

A cracked polymer templated metal network as a transparent conducting electrode for ITO-free organic solar cells†

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We report a highly transparent, low resistance Ag metal network templated by a cracked polymer thin film and its incorporation in an organic solar cell. The performance of this scalable metallic network is comparable to that of conventional ITO electrodes. This is a general approach to replace ITO in diverse thin film devices.

Organic solar cells (OSCs) are an attractive option for large area and inexpensive production of modules. This is because OSCs are well suited to low cost manufacturing due to simple processing¹ steps that are easily translated to roll-to-roll mass production² and lead to near commercialization. In the recent past, extensive research has been carried out on the active semiconductor ingredients³ and electrode optimization⁴ in order to improve the OSC performance,⁵ which is typically poorer compared to other types of solar cells such as dye sensitized solar cells⁶ and silicon based solar cells.^{7,8} Nonetheless, OSCs occupy a unique position in the photovoltaic roadmap as solid state, flexible, environmentally benign and ultralight large area devices.⁹

In OSCs, indium tin oxide (ITO) is the most commonly used transparent conducting electrode. Best ITO films exhibit a transmittance of 92% in the visible region and a sheet resistance of 10 ohm \square^{-1} .¹⁰ Indium is scarce and expensive;¹¹ ITO requires high temperature processing, is brittle and develops cracks on flexible substrates.¹² In order to address these issues, there has been much effort towards alternative electrodes, as reported in the literature. Graphene,¹³ carbon nanotubes (CNTs),^{14,15} and Ag nanowires^{16,17} have been proposed as alternatives to ITO. Graphene has outstanding optical properties, but a limited sheet resistance of 30 ohm \square^{-1} .¹³ The CNT

networks also exhibit high sheet resistance and in addition, are less stable under ambient conditions.^{14,18} Ag nanowire networks show relatively superior performance in terms of transmittance and sheet resistance.¹⁷ But like any other network produced from pre-synthesized nanowires (tubes), they suffer from contact resistance at innumerable crossbar junctions and high roughness.^{19,20} Further, the redundant wires/tubes in the network can short the OSCs, which restricts its use in roll-to-roll fabrication. Often, they require an extra treatment, such as mechanical pressing,²¹ thermal treatment²⁰ or poly(3,4-ethylenedioxythiophene) poly(styrenesulfonate) (PEDOT:PSS) coating to improve the performance.¹⁹

Metal grids offer attractive alternatives to the above TCEs.^{22–30} Indeed being free of junctions, they exhibit high performance in terms of transmittance and sheet resistance, and have been successfully used in optoelectronic devices including OSCs. Typically, they are produced by patterning using lithographic techniques such as photolithography,²² soft lithography,²³ phase shift lithography²⁴ and nanoimprint lithography.²⁵ A technologically relevant and simple alternative is the direct printing of grids using silver nanoparticle inks either by ink-jet printing^{26,27} or laser printing^{27–29} or flexographic and thermal imprinting.³⁰ These recipes for Ag grids result in relatively large feature sizes. Such printed patterns are often used as current collecting grids in diverse applications.

Recently, metal meshes have been fabricated based on cracked TiO₂ templates and their application in touch screens has been shown.^{31,32} Some of us³³ have shown, very recently, selective deposition of Cu by electroplating in the cracked regions of a polymer template to produce Cu mesh based TCEs. These cracked template methods are highly scalable but need to be tested for their applicability and device integration in thin film organic devices such as OSCs, OFETs, *etc.* One of the basic challenges is to obtain high light transmission maintaining low sheet resistance for such TCEs. Additionally, in thin film devices, the uniformity and connectivity of such meshes are very critical for the reproducibility of such devices.

In this study, we have explored the feasibility of cracking as a tool for the preparation of such metal meshes from inexpensive

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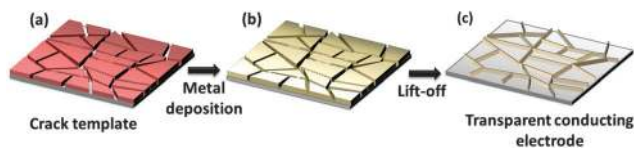


Fig. 1 Schematic illustration of transparent conducting electrode fabrication (a) cracked template (b) deposition of metal on a cracked template and (c) lift-off of the template giving rise to the Ag network based TCE.

polymer templates at room temperature and integrating them as TCEs in bulk heterojunction OSCs. Specifically, using a seamless Ag network in the form of a mesh as a replacement for ITO electrodes, several OSCs having an inverted geometry have been made, which exhibited performance comparable to those produced using ITO. We like to note that this work emphasizes the suitability and reproducibility of such an ITO-free TCE in OSCs and not specifically the optimization of efficiency of any kind of device.

The process flow for templating is shown schematically in Fig. 1. First, a crack layer is produced by drying an acrylic based colloidal dispersion on a glass substrate wherein highly interconnected cracks are spontaneously obtained. These U-shaped grooves are complete cracks down to the substrate to be distinguished from incomplete cracks.³⁴ Using the cracked template, the metal (Ag, 55 nm) is deposited by vacuum evaporation and subsequently, the template is washed away with chloroform.

As shown in Fig. 2a, the Ag mesostructures on glass appear to be well interconnected throughout the network. The metal fill factor is estimated to be $\sim 20\%$ with a structural width of $\sim 2 \mu\text{m}$ and an average cell size (spacing between the Ag structures) of 20 to 60 μm . The optical profilometric image ($1.2 \times 0.8 \text{ mm}^2$) in Fig. 2b not only reveals the connectivity of the Ag network over a large area but also shows its seamless nature. The surface roughness of the network is estimated to be $\sim 5 \text{ nm}$

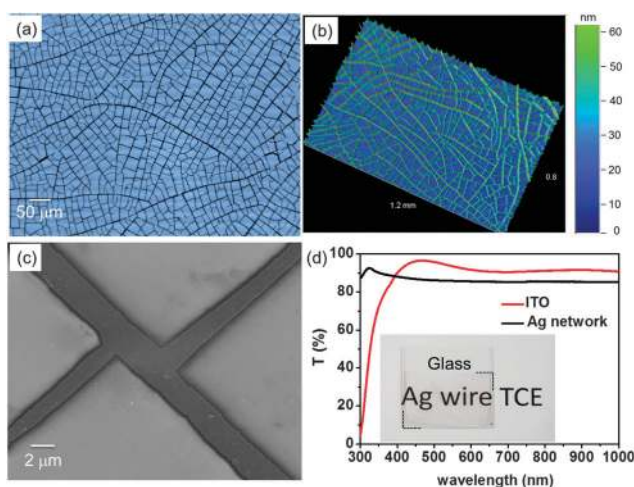


Fig. 2 Characterization of the Ag network derived TCE: (a) the optical microscope image (transmission mode), (b) the optical profiler image in a 3D view, (c) the SEM image of a network junction and (d) comparison of transmittance spectra of Ag metallic network and ITO. The inset shows the photograph of a Ag network TCE on a glass substrate.

while the peak-to-valley roughness, which corresponds to the network thickness, is 55 nm. The SEM image in Fig. 2c shows a junction where the network surface continues to be smooth and the junction itself is seamless unlike crossbar junctions commonly seen in network TCEs made from pre-synthesized Ag nanowires.²¹ Accordingly, the sheet resistance was found to be $\sim 10 \text{ ohm } \square^{-1}$, which is rather low considering the thickness of only 55 nm. The transmittance of the Ag network was $\sim 86\%$ at 550 nm (see Fig. 2d) and the overall transmittance is comparable to that exhibited by ITO but, importantly, extends down to the UV region (see the photograph in the inset). ITO, on the other hand, is a good UV absorber.

In the next step of solar cell fabrication, the ZnO barrier layer was brought in by decomposing a zinc acetate film at $150 \text{ }^\circ\text{C}$ in air (see ESI† for experimental details).³⁵ For a proposed thickness of 45 nm of ZnO, the obtained layer was non-uniform and discontinuous (Fig. 3a) as evident from the dark and grey regions, the latter corresponding to the ZnO covered network. Beiley *et al.* described previously the fabrication of a Ag nano-wire–ZnO nanoparticle composite to obtain a semi-transparent top electrode and achieved a good power conversion efficiency (PCE) of 4.3% in PBDTTPD/PC₇₀BM devices.³⁶ Based on this fact, we increased the thickness of the ZnO layer to 95 nm, 135 nm and 230 nm. We observed an increasing degree of covering of the metal network with the increasing thickness of ZnO. For example, in the SEM image of the 135 nm ZnO layer shown in Fig. 3b, the ZnO film is seen to be uniformly submerging the entire Ag network. This is found to be the optimal ZnO thickness required to submerge the 55 nm thick Ag network so as to have reproducible devices without shorting.

Four bottom-illuminated inverted photovoltaic cells (5 devices in each) consisting of poly-3-hexylthiophene (P3HT) and phenyl-C61-butyric acid methyl ester (PCBM) were fabricated on a glass substrate, as shown schematically in Fig. 4a using a Ag network derived TCE. The bar graphs of solar cell parameters (V_{oc} , J_{sc} , FF and PCE) for all four cells with varying thicknesses of the ZnO barrier layer are shown in Fig. 4b. In cell 1, all five devices with a 45 nm ZnO layer were defective due to shorting. For cell 2 with a 95 nm ZnO layer, only two devices were working with moderate efficiencies. On the other hand, for cells 3 and 4 with 135 and

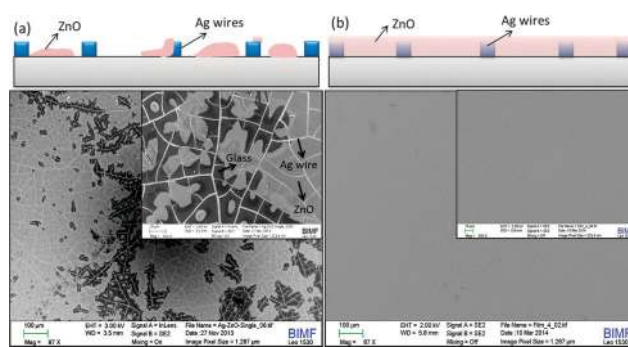


Fig. 3 SEM images and schematic illustrations of the Ag network TCE covered with ZnO layer having a thickness of (a) 45 nm and (b) 135 nm. For more SEM images, see the ESI,† Fig. S1.



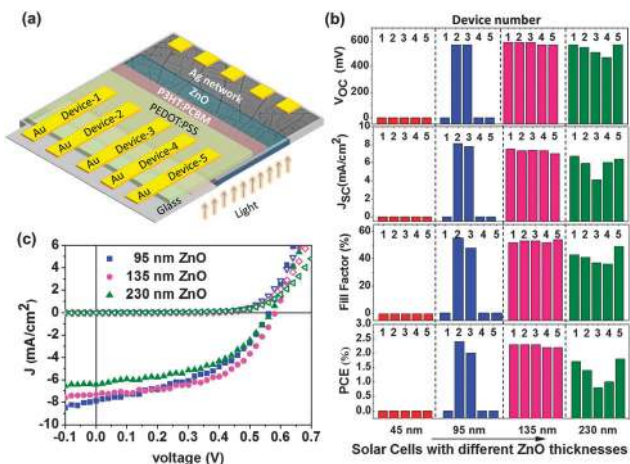


Fig. 4 (a) Schematic illustration of the inverted P3HT-PCBM solar cell with Ag network TCE, (b) bar graph of cell parameters for solar cells of different ZnO layer thicknesses (45, 95, 135 and 230 nm), (c) examples of J - V characteristics of solar cells with different ZnO layer thicknesses in the dark (open symbols) and in light (filled symbols). Note that the best performing solar cell is chosen from each cell.

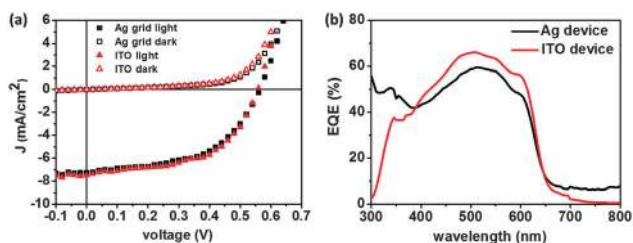


Fig. 5 Comparison of (a) J - V characteristics and (b) EQE for the optimized Ag network and ITO devices.

230 nm ZnO barrier layer thicknesses, respectively, all five devices were found to be functioning. Significantly, the five devices with a 135 nm ZnO layer (cell 3), showed uniform performance with an average efficiency of $2.26 \pm 0.05\%$, while those with a 230 nm ZnO layer exhibited some variations in performance with lower efficiency ($1.34 \pm 0.43\%$, see Fig. 4b).

The decrease in the cell efficiency for higher ZnO thicknesses can be attributed to higher series resistance of the cell leading to lower current density and fill factor values (see Fig. 4b).³⁵ Indeed, the variations in solar cell parameters among the five devices exhibit a similar trend. The J - V characteristics of typical solar cells with different ZnO layer thicknesses (95 nm, 135 nm, and 230 nm) are shown in Fig. 4b (see the ESI,† Fig. S2 for J - V characteristics of all working devices).

From Fig. 5a, we can compare the J - V characteristics of a typical device fabricated using a Ag network (from cell 3) with

that produced using ITO as a TCE (ITO: $T = 93\%$ and $R_s = 16 \text{ ohm } \square^{-1}$). The Ag network cell has an optimum ZnO layer thickness of 135 nm. For ITO devices, the thickness of the ZnO blocking layer was kept at 45 nm which is optimum for an inverted solar cell.³⁵ All other process parameters such as the active layer thickness, top electrode, annealing temperature, *etc.* were kept the same for all the devices. It is clear from the plot that the Ag network TCE based solar cell follows a similar trend, as the ITO based cell and the derived parameters are quite comparable (see Table 1). Thus, the efficiency of the Ag network TCE based solar cell was 2.14%, while that obtained for ITO based cells under similar ambient conditions was 2.27%.

These results presented here clearly demonstrate the potential of the Ag network based TCE as an alternative to ITO in OSCs. However, a slightly thicker ZnO layer is required to cover the Ag network thickness. Since the ZnO is filled in the interstitial domains between the metallic network, it improves the charge collection properties, which is similar to coating a PEDOT:PSS layer as reported for conventional metal grids.¹⁹ The external quantum efficiency (EQE) measurements (Fig. 5b) show that the Ag network TCE based cell exhibits slightly less values in the visible region compared to those derived from ITO cells, but in the UV region, the former excels which makes the overall performance similar for both cells. This can be clearly understood from the differences in transmission of the respective electrodes in the UV region (Fig. 2d). The same information can be inferred by comparing the absorption spectra of the two solar cells (ESI,† Fig. S3) with their EQE spectra.

The method developed in this study for the TCE and associated OSC fabrication has several merits. While the present study has focused on only one type of metallic network, its scope of application can be easily extended to other thin layer devices by varying the template thickness and other parameters in the initial stages of crack template formation. Thus, it is possible to obtain TCEs with different network thicknesses and connectivity and importantly with different metals. Here, Ag was used as a typical example since the work function is favorable for P3HT-PCBM solar cells having an inverted geometry. However, it can be replaced with other metals such as Cu, Al, and Pt depending on the need of application. The metal network based TCE used in the present study works as a current collecting grid at a tens of micrometer scale with a metal fill factor of 20%.

Conclusions

In conclusion, we demonstrated the incorporation of a highly transparent and highly conducting Ag metallic network obtained using a cracked polymer template, in a thin film organic solar cell for the first time. The performance of this

Table 1 Summary of the solar cell parameters corresponding to J - V characteristics shown in Fig. 5a. Note, that a Ag grid device has a 135 nm thick ZnO layer, whereas an ITO device has a ZnO layer of only 45 nm

TCE	V_{oc} (V)	J_{sc} (mA cm^{-2})	FF (%)	PCE (%)	R_{sh} ($\Omega \text{ cm}^{-2}$)	R_s ($\Omega \text{ cm}^{-2}$)@ V_{oc}	R_s ($\Omega \text{ cm}^{-2}$)@ P_{max}
Ag grid	0.57	7.2	51.8	2.14	421	8.3	49.7
ITO	0.55	7.5	55.2	2.27	413	6.7	48.6



highly scalable metal network as a TCE is comparable to that of conventional ITO electrodes. This is a general approach to replace ITO in diverse thin film devices. The crack template approach is universal for any type of metal or substrate material.

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