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In-situ grown organic nanofibers have been prepared on metal electrodes patterned by electron beam lithography (EBL). A systematic investigation shows that the light emission from these nanofibers driven by an AC gate voltage depends non-linearly on the amplitude of the AC gate voltage and linearly on the frequency of the gate voltage, which indicates that a model involving space charge field (SCF) assisted electron tunneling can be applied. The photoluminescence spectra of para-hexaphenylene (p6P) and α -sexithiophene (6T) nanofibers illustrate that the emission color of the in-situ grown nanofibers can be tuned by depositing two types of discontinuous organic layers on the same platform.











The light emission intensity from a nanofiber based device depends nonlinearly on the AC voltage amplitude and linearly on the AC gate voltage frequency, which is similar to the dependence obtained from a thin film based device ^[1]. This indicates that a model based on SCF assisted electron tunneling can be applied ^[2].



The light emission obtained by grounding both metal electrodes and applying an AC voltage to the gate indicates that there is no charge carrier transporting between two metal electrodes; rather, the light emission is due to the subsequent injection of holes and electrons from the same metal electrode requiring a sufficiently high field strength to be provided at the electrode edge by the gate voltage.

[1]: X. Liu et al. "AC-biased organic light-emitting field-effect transistors from naphthyl end-capped oligothiophenes" Org. Electron. Vol 11, p. 1096-1102 (2010)

[2]: X. Liu et al. "Charge-carrier injection assisted by space-charge field in AC-driven organic light –emitting transistors" Org. Electron. Vol 12, p. 1724-1730 (2011)

- (a) During the negative half-period of the gate voltage, holes inject from the Au electrode due to the electric field and form a positive space charge field;
- During the positive half-period of the gate voltage, electrons tunnel out assisted (b) by the positive space charge field, recombining with the holes injected in the previous step.

Bi-layer nanofiber device

The fluorescence microscope images show a light-emitting device consisting of 3.5 nm p6P in-situ grown nanofibers (a), and the same device after being deposited a second layer with 7 nm 6T nanofibers (b). The inset SEM images illustrate that the 6T nanofibers can be well distinguished from the previously deposted p6P nanofibers.



The photoluminescence spectra suggest a reliable 0.8 method to mix clean colors



--p6P

 $-\star - p6P+6T$

Summary

In-situ grown p6P/6T nanofibers on nanoscale electrodes have been used to investigate AC-driven light emission. The emission color of the nanofibers can be tuned by depositing two types of discontinuous organic layers on the same platform.

Future work

Optimize the nanofiber in-situ growth on nanoscale electrodes in order to achieve a higher growth density. Investigate the influence of the thickness of the two materials on the resulting spectrum.