Mطالعه استحالت ساختاری پروسکایت های ترکیب هالوژنی

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چکیده - مواد پروسکایتی به واسطه خواص ایتوالکترونیکی خارق العادّاتی که دارند، در طی سال ها اخیر مورد توجه بسیاری قرار گرفته‌اند. به‌طور خاص، سلول‌های خورشیدی پروسکایتی با بذورهای پاتریکی ایجادار طولیانی مدتی این مواد است. در این پژوهش نحوه ترکیب پروسکایت‌ها با استفاده از دو ذوب بدی سرب و مبلی آلومینیوم کلرید بررسی شد. بدی سرب با استفاده از سرب راهی نشانی به روش ترکیبی پروسکایت کدرنگ شده و به‌طور بدی سرب در عرض‌های مشابه به کمک الگوهای بشری که به‌طور کلی در شرایط مختلف قرار دارد. بیانیه به دست آمده با روش‌های مشخصه‌بایی الگوهای بشری مشخصه‌بایی و همچنین طیف جذب نوری نشان دهنده تغییر مواد ترکیبی در زمان‌های مختلف قرار گرفته است.

کلید واژه - پروسکایت، سلول خورشیدی، استحالت ساختاری، روی جند مرحله ای

Study of Structural Phase Transformation in Halide Pervoskites

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Abstract - Perovskite materials have gained a lot of attention because of their extraordinary optoelectronics properties. Solar cells based on these materials have reached power conversion efficiency of above 20% in a short period of time. However, there has still work to do about their stability and durability. In this work we demonstrate the effect of CH3NH3Cl on PbI2 to produce a perovskite layer. PbI2, the first precursor, was deposited using common spin coating method mentioned in the literature. The deposited layer was exposed to CH3NH3Cl solution. XRD and UV-Visible characterizations were done. The devices fabricated using these layers have shown power conversion efficiency of 5.4%.

Keywords: Solar cells, Perovskites, Sequential deposition, Phase transformation
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1 Introduction

Metal Halide perovskites due to their excellent optoelectronics properties have emerged as a potential absorber layer for thin film solar cells[1–3]. Their high carrier mobility and high optical absorption are the main reasons of their excellent photovoltaic properties[4], [5]. Solar cells fabricated using perovskite materials have reached the efficiencies above 20% recently[6]. Moreover the main advantages of these materials over other photovoltaic technologies are their low cost and facile deposition methods[5], [7]. But still, they lack a long durability and also presence of lead in their structure has caused environmental concerns[5], [8].

Methyl ammonium lead triiodide (MAPbI$_3$) is the main perovskite material that is used as light absorber for solar cells[5], [9]. Mixed halide perovskites are also used to tune the bandgap and durability of fabricated devices and [10–12]. Bromine as another halide has been added to the structure of perovskite[10]. Increased bandgap in comparison to MAPbI$_3$ is the first consequence of adding bromine in the perovskite structure[10]. Another effect of bromine is increasing the stability of perovskite structure. It has been shown by Noh and et al that perovskites with a certain amount of bromine in their structure are more resistant to humidity and other environmental effects[10].

But chlorine as another halide has a different behaviour in perovskite structure. The first solar cell using perovskite as light absorber in the work of Snaitth and co-workers, used MAPbI$_3$,Cl$_x$ [11]. Later studies on MAPbI$_3$,Cl$_x$ have showed a diffusion length longer than 1µm[4]. Yang and co-workers showed that chlorine is escaping the structure during the formation on perovskite layer and it only helps to enlarge the grain boundaries [13].

In this work, we have used methyl ammonium chloride (MACl) to convert PbI$_2$ to perovskite layer. Perovskite layers showed a two phase structure at short times of dipping but for the long times a single phase structure of mixed halide perovskite is fabricated. The best cell fabricated using this method showed a power conversion efficiency (PCE) of 5.4% .

![Figure 1: Fabricated device structure](image)

2 Experimental Details

2.1 Preparing compact layer

FTO-coated glass substrates were patterned by Zn powder and 2M HCl solution etching. The
patterned FTO substrates were cleaned by soap-deionized water solution, followed by ultrasonication at 50°C deionized water, ethanol, and isopropanol, and then subject to an O3/ultraviolet treatment for 20 min to room temperature. First a ltci-TiO2 deposited on FTO and after treated with 200mM TiCl4 at 70° C for 20 minutes[14].

2.2 Deposition of perovskite layer

PbI2 layer was deposited by spin coating of 1.25M solution of PbI2 in N,N-dimethylformamide (DMF). The layers are dipped into a MACl precursor with a concentration of 10mg/mL for different times. It is followed by a thermal annealing of 100° C on a hotplate for 10 minutes.

2.3 Deposition of Spiro-OMeTAD/Au

Spiro-OMeTAD-based hole transporting layer was deposited using preparation method mentioned in previous works [9]. Au as a back contact was deposited by thermal evaporation. The fabricated layer is 100nm thick.

3 Results and Discussion

Fabricating perovskite layer with the sequential steps method is a well-known and common method to produce a uniform and low pin-hole layer[3]. In this regard, we used MACl for the second precursor to convert the first precursor to perovskite. Fabricated perovskite layer in the first time due to the figure 2 has shown a two phase structure. We can see in the XRD results that there are two peaks for perovskite layer. One in 14.1 for the MAPbI3 and one in 15.6 for the mixed perovskite layer. But as we increase the dipping time the peak relating to MAPbI3 is disappearing. Moreover, the main peak in the 12.6 is related to remained PbI2 layer. But as shown in figure 2(black curve), by increasing the time of dipping all the PbI2 is converted to perovskite. XRD results also indicate a high crystalline mixed perovskite layer.

One of the main characteristics for absorber layers in solar cells are their absorption spectrum. MAPbI3 absorbs light from 1.55 eV photon energy and higher[15]. Fabricated perovskite layers have shown two absorption onset, indicating their two phase structure.

4 Conclusion

Fabricated devices using the two phase perovskite layers with the normal device architecture, has reached a PCE of 5.4% and open circuit voltage of 1.02 volts.
A sequential method with a new second precursor was studied. For short periods of dipping in second precursor a two phase perovskite structure was fabricated. As we increased the time of dipping the MAPbI₃ disappeared. Devices fabricated using the two phase perovskite layer has shown a PCE of 5.4% and open circuit voltage of 1.02 volts.

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References


