

From molecules in solution to molecules on surfaces – using supramolecular dyads to form functional self-assembled networks on graphene

Quentin Fernez, Shiva Moradmand, Michele Mattera, William Djampa-Tapi, Céline Fiorini-Debuisschert, Fabrice Charra, David Kreher, Fabrice Mathevet, Imad Arfaoui, Lydia Sosa Vargas

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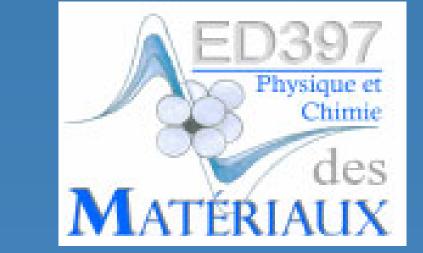
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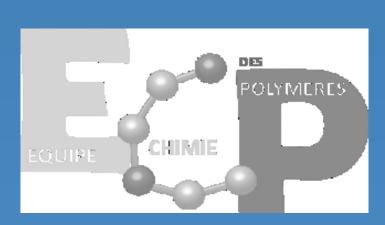




From molecules in solution to molecules on surfaces – using supramolecular dyads to form functional self-assembled networks on graphene



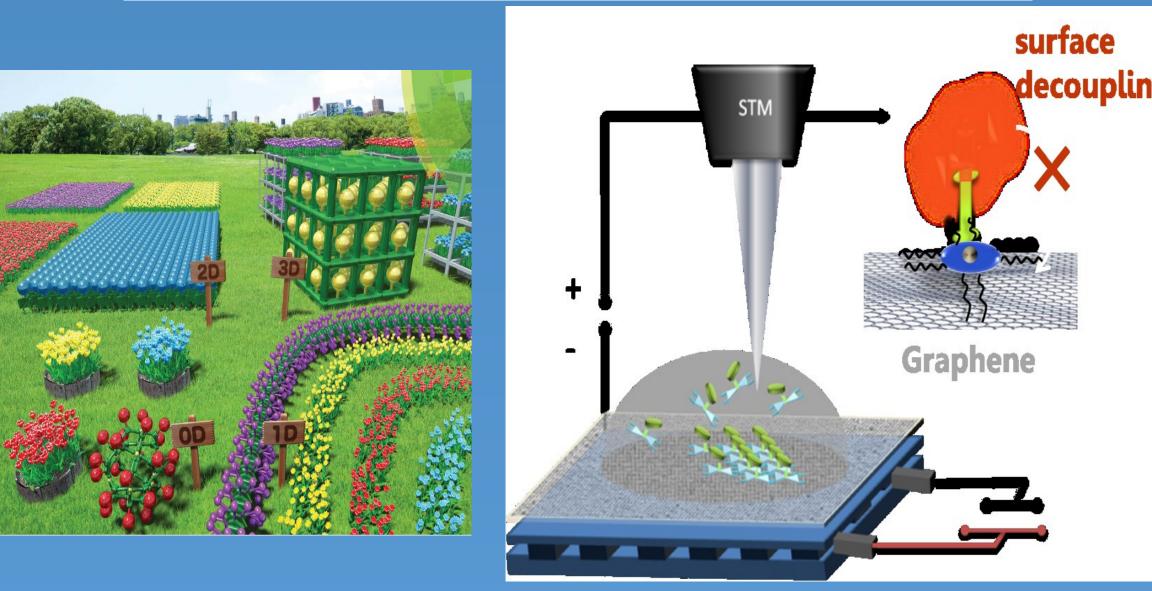
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 $Ar-NH_2 =$

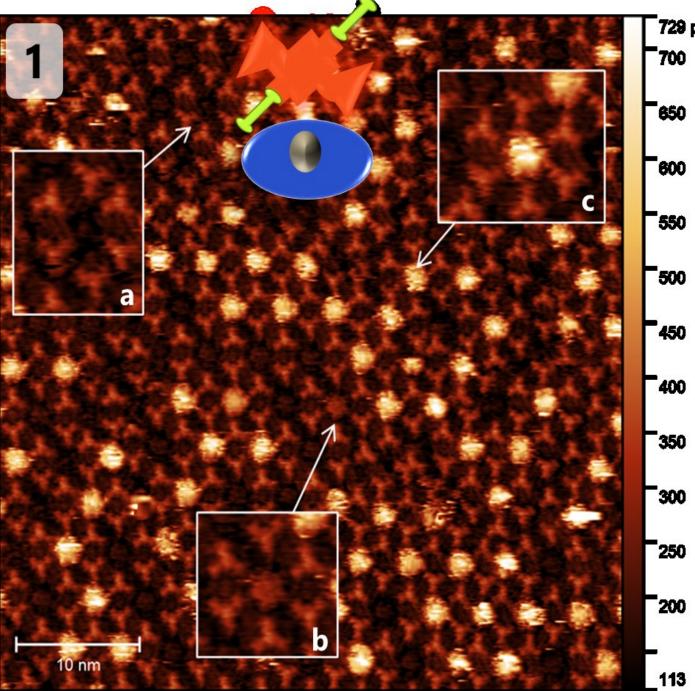
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INTRODUCTION and GOALS

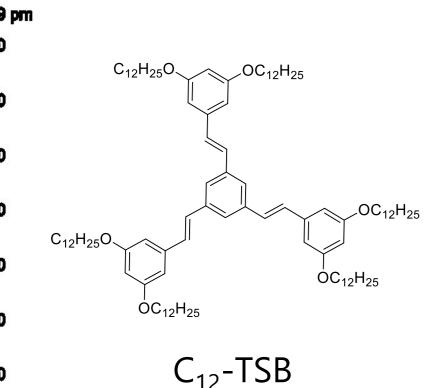


SELF-ASSEMBLY PROPERTIES

STM images are monolayers of several molecular pedestals on Highly Oriented Pyrolytic Graphite (HOPG) surfaces.



STM image: Bias: -1,0 V, Current: 25 pA, Solid-liquid interface in phenyloctane



- **Host-guest network**
- **Orientation** of PDI with respect to ZnPc and HOPG surface is **unknown**
- After deposition of ZnPc-PDI (1-1) in C_{12} -TSB host network: 3 different signals can be identified: the empty TSB pore (a), The TSB pore + ZnPc (b) and TSB pore hosting a ZnPc-PDI (c)

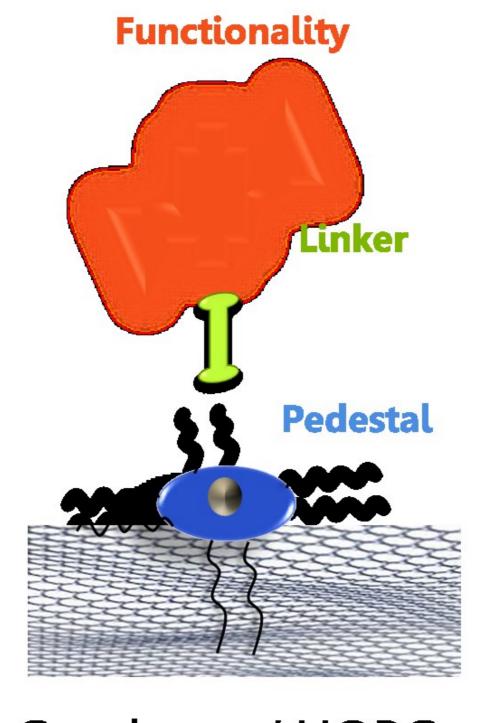
To develop new functional materials, nanoscopic scale architecture control is necessary¹. As a bottom-up synthesis strategy, molecular self-assemblies on surface have been explored, and have demonstrated good control of patterning on various surfaces². From these surfaces; graphene is of interest due to its versatility and potential in future semiconductive devices³. The work presented here describes the functionalization of a graphenoid surface via a supramolecular approach.

We aim to synthesis of versatile and functionalized building-blocks which self-organize on graphenoid surfaces. We are mainly interested in :

- Tuning the surface self-assembly via the functionalisation of the molecular building blocks
- Grafting a fluorescent functionality decoupled from the surface
- Studying by Scanning Tunnelling Microscopy (STM) the topology of the molecular 2D-assemblies

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DESIGN STRATEGY



Graphene / HOPG

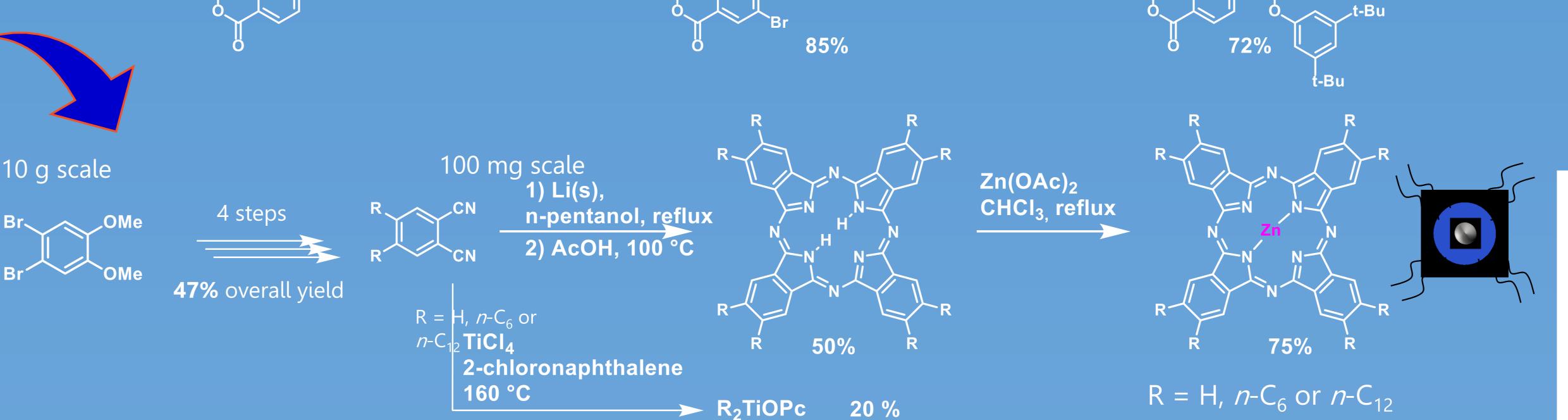
These 'smart' building-blocks were designed with the aim to develop a homogeneous, functional, integrated self-assembly on a graphenoid surface. With this molecular design strategy we are able to **fine-tune** the self-assembly, **orientation** and **functionality** via modification of:

Functionality: a perylene (PDI); with steric hindrance to avoid its assembly on the surface. It is a strongly fluorescent component. A donor/acceptor unit could replace it.

Linker: a pyridyl derivative; decouple the functionality from the surface. Orientation and distance can be controlled by modifying it.

Pedestal: a phthalocyanine (Pc); functionalized on peripheral positions to enable the supramolecular self-assembly on the surface. A nonfunctionalized phthalocyanine was also used in host guest systems.

SYNTHESIS



CONCLUSIONS

Alkylated ZnPcs were synthesized as well as several **PDIs** with **different linkers**.

In solution, the association of **PDI with ZnPcs** comes from π -stacking. On the contrary, the association of PDI with ZnPs comes from **metal-ligand interactions**.