SUPPORTING INFORMATION

Enhanced Photostability of Chlorophyll-*a* using Gold Nanoparticles as Efficient Photoprotector

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Fig. S-1 XPS survey spectra of Chla, Chla-AuNPs, irradiated Chla and irradiated Chla-AuNPs.

High-resolution C1s and N1s XPS spectra for gold nanoparticles (AuNPs) are shown in Fig.S-2 and S-3, respectively. The C1s region of AuNPs comprises three distinct peaks. The peak appearing at 285 eV, is assigned to C-C and C-H from tetraoctylammonium bromide (TOAB), the stabilizing agent used in the preparation of AuNPs. The second peak at 286.07 eV is ascribed to C-N of quaternary amine from TOAB, while a smaller peak at higher binding energy (287.17 eV) is probably due to an impurity.



Fig. S-2 C 1s narrow scan XPS spectra for Chla and Chla-AuNPs.



Fig. S-3 O 1s narrow scan XPS spectra for Chla and Chla-AuNPs.



Fig. S-4 C1s narrow scan XPS spectrum for AuNPs.

Kinetics of Photochemical reaction

The photodegradation of Chla in presence of O₂ can be described by following equation

$$Chla + O_2 \xrightarrow{h\nu} Chla(O_2) \tag{1}$$

The photodegradation rate of Chl*a* was monitored by recording the absorbance at 665 nm as a function of irradiation time, and is:

$$-\frac{d[Chla]}{dt} = k[Chla]^*[O_2]$$
⁽²⁾

Where [Chl*a*] denotes the concentration of Chl*a*, $[O_2]$ denotes the concentration of O_2 , *t* is the reaction time, and *k* is the rate constant. As the $[O_2]$ remains constant as the reaction proceeds, the reaction can be considered pseudo-first-order because it depends on the concentration of [Chl*a*] only.

Thus, eq. (2) can be written as follows:

$$-\frac{d[Chla]}{dt} = k^{\cdot}[Chla]$$
(3)
where $k^{\cdot} = k[O_2]$

Solving eq. (3), one obtains eq. (4)

$$\ln([Chla]_{t}) = -k't + \ln([Chla]_{0})$$
(4)

Where $[Chla]_0$ is the initial concentration of Chla and $[Chla]_t$ is the concentration of Chla at any time t. A plot of $ln([Chla]_t)vs$ t should yield a straight line with a slope of k'; a typical plot is shown in the inset of Fig. 4.