

# Charge Transport in van der Waals Thin Films, the case of Reduced Graphene Oxide from single flake to thin film

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Large area van der Waals (vdW) thin films are assembled materials consisting of a network of randomly stacked nanosheets. The multiscale structure and the two dimensional (2D) nature of the building block mean that interfaces naturally play a crucial role in the charge transport of such thin films. While single or few stacked nanosheets (i.e., vdW heterostructures) have been the subject of intensive works, [1-3] little is known about how charges travel through multilayered, more disordered networks. [4] Here, we report a comprehensive study of a prototypical system given by networks of randomly stacked reduced graphene oxide 2D nanosheets, whose chemical and geometrical properties can be controlled independently [5], permitting to explore networks ranging from a single nanosheet to some billions forming thin films, with room temperature resistivity spanning from  $10^{-5}$  to  $10^{-1} \Omega \cdot m$ .

Most studies give an ambiguous interpretation of charge transport phenomena by qualitative plot of resistance vs temperature, while a univocal interpretation of the charge transport phenomenon is still object of debate: Efros-Shklovskii variable-range hopping (ES-VRH) or 2D Mott variable-range hopping (2D-VRH). Based on analysis of the reduced activation energy  $W(T)$  [6], we systematically observed a clear transition between two different regimes: Efros-Shklovskii variable-range hopping (ES-VRH) below a critical temperature  $T^*$  and power law behavior above.

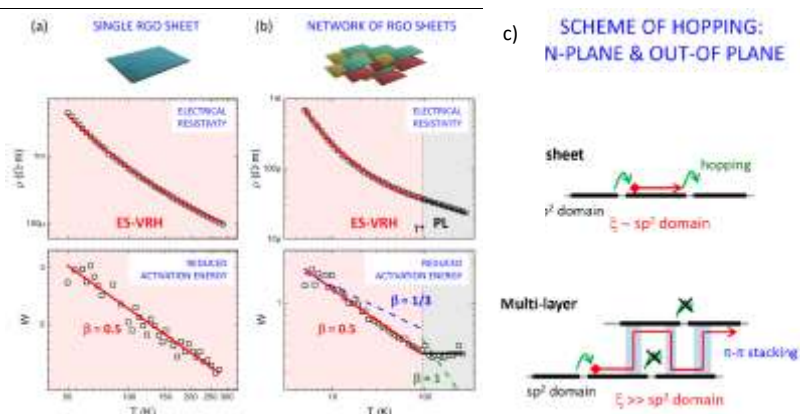
First, we demonstrate that the two regimes are strongly correlated with each other, both depending on the charge localization length  $\xi$ , calculated by the ES-VRH model, which corresponds to the characteristic size of overlapping  $sp^2$  domains belonging to different nanosheets. Thus, we propose a microscopic model describing the probability that charges circumvent the hopping barriers increases with film thickness, with a corresponding increase in the effective delocalization of the electronic states up to the micron scale [7].

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## References

- [1] Joung et al., Physical Review B, 86(23), 235423 (2012)
- [2] Gómez-Navarro et al., Nano Letters, 2007. 7(11): p. 3499-3503.
- [3] Mattevi et al., Advanced Functional Materials, 2009. 19(16): p. 2577-2583.
- [4] Kim et al., The Journal of Physical Chemistry C, 2015. 119(51): p. 28685-28690.
- [5] Kovtun A., et al., Carbon 143 268-275 (2019)
- [6] Zabrodskii et al., Zh. Eksp. Teor. Fiz, 1984. 742: p. 425-33.
- [7] Kovtun, A., et al., ACS Nano 2021, 15, 2654–2667

## Figures



**Figure 1:** Resistivity vs Temperature of single sheet rGO (a) and thin film (b). Charge transport scheme(c).