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Oubaha, Hamid, Demitri, Nicola, Rault-Berthelot, Joëlle, Dubois, Philippe, Coulembier, Olivier and Bonifazi, Davide 2019. Photoactive boron-nitrogen-carbon hybrids: from azo-borazines to polymeric materials. *Journal of Organic Chemistry* 84 (14) , pp. 9101-9116. 10.1021/acs.joc.9b01046

Publishers page: <http://dx.doi.org/10.1021/acs.joc.9b01046>

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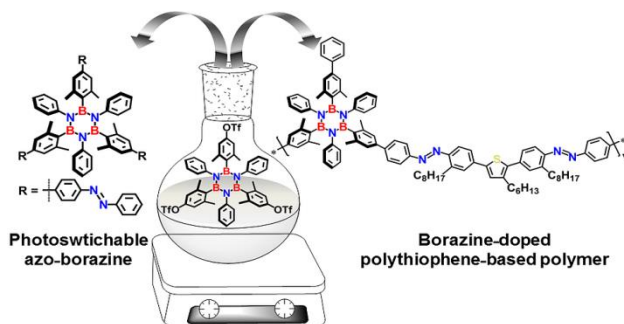


Photoactive boron-nitrogen-carbon hybrids: from azo-borazines to polymeric materials

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KEYWORDS: azobenzene, borazine, molecular switches, *trans/cis* isomerization, cross-coupling, polymers, polythiophene, solvatochromism.



ABSTRACT: In this paper, we describe the synthetic routes for preparing a novel switchable BNC-based chromophore, composed of a borazine core peripherally functionalized with azobenzene moieties. Capitalizing on the Pd-catalyzed *Suzuki* cross-coupling reaction between a *tris*-triflate borazine and an organoboron azobenzene derivative, a photoswitchable azo-borazine derivative was successfully prepared. The molecule showed reversible *E/Z* photoisomerization upon irradiation at the maximum of the intense π - π^* absorption feature (360 nm). X-Ray crystallographic investigations revealed a non-planar orientation of the three azobenzene moieties and the *trans* configuration of the $-N=N-$ bonds. Building on the synthetic versatility of the borazine-azobenzene derivative, we used this photoactive scaffold to engineer soluble BN-doped polythiophene polymers. Photophysical characterization performed in solvents of different polarity, suggested that the polymer undergoes intramolecular charge-transfer (ICT).

INTRODUCTION

Stimuli-responsive molecules and materials capable of undergoing reversible structural changes are of great interest for future technology.¹ A typical example is the *E/Z* photoisomerization of azobenzenes, which is characterized by an irradiation-induced *Z*→*E* isomerization between the thermodynamically favored, stretched *E*-isomer and its *Z*-isomer. The photoisomerization is accompanied by a dramatic change in electronic structure, geometric shape, and polarity, resulting into the spatiotemporal control of the conformational and overall structural properties of molecular systems.² Notable applications include their integration into architectures expressing functional properties in materials science,³ biology⁴ and molecular recognition.^{1c, 5}

The replacement of C=C units by more polarized BN couples is a versatile functionalization strategy to tailor the optoelectronic characteristics of organic π -systems without significant structural alteration of the nanostructure.⁶ The BN/CC isosterism concept goes back to the seminal discovery of the borazine ($\text{H}_3\text{B}_3\text{N}_3\text{H}_3$) by *Stock* and *Pohland* in 1926.⁷ The borazine core is commonly referred to as the “inorganic benzene”, owing to its *iso*-electronic and *iso*-structural relationships with benzene.⁷ Accordingly, borazine and its derivatives are valuable scaffolds to be inserted as doping units in graphitic-based carbon materials to tailor their optoelectronic characteristics⁸ and self-assembly properties on surfaces.^{9, 23} Important examples include dendritic borazine-doped polyphenylenes,¹⁰ borazine-doped coronenes¹¹ and borazino-based polymers.^{10, 12} Not only borazine-based materials, but also hybrid BNC-based architectures are attracting increasing consideration due to their thermal stability, mechanical robustness and tunable optoelectronic properties.¹³ In this respect, the incorporation of BN units into π -conjugated polymers is emerging as an evident route to broaden the scope of organic electronic materials.^{6d, 13b, 14} For their part, π -conjugated polymers have revolutionized the field of optoelectronics¹⁵ and are nowadays used to produce low-cost, large-area, and flexible electronic devices.¹⁶ In these materials, the tailoring of the LUMO/HOMO energy levels for a given application is of high importance.^{15d, 17} Among the different strategies, the incorporation of heteroatomic units into the polymer structures such as boron and nitrogen atoms revealed to be a powerful strategy to prepare isostructural molecular^{6c, 11, 18} and polymeric materials^{6d, 14a, 19} bearing strong local dipole moments.^{14a, 14b, 20}

Owing to the possibility of functionalizing the borazine core with different groups on the aryl substituents at the *N* and *B* atoms,^{6e, 7d, 21} and to the easy polymerization of thiophenes for producing low-cost and flexible electronic devices, we have envisaged to employ the borazine (B_3N_3) as the core skeleton and azobenzene as photo-responsive unit to engineer new photoswitchable molecular BNC hybrid materials (**Fig. 1**). Considering the conductive properties of poly(3-hexylthiophene) (P3HT),²² the 3-hexylthiophene scaffold was selected as monomeric unit to develop the polymer material. Also, we were not unmindful to exploit the side alkyl chains to enhance the solubility of the final material (**Fig. 1b**).

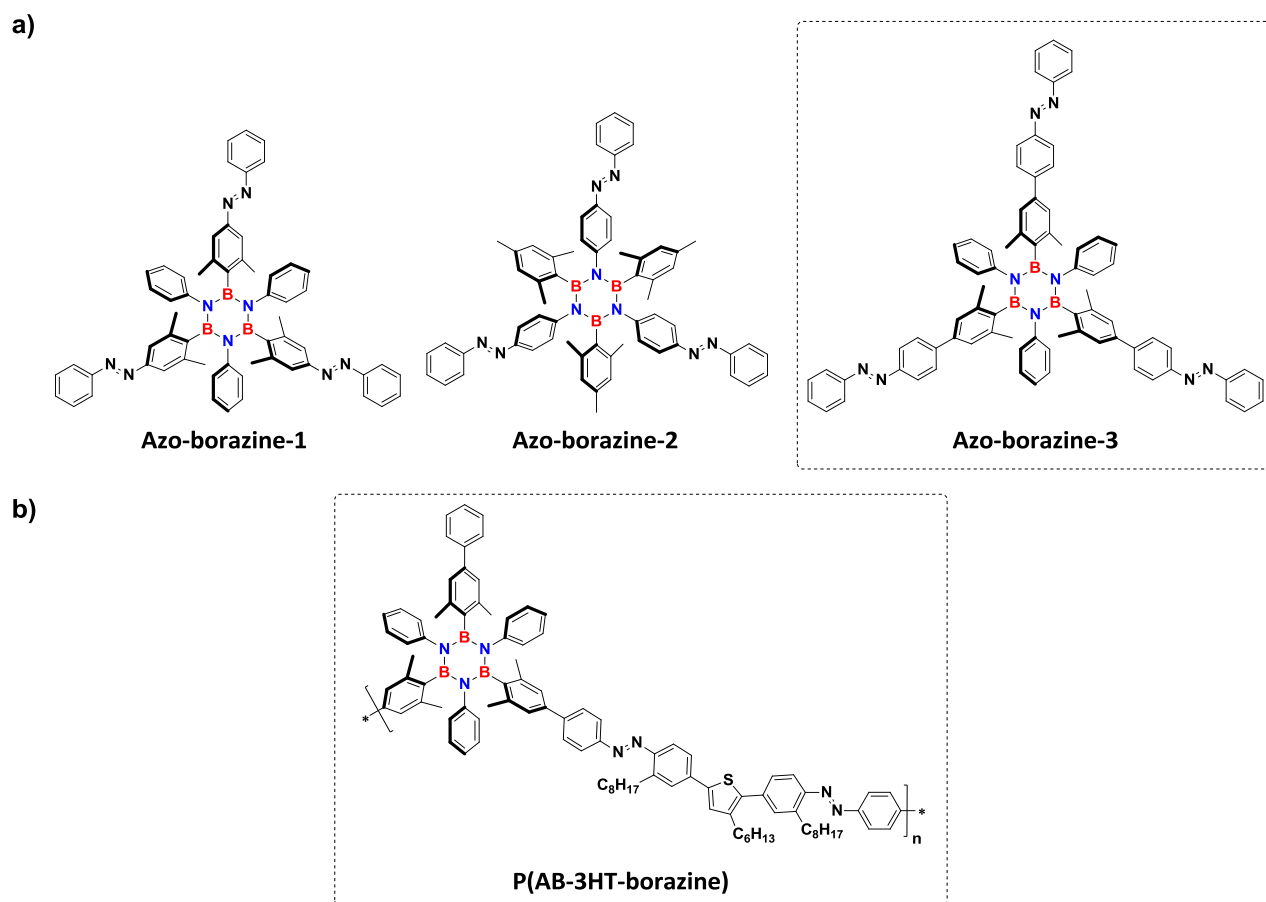
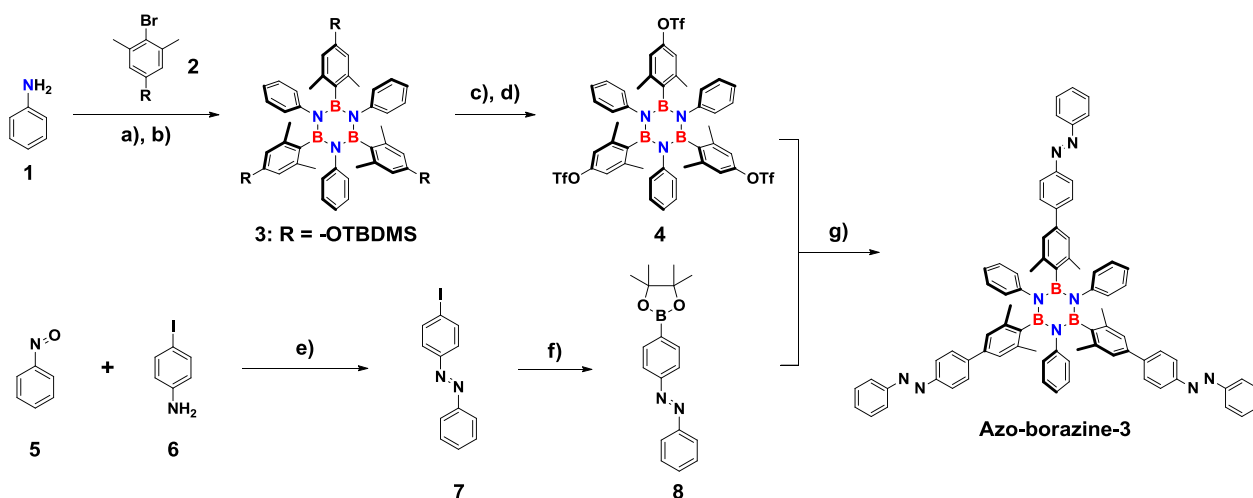


Figure 1. Molecular structures of the targeted borazine-based BNC molecules and hybrid polymers.

RESULTS AND DISCUSSION

Synthesis the azo-borazine core. Three classes of derivatives were targeted as photoactive BNC molecular scaffold: **azo-borazine-1**, **azo-borazine-2**, and **azo-borazine-3** (**Fig. 1a**). Due to the chemical sensitivity of the B_3N_3 core under the reaction conditions to form azobenzene moiety, **azo-borazine-1** and **azo-borazine-2** could not be prepared (see SI for the attempts). Thus, we decided to follow a convergent synthetic pathway, in which a precursor B_3N_3 core is reacted with azobenzene moieties, avoiding the need to use strong oxidizing conditions to form the azo-functional group. It is following this approach, that we prepared **azo-borazine-3** reacting a suitable aryl halide or related electrophile (*i.e.*, $ArOTf$) with an organoboron derivative of a given azobenzene under the *Suzuki* cross-coupling reaction conditions (**Scheme 1**). Specifically, the *tert*-butyldimethylsilyl (TBDMS)-protected borazine derivative **3** was prepared following a modified protocol^{8, 23} of that reported by Groszos and Stafiej²⁴ and later by Yamaguchi.²¹ The synthesis started with the reaction of aniline **1** with BCl_3 under refluxing conditions gave intermediate tri-*B*-chloro-tri-*N*-phenylborazine that, upon reaction with TBDMS-protected $ArLi$, could be transformed into TBDMS-protected borazine derivative **3** in 49% yield (reaction a-b). Removal of the TBDMS protecting group with TBAF and successive esterification with Tf_2O in pyridine, led to key intermediate *tris*-triflate borazine **4** in 75% yield (reactions c-d).¹⁰ White suitable crystals were obtained from evaporation of a solution containing either compound **3** (**Fig. 2a**) or compound **4** (**Fig. 2b**) in $CHCl_3$ belonging to the *P-1* and *P2₁/n* space groups, respectively. As expected, the X-ray molecular structures show the quasi-orthogonal arrangement of the aryl moieties, which place the methyl substituents atop the *B* atoms, sterically shielding the electrophilic *B*-centers.²⁵ In parallel, the synthesis of the organoboron derivative of

azobenzene **8** was performed through the *Mills* reaction, starting from commercially available nitrosobenzene **5** and 4-iodoaniline **6** followed by *Miyaura* borylation reaction (**Scheme 1**, reactions e-f). Final *Suzuki* cross-coupling reaction between borazine **4** and azobenzene-boronic ester **8** yielded desired **azo-borazine-3** in 83% yield (**Scheme 1**, reaction g).



Scheme 1. Synthetic approach toward the preparation of azobenzene-borazine-based BNC hybrid materials. Reagents and conditions: (a) BCl_3 , toluene, reflux, 18 h; (b) **2**, *tert*-BuLi, THF, -84°C to r.t., 16 h, 49%; (c) TBAF, THF, 0°C , 2 h; (d) Tf_2O , pyridine, r.t., 16 h, 75%; (e) AcOH, r.t., 18 h, 81%; (f) B_2Pin_2 , KOAc, $[\text{Pd}(\text{dppf})\text{Cl}_2\cdot\text{CH}_2\text{Cl}_2]$, DMF, 80%; (g) K_2CO_3 , $[\text{Pd}(\text{PPh}_3)_4]$, 1,4-dioxane: H_2O (3:1 v/v), 105°C , 18 h, 83%. TBAF: *tetra*-butylammonium fluoride. Tf_2O : *tri*-fluoromethanesulfonic anhydride. B_2Pin_2 : 4,4,4',4',5,5',5',5'-Octamethyl-2,2'-*bi*-1,3,2-dioxaborolane. $[\text{Pd}(\text{dppf})\text{Cl}_2\cdot\text{CH}_2\text{Cl}_2]$: [1,1'-bis(diphenylphosphino)ferrocene dichloropalladium(II)] dichloromethane adduct.

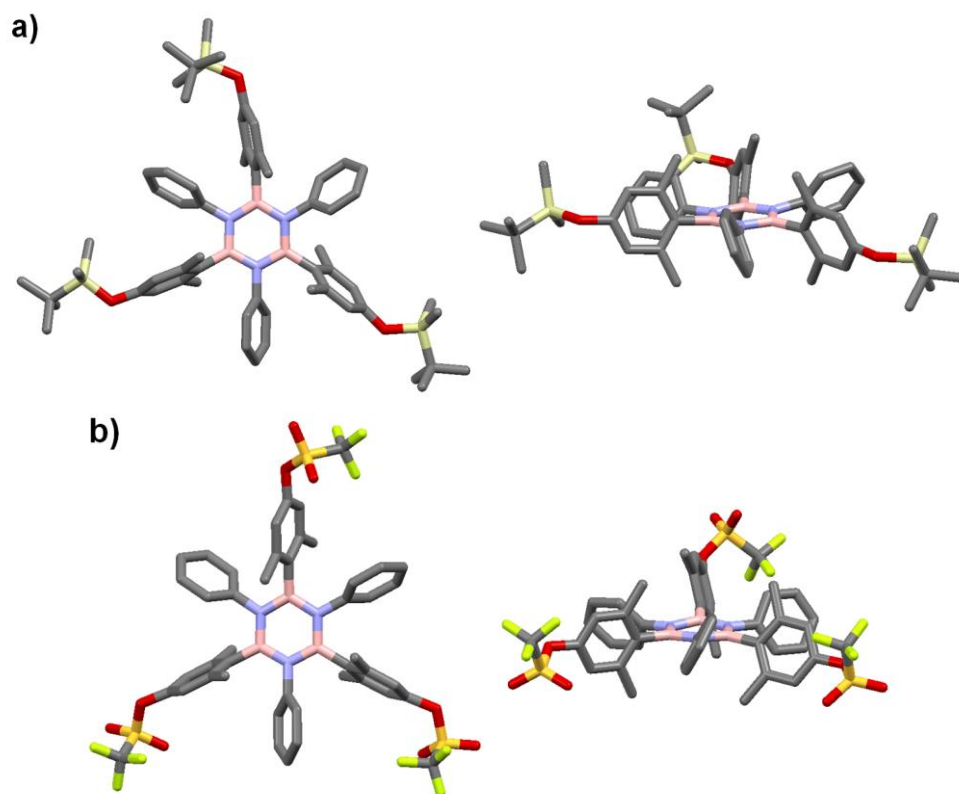


Figure 2. X-ray determined molecular structure of **a)** compound **3** and **b)** compound **4** (Atom colors: blue N, pink B, red O, yellow Si, orange S, light green F, gray C; hydrogen atoms are omitted for clarity). Space group: **a)** $P-1$ **b)** $P2_1/n$.

Azo-borazine-3 derivative was characterized using NMR, IR, and MALDI-ToF spectrometry. At first, **azo-borazine-3** was identified by MALDI-ToF through the detection of the peak corresponding to the molecular ion at m/z 1162.6010 ($[M+H]^+$) (calcd for $C_{78}H_{67}B_3N_9^+$ 1162.5793). 1H NMR analysis further confirmed the chemical structure (**Fig. 3**) and the ^{11}B NMR spectrum showed a broad peak at 36.0 ppm (inset **Fig. 3**), consistent with a boron atom substituted by two *N*-atoms and linked to one carbon atom. Finally, IR spectrum revealed an intense band at 1353 cm^{-1} , characteristic of the B-N stretching in a borazine core.⁸

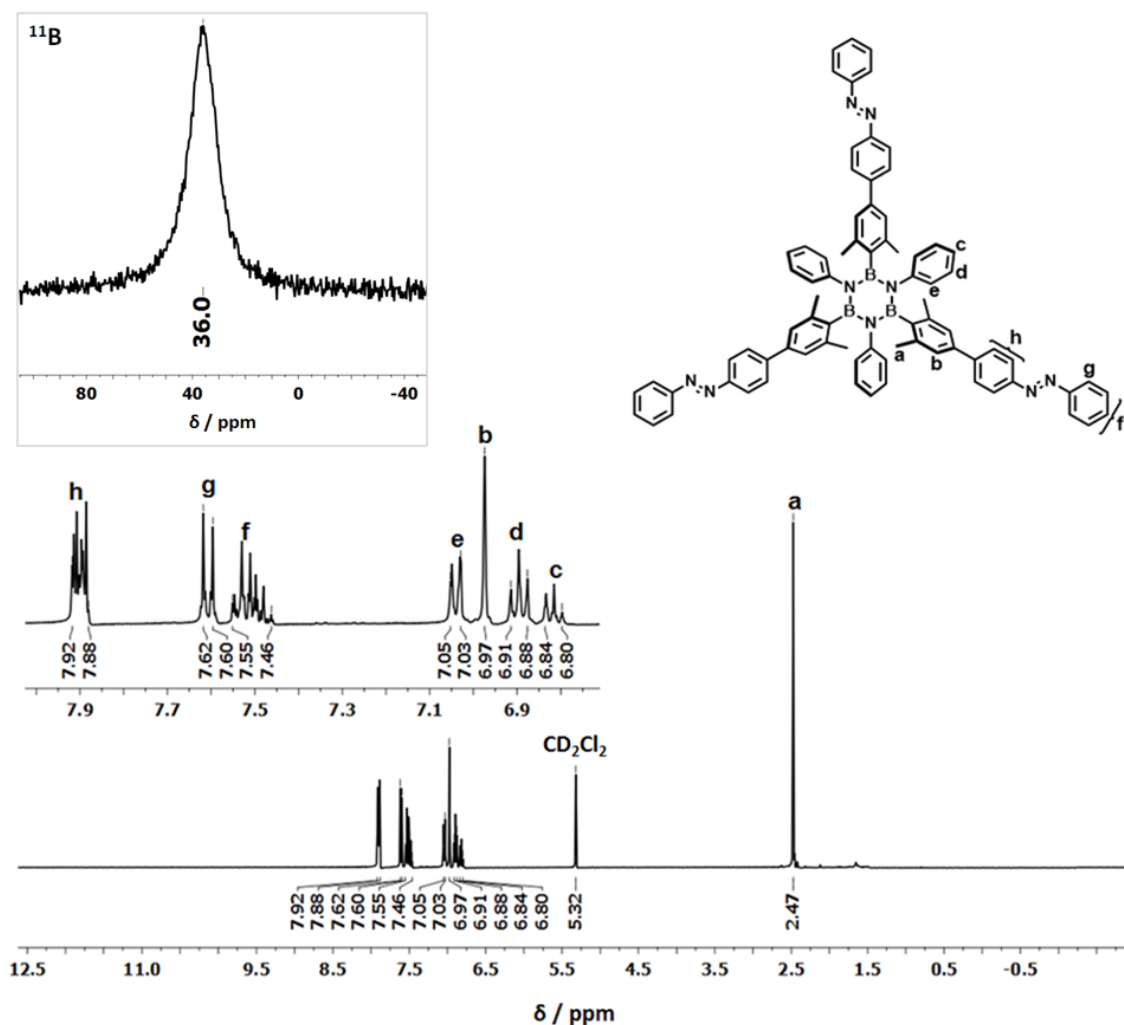


Figure 3. 1H NMR spectrum of **azo-borazine-3** (CD_2Cl_2 , 400 MHz, 25 °C). Inset: a zoom of the region 6.60–8.00 ppm and ^{11}B NMR spectrum ($CDCl_3$, 128 MHz, 25 °C).

Small needle-like orange crystals of **azo-borazine-3** suitable for X-ray diffraction were grown by vapor diffusion of hexane to a solution of **azo-borazine-3** in CH_2Cl_2 . Similarly to borazine **4**, the X-ray molecular structure shows the quasi-orthogonal arrangement of the aryl moieties. The X-ray analysis confirms the *trans* configuration of the $-N=N-$ functional groups and reveals the non-planar arrangement of the three aryl rings at the solid state (**Fig. 4a**). A deviation from orthogonal arrangement of the aryl substituents and the borazine ring is clearly observed with inter-planar angles of 122.6° between the borazine ring and phenyl groups, and of 108.9° between both phenyl and xylyl groups. As previously observed in the literature,^{8, 25} the intra-annular distance values between boron and nitrogen atoms are between 1.43 and 1.46 Å, with the internal angle of the borazine cycle between 122.7° and 117.03° (BNB and NBN, respectively).

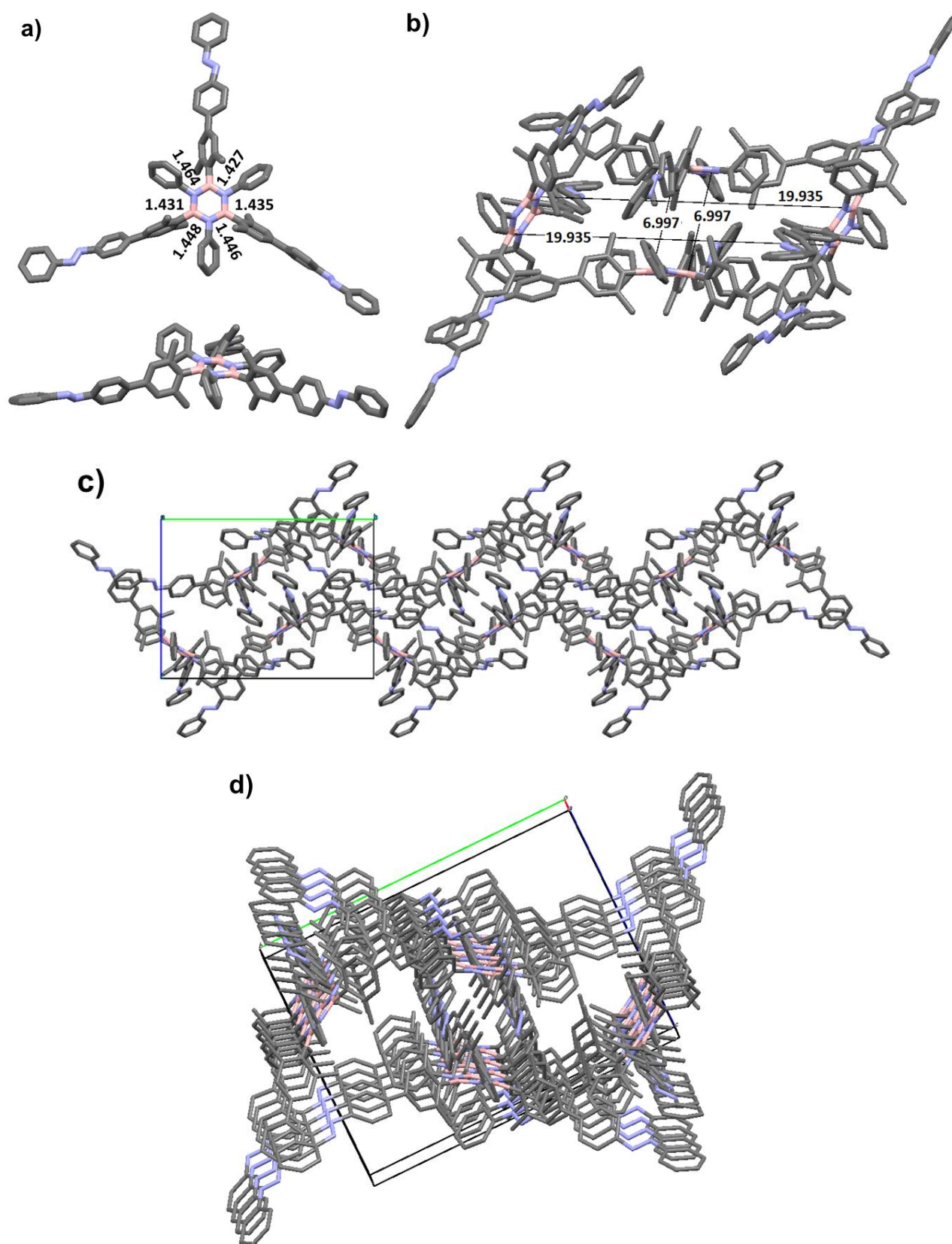


Figure 4. (a) Frontal (**top**) and side (**bottom**) views of the X-ray crystal structure of **azo-borazine-3** (with the B-N distances in Å), (b) packing diagram viewed along the *a* axis and (c) Crystal packing, along the *a* crystallographic direction, showing zigzag stacking shape of the molecules. (Atom colors: blue N, pink B, gray C; hydrogen atoms are omitted for clarity). Space group: *P21/c*.

Notably, the azobenzene moieties in the crystal exist in a twisted shape with respect to the *B*-Aryl groups (**Fig. 4a**). Interestingly, the crystal packing four **azo-borazine-3** molecules stack in parallelepiped-like fashion. The borazine cores constituting the four faces of the arrangement, with each *B*-center facing the *N*-center forming layers separated by a distance of 6.99 and 19.93 Å (**Fig. 4b**). Each layer is formed by borazine molecules lying side by side on the same plane and separated by a distance of 20.22 Å (**Fig. 4c**) and the adjacent layers are assembled in a zigzag-like stacking forming a 3D porous assembly (**Fig. 4d**).

Photo- and thermal-isomerizations of the azo-borazine-3. The photo-isomerization reaction between the *E*- and *Z*-isomers of **azo-borazine-3** was studied in toluene ($5 \cdot 10^{-6}$ M) at r.t. (**Fig. 5**). The molecule exhibits two UV absorption bands at 360 nm and 450 nm, which are attributed to $\pi\text{-}\pi^*$ (of the *E*-isomer) and $n\text{-}\pi^*$ (of the *Z*-isomer) electronic transitions, respectively.^{2b, 26} When irradiated at 360 nm, remarkable changes are observed in the absorption spectra of the **azo-borazine-3** (**Fig. 5a**), with an intense and fast increase of a new band centered at around 290 nm and a substantial increase of the intensity of the $n\text{-}\pi^*$ band at 450 nm (inset of **Fig. 5a**) with a concomitant disappearance of the $\pi\text{-}\pi^*$ absorption band. Neat isosbestic points are maintained throughout the photoradiation experiments, showing the clean occurrence of the nearly quantitative *E*→*Z* isomerization after 3 min of irradiation, while no evolution in the UV-Vis absorption was observed even after 20 min of irradiation at 360 nm. On the other hand, irradiation at 450 nm causes the *Z*→*E* isomerization, as observed by the simultaneous decrease and increase of the $n\text{-}\pi^*$ and $\pi\text{-}\pi^*$ absorption bands, respectively. The full restoring of the *E*-isomer was not observed even after 23 min of irradiation at 450 nm (**Fig. 5b**).

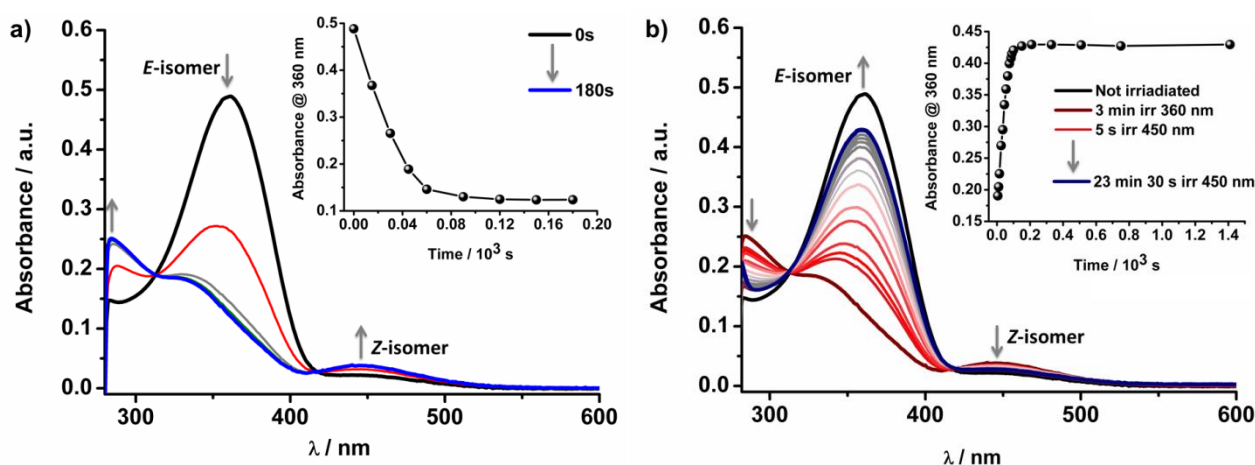


Figure 5. Reversible change in the absorption spectra of a $5 \cdot 10^{-6}$ M solution of **azo-borazine-3** in toluene under irradiation at (a) 360 nm and (b) 450 nm at r.t. (23 °C). Inset: UV absorption at 360 nm of **azo-borazine-3** in toluene as a function of irradiation time.

Repetitive photo-induced switching cycles evidenced an excellent fatigue resistance and remarkable photostability of **azo-borazine-3** (**Fig. 6**). As expected, the borazine core has little influence on the photochemical properties, with the **azo-borazine-3** showing the typical *E/Z* switching behavior of standard azobenzenes.^{2b, 27} The UV absorbing BN-core does not interfere with the excitation properties of the azobenzene moieties. The thermal *Z*→*E* isomerization (obtained after irradiation at 360 nm for 3 min) was realized by measuring the absorbance of a $5 \cdot 10^{-6}$ M solution of **azo-borazine-3** in toluene at different temperatures (**Fig. 7a**). As shown in **Figure 7**, the

thermal $Z \rightarrow E$ interconversion occurs very slowly, and the spectral profile of the E -isomer is fully recovered only upon heating at 100 °C.

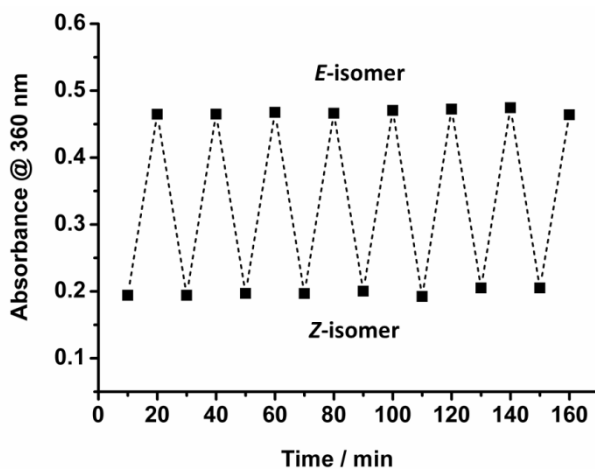


Figure 6. Repetitive photoinduced switching cycles of the **azo-borazine-3** obtained by alternating irradiations at 360 and 450 nm ($5 \cdot 10^{-6}$ M solution in toluene at 23 °C).

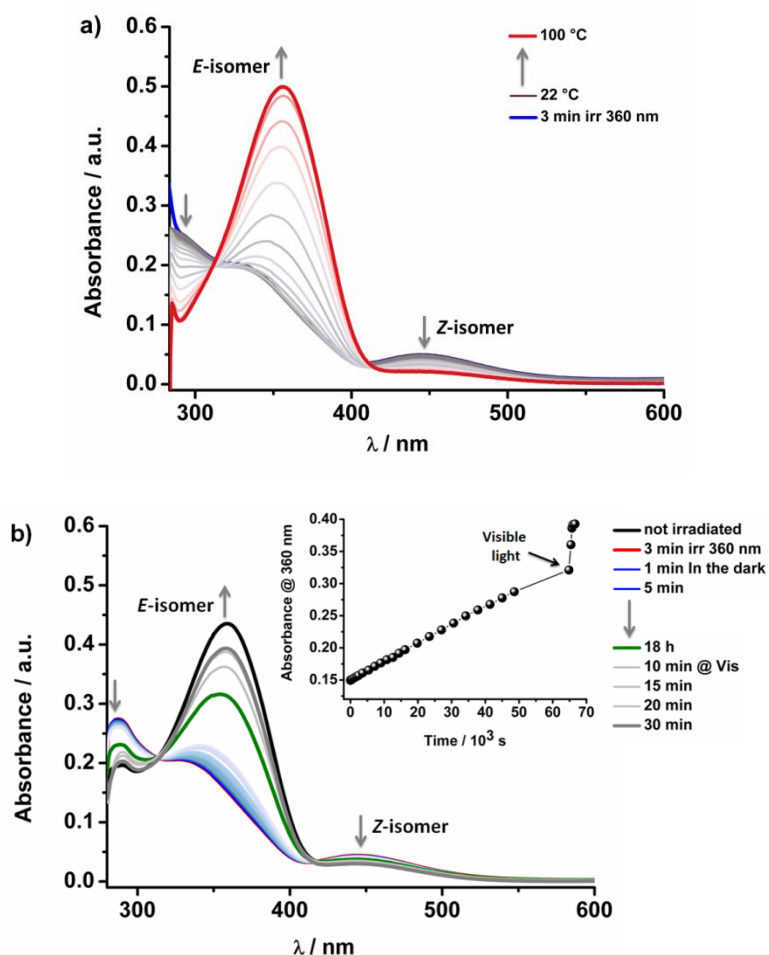


Figure 7. UV-Vis measurements of the thermal $Z \rightarrow E$ isomerization of **azo-borazine-3** after sample irradiation at 360 nm ($5 \cdot 10^{-6}$ M in toluene) (a) and at r.t. in the dark (b).

Self-assembly behavior of azo-borazine-3. To investigate the aggregation properties of **azo-borazine-3**, a THF/water mixture solvent was used to trigger the self-assembly into nanoparticles. Particles were formed by

dissolving the **azo-borazine-3** in THF containing various fractions (f_w) of water at a concentration of $4.6 \cdot 10^{-6}$ M under sonication at 40 °C for 15 min. As shown in **Figure 8a**, in pure THF the absorption spectra of the **azo-borazine-3** consist of a strong absorption peak at 360 nm and a very weak band around 450 nm attributed to $\pi-\pi^*$ and $n-\pi^*$ electronic transitions of the azobenzene unit, respectively. Upon increasing the water fraction, the absorption spectra broadened and showed a hypochromic effect in intensity, suggesting the formation of aggregates.²⁸ Similarly, emission measurements were recorded for the **azo-borazine-3** under excitation at 360 nm (**Fig. 8b**). **Azo-borazine-3** in THF exhibits an emission maximum at 406 nm with a fluorescence quantum yield (Φ_f) of 0.82%. An appreciable enhancement of the emission intensity was observed with increasing fractions of water, with the maximum being at 406 nm ($\Phi_f = 3.28\%$, $f_w = 95\%$). In these compounds, intramolecular rotations play a crucial role to populate the non-radiative decay channels for the excited states and enhance their rates.²⁹ The free intramolecular torsional motion in solution favours the non-radiative decays that are the responsible for the weak emission in THF.³⁰ However, these torsional motions are hindered in the aggregated state. The enhancement of the emission in the aggregated state also suggested a suppression of the concentration quenching in the aggregated state by amplified intermolecular distances resulting from the partially distorted structure of the fluorogenic units.³¹

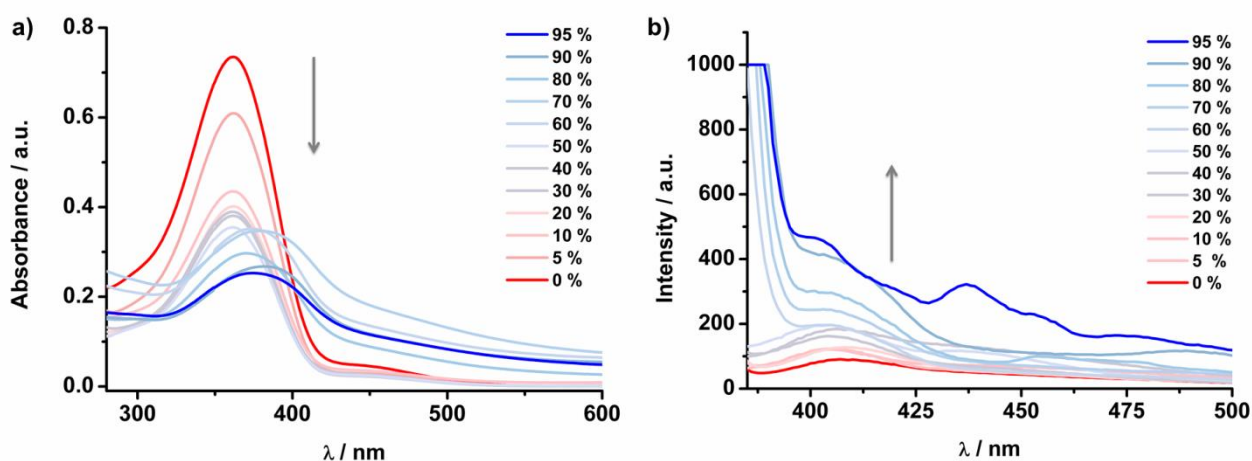


Figure 8. (a) Absorption and (b) fluorescence ($\lambda_{\text{excitation}} = 360$ nm) spectra of the **azo-borazine-3** in THF with increasing water contents (f_w %) recorded at r.t. (The final concentration was $4.6 \cdot 10^{-6}$ M).

Then, we investigated the morphology of the as-formed aggregates using transmission electron microscope (TEM). Spherical aggregates with different diameters from 90 nm to ~ 600 nm were obtained for samples containing **azo-borazine-3** prepared in a THF/H₂O mixture ($f_w = 95\%$) (**Fig. 9a-d**). Notably, some of the particles stick together to form large aggregates (**Fig. 9e**). Dynamic light scattering (DLS) measurements (**Fig. 9f**) performed on the same sample confirmed the presence of aggregates displaying hydrodynamic diameters ranging from 110 ± 20 nm to 295 ± 150 nm, together with larger particles with hydrodynamic diameters of 4 ± 1 μm . Since the formation of the nanoparticles involves stepwise addition of a poor solvophobic solvent (H₂O) to a stable solution of the **azo-borazine-3** in THF, the solubility of the molecule is gradually reduced upon addition of water, leading to the formation of aggregates. Larger nanoparticles are formed either through attractive *VdW* or dipolar forces. As previously suggested by *Ajayaghosh* and *co-workers* for azobenzene-based nanodots,^{3g} the growth of these aggregates could also be attributed to a light-driven “*Ostwald ripening*”-type mechanism.³²

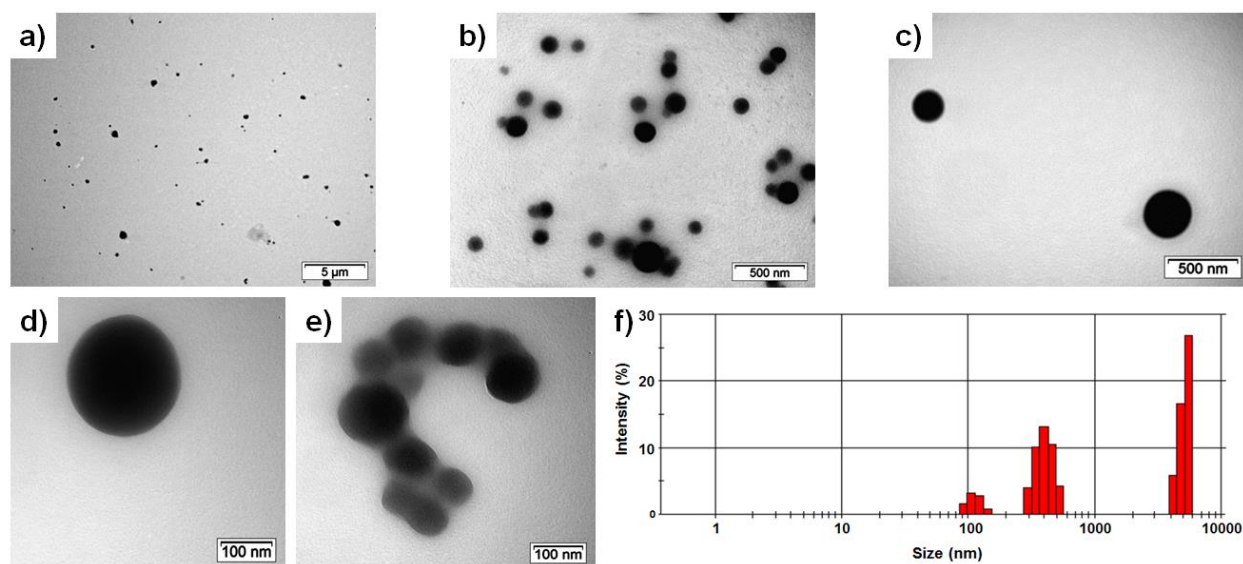
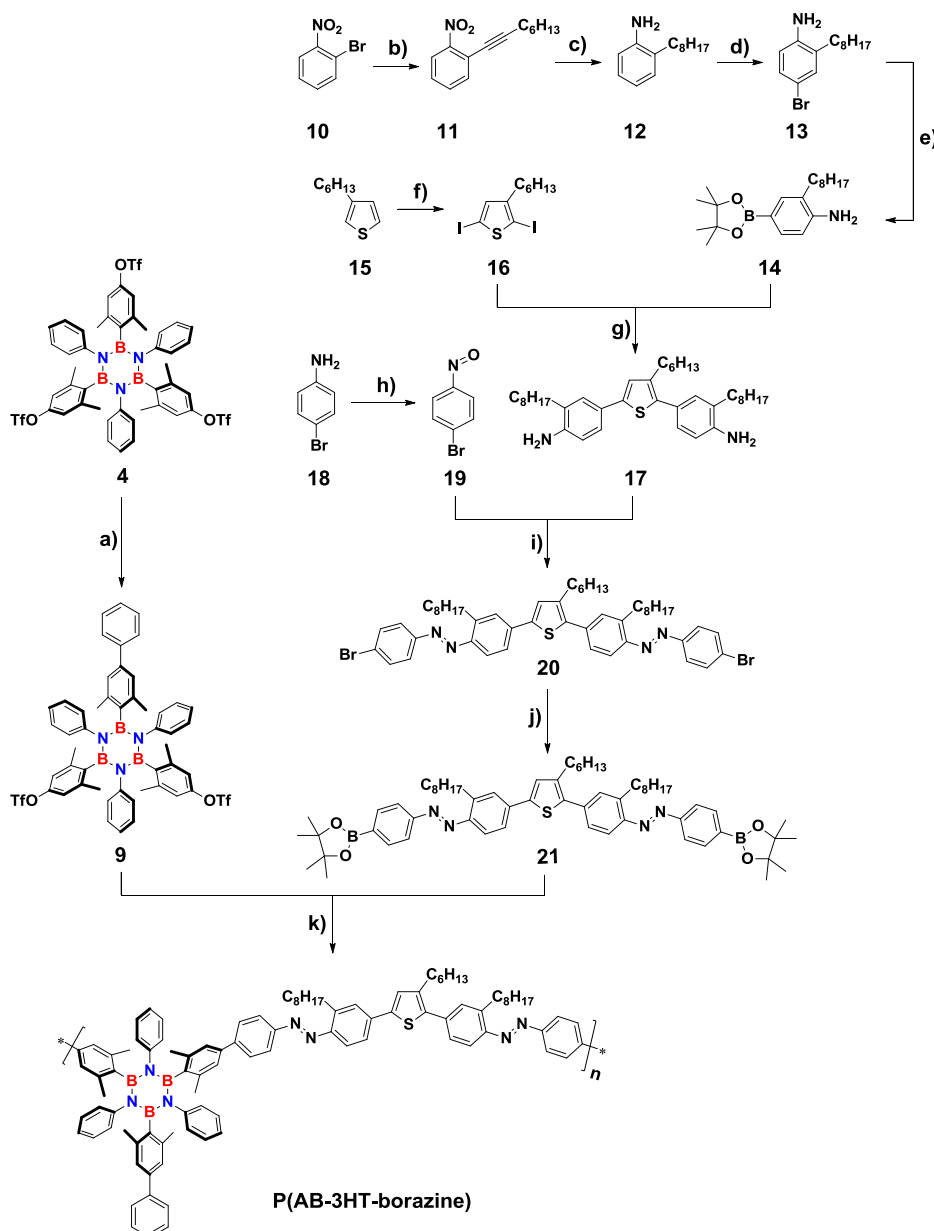


Figure 9. (a-e) TEM images of the **azo-borazine-3** prepared from THF solution with 95% of water and (f) average size of the particles as measured by dynamic light scattering ($4.6 \cdot 10^{-6}$ M, $f_w = 95\%$).

Synthesis of the borazine-doped polythiophene-based polymer. For the application of azo-borazine as building block in polymers main chain, the key issue is to introduce reactive groups on repetitive units. Therefore, we synthesized *bis*-triflate borazine derivative (*bis*-OTf-borazine) **9** starting from the *tris*-OTf-borazine **4** and azobenzene-functionalized thiophene derivative ((*1E,1'E*)-2,2'-((3-hexylthiophene-2,5-diyl)bis(2-octyl-4,1-phenylene))bis(*1*-(4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)phenyl)diazene) (AB-3HT-(BPin)₂) **21** as monomers. *Bis*-OTf-borazine **9** was prepared in 30% yield via *Suzuki* cross-coupling reaction between *tris*-OTf-borazine **4** and phenylboronic acid (**Scheme 2**, reaction a). In parallel, AB-3HT-(BPin)₂ **21** monomer was prepared starting from *1*-bromo-2-nitrobenzene **10** and *1*-octyne through *Sonogashira* cross-coupling reaction leading to *1*-nitro-2-(oct-1-yn-1-yl)-benzene **11** in 90% yield (**Scheme 2**, reaction b). Compound **11** was quantitatively reduced using Pd-catalyzed hydrogenation yielding compound **12** (**Scheme 2**, reaction c). A regio-selective bromination of **12** by pyridinium hydrobromide perbromide gave 4-bromo-2-octylaniline **13** in 71% yield (**Scheme 2**, reaction d). *Miyaura* borylation of compound **13** in *1,4*-dioxane provided 2-octyl-4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)aniline **14** in 50% yield (**Scheme 2**, reaction e). Compound **14** was then coupled to 3-hexyl-2,5-diiodothiophene **16**, prepared in 90% yield from 3-hexylthiophene **15** (**Scheme 2**, reaction f), via *Suzuki* cross-coupling reaction yielding (4,4'-((3-hexylthiophene-2,5-diyl)bis(2-octylaniline)) **17** in 30% yield (**Scheme 2**, reaction g). Compound **17** was reacted with *1*-bromo-4-nitrosobenzene **19**, prepared from 4-bromoaniline **18** (**Scheme 2**, reaction h), using *Mills* reaction conditions³³ giving access to (*1E,1'E*)-2,2'-((3-hexylthiophene-2,5-diyl)bis(2-octyl-4,1-phenylene))bis(*1*-(4-bromophenyl)diazene) AB-3HT-Br₂ **20** in 45% yield (**Scheme 2**, reaction i). *Miyaura* borylation of compound **20** led to the desired monomer AB-3HT-(BPin)₂ **21** in 65% yield (**Scheme 2**, reaction j). Finally, the synthesis of the **P(AB-3HT-borazine)** polymer was carried out using a standard *Suzuki* cross-coupling polycondensation reaction (**Scheme 2**, reaction k). The molecular weight estimated by gel permeation chromatography (GPC) relative to polystyrene standard in THF (+2% Et₃N) at 30 °C was 13.2 kDa with a dispersity (\bar{M}_w) of 1.92 and a degree of polymerization (DP) of 9.



Scheme 2. Synthetic routes for preparing **P(AB-3HT-borazine)** polymer. Reagents and conditions: (a) Phenylboronic acid, $[\text{Pd}(\text{PPh}_3)_4]$, K_2CO_3 , 1,4-dioxane:water (3:1 v/v), 100 °C, 8 h, 30 %; (b) 1-octyne, $[\text{Pd}(\text{PPh}_3)_4]$, CuBr, Et_3N , 90 °C, 15 h 90%; (c) H_2 (1 atm), Pd/C (10 wt.%), EtOH, r.t., 6 h, 95%; (d) pyridinium hydrobromide perbromide, THF, r.t., 1 h 45, 71%; (e) B_2Pin_2 , $[\text{Pd}(\text{dppf})\text{Cl}_2 \cdot \text{CH}_2\text{Cl}_2]$, KOAc, 1,4-dioxane, 90 °C, 24 h, 50%; (f) NIS, AcOH: CHCl_3 (3:4 v/v), r.t., 24 h, 90%; (g) $[\text{Pd}(\text{PPh}_3)_4]$, K_2CO_3 , 1,4-dioxane: H_2O (3:1), 105 °C, 24 h, 30%; (h) oxone, $\text{CH}_2\text{Cl}_2 \cdot \text{H}_2\text{O}$ (1:3 v/v), r.t., 3 h, quant.; (i) AcOH:EtOAc (1:1 v/v), 40 °C, 36 h, 45%; (j) B_2Pin_2 , $[\text{Pd}(\text{dppf})\text{Cl}_2 \cdot \text{CH}_2\text{Cl}_2]$, AcOK, 1,4-dioxane, 90 °C, 24 h, 65%; (k) $[\text{Pd}(\text{PPh}_3)_4]$, K_2CO_3 , toluene:water (2:0.2 v/v), 100 °C, 48 h, 60%. NIS: *N*-iodosuccinimide.

The polymer is soluble in common organic solvents such as CH_2Cl_2 , chloroform, THF, toluene and insoluble in acetone, methanol and *n*-heptane. ^1H NMR spectrum of **P(AB-3HT-borazine)** polymer in CDCl_3 (**Fig. 10**) showed all characteristic peaks of its repeating units. The low-field shifting of the aromatic proton resonance of the (OTf)-aryl group of the borazine unit from 6.49 ppm in the starting material (*bis*-OTf-borazine) to 6.92 ppm in the polymer structure, suggests that the desired polymer has been formed. As revealed by thermogravimetric analysis (TGA) (Supporting Information, **Fig. S25**), **P(AB-3HT-borazine)** polymer exhibits good thermal stability with a mass loss of 5 wt.% starting at 203 °C, and a loss of about 30 wt.% at 351 °C. On the other hands, DSC analysis performed from -25 to 200 °C showed a glass transition (T_g) at about 112 °C. Moreover, no melting

and crystallization peaks were detected, suggesting the amorphous nature of the polymer (Supporting Information, Fig. S26).

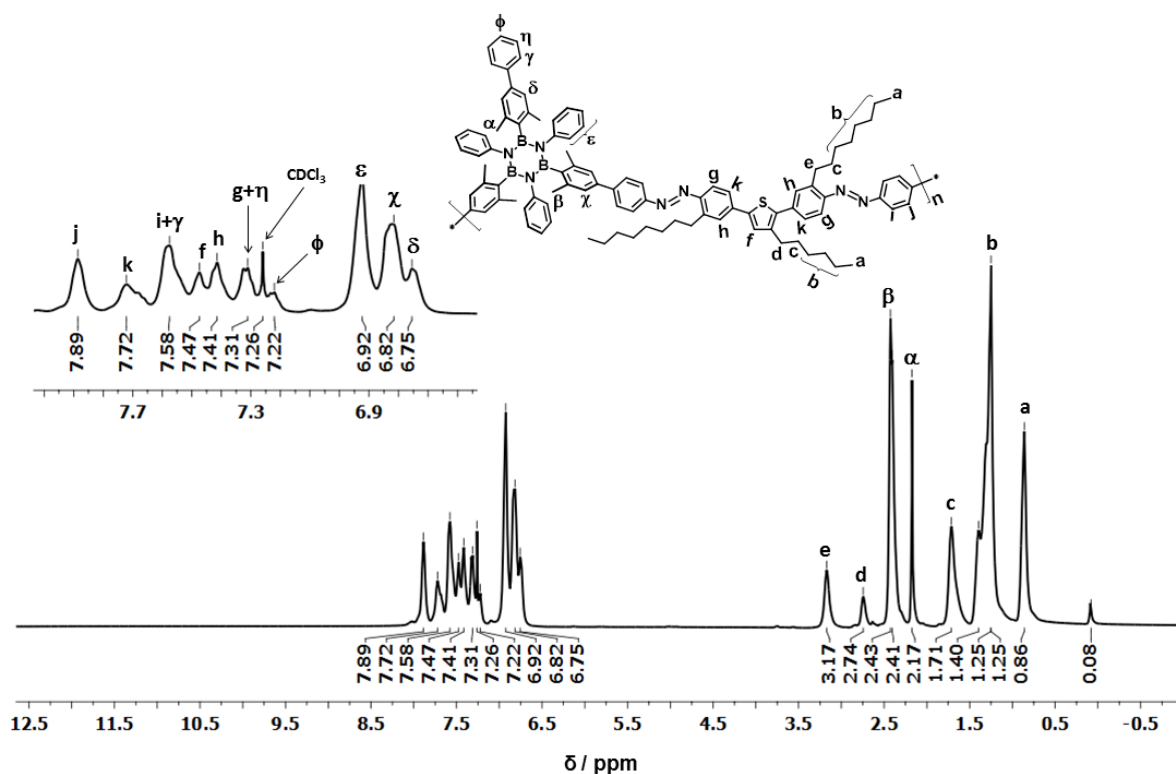


Figure 10. ^1H NMR spectrum of **P(AB-3HT-borazine)** polymer (CDCl_3 , 500 MHz, 25 $^\circ\text{C}$).

The absorption spectra of monomers *bis*-OTf-borazine **9**, AB-3HT-(BPIn) $_2$ **21** as well as **P(AB-3HT-borazine)** polymer in solutions and in thin films are depicted in **Figure 11**.

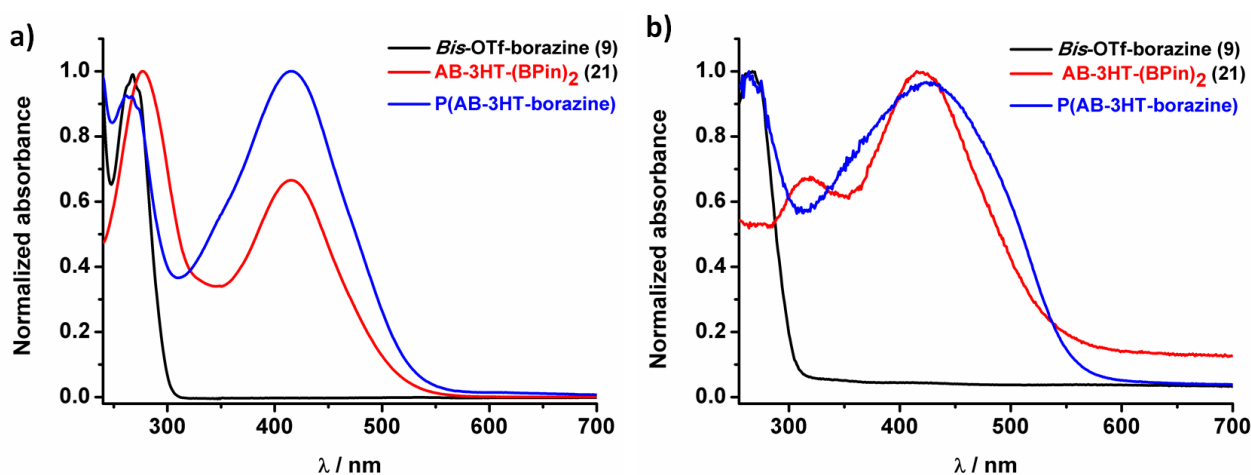


Figure 11. Normalized absorption spectra of monomers **9**, **21** and **P(AB-3HT-borazine)** polymer in CHCl_3 solutions (**a**) and in thin films (**b**).

Absorption spectra of borazine unit either in solution or in film gave predominately one band at 268 nm and 266 nm, respectively, corresponding to S_0 - S_1 transition. Absorption spectra of thiophene derivative unit **29** in solution showed two absorption maxima at 276 nm and 416 nm, corresponding to the S_0 - S_2 and S_0 - S_1 transitions, respectively. However, these bands are bathochromic shifted in the solid state. The **P(AB-3HT-borazine)**

polymer showed dual absorption bands. The high-energy band at around 268 nm develops from overlapped maximum spectra of S_0 - S_2 transitions of the borazine and the thiophene moieties, while, the low-energy band (320-550 nm) is ascribed to the absorption of the thiophene derivative unit. Furthermore, no other bands have been appeared at high wavelengths suggesting that no separate charge transfer occurred at the ground state. The band gap (E_g^{opt}) of **P(AB-3HT-borazine)** polymer calculated from the absorption onsets (548 nm) in the solid state is 2.27 eV, which is close to that of AB-3HT-(BPIn)₂ derivative (2.30 eV). Further, we have recorded emission spectra of the monomers (**9** and **21**) and the polymer (**P(AB-3HT-borazine)**) in CHCl₃ (λ_{exc} = 268 nm, 276 nm and 416 nm, **Fig. 12**). A well-resolved peak at 330 nm corresponding to the S_1 - S_0 transition was observed when the borazine unit **9** is excited at 268 nm, whereas thiophene monomer **21** predominately gave two emission maxima at 400 nm and 460 nm (λ_{exc} = 276 nm), attributing to S_2 - S_0 and S_1 - S_0 electronic transitions, respectively. An emission maximum was obtained at 475 nm when excited at 416 nm, corresponding to S_1 - S_0 electronic transition. On the other hand, broad emission spectra were obtained when the polymer is excited at 268 nm and 276 nm, with a maximum at 330 nm. The broad nature of the spectra obtained for the polymer is attributed to the contribution of the borazine and the thiophene moieties. The excitation of polymer at 416 nm gave a weakly-emissive band between 560 nm and 550 nm. In addition to the broad nature of the fluorescence spectra, the emission intensities are decreased compared to those of the reference borazine and thiophene derivatives alone, suggesting the occurrence of an intramolecular charge-transfer (ICT).

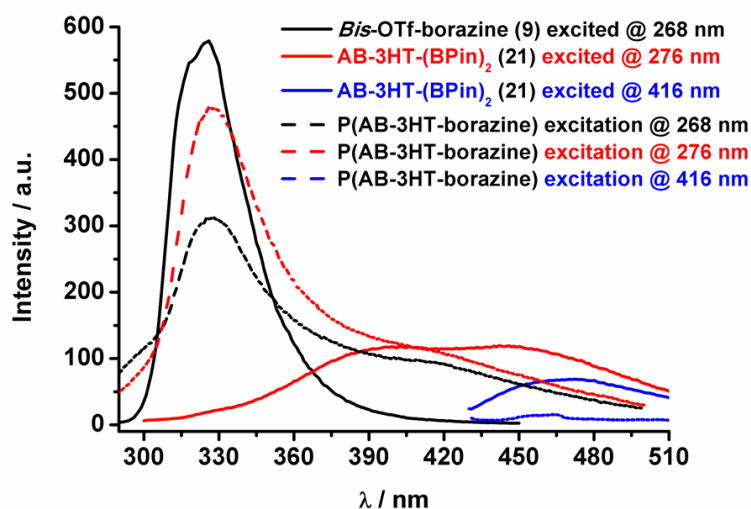


Figure 12. Emission spectra of *bis*-OTf-borazine **9**, AB-3HT-(BPIn)₂ **21** and **P(AB-3HT-borazine)** polymer recorded at r.t. in CHCl₃ (λ_{exc} = 268 nm, 276 nm and 416 nm).

The photo-switching properties of the obtained **P(AB-3HT-borazine)** polymer was studied in CHCl₃ (0.025 g/L) at r.t. When irradiated at the maximum absorption band (440 nm) no significant changes in the absorption band were noticed (**Fig. 13**), suggesting that the polymer does not undergo any photo-induced switching process. Considering that the band corresponding to the π - π^* transition of the azo-group around 360 nm is veiled by the intense S_0 - S_1 transition band of the thiophene derivative unit, one can hypothesize that the lack of a selective excitation of the azobenzene moieties is the main reason for the unsuccessful polymer switching. Also, it cannot be excluded that an energy transfer from the excited state of the azobenzene to the oligothiophene could occur as well, quenching the photoisomerization reaction.

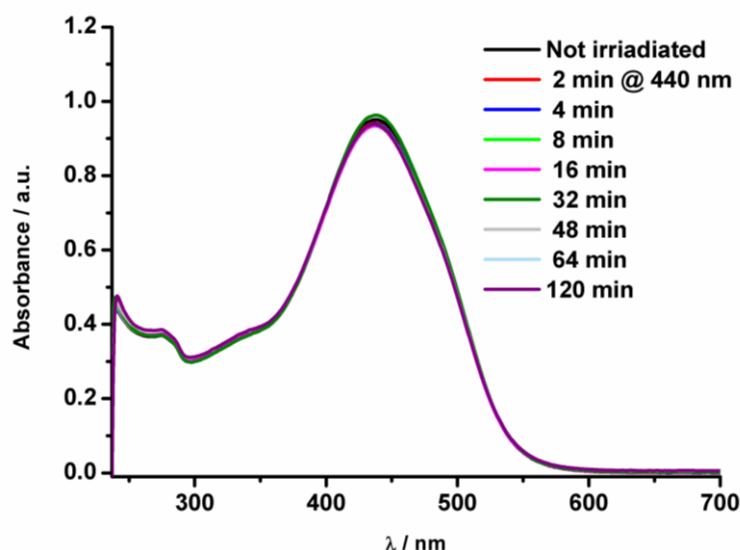


Figure 13. UV change in the absorption spectra of a 0.025 g/L solution of **P(AB-3HT-borazine)** polymer in CHCl_3 under irradiation at 440 nm at r.t.

The good solubility of the **P(AB-3HT-borazine)** polymer in a variety of organic solvents allowed a detailed study of its solvatochromic behavior. Thus, in order to get insight into the ICT nature of the **P(AB-3HT-borazine)** polymer, the solvent-dependent absorption and emission profiles (**Fig. 14**) were recorded in cyclohexane (dielectric permittivity $\epsilon = 2.02$), toluene ($\epsilon = 2.38$), THF ($\epsilon = 7.58$), CHCl_3 ($\epsilon = 4.81$), MeOH ($\epsilon = 32.7$), MeCN ($\epsilon = 37.5$) and DMSO ($\epsilon = 46.7$). A slight red shift was observed in the absorption band maxima accompanied by a broadening of the spectra when passing from nonpolar to polar solvents (**Fig. 14a**).³⁴ However, changes in the emission spectra are more pronounced (**Fig. 14b**), with the polymer emitting at 660 nm and 720 nm. It is worthy to note that no significant differences were found between protic and aprotic solvents, suggesting that the hydrogen-bonding interaction is not responsible for the red shift. The significant solvent-induced fluorescence quenching could be explained either by an excited-state photo-induced intramolecular charge-transfer (ICT) or by internal energy transfer (IET) or even both phenomena taking place between the borazine and thiophene units.

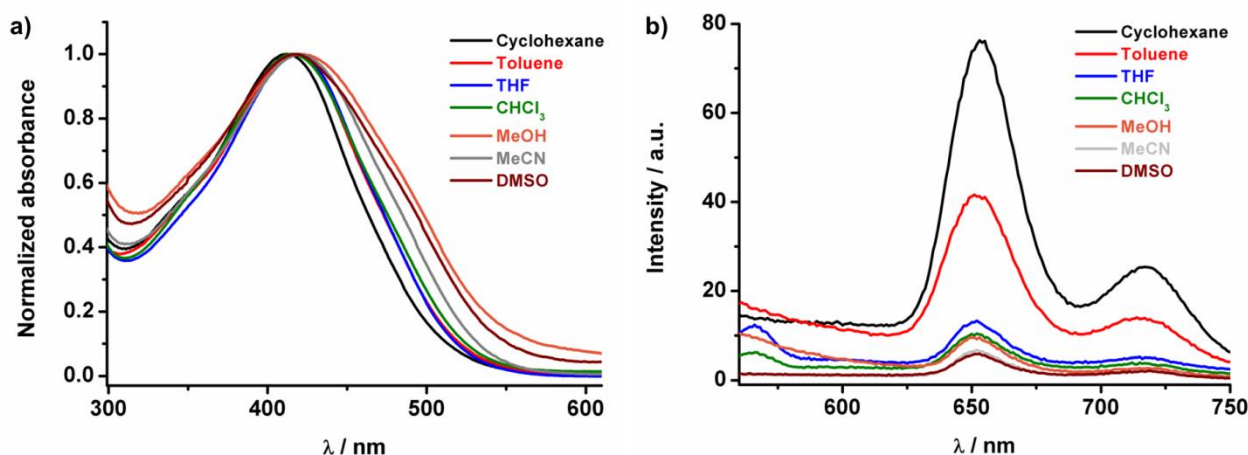


Figure 14. Absorption (a) and emission (b) spectra of the **P(AB-3HT-borazine)** polymer in various solvents of different polarities ($\lambda_{\text{exc}} = 416 \text{ nm}$).

Cyclic voltammetry (CV) was used to investigate the electrochemical properties of the polymer **P(AB-3HT-borazine)**, along with that of its monomeric precursors *bis*-OTf-borazine **9** and AB-3HT-(BPin)₂ **21** in a solution of *n*-Bu₄NPF₆ (0.2 M) in CH₂Cl₂. As shown in **Figure 15**, the oxidation of borazine **9** is observed at a more anodic potential than that of the polymer and the thiophene **21**. Its oxidation peak is observed around 1.6 V and its HOMO is determined at -5.8 eV from its onset oxidation potential $E_{\text{onset}}^{\text{ox}}$ measured at 1.4 V. No reduction was observed before the reduction of the electrolytic medium, suggesting that this borazine has a LUMO energy level higher than -1.9 eV. The CV recorded for the AB-3HT-(BPin)₂ **21**, presents reversible oxidation and reduction processes with E_{ox}^1 at 1.3 V and E_{red}^1 at -1.39 V. From the $E_{\text{onset}}^{\text{ox}} / E_{\text{onset}}^{\text{red}}$, the HOMO and LUMO energy levels of AB-3HT-(BPin)₂ **21** were determined to be at -5.58 eV and -3.17 eV, respectively, resulting in an electrochemical bandgap of 2.41 eV. The CV trace of the **P(AB-3HT-borazine)** polymer exhibited a weakly reversible oxidation (at a lowest potential value than that of the two precursors) and an irreversible reduction wave with E_{ox}^1 and E_{red}^1 at 1.24 V and -1.78 V, respectively. The HOMO and LUMO energy levels were calculated to be -5.36 eV and -3.08 eV, respectively, from their onset oxidation/reduction potential values. The electrochemical band gap was calculated to be 2.28 eV, which is slightly lower than that observed for the thiophene precursors, indicating a higher conjugation length in the polymer (see also the optical band gap of 2.27 eV).

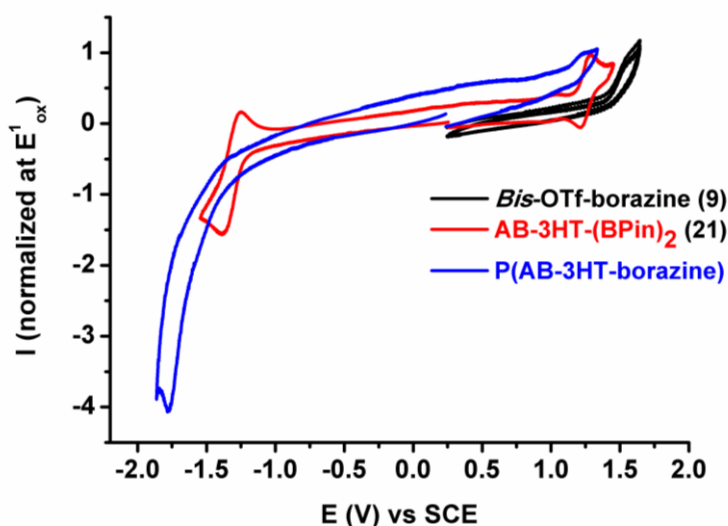


Figure 15. Cyclic voltammograms (CV) recorded in CH₂Cl₂, *n*-Bu₄NPF₆ (0.2 M), using a platinum disk working electrode (\varnothing 1 mm) at a scan rate of 100 mV/s, Al₂O₃ was added to the electrolytic medium to remove water traces. CV traces of *bis*-OTf-borazine **9** in black, AB-3HT-(BPin)₂ **21** in red and polymer **P(AB-3HT-borazine)** in blue.

CONCLUSIONS

Owing to the possibility of functionalizing borazine cores with different groups on the aryl substituents at the *N* and *B* atoms, in this paper we reported the use of the borazine core as a modular scaffold to prepare photoactive hybrid BNC by means of the Pd-catalysed *Suzuki* cross-coupling reaction from the versatile *tris*-OTf-borazine derivative (**4**). Following this approach, we developed molecular BNC hybrid molecular scaffolds that, decorated with three azobenzene moieties, displayed reversible *E/Z* photoisomerization. The chemical structure was unveiled by X-ray diffraction analysis. The photoswitching properties of **azo-borazine-3** have been investigated by means of UV/VIS spectroscopy and exhibits efficient reversible *E/Z* isomerization in toluene. To further

explore the chemical versatility of the borazine core, the new BN-doping polythiophene-type polymer **P(AB-3HT-borazine)** incorporating the borazine core, azobenzene and 3HT skeletons was also developed through *Suzuki*-type cross-coupling polycondensation reaction. Thermal, optical and electrochemical properties were investigated. In particular, emission studies in solvents of different polarity, suggested that the polymer undergoes intramolecular charge-transfer (ICT). This work opens the gate of applying the rapidly developing borazine chemistry as a new toolbox for engineering functional materials with tailored and exploitable optoelectronic properties.

EXPERIMENTAL SECTION

General Experimental Methods. Adsorption silica chromatography columns (SCC) were performed using Merck Gerduran silica gel 60 (particle size 40 - 60 μm). NMR spectra (^1H , ^{13}C NMR, ^{11}B and ^{19}F NMR) were recorded on a 400 MHz (Jeol JNM EX-400) or 500 MHz NMR (Jeol JNM ECZR-500) spectrometer. Chemical shifts are reported in parts per million (ppm) using the solvent residual signal as an internal reference (CDCl_3 $\delta_{\text{H}} = 7.26$ ppm, $\delta_{\text{C}} = 77.16$ ppm; CD_2Cl_2 $\delta_{\text{H}} = 5.32$ ppm, $\delta_{\text{C}} = 54.00$ ppm; $(\text{CD}_3)_2\text{SO}$ $\delta_{\text{H}} = 2.50$ ppm, $\delta_{\text{C}} = 39.52$ ppm). For ^{11}B , chemical shifts are reported in ppm downfield from $\text{BF}_3 \cdot \text{OEt}_2$ as internal reference, and analyses were performed in quartz tubes. Resonance multiplicity is described as s (singlet), d (doublet), t (triplet), q (quartet), quin (quintet), m (multiplet), br (broad signal), dd (doublet of doublets). Carbon spectra were acquired with a complete decoupling for the proton. All spectra were recorded at 25 $^\circ\text{C}$ unless specified otherwise. Infrared spectra (IR) were recorded on a Perkin-Elmer Spectrum II FTIR System UATR, mounted with a diamond crystal. Selected absorption bands are reported in wavenumber (cm^{-1}). (i) ESI mass spectra (HRMS) were performed on a Waters QToF2 or on a Bruker maXis Q-ToF mass spectrometer operating in positive mode at Centre de spectrométrie de masse of University of Mons and the “Fédération de Recherche” ICOA/CBM (FR2708) platform of Orléans in France. (ii) High-resolution MALDI mass spectra (HRMS) were performed on a Waters QToF Premier mass spectrometer at Centre de spectrométrie de masse of University of Mons. Melting points (Mp) were measured on a Büchi Melting Point B-545 apparatus in an open capillary and have not been corrected. Light irradiation experiments were performed with monochromatic light produced with a LOT Oriel equipment, consisting of a 1000 W Xe lamp (LSB551 ozone free) connected via an axial optical path to a computer-controlled monochromator MSH-300. The 180° optical path was composed of a condenser lens, which collimated the light emitted from the source, and a focusing lens aligned to the monochromator entrance slit. The entrance and the exit slits of the monochromator were adjusted in order to have a 20 nm bandwidth output light. The sample was placed in the collimated beam of the monochromatic light. UV-vis absorption spectra were recorded on a Varian Cary 5000 UV-Vis-NIR spectrophotometer, using quartz cells with a path length of 1.0 cm. Emission spectra were recorded on a Cary-Varian Eclipse fluorescence spectrophotometer recorded using quartz cuvettes. All photophysical experiments were carried out in air-equilibrated solutions otherwise stated. Data collections were performed at the X-ray diffraction beamline (XRD1) of the Elettra Synchrotron, Trieste (Italy)³⁵, using a Pilatus 2M hybrid-pixel area detector (DECTRIS Ltd., Baden-Daettwil, Switzerland). Complete datasets were collected at 100 K (nitrogen stream supplied through an Oxford Cryostream 700) with a monochromatic wavelength of 0.700 Å through the rotating crystal method. The crystals of compounds were dipped in N-paratone and mounted on the goniometer head with kapton loops (MiTeGen, Ithaca, USA). Complete datasets for **4** and **3** have been obtained

merging two different data collections done on the same crystal, mounted with different orientations. The diffraction data were indexed, integrated and scaled using XDS.³⁶ The structures were solved by direct methods implemented in the Sir2014 code.³⁷ Fourier analysis and refinement were performed by the full-matrix least-squares based on F^2 implemented in SHELXL-2018/3.³⁸ The Coot program was used for modeling.³⁹ Anisotropic thermal motion was then applied to all atoms with occupancy greater than 75%. Hydrogen atoms were included at calculated positions with isotropic $U_{\text{factors}} = 1.2 \cdot U_{\text{eq}}$ or $U_{\text{factors}} = 1.5 \cdot U_{\text{eq}}$ for methyl groups (U_{eq} being the equivalent isotropic thermal factor of the bonded non hydrogen atom). Restraints on bond lengths, angles and thermal motions (DFIX, DANG, SIMU and DELU) have been applied on disordered fragments. Essential crystal and refinement data are reported in the supporting information document (**Table S1**). Size exclusion chromatography (SEC or GPC) was performed in THF (with 2% triethylamine added) at 35 °C using a Polymer Laboratories liquid chromatograph equipped with a PL-DG802 degasser, an isocratic HPLC pump LC 1120 (flow rate = 1 mL.min⁻¹), a Marathon autosampler (loop volume = 200 µL, solution conc. = 1 mg.mL⁻¹), a PL-DRI refractive index detector and three columns: a PL gel 10 µm guard column and two PL gel Mixed-B 10 µm columns (linear columns for separation of MWPS ranging from 500 to 106 daltons). Poly(styrene) standards were used for calibration. Cyclic voltammetry was recorded against saturated calomel electrode (SCE) in CH₂Cl₂-*n*-Bu₄NPF₆ (0.2 M) using a platinum disk working electrode (Ø 1.0 mm) at a scan rate of 100 mV.s⁻¹. Al₂O₃ was added to the electrolytic medium to avoid any water traces. Chemicals were purchased from Sigma Aldrich, Acros Organics, Fluorochem, TCI, VWR, carbosynth, and ABCR, and were used as received, unless otherwise stated. Solvents were purchased from Sigma Aldrich, Acros Organics and VWR, and deuterated solvents were purchased from Eurisotop. THF was distilled from Na/benzophenone and stored on activated 3 Å molecular sieves 24 hours prior to use and toluene were distilled from CaH₂ and stored on activated 3 Å molecular sieves 24 hours prior to use. Aniline was purified by distillation from Sn powder, followed by distillation from Zn powder, followed by distillation from CaH₂ under inert atmosphere, yielding a transparent colorless liquid stored protected from light under argon. Anhydrous conditions, when necessary, were achieved by keeping all glassware in the oven at 120 °C overnight and then allowed to cool down under a vacuum followed by drying the two-neck flask by flaming with a heat gun under a 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 rubber stoppers used to close the flask's necks.

4-bromo-3,5-di-methylaniline (5'). To a solution of distilled *3,5-di-methylaniline 4'* (1.2 mL, 10 mmol) in anhydrous DMF (10 mL) was added dropwise a solution of *N*-bromosuccinimide (NBS) (1.8 g, 10 mmol) in anhydrous DMF (10 mL) at 0 °C. The reaction mixture was stirred at r.t. for 24 h. The crude is extracted with CH₂Cl₂, and the organic phases are dried over MgSO₄ and evaporated yielding a brown solid, purified by column chromatography (cyclohexane/EtOAc 0 to 20 %) to the desired product **5'** as a white solid (1.6 g, 80%). Mp: 72.7 – 74.1 °C. ¹H NMR (CDCl₃, 400 MHz): δ 2.32 (s, 6 H), 3.54 (s, 2 H), 6.44 (s, 2 H). ¹³C{¹H} NMR (CDCl₃, 101 MHz): δ 24.0, 115.2, 116.0, 139.0, 145.1. FTIR (UATR) ν (cm⁻¹): 3403, 3307, 2953, 2916, 1623, 1586. HRMS (EI-ToF) m/z: [M]⁺ Calcd for C₈H₁₀BrN⁺; 198.9997; Found, 198.9998.

N,N'-di-allyl-4-bromo-3,5-di-methylaniline (6') and *N,N'-di-benzyl-4-bromo-3,5-di-methylaniline (7')*. *4-bromo-3,5-dimethylaniline* (1.001 g, 5 mmol), NaHCO₃ (1.2605 g, 15 mmol) and sodium dodecyl sulfate (SDS) (20 mg) were dispersed in a mixture of EtOH and H₂O (2:5 v/v) (30 mL). R-Br (R= Allyl or R= Bn) (1 mL, 11.6

mmol) was added dropwise to the mixture. The reaction was carried out at 80 °C for 16 h. The reaction mixture was diluted with CH₂Cl₂, washed with brine and water, dried over MgSO₄ and evaporated to yield brownish oil. Column chromatography (cyclohexane) purification afforded the desired product **6'** and **7'** as transparent oil (1.34 g, 96%) and white solid (1.52 g, 80%), respectively.

N,N'-diallyl-4-bromo-3,5-dimethylaniline (**6'**). ¹H NMR (CDCl₃, 400 MHz): δ 2.43 (s, 6 H), 3.95 (m, 4 H), 5.22–5.26 (m, 4 H), 5.91 (m, 2 H), 6.52 (s, 2 H). ¹³C{¹H} NMR (CDCl₃, 101 MHz): δ 24.4, 52.8, 112.5, 114.1, 116.1, 133.8, 138.38, 147.4. FTIR (UATR) ν (cm⁻¹): 3078, 2917, 2979, 1642, 1588, 1476, 1376, 1417, 1279, 1199, 994, 918, 823, 737. HRMS (ESI-ToF) m/z: [M]⁺ Calcd for C₁₄H₁₈BrN⁺, 279.0623; Found 279.0620.

N,N'-di-benzyl-4-bromo-3,5-di-methylaniline (**7'**). Mp: 119.2–121.4 °C. ¹H NMR (CDCl₃, 400 MHz): δ 2.29 (s, 6 H), 4.59 (s, 4 H), 6.49 (s, 2 H), 7.21 (m, 4 H), 7.35 (m, 6 H). ¹³C{¹H} NMR (CDCl₃, 101 MHz): δ 24.4, 54.0, 112.5, 114.6, 126.8, 127.1, 128.8, 138.4, 138.7, 148.0. FTIR (UATR) ν (cm⁻¹): 3026.59, 2917.93, 1591.86, 1200.17, 1073.88, 1026.88. HRMS (ESI-ToF) m/z: [M]⁺ Calcd for C₂₂H₂₂BrN⁺, 379.0936; Found 379.0935.

Allyl-borazine (9') and *Benzyl-borazine (10')*. Following the general procedure for borazine synthesis⁹: aniline (0.28 mL, 3.0 mmol) was used in anhydrous toluene (8 mL) with BCl₃ (1 M in heptane, 2.7 mL, 2.64 mmol). Ar-Li is prepared as nucleophile from *N,N'*-di-allyl-4-bromo-3,5-di-methylaniline **6'** or *N,N'*-di-benzyl-4-bromo-3,5-di-methylaniline **7'** (0.681 g, 2.42 mmol) in anhydrous THF (10 mL) with *tert*-BuLi (1.7 M in pentane, 2.92 mL, 4.96 mmol). The yellow suspension was stirred at r.t. for 16 h. Solvents were removed under reduced pressure giving brown solid. Precipitation in cold MeOH yielded the desired compound Allyl-borazine **9'** and Benzyl-borazine **10'** as white solids (0.32 g, 35%) and (0.84 g, 70%), respectively.

Allyl-borazine (9'). Mp: 195.5–196.9 °C, ¹H NMR (CDCl₃, 400 MHz): δ 2.14 (s, 18 H), 3.65 (d, 12 H), 4.92–5.02 (q, 12 H), 5.65 (m, 6 H), 5.93 (s, 6 H), 6.67–6.75 (m, 15 H). ¹³C{¹H} NMR (CDCl₃, 101 MHz): δ 23.6, 52.4, 111.0, 115.9, 123.6, 126.5, 127.6, 134.7, 138.4, 146.9, 148.2. Signal for carbon atom attached to boron atom is missing due to quadrupolar relaxation. ¹¹B NMR (CDCl₃, 128 MHz): δ 36.2 (br). FTIR (UATR) ν (cm⁻¹): 3024, 2913, 1642, 1598, 1491, 1350. HRMS (MALDI-ToF) m/z: [M]⁺ Calcd for C₆₀H₆₉B₃N₆⁺, 906.5863; Found 906.5854.

Benzyl-borazine (10'). Mp: 229.8–231.8 °C. ¹H NMR (CDCl₃, 400 MHz): δ 2.15 (s, 18 H), 4.35 (s, 12 H), 6.03 (s, 6 H), 6.87 (m, 15 H), 7.02–7.04 (m, 12 H), 7.19–7.28 (m, 18 H). ¹³C{¹H} NMR (CDCl₃, 101 MHz): δ 23.7, 53.8, 110.7, 124.1, 126.9, 127.1, 127.4, 127.7, 128.67, 139.1, 139.5. Signal for carbon atom attached to boron atom is missing due to quadrupolar relaxation. ¹¹B NMR (CDCl₃, 128 MHz): δ 36.1 (br). FTIR (UATR) ν (cm⁻¹): 3057, 1600, 1492, 1394, 1351, 1273, 1202, 1170, 1028, 964, 845, 728, 697. HRMS (ESI-ToF) m/z: [M+H]⁺ Calcd for C₈₄H₈₂B₃N₆⁺, 1207.6875; Found 1207.6841.

Tri-amino-borazine (11'). Benzyl-borazine (**10'**) (797.7 mg, 0.66 mmol) was dissolved in a mixture of CH₂Cl₂ and MeOH (85:15 v/v) (30 mL). Pd(OH)₂/C (20 %, 161.1 mg, 1.14 mmol) was added to the solution. The suspension was hydrogenated with H₂ (1 atm) at r.t. for 24 h. The crude was filtrated on celite pad and evaporated to a brown residue purified on neutral alumina plug (MeOH) yielding tri-amino-borazine **11'** as a white powder (0.42 g, 95 %). Mp: decomposition before melting around 150 °C. ¹H NMR (CDCl₃, 400 MHz): δ 2.19 (s, 18 H), 6.00 (s, 6 H), 6.67 (t, 3 H), 6.74 (t, 6 H), 6.86–6.88 (d, 6 H). ¹³C{¹H} NMR (CDCl₃, 101 MHz): δ 23.6, 114.8,

125.1, 127.7, 128.4, 139.6, 146.6, 148.2. Signal for carbon atom attached to boron atom is missing due to quadrupolar relaxation. ^{11}B NMR (CDCl_3 , 128 MHz): δ 36.9 (br). FTIR (UATR) ν (cm^{-1}): 2913, 1599, 1491, 1491, 1452, 1355, 1307, 1164, 1074, 1026, 842, 746, 699. HRMS (ESI-ToF) m/z : $[\text{M}]^+$ Calcd for $\text{C}_{42}\text{H}_{45}\text{B}_3\text{N}_6^+$, 666.3985; Found 666.4021.

(4-bromo-3,5-dimethylphenoxy)(tert-butyl)dimethylsilane (2). 4-bromo-3,5-dimethylphenol (6.28 g, 31.20 mmol) and imidazole (6.35 g, 93.3 mmol) were dissolved in DMF (80 mL) and *tert*-butyldimethylsilyl chloride (TBDMSCl) (6.6 g, 43.68 mmol) was added. The reaction was carried out at 50 °C for 16 h. The reaction mixture was cooled down to r.t. and extracted with CH_2Cl_2 (3 x 60 mL). The organic phase was dried over MgSO_4 and evaporated under reduced pressure at 70 °C to remove residues of TBDMSCl yielding molecule **2** as translucent oil (9.25 g, 94%). ^1H NMR (CDCl_3 , 400 MHz): δ 0.21 (s, 6 H), 1.00 (s, 9 H), 2.37 (s, 6 H), 6.60 (m, 2 H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 100 MHz): δ -4.3, 18.3, 24.0, 25.8, 119.1, 120.0, 139.1, 154.2. FTIR (UATR) ν (cm^{-1}): 2955, 2929, 2858, 1582, 1464, 1410, 1362, 1321, 1252, 1167, 1031, 1020, 978, 834, 779, 696, 673. HRMS (ESI-ToF) m/z : $[\text{M}+\text{H}]^+$ Calcd for $\text{C}_{14}\text{H}_{24}\text{BrOSi}^+$, 315.0774; Found 315.0771. Characterization is in accordance with data reported in the literature.¹⁰

Tri-B-(2,6-dimethyl-4-(O-TBDMS)-phenyl)-tri-N-phenyl-borazine (3). Aniline (0.61 mL, 6.6 mmol) was used in anhydrous toluene (8 mL) with BCl_3 (1 M in heptane, 8 mL, 7.92 mmol). Ar-Li is prepared as nucleophile from 4-bromo-3,5-dimethylphenoxy(*tert*-butyl)dimethylsilane **2** (2.46 g, 7.8 mmol) in anhydrous THF (20 mL) with *tert*-BuLi (1.7 M in pentane, 10.1 mL, 17.16 mmol). The yellow suspension was stirred at r.t. for 16 h. Solvents were removed under reduced pressure affording pale yellow solid. The mixture was poured in water (30 mL) and extracted with CH_2Cl_2 (3 x 30 mL). The organic phase was dried over MgSO_4 and evaporated. The crude was precipitated from cold MeOH to yield molecule **3** as white powder (1.1 g, 49%). Mp: 205-207 °C. ^1H NMR (CDCl_3 , 400 MHz): δ -0.04 (s, 18 H), 0.84 (s, 27 H), 2.19 (s, 18 H), 6.06 (m, 6 H), 6.68-6.79 (m, 15 H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 100 MHz): δ -4.4, 18.3, 23.1, 25.9, 117.7, 124.12 126.8, 127.2, 139.0, 146.3, 154.6. Signal for carbon atom attached to boron atom is missing due to quadrupolar relaxation. ^{11}B NMR (CDCl_3 , 128 MHz): δ 36.4 (br). FTIR (UATR) ν (cm^{-1}): 3040, 2929, 1597, 1491, 1463, 1357, 1251, 1151, 1075, 1038, 1005, 968, 905, 866, 848, 836, 814, 736, 684. HRMS (ESI-ToF) m/z : $[\text{M}+\text{H}]^+$ Calcd for $\text{C}_{60}\text{H}_{85}\text{B}_3\text{N}_3\text{O}_3\text{Si}_3^+$, 1012.6172; Found 1012.6193. Characterization is in accordance with data reported in the literature.¹⁰

Tris-OTf-borazine (4). Tri-B-(2,6-dimethyl-4-(O-TBDMS)-phenyl)-tri-N-phenyl-borazine **3** (736.1 mg, 0.73 mmol) was dissolved in anhydrous THF (20 mL). TBAF (1 M in THF, 0.8 mL, 0.803 mmol) was added dropwise at 0 °C, a white precipitate was formed in a yellowish solution. The reaction mixture was stirred at r.t. for 2 h. The precipitate was then filtrated off and washed with THF (3 x 30 mL). The obtained solid was dried under vacuum and used in the followed step without further purification. The obtained off-white solid from the step one (478.8 mg, 0.71 mmol) was dissolved in pyridine (30 mL) and triflic anhydride (Tf_2O) (12 eq., 1.6 mL, 8.52 mmol) was added dropwise at 0 °C. The brownish solution was stirred at r.t. for 15 h. Solvents were removed under reduced pressure giving brown solid extracted with CH_2Cl_2 (3 x 50 mL). Purified by column chromatography (cyclohexane/ CH_2Cl_2 , 8:2 v/v) affording the desired compound **4** as a white powder (0.583 g, 75%). Mp: 236-238 °C. ^1H NMR (CDCl_3 , 400 MHz): δ 2.32 (s, 18 H), 6.49 (s, 6 H), 6.75-6.77 (m, 6 H), 6.80-6.82 (m, 9 H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 100 MHz): δ 23.2, 27.0, 118.1, 120.3, 125.5, 126.4, 127.5, 140.4, 144.8, 149.2. Signal for

carbon atom attached to boron atom is missing due to quadrupolar relaxation. ^{19}F NMR (CDCl_3 , 376 MHz): δ -72.76 (s). ^{11}B NMR (CDCl_3 , 128 MHz): δ 35.7 (br). FTIR (UATR): ν (cm^{-1}) 2924, 1590, 1492, 1419, 1364, 1240, 1207, 1121, 1013, 946, 845, 813, 769, 702. HRMS (ESI-ToF) m/z : $[\text{M}+\text{H}]^+$ Calcd for $\text{C}_{45}\text{H}_{40}\text{B}_3\text{F}_9\text{N}_3\text{O}_9\text{S}_3^+$, 1066.2057; Found 1066.2067. Characterization is in accordance with data reported in the literature.¹⁰

(E)-1-(4-iodophenyl)-2-phenyldiazene (**7**). Nitrosobenzene **5** (3.0 g, 28 mmol) and 4-iodoaniline **6** (6.13 g, 28.0 mmol) were dissolved in acetic acid (50 mL). The mixture was stirred at r.t. for 18 h. The reaction was quenched by water (100 mL) and the orange precipitate was filtered off and washed with water. The crude was purified by column chromatography (cyclohexane) affording the desired molecule **7** as an orange solid (7.0 g, 81%). Mp: 98.2-100.4 °C. ^1H NMR (DMSO, 400 MHz): δ 7.59-7.62 (m, 3 H), 7.67-7.69 (m, 2 H), 7.89-7.92 (m, 2 H), 7.98-8.00 (m, 2 H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 100 MHz): δ 97.8, 123.1, 124.6, 129.3, 131.5; 138.5, 152.0, 152.6. FTIR (UATR) ν (cm^{-1}): 3060, 3041, 2252, 1917, 1744, 1576, 1565, 1474, 1444, 1393, 1297, 1221, 1153, 1070, 1051, 1017, 928, 907, 834, 731, 705, 689, 627. HRMS (ESI-ToF) m/z : $[\text{M}+\text{H}]^+$ Calcd for $\text{C}_{12}\text{H}_{10}\text{N}_2\text{I}^+$, 308.9883; Found 308.9882.

(E)-1-phenyl-2-(4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)phenyl)diazene **(8)**.

(E)-1-(4-iodophenyl)-2-phenyldiazene **7** (1.6 g, 5.2 mmol), B_2Pin_2 (1.98 g, 7.8 mmol), KOAc (1.02 g, 10.4 mmol) and $[\text{Pd}(\text{dppf})\text{Cl}_2\cdot\text{CH}_2\text{Cl}_2]$ (0.1272 g, 0.156 mmol) were suspended in anhydrous DMF (degassed by 3 freeze-to-thaw cycles) (20 mL). The reaction was carried out under argon at 90 °C for 18 h. After cooling down to r.t., the reaction mixture was diluted with CH_2Cl_2 (60 mL), washed with brine and H_2O , dried over MgSO_4 and evaporated to a brown dark orange solid purified by column chromatography (cyclohexane/ CH_2Cl_2 , from 0 to 20%) yielding the desired compound **8** as an orange solid (1.3 g, 80%). Mp: 93.5-94.7 °C. ^1H NMR (CD_2Cl_2 , 400 MHz): δ 1.36 (s, 12 H), 7.51-7.56 (m, 3 H), 7.88-7.95 (m, 6 H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CD_2Cl_2 , 100 MHz): δ 25.1, 84.5, 122.3, 123.3, 129.5, 131.7, 135.9, 153.1, 154.7. ^{11}B NMR (CD_2Cl_2 , 128 MHz): δ 29.6 (s). FTIR (UATR) ν (cm^{-1}): 3063, 2982, 2926, 1602, 1569, 1502, 1467, 1444, 1397, 1357, 1332, 1302, 1274, 1140, 1088, 1000, 963, 855. HRMS (ESI-ToF) m/z : $[\text{M}+\text{H}]^+$ Calcd for $\text{C}_{18}\text{H}_{22}\text{BN}_2\text{O}_2^+$, 309.1769; Found 309.1773.

Azo-borazine-3. *Tris*-OTf-borazine **4** (295.5 mg, 0.277 mmol), K_2CO_3 (462.0 mg, 3.328 mmol), compound **8** (341.9 mg, 1.114 mmol) and $[\text{Pd}(\text{PPh}_3)_4]$ (19.2 mg, 16.6×10^{-3} mmol) were suspended in a mixture of 1,4-dioxane and water (3:1 v/v) (16 mL) (degassed with 5 freeze-pump-thaw cycles) in a flame-dried 50-mL Schlenk flask. The reaction was carried out at 105 °C for 18 h. The reaction mixture was cooled down to r.t., diluted with CH_2Cl_2 (100 mL), filtrated on celite, washed with brine (100 mL) and H_2O (3 x 100 mL). The organic phase was dried over MgSO_4 . The crude was purified by column chromatography (cyclohexane/ CH_2Cl_2 , from 0 to 10%) affording the desired compound **azo-borazine-3** as an orange solid (0.267 g, 83%). Mp: 270-272 °C. ^1H NMR (CD_2Cl_2 , 400 MHz): δ 2.47 (s, 18 H), 6.80-6.84 (t, 3 H), 6.88-6.91 (t, 6 H), 6.97 (s, 6 H), 7.03-7.05 (d, 6 H), 7.46-7.55 (m, 9 H), 7.60-7.62 (d, 6 H), 7.88-7.92 (m, 12 H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 100 MHz): δ 25.0, 84.2, 93.9, 122.1, 122.9, 123.0, 123.1, 123.1, 123.6, 128.0, 129.2, 129.2, 129.3, 131.1, 131.3, 135.8. Signal for carbon atom attached to boron atom is missing due to quadrupolar relaxation. ^{11}B -NMR (CDCl_3 , 128 MHz): δ 35.98 (br). FTIR (UATR) ν (cm^{-1}): 3039, 2915, 1598, 1546, 1490, 1454, 1422, 1353, 1305, 1292, 1223, 1154, 1072, 1011, 838, 792. HRMS (MALDI-ToF) m/z : $[\text{M}+\text{H}]^+$ Calcd for $\text{C}_{78}\text{H}_{67}\text{B}_3\text{N}_9^+$, 1162.5793; Found 1162.6010.

Bis-OTf-borazine (**9**). *Tris*-OTf-borazine **4** (200.5 mg, 0.202 mmol), K₂CO₃ (33.6 mg, 0.242 mmol), phenylboronic acid (8.2 mg, 67 x 10⁻³ mmol) and [Pd(PPh₃)₄] (7.2 mg, 6.06 x 10⁻³ mmol) were suspended in a mixture of 1,4-dioxane and water (3:1, v/v) (12 mL) (degassed with 5 freeze-pump-thaw cycles) in a flame-dried 50-mL Schlenk flask. The reaction was carried out at 105 °C for 8 h. The reaction mixture was cooled down to r.t., diluted with CH₂Cl₂ (100 mL), filtered on celite, washed with brine (100 mL) and H₂O (3 x 100 mL). The organic phase was dried over MgSO₄, evaporated to a brownish solid and purified by column chromatography (cyclohexane/CH₂Cl₂, from 0 to 10%) yielding the desired compound **9** as a white solid (0.06 g, 30%). Mp: 180.4-182.2 °C. ¹H NMR (CD₂Cl₂, 400 MHz): δ 2.33 (s, 18 H), 6.49 (s, 4 H), 6.75-6.83 (m, 17 H), 7.21 (t, 1 H), 7.30 (t, 2 H), 7.38-7.40 (d, 2 H). ¹³C{¹H} NMR (CDCl₃, 101 MHz): δ 23.3, 23.4, 118.0, 124.4, 125.1, 125.3, 126.5, 126.7, 126.8, 126.9, 127.3, 127.5, 128.6, 138.0, 139.8, 140.4, 141.2, 145.1, 145.3, 149.2. ¹⁹F NMR (CDCl₃, 376 MHz): δ -72.75. ¹¹B NMR (CDCl₃, 128 MHz): δ 35.2 (br). FTIR (UATR) ν (cm⁻¹): 3040, 2923, 2854, 1590, 1492, 1454, 1419, 1363, 1310, 1240, 1208, 1142, 1120, 1075, 1013, 945, 908, 869, 845, 794, 749, 701, 582. HRMS (MALDI-ToF) m/z: [M+H]⁺ Calcd for C₅₀H₄₅B₃N₃F₆O₆S₂⁺, 994.2969; Found 994.3002.

1-nitro-2-(oct-1-ynyl)benzene (**11**). *1*-bromo-2-nitrobenzene **10** (2.5 g, 12.5 mmol), *1*-octyne (1.73 g, 2.4 mL, 12.5 mmol), [Pd(PPh₃)₄] (288 mg, 0.25 mmol) and CuBr (73 mg, 0.5 mmol) were dissolved in a freshly distilled and degassed with 3 freeze-pump-thaw cycles Et₃N (50 mL) a flame-dried 100-mL Schlenk flask. The reaction was carried out at 90 °C under argon atmosphere for 15 h. The reaction mixture was cooled down to r.t., diluted with 300 mL of Et₂O, washed with a saturated solution of NH₄Cl (3 x 100 mL) and water (3 x 100 mL), dried over MgSO₄, and evaporated to a dark oil. The crude product was purified by column chromatography (*n*-heptane/EtOAc, from 0 to 20%) yielding the desired compound **11** as a brownish oil (2.6 g, 90%). ¹H NMR (CDCl₃, 500 MHz): δ 0.89 (t, 3 H), 1.29-1.34 (m, 4 H), 1.46 (m, 2 H), 1.62 (m, 2 H), 2.45 (t, 2 H), 7.36 (t, 1 H), 7.49 (t, 1 H), 7.54 (d, 1 H), 7.92 (d, 1 H). ¹³C{¹H} NMR (CDCl₃, 126 MHz): δ 14.1, 19.9, 22.6, 28.4, 28.6, 31.4, 76.0, 99.5, 119.4, 124.4, 127.9, 132.6, 134.8, 150.1. FTIR (UATR) ν (cm⁻¹): 2954, 2929, 2858, 2228, 1608, 1568, 1524, 1480, 1466, 1439, 1429, 1262, 851, 783, 744, 655. LRMS (EI-ToF) m/z: [M]⁺ Calcd for C₁₄H₁₇NO₂⁺, 231.1259; Found 231.131. Characterization is in accordance with data reported in the literature.⁴⁰

2-octylaniline (**12**). *1*-nitro-2-(oct-1-ynyl)-benzene **11** (2.24 g, 9.7 mmol) and Pd/C (10 %) (222.4 mg, 0.204 mmol) were suspended in EtOH (80 mL). The reaction mixture was hydrogenated with H₂ (1 atm) at r.t. for 6 h. The crude was filtrated on celite .and evaporated to brown oil (1.9 g, 95%). ¹H NMR (CDCl₃, 500 MHz): δ 0.92 (t, 3 H), 1.31-1.43 (m, 10 H), 1.64 (m, 2 H), 2.49 (t, 2 H), 3.62 (s, 2 H), 6.68-6.70 (d, 1 H), 6.76 (t, 1 H), 7.03-7.07 (m, 2 H). ¹³C{¹H} NMR (CDCl₃, 100 MHz): δ 14.2, 22.7, 28.8, 29.4, 29.6, 29.8, 31.4, 32.0, 115.6, 118.7, 126.8, 127.0, 129.4, 144.1. FTIR (UATR) ν (cm⁻¹): 3468, 3379, 3022, 2954, 2853, 1619, 1583, 1526, 1456, 1272, 1156. LRMS (ESI-ToF) m/z: [M]⁺ Calcd for C₁₄H₂₃N⁺, 205.1825; Found 205.1083. Characterization is in accordance with data reported in the literature.⁴⁰

4-bromo-2-octylaniline (**13**). A solution of pyridinium hydrobromide perbromide (2.33 g, 7.3 mmol) in THF (50 mL) was added dropwise over 50 min to 2-octylaniline **12** (1.50 g, 7.3 mmol) in THF (50 mL). After completion of the addition, the solution was stirred at r.t. for 1 h. The solution was filtered to remove the formed pyridinium hydrobromide salt. Then, the solution was treated with a saturated solution of NaHCO₃ (3 x 100 mL) to remove any excess of bromide, washed with water (3 x 100 mL), dried over MgSO₄ and evaporated to a dark oil. The

crude product was purified by column chromatography (*n*-heptane/EtOAc, from 0 to 30%) yielding the desired compound **13** as a brownish oil (1.47 g, 71%). ¹H NMR (CDCl₃, 500 MHz): δ 0.91 (t, 3 H), 1.30-1.42 (m, 10 H), 1.60 (m, 2 H), 2.43 (t, 2 H), 3.61 (s, 2 H), 6.53 (d, 1 H), 7.11-7.12 (dd, 1 H), 7.15 (d, 1 H). ¹³C{¹H} NMR (CDCl₃, 100 MHz): δ 13.2, 21.8, 27.5, 28.3, 28.6, 28.7, 30.2, 31.0, 109.4, 116.0, 128.2, 128.5, 130.9, 142.2. FTIR (UATR) ν (cm⁻¹): 3475, 3387, 2953, 2853, 1619, 1488, 1409, 1278, 1077. MS (GC-MS) *m/z*: [M]⁺ Calcd for C₁₄H₂₂BrN⁺, 283.0936; Found 283.12. Characterization is in accordance with data reported in the literature.⁴⁰

2-octyl-4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)aniline (**14**). 4-bromo-2-octylaniline **13** (1.32 g, 4.62 mmol), B₂Pin₂ (1.41 g, 5.54 mmol), KOAc (1.36 g, 13.86 mmol) and [Pd(dppf)Cl₂·CH₂Cl₂] (100 mg, 0.14 mmol) were suspended in 1,4-dioxane (degassed by 3 freeze-to-thaw cycles) (45 mL). The reaction was carried out at 85 °C for 16 h. After cooling down to r.t. the reaction mixture was diluted with EtOAc (100 mL), filtered over celite, washed with brine (2 x 100 mL) and H₂O (3 x 100 mL), dried over MgSO₄ and evaporated to black oil. The crude product was purified by column chromatography (EtOAc/*n*-heptane, from 0 to 20%) yielding the molecule **14** as a brownish oil (0.765 g, 50%). ¹H NMR (CDCl₃, 500 MHz): δ 0.88 (t, 3 H), 1.26-1.32 (m, 22 H) (integration 34 H is due to the presence of pinacolborane 12H), 1.61 (m, 2 H), 2.49 (t, 2 H), 3.81 (s, 2 H), 6.64-6.66 (d, 1 H), 7.49 (m, 2 H). ¹³C{¹H} NMR (CDCl₃, 101 MHz): δ 14.2, 22.8, 25.1, 29.0, 29.4, 29.6, 30.0, 31.5, 32.0, 77.0, 77.6, 77.7, 83.3, 83.6, 114.7, 125.8, 134.2, 136.5, 147.4. ¹¹B NMR (CDCl₃, 128 MHz): δ 29.4 (s). FTIR (UATR) ν (cm⁻¹): 3473, 3365, 2925, 1621, 1605, 1418, 1353, 1325, 1303, 1165, 1123, 1106, 1090, 964, 856. HRMS (ESI-ToF) *m/z*: [M+H]⁺ Calcd for C₂₀H₃₄BNO₂⁺, 332.2755; Found 332.2760. Characterization is in accordance with data reported in the literature.⁴⁰

3-Hexyl-2,5-diiodothiophene (**16**). A solution of *N*-Iodosuccinimide (NIS) (14.04 g, 62.4 mmol) in CHCl₃ (30 mL) was added dropwise to a solution of 3-hexylthiophene **15** (5.3 mL, 29.71 mmol) in a mixture of acetic acid and CHCl₃ (30:10 mL). The reaction mixture was stirred in dark at r.t. for 24 h. The reaction mixture was diluted with CH₂Cl₂ (100 mL), quenched with a saturated solution of Na₂S₂O₃ (50 mL), washed with H₂O (3 x 100 mL), dried over MgSO₄ and evaporated to light pink oil. The crude product was purified by column chromatography (cyclohexane) affording the desired compound **16** as a colorless oil (11.2 g, 90%). ¹H NMR (CDCl₃, 400 MHz): δ 0.92 (t, 3 H), 1.32 (m, 6 H), 1.55 (m, 2 H), 2.50 (t, 2 H), 6.90 (s, 1 H). ¹³C{¹H} NMR (CDCl₃, 101 MHz): δ 14.3, 22.7, 28.9, 30.0, 31.7, 32.1, 76.1, 77.2, 137.8, 149.5. FTIR (UATR) ν (cm⁻¹): 2953, 2923, 2853, 1525, 1464, 1395, 1190, 982, 907, 827, 724, 695, 649, 468. HRMS (ESI-ToF) *m/z*: [M+H]⁺ Calcd for C₁₀H₁₅I₂S⁺, 420.8978; Found 420.8961. Characterization is in accordance with data reported in the literature.⁴⁰

(4,4'-(3-hexylthiophene-2,5-diyl)bis(2-octylaniline)) (**17**).
2-octyl-4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)aniline **14** (520.1 mg, 1.6 mmol), *3-hexyl-2,5-diiodothiophene* **16** (300.2 mg, 0.71 mmol) and K₂CO₃ (790 mg, 5.72 mmol) were dissolved in a mixture of 1,4-dioxane and H₂O (9:3 v/v) (12 mL). The reaction mixture was degassed by three freeze-pump-thaw cycles. Then, [Pd(PPh₃)₄] (46.2 mg, 0.04 mmol) was added. The reaction was carried out at 105 °C for 24 h. The mixture was cooled down to r.t., diluted with CH₂Cl₂ (50 mL), filtered over celite, washed with brine (2 x 100 mL) and water (2 x 100 mL), dried over MgSO₄, evaporated and purified by column chromatography (cyclohexane/EtOAc, from 0 to 10%) yielding the desired compound **17** as yellow viscous oil (0.122 g, 30%). ¹H NMR (CDCl₃, 400 MHz): δ 0.89-0.90 (t, 9 H), 1.30-1.45 (m, 24 H), 1.66 (m, 6H), 2.52 (t, 4 H), 2.63 (t, 2 H), 3.70

(s, 4 H), 6.68 (q, 2 H), 7.03 (s, 1 H), 7.13 (s, 2 H), 7.28-7.30 (m, 2 H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 101 MHz): δ 14.2, 22.8, 22.8, 28.8, 28.9, 29.2, 29.4, 29.5, 29.7, 29.7, 29.8, 29.9, 31.2, 31.4, 31.5, 31.9, 32.0, 115.5, 115.9, 123.8, 124.4, 125.4, 125.8, 126.9, 127.0, 127.3, 127.8, 130.4, 136.5, 138.4, 141.9, 143.4, 143.6. FTIR (UATR) ν (cm^{-1}): 3380, 2954, 2923, 2853, 1619, 1497, 1464, 1377, 1283, 1156, 907, 816, 732, 646, 532. LRMS (ESI-ToF) m/z : $[\text{M}+\text{H}]^+$ Calcd for $\text{C}_{38}\text{H}_{59}\text{N}_2\text{S}^+$, 575.4393; Found 575.4331. Characterization is in accordance with data reported in the literature.⁴⁰

l-bromo-4-nitrosobenzene (**19**). A solution of Oxone® (7.2 g, 11.7 mmol) in water (48 mL) was added to a solution of freshly recrystallized 4-bromoaniline (1.04 g, 5.84 mmol) in CH_2Cl_2 (12 mL). The mixture was stirred at r.t. for 4 h. The reaction mixture was extracted with CH_2Cl_2 (3 x 20 mL) to give the desired compound **19** as a pale green solid that was used in the next step without further purification (2.15 g, quant.). ^1H NMR (CDCl_3 , 400 MHz): δ 7.78 (s, 4 H).

(*1E,1'E*)-2,2'-((3-hexylthiophene-2,5-diyl)bis(2-octyl-4,1-phenylene))bis(1-(4-bromophenyl)diazene) (**20**).

l-bromo-4-nitrosobenzene **19** (2.14 g, 11.51 mmol) and compound **17** (1.51 g, 2.63 mmol) were dissolved in a mixture of AcOH and EtOAc (1:1 v/v) (40 mL). The reaction mixture was stirred at 45 °C for 36 h. The mixture was cooled down to r.t., diluted with CH_2Cl_2 (50 mL), washed with water (3 x 100 mL), dried over MgSO_4 , evaporated and purified by column chromatography (cyclohexane/ CH_2Cl_2 , from 0 to 30%) yielding the desired compound **20** as a viscous orange oil (1.08 g, 45%). ^1H NMR (CDCl_3 , 500 MHz): δ 0.87 (t, 9 H), 1.25-1.42 (m, 26 H), 1.71 (m, 6 H), 2.75 (t, 2 H), 3.14-3.19 (q, 4 H), 7.34 (s, 1 H), 7.39-7.41 (dd, 1 H), 7.48 (d, 1 H), 7.52-7.54 (dd, 1 H), 7.60 (d, 1 H), 7.65-7.67 (m, 4 H), 7.72-7.75 (d, 2 H), 7.79-7.82 (m, 4 H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 126 MHz): δ 14.2, 14.3, 22.8, 22.8, 29.4, 29.4, 29.6, 29.6, 29.7, 31.2, 31.6, 31.8, 31.8, 32.0, 32.3, 32.4, 115.8, 116.2, 123.8, 124.5, 124.6, 125.2, 125.3, 127.1, 127.3, 127.4, 131.1, 132.5, 132.5, 137.1, 137.7, 138.1, 141.1, 142.1, 143.8, 144.4, 149.2, 149.3, 151.9, 152.0. FTIR (UATR) ν (cm^{-1}): 2953, 2922, 2853, 1905, 1596, 1574, 1476.67, 1453, 1411, 1377, 1297, 1260, 1222, 1185, 1149, 1117, 1066, 1007, 886, 832, 722, 704. HRMS (ESI-ToF) m/z : $[\text{M}+\text{H}]^+$ Calcd for $\text{C}_{50}\text{H}_{63}\text{Br}_2\text{N}_4\text{S}^+$, 909.3135; Found 909.3126.

(*1E,1'E*)-2,2'-((3-hexylthiophene-2,5-diyl)bis(2-octyl-4,1-phenylene))bis(1-(4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)phenyl)diazene) (AB-3HT-(BPin)₂) (**21**). Compound **20** (1.00 g, 1.1 mmol), B_2Pin_2 (1.12 g, 4.41 mmol), KOAc (0.6477 g, 6.6 mmol) and $[\text{Pd}(\text{dppf})\text{Cl}_2 \cdot \text{CH}_2\text{Cl}_2]$ (54 mg, 0.066 mmol) were suspended in 1,4-dioxane (degassed by 3 freeze-to-thaw cycles) (45 mL). The reaction was carried out at 90 °C for 24 h. After cooling down to r.t. the reaction mixture was diluted with CH_2Cl_2 (100 mL), filtered over celite to remove the Pd catalyst, washed with brine (2 x 100 mL) and H_2O (3 x 100 mL), dried over MgSO_4 and evaporated to a black oil purified by column chromatography (cyclohexane/ CH_2Cl_2 , from 0 to 10%) to yield the desired compound **21** as orange viscous oil (0.72 g, 65%). ^1H NMR (CDCl_3 , 400 MHz): δ 0.87 (t, 9 H), 1.27-1.38 (m, 50 H), 1.73 (m, 6 H), 2.75 (t, 2 H), 3.18 (q, 4 H), 7.34 (s, 1 H), 7.40-7.42 (dd, 1 H), 7.49 (d, 1 H), 7.53-7.55 (dd, 1 H), 7.61 (d, 1 H), 7.73-7.76 (m, 2 H), 7.89-7.92 (m, 4 H), 7.96-7.99 (m, 4 H). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , 100 MHz): δ 14.2, 14.2, 22.8, 22.8, 25.0, 25.2, 27.0, 27.1, 29.4, 29.4, 29.4, 29.6, 29.6, 29.7, 29.7, 30.3, 31.2, 31.8, 32.0, 32.3, 32.4, 43.6, 83.6, 84.2, 84.2, 115.8, 116.1, 122.2, 123.7, 127.0, 127.3, 127.3, 131.1, 135.8, 135.8, 137.0, 137.6, 138.1, 141.0, 142.1, 143.7, 144.4, 149.5, 149.6, 155.0, 155.1. Signals for the two carbon atoms attached to boron atoms are missing due to quadrupolar relaxation. ^{11}B NMR (CDCl_3 , 128 MHz): δ 29.3 (s). FTIR (UATR) ν (cm^{-1}): 2923, 2853,

1599, 1571, 1501, 1480, 1456, 1396, 1355, 1324, 1272, 1216, 1186, 1124, 1142, 1086, 1014, 962, 886, 858, 848, 830, 736, 668, 653, 578, 540. HRMS (ESI-ToF) m/z : $[M+H]^+$ Calcd for $C_{62}H_{87}B_2N_4O_4S^+$, 1005.6668; Found 1005.6706.

***P(AB-3HT-borazine)* polymer.** Bis-OTf-borazine **9** (100.00 mg, 0.1 mmol), compound **21** (101.06 mg, 0.1 mmol), K_2CO_3 (1.66 g, 1.2 mmol) were suspended in a mixture of toluene and water (2:0.2 v/v). The reaction mixture was degassed by three freeze-pump-thaw cycles. Then, $[Pd(PPh_3)_4]$ (17.33 mg, 0.015 mmol) was added. The reaction was vigorously stirred at 100 °C for 48 h. The deep-orange reaction mixture turned gradually into brown accompanied with the precipitation of a black solid. **P(AB-3HT-borazine)** polymer was obtained by purification via Soxhlet extraction for 12 h using $CHCl_3$ followed by precipitation in MeOH. The orange-like polymer was finally dried under reduced pressure at 50 °C for 24 hours in 60% yield. Size exclusion chromatography (SEC) relative to polystyrene standard in THF (+2% Et_3N) at 30 °C: M_n = 13.2 kDa, M_w = 25.0 kDa, D_M = 1.92, DP = 9. 1H NMR ($CDCl_3$, 500 MHz): δ 0.86 (br, 9 H), 1.25-1.40 (br, 50 H), 1.71 (s, 6 H), 2.17 (s, 6 H), 2.41-2.43 (br, 12 H), 2.74 (s, 2 H), 3.17 (s, 4 H), 6.75 (s, 1 H), 6.82 (s, 4 H), 6.92 (s, 15 H), 7.22 (s, 1 H), 7.31 (br, 4 H), 7.41 (s, 2 H), 7.47 (s, 1 H), 7.58 (br, 6 H), 7.72 (br, 2 H), 7.89 (s, 4 H).

ASSOCIATED CONTENT

Supporting Information

Synthetic schemes, copies of NMR spectra and thermal characterization data. This material is available free of charge via the Internet at <http://pubs.acs.org>.

CCDC 1906843, 1906842 and 1906844 contain the supplementary crystallographic data for compounds **3**, **Azo-borazine-3** and **4**. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via <https://www.ccdc.cam.ac.uk/structures>.

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Notes

The authors declare no competing financial interest.

ACKNOWLEDGEMENT

The authors thank the Universities of Namur and Mons for the financial support to H.O. doctoral fellowship. D.B. thanks the Walloon Region for the ‘Flycoat’ and ‘TUBOLED’ POC projects, the Science Policy Office of the Belgian Federal Government (BELSPO-IAP 7/05 project) and School of Chemistry at Cardiff University. O.C. is Research Associate for the F.R.S. -FNRS.

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