

A highly efficient organic sensitizer for dye-sensitized solar cells†

Suyoung Hwang,^a Jung Ho Lee,^a Chanmoo Park,^a Hoinglae Lee,^a Chaekyu Kim,^a Chiyong Park,^a Mi-Hyeon Lee,^b Wanin Lee,^b Jihee Park,^c Kyungkon Kim,^c Nam-Gyu Park^{*c} and Chulhee Kim^{*a}

Received (in Cambridge, UK) 28th June 2007, Accepted 22nd August 2007

First published as an Advance Article on the web 14th September 2007

DOI: 10.1039/b709859f

We have synthesized a highly efficient organic dye for a dye-sensitized solar cell; the overall solar-to-energy conversion efficiency was 9.1% at AM 1.5 illumination (100 mW cm^{-2}); short-circuit current density (J_{sc}) = 18.1 mA cm^{-2} , open circuit photovoltage (V_{oc}) = 743 mV and fill factor (ff) = 0.675.

Dye-sensitized solar cells (DSSCs) have been a target of intensive research in recent years due to their capability to convert solar light into electricity with low materials and production costs.^{1,2} One of the key material issues in DSSCs is the photosensitizer, which can exhibit high efficiencies in the conversion of solar energy into electricity. Ru complex photosensitizers such **N3**, **N719** and Black dyes exhibit efficiencies higher than 10%.^{1,2} The performance of DSSCs based on organic dyes have also recently been remarkably improved. Therefore, it is expected that organic dyes could compete with Ru complexes in the near future.³

Organic dyes have many advantages as photosensitizers, such as large molar extinction coefficient, control of absorption wavelength, facile design and synthesis, and lower cost than Ru complexes.⁴ Efficient organic dyes such as coumarin, indoline, phthalocyanine and conjugated oligo-ene dyes have been reported.^{3–7} However, as yet, the use of organic dyes for DSSCs has a limitation due to their relatively low efficiency compared Ru complexes. Herein, we report a highly efficient dye which exhibits 9.1% overall solar-to-electric energy conversion efficiency.

As shown in Fig. 1, we have prepared the organic dye **TA-St-CA**, which contains a π -conjugated oligo-phenylenevinylene unit with an electron donor–acceptor moiety for intramolecular charge transfer and a carboxyl group as an anchoring unit for the attachment of the dye onto TiO_2 nanoparticles.†

The UV-vis spectrum of **TA-St-CA** in Fig. 2 shows an absorption maximum at 386 nm in ethanol ($\epsilon = 31600 \text{ M}^{-1} \text{ cm}^{-1}$).

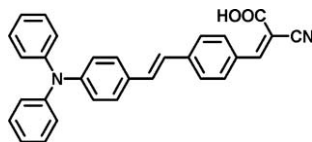


Fig. 1 Molecular structure of **TA-St-CA** dye.

^aDepartment of Polymer Science and Engineering, Inha University, Incheon 402-751, Korea. E-mail: chk@inha.ac.kr; Fax: +82 32-865-5178; Tel: +82 32-860-7487

^bDepartment of Chemistry, Inha University, Incheon 402-751, Korea

^cCenter for Energy Materials, Korea Institute of Science and Technology (KIST), Seoul 136-791, Korea. E-mail: npark@kist.re.kr; Fax: +82 2-958-5309; Tel: +82 2-958-5365

† Electronic supplementary information (ESI) available: Detailed synthetic procedures and spectral characterization. See DOI: 10.1039/b709859f

The HOMO level of the dye was estimated from its oxidation potential by using cyclic voltammetry, as shown in Fig. 3. The measurement was carried out using a Pt electrode in acetonitrile solution containing 0.1 M tetrabutylammonium tetrafluoroborate as an electrolyte. A standard ferrocene/ferrocenium (Fc/Fc^+) system was used to calibrate the redox peak. The HOMO level was determined from the oxidation potential value with respect to ferrocene using an Ag/AgCl reference electrode. The HOMO level of **TA-St-CA** was -5.2 eV . The band gap of **TA-St-CA** was 2.4 eV , which was estimated from the absorption edge of the absorption spectrum. The LUMO of **TA-St-CA**, calculated from the HOMO and optical band gap, was -2.8 eV , which provides a favorable energy matching with the conduction level of TiO_2 particles for the electron transfer from the LUMO of the dye to the conduction band of TiO_2 . The HOMO, LUMO and band gap energy results are summarized in Table 1.⁸

Fig. 4 shows the optimized molecular structure and frontier molecular orbitals of **TA-St-CA**. At the HOMO of the dye, electrons are homogeneously distributed in the π -conjugated oligo-phenylenevinylene unit and the triphenylamine electron donor

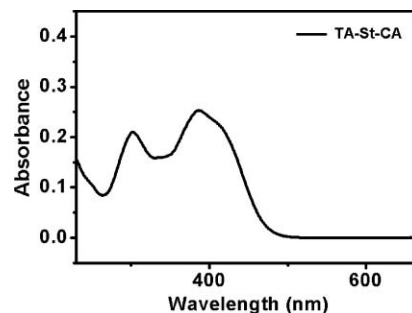


Fig. 2 Absorption spectrum of **TA-St-CA** in ethanol.

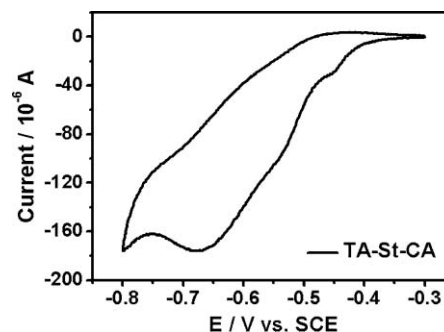


Fig. 3 Cyclic voltammogram of **TA-St-CA** measured at a sweep rate of 20 mV sec^{-1} .

Table 1 Electrochemical properties and photovoltaic performance of TA-St-CA

Dye	$E_{\text{HOMO}}/\text{eV}^a$	$E_{\text{bandgap}}/\text{eV}^b$	$E_{\text{LUMO}}/\text{eV}^c$	$J_{\text{sc}}/\text{mA cm}^{-2}$	V_{oc}/mV	ff	η (%)
TA-St-CA	−5.2	2.4	−2.8	18.1	743	0.675	9.1
N719	−5.6 ^d	2.6 ^d	−3.0 ^d	19.9	769	0.657	10.1

^a Estimated from the oxidation potential, determined by cyclic voltammetry. ^b Estimated from the edge of absorption spectrum. ^c Calculated from the band gap and HOMO value. ^d See ref. 8.

moiety. However, at the LUMO level, with electronic excitation, intramolecular charge transfer induced electron movement from the donor site to the acceptor moiety of the dye. Therefore, excited electrons can be injected into the conduction band of TiO_2 via the carboxyl group adjacent to the cyano group.

A TA-St-CA DSSC was prepared, where a TiO_2 film was composed of a double layer, with a $\sim 5 \mu\text{m}$ thick 400 nm particle film over-coated onto a 10 μm thick 20 nm particle film on FTO glass (Pilkington TEC8, $\sim 8 \Omega/\text{sq}$). The annealed TiO_2 film was further treated with 0.2 M TiCl_4 for 22 h at ambient temperature. Dye adsorption was performed with a 0.5 mM TA-St-CA solution in ethanol. For comparison, a N719 DSSC was prepared by the same procedure. Photocurrent and voltage were measured using a solar simulator equipped with a 1000 W ozone-free Xenon lamp and AM 1.5 G filter (Oriel), where the light intensity was adjusted with an NREL-calibrated Si solar cell with a KG-5 filter to 1 sunlight intensity (100 mW cm^{-2}).

The photocurrent density–photovoltage (I – V) curve is shown in Fig. 5. The short circuit current density (J_{sc}) and open circuit voltage (V_{oc}) of TA-St-CA were 18.1 mA cm^{-2} and 743 mV, respectively, with a fill factor (ff) of 0.675. The overall conversion efficiency (η) for TA-St-CA was 9.1%, under which conditions the efficiency of N719 was 10.1%.

Fig. 6 shows the incident photon-to-current conversion efficiency (IPCE) data for TA-St-CA, where the IPCE data was collected at a low chopping speed of 10 Hz under a monochromatic beam generated by a 75 W Xenon light source. The IPCE value in the range 400 to 550 nm reached around 80%, which is higher than that for N719. The red-shift in the IPCE of TA-St-CA in Fig. 6, compared to the absorption spectrum of TA-St-CA in Fig. 1, can be ascribed to aggregation of the dye on titania.

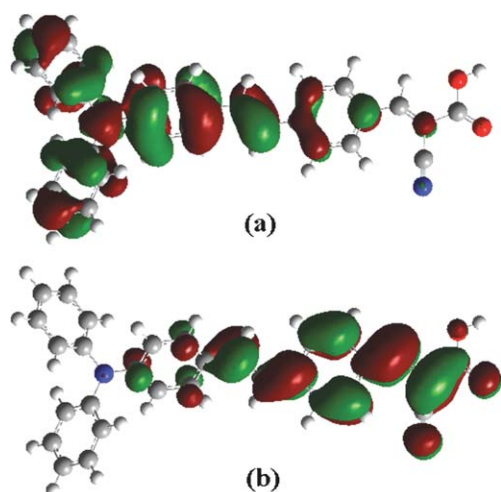


Fig. 4 The frontier (a) HOMO and (b) LUMO orbitals of TA-St-CA optimized with TD-DFT at the B3LYP/6-31+G(d) level.

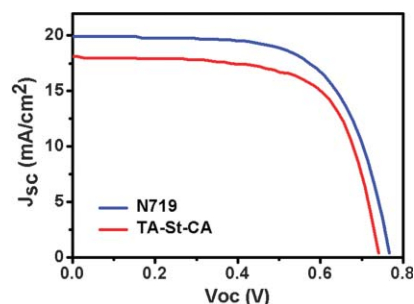


Fig. 5 Photocurrent density–photovoltage curve of TA-St-CA under AM 1.5 G radiation (100 mW cm^{-2}) (0.7 M *N*-methyl-*N*-butyl imidazolium iodide, 0.2 M LiI, 50 mM iodine, 0.5 M *tert*-butylpyridine in acetonitrile/3-methoxypropionitrile (50 : 50 V/V)).

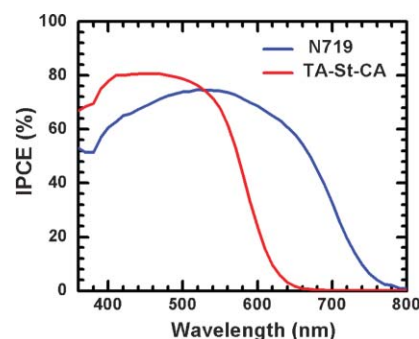


Fig. 6 IPCE spectrum for a DSSC based on TA-St-CA.

We are currently investigating the solar cell performance of π -conjugated donor–acceptor sensitizers having strong absorptions with wavelengths longer than 600 nm to achieve higher efficiencies. In addition, a tandem cell with TA-St-CA and N719 is under investigation because TA-St-CA and N719 showed good performance over different wavelength ranges. The detailed cell structure and photovoltaic characteristics will be published elsewhere.

This work was supported by a KIST internal project. We thank Dr Jae Woong Yu for help with the experiments.

Notes and references

- (a) B. O'Regan and M. Grätzel, *Nature*, 1991, **353**, 737; (b) M. Grätzel, *Photochem. Photobiol.*, 2003, **4**, 145; (c) M. Grätzel, *Photochem. Photobiol.*, 2004, **164**, 3.
- (a) M. K. Nazeeruddin, S. M. Zakeeruddin, R. Humphry-Baker, M. Jirousek, P. Liska, N. Vlachopoulos, V. Shklover, C. Fischer and M. Grätzel, *Inorg. Chem.*, 1999, **38**, 6298; (b) A. Hagfeldt and M. Grätzel, *Chem. Rev.*, 1995, **95**, 49; (c) A. Hagfeldt and M. Grätzel, *Acc. Chem. Res.*, 2000, **33**, 269; (d) M. Grätzel, *Nature*, 2001, **414**, 338; (e) M. K. Nazeeruddin, P. Péchy, T. Renouard, S. M. Zakeeruddin, R. Humphry-Baker, P. Comte, P. Liska, L. Cevey, E. Costa, V. Shklover, L. Spiccia, G. B. Deacon, C. A. Bignozzi and M. Grätzel, *J. Am. Chem. Soc.*, 2001, **123**, 1613.

- 3 (a) S. Ferrere, A. Zaban and B. A. Gregg, *J. Phys. Chem. B*, 1997, **101**, 4490; (b) K. Hara, T. Horiguchi, T. Kinoshita, K. Sayama, H. Sugihara and H. Arakawa, *Sol. Energy Mater. Sol. Cells*, 2000, **64**, 115; (c) K. Sayama, K. Hara, N. Mori, M. Satsuki, S. Suga, S. Tsukagoshi, Y. Abe, H. Sugihara and H. Arakawa, *Chem. Commun.*, 2000, 1173; (d) K. Sayama, K. Hara, S. Tsukagoshi, Y. Abe, Y. Ohga, A. Shinpo, S. Suga and H. Arakawa, *New J. Chem.*, 2001, **25**, 200; (e) S. Kim, J. K. Lee, S. O. Kang, J. Ko, J.-H. Yum, S. Fantacci, F. D. Angelis, D. D. Censo, M. K. Nazeeruddin and M. Grätzel, *J. Am. Chem. Soc.*, 2006, **128**, 16701.
- 4 (a) K. Hara, Z. S. Wang, T. Sato, A. Furube, R. Katoh, H. Sugihara, Y. Dan-oh, C. Kasada, A. Shinpo and S. Suga, *J. Phys. Chem. B*, 2005, **109**, 15476; (b) K. Hara, T. Sato, R. Katoh, A. Furube, Y. Ohga, A. Shinpo, S. Suga, K. Sayama, H. Sugihara and H. Arakawa, *J. Phys. Chem. B*, 2003, **107**, 597; (c) D. P. Hagberg, T. Edvinsson, T. Marinado, G. Boschloo, A. Hagfeldt and L. Sun, *Chem. Commun.*, 2006, 2245.
- 5 (a) K. Hara, Y. Tachibana, Y. Ohga, A. Shinpo, S. Suga, K. Sayama, H. Sugihara and H. Arakawa, *Sol. Energy Mater. Sol. Cells*, 2003, **77**, 89; (b) K. Hara, M. Kurashige, S. Ito, A. Shinpo, S. Suga, K. Sayama and H. Arakawa, *Chem. Commun.*, 2003, 252; (c) K. Hara, Y. Dan-oh, C. Kasada, Y. Ohga, A. Shinpo, S. Suga, K. Sayama and H. Arakawa, *Langmuir*, 2004, **20**, 4205; (d) K. Hara, T. Sato, R. Katoh, A. Furube, T. Yoshihara, M. Kurashige, S. Ito, A. Shinpo, S. Suga and H. Arakawa, *Adv. Funct. Mater.*, 2005, **15**, 246; (e) T. Horiuchi, H. Miura, K. Sumioka and S. Uchida, *J. Am. Chem. Soc.*, 2004, **126**, 12218; (f) M. Liang, W. Xu, F. Cai, P. Chen, B. Peng, J. Chen and Z. Li, *J. Phys. Chem. C*, 2007, **111**, 4465.
- 6 T. Kitamura, M. Ikeda, K. Shigaki, T. Inoue, N. A. Anderson, X. Ai, T. Lian and S. Yanagida, *Chem. Mater.*, 2004, **16**, 1806.
- 7 D. P. Hagberg, T. Edvinsson, T. Marinado, G. Boschloo, A. Hagfeldt and L. Sun, *Chem. Commun.*, 2006, 2245.
- 8 M. K. Nazeeruddin, F. D. Angelis, S. Fantacci, A. Selloni, G. Viscardi, P. Liska, S. Ito, B. Takeru and M. Grätzel, *J. Am. Chem. Soc.*, 2005, **127**, 16835.

Find a SOLUTION

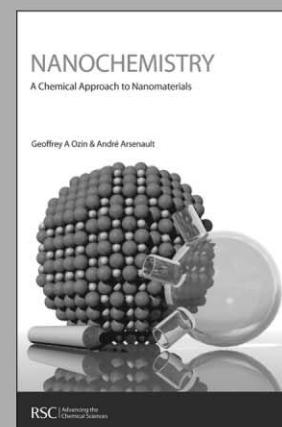
... with books from the RSC

Choose from exciting textbooks, research level books or reference books in a wide range of subject areas, including:

- Biological science
- Food and nutrition
- Materials and nanoscience
- Analytical and environmental sciences
- Organic, inorganic and physical chemistry

Look out for 3 new series coming soon ...

- RSC Nanoscience & Nanotechnology Series
- Issues in Toxicology
- RSC Biomolecular Sciences Series



RSC | Advancing the
Chemical Sciences

www.rsc.org/books