A Study of perovskite solar cell with Fe3+/Ga3+ doped TiO2 layer

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Abstract

Recently, organic-inorganic lead halide perovskite solar cells (PSCs) have emerged as an extremely attractive photovoltaic device. Perovskite has a structure of a large number of solar cells, which is very promising for the rapid commercialization of large-scale, low-cost solar cells. PSCs have many advantages as it uses simple processing techniques and inexpensive raw materials. The conduction band of TiO2 used in many PSCs coincides well with the conduction band of the perovskite active layer. In addition, TiO2 has excellent stability and conductivity and can be obtained by various deposition methods at a low cost. However, TiO2 traps electrons in oxygen vacancies and Ti lattice sites, which leads to recombination, which in turn reduces the efficiency of the device. In this study, we investigated the effects of Fe³⁺, Ga³⁺ ions on the efficiencies of PSCs by doping them in a compact and mesoporous layer of TiO2. Fe3+, Ga3+ with an atomic radius similar to that of Ti⁴⁺ can replace Ti cations, and Fe³⁺, Ga³⁺ doped TiO₂ is considered to be a better candidate for photoelectric performance of PSCs.

1. Introduction

During the last three decades, the development of renewable energy technologies has received substantial attention to overcome the effects of energy crisis, environmental pollution, and global warming. Among the various renewable and sustainable energy resources, solar power is the most remarkable, because of its abundance, global availability, and cleanliness. [1] Recently, organic-inorganic lead halide perovskite solar cells (PSCs) have emerged as an extremely attractive photovoltaic device. [2-5] Perovskite has a structure of a large number of solar cells, which is very promising for the rapid commercialization of large-scale, low-cost solar cells. PSCs have many advantages as it uses simple processing techniques and inexpensive raw materials. Perovskite materials have the general formula of ABX, where A is generally methyl ammonium (CH₃NH³⁺ or MA), B is a metal ion, such as Pb or Sn, and X represents a halogen ion, such as I, Cl, or Br. A distinct advantage of lead-based perovskites (i.e., MAPbX₃) is that their band gap can be easily tuned from 1.2 to 2.3 eV by varying the composition of cations and anions. [6-7] The conduction band of TiO₂ used in many PSCs coincides well with the conduction band of the perovskite active layer. In addition, TiO₂ has excellent stability and conductivity and can be obtained by various deposition methods at a low cost. In this regard, in many PSCs, TiO2 is widely used as the ETL owing to its excellent electronic properties.[8-11] However, TiO₂ traps electrons in oxygen vacancies and Ti lattice sites, which leads to recombination, which in turn reduces the efficiency of the device. Therefore, the generation of oxygen vacancies and reduction in the number of defects of TiO₂ facilitate the development of stable PSCs. Many studies have shown that the efficiency can be effectively improved by reducing the recombination of TiO₂ by doping appropriate metal ions.

In this study, we investigated the effects of Fe³⁺, Ga³⁺ ions on the efficiencies of PSCs by doping them in a compact and mesoporous layer of TiO₂. Fe³⁺, Ga³⁺ was dissolved in ethanol and mixed with a TiO₂ solution in various ratios. The metal ion can effectively enhance the electron transfer of TiO₂. Fe³⁺ with an atomic radius similar to that of Ti⁴⁺ can replace Ti cations, and Fe³⁺, Ga³⁺ doped TiO₂ is considered to be a better candidate for photoelectric performance of PSCs. Upon the doping with Fe³⁺, the conductivity increased and the recombination decreased, which increased the efficiency of the PSC.

2. Results & discussion

The XRD measurement is performed to reveal the lattice structure. As shown in Figure 1 (a), the Ga^{3+} , Fe^{3+} -doped TiO_2 layer consists of the anatase phase, as the Ga^{3+} , Fe^{3+} ion radius is very similar to that of Ti^{4+} and thus the Fe^{3+} dopant cannot change the lattice of TiO_2 .

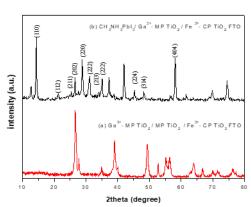


Fig. 1 XRD patterns of MAPbI₃, Fe³⁺, Ga³⁺ - TiO₂.

Figure 1 (b) shows the XRD pattern of $TiO_2/CH_3NH_3PbI_3$ with peaks at 13.8, 19.7, 23.1, 24.2, 28.1, 31.5, 40.3, and 42.8° attributed to the (110), (112), (211), (222), (224), and (314) lattice planes, respectively.

Figure 2 (a) is a mesoporous TiO₂ layer made of diluted 18NR-T. Figure 2 (b) is a cross-sectional view of a mp-TiO₂/

Ga-doped mp-TiO₂ layer prepared by spin-coating a mesoporous 18NR-T. Figure 2 (b) is a cross-sectional view of a mp-TiO₂/Ga-doped mp-TiO₂ layer prepared by spin-coating a mesoporous TiO₂ layer prepared by adding gallium nitrate to a solution of the same concentration. figure 2(C) is a cross-sectional view of the PSC produced, and CP-TiO₂ layer was prepared by mixing iron nitrate and TiO₂ sol. The Fe³⁺ doped TiO₂ / mesoporous TiO₂ / Ga³⁺ doped TiO₂ / perovskite / HTL / electrode structures were fabricated.

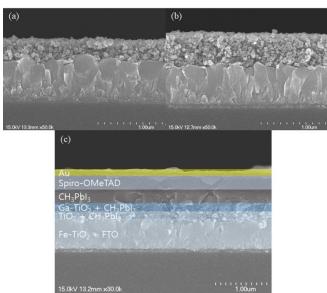


Fig. 2 Scanning Electron Microscope images of (a) mesoporous TiO₂ layer, (b) Gallium doped mesoporous TiO₂ layer on mesoporous TiO₂ layer and (c) Fabricated perovskite solar cell.

Figure 3 is a graph of the absorbance of the TiO₂ layer. It can be seen that the bandgap changes when iron is doped and when gallium is doped, which suggests that a valence band can be distributed between the FTO and the perovskite layer.

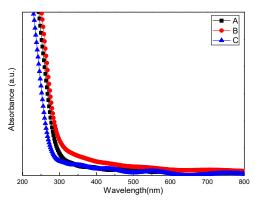


Fig. 3 The absorbance graph of (a) mesoporous TiO_2 layer, (b) Fe doped TiO_2 layer and (c) Ga doped TiO_2 layer.

The J-V curve in Figure 4 shows that the PSCs photovoltaic properties are improved. The corresponding photovoltaic properties are summarized in Table I. Oxygen vacancies or Ti lattice sites are converted to Ti³⁺ by stoichiometric defects in the TiO₂ lattice. Therefore, the Ti³⁺ lattice defects can induce low energy levels and the conduction band can trap electrons

and reduce the electron transport efficiency, leading to recombination. The increase in open-circuit voltage (V_{oc}) is worth discussing. When TiO_2 is doped with metal cations, the TiO_2 energy level is changed and the V_{oc} value is increased.

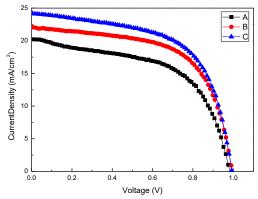


Fig. 4 J–V curves of the PSCs of (a) CP TiO₂/MP TiO₂, (b) Fe³⁺-CP TiO₂/MP TiO₂, (c) Fe³⁺-CP TiO₂/Ga³⁺-MP TiO₂.

Table I Current density-voltage characteristics of best performing

perovskite solar cell				
	V_{oc}	J_{sc}	FF	η
	(V)	(mA/cm^2)	(%)	(%)
(a)	0.978	24.25	48.37	11.47
(b)	1.010	23.18	60.19	13.84
(c)	0.993	24.16	59.96	14.39

3. Conclusions

We fabricated PSCs based on MAPbI₃ using Fe³⁺ and Ga³⁺. The TiO₂ layer doped with Fe³⁺ and Ga³⁺ has a uniform surface, and electron recombination is reduced to improve electrical conductivity. The PSC exhibited a short-circuit current density (J_{sc}) of 24.16 mA/cm², V_{oc} of 0.993 V, FF of 59.96%, and efficiency (η) of 14.39%. Therefore, the use of the Fe³⁺, Ga³⁺-doped TiO₂ layer is a promising approach for fabrication of high-performance PSCs.

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