

Optical properties. of PZN-PT nanoparticles thin layers for PV applications

R Ndioukane, D. Kobor, L. Motte, J. Solard

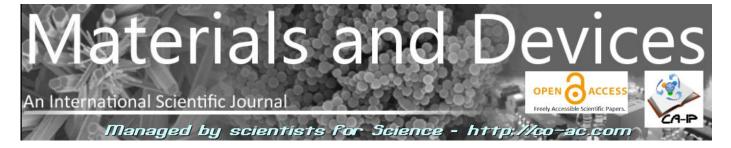
▶ To cite this version:

R Ndioukane, D. Kobor, L. Motte, J. Solard. Optical properties. of PZN-PT nanoparticles thin layers for PV applications. Materials and Devices, Collaborating Academics – International Press, 2019, 4 (1), 10.23647/ca.md20191502. hal-02151288

HAL Id: hal-02151288 https://hal.archives-ouvertes.fr/hal-02151288

Submitted on 7 Jun2019

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Article type: Conf.-Conference paper (CASAMANSUN)

Optical properties of PZN-PT nanoparticles thin layer on ITO glass for photovoltaic application

R. Ndioukane (1), D. Kobor (1), L. Motte (2), J. Solard (3)

(1) Laboratoire de Chimie et de Physique des Matériaux, Assane Seck University of Ziguinchor, Senegal

(2) Laboratory for Vascular Translational Science, Paris 13 University, France

(3) Centrale de proximité en nanotechnologies de Paris Nord, Institut Universitaire de Technologie de Villetaneuse, France

Corresponding author: r.ndioukane1532@zig.univ.sn

RECEIVED: 14 september 2018 / RECEIVED IN FINAL FORM: 12 february 2019 / ACCEPTED: 15 february 2019

Abstract: In this work, undoped and Mn doped Pb(Zn_{1/3}Nb_{2/3})O₃-4.5PbTiO₃ nanoparticles were dispersed in biopolymer and in mixed biopolymer + pentacene as active layer and deposited by spin coating on ITO glasses. Morphological, optical and electrical properties of these layers were investigated. SEM images show the superposition of different deposited layers ITO/TiO2/PZN-PT-np-biopolymer on glass substrate with thicknesses of 1.600 µm, 1.505 µm and 1.765 µm respectively. The absorbance value in UV visible for pentacene layer increases from 75 % to 99 % while the transmittance for ITO glasses diminishes from more than 80 % to 2 %. Optical gaps of ITO, TiO₂, PZN-PT nanoparticles are respectively 3.75 eV, 3.2 eV and 3.15 eV. Pentacene deposition reduced the gap to 1.65 eV for undoped sample and 1.60 eV for the doped ones. Intermediate gaps (2.3 eV and 2.6 eV for undoped sample and 2.15 eV and 2.7 eV for doped sample) were observed. Photoluminescence performed between 450 nm and 750 nm confirms « these intermediate gaps ».

Keywords: PEROVSKITE, NANOPARTICLES, THIN FILM, OPTICAL PROPERTIES, BAND GAP

Cite this article: Rémi Ndioukane, Diouma Kobor, Laurence Motte and Jeanne Solard, *OAJ Materials and Devices, Vol 4 (1), 1502 (2019) – DOI:* 10.23647/ca.md20191502

OAJ Materials and Devices, Vol 4 (1), 1502 (2019) – DOI: 10.23647/ca.md20191502

Introduction

Among all the solution-processed thin film optoelectronic material technologies investigated over the past 2 decades, organic-inorganic (or hybrid) methylammonium lead triiodide (MAPbI3) perovskites have emerged as clear frontrunners with proof-of-concept high performance devices demonstrated for a broad range of applications [1-8]. The MAPbI3-based photovoltaic devices have demonstrated constantly increasing power conversion efficiency (PCE), which now exceeds 22% [9-10] and is steadily approaching that of single-junction monocrystalline Silicon (c-Si) solar cells. Although the record for the highest efficiency perovskite solar cell was achieved using a mesoporous titania (TiO₂)-based architecture [10-12] photovoltaic devices employing a simple planar architecture are closing in with the highest reported efficiency of 20% [6, 13, 14]. Despite this breakthrough, hybrid lead-halide perovskites are known to degrade due to moisture and heat, upon prolonged exposure to light and are prone to ion or halide vacancy migration, leading to unstable operation of photovoltaic devices. Great efforts have been put forth for achieving highly efficient planar solar cells by exploring for potential ideal contact layers [15]. In addition, it is now realized that the next steps in advancing hybrid perovskitebased materials toward a viable photovoltaic technology will require simultaneously improving both the overall efficiency and also intrinsic stability. To overcome such difficulties, we oriented our research to inorganic PZN-PT perovskite materials with excellent and stable properties compared to organic-perovskite ones. However, despite their the excellent properties, one of the greatest difficulties to integrate widely such materials in electronic devices is to achieve them in thin films form because of their incongruent melting property. In this paper, undoped and Mn doped PZN-4.5PT nanoparticles were dispersed in biopolymer and in mixed biopolymer + pentacene as active layer and deposited by spin coating on ITO glasses. After the successful deposition of the thin film, morphological, optical and electrical properties were investigated.

Experimental procedure

The quality of the layers depends strongly on the state of the materials surfaces and their interfaces, whether organicorganic or metal-polymer. Thus a chemical treatment of the ITO substrate surface is necessary to eliminate the contaminating elements. For this, we combined the ultrasonic method with piranha to clean substrates. The piranha solution is composed of H_2SO_4 96 % (30 ml) and H_2O_2 (10 ml). After cleaning, the different layers, namely titanium oxide (TiO₂), undoped and Mn doped PZN-4.5PT nanoparticles dispersed in a biopolymer and pentacene were successively deposited by spin coating. In this study we used a natural biopolymer containing polysaccharides with short side chains. This polymer can take several conformations that allow it to adapt or intercalate depending on the environment.

Colloidal suspension of TiO₂

The titanium oxide used in this study is a finished product

marketed by Sigma-Aldrich (CAS number 13463-67-7). The suspension is obtained by grounding TiO2 powder in a mortar and adding a few drops of acetic acid giving a white paste. The as prepared paste is deposited on ITO glass substrate at room temperature using a spin coater Midas 1200D at 3500 rpm with an initial acceleration of 5 seconds and an operating time of 2 min.

PZN-PT nanoparticles thin film fabrication

Undoped and Mn doped PZN-4.5PT single crystals synthesized by the flux method [16] were grounded in a mortar to obtain a very fine powder (TEM characterizations revealed nanoparticles of the nanometer order and studies (not shown here) have confirmed that the population of spherical particles having a diameter of 30 nm was more representative). The grounded powder was dispersed in biopolymer solution. To obtain a homogeneous film, spin coating process was carried out at room temperature at 3500 rpm with an initial acceleration of 5 seconds and an operating time of 10 min. The biopolymer is used to disperse undoped and Mn doped PZN-4.5PT perovskite nanoparticles but also as a hole-injecting layer through the organic layer/ITO interface. Secondly it serves buffer layer to prevent diffusion of oxygen and indium to active layer, these impurities can act as exciton recombination centers or photogenerated traps carriers.

Results and discussion

SEM images

Surface morphology was observed by using an electron beam lithography system Pioneer Raith model in C(PN)2 (Paris 13 University). Observation of film surfaces in a scanning electron microscope makes it possible to verify the homogeneity of films, grains shapes and aggregates as well as the qualitative analysis of layers.

SEM images of samples composed of ITO/TiO2/PZN-PT-npbiopolymer films (DKRN-ITO2), ITO/TiO2/PZN-PT+1%Mn-npbiopolymer (DKRN-ITO4), ITO/TiO2/PZN-PT-npbiopolymer/pentacene (DKRN-ITO7), ITO/TiO2/PZN-PT+1%Mn-np-biopolymer/pentacene (DKRN-ITO8) deposited at room temperature on glass substrates are shown respectively in figure 1. DKRN-ITO2 and DKRN-ITO4 (Fig. 1a and 1b) shows a homogeneous, sparse and porous surface, showing white gray coloured non-regular agglomerated grains with an average size of 30 nm. DKRN-ITO7 and DKRN-ITO8 samples (Fig. 1c and 1d), developed in the same conditions with the only difference of adding a pentacene layer, have uniform, compact and homogeneous slightly black surfaces. This suggests that pentacene leads to a coalescence of the grains by diffusion through the pores. The morphology of the surface is initially related to the energy of the particles. Indeed, when the particle on the substrate acquires a great energy, it will have a large surface diffusion length. Then, this allows it to reach the sites energetically favorable and thus to

OAJ Materials and Devices, Vol 4 (1), 1502 (2019) - DOI: 10.23647/ca.md20191502

complete the possible gaps, reducing the irregularities of the surface. In the other case, the particles deposited on the substrate in contact with the air having lost a large part of the energy, their diffusion length will be very short. Therefore, the morphology of the surface will reflect a part of the morphology of the substrate but with greater irregularities due to the shading effect, and secondly the nature of the deposited material [17].

Observation in scanning electron microscope in transverse mode, allowed us to see the disposition of the various deposited layers but also to determine their thickness. For example, in figure 2a showing the SEM image in transverse mode of the DKRN-ITO2 sample composed of the ITO / TiO2 / np-PZN-4.5PT-biopolymer layers, we obtained thicknesses of 1.600 μ m, 1.505 μ m and 1.765 μ m respectively for ITO, TiO₂ and np-PZN-PT-biopolymer. This allowed us to confirm the value of the thickness of the ITO given by the supplier (1.600 μ m).

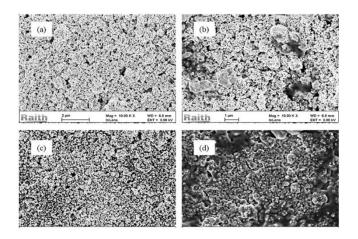


Figure 1: SEM Images of thin film (a) DKRN-ITO2, (b) DKRN-ITO4, (c) DKRN-ITO7 and (d) DKRN-ITO8

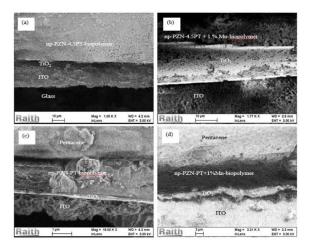


Figure 2: SEM Images in transversal mode (a) DKRN-ITO2, (b) DKRN-ITO4, (c) DKRN-ITO7 and (d) DKRN-ITO8

Thicknesse (µm)	ITO	TiO ₂	Biopolymer	Pentacene
DKRN-ITO2	1,600	1,505	1,765	-
DKRN-ITO4	1,600	2,401	1,667	-
DKRN-ITO7	1,600	1,709	1,680	1,431
DKRN-ITO8	1,600	1,691	1,582	1,322

Table 1: Thickness values of different layers

Optical properties

Optical properties of thin films were investigated using an UV-Visible spectrophotometer with an integrating sphere model LAMBDA 950S in IM2NP (Marseille). Measurements are made with a pitch of 5 on a wavelength spectrum ranging from 250 and 1500 nm.

Transmittance and absorption

Typical transmission spectra obtained for the different deposited layers at room temperature are shown in figure 3a. For the ITO reference sample composed of ITO layer only, there is a strong transmission of more than 80 % throughout the visible band. This confirms the transmission value given by the supplier. This transmission decreases with the deposition respectively of TiO2/np-PZN-4.5PT-biopolymere, TiO2/np-PZN-4.5PT+1%Mn-biopolymer, TiO2/np-PZN-4.5PT-biopolymere/pentacene and TiO2/np-PZN-4.5PT+1 %Mn-biopolymer/pentacene.

For the different deposits, the general patterns of the spectra are similar. Two transmission domains can be distinguished according to the wavelength:

- A domain (λ < 800 nm) characterized by high absorption and low transmission of the different layers, which corresponds to the absorption of the layer in question.

- A relatively high transmission domain for λ range of 800 - 1500 nm, it increases sharply and tends to values between 30 and 56 % for the DKRN-ITO2 sample, 26 and 47 % for the DKRN-ITO4 sample, 25 and 45 % for the DKRN-ITO7 sample, 45 and 70 % for the DKRN-ITO8 sample. Authors in the literature, such as D. Kobor [18] report transmission values of 25 and 45 % for wavelengths between 600 and 900 nm of PZN-4.5PT single crystals. This confirms the values obtained in this study, although we can also consider the influence of the organic layers of biopolymer and pentacene which tend to attenuate this transparency. In figure 3b, it can be seen that the addition of the pentacene layer greatly increases the absorption throughout the visible spectrum. It is deduced that

OAJ Materials and Devices, Vol 4 (1), 1502 (2019) - DOI: 10.23647/ca.md20191502

pentacene increases the absorption band in the visible (~ 90 %). Indeed, pentacene is a semiconductor that absorbs well in the visible and authors like A. Skaiky [19] have shown it. This is a good thing for photovoltaic applications. On the other hand, the biopolymer seems to have less influence on the absorption compared to pentacene.

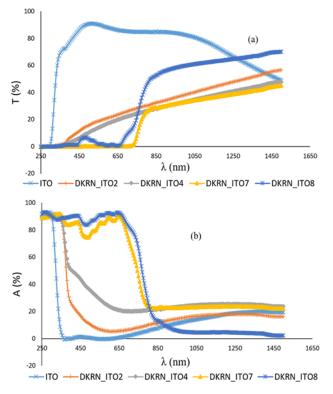


Figure 3: (a) transmittance and (b) absorbance according to the UV-Visible-NIR wavelength for different samples

Determination of band gap

To calculate the optical gap Eg, we used Tauc relation [20].

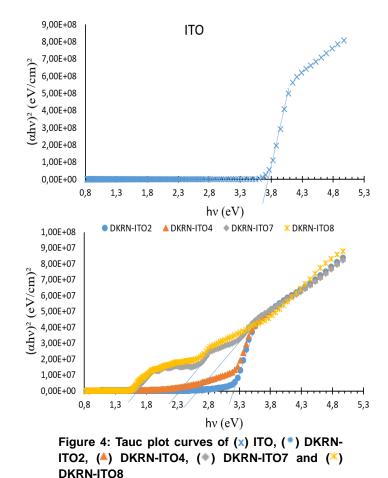
$$\alpha h \nu = A (h \nu - E_g)^n \tag{1}$$

Where A is a constant, hv is photon energy, E_g is the allowed energy gap, $n = \frac{1}{2}$ for allowed direct transition, α is the absorption coefficient and n = 2 for allowed indirect transition.

Thus, from the value of the absorption coefficient and by plotting the curve of $(ahv)^2$ as a function of the photons energy, we have determined the optical gap Eg: it suffices to draw the slope in the region of absorption as shown in figure 4 and the intersection of this line with the abscissa axis (for ahv = 0) gives directly Eg. The values found are shown in table 2.

For the ITO reference sample, the value of the gap found (3.75 eV) is in perfect correlation with that found by G. Fatma-Zohra [21] (3.80 eV). For the DKRN-ITO2 and DKRN-

ITO4 samples, the gap remains high with values of 3.20 eV and 3.15 eV that can respectively correspond to the TiO₂ gap that is 3.02 eV according to Schon [22] and PZN-PT gap equal to 3.04 eV according to He Chong-Jun [23]. Pentacene deposition reduced the gap to 1.65 eV for the DKRN-ITO7 sample and 1.60 eV for DKRN-ITO8. We also note the presence of intermediate gaps (2.30 eV and 2.60 eV for DKRN-ITO7 and 2.15 eV and 2.70 eV for DKRN-ITO8). These intermediate gaps could be due to the heterogeneity of the layers (presence of TiO₂, nanoparticles of PZN-4.5PT undoped or doped Mn, biopolymer, pentacene). These values could correspond respectively to the gap of the pentacene layer (1.65 and 1.60 eV), the combination of the biopolymer with pentacene (2.30 and 2.15 eV) and TiO₂ (2.60 and 2.70 eV). It appears from the results that pentacene tends to influence even the gap of the inner layers (TiO2, nanoparticles and biopolymer) whose absorption is considerably reduced. The results are in perfect correlation with the curves of figure 3.



OAJ Materials and Devices, Vol 4 (1), 1502 (2019) – DOI: 10.23647/ca.md20191502

Samples	E _{g1} (eV)	E _{g2} (eV)	E _{g3} (eV)
ITO	3,75	-	-
DKRN-ITO2	3,20	-	-
DKRN-ITO4	3,15	-	-
DKRN-ITO7	1,65	2,30	2,60
DKRN-ITO8	1,60	2,15	2,70

Table 2: Gap values of different samples

Photoluminescence

To confirm the effect of pentacene on the reduction of the optical gap, we investigate photoluminescence on samples DKRN-ITO7 and DKRN-ITO8. Figure 5 shows the photoluminescence (PL) spectra obtained. We observe a first emission peak at 2.3 eV and a second peak at 2.5 eV for DKRN-ITO7 while for DKRN-ITO8 we have a peak at 2.2 eV and another at 2.6 eV. This confirms the values of the intermediate gaps found by the Tauc method. Photoluminescence was performed between 450 nm and 750 nm. This does not allow observing the peak of 1.6 eV revealed by the method of Tauc. Indeed, this peak would probably appear in the infrared.

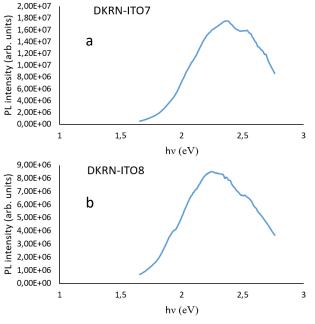


Figure 5: Photoluminescence of (a) DKRN-ITO7 and (b) DKRN-ITO8

Conclusion

PZN-4.5PT nanoparticles thin layers were successfully prepared using a biopolymer. SEM images revealed the superposition of different deposited layers. It shows uniform, compact and homogeneous surface with deposition of pentaconene layer. Optical characterization by transmittance and absorbance measurements allowed us to observe the influence of the layers on absorption and optical gap. It also made it possible to follow the evolution of the gap according to the nature of the deposited layer. Pentacene has been very successful in improving photovoltaic properties.

Acknowledgements:

This work is supported by Agence Universitaire de la Francophonie (AUF).

Complementary informations on authors:

Rémi Ndioukane: r.ndioukane1532@zig.univ.sn, Research Gate (Rémi Ndioukane), LinkedIn (REMI NDIOUKANE)

Diouma Kobor: <u>dkobor@univ-zig.sn</u>, Research Gate (Diouma Kobor)

Laurence Motte: lmotte2009@hotmail.fr, Research Gate (L. Motte)

Jeanne Solard: jeanne.solard@iutv.univ-paris13.fr, Research Gate (Jeanne Solard)

REFERENCES

1. M. M. Lee, J. Teuscher, T. Miyasaka, T. N. Murakami, H. J. Snaith, Science, vol.338, p 643-647(2012)

OAJ Materials and Devices, Vol 4 (1), 1502 (2019) - DOI: 10.23647/ca.md20191502

- J. Burschka, N. Pellet, S. J. Moon, R. Humphry-Baker, P. Gao, M. K. Nazeeruddin and M. Grätzel, Nature, vol.499, p 316-320 (2013)
- C. C. Stoumpos, C. D. Malliakas, J. A. Peters, Z. Liu, M. Sebastian, J. Im, M. G. Kanatzidis, Crystal Growth and Design, vol.13, p 2722-2727(2013).
- 4. N. J. Jeon, J. H. Noh, Y. C. Kim, W. S. Yang, S. Ryu, & S. I. Seok, Nature Materials, Vol.13 p 897–903 (2014).
- Z-K. Tan Z, R. S. Moghaddam, M. L. Lai, P. Docampo, R. Higler, F. Deschler, R. H. Friend, Nature Nanotechnology, Vol.9, p 687-692 (2014)
- 6. H. Zhou, Q. Chen, G. Li, S. Luo, T-b. Son, H. S. Duan, Y. Yang, Science, vol.345, p542–546 (2014)
- 7. Y. Fang, Q. Dong, Y. Shao, Y. Yuan, & J. Huang, Nature Photonics, vol.9, p 679–686 (2015)
- 8. H. Zhu, Y. Fu, F. Meng, X. Wu, Z. Gong, Q. Ding, X-Y. Zhu, Nature Materials, vol.14, p 636-642 (2015)
- 9. M. A. Green, K. Emery, Y. Hishikawa, W. Warta, & E. D. Dunlop, Progress in Photovoltaics. Research and Applications vol.24, p 905-913 (2016)
- M. Saliba, T. Matsui, J-Y. Seo, K. Domanski, J. P. Correa-Baena, M. K. Nazeeruddin, M. Grätzel, Energy & Environmental Science, vol.9, p 1989-1997 (2016)
- 11. W. S. Yang, J. H. Noh, N. J. Jeon, Y. C. Kim, S. Ryu, J. Seo & S. I. Seok, Science, vol.348, p 1234-1237 (2015)
- 12. D. Y. Son D, J. W. Lee, Y. J. Choi, I. H. Jang, S. Lee, P. J. Yoo, N. G. Park, Nature Energy, vol.1, p 16081 (2016)
- 13. C. Bi, Q. Wang, Y. Shao, Y. Yuan, Z. Xiao, & J. Huang, Nature Communications, vol.6 (2015)
- 14. M. Yang, T. Zhang, P. Schulz, Z. Li, G. Li, D. H. Kim, Y. Zhao, Nature Communications, vol.7 p 12305 (2016)
- 15. W. Nie, G. Gupta, B. K. Crone, F. Liu, D. L. Smith, P. P. Ruden, A. D. Mohite, Advanced Science, vol.2, p 1500024 (2015)
- 16. F. Dupret and N. Van Den Bogaert, Growth Mechanisms and Dynamics, Part B. In: Hurle D.T.J, Ed., Handbook of Crystal Growth 2, North-Holland, New York (1994)
- 17. S. J. Chen, Y. C. Liu, J. G. Ma, D. X. Zhao, Z. Z. Zhi, Y. M. Lu, X. W. Fan, Journal of Crystal Growth, vol.240 p 467–472 (2002)
- 18. D. Kobor, Synthesis and Characterization of PZN-4.5PT single crystals by Flux method, Thesis, INSA-Lyon (2005)
- A. Skaiky (2013), Elaboration, characterization and modeling of pentacene-based transistors: Application to organic electronic circuits, Thesis, University of Limoges (2013)
- 20. J. Tauc, The Optical Properties of Solids, Academic Press, New York (1966)
- 21. G. Fatma-Zohra, Elaboration and characterization of thin films of ZnO: Al and ZnO: Al / TiO2, Thesis, University of M'hamed Bougara-Boumerdes (2012)
- 22. J. H. Schon and B. Batlogg, Journal of Applied Physics, vol.89, p 336-342 (2001)
- 23. C. J. He, X. D. Fu, F. Xu, J. M. Wang, K. J. Zhu, C. L. Du & Y. W. Liu, Chinese Physics B vol.21, p 054207 (2012)

Important: Articles are published under the responsability of authors, in particular concerning the respect of copyrights. Readers are aware that the contents of published articles may involve hazardous experiments if reproduced; the reproduction of experimental procedures described in articles is under the responsability of readers and their own analysis of potential danger.

Reprint freely distributable – Open access article

Materials and Devices is an Open Access journal which publishes original, and **peer-reviewed** papers accessible only via internet, freely for all. Your published article can be freely downloaded, and self archiving of your paper is allowed and encouraged!

We apply « **the principles of transparency and best practice in scholarly publishing** » as defined by the Committee on Publication Ethics (COPE), the Directory of Open Access Journals (DOAJ), and the Open Access Scholarly Publishers Organization (OASPA). The journal has thus been worked out in such a way as complying with the requirements issued by OASPA and DOAJ in order to apply to these organizations soon.

Copyright on any article in Materials and Devices is retained by the author(s) under the Creative Commons (Attribution-NonCommercial-NoDerivatives 4.0 International (CC BY-NC-ND 4.0)), which is favourable to authors.



Aims and Scope of the journal : the topics covered by the journal are wide, Materials and Devices aims at publishing papers on all aspects related to materials (including experimental techniques and methods), and devices in a wide sense provided they integrate specific materials. Works in relation with sustainable development are welcome. The journal publishes several types of papers : A: regular papers, L : short papers, R : review papers, T : technical papers, Ur : Unexpected and « negative » results, Conf: conference papers.

(see details in the site of the journal: http://materialsanddevices.co-ac.com)

We want to maintain Materials and Devices Open Access and free of charge thanks to volunteerism, the journal is managed by scientists for science! You are welcome if you desire to join the team!

Advertising in our pages helps us! Companies selling scientific equipments and technologies are particularly relevant for ads in several places to inform about their products (in article pages as below, journal site, published volumes pages, ...). Corporate sponsorship is also welcome!

Feel free to contact us! contact@co-ac.com

OAJ Materials and Devices, Vol 4 (1), 1502 (2019) – DOI: 10.23647/ca.md20191502