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Advanced Characterization of Pillared Graphene-Based Materials for Supercapacitors

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Materials for supercapacitors have to display large surface area to maximize the density of adsorbed ions [1], but they must also possess a hierarchical porosity (Fig. 1) to favour ions diffusion. Graphenederived materials are studied as they –theoretically- combine all these porosity-related parameters. However experimentally the achieved capacitance remains limited because graphene sheets tend to re-aggregate. Our group has selected a method to avoid this issue by creating an expanded graphene structure by bridging and separating the sheets using aliphatic diamine pillars. These assemblies are called Pillared Graphene Materials (PGM). Electrochemical results showed that these PGM display higher capacitance than reference non pillared materials [2,3]. However, it was not possible to correlate these storage performances improvements to adsorption active surface area determined with N₂ gas sorption (BET), which questions the suitability of this technique for measuring the full porosity of these samples. Therefore, to further correlate storage performances to structure properties, we started using small angle X-ray/neutron scattering (SAXS/SANS) techniques which are perfectly suited to characterize the sample structure, and reveal mesoscale porosities (~20 to 140 nm).

In the frame of this presentation, we will show how SAXS/SANS have been applied to pristine materials but also to specifically formulated samples (pellet, electrodes) in order to highlight structural differences arising from synthesis/formulation protocols or from molecular pillaring. Reference samples morphologies were also shown to be very dependent of the synthesis method.

References

- [1] M. Salanne, et al, Nat. Energy 2016.
- [2] Banda, et al, J. Power Sources 2017.
- [3] Banda et al., ACS Nano, 13 (2019).

Figures

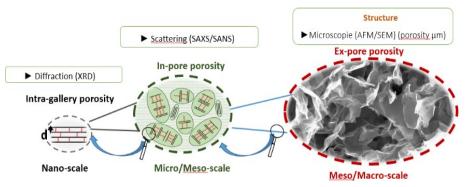


Fig. 1 Conceptual scheme of the different level of porosity in pillared graphene materials