

# Posters

## IMPROVEMENT OF POWER OUTPUT IN ORIENTED PHOTOSYSTEM I-BASED NANODEVICES

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Photosystem I (PSI) pigment-protein complex with its an internal quantum yield close to unity for generation of the primary charge separation state makes this complex one of the most efficient light-harvesting molecular machines for application in energy-converting devices. One of the requirements for efficient application of this biological macromolecule in such devices is the appropriate interface assembly between the photoactive protein and the electrode that ensures optimal electronic communication between both modules.

In this study, the conductive interface between the PSI layer and graphene electrode was provided by an organic self-assembled monolayer (SAM) composed of  $\pi$ - $\pi$  stacked pyrene functionalized with nitrilotriacetic acid chelated to Ni<sup>2+</sup> cations (Ni-NTA). Upon specific conjugation of His<sub>6</sub>-tagged cytochrome *c*<sub>553</sub> (cyt *c*<sub>553</sub>) with the graphene-pyrene-Ni-NTA FTO electrode, the PSI photoactive layer was incorporated in an oriented manner, i.e. with its donor (P700) side towards the electrode, taking the advantage of the molecular interaction between PSI and cyt *c*<sub>553</sub>. The photoelectrochemical analysis showed mediatorless generation of cathodic photocurrents on the FTO-graphene conductive substrate, even at an open circuit potential, which were further increased upon application of negative overpotential. These results indicate the production of the specific orientation of PSI resulting in the improved direct electron transfer and photocurrent output.

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## MOLECULAR MECHANISMS OF PHOTOADAPTATION OF THE RED ALGAL PHOTOSYNTHETIC APPARATUS

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The monomeric PSI-LHCI supercomplex from an extremophilic red microalga *Cyanidioschyzon merolae* exhibits structural traits of both cyanobacterial and eukaryotic counterparts that places it as an intermediate evolutionary link between them [1, 2]. Our research on extremophilic red algae, led us to the discovery of the molecular mechanisms underlying unprecedented robustness of the *C. merolae* PSI-LHCI supercomplex upon its exposure to harsh conditions. By the combined use of biochemical, spectroscopic, mass spectrometry and electron microscopy/single particle analyses we characterised three such mechanisms, whose biochemical principles make them independent both at the metabolic and time response levels [3]. The first mechanism consists of the accumulation of a photoprotective carotenoid, zeaxanthin, in both LHCI antenna and PSI reaction centre. The second mechanism involves the structural and functional remodelling of the LHCI antenna. The third mechanism involves the presence of the two PSI-LHCI isomers whose stoichiometric ratio alters in response to dynamic environmental conditions. This process is accompanied by dissociation of the PsaK core subunit, as shown by the quantitative mass spectrometry analysis [3]. Our experiments show that the low light-treated *C.*

*merolae* PSI-LHCI complex incorporates up to 8 Lhc antennae subunits which are organized as 2 rows on the PsaF/PsaJ side of the core complex. The spectroscopic and biochemical analyses show no evidence of functional coupling of the phycobilisomes with the PSI-LHCI supercomplex, suggesting that such an interaction does not occur under our experimental conditions [3] in agreement with other studies. [4, 5]

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