Pt-Free Counter Electrodes with Carbon Black and 3D Network Epoxy Polymer Composites

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Carbon black (CB) and a 3D network epoxy polymer composite, representing dual functions for conductive corrosion protective layer (CCPL) and catalytic layer (CL) by the control of CB weight ratio against polymer is developed. Our strategy provides a proper approach which applies high catalytic ability and chemical stability of CB in corrosive triiodide/iodide (I$_3^−$/I$^−$) redox electrolyte system. The CB and a 3D network epoxy polymer composite coated on the stainless steel (SS) electrode to alternate counter electrodes in dye sensitized solar cells (DSSCs). A two-step spray pyrolysis process is used to apply a solution containing epoxy monomers and a polyfunctional amine hardener with 6 wt% CB to a SS substrate, which forms a CCPL. Subsequently, an 86 wt% CB is applied to form a CL. The excellent catalytic properties and corrosion protective properties of the CB and 3D network epoxy polymer composites produce efficient counter electrodes that can replace fluorine-doped tin oxide (FTO) with CCPL/SS and Pt/FTO with CL/CCPL/SS in DSSCs. This approach provides a promising approach to the development of efficient, stable, and cheap solar cells, paving the way for large-scale commercialization.

For more than two decades, dye-sensitized solar cells (DSSCs) have been intensively investigated in industry and academia as a viable alternative to conventional silicon-based photovoltaic cells\textsuperscript{1–10}. A typical DSSC comprises a dye-sensitized titanium dioxide (TiO$_2$) nanocrystalline photoanode, an electrolyte that provided a redox couple (I$_3^−$/I$^−$), and a counter electrode (CE)\textsuperscript{11}. Significant research efforts have been applied toward optimizing each component by modifying the photoanodes, developing alternative dyes, improving the redox couple, and introducing structural changes, to yield highly efficient DSSCs\textsuperscript{12–19}. Pt-coated fluorine-doped tin oxide (FTO) is typically used as the CE because of its excellent catalytic activity toward reducing I$_3^−$. Unfortunately, Pt-coated CE is typically produced through high-temperature hydrolysis processes that are incompatible with flexible conducting polymer electrodes\textsuperscript{20–24}. Furthermore, Pt corrodes in electrolytes containing iodide to generate PtI$_4$\textsuperscript{24–26}. The Pt metal catalyst coating has been replaced with other materials, such as titanium nitrides\textsuperscript{10,27}, cobalt sulfide\textsuperscript{28}, and carbon derivatives\textsuperscript{29–33}. Carbon derivatives are particularly attractive, as they are abundant, low-cost, and provide high catalytic activities while remaining chemically stable in the presence of iodine redox couples\textsuperscript{32–36}. Carbon black (CB) has been used on a large scale, for example in printing toners, reinforcing additives in automobile tires, and as conductive fillers in plastics, elastomers, and films\textsuperscript{37}. CB is a conductive material with good catalytic activity for the reduction of triiodide\textsuperscript{38}. Carbonaceous materials present active catalytic sites at their edges\textsuperscript{39}. Therefore, CB, which have many edges, may be more active than highly structured carbon materials, such as graphites, graphenes, and carbon nanotubes\textsuperscript{40–41}. CB powders cannot retain their shapes on a substrate; thus, they must be mixed with other binder materials, such as polymers or a TiO$_2$ slurry containing organic surfactants and binders\textsuperscript{39,35,42}. For example, Grätzel et al.\textsuperscript{40} and Jun et al.\textsuperscript{43}, independently employed a CB/TiO$_2$ slurry system in DSSC electrodes. Recently, Ho et al. reported the use of a CB/polymer composite CE comprising poly(3,4-ethylenedioxythiophene)\textsuperscript{44}. We, herein, extend the approaches introduced in these works by examining three-dimensional (3D) network polymers formed by polymerized monomers that had been cross-linked (XL) \textit{in-situ} with CB to robustly fix the CB powders onto a substrate and act as a binder for the CB powders. The 3D network polymers are composed of an epoxy monomer and a polyfunctional amine hardener. The chemically XL epoxy polymer provides strong mechanical properties, chemical resistance, thermal resistance, and adhesive properties\textsuperscript{45,46}.

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Here, we describe the development of novel stainless steel (SS) CEs coated with a composite of CB and 3D networked polymers to replace conventional Pt CEs on FTO glass. Previous studies have reported the development of FTO glass alternates for the fabrication of flexible and bendable electrodes. SSs are highly conductive and have a work function (~4.4 eV) that is appropriate for use as a CE in DSSCs. SSs are cost-effective materials compared with FTO glasses. Tan et al. and Kang et al. independently reported the development of an SS mesh and an SS substrate-based CE, respectively. However, the previous reports did not comment on the corrosion problems inherent to these materials and not show the long-term stability of the resulting DSSCs. Corrosion is a key technical feature that must be addressed in practical applications of SS CEs in DSSCs. In this work, we report a CB-coated SS CE with highly thermal stability, thus without corrosion problems. The photovoltaic performance for the device using this CB-coated SS CE remained nearly constant after 30 days of storage at 65 °C. Meanwhile, a reference device consisting of Pt/FTO decreased up to almost 50% of the photovoltaic performance. We successfully coated both a conductive corrosion protective layer (CCPL) and a catalytic layer (CL) with a composite of CB and 3D network polymers onto an SS substrate using spray pyrolysis, as illustrated in Fig. 1a. The weights ratio of the CB incorporated into the polymer matrices of the CCPL and CL were 6 and 86 wt%, respectively. The CCPL formed a compact dense structure that prevented corrosion due to electrolyte penetration, whereas the CL formed a nanoporous structure that provided a large surface area for increased catalytic activity.

Results
Percolation concentration of CB in CCPL. We optimized a coating composition containing CB and a polymer for use as a CCPL by preparing 4 samples (3, 6, 12, and 20 wt%), and we investigated their morphology, conductivity, corrosion protection properties, and electrical and catalytic properties. Figure 1b,c shows field emission scanning electron microscopy images (FE-SEM) of the CCPL layer cross-section and top surface, respectively. Smooth films with 2.4–2.5 μm thick were obtained, and the CB are well-dispersed within the 3D XL polymer matrix. The composites prepared with more than 6 wt% CB displayed well-established vertical interconnectivity among the CB due to the in-situ 3D XL polymer matrix, which appeared to improve as the CB content increase. The CB film did not detach from the composite during the pressure sensitive tape (PST) test. These results indicated that the XL polymers acted as an excellent binder for the CB. An efficient charge transport was obtained in the compact CB composites, and the in-situ generated 3D XL polymers guaranteed interconnections among the CB. Samples in which the CB were separated as a result of the large polymer content yield poor charge transport.
through the conducting CB network. Good mechanical properties, including adhesion, chemical resistance, and corrosion protection, were obtained at optimal composite compositions (e.g., at the percolation concentration) without incurring a loss in the conductivity. The resistivities of the four composites using a 4-point probe method (see the inset in Fig. 2) were measured and the conductivities were calculated (Table 1).

The conductivity of the composite prepared 3 wt% CB was 0.081 S cm$^{-1}$, almost one order of magnitude lower than the conductivity of a film prepared with 6 wt% CB (0.85 S cm$^{-1}$). Further increases in the CB content only increased the conductivity to 0.96 S cm$^{-1}$ at 20 wt% CB. Figure 2 shows the CB concentration-dependent conductivity. A critical point indicative of the percolation concentration was reached at 6 wt% CB. Therefore, we employed 6 wt% CB in the CCPL tested in this work.

Chemical resistance and corrosion protective properties of the CCPL. We next investigated the chemical resistance and corrosion protective properties of the CCPL (6 wt%) in the presence of an electrolyte solution and compared these properties to those obtained using a bare SS substrate dipped in an electrolyte containing 0.03 M iodine and additives (t-butylyl pyridine (tBP) and lithium bis(trifluoromethanesulfonyl)imide (LiTFSI)) at 65 °C. The bare SS substrate was completely corroded within a day, as indicated by the observed color change (Fig. 3a). On the other hand, no corrosion was observed at the CCPL surface, even after 30 days (Fig. 3b). We attempted to compare the bare SS surfaces before and after the soaking test by removing the CCPL layer, but the adhesion between the SS and CCPL layer was too robust to permit removal of the CCPL. These results demonstrated that the CCPL do not react with the electrolyte, and the CCPL effectively prevented the liquid electrolyte from reaching the SS. A 6 wt% CB solution provided the percolation concentration in our system which provided good adhesive properties, good chemical resistance and good corrosion protection ability without a loss in conductivity. In addition, the CCPL/SS electrode exhibited high flexibility due to the high ductility of a composite of CB and 3D network polymers on an SS substrate. Figure 3c shows no cracks under bending deformation of more than 100 times.

Redox activity of CCPL. Devices having a compact CCPL layer were firstly fabricated to evaluate the catalytic properties of the CCPL because the electrode (CCPL/SS) prepared with 6 wt% CB exhibited a reasonably high conductivity. In addition, we prepared a control CE based on a Pt-coated CCPL layer (Pt/CCPL/SS) for comparison with the CCPL/SS electrode. Figure 4a shows the J–V characteristics of the devices prepared with the CCPL/SS and Pt/CCPL/SS CEs. The devices employing the CCPL CE exhibited a very poor photovoltaic performance with a short-circuit current density ($J_{SC}$) of 5.5 mA cm$^{-2}$, an open-circuit voltage ($V_{OC}$) of 0.53 V, a fill factor (FF) of 14.9%, and a power conversion efficiency (PCE) of 0.43%. These values were lower than those obtained from the device employing the Pt/CCPL/SS CE ($J_{SC}$ = 17.7 mA cm$^{-2}$, $V_{OC}$ = 0.70 V, FF = 59.4%, and PCE = 7.3%). These results indicated that the compact CCPL CE could not function as both a corrosion protective layer and as a catalytic layer due to the insufficient contact area between the CB nanoparticles and I$_3^{-}$. A good photovoltaic

Table 1. Thickness (μm), resistivity (Ohm), sheet resistance (Ohm/□), and conductivity values obtained from the CCPL/SS substrates prepared with 3, 6, 12 or 20 wt% CB. $^a$Values were obtained from the average heights measured from the cross-sectional FE-SEM images. $^b$Determined using the 4-point probe method. $^c$ Sheet resistance = correction factor $\times$ resistivity obtained from the 4-point probe method. The value of the correction factor was 4.532, the conducting film thickness was less than the distance of the spacing between the probe tips (1000 μm), and the edges of the film (2.5 cm) were separated from the measurement point by more than 4 times the distance between the probe tips.

![Figure 2. The resistivity and conductivity as a function of the wt% CB.](image-url)
performance was obtained from the device prepared with the Pt/CCPL/SS CE, demonstrating that FTO can be replaced with a CCPL/SS substrate. These results suggested that the electron transport from the Pt layer to the SS through the interconnected CB in the CCPL was highly efficient. Figure 4b shows the electrochemical impedance spectroscopy (EIS) results of CCPL/SS and Pt/CCPL/SS CEs in symmetric cells consisting of two identical CEs at 0 V under dark conditions.
The impedance spectrum of the Pt/CCPL/SS CE was dominated by the $I_3^-/I^-$ redox reaction at Pt and the charges transfer to the SS through the CB in the CCPL. Assuming that the Pt/CB and CB/SS interfaces were characterized as non-Ohmic and Ohmic contacts, respectively, two different interfaces were expected to be present: the electrolyte/Pt and Pt/CB interfaces. Indeed, two semicircles were observed in the spectra. The small semicircle at high frequencies and the large semicircle at low frequencies (in the dark, under a forward bias) were attributed to electron transfer (series resistance, $R_S = \text{ca. 11 } \Omega \text{cm}^2$) at the Pt/CB interface and the redox reaction (charge transfer resistance, $R_{CT} = \text{ca. 40 } \Omega \text{cm}^2$) at the electrolyte/Pt interface, respectively. These results indicated that the catalytic activity of Pt is good, although electrons were more easily transported from Pt to the CB than from Pt to the electrolyte during catalysis. A symmetric cell consisting of two CCPL/SS CEs exhibited only one semicircle characterized by $10,500 \Omega \text{cm}^2$ at the electrolyte/CB interface due to the Ohmic contact at the CB/SS interface; however, the $R_{CT}$ value is large, indicating poor catalytic properties at the CCPL CE. This result agree well with the poor photovoltaic performances of the devices prepared with only the CCPL CE, suggesting that a catalytic layer (CL) with a nanoporous structure was required to increase the surface area at the electrolyte/CB interface.

Adhesive property of the CL. The CB content was increased (and the polymer content decreased) to increase the surface area of the CB. This could decrease the adhesion properties of the CB on the 3D XL polymer matrix. The maximum CB content at which the CB did not detach from the surface of the CL was identified by conducting a pull-off adhesion strength test using a PST to evaluate the adhesive properties of the CLs having CB contents up to 95 wt%. Figure S2 shows representative photographs of the pull-off adhesion strength test results obtained from the two CLs containing 86 or 90 wt% CB. The detachment of CB on the surfaces of the PST was not observed until an 86 wt% CB is used in the CLs. A greater number of CB were found to detach at concentrations beyond this threshold. The detached CB were clearly visible on the PST at and above a 90 wt% CB in the CL, as indicated by the yellow dotted circle in the photograph of the PST surface. These results were used to determine a CB concentration of 86 wt% in the optimized CL on the CCPL.

Electrochemical catalytic ability of the CL. In order to investigate the electrochemical catalytic abilities of the CL, cyclic voltammetry (CV) measurement was performed using a three-electrode electrochemical system with a scan rate of 30 mV/s. The working electrodes were CL/CCPL/SS electrodes with 40, 55 or 86 wt% CB in the CLs. Pt/FTO and CCPL/SS electrodes also tested for comparison. A Pt wire and Ag/AgCl were used as a counter electrode and a reference electrode, respectively, in the acetonitrile solution. As shown in Fig. 5a, each CV curve presents one pair of redox peaks corresponded to the reaction of $I_3^- + 2e^- \rightarrow 3I^-$. An anodic peak is observed at higher potentials due to the reduction of the electrolyte at Pt, while a cathodic peak is observed at lower potentials due to the oxidation of the electrolyte at Pt. The anodic and cathodic peak currents increased with increasing CB content in the CL, indicating an increase in the catalytic activity of Pt. The peak currents were highest for the CL containing 86 wt% CB, suggesting that a CL with a nanoporous structure is required to increase the surface area at the electrolyte/CB interface.

Figure 5. (a) Cyclic voltammograms obtained from the oxidation and reduction of the $I_3^-/I^-$ redox couple using the CCPL/SS, CL(40, 55 or 86 wt%)/CCPL/SS or Pt/FTO electrodes, at a scan rate of 30 mV/s. (b) Anodic and cathodic peak currents for 20 consecutive CVs of the Pt/FTO electrode and the CL(86wt%)/CCPL/SS electrode. (c) Electrochemical impedance spectra of the CL/CCPL/SS ECs prepared with 40%, 55%, or 86% CB in symmetric cells consisting of two identical CEs at 0 V under dark conditions. (d) Representative J–V curves measured in devices prepared using CL/CCPL/SS ECs with 40%, 55%, or 86% CB, and a Pt/FTO CE as a control.
peak current density ($J_{pp}$) and a cathodic peak current density ($J_{pa}$) corresponded to the oxidation of the iodide ions and the reduction of the tri-iodide ions, respectively. The oxidation/reduction peaks for the Pt/FTO electrode were shown at 0.352 V/−0.210 V with peak to peak separation potential ($E_{pp}$) of 0.562 V and $J_{pp}/J_{pa}$ values were 1.211 mA cm$^{-2}$/−0.700 mA cm$^{-2}$. The narrower $E_{pp}$ and higher $J_{pa}$ indicated better electrochemical catalytic activity. In the CCPL/SS electrode, very small amounts of currents flew through the surface of CCPL which had a compact dense structure with a small surface area. The small amounts of currents account for a high $R_{ct}$ value obtained from EIS measurement. CV results of electrodes with increased CB weight ratio (40, 55 or 86 wt%) in the CL presented decreased $E_{pp}$ and enlarged $J_{pa}$ values, indicating effective electrochemical catalytic abilities. The increase of weight ratio of CB in the CL resulted in a higher surface area, where redox reaction occurred, inducing the smaller $R_{ct}$. The $E_{pp}$, $J_{pa}$ and $J_{pc}$ values of the CL(86wt%)/CCPL/SS electrode were 0.576 V, 1.206 mA cm$^{-2}$ and −0.701 mA cm$^{-2}$, respectively. This alternative electrode exhibited the efficient charge transfer ability, which was comparable with the Pt/FTO electrode by the decline of $R_{ct}$. Figure 5b shows the $J_{pa}$ and $J_{pc}$ of the Pt/FTO electrode and the CL(86wt%)/CCPL/SS electrode during the 20 consecutive cycle scans. The $J_{pc}$ values of the CL/CCPL/SS electrode were almost maintained steadily, which revealed the excellent electrochemical stability in the $I_{1}^-/I^-$ based electrolyte system.

### Photovoltaic performances and impedance analysis of DSSCs

As demonstrated above, the electrochemical impedance spectroscopy (EIS) measurements obtained from the symmetric cells established the criteria for the CL surface activity. Figure 5c shows the EIS spectra obtained from symmetric cells consisting of CL/CCPL/SS CEs with 40, 55 or 86 wt% CB in the CLs. One semicircle corresponded to Ohmic contact between the CB in the CL and the CCPL. The resistance of the CL (40 wt%) due to the catalytic activity at the electrolyte/CB interface was $R_{ct} = 119 \Omega$ cm$^2$, much smaller than the resistance obtained from the CCPL (10,500 Ω cm$^2$ for 6 wt% CB). This value decreased as the CB concentration increase. $R_{ct}$ for the CB (86 wt%) is 41 Ω cm$^2$, almost equal to the value obtained from Pt/CCPL (40 Ω cm$^2$, Table 1).

Photocells employing CL/CCPL/SS CEs having 40, 55, or 86 wt% CB in the CLs were fabricated, and the $J$-V characteristics under simulated air mass 1.5 global (AM 1.5G) solar irradiation were measured. The photovoltaic parameters, measured resistance values and catalytic activities are summarized in Table 2, and the representative $J$-$V$ characteristics are presented in Fig. 5d. The photovoltaic performances of the devices followed the trends in the $R_{ct}$ values, indicating that the catalytic activity at the electrolyte/CB interface could be enhanced by increasing the CB content in the CL of the CCPL/SS CE. As the CB content in the CL increased, the FF values increased from 40.3 to 60.6% due to the enhanced catalytic activity, which minimized the recombination reaction between photoinduced electrons and $I_1^-$. The improved catalytic activity affected the $V_{oc}$ values according to the equation: $V_{oc} \sim (nkt/q) \times \ln(I_{oc}/I_{j})$, where, $n$ is the device ideality factor, $k$ is the Boltzmann constant, $T$ is the temperature in Kelvin, $q$ is the fundamental charge, and $I_{j}$ is the saturation current density$^{33,34}$.

The best performance was obtained from the device prepared with a CL(86wt%)/CCPL/SS CE, yielding $J_{sc} = 14.9$ mA cm$^{-2}$, $V_{oc} = 0.68$ V, FF = 60.6%, and PCE = 6.1%. With the exception of $J_{sc}$, these values were equal to those obtained from a device prepared with a Pt/CCPL/SS CE ($J_{sc} = 17.7$ mA cm$^{-2}$, $V_{oc} = 0.70$ V, FF = 59.4%, and PCE = 7.3%) as summarized in Table 1. In general, the $J_{sc}$ values could be estimated using the equation: $J_{sc} = q \times \eta_{lh} \times \eta_{inj} \times \eta_{cc} \times I_{oc}$ where $q$ is the fundamental charge of an electron, $I_{oc}$ is the intensity of the incident light, $\eta_{lh}$ is the light harvesting efficiency, $\eta_{inj}$ is the electron injection efficiency from the excited dye molecules to the TiO$_2$ conduction band, and $\eta_{cc}$ is the charge collection efficiency of the injected electrons at the transparent conductive oxide (TCO) layer. Assuming that $I_{oc}$, $\eta_{lh}$, and $\eta_{inj}$ were equal in the two devices, different $J_{sc}$ values could account for the variations in the quantity of dye molecules adsorbed onto the electrodes, as reflected in $\eta_{cc}$, although we employed identical experimental conditions (Figure S3).

| Electrodes             | $R_{ct}$ ($\Omega$ cm$^2$) | $J_{sc}$ (mA cm$^{-2}$) | $V_{oc}$ (V) | FF (%) | PCE (%) |
|------------------------|-----------------------------|--------------------------|--------------|--------|---------|
| CCPL(6 wt%)/SS         | 10,500                      | 5.5                      | 0.53         | 14.9   | 0.43    |
| Pt/CCPL(6 wt%)/SS      | 40                          | 17.7                     | 0.70         | 59.4   | 7.3     |
| CL(40 wt%)/CCPL/SS     | 119                         | 12.7                     | 0.62         | 40.3   | 3.2     |
| CL(55 wt%)/CCPL/SS     | 68                          | 14.8                     | 0.64         | 42.7   | 4.0     |
| CL(86 wt%)/CCPL/SS     | 41                          | 14.9                     | 0.68         | 60.6   | 6.1     |
| Pt/FTO                 | --                          | 15.4                     | 0.74         | 62.9   | 7.1     |

Table 2. Summary of the catalytic resistance values obtained from various electrodes and their photocurrent–voltage characteristics$^{a}$ under AM 1.5 illumination. $^{a}$10 devices prepared using each CE.

$^{b}$The catalytic resistances ($R_{ct}$) were calculated by fitting to an equivalent circuit (the models are illustrated in Figure S1).
The DSSC prepared using the CL/CCPL/SS CE also displayed a high photovoltaic performance (PCE) in Pt/FTO can be replaced with CL/CCPL/SS. Further performance enhancements may be realized by carefully no corrosion problems. Furthermore, these results demonstrated that FTO can be replaced with CCPL/SS and (PCE may have accounted for the increased $E_{\text{redox}}$ due to changes in the redox species concentration at the surface of the dye-sensitized TiO$_2$ photoanode.

We previously observed that $I_3^-$ molecules became entrapped in the highly porous network polymer matrix at a TiO$_2$ photoanode, thereby increasing $V_{OC}$. Here, the opposite conditions were present because we employed a porous 3D network polymer matrix at the CE. These conditions were consistent with the measured decrease in $V_{OC}$ as the polymer content in the CLs increased, which increased the number of sites available to trap $I_3^-$. Nevertheless, the PCE of the device prepared with the CL(86 wt%)/CCPL/SS electrode reached a PCE that is 86% of the PCE obtained from a device prepared with Pt/FTO. Our results suggest that FTO could be replaced with CCPL/SS, but also that Pt/FTO could be replaced with CL/CCPL/SS.

In conclusion we successfully fabricated a novel flexible and cost-effective stainless steel (SS) counter electrode from a composite composed of CB and 3D network polymers to function as a CCPL and CL. The composite film, which provided a catalytic layer and acted as a conductive corrosion protective film, was prepared using a mixture of CB, cross-linkable epoxy monomers, and polyamine hardener and was applied using the spray pyrolysis method. The SS substrate acted as an efficient charge collecting electrode. The CCPL (6 wt% of CB) was chemically stable against electrolytes and successfully prevent the penetration of electrolytes, demonstrating excellent corrosion protection properties. The Pt/CCPL/SS CE was successfully substituted for a conventional Pt/FTO CE and displayed a photovoltaic performance (PCE = 7.3%) that was equivalent to that obtained from Pt/FTO (PCE = 7.1%). The CL (86% of CB) provided an effective substitute for the conventional catalytic material (Pt).

The DSSC prepared using the CL/CCPL/SS CE also displayed a high photovoltaic performance (PCE = 6.1%). We demonstrated that the CB-coated SS CE (CL/CCPL/SS CE) exhibited highly thermal stability, thus there were no corrosion problems. Furthermore, these results demonstrated that FTO can be replaced with CCPL/SS and Pt/FTO can be replaced with CL/CCPL/SS. Further performance enhancements may be realized by carefully optimizing the device architecture.

Methods

Chemicals and materials. Stainless steel (SS) was provided from POSCO. Carbon black (CB) powders purchased from Beilum Carbon Chemical Limited. Polyethyleneimine (branched, average Mw ~25,000 by light scattering method, average Mn ~10,000 by GPC), trimethylolpropane triacyclidic ether, tert-butanol, iodine (I$_2$), guanidinium thiocyanate (GuSCN), 4-tert-butylpyridine (tBP), acetonitrile (AN), and valeronitrile (VN) were purchased from Sigma-Aldrich. Fluorine-doped SnO$_2$ (FTO) glass, Ti-nanoxide T/SP, Ti-nanoxide R/SP, cis-disothiocyanato-bis(2,2'-bipyridyl-4,4'-dicarboxylato) ruthenium(II) bis(tetrabutyl-ammonium) (N719), 1-butyl-3-methylimidazolium iodide (BMI) were purchased from Solaronix SA. Spray gun (AIR BRUSH KIT, ABS-130) was purchased from Bluebird.

Preparation of the stainless steel counter electrode with a composite of CB and cross linked (XL) 3D network polymers. A SS substrate was cleaned using water and ethanol, then dried using air blow. The cleaned SS was placed onto the hotplate at 120°C. Trimethylolpropane triacyclidic ether and polyethyleneimine was dissolved in ethanol. Various wt% (3, 6, 12, 20, 40, 55, 86 and 90 wt%) of CB was dispersed in the solution through ultrasonication (750 W, Sonics & Materials, Inc, Newtown, CT, USA) for 10 min. Firstly, the mixed solution with small wt% of CB to polymer was sprayed onto the SS substrate to form a CCPL. Subsequently, the mixed solution with large wt% of CB to polymer was sprayed onto a CL. Each spray pyrolysis step was taken for 5 minutes in speed of 2 mL min$^{-1}$. In-situ cross-linking network polymerization instantly occurred as soon as the mixed solution sprayed at the hot SS substrate or the hot CCPL.

References

1. Ito, S. et al. Fabrication of thin film dye sensitized solar cells with solar to electric power conversion efficiency over 10%. Thin Solid Films 516, 4613–4619 (2008).
2. Chang, J. et al. Co-sensitization promoted light harvesting for organic dye-sensitized solar cells using unsymmetrical squaraine dye and novel pyrenoimidazole-based dyes. J. Power Sources 240, 779–785 (2013).
3. J. Lau, G. P. S. et al. Robust High-performance Dye-sensitized Solar Cells Based on Ionic Liquid-sulfolane Composite Electrolytes. Sci. Rep. 5, 18158 (2015).
4. Ramasamy, E. & Lee, J. Ferrocene-derivatized ordered mesoporous carbon as high performance counter electrodes for dye-sensitized solar cells. Carbon 48, 3715–3720 (2010).
5. Choi, J., Kang, G. & Park, T. A Competitive Electron Transport Mechanism in Hierarchical Homogeneous Hybrid Structures Composed of TiO$_2$, Nanoparticles and Nanotubes. Chem. Mater. 27, 1359–1366 (2015).
6. Buhbut, S. et al. Controlling dye aggregation, injection energies and catalytic recombination in organic sensitzer based dye cells using a single electrolyte additive. Energ. Environ. Sci. 6, 3046–3053 (2013).
7. Yoon, J., Kang, D. K., Won, J., Park, J. Y. & Kang, Y. S. Dye-sensitized solar cells using ion-gel electrolytes for long-term stability. J. Power Sources 201, 395–401 (2012).
8. Kwon, Y. S. et al. Exploring the heterogeneous interfaces in organic or ruthenium dye-sensitized liquid- and solid-state solar cells. ACS Appl. Mater. Inter. 4, 3141–3147 (2012).
9. Kwon, J. et al. Nanopatterned conductive polymer films as a Pt, TCO-free counter electrode for low-cost dye-sensitized solar cells. Nanoscale 5, 7838–7843 (2013).
10. Yoo, B. et al. Titanium nitride thin film as a novel charge collector in TCO-less dye-sensitized solar cell. J. Mater. Chem. 21, 3077–3084 (2011).
11. O’Regan, B. & Grätzel, M. A low-cost, high-efficiency solar cell based on dye-sensitized colloidal TiO₂ films. Nature 353, 737–740 (1991).
12. Maitra, M. W. et al. A dye sensitized solar cell using natural counter electrode and natural dye derived from mangosteen peel waste. Sci. Rep. 5, 15230 (2015).
13. Yeh, M. H. et al. A low-cost counter electrode of ITO glass coated with a graphene/Nafion® composite film for use in dye-sensitized solar cells. Carbon 50, 4192–4202 (2012).
14. Park, S. H., Lim, J., Song, I. Y., Lee, J. R. & Park, T. Physically Stable Polymer-Membrane Electrolytes for Highly Efficient Solid-State Dye-Sensitized Solar Cells with Long-Term Stability. Adv. Energ. Mater. 4, 201300489 (2014).
15. Kim, H. S., Ko, S. B., Jang, I. H. & Park, N. G. Improvement of mass transport of the [Co(bpy)₃]²⁺/III redox couple by controlling nanostructure of TiO₂ films in dye-sensitized solar cells. Chem. Commun. 47, 12637–12639 (2011).
16. Song, I. Y., Kwon, Y. S., Lim, J. & Park, T. Well-defined all conducting block copolymer bilayer hybrid nanostructure: selective positioning of lithium ions and efficient charge collection. ACS Nano 8, 6893–6901 (2014).
17. Dor, S., Grinis, L., Rhule, S. & Zaban, A. Electrochemistry in Mesoporous Electrodes Influence of Nanoporosity on the Chemical Potential of the Electrolyte in Dye-Sensitized Solar Cells. J. Phys. Chem. C 113, 2022–2027 (2009).
18. Lim, J., Kwon, Y. S. & Park, T. Effect of coadsorbent properties on the photovoltaic performance of dye-sensitized solar cells. Chem. Comm. 47, 4147–4149 (2011).
19. Han, H. G. et al. Ultralast Fabrication of Flexible Dye-Sensitized Solar Cells by Ultrasonic Spray-Coating Technology. Sci. Rep. 5, 14645 (2015).
20. Chang, Q. et al. Nanoarchitecture of variable sized graphene nanosheets incorporated into three-dimensional graphene network for dye sensitized solar cells. Carbon 85, 185–193 (2015).
21. Kay, A. & Graetzel, M. Artificial photosynthesis. 1. Photosensitization of titania solar cells with chlorophyll derivatives and related natural porphyrins. J. Phys. Chem. 97, 6272–6277 (1993).
22. Tranclík, J. E., Barton, S. C. & Hone, J. Transparent and Catalytic Carbon Nanotube Films. Nano Letters 8, 982–987 (2008).
23. Xia, J., Masaki, N., Jiang, K. & Yanagida, S. The influence of doping ions on poly(3,4-ethylenedioxythiophene) as a counter electrode of a dye-sensitized solar cell. J. Mater. Chem. 17, 2845 (2007).
24. Murakami, T. N. & Grätzel, M. Counter electrodes for DSC: Application of functional materials as catalysts. Inorg. Chem. Acta. 361, 572–580 (2008).
25. Olsen, E., Hagen, G. & Lindquist, S. E. Dissolution of platinum in methoxy propionitrile containing LiI/I₂. J. Electrochem. Soc. 157, 988–992 (2010).
26. Cai, F., Liang, J., Tso, Z., Chen, J. & Xu, R. Low-Pt-loading acetylene-black cathode for high-efficient dye-sensitized solar cells. J. Power Sources 177, 631–636 (2008).
27. Kaskel, S., Schlichte, K. & Kratzke, T. Catalytic properties of high surface area titanium nitride materials. J. Mol. Catal. A Chem. 208, 291–298 (2004).
28. Wang, M. et al. CoS Supersedes Pt as Efficient Electrocatayst for Triiodide Reduction In Dye-Sensitized Solar Cells. J. Am. Chem. Soc. 131, 15976–15977 (2009).
29. Lee, B., Buchholz, D. B. & Chang, R. P. H. An all carbon counter electrode for dye sensitized solar cells. Energ. Environ. Sci. 5, 6941–6952 (2012).
30. Huang, Z. et al. Application of carbon materials as counter electrodes of dye-sensitized solar cells. Electrochem. Commun. 9, 596–598 (2007).
31. Lee, W. J., Ramasamy, E., Lee, D. Y. & Song, J. S. Efficient dye-sensitized solar cells with catalytic multiwall carbon nanotube counter electrodes. ACS Appl. Mater. Inter. 1, 1145–1149 (2009).
32. Ito, S. & Mikami, Y. Porous carbon layers for counter electrodes in dye-sensitized solar cells: Recent advances and a new screen-printing method. Pure Appl. Chem. 83, 2089–2106 (2011).
33. Costa, R. D., Lodermeyer, F. & Guldi, D. M. Recent advances in multifunctional nanocarbons used in dye-sensitized solar cells. Environ. Environ. Sci. 7, 1281–1296 (2014).
34. Veerapan, G., Bojan, K. & Rhee, S. W. Amorphous carbon as a flexible counter electrode for low cost and efficient dye sensitized solar cell. Renew. Energy 41, 383–388 (2012).
35. Kay, A. & Grätzel, M. Low cost photovoltaic modules based on dye sensitized nanocrystalline titanium dioxide and carbon powder. Sol. Energ. Mat. Sol. C 44, 99–117 (1996).
36. Chang, L. H. et al. A graphene-multi-walled carbon nanotube hybrid supported on fluorinated tin oxide as a counter electrode of dye-sensitized solar cells. J. Power Sources 222, 518–525 (2013).
37. Donnet, J. B. Nano and microcomposites of polymers elastomers and their reinforcement. Compos. Sci. Technol. 63, 1085–1088 (2003).
38. Wrobłowa, H. S. & Saunders, A. Flow-through electrodes: II. The I₁/I⁻ redox couple. Electroanal. Chem. Interfacial. Electrochem. 42, 329–346 (1973).
39. Li, P. et al. High-performance and low platinum loading Pt/Carbon black counter electrode for dye-sensitized solar cells. Sol. Energy 83, 845–849 (2009).
40. Murakami, T. N. et al. Highly Efficient Dye-Sensitized Solar Cells Based on Carbon Black Counter Electrodes. J. Electrochem. Soc. 153, A2255–A2261 (2006).
41. Kamau, G. N. Surface preparation of glassy carbon electrodes. Anal. Chim. Acta. 207, 1–16 (1988).
42. Fan, B., Mei, X., Sun, K. & Ouyang J. Conducting polymer/carbon nanotube composite as counter electrode of dye-sensitized solar cells. Appl. Phys. Lett. 93, 143103 (2008).
43. Lim, J., Ryu, S. Y., Kim, J. & Jun, Y. A study of TiO₂/carbon black composition as counter electrode materials for dye-sensitized solar cells. Nanoscale Res. Lett. 8, 227 (2013).
44. Chen, P. W. et al. Dye-sensitized solar cells with low-cost catalytic films of polymer-loaded carbon black on their counter electrode. RSC Adv. 3, 5871–5881 (2013).
45. González-Dominguez, J. M., Díez-Pascual, A. M., Ansón-Casaos, A., Gómez-Fatou M. A. & Martínez M. Epoxy composites with covalently anchored amino-functionalized SWNTs: Towards the tailoring of physical properties through targeted functionalization. J. Mater. Chem. 21, 14948–14958 (2011).
46. Huang, J. et al. Thermomechanical properties of polyimide-epoxy nanocomposites from cubic silsesquioxane epoxides. J. Mater. Chem. 14, 2858–2863 (2004).
47. Jun, Y., Kim, J. & Kang, M. G. A study of stainless steel-based dye-sensitized solar cells and modules. Sol. Energ. Mat. Sol. C 91, 779–784 (2007).
48. Li, H. et al. Highly flexible, low-cost, all stainless steel mesh-based dye-sensitized solar cells. Nanoscale 6, 13203–13212 (2014).
49. Wilson, R. G. Vacuum Thermionic Work Functions of Polycrystalline Be, Ti, Cr, Fe, Ni, Cu, Pt, and Type 304 Stainless Steel. J. Appl. Phys. 37, 2261–2267 (1966).
50. Huang, X. et al. Stainless steel mesh-based flexible quasi-solid dye-sensitized solar cells. Sol. Energ. Mat. Sol. C 94, 1005–1010 (2010).
51. Balasigam, S. K., Kang, M. G. & Jun, Y. Metal substrate based electrodes for flexible dye-sensitized solar cells: fabrication methods, progress and challenges. Chem. Comm. 49, 1147–1157 (2013).
52. Huang, J. C. Carbon black filled conducting polymers and polymer blends. Adv. Polym. Tech. 21, 299–313 (2002).
53. Lim, J., Kim, T. & Park, T. Fast cascade neutralization of an oxidized sensitizer by an in situ-generated ionic layer of I- species on a nanocristalline TiO2 electrode. Energ. Environ. Sci. 7, 4029–4034 (2014).

54. Huang, S. Y., Schlichthorn, G., Nozik, A. J., Gratzel, M. & Frank, A. J. Charge Recombination in Dye-Sensitized Nanocrystalline TiO2 Solar Cells. J. Phys. Chem. B 101, 2576–2582 (1997).

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**Author Contributions**

T.P. and G.K. designed the study. G.K. executed the experiments. J.C. carried out the scanning electron microscope. T.P. and G.K. wrote the manuscript. All authors read and approved the final manuscript.

**Additional Information**

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