

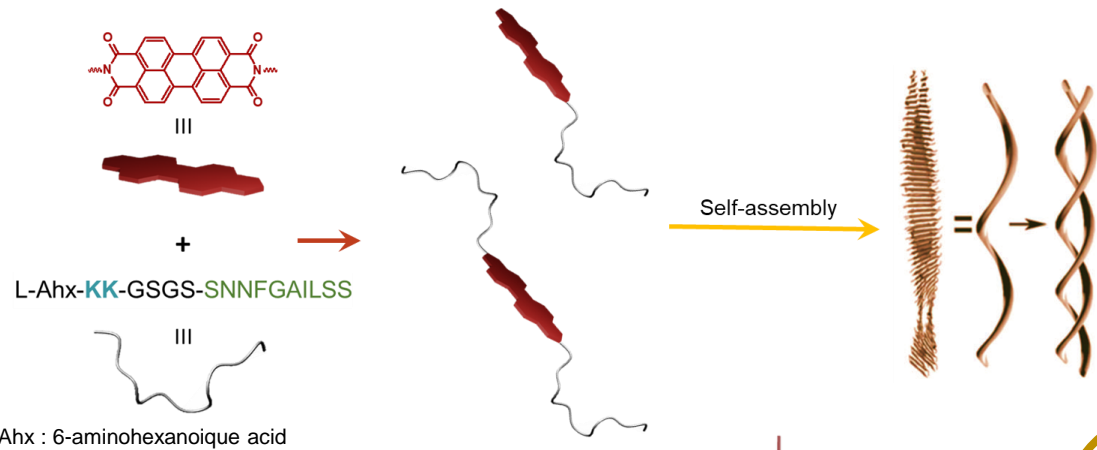
# Synthesis and self-assembly of perylene diimide-conjugated peptides

Nadjib Kihal<sup>1,2</sup>, Ali Nazemi<sup>1</sup> and Steve Bourgault<sup>1,2</sup>

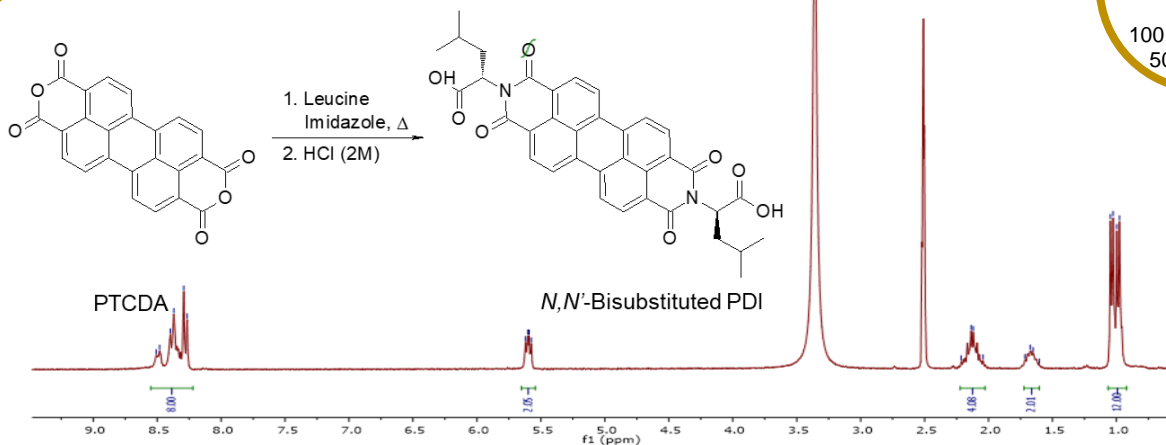
<sup>1</sup> Department of Chemistry, Université du Québec à Montréal, QC, Canada. // <sup>2</sup> Quebec Network for Research on Protein Function, Engineering and Applications, PROTEO

## Introduction

Control the directionality of PDI self-assembly into defined nanostructures remains problematic. We exploited the properties of amyloid peptides to design nanofilaments functionalized with PDI.



## Design perylene diimide-conjugated peptides

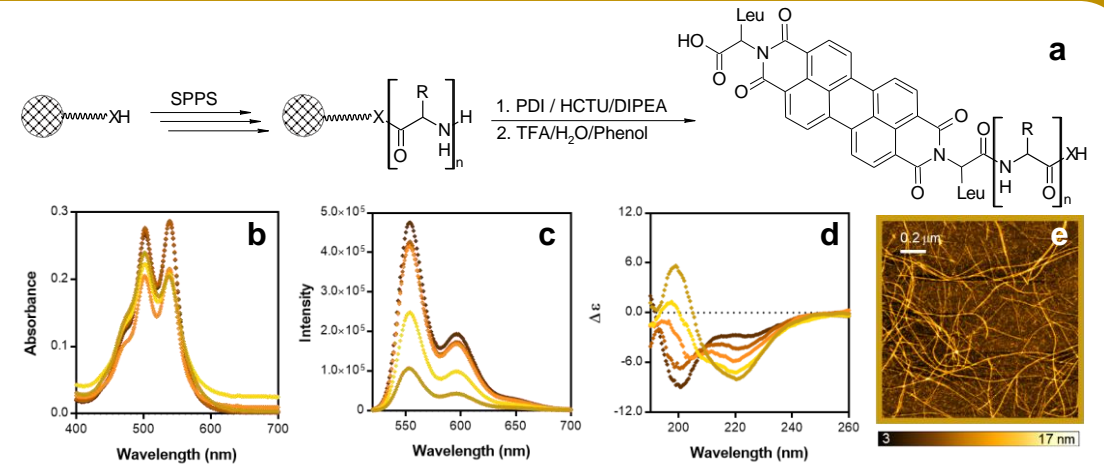


\* PTCDA : Perylenetetracarboxylic dianhydride  
\* PDI : Perylediimide

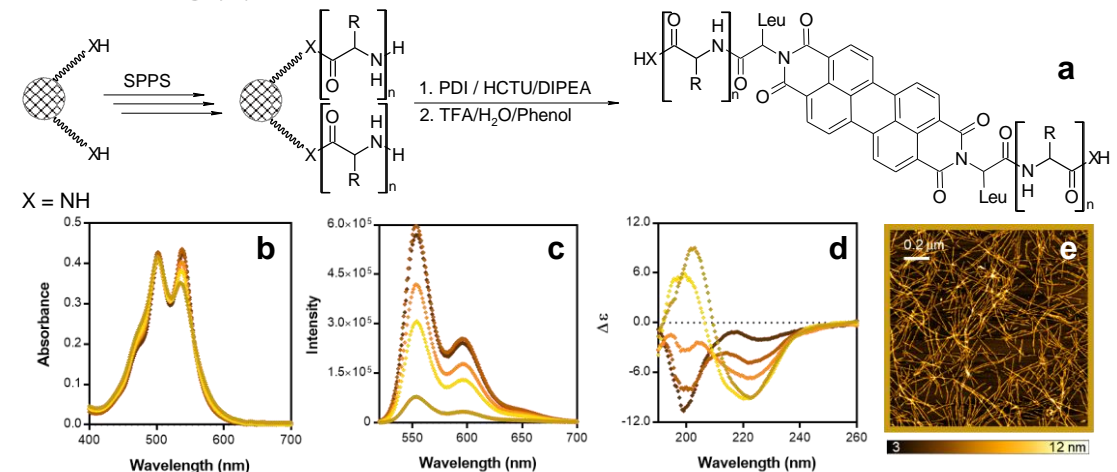
Synthesis of  $N,N'$ -bisubstituted PDI and  $^1\text{H-NMR}$  spectra (400 MHz,  $d_6$ -DMSO).

pH 7.4  
100 % TrisHCl  
500  $\mu\text{M}$  / rt

I



II



\* SPPS : Solid Phase Peptide Synthesis

Synthesis (a), UV-vis (b), fluorescence emission (c), ultraviolet range CD (d) spectra and AFM images (e) of PDI-I<sub>20-29</sub> (I) and PDI-[I<sub>20-29</sub>]<sub>2</sub> (II) incubated in Tris Buffer, pH 7.4, at a concentration of 500  $\mu\text{M}$  and under continuous agitation at 0 h, and after 24 h, 48 h, 72 h, and 1 week incubation.

## Conclusion

Defined nanostructures functionalized with PDI can be obtained from amyloid peptide building blocks, opening to novel applications in bioimaging, photodynamic therapy and bioelectronic.