



# PHOTOSTABLE AND BIOCOMPATIBLE CARBON DOTS FROM CITRIC ACID FOR BIOIMAGING

F. Fiori<sup>1,2</sup>, F. Olia<sup>1</sup>, L. Stagi<sup>1</sup>, L. Malfatti<sup>1,3</sup>, S. Ledda<sup>4</sup>, P. Innocenzi<sup>1,3</sup>

<sup>1</sup>Laboratory of Materials Science and Nanotechnology (LMNT), Department of Biomedical Sciences, University of Sassari, Sassari, Italy

<sup>2</sup>Department of Chemistry, Biology and Biotechnologies, University of Perugia, Perugia, Italy

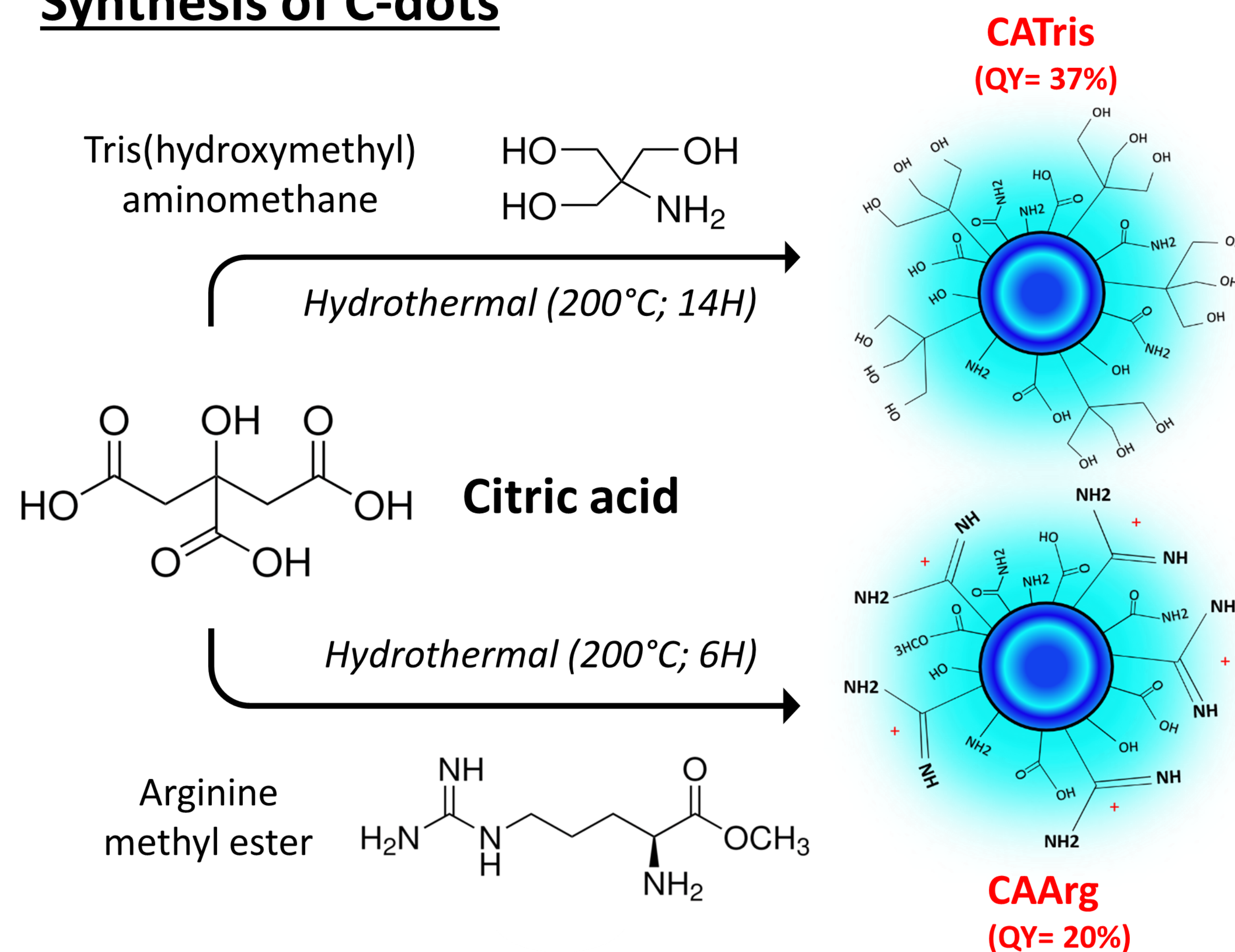
<sup>3</sup>National Interuniversity Consortium of Materials Science and Technology (INSTM), Florence, Italy

<sup>4</sup>Department of Veterinary Medicine, University of Sassari, Sassari, Italy

## INTRODUCTION

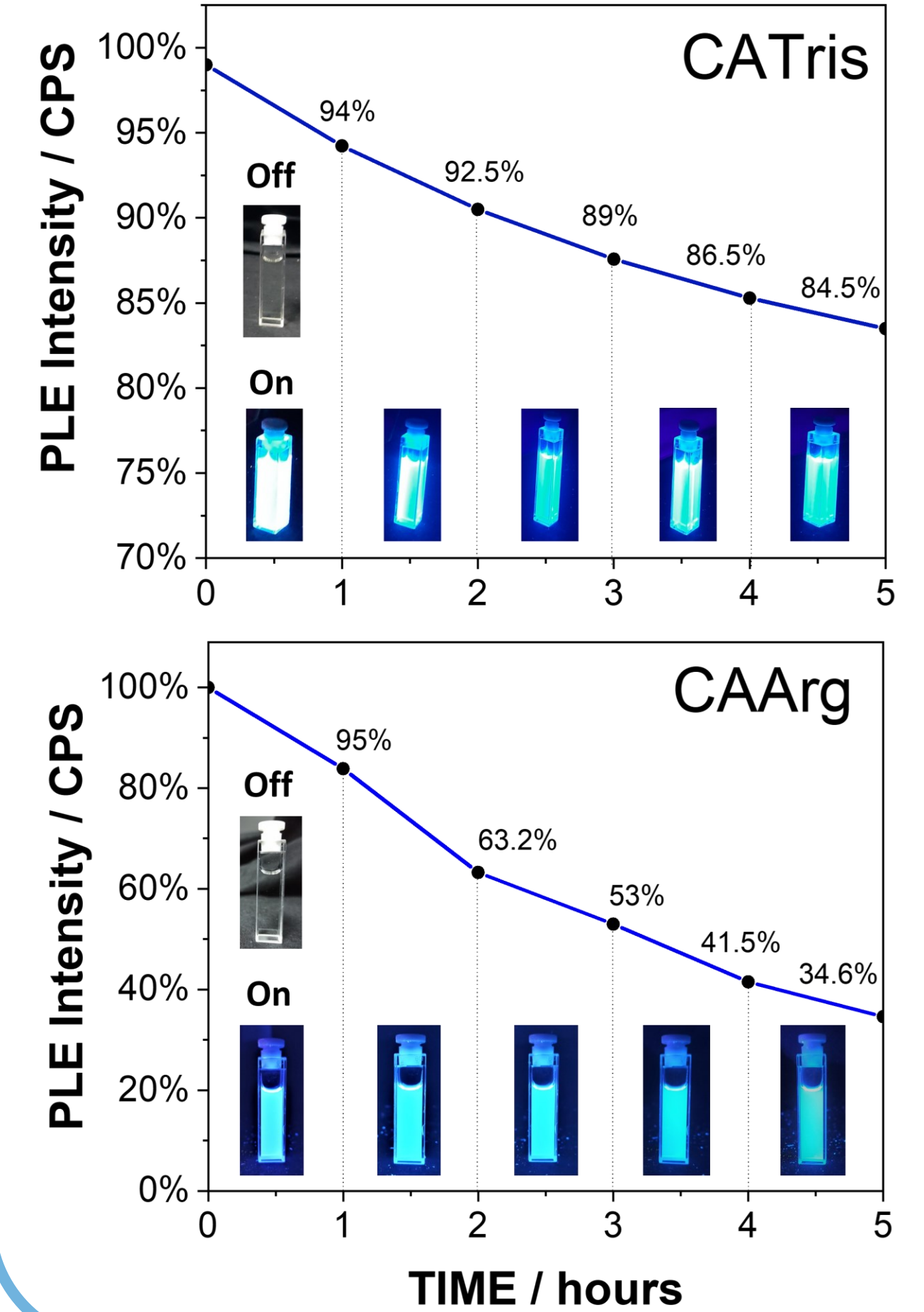
Carbon dots (**C-dots**) are nanoparticles obtained by the assembly of simple organic molecules and characterized by a strong fluorescence emission, photostability and low cytotoxicity. In this work<sup>1</sup>, two C-dots were synthesized from **Citric acid**, which represents a common source to form a carbonaceous core<sup>2</sup>. Tris (hydroxymethyl) aminomethane was chosen for **CATris** because of the possible formation of a dendritic structure associated with photostability properties<sup>3</sup>. While Arginine, was selected for **CAArg** for its guanidinium group that is characterized by strong interactions with cell membranes and thus efficient uptake<sup>4</sup>.

## Synthesis of C-dots



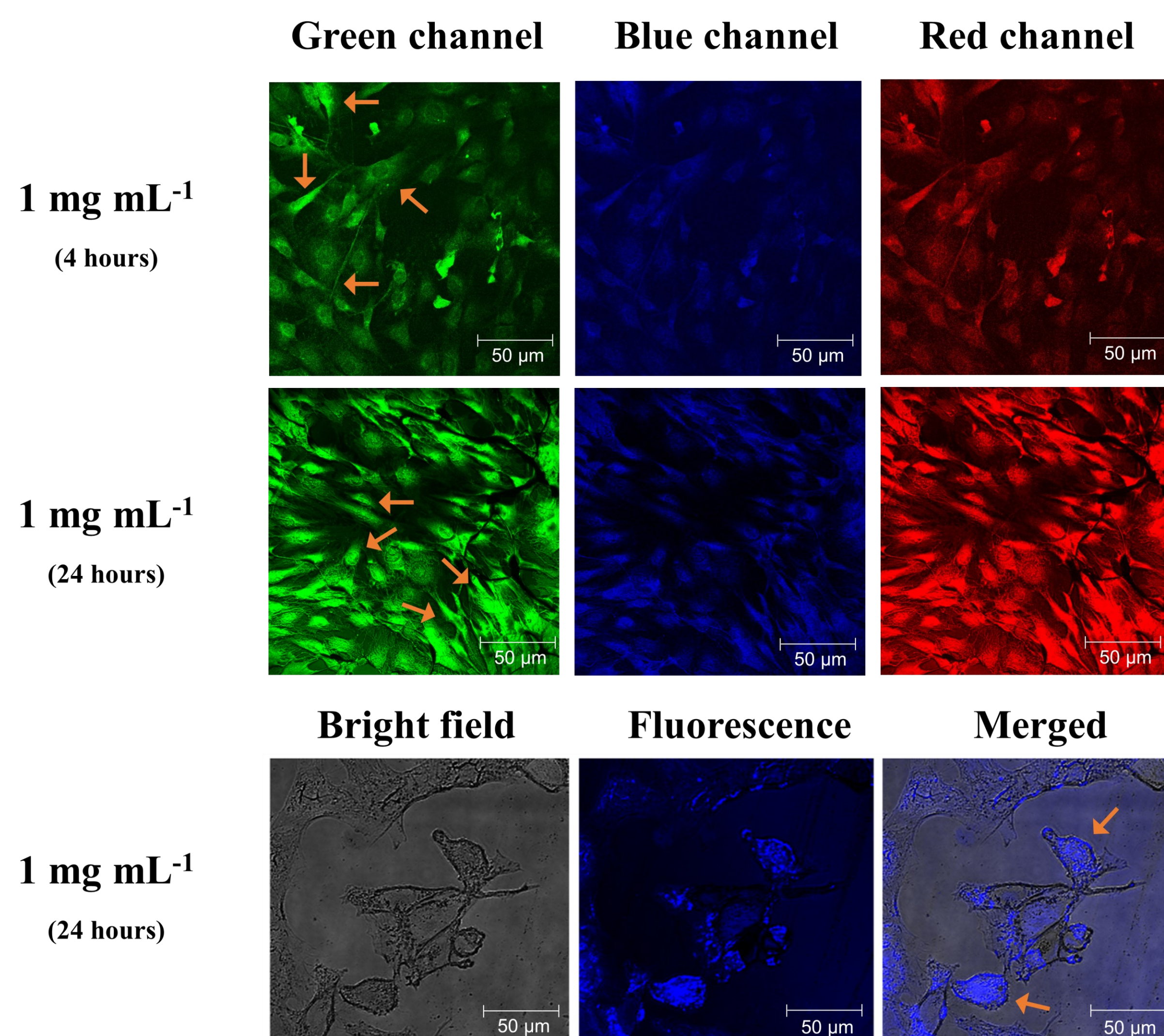
## PHOTOSTABILITY

$\lambda_{ex} = 365 \text{ nm}$

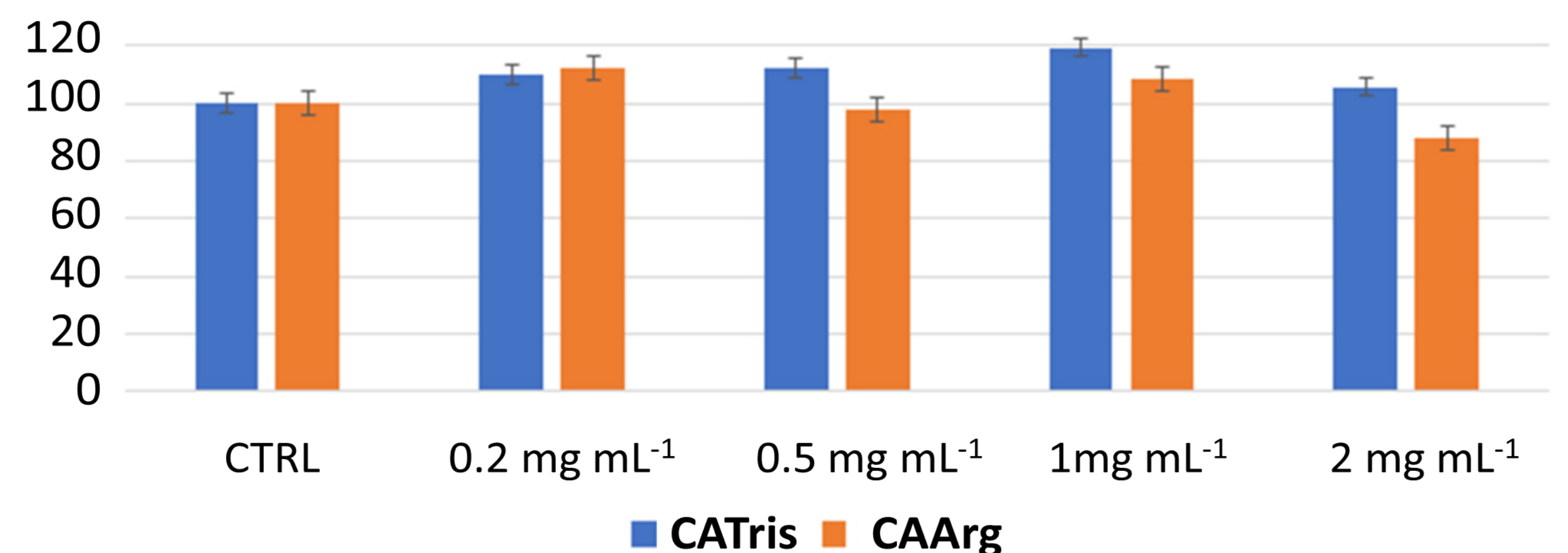


## FLUORESCENCE CONFOCAL MICROSCOPY

images of fibroblast with internalized CATris



## CELL VIABILITY ASSAY: (MTT - % of control) evaluated after 24 hours exposition of Fibroblast Cells to different concentrations of C-dots



## CONCLUSIONS

The present study investigated the cellular uptake of two types of **C-dots** obtained from citric acid. **CATris** displayed a better optical performance over **CAArg** and a higher photostability under UV irradiation. The biocompatibility assay showed that up to the concentration of 2 mg mL<sup>-1</sup>, both C-dots can be safely incubated in fibroblast cell cultures without any cytotoxicity. Finally, during the fluorescence confocal microscopy experiments, C-dots mainly localized in the cytoplasm, with some nanoparticles also detected in the nucleus, proving their suitability for **bioimaging** applications.

## REFERENCES

- Fiori, F. *et al.* Highly Photostable Carbon Dots from Citric Acid for Bioimaging. *Materials* **2022**, 15, 2395
- Ludmerczki, R. *et al.* Carbon Dots from Citric Acid and its Intermediates Formed by Thermal Decomposition. *Chem. A Eur. J.* **2019**, 25, 11963–11974
- Liu, Y. *et al.* Highly fluorescent nitrogen-doped carbon dots with excellent thermal and photo stability applied as invisible ink for loading important information and anti-counterfeiting. *Nanoscale* **2017**, 9, 491–496
- Fu, C. *et al.* Arginine-modified carbon dots probe for live cell imaging and sensing by increasing cellular uptake efficiency. *Mater. Sci. Eng. C* **2017**, 76, 350–355