

Solar-driven photocatalytic degradation of amoxicillin using TiO₂/Carbon Quantum Dots nanocomposites immobilized in electrospun polycaprolactone fibers

Beatriz Valadares^{1,2}, Valentina Silva³, Rosa Baptista⁴, Bernardo Almeida⁴, Diana Lima^{3,5}, Vânia Calisto³, Goreti Pereira³

¹Department of Chemistry, University of Aveiro, 3810-193 Aveiro, Portugal.
beatriz.valadares@ua.pt

²Polytechnic Institute of Coimbra, Coimbra Health School, Rua 5 de Outubro – S. Martinho do Bispo, Apartado 7006, 3046-854 Coimbra, Portugal

³Department of Chemistry and CESAM, University of Aveiro, 3810-193 Aveiro, Portugal,

⁴Centre of Physics of Minho and Porto Universities (CF-UM-UP), Laboratory for materials and Emergent Technologies (LAPMET), University of Minho, 4710-057 Braga, Portugal.

⁵H&TRC - Health & Technology Research Center, Coimbra Health School, Polytechnic University of Coimbra, 3045-043, Coimbra, Portugal

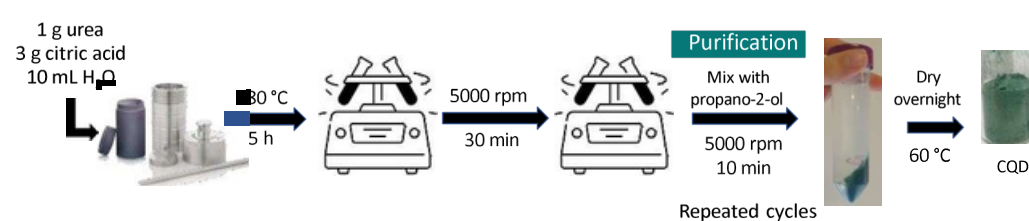
INTRODUCTION

The persistence of antibiotics in water contributed to antimicrobial resistance, as conventional treatments fail to fully remove them. Solar-driven photocatalysis offers a sustainable way to degrade these contaminants. TiO₂ modified with Carbon Quantum Dots (CQDs) enhances light absorption, while immobilization in polycaprolactone (PCL) nanofibers allows easy recovery and reuse. This work aims to develop and evaluate TiO₂/CQDs photocatalysts embedded in electrospun PCL nanofibers for the solar-assisted degradation of amoxicillin. These study focuses on optimizing photocatalytic performance and demonstrating the potential of this composite material for sustainable water treatment applications.

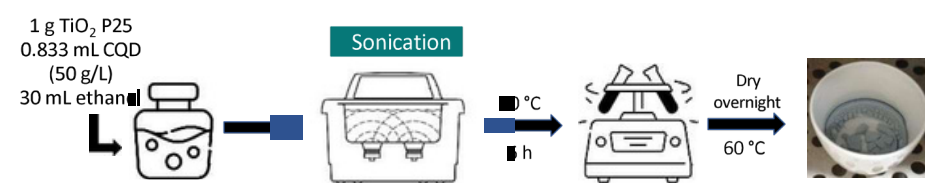
METHODOLOGY

Carbon Quantum Dots Synthesis

CQD were synthesized by a simple hydrothermal treatment

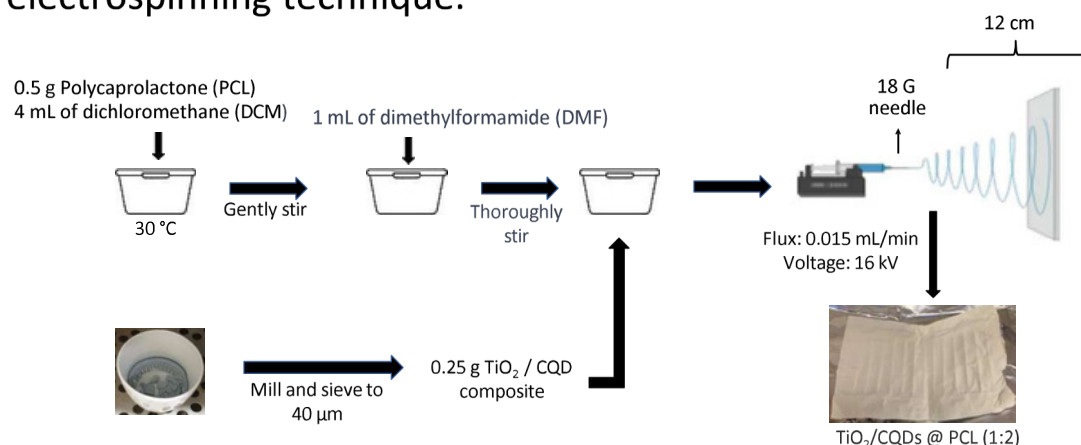


TiO₂/CQD nanocomposite synthesis



Incorporation of TiO₂/Carbon Quantum Dots into Polycaprolactone nanofibers

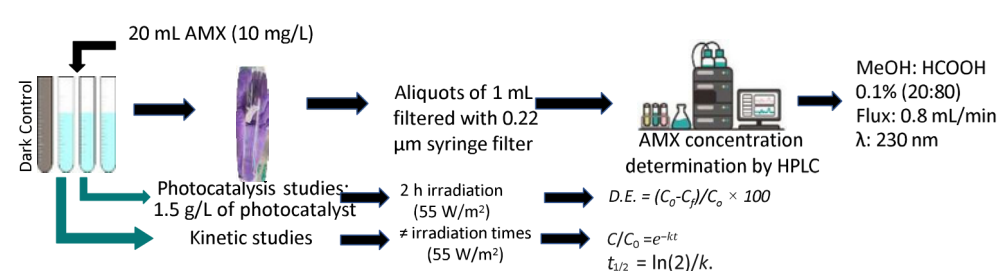
The composite was then incorporated into a matrix of polycaprolactone (PCL) biopolymer nanofibers using the electrospinning technique.



The electrospun fibers were analyzed by SEM (Scanning Electron Microscopy) operating at an acceleration voltage of 10 kV. To prepare the nanofibers for analysis, they were deposited onto a silica surface, pre-coated with a 10 nm thick Au-Pd (80-20 weight %) film.

Photodegradation studies

The nanofibers were tested for the photocatalysis of 10 mg/L of AMX in phosphate buffer (PBS) 0.001 mol/L, pH 8.0, using 1.5 g/L of photocatalyst and irradiation in a solar radiation simulator during 2 h.



RESULTS & DISCUSSION

•Characterization of nanofibers-TiO₂/CQD @ PCL (1:2) by microscopy

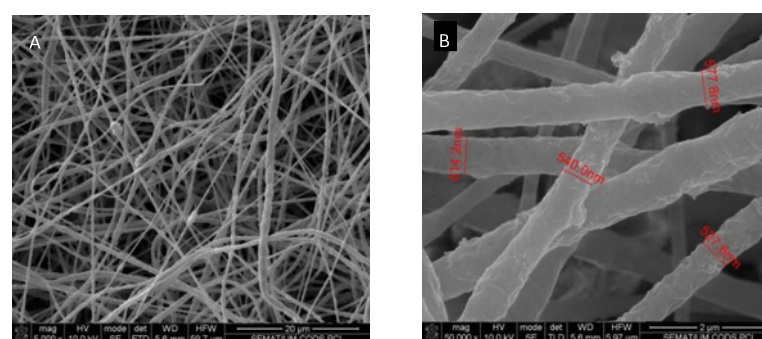
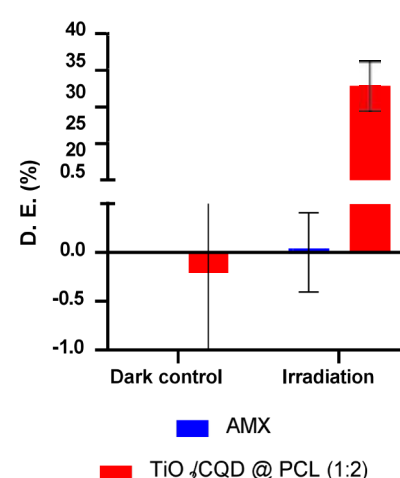


Figure 1. Microscopy images of TiO₂/CQD @ PCL (1:2) in SEM with (A) 5000 x and (B) 50000 x of magnification, respectively.

•Photodegradation studies

Photolysis vs Photocatalysis



Kinetics

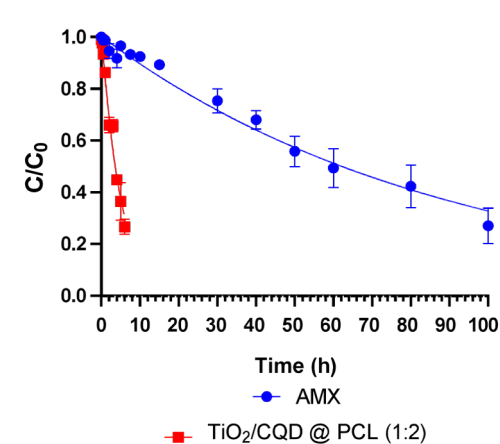


Figure 2. (A) Preliminary results for the application of 1.5 g/L of TiO₂/CQD @ PCL (1:2) for the removal of 10 mg/L of AMX and (B) Kinetics of photodegradation of AMX in presence and absence of photocatalyst.

Table 1. Data on pseudo-first order rate constants (k (h⁻¹)), determination coefficient (R^2) and half-life times ($t_{1/2}$ (h)) obtained for with and without photocatalyst under simulated solar radiation for 10 mg/L AMX photodegradation.

Sample	$k \pm SD$ (h ⁻¹)	r^2	$t_{1/2} \pm SD$ (h)
Photolysis	0.0112 ± 0.0002	0.9518	62 ± 1
Photocatalysis	0.201 ± 0.006	0.9651	3.45 ± 0.03

Increase of 18 TIMES AMX removal!

CONCLUSION & FUTURE WORK

The presented results constitute a promising and sustainable treatment for antibiotics' efficient removal from water. The electrospun PCL nanofibers doped with TiO₂/CQD nanocomposites afforded a simple and easy recoverable photocatalyst. Next in line, the continuation of the optimization of the photocatalyst application will be studied, various cycles of application will be performed, and the stability of the nanofiber will be evaluated.

ACKNOWLEDGEMENTS

Acknowledgements: This work is funded by national funds through FCT – Fundação para a Ciência e a Tecnologia I.P., under the project/grant UID/50006 + LA/P/0094/2020 (doi.org/10.54499/LA/P/0094/2020). Valentina Silva thanks FCT support for the individual PhD grant (2022.10472.BD, https://doi.org/10.54499/2022.10472.BD). This research was also funded by FCT through FEDER (European Fund for Regional Development)-COMPETE-QREN-EU (ref. UID/FIS/04650/2013 and UID/FIS/04650/2019). Diana L.D. Lima thanks support by Fundação para a Ciência e a Tecnologia (FCT) to the SOLCAT research project (2023.12723.PEX, https://doi.org/10.54499/2023.12723.PEX).