

P. WANG¹
S.M. ZAKEERUDDIN^{1,✉}
M. GRÄTZEL^{1,✉}
W. KANTLEHNER²
J. MEZGER²
E.V. STOYANOV²
O. SCHERR²

Novel room temperature ionic liquids of hexaalkyl substituted guanidinium salts for dye-sensitized solar cells

¹ Laboratory for Photonics and Interfaces, Swiss Federal Institute of Technology, CH 1015, Lausanne, Switzerland

² FH Aalen, FB Chemie/Organische Chemie, 73430 Aalen, Germany

Received: 4 September 2003/Accepted: 9 December 2003
Published online: 5 March 2004 • © Springer-Verlag 2004

ABSTRACT A novel family of room temperature ionic liquids, *N, N*-diethyl-*N', N'*-dipropyl-*N''*-hexyl-*N''*-methylguanidinium iodide (SGI) and *N, N, N', N'*-tetramethyl-*N''*, *N''*-dipentyl-guanidinium tricyanomethanide (SGTM) were designed and synthesized. Due to the strong charge delocalization on the tricyanomethanide anion and, thus, weaker ion-pairing, SGTM has a lower viscosity than SGI salt that has iodide as an anion. SGI was successfully used as an iodide resource for dye-sensitized nanocrystalline solar cells. The device with a solvent-free, SGI-based electrolyte achieved a 5.9% power conversion efficiency under an air mass 1.5 incident light of 9.47 mW/cm².

PACS 82.45.Gj; 82.45.Vp; 82.47.Jk

1 Introduction

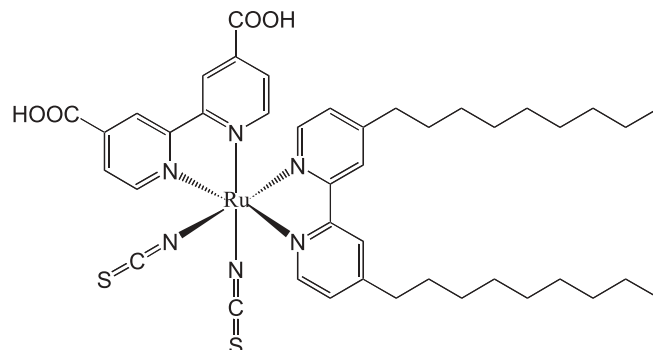
Room temperature ionic liquids have attracted growing interest in recent years owing to their unique chemical and physical properties. It is well known that cations like ammonium, phosphonium, sulphonium, pyridinium, pyrrolidinium and imidazolium can form room temperature ionic liquids with a variety of anions [1–4]. Ionic liquids as promising green solvents have been extensively studied for applications such as catalysis, organic synthesis, batteries and photovoltaics due to their chemical and thermal stability, nonflammability, nonvolatility, high ionic conductivity and possible wide electrochemical window. Prototype guanidinium iodide is a solid salt at room temperature due to the planar structure of the cation and the strong ion pairing. However, asymmetric substitution of hydrogen atoms on nitrogen with different alkyl groups has reduced the melting point drastically to form a room temperature ionic liquid (SGI). *N, N, N', N'*-tetramethyl-*N''*, *N''*-dipentylguanidinium chloride is a solid at room temperature, while its tricyanomethanide salt (SGTM) is a liquid with low viscosity, which can be ascribed to the reduced ion-pairing effect due to the strong charge delocalization on the

tricyanomethanide anion. This is the first time that room temperature ionic liquids with guanidinium as a cation and tricyanomethanide as an anion have been reported. Importantly, SGI can be used as an iodide resource for dye-sensitized nanocrystalline solar cells [5–12]. Earlier, hexamethyl substituted guanidinium tricyanomethanide was reported to be a solid at room temperature, and the synthetic procedure used was time consuming, expensive and complicated [13]. Here, we report a simple synthesis of hexaalkyl substituted guanidinium salts starting from cheap materials. While preparing the manuscript, new room temperature ionic liquids based on the tetra-alkyl-dimethylguanidinium cations were reported [14].

2 Experimental

2.1 Reagents

The amphiphilic Z-907 dye [*cis*-Ru(H₂dcbpy)(dncpy)(NCS)₂], where the ligand H₂dcbpy is 4,4'-dicarboxylic acid-2,2'-bipyridine and dncpy is 4,4'-dinonyl-2,2'-bipyridine] was synthesized as described in our previous paper [15] and its chemical structure is shown in Scheme 1. 400-nm-sized TiO₂ particles were received as a gift from CCIC (Japan). *N*-methylbenzimidazole (NMBI) was purchased from Aldrich and recrystallized from diethyl ether. 3-methoxypropionitrile (MPN, Fluka) was distilled before use. Other chemicals were used as received.



Scheme 1

✉ Fax: +41-21/693 4111,
E-mail: shaik.zakeer@epfl.ch; michael.gratzel@epfl.ch

2.2 Preparation of *N, N*-diethyl-*N', N'*-di-*n*-propylurea

To a solution of 106 g (1.05 mol) of di-*n*-propyl amine and 106 g (1.05 mol) of triethylamine in 500 ml anhydrous acetonitrile, 135.6 g (1.0 mol) of *N, N*-diethylcarbamoyl chloride was added drop wise at -10°C and stirred at this temperature for 16 h. The precipitated triethylamine hydrochloride was filtered off and the solvent was evaporated under reduced pressure. The viscous residue was dissolved in 400 ml ether. The solution was kept at $+2^{\circ}\text{C}$ for 16 h. The separated hydrochloride was filtered off and the ether was removed in a rotary evaporator. The *N, N*-diethyl-*N', N'*-di-*n*-propylurea was obtained by distillation under reduced pressure using a 25 cm vigreux-columnne. Yield: 92%. bp: $106\text{--}110^{\circ}\text{C}/12\text{ Torr}$. n_{D}^{20} : 1.4496. $^1\text{H NMR}$ (CDCl_3 , 250 MHz) δ_{H} : 0.87 (t, $J = 7.4\text{ Hz}$, 6H, $\text{N}(\text{CH}_2)_2\text{--CH}_3$), 1.10 (t, $J = 7.1\text{ Hz}$, 6H, $\text{N--CH}_2\text{--CH}_3$), 1.53 (m, 4H, $\text{NCH}_2\text{--CH}_2\text{--CH}_3$), 3.06–3.20 (m, 8H, N--CH_2). $^{13}\text{C NMR}$ (CDCl_3 , 62.5 MHz) δ_{C} : 11.43, 13.26 (CH_3), 21.21 ($\text{N--CH}_2\text{--CH}_2$), 42.04 ($\text{NCH}_2\text{--CH}_3$), 49.92 ($\text{N--CH}_2\text{--CH}_2$), 165.30 (C=O). IR (direct, cm^{-1}): 2963, 2926, 2888 (CH_3 , CH_2), 1641 (C=O). $\text{C}_{11}\text{H}_{24}\text{N}_2\text{O}$ (200.33) calc.: C, 65.95; H, 12.08; N, 13.98 (%). Found: C, 65.63; H, 12.07; N, 14.13 (%).

2.3 Preparation of *N, N*-diethyl-*N', N'*-di-*n*-propyl-*N''*-hexylguanidine

To a solution of 200 g (1.0 mol) *N, N*-diethyl-*N', N'*-di-*n*-propylurea in 200 ml anhydrous acetonitrile, 110.0 g (1.11 mol) of gaseous phosgene were introduced with stirring at -5°C . The reaction mixture was stirred for 6 h at the same temperature and additionally for 24 h at $+18^{\circ}\text{C}$. In order to remove the excess of phosgene, 10 ml from the solvent was distilled off under reduced pressure (200 Torr). A mixture of 101 g (1.0 mol) hexyl amine and 101 g (1.0 mol) of triethylamine was added drop wise so that the reaction mixture began to reflux. After 1 h at room temperature, the solvent was removed. To the formed mixture of salts was added at 0°C with intensive stirring a solution of 40 g (1.0 mol) of sodium hydroxide in 100 ml water. The triethylamine and the water were removed by evaporation, and the residue was dissolved in hot acetonitrile (200 ml). The solution was filtered and the filtrate was concentrated under reduced pressure to create the crude product. Then, a solution of 12 g (0.3 mol) sodium hydroxide in 100 ml of water was added drop wise at 0°C with intensive stirring to a suspension of 32 g (0.1 mol) of the crude *N, N*-diethyl-*N', N'*-di-*n*-propyl-*N''*-hexylguanidinium chloride in 100 ml of diethyl ether. After 20 min, the ether layer was separated and the water phase was extracted with 50 ml diethyl ether once. The combined organic extracts were dried over potassium carbonate for 1 h, the drying agent was filtered off and the solvent was removed. The residue was distilled under reduced pressure using a 25 cm vigreux-columnne. Yield: 81%. bp: $86^{\circ}\text{C}/10^{-2}\text{ Torr}$. n_{D}^{20} : 1.4626. $^1\text{H NMR}$ (CDCl_3 , 250 MHz) δ_{H} : 0.80–0.99 (m, 9H, $\text{N}[(\text{CH}_2\text{--CH}_2\text{--CH}_3)_2]$, $\text{N--}(\text{CH}_2)_5\text{--CH}_3$), 1.02–1.05 (m, 6H, $\text{N}[(\text{CH}_2\text{--CH}_3)_2]$), 1.29–1.37 (m, 6H, $\text{N}(\text{CH}_2)_2\text{--}(\text{CH}_2)_3\text{--CH}_3$), 1.40–1.54 (m, 6 H, $\text{N}(\text{CH}_2)\text{--}(\text{CH}_2)\text{--}(\text{CH}_2)_3\text{--CH}_3$, $\text{N}[(\text{CH}_2\text{--CH}_2\text{--CH}_3)_2]$, 2.88–2.98 (m, 4H, N--CH_2), 2.91–3.16 (m, 6H, N--CH_2). $^{13}\text{C NMR}$ (CDCl_3 , 62.5 MHz) δ_{C} : 11.49, 11.61, 12.97, 13.62, 14.12 (CH_3), 20.91, 21.73, 22.78,

22.80, 31.91, 31.96, 41.64, 42.54, 49.48, 51.08, 51.13 (CH_2), 158.74 (CN_3). IR (direct, cm^{-1}): 2956, 2926, 2855 (CH_3 , CH_2), 1613 (C=N). $\text{C}_{17}\text{H}_{37}\text{N}_3$ (283.50) calc.: C, 72.02; H, 3.15; N, 14.82 (%). Found: C, 71.03; H, 12.91; N, 15.00 (%).

2.4 Preparation of *N, N*-diethyl-*N', N'*-di-*n*-propyl-*N''*-hexyl-*N''*-methylguanidinium iodide (SGI)

4.3 g (0.03 mol) of methyl iodide were added to a stirred solution of 8.5 g (0.02 mol) *N, N*-diethyl-*N', N'*-di-*n*-propyl-*N''*-hexylguanidine in 5 ml anhydrous acetonitrile. After stirring at room temperature for 12 h, the excess of methyl iodide and the solvent were removed under reduced pressure. Yield: 96%. n_{D}^{20} : 1.5370. $^1\text{H NMR}$ (CDCl_3 , 250 MHz) δ_{H} : 0.86–1.02 (m, 9H, $\text{N}(\text{CH}_2\text{--CH}_2\text{CH}_3)_2$, $\text{N--}(\text{CH}_2)_5\text{--CH}_3$), 1.21–1.34 (m, 12H, $\text{N}(\text{CH}_2\text{--CH}_3)_2$, $\text{N--}(\text{CH}_2)_2\text{--}(\text{CH}_2)_3\text{--CH}_3$), 1.45–1.77 (m, 6H, $\text{N}(\text{CH}_2\text{--CH}_2\text{--CH}_3)_2$, $\text{N--}(\text{CH}_2\text{--CH}_2\text{--}(\text{CH}_2)_3\text{--CH}_3$), 2.98–3.46 (m, 10H, $\text{N}(\text{CH}_2\text{--CH}_2\text{--CH}_3)_2$, $\text{N}(\text{CH}_2\text{--CH}_3)_2$, $\text{N--CH}_2\text{--}(\text{CH}_2)_4\text{--CH}_3$), 3.11 (s, 3H, N--CH_3). $^{13}\text{C NMR}$ (CDCl_3 , 62.5 MHz) δ_{C} : 11.39, 11.53, 12.99, 13.58, 13.93 (CH_3), 21.00, 21.33, 22.51, 26.58, 27.27, 31.30 (CH_2), 39.70 (N--CH_3), 44.25, 51.62, 51.72, 53.20, 53.28 (CH_2), 163.84 (CN_3). IR (direct, cm^{-1}): 2956, 2928, 2865 (CH_3 , CH_2), 1534 (C--N). $\text{C}_{18}\text{H}_{40}\text{N}_3\text{I}$ (425.44) calc.: C, 50.82; H, 9.48; N, 9.88; I, 29.83 (%). Found: C, 50.17; H, 9.57; N, 9.64; I, 29.47 (%).

2.5 Preparation of *N, N, N', N'*-tetramethyl-*N''*, *N''*-dipentylguanidinium tricyanomethanide (SGTM)

Solutions of 6.5 g (0.05 mol) potassium tricyanomethanide [16] in 60 ml acetonitrile and 10.4 g (0.03 mol) *N, N, N', N'*-tetramethyl-*N''*, *N''*-dipentylguanidinium chloride [17, 18] in 10 ml acetonitrile were mixed and stirred at room temperature for 18 h. The precipitated potassium chloride was filtered off and the solvent was removed under reduced pressure. To the residue was added 50 ml dichloromethane and kept at $+2^{\circ}\text{C}$ for 16 h. The mixture was filtered and the filtrate was evaporated in a rotary evaporator. Yield: 8.7 g (84%). n_{D}^{20} : 1.4958. $^1\text{H NMR}$ (CDCl_3 , 250 MHz) δ_{H} : 13.66 (t, $J = 6.8\text{ Hz}$, 6H, $\text{N}[(\text{CH}_2)_4\text{--CH}_3]_2$), 1.22–1.39 (m, 8H, $\text{N}[(\text{CH}_2)_2\text{--}(\text{CH}_2)_2\text{--CH}_3]_2$), 1.42–1.47 (m, 2H, $\text{N}[(\text{CH}_2)\text{--}(\text{CH}_2)\text{--}(\text{CH}_2)_2\text{--CH}_3]_2$), 1.50–1.67 (m, 2H, $\text{N}[(\text{CH}_2)\text{--}(\text{CH}_2)\text{--}(\text{CH}_2)_2\text{--CH}_3]_2$), 2.99 (s, 6H, $\text{N}(\text{CH}_3)_2$), 3.02 (s, 6H, $\text{N}(\text{CH}_3)_2$), 3.06–3.27 (m, 4H, $\text{N}(\text{CH}_2\text{--}(\text{CH}_2)_3\text{--CH}_3)_2$. $^{13}\text{C NMR}$ (CDCl_3 , 63 MHz) δ_{C} : 6.28 [$\text{C}(\text{CN})_3$], 13.91 (CH_3), 22.29, 27.40, 28.82 (CH_2), 40.40, 40.44 [$\text{N}(\text{CH}_3)_2$], 49.81 (N--CH_2), 121.79 ($\text{C}\equiv\text{N}$), 163.28 (CN_3). IR (direct, cm^{-1}): 2951, 2931, 2865 (CH_3 , CH_2), 2151 ($\text{C}\equiv\text{N}$), 1585, 1561 (C--N). $\text{C}_{19}\text{H}_{34}\text{N}_6$ (346.28) calc.: C, 65.86; H, 9.89; N, 24.25(%). Found: C, 65.77; H, 9.97; N, 23.81 (%).

2.6 Viscosity measurements

Viscosities were measured with an ARES (Advanced Rheometric Expansion System, Rheometrics) rheometer in oscillating shear with parallel plates of 25 mm in diameter. The thickness of the specimens was about 0.6 mm. The

oscillating shear measurements were performed in the linear viscoelastic regime as determined from strain sweeps at constant shear rate.

2.7 Electrochemical measurements

A computer controlled Autolab P20 electrochemical workstation (Eco Chimie, Netherlands) was employed for voltammetric measurements. A two-electrode electrochemical cell, consisting of a 5.0 μm radius Pt ultramicroelectrode (Bioanalytical Systems, Inc., USA) as a working electrode and a Pt foil as a counter electrode, was used for the diffusion coefficient measurements of triiodide and iodide. A CDM210 conductivity meter (Radiometer Analytical, France) was employed to measure the conductivities of our electrolyte at various temperatures. The CDC749 conductivity cell (Radiometer Analytical, France) with a nominal cell constant of 1.70 cm^{-1} was calibrated with 0.1 M KCl aqueous solution prior to experiments. A DT Hetotherm cycle heat pump (Heto, Denmark) was used to control the temperature of electrolyte.

2.8 Fabrication of dye-sensitized solar cells

A screen-printed double layer of TiO_2 particle was used as photoanode. A 10 μm thick film of 20 nm sized TiO_2 particles was first printed on the fluorine-doped SnO_2 conducting glass electrode and further coated by 4 μm thick second layer of 400 nm light scattering anatase particles. The detailed fabrication procedures for double-layer TiO_2 nanocrystalline photoanodes and solar cells were described in our previous work [19]. The TiO_2 electrodes were sensitized by immersing them into a dye solution consisting of 0.3 mM Z-907 dye in acetonitrile and *t*-butanol (volume ratio: 1 : 1) at room temperature for 12 h. The electrodes were separated by a 35 μm thick Bynel hot-melt ring (DuPont, USA) and sealed up by heating. The internal space was filled with electrolyte using a vacuum pump to produce device A and B with electrolytes A and B, respectively. Electrolyte A contains 0.5 M *N*-methyl-benzimidazole (NMBI) and 0.5 M I_2 in SGI. Electrolyte B is composed of 0.6 M SGI, 0.5 M NMBI and 0.1 M I_2 in 3-methoxypropionitrile (MPN). In these electrolytes, NMBI is used as base. Additionally, electrolyte B is mainly based on organic solvent, and SGI is used as iodide resource. It should be noted that the added iodine would react with the excess iodide to form triiodide. The cells with electrolytes A and B are denoted as device A and B, respectively.

2.9 Photoelectrochemical measurements

A 450 W xenon light source (Oriel, USA) was used to give an irradiance of 100 mW cm^{-2} (the equivalent of one sun at AM 1.5) at the surface of solar cells. The spectral output of the lamp was matched in the region of 350–750 nm with the aid of a Schott K113 Tempax sunlight filter (Präzisions Glas & Optik GmbH, Germany), so as to reduce the mismatch between the simulated and the true solar spectrum to less than 2%. Various incident light intensities were regulated with neutral wire mesh attenuators. The current-voltage characteristics of the cell under these conditions were obtained by applying external potential bias to the cell and measuring the generated

photocurrent with a Keithley model 2400 digital source meter (Keithley, USA). This process was fully automated using Wavemetrics software (<http://www.wavemetrics.com/>).

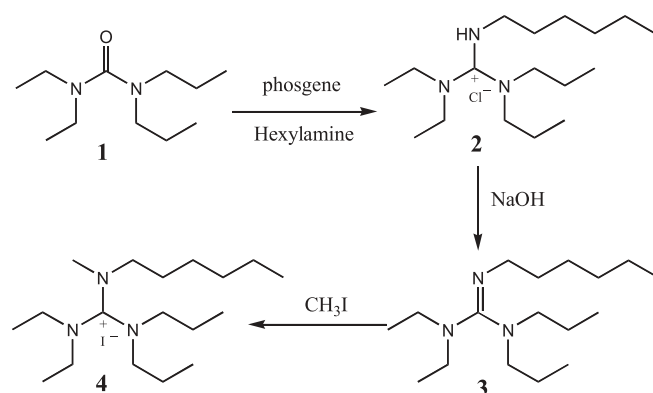
3 Results and discussions

3.1 Synthesis

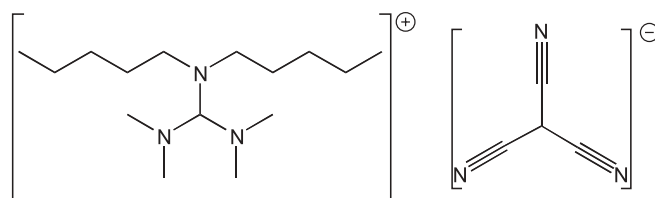
The synthesis of *N,N*-diethyl-*N',N'*-dipropyl-*N''*-hexyl-*N''*-methylguanidinium iodide (**4**, SGI) is described in Scheme 2 and *N,N,N',N'*-tetramethyl-*N''*,*N''*-dipentylguanidinium tricyanomethanide (SGTM, Scheme 3) was prepared by anion exchange reaction of *N,N,N',N'*-tetramethyl-*N''*,*N''*-dipentylguanidinium chloride with potassium tricyanomethanide.

3.2 Viscosity

The electrostatic attraction force between cations and anions, the steric packing of ions, the tendency to form hydrogen bonding and the strength of Van der Waals interaction essentially determine the viscosity of ionic liquids. In the case of the ionic liquid SGI, asymmetric substitution of the protons on the nitrogen of the guanidinium iodide with different alkyl groups results in a room temperature ionic liquid with viscosity of 3330 cPs, which is likely caused by destabilization of the crystalline state. The viscosity of ionic liquid also depends on the charge densities of ions, which determine the strength of electrostatic attraction force between anions and cations. *N,N,N',N'*-tetramethyl-*N''*,*N''*-dipentylguanidinium chloride is a solid due to strong ion-pairing. Replacing the chloride with tricyanomethanide anion, which has a very low negative charge density due to delocalisation over three cyanide groups, forms a less viscous ionic liquid with a viscosity of 88 cPs. Even though the cation



Scheme 2



Scheme 3

is not exactly same in case of SGI and SGTM, the viscosity decreases from 3330 cPs to 88 cPs by changing the anions from iodide to tricyanomethanide. The tricyanomethanide ion has a lower negative charge density compared with iodide in reducing the coulombic interactions with the guanidinium cation. It is interesting that ionic liquids having relatively low viscosity can be prepared from guanidinium cations and tricyanomethanide anions.

3.3 Conductivity

The conductivity (σ) of pure SGI and solvent-free electrolyte A was first examined. It provides information on the mobility of the ions and ion-pairing phenomenon, which are expected to influence the photovoltaic performance and in particular the fill factor of the solar cell. Mechanistically, when an ionic transport process involves an intermolecular ion hopping, the conductivity will be determined by the thermal hopping frequency, which leads to an Arrhenius conductivity-temperature behaviour. As is apparent from the inset of Fig. 1, the Arrhenius equation cannot be used to describe the conductivity-temperature behaviour of the electrolyte. A better fit to the data (Fig. 1) is obtained by the Vogel–Tammann–Fulcher (VTF) equation [20].

$$\sigma(T) = AT^{-1/2} \exp[-B/(T - T_0)] \quad (1)$$

In (1), A and B are constants; T is the absolute temperature, while T_0 is the glass-transition temperature. The VTF conductivity-temperature behaviour indicates there is still a strong ion pairing in pure SGI and electrolyte A. The higher conductivity of electrolyte A than pure SGI at low temperature is due to less ion pairing since the negative charge density of triiodide is lower compared with iodide. However, the conductivity difference between SGI and electrolyte A becomes less at high temperature, indicating a higher freedom of ions in pure SGI as the temperature increases.

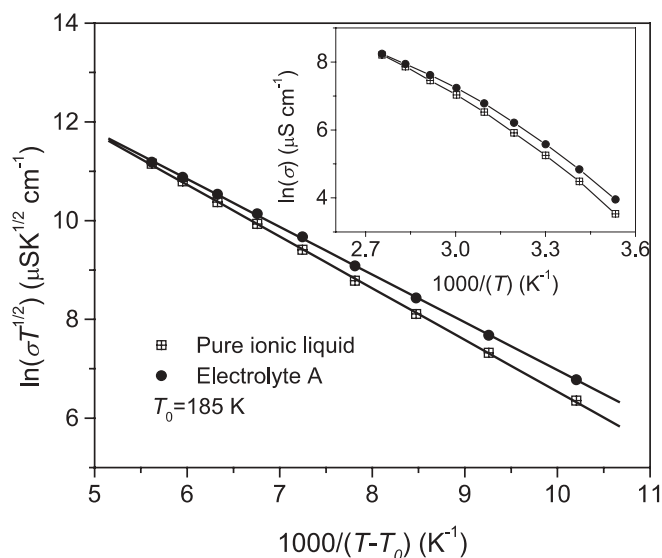


FIGURE 1 Plots of conductivity-temperature data in the VTF coordinates for pure SGI and electrolyte A. The inset is Arrhenius plots of conductivity-temperature data

3.4 Photovoltaic measurements

Figure 2 presents typical photocurrent density-voltage curves under simulated air mass (AM) 1.5 sunlight illumination. At an intensity of 1.47 mW/cm², device A with solvent-free ionic liquid electrolyte has a short-circuit photocurrent density (J_{sc}) of 0.244 mA/cm², open-circuit voltage (V_{oc}) of 670 mV, fill factor (ff) of 0.73, and an overall energy conversion efficiency (η) of 7.4%. The corresponding device parameters (J_{sc} , V_{oc} , ff , and η) under an irradiance of 9.47 mW/cm² are 1.26 mA/cm², 710 mV, 0.628, and 5.9%, respectively. However, if the intensity exceeds 50 mW/cm², the conversion efficiency of the device declines due to mass transport limitations. Thus, the photocurrent response of device A depends in a nonlinear fashion on the light intensity that is caused by the high viscosity of SGI, reducing the triiodide diffusion coefficient. Photovoltaic parameters (J_{sc} , V_{oc} , FF , and η) for device B at an irradiance of 100 mW/cm² are 13.1 mA/cm², 747 mV, 0.63, and 6.2%. Efficiencies of device B under sunlight of 10 and 50 mW/cm² are 7.4 and 7%, respectively. Thus, the performance of device B is similar to that of a solar cell [15] based on a similar electrolyte containing 1,2-dimethyl-3-propylimidazolium iodide instead of SGI.

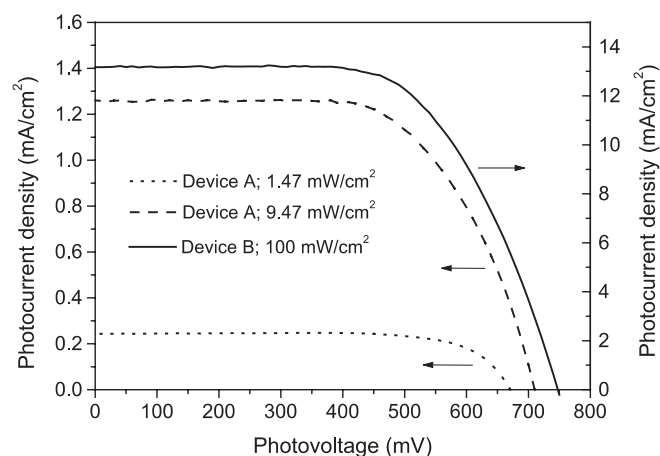


FIGURE 2 Photocurrent density-voltage characteristics of device A and B at different incident light intensities. Cell active area: 0.152 cm²

3.5 Diffusion and its implication to device performance

Diffusion of iodide and triiodide in electrolytes A and B were measured to reveal the limiting step of devices A and B at room temperature. The steady-state voltammograms of electrolytes A and B at the scan rate of 10 mV/s are shown in Fig. 3. The apparent diffusion coefficients (D_{app}) of iodide and triiodide can be calculated from anodic and cathodic steady-state currents (I_{ss}) according to the following equation [21, 22].

$$I_{ss} = 4ncaFD_{app} \quad (2)$$

Where n is the electron transfer number per molecule, F is the Faraday constant and c is the bulk concentration of electroactive species. The calculated diffusion coefficients of triiodide and iodide in electrolyte A are 3.8×10^{-8} and

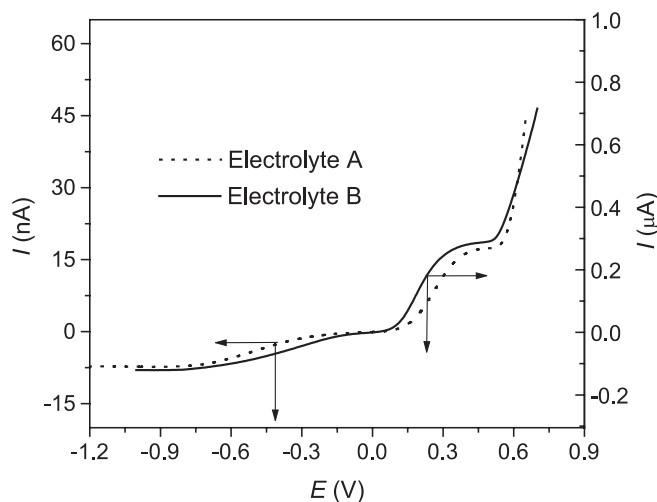


FIGURE 3 Steady-state voltammograms for a Pt ultramicroelectrode in electrolyte A and B. Scan rate: 10 mV/s

$6.9 \times 10^{-8} \text{ cm}^2 \text{ s}^{-1}$, respectively, which are about five times lower than in solvent-free, 1-methyl-3-propylimidazolium iodide based ionic liquid electrolyte [10]. Furthermore, the possible limiting photocurrent density (J_{lim}) in device A with this electrolyte can be derived by (3) to be 5 mA/cm^2 .

$$J_{\text{lim}} = 6\varepsilon_p F D c \frac{1 + \frac{b}{\varepsilon_p l}}{\left[1 + 3\varepsilon_p \frac{b}{l} + 1.5 \left(\frac{b}{l}\right)^2\right]} \times l \quad (3)$$

where D and c are the diffusion coefficient and concentration of triiodide; ε_p and b are the porosity (63%) and thickness of nanocrystalline film (10 μm); l is the electrolyte thickness (10 μm) from photoanode to counter electrode [23]. Thus, we believe that the poor performance of device A at high light intensities is due to the slow diffusion of triiodide in electrolyte A. On the other hand, the calculated diffusion coefficients of triiodide and iodide in electrolyte B are 2.5×10^{-6} and $5.8 \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$, respectively. Thus, the fast diffusion of triiodide in electrolyte B allows for a limiting photocurrent density of 65.8 mA/cm^2 . However, the dependence of J_{sc} of device B on incident photon intensity is still not linear, indicating that its limiting step is due to dye regeneration or photo-induced electron injection rather than the diffusion of triiodide in electrolyte B [12, 15].

4 Conclusions

In summary, novel room temperature ionic liquids, N, N -diethyl- N' , N' -dipropyl- N'' -hexyl- N'' -methylguanidinium iodide (SGI) and N, N, N' , N' -tetramethyl- N'' , N'' -dipentylguanidinium tricyanomethanide (SGTM), have been designed and synthesized. By introducing more delocalized tricyanomethanide anion, the viscosity of SGTM is 38

times larger than SGI. Devices with solvent-free, ionic liquid electrolyte yielded 5.9% power conversion efficiency under an air mass 1.5 incident light of 9.47 mW/cm^2 . The slow diffusion of triiodide in solvent-free, SGI-based electrolyte was proposed as the limiting step for high power application. Furthermore, the organic solvent-based electrolyte containing SGI as iodide resource gave a good photovoltaic performance as that obtained with an imidazolium salt. Its non-linear photovoltaic behaviour may be related to the kinetics of dye regeneration or photo-induced electron injection. Further studies are underway to understand the effect of electrolyte composition on device efficiencies and, thus, to design new guanidinium ionic liquids with lower hydrodynamic viscosity for better device performance.

ACKNOWLEDGEMENTS We are grateful to the Swiss National Science Foundation and the European Office of U.S. Air Force for financial support under Contract No. F61775-00-C0003 and to Marlene Rodlert (LTC, EPFL) for the viscosity measurements.

REFERENCES

- 1 T. Welton: *Chem. Rev.* **99**, 2071 (1999)
- 2 P. Wasserscheid, W. Keim: *Angew. Chem. Int. Ed.* **39**, 3773 (2000)
- 3 R. Sheldon: *Chem. Commun.*, 2399 (2001)
- 4 J. Dupont, R.F. De Souza, P.A.Z. Suarez: *Chem. Rev.* **102**, 3667 (2002)
- 5 B. O'Regan, M. Grätzel: *Nature* **353**, 737 (1991)
- 6 M. Grätzel: *Nature* **414**, 338 (2001)
- 7 N. Papageorgiou, Y. Athanassov, M. Armand, P. Bonhôte, H. Pettersson, A. Azam, M. Grätzel: *J. Electrochem. Soc.* **143**, 3099 (1996)
- 8 H. Matsumoto, T. Matsuda, T. Tsuda, R. Hagiwara, Y. Ito, Y. Miyazaki: *Chem. Phys. Lett.* **26** (2001)
- 9 W. Kubo, T. Kitamura, K. Hanabusa, Y. Wada, S. Yanagida: *Chem. Commun.* **374** (2002)
- 10 P. Wang, S.M. Zakeeruddin, P. Comte, I. Exnar, M. Grätzel: *J. Am. Chem. Soc.* **125**, 1166 (2003)
- 11 S. Murai, S. Mikoshiba, H. Sumino, T. Kato, S. Hayase: *Chem. Commun.* **1534** (2003)
- 12 P. Wang, S.M. Zakeeruddin, J.E. Moser, M. Grätzel: *J. Phys. Chem. B* **107**, 13280 (2003)
- 13 Y.A. Pentin, D.I. Makhon'kov, L.Y. D'yachkova, I.I. Baburina, N.S. Zefirov: *Zh. Obshch. Khim.* **48**, 1850 (1978)
- 14 N.M.M. Mateus, L.C. Branco, N.M.T. Lourenço, C.A.M. Afonso: *Green. Chem.* **5**, 347 (2003).
- 15 P. Wang, S.M. Zakeeruddin, J.E. Moser, M.K. Nazeeruddin, T. Sekiguchi, M. Grätzel: *Nat. Mater.* **2**, 402 (2003)
- 16 J. Falbe: *Methoden der Organischen Chemie* (Georg Thieme Verlag, Stuttgart 1985)
- 17 W. Kantelehner, E. Haug, W.W. Mergen, P. Speh, T. Maier, J.J. Kapasakalidis, H.-J. Bräuner: *Synthesis* (Stuttgart), 904 (1983)
- 18 W. Kantelehner, E. Haug, W.W. Mergen, P. Speh, T. Maier, J.J. Kapasakalidis, H.-J. Bräuner und H. Hagen: *Liebigs Ann. Chem.* **108** (1984)
- 19 P. Wang, S.M. Zakeeruddin, P. Comte, R. Charvet, R. Humphry-Baker, M. Grätzel, M: *J. Phys. Chem. B* **107**, 14336 (2003)
- 20 G.Y. Gu, S. Bouvier, C. Wu, R. Laura, M. Rzeznik, K.M. Abraham: *Electrochim. Acta* **45**, 3127 (2000)
- 21 P. Wang, S.M. Zakeeruddin, I. Exnar, M. Grätzel: *J. Chem. Soc., Chem. Commun.* **2972** (2002)
- 22 R. Kawano, M. Watanabe: *Chem. Commun.* **330** (2003)
- 23 N. Papageorgiou, M. Grätzel, P.P. Infelta: *Sol. Energy Mater. Sol. Cells* **44**, 405 (1996)