Advances in Conversion Efficiency and Thermal Stability of the Perovskite-based Solar Cell: Review

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Abstract. This paper presents a small review on the technological advances made on the perovskite-based solar cell. Through this summary of the results of the research on perovskite, the reader will have an overview of the perovskite material, the different structures of a perovskite solar cell, and the opto-electrical properties of such cell as well as the electrical models used in its simulation. Finally, the paper presents in a very brief way the challenges that this technology will have to overcome before finding its place in the photovoltaic market.

Keywords: perovskite solar cell, review, thermal stability.

Introduction

Among the photovoltaic cell technologies, perovskite solar cell technology has become probably the hottest topic in photovoltaics as can be seen in the number of publications and conference topics on the subject. In 2009, Miyasaka et al. developed a perovskite-based cell with an efficiency of 2.2%. But, by replacing Boron with Iodine, the efficiency improved to 3.8%. Ten years later, the confirmed efficiency of the single junction cell of a perovskite-based cell was 21%. In 2020, a perovskite-based solar cell with a conversion efficiency of 25.5% was achieved by Jeong M. and all. Several models of perovskite-based solar cell structure, elaboration and optimization method and simulation models have been proposed in the published studies. In spite of the visible advances in terms of structure stability and conversion performance, much research effort remains to be done to ensure the thermal stability of this cell which will guarantee its commercial success. The objective of this paper is to highlight these advances and to review the manufacturing and simulation models of the perovskite cell in order to bring together the similarities in the methods and structure that will ensure its thermal stability.

Structure of the perovskite solar cell

Perovskite mineral is used as a photovoltaic solar cell absorber. It can be elaborated from a variety of materials and different synthesis methods. Perovskite was discovered in 1839 in the Ural Mountains in Russia and named after the Russian mineralogist L.A.Perovskite [1]. The chemical formula of this mineral is CaTiO3 (calcium titanium oxide). Compounds that have a similar structure to CaTiO3 (ABX3) are called perovskites. In general, in the ABX3 structure of perovskite, A is a large monovalent cation that occupies the cubooctahedral sites in a cubic space. B is a small divalent metal cation that occupies the octahedral sites and X is an anion (typically a halogen, however, X can be oxygen, carbon or nitrogen). The
structure of perovskite and its stability is quantified by two crucial parameters: the tolerance factor \((t)\) given by equation 1 and the octahedral factor \((\mu)\).

\[
t = \frac{(R_A + R_X)}{\sqrt{2(R_B + R_X)}}
\]

This factor is a ratio of the ionic radius of the divalent cations \(R_A, R_B\) and the radius of the anion \(R_X\). These two parameters are generally between 0.81 and 1.11 for \(t\) and then 0.44 and 0.90 for \(\mu\) [1]. The perovskite-based solar cell has a variety of architecture. As in other photovoltaic cell technologies, the perovskite cell has an electron transport layer (ETL) and a hole transport layer (HTL) in addition to the perovskite absorber layer. Thus, the configuration of the cell is very crucial to expect a high-performance perovskite cell. From the literature, two configurations stand out for perovskite based cells: the planar configuration and the mesoporous configuration. Each configuration can be elaborated according to the conventional structure (N-i-P) or the inverse structure (P-i-N). The following figures illustrates its two configurations.

Figure 1. The perovskite-based solar cell configuration: a) the planar configuration and b) the mesoporous configuration.

The evolution of this technology is meteoric with a yield that has evolved from 3.3% in 2009 to 25% in 2020 (figure 2).

Figure 2. The best annual yield of solar cells based on perovskite.
This performance is partly due to the understanding in the structure of the layers of this cell, the treatment reserved to their interface and the interest that this technology arouses in research. The following table summarizes the different structures of the perovskite-based cell studied in the literature. It is necessary to recall that the choice of a structure aims at the improvement and the stability of the existing cell.

### Table 1. The different structures of the perovskite-based cell studied in the literature.

| Structures | PCE  | Stability time | Ref |
|------------|------|----------------|-----|
| FTO / TiO$_2$ - Cl / MAPbI$_3$ / Spiro-OMeTAD / Au | 21% | 2000 h | [3] |
| FTO / Doped C$_{60}$ / mixed perovskite / Spiro-OMeTAD / Au | 17% | 650 h | [4] |
| Glass / ITO / PEDOT:PSS / MAPbI$_3$ / PCBM / EFGnP$_5$ - F / Al | 14.3% | 30 days | [5] |
| FTO / LBSO / MAPbI$_3$ / PTAA / Au | 21.2% | 120 h | [6] |
| FTO / TiO$_2$ / MAPbI$_3$ / PTAA / Au | 19.6% | 120 h | |
| Glass / FTO / c-TiO$_2$ / mp-TiO$_2$ / mixed perovskite / Spiro-OMeTAD-SWCNT | 15% | 580 h | |
| FTO / c-TiO$_2$ / m-TiO$_2$ / perovskite / Spiro-OMeTAD / Au | 14.6% | 300 h | [7] |
| FTO / c-TiO$_2$ / m-TiO$_2$ / perovskite / ZrO$_2$ / Carbon | 11.9% | 12000 h | [8] |
| FTO / c-TiO$_2$ / m-TiO$_2$ / Cs$_3$M / HTL / Au | 21.2% | 250 h | [8] |
| FTO / c-TiO$_2$ / m-TiO$_2$ / Cs$_3$M / HTL / Au | - | 250 h | |
| ITO / cp-TiO$_2$ / ms-TiO$_2$ / perovskite / PTAA / Au | 20.6% | 160 h | [9] |
| ITO / cp-TiO$_2$ / ms-TiO$_2$ / perovskite / Spiro-mF / Au | 24.8% | 500 h | [10] |
| ITO / cp-TiO$_2$ / ms-TiO$_2$ / perovskite / Spiro-oF / Au | - | - | |
| FTO / bl-TiO$_2$ / mp-TiO$_2$ / MAPb$_{1-x}$Br$_x$ / PTAA / Au | 16.2% | - | [11] |
| FTO / PEDOT:PSS / MAPb$_{1-x}$Cl$_x$ / PCBM / Al | 17.7% | - | [12] |
| ITO / PEIE / TiO$_2$ / perovskite / Spiro-OMeTAD / Au | 19.3% | - | [13] |
| FTO / bl-TiO$_2$ / mp-TiO$_2$ / FAPbI$_3$ / PTAA / Au | 20.2% | - | [14] |

If these different structures of the perovskite cell are inexpensive to develop, it should be noted that the structural, electrical and optical properties of the perovskite cell degrade when exposed to the ambient air. This degradation, which is one of the major challenges to overcome if the technology is to appear on the photovoltaic market, is simply due to the reaction of the perovskite layer with the oxygen in the air and with water vapor.

### Electrical and optical properties of the perovskite solar cell

The optical band gap of the halide-based perovskite materials used is about 1.6 eV. The absorption of these materials covers only a fraction of the ultraviolet and visible light; 45% to 50% of the entire solar spectrum [15] [16].

However, the absorption range of single crystal perovskite is red-shifted [17]. The absorption in single crystal perovskite starts from 850 nm while it starts at 780 nm for polycrystalline thin films [18].

This absorption shift in the single crystal is due to the enhanced use of below-bandgap absorption attributed mainly to the indirect-bandgap absorption transition with a bandgap of 60 meV smaller than the direct bandgap. The absorption coefficient of the below bandgap is small compared to that of the above-gap transition making the below bandgap absorption negligible in polycrystalline thin films but obvious in thick single crystals [19].

Z. Cheu showed that the absorption coefficient of single crystal perovskite is an order of magnitude higher than those of conventional ruthenium dyes. Liu et al., have shown that perovskite material is an excellent absorber in the visible spectrum, but very transparent in the near infrared spectrum [19].
Blessing E. et al [20], observed similar results to those demonstrated by Aharon et al [21], concerning the improvement of the absorption of the perovskite cell and the decrease of the optical conductivity when the volumetric ratio of methylammonium lead bromine introduced in a fixed volume of MAPbI3 increases. The knowledge of the different optical properties and absorption limits and the search for methods to extend the absorption range will contribute to the improvement of the performance of the perovskite technology. This quest for performance improvement also involves simulation studies using electrical models of the solar cell.

**Electrical model of the perovskite cell**

The current-voltage characteristic of a cell does not provide reasonable insight into the actual mechanisms of charge transport, recombination and storage. As such, an evaluation of the response of a solar cell to a small perturbation of various real inputs such as voltage or light in the frequency domain of the cell remains an essential method. These mechanisms are summarized in the form of an equivalent circuit that describes the actual physical processes in terms of the passive electrical element and the voltage and current sources. The perovskite cell has been treated theoretically as an ordinary direct band gap semiconductor [22]. Therefore, it is considered as a simple p-i-n diode governed by the equivalent circuit equations of the said diode. Many authors have used the one-diode (figure 3a) and two-diode models for the equivalent circuit of the perovskite-based solar cell [23] [24]. In addition to these commonly encountered models, other models often used for organic cells have been used to model perovskite solar cell.

Huang et al. have developed for a perovskite-based solar cell, an equivalent circuit model very close to the one developed by Mazhai for organic cells. The model is shown in Figure 3b.

Ebadi et al, proposed an equivalent circuit model for the perovskite cell including electrical components and ionic components [25] (figure 3c).

![Figure 3](image-url) **Figure 3.** Electrical model of perovskite solar cell: (a) one diode equivalent circuit model, Figure (b) The solar cells' lumped-parameter equivalent circuit model proposed by Mazhari [26] and (c) Equivalent circuit for the fitting photo-voltage rise and decay plots [25]
Future challenges in perovskite solar cell

The commercialization of the perovskite-based solar cell remains closely linked to the improvement and stability of its performance. If its efficiency reached 25% in 2020, the future challenge of the cell remains its stability.

J. Zhang et al., have shown that the Spiro-OMeTAD hole transport layer used in the elaboration of the perovskite cell contributes to its poor stability when exposed to ambient air [27].

Much effort has been made to develop hole transport layer to replace the Spiro-OMeTAD layer [28] [29] [30]. This layer could be replaced by a Cu:NiOx layer. However, Damian et al., have shown in their study that using Cu:NiOx as the hole transport layer results in a huge drop in open circuit voltage and short circuit current. On the other hand, by using Cu:NiOx/PTAA, the quality of the cell improves as well as its efficiency [31]. In 2020, Mingyu J. et al., replaced the Spiro-OMeTAD hole transport layer with two fluorinated isomers. This process resulted in a perovskite-based cell with 24.8% efficiency stable at 87% of its efficiency after 500 hours under humidity conditions without encapsulated [10]. Based on this observation, Yanjie Wu sought to improve the stability of the perovskite-based cell and the mobility of the holes by replacing this layer with a MoO3 layer. In addition, they sought to improve the performance of the cell through the extension of the absorption range to the infrared. For this purpose, a PBDB-TF:BTP-4Cl bulk-heterojunction layer was integrated into the cell [32].

In 2018, Constantina E. et al., developed a perovskite cell with ambition to improve its performance and stability. Thus, by varying the cations (A= Cs, FA, MA) and halogen (X= I, Br, Cl) in the perovskite APbX3, the authors noted not only an improvement in efficiency to 17%, but also a high stability of the cell, low presence of impurity, and low hysteresis when Cesium is used as a cation [33].

The other challenge in perovskite-based cell research is the substitution of Pb in perovskite with a non-toxic metal. The metals of group 14 of the Mendeleyev table are possible candidates, but their major problem is their chemical instability in the required oxidation state [34]. Considering the enthusiasm that researchers have for this technology, it is certain that in a few years to come, these barriers will be lifted.

Conclusion

There is no doubt about the performance that perovskite cell can achieve, which rivals crystalline silicon with 25.5% efficiency in 2020. The major future challenge facing research on perovskite before its commercialization is the stability of its current structure in outdoor conditions, as it reacts with oxygen in the air and water vapor.

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