## **Supporting information**

## Biodegradable and drug-eluting inorganic composites based on mesoporous zinc oxide for urinary stent applications

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**Figure S1.** Picture of sample polyHEMA@ZnO\_1% which includes the silicon rubber mold and plates of Teflon.

The nitrogen sorption measurement allows for the calculation with the Brunauer-Emmet-Teller (BET) model of the specific surface area, about 19.58 m<sup>2</sup>/g. It also resulted some mesopourous-sized porosities of about 4 nm in diameter, calculated by Density Functional Theory (DFT) model applied to the equilibrium desorption branch of the isotherm. These pores, as well as the flower-like morphology, are responsible for the relatively high surface area and also act as preferential adsorption sites of the drugs.



**Figure S2.** (**a**) Nitrogen sorption isotherm with indication of the calculated BET surface area and (**b**) DFT pore size distribution of the mesoporous ZnO flower-like microparticles.



**Figure S3.** Morphological analysis of poly(HEMA-co-AA)@ZnO composite samples incorporating different ZnO amounts: (a) ZnO\_0.1 wt%; (b) ZnO\_1 wt%.



**Figure S4.** EDX results obtained for (**a**) polyHEMA@ZnO\_0.1% and (**b**) polyHEMA@ZnO\_1%. Each table summarizes the % atomic weight of each detected element. The detection of Pt is due to metallic coating of the samples needed for FESEM imaging.



Figure S5. FT-IR spectrum of mesoporous ZnO flower-like powders.



**Figure S6.** Concentration of zinc cations released from ZnO-based samples in cell culture medium (DMEM ,10 % fetal bovine serum) at different incubation times.





(c)



(**d**)











(**g**)



(h)



**Figure S7.** FT-IR spectra in case of Diclofenac and Ibuprofen release: (**a**,**b**) polyHEMA; (**c**,**d**) polyHEMA@ZnO\_0.1%; (**e**,**f**) polyHEMA@ZnO\_1%; (**g**,**h**) poly(HEMA-co-AA); (**i**,**j**)poly(HEMA-co-AA)@ZnO\_1%.