The Role of Graphene in Perovskite Solar Cells

A. Agresti¹

S. Pescetelli¹, F. Gabelloni², A. Abate³, E. Burzi³, F. Biccari³, M. Gurioli³, E. Kymakis⁴, F. Bonaccorso⁵, A. Vinattieri³, A. Di Carlo¹.

1 C.H.O.S.E. (Centre for Hybrid and Organic Solar Energy), Department of Electronic Engineering, University of Rome Tor Vergata, Via del Politecnico 1, I-00133 Rome, Italy.

2 Dept. of Physics and Astronomy, University of Florence and LENS, Via Sansone 1, I–50019 Sesto Fiorentino (FI), Italy.

3 Adolphe Merkle Institute, Chemin des Verdiers, CH-1700 Fribourg, Switzerland.

4 Center of Materials Technology and Photonics & Electrical Engineering Department School of Applied Technology, Technological Educational Institute (T.E.I) of Crete Heraklion, 71 004 Crete, Greece.

5 Istituto Italiano di Tecnologia, Graphene Labs, Via Morego 30, I-16163 Genova, Italy.

antonio.agresti@uniroma2.it

The impressive power conversion efficiency (PCE=22.1% [1]) reported for perovskite solar cells (PSCs) has been reached thanks tremendous efforts made to the in growth process, improving the the morphology of the active film and the energy level matching between the cell's constituent layers. Despite that, record PCE is yet far from the theoretical predicted efficiency limit (approaching 31%). Notably, losses due to interfacial recombination negatively affect the charge injection at perovskite/transporting layer interface. Similarly, poor charge transport in electron (ETL) and hole (HTL) transporting layers severely limits charge collection at the electrodes. Moreover, the stability of PSCs still remains an open issue to be addressed. In fact, the poor air (H₂O and O_2 adsorption), photo, and thermal stability could dramatically restrict the future commercial exploitation of PSCs.

In this work, graphene and related twodimensional materials (GRMs) have been introduced in the device structure in order to improve the charge injection and/or collection at the electrodes and the device's long-term stability under real working conditions [2].

On one hand, the insertion of graphene flakes into the mesoporous-TiO₂ layer (mTiO₂) and of lithium-neutralized graphene oxide (GO-Li) as interlayer (see Fig.1) showed enhanced conversion efficiency and stability on both small and large area devices, by demonstrating the crucial role of graphene interface engineering (GIE) at mTiO₂/perovskite interface [3]. In particular, GIE is here demonstrated as an effective way to control the crystalline quality of the perovskite active material by means of continuous wave and picosecond timeresolved optical spectroscopy. More in detail, by exciting the graphene-based sample with a low-intensity radiation (2.06 eV) at T=11K, we point-out that the densities of traps at mTiO₂/perovskite interface are lower with respect to the substrate. The reference observed behaviour proves the high quality of the perovskite film when graphene-modified ETL is employed as substrate On the other hand, the application of graphene-based oxides (GBOs) at PSC counter-electrode is here demonstrated as an effective way to prevent the ions diffusion from perovskite into the hole transporting material. The ions blocking

capability of GBOs has been verified by intensity modulated photocurrent spectroscopy (IMPS). The stability improvement gained by the application of GIE in PSCs opens a promising route for further scale-up and stabilize the PSC technology, towards its commercialization.

References

- [1] http://www.nrel.gov/pv/assets/image s/efficiency_chart.jpg
- [2] A. Agresti et al., ChemSusChem, 9, (2016),2609
- [3] A. Agresti et al., ACS Energy Letters, 2, (2017) 279