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Divanillin based polyazomethines: toward bio-based and metal-free \(\pi\)-conjugated polymers

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Polyazomethines, Divanillin, Carbazole

ABSTRACT: Divanillin was synthesized in high yield and purity using Laccase from Trametes versicolor. It was then polymerized with Benzene-1,4-diamine and 2,7-diaminocarbazole to form polyazomethines. Polymerizations were performed under micro-wave irradiation and without transition-metal based catalyst. These bio-based conjugated polyazomethines present a broad fluorescence spectrum ranging from 400 to 600 nm. Depending on the co-monomer used, polyazomethines with molar masses around 10 kg/mol and with electronic gaps ranging from 2.66 to 2.85 eV were obtained. Furthermore, TD-DFT calculations were performed to corroborate experimental results.

1. Introduction.

Thanks to recent works on the selective oxidative coupling of phenolic molecules via enzymatic catalysis, a bio-based platform of lignin-derived aromatics is now available. 1,2

Among them, vanillin has a large potential as it is one of the only phenolic compounds industrially available from biorefinery. 3 Vanillin can be easily dimerized from C-C coupling using an environmental-friendly process as was demonstrated by Llevot et al. 4 Such divanillin was used as a bio-based aromatic building block for the design of semi-aromatic polymers exhibiting different thermo-mechanical properties. 5 However, divanillin monomer has also the potential to be valorized in the field of organic electronics. 6 Indeed, in addition to the two phenolic functions, such derivative possesses two aldehyde moieties that can be involved in condensation reactions, for instance with amino groups, leading to polyazomethines. Polyazomethines exhibit many advantages as they can be obtained through metal-free polycondensation with water as the only by-product. 7–12 Moreover, polyazomethines can be synthesized under micro-wave irradiation which brings the benefit of being faster and more efficient than conventional heating. 13,14 \(\pi\)-conjugated polyazomethines present a good thermal resistance and are air-stable. 15,16 They possess PPV-like electronic properties, can be tuned to exhibit a high absolute fluorescence quantum yield when doped and exhibit a good chemical and electrochemical resistance. 9,15–22 Interestingly, such polymers could be prepared by selecting diamino-functionalized monomers versus bis-aldehyde divanillin.

In this work, we describe the synthesis of a divanillin monomer substituted by alkyl chains and its subsequent polycondensation with 1, 4-benzene diamine or 2,7-carbazole diamine. The optical and electrochemical properties of the so-formed polyazomethines were also investigated.

2. Results and discussion.

The 6,6-Dihydroxy-5,5-dimethoxy-[1,1-biphenyl]-3,3-dicarboxaldehyde (Divanillin or DV) was synthesized from vanillin at room temperature using the procedure already described by us (Scheme 1). 1 In \(^1\)H NMR, (Figure S11); the signal at 9.7 ppm corresponds to the two aldehyde protons and the signal at 7.5 and 7.2 ppm to the aromatic ones. The signal at 3.8 ppm corresponds to the methoxy groups. The structure of DV was also attested by ATR-FTIR (Figure S12).

In order to improve the solubility of DV and polyazomethines thereof in classical organic solvents, the latter was alkylated. 9,13,14Etherification of DV and polyazomethines thereof was performed in DMSO in the presence of KOH and 2-ethylhexyl bromide to give Divanillin Ethyl Hexylated or DVEH (Scheme 1). The alkylation reaction was assessed by \(^1\)H NMR spectroscopy and ATR-FTIR (Figure S13 and S14); NMR signals around 3.81 ppm and the signals between 0.69 and 1.39 ppm confirmed the completion of alkylation reaction.
The divanillin-based monomer (DVEH) was polymerized at the stoichiometry ratio 1:1 both with commercially available benzene-1,4-diamine and 2,7-diaminocarbazole to give respectively P1 and P2 (synthesized according to the literature) \(^{(3)}\) (Scheme 1). The polymerization reaction being performed under micro-wave irradiation which is usually faster than conventional heating. \(^{(25,26)}\) Silica was added to the reaction medium to remove water and displace the equilibrium towards the polymer formation. P1A and P2 were obtained after 4 hours of reaction and purified by precipitation in methanol. Interestingly, the use of silica allowed a simple recovery process of the polyazomethine through filtration before precipitation. However, silica is not necessary to carry out the polymerization reaction. Indeed, the synthesis of the polyazomethine was also performed without silica, in just 5 minutes instead of 4h (P1B). The crude polyazomethine was then dissolved in methylene chloride; methanol was added and then the solvent was evaporated using a rotary evaporator. The powder obtained was rinsed with methanol, giving the final product P1C.

The polyazomethines were characterized by NMR and SEC. \(^{1}H\) NMR signal at 8.5 ppm clearly attested the formation of azomethine bond (-CH=N-) (Figure SI5 and SI6). All the molecular characteristics, thermal and optical properties of DVEH-based polyazomethines are reported in Table 1.

![Scheme 1: Alkylation of divanillin with ethylhexyl side chains (DVEH) and polymerization with p-phenylenediamine (P1) and diamino carbazole (P2).](image)

SEC traces show discrete peaks at low molar masses, according to a polycondensation pathway with heavy monomers (Figure SI7 and SI8). Both polymers have a dispersity close to two, which is consistent with a step-growth mechanism (at high conversion). The two families of polyazomethines present apparent number-average molar masses \(M_n\) of 10600 g.mol\(^{-1}\) and 6200 g.mol\(^{-1}\) respectively for P1A and P2, for a reaction time of four hours. For the polyazomethines P1, similar molar mass was obtained in just 5 minutes of reaction, after purification. Interestingly, post-condensation was probably occurring during the purification, resulting in a higher molar mass (10000 g/mol with 70% yield) compared to the crude. It confirms that post-condensation occurs during solvent evaporation. Thermal properties of the polyazomethines were determined by thermogravimetric analysis under \(N_2\) atmosphere at a heating rate of 10°C per minute. All polyazomethines present a good thermal stability with a degradation temperature around 405°C (Figure SI9 and SI10). The diamine used does not have a significant impact on the thermal stability. In addition, differential scanning calorimetry (DSC) analyses carried out between -80°C and 200°C did not show any specific transition neither glass transition nor crystallinity behavior, indicating the very rigid structure of these polyazomethines and their probable amorphous state (see Figure SI11 and SI12).

All the polyazomethines were characterized by UV-vis absorption and fluorescence spectroscopy analyses. As shown in Fig. 1, the carbazole-based polyazomethine main absorption band is red-shifted as compared to the phenyl-based derivative, from 364 to 390 nm. This behavior could be due to a stronger electron-rich/electron-poor couple in the case of the carbazolylene derivative or to a more planar conformation increasing the conjugation length. It is even amplified in fluorescence spectra where the shift can reach 150 nm. This is coherent with the higher molar attenuation coefficient of carbazole compared to the one of the bisvanillin. \(^{(27)}\) Both P1 and P2 have a weak emission, with a fluorescence quantum yield below 2% in solution.

![Figure 1. Absorption spectra (top) and emission spectra (excited at 360 nm, bottom) of the two DVEH-based polyazomethines P1A and P2 in methylene chloride](image)
Electrochemical features of these polyazomethines were determined by cyclic voltammetry using dichloromethane solutions of tetrabutylammonium hexafluorophosphate (TBAPF₆) as electrolyte, silver as reference electrode and platinum for the working and counter-electrodes. Finally, the material was solubilized in the electrolyte solution at a concentration of 0.1 g.L⁻¹ (Figures SI13 and SI14). The values of the energy levels are indicated in Table 2.

The slight differences between the electrochemical and optical band-gaps can be explained by the fact that the redox peaks result from localized sites rather than from the conjugated backbone.²⁸ For the 1, 4-diaminobenzene-based polyazomethines, both optical and electrochemical gap are comparable within experimental errors, with a value around 2.67 eV for the electrochemical gap and 2.96 eV for the optical one.

**Table 1. Molar masses, thermal and optical properties of divanillin-based polyazomethines.**

| Polyazomethines | Co-monomer of DVEH | Experimental conditions | Mₙ b (g.mol⁻¹) | Mₚ b (g.mol⁻¹) | Db | D𝑝 b | Tb c | λₘₐₓsol(λₘₐₓfilm) [nm] | λₘₐₓsol (λₘₐₓfilm) [nm] |
|-----------------|---------------------|------------------------|----------------|----------------|-----|------|-----|------------------------|------------------------|
| PtA             | NH₂-Ph-NH₂          | 4h, silica, purified   | 10600          | 21900          | 2.1 | 18   | 395 | 285-365 (290-370)       | 430 (460)              |
| PtB             | NH₂-Ph-NH₂          | 5min, no silica, crude | 5700           | 11500          | 2   | 9    |     | 285-365 (290-370)       |                       |
| PtC             | NH₂-Ph-NH₂          | 5min, no silica, purified | 10000         | 20000          | 2   | 17   |     |                        |                       |
| P₂              | NH₂-Cbz-NH₂         | 4h, silica, purified   | 6200           | 15400          | 2.5 | 7    | 401 | 270-390 (ND)            | 580 (ND)              |

* Under micro-wave irradiation at 130°C in toluene with APTS as a catalyst b Determined by Size Exclusion Chromatography (SEC) relative to polystyrene standards in THF at 40°C. c Decomposition temperature at 20% weight loss, evaluated under N₂, at a heating rate of 10 °C/min. d Determined in dichloromethane solution, e Determined on films obtained by drop-casting on quartz.

**Table 2. Physical properties of divanillin based polyazomethines.**

| Polyazomethine | HOMO [eV] a | LUMO [eV] a | E_{g}^{el}(eV) b | E_{g}^{opt}(eV) c | HOMO d | LUMO d | E_{g} d |
|----------------|------------|------------|------------------|------------------|--------|--------|--------|
| PTA            | - 5.21     | - 2.36     | 2.85             | 3.02             | -5.50  | -1.911 | 3.58   |
| P₂             | - 5.18     | - 2.50     | 2.68             | 2.85             | -5.203 | -1.808 | 3.39   |

* a Estimated from the oxidation and reduction potentials measured by cyclic voltammetry in CH₃Cl₃ solution. b Calculated by difference between oxidation and reduction potentials. c Calculated by Tauc method. d Calculated by B3LYP/6-31 (d,p) level of theory

In the case of carbazole-based polyazomethine, these values reach respectively 2.66 eV and 2.85 eV. These gap reductions are coherent with the red-shift observed in both absorption and emission spectra.

DFT calculations were carried out on DV-Ph and DV-Cbz derivatives at B3LYP/6-31(d,p) level of theory with chloroform as implicit solvent (CPCM). Different angles in both molecules were first estimated (Figures SI15 and SI16 and Table SI1). The choice of DFT functional employed is based on the closeness of calculated electronic bandgap (E₉ = E_{HOMO} - E_{LUMO}) of monomers (as a function of different DFT functional employed, see Table SI2 for more details) with respect to experimental values determined by cyclic voltammetry (Table SI3). DV-Cbz derivative shows lower E₉ compared to DV-Ph derivative as reported in Table 2.

Furthermore, Khun’s model was employed to compute the electronic band gap at polymer limit (Equation 1).

\[
E_{g} = E_{0} \left(1 + D_{k} \cos \left(\frac{\pi}{n+1}\right)\right)
\]

When n→∞, n is the number of monomer units in the oligomer, E₀ corresponds to electronic transition energy when N = 1, N being number of double bonds along the shortest conjugated pathway between terminal carbon atoms for oligomers of size n (Figure SI17). Dₖ is the force constant that represent the strength of coupling between single and double bonds, implicitly linked to the efficiency of connection between the units in an oligomer. By plotting E₉ as a function of number of double bonds (N), it exhibits higher π-conjugation of DV-Cbz derivative suggesting a better delocalization of π electrons b with respect to
DV-Ph derivatives with values of $D_k$ as 0.818 and 0.748 respectively. (Table SI4). These results are in agreement with experimental data.

3. Conclusion.
The synthesis of different polyazomethines embedding di-vanillin bio-based platform was performed. Divanillin was synthesized and alkylated, then polymerized with two different diamines: 1. 4-benzene diamine and an alkylated 2,7-diaminocarbazole. The polymerization was performed under micro-wave irradiation, without any metallic catalyst and with water as the only by-product. Remarkably, all polyazomethines present a broad emission at the beginning of the visible range. The carbazole-based polyazomethine is red-shifted compared to the phenyl-based one, in the case of both absorption and emission. This bathochromic shift is expected as the carbazole moiety is more electron-donating than the phenyl one. Interestingly, the methodology followed meets some challenges for a "green chemistry" such as lowering the number of steps, avoiding transition metal catalyst and use of bio-based monomers to quote a few. In addition, these new polyazomethines have promising properties (confirmed by TD-DFT calculations) for opto-electronic applications like OLED for example.

These encouraging results pave the way for a whole family of bio-based π-conjugated polyazomethines. Optical properties can be enhanced by optimizing the structure and/or by playing on doping. The extension to other π-conjugated moieties is currently investigated. Furthermore, for a better understanding of optical and electrical properties, small model trimers and dimers will be isolated and fully characterized. All these new development will be discussed in forthcoming papers.

4. Experimental Section

Materials.
Vanillin (>97%), Laccase from Trametes Versicolor, and 2-ethyl hexyl bromide (95%) were obtained from Sigma-Aldrich. Para-toluene sulfonic acid (PTSA, 99%) was purchased from TCI. Silica gel (pore size 60 Å, 230-400 mesh particle size, particle size 40-63 µm) was obtained from Honeywell Fluka. All products and solvents (reagent grade) were used as received except otherwise mentioned. The solvents were of reagent grade quality and were purified wherever necessary according to the methods reported in the literature. Flash chromatography was performed on a Grace Revelers apparatus, employing silica cartridges from Grace. Cyclohexane: ethyl acetate gradients were used as eluents. The detection was performed through ELSD and UV detectors at 254 nm and 280 nm. The reactions under micro-wave irradiation were performed on a Discover-SP from CEM, with the temperature measured by infrared; the power of the apparatus is constantly adjusted to reach and then stay at the set temperature.

Characterization techniques.
1H, 13C and 1H-13C HSQC NMR measurements were performed with a Bruker Avance 400 spectrometer (400.20 MHz and 100.63 MHz for 1H and 13C, respectively) at room temperature using deuterated solvent. IR spectra were recorded with Bruker Tensor 27 spectrometer using a 0.6 mm-diameter beam. Samples were analyzed with the attenuated total reflection (ATR) method. High resolution mass spectroscopy analyses were performed on an AutoSpec-Waters spectrometer (EI). Optical absorption spectra were obtained with a UV-visible spectrophotometer (UV-3600, Shimadzu). Photoluminescence spectra were obtained from a spectrofluorometer (Fluoromax-4, Horiba Scientific).

Molar masses of polymers P1A, P1B and P1C measured in THF were determined by size exclusion chromatography (SEC) using a three-columns set of Resipore Agilent: one guard column Resipore Agilent PL113-1300, then two column Resipore Agilent PL113-6300, connected in series, and calibrated with narrow polystyrene standards from polymer Laboratories using both refractometric (GPS 2155) and UV detectors (Viscotek). THF was used as eluent (0.8 mL/min) and trichlorobenzene as a flow marker (0.15%) at 30°C. Molar masses of polymer P2 measured in THF were determined by size exclusion chromatography (SEC) using a three-columns set of TSK gel TOSOH (G2000, G3000, G4000 with pore sizes of 20, 75, and 200Å respectively, connected in series) calibrated with narrow polystyrene standards from polymer Laboratories using both refractometric and UV detectors (Varian). THF was used as eluent (1 mL/min) and trichlorobenzene as a flow marker at 40°C. Molar masses of polymer samples in CHCl3, were measured by SEC at 40°C with THF as eluent, using a Viscotek VE2001-GPC. Polymer laboratories-Varian (one guard column and three columns based on cross-linked polystyrene, pore size = 200 Å, 75 Å and 20 Å), and PS standards were used for calibration.

TGA have been performed on a TA-Q50, from 25°C to 600 - 700 °C with a heating of 10°C.min-1 under nitrogen flow. DSC analysis have been performed on a TA instrument, under Helium flow, with an LN2 cooling and modulated with +/- 0.64 °C every 60 seconds. Sample were heated at 10°C.min-1 and cooled down at 5°C.min-1.

Electrochemical measurements and HOMO-LUMO calculation were performed in solution. A solution of 0.1 g.L-1 of the investigated polymer in CH2Cl2 with 0.1 M tetrabutylammonium hexafluorophosphate (TBAPF6) as electrolyte was prepared. Cyclic voltammetry (CV) measurements were then performed on the solution using silver wire as reference electrode and platinum for the working and counter electrodes. A solution of ferrocene (1 mM in the same solvent) was prepared in the same conditions and the redox potential of Fc/Fc+ vs Ag (EFc/Fc+ Ag) was measured, to be used as a reference for calibration.
Synthesis of 6,6′-Dihydroxy 5,5′dimethoxy-[1,1′biphenyl]-3,3′dicarboxaldehyde (Divanillin or DV).

A solution of vanillin in acetone was added to acetate buffer saturated in oxygen with laccase from Trametes Versicolor. In these conditions, the dimer formed precipitates and can be recovered by simple filtration. The filtrate has then simply been reloaded in vanillin and oxygen to start again the synthesis. Yield: 85%

1H-NMR (400.20 MHz, (CD3)2SO, ppm): δ 9.69 (s, 2H); 7.57 (d, J=1.9 Hz, 2H); 7.16 (d, J=1.9 Hz, 2H); 3.76 (s, 6H).

13C-NMR (100.63 MHz, (CD3)2SO, ppm): 191.2; 153.6; 152.1; 132.0; 128.6; 109.7; 75.2; 55.9; 40.4; 30.3; 29.1; 23.5; 22.9; 14.1; 11.0.

FT IR (ATR, cm⁻¹): ν= 2924, 2851, 1697, 1581, 1508, 1464, 1264, 1128, 1049, 970, 872, 788, 653

SEC, 1H NMR and TGA traces are available in Supporting Information.

Synthesis of 6,6′-bis-2-ethylhexyl-5,5′dimethoxy-[1,1′biphenyl]-3,3′dicarboxaldehyde (DVEH).

In a dried and nitrogen flushed 100 mL glassware, divanillin (2 g, 6.62 mmol) was solubilized in 20 mL of previously dried DMSO. Then KOH (0.89 mg, 15.9 mmol) was added to the reaction mixture which was heated at 80°C for 2 hours. Then, 2.2 equivalent of the 2-ethylhexyl bromide were added to the reaction mixture and heated for 12 more hours. The reaction mixture was then poured into 300 mL of water and extracted with 100 mL of diethyl ether, 3 times. The crude product recovered after the diethyl ether evaporation, was purified with flash chromatography in a mixture of cyclohexane/ethyl acetate (95/5).

Characterization of 6,6′-bis-2-ethylhexyl-5,5′dimethoxy-[1,1′biphenyl]-3,3′dicarboxaldehyde (DVEH).

An NMR, FTIR, SEC, DSC and cyclic voltamograms.

Supporting Information.

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SUPPORTING INFORMATIONS

Computational details (methods) and all supplementary characterization data (NMR, FTIR, SEC, DSC and cyclic voltamograms).
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Abbreviations

DMSO, Dimethyl sulfoxide; DSC, Dynamic scanning calorimetry; HOMO, Highest occupied molecular orbital; LUMO, Lowest unoccupied molecular orbital; NMR, Nuclear Magnetic Resonance

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Supporting Information

Divanillin-based polyazomethines: toward bio-based and metal-free π-conjugated polymers

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Computational Details

1. Electronic and optical properties of monomers

Relaxed potential energy surface scans were first performed on monomer units of DV-Cbz and DV-Ph to sample the possible stable conformations with respect to the torsional angles around single bonds. Calculations were performed at PBE0/6-31(d,p) level of theory with chloroform as implicit solvent using CPCM implicit solvation model. Three dihedral angles were considered for each derivative, θ₁, θ₂ and θ₃ as shown in Figure S15. Corresponding energy scans are presented in Figure S16.

The relative energies (ΔE in kcal/mol) of the various stable conformers obtained from the Potential Energy Surface (PES) scans are collected in Table S1.

In a second step, DV-Ph and DV-Cbz derivatives were fully optimized by starting from the lowest-energy conformer obtained from the PES scans. Optimizations were carried out at the DFT level using the 6-31(d,p) basis set with various exchange-correlation functionals (XCFs) and chloroform as implicit solvent. Frequency calculations were then performed to ensure proper optimization of the geometries. Table S2 reports the HOMO and LUMO energies (in eV), as well as the electronic band gap (Eg = ELUMO − EHOMO) of both derivatives, for a selection of exchange-correlation functionals (XCFs). Irrespective of the DFT functional employed, EG−DV-Cbz< EG−DV-Ph.

Furthermore, vertical transition energies (EVert) were computed at the time-dependent DFT (TD-DFT) level using the same XCFs and basis set as those employed in geometry optimizations. Consistently with the electronic band gap, results collected in Table S2 also show that EVert−DV-Cbz< EVert−DV-Ph, irrespective of the choice of XCF selected.

2. Electronic and optical properties of increasing-size oligomers

Similar DFT and TD-DFT calculations were then carried out on increasing-size oligomers of both derivatives, using the B3LYP and CAM-B3LYP functionals. B3LYP was selected since it provides good estimate of the HOMO and LUMO energy levels compared to those deduced experimentally from cyclic voltammetry measurements, although such comparisons are highly approximate. CAM-B3LYP was selected because it is known to improve the description of low-lying charge-transfer excited states in push-pull dyes [81]. Electronic band gaps (E₉) and vertical transition energies (EVert) as a function of chain length of the oligomers are gathered in Table S3. Irrespective of the increase in chain length, EG−DV-Cbz< EG−DV-Ph.
3. Electronic and optical gaps at polymer limit

Finally, the size-converged electronic and optical gaps were evaluated using a fitting procedure based on the Khun’s model, which is based on linear coupling of double bonds (harmonic oscillators) that contribute to electronic transitions.\[^{[S2, S3]}\] In this procedure, the electronic gap or vertical transition energies reported in Table S3 are first plotted against 1/N, where N is the number of double bonds along the shortest conjugated pathway connecting the terminal carbon of the oligomers. Then, a linear regression can be obtained based on Khun’s equation (Equation 1) to extract energy gaps at polymer limit:

\[
E_g = E_0 \sqrt{1 + D_k \cos \left( \frac{\pi}{N+1} \right)}
\]  

In the above equation, \( E_0 \) and \( D_k \) are the fitting parameters. \( E_0 \) corresponds to the energy gap when \( N = 1 \) and \( D_k \) is a force constant that represents the strength of coupling between single and double bonds in a given oligomer, entailing that \( D_k \) is implicitly linked to the efficiency of the \( \pi \)-conjugation between consecutive monomeric units (higher the value of \( D_k \), higher the delocalization of \( \pi \) electrons).\[^{[S4]}\]

The evolution of the electronic and optical gaps as a function of 1/N, calculated at the B3LYP/6-31G(d) and CAM-B3LYP/6-31G(d) levels, are shown in Figure S17. The size-converged electronic and optical gaps, as well as the optimized \( E_0 \) and \( D_k \) parameters, are reported in Table S4. The two levels of calculation predict that the DV-Cbz derivative shows lower electronic and optical gaps at polymer limit than the DV-Ph derivative. This relative ordering both originates from a lower value of the band gap of the DV-Cbz monomer (see section 1), and from a more efficient \( \pi \)-conjugation between consecutive units, as indicated by the larger \( D_k \) values.

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\[^{[S4]}\] Wykes, M.; Milin-Medina, B.; Gierschner, J. Computational engineering of low bandgap copolymers, *Frontiers in Chemistry*, 2013, 1, 35.
Figure S1. $^1$H-NMR (top) and $^{13}$C-NMR (bottom) spectra of DV (400.20 MHz and 100.63 MHz respectively, in (CD$_3$)$_2$SO)
Figure S2. ATR-FTIR spectrum of DV

Figure S3. $^1\text{H}-^{13}\text{C}$ HSQC NMR spectrum of DVEH (400.20 MHz and 100.63 MHz respectively, in CDCl$_3$)
Figure S4. ATR-FTIR spectrum of DVEH

Figure S5. $^1$H-NMR spectrum of P1A (400.20 MHz, in CDCl$_3$)
Figure S6. $^1$H-NMR spectrum of P2 (400.20 MHz, in CDCl$_3$)

Figure S7. SEC traces of P1A, P1B and P1C (in THF, polystyrene standard)
Figure S8. SEC trace of P2 (in THF, polystyrene standard)

Figure S9. TGA trace of P1A
Figure S10. TGA trace of P2

Figure S11. DSC trace of P1A
Figure S12. DSC trace of P2

Figure S13. Cyclic voltammograms (left-reduction, right-oxidation) of P1A in CH₂Cl₂ solution
Figure S14. Cyclic voltammograms (left-reduction, right-oxidation) of P2 in CH₂Cl₂ solution

Figure S15. Dihedrals considered for the relaxed potential energy surface scans in monomers of (left) DV-Ph and (right) DV-Cbz derivatives

[a] Dihedral: θ₁  [b] Dihedral: θ₂  [c] Dihedral: θ₃
**Figure S16.** Relaxed potential energy surface scans with respect to dihedrals $\theta_1$, $\theta_2$ and $\theta_3$ for monomers of **DV-Ph** (Ph: Black - Circles) and **DV-Cbz** (Cbz: Red - Squares)

![Graphs showing relaxed potential energy surface scans for DV-Ph and DV-Cbz monomers.]

[a] B3LYP: $E_G$  [b] CAM-B3LYP: $E_G$  [c] CAM-B3LYP: $E_{Vert}$

**Figure S17.** Evolution with chain length of the electronic gap of increasing-size **DV-Cbz** (red) and **DV-Ph** (black) oligomers, as calculated at the (a, left) B3LYP/6-31G(d) and (b, centre) CAM-B3LYP/6-31G(c) levels. The right panel reports the evolution of the vertical transition energy ($E_{Vert}$) calculated at the CAM-B3LYP/6-31G (c) level. Dotted lines are Khun fits

| Derivative / Dihedral | $\theta_1$ | $\theta_2$ | $\theta_3$ | $\Delta E$ (kcal/mol) |
|-----------------------|------------|------------|------------|-----------------------|
| **DV-Ph**             |            |            |            |                      |
| 40.0                  | 177.48     | 126.39     |            | 1.020                 |
| 140.0                 | 177.53     | 126.31     |            | 1.149                 |
| 39.55                 | 0.0        | 126.51     |            | 1.864                 |
| 39.89                 | 180.0      | 126.84     |            | 1.036                 |
| **40.08**             | **176.98** | **60.0**   |            | **0.0**               |
| 39.73                 | 176.96     | 120.0      |            | 1.072                 |
| **DV-Cbz**            |            |            |            |                      |
| 40.0                  | 177.61     | 126.27     |            | 0.937                 |
| 140.0                 | 176.77     | 125.32     |            | 0.937                 |
| 40.81                 | 0.0        | 126.67     |            | 1.081                 |
| 41.11                 | 180.0      | 125.18     |            | 1.895                 |
| **39.30**             | **177.69** | **60.0**   |            | **0.0**               |
| 40.18                 | 176.59     | 120.0      |            | 1.098                 |
Table S1. Stable conformers obtained from the PES scans for $\theta_1$, $\theta_2$ and $\theta_3$ and relative energies ($\Delta E$ in kcal/mol)

| Functional | DV-Ph | | DV-Cbz | | ΔE_{(Phnl-Crbz)} |
|------------|-------|------|------|------|----------------|
|            | HOMO  | LUMO | $E_G$ | $\lambda_{\text{max}}$ | $E_{\text{Vert}}$ | HOMO  | LUMO | $E_G$ | $\lambda_{\text{max}}$ | $E_{\text{Vert}}$ | ΔE_{G} | ΔE_{Vert} |
| PBE0       | -5.952| -1.643| 4.309 | 352 | 3.516 | -5.631| -1.622| 4.009 | 376 | 3.294 | 0.300 | 0.222 |
| B3LYP      | -5.625| -1.753| 3.872 | 394 | 3.375 | -5.329| -1.702| 3.627 | 394 | 3.104 | 0.245 | 0.235 |
| CAM-B3LYP  | -7.037| -0.480| 6.556 | 315 | 3.930 | -6.632| -0.483| 6.149 | 329 | 3.766 | 0.407 | 0.164 |
| wB97XD     | -7.599| -0.051| 7.548 | 313 | 3.951 | -7.220| -0.065| 7.155 | 323 | 3.834 | 0.393 | 0.117 |
| M06HF      | -8.797| -0.446| 8.350 | 301 | 4.107 | -8.286| -0.353| 7.933 | 304 | 4.071 | 0.417 | 0.036 |
| M062X      | -6.964| -0.827| 6.137 | 319 | 3.886 | -6.597| -0.861| 5.736 | 333 | 3.716 | 0.401 | 0.170 |

Table S2. Electronic ($E_G$) and optical ($E_{\text{Vert}}$) gaps (in eV), as well as maximum absorption wavelengths (in nm) of DV-Cbz and DV-Ph derivatives calculated at the TD-DFT level using various XCFs with the 6-31(d,p) basis set. $\Delta E_{(Phnl-Crbz)}$ corresponds to differences between the electronic/optical gaps of the two derivatives

| Functional | Unit | DV-Cbz | | | | DV-Cbz | | | | | | ΔE_{G(Phnl-Crbz)} |
|------------|-----|--------|---|---|---|--------|---|---|---|---|---|---|
| [a] B3LYP  | Monomer | -5.625 | -1.753 | 3.872 | | -5.329 | -1.702 | 3.627 | | 0.245 |
|            | Dimer | -5.516 | -1.894 | 3.621 | | -5.224 | -1.795 | 3.429 | | 0.192 |
|            | Trimer | -5.507 | -1.909 | 3.597 | | -5.214 | -1.803 | 3.411 | | 0.186 |
|            | Tetramer | -5.500 | -1.911 | 3.588 | | -5.203 | -1.808 | 3.395 | | 0.190 |

Table S3. Electronic ($E_G$) and optical ($E_{\text{Vert}}$) gaps (in eV), as well as maximum absorption wavelengths (in nm) of increasing-size DV-Cbz and DV-Ph derivatives, as calculated at the [a] B3LYP/6-31G(d) and [b] CAM-B3LYP/6-31G(d) levels. $\Delta E_{G(Phnl-Crbz)}$ corresponds to differences between the electronic/optical gaps of the two derivatives

| Functional | Unit | DV-Ph | | | | DV-Cbz | | | | | | ΔE_{G(Phnl-Crbz)} |
|------------|-----|--------|---|---|---|--------|---|---|---|---|---|---|
| [b] CAM-B3LYP | Monomer | -7.037 | -0.480 | 6.556 | 315 | 3.930 | -6.632 | -0.483 | 6.149 | 329 | 3.766 | 0.407 | 0.164 |
|            | Dimer | -6.879 | -0.623 | 6.248 | 334 | 3.712 | -6.553 | -0.580 | 5.973 | 342 | 3.623 | 0.275 | 0.089 |
|            | Trimer | -6.862 | -0.634 | 6.227 | 336 | 3.688 | -6.535 | -0.594 | 5.941 | 343 | 3.607 | 0.286 | 0.081 |
|            | Tetramer | -6.857 | -0.642 | 6.210 | 337 | 3.679 | -6.529 | -0.601 | 5.928 | 344 | 3.596 | 0.285 | 0.083 |

Table S4. Electronic band gap and vertical transition energies at the polymer limit ($E_g$ in eV), and optimized $E_0$ and $D_k$ parameters (in eV, extracted from Equation 1) for DV-Cbz and DV-Ph derivatives.