Emphasizing the Operational Role of a Novel Graphene-based Ink into High Performance Inverted Ternary Organic Solar Cell



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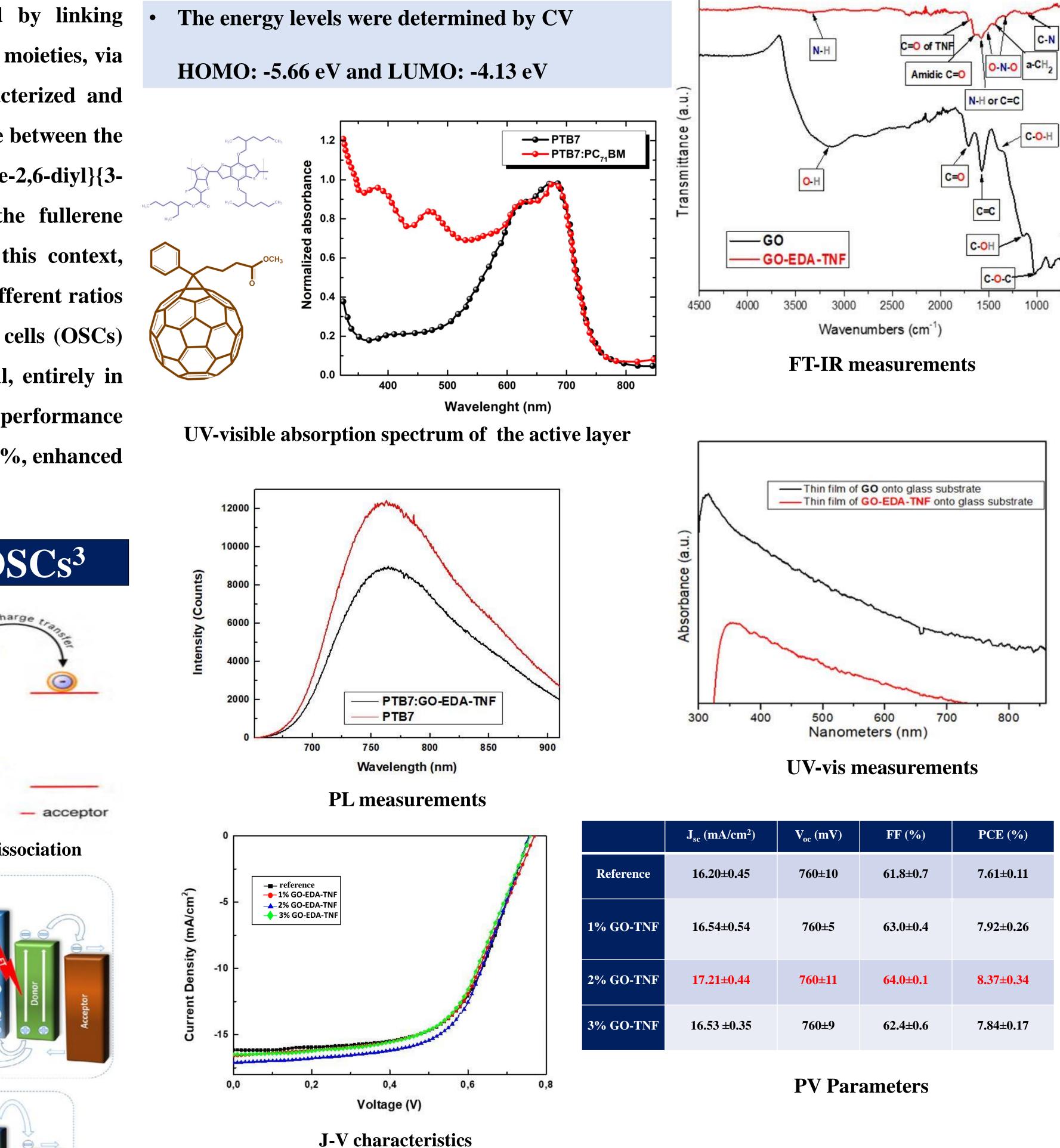
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Abstract

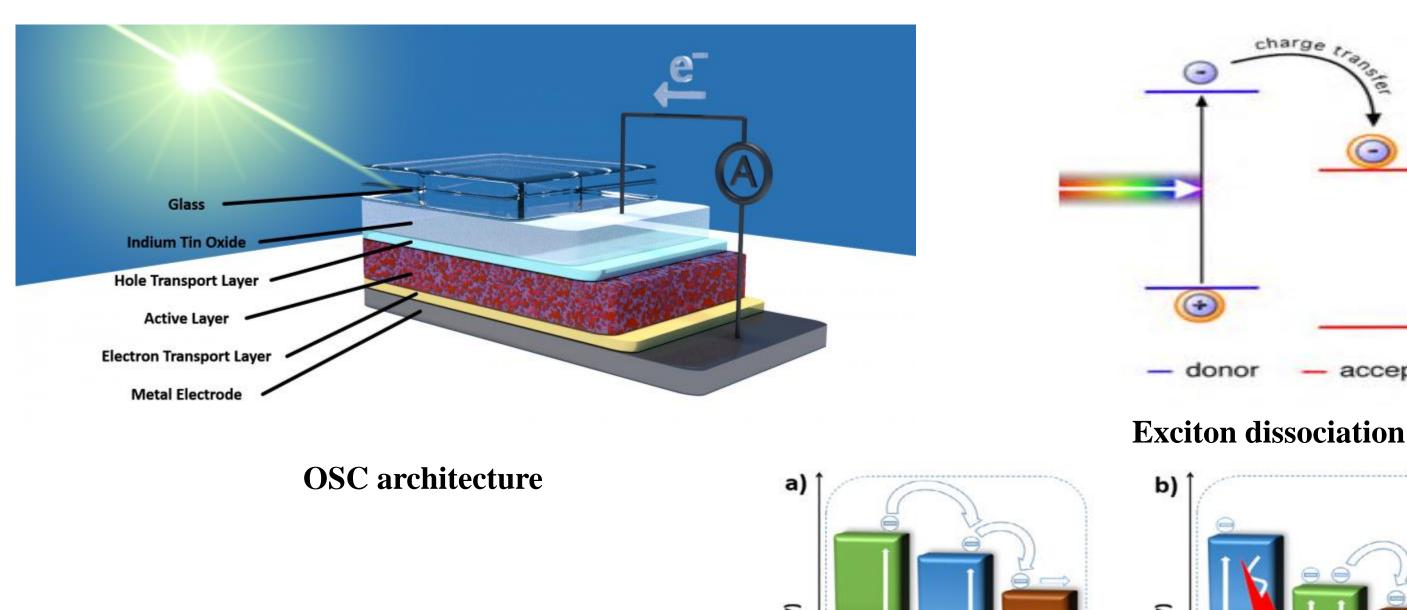
A novel solution-processed graphene-based nanomaterial was synthesized by linking graphene oxide (GO) with 2,5,7-trinitro-9-fluorenone-4-carboxylic acid (TNF) moieties, via simple synthetic procedures.^{1,2} The yielded GO-TNF was thoroughly characterized and proved that it presents favorable energy levels to function as a bridge structure between the polymer donor poly({4,8-bis[(2-ethylhexyl)oxy]benzo[1,2b:4,5b']dithiophene-2,6-diyl}{3fluoro-2-[(2ethylhexyl)carbonyl] thieno [3,4b]thiophenediyl}) (PTB7) and the fullerene

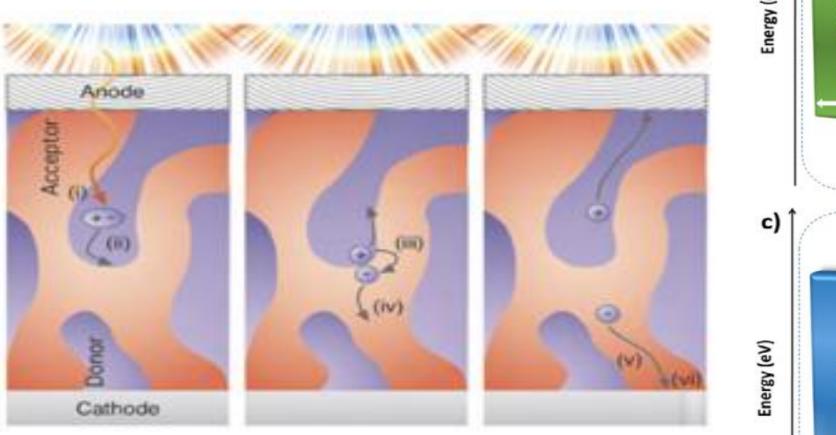
Characterization



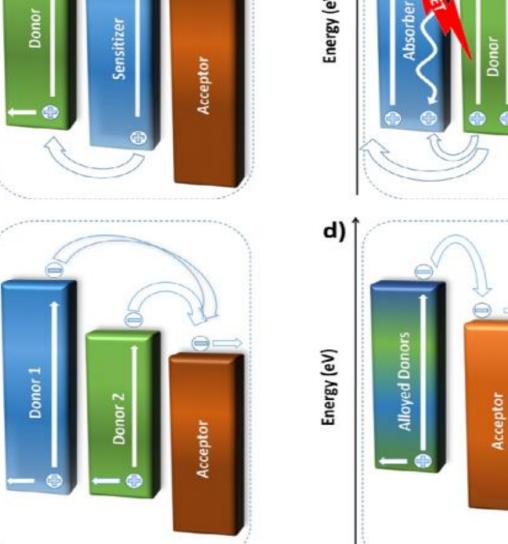
derivative acceptor [6,6]-phenyl-C₇₁-butyric-acid-methylester (PC₇₁BM). In this context, **GO-EDA-TNF** ink was incorporated within the binary photoactive layer, in different ratios (1-3% ratio to the polymer), for the effective inverted ternary organic solar cells (OSCs) fabrication of the structure ITO/PFN/PTB7:GO-EDA-TNF:PC₇₁BM/MoO₃/Al, entirely in inert atmosphere. The addition of 2% GO-TNF ink led to an overall device performance improvement, achieving a champion power conversion efficiency (PCE) of 8.71%, enhanced by ~10%, compared to the reference cell.

Basic Operational Principles of Ternary OSCs³



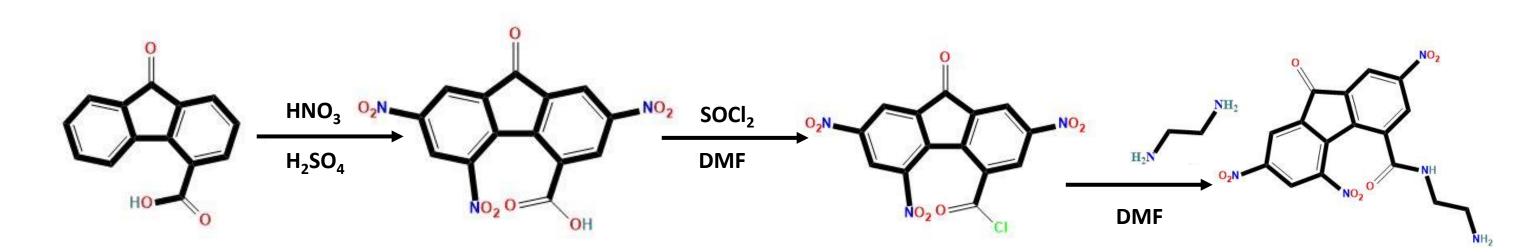


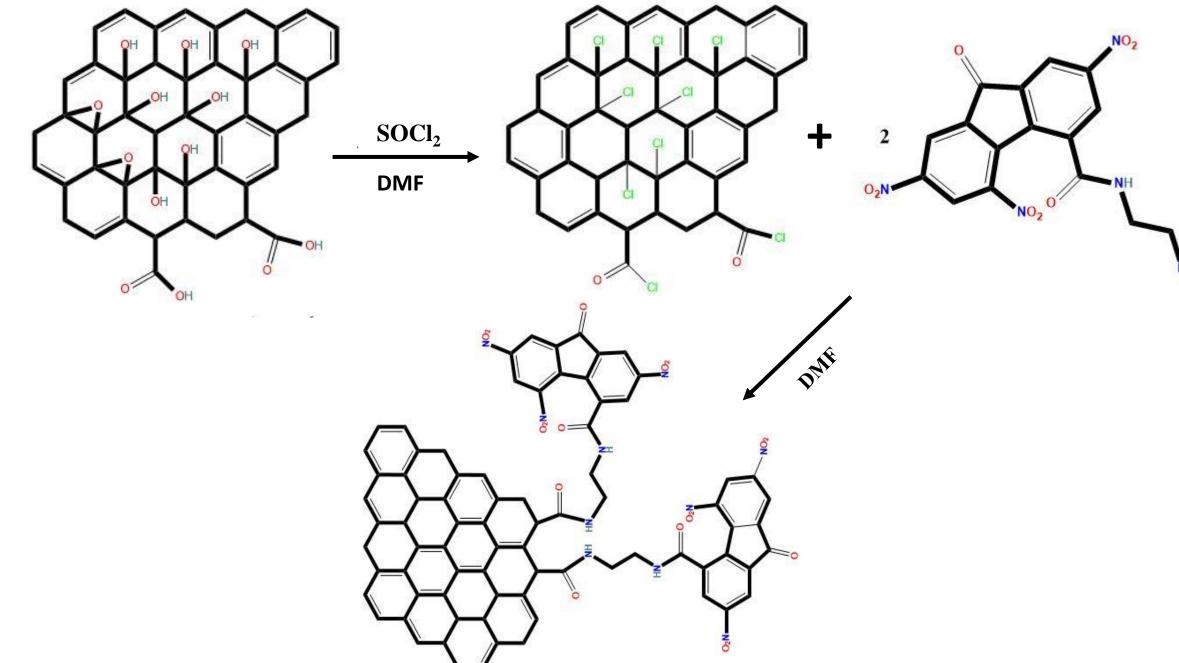
Exciton dissociation mechanism in the active layer

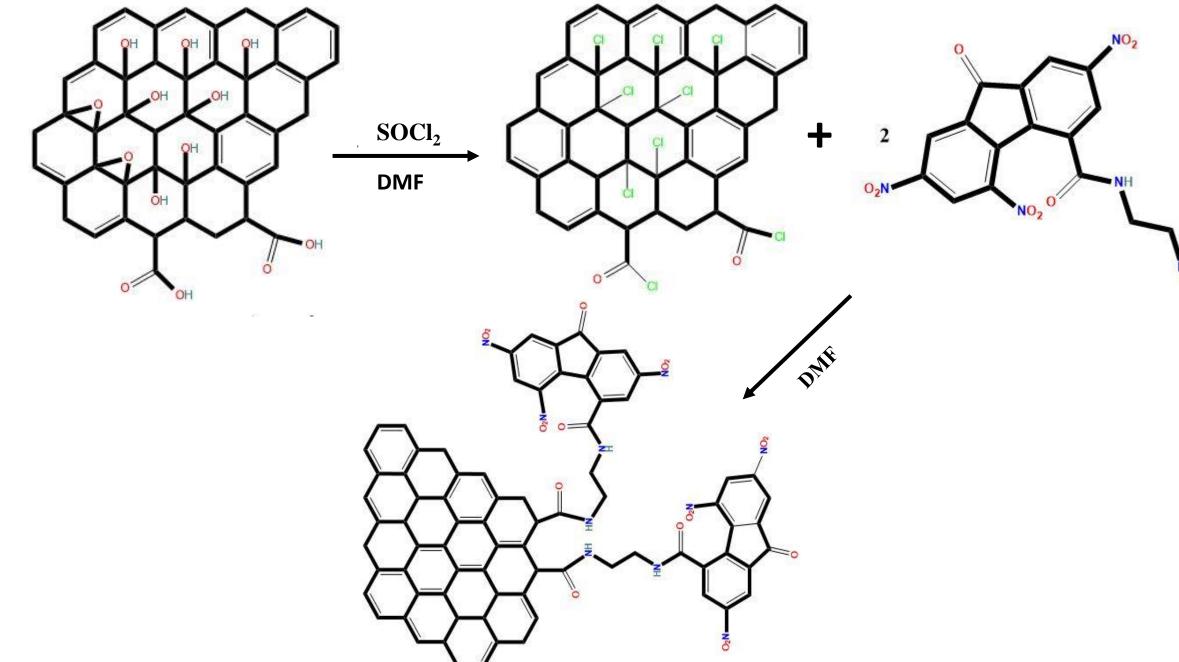


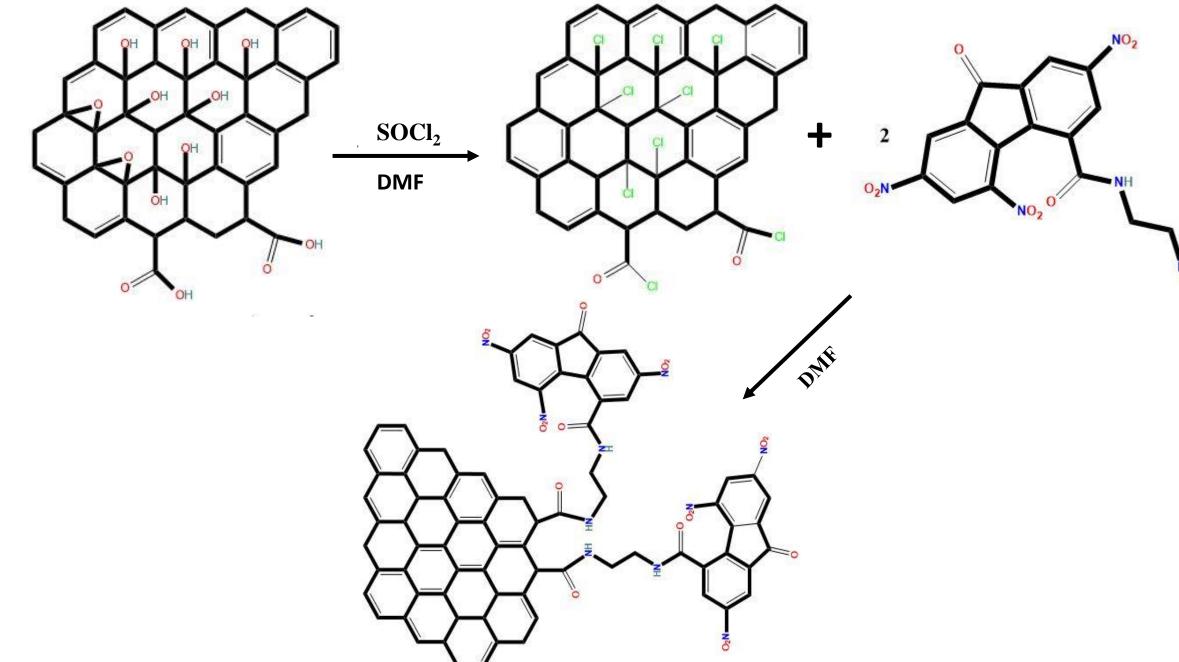
Operational mechanisms of the third component in the active layer of ternary OSCs

GO-TNF synthetic procedure









Conclusions – Future work

In this work, we have successfully prepared a novel graphene-based ink for the incorporation in high performance organic solar cells. The linkage of TNF moieties with graphene oxide (GO) was conducted via the creation of amidic bonds and improved the properties of GO, by reducing its bandgap. The PV performance was very promising, and we intend that it will be further improved in the near future.

References

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We thank Dr. G. Kenanakis and Prof. A. G. Coutsolelos' group for ATR FT-IR and CV measurements, respectively. This research is co-financed by Greece and the European Union (European Social Fund- ESF) through the **Operational Programme «Human Resources Development, Education and Lifelong Learning» in the context of the** project "Reinforcement of Postdoctoral Researchers" (MIS-5001552), implemented by the State Scholarships Foundation (IKY), Grant No. 13992.



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