

Dynamic Rapid Synthesis of Bis(2,2'-bipyridine)nitrate Zinc (II) Nitrate Using a Microwave Method and its Application to Dye-Sensitized Solar Cells (DSSC)

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This study examined the synthesis of the crystal structure of bis(2,2'-bipyridine)nitrate zinc (II) nitrate, $[\text{Zn}(\text{bipy})_2(\text{NO}_3)]^+\text{NO}_3^-$ using a microwave treatment at 300 W and 60 Hz for the application to dye-sensitized solar cells. The simulated complex structure of the complex was optimized with the density functional theory calculations for the UV-vis spectrum of the ground state using Gaussian 03 at the B3LYP/LANL2DZ level. The structure of the acquired complex was expected a penta-coordination with four nitrogen atoms of bipyridine and the oxygen bond of the NO_3^- ion. The reflectance UV-vis absorption spectra exhibited two absorptions (L-L transfers) that were assigned to the transfers from the ligand (σ , π) of NO_3^- to the ligand (σ^* , π^*) of pyridine at around 200 - 350 nm, and from the non-bonding orbital (n) of O in NO_3^- to the p-orbital of pyridine at around 450 - 550 nm, respectively. The photoelectric efficiency was approximately 0.397% in the dye-sensitized solar cells with the nanometer-sized TiO_2 at an open-circuit voltage (Voc) of 0.39 V, a short-circuit current density (Jsc) of 1.79 mA/cm², and an incident light intensity of 100 mW/cm².

Key Words: Bis(2,2'-bipyridine)nitrate zinc (II) nitrate, Microwave treatment, Dye-sensitized solar cells, Photoelectric efficiency

Introduction

Dye-sensitized solar cells (DSSC) have extensively been studied because of their attractive advantages, such as their low cost, less toxic manufacturing process, easy scale-up, light weight and potential use of flexible panels, compared to conventional p-n junction devices.^{1,2} An important element of DSSC materials is the dye compound, which consists of conjugated π -electrons and must exhibit the following four properties: 1) Electrons must be generated from the HOMO to the LUMO ($\delta \rightarrow \pi$ electron transition) of the dye molecules through light absorption, and injected to the LUMO at TiO_2 electrode. 2) All light in the visible area must be absorbed. 3) The chemical combination of the solid oxide (TiO_2) and the dye molecule must be strong. 4) The dye compound must be stable to light and heat. Particularly, the Ru complex is a typical example of a commercial dye that possesses all of the required properties for application in DSSCs.³⁻⁵ The high efficiency of the Ru complex arises because the electrons of both the singlet excited state and the excited triplet state can be injected into the conduction band of the semiconductor electrode (TiO_2). Especially, Nazeeruddin *et al.*⁶ studied a high molar extinction coefficient charge transfer ruthenium sensitizer $[\text{Ru}(\text{4,4'-dicarboxylic acid-4'-carboxylate-2,2'-bipyridine})(\text{4,4'-di-(2-(3,6-dimethoxyphenyl)ethenyl)-2,2'-bipyridine})(\text{NCS})_2]$ that was anchored onto a nanocrystalline TiO_2 films. This sensitizer achieved a very efficient sensitization in the visible region under AM 1.5 sunlight at a short-circuit photocurrent density of 18.84 mA/cm², an open-circuit voltage was 783 mV and a fill factor of 0.73, corresponding to an overall conversion efficiency of 10.82%. On the other hand, Bignozzi *et al.*⁷ reported the preparation and photoelectrochemical characterization of a red sensitive osmium complex, containing 4,4,4-tricarboxy-2,2,6,2-terpyridine and cyanide ligands. The

photochemical stability was qualitatively investigated and was better than the best Ru-based sensitizer to date, with a maximum monochromatic incident photon-to-current conversion efficiency (IPCE) of 85%, a Jsc of 9.59 mA cm⁻², a Voc of 0.552 V, and a FF of 0.54, corresponding to an overall conversion efficiency of 2.86% under AM 1.5 sunlight. Additionally, Chan *et al.*⁸ researched a series of sublimely substituted chlorotri-carbonyl bis(phenylimino)acenaphthene rhenium(I) complexes that were used in the fabrication of photovoltaic devices. The best power conversion efficiency of these devices was 1.29% under an illumination of AM1.5 simulated solar light. In these examples, the dyes were composed of novel high-cost metals, such as Ru, Re, Ir and Os. In contrast, the use of low-cost dyes will hasten the commercial availability of DSSCs. Therefore, this study attempted to obtain stable transition metal complexes at a low cost and apply them to the dyes in DSSCs. Over the last few years, considerable efforts have been focused on the design and synthesis of zinc complexes, which have attracted extensive interest in many fields, such as blue or green emitting materials in LEDs (light emitting diodes).⁹⁻¹¹ Particularly, Zhang *et al.*¹⁰ fabricated a ZnO LED by depositing undoped n-type ZnO onto a p-type layer. The turn-on voltage of this LED was about 3.70 V at 100K, which approached the band gap of ZnO, and the electroluminescence spectra exhibited two bands at 423 and 523 nm. On the other hand, Hong *et al.*¹¹ showed that a device containing 20 wt % of a ZnO incorporated hybrid material (2 wt %) emitted nearly pure white light and a total luminous flux of 1.80 lm at an operating voltage of 20 mA. The lifetime measurement data of the fabricated device containing the polymer-ZnO hybrid materials exhibited significant improvements over its pure counterpart because of the "caging effect" of the ZnO shell, which reduced the self-quenching of the polymer molecules in the core.

Despite their great light-emitting ability, research has not focused on the use of Zn-complexes in solar cases. Therefore, in this study, the zinc (II) complex was easily and stably synthesized through the formation of bipyridine chelate ligands on the metal center. A microwave thermal method at 300 W and 60 Hz was used the new synthesis approach. This method differed from the reflux method that was conducted under room temperature and atmosphere pressure. A perfect crystallization and a rapid synthesis time were expected from the microwave method. The crystal structure and the optical properties of the synthesized complex were analyzed using the X-ray photo, FT-IR, UV-vis, and photoluminescence spectra. Finally, this complex was examined for DSSC applications.

Experimental

Synthesis of bis(2,2'-bipyridine)nitrate zinc (II) nitrate. The Zn (II) complex was prepared using the microwave thermal method. As shown in Figure 1, Zn(NO₃)₂·6H₂O (Aldrich Co, 99%) and 2,2'-bipyridine (Wako Co, 99.0%) were added to a round 500 mL flask at a molar ratio of approximately 1:3 with 120 mL of distilled water, and the mixture was stirred for 2 h. The final solution was transferred into a microwave thermal reactor [Jung Hwa MQD; at Yeungnam University, 0 - 700 watt with water solvent, 60 Hz fixed] and treated at 300 watt and 60 Hz for 80 minutes. Microwave synthesis technology is the most effective design for applying microwave energy to a sample. It delivers true continuous power to the sample to ensure reproducible reaction conditions and results, regardless of volume, geometry, or changes in physical properties. The system is compact and easy-to-use, yet powerful enough to drive the most difficult reactions to completion in minutes. Faster reactions (10 - 1,000 times faster) provide increased yields (10 - 30% higher on average) and improved selectivity compared to the conventional thermal systems. Optimizing reactions in fewer steps gives more time to explore creativity. Microwave-enhanced chemistry has the added benefit of using much less solvent than traditional methods, resulting in significantly less waste and expense. A flowing coolant was applied to the top of the beaker in order to prevent the reaction mixture from

rapidly boiling. The solid zinc complex was separated from the solution after the reaction, and a pale pink Zn (II) complex was attained after drying at 353 K.

Characterization. The ground state geometry of bis(2,2'-bipyridine)nitrate zinc (II) nitrate was assigned using a computational method and optimized using the Density functional theory (DFT) calculations with Gaussian 03¹² at 298 K in the gas phase at the B3LYP/LANL2DZ level. The UV-vis spectrum of the ground state was obtained using the TDDFT at B3LYP/LANL2DZ. The X-ray photon spectroscopy (XPS) measurements of C1s, N1s, O1s, and Zn2p in bis(2,2'-bipyridine)nitrate zinc (II) nitrate were recorded using an AXIS-NOVA (Kratos Inc., at the Korea Basic Institute Jeonju center, South Korea) system that was equipped with a non-monochromatic AlK α (1486.6 eV) X-ray source. The powders were pelleted at 1.2×10^4 kPa for 1 min, and the 1.0-mm pellets were stored overnight in a vacuum (1.0×10^{-7} Pa) in order to remove any water molecules from the surface prior to the measurements. The experiments were conducted using a 200-W source power with an angular acceptance of $\pm 5^\circ$. The analyzer axis made a 90° angle with the specimen surface. A Shirley function was used to remove the background in the XPS data analysis, and the signals were fitted using the mixed Lorentzian-Gaussian curves. The UV-vis spectra for bis(2,2'-bipyridine)nitrate zinc (II) nitrate were obtained using a Cary 500 spectrometer with a reflectance sphere over the spectral range from 200 to 800 nm. The FT-IR spectra (FT-IR-8100M, Shimadzu Co., at Yeungnam University, South Korea) for bis(2,2'-bipyridine)nitrate zinc (II) nitrate were recorded on a Mattson 1000 spectrometer using the diffused reflectance method. The powdered sample was mixed with CaF₂ and pressed into pellet form. The scan range for the measurements ranged from 400 to 3600 cm⁻¹, and 50 scans were accumulated to obtain a resolution of 4.0 cm⁻¹. Bis(2,2'-bipyridine)nitrate zinc (II) nitrate underwent fluorescence (FL) spectroscopy (Model FP-777, Jasco, at Yeungnam University, South Korea) in order to examine the number of photo-excited electron hole pairs. The target wavelength was 325 nm, and 0.5 mM Zn-complex was diluted in acetonitrile. The cyclic voltammetry data were obtained using a BAS 100B at room temperature and a scan rate of 100 mV/s with 0.1 M KCl (0.2 mM Zn-complex) as the supporting electrolyte a platinum wire as the working and counter electrodes, and Ag/AgCl as the reference electrode. Ferrocene was used as the internal reference, and the observed potentials of the redox couples were referenced to ferrocene.

Manufacturing dye-sensitized solar cells (DSSC). A slurry was produced by mixing 5.0 g nanometer-sized TiO₂ powder with 10 mL alcohol after sonication for 24 h at 1200 W/cm² in order to prepare the TiO₂ thin film (Degusa P-25, 50 - 70 nm). TiO₂ was coated onto a fluorine-doped, SnO₂ conducting glass plate (Hartford FTO, ~ 30 ohm/cm², 80% transmittance in the visible region) twice using the squeeze printing technique in order to fabricate the thin film with an approximate thickness of 10.0 μ m. The film was heat treated at 723 K for 30 minutes in order to remove the alcoholic solvent. The P-25 TiO₂ thin film electrode was immersed in a 3.0×10^{-4} M bis(2,2'-bipyridine)nitrate zinc (II) nitrate dye solution at room temperature for 24 h, rinsed with anhydrous ethanol and dried. The Pt-coated

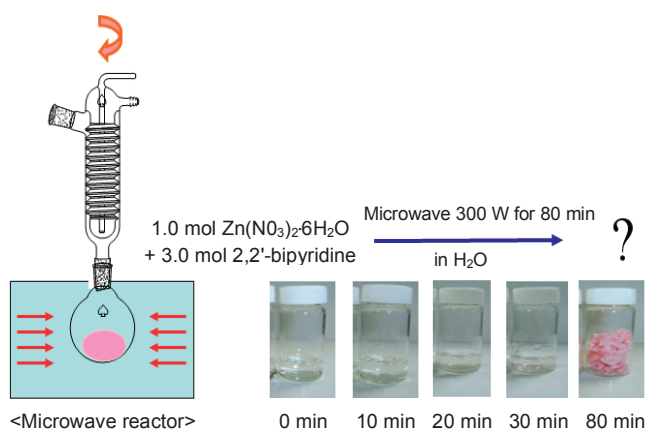


Figure 1. Preparation of bis(2,2'-bipyridine)nitrate zinc (II) nitrate using the microwave method.

FTO electrode was placed over the dye-adsorbed TiO₂ electrode, and the edges of the cell were sealed with a sealing sheet (PECHM-1, Mitsui-Dupont Poly Chemical). The redox electrolyte consisted of 0.50 mol KI, 0.05 mol I₂, and 0.5 mol 4-*tert*-butylpyridine as the solvent. The photocurrent-voltage (I-V) curves were used to calculate the J_{sc}, V_{oc}, FF, and overall conversion efficiency of the DSSC. The I-V curves were measured under white light irradiation from a xenon lamp (max. 150 W, Newport), with an incident light intensity and an active cell area of 100 mW/cm² and 0.40 (0.8 × 0.5) cm², respectively.

Results and Discussion

Characterization of bis(2,2'-bipyridine)nitrate zinc (II) nitrate. Unfortunately, the synthesized compound could not be re-crystallized in this study. Therefore, the structure was attained using a computational method and optimized using the Density Functional Theory (DFT) calculations, with Gaussian 03 at the B3LYP/LANL2DZ level. The UV-vis spectrum of the ground state was obtained using the TDDFT at B3LYP/LANL2DZ. First, the synthesized complex was assumed to have a hexa- or penta-coordination, and then the theoretical UV-vis spectrum was simulated using TDDFT. The spectrum in Figure 2A confirmed that the structure was penta-coordinated, which matched the actual analyzed UV-vis spectrum in Figure 2B. In Figure 2A, the two special absorptions at 429 and 441 nm were assigned to the transfers from the non-bonding ligand (n, HOMO) of NO₃ to the p* ligands (LUMO) of pyridine in penta-coordinated structure. However, the spectrum did not correspond to a hexa-coordination, having the four bipyridine N atoms and chelated NO₃. The signal refinement and the selected spectra for the penta-coordinated complex, bis(2,2'-bipyridine)nitrate zinc (II) nitrate, are given in Appendix 1. According to the simulation in Figure 2A, the Zn atom of the title complex (II) had a distorted octahedral coordination and was linked by the four bipyridine N atoms. One of the O-atom donors lied further from the Zn atom because of the pseudo Jahn-Teller distortions, without chelated NO₃. The Zn atom exhibited a stereochemistry with a pseudo-C₂ symmetry (perfectly hexa-coordination) bisecting the NO₃ ligand and passing between the bipyridine ligands. The computational results confirmed that the geometry of the [Zn(bipy)₂(NO₃)⁺] complex was a penta-coordination with the NO₃-ligated structure. On the other hand, Figure 2B shows the actual reflectance UV-vis absorption spectrum in the wavelength range from 200 - 800 nm for the [Zn(bipy)₂(NO₃)⁺] species. Two types of absorption band, which were assigned to L-L [p[σ, π] (NO₃) → p*[σ*, π*] (NO₃) or p(bpy) → p*(bpy)] transitions were localized at 230, 265, and 320 nm in both the bipyridine compound and the synthesized [Zn(bipy)₂(NO₃)⁺] complex. Otherwise, the main spectrum of the colored complexes, which was assigned to Zn → Zn d-d transfer transitions, was not observed because Zn (II) had a full filled d-orbital. Similar to the theoretical spectrum in Figure 2A, the two absorption bands, which were assigned to the transfers from the non-bonding ligand (n, HOMO) of NO₃ to the p* ligands (LUMO) of pyridine at around 450~550 nm, were also absorbed in the penta-coordinated structure, [Zn(bipy)₂(NO₃)⁺] complex.

Figure 3 presents the typical survey and high-resolution

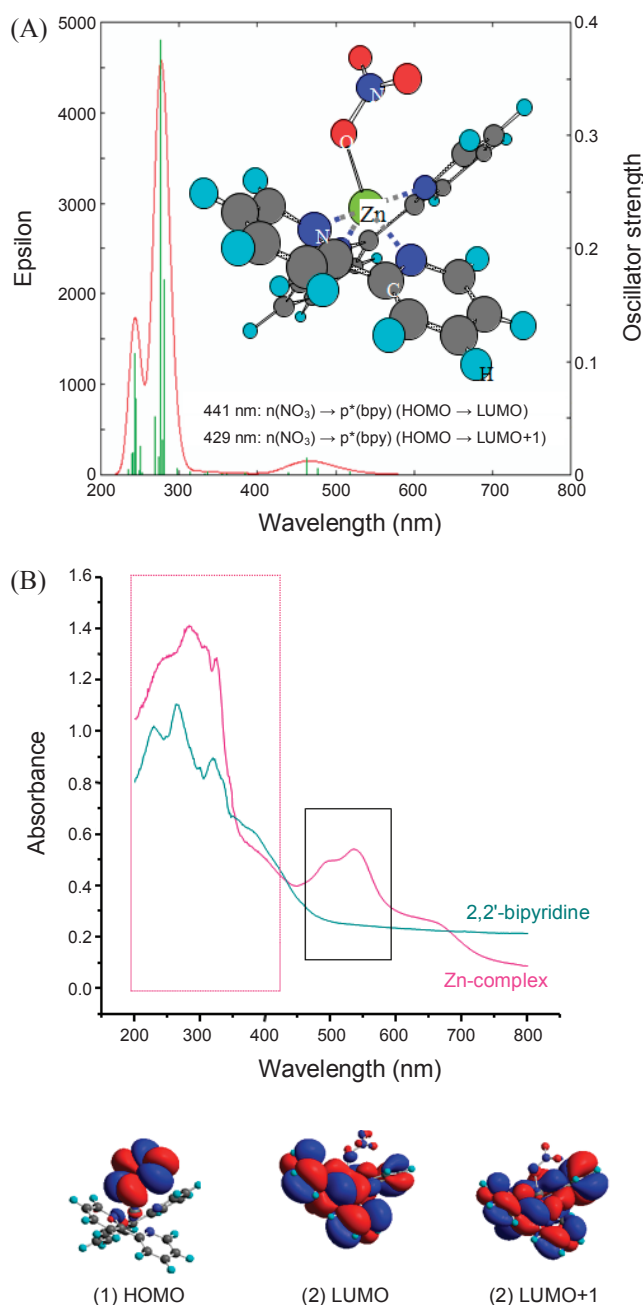


Figure 2. The expected structure (inside) of bis(2,2'-bipyridine)nitrate zinc (II) nitrate, and the (A) theoretical and (B) actual reflectance UV-vis absorption spectra of bis(2,2'-bipyridine)nitrate zinc (II) nitrate.

spectra from the quantitative XPS analysis of the [Zn(bipy)₂(NO₃)⁺] complex. The survey spectra contained the C1s, N1s, O1s and Zn2p peaks. The 2p_{3/2} and 2p_{1/2} spin-orbital photoelectrons of the Zn-centered metal that were located at binding energy of 1022 - 1023 and 1045 - 1047 eV, respectively, were assigned to the Zn²⁺ component.¹³ The O1s spin-orbital photoelectron of NO₃ was located at binding energy of 532 - 533 eV, and, N1s peak at 399.8 eV was assigned to cyanide in bipyridine. Finally, the C1s spin-orbital photoelectron was located at a binding energy of 285 eV and was assigned to the carbide in bipyridine. These results implied that the [Zn(bipy)₂(NO₃)⁺] complex was perfectly synthesized in this study.

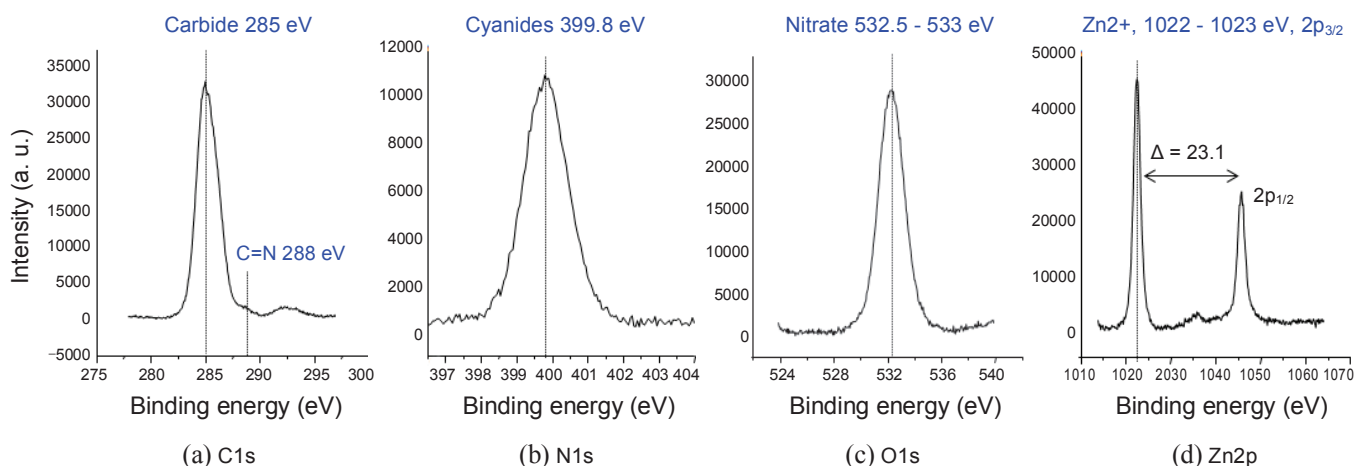


Figure 3. XPS curves of bis(2,2'-bipyridine)nitratocopper (II) nitrate.

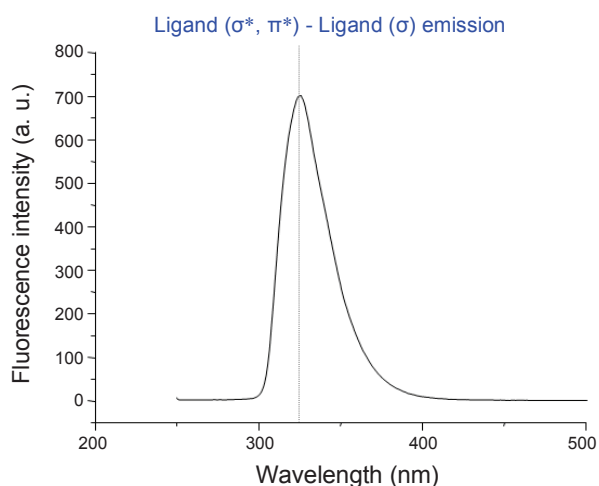


Figure 4. FL spectrum of bis(2,2'-bipyridine)nitratocopper (II) nitrate.

The FL curve suggested that the electrons in the HOMO of bis(2,2'-bipyridine)nitratocopper (II) nitrate were transferred to the LUMO, and then the excited electrons were stabilized through photoemission, in Figure 4. In general, the FL intensity increases with increasing number of emitted electrons, resulting from the recombination between the excited electrons and holes, and, consequently, a decrease in the photo activity. Therefore, there is a strong relationship between the FL intensity and the photo activity. In particular, the FL intensity greatly decreases when a metal captures excited electrons or exhibits conductivity, which is known as the relaxation process. Only one curve in the $[\text{Zn}(\text{bipy})_2(\text{NO}_3)]^+$ complex exhibited a yellow/green emission at 325 nm. The band broadening was attributed to the overlapped emissions from the higher and lower excited states to the ground states, corresponding to the ligand-to-ligand transitions because of the effects on the emission energy from the free ligand to the complex.¹⁴ The HOMO of the complex was a non-bonding orbital, which was localized on the non-coordinating oxygen atom, or otherwise, the HOMO was the carbon atom in the bipyridyl ring. Generally, the energy of the HOMO level of the complex is much lower than the free ligand. The LUMO of the complex was a p-orbital, consisting mostly of

atomic orbitals from one of the bipyridyl rings. The role of the metal ion was to increase the co-planarity and the conformational rigidity in the molecule structure and decrease the $n \rightarrow p$ energy gap.

The oxidation potentials were measured using cyclic voltammetry in distilled water solutions of the 0.2 mM complex with glassy carbon as the working electrode and 0.1 M KCl as the supporting electrolyte, in Figure 5. The $\text{Zn}(0) \rightarrow \text{Zn}(\text{II})$ redox process was reversible. The voltages of the HOMO and LUMO in $[\text{Zn}(\text{bipy})_2(\text{NO}_3)]^+$ were -4.11 and -1.63 eV, respectively. Therefore, the expected band gap was 2.48 eV. Additionally, the progressive shift in $[\text{Zn}(\text{bipy})_2(\text{NO}_3)]^+$ at lower potential values upon the deprotonation of the NO_3 groups was in agreement with the destabilization effects of the negative NO_3 charge on the bipy p-orbital. Therefore, the back-donation from the metal was not as effectively accepted. The electron transfer diagram was proposed in Figure 5 based on these results. When a dye molecule absorbs light, the electrons of the highest occupied molecular orbital (HOMO) orbital are excited to an electronically excited state, corresponding to the lowest unoccupied molecular orbital (LUMO) orbital. The excited dye molecule injected an electron into the conducting band of the TiO_2 film. The oxidized dye was restored by an electron donation from the reducing ions in the electrolyte, which is usually an organic solvent containing a redox system. The donated electron was in turn regenerated through the reduction of the conjugated ions in the electrolytes. The circuit was completed by the electron migration through an external load.¹⁵⁻¹⁷ The location of the dye LUMO energy position must be higher than the LUMO energy position of TiO_2 . Fortunately, in this study, the LUMO of the synthesized $[\text{Zn}(\text{bipy})_2(\text{NO}_3)]^+$ complex was located at a higher position than TiO_2 , resulting in a successful photo-voltaic cycle.

Figure 6 shows the interfacial binding energy between the dye molecule, $[\text{Zn}(\text{bipy})_2(\text{NO}_3)]^+$ complex and the surface of the modified TiO_2 , which was examined using FT-IR spectroscopy. Generally, the efficiency of the charge injection process is strongly dependent on the bonding structure of the dye molecules that are adsorbed on the TiO_2 film. Additionally, the electron transfer in a DSSC is strongly affected by the electrostatic

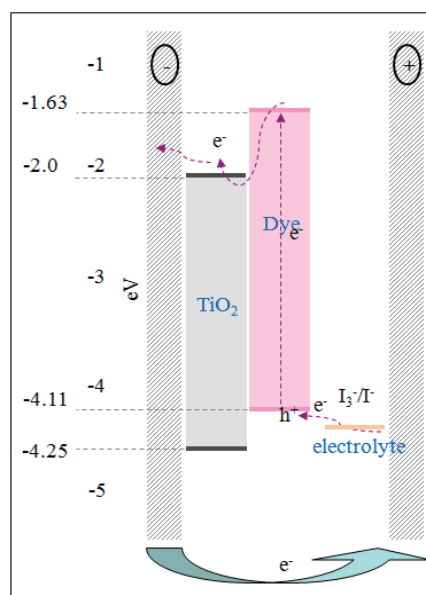
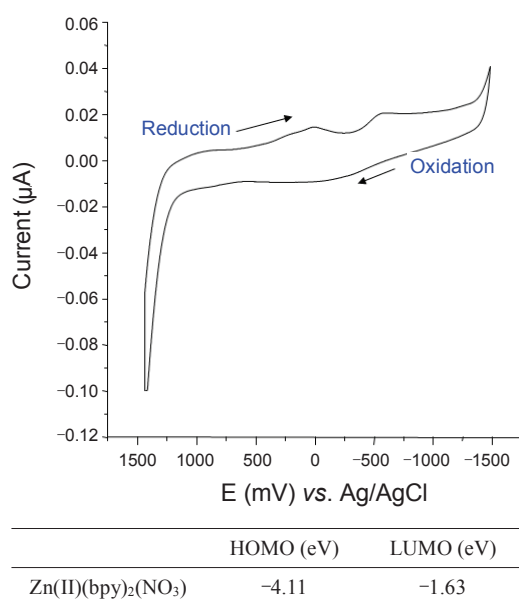


Figure 5. Expected electron transfer diagram of the DSSC with bis(2,2'-bipyridine)nitrate zinc (II) nitrate, and the cyclic voltammetry of bis(2,2'-bipyridine)nitrate zinc (II) nitrate in distilled water solutions of the 0.2 mM complex using a glassy carbon as the working electrode and 0.1 M KCl as the supporting electrolyte.

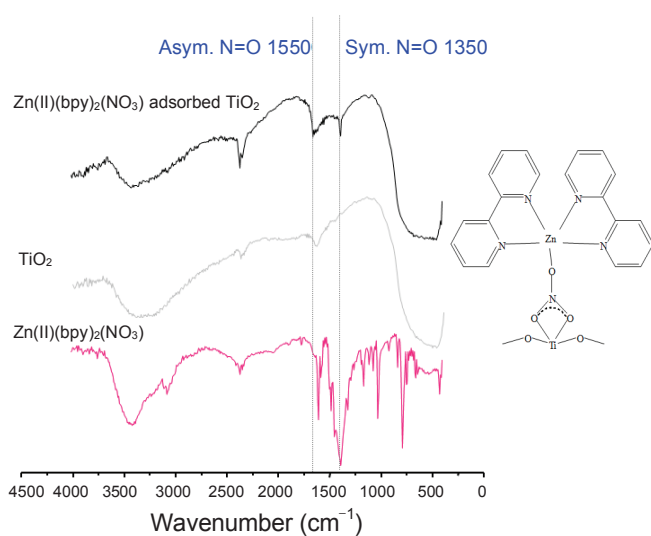


Figure 6. FT-IR spectroscopy of bis(2,2'-bipyridine)nitrate zinc (II) nitrate and an adsorbed model between the surface of TiO₂ and the Zn(II) complex.

and chemical interactions between the TiO₂ surface and the adsorbed dye molecules. Regarding the specific adsorption for FT-IR,¹⁸ the IR spectra exhibited absorptions at 1600 - 1500 and 1390 - 1300 cm⁻¹, which were assigned to the C=C and N=O (-NO₂) stretching modes, respectively. After the dye-adsorption, the N=O band decreased because it was transferred to the N-O stretching mode because of the bidentate coordination of the [Zn(bipy)₂(NO₃)₂]⁺ dye on the surface of the TiO₂ films. The bond between NOO⁻ and the surface of the TiO₂ film was assumed to be strong because of the perfect bidentate linkage. Furthermore, the IR spectrum of the dye-TiO₂ exhibited a broad band at around 500 cm⁻¹, which was assigned to metal-O₂ corresponding to the new Ti-O bond between the O of NOO⁻ and

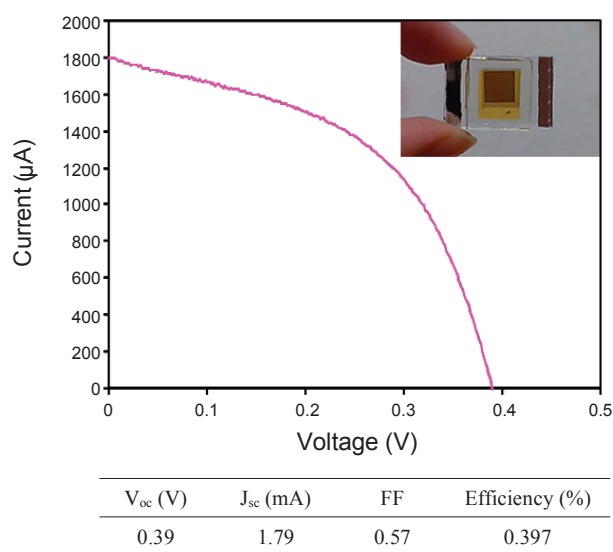


Figure 7. Photocurrent-voltage (I-V) curves of TiO₂ that was adsorbed with bis(2,2'-bipyridine)nitrate zinc (II) nitrate.

the Ti atom. Therefore, the adsorbed model between the surface of TiO₂ and the [Zn(bipy)₂(NO₃)₂]⁺ dye was proposed on the right in Figure 6.

Energy conversion efficiency on DSSC. The photoelectric properties were measured using a voltmeter and ampere meter (Model 2000, Keithley) with a variable load. A 150 W illuminant Xenon lamp was employed as the radiation source at an AM-1.5 radiation angle. The light intensities were measured using a power analyzer and a thermal smart-sensor. The FF and the solar energy conversion efficiency (η) were calculated using Equations (1) and (2), respectively.^{19,20}

$$FF = I_{\max} \times V_{\max} / I_{sc} \times V_{oc} \quad (1)$$

$$\eta (\%) = P_{\text{out}} / P_{\text{in}} \times 100 = I_{\text{max}} \times V_{\text{max}} / P_{\text{in}} \times 100 \\ = I_{\text{sc}} \times V_{\text{oc}} \times FF \quad (2)$$

Figure 7 shows the I-V curves of TiO₂ that was adsorbed with the [Zn(bipy)₂(NO₃)⁺] complex. The FF, V_{oc}, J_{sc}, and overall energy efficiency were determined using the above equations. The TiO₂/[Zn(bipy)₂(NO₃)⁺] DSSC had a V_{oc} of 0.39 V and a J_{sc} of 1.79 mA/cm² at an incident light intensity of 100 mW/cm². The power conversion efficiency of the TiO₂/[Zn(bipy)₂(NO₃)⁺] DSSC was 0.397%. Unexpectedly, the prepared [Zn(bipy)₂(NO₃)⁺] dye was well-adsorbed on the Ti in the TiO₂ film, despite the absence of the COO⁻ group in the bipyridine ring, because of the attachment of two O atoms in NO₃ to the Ti metal. The efficiency was smaller than the efficiency of the Ru-complex (about 3.0 - 5.0%, N719), which is a commonly dye in DSSCs. The efficiency resembled the photovoltaic efficiency on DSSCs that were assembled using Fe-complex dyes.²¹ However, the photovoltaic efficiency could potentially be greatly enhanced in future research through the modification of the bipyridine group with electron donating or withdrawing ligands so that it stably combines with the transition metals.

Conclusions

The microwave method, is a simple and dynamic rapid synthetic procedure that was used to prepare the [Zn(bipy)₂(NO₃)⁺] complex with a partially visible absorption region, *via* a perfectly reversible electrochemical process. The density functional theory calculations revealed that the acquired complex was supposed to a penta-coordination with four nitrogen atoms of bipyridine and the oxygen bond of the NO₃⁻ ion. The FL spectra of the complex displayed a yellow/green luminescence in the acetonitrile solvent. The reflectance UV-vis absorption spectra for the complex exhibited three absorptions that were assigned to two types of L-L transfers at around 200 - 350 nm and 450 - 550 nm. The IR spectra suggested that two O atoms of NO₃⁻ were bonded to the surface of TiO₂. The photoelectric efficiency was approximately 0.397% in the DSSC with the nanometer-sized TiO₂ at a V_{oc} of 0.39 V, a J_{sc} of 1.79 mA/cm², and a fill factor of 0.57.

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References

1. Chou, C. S.; Yang, R. Y.; Yeh, C. K.; Lin, Y. J. *Powder Technol.* **2009**, *194*, 95.

2. An, H.; Xue, B.; Li, D.; Li, H.; Meng, Q.; Guo, L.; Chen, L. *Electrochem. Commun.* **2006**, *8*, 170.
3. Song, H. K.; Park, Y. H.; Han, C. H.; Jee, J. G. *J. Ind. Eng. Chem.* **2009**, *15*, 62.
4. Li, X.; Gui, J.; Yang, H.; Wu, W.; Li, F.; Tian, H.; Huang, C. *Inorg. Chim. Acta* **2008**, *361*, 2835.
5. Christiana, A. M.; Irene, V.; Athanassios, I. P.; Polycarpus, F. J. *Photochem. Photobiol. A* **2007**, *191*, 6.
6. Nazeeruddin, Md. K.; Bessho, T.; Cevey, Le.; Ito, S.; Klein, C.; De Angelis, F.; Fantacci, S.; Comte, P.; Liska, P.; Imai, H.; Graetzel, M. J. *Photochem. Photobiol. A* **2007**, *185*, 331.
7. Argazzi, R.; Larramona, G.; Contado, C.; Bignozzi, C. A. *J. Photochem. Photobiol. A* **2004**, *164*, 15.
8. Mak, C. S. K.; Wong, H. L.; Leung, Q. Y.; Tam, W. Y.; Chan, W. K.; Djurišić, A. B. *J. Organomet. Chem.* **2009**, *694*, 2770.
9. Kalisch, H.; Hamadeh, H.; RuK land, R.; Krysa, J.; Berntgen, A.; Heuken, M. *J. Cryst. Growth* **2000**, *214/215*, 1163.
10. Zhanga, Z. Z.; Wei, Z. P.; Lu, Y. M.; Shen, D. Z.; Yao, B.; Li, B. H.; Zhao, D. X.; Zhang, J. Y.; Fan, X. W.; Tang, Z. K. *J. Cryst. Growth* **2007**, *301-302*, 362.
11. Uthirakumar, P.; Lee, Y. S.; Suh, E. K.; Hong, C. H. *J. Lumin.* **2008**, *128*, 287.
12. GAUSSIAN03, Frisch, M. J.; Trucks, G. W.; Schlegel, H. B.; Scuseria, G. E.; Robb, M. A.; Cheeseman, J. R.; Montgomery, J. A., Jr.; Vreven, T.; Kudin, K. N.; Burant, J. C.; Millam, J. M.; Iyengar, S. S.; Tomasi, J.; Barone, V.; Mennucci, B.; Cossi, M.; Scalmani, G.; Rega, N.; Petersson, G. A.; Nakatsuji, H.; Hada, M.; Ehara, M.; Toyota, K.; Fukuda, R.; Hasegawa, J.; Ishida, M.; Nakajima, T.; Honda, Y.; Kitao, O.; Nakai, H.; Klene, M.; Li, X.; Knox, J. E.; Hratchian, H. P.; Cross, J. B.; Bakken, V.; Adamo, C.; Jaramillo, J.; Gomperts, R.; Stratmann, R. E.; Yazyev, O.; Austin, A. J.; Cammi, R.; Pomelli, C.; Ochterski, J. W.; Ayala, P. Y.; Morokuma, K.; Voth, G. A.; Salvador, P.; Dannenberg, J. J.; Zakrzewski, V. G.; Dapprich, S.; Daniels, A. D.; Strain, M. C.; Farkas, O.; Malick, D. K.; Rabuck, A. D.; Raghavachari, K.; Foresman, J. B.; Ortiz, J. V.; Cui, Q.; Baboul, A. G.; Clifford, S.; Cioslowski, J.; Stefanov, B. B.; Liu, G.; Liashenko, A.; Piskorz, P.; Komaromi, I.; Martin, R. L.; Fox, D. J.; Keith, T.; Al-Laham, M. A.; Peng, C. Y.; Nanayakkara, A.; Challacombe, M.; Gill, P. M. W.; Johnson, B.; Chen, W.; Wong, M. W.; Gonzalez, C.; Pople, J. A. *Gaussian 03; Revision C.02; Gaussian Inc.: Wallingford, CT, 2004.*
13. Mouider, J. F.; Stickle, W. F.; Sobol, P. E.; Bomben, K. D. *Handbook of X-ray Photoelectron Spectroscopy*; Chastain, J., Ed.; Perkin-Elmer Corporation: 1992.
14. Huang, L.; Zhong, A. G.; Chen, D. B.; Liang, H. D. *J. Mol. Struct.* **2009**, *922*, 135.
15. Gagliardi, S.; Giorgi, L.; Giorgi, R.; Lisi, N.; Dikonimos, M. T.; Salernitano, E.; Rufoloni, A. *Superlattices Microst.* **2009**, *46*, 2058.
16. Philipp, W.; Markus, Z.; Christian, S.; Marko, B.; Ursa, O. K.; Dirk, G.; Peter, W.; Andreas, H.; Heiner, J. G. *J. Photochem. Photobiol. A* **2008**, *197*, 25.
17. Lee, W.; Ramasamy, E.; Lee, D.; Song, J. *Sol. Energ. Mat. Sol. C* **2008**, *92*, 814.
18. AkyuEza, S.; AkyuEz, T.; Ozer, N. M. *J. Mol. Struct.* **2001**, *565-566*, 493.
19. Aziz, M. S. *Solid State Electron.* **2008**, *52*, 1145.
20. El-Nahass, M. M.; Zeyada, H. M.; Abd-El-Rahmana, K. F.; Darwish, A. A. *Sol. Energ. Mat. Sol. C* **2007**, *91*, 1120.
21. Jayaweera, P. M.; Palayangoda, S. S.; Tennakone, K. *J. Photochem. Photobiol. A* **2001**, *140*, 173.