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## A NOVEL CONJUGATED DONOR-ACCEPTOR SYSTEM BASED ON TETRATHIAFULVALENE MERGING PYRENE UNIT: SYNTHESIS, PHYSICAL PROPERTIES AND THEORETICAL CALCULATIONS

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**Abstract** – A novel conjugated donor-acceptor compound (**1**) based on tetrathiafulvalene has been synthesized and fully characterized. Intramolecular charge transfer (ICT) in compound **1**, arising from HOMO-LUMO singlet transitions, has been experimentally evidenced through absorption/fluorescence spectra and theoretical calculations.

### INTRODUCTION

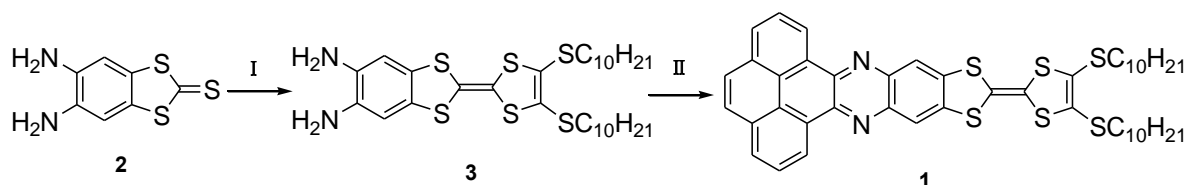
Among all kinds of molecular electron donor moieties, tetrathiafulvalene (TTF) and its derivatives have been received much attention as strong  $\pi$ -electron donors which are capable of forming persistent cation radical and dication species sequentially.<sup>1</sup> The molecular systems of D-A (donor-acceptor) compounds have been widely investigated especially in the last few years because of their potential applications in organic conducting materials,<sup>2</sup> molecular electronics,<sup>3</sup> optoelectronics,<sup>4</sup> photovoltaics,<sup>5</sup> and nonlinear optics.<sup>6</sup> Undoubtedly, the important feature in such compounds is the property of intramolecular charge transfer (ICT), which clearly depends on the nature of the linkage between the redox active units. In practice, D and A components are usually linked by saturated flexible or rigid  $\sigma$  spacers (D- $\sigma$ -A).<sup>7</sup> Thereby, the structural D and A moieties on which the HOMO and LUMO are localized and kept apart from each other, respectively. This hinders their strong coupling in the electronic ground and excited states. However, only a few compounds described so far have a direct link between the TTF unit and the acceptor, such as TTF-dipyridophenazine,<sup>8</sup> TTF-phenanthroline,<sup>9</sup> TTF-TCNQ type bithienoquinoxaline,<sup>10</sup>

TTF-perylenediimide,<sup>11</sup> thus leading to a more efficient intramolecular charge transfer and planar structures.

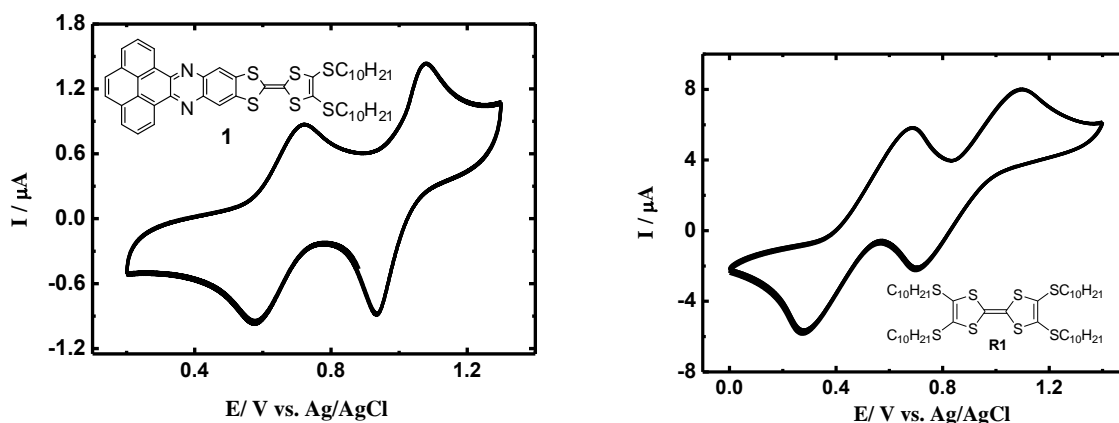
Pyrene skeleton has been attracted much interest for its electron-accepting and unique luminescence properties.<sup>12</sup> As a continuous work,<sup>8,13</sup> this paper presents the synthesis of a conjugated D-A compound **1** (Scheme 1), containing tetrathiafulvalene and pyrene unit, together with experimental and theoretical evidences for intramolecular charge transfer.

## RESULTS AND DISCUSSIONS

The synthetic<sup>8,14</sup> pathway of compound **1** is outlined in Scheme 1. In order to assure solubility, the TTF with dithiodecyl chains is employed. Compound **1** can be synthesized via direct condensation of pyrene-4,5-dione with 5,6-diamino-2-[4,5-bis(decylthio)-1,3-dithio-2-ylidene]-benzo[d]-1,3-dithiole (**3**) in AcOH. <sup>1</sup>H NMR spectroscopy, MS spectrometry and elemental analysis confirmed the structure of the D-A conjugated system.



**Scheme 1.** Synthetic route of compound **1**. (I) 4,5-bis-(decylthio)-1,3-dithiole-2-one (2 equiv), triethylphosphite/toluene, 120 °C, 35%; (II) pyrene-4,5-dione (1 equiv), AcOH, refluxed, 90%.

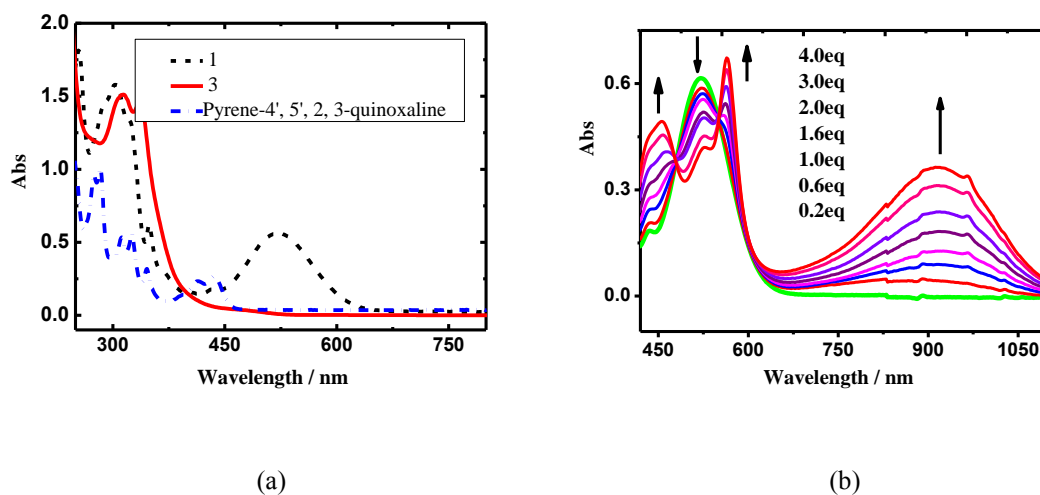


**Figure 1.** Cyclic voltammograms of **1** and **R1** in CH<sub>2</sub>Cl<sub>2</sub> (1.2 × 10<sup>-5</sup> M, scanning rate 100 mV/s) with platinum wires as working and counter electrodes, respectively, Ag/AgCl as reference electrode, and *n*-Bu<sub>4</sub>NPF<sub>6</sub> (0.1M) as supporting electrolyte.

The electrochemical properties of compound **1** are investigated by cyclic voltammetry together with reference compound **R1** for comparison (Figure 1). By comparing with compound **R1**, the first oxidation

potential of compound **1** is shifted from 0.48 to 0.65 V, and the second oxidation potential is shifted from 0.80 to 1.0 V. However, the oxidation waves of pyrene-4',5',2,3-quinoxaline and the reduction waves are absent maybe due to the measuring limitation in our experimental condition. These results can be rationalized in terms of electronic interaction between donor and acceptor moieties within the conjugated D-A system. Since TTF functional group of the molecule is involved in a delocalization path with pyrene-4',5',2,3-quinoxaline group, strong intramolecular electron interaction made the electron donating and electron accepting ability of molecule **1** decrease dramatically.

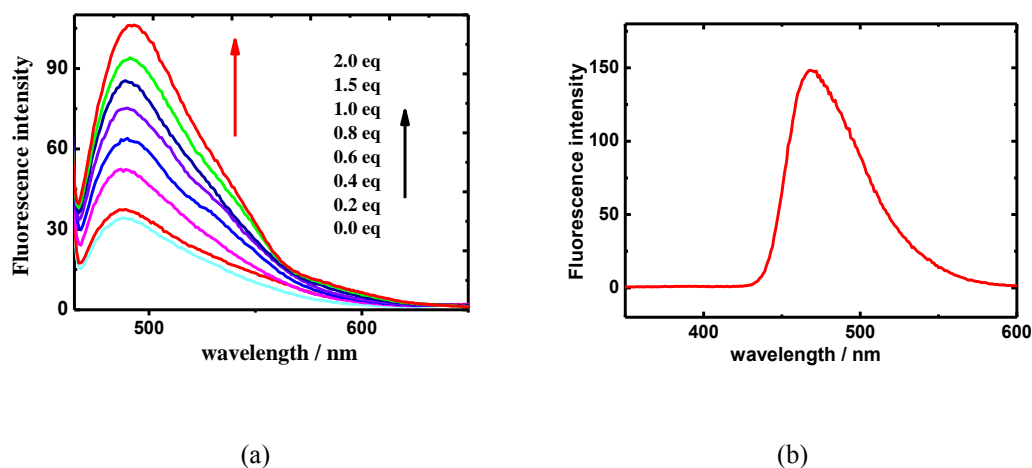
Figure 2(a) (green line) shows the UV-Vis absorption spectra of compound **1**. By comparison with the spectra of compound **3** and pyrene-4',5',2,3-quinoxaline, new electronic transitions (450-650nm, centered at 519 nm) can only be observed in **1**, which can be attributed to ICT transitions from donor to acceptor. The broad absorption band in this compound suggests a good electronic delocalization throughout the whole D-A system.<sup>15</sup> By oxidation (Figure 2(b)), the generated radical cation  $\mathbf{1}^{+\cdot}$  and dication  $\mathbf{1}^{2+}$  were characterized by the disappearance of the broad ICT band at 519 nm and the emergence of new bands centered at 454, 564 and 915 nm. The low-energy absorption at 915 nm is now assigned to an intrinsic absorption of the TTF radical cation ( $\text{TTF}^{+\cdot}$ ), which is red-shifted compared with those of other TTF derivatives.<sup>16</sup>



**Figure 2.** (a) Absorption spectra of compound **1** ( $1.7 \times 10^{-5}$  M in  $\text{CH}_2\text{Cl}_2$ ), **3** ( $5.7 \times 10^{-5}$  M in  $\text{CH}_2\text{Cl}_2$ ) and pyrene-4',5',2,3-quinoxaline ( $3.6 \times 10^{-5}$  M in  $\text{CH}_2\text{Cl}_2$ ). (b) Absorption spectra of compound **1** ( $1.7 \times 10^{-5}$  M) in  $\text{CH}_2\text{Cl}_2$  with the increasing amounts of  $\text{Fe}^{3+}$  [ $\text{Fe}(\text{ClO}_4)_3 \cdot 6\text{H}_2\text{O}$ ,  $1.3 \times 10^{-3}$  M, in MeOH].

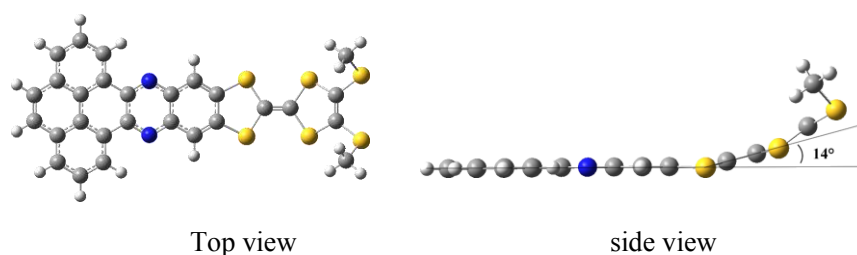
As comparing with pyrene-4',5',2,3-quinoxaline, compound **1** shows much weaker fluorescence (Figure 3). The fluorescence of compound **1** is quenched maybe due to the photo-induced electron transfer between TTF and pyrene-4',5',2,3-quinoxaline unit.<sup>17</sup> After addition of  $\text{Fe}^{3+}$ , then the TTF unit in compound **1** is oxidized to its corresponding cation and further lead to the photo-induced electron transfer

between TTF and pyrene-4',5',2,3-quinoxaline unit is arrested, then the fluorescence of compound **1** starts to increase. The result suggesting that compound **1** is one of good candidates for redox-driven fluorescence molecular switch and maybe has some potential applications in future molecular devices.<sup>18</sup>

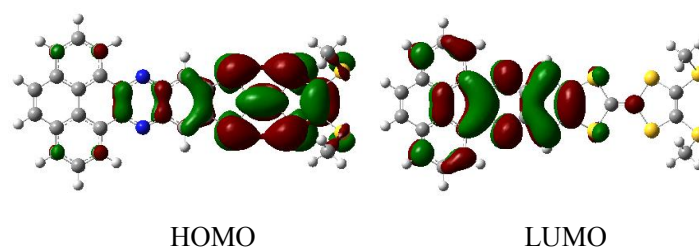


**Figure 3.** (a) Fluorescence spectra of **1** ( $1.7 \times 10^{-5}$  M) in  $\text{CH}_2\text{Cl}_2$  with the increasing amounts of  $\text{Fe}^{3+}$  [ $\text{Fe}(\text{ClO}_4)_3 \cdot 6\text{H}_2\text{O}$ ,  $1.3 \times 10^{-3}$  M, in MeOH]. (b) Fluorescence spectra of pyrene-4',5',2,3-quinoxaline ( $3.6 \times 10^{-5}$  M in  $\text{CH}_2\text{Cl}_2$ ).

To get further insights into the molecular structure and electronic distribution of the D-A system, density functional theory (DFT) calculations are performed in gas phase on a B3LYP/6-31 G (d, p) level with Gaussian 09.<sup>19</sup> The peripherally longer alkyl groups are removed for simplification as they are not expected to have significantly affects on the conformation and energy levels. The simplified molecule is labeled as model **1**.



**Figure 4.** Optimized geometry structure of model **1**



**Figure 5.** Frontier molecular orbitals of model **1**

The optimized structure afforded by the calculations is displayed in Figure 4, which shows that the whole molecule is nearly in an identical plane with a dihedral angle between pyrene-4',5',2,3-quinoxaline and TTF ring as approximate 14°. The frontier molecular orbitals are given in Figure 5. For the HOMO of model **1**, the orbital is mainly located over the TTF moiety with small contribution of pyrene-4',5',2,3-quinoxaline group; while for the LUMO, the orbital is mainly located on pyrene-4',5',2,3-quinoxaline group, both in good agreement with their expected electron-donor and electron-acceptor properties. This is a prerequisite to ICT transition and the positive shift of the oxidation potentials of the donor subunit.<sup>20</sup>

In summary, a novel D-A compound (**1**) based on tetrathiafulvalene has been synthesized and characterized. Cyclic voltammetry experiment demonstrates that the oxidation potentials of compound **1** are higher than those of **R1**, which suggests a good electronic communication between the donor and acceptor. Intramolecular charge transfer (ICT) in compound **1**, arising from HOMO-LUMO singlet transitions, has been experimentally evidenced through UV-Vis spectra and theoretical calculations. The fluorescence properties of compound **1** suggest that this system has attractive potential application as molecular switch.

## EXPERIMENTAL

### 1. General

Unless otherwise indicated, all reagents were purchased from commercial sources and used without additional purification. All reactions were performed under N<sub>2</sub>. Toluene and triethylphosphite were dried and distilled over Na. Compound **2**,<sup>8</sup> pyrene-4,5-dione<sup>14</sup> were prepared as previously described. <sup>1</sup>H NMR spectra were obtained on a Bruker AC 300 spectrometer operating at 300 MHz; Chemical shifts  $\delta$  were calibrated against TMS as an internal standard. Mass spectrum was recorded on a FTMS 4.7T BioAPEX II for MALDI ionisation method. Elemental analyses were performed on an EA1110 Elemental Analyzer CHN Carlo Erba Instruments. FT-IR data were collected on a Perkin Elmer One FT-IR spectrometer. Absorption spectra were measured using a Hitachi (model U-3010) UV-Vis spectrophotometer. Fluorescence measurements were carried out using a Hitachi (model F-4500) spectrophotometer in a 1 cm quartz cell. Cyclic voltammetry measurements were carried out on a CHI660 system with platinum as the working and counter electrodes, Ag/AgCl as the reference electrode and *n*-Bu<sub>4</sub>NPF<sub>6</sub> as the supporting electrolyte.

### 2. Synthesis of 4',5'-bis(decylthio)tetrathiafulvenyl-3,4-(pyrene-4',5',2,3-quinoxaline) **1**:

A solution of compound **3** (100 mg, 0.16 mmol) and pyrene-4,5-dione (36.9 mg, 0.23 mmol) in AcOH (20 mL) was refluxed for 3 h under N<sub>2</sub>. After filtration, the precipitate was collected and purified by

chromatography (CH<sub>2</sub>Cl<sub>2</sub>) to give compound **1** as a mauve solid (123 mg, 90%). <sup>1</sup>H NMR (300MHz, CDCl<sub>3</sub>) δ: 0.86 (T, *J* = 6.6, 6H, 2×CH<sub>3</sub>), 1.26 (M, 24H, 12×CH<sub>2</sub>), 1.43 (M, 4H, 2×CH<sub>2</sub>), 1.60 (M, 4H, 2×CH<sub>2</sub>), 2.86 (T, *J* = 7.4, 4H, 2×SCH<sub>2</sub>), 8.00 (S, 2H, aromatic), 8.25 (M, 2H, aromatic), 8.05 (M, 4H, aromatic), 9.42 (T, 2H, aromatic). Selected IR data (cm<sup>-1</sup>, KBr pellet): 2916, 2849, 1624, 1441, 1424, 1352, 1205, 1168, 1103, 858, 831, 718; MS (MALDI-TOF) calcd for C<sub>46</sub>H<sub>52</sub>N<sub>2</sub>S<sub>6</sub>: 824.25, found 824.25 [M]<sup>+</sup>; Anal. Calcd for C<sub>46</sub>H<sub>52</sub>N<sub>2</sub>S<sub>6</sub>: C, 66.94; H, 6.35; N, 3.39. Found: C, 67.29; H, 6.50; N, 3.26.

### 3. Synthesis of 5,6-diamino-2-(4,5-bis(decylthio)-1,3-dithio-2-ylidene)benzo[d]-1,3-dithiole **3**:

Triethylphosphite (15 mL) was added to a solution of 4,5-bis(decylthio)-1,3-dithiole-2-one (214 mg, 1.0 mmol) and 5,6-diaminobenzene-1,3-dithiole-2-thione (**2**) (926 mg, 2.0 mmol) in toluene (15 mL) under N<sub>2</sub>. The mixture was heated to 120 °C and stirred for 3 h. After the reaction, the excess solvent was removed under vacuum to afford an oily red residue, which was subjected to chromatography (silica gel, CH<sub>2</sub>Cl<sub>2</sub>/EtOAc (1/8, v/v)). The crude product was further purified by recrystallization from EtOAc to give compound **3** as a yellow solid (220 mg, 35%). mp 100-102 °C; <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>): δ = 0.78-0.83 (t, 6H), 1.19-1.35 (m, 28H), 1.47-1.54 (m, 4H), 2.78-2.83 (t, 4H), 4.68 (s, 4H), 6.52 (s, 2H); Selected IR data (cm<sup>-1</sup>, KBr pellet): 3378, 2955, 2919, 2850, 1483, 1291, 859; EI-MS *m/z* (relative intensity): calcd for C<sub>30</sub>H<sub>48</sub>N<sub>2</sub>S<sub>6</sub>: 628, found 628 [M]<sup>+</sup>; Anal. Calcd for C<sub>30</sub>H<sub>48</sub>N<sub>2</sub>S<sub>6</sub>: C, 57.28; H, 7.69; N, 4.45. Found: C, 57.45; H, 7.77; N, 4.36.

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## REFERENCES

- (a) D. F. Perepichka, M. R. Bryce, E. J. L. McInnes, and J. P. Zhao, *Org. Lett.*, 2001, **3**, 1431; (b) G. Cosquer, F. Pointillart, Y. Le Gal, S. Golhen, O. Cador, and L. Ouahab, *Dalton Trans.*, 2009, 3495; (c) M. Bendikov, F. Wudl, and D. F. Perepichka, *Chem. Rev.*, 2004, **104**, 4891; (d) F. Dumur, N. Gautier, N. Gallego-Planas, Y. Şahin, E. Levillain, N. Mercier, P. Hudhomme, M. Masino, A. Girlando, V. Lloveras, J. Vidal-Gancedo, J. Veciana, and C. Rovira, *J. Org. Chem.*, 2004, **69**, 2164; (e) N. Gautier, F. Dumur, V. Lloveras, J. Vidal-Gancedo, J. Veciana, C. Rovira, and P. Hudhomme, *Angew. Chem. Int. Ed.*, 2003, **42**, 2765; (f) B. T. Zhao, W. B. Guo, and P. Z. Hu, *Heterocycles*, 2010, **81**, 1661.
- M. Iyoda, M. Hasegawa, and Y. Miyake, *Chem. Rev.*, 2004, **104**, 5085.

3. (a) J. K. Sørensen, M. Vestergaard, A. Kadziola, K. Kilså, and M. B. Nielsen, [Org. Lett., 2006, 8, 1173](#); (b) D. Canevet, M. Salle, G. Zhang, D. Zhang, and D. Zhu, [Chem. Commun., 2009, 2245](#); (c) K. A. Nielsen, G. H. Sarova, L. Martín-Gomis, F. Fernández-Lázaro, P. C. Stein, L. Sanguinet, E. Levillain, J. L. Sessler, D. M. Guldi, Á. Sastre-Santos, and J. O. Jeppesen, [J. Am. Chem. Soc., 2007, 130, 460](#).
4. (a) D. C. Magri, [New J. Chem., 2009, 33, 457](#); (b) H. Imahori, [Org. Biomol. Chem., 2004, 2, 1425](#).
5. (a) N. Martín, L. Sánchez, M. Á. Herranz, B. Illescas, and D. M. Guldi, [Acc. Chem. Res., 2007, 40, 1015](#); (b) S. Wenger, P.-A. Bouit, Q. Chen, J. I. Teuscher, D. D. Censo, R. Humphry-Baker, J.-E. Moser, J. L. Delgado, N. Martín, S. M. Zakeeruddin, and M. Grätzel, [J. Am. Chem. Soc., 2010, 132, 5164](#).
6. (a) B. Insuasty, C. Atienza, C. Seoane, N. Martín, J. Garín, J. Orduna, R. Alcalá, and B. Villacampa, [J. Org. Chem., 2004, 69, 6986](#); (b) M. González, J. L. Segura, C. Seoane, N. Martín, J. Garín, J. Orduna, R. Alcalá, B. Villacampa, V. Hernández, and J. T. López Navarrete, [J. Org. Chem., 2001, 66, 8872](#); (c) A. I. de Lucas, N. Martín, L. Sánchez, C. Seoane, R. Andreu, J. Garín, J. Orduna, R. Alcalá, and B. Villacampa, [Tetrahedron, 1998, 54, 4655](#); (d) R. Andreu, A. I. de Lucas, J. Garín, N. Martín, J. Orduna, L. Sánchez, and C. Seoane, [Synth. Met., 1997, 86, 1817](#); (e) J. F. Lamère, I. Malfant, A. Sournia-Saquet, P. G. Lacroix, J. M. Fabre, L. Kaboub, T. Abbaz, A.-K. Gouasmia, I. Asselberghs, and K. Clays, [Chem. Mater., 2007, 19, 805](#).
7. (a) Y. Zhang, L.-Z. Cai, C.-Y. Wang, G.-Q. Lai, and Y.-J. Shen, [New J. Chem., 2008, 32, 1968](#); (b) J. Baffreau, F. Dumur, and P. Hudhomme, [Org. Lett., 2006, 8, 1307](#); (c) E. Tsiperman, J. Y. Becker, V. Khodorkovsky, A. Shames, and L. Shapiro, [Angew. Chem. Int. Ed., 2005, 44, 4015](#); (d) D. F. Perepichka, M. R. Bryce, C. Pearson, M. C. Petty, E. J. L. McInnes, and J. P. Zhao, [Angew. Chem. Int. Ed., 2003, 42, 4636](#); (e) R. B. Hou, H. Qiu, T. Chen, and B. Z. Yin, [Heterocycles, 2009, 78, 1799](#).
8. C. Jia, S.-X. Liu, C. Tanner, C. Leiggenger, A. Neels, L. Sanguinet, E. Levillain, S. Leutwyler, A. Hauser, and S. Decurtins, [Chem. Eur. J., 2007, 13, 3804](#).
9. (a) C. Goze, N. Dupont, E. Beitler, C. Leiggenger, H. Jia, P. Monbaron, S.-X. Liu, A. Neels, A. Hauser, and S. Decurtins, [Inorg. Chem., 2008, 47, 11010](#); (b) L. K. Keniley, L. Ray, K. Kovnir, L. A. Dellinger, J. M. Hoyt, and M. Shatruk, [Inorg. Chem., 2010, 49, 1307](#).
10. X. Guégano, Alexander L. Kanibolotsky, C. Blum, S. F. L. Mertens, S.-X. Liu, A. Neels, H. Hagemann, P. J. Skabara, S. Leutwyler, T. Wandlowski, A. Hauser, and S. Decurtins, [Chem. Eur. J., 2009, 15, 63](#).
11. M. Jaggi, C. Blum, N. Dupont, J. Grilj, S.-X. Liu, J. Hauser, A. Hauser, and S. Decurtins, [Org. Lett., 2009, 11, 3096](#).

12. Z. Ning, Q. Zhang, W. Wu, H. Pei, B. Liu, and H. Tian, *J. Org. Chem.*, 2008, **73**, 3791.
13. (a) C. Jia, D. Zhang, W. Xu, and D. Zhu, *Org. Lett.*, 2001, **3**, 1941; (b) C. Jia, D. Zhang, W. Xu, X. Shao, and D. Zhu, *Synth. Met.*, 2003, **137**, 1345; (c) C. Jia, D. Zhang, C.-M. Liu, W. Xu, H. Hu, and D. Zhu, *New J. Chem.*, 2002, **26**, 490; (d) C. Jia, S.-X. Liu, C. Tanner, C. Leiggenger, L. Sanguinet, E. Levillain, S. Leutwyler, A. Hauser, and S. Decurtins, *Chem. Commun.*, 2006, 1878; (e) C. Jia, S.-X. Liu, A. Neels, G. Labat, and S. Decurtins, *Polyhedron*, 2005, **24**, 3032; (f) C. Jia, S.-X. Liu, C. Ambrus, A. Neels, G. Labat, and S. Decurtins, *Inorg. Chem.*, 2006, **45**, 3152.
14. J. Hu, D. Zhang, and F. W. Harris, *J. Org. Chem.*, 2005, **70**, 707.
15. Y.-P. Zhao, L.-Z. Wu, G. Si, Y. Liu, H. Xue, L.-P. Zhang, and C.-H. Tung, *J. Org. Chem.*, 2007, **72**, 3632.
16. (a) Y. Zeng, G. Zhang, D. Zhang, and D. Zhu, *J. Org. Chem.*, 2009, **74**, 4375; (b) H. Wu, D. Zhang, L. Su, K. Ohkubo, C. Zhang, S. Yin, L. Mao, Z. Shuai, S. Fukuzumi, and D. Zhu, *J. Am. Chem. Soc.*, 2007, **129**, 6839.
17. G. Zhang, D. Zhang, X. Guo, and D. Zhu, *Org. Lett.*, 2004, **6**, 1209.
18. S. Leroy-Lhez, J. Baffreau, L. Perrin, E. Levillain, M. Allain, M.-J. Blesa, and P. Hudhomme, *J. Org. Chem.*, 2005, **70**, 6313.
19. R. A. Gaussian 09, M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, J. M. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, O. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski, and D. J. Fox, Gaussian, Inc., Wallingford CT, 2009.
20. Q.-Y. Zhu, L.-B. Huo, Y.-R. Qin, Y.-P. Zhang, Z.-J. Lu, J.-P. Wang, and J. Dai, *J. Phys. Chem. B*, 2010, **114**, 361.