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Tanguy Jousselin-Oba, Masashi Mamada, Jérôme Marrot, Antoine Maignan, Chihaya Adachi, et al.. Excellent Semiconductors Based on Tetracenotetracene and Pentacenopentacene: From Stable Closed-Shell to Singlet Open-Shell. *Journal of the American Chemical Society*, 2019, 141 (23), pp.9373-9381. 10.1021/jacs.9b03488 . hal-02347038

HAL Id: hal-02347038

<https://hal.science/hal-02347038>

Submitted on 5 Nov 2019

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Excellent Semiconductors Based on Tetraceno-Tetracene and PentacenoPentacene: From Stable Closed-Shell to Singlet Open-Shell.

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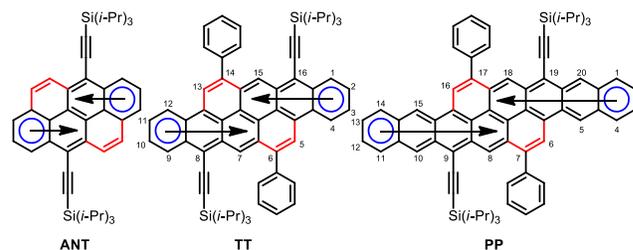
Supporting Information Placeholder

ABSTRACT: Designing stable open-shell organic materials through the modifications of the π -topology of molecular organic semiconductors have attracted recently considerable attention. However, their uses as an active layer in organic field-effect transistors (OFETs) are very limited and the obtained hole and electron charge mobilities are around $10^{-3} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$. Herein, we disclosed the synthesis of two materials so-called tetraceno-tetracene (**TT**) and pentaceno-pentacene (**PP**) which have low band gap of 1.79 and 1.42 eV, respectively. Their ground state natures have been investigated by different experiments including steady state absorption, electron spin resonance, superconducting quantum interfering device (SQUID) and variable temperature NMR along with DFT calculations. **TT** and **PP** have closed-shell and singlet open-shell structures in their ground state, respectively, and possess high stability. Their biradical characteristics were found to be 0.50 and 0.64. The origin of the open-shell character of **PP** is related to the concomitant opening of two tetracenes with the recovering of two extra aromatic sextets and a small HOMO-LUMO energy gap (gap < 1.5 eV). Thanks to the high stability, thin film OFET devices could be fabricated. In TG-BC configuration **PP** shows remarkably high hole mobility of $1.4 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ while **TT** exhibits a hole mobility of $0.77 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$. In configuration of BG-TC, ambipolar behaviors for both were obtained with hole and electron mobilities of 0.21 and $0.01 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ for **PP** and 0.14 and $0.006 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ for **TT**.

INTRODUCTION

Acenes composed of laterally condensed benzenoid rings are a special class of polycyclic aromatic hydrocarbons (PAHs) that have been intensively studied in the last two decades.¹ Functionalized acenes up to five benzene rings have been of interest in potential applications going from organic light-emitting diodes (OLEDs), organic photovoltaic devices to organic field-effect transistors (OFETs).² For this latter application, the benchmark of soluble acenes is 6,13-bis(*triisopropylsilyl*ethynyl)pentacene (**TIPS-PEN**) that showed excellent compatibility between stabil

Chart 1. Clar's structures of the closed-shell form of TIPS-Anthanthrene, TT and PP.



ity and solubility and exhibited good charge transport.³ Since the mobilities in OFETs depends on the packing in the solid phase and the reorganization energy that decreases with the acene length, longer acenes containing more than five rings were particularly promising for optoelectronic devices.⁴ Although in the past decade, the synthesis of soluble hexacene, heptacene and nonacenes functionalized with silyl acetylenes was achieved, their stabilities in solution were detrimental to show their intrinsic properties.⁵ Indeed, nowadays, it is clear that longer acenes should be regarded as singlet open-shell PAHs in the ground state.⁶ In recent years, a large number of delocalized biradical PAHs that can be written with a Kekulé structure including bisphenalenyls, zethrenes, indenoacenes, anthenes, peri-tetracene and rylene ribbons have been disclosed.⁷⁻¹³ Compared to longer acenes, the stability of those biradicals are genuinely higher due to the presence of at least two aromatic sextets. For most biradical PAHs, the reactive sites are functionalized with bulky groups which are not really excellent groups to promote good π -stacking in the solid state, leading to charge mobilities of around $10^{-3} \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ at best in OFETs.^{8e, 10e, 10h} Finding a chemical structure combining a biradical character in the ground state, high stability and suitably functionalized to promote high π -overlap in the solid state is highly appealing for the fabrication of ambipolar OFET devices.

In the recent years, some groups including us have been interested on two dimensional (2D) PAHs based on tetraceno-tetracene derivatives (**TT**) that correspond to two tetracenes fused at the

zigzag edge in different manners.¹⁴ They have shown similar optoelectronic properties like **TIPS-PEN** associated with high charge mobility and stability. The enhanced stability has been associated to the presence of two aromatic sextets in the molecule, leading to high distortion energy in interaction with oxygen.¹⁵ Indeed, the change of aromaticity from **TT** to its oxidized form is globally comparable to that observed from anthracene to its oxidized form.¹⁶ Since **TT** can be regarded as a dibenzoanthanthrene and **TT** functionalized with only TIPS-acetylene groups in the 8,16 positions has an absorption maximum (λ_{max}) at 648 nm, corresponding to a bathochromic shift of 153 nm (0.59 eV) in comparison with TIPS-anthranthrene **ANT** (495 nm),^{14d, 17} the corresponding TIPS-dinaphtho-anthranthrene so called pentaceno-pentacene **PP** as shown in Chart 1 should have a λ_{max} around 800 nm, absorption behavior in between alkylsilylhexacene (740 nm) and alkylsilylheptacene (840 nm).^{5a} Since longer acenes are singlet open-shell in the ground state and **PP** is an acene-like molecule, if the energy gap HOMO-LUMO is a critical factor for the occurrence of the biradical character as shown recently by Wu, et al. on their study of rylene ribbons, **PP** should be a singlet biradical and could be an excellent organic semiconductor.¹³

Herein, we report an efficient synthesis of **PP** and its smaller homologue **TT** for comparison and discuss their optoelectronic properties and stabilities and their ground-state electronic structures studied by different experiments and density functional theory (DFT) calculations and finally their charge transport behaviors in OFETs.

Pristine **PP** has been recently prepared using a simple methodology so-called dehydrative π -extension (DPEX) from the cyclisation of pyrene functionalized with two 2-naphthaldehyde derivatives.¹⁸ However due to the insolubility of the material, no optoelectronic properties have been reported.

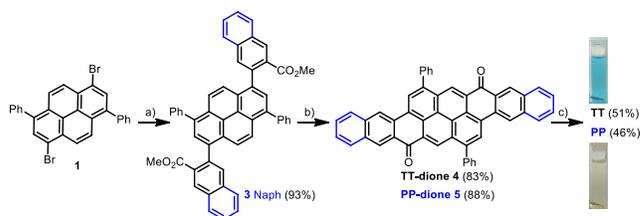
TT and **PP** have been functionalized with TIPS-acetylene in the most reactive positions as recently shown in **TT** derivatives, to ensure stability and solubility and to promote good π -stacking in solid state.^{14b} The phenyl groups are here not only to avoid the competition between 1,4- and 1,2-addition on the carbonyl groups of the TIPS-acetylene groups at the last stage of the preparation but also to be eventually beneficial for the solubility and the packing issues.^{14c-d}

RESULTS AND DISCUSSION

The synthetic path for the preparation of **TT** and **PP** derivatives is depicted in Scheme 1 and follows the strategy used previously for the synthesis of **TT** derivatives.^{14b} From 1,6-dibromo-3,8-diphenylpyrene **1**, cross-coupling reaction with phenyl- or naphthylboronic acid pinacol ester derivatives afforded diester **2** and **3** respectively in yield above 90%. The TT-dione **4** and the PP-dione **5** were obtained in excellent yield via a double intramolecular acylations using TfOH as activator.¹⁸ Addition of the lithium (triisopropylsilyl)acetylide to either **4** and **5** followed by SnCl₂ mediated reduction of the diol intermediates afforded the desired **TT** as dark blue solid and **PP** as dark green solid with yield around 50%. The structure of **TT** was confirmed by X-ray diffraction and the purity of samples was checked by ¹H NMR, MALDI-TOF measurement in which only one peak assigned to molecular ion of the materials was observed and by elemental analysis for **PP** (see Supporting information). For this latter analysis, the expected values for carbon and hydrogen are 87.40% and 6.93% and the found values are 87.16% and 6.96%.

Optical and Electrochemical properties: **TT** and **PP** are blue and light green in chloroform solution, respectively. The UV/Vis absorption spectrum of **TT** shows typical acene-like vibronic feature with three distinct bands at 559, 608, and 662 nm (λ_{max}) that increase in intensity going towards the low energy of the

Scheme 1. Synthetic path of **TT** and **PP**.



(a) Ester boronic derivatives, Pd₂(dba)₃, S-Phos, K₃PO₄, toluene/water, 100°C; (b) TfOH, 1,2-dichloroethane, 90°C; (c) 1) Lithium TIPS-acetylide, THF, 0°C, 2) SnCl₂/HCl for **TT** and SnCl₂, toluene, r.t. for **PP**

spectrum (Figure 1 (a)). In the similar manner, the typical three bands are observed for **PP** with λ_{max} found at 806 nm which is red-shifted of 142 nm (0.33 eV) compared to that of **TT** and λ_{max} appears between hexacene (740 nm) and heptacene (840 nm) functionalized with silyl-acetylene groups as expected.^{5a} Like functionalized heptacene derivatives and most of singlet biradicals, after λ_{max} , a weak and broad transition is observed that extends up to 900 nm.^{5b-c} This small transition that could be attributed to the HOMO, HOMO → LUMO, LUMO double excitation, is the signature that **PP** has a significant contribution of the singlet biradical form in the ground state.^{20, 7a} It is noteworthy that the molar absorption coefficient (ϵ) at λ_{max} of **TT** (94000 L mol⁻¹ cm⁻¹) is 1.27 larger than the value obtained for **PP** (74000 mol⁻¹ cm⁻¹), indicating that **TT** has a higher overall aromaticity than **PP**. Indeed, in a similar manner, TIPS-tetracene (32000 L mol⁻¹ cm⁻¹) has an ϵ value 1.23 higher than **TIPS-PEN** (26000 L mol⁻¹ cm⁻¹).²¹

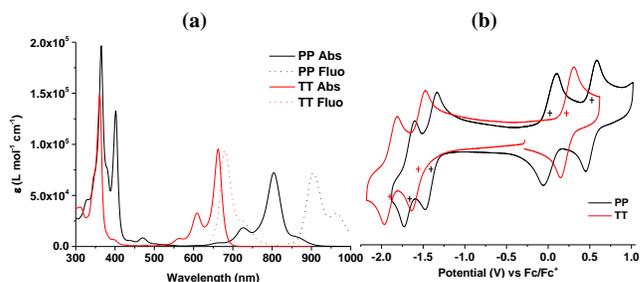


Figure 1. (a) Absorption and emission spectra in chloroform and (b) Cyclic voltammetry of **TT** and **PP** in chlorobenzene.

TT exhibits significant fluorescence properties in chloroform solution and emits at 679 nm with the fluorescence quantum yield of 63% and the total decay lifetime of 7.6 ns. As for low band gap materials with a significant contribution of the singlet biradical form in the ground state, **PP** shows very weak fluorescence at 904 nm when excited at 806 nm, with the fluorescence quantum yield of 0.8% and the total decay life time of 4.3 ns. It is noteworthy that the fluorescence spectrum obtained using an excitation wavelength of 365 nm, on top of the band at 904 nm, exhibits two weak fluorescence bands at 460 nm and 670 nm that increase strongly in intensity with the number of scans (see Supporting Information). These results indicate that **PP** degrades readily to two different oxidation products upon high energy irradiation.

Electrochemistry: Cyclic voltammetry (CV) was performed to investigate the electrochemical behaviors and probe the HOMO/LUMO energy levels of materials. The measurements were performed in chlorobenzene and half-wave potentials were determined relative to Fc/Fc⁺. CVs of materials are shown in Figure 1 (b). Both **TT** and **PP** show amphoteric redox behaviors and exhibit two reversible reductions and two reversible oxidations for **PP** and only one reversible oxidation for **TT**. Half-wave

potentials of the first oxidation/reduction peaks were at 0.23/−1.56 for **TT** and 0.02/−1.40 V for **PP**. From these values, the HOMO/LUMO energy levels were found to be −5.03/−3.24 eV for **TT** and −4.82/−3.40 eV for **PP**. Thus, the electrochemical energy gaps were estimated to be 1.79 and 1.42 eV, respectively. As expected increasing the number of benzenoid rings in linear fashion reduces the HOMO-LUMO band gap by stabilizing the LUMO and destabilizing the HOMO energy levels at the same time. The ionization potential for the thin-film of **PP** determined using a photoelectron yield spectroscopy was −4.87 eV, which is in good agreement with the HOMO level measured by CV (See Supporting Information).

Stability experiments of the materials in saturated air-chloroform solution in lightening laboratory were performed and compared to **TIPS-PEN** by monitoring the change of UV/Vis absorption over time (Figure 2, see Supporting Information). **TT** has a half-life of 6.5 days and it is about 3.8 times more stable than **TIPS-PEN** (1.7 days). As **TT** and **TIPS-PEN** have absorption bands in the same region of the spectrum and the ϵ value of **TT** is about 3.6 higher than that of **TIPS-PEN**, **TT** is intrinsically 14 times more stable than the **TIPS-PEN**. Remarkably, **PP** is about 3 times more stable than **TIPS-PEN** with a half-life of 4.9 days that increases to 19 days in presence of air without light.

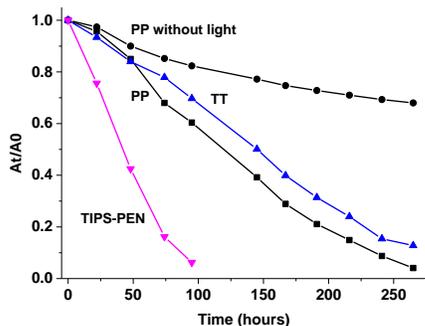


Figure 2. Change of UV/Vis absorption (followed at their own λ_{\max}) over time in chloroform (20 μm) of **TT** and **PP** compared to **TIPS-PEN**.

VT ^1H NMR, ESR and SQUID measurements: Since the UV/Vis spectrum of **PP** shows that the singlet biradical state contributes in the ground state, variable temperature (VT) ^1H NMR spectra were performed in 1,2-dichlorobenzene- d_4 from 0 to 140 $^\circ\text{C}$ (only shown up to 80 $^\circ\text{C}$, Figure 3 (a), see Supporting Information).

The assignments of the ^1H NMR signals were based on COSY and NOESY experiments recorded at 20 $^\circ\text{C}$ (see Supporting Information). The two singlet signals at 9.72 ppm and 9.67 ppm were assigned to protons H8, H18 and H10, H20 respectively based on the presence of dipolar contacts with protons of the TIPS groups and a weak dipolar contact with protons of the phenyl groups for H8 and H18. The two other singlet signals at 9.53 ppm and 9.21 ppm were assigned to H5, H15 and H6, H16 respectively. Even though protons H6, H16 are in close proximity with protons of the phenyl groups, no dipolar contact was observed. Therefore, we assume that the most deshielded signal should be assigned to protons H5, H15 since they are located at the zigzag edge of a ring which is delocalized in nature and the most shielded one assigned to protons H6, H16 since they are located at the side of the molecule. The protons of the phenyl groups were easily identified at 8.09, 7.76 and 7.67 ppm for the protons in ortho, meta and para of the substitution respectively. The most deshielded signal centered at 7.51 ppm was attributed to protons H2, H12 and H3, H13 since they are at the side of the molecule.

Their neighbors protons H1, H11 and H4, H14 were found to be at 8.18 ppm and 8.15 ppm respectively based on COSY and the presence of a dipolar contact between H1, H11 and the protons of the TIPS.

The VTR experiments show that at 0 $^\circ\text{C}$, all the signals are sharp and as the temperature increases, the signals of protons H8, H18 and H10, H20 and H1, H11 broaden and disappear completely at 80 $^\circ\text{C}$. The signals of protons H5, H15 and H2, H12 and H3, H13 broaden but do not collapse even at 140 $^\circ\text{C}$ (see Supporting Information). The signal of protons H6, H16 and the three signals related to the protons of the phenyl groups are not much altered. All the signals are recovered when the sample is cooled back to r.t indicating that **PP** is stable when heated in solution. The ^1H NMR signal broadening by increasing temperature is a typical phenomenon for singlet open-shell PAHs and results from a thermally accessible triplet biradical form which is slightly higher in energy than the singlet biradical one. Hence, the spin distribution of the triplet biradical of **PP** (Figure 3 (b)) was calculated and indicates that the spin density is highly localized at the zigzag edge. The protons that hold the highest spin density collapse in VT ^1H NMR experiment whereas the other protons with less spin density or with no density at all broaden or are not much altered.

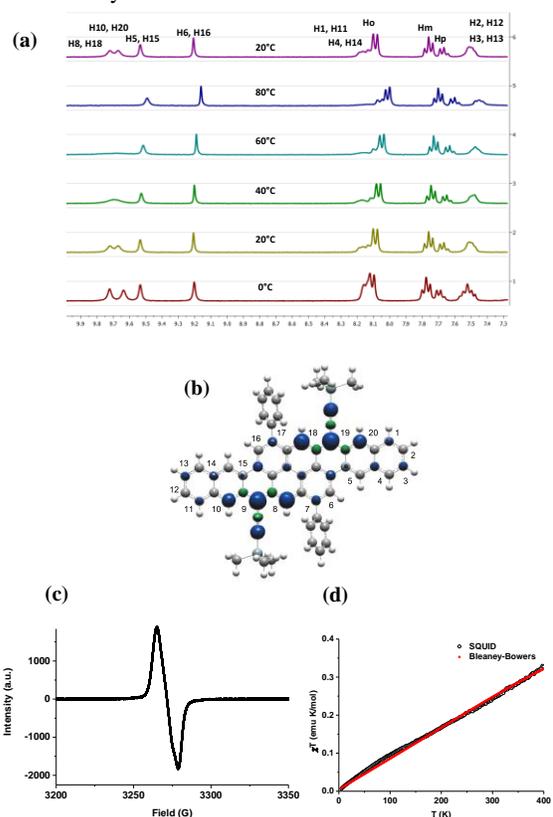


Figure 3. (a) VT ^1H NMR spectra (aromatic region) of **PP** in *o*-DCB- d_4 . The signal assignments referred to the structure shown in Chart 1; (b) Spin density of the triplet biradical of **PP**; (c) ESR spectrum of **PP** in powder at r.t.; (d) $\chi_m T$ versus T plot for **PP** in the solid state in the temperature range 5–400K from SQUID measurements.

In solid state, **PP** exhibits a ESR signal at $g = 2.0026$ at room temperature (Figure 3 (c)) which is typical for singlet open-shells with triplet state thermally accessible. The superconducting quantum interference device (SQUID) measurement at the temperature range of 5–400K shows that the magnetic susceptibility ($\chi_m T$) decreases with a reduction of temperature (Figure 3 (d)). Fitting the data by using Bleaney-Bowers equation gave a singlet–triplet energy gap (ΔE_{S-T}) of −4.85 kcal/mol ($2J = -$

1696 cm⁻¹).^{23, 24} Similar $\chi_m T$ versus T plots and ΔE_{S-T} values were obtained recently with biradical picene derivatives.^{10h}

Theoretical Calculations: In order to have a deeper insight into the electronic nature of **PP**, density functional theory (DFT) analysis was performed using Gaussian 16 except for anisotropy of the current induced density (ACID) plot and nucleus-independent chemical shift (NICS) calculations (see Supporting Information for details).^{25, 26} The biradical character was estimated from the electron occupancies of the frontier natural orbitals through the Yamaguchi's scheme (y_0) at UHF/6-31G(d,p) level of theory.²⁷ For comparison, the biradical character of **PEN**, hexacene (**HEX**) and heptacene (**HEPT**) and their homologues functionalised with TIPS-acetylene were also calculated (Table 1, see Supporting information). For the calculation, the TIPS groups were replaced by TMS groups. For all systems under studies, the TMS-acetylene groups increase the biradical character by about 0.04–0.06 due to the extra resonance afforded by each spin and reduces the energie difference between singlet and triplet open-shell forms. The calculations predict that pristine **TT** and **PEN** and theirs functionalized homologues have a closed-shell ground state whereas **PP** and higher acenes have an open-shell ground state. **PP** has a biradical character y_0 of 64%, value in between of **TMS-HEX** (58%) and **TMS-HEPT** (68%). Indeed, the biradical character increases while the band gap and the singlet-triplet energy gap decrease.

Table 1. DFT calculation of the biradical character (y_0)

Compound	y_0^a	$\Delta E(\text{OS} - \text{CS})^b$	ΔE_{S-T}^b	$\langle s^2 \rangle^b$
native TT	0.45	0	-17.19	0
TT	0.50	0	-13.86	0
native PP	0.60	-0.109	-8.43	0.265
PP	0.64	-0.374	-6.54	0.460
PEN	0.41	0	-18.00	0
TMS-PEN	0.47	0	-13.55	0
TMS-HEX	0.58	-0.256	-8.40	0.373
TMS-HEPT	0.68	-1.898	-5.46	0.828
TMS-DNP	0.57	0	-9.50	0

^a at UHF/6-31G(d,p) level of theory; ^b at B3LYP/6-311G(d,p) level

The calculated singly occupied molecular orbital (SOMO) profiles of the α and β spins of the singlet biradical form of **PP**, shown in Figure 4 (a) are highly delocalized and the spin density map indicates that the largest spin densities are typically found at the zigzag edge (Figure 4 (b)). The couple of carbons C9 and C19 (0.188) holds the largest spin density, followed by the carbons C8 and C18 (0.158), and then by the carbons C10 and C20 (0.104). The carbons C9 and C19 are protected with TIPS-acetylene and the latter bears appreciable spin density (0.099) and so takes part to the delocalization of the spin as mentioned above. These results indicate that **PP** should be considered as a resonance hybrid structure between the closed-shell form and six open shell resonance forms having the highest thermodynamic stabilities by keeping four aromatic sextets as shown in Figure 4 (c).

The origin of the open-shell nature of **PP** is interesting. As shown in Chart 1, the Clar's structures of **ANT**, **TT** and **PP** possess two aromatic sextets that are delocalized over two, three and four rings respectively. These representations correspond to three Clar's resonance forms for **ANT**, to eight Clar's resonance forms out of ten possible for **TT** and to fifteen out of nineteen possible for **PP** (see Supporting information). In contrast to **ANT** that cannot hold an aromatic sextet in the rings marked in red according to Clar's sextet rule, the missing two structures for **TT** and four for **PP** correspond to those where one aromatic sextet would

be localized on that rings that can be neglected (2/10 for **TT** and 4/19 for **PP**) (See Supporting Information).²⁸

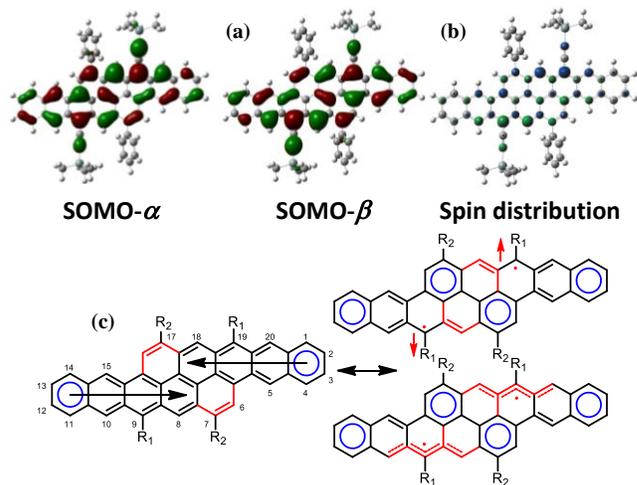


Figure 4. (a) Calculated SOMOs for the α and β electrons (UCAM-B3LYP) and (b) spin density distribution of the singlet biradical of **PP** with TMS group for simplification; (c) Resonance between the closed-shell and the open-shell forms of **PP**.

These representations would suggest that the rings marked in red should have a bond with a high degree of double character (shortest bond length) and the remaining bonds at the zigzag edge should be similar to naphthalene for **ANT**, anthracene for **TT** and tetracene for **PP**.

The theoretical bond lengths of **ANT**, **TT** and **PP** along with those of naphthalene, anthracene and tetracene are depicted in Figure 5. The bond lengths at the zigzag edge of the acenes show bond length alternation and the double character of bond **a** increases with the acene length going from 1.373 Å for naphthalene to 1.367 Å for anthracene and 1.363 Å for tetracene. For the peri-fused systems, as expected the bond lengths of bonds **a** are the shortest ranging from 1.355 Å for **ANT** to 1.358 Å for **PP** indicating their high degree of double bond character. The remaining bond lengths of **ANT**, **TT** and **PP** at the zigzag edge are very similar to naphthalene, anthracene and tetracene respectively except for the bonds **c** and **d**. Since these bonds are very similar for the three peri-fused systems, we can assume that these bond lengths are governed to some extent by the geometry of the molecules.

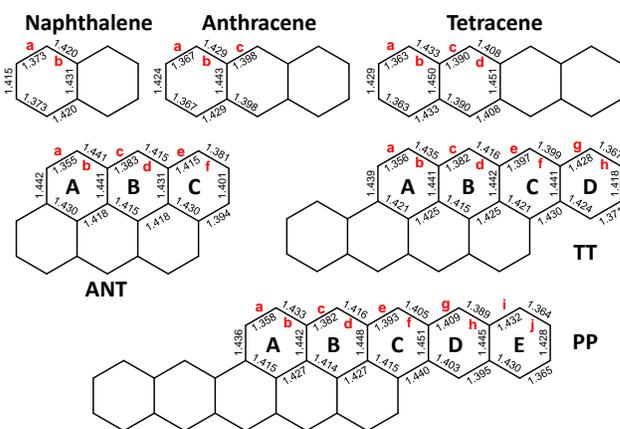


Figure 5. Theoretical bond lengths of naphthalene, anthracene, tetracene, pristine **ANT**, **TT** and **PP**.

The Clar's representations of the three peri-fused systems are also in agreement with the calculation of NICS using the NICS-

XY scan and π -only methods and the corresponding anisotropy of the current induced density (ACID) plot (Figure 6).²⁵⁻²⁶

For all systems, rings A sustain the lowest diatropic ring current with NICS(1.7)_{zz} values ranging from -10.0 to -10.4 ppm indicating that rings A are not delocalized rings. The NICS(1.7)_{zz} values of rings B and C for **ANT**, rings B, C and D for **TT** and rings B, C, D and E for **PP** are similar to those obtained for naphthalene, anthracene and tetracene respectively (see Supporting Information). The acid plots indicate that the diatropic ring current is mainly delocalized at the zigzag edge and all the rings sustain a diatropic ring current (red arrows) like for delocalized rings except for rings A for which the diatropic ring current (blue arrows) is rather related to a conjugation. There is also a transannular ring current (black arrow) below rings A due to the fact that the rings below are delocalized rings in nature confirming the Clar's representation of the three peri-fused systems as shown in Chart 1.

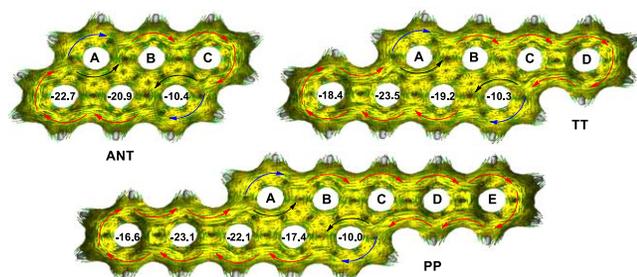


Figure 6. NICS(1.7)_{zz} calculations and acid plots of closed-shell forms of **ANT**, **TT** and **PP**.

Accordingly, the closed-shell of **PP** can be considered as two tetracene fused at the face and bridged by two ethylenic spacer. Therefore, the open-shell character of **PP** occurs due to the concomitant opening of two tetracenes with the recovering of two extra aromatic sextets.

In the recent literature, there are two open shell systems namely peri-tetracene (**PT**) and its opened homologues tetrabenzoo[*a,f,j,o*]perylene (**TBP**) that involve the opening of two tetracenes (Figure 7).¹² However, in these cases, the opening of the two tetracenes leads to the gain of three aromatic sextets instead of two indicating that the change of aromaticity between the closed- and the open-shell for them is higher than for **PP**. In contrast, for the isomer of **TBP** namely dinaphtho[*a,j*]perylene (**DNP**), the change of aromaticity between the closed- and the open-shell is lower than for **PP** since the biradical form has only one extra aromatic sextet (Figure 7).²⁹ Consequently, **DNP** is a closed-shell system in the ground state according to DFT calculation even though **DNP** has a biradical character y_0 of 57 % very closed to that of **TIPS-HEX** (58%) (see Table 1). The difference between **DNP** and **TIPS-HEX** is the absorption behavior. **DNP** has a λ_{\max} at 719 nm while **TIPS-HEX** absorbs at 740 nm. Accordingly, it seems that for benzenoid PAHs functionalized with TIPS-acetylene, the absorption cut-off should be around 740 nm to get a singlet biradical in the ground state.

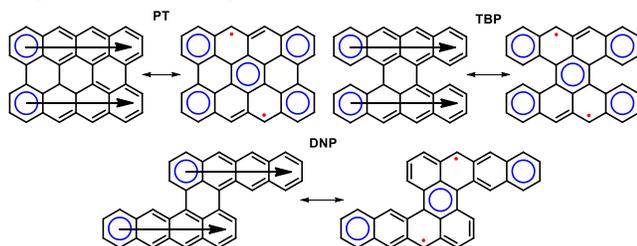


Figure 7. Resonance forms of **PT**, **TBP** and **DNP**.

It is noteworthy to know that for rylene ribbons, the concomitant opening of six naphthalenes is necessary to get an open-shell character while for anthrenes, the open-shell character occurs with the opening of three anthracenes with the formation of three extra aromatic sextets.^{13, 11a}

X-ray Crystallographic Analysis: Single crystals of **TT** were obtained by slow evaporation from a chloroform solution. The crystal has a triclinic symmetry with *P*-1 space group. The molecule is not perfectly planar but slightly wavy due to the fact that at the bay region, the torsion angle between carbons 4, 4a, 4b and 5 is about -9.36° (Figure 8). Consequently, the experimental bond lengths are not strictly comparable to that of the calculated ones. Nevertheless, the bond lengths at the zigzag edge show bond length alternation. Even though the bond lengths of bonds **a** (1.362 Å) and **h** (1.363 Å) are nearly identical in contrast to the calculated ones (1.356 Å for **a** and 1.367 Å for **h**), the bond **b** is much longer (1.446 Å) suggesting that the bonds **a** and **b** belong to a ring where the π -electrons are not delocalized but rather fixed. Moreover, even though the bonds lengths of bonds **e** (1.419 Å) and **f** (1.417 Å) are longer than that of the calculated ones (1.397 Å for **e** and 1.399 Å for **f**), they are also nearly identical.

Amazingly, like **TIPS-PEN**, **TT** adopts a 2-D “bricklayer” arrangement even if the TIPS groups are not in the middle of the π -skeleton (Figure 8).^{3a} The phenyl groups help at the arrangement of the molecules as they are in the middle of two TIPS of adjacent molecules. Due to the inversion center located at the middle of the stacked dimer, there are two alternating intermolecular center-to-center distances of 7.63 and 9.67 Å between adjacent molecules. The corresponding interplanar distances are 3.53 and 3.43 Å, respectively.

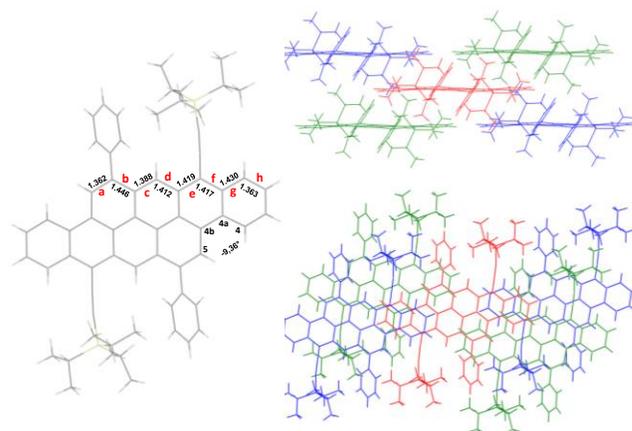


Figure 8. Experimental selected bond lengths and molecular packing of **TT**.

OFET Devices: Thanks to the stability of **TT** and **PP**, OFETs in solution-processed thin films were fabricated (see Supporting Information for details). Although organic semiconductors are intrinsically ambipolar and should be able to transport both holes and electrons, however, most PAHs including pentacene, only show unipolar p-type transport behavior in the standard device configuration using gold as the source and drain electrodes.³⁰ In contrast, the small energy gap and the amphoteric redox character of **TT** and **PP** allow both carrier injection and an ambipolar charge transport in bottom-gate/top-contact (BG/TC) devices. A BG/TC device with the **PP** layer formed by the solution-shearing method showed balanced hole ($0.07 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$) and electron ($0.02 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$) mobilities (Figure 9), which are more than one-order of magnitude higher than those previously reported for most biradical PAHs.^{8e, 10e, 10h} Since the mobilities of small molecules are strongly related to the crystallinity of the films, which depends on the preparation method such as spin-coating, drop-casting, dip-coating or solution-shearing, we observed a better hole mobility

for the drop-casted **PP** devices ($\mu_h = 0.21 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ and $\mu_e = 0.01 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$) owing to the larger crystalline domains (See Supporting Information). The corresponding **TT** devices also showed ambipolar transport with slightly lower mobilities ($\mu_h = 0.14 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ and $\mu_e = 0.006 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$). Further optimization resulted in better hole mobilities up to $1.4 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ and $0.77 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ for **PP** and **TT**, respectively, in TG/BC devices with CYTOP, which has a low dielectric constant and low interface trap sites. The increase of the hole mobilities in the large crystalline films makes observing the relatively low electron transport difficult. Since a single crystal of **PP** could not be obtained, the potential of **PP** has not been completely elucidated. However, the mobility of **PP** is much higher than that of polycrystalline **TIPS-PEN** and many other PAHs fabricated by standard solution processes. These results proved that low band gap materials based on acenoacenes with closed- and open-shell structures are excellent organic semiconductors.

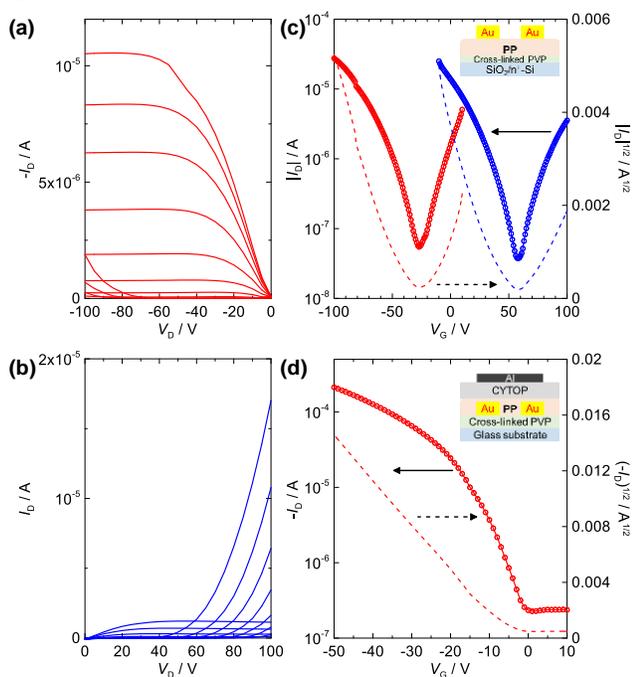


Figure 9. OFET devices of **PP**. (a) and (b) Output and (c) Transfer characteristics of a BG/TC device with a solution-sheared **PP** layer; (d) Transfer characteristics of a TG/BC device with a drop-casted **PP** layer.

CONCLUSION

We have reported the efficient synthesis of two peri-fused materials **TT** and **PP**. Suitably functionalized with TIPS-acetylene at the most reactive site and the presence of phenyl groups at the zigzag edge, **TT** and **PP** are low band gap materials of 1.79 and 1.42 eV, respectively, and exhibit remarkable high stability under ambient condition. Thanks to different experimental techniques and DFT calculations, we have shown that **TT** and **PP** have a closed-shell and open-shell configuration in the ground state, respectively. The open shell character of **PP** is related to the concomitant opening of two tetracenes with the recovering of two extra aromatic sextets and a small HOMO-LUMO energy gap (gap < 1.5 eV). Thanks to the high stability, thin films OFETs could be fabricated. In TG-BC configuration, **PP** shows remarkable hole mobility of $1.4 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ while **TT** exhibits a hole mobility of $0.77 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$. In configuration BG-TC, ambipolar behaviors for both were obtained with hole and electron mobilities of 0.21 and $0.01 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ for **PP** and 0.14 and $0.006 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ for **TT**. Since **PP** is a singlet biradical in the ground state and exhibits high stability and high mobility, it could be one of the

rare open shell materials suitable to be used in spintronics.^{7a, 31} The synthesis of other pentacenopentacene isomers and longer homologues is underway in our laboratory.

ASSOCIATED CONTENT

Supporting Information

Synthesis, characterization, crystal data, device fabrication and theoretical calculations. The Supporting Information is available free of charge on the ACS Publications website.

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Notes

The authors declare no competing financial interests.

ACKNOWLEDGMENT

M.F. and A.Y. acknowledge financial support from the Agence Nationale de la Recherche ANR-16-CE07-0024 (GATE). T.J.-O. has been an International Research Fellow of the Japanese Society for the Promotion of Science (JSPS) for three months through the JSPS Summer Program and acknowledges CNRS for PhD grant. M.M. and C.A. acknowledge financial support from JST ERATO Grant No. JPMJER1305 and JSPS KAKENHI Grant No. JP19H02790. The authors thank Dr. W. J. Potscavage, Jr., for assistance in data processing.

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