S.; Masoodi, H. R. Theoretical study of the influence of cation- π and anion- π interactions on some NMR data of borazine complexes. *Chem. Phys. Lett.* **2015**, *629*, 46–52.
- (27) Yang, Y.; Mehta, M. A.; Fujinami, T.; Inoue, T. Ionic conductivity and interfacial properties of polymer electrolytes based on PEO and boroxine ring polymer. *J. Appl. Polym. Sci.* **2002**, *84*, 17–21.
- (28) Tao, R.; Fujinami, T. Improvement of electrochemical properties of PEO-LiTFSI electrolyte by incorporation of boroxine polymers with different backbone lengths. *J. Appl. Electrochem.* **2005**, *35*, 163–168.
- (29) Resendiz-Lara, D. A.; Stubbs, N. E.; Arz, M. I.; Pridmore, N. E.; Sparkes, H. A.; Manners, I. Boron-nitrogen main chain analogues of polystyrene: poly(B-aryl)aminoboranes via catalytic dehydrocoupling. *Chem. Commun.* **2017**, *53*, 11701–11704.
- (30) Resendiz-Lara, D. A.; Whittell, G. R.; Leitao, E. M.; Manners, I. Catalytic Synthesis, Characterization, and Properties of Polyamino-borane Homopolymers and Random Copolymers. *Macromolecules* **2019**, *52*, 7052–7064.
- (31) Staubitz, A.; Presa Soto, A.; Manners, I. Iridium-Catalyzed Dehydrocoupling of Primary Amine-Borane Adducts: A Route to High Molecular Weight Polyaminoboranes, Boron-Nitrogen Analogues of Polyolefins. *Angew. Chem., Int. Ed.* **2008**, *47*, 6212–6215.
- (32) Lorenzo-García, M. M.; Bonifazi, D. Renaissance of an Old Topic: From Borazines to BN-doped Nanographenes. *Chimia* **2017**, *71*, 550–557.
- (33) Nagasawa, K. Borazines Stable to Hydrolysis. *Inorg. Chem.* **1966**, *5*, 442–445.
- (34) Fasano, F.; Dosso, J.; Bezzu, C. G.; Carta, M.; Kerff, F.; Demitri, N.; Su, B. L.; Bonifazi, D. BN-Doped Metal–Organic Frameworks: Tailoring 2D and 3D Porous Architectures through Molecular Editing of Borazines. *Chem.—Eur. J.* **2021**, *27*, 4124–4133.
- (35) Uchamaru, Y.; Koda, N.; Yamashita, H. Thermally stable borazine-based polymer. *Phosphorus, Sulfur, Silicon Relat. Elem.* **2001**, *169*, 231–234.
- (36) Watanabe, H.; Totani, T.; Yoshizaki, T. Preparation and Conformation of B-Triethynylborazine Derivatives. *Inorg. Chem.* **1965**, *4*, 657–660.
- (37) Lorenzo-García, M. M.; Bonifazi, D. Renaissance of an old topic: From borazines to BN-doped nanographenes. *Chimia* **2017**, *71*, 550–557.
- (38) Kervyn, S.; Fenwick, O.; Di Stasio, F.; Shin, Y. S.; Wouters, J.; Accorsi, G.; Osella, S.; Beljonne, D.; Cacialli, F.; Bonifazi, D. Polymorphism, fluorescence, and optoelectronic properties of a borazine derivative. *Chem.—Eur. J.* **2013**, *19*, 7771–7779.
- (39) Groszós, S. J.; Stafiej, S. F. Organoboron Compounds. I. A New Synthesis of B-Trialkyl and Triaryl-N-triphenylborazoles. *J. Am. Chem. Soc.* **1958**, *80*, 1357–1360.
- (40) Wakamiya, A.; Ide, T.; Yamaguchi, S. Toward π -conjugated molecule bundles: Synthesis of a series of B,B',B''-trianthryl-N,N',N''-triarylborazines and the bundle effects on their properties. *J. Am. Chem. Soc.* **2005**, *127*, 14859–14866.

- (41) Brown, S. P.; Heyes, S. J.; Wimperis, S. Two-dimensional MAS multiple-quantum NMR of quadrupolar nuclei. Removal of inhomogeneous second-order broadening. *J. Magn. Reson., Ser. A* **1996**, *119*, 280–284.
- (42) Perrault, C. M.; Juncker, D.; Eon, P. H. Preparation and shear modulus of polyacrylamide gels as nerve cell culture. *AIP Conf. Proc.* **2008**, *1027*, 615–617.
- (43) Moins, S.; Martins, J. C.; Krumpmann, A.; Lemaure, V.; Cornil, J.; Delbosc, N.; Decroly, A.; Dubois, P.; Lazzaroni, R.; Gohy, J.-F.; Coulembier, O. Potential of polymethacrylate pseudo crown ethers as solid state polymer electrolytes. *Chem. Commun.* **2017**, *53*, 6899–6902.
- (44) Stephan, A. M. Review on gel polymer electrolytes for lithium batteries. *Eur. Polym. J.* **2006**, *42*, 21–42.
- (45) Xue, Z.; He, D.; Xie, X. Poly(ethylene oxide)-based electrolytes for lithium-ion batteries. *J. Mater. Chem. A* **2015**, *3*, 19218–19253.
- (46) Huh, P. H.; Choi, M. G.; Jo, J. J.; Lee, J. K.; Lee, J. O.; Yang, W. Effect of salt concentration on the glass transition temperature and ionic conductivity of poly(ethylene glycol)-polyurethane/LiClO₄ complexes. *Macromol. Res.* **2004**, *12*, 422–426.
- (47) Rodrigues, L. C.; Barbosa, P. C.; Silva, M. M.; Smith, M. J. Electrochemical and thermal properties of polymer electrolytes based on poly(epichlorohydrin-co-ethylene oxide-co-allyl glycidyl ether). *Electrochim. Acta* **2007**, *53*, 1427–1431.
- (48) Stolwijk, N. A.; Heddier, C.; Reschke, M.; Wiencierz, M.; Bokeloh, J.; Wilde, G. Salt-concentration dependence of the glass transition temperature in PEO-NaI and PEO-LiTFSI polymer electrolytes. *Macromolecules* **2013**, *46*, 8580–8588.
- (49) Terada, S.; Mandai, T.; Suzuki, S.; Tsuzuki, S.; Watanabe, K.; Kamei, Y.; Ueno, K.; Dokko, K.; Watanabe, M. Thermal and electrochemical stability of tetraglyme-magnesium bis-(trifluoromethanesulfonyl)amide complex: Electric field effect of divalent cation on solvate stability. *J. Phys. Chem. C* **2016**, *120*, 1353–1365.
- (50) Karan, N.; Pradhan, D.; Thomas, R.; Natesan, B.; Katiyar, R. S. Solid polymer electrolytes based on polyethylene oxide and lithium trifluoro-methane sulfonate (PEO-LiCF₃SO₃): Ionic conductivity and dielectric relaxation. *Solid State Ionics* **2008**, *179*, 689–696.
- (51) Sakakibara, T.; Kitamura, M.; Honma, T.; Kohno, H.; Uno, T.; Kubo, M.; Imanishi, N.; Takeda, Y.; Itoh, T. Cross-linked polymer electrolyte and its application to lithium polymer battery. *Electrochim. Acta* **2019**, *296*, 1018–1026.
- (52) Naiwi, T. S. R. T.; Aung, M. M.; Ahmad, A.; Rayung, M.; Su'ait, M. S.; Yusof, N. A.; Lae, K. Z. W. Enhancement of plasticizing effect on bio-based polyurethane acrylate solid polymer electrolyte and its properties. *Polymers* **2018**, *10*, 1142.
- (53) Ibrahim, S.; Ahmad, A.; Mohamed, N. S. Characterization of novel castor oil-based polyurethane polymer electrolytes. *Polymers* **2015**, *7*, 747–759.
- (54) Michael, M. S.; Jacob, M. M. E.; Prabakaran, S. R. S.; Radhakrishna, S. Enhanced lithium ion transport in PEO-based solid polymer electrolytes employing a novel class of plasticizers. *Solid State Ionics* **1997**, *98*, 167–174.
- (55) Rathod, S. G.; Bhajantri, R. F.; Ravindrachary, V.; Pujari, P. K.; Nagaraja, G. K.; Naik, J.; Hebbar, V.; Chandrappa, H. Temperature-dependent ionic conductivity and transport properties of LiClO₄-doped PVA/modified cellulose composites. *Bull. Mater. Sci.* **2015**, *38*, 1213–1221.
- (56) Guo, Z.; Zhang, Y.; Dong, Y.; Li, J.; Li, S.; Shao, P.; Feng, X.; Wang, B. Fast Ion Transport Pathway Provided by Polyethylene Glycol Confined in Covalent Organic Frameworks. *J. Am. Chem. Soc.* **2019**, *141*, 1923–1927.
- (57) Carbone, L.; Gobet, M.; Peng, J.; Devany, M.; Scrosati, B.; Greenbaum, S.; Hassoun, J. Polyethylene glycol dimethyl ether (PEGDME)-based electrolyte for lithium metal battery. *J. Power Sources* **2015**, *299*, 460–464.
- (58) Yang, P.; Liu, L.; Li, L.; Hou, J.; Xu, Y.; Ren, X.; An, M.; Li, N. Gel polymer electrolyte based on polyvinylidene fluoride-co- hexafluoropropylene and ionic liquid for lithium ion battery. *Electrochim. Acta* **2014**, *115*, 454–460.
- (59) Craciun, L.; Ho, D. M.; Jones, M.; Pascal, R. A. A carboranylpentaphenylbenzene. *Tetrahedron Lett.* **2004**, *45*, 4985–4987.
- (60) Harrity, J. P.; Moore, J. E.; York, M. A metal-free cycloaddition approach to highly substituted aromatic boronic esters. *Synlett* **2005**, 860–862.
- (61) Ester, C.; Maderna, A.; Pritzkow, H.; Siebert, W. Syntheses, structures, and reactivity of hexaborylbenzene derivatives. *Eur. J. Inorg. Chem.* **2000**, 1177–1184.

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