244 Spectroscopic study of cadmium Quantum Dots biosynthesized by *Escherichia coli* using taurine as sulphur source.

V. Durán-Toro^{1,2} and J. M. Pérez-Donoso¹

1- Bionanotechnology and Microbiology Lab, Center for Bioinformatics and Integrative Biology (CBIB), Universidad Andres Bello, Republica 239, Santiago, Chile

2- Facultad de Ciencias Químicas y Farmacéuticas, Universidad de Chile, Sergio Livingstone Pohlhammer 1007, Santiago, Chile.

E-mail address corresponding author: vicente.mdt@gmail.com

Generation of cadmium sulphide (CdS) quantum dots (QDs) using *E. coli* cultures is a novel and promising alternative to traditional methods of synthesis. This phenomenon depends of the sulphide sources used in the process [1]. To date, no synthesis has been reported using taurine as sulphur source. Based on this, we report the characterization of CdS QDs synthesized using taurine as the main sulphur source by *E. coli* cultures.

In the present work we evaluated the changes in culture fluorescence during cadmium incubation as a preliminary screening for NPs formation [2]. A color change from green to red in pellets fluorescence was detected after 12 h incubation indicating the generation of cadmium based QDs as reported before [2].

Afterward, we performed a multispectral confocal microscopic analysis in order to evaluate the location an in situ properties of the NPs. An intense green fluorescence emission in both poles of E. coli cells treated during 10 h with cadmium was found indicating the intracellular formation of ODs. Finally; in-situ fluorescence spectrum analysis was determined and reported for biosynthetic cells for the first time. Interesting, a maximum emission wavelength (λ_{em}) of 500 nm is reported confirming the biosynthesis screening. Finally, Fig. 1a shows the absorbance spectrum of NPs purified from bacterial cells. A smooth disturbance in the range of 350-390 nm correlate with the plasmonic behavior described for biosynthesized CdS QDs [2]. Furthermore, fluorescence emission spectrum (Fig. 1b) shows a λ_{em} at 500 (green) and 540 (orange) nm for samples incubated with cadmium for 10 and 24 h, respectively. The change in the λ_{em} confirms the size growth of the NPs, a classical behaviour characterizing QDs. Additionally, the maximum excitation wavelength (λ_{ex}) do not vary in an incubationtime depending form, which is also characteristic of QDs.

Based in these results we confirmed the generation of Cd QDs using taurine by *E. coli*, demonstrating the potentiality of taurine as sulphur source during biological synthesis of NPs of different colours.



Fig. 1 Spectroscopic characterization of biosynthesized cadmium fluorescent NPs. 1a., Absorbance spectrum of biosynthesized NPs incubated after 10 and 24 h with cadmium. 1b., Emission fluorescence spectrum of biosynthesized NPs after 10 and 24 h of cadmium incubation. 1c., Excitation fluorescence spectrum of biosynthesized NPs. 1d., Biosynthesized NPs after 0 (control), 10 and 24 h of metal incubation exited at 365 nm.

[1] Mandal, D., Bolander, M. E., ... & Mukherjee, P. Appl Microbiol Biotechnol. **69**, (2006).

[2] Monrás, J. P., Díaz, V., ... & Pérez-Donoso, J. M. PloS one. 7, e48657, (2012).