

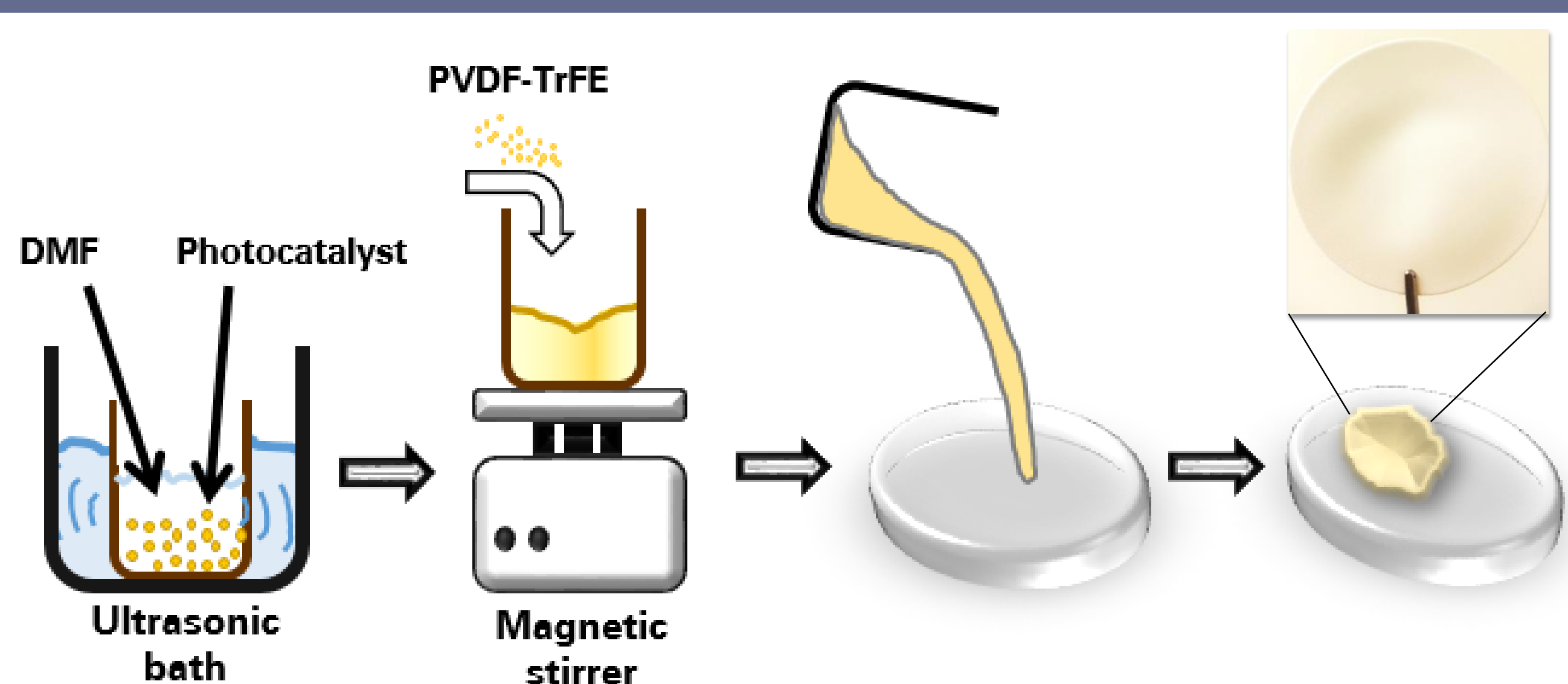
Introduction

Pharmaceuticals in the environment are of major public health concern due to their potential toxicity to humans and other living organisms [1]. When consumed, pharmaceuticals are excreted into the sewer network and reach the WWTP [2], [3], [4]. The inability of the WWTPs to effectively eliminate these pollutants makes it necessary to find alternative methods. Therefore, research on Advanced Oxidation Processes (AOPs) has been immense [5]. Among several AOPs, heterogeneous photocatalysis arises as a promising approach regarding water pollution mitigation as it only requires a source of energy and a photocatalyst [6], producing reactive hydroxyl radicals that degrade different organic contaminants in water.

The photocatalysts can be employed either in suspension or immobilized. Immobilized photocatalysts show an ease of recovery and great potential for reutilization [4], [7]. An efficient attachment of the photocatalyst to the substrate and a production method that does not reduce the efficiency of the catalytic properties are paramount for application. Polymers are good materials to immobilize the photocatalysts because of their high mechanic stability and durability [8].

In this work, 8 % wt. P25 TiO₂ nanoparticles were immobilized in PVDF-TrFE by solvent casting. One of the problems arising from the use of PVDF-TrFE is its low wettability which limits the interaction between the pollutant and the photocatalytic material. The nanocomposites were characterized by SEM/EDX, FTIR-ATR and contact angle, and their photocatalytic properties and reusability assessed on the degradation of four compounds: bisphenol A, ciprofloxacin, ibuprofen and methylene blue.

Figure 1: Scheme of the preparation method of the nanostructured materials



Aim of the project

Synthesis and reusability of nanostructured materials to use on the degradation of water pollutants via photocatalysis using TiO₂ and UV.

Methods

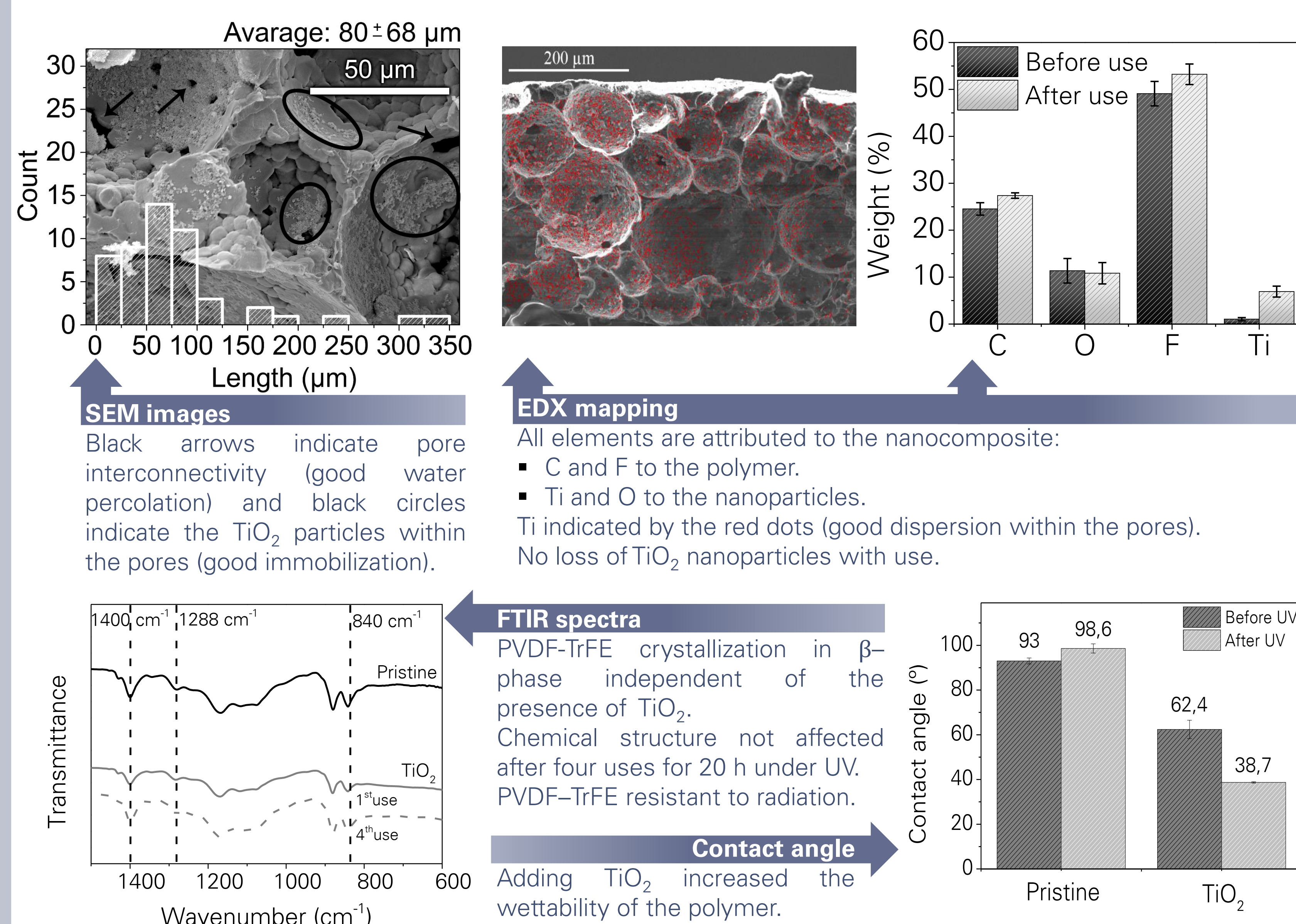
- 8 % wt. TiO₂ P25 immobilized in PVDF-TrFE (2.5 x 10 cm)
- 50 ml containing each pollutant were exposed for 5 hr to UV (365 nm) at 1.8–1.9 mW cm⁻²
- Nanocomposites were reused 3 times
- The degradation of the pollutants was determined by the Langmuir-Hinshelwood model:

$$\ln \frac{C}{C_0} = -kt$$

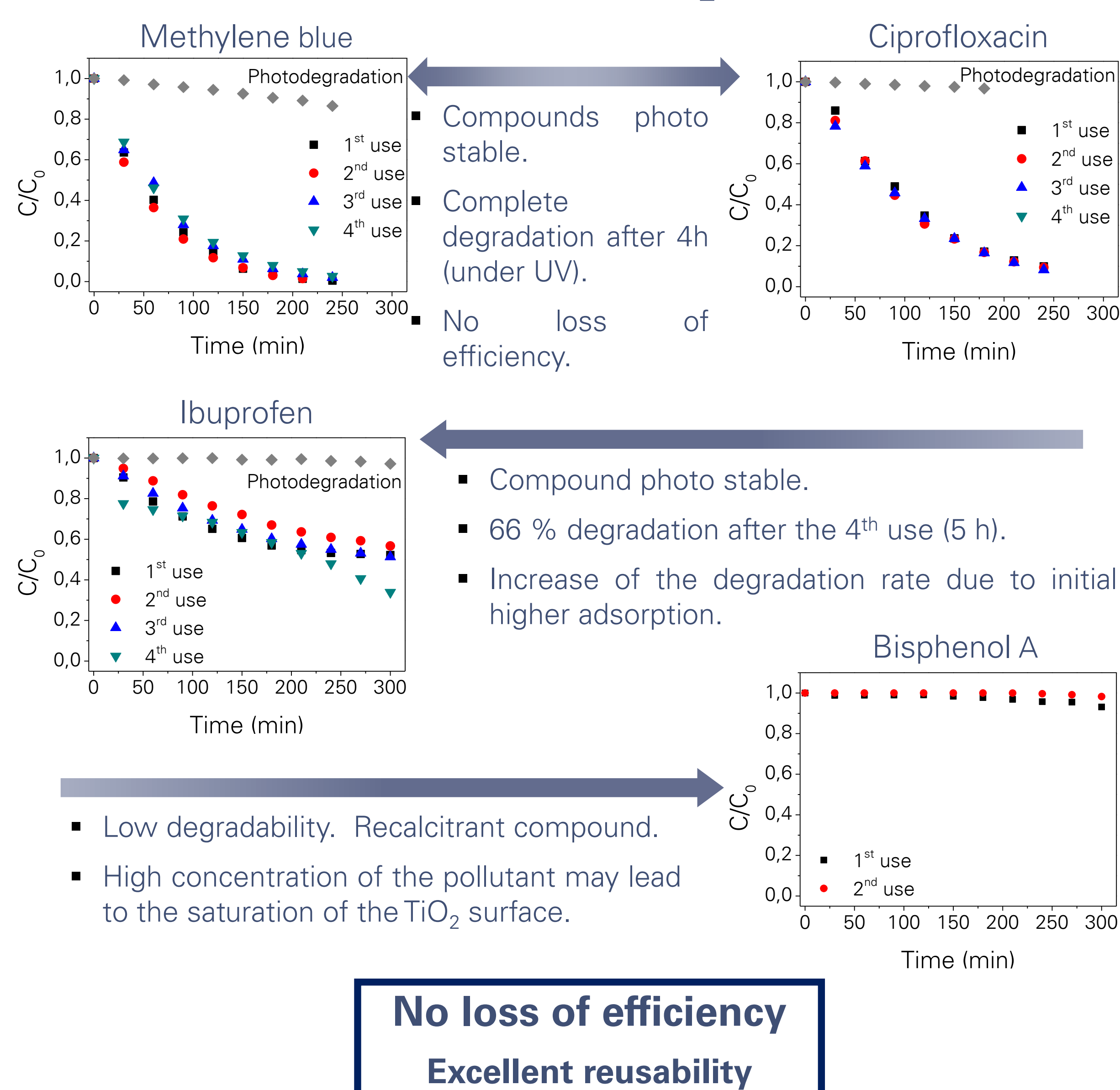
C_0 - initial concentration of the compound
 C - concentration of the compound at time t

Results and Discussion

Characterization of the TiO₂/PVDF-TrFE nanocomposites



Photocatalytic activity of the TiO₂/PVDF-TrFE under UV



Conclusions

- The produced nanocomposites exhibited a highly porous structure and the addition of TiO₂ increased the hydrophilicity of the nanocomposites.
- The polymeric structure was not affected by the addition of TiO₂ and it was kept intact after four uses (total time of 20 h) under UV.
- The tested compounds were totally or partially degraded by the produced material under UV, except bisphenol A, which is a recalcitrant compound.
- The reutilization of the nanocomposites showed no loss of efficiency. EDX confirmed the presence of the TiO₂ after 4 uses.
- These nanocomposites are a cost-effective nanomaterial for environmental remediation.

Acknowledgements

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References

- [1] M. Isidori et al., Environment international, vol. 35, no. 5, pp. 826–829, 2009.
- [2] S. Teixeira et al., Journal of Environmental Chemical Engineering, vol. 4, no. 1, pp. 287–292, Mar. 2016.
- [3] S. Mompelat, B. Le Bot, and O. Thomas, Environment international, vol. 35, no. 5, pp. 803–814, 2009.
- [4] V. Homem and L. Santos, Journal of Environmental Management, vol. 92, no. 10, pp. 2304–2347, 2011.
- [5] M. Klavarioti, D. Mantzavinos, and D. Kassinos, Environment International, vol. 35, no. 2, pp. 402–417, 2009.
- [6] S. Ahmed et al., Desalination, vol. 261, no. 1–2, pp. 3–18, 2010.
- [7] L. Pereira et al., Photochemistry and Photobiology, vol. 89, no. 1, pp. 33–39, 2013.
- [8] S. Singh, H. Mahalingam, and P. K. Singh, Applied Catalysis A: General, vol. 462–463, pp. 178–195, 2013.