CH$_3$NH$_3$PbI$_3$ IV Output Parameters Degradation Investigation

M.N. Orlova$^1$, S.I. Didenko$^1$, D.S. Saranin$^1$, O.I. Rabinovich$^1$, A.Y. Krukov$^2$, A.V. Kolesnikov$^2$

$^1$ NUST "MISiS", 4, Leninshiy pros., 119040 Moscow, Russia
$^2$ D.I. Mendeleev University of Chemical Technology of Russia, Miusskay sq, Moscow, Russia

(Received 10 July 2016; published online 29 November 2016)

Organic photovoltaics, based on hybrid inorganic organic optoelectronic perovskites, with structure alkali-metal-halide are the newest emerging technology in the third generation development. Despite tremendous efficiency records, more than 21%, optoelectronic perovskites’ instability prevents their commercialization and mass production. Issues with degradation are caused by various types of environmental influences. The main issues with stability and power loss in devices are linked to moisture, oxygen, temperature, and light-induced structure defects. Initial measurements are taken after long term debugging with minimal aggressive exposure to environmental conditions. In this case, preliminary degradation studies begin from measurements of light-induced effects. In this work, we will present the main trends in degradation of external characteristics during common I-V measurements, in the order of parameters which were effected the least by environmental factors. This investigation was made on fixed CH$_3$NH$_3$PbI$_3$ solar cells with standard 1.5 AM testing and initial efficiencies more than 8%.

Keywords: Organic photovoltaics, Perovskite, Power conversation efficiency, Solar cells.

DOI: 10.21272/jnep.8(4(1)).04004 PACS numbers: 00.05.Tp, 85.60.Jb

1. INTRODUCTION

Third generation of photovoltaic mostly is presented by organic solar cells, organic photovoltaics (OPV) and emerging hybrid perovskites (PS), both of devices are thin film cells. Last 5 years in this field showed big jump in growth of power conversion efficiency (PCE) up to 13% for OPV and over 20% for PS [1-5]. Since beginning XXI century OPV achieved traditional silicone based cells, in its turn commercialized amorphous Si:H (~ 13%) and on the other hand PS cells moves breakthrough towards to crystalline Si, CIGS (Copper Indium Gallium diSelenide) and CdTe, to more than 20% PCE level and now are close to Shockley-Queasier limit [6-10].

Based on competitive laboratory results next development step requires up – scaling and stable work of devices, here scientific area expands to many directions: roll-to-roll deposition techniques, resistance for environment moisture and oxygen with suitable encapsulation, long-term under light stability [11-12].

At least one direction of research is one of the main keys for PS cells development, because physics of elementary processes is still not determined in details. But though basic origins and questions of light-induced degradation are formulated:

1. Degradation due to trapped charges at abrupt organic/cathode interface;
2. Bond breaking, caused by hot polarons/hot excitons;
3. Bipolarons near cathode- semiconductors interface;
4. UV light affecting.

Firstly, it’s necessary to note, that accordingly to PS advantage in film processing – low formation energy between anion (for example MAI – methyl ammonium iodide – CH$_3$NH$_3$I) and cation (for example PbF$_2$), it is easy to make, but in opposite easy to pull apart by heat, electric field and light (eq.1) [13].

\[
2I \frac{\leftrightarrow}{\text{underlight}} L_s + 2e^- \quad (1a)
\]

(on the donor – acceptor interface)

\[
3\text{CH}_3\text{NH}_3^+ \leftrightarrow 3\text{CH}_3\text{NH}_3^+ \uparrow +3\text{H}^+ \quad (1b)
\]

\[
\Gamma + I_s + 3\text{H}^+ + 2\eta \leftrightarrow 3\text{H} \uparrow \quad (1c)
\]

Equation 1 showed only one of possible routes of perovskite dissociation during degradation. Main factor of chemical structure stability lies in optimal crystal compatibility of composing atoms radiuses. Perovskite structure is ABX$_3$ (A-alkali, B- metal (Ph, Sn, Ge), X- halide). Goldschmidt in 1927 formulated special stability factor, depending on atom radius ratio (Eq.2):

\[
t = \frac{r_A + r_X}{\sqrt{(r_B + r_X)}} \quad (2)
\]

For most stable molecules, this factor is 1, while for perovskites it is nearly 0.8. For summary of this description of perovskite instability, report about long-term study shows decreasing of peak characteristics after few days or weeks (depends on structure and composition) and degradation measurement are no longer, then 10 000 hours. Although, promising progress is reported in Snaith and Graetzel’s groups in different works with using of metal oxides, mesoscopic structures, small radius cations doping [14-15].

Influence of degradation on J-V parameters doesn’t decrease all parameters at the same time. Absorption may be stable, while quantum efficiency decreases, exciton quenching by polarons creates trap-states and subbandgaps [16]. In this work, series of periodical measurements on widely-known MA$\text{PbI}_3$ PS solar cell under standard 1.5 AM light conditions was made. Light-induced degradation firstly decreases quantum efficiency with result fall, maximum power decrease was showed. Than C-H bonds starts to break, that causes shunt resistance losing and leaks increasing. Finally dissociation of perovskites leads to absorption loss and impairment of cathode- semiconductor interface, as result – J_sc fall and R_sh rise.

2077-6772/2016/8(4(1))04004(4) 04004-1 © 2016 Sumy State University
2. EXPERIMENTAL PROCEDURES

Solar cells based on perovskite CH$_3$NH$_3$PbI$_3$ were produced with planar structure (Fig. 1).

Transmission of PbI$_2$ and MAI (both DYESOL production) previously dissolved in GBL at 1 ppm. Perovskite solution was mixed in 1,2 M concentration using PbI$_2$ and MAI (both DYESOL production) previously dissolved in GBL. DMSO (7:3 by volume), PCBM acceptor solution was prepared in anhydrous chlorobenzene in 10 mg/ml concentration. ITO pixelated substrates (Lumtec, 10 Ohm / □) were cleaned by next steps in ultrasonic bath (20 minutes on each step) – Alkaline detergent 1%, deionized water (18 Mohms), acetone, toluene, methanol, isopropanol (all ME grade) and final UV exposure during 15 minutes for surface activation.

Firstly PEDOT : PSS 1.1%, water solution (Heraeus Clevios) was spincoated on ITO substrates with 3000 rpm speed during 60 sec to grow 30 nm layer, then it was annealed at 150 °C during 10 minutes. After it perovskite film was formed with 2 steps spincoating, from 20 sec at 1000 rpm to 20 sec at 5000 rpm and toluene dripping at last step, annealing was done at 100 °C (10 minutes). PCBM (Solene) layer was spin-coated at 1000 rpm during 40 sec. Finally Al cathode layer was thermally evaporated to achieve 100 nm thickness at 2×10$^{-6}$ Torr vacuum level. Encapsulation was done with Norlan 65 pixel square was obtained as 0.084 cm$^2$ square. I-V measurement was done by Newport Thermo Oriel solar simulator at 1.5 AM spectrum, 100 mW/cm$^2$ irradiation level and by using Keithley 2400 source-measure unit.

3. RESULTS AND DISCUSSION

Firstly initial I-V measurements were done for 4 samples for fixing “as produced” results (Fig. 2 and Table 1), then best sample (№ 3) was chosen for I-V parameters degradation study.

After initial testing sample № 3 was measured periodically with 12 hours interval, and it was clearly observed, that $U_{oc}$ and $J_{sc}$ were staying constant while fill factor (FF) is essentially a measure of quality of the solar cell start its decrease from 0.72 to 0.54 (loss of 33 %) in cause of $R_s$ increasing up to 15 Ohm·cm$^{-2}$ – cathode – PCBM interface degradation.

After initial testing sample № 3 was measured periodically with 12 hours interval, and it was clearly observed, that $U_{oc}$ and $J_{sc}$ were staying constant while FF start its decrease from 0.72 to 0.54 (loss – 33 %) in cause of $R_s$ rise up to 15 Ohm·cm$^{-2}$ – cathode – PCBM interface degradation.

Next 36 hours of measurements were obtained (fig. 3, 4. and tab. 2, 3) with continue decreasing of FF, $R_{sh}$ and $R_s$ and starting degradation of $V_{oc}$ and $J_{sc}$, that characterizes by quantum efficiency decrease and absorbance of perovskite layer and perovskite – PCBM interface.

Thereby it is forming a clear trend for solar cell degradation without encapsulation the complete degradation is observed for 3 days, whereas for the encapsulated solar cells their efficiency loss during 6 weeks was from 3 to 15 %. Overall the curve of efficiency change versus time is presented in figure 3.

Tendencies I-V output parameters have clear order in decreasing of I-V filling area – preliminary power loosing (firstly from 0.72 to 0.54 during during first 36 hours). Then perovskite cells lost their quantum efficiency, that’s causes $J_{sc}$ decreasing (after 72 hours of operation) and finally perovskite molecules lost their stability – decreasing of $V_{oc}$. This affecting is proportional to decreasing of $R_{sh}$ and increasing $R_s$ respectively to degradation of PCBM – Al interface, in its turn it
was obtained in higher value of leaks in dark measurements and degradation of Al cathode-acceptor PCBM interface – bigger slope of UI curve in 1 quadrant in linear region.

Sufficiently high open circuit voltage for the solar cells on an organic basis - more than 0.6 V is due to the low position of the HOMO level donor relatively to the PEDOT: PSS hole-transport layer. This is due to its lower position levels HOMO / LUMO = −5.4 eV / −3.6 eV, and the width of the band gap 1.8 eV. This is value of PCBM increases the open circuit voltage of solar cells based on perovskite to 0.8 V. When working in air the moisture saturation and donor and acceptor oxidation occur, result increased and their specific surface resistance increases. The contacts of the junction metal-semiconductor and semiconductor – a hole-transport PEDOT: PPS layer become high-resistance, and in the heterojunction dark current leakage increases. The effect of forming a solar cell structure under such conditions is clearly seen on the I-V curve. By the type of graph it can be determined the noticed degradation points: Large slope of the curve in the current-voltage characteristics in the first quarter is characterized by high series resistance $R_s$, which reached 100 Ohm-cm$^2$. Large slope of the curve in the third I-V quarter is characterized by the dark current leakage and the decreasing in shunt resistance $R_{sh}$ 1000 Ohm-cm$^2$. The general graph is characterized by insufficient slope of exponential growth, reduced the corresponding values of $R_s$ and $R_{sh}$. The value of leakage is inversely proportional to the shunt resistance $R_{sh}$, which, in turn, should be as large as possible. In the real structures, $R_{sh}$ is characterized by the quality of p-n junction and its isolation from the electrodes to prevent punctures. However, at the same time there was a voltage drop to a value ~ 0.4 B, followed by a decrease in the filling factor. This effect should be related to the deterioration of the contact phases of the donor and acceptor. The process described above gave the expected results, the growth of the main values of output parameters. The output characteristics of the sample № 3 showed the highest efficiency 8.23 %. Besides the standard I–V removal from the solar cell pixel, it was also carried measurements out through the optical aperture calibrated mask with a diameter of 1000 microns. This type I-V measurement on the area equal to 0.9075 cm$^2$ (equivalent circle area with a diameter of 1 mm) allows fixing inhomogeneity in the element microstructure of the on the pixel area thus it becomes possible to determine higher or lower performance results which solar cell is able to produce.

### 4. SUMMARY

The I-V investigation show that output parameters trend have clear order in decreasing of I-V filling area due to preliminary power loss. Then perovskite cells lost their quantum efficiency, that’s causes $J_{sc}$ decreasing and finally perovskite molecules lost their stability – decreasing of $V_{oc}$. This influence is proportional to $R_{sh}$ decreasing and $R_s$ rising, respectively, in its turn it was obtained in higher value of leaks in dark measurements and degradation of Al cathode-acceptor PCBM interface – larger slope of I-V curve in first quadrant in linear region.
region. Degradation occur a lot of goals for scientists, firstly of course it’s molecule stability – doping of perovskite is main trend with two direction – dopants to methyl amin, such as form amide, introducing of small molecule cations – Cs, Ga, Cu an others. This improvements should lift up tolerance factor and finally stabilize operation of devices, that can be checked with transient measurements at power point and optical studies – internal quantum efficiency stable as absorbance level.

ACKNOWLEDGEMENTS

This work was supported by grant of Federal Target Program "Research and development in priority areas of Russian scientific and technological complex for 2014-2020", on 22 August 2014 № 14.574.21.0096 (unique identifier RFMEFI57414X0096).

REFERENCES